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Matheus de Andrade Cruz

ADVANCEMENTS TOWARDS FOSSIL ENERGY SUSTAINABILITY: CO<sub>2</sub>-RICH  
NATURAL GAS PROCESSING, MONETIZATION OF FLUE-GAS DESULFURIZATION  
RESIDUES, AND DECARBONIZATION VIA PHASE-CHANGING ABSORPTION

Rio de Janeiro  
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Doctoral thesis presented to the Environmental Engineering Program, Escola Politécnica & Escola de Química, from Universidade Federal do Rio de Janeiro, as part of the requirements for obtaining a Doctor of Science degree in Environmental Engineering.

Advisors: Ofélia de Queiroz F. Araújo, PhD  
José Luiz de Medeiros, DSc.

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*To the memory of my father Marcelo Cruz.  
To my mother Cristina, my brother Davi  
and my wife Daniele.*

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## RESUMO

CRUZ, M. de A. **Avanços para a sustentabilidade de energias fósseis: processamento de gás natural rico em CO<sub>2</sub>, monetização de resíduos de dessulfurização de gases exaustos e descarbonização por absorção com solventes bifásicos.** Tese (Doutorado em Engenharia Ambiental), Programa de Engenharia Ambiental, Escola Politécnica & Escola de Química, Universidade Federal do Rio de Janeiro, 2020. Orientadores: Ofélia de Queiroz Fernandes Araújo e José Luiz de Medeiros.

Visando aumento de eficiência energética e mitigação de impactos ambientais na cadeia de valor de energia fóssil são desenvolvidas/avaliadas novas tecnologias, principalmente de geração/utilização de energia. Assim, essa tese se desdobra em três linhas de pesquisa. Primeiramente, a produção offshore de petróleo e gás tem um histórico de baixa eficiência energética/exergética em relação à produção/utilização de energia no processamento de gás rico em CO<sub>2</sub>. Assim, novos conceitos envolvendo compressores centrífugos são avaliados: (i) resfriamento primário a 4°C com águas profundas, baixando a temperatura da água de resfriamento inter-estágio, com conseqüente redução do consumo energético de compressores; (ii) substituição de compressores centrífugos superdimensionados com ineficientes recírculos *anti-surge* por compressores menores em paralelo sem recírculo, que reduzem drasticamente a destruição exergética nesses sistemas ao longo de campanhas com carga de gás decrescente. Na segunda vertente do trabalho, é abordada a questão de termoelétricas a carvão que produzem resíduos de dessulfurização de gases exaustos (FGD, do inglês *Flue-Gas Dessulfurization*) fora de especificação para aproveitamento industrial, sendo destinados a aterros. Foi avaliado tratamento para comercialização de resíduos de FGD como aditivo para cimento. Em terceiro lugar, aborda-se a remoção de CO<sub>2</sub> de gases exaustos por soluções aquosas de alcanolaminas, que implica penalidade energética a termoelétricas. São realizados experimentos de captura de CO<sub>2</sub> com solventes bifásicos, desenvolvidos para redução do consumo de energia do processo. No processamento offshore de gás rico em CO<sub>2</sub>, é demonstrado que a utilização de captura profunda de água do mar reduz em 2% a 5% o consumo de energia/gás combustível e emissões de CO<sub>2</sub> e em 9,5% o custo dos principais equipamentos de processo. Ademais, compressores com recírculo *anti-surge* mantêm o consumo de energia/gás combustível alto durante toda a campanha de produção, resultando em eficiências exergéticas em torno de 49%/83% entre 25%/100% de carga de gás, enquanto compressores menores em paralelo sem recírculo reduzem a energia/consumo de combustível proporcionalmente à carga de gás, mantendo essa eficiência entre 80% e 88% e eliminando uma turbina a gás. Para usinas a carvão de 360MW, o custo nivelado de energia é reduzido em ~3% com a comercialização de resíduos de FGD. Uma avaliação *gate-to-gate* de impactos ambientais revela que a comercialização do resíduo é ambientalmente menos prejudicial do que dispô-los em aterros. Finalmente, experimentos em uma planta-piloto de pequena escala para captura de CO<sub>2</sub> indicam que soluções aquosas de monoetanolamina/propanol são promissores solventes bifásicos. É apresentado o projeto de uma planta-piloto de maior escala para testes de longa duração com solventes bifásicos.

Palavras-chave: Processamento de gás natural rico em CO<sub>2</sub>; Análise exergética; Gases exaustos; Absorção por solventes bifásicos; Resíduos de FGD;.

## ABSTRACT

CRUZ, M. de A. **Advancements Towards Fossil Energy Sustainability: CO<sub>2</sub>-Rich Natural Gas Processing, Monetization of Flue-Gas Desulfurization Residues, and Decarbonization via Phase-Changing Absorption.** Thesis (Doctorate in Environmental Engineering), Programa de Engenharia Ambiental, Escola Politécnica & Escola de Química, Universidade Federal do Rio de Janeiro, 2020. Advisors: Ofélia de Queiroz Fernandes Araújo and José Luiz de Medeiros.

Energy-efficiency and impact mitigation along the fossil-energy value chain drive the development/assessment of new technologies to improve the generation/utilization of fossil-energy. Bearing in mind those targets, this thesis unfolds in three research lines. Firstly, offshore oil/gas production in tropical deep-waters has a track record of low energy/exergy efficiencies regarding gas-fired power production/utilization in CO<sub>2</sub>-rich gas processing. Here, new concepts for better utilization of centrifugal compressors are assessed: (i) 4°C deep seawater primary-cooling lowering cooling-water temperature, consequently reducing power consumption of multistage intercooled compression; (ii) substitution of large centrifugal compressors with inefficient anti-surge recycles by multiple-paralleled smaller compressors dramatically reducing exergy destruction in compression systems along offshore campaigns with decreasing gas-load. Secondly, coal-fired power plants produce problematic flue-gas desulfurization (FGD) solid-residues often destined to landfills. Here, monetization was assessed for FGD solids as cement additives. Thirdly, aqueous-amine flue-gas decarbonation entails energy-penalty for power plants. Carbon capture experiments with phase-changing absorption solvents are conducted aiming at energy-penalty reduction. On offshore CO<sub>2</sub>-rich gas processing, it is shown that deep-seawater utilization lowers 2%-5% power/fuel-gas consumption and CO<sub>2</sub> emissions, and 9.5% of equipment investment. Moreover, large compressors with anti-surge recycle keep power/fuel-gas consumptions high throughout entire rig campaign entailing exergy efficiencies around 49%/83% at 25%/100% gas-loads, while multiple-paralleled smaller compressors reduce power/fuel-gas consumptions proportionally to gas-load keeping exergy efficiency around 80%-88% throughout campaign, eliminating one gas-turbine. For 360MW coal-fired power plants it is shown that the levelized-cost of energy is reduced ~3% thanks to commercializing FGD residues. Gate-to-gate environment impact assessment unveils upgrading/commercializing as environmentally better than landfilling FGD residues. Finally, experiments on a small decarbonation pilot-plant indicate aqueous-monoethanolamine-propanol solutions as promising CO<sub>2</sub> capture phase-changing solvents. The design of a larger pilot-plant intended to perform long-run trials with phase-changing solvent is presented.

Keywords: CO<sub>2</sub>-rich natural gas processing; Exergy analysis; Flue-gas; Phase-changing absorption; FGD residues.

## ABBREVIATIONS

<b>AGWA</b>	Acid gas-water-amine
<b>AMP</b>	2-amino-2-methylpropanol
<b>BAT</b>	Best available technologies
<b>BOE</b>	Barrels of oil equivalent
<b>CAPEX</b>	Capital expenditure
<b>CCP</b>	Coal combustion products
<b>CCS</b>	Carbon capture and storage
<b>CCUS</b>	Carbon capture utilization and storage
<b>CW</b>	Cooling-water
<b>DEA</b>	Diethanolamine
<b>DSW</b>	Deep seawater
<b>EKC</b>	Environmental Kuznets curves
<b>EOR</b>	Enhanced oil recovery
<b>EPCI</b>	Engineering, procurement, construction and installation
<b>EROI</b>	Energy return on investment
<b>ERR</b>	Energy return ratios
<b>FBR</b>	Fluidized bed reactor
<b>FCI</b>	Fixed Capital Investment
<b>FGD</b>	Flue-gas desulfurization
<b>FGDR</b>	Flue-gas desulfurization residues
<b>FPSO</b>	Floating production storage and offloading
<b>GDP</b>	Gross domestic product
<b>GHG</b>	Greenhouse gases
<b>GOR</b>	Gas to oil ratio
<b>GT</b>	Gas-turbines
<b>HCDP</b>	Hydrocarbon dew point
<b>LCC</b>	Life cycle cost
<b>LCOE</b>	Levelized cost of electricity
<b>MDEA</b>	Methyldiethanolamine
<b>MEA</b>	Monoethanolamine
<b>NER</b>	Net energy ratio
<b>NG</b>	Natural gas
<b>NGCC</b>	Natural gas combined cycle
<b>O&amp;G</b>	Oil and gas
<b>OPEX</b>	Operational expenditure
<b>P&amp;ID</b>	Process and instrumentation diagram
<b>PCAS</b>	Phase-changing absorption solvents
<b>PCC</b>	Pulverized coal combustion
<b>PM</b>	Particulate matter
<b>SD-FGDR</b>	Semi-dry flue-gas desulfurization residue
<b>SDG</b>	Sustainable development goals
<b>TETA</b>	Triethylenetetramine
<b>TMPDA</b>	Tetramethyl-1,3-propanediamine
<b>TPES</b>	Total primary energy supply
<b>VLCC</b>	Very large crude carriers
<b>WDP</b>	Water dew point
<b>WOR</b>	Water to oil ratio

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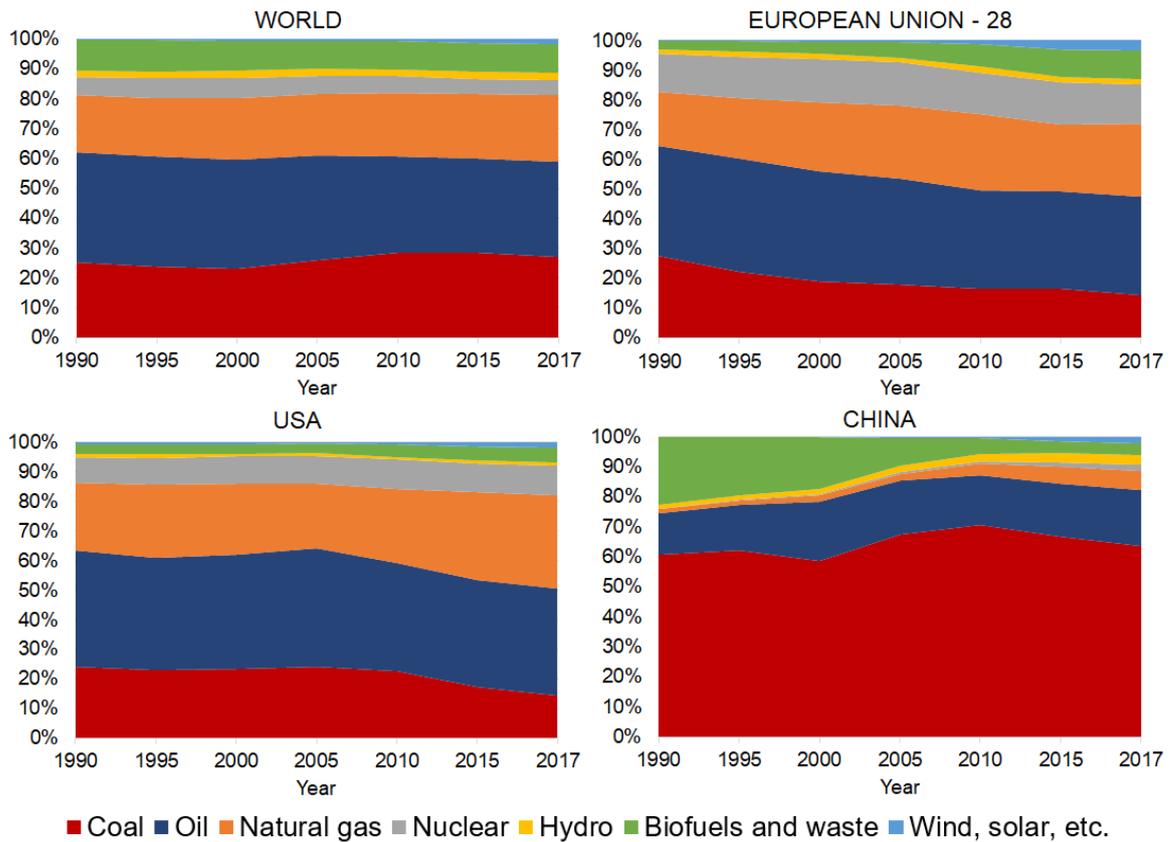
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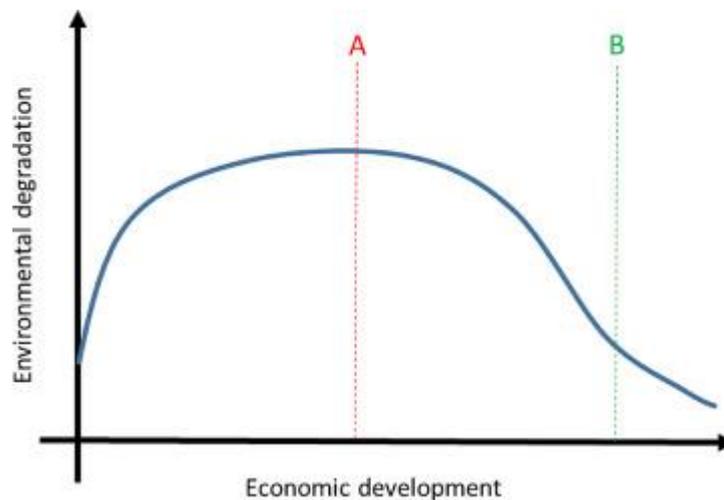
## 1. INTRODUCTION

The United Nations established universal access to affordable, reliable and sustainable energy as a goal of the 2030 Agenda for Sustainable Development (United Nations, 2015). Nevertheless, the energy sector has been considered the major responsible for climate changes, accounting for 80% of CO<sub>2</sub> emissions (IEA, 2019a). Furthermore, the extraction, processing and combustion of fossil fuels cause deterioration of air quality, depletion of natural resources among other negative environmental impacts. Since the developed economies are responsible for the major energy demand, they have been leading efforts to replace environmentally non-friendly energy sources (e.g. coal) by natural gas and renewable energy (wind and solar). Additionally, carbon capture utilization and storage (CCUS) technologies are expected innovations in clean-energy grids. However, CCUS deployments and grid transition to renewables are low-paced, comparatively to climate targets – fossil fuels remain responsible for 80% of total primary energy supply (TPES) worldwide, with shares varying across regions.

Frequently, in American countries, the energy demand of the transport sector is predominant and oil share is higher, while, in Asia, heat and power generation is responsible for most of the energy demand, prevailing coal in the energy matrix (IEA, 2020), as depicted in Fig 1.1. This difference in the share of coal-fired energy between developed and developing economies is explained through the Environmental Kuznets Curves (EKC) theory (Stern, Common and Barbier, 1996). In the early '90s, several authors fitted empirical data on environmental degradation versus national income. Most of the plots revealed an inverted U-pattern, as illustrated in Fig. 1.2. According to EKC hypothesis, environmental degradation increases with economic growth only in the least developed economies; at a certain degree of development, an inflection point exists (point A in Fig 1.2) and incomes are reverted in environmental protection. From this stage on, the detrimental effects of economic growth are surpassed by environmental improvements. From point B of Fig. 1.2 onwards, sustainable development is achieved.



**Figure 1.1. Total Primary Energy Supply of the World, European Union, United States of America and China. Graphs are based on data from IEA (2020).**

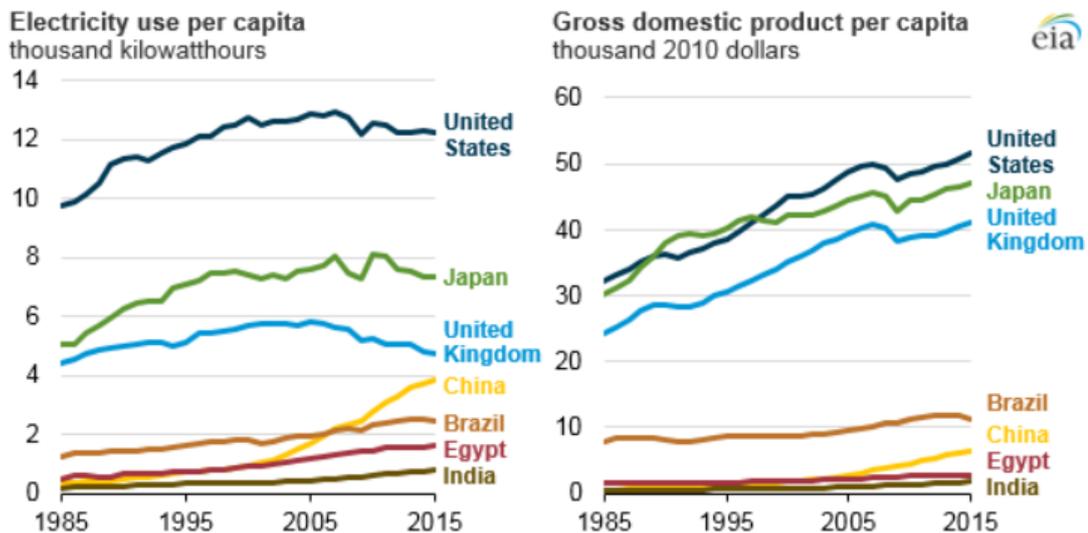


**Figure 1.2. The Environmental Kuznet Curve. Line A - inflection point. Line B - sustainable stage of economic development. Source: Vallero and Shulman (2019)**

Generally, the level of development of cities and countries is expressed by economic metrics, like gross domestic product (GDP), gross national income (GNI) or purchase power parity (PPP). Environmental degradation is usually expressed as a release rate or concentration of

pollutants in natural systems, e.g. particulate matter (PM) concentration or SO<sub>2</sub> flow rate of emissions in the air. However, many other metrics exist to represent the level of environmental degradation and economic development of a region or nation. In fact, sensitivities to localization (country, city) and time (year) do not empirically support the inverted U-patterned of EKC, and several relevant air pollutants do not follow EKC pattern (Harbaugh, Levinson and Wilson, 2002).

Destek and Sarkodie (2019) investigated the relationship between economic growth, energy consumption, financial development and ecological footprint from 1977 to 2013 in 11 developing economies. The results revealed that, although for certain countries the EKS shape is confirmed, for energy-intensive economies, like China and India, a U-shaped relationship occurred. The authors also found a one-way causality from energy consumption to ecological footprint. Energy is a major driver of economic development but also of environmental degradation. Seeking for economic growth and job creation least developed nations need to increase energy production while avoiding pollution, climate changes and waste generation. EIA (2017) data (Fig. 1.3) corroborates with the findings of Destek and Sarkodie (2019).

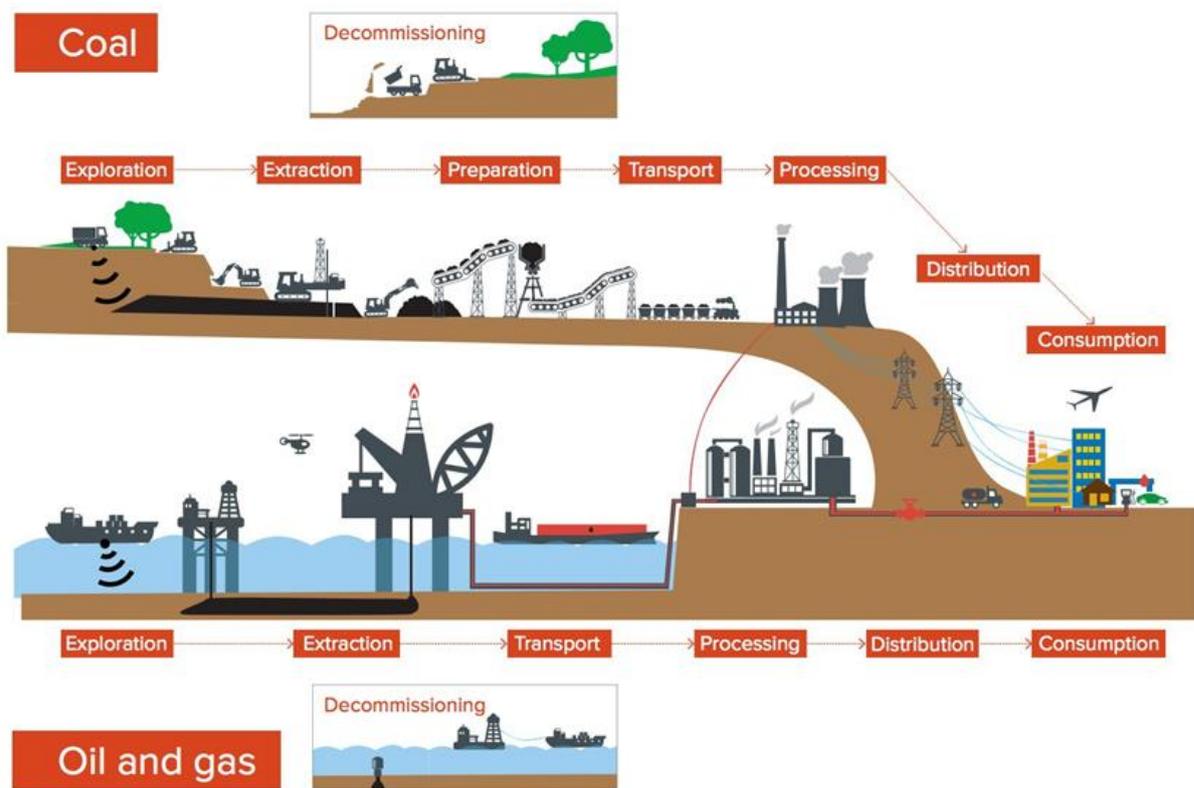


**Figure 1.3. Gross Domestic Product and Electricity Use (2011 – 2015) (EIA, 2017)**

As an example, in 2008 India had the highest air pollution in the world (Somani, 2013). Not coincidentally, the Indian economy has grown 6.14% per year between 1980 and 2008. India's TPES and energy use per capita increased quickly along these 3 decades pushing the environmental degradation. Fig. 1.3 shows that the developed countries were able to keep GDP growth while reducing the electricity use per-capita over the last 5 years. Differently,

developing countries have GDP growth followed by increased electricity use and, consequently, higher environmental degradation.

This scenario shows that fossil fuels will prevail in the TPES in the short to mid-term, demanding technologies to tackle climate change and other negative environmental impacts along the entire fossil-energy value chain, illustrated in Fig. 1.4. Emissions reduction should target reductions of the energy intensity of processes through enhancements of energy efficiency, mitigation of pollutants release and solid wastes management.



**Figure 1.4. Fossil Fuels Value Chain. (InfrastructureUSA, 2015)**

In this context, this thesis approaches technological advances in three different fronts within the fossil-energy chain: offshore primary processing of oil and gas; solid waste management of coal-fired power plants and carbon capture from flue-gases, i.e. post-combustion capture. It is worth noting that although post-combustion technologies are mostly employed in power plants it is applicable to other steps of the fossil-energy value chain, e.g. oil production and refining.

## 1.1. Objectives

Looking for enhancing sustainability of the fossil-energy supply, the objective of this thesis is to assess the technical, economic and/or environmental aspects of three technologies to reduce environmental impacts, respectively three research lines – R1, R2 and R3. This research aims at answering the questions: (i) Are the investigated technologies technically feasible? (ii) Are the technologies advantageous economic and/or environmentally? (iii) Are there bottlenecks or technological challenges to the commercial deployment of the technologies?

### 1.1.1. Specific Goals

#### ***R1 - Offshore Processing of CO<sub>2</sub>-Rich Natural Gas***

The goal of this research line is to develop and analyze two process modifications in a Floating Production Storage and Offloading (FPSO) unit processing CO<sub>2</sub>-rich natural gas: (R1.1) the use of deep seawater (DSW) as primary cooling utility and (R1.2) parallel-compressors scheme to avoid energy expenditure with anti-surge recycles. The main targets of these innovations are the assessment of technical and economic feasibility of the proposed modifications and the resulting reduction of CO<sub>2</sub> emissions. The implementation of an exergy analysis methodology customized to gas processing is a secondary specific objective, inherent to the topic (R1.2).

R1.1 objective is to evaluate the effects of using DSW as an indirect cooling utility in an FPSO. This alternative is compared to the traditional process, which uptakes warmer surface seawater. The comparison addresses energy consumption, economics and CO<sub>2</sub> emissions. More specifically, it seeks to compare:

- Compressor, gas-turbines (GT) and cooling-water (CW) pumping power;
- Heat exchangers duty;
- Equipment weight, equipment cost and revenue from natural gas (NG) production;
- Fuel-gas consumption and CO<sub>2</sub> emissions along the FPSO lifespan;

R1.2 aims to compare the exergy efficiency and power demand of an FPSO operating at peak and partial gas-loads using two process schemes. One with smaller parallel compressors and

variable-speed drive versus the traditional design, with anti-surge recycle. Detailed comparisons are:

- Exergy efficiency;
- Energy intensity, in barrels of oil equivalent (BOE) burned per BOE produced;
- Carbon dioxide intensity, in tons of CO<sub>2</sub> emitted per BOE produced;
- Fixed Capital Investment (FCI) and weight of topside equipment resulting from the process modification.

### ***R2 - Desulfurization Residues from Coal-Fired Power Plants***

This line aims to investigate the technical, economic and environmental aspects of implementing a thermal treatment of semi-dry Flue-Gas Desulfurization residue (SD-FGDR), a solid by-product contaminated with calcium sulfite (CaSO<sub>3</sub>). The proposed technology is conceived to avoid landfills occupied by coal ashes mixed with SD-FGDR. This environmental liability is transformed into a resource, used to produce cement. More detailed objectives are:

- Design and construction of a pilot-plant using a fluidized bed reactor (FBR), projected to oxidize CaSO<sub>3</sub> to CaSO<sub>4</sub> (gypsum). This reactor is used to obtain process parameters needed for scale-up of the technology.
- Economic feasibility analysis of the treatment retrofitted to a coal-fired power plant, including calculation of the final Levelized Cost of Electricity (LCOE).
- Comparison of the environmental performance of the power plant with and without the SD-FGDR treatment.

### ***R3 - CO<sub>2</sub> Capture from Flue-Gases by Phase-Changing Absorption Solvents***

The primary objective of this research line is to confirm the potential of energy saving of using phase-changing absorption solvents (PCAS) to capture CO<sub>2</sub> from flue-gases. To fulfill this goal, three major steps are pursued:

- Literature review, selection and preliminary tests of phase-changing absorption solvents (PCAS). At laboratory-scale, basic properties of the selected PCAS are analyzed, e.g. liquid phases behavior, CO<sub>2</sub> loading, density and viscosity.

- Design and construction of a bench-plant for solvent screening, that operates in batch-mode and enables fast absorption and desorption tests, using small volumes of solvent.
- Design of a pilot-plant, that operates in continuous-mode. This pilot-plant has the same layout of an industrial plant and would provide important process parameters, like the energy penalty (GJ/t of CO<sub>2</sub> captured). Future data produced during long run tests can be used to support process simulation and next scale-up design of the technology.

## 1.2. Justification

Given the potentially catastrophic impacts of climate change, CO<sub>2</sub> emissions have been the focus of environmental and energy-related developments. However, the fossil-energy value chain is responsible for several other negative environmental impacts, to the air, soil and water. It is important to develop and improve the technologies of extraction, processing and conversion of fossil fuels to reduce the environmental footprint of this energy source. The three proposed research lines – R1, R2 and R3 – are justified in the present section.

### *R1 - Offshore Processing of CO<sub>2</sub>-Rich Natural Gas*

Given the forecast of 25% increase on energy demand and replacement of coal by NG, the Oil & Gas industry will be in charge of 50% of the global TPES until 2040 (IEA, 2018a). According to the International Association of Oil and Gas Producers (IOGP, 2019), in 2018 the major oil and gas companies produced 1943 MMt of hydrocarbons (HC), with 55.2% of them came from offshore fields. This information justifies the choice and shows the relevance of the approached research lines.

Fossil fuels are believed to contribute to the global energy mix for decades to come, and to be the major energy source in the short to mid-term. Oil and gas (O&G) production from offshore reserves has emitted 115t of greenhouse gases (GHG) per 1000t of HC produced, totalizing 223.4 MMt CO<sub>2e</sub> in 2018. The IOGP (2019) informs that 71% of the CO<sub>2</sub> emissions (excluded methane) from the O&G industry are related to internal fuel combustion for energy production. Thus, given the environmental concerns and a carbon regulation scenario, it is of utmost importance to improve energy efficiency of O&G operations to reduce CO<sub>2</sub> emissions.

The Brazilian Pre-salt has challenging design conditions, such as the high gas to oil ratio (GOR) and CO<sub>2</sub>-rich associated gas. These issues render the primary processing of associated

gas more energy-intensive than in conventional fields. There requires CO<sub>2</sub> separation and injection for Enhanced Oil Recovery (EOR) and dispatch of preconditioned gas to onshore processing units, located 150-300km away, demanding high pressures, frequently, near 25000 kPa. These peculiarities strongly impact the cost and energy consumption of the FPSO gas processing plant, and drives research line R1.

## ***R2 - Desulfurization Residues from Coal-Fired Power Plants***

In the next decades, coal will have a major participation in the TPES of developing countries, especially in China and India. Consequently, post-combustion technologies to mitigate air pollution are demanded, notably for sulfur oxides, nitrous oxides and carbon dioxide removal from flue-gas. The first, Flue-gas Desulfurization (FGD), is a fingerprint of coal-fired power plants. Ash yield from coal combustion is between 3% to 49% (Pierce and Dennen, 2009) and some unit operations of power plants also produce solid by-products. Therefore, coal-fired power generation produces a massive amount of the so-called Coal Combustion Products (CCP). In 2017, the total production of CCP was around 1.1 billion metric tons (Harris, Heidrich and Feuerborn, 2019) The majority of CCPs (~64% in 2016) are considered resources and have several large-scale applications. If not utilized, they are piled up in landfills, becoming an environmental liability. Flue-gas desulfurization residues (FGDR) produced by coal-fired power plants can be used as raw materials, e.g. for civil and agricultural applications. However, some pollutants contained in the FGDR might contaminate the local environment, hindering their material reuse. (Phoungthong et al., 2018).

The FGD technology influences the residue composition and potential of utilization. The SD-FGDR is rich in sulfite (SO<sub>3</sub><sup>-</sup>), which limits its use. The ASTM Standard C618 (ASTM, 2015), for example, impose a limit on the SO<sub>3</sub><sup>-</sup> content to use fly-ash as a cement additive. Therefore, it is not possible to utilize fly-ash as a cement additive when it is mixed to SD-FGDR. Alternative uses of these contaminated ashes are limited. Frequently, large landfills are used to stock the contaminated coal ashes.

Around 40% to 60% of the desulfurization market runs in favor of SD-FGD and the rest uses the wet technology (Blankinship, 2005). As a result, the amount of SD-FGDR produced around the world is very expressive and justifies the research line R2.

### ***R3 - CO<sub>2</sub> Capture from Flue-Gases by Phase-Changing Absorption Solvents***

The increase of the CCUS capacity would make the necessity of fossil fuels cut less aggressive when aiming to meet the Paris Agreement global warming goals (Foster and Elzinga, 2015). Without CCUS the fossil energy production must be reduced by 50% until 2040. Considering a scenario where CCUS capacity reaches 7 Gt CO<sub>2</sub>/y until 2040 and 11 Gt CO<sub>2</sub>/y until 2100, this would be a reduction of 34% on fossil-energy cuts (Copenhagen Economics, 2017). In a more optimistic scenario, the Global CCS Institute (2016) predicts that a capacity of 4 GtCO<sub>2</sub>/y would be sufficient to keep global warming below 2°C. Anyway, CCUS is extremely important to limit global warming. However, the CCUS capacity barely reached 40 MMt/y in 2019 (Global CCS Institute, 2019).

The chemical absorption is currently the more mature technology for large scale post-combustion carbon capture. However, the energy spent to regenerate the solvent – the energy penalty – is considered a major hindrance to the deployment of this technology. The PCAS emerged as an opportunity to reduce this energy penalty and contribute to turning CCUS more economically feasible, increasing its installed capacity. These facts and arguments justify research line R3.

### **1.3. Outline of the Thesis**

This thesis is organized as an assemblage of published articles (Chapters 3, 5 and 6), a submitted article, currently under peer review (Chapter 4), and a conference paper (Chapter 7). A pilot-plant project is included as an additional section of Chapter 7. The research lines R1, R2 and R3 are developed according to the Table 1.1.

**Table 1.1. Correlation between scientific production and thesis content.**

Research Line	Chapters	References
(R1) Offshore Processing of CO <sub>2</sub> -Rich Natural Gas	3 4	(Cruz, Araújo and de Medeiros, 2018) Under review. Code <i>APEN-D-20-00835</i>
(R2) Desulfurization Residues from Coal-Fired Power Plants	5 6	(Cruz et al., 2017) (Cruz et al., 2018)
(R3) CO <sub>2</sub> Capture from Flue-Gases by Phase-Changing Absorption Solvents	7	(Cruz et al., 2019)

Figure 1.5 shows where each research line is supposed to be applied along the fossil energy value chain. R3, identified by blue asterisks in the Figure 1.5, is related to CCUS and could be applied to multiple steps of coal, oil and gas value chains.

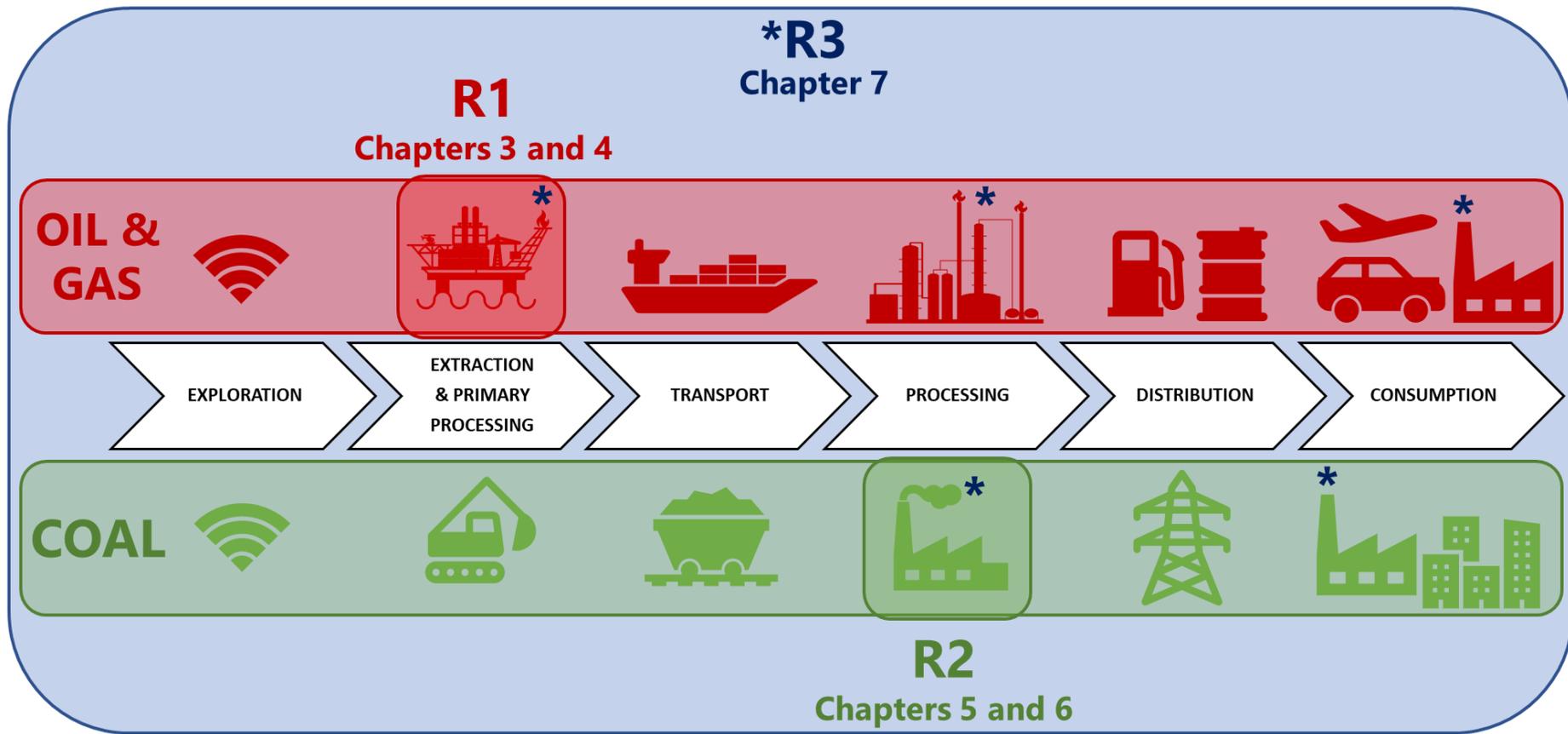


Figure 1.5. Distribution of Research Lines and Chapters of the Thesis Along the Fossil Fuels Value Chain.

Chapter 2 presents a literature review, approaching an overview of each proposed research line. This information supplies a macro perspective, that contextualizes and support the more focused literature review addressed in each chapter. Firstly, referring to R1, general information about the offshore O&G sector, CO<sub>2</sub>-rich fields, topside technologies and environmental constraints are presented. Secondly, referring to R2, a compilation of relevant information concerning coal-fired power plants and SD-FGD technologies are presented. Finally, general aspects of traditional post-combustion carbon capture and an extended review about PCAS are presented, complementing research line R3.

In Chapter 3, the article by Cruz, Araújo and De Medeiros (2018) is integrally reproduced, as published. It presents a comparative analysis of using colder DSW uptakes as a primary cooling utility in a FPSO against the traditional surface seawater uptake. The proposed process modification is compared to the base case in terms of power consumption, CO<sub>2</sub> emissions and detailed equipment sizing and cost estimation. Possible extra revenue from natural gas in consequence of fuel gas savings are also estimated.

Chapter 4 contemplates a process design innovation in an offshore CO<sub>2</sub>-rich natural gas processing scenario. The use of smaller parallel compressors and variable-speed drive is compared to the traditional layout with anti-surge recycle, at peak and partial gas-loads. A customized exergy analysis methodology is implemented to support the assessment. Processes are compared in terms of exergy efficiency, investment, footprint and emissions.

In Chapter 5, the published article of Cruz et al. (2017) is integrally reproduced. The study addresses a SD-FGDR treatment unit, to promote dry oxidation of calcium sulfite to calcium sulfate. A Brazilian coal-fired power plant facing decision-making process on SD- FGDR destination is regarded as the case study. The main equipment is sized and scaled-up based on the pilot-plant process parameter and patents of similar processes. An economic assessment is performed, including capital, operational and maintenance costs, residue revenue and LCOE. This information is used to determine the impact that the SD-FGDR treatment would imply on the electricity price if it were applied to the analyzed power plant.

In Chapter 6, the article of Cruz et al. (2018) is reproduced. To support decision-making on process configurations to monetize the SD-FGDR, a gate-to-gate assessment of potential environmental impacts is performed. Three scenarios are considered: BASE - the standard power plant, CASE I – the base plant with SD-FGDR treatment, CASE II – bypass of desulfurization system.

Chapter 7 is divided into two subsections. Section 7.1 reproduces the conference paper SDEWES2019.0276 (Cruz et al., 2019), entitled Chemical Absorption of CO<sub>2</sub> from Flue-Gases: Experiments with Phase Changing Solvents in a Bench Scale Plant, presented at the 14<sup>th</sup> Conference on Sustainable Development of Energy, Water and Environment Systems – Dubrovnik, 2019. This paper presents the state-of-the-art concerning PCAS and selects three of them to be tested in laboratory and in a solvent screening plant. The plant design is briefly presented in the article. One solvent, based on monoethanolamine and 1-propanol, was considered more suitable. Section 7.2 presents the process description, process flow diagram, P&ID, main equipment design and details about the control and automation system of a pilot-plant. The plant is designed to operate in continuous mode, aiming to support the scale-up of the technology to industrial applications.

Chapter 8 encompasses all the research lines presenting and discussing the overall and specific conclusions, findings and highlights of the thesis.

Appendices A to G unveil front pages and a complete bibliography of the publications of the Chapters 3 to 7. When existent, supplementary materials of each chapter are included in the Appendix H.

Appendices F and G unveil front pages and bibliography of further publications and co-authorships.

Appendix I presents design details, process specification, data sheets, draws and pictures of the main equipment of the pilot plant presented in Chapter 7, Section 7.2.

## **1.4. References of Chapter 1**

ASTM, 2015. C618-15, Standard Specification for Coal Fly Ash and Raw or Calcined Natural Pozzolan for Use. Annu. B. ASTM Stand. <https://doi.org/10.1520/C0618-15>

Blankinship, S., 2005. Looking for a Good Scrubbing: Today's FGD Technology [WWW Document]. Power Eng. URL <http://www.power-eng.com/articles/print/volume-109/issue-9/features/looking-for-a-good-scrubbing-todayrsquos-fgd-technology.html> (accessed 3.15.17).

Copenhagen Economics, 2017. The future of fossil fuels: How to steer fossil fuels use in a transition to a low-carbon energy system. An analysis of fossil fuels trajectories in low-carbon scenarios.

- Cruz, M. de A., Araújo, O.Q.F., de Medeiros, J.L., 2018. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. *J. Nat. Gas Sci. Eng.* 56, 193–211. <https://doi.org/10.1016/j.jngse.2018.06.011>
- Cruz, M. de A., Araújo, O.Q.F., de Medeiros, J.L., de Castro, R.D.P.V., Ribeiro, G.T., de Oliveira, V.R., 2017. Impact of solid waste treatment from spray dryer absorber on the levelized cost of energy of a coal-fired power plant. *J. Clean. Prod.* 164. <https://doi.org/10.1016/j.jclepro.2017.07.061>
- Cruz, M. de A., Musse, A.P.S., de Medeiros, J.L., Araújo, O.Q.F., 2019. SDEWES2019.0276 Chemical Absorption of CO<sub>2</sub> from Flue Gases: Experiments with Phase Changing Solvents in a Bench Scale Plant, 14th Conference on Sustainable Development of Energy, Water and Environment Systems. Faculty of Mechanical Engineering and Naval Architecture, Zagreb.
- Cruz, M.D.A., de Castro, R. de P.V., Araújo, O. de Q.F., de Medeiros, J.L., 2018. Environmental Performance of a Solid Waste Monetization Process Applied to a Coal-Fired Power Plant with Semi-Dry Flue Gas Desulfurization. *J. Sustain. Dev. Energy, Water Environ. Syst.* 7, 506–520. <https://doi.org/http://dx.doi.org/10.13044/j.sdewes.d6.0251>
- Destek, M.A., Sarkodie, S.A., 2019. Investigation of environmental Kuznets curve for ecological footprint: The role of energy and financial development. *Sci. Total Environ.* 650, 2483–2489. <https://doi.org/https://doi.org/10.1016/j.scitotenv.2018.10.017>
- Foster, S., Elzinga, D., 2015. The Role of Fossil Fuels in a Sustainable Energy System. *UN Chron.* LII.
- Global CCS Institute, 2019. The Global Status of CCS: 2019, Global CCS Institute. Melbourne, Australia. [https://doi.org/10.1007/springerreference\\_15392](https://doi.org/10.1007/springerreference_15392)
- Global CCS Institute, 2016. The Global Status of CCS: 2016. Summary Report. Global CCS Institute, Docklands, AU.
- Harbaugh, W.T., Levinson, A., Wilson, D.M., 2002. Reexamining the Empirical Evidence for an Environmental Kuznets Curve. *Rev. Econ. Stat.* 84, 541–551. <https://doi.org/10.1162/003465302320259538>
- Harris, D., Heidrich, C., Feuerborn, J., 2019. Global aspects on Coal Combustion Products [WWW Document]. *Coaltrans Conf.* URL <https://www.coaltrans.com/insights/article/global-aspects-on-coal-combustion-products> (accessed 2.4.20).
- IEA, 2020. Data & Statistics - IEA [WWW Document]. Int. Energy Agency. URL [https://www.iea.org/data-and-statistics?country=EU28&fuel=Energy supply&indicator=Total primary energy supply \(TPES\) by source](https://www.iea.org/data-and-statistics?country=EU28&fuel=Energy supply&indicator=Total primary energy supply (TPES) by source) (accessed 1.31.20).
- IEA, 2019. CO<sub>2</sub> emissions from fuel combustion. France.
- IEA, 2018. 2018 World Energy Outlook: Executive Summary. *Oecd/Iea.*

- InfrastructureUSA, 2015. Fossil Fuel Subsidy Reform: From Rhetoric to Reality [WWW Document]. InfrastructureUSA. URL <https://www.infrastructureusa.org/fossil-fuel-subsidy-reform-from-rhetoric-to-reality/> (accessed 3.2.20).
- IOGP, 2019. IOGP Environmental performance indicators - 2018 data - Atmospheric emissions Int. Assoc. Oil Gas Prod. URL <https://data.iogp.org/Environment/Emissions> (accessed 2.4.20).
- IPCC, 2014. Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, Intergovernmental Panel on Climate Change. IPCC, Geneva, Switzerland.
- Phoungthong, K., He, P.-J., Shao, L.-M., Zhang, H., 2018. Leachate phytotoxicity of flue gas desulfurization residues from coal-fired power plant. *Environ. Sci. Pollut. Res.* 25, 19808–19817. <https://doi.org/https://doi.org/10.1007/s11356-018-2207-8>
- Pierce, B.S., Dennen, K.O., 2009. The National Coal Resource Assessment Overview. U.S. Geol. Surv. Prof. Pap. 1625, 402.
- Somani, A.K., 2013. Environmental Tax Reform and Economic Welfare. Harvard.
- Stern, D.I., Common, M.S., Barbier, E.B., 1996. Economic growth and environmental degradation: The environmental Kuznets curve and sustainable development. *World Dev.* 24, 1151–1160. [https://doi.org/10.1016/0305-750X\(96\)00032-0](https://doi.org/10.1016/0305-750X(96)00032-0)
- United Nations, 2015. Transforming our World: The 2030 Agenda for Sustainable Development. New York. <https://sustainabledevelopment.un.org/post2015/transformingourworld>.
- U.S. EIA, 2017. Link between growth in economic activity and electricity use is changing around the world. U.S. Energy Inf. Adm. URL <https://www.eia.gov/todayinenergy/detail.php?id=33812> (accessed 1.31.20).
- Vallero, D.A., Shulman, V., 2019. Chapter 1 - Introduction to Waste Management☆, in: Letcher, T.M., Vallero, D.A.B.T.-W. (Second E. (Eds.), . Academic Press, 3–14. <https://doi.org/https://doi.org/10.1016/B978-0-12-815060-3.00001-3>
- Yue, X.L., Gao, Q.X., 2018. Contributions of natural systems and human activity to greenhouse gas emissions. *Adv. Clim. Chang. Res.* 9, 243–252. <https://doi.org/10.1016/j.accre.2018.12.003>

## 2. LITERATURE REVIEW

### 2.1. R1 – Offshore Primary Processing of CO<sub>2</sub>-Rich Natural gas

In 2015, members of the United Nations adopted the Sustainable Development Goals (SDGs) established in the 2030 Agenda for Sustainable Development (United Nations, 2015). The agenda, including SDGs, comprehends a global action plan for social inclusion, environmental sustainability and economic development. The IPIECA, a nonprofit association that promotes the continuous improvement in the O&G sector, published an Atlas (IPIECA, 2017) that lists the positive and negative impacts of this industry within the SDGs. This Atlas also shows initiatives of the sector to contribute to achieving each SDGs. Among several actions, two of them are related to the research line R1 developed in this thesis. Regarding the SDG 7 (affordable and clean energy) the atlas suggests: “7. *By 2030, double the global rate of improvement in energy efficiency; 7.a By 2030, enhance international cooperation to facilitate access to clean energy research and technology, including renewable energy, energy efficiency and advanced and cleaner fossil-fuel technology, and promote investment in energy infrastructure and clean energy technology* (IPIECA, 2017, p. 37).

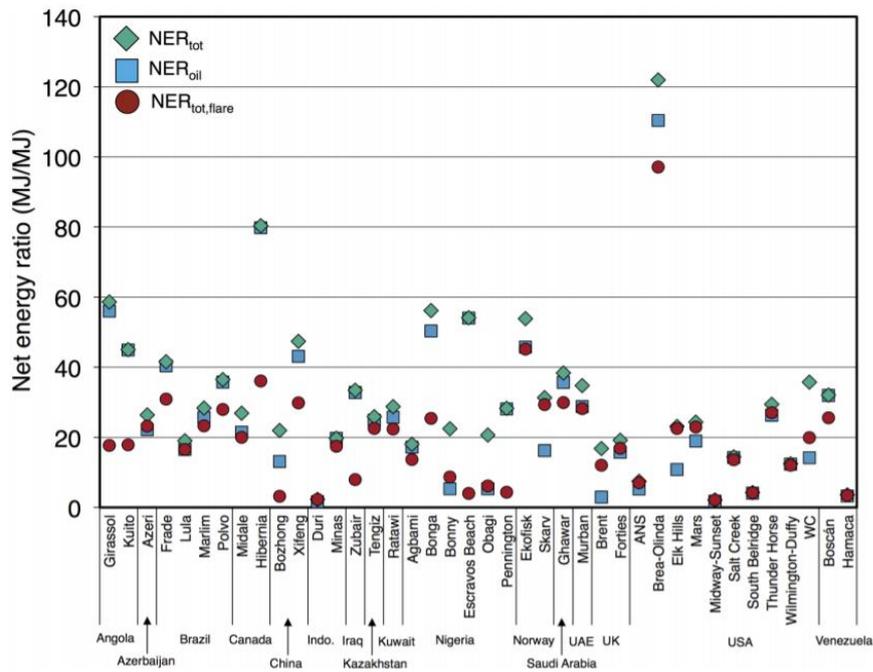
According to IOGP (2019), 71% of the CO<sub>2</sub> emissions (excluded methane) of the O&G sector are related to internal energy production. In 2011, extraction and transformation of hydrocarbons consumed 6.9% of the energy produced from themselves (IPIECA, 2017) and 3% to 4% of the global TPES (Masnadi et al., 2018). Based on these facts, the IPIECA highlights the importance of improving energy efficiency of O&G production operations, as an alternative to mitigate energy scarcity and GHG emissions worldwide. In this direction, oil companies have worked along the last decades to reduce their flaring, venting and fugitive emissions. Further efforts are needed to reduce energy losses and increase energy efficiency.

#### 2.1.1. Energy and Carbon Intensity of Oil & Gas Production

The oil industry targets to profit from the exploration, production and delivery of energy-carrying substances separated from crude oil – the “energy target” – and supply the net energy produced to society. To discover, extract, process, transport and refine oil into products consumes a large fraction of the energy available in the primary source. Meeting the “energy

target”, it is important to measure the energy productivity along the energy supply chain. Amongst many proposed energy return ratios (ERR), energy return on investment (EROI) and net energy ratio (NER) are commonly used to give assessing productivity of energy supply chain. The ERR is sensitive to the primary energy source characteristics. In the O&G industry, relevant characteristics, among others, are GOR, water to oil ratio (WOR), oil density (e.g. °API) and water depth, demanding a bottom-up engineering-based analysis of oilfield operations (Brandt et al., 2015).

Brandt et al. (2015) performed a bottom-up analysis of 40 oil fields around the world. The study employed 3 types of NER:  $NER_{oil}$ ,  $NER_{tot}$  and  $NER_{tot,flare}$ . The first considers only the energy consumed by oil-related operations; the second includes NG, condensates and internal electricity production operations; and the last includes gas flaring. The production-weighted mean  $NER_{tot}$  for all fields is 32.5 MJ/MJ and varied within the range of 10–35 MJ/MJ, excluding fields using thermal EOR. Fig 2.1 shows the regional variation of NERs.



**Figure 2.1. Net energy ratio ( $NER_{tot}$  and  $NER_{oil}$ ) of oilfields. Source: Brandt et al. (2015).**

From Fig 2.1 it can be noticed that the pre-salt Lula field has a lower  $NER_{tot}$  compared to the other Brazilian fields (Marlim, Polvo and Frade). Pre-salt has an exceptionally large GOR and exhibits  $CO_2$ -rich associated NG. Additionally, flaring or venting NG or  $CO_2$  is prohibited by local regulations, to mitigate GHG emissions. Consequently, the current pre-salt O&G operation is to separate  $CO_2$  from  $CO_2$ -rich NG, compress the  $CO_2$ -rich streams and used as

EOR fluid. The pre-treated NG is compressed and transported to onshore processing facilities. The primary conditioning of the CO<sub>2</sub>-rich NG reduces the ERRs of the pre-salt oilfields. When flaring is included, many fields present expressively lower NERs. Table 2.1 shows the influence of oilfield characteristics separated by groups. Fields with high WOR and depth presented lower NER<sub>tot</sub> because of the energy required by lifting; heavy and ultra-heavy oil have even lower NERs, due to the use of thermal recovery; and the thermal recovery subgroup presented the lowest NER<sub>tot</sub>. (Brandt et al., 2015).

**Table 2.1. Total Net Energy Ratio of oilfields. Adapted from Brandt et al. (2015)**

	<b>N° of Fields</b>	<b>Production-Weighted Mean</b>
All fields	40	32.5
High WOR (> 10 bbl water/bbl oil)	6	12.3
Deep (10000ft < depth <15,000ft)	12	29.7
Ultra-deep (depth >15,000ft)	3	22.3
Old (> 40 years old fields)	15	35.9
Heavy oil (15° < API gravity < 22.5°)	10	17.7
Ultra-Heavy oil (API gravity < 15°API)	3	10.6
Thermal EOR	3	2.8

Masnadi et al. (2018) developed and applied a methodology to determine the carbon intensity and well-to-refinery life cycle emissions of 8966 oil fields in 90 countries (98% of oil and condensate production in 2015). The study accounts for GHG emissions of main upstream activities (exploration, drilling and development, production and extraction, surface processing, and transport to the refinery inlet). The summary of this study is illustrated in Fig. 2.2. Given the impossibility of sudden cut on the global oil and gas consumption – unforeseen pandemic crisis of 2020<sup>1</sup>, Masnadi et al. (2018) highlighted three main strategies to reduce the carbon intensity of the upstream activities: resource management, resource prioritization, and innovative technologies. The last one is explored through the research line R1 of this thesis, taking the CO<sub>2</sub>-rich fields of Pre-salt as a scenario.

<sup>1</sup>In the 2020 pandemic, oil and gas suffered a sudden cut.

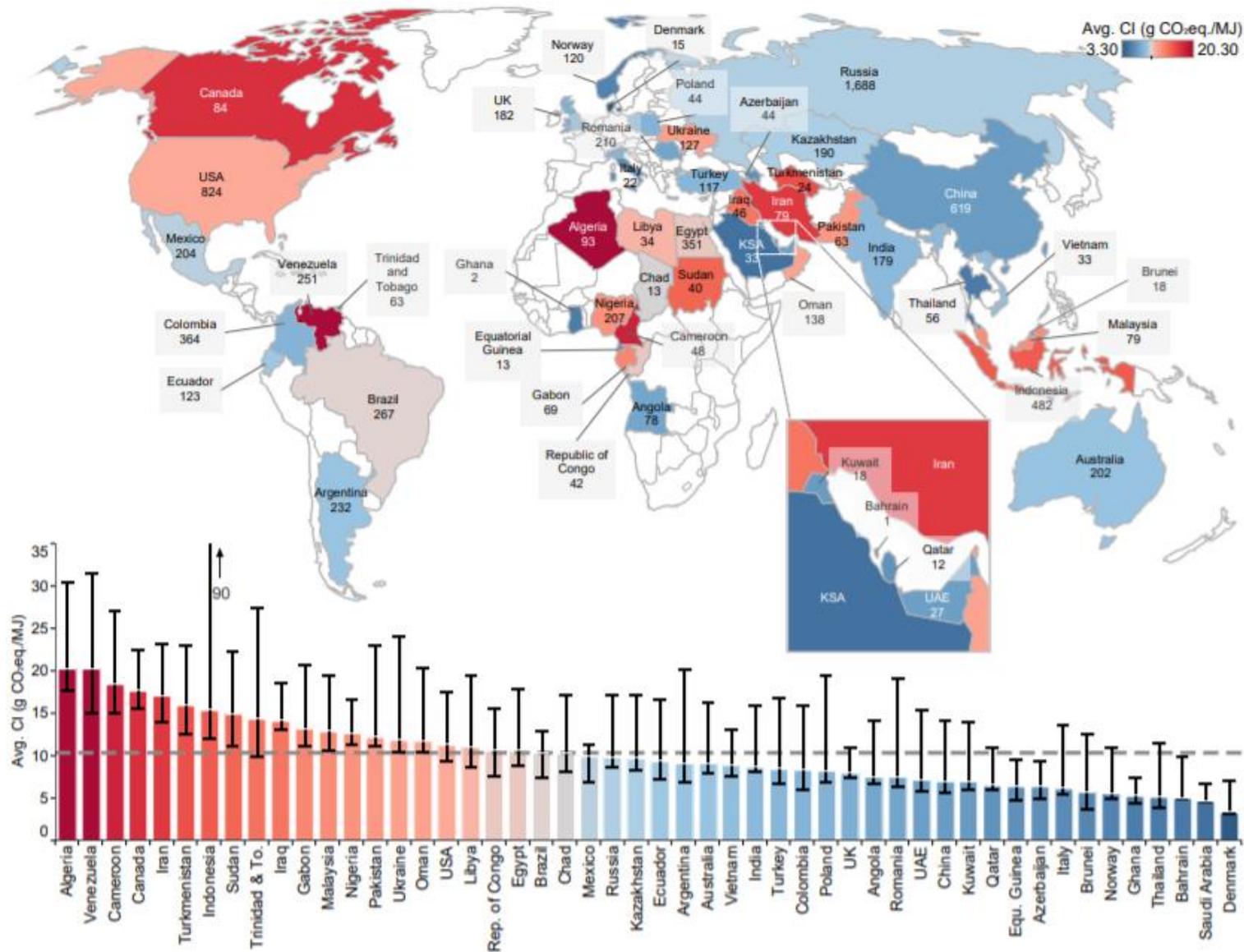


Figure 2.2. Global upstream crude oil carbon intensity (2015). (Masnadi et al., 2018)

### 2.1.2. CO<sub>2</sub>-rich fields implication on FPSO design and energy intensity

The Pre-salt region holds the largest Brazilian oil fields. The fields are located between 150km – 300km far from the coast and the water depth ranges around 1500m – 3000m (ultra-deep waters). The total proven reserves of Petrobras reached 9590 MMboe in 2019. This data did not take into account the giant field of Búzios with reserves estimated in 10000 MMboe). (Petrobras, 2020) In January of 2020, the Pre-salt production reached 2150 MMbbl/d of oil and 84572 MMSm<sup>3</sup>/d of NG, totalizing 2682 MMboe/d. This production corresponded to 66.4% of the Brazilian production of oil and gas (ANP, 2020).

Because of the high GOR and CO<sub>2</sub> content found in pre-salt reservoirs, to produce oil a massive flow rate of CO<sub>2</sub>-rich gas must be separated and treated, enabling injection of the CO<sub>2</sub>-rich stream (EOR) and export of the surplus of pre-treated NG. These constraints impose unusual state-of-the-art production practices like strict control of water dew point (WDP, < 1ppm) and hydrocarbon dew point (HCDP, < 1000ppm); CO<sub>2</sub> separation to achieve a concentration < 3% on NG, and compression to high pressures (25000kPa to 55000kPa). These features demanded new technologies and improvements in the FPSO design and raised the energy demand, carbon intensity and topside equipment weight and cost to unprecedented levels. (Araújo et al., 2017a)

Amongst several advantages, FPSOs are the preferred alternative for remote fields because of its high oil storage capacity and wide deck area to support topside facilities. The largest FPSOs are converted from very large crude carriers (VLCC). These converted FPSOs usually produce up to 180000 bbl/d of oil and processes up to 11 MMSm<sup>3</sup>/d of gas with topsides weighing up to 35000t (MODEC, 2019). However, the oil and gas industry demands, especially from pre-salt operators, keep increasing in terms of larger and heavier topsides; greater oil storage and gas processing capacity, accommodations space and extended lifetime (NS Energy, 2019). The increased complexity of primary processing operations also increased the cost of the FPSOs, penalizing offshore production projects. Therefore, the main FPSO suppliers were forced to replace VLCCs for improved and larger hulls.

Concerning market competitiveness, Engineering, Procurement, Construction and Installation (EPCI) companies developed new standardized hulls and topside modules. Standardizing resulted on savings in the supply chain and construction phases shortening the delivery time

and investment cost of the FPSO (SBM Offshore, 2019), provided financial gains from the earlier start of operations, greater safety and quality.

MODEC (2019) launched the M350 hull, addressing shipbuilding standards:

- Dimensions: 350 m long, 64 m wide, 33 m molded depth;
- Total oil storage capacity of 350000 m<sup>3</sup> (2.2 MMbbl);
- 20% larger deck area for topsides, compared to a VLCC, supporting up to 50,000t;
- Increased safety and lifetime of 25 years due to double hull (double sides and double bottom walls);
- Accommodation for 160 workers, and helicopter parking.

MODEC recently signed a sales and purchase agreement to supply the largest FPSO ever delivered to Brazil, Fig. 2.3. The FPSO will be the second application of the M350 hull. This vessel will be deployed at Bacalhau, Block BM-S-8, in the Brazilian pre-salt., with capacity of oil 220,000 bbl oil/d, 2,000,000 bbl of storage capacity and gas processing plant capacity of 15 MMSm<sup>3</sup>/d, almost threefold higher than the average capacity of an FPSO converted from VLCC (MODEC, 2020).



**Figure 2.3. Preliminary Layout of FPSO “Bacalhau” (MODEC, 2020).**

Following the same strategy of MODEC, SBM launched the FAST4WARD standardized design (SBM Offshore, 2019), shown in Fig. 2.4, with the following features:

- New-built hull designed for a 30-year lifetime and oil storage capacity of up to 2.3 MMbbl.

- 13% larger deck space compared to a VLCC, with 30% more topside footprint, accommodating up to 50,000t.



**Figure 2.4. SBM FAST4WARD FPSO (SBM Offshore, 2019).**

Although the new hull designs provide more space and weight to accommodate topside equipment, the extra energy expended to treat extremely high flow rates of CO<sub>2</sub>-rich associated gas remains an environmental challenge. For layout simplification and footprint saving, most FPSOs use simple gas turbines to generate power and heat. This generation scheme has low-efficiency and contributes to increasing the carbon intensity of O&G production. Another villain of offshore production sustainability is the use of oversized compressors to match design gas-loads. Most of the time, the gas processing plant operates at partial load and much energy is wasted in anti-surge recycle loops, especially under reduced gas production. A more specific literature review, regarding the investigated technologies to reduce the energy and carbon intensity of the primary processing of CO<sub>2</sub>-rich NG is presented in chapters 3 and 4.

## **2.2. R2 – Desulfurization of Flue-Gases from Coal-Fired Power Plants**

### **2.2.1. Coal-Fired Power Plants**

Coal-fired power plants supplied 41% of the electricity demand worldwide in 2016 (WCA, 2017). In the last 20 years, Asia was responsible for 90% of the increase in the global coal-fired capacity. These power plants have an average of 12 years of operation and potentially longer operational lifetime ahead (IEA, 2019b). Coal is responsible for around 25% of the global TPES, but, in some developing nations, coal share is considerably higher, e.g. > 60% in China

(IEA, 2020). Besides GHG (e.g. CO<sub>2</sub> and N<sub>2</sub>O), combustion of fossil fuels releases SO<sub>2</sub>, NO<sub>x</sub>, CO, particulate matter (PM), heavy metals, halide compounds, and dioxins into the atmosphere. SO<sub>2</sub> is formed if sulfur is present in the fuel.

Treated NG is often free of sulfur, coal has a significant content of this element (European Commission, 2006). SO<sub>2</sub> emissions cause detrimental impacts on the environment and to human health. Exposure to high concentrations of SO<sub>2</sub> is linked to respiratory and cardiovascular diseases. SO<sub>2</sub> also leads to acid deposition in the environment and consequent acidification of water bodies and damage to natural vegetation, crops, buildings and monuments (Srivastava and Jozewicz, 2001).

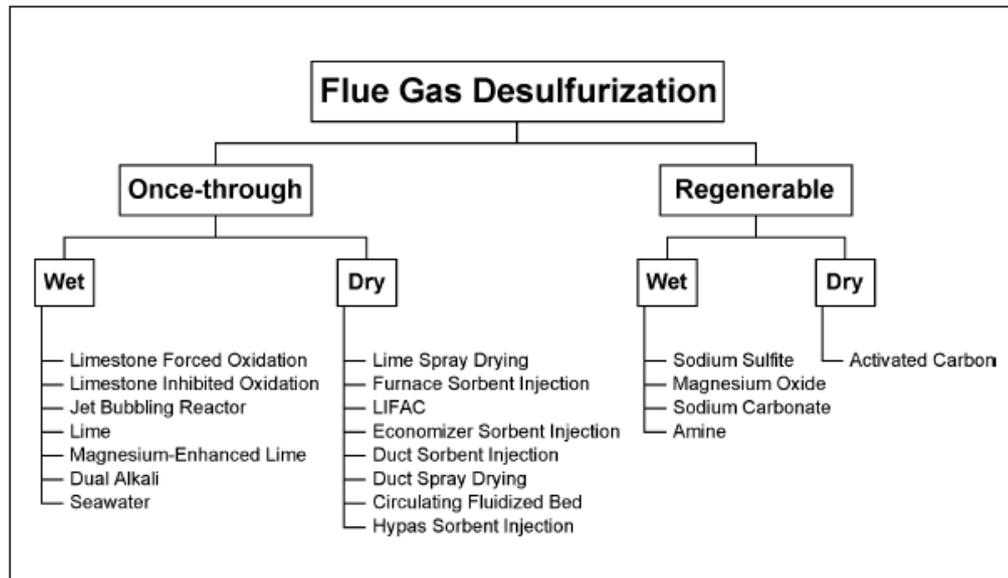
In the face of these impacts, most countries adopted strict limits on SO<sub>2</sub> emissions from power plants. The United States launched initiatives like the Clean Air Act Amendments (CAAA), which includes the Acid Rain SO<sub>2</sub> Reduction Program, focused on reducing SO<sub>2</sub> emissions from thermal power stations (Srivastava and Jozewicz, 2001). To comply with the SO<sub>2</sub> emissions limit, besides using low-sulfur fuel, FGD is necessary. In the European Union, the use of FGD technologies is mandatory for plants with more than 100 MW of capacity. SO<sub>2</sub> emissions limit and recommended FGD technologies are shown in Fig. 2.5 (European Commission, 2006).

Capacity (MW <sub>e</sub> )	SO <sub>2</sub> emission level (mg/Nm <sup>3</sup> )						BAT to reach these levels
	Coal and lignite		Peat		Liquid fuels for boilers		
	New plants	Existing plants	New plants	Existing plants	New plants	Existing plants	
50 – 100	200 – 400* 150 – 400* (FBC)	200 – 400* 150 – 400* (FBC)	200 – 300	200 – 300	100 – 350*	100 – 350*	Low sulphur fuel or/and FGD (dsi) or FGD (sds) or FGD (wet) (depending on the plant size). Seawater scrubbing. Combined techniques for the reduction of NO <sub>x</sub> and SO <sub>2</sub> . Limestone injection (FBC).
100 – 300	100 – 200	100 – 250*	200 – 300 150 – 250 (FBC)	200 – 300 150 – 300 (FBC)	100 – 200*	100 – 250*	
>300	20 – 150* 100 – 200 (CFBC/ PFBC)	20 – 200* 100 – 200* (CFBC/ PFBC)	50 – 150 50 – 200 (FBC)	50 – 200	50 – 150*	50 – 200*	
Notes: FBC: Fluidised bed combustion PFBC: Pressurised fluidised bed combustion FGD(sds): Flue-gas desulphurisation by using a spray dryer FGD(dsi): Flue-gas desulphurisation by dry sorbent injection * Some split views appeared in these values and are reported in Sections 4.5.8 and 6.5.3.3 of the main document.							

**Figure 2.5. European Union SO<sub>2</sub> emissions limit and recommended technologies (European Commission, 2006).**

## 2.2.2. Flue-gas Desulfurization Systems

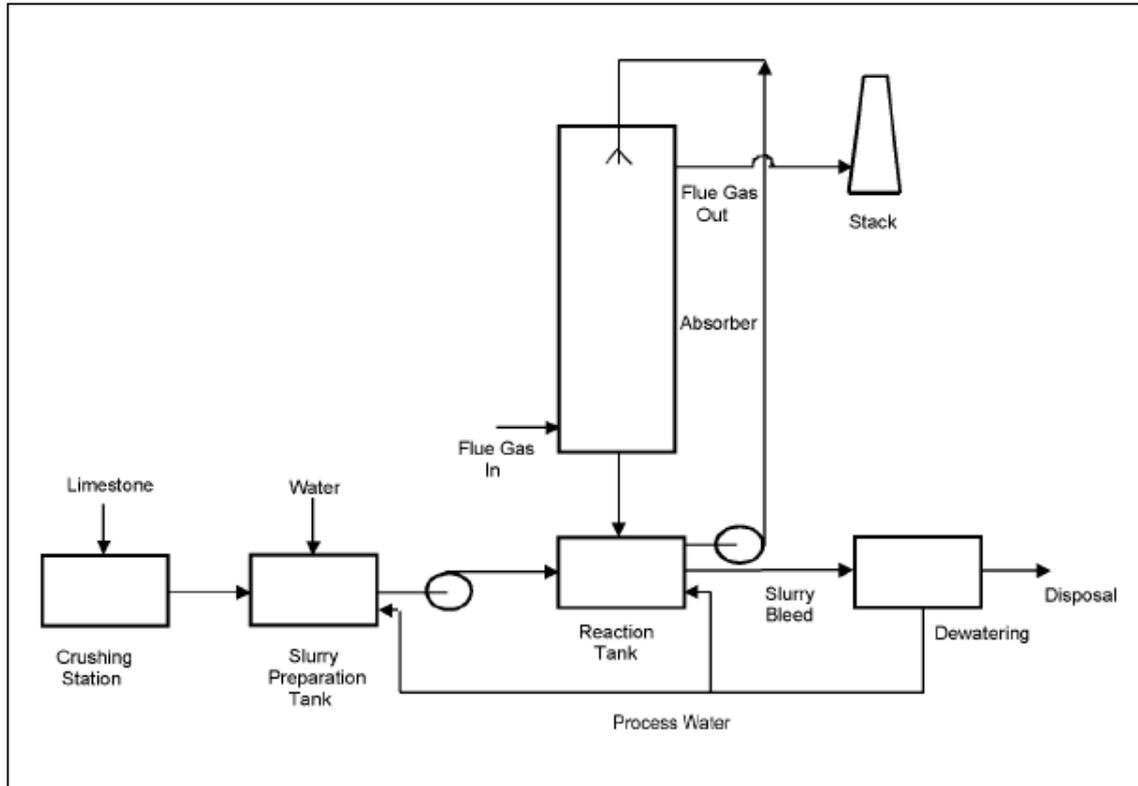
Srivastava and Jozewicz (2001) classified the FGD technologies according to the sorbent destination after reacts with SO<sub>2</sub>, as shown in Fig 2.6. Once-through systems consider spent sorbent as waste or byproduct. Regenerable processes have an additional step to remove SO<sub>2</sub> from the spent sorbent. Usually, SO<sub>2</sub> is used to produce sulfuric acid or sulfur.



**Figure 2.6. Classification of FGD technologies(Srivastava and Jozewicz, 2001).**

The Best Available Technologies (BAT) for desulfurization are wet and spray dry scrubbing, with a market share of more than 90%. Wet-FGD has SO<sub>2</sub> removal rate of 92 – 98 % while spray dry scrubber (a SD-FGD technology) has a slightly lower reduction rate of 85 – 92 % (European Commission, 2006). The selection of the FGD technology must consider several factors like power plant capacity, sulfur content of the fuel, capital expenditure (CAPEX), operational expenditure (OPEX), and space availability, sorbent material and water (Ma et al., 2000). In general, SD-FGD is adopted for power plants with less than 300 MW of thermal capacity. Wet-FGD is not considered for plants with less than 100 MW of capacity, due to economic unfeasibility (European Commission, 2006).

The wet-FGD, shown in Fig. 2.7, is the most widely used FGD process, holding more than 80% of the market share. It has a higher performance and lower cost compared to other available technologies. However, its specific water consumption is higher, reaching 250 l/MWh in a subcritical plant. (Carpenter, 2012) Regarding the solid byproducts, the wet-FGD has the advantage to form gypsum (CaSO<sub>4</sub>), which has several industrial applications.



**Figure 2.7. Wet FGD Process (Srivastava and Jozewicz, 2001).**

Eq. 2.2.1 is the overall reaction of the wet-FGD system, occurring in 4 steps (Eqs. 2.2.1a to 2.2.1e).



*SO<sub>2</sub> Absorption:*



*Limestone dissolution:*



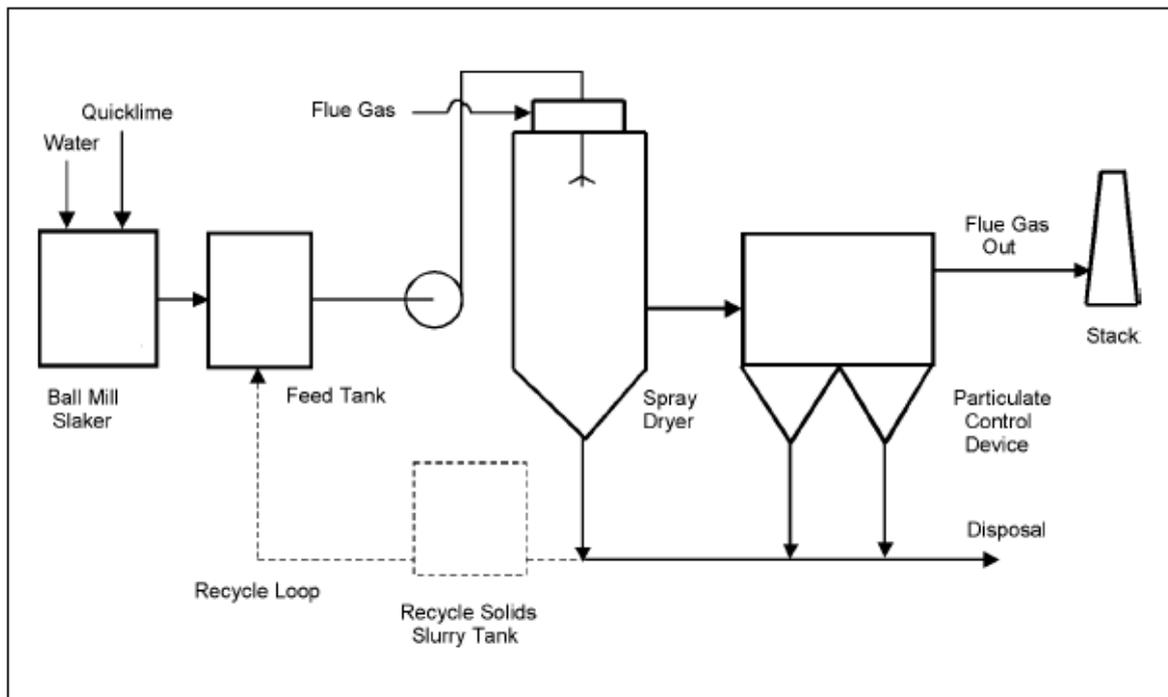
*Oxidation:*



*Precipitation:*



SD-FGD is usually applicable for small to medium-sized power plants (Dehghani and Bridjanian, 2010). This technology, illustrated in Fig. 2.8, trades off efficiency, water use and cost (Sage and Ford, 1996).



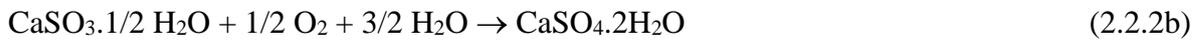
**Figure 2.8. Semi-dry FGD Process (Srivastava and Jozewicz, 2001).**

The water demand of SD-FGD is 60% lower compared to the humid route. This characteristic makes the semi-dry alternative more attractive in regions under water scarcity, such as the Northeast of Brazil and the Western of the USA. The semi-dry technology installed capacity ranks second worldwide, corresponding to approximately 15% of market share (Carpenter, 2012). However, the application of SD-FGD is limited to low sulfur coal (0% to 3%) (Funch-Jensen and Rubner-Petersen, 2007)

Usually, SD-FGD units have spray dryer vessels where the hot flue-gas contacts a mist of atomized fresh lime slurry (Hill and Zank, 2000). Simultaneous heat and mass transfers remove  $\text{SO}_2$  from the flue-gas, according to the Eq. 2.2.2a, and dry the products. The final product is a powder, named desulfurization residue (Dehghani and Bridjanian, 2010).



Because the reaction occurs in an atmosphere poor in oxygen and water, instead of forming gypsum, calcium sulfite ( $\text{CaSO}_3$ ) predominates. Just a small fraction of calcium sulfite should be oxidized to gypsum, according to the Eq. 2.2.2b.



Research line R2 develops a technology for utilization of desulfurization residue targeting its use in the cement industry.

## **2.3. R3 – CO<sub>2</sub> Capture from Flue-Gases by Phase-Changing Solvents**

### **2.3.1. Greenhouse Gases Emissions from Fossil Energy**

From 2010 to 2017, the energy sector (electricity, heat generation and transport) accounted for more than two-thirds of total anthropogenic carbon dioxide emissions. The other third was caused by industry and building consumption (IEA, 2019a).

The Global Carbon Budget 2019 (Friedlingstein and Al, 2019) estimates that, from 1850 and 2018, 645 Gt of carbon (GtC) were emitted by human activities. Fossil fuels contributed with almost 66% of these emissions, as depicted in Fig 2.9. In 2018, the GHG emissions from fossil-fueled energy reached 10 GtC/y (Friedlingstein and Al, 2019). The environmental impact of electricity and heat consumption depends on the energy source. Developed countries' share of clean energy from renewable sources (e.g. wind and solar) are increasing lately. However, developing countries, especially China, are still dependent on fossil and low-efficient energy production. The result of such disparity is shown in Fig. 2.10.

CO<sub>2</sub> emissions reached 32.8 billion tons in 2017, increasing after three years of stability. In 2018, the increase accelerated due to global economic growth and a slower pace of penetration of renewables (IEA, 2019a). Atmospheric carbon dioxide concentration reached 411 ppm in December of 2019 (NOAA, 2020), and is currently 40% higher than in the pre-industrial era.



In 2015, the Paris Agreement established targets of maximum temperature increase of 1.5°C to 2°C until the end of the century, compared to the pre-industrial era (UNFCCC, 2016). These targets resulted in a “*carbon budget*”, which refers to the amount of carbon that could be released until 2100 given the considered temperature target. According to IPCC (2014), the *carbon budget* is around 2900 GtCO<sub>2e</sub> and 65% of this budget was already consumed by 2011, and the proven fossil fuel reserves exceed the carbon budget by 3 to 6 times. For a 1.5°C target, 200 GtCO<sub>2e</sub> could be released to the atmosphere; hence, the current GHG emissions rate should be halved until 2040, despite the global energy demand being expected to increase by 30% to 75% in this period. Clearly, a fast transition from current carbon-intensive energy to a low carbon system (zero or negative emissions) is required (Copenhagen Economics, 2017).

Increased energy efficiency is essential for a quick, feasible and less economically damaging transition (IEA, 2018b). However, most of the current conventional energy production systems are optimized from the process perspective. Opportunities for feasible energy efficiency increase are scarce. Renewable energy sources are claimed to be the ultimate solution to global warming, and its share has been growing worldwide, especially in the European Union. Despite the great environmental advantages, currently, a 100% renewable grid is technically unfeasible. Not to mention the economic issues, the inherent intermittency of wind and solar energies and low energy storage capacity are technical hindrances to renewables’ growth. Thus, fossil fuels are expected to remain in the global energy matrix in the mid-term and CCUS is required to achieve negligible or negative emission, as required to meet Paris Agreement goals. (Global CCS Institute, 2019)

### 2.3.2. Post-Combustion Carbon Capture by Chemical Absorption

The International Energy Agency considers retrofit with CCUS as one of three options to cut GHG emissions from the current 2080 GW of coal-fired installed power capacity. The other options are retrofit with biomass co-firing equipment or prematurely retire coal-based power systems (IEA, 2019b), being these two alternatives more challenging technically and harmful economically. Over the last 20 years, CCUS has evolved from an option to a necessity to mitigate climate change (Global CCS Institute, 2019).

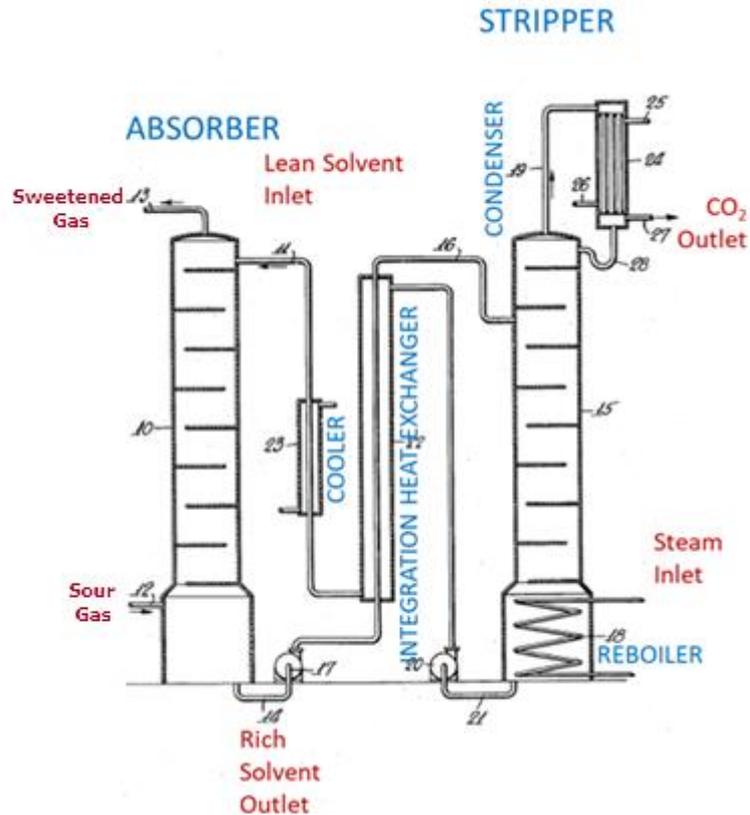
Much of the useful energy (heat and power) production is centralized in thermal power plants, where fossil-fuel combustion produces gases – flue-gases, with contaminants, including CO<sub>2</sub>. The flue-gases are treated and emitted to the atmosphere through chimneys. Flue-gases from

power plants usually have low pressure ( $< 150$  kPa) and moderate temperature ( $50^{\circ}\text{C} - 100^{\circ}\text{C}$ ). Post-combustion carbon capture technologies aim at mitigating  $\text{CO}_2$  emissions from flue-gases before discharging it into the atmosphere. Chemical or physical absorption are the technologies closest to full-scale availability, favoring retrofitting (Araújo and de Medeiros, 2017). However, there are emerging separation technologies, e.g. membrane contactors (de Medeiros et al., 2013; Liu et al., 2020), electrochemical membranes (Tong et al., 2015), or hybrid systems (Frimpong et al., 2019).

Gas-liquid absorption is a preferred choice for post-combustion  $\text{CO}_2$  capture because reactive solvents efficiently separate highly diluted  $\text{CO}_2$  from flue-gases (Budzianowski, 2016). One of the most recognized and well-established technology for post-combustion carbon capture is chemical absorption with aqueous alkanolamines solutions. Primary, secondary, tertiary and hindered alkanolamines are used in Acid Gas-Water-Amine (AGWA) systems, commonly: monoethanolamine (MEA), diethanolamine (DEA), methyldiethanolamine (MDEA), and 2-amino-2-methylpropanol (AMP), respectively. (de Medeiros, Barbosa and Araújo, 2013)

Modeling of AGWA systems is complex and computationally demanding. The models must deal with nonequilibrium ionic chemical reactions, and heat/mass transfers across vapor-liquid interfaces. Frequently, the necessary physical properties are unavailable and the ions generated by weak dissociations are unknown. High pressures, concentrations, and loadings, make the use of idealities, such as ideal law for gas, inaccurate (de Medeiros, Barbosa and Araújo, 2013). These issues must be circumvented using the methodology of de Medeiros, Barbosa and Araújo (2013), which incorporates molecular species into a chemical equilibrium theory framework using cubic equations of state.

One of the first processes for removing acid gases ( $\text{CO}_2$  and  $\text{H}_2\text{S}$ ) from NG with MEA was proposed and patented by Bottoms (1930). This process, shown in Fig. 2.10, is also useful to remove  $\text{CO}_2$  from flue-gases.

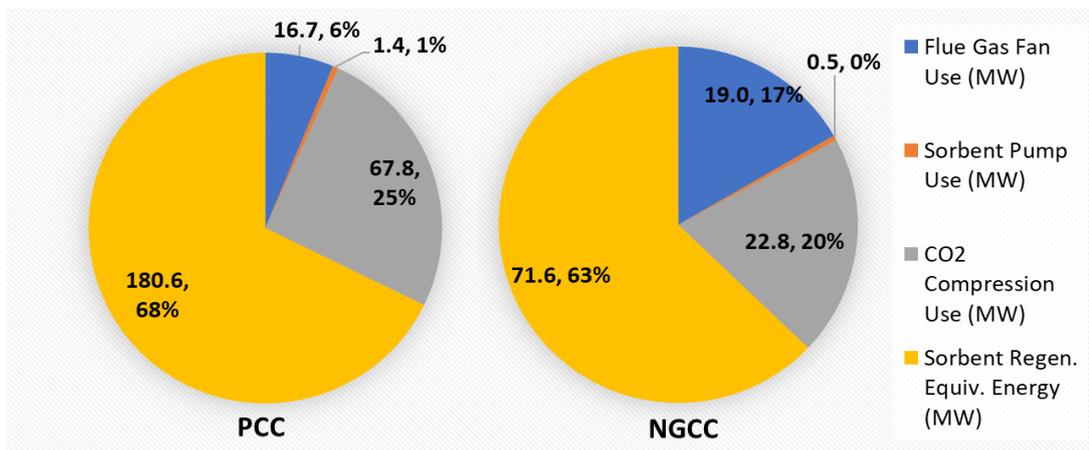


**Figure 2.10. Gas-Sweetening Process using Monoethanolamine. Adapted from Bottoms (1930)**

The main drawback of the chemical absorption processes using solvents in a closed loop, as in Fig. 2.10, is the energy penalty for solvent regeneration. This penalty is pointed out as the major obstacle to the widespread deployment of full-scale applications of carbon capture by reactive absorption, especially in coal and NG power plants (Boot-Handford et al., 2014). The energy penalty is defined as the energy spent to capture a certain amount of CO<sub>2</sub> from a process stream, usually expressed as GJ/t of CO<sub>2</sub> removed. According to Knudsen et al. (2009), the use of steam from the Rankine cycle to regenerate the solvent at the stripper column consumes between 3.6 and 3.8 GJ/t of CO<sub>2</sub> captured. Carbon Capture and Storage (CCS) can increase approximately 70% the life cycle cost (LCC) of fossil-fueled electricity and cement and, similarly 40% of steel LCC (Zhang et al., 2013). The Bottom's process can no longer be considered the benchmark for CO<sub>2</sub> capture but most of the chemical absorption technologies are based on this concept. Reduced CO<sub>2</sub> loading (kgCO<sub>2</sub>/kg of solvent), thermal and oxidative degradation, corrosivity and evaporation losses are often reported as other weaknesses of traditional solvents for CO<sub>2</sub> capture.

In coal-fired power plants, there is a linear relationship between regeneration energy and the overall efficiency of the electricity generation. It is estimated that each GJ/t of CO<sub>2</sub> translates

into a 2% decrease in the heat ratio or global efficiency of the plant. Considering thermal plants with efficiency between 40% and 60%, an energy penalty of 5% causes an increase of approximately 10% in fuel consumption. The resulting impact on the electricity price is considerable (Goto, Yogo and Higashii, 2013). More precise quantification of the effect of retrofitting a 500MW Pulverized Coal Combustion plant was performed in the present research using the software IECM v 11 (Rubin, 2018) shows that adding CCS with MEA increased the LCOE by 115% (from 61 \$/MWh to 131 \$/MWh). The regeneration consumption is 180 MW. Considering a NG combined cycle (NGCC) with the same nominal capacity, the LCOE increased 52% (from 66 \$/MWh to 100 \$/MWh), with a consumption of 72 MW from the solvent regeneration system. In both cases, most of the CCS energy penalty came from the reboiler consumption of steam, in the solvent regeneration column, as shown in Fig. 2.11.



**Figure 2.11. CCS energy penalty of Pulverized Coal Combustion and Natural Gas Combined Cycle. Data obtained by the author from simulations using the software IECM v 11 (Rubin, 2018).**

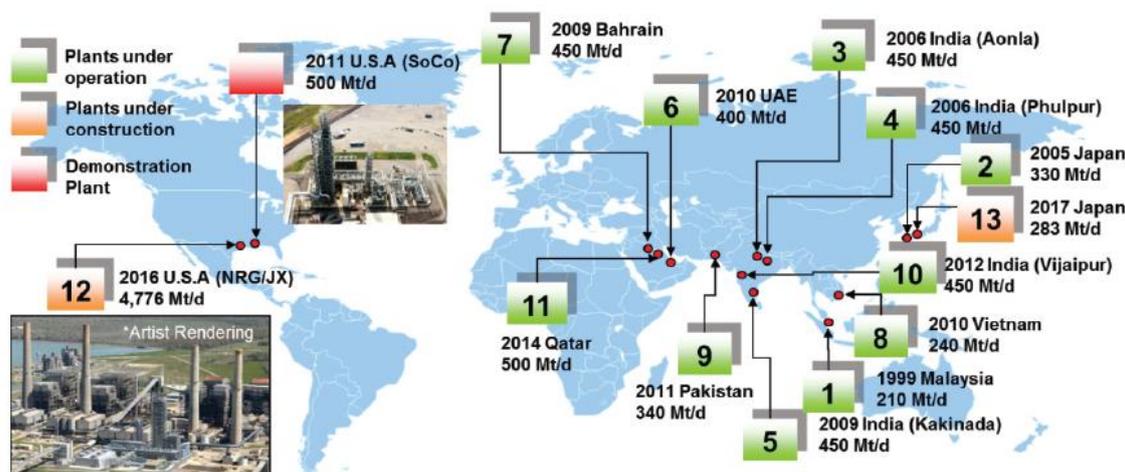
These preliminary results motivated the R3 research line, and other developments of solvents and process designs to lower regeneration energy requirements, e.g.: advanced solvents (Goto, Yogo and Higashii, 2013); inter-stage cooling and heating in the absorber and regenerating columns, respectively (Frailie et al., 2013); energy integrations; exhaust gas recycling (Li et al., 2011), among other improvements (Park et al., 2016; Rezazadeh et al., 2017; Sachde; Rochelle, 2014; Zhang; Rochelle, 2014). Industries from the chemical and energy sectors developed solvents and processes that reduced the energy penalty for regeneration by up to 43%, compared to the classic benchmark (MEA 30% - 3.7 GJ/tCO<sub>2</sub>), as shown in Table 2.2. Mitsubishi Heavy Industries owns one of the most modern and mature processes, the KM CDR (Kansai Mitsubishi Carbon Dioxide Recovery Process), in pilot and industrial scale in several applications, as shown in Figure 2.12. The Petra Nova joint venture adopted de KM-CDR and

is currently the world's largest CCS plant. This plant has capacity to capture 4776 t/d of CO<sub>2</sub>, with EOR injection 130 km apart. This capture flow ratio is equivalent to the flue-gas production of a 240 MW power plant (Miyamoto et al., 2017).

**Table 2.2. Commercially available chemical absorption CCS technologies.**

Process	Company	Solvent	Energy Penalty	Reference
Cansolv	Shell Royal Dutch	DC-103	2.33 GJ/t CO <sub>2</sub>	Singh and Stéphenne (2014)
KM CDR	Mitsubishi Heavy Industry	KS-1	2.29 GJ/t CO <sub>2</sub>	Kadono et al. (2013)
Econamine FG+	Fluor Corporation	Amine-Based	3.00* GJ/t CO <sub>2</sub>	Mathias, Reddy and O'Connell (2009)

\*3.00 GJ/t of steam for reboiler + 2.75 GJ/t for NH<sub>3</sub> refrigeration (gas compression cycle).



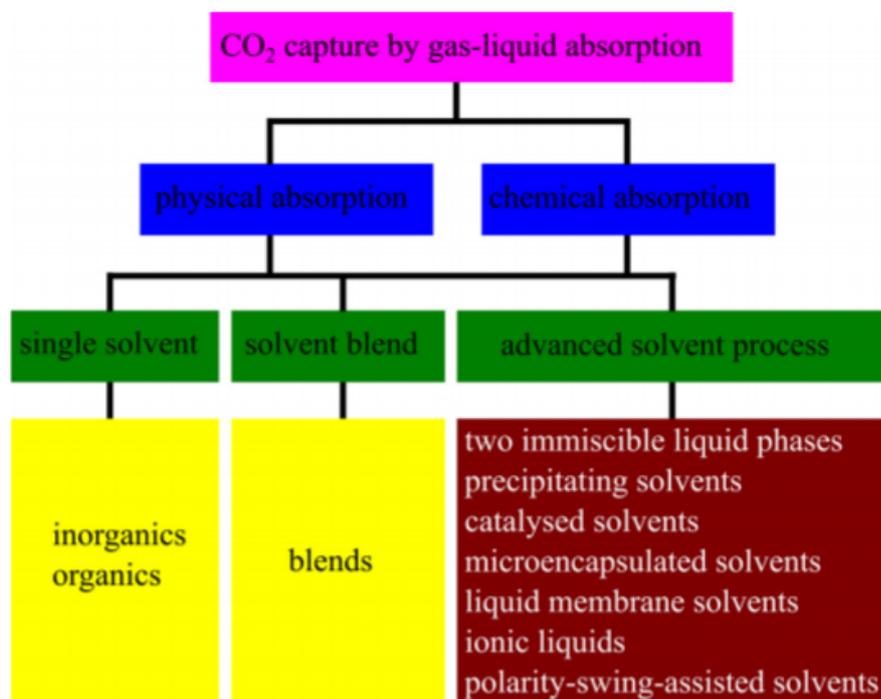
**Figure 2.12. KM-CDR applications worldwide (Miyamoto et al., 2017).**

Current conventional chemical absorption processes are highly optimized in terms of layout and energy integration. Process design improvement opportunities are scarce. Developments on new solvents, like PCAS, aims at reducing the inherent energy penalty of chemical absorption processes. Such solvents would contribute to the faster deployment of CCUS in industrial scale worldwide, meeting sustainable development goals.

### 2.3.3. Phase-Changing Absorption Solvents for CO<sub>2</sub> Capture

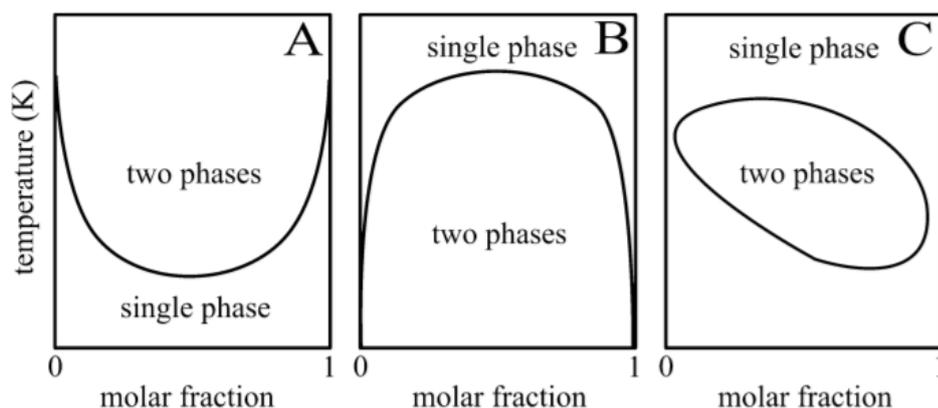
PCAS are considered advanced solvents for gas-liquid CO<sub>2</sub> absorption. An advanced solvent has a physical or chemical characteristic that enhances the energy performance of a CO<sub>2</sub> capture process. Often, the energy spent with solvent regeneration, using stripping columns, is reduced. Post-combustion processes using advanced solvents could significantly reduce the decarbonization energy penalty, benefiting the energy and other carbon-intensive sectors

(Budzianowski, 2016). Fig 2.13 shows a gas-liquid absorption selection tree, where PCAS (red box, as two immiscible liquid phases) is classified as an advanced solvent. The fundamental property that differentiates a PCAS from other solvent is the formation of immiscible liquid phases triggered by temperature and/or CO<sub>2</sub> loading of the solvent. The energy saving results from the reduced mass of solvent sent to the stripper, considering that only one of the immiscible phases is rich in CO<sub>2</sub>. The reboiler duty and possibly the footprint of the stripper column are reduced, resulting in CAPEX and OPEX savings (Coulier et al., 2017; Liebenthal et al., 2013).

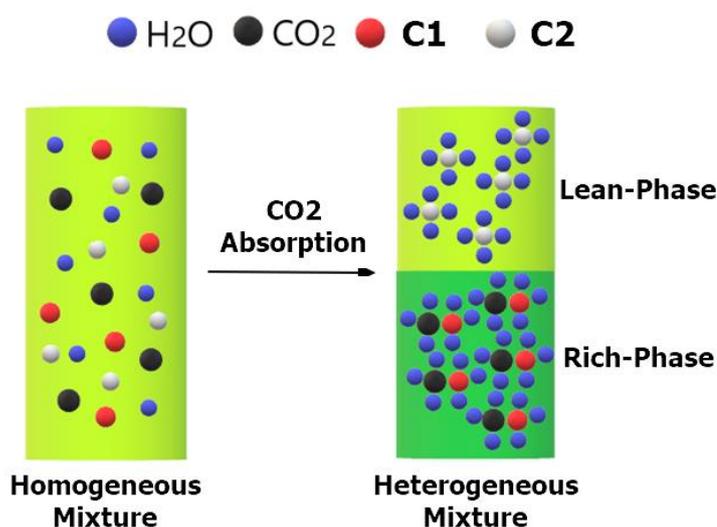


**Fig 2.13. Gas-liquid absorption solvents for CO<sub>2</sub> capture. (Budzianowski, 2016).**

Three possible scenarios of immiscible liquid phase formation are shown in Fig. 2.14, as a function of temperature. Similar behavior may occur as a function of CO<sub>2</sub> loading instead of temperature. Fig 2.15 illustrates a generic process of phase-split promoted by CO<sub>2</sub> absorption.



**Fig 2.14. Possibilities of immiscible liquid phases formation. (A) lower critical temperature, (B) upper critical temperature, and (C) upper and lower critical temperatures (Budzianowski, 2016).**



**Fig 2.15. Liquid-liquid phase-split promoted by CO<sub>2</sub> capture.**

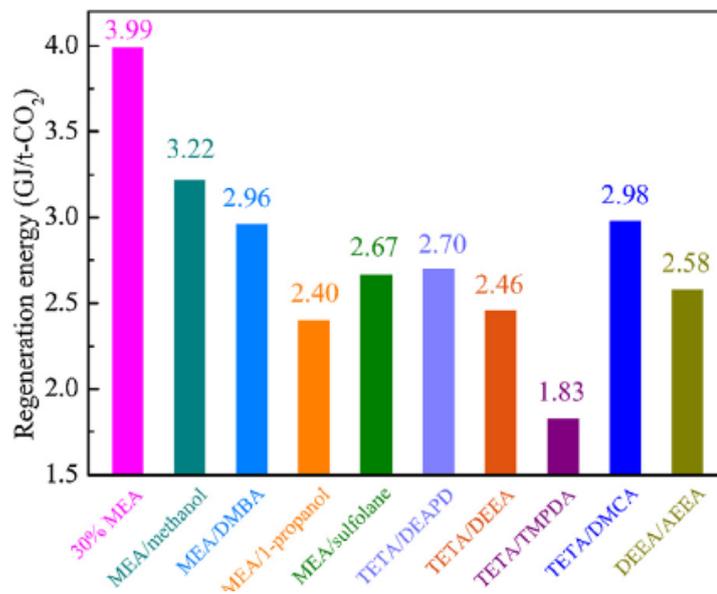
A suitable PCAS should have the following behavior (Budzianowski, 2016):

- Operate homogeneously under scrubbing temperatures.
- Operate as a biphasic mixture (two immiscible liquid phases) under stripping temperatures or after saturation with CO<sub>2</sub>.
- Form a CO<sub>2</sub>-lean and a CO<sub>2</sub>-rich phase, easily separable by gravity.
- Preferably, the volume of the CO<sub>2</sub>-rich phase should be smaller than the volume of the lean phase.

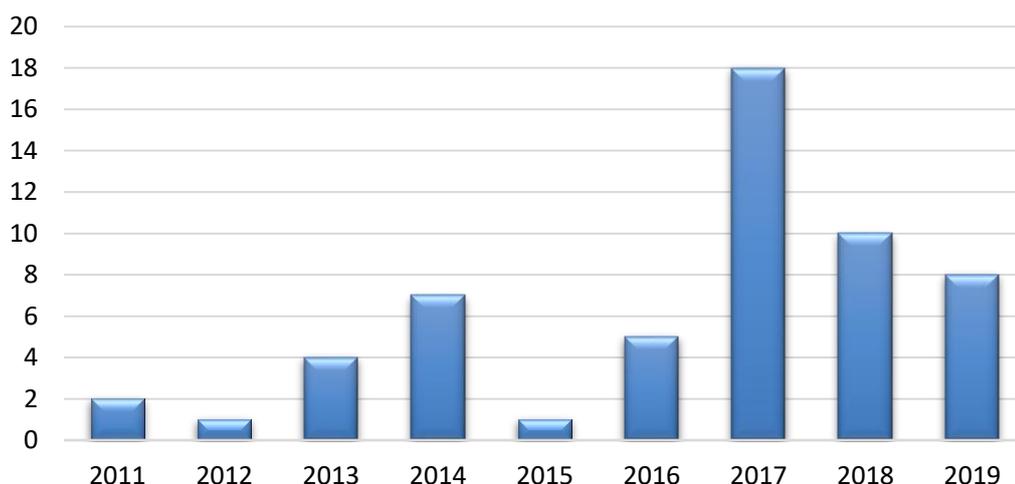
Wang et al. (2019) estimated the regeneration energy of several PCAS. The authors considered a 300MW Pulverized Coal Combustion (PCC) power plant using CCS with MEA 30% (w/w)

as the baseline. The results are depicted in Fig. 2.16. The PCAS based on triethylenetetramine (TETA) and Tetramethyl-1,3-propanediamine (TMPDA) presented regeneration energy 54% lower than the baseline. MEA/1-propanol ranked second, with a 40% reduction. These results denote the potential benefits of the full-scale application of PCAS on the energy sector. However, the results of Wang et al. (2019) are mostly based on simulation, using Aspen Plus. The lack of information concerning the simulation setup, especially the thermodynamic model calibration, poses a high degree of uncertainty in the accuracy of the energy penalty reported.

A systematic review was performed to determine the state-of-the-art on PCAS. The following group of keywords was chosen as input on Google Scholar: (*"liquid-liquid phase separation" OR "phase transition" OR "de-mixing" OR "phase change solvent" OR "biphasic solvent" OR "thermomorphic"*) AND (*"CO<sub>2</sub> capture" OR "CO<sub>2</sub> absorption" OR "carbon dioxide capture" OR "carbon dioxide absorption"*). Results were limited to the years 2010 to 2019 and totaled 5.570 articles, and 56 articles were considered more relevant to in depth analysis. Most of the selected articles were published on indexed scientific journals, with high impact factors. An overview of the selected articles reveals that Chinese institutions are responsible for 43% of published papers, followed from France, with 11% of the production, as shown in Fig. 2.17.

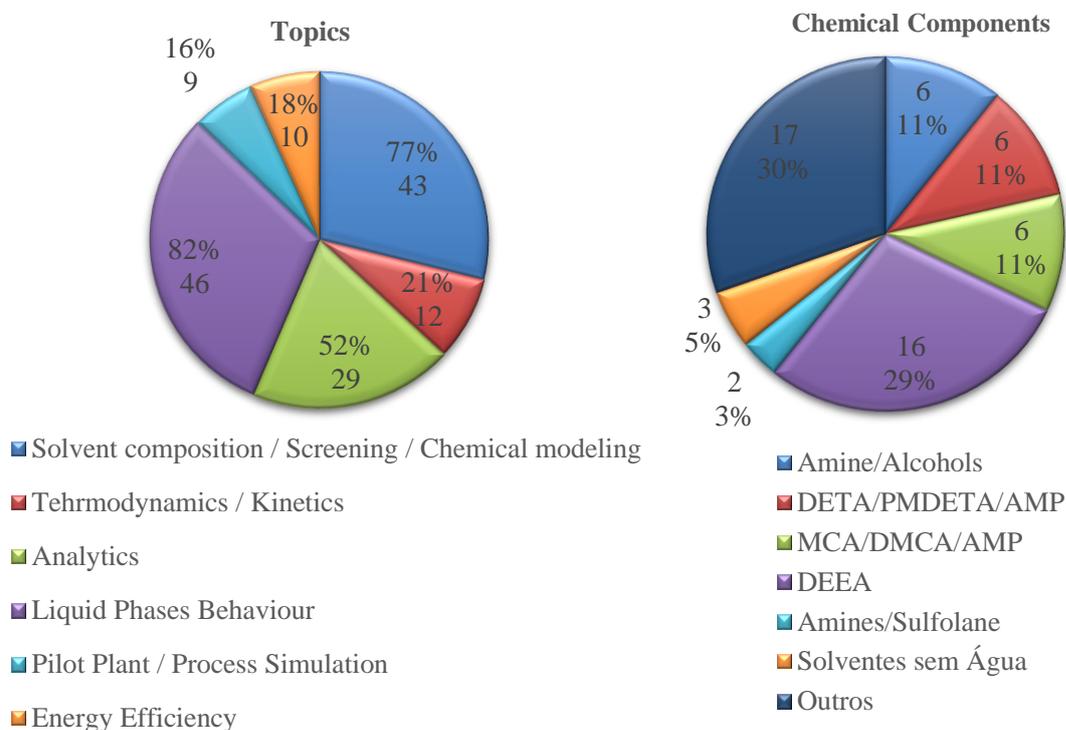


**Fig 2.16. Comparison of the regeneration heat of MEA-based and other phase-changing absorption solvents (Wang et al., 2019).**



**Figure 2.17. Publications on phase-changing absorption solvents from 2010 to 2019.**

The review shows a peak of publications on PCAS in 2017 (18 articles) followed by 10 publications in 2018 and 8 until April 2019. Those numbers reflect the relevance and interest on the research line R3 approached in this thesis. Fig. 2.18 presents the most studied topics and chemical components found in the selected articles on PCAS. Most of the selected articles deals with liquid phases behavior and chemical composition of PCAS, and are mostly on laboratory evaluations, rarely targeting pilot plant experiments and process modeling and simulation, denoting the early stage of development on PCAS.



**Figure 2.18. Main topics and chemical components related to phase-changing absorption solvents for CO<sub>2</sub> capture.**

## 2.4. References of Chapter 2

- ANP, 2020. Boletim da Produção de Petróleo e Gás Natural - Janeiro 2020 [WWW Document]. URL <http://www.anp.gov.br/arquivos/publicacoes/boletins-anp/producao/2020-01-boletim.pdf> (accessed 3.4.20).
- Araújo, O. de Q.F., de Medeiros, J.L., 2017. Carbon capture and storage technologies: present scenario and drivers of innovation. *Curr. Opin. Chem. Eng.* 17, 22–34. <https://doi.org/10.1016/j.coche.2017.05.004>
- Araújo, O. de Q.F., Reis, A. de C., de Medeiros, J.L., Nascimento, J.F. do, Grava, W.M., Musse, A.P.S., 2017. Comparative analysis of separation technologies for processing carbon dioxide rich natural gas in ultra-deepwater oil fields. *J. Clean. Prod.* 155, 12–22. <https://doi.org/10.1016/j.jclepro.2016.06.073>
- Boot-Handford, M.E., Abanades, J.C., Anthony, E.J., Blunt, M.J., Brandani, S., Mac Dowell, N., Fernández, J.R., Ferrari, M.-C., Gross, R., Hallett, J.P., Haszeldine, R.S., Heptonstall, P., Lyngfelt, A., Makuch, Z., Mangano, E., Porter, R.T.J., Pourkashanian, M., Rochelle, G.T., Shah, N., Yao, J.G., Fennell, P.S., 2014. Carbon capture and storage update. *Energy Environ. Sci.* 7, 130–189. <https://doi.org/10.1039/C3EE42350F>
- Bottoms, R.R., 1930. Process for separating acidic gases. US17883901A.
- Brandt, A.R., Sun, Y., Bharadwaj, S., Livingston, D., Tan, E., Gordon, D., 2015. Energy Return on Investment (EROI) for Forty Global Oilfields Using a Detailed Engineering-Based Model of Oil Production. *PLoS One* 10, e0144141.
- Budzianowski, W.M., 2016. Explorative analysis of advanced solvent processes for energy efficient carbon dioxide capture by gas–liquid absorption. *Int. J. Greenh. Gas Control* 49, 108–120. <https://doi.org/10.1016/J.IJGGC.2016.02.028>
- Carpenter, A.M., 2012. Low water FGD technologies. IEA Clean Coal Center.
- Copenhagen Economics, 2017. The future of fossil fuels: How to steer fossil fuels use in a transition to a low-carbon energy system. An analysis of fossil fuels trajectories in low-carbon scenarios.
- Coulier, Y., Lowe, A.R., Coxam, J.-Y., Ballerat-Busserolles, K., 2017. Thermodynamic Modeling and Experimental Study of CO<sub>2</sub> Dissolution in New Absorbents for Post-Combustion CO<sub>2</sub> Capture Processes. *ACS Sustain. Chem. Eng.* acssuschemeng.7b03280. <https://doi.org/10.1021/acssuschemeng.7b03280>
- de Medeiros, J.L., Barbosa, L.C., Araújo, O.D.Q.F., 2013a. Equilibrium approach for CO<sub>2</sub> and H<sub>2</sub>S absorption with aqueous solutions of alkanolamines: Theory and parameter estimation. *Ind. Eng. Chem. Res.* 52, 9203–9226. <https://doi.org/10.1021/ie302558b>
- de Medeiros, J.L., Nakao, A., Grava, W.M., Nascimento, J.F., Araújo, O. de Q.F., 2013b. Simulation of an offshore natural gas purification process for CO<sub>2</sub> removal with gas-liquid contactors employing aqueous solutions of ethanolamines. *Ind. Eng. Chem. Res.* 52, 7074–7089. <https://doi.org/10.1021/ie302507n>

- Dehghani, A., Bridjanian, H., 2010. Flue Gas Desulfurization Methods To Conserve the Environment. *Pet. Coal*.
- European Commission, 2006. Integrated Pollution Prevention and Control Reference Document on Best Available Techniques for Large Combustion Plants, Integrated Pollution Prevention and Control.
- Frailie, P.T., Madan, T., Sherman, B.J., Rochelle, G.T., 2013. Energy performance of advanced stripper configurations. *Energy Procedia* 37, 1696–1705. <https://doi.org/10.1016/j.egypro.2013.06.045>
- Friedlingstein, P., Al, E., 2019. Global Carbon Budget 2019. *Earth Syst. Sci. Data* 11, 1783–1838. <https://doi.org/https://doi.org/10.5194/essd-11-1783-2019>
- Frimpong, R.A., Irvin, B.D., Nikolic, H., Liu, K., Figueroa, J., 2019. Integrated hybrid process for solvent-based CO<sub>2</sub> capture using a pre-concentrating membrane: A pilot scale study. *Int. J. Greenh. Gas Control* 82, 204–209. <https://doi.org/10.1016/J.IJGGC.2019.01.016>
- Funch-Jensen, A., Rubner-Petersen, M., 2007. Semi-dry scrubber technologies: Spray Drying Absorption, in: *ESKOM Scrubber Seminar*. World Pollution Control Association, Copenhagen, DN, p. 50.
- Global CCS Institute, 2019. *The Global Status of CCS: 2019*, Global CCS Institute. Melbourne, Australia. [https://doi.org/10.1007/springerreference\\_15392](https://doi.org/10.1007/springerreference_15392)
- Goto, K., Yogo, K., Higashii, T., 2013. A review of efficiency penalty in a coal-fired power plant with post-combustion CO<sub>2</sub> capture. *Appl. Energy* 111, 710–720. <https://doi.org/10.1016/J.APENERGY.2013.05.020>
- Hill, F.F., Zank, J., 2000. Flue gas desulphurization by spray dry absorption. *Chem. Eng. Process. Process Intensif.* [https://doi.org/10.1016/S0255-2701\(99\)00077-X](https://doi.org/10.1016/S0255-2701(99)00077-X)
- IEA, 2020. *Data & Statistics - IEA [WWW Document]*. Int. Energy Agency. URL [https://www.iea.org/data-and-statistics?country=EU28&fuel=Energy supply&indicator=Total primary energy supply \(TPES\) by source](https://www.iea.org/data-and-statistics?country=EU28&fuel=Energy supply&indicator=Total primary energy supply (TPES) by source) (accessed 1.31.20).
- IEA, 2019a. *CO<sub>2</sub> emissions from fuel combustion [WWW Document]*. URL [https://webstore.iea.org/download/direct/2505?fileName=CO<sub>2</sub>\\_Emissions\\_from\\_Fuel\\_Combustion\\_2019\\_Overview.pdf](https://webstore.iea.org/download/direct/2505?fileName=CO2_Emissions_from_Fuel_Combustion_2019_Overview.pdf)
- IEA, 2019b. *World Energy Outlook - Executive Summary [WWW Document]*. Int. Energy Agency. URL [www.iea.org](http://www.iea.org)
- IEA, 2018. *Perspectives for the Energy Transition: The Role of Energy Efficiency [WWW Document]*. URL <http://www.iea.org/publications/freepublications/publication/Perspectives for the Energy Transition - The Role of Energy Efficiency.pdf> (accessed 7.2.18).
- IOGP, 2019. *IOGP Environmental performance indicators - 2018 data - Atmospheric emissions [WWW Document]*. Int. Assoc. Oil Gas Prod. URL <https://data.iogp.org/Environment/Emissions> (accessed 2.4.20).

- IPCC, 2014. Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, Intergovernmental Panel on Climate Change. IPCC, Geneva, Switzerland.
- IPIECA, 2017. Mapping the Oil and Gas Industry To the Sustainable Development Goals: an Atlas [WWW Document]. URL [http://www.ipieca.org/media/3093/mapping\\_og\\_to\\_sdg\\_atlas\\_lr\\_2017.pdf](http://www.ipieca.org/media/3093/mapping_og_to_sdg_atlas_lr_2017.pdf)
- Kadono, K., Suzuki, A., Iijima, M., Ohishi, T., Tanaka, H., Hirata, T., Kondo, M., 2013. New Energy Efficient Processes and Newly Developed Absorbents for Flue Gas CO<sub>2</sub> Capture. *Energy Procedia* 37, 1785–1792. <https://doi.org/10.1016/J.EGYPRO.2013.06.055>
- Knudsen, J.N., Jensen, J.N., Vilhelmsen, P.J., Biede, O., 2009. Experience with CO<sub>2</sub> capture from coal flue gas in pilot-scale: Testing of different amine solvents. *Energy Procedia* 1, 783–790. <https://doi.org/10.1016/j.egypro.2009.01.104>
- Li, H., Haugen, G., Ditaranto, M., Berstad, D., Jordal, K., 2011. Impacts of exhaust gas recirculation (EGR) on the natural gas combined cycle integrated with chemical absorption CO<sub>2</sub> capture technology. *Energy Procedia* 4, 1411–1418. <https://doi.org/10.1016/j.egypro.2011.02.006>
- Liebenthal, U., Di D. Pinto, D., Monteiro, J.G.M.S., Svendsen, H.F., Kather, A., 2013. Overall process analysis and optimisation for CO<sub>2</sub> Capture from coal fired power plants based on phase change solvents forming two liquid phases. *Energy Procedia* 37, 1844–1854. <https://doi.org/10.1016/j.egypro.2013.06.064>
- Liu, B., Tang, C., Li, X., Wang, B., Zhou, R., 2020. High-performance SAPO-34 membranes for CO<sub>2</sub> separations from simulated flue gas. *Microporous Mesoporous Mater.* 292, 109712. <https://doi.org/10.1016/J.MICROMESO.2019.109712>
- Ma, X., Kaneko, T., Tashimo, T., Yoshida, T., Kato, K., 2000. Use of limestone for SO<sub>2</sub> removal from flue gas in the semidry FGD process with a powder-particle spouted bed. *Chem. Eng. Sci.* [https://doi.org/10.1016/S0009-2509\(00\)00090-7](https://doi.org/10.1016/S0009-2509(00)00090-7)
- Masnadi, M.S., El-Houjeiri, H.M., Schunack, D., Li, Y., Englander, J.G., Badahdah, A., Monfort, J.-C., Anderson, J.E., Wallington, T.J., Bergerson, J.A., Gordon, D., Koomey, J., Przesmitzki, S., Azevedo, I.L., Bi, X.T., Duffy, J.E., Heath, G.A., Keoleian, G.A., McGlade, C., Meehan, D.N., Yeh, S., You, F., Wang, M., Brandt, A.R., 2018. Global carbon intensity of crude oil production. *Science* (80-. ). 361, 851–853. <https://doi.org/10.1126/science.aar6859>
- Mathias, P.M., Reddy, S., O’Connell, J.P., 2009. Quantitative evaluation of the aqueous-ammonia process for CO<sub>2</sub> capture using fundamental data and thermodynamic analysis. *Energy Procedia* 1, 1227–1234. <https://doi.org/10.1016/J.EGYPRO.2009.01.161>
- Miyamoto, O., Maas, C., Tsujiuchi, T., Inui, M., Hirata, T., Tanaka, H., Yonekawa, T., Kamijo, T., 2017. KM CDR Process™ Project Update and the New Novel Solvent Development. *Energy Procedia* 114, 5616–5623. <https://doi.org/10.1016/J.EGYPRO.2017.03.1700>
- MODEC, 2020. MODEC Awarded Contract by Equinor to Supply FPSO for offshore field in Brazil [WWW Document]. Modec Inc. URL

- [https://www.modec.com/news/2020/20200130\\_pr\\_bacalhau\\_en.html](https://www.modec.com/news/2020/20200130_pr_bacalhau_en.html) (accessed 3.4.20).
- MODEC, 2019. M350 NEW BUILT FPSO HULL [WWW Document]. URL [https://www.gastechevent.com/media/41272/m350\\_e\\_brochure.pdf](https://www.gastechevent.com/media/41272/m350_e_brochure.pdf) (accessed 3.6.20).
- NOAA, 2020. ESRL Global Monitoring Division - Global Greenhouse Gas Reference Network [WWW Document]. US Dep. Commer. Earth Syst. Res. Lab. URL <https://www.esrl.noaa.gov/gmd/ccgg/trends/> (accessed 2.2.20).
- NS Energy, 2019. MODEC unveils two next-generation new built FPSO hulls [WWW Document]. NS Energy Bus. URL <https://www.nsenergybusiness.com/news/modec-built-fps-hulls/> (accessed 3.6.20).
- Park, T., Bae, J., Lee, C.J., Lee, J.M., 2016. A Sequential Method for Determining Optimal Stripper Pressure and Terminal Pressure in CO<sub>2</sub> Capture and Liquefaction Process Using MEA. *IFAC-PapersOnLine* 49, 657–662. <https://doi.org/10.1016/j.ifacol.2016.07.250>
- Petrobras, 2020. Fechamos o ano de 2019 com ótimo desempenho operacional [WWW Document]. Blog Fatos e Dados. URL <https://petrobras.com.br/fatos-e-dados/fechamos-o-ano-de-2019-com-otimo-desempenho-operacional.htm> (accessed 3.4.20).
- Rezazadeh, F., Gale, W.F., Rochelle, G.T., Sachde, D., 2017. Effectiveness of absorber intercooling for CO<sub>2</sub> absorption from natural gas fired flue gases using monoethanolamine solvent. *Int. J. Greenh. Gas Control* 58, 246–255. <https://doi.org/10.1016/j.ijggc.2017.01.016>
- Rezazadeh, F., Gale, W.F., Sachde, D., Rochelle, G.T., 2014. Absorber intercooling configurations using aqueous piperazine for capture from sources with 4 to 27% CO<sub>2</sub>. *Energy Procedia* 63, 1637–1656. <https://doi.org/10.1016/j.egypro.2014.11.174>
- Rubin, E.S., 2018. Integrated Environmental Control Model [WWW Document]. Carnegie Mellon. URL <https://www.cmu.edu/epp/iecm/index.html> (accessed 7.26.18).
- Sage, P.W., Ford, N.W.J., 1996. Review of sorbent injection processes for low-cost sulphur dioxide control. *Proc. Inst. Mech. Eng. Part A J. Power Energy*. [https://doi.org/10.1243/PIME\\_PROC\\_1996\\_210\\_031\\_02](https://doi.org/10.1243/PIME_PROC_1996_210_031_02)
- SBM Offshore, 2019. SBM Offshore-Fast4Ward ® [WWW Document]. URL <https://www.sbmoffshore.com/wp-content/uploads/2019/09/SEPT-2019-FAST4WARD-BROCHURE.pdf> (accessed 3.4.20).
- Singh, A., Stéphenne, K., 2014. Shell Cansolv CO<sub>2</sub> capture technology: Achievement from First Commercial Plant. *Energy Procedia* 63, 1678–1685. <https://doi.org/10.1016/J.EGYPRO.2014.11.177>
- Srivastava, R.K., Jozewicz, W., 2001. Flue gas desulfurization: The state of the art. *J. Air Waste Manag. Assoc.* <https://doi.org/10.1080/10473289.2001.10464387>
- Tong, J., Zhang, L., Han, M., Huang, K., 2015. Electrochemical separation of CO<sub>2</sub> from a simulated flue gas with high-temperature ceramic-carbonate membrane: New observations. *J. Memb. Sci.* <https://doi.org/10.1016/j.memsci.2014.12.017>

- UNFCCC, 2016. Report of the Conference of the Parties, Addendum Part two: Action taken by the Conference at its twenty-first session [WWW Document]. URL <http://unfccc.int/resource/docs/2015/cop21/eng/10a01.pdf> (accessed 7.14.16).
- United Nations, 2015. Transforming our World: The 2030 Agenda for Sustainable Development [WWW Document]. URL [https://sustainabledevelopment.un.org/content/documents/21252030\\_Agenda\\_for\\_Sustainable\\_Development\\_web.pdf](https://sustainabledevelopment.un.org/content/documents/21252030_Agenda_for_Sustainable_Development_web.pdf) (accessed 3.5.20).
- Wang, R., Liu, S., Wang, L., Li, Q., Zhang, S., Chen, B., Jiang, L., Zhang, Y., 2019. Superior energy-saving splitter in monoethanolamine-based biphasic solvents for CO<sub>2</sub> capture from coal-fired flue gas. *Appl. Energy* 242, 302–310. <https://doi.org/10.1016/J.APENERGY.2019.03.138>
- WCA, 2017. Coal & Electricity [WWW Document]. World Coal Assoc. URL <https://www.worldcoal.org/coal/uses-coal/coal-electricity> (accessed 3.14.17).
- Zhang, J., Qiao, Y., Wang, W., Misch, R., Hussain, K., Agar, D.W., 2013. Development of an energy-efficient CO<sub>2</sub> capture process using thermomorphic biphasic solvents. *Energy Procedia* 37, 1254–1261. <https://doi.org/10.1016/j.egypro.2013.05.224>
- Zhang, Y., Rochelle, G.T., 2014. Absorber Performance with High CO<sub>2</sub>. *Energy Procedia* 63, 1329–1338. <https://doi.org/10.1016/j.egypro.2014.11.142>



### **3. DEEP SEAWATER INTAKE FOR PRIMARY COOLING IN TROPICAL OFFSHORE PROCESSING OF NATURAL GAS WITH HIGH CARBON DIOXIDE CONTENT: ENERGY, EMISSIONS AND ECONOMIC ASSESSMENTS**

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CRUZ, M. DE A.; ARAÚJO, O. DE Q. F.; DE MEDEIROS, J. L. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. **Journal of Natural Gas Science and Engineering**, v. 56, n. June, p. 193–211, 2018.

#### **Abstract**

In deepwaters offshore oil-gas rigs, centrifugal compressor trains are major power consumers, requiring intercoolers conventionally designed assuming surface seawater for primary cooling, limiting compressor inlet gas temperatures to 40°C at tropical sites. On the other hand, at tropical deepwaters the available deep seawater at 4°C can be exploited to reduce compression power – nearly proportional to inlet gas absolute temperature – entailing energy, economic and environmental benefits. This work considers a new primary cooling for deepwaters offshore platforms based on deep seawater (DSW) intake at 4°C from depths around 900 m, reducing the outlet temperature of intercoolers to 12°C. DSW intake alternative is assessed in terms of power consumption, CO<sub>2</sub> emissions and economy employing detailed equipment sizing and cost estimation. Depending on gas flow rate, it is shown that DSW intake lowers compressors power up to 9.2%, besides several indirect benefits: elimination of one CO<sub>2</sub> compressor; 30% less heat transfer areas; 4.5% less fuel gas consumption; 4% less gas turbines power; 9.5% (15 MMUS\$) less investment; 14.4% (226 t) less topside weight, while making refrigeration unnecessary for dew point adjustment. DSW intake also entails 5% more efficient energy usage and 9327 tCO<sub>2</sub>/y less emissions, boosting economic performance under carbon taxation.

**Keywords:** Deep seawater intake; Deepwater oil production; Offshore natural gas processing; FPSO; Gas compression; Energy usage efficiency.

Supplementary Materials for this chapter are found in the Appendix H, Section A.

## Abbreviations

<i>CAPEX</i>	Capital Expenditures
<i>CW</i>	Cooling-Water
<i>DSW</i>	Deep Seawater
<i>EIA</i>	Environmental Impacts Assessment
<i>EOR</i>	Enhanced Oil Recovery
<i>FLNG</i>	Floating Liquefied Natural Gas Plant
<i>FPSO</i>	Floating Production, Storage and Offloading
<i>GHG</i>	Greenhouse Gas
<i>GOR</i>	Gas-to-Oil Ratio
<i>GT</i>	Gas Turbine
<i>HCDP</i>	Hydrocarbon Dew-Point
<i>HCDPA</i>	Hydrocarbon Dew-Point Adjustment
<i>HDPE</i>	High Density Polyethylene
<i>HHV</i>	Higher Heating Value
<i>LHV</i>	Lower Heating Value
<i>MP</i>	Membrane Permeation
<i>NG</i>	Natural Gas
<i>NPSH</i>	Net Positive Suction Head
<i>OPEX</i>	Operational Expenditures
<i>OTEC</i>	Offshore Thermal Energy Conversion
<i>PFD</i>	Process Flow Diagram
<i>PR-EOS</i>	Peng-Robinson Equation of State
<i>RH</i>	Relative Humidity
<i>SW</i>	Seawater
<i>TSA</i>	Temperature Swing Adsorption
<i>US\$</i>	US Dollar
<i>VLCC</i>	Very Large Crude Carrier
<i>WDPA</i>	Water Dew Point Adjustment
<i>WHRU</i>	Waste Heat Recovery Unit
<i>WOR</i>	Water-to-Oil Ratio.

## Nomenclature

$A_0$	Effective outside heat transfer surface ( $m^2$ )
$bb/d$	Barrels per day
$CO_{2Eq}$	Carbon dioxide equivalent
$F$	<i>LMTD</i> correction factor
$MW$	Molecular Mass
$MMSm^3/d$	Millions of Standard $m^3$ (293 K, 101.33 kPa) per day
$n$	Polytropic exponent
$N$	Number of compression stages
$P$	Absolute pressure (kPa, bar)
$q$	Mass flow rate (kg/h)
$Q$	Heat duty (kW)
$R$	Ideal gas constant (8.314 kJ/kmol.K)
$r_p$	Compression ratio
$T$	Absolute temperature (K)
$U$	Overall heat transfer coefficient (kW/ $m^2$ .K)

$W$	Brake horsepower (kW)
$Z$	Compressibility factor

## Greek Symbols

$\eta_p$	Polytropic efficiency
$\Delta P$	Head loss (kPa)
$\Delta T_M$	Corrected mean temperature difference (K)

### 3.1. Introduction

Since 2000, offshore fields respond for  $\approx 30\%$  of the world production of oil and natural gas (Rui et al., 2017). Due to still modest competitiveness of renewable energy sources, fossil sources will continue to play significant role in global energy matrix in the short to mid-term, especially natural gas (NG). However, oil price decline and new climate change mitigation policies – e.g. carbon taxation – have been a challenge to this industry. CO<sub>2</sub> emission taxation is already a reality in many countries, with Sweden imposing the highest tax of 140 US\$/t (IEA, 2016). According to the International Association of Oil and Gas Producers (IOGP, 2016) main oil and gas companies emitted 280 Mt CO<sub>2Eq</sub> of greenhouse gases (GHG) in 2015, of which 68% is related to fuel combustion for in-place energy production. Thus, energy usage efficiency and CO<sub>2</sub> emissions are becoming not solely an environmental, but also, an economic issue for oil and gas producers. As energy usage efficiency and CO<sub>2</sub> emissions are inversely interrelated concepts, using processing strategies with higher energy usage efficiency implies lowering CO<sub>2</sub> emissions, alleviating the environmental burden of oil and gas industries. In other words, there is no option to the carbon fossil industry but questing for better energy usage efficiency.

#### 3.1.1. Offshore Oil and Gas Processing: Improving Efficiency of Energy Usage

New developments on offshore oil and gas primary processing represent opportunities to, cumulatively, improve efficiency of energy usage, reduce GHG emissions, lower topsides footprint and, consequently, reduce costs. All these effects contribute to increase the economic and environmental feasibility of offshore oil and gas production. The literature presents several recent works on energy assessment and optimization of primary processing on offshore oil and gas rigs, comprising measures to improve energy usage efficiency (Nguyen et al., 2016a),

better power generation schemes with organic cycles (Pierobon et al., 2013), air-bottoming cycles (Pierobon and Haglind, 2014), steam-bottoming cycles (Nguyen et al., 2014a), heat-exchanger network optimization for minimum energy consumption (Pierobon et al., 2013; Pierobon and Haglind, 2014; Nguyen et al., 2014a) and offshore power production via combined cycles (Rivera-Alvarez et al., 2015). To improve energy usage efficiency of the process as a whole, including carbon capture units such as post-combustion amine plants, Nguyen et al. (2014b; 2016b) constructed multi-objective frameworks for optimizing CO<sub>2</sub> mitigation alternatives and platform lifecycle, using as working scenario an ending-life oil and gas offshore platform in the Norwegian North Sea.

As easily verified in these works, offshore oil and gas production follows a standard practice of centralizing multiple production wells in a single offshore processing unit, with topside equipment designed to separate and process oil, gas and water (Asibor et al., 2013). At ultra-deep waters distant from coast, Floating Production Storage and Offloading (FPSO) units configure the preferred choice of platform type (Gallo et al., 2017; Araújo et al., 2017; Ataújo and de Medeiros, 2017). Several works (Rivera-Alvarez et al., 2015; Gallo et al., 2017; Araújo et al., 2017; Voldsund et al., 2013; Nguyen et al., 2013; Teixeira et al., 2016) performed steady-state energy and exergy analysis of gas and oil processing on FPSOs and other offshore rigs, unveiling the common fact that compressors and gas turbines (GT) are major power sinks and power sources where exergy is mostly destroyed (Voldsund et al., 2014). This is particularly true for centrifugal compressors operating at partial load, when Joule-Thomson depressurizing recycles are used to sustain gas flow rate at sufficient levels to prevent surge (GPSA, 2004), despite the great rate of exergy destruction entailed by such anti-surge strategies.

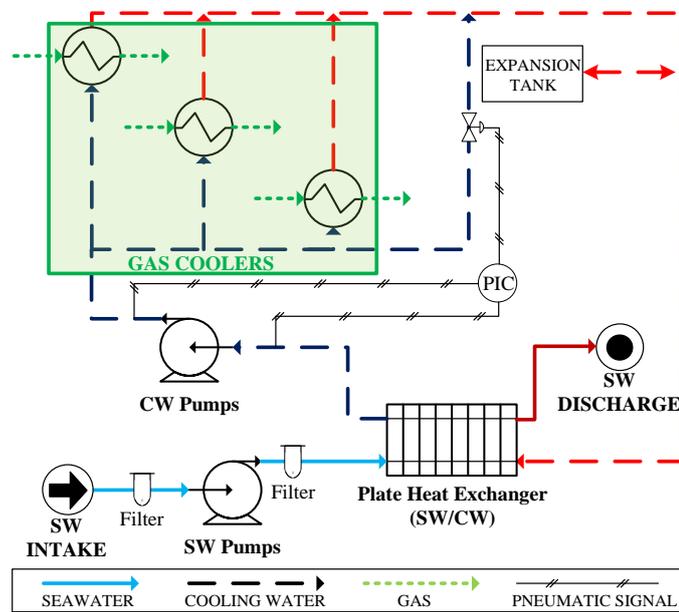
### 3.1.2. Deep Seawater Intake at Tropical Latitudes

Based on this last fact, the present work assesses the potential increase of energy usage efficiency of offshore gas processing by employing deep seawater (DSW) intake for primary cooling of a FPSO operating on deepwaters at tropical sites. As compressor machinery is dominant in gas processing FPSOs Eq (3.1), which is commonly used (GPSA, 2004) to calculate brake horsepower of centrifugal compressors, can demonstrate the interrelationship between temperature of primary cooling and FPSO energy usage efficiency, where  $r_p$ ,  $n$ ,  $\eta_p$ ,  $W$ ,  $q$ ,  $Z$ ,  $R$ ,  $T_l$ ,  $MW$  respectively represent compression ratio, polytropic exponent, polytropic efficiency, brake horsepower (kW), gas flow rate (kg/h), average compressibility factor, ideal

gas constant (8.314 kJ/kmol.K), gas inlet absolute temperature (K) and gas molar mass (kg/kmol). Eq (3.1) shows that the power of centrifugal compressors is nearly proportional to the gas inlet absolute temperature, which is indirectly linked to the primary cooling temperature.

$$W = \frac{q.Z.R.T_1 \left\{ r_p^{\frac{n-1}{n}} - 1 \right\}}{3600.\eta_p.MW.(n-1)/n} \quad (3.1)$$

On a FPSO the primary cooling source is seawater (SW), as displayed in Fig. 3.1, where it is seen that SW operates in open loop. It is aspirated by SW pumps, passes through plate heat exchangers and returns to the ocean. In the plate exchangers, SW absorbs heat from the closed-loop cooling water (CW) coming warm from process heat exchangers. The cooled CW is then returned to the process plant by CW pumps. The main CW demanding units are intercoolers and aftercoolers of gas compression trains, whose heat duties have the same magnitude of the powers of the respective precedent compression stages. Therefore, inlet SW temperature indirectly imposes a lower bound on the temperature of gas leaving intercoolers and aftercoolers, affecting total compression power and energy usage efficiency of FPSOs.

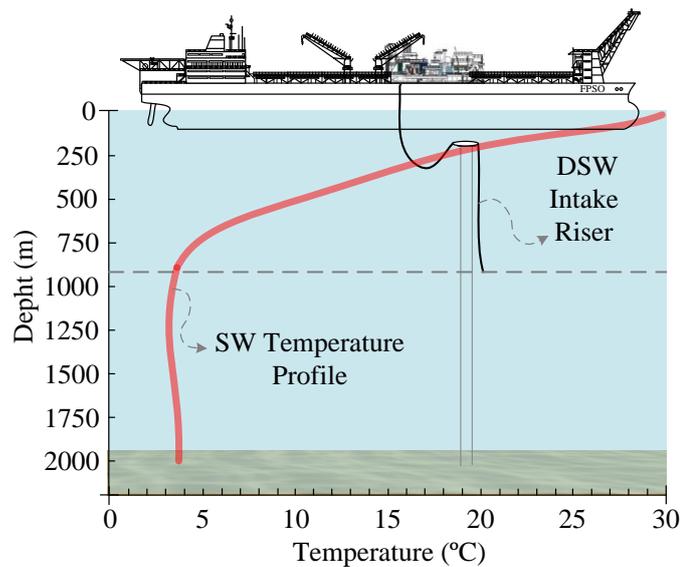


**Figure 3.1. FPSO CW circuit cooled by SW intake**

Shallow-water offshore platforms were also studied assuming SW intake at 8°C for primary cooling (Nguyen et al., 2016b; Pierobon et al., 2013; Pierobon and Haglind, 2014; Nguyen et al., 2014a; Voldsund et al., 2014). The underlying reason is that such units are situated at high

latitudes (e.g. Norwegian North Sea). On the other hand, at tropical or sub-tropical deepwaters – e.g. Gulf of Mexico, South Atlantic Brazilian Pre-Salt and South-East Asia – the surface SW is much hotter, attaining 25°C to 32°C in the summer. This entails lowest CW temperature near to 35°C, implying that hot compression gases can be cooled down to 40°C (313 K) only.

On the other hand, at tropical deepwaters, SW temperature falls continuously becoming almost constant beyond 800 m of depth and attaining a seabed temperature around 4°C as shown in Fig. 3.2 for tropical latitudes of the South Atlantic on Pre-Salt basin. Thus, DSW intake below 900 m of depth is less cost-effective, implying that the layer between 800 m and 900 m of depth is established as the best temperature-cost compromise for positioning DSW intake (Rogez, 2012). This demonstrates that DSW intake is more relevant for FPSOs located at tropical seas, where differences of up to 28°C can be observed between surface SW and DSW. On the other hand, on high latitude seas the maximum temperature difference from surface SW to DSW is less than  $\approx 8^\circ\text{C}$ , entailing that DSW intake is less interesting in such cases.



**Figure 3.2. DSW intake and thermal profile of deepwaters at tropical latitudes in the South Atlantic**

DSW intake at 4°C allows to cool down CW to 7°C and, consequently, compressor outlet gas to 12°C (285K), excepting in cases where there is risk of gas hydrate formation and/or of attaining lower bound of equipment working temperature. Eq. (3.1) shows that a decrease of 28K (e.g. from 313 K to 285 K) of gas inlet temperature reduces compression power approximately by 9%, which can be achieved by adopting DSW intake to cool the CW circuit. Additionally,

DSW intake leads to other indirect benefits: (i) elimination of refrigeration for Hydrocarbon Dew Point Adjustment (HCDPA); (ii) reduction of area and weight of heat exchangers; (iii) decrease of gas turbines power; (iv) decrease of fuel gas consumption; (v) elimination of one stage in CO<sub>2</sub> compression train (i.e., only three stages instead of the usual four); and (vi) reduction of the dehydration load of Temperature Swing Adsorption (TSA) units for Water Dew Point Adjustment (WDPA).

### 3.1.3. Present Work

The literature shows a growing interest in optimization of energy usage efficiency of offshore platforms. In this context, DSW intake is a factor to be considered for better energy usage efficiency of offshore rigs as it can effectively reduce compressor power on tropical deep-waters. Despite this, there are only a handful of literature works that studied DSW intake for obtaining resources and improving energy usage efficiency of plants, where only a fraction of them considered offshore gas processing applications. Rogez (2012) analyzed high capacity ( $\approx 30,000$  m<sup>3</sup>/h) 800 m depth DSW intake at 5°C to floating liquefied NG (FLNG) platforms and offshore thermal energy conversion (OTEC) systems, demonstrating its theoretical feasibility, but without field validation. Wei et al. (1980) discussed DSW for mariculture and nuclear power plant cooling. Blomster and Stanimirov (2004) considered the manufacture of  $\approx 600$ m polyethylene pipes for DSW applications. Petkovic et al. (1993) analyzed flexible or rigid ducts for suctioning 500 m depth DSW at 7°C. These studies were motivated by expected gains achieved with DSW intake in terms of energy, weight and costs, but did not attempt quantitative assessment of the technology.

The present work contributes to filling this gap. It is presented a quantitative assessment of the effects of using DSW intake in terms of power consumption, economic responses (capital investment and revenues) and CO<sub>2</sub> emissions of a typical oil and gas processing FPSO operating on the Brazilian Pre-Salt. The choice of this scenario is relevant for implementation of DSW alternative, due to the stringent design conditions of oil and gas FPSO's in this area, such as unusual high gas to oil ratio (GOR) and high %CO<sub>2</sub> in raw NG. These conditions demand proper CO<sub>2</sub> separation, CO<sub>2</sub>-rich fluid dispatch to enhanced oil recovery (EOR) and NG exportation to onshore plants (Araújo and de Medeiros, 2017). Compression of CO<sub>2</sub>-rich fluid to EOR and exportation gas, both at high pressures, has strong impact on capital investment and power consumption of the FPSO. In this context, the potential of innovation of

DSW intake was evaluated comparing conventional surface SW intake with DSW intake, in terms of the following items of performance: (i) power of compressors, GTs, CW pumps and SW/DSW pumps; (ii) gas processing plant layout; (iii) equipment weight, capital expenditures (CAPEX) and NG exportation revenues; (iv) power demand, fuel gas consumption and CO<sub>2</sub> emissions along project life; (v) energy usage efficiency of FPSO gas processing plant; and (vi) sensitivity analysis in carbon taxation scenarios.

### 3.2. Methods

Table 3.1 presents the two considered scenarios of SW/DSW intake for assessment of impacts on FPSO gas processing plant.

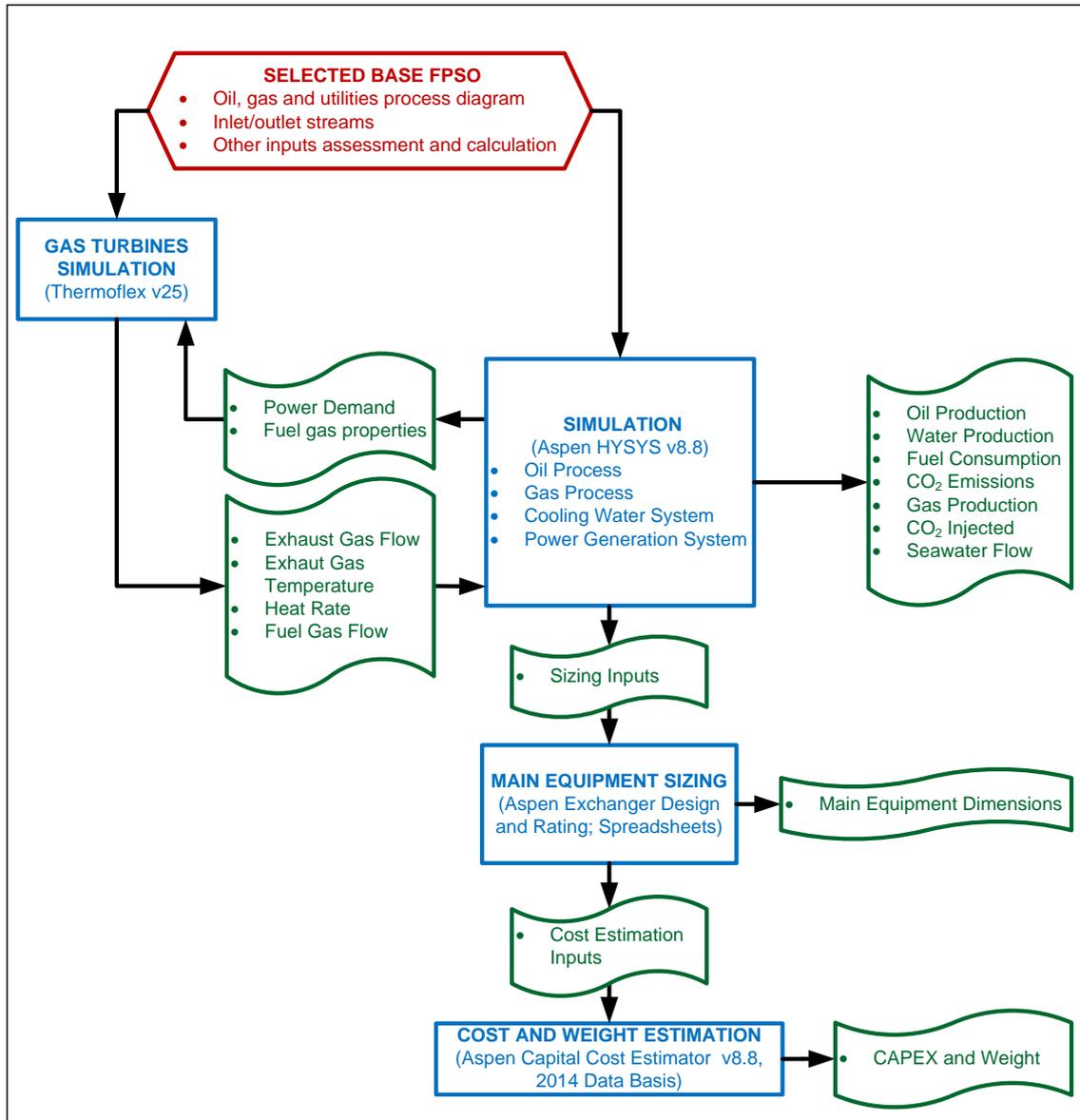
**Table 3.1. Scenarios for comparison of SW intake schemes**

	BASE-CASE	DSW-CASE
SW Intake Temperature	32°C	4°C
Cold CW Temperature	35°C	7°C
CW $\Delta T$	20°C	20°C
Hot CW Temperature	55°C	27°C
SW Outlet Temperature	<40°C	<12°C
Cooled Gas Temperature	40°C	12°C

CW  $\Delta T$ : CW change of temperature

The procedure depicted in Fig. 3.3 was applied to the FPSO gas processing plant under traditional surface SW intake as Base-Case and the alternative DSW-Case using DSW intake. The design gas processing capacity is considered 5 MMSm<sup>3</sup>/d. Both cases were evaluated under five gas processing loadings – 100%, 75%, 50% and 25% of full design capacity – resulting in ten simulation cases.

Anti-surge recycle loops were adjusted to maintain the design gas flow rate through compressors ( $\approx 5$  MMSm<sup>3</sup>/d). Specific simulation tools were employed considering the diversity of FPSO unit operations for oil, gas and utilities. The oil and gas processing plants – including CW/SW circuits and GTs – were simulated with HYSYS 8.8 (Aspentech) using Peng-Robinson Equation of State (PR-EOS) for thermodynamic modeling. GT simulations were validated with Thermoflex (Thermoflow Inc.), employing 25 simulations of the selected GT at same capacity and ambient conditions of temperature, pressure and air humidity. Equipment sizing and cost estimation were executed with HYSYS 8.8, Aspen Exchanger Design and Rate 8.8 and Aspen Capital Cost Estimator 8.8 (Aspentech).



**Figure 3.3. Procedure flowchart for this study**

### 3.2.1. Base-Case Definition

The operation conditions of FPSO *Cidade de Paraty* were chosen to define the Base-Case in this study. It operates at Brazilian Pre-Salt since 2013 with processing capacity of 5 MMSm<sup>3</sup>/d of gas and 100,000 bbl/d of oil. Information concerning this FPSO was extracted from the Environmental Impacts Assessment (EIA) of the Activity of Production and Flow of Oil and Gas from Santos Basin Pre-Salt Pole – Stage 1 (Petrobras, 2013), from now on simply referred as EIA. Real operational and inlet data from EIA are available in the Internal Appendix IA, of chapter 3 (Tables IA.1, IA.2 and IA.3, Figs. IA.1 and IA.2).

FPSOs operating on Pre-Salt fields are not allowed to flare associated gas. Therefore, gas processing plants contemplates CO<sub>2</sub> removal from NG, destination of the CO<sub>2</sub>-rich fluid to EOR, and exportation of treated NG through pipelines to onshore plants. This imposes high compression power demand, configuring an application niche for DSW alternative. Fig. IA.1 depicts the block diagram of the *Cidade de Paraty* topside processing plant, including CW system. The gas production profile along project life is supplied by EIA (Petrobras, 2013) in Fig. IA.2, where the gas profile of %CO<sub>2</sub> is an estimated curve built with the only two values, initial and final, provided by EIA, respectively, 8% mol and 55% mol. In the real situation, the continuous increase of CO<sub>2</sub> content of associated gas affects equally both DSW-Case and Base-Case: flow rate of re-injection compression trains (C-700 and C-600) increases, while the flow rate of NG exportation compressors (C-500) decreases. Due to such continually changing conditions and lack of all necessary data, the rigorous evaluation of this effect was not attempted in this work. Instead, the design %CO<sub>2</sub> of raw NG was herein assumed constant at 15% mol, the time-average %CO<sub>2</sub> of the raw gas profile in Fig. IA.2. This way the complexity of the problem and the number of explored cases could be kept at workable levels. The average gas plant feed is shown in Table IA.1.

### 3.2.2. Oil Processing Plant

The oil processing plant was simulated to obtain the flow rate, composition, temperature and pressure of the raw gas that feeds the gas processing plant. HYSYS extension GOR Adjustment was used for crude feed modeling. Flow rates, compositions, pressure and temperature of oil, gas and water streams in Table IA.1 were used as inputs. Three-phase crude arrives from thirteen risers after 2223 m of sub-sea elevation change from the reservoir at 41368 kPa and 40°C. Gas streams were separated from the crude feed at three pressures: 1850 kPa, 667.7 kPa and 250 kPa. GOR and water-to-oil ratio (WOR) targets were set to 268.2 Sm<sup>3</sup>/m<sup>3</sup> and 0.02537 m<sup>3</sup>/m<sup>3</sup>, respectively, to achieve the predicted flows of oil, gas and water considering the year (Fig. IA.2) of maximum gas production of 4.750184 MMSm<sup>3</sup>/d with 111462 bbl/d of oil and 2906 bbl/d of water. Oil plant process flow diagram (PFD) for simulation and GOR Adjustment settings are in Fig. A1.1 (Supplement A1 of Supplementary Materials at Appendix H).

### 3.2.3. Gas Processing Plant Simulation: Base-Case

Fig. 3.4 is a simplified PFD of the gas processing plant for the Base-Case. The simulation PFD is shown in Fig. A1.3 (Supplement A1 of Supplementary Materials at Appendix H) Gas streams 103, 206 and 201, coming from the high, medium and low-pressure oil-gas separators, feed the gas plant. Table 3.2 summarizes data of compressor sets.

**Table 3.2. Centrifugal compressors summary: Base-Case**

PFD AREA	100	200	500	600	700	900
Compressor Service	Main	Gas Recovery	Gas Export	CO <sub>2</sub>	Injection	C <sub>3</sub> Cycle
Number of Stages	1	2	2	4	1	1
Inlet Pressure (kPa)	1800	250	4500	400	25000	476.6
Discharge Pressure (kPa)	5200	1800	25000	25000	55000	1738
Gas cooler 1 ΔP (kPa)	50	25	50	25	50	-
Gas cooler 2 ΔP (kPa)	-	50	50	25	-	-
Gas cooler 3 ΔP (kPa)	-	-	-	50	-	-
Gas cooler 4 ΔP (kPa)	-	-	-	50	-	-
Pressure Ratio/Stage	2.9167	2.7708	2.3649	2.8385	2.2020	3.6466
1 <sup>st</sup> stage actual flow (m <sup>3</sup> /h)	12027	1317	2894	12939	313	1885
2 <sup>nd</sup> stage actual flow (m <sup>3</sup> /h)	-	1848	1110	4728	-	-
3 <sup>rd</sup> stage actual flow (m <sup>3</sup> /h)	-	-	-	1583	-	-
4 <sup>th</sup> stage actual flow (m <sup>3</sup> /h)	-	-	-	467	-	-
1 <sup>st</sup> stage polytropic efficiency	82%	74%	77%	83%	57%	75%
2 <sup>nd</sup> stage polytropic efficiency	-	75%	73%	79%	-	-
3 <sup>rd</sup> stage polytropic efficiency	-	-	-	74%	-	-
4 <sup>th</sup> stage polytropic efficiency	-	-	-	70%	-	-

Eq. (3.2) is used to calculate the compression ratio  $r_P$  of a compression stage belonging to a  $N$ -staged compressor train, inlet pressure  $P_0$  and final discharge pressure  $P_N$ , where  $\Delta P_i$  represents the head loss in the  $i^{th}$  intercooler. The polytropic efficiency is determined via GPSA (2004), which presents efficiency of centrifugal compressors for several flow rate ranges. These data allowed building the correlation of Fig. A1.4 (Supplement A1 of Supplementary Materials at Appendix H) to predict polytropic efficiency from gas flow rate.

$$P_0 * r_P^N - \sum_{i=1}^N \Delta P_i * r_P^{N-i} - P_N = 0 \quad (3.2)$$

Five simulations were executed. The first is a simulation for equipment sizing at design conditions adopting maximum compressor operational flow rates from EIA (Fig. IA.1). The four additional simulations assume 100%, 75%, 50% and 25% of the total gas plant capacity. The 100% capacity corresponds to 4.75 MMSm<sup>3</sup>/d of gas flow rate referring to the year of maximum gas production in Fig. IA.2. Flow rates of anti-surge recycles were adjusted to keep design flow rates of all compression stages.

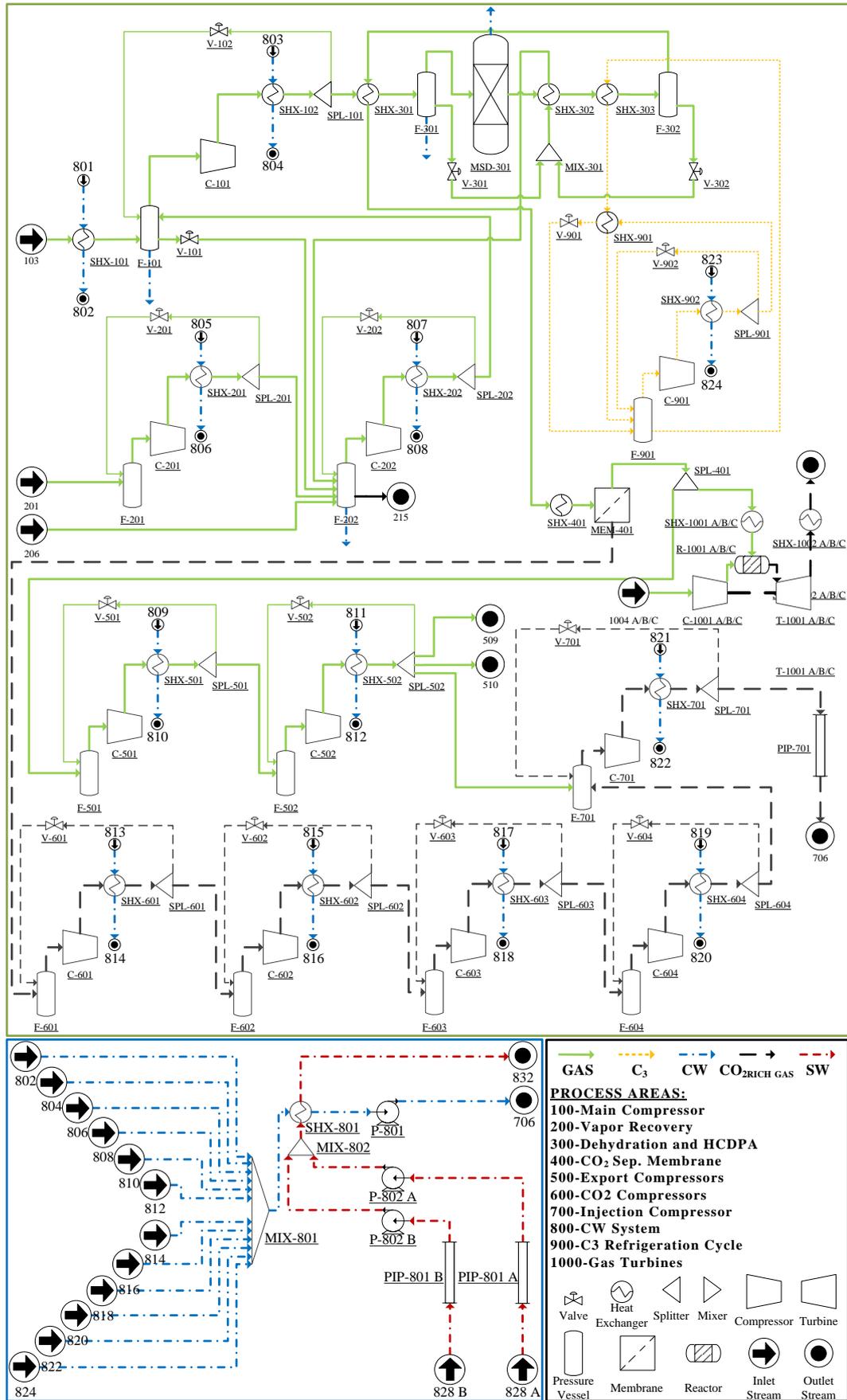


Figure 3.4. PFD of gas processing plant: Base-Case

Heat exchangers specifications, including CW and gas temperatures, are shown in Table A1.3, (Supplement A1, Suppl. Mat.). A propane refrigeration cycle cooled by CW was used for hydrocarbon dew-point adjustment (HCDPA). For a HCDP of 10°C, the propane evaporator temperature was 0°C with 476.6 kPa of dew pressure, while the condenser temperature was 50°C with bubble pressure of 1738 kPa. CO<sub>2</sub> separation was executed by a membrane permeation (MP) unit, which was the same in both Base-Case and DSW-Case, eliminating the need to calculate its contribution to costs and equipment weight in the comparative analysis. Despite MP units can be installed as customized user operations in HYSYS PFDs for designing MP area (Arinelli et al., 2017), here MP CAPEX has no importance, so that the MP unit was simulated via a simple stream calculator using as specifications the feed stream and the desired %CO<sub>2</sub> in retentate (5%mol) and in permeate (56.6%mol) as in EIA (Petrobras, 2013). Assuming permeation of only CO<sub>2</sub> and CH<sub>4</sub>, compositions and flow rates of permeate and retentate were determined by mass balances, while the respective temperatures followed via energy balance.

The CW circuit was designed considering gas processing at 100% capacity, giving the highest CW consumption, with most anti-surge loops disabled, except in the compressor train of CO<sub>2</sub> rich fluid to EOR. In this case the flow rate of EOR compressors was lower than their specific design value, corresponding to all processed gas being injected to EOR due to inexistent subsea pipelines.

#### 3.2.4. Gas Processing Plant Simulation: DSW-Case

The approach of the Base-Case was adapted according to the particularities of DSW-Case simplified PFD in Fig. 3.5, which corresponds to the simulation PFD in Fig. A1.5 (Supplement A1 of Supplementary Materials at Appendix H). Due to the lower CW temperature the process was modified: (i) Propane refrigeration cycle (Area 900) was unnecessary as a mere cooler attains the HCDPA target using CW at 7°C; (ii) gas at 12°C from intercoolers allowed to use only three CO<sub>2</sub> compression stages (Area 600), without overpassing compressor threshold temperature of ~180°C; (iii) at partial load (75%, 50%, 25% of maximum gas capacity), the anti-surge control is activated to keep compressor design flow rates.

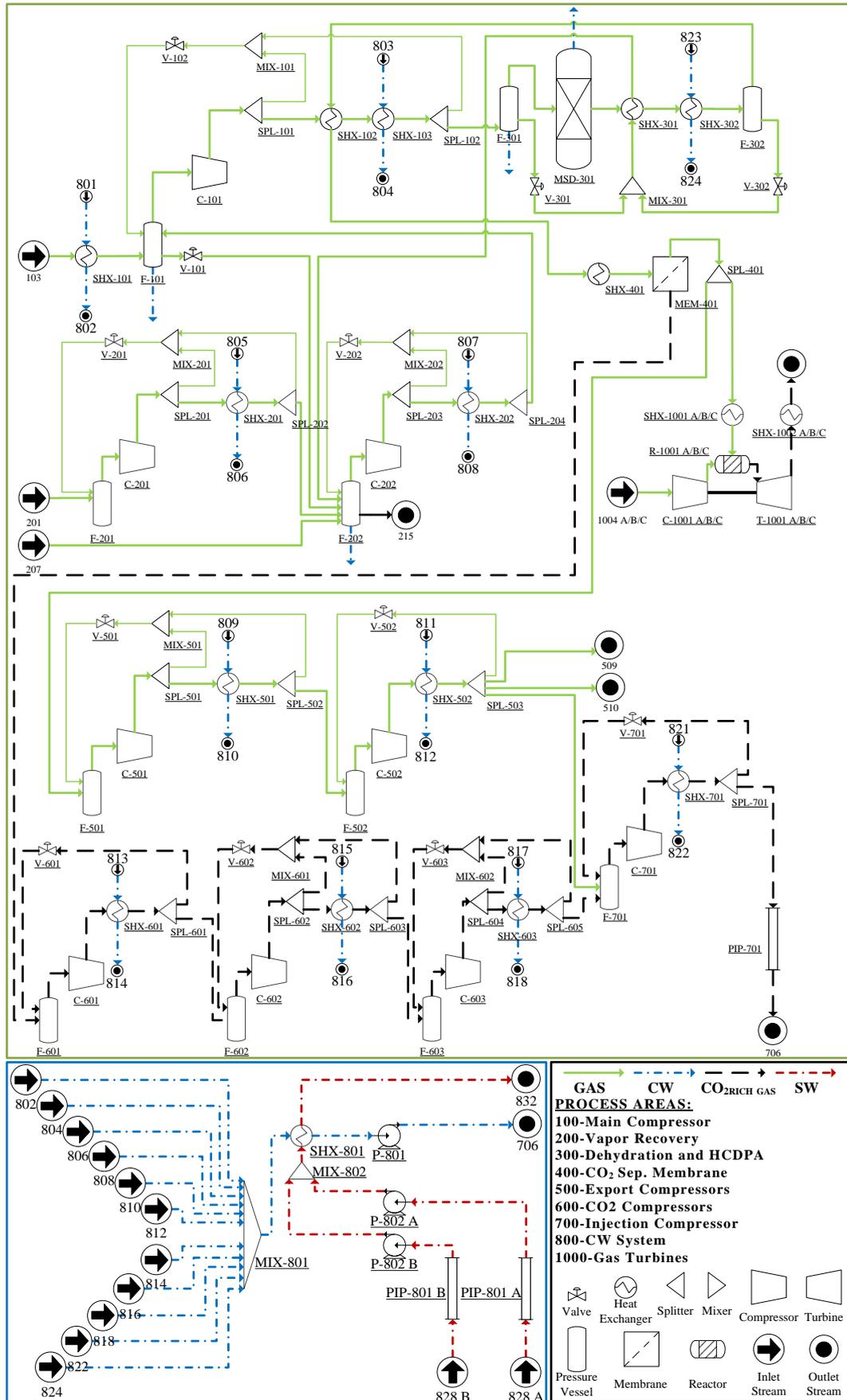


Figure 3.5. PFD of gas processing plant: DSW-Case

However, Joule-Thomson effects in anti-surge valves resulted in very low temperatures in some suction lines. That could lead to hydrate formation and freezing for non-dehydrated gas (e.g. compressors C-101, C-201 and C-202 in Fig. 3.5), besides issues of low carbon steel resistance. Hydrate and freezing conditions were detected via HYSYS Hydrate-Analysis tool, so that whenever hydrate conditions were detected, hot-bypasses (i.e. recycling hot gas from compressor outlet before intercooling) set the temperature after anti-surge valves to at least 3°C above hydrate temperature. Temperatures below 0°C were detected in compressors C-501, C-602 and C-603, and hot-bypasses were used to set suction temperature to 5°C, also avoiding expensive stainless-steel compressors. Anti-surge recycles and hot by-passes were adjusted to reach design flow rates and minimum temperatures of compression stages. Table 3.3 shows compressors parameters of the DSW-Case with supplementary information in Tables A1.4 to A1.7 (Supplement A1, Suppl. Mat.). Heat exchangers specifications – including CW and gas temperatures – are in Table A1.8 (Supplement A1, Suppl. Mat.)

**Table 3.3. Centrifugal compressors summary: DSW-Case**

	Process Area				
	100	200	500	600	700
Compressor Service	Main	Gas Recovery	Gas Export	CO <sub>2</sub>	Injection
Number of Stages	1	2	2	3	1
Inlet P (kPa)	1800	250	4500	400	25000
Discharge P (kPa)	5200	1800	25000	25000	55000
Gas cooler 1 ΔP (kPa)	50	25	50	25	50
Gas cooler 2 ΔP (kPa)	-	50	50	25	-
Gas cooler 3 ΔP (kPa)	-	-	-	50	-
Pressure Ratio/Stage	2.9167	2.7708	2.3649	3.9974	2.2020
1 <sup>st</sup> stage actual flow (Sm <sup>3</sup> /h)	10437	1317	2894	12888	313
2 <sup>nd</sup> stage actual flow (Sm <sup>3</sup> /h)	-	1431	835	2943	-
3 <sup>rd</sup> stage actual flow (Sm <sup>3</sup> /h)	-	-	-	581	-
1 <sup>st</sup> stage polytropic efficiency	82%	74%	77%	83%	57%
2 <sup>nd</sup> stage polytropic efficiency	-	74%	72%	77%	-
3 <sup>rd</sup> stage polytropic efficiency	-	-	-	71%	-

### 3.2.5. Gas Turbines (GT) Simulation

The Base-Case uses four aero-derivate GT model GE LM 2500, a spare included. Each GT has a dedicated waste heat recovery unit or WHRU (Teixeira et al., 2016; Nguyen et al., 2016b) fed with the respective exhaust gas. GT is simulated integrating an adiabatic air compressor, a combustion reactor and an adiabatic expander, as shown in Area 1000 of Figs. 3.4 and 3.5. GT simulation model in HYSYS was calibrated against GT simulation with Thermoflex 25 by adjusting compressor and expander adiabatic efficiencies to match power and air/fuel ratio to match exhaust temperature. At operation conditions of the Base-Case and the DSW-Case GT simulations were validated against Thermoflex results. Due to lower power demand of DSW-

Case, a smaller GE LM2500 GT model was prescribed, with 26.7 MW gross power (ISO conditions) – instead of the 28.2 MW GT model of Base-Case – burning fuel gas of nearly same composition of Base-Case in Table A.2. Table 3.4 summarizes GT information at three conditions.

**Table 3.4. Selected gas turbine (GT) data**

CONDITION	BASE-CASE			DSW-CASE		
	Iso	Design	Operation	Iso	Design	Operation
Fuel	Methane	Fuel Gas	Fuel Gas	Methane	Fuel Gas	Fuel Gas
Altitude (m)	0	0	0	0	0	0
P (kPa)	101.33	101.33	101.33	101.33	101.33	101.33
T (°C)	15	23	30	15	23	30
Relative Humidity	60%	87%	77%	60%	87%	77%
Filter ΔP (kPa)	0.0	1.0	1.0	0.0	1.0	1.0
Exhaust Duct ΔP (kPa)	0.0	0.5	0.5	0.0	0.5	0.5
Net power (MW) <sup>1</sup>	28.176	23.536	25.78	26.718	22.172	24.337
Compressor Efficiency <sup>2</sup>	83.9%	83.9%	84.2%	86.0%	85.2%	85.7%
Expander Efficiency <sup>2</sup>	88.0%	88.0%	88.0%	87.5%	87.5%	87.5%
Fuel Gas Flow (kg/s) <sup>1</sup>	1.53	1.537	1.648	1.459	1.46	1.571
Inlet Air Flow (kg/s) <sup>1</sup>	85.85	76.27	80.85	82.25	75.55	77.06
Fuel LHV (kJ/Sm <sup>3</sup> ) <sup>3</sup>	32824	40737	40737	32824	40750	40750
Heat Rate (kJ/kWh) <sup>1</sup>	9622	10213	9998	9676	10299	10096
GT Efficiency (Yield) <sup>1</sup>	37.4%	35.2%	36.0%	37.2%	35.0%	35.7%
Combustion T (°C) <sup>2</sup>	1192	1201	1196	1170	1183	1179
Exhaust T (°C) <sup>1</sup>	513	523.7	519.5	512.4	525.1	521.9
Exhaust Flow (kg/s) <sup>1</sup>	86.66	77.0	81.6	83.07	73.26	77.83
Fuel Consumption (MMSm <sup>3</sup> /d) <sup>1</sup>	0.198	0.142	0.152	0.1890	0.1345	0.1447
CO <sub>2</sub> Emissions (t/d) <sup>1</sup>	366	354	380	349	337	362

1 - From GE LM2500 GT simulation with Thermoflex

2 - From GT simulation with HYSYS

3 - Table IA.2

The adiabatic efficiency of GT air axial compressor varies from 78% to 87%, while the adiabatic efficiency of GT expander depends on temperature in the combustion chamber, varying from 84% at 1200°C to 92% at 1288°C (Boyce, 2002). Air temperature and relative humidity (RH) affect GT power output and were considered the same in HYSYS and Thermoflex runs, respectively as 23°C and 87% according to EIA – as design premise, the maximum air temperature and minimum RH were 30°C and 77%. This effect is represented in Fig. A1.6 (Supplement A1 of Supplementary Materials at Appendix H) as a power surface of GE LM2500 versus RH and ambient temperature.

### 3.2.6. SW Intake and CW Circuit

For the Base-Case (Fig. 3.4) the total CW flow rate through the SW/CW plate exchanger SHX-801 was obtained adding all CW exchanger outlets (streams 802 to 824), while SW was fed from two headers (PIP-801 A/B). The CW system was designed to operate at 100% of gas processing capacity of 5.0 MMSm<sup>3</sup>/d at 735.5 kPa of discharge pressure of pumps P-801 and P-802A/B. SW flow rate was calculated by a controller (SVC-801) setting HX-801 outlet temperature to 40°C, the SW disposal limit temperature according to local legislation (CONAMA, 2011). The NPSH is a critical issue in designing SW intake pipes as insufficient suction head leads to cavitation. Table 3.5 details the design of SW intake pipes for the Base-Case. The DSW-Case was designed with same parameters excepting the final DSW temperature of 11°C (Table 3.1) and the intake pipe length of 900 m giving 1550 m of total equivalent pipe length instead the 380 m of the Base-Case.

**Table 3.5. SW intake pipe sizing: Base-Case**

Total Equivalent Pipe Length (m) <sup>1</sup>	380
Net Elevation to Sea Level (m) <sup>2</sup>	-2.0
Absolute Pipe Roughness (mm) <sup>3</sup>	0.0015
Max Allowed $\Delta P$ (kPa) <sup>4</sup>	60
Suction Filter $\Delta P$ (kPa)	30
Max Flow Velocity (m/s)	2.0
External Heat Transfer	Isothermal
Max SW Flow Rate per Pipe (m <sup>3</sup> /h) <sup>5</sup>	2000

1 - SW intake at 30m depth and 350m of equivalent length;

2 - Fig. A1.7 (Supplement A1 of Supplementary Materials at Appendix H); 3 - HDPE pipes;

4 – Rogez (2012); 5 - Limited by pump maximum flow

### 3.2.7. Equipment Sizing

Equipment sizing was performed to allow cost and weight estimations. Reducing CW temperature strongly affects compressors, HCDPA, NG dehydration TSA units, GTs, pumps, NG intercoolers and aftercoolers. Heat exchangers demand special attention as they had to be resized due to DSW intake effects on flow rate and temperature of inlet and outlet process streams. Heat exchangers were designed (TEMA, 2007) with Eq. (3.3), where  $A_o$  is the effective heat transfer surface (m<sup>2</sup>),  $Q$  is the heat duty (kW),  $U$  is the overall heat transfer coefficient (kW/m<sup>2</sup>.K) and  $\Delta T_M$  is the corrected mean temperature difference (K) calculated via Eq. (3.4) for constant  $U$ , where  $LMTD$  is the log mean temperature difference (K),  $F$  is the

correction factor dependent of exchanger TEMA type and tube passes, and *GTTD* and *LTTD* are greater and lower terminal temperature differences (K).

$$A_o = Q / (U * \Delta T_M) \quad (3.3)$$

$$\Delta T_M = LMTD * F = \left( \frac{GTTD - LTTD}{\ln(GTTD / LTTD)} \right) * F \quad (3.4)$$

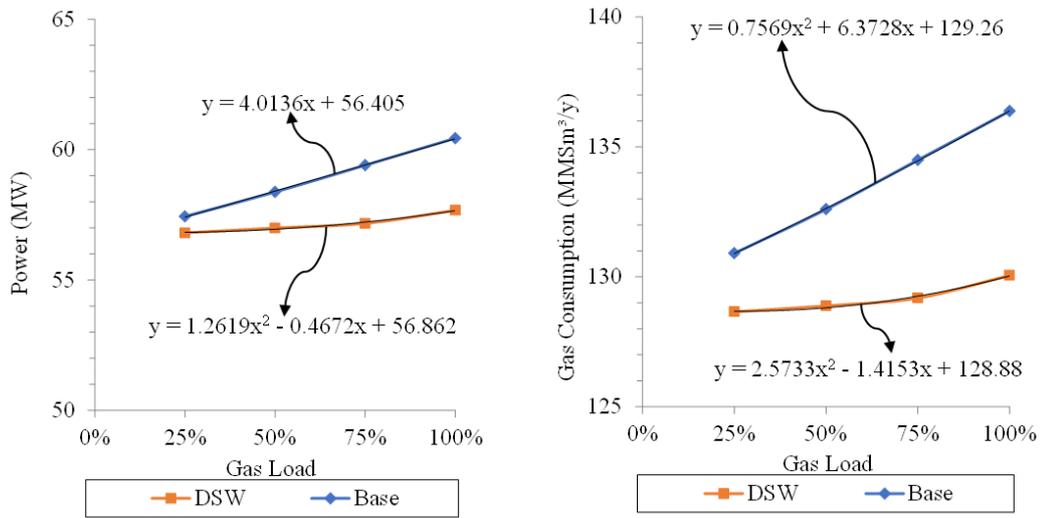
All variables in Eq. (3.3) are affected by changes of flow rate and inlet/outlet temperatures. Furthermore, construction materials, pressure rating and other parameters could also vary, impacting exchanger costs. Therefore, detailed heat exchanger sizing was necessary for evaluation of DSW intake impacts. Rigorous Aspen Exchanger Design and Rating v8.8 was used to size shell-and-tube exchangers and CW-SW plate exchangers, with all necessary information in Tables S2-1, S2-2, S2-3 and A2.4 (Supplement A2, Suppl. Mat.). In DSW-Case, the low temperature of compressed gas allows 6.4% reduction of water content in the NG feed to the molecular sieve TSA dehydration unit, entailing bed size reductions and/or extending TSA cycle, both advantageous aspects that were disregarded in the present analysis.

### 3.2.8. Equipment Costs and Weights

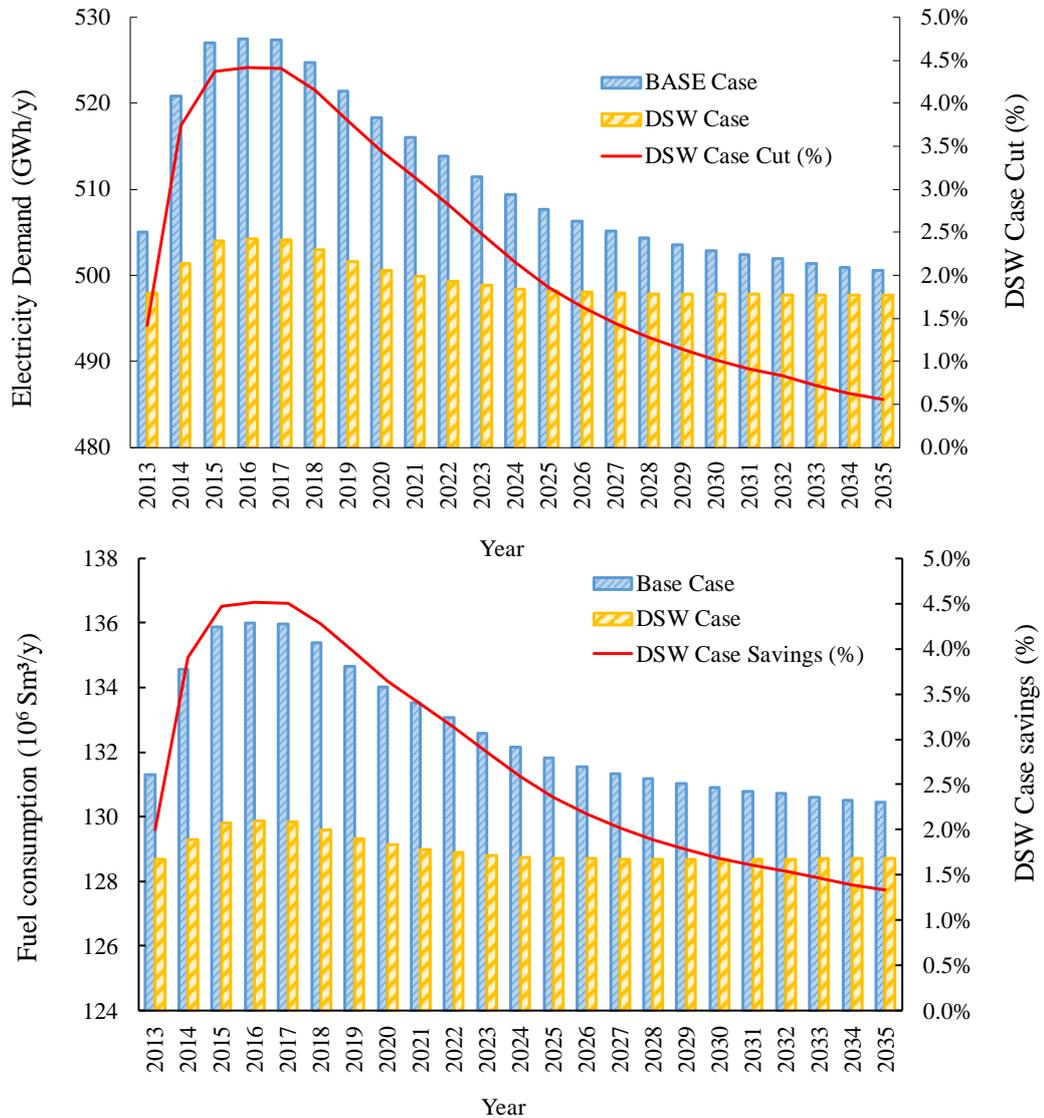
CAPEX and weight of equipment were estimated via Aspen Capital Cost Estimator 8.8 (ASPENTECH, 2014 data basis). Detailed information for such estimates is available in Tables A3.1 to A3.12 (Supplement A3, Suppl. Mat.).

### 3.2.9. Power and Fuel Gas Consumptions

The FPSO total power demand was based on plant simulation results and on complementary FPSO power consumption data from Martins et al. (2014) in Table A1.10 (Supplement A1, Suppl. Mat.). Results allowed calculation of fuel gas consumption at 100%, 75%, 50% and 25% of gas processing capacity, which were fitted against percentage of gas processing load in Fig. 3.6. These correlations were used to predict FPSO annual consumptions of power and fuel gas as shown in Fig. 3.7, assuming 8760 hours per year and process load between 18% and 100% of gas processing capacity according to raw gas feeds forecasted along 23 years of FPSO operation in Fig. IA.2. Tables IA.1 and IA.2 display the average compositions of raw NG and fuel gas, respectively.



**Figure 3.6. Power demand (y) and fuel gas consumption (z) vs gas processing load (x) for Base-Case and DSW-Case**



**Figure 3.7. Predicted FPSO consumptions of electricity and fuel gas**

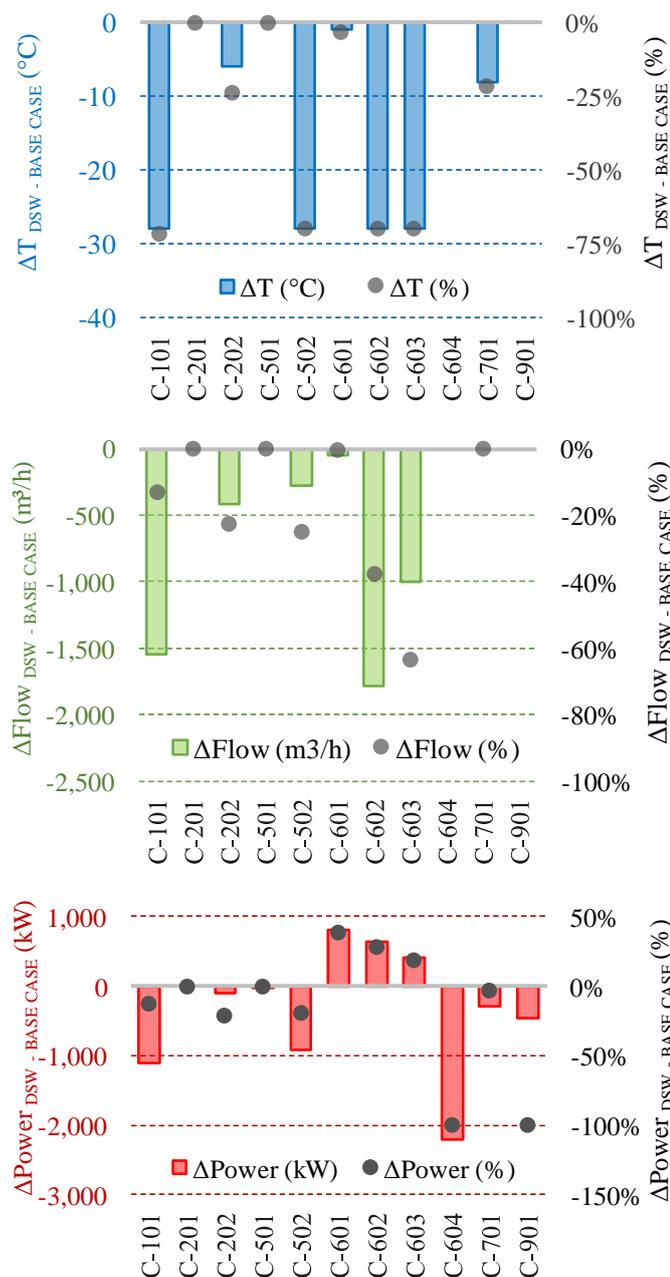
### 3.3. Results and Discussion

Simulations, equipment sizing and economic analysis generated a massive amount of data that is organized and used to calculate secondary information and final results. A selection of the most significant results is presented in this section. Complementary results are available in Supplement A4, Suppl. Materials.

#### 3.3.1. Process Simulation Results

##### 3.3.1.1. Compressors

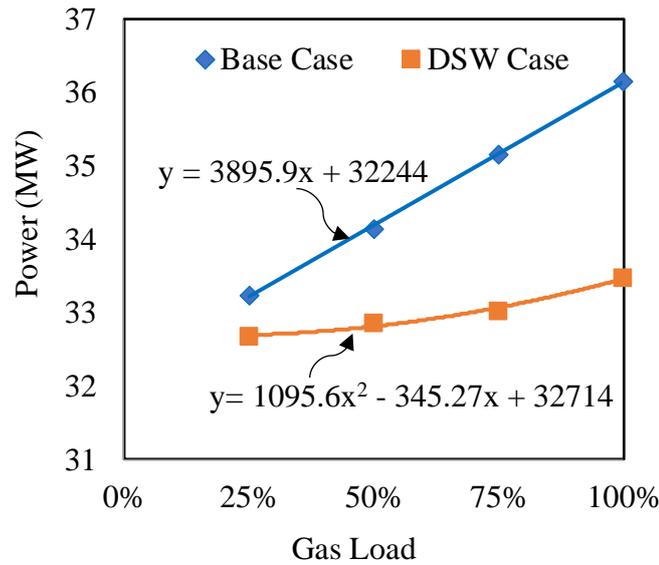
DSW intake at 4°C decreased total compression design power by 9.0% (3.217 MW), as shown in Table A4.1 (Supplement A4 of Supplementary Materials at Appendix H). However, larger variations of power were found for some compressors (Fig. 3.8), as compressor power does not depend solely on inlet gas temperature. Compressor C-901 was excluded due to elimination of the refrigeration cycle for HCDPA (i.e. 100% power reduction). Furthermore, substitution of surface SW intake (Base-Case) by DSW intake reduced one stage of the CO<sub>2</sub> rich gas compression train (C-600) (i.e. 100% power reduction of C-604). Compression ratio per stage moved from 2.8 to 4.0, increasing stage power from 18% up to 38%, compared to the same stages of the Base-Case. As a rule, the higher the number of intercooled compression stages, the lower the power of the compression train as a whole (as it approaches the isothermal compression limit) and the higher the investment – as the cost of  $N$  stages grows with  $\approx N*(Power/N)^{0.6}$ . Prescribing three stages in C-600 instead of four in DSW-Case entails an increase in the power of each stage for two reasons: (i) higher stage compression ratio, and (ii) higher temperature increase per stage. But, as a whole, the design power of C-600 was reduced by 4% or 0.35 MW. This was possible because the lower number of stages was compensated by the lower temperature of gas inlets in virtue of DSW primary cooling.



**Figure 3.8. Variation of compressor performances from Base-Case to DSW-Case**  
**\*C-604 and C-901 power reductions of 100% mean they are absent in DSW-Case**

Compressors C-202 and C-502 benefited from reduced inlet flow rate and temperature achieving  $\approx 20\%$  of power reduction at full gas load. Comparison of compressor powers at partial loads is shown in Figs. S4-1 and A4.2 (Supplement A4 of Supplementary Materials at Appendix H). Fig. 3.9 demonstrates that the total power savings achieved in DSW-Case decreases as raw NG feed flow rate decreases, since anti-surge flow rates increase. Also, gas cooling from Joule-Thomson effect in anti-surge recycling valves limits the benefits of colder CW.

Additionally, compressors exhibited different power behavior due to process constraints in DSW-Case: gas hydrate and ice formations and minimum operating temperature of compressor materials. Thus, DSW-Case requires hot-bypasses in some compressors, to avoid such low temperature issues. From 25% to 100% of gas processing capacity, DSW intake entails 3% to 9.2% of compressor power savings.



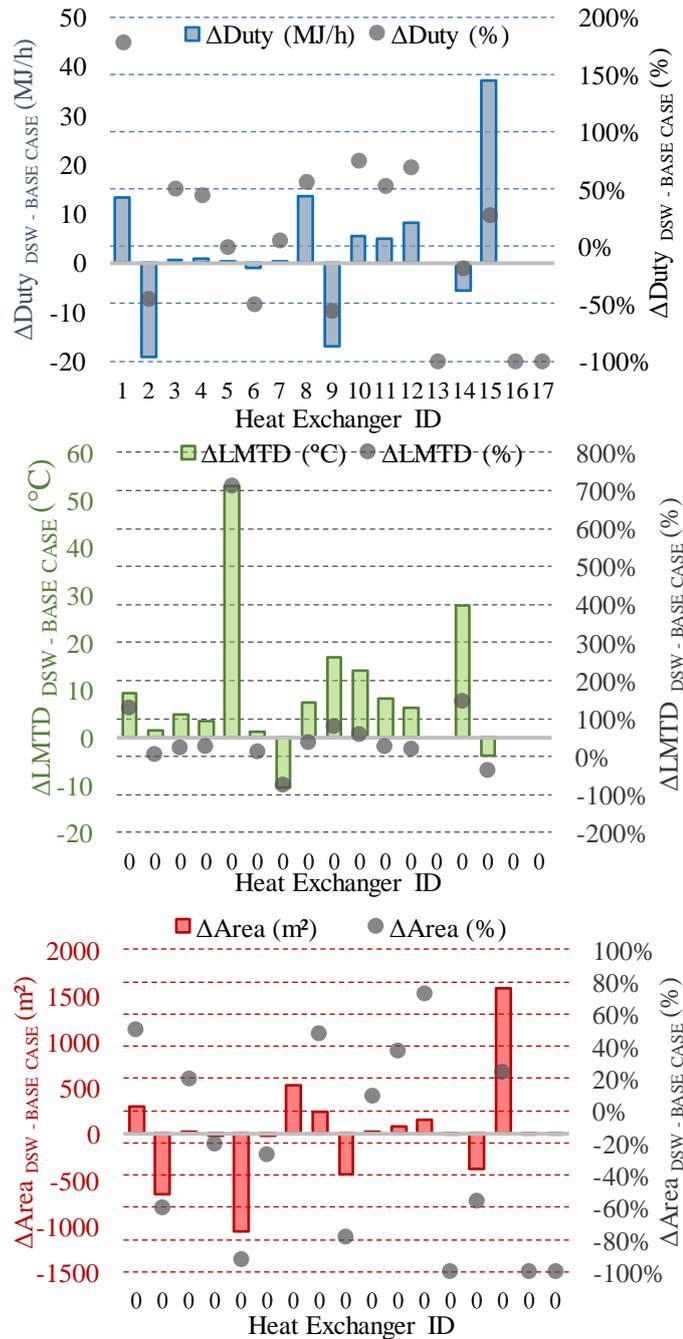
**Figure 3.9. Total compressors power (y, MW) vs gas processing load (x, %).**

As %CO<sub>2</sub> in associated gas increases along project lifetime, it is expected that flow rates of injection gas (C-700) and CO<sub>2</sub> rich gas (C-600) compression trains increase, accompanied by a decrease of the flow rate of NG exportation compressor C-500. On one hand, due to elimination of one compressor stage, the benefit of DSW intake on total design power of C-600 is modest (4% or 0.35 MW) relatively to Base-Case. On the other hand, NG compressors (C-500) running with extra gas cooling from DSW reached ≈1.0 MW lower design power. Thus, if %CO<sub>2</sub> increases in the raw NG feed, the advantage of DSW intake decreases. Nevertheless, if the original four stages of C-600 are maintained, the increase of %CO<sub>2</sub> in associated gas would favor DSW intake.

### 3.3.1.2. Heat Exchangers

Fig. 3.10 shows the differences in terms of duty, *LMTD* and area of heat exchangers, with exchanger IDs provided in Table A.3. Detailed exchanger results are shown in Table A4.2 (Supplement A4, Suppl. Mat.). A reduction of 6% in total heat exchangers duty was achieved

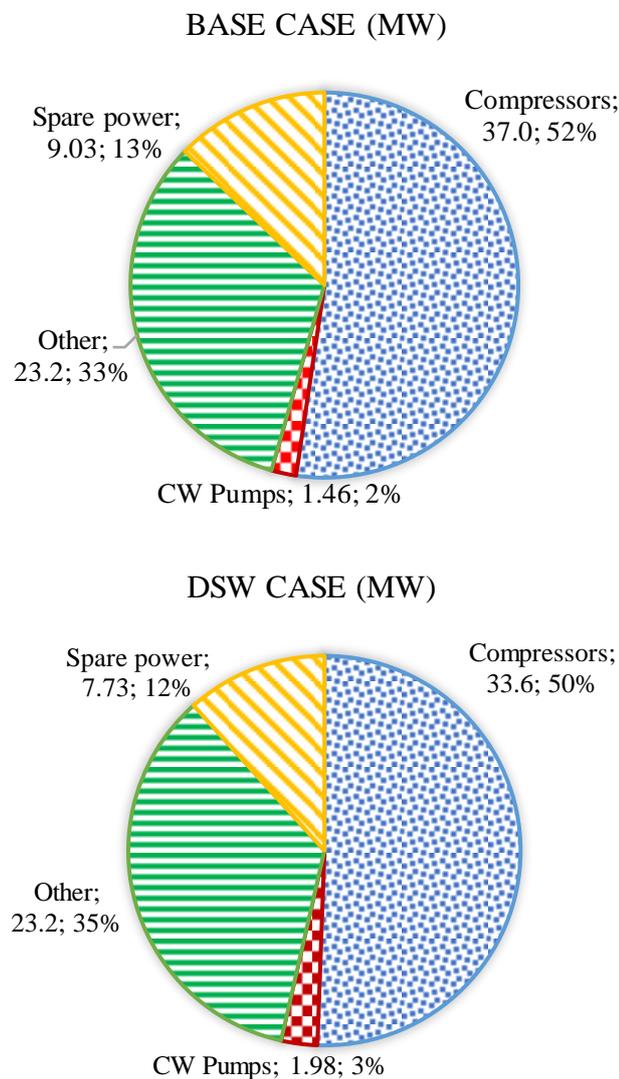
in the DSW-Case, but the total heat exchange area reduction was almost negligible (1.6%). But considering shell-and-tube exchangers apart from SW plate exchangers, the respective area reduction reaches 29%. This entails an expressive decrease in weight and costs, because shell-and-tube exchangers for high-pressure gas have higher specific cost (\$/m<sup>2</sup>) and weight (kg/m<sup>2</sup>) relatively to low-pressure SW plate exchangers.



**Figure 3.10. Variations of heat exchanger performances from Base-Case to DSW-Case**  
 \*Exchangers 13, 16 and 17 with 100% of area and duty reductions were absent in DSW-Case

### 3.3.1.3. Gas Turbines (GTs)

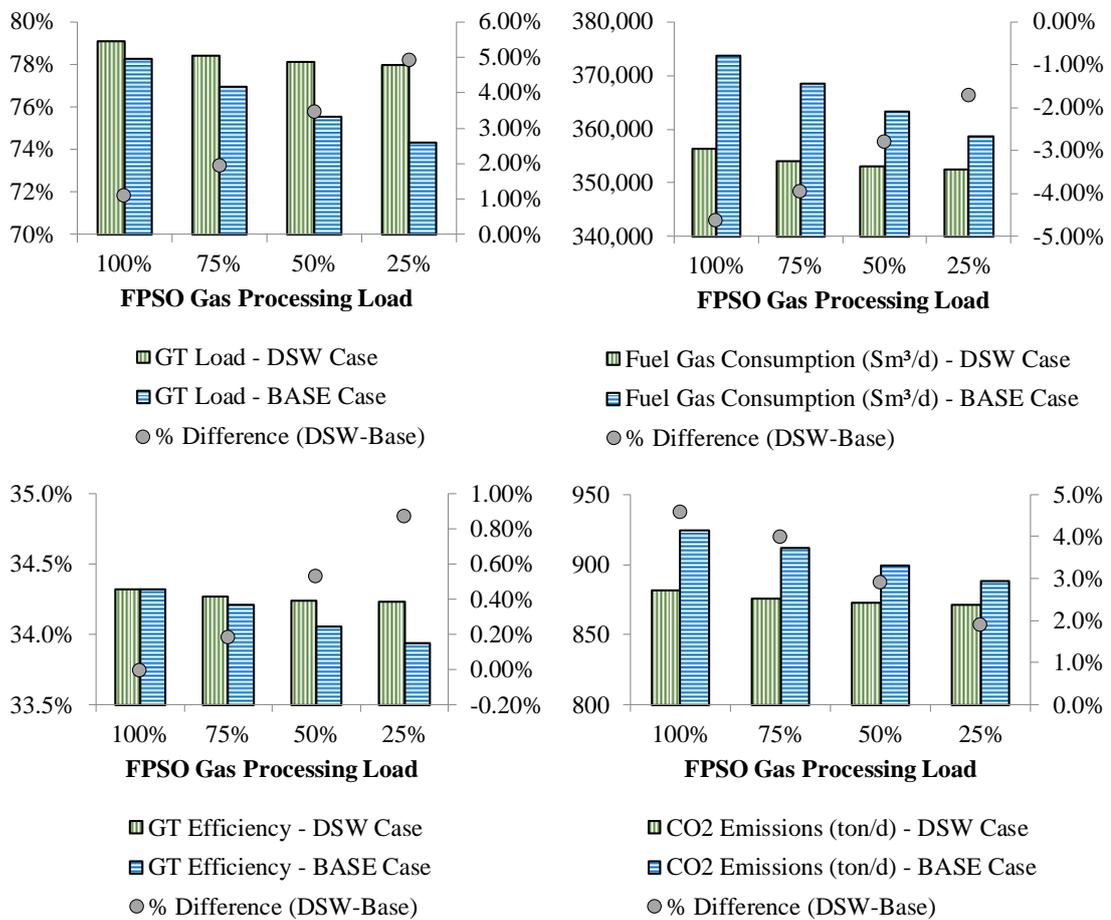
Total power demands were calculated as 61.6 MW and 58.8 MW for Base-Case and DSW-Case, respectively. Considering three operating GTs, the average power per GT was 20.5 MW for Base-Case and 19.6 MW for DSW-Case. Fig. 3.11 shows the calculated power demand of compressors, CW pumps and other FPSO consumers of power. The spare power corresponds to the idle share of total generation capacity, considering the three selected GT power generation at design condition (30°C and 94% RH), i.e. 3 x 23.5 MW for Base-Case and 3 x 22.2 MW for DSW-Case.



**Figure 3.11. FPSO power consumers (MW) by equipment category**

GTs simulation with HYSYS reproduced the same performance of the respective simulations with Thermoflex 25 for Base-Case and DSW-Case. An example of validation for full gas load

at design conditions (air at 23°C and 87% RH) is shown in Table A4-3 (Supplement A4 of Supplementary Materials at Appendix H) and GT performance comparisons at partial load are depicted in Figs. A4.2 and A4.3 (Supplement A4 of Supplementary Materials at Appendix H). Fig. 3.12 summarizes the most relevant results. It is shown that GTs of DSW-Case operate with loads from 1% to 5% greater than for the Base-Case, but the effect on efficiency is negligible at 100% gas load and reaches less than 1% for 25% gas load. CO<sub>2</sub> emissions are directly proportional to fuel gas consumption, presenting reductions between 1.5% and 4.5% depending on % gas load.



**Figure 3.12. Gas turbine (GTs) relative performances of DSW-Case and Base-Case vs % gas plant load**

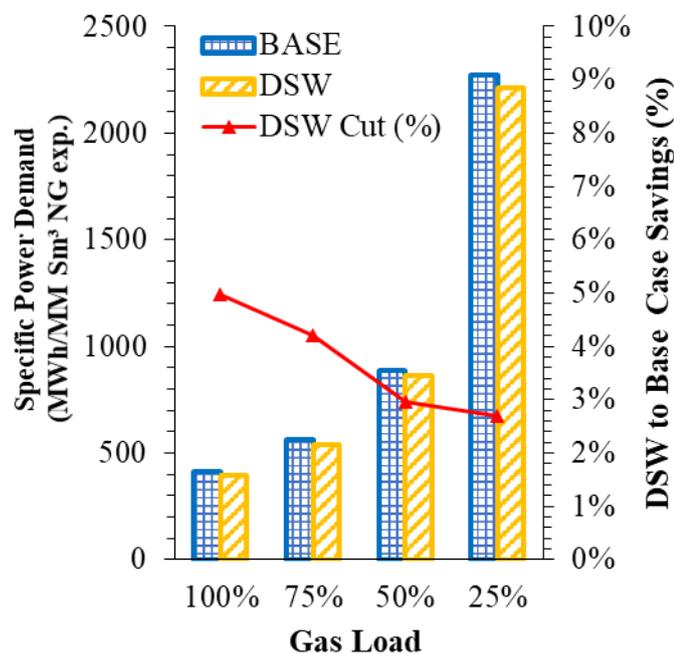
### 3.3.1.4. Pumps

The design SW flow rate of the DSW-Case was 31% greater than Base-Case. Nevertheless, this result is almost irrelevant because, as shown in Fig. 3.11, pumps are responsible for less than

3% of total FPSO power consumption. A summary of pumps results is in Table A4.4 (Supplement A4, Suppl. Mat.).

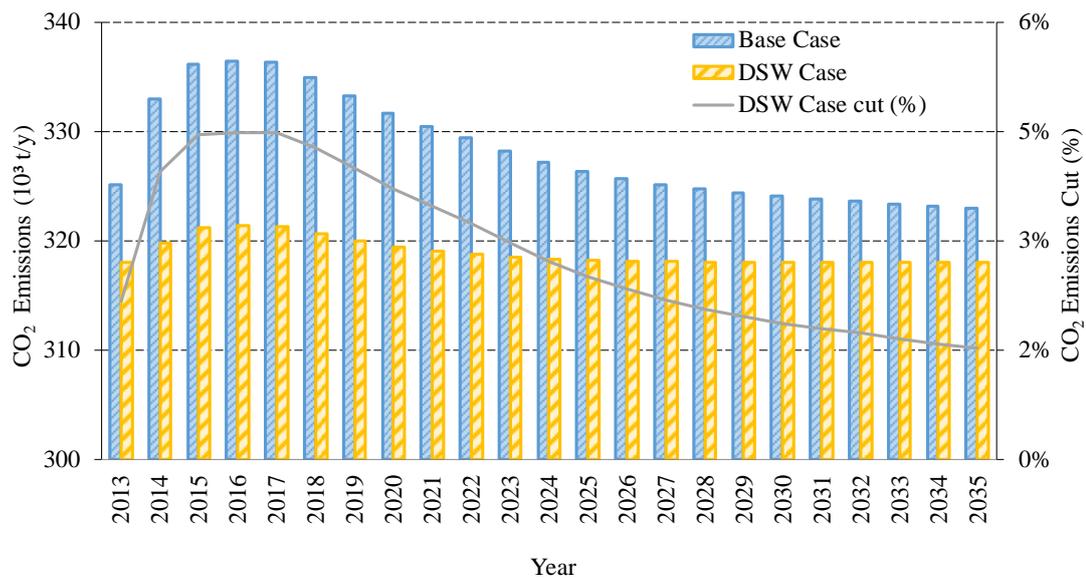
### 3.3.2. Energy Usage Efficiency and CO<sub>2</sub> Emissions

The energy usage efficiency of the gas plant is compared in terms of MWh/Sm<sup>3</sup> of treated NG exported. From 25% up to 100% of gas load, the DSW-Case achieved an increase from 2.7% up to 5.0% in energy usage efficiency as shown in Fig. 3.13.

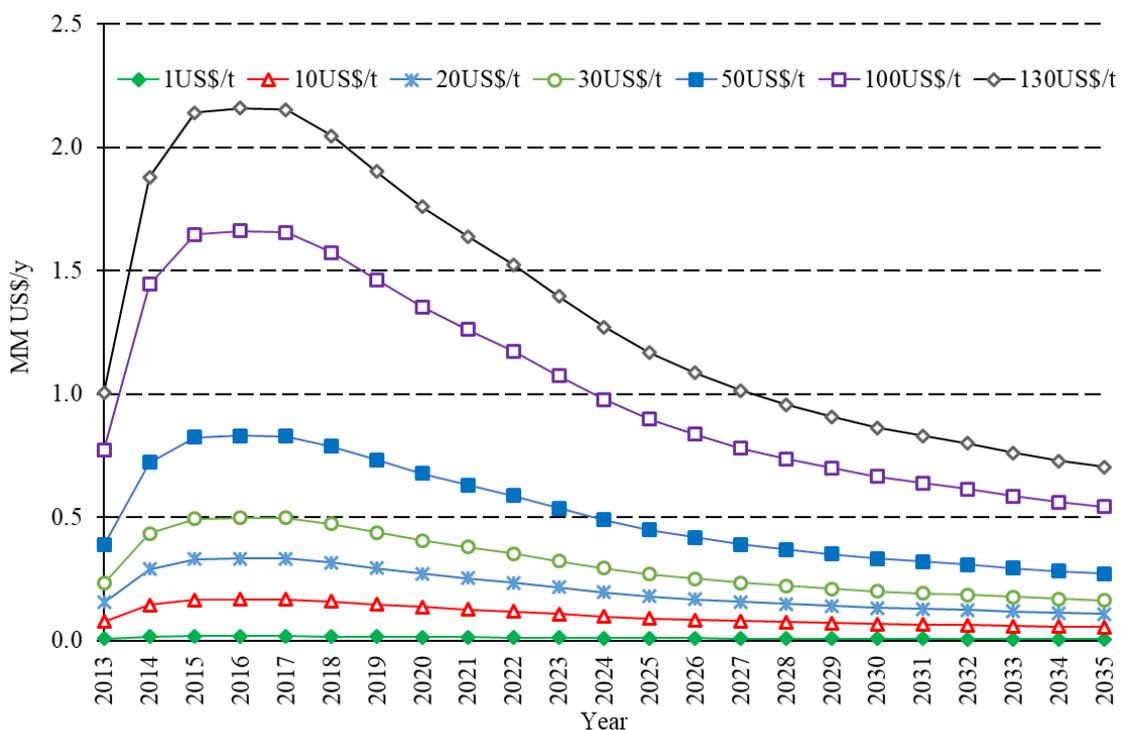


**Figure 3.13. FPSO Specific Power Demand (MWh/MM Sm<sup>3</sup> NG exported) and its % savings in DSW-Case relative to Base-Case vs gas plant % load**

Fig. 3.14 shows the comparison of CO<sub>2</sub> emissions for DSW-Case and Base-Case along the FPSO lifetime. DSW-Case emits 1.5% to 4.2% less CO<sub>2</sub> than Base-Case, depending on the year of operation. An average emission reduction of 9327 tCO<sub>2</sub>/y was achieved. The total reduction for 23 years of operation is 214,500 tCO<sub>2</sub>, or 3% of Base-Case emissions. Besides making FPSO operation more environmentally sound, DSW intake can be more profitable in case of carbon taxation as shown in the sensitivity analysis of Fig. 3.15, where the range of carbon taxes is based on World Energy Outlook 2016 (IEA, 2016). The annual extra savings due to lower carbon emissions of DSW-Case were negligible for 1 US\$/tCO<sub>2</sub> taxation but could become relevant for higher carbon taxes. Considering in Fig. 3.15 the maximum carbon taxation (130 US\$/t) and gas production, the savings could reach 2.1 MMUS\$/y.



**Figure 3.14. FPSO CO<sub>2</sub> emissions along project lifecycle**



**Figure 3.15. Sensitivity of avoided carbon taxation costs (MMUS\$/y) with carbon tax along FPSO lifecycle with DSW-Case**

### 3.3.3. Equipment Weight and Capital Costs

Equipment weight and CAPEX were estimated with ASPEN Capital Cost Estimator 8.8. A summary of results is presented in Table 3.6. Detailed results of equipment weight and CAPEX are available for Base-Case and DSW-Case in Tables A4.5 to A4.10, (Supplement A4, Suppl.

Mat.). CAPEX savings in the DSW-Case does not include the potential reduction of FPSO hull cost due to reducing topside weight. Additionally, an extra income of 6.2 MMUS\$ (net present value) is achieved by the DSW-Case, considering extra NG revenues due to lower fuel gas consumption. Total savings correspond to 1.33% of FPSO total fixed investment estimated as 1.6 Billion US\$, while the estimated CAPEX reduction by DSW intake is ~15 MMUS\$. It is worth noting that DSW intake may become less attractive according to the magnitude of its lifecycle cost – CAPEX plus all maintenance costs of DSW intake piping and connections. Hence, the obtained savings in virtue of DSW intake of ~21 MMUS\$ considering CAPEX and OPEX of the gas plant, may be overshadowed by the lifecycle cost of the intake piping itself, in which case the DSW intake solution would be rendered economically unfeasible.

A rough estimative of (non-installed) piping cost for DSW intake at 900 m of depth is ~1.0 MMUS\$ for two parallel 26” x 1200 m risers (710 mm nominal diameter) made of high density polyethylene (HDPE 100), with nominal pressure class 1600 kPa (PN 16) and standard diameter ratio 11 (SDR 11), which is rated at ~US\$ 440/m. However, the real lifecycle cost depends on the installed structure to keep the risers in the right position (buoys, anchors, cables) and its maintenance expenses. Moreover, the challenge to implement the new technology goes beyond economic calculations. Intake design, pipelines lifetime, installation and maintenance are determinant aspects to prove the technical economic feasibility of DSW intake technology.

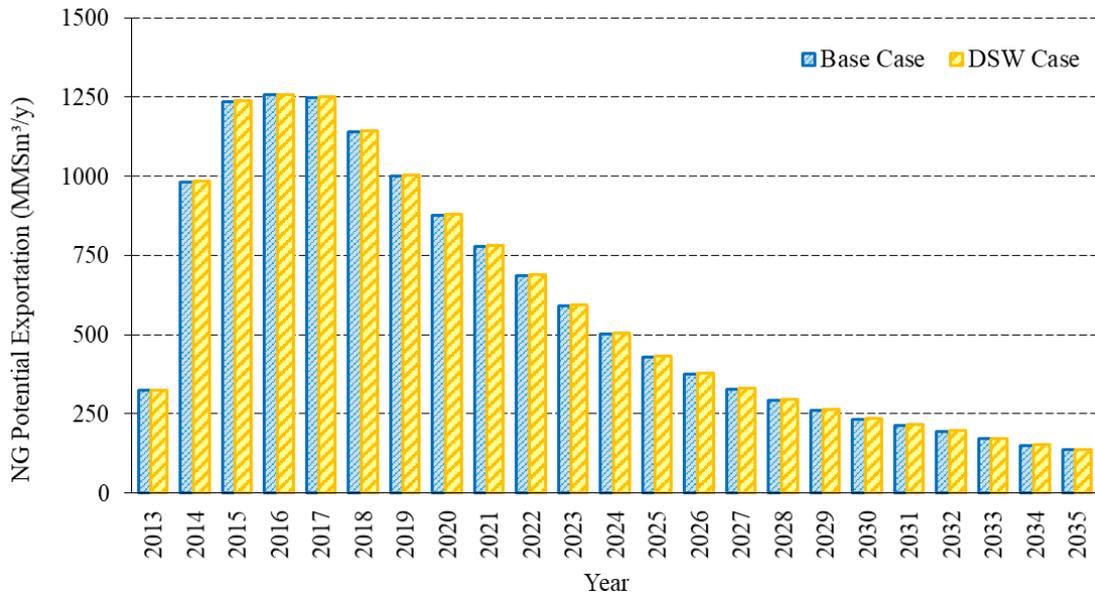
**Table 3.6. Summary of equipment weight and CAPEX results**

Equipment	Total Weight (t)		$\Delta$ (t)	$\Delta$ (%)	CAPEX (MM US\$)		$\Delta$ (MMUS\$)	$\Delta$ (%)
	BASE CASE	DSW CASE			BASE CASE	DSW CASE		
Compressors	349	306	43	-14	80.2	77.2	-2.93	-3,7
Exchangers	529	357	150	-42	39.2	34.4	-4.81	-12
GTs	902	858	44	-5.1	36.9	29.2	-7.69	-21
CW Pumps	18	21	-3.4	16	1.97	2.39	0.42	22
TOTAL	1798	1572	-226	-14	158	143	-15	-9.5

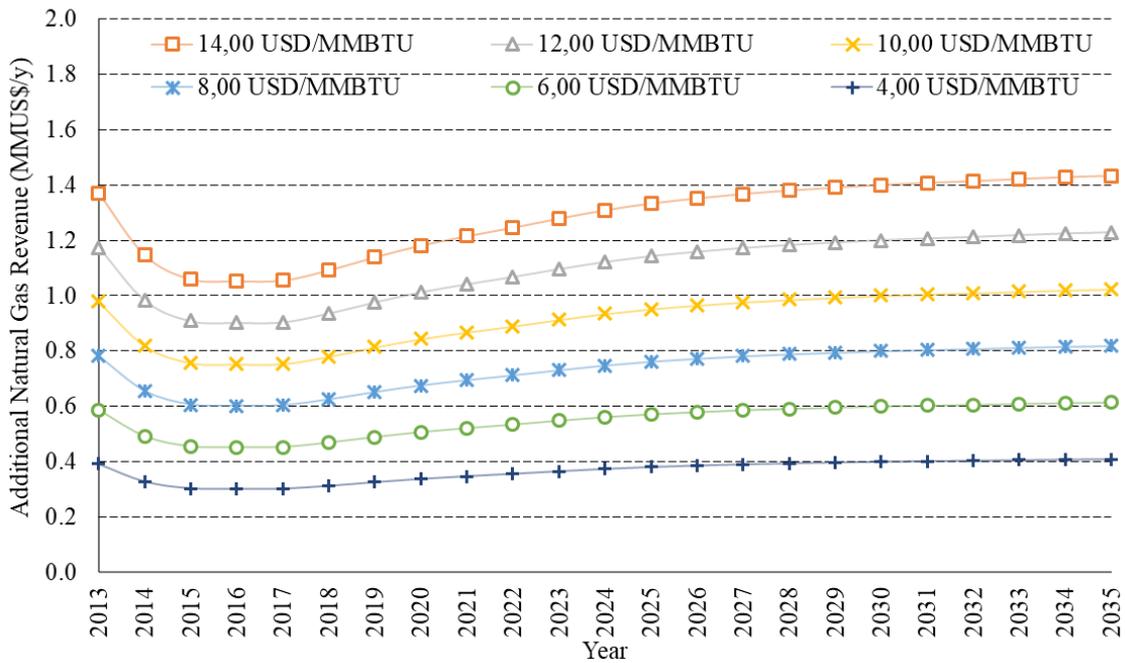
### 3.3.4. Extra NG Revenues

As seen in the DSW-Case, DSW intake can lower fuel gas consumption and increase NG exportation from 0.2% to 2.0% on FPSO lifecycle according to % gas load. However, the actual additional NG revenues are lower than the resulting reduction of fuel gas burned, because part of the additional treated gas is reinjected due to unavailability of sub-sea pipelines

whose implementation normally follows a different scheduling. Nevertheless, DSW intake could allow extra revenues according to the availability of pipeline capacity to export such extra gas. This potential extra NG exportation effect is shown in Fig. 3.16. Fig. 3.17 depicts a sensitivity analysis of the potential extra annual revenues from NG exportation for NG price ranging from 4 US\$/MMBTU to 14 US\$/MMBTU.



**Figure 3.16. NG potential exportation of Base-Case and DSW-Case along FPSO lifecycle**



**Figure 3.17. Sensitivity of extra NG annual revenues (MMUS\$/y) with NG price along FPSO lifecycle**

### 3.4. Conclusions

This work investigated the gains of power consumption, economic responses and CO<sub>2</sub> emissions as conventional surface SW intake is replaced by deep seawater (DSW) intake at 900 m of depth for use as primary cooling of FPSOs processing CO<sub>2</sub> rich NG at tropical deepwaters. The investigation consisted in comparing a Base-Case gas plant with conventional surface SW intake against the proposed alternative DSW-Case gas plant with DSW intake. The Base-Case corresponds to a selected FPSO (*Cidade de Paraty*) operating in Brazil Pre-Salt basin and processing a nominal flow rate of 5 MMSm<sup>3</sup>/d of raw NG with average CO<sub>2</sub> content of 15%mol. The resulting variations of gas processing layout, electricity generation capacity, energy usage efficiency and CO<sub>2</sub> emissions were assessed using rigorous thermodynamic simulation and specific engineering software for detailed process design and cost analysis. Results showed reduction of CO<sub>2</sub> emissions and increase in energy usage efficiency from 2.7% to 5.0%, depending on % gas processing load, where the highest value corresponds to full gas processing load.

Equipment sizing and costs were estimated for Base-Case and DSW-Case and compared. DSW intake promotes modest CAPEX and topside weight savings if compared to total FPSO cost and weight, but results are expressive considering only the gas processing and electricity generation modules. Furthermore, DSW intake also leads to other indirect advantages like less fuel gas consumption, less CO<sub>2</sub> emission, less FPSO power demand, elimination of the refrigeration cycle for hydrocarbon dew-point adjustment (HCDPA) and 6% reduction of water content in the gas feed to dehydration TSA units for water dew-point adjustment (WDPA), which could result in extended TSA cycle and/or smaller molecular sieve bed, lowering TSA weight and CAPEX. Modifications on gas processing capacity, HCDPA and WDPA systems, NG CO<sub>2</sub> content, exportation/injection ratio and other processing characteristics would lead to different DSW intake impacts on FPSO energy usage efficiency, CAPEX, operational costs and equipment weight.

Although the study has reached its main goals, there were some limitations of scope due to the huge size of the problem and the already massive calculations to report. Firstly, the effect of the continuous increase of CO<sub>2</sub> content in the associated gas along project lifetime was disregarded; i.e. the analysis only considered the average lifetime value of 15%mol CO<sub>2</sub> in the raw NG. Secondly, for the same reason, it was not possible to investigate DSW intake effects

over the usual range of FPSO gas processing capacities and CO<sub>2</sub> content in the feed gas, e.g., 2.0 – 12.0 MMSm<sup>3</sup>/d and 5% mol - 80% mol, respectively. Thirdly, the impact of DSW intake on gas dehydration units using expensive TSA cycles was detected as advantageous, but was not adequately measured. In the same way, it was not assessed the potential impact of DSW intake on certain specific gas processing alternatives such as HCDPA via Joule-Thomson expansion or supersonic separator instead of propane refrigeration; CO<sub>2</sub> separation via chemical absorption or supersonic separator or cryogenic distillation instead of MP, or even hybrid capture processes (e.g., chemical absorption coupled to MP) instead of pure MP.

Additionally, the DSW flow along the intake piping was considered isothermal at 4°C, which is not realistic as some heavy insulation would be necessary on the 900 m risers, impacting DSW intake costs. Nevertheless, it is straightforward to estimate the profile of temperature along the intake risers in order to obtain DSW temperature at the inlet of the plate heat exchanger (DSW/CW). Another aspect has to do with the lack of specificity of the cost estimation method for offshore plants and ultra-high pressures (> 20000 kPa), probably resulting in underestimated equipment fixed investment (CAPEX). In connection to this point, lifecycle cost evaluation of the installed piping for DSW intake must also be performed in order to determine the ultimate feasibility of DSW intake technology.

By last, the feasibility of DSW intake should be also investigated within a continuous or mixed-integer non-linear optimization framework so that certain features that were assumed pre-defined and constant in this study could vary in order to seek optimum values, configurations and dimensioning. For instance, the present analysis showed that certain FPSO units become problematic if cooled with cold CW at 7°C, being preferable to use traditional CW with conventional thermal range 35°C-55°C. Therefore a possible optimization formulation would consider two independent CW circuits – 35°C-55°C CW and 7°C-27°C CW, the latter requiring more expensive lines with insulation, etc – whose service heat loads, allocation points, exchanger/pump dimensioning and circulation flow rates are continuous or mixed-integer decision variables to be sought. Opportunely, two independent seawater intakes could also be considered as primary cooling – conventional SW and DSW – whose flow rates and intake dimensioning (e.g. riser diameters and pumps) are also passive of mixed-integer optimization. In this case, it is conceivable to let free the exiting temperature of DSW in the plate exchanger to reach the maximum environmental limit of 40°C for disposal (counterpointing the narrow DSW thermal range stipulated in Table 3.1), this way reducing DSW flow rate and the cost of

deep-intake piping, insulations, pumps and footprint for the same total heat load. Relaxing the exiting temperature of DSW seems a reasonable point to be questioned by optimizations as it has some thermodynamic support in the context of exergy analysis. In the present study DSW is returned to the sea at 11°C only (Table A1.8, Supplement A1, Suppl. Mat.) with a huge flow rate of 1.64 m<sup>3</sup>/s (Table A4.4, Supplement A4, Suppl. Mat.). This represents a valuable flow of exergy (relative to reference SW environment at 32°C) being wasted in the present DSW implementation. This flow of wasted exergy could be reduced by returning a lower flow rate of hotter DSW to the sea at the expenses of using larger heat exchangers in the process.

### 3.5. References of Chapter 3

Araújo O de QF, Reis A de C, de Medeiros JL, Nascimento JF, Grava WM, Musse APS. Comparative analysis of separation technologies for processing carbon dioxide rich natural gas in ultra-deepwater oil fields. *J. Clean. Prod.* 2017; 155:12–22, doi:10.1016/j.jclepro.2016.06.073.

Araújo O de QF and de Medeiros JL. Carbon capture and storage technologies: present scenario and drivers of innovation. *Current Opinion in Chemical Engineering* 2017, 17, 22–34, doi 10.1016/j.coche.2017.05.004

Arinelli LO, Trotta TAF, Teixeira AM, de Medeiros JL, Araújo O de QF. Offshore Processing of CO<sub>2</sub> Rich Natural Gas with Supersonic Separator versus Conventional Routes. *J. of Nat. Gas Sci. and Eng.* 2017, 46, 199-221, doi 10.1016/j.jngse.2017.07.010

Asibor E, Marongiu-Porcu M, Economides MJ. Oil production optimization without natural gas constraints: The harvesting of upstream natural gas. *J. Nat. Gas Sci. Eng.* 2013; 15:59–68, doi:10.1016/j.jngse.2013.09.004.

Blomster TJ, Stanimirov M. Continuously extruded long length polyethylene pipes for seawater intakes and marine outfalls. *Desalination* 2004; 166:275–286, doi:10.1016/j.desal.2004.06.082.

Boyce MP. *GasTurbine Engineering Handbook*. 2nd Ed. Houston, Gulf Professional Publishing, 2002.

CONAMA. Resolução CONAMA 430/2011. DOU 2011:9.

Gallo WLR, Gallego AG, Acevedo VL, Dias R, Ortiz HY, Valente BA. Exergy analysis of the compression systems and its prime movers for a FPSO unit. *J. Nat. Gas Sci. Eng.* 2017; 44:287–298, doi:10.1016/j.jngse.2017.04.023.

GPSA. *Engineering Data Book*. 20th Ed., Gas Processors Suppliers Association, 2004.

IEA. *World Energy Outlook 2016*. 2016.

IOGP. *Environmental Performance Indicators - 2015 data*. London: 2016.

- Martins PM, Delfino B, Fachini R. Projeto de Sistemas Oceânicos II. FPSO Campo de Libra. UFRJ 2014. [http://www.deno.oceanica.ufrj.br/deno/prod\\_academic/relatorios/2014/RafaelFachini+BrunoDelfino/relat2/relat2\\_completo.htm](http://www.deno.oceanica.ufrj.br/deno/prod_academic/relatorios/2014/RafaelFachini+BrunoDelfino/relat2/relat2_completo.htm) (accessed January 1, 2017).
- Nguyen T-V, Pierobon L, Elmegaard B, Haglind F, Breuhaus P, Voldsund M. Exergetic assessment of energy systems on North Sea oil and gas platforms. *Energy* 2013; 62:23–36, doi:10.1016/j.energy.2013.03.011.
- Nguyen T-V, Tock L, Breuhaus P, Maréchal F, Elmegaard B. Oil and gas platforms with steam bottoming cycles: System integration and thermoenviromonic evaluation. *Appl. Energy* 2014; 131:222–237, doi:10.1016/j.apenergy.2014.06.034.
- Nguyen T-V, Fülöp TG, Breuhaus P, Elmegaard B. Life performance of oil and gas platforms: Site integration and thermodynamic evaluation. *Energy* 2014; 73, doi:10.1016/j.energy.2014.06.021.
- Nguyen T-V, Voldsund M, Breuhaus P, Elmegaard B. Energy efficiency measures for offshore oil and gas platforms. *Energy* 2016; 117:325–40, doi:10.1016/j.energy.2016.03.061.
- Nguyen T-V, Tock L, Breuhaus P, Maréchal F, Elmegaard B. CO<sub>2</sub>-mitigation options for the offshore oil and gas sector. *Appl. Energy* 2016; 161:673–694, doi:10.1016/j.apenergy.2015.09.088.
- Petkovic MAL, Sattamini SR, Moraes CAC, Cavalcanti GN. Sistema de captação de água a grandes profundidades para plataformas semi-submersíveis de produção. PI9102045A, 1993.
- Petrobras. Polo Pre-Sal - Bacia de Santos - Etapa 1. IBAMA 2013. <http://licenciamento.ibama.gov.br/Petroleo/Producao/Producao - Polo Pre-Sal - Bacia de Santos - Etapa 1 - Petrobras/> (accessed July 4, 2017).
- Pierobon L, Nguyen T Van, Larsen U, Haglind F, Elmegaard B. Multi-objective optimization of organic Rankine cycles for waste heat recovery: Application in an offshore platform. *Energy* 2013; 58, doi:10.1016/j.energy.2013.05.039.
- Pierobon L, Haglind F. Design and optimization of air bottoming cycles for waste heat recovery in off-shore platforms. *Appl. Energy* 2014; 118, doi:10.1016/j.apenergy.2013.12.026.
- Rivera-Alvarez A, Coleman MJ, Ordonez JC. Ship weight reduction and efficiency enhancement through combined power cycles. *Energy* 2015; 93, doi:10.1016/j.energy.2015.08.079.
- Rogez F. Deep large seawater intakes: a common solution for Floating LNG in oil & gas industry and OTEC in marine renewable energy. 4th Int. Conf. Ocean Energy, 2012, 1–6.
- Rui Z, Li C, Peng F, Ling K, Chen G, Zhou X. Development of industry performance metrics for offshore oil and gas project. *J. Nat. Gas Sci. Eng.* 2017; 39:44–53, doi:10.1016/j.jngse.2017.01.022.
- Teixeira AM, Arinelli LO, de Medeiros JL, Araújo OQF. Exergy Analysis of monoethylene glycol recovery processes for hydrate inhibition in offshore natural gas fields. *J. of Nat. Gas Sci. and Eng.* 2016, 35, 798-813, doi 10.1016/j.jngse.2016.09.017

TEMA. Standards of the Tubular Exchanger Manufacturers Association. 9th ed. Tarrytown, NY: Tubular Exchanger Manufacturers Association, INC.; 2007.

Voldsund M, Ertesvag IS, He W, Kjelstrup S. Exergy analysis of the oil and gas processing on a north sea oil platform a real production day. Energy 2013; 55:716–727, doi:10.1016/j.energy.2013.02.038.

Voldsund M, Nguyen T-V, Elmegaard B, Ertesvåg IS, Røsjorde A, Jøssang K, et al. Exergy destruction and losses on four North Sea offshore platforms: A comparative study of the oil and gas processing plants. Energy 2014; 74:45–58, doi:10.1016/j.energy.2014.02.080.

Wei C, Huang B-J, Kong M-S. Engineering Analysis of Pumping Cold Deep Nutrient-Rich Seawater for Mariculture and Nuclear Power Plant Cooling. Ocean. Eng. 1980; 7:501–520.

### 3.6. Appendix 3A: *Cidade de Paraty* Plant, Gas Production, Inlet Streams, Fuel Gas and Exchanger IDs

**Table 3A.1. Gas plant inlet streams for simulation from EIA**

	OIL	GAS	WATER
Temperature (°C)	40	40	40
Pressure (kPa)	41368	41368	41368
Composition (mol fraction)			
CO <sub>2</sub>	0.0000	0.1513	0.0000
H <sub>2</sub> S	0.0000	0.0000	0.0000
N <sub>2</sub>	0.0000	0.0051	0.0000
Methane	0.0000	0.6396	0.0000
Ethane	0.0000	0.0924	0.0000
Propane	0.0050	0.0591	0.0000
i-Butane	0.0019	0.0099	0.0000
n-Butane	0.0081	0.0207	0.0000
i-Pentane	0.0065	0.0052	0.0000
n-Pentane	0.0122	0.0075	0.0000
C <sub>6</sub> *	0.0295	0.0063	0.0000
C <sub>7</sub> *	0.0559	0.0009	0.0000
C <sub>8</sub> *	0.0777	0.0016	0.0000
C <sub>9</sub> *	0.0658	0.0005	0.0000
C <sub>10</sub> *	0.0604	0.0000	0.0000
C <sub>11</sub> *	0.0405	0.0000	0.0000
C <sub>12</sub> *	0.0537	0.0000	0.0000
C <sub>13</sub> *	0.0488	0.0000	0.0000
C <sub>14</sub> *	0.0436	0.0000	0.0000
C <sub>15</sub> *	0.0308	0.0000	0.0000
C <sub>16</sub> *	0.0339	0.0000	0.0000
C <sub>17</sub> *	0.0224	0.0000	0.0000
C <sub>18</sub> *	0.0236	0.0000	0.0000
C <sub>19</sub> *	0.0224	0.0000	0.0000
C <sub>20</sub> +	0.3573	0.0000	0.0000
H <sub>2</sub> O	0.0000	0.0000	1.0000
MM C <sub>20</sub> +		500	
Density C <sub>20</sub> +		0.9496	

**Table 3A.2. Fuel Gas Properties (Base-Case from EIA)**

	Base Case	DSW Case
Pressure (kPa)	3500	3500
Temperature (°C)	22.5	22.5
LHV (kJ/kg)	43487	43486
HHV (kJ/kg)	47848	47847
Molar Mass (kg/kgmol)	22.91	22.92
COMPOSITION (Mol Fraction)		
N <sub>2</sub>	0.00672	0.00673
CO <sub>2</sub>	0.05000	0.05000
Methane	0.69910	0.70100
Ethane	0.14380	0.14000
Propane	0.06802	0.06966
n-Butane	0.01684	0.01687
n-Pentane	0.00567	0.00583
Hexane	0.00093	0.00099
Isobutane	0.00892	0.00892

**Table 3A.3. Heat Exchangers IDs in Fig. 3.10**

Base-Case		DSW-Case	
ID in Fig. 3.10	TAG	ID in Fig. 3.10	TAG
1	SHX-101	1	SHX-101
2	SHX-102	2	SHX-103
3	SHX-201	3	SHX-201
4	SHX-202	4	SHX-202
5	SHX-301	5	SHX-102
6	SHX-302	6	SHX-301
7	SHX-303	7	SHX-302
8	SHX-501	8	SHX-501
9	SHX-502	9	SHX-502
10	SHX-601	10	SHX-601
11	SHX-602	11	SHX-602
12	SHX-603	12	SHX-603
13	SHX-604	14	SHX-701
14	SHX-701	15	SHX-801
15	SHX-801		
16	SHX-901		
17	SHX-902		

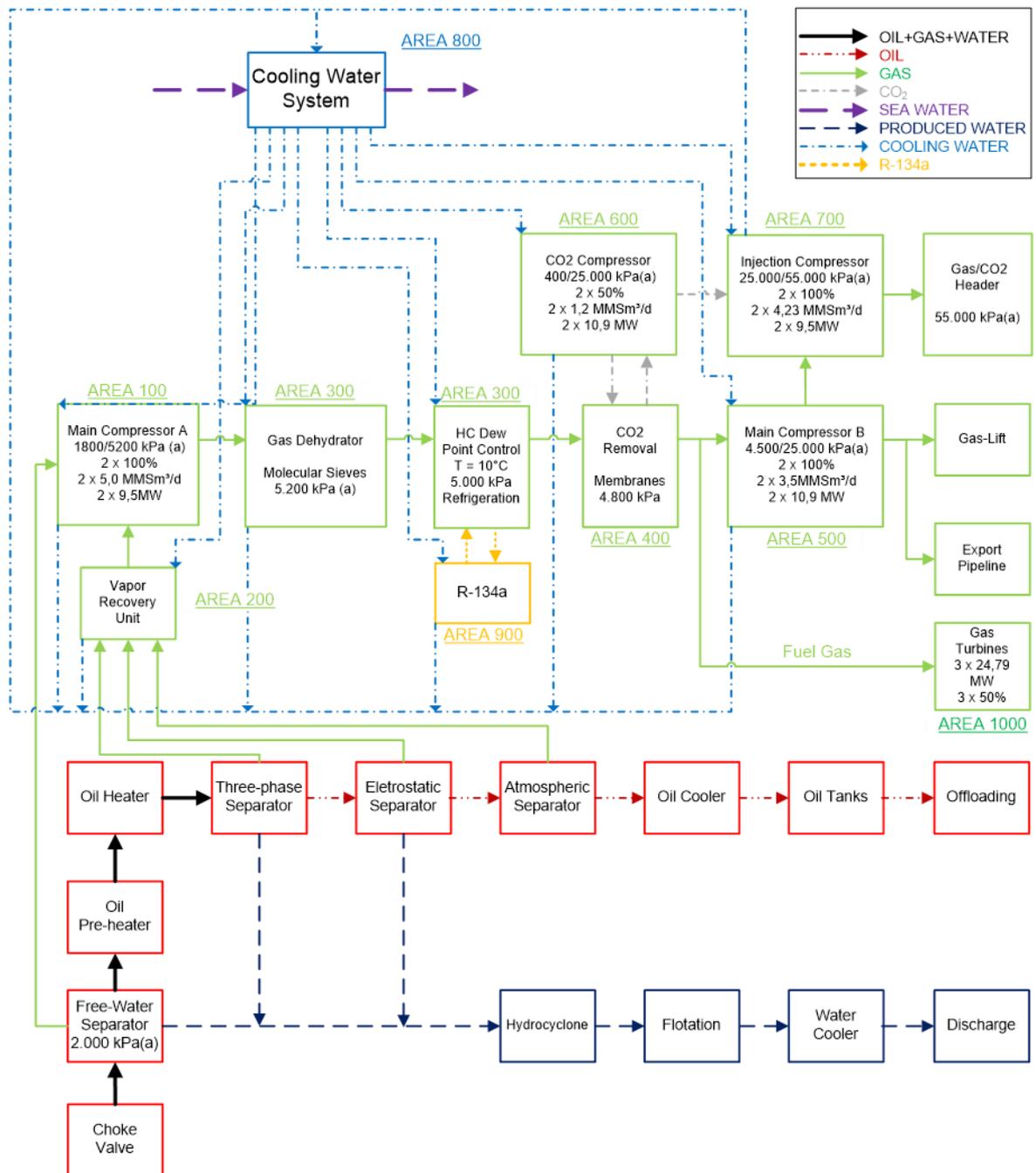
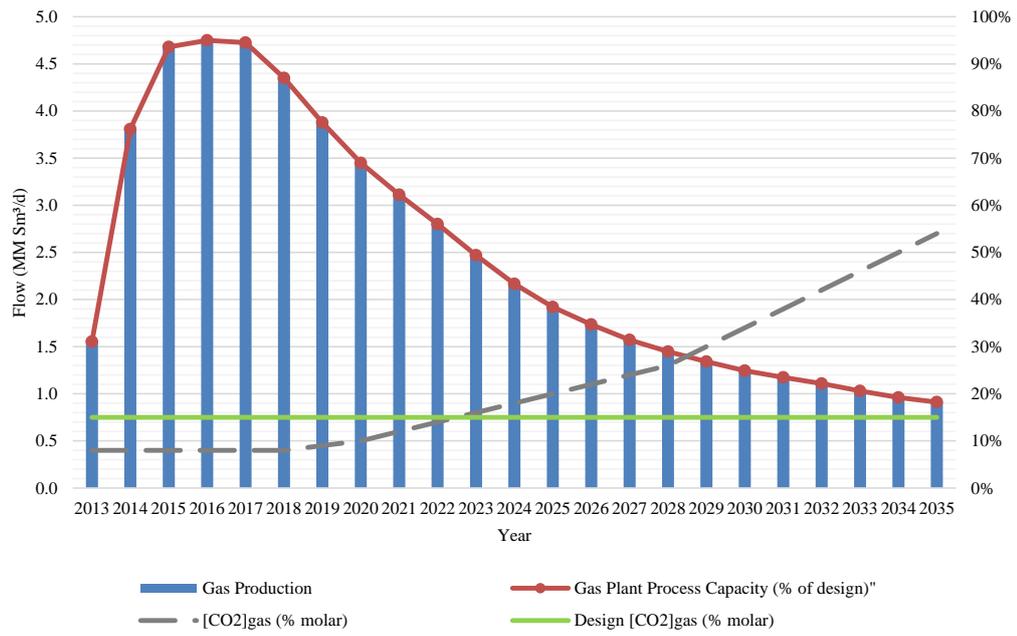


Figure 3A.1. Block diagram of topside plant of FPSO Cidade de Paraty



**Figure 3A.2. Forecasted gas production**  
 (%CO<sub>2</sub> curve estimated with only initial 8%mol and final 55%mol from EIA)



## 4. EXERGY, ENERGY AND EMISSIONS ANALYSIS OF COMPRESSORS SCHEMES IN OFFSHORE RIGS: CO<sub>2</sub>-RICH NATURAL GAS PROCESSING

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### **Abstract**

Deepwater oil and associated gas productions resort to floating rigs operating at continuously decreasing gas-loads during the last three quarters of the field campaign. As centrifugal compressors are sized at maximum loads, anti-surge recycles are used making operation inefficient in terms of power consumption and emissions per oil barrel produced. Smaller paralleled compressors and variable-speed drivers are investigated at peak and partial gas-loads and compared to traditional anti-surge recycle designs in terms of exergy efficiency, investment, footprint and emissions. Oversized compressors with anti-surge recycles result in almost constant power consumption along process lifespan, regardless the gas-load, increasing fuel and CO<sub>2</sub> intensities as gas-load decreases and attaining exergy efficiencies of 49% and 83% at 25% and 100% gas-loads, respectively. On the other hand, with variable-speed drivers and smaller paralleled compressors, power consumption becomes proportional to gas-load with exergy efficiencies always between 80% and 88%, and attaining 11% and 39% less power consumptions at 100% and 25% gas-loads. Moreover, CO<sub>2</sub> intensity and investment are, respectively 34% and 3% less than in traditional layouts with oversized compressors. These savings resulted from eliminating a gas turbine thanks to lower power demand when no anti-surge recycles are used.

**Keywords:** Offshore gas processing; CO<sub>2</sub>-rich natural gas; Centrifugal compressors; Anti-surge recycle; Exergy analysis; CO<sub>2</sub> emissions.

Supplementary Materials for this chapter are found in the Appendix H, Section B.

## Abbreviations

BOE Equivalent Oil Barrel; bbl/d Barrels per Day; CW Cooling-Water; EOR Enhanced Oil Recovery; FPSO Floating Production, Storage & Offloading; GT Gas-Turbine; HCDPA Hydrocarbon Dew-Point Adjustment; MMSm<sup>3</sup>/d Million Standard m<sup>3</sup> per Day; MP Membrane-Permeation; MPSC Multiple-Paralleled Smaller Compressors; MMUSD Million US Dollars; NG Natural Gas; PHW Pressurized-Hot-Water; PR-EOS Peng-Robinson Equation-of-State; RER Reference Environmental Reservoir; RPM Revolutions per Minute; SW Seawater; SSLC Single-Shaft Larger Compressors; WDPA Water Dew-Point Adjustment; VRU Vapor-Recovery Unit; VSD Variable-Speed Driver.

## Nomenclature

$\dot{B}$	<i>Exergy flow rate (kW)</i>
$CO_{2eq}$	<i>CO<sub>2</sub> equivalent</i>
$F_j$	<i>j<sup>th</sup> feed flow rate (kmol/s)</i>
$FCI$	<i>Fixed Capital Investment (MMUSD)</i>
$\bar{H}$	<i>Molar enthalpy (kJ/kmol)</i>
$K_j$	<i>j<sup>th</sup> product flow rate (kmol/s)</i>
$LMTD$	<i>Log-Mean temperature difference (°C)</i>
$nc, nfs, nps$	<i>Numbers of components/feeds/products</i>
$nwi, nwe$	<i>Numbers of imported/exported powers</i>
$P, Q$	<i>Pressure (bar, kPa), Heat duty (kW)</i>
$R$	<i>Ideal gas constant (<math>R=8.314*10^{-5}bar.m^3/mol.K</math>)</i>
$\bar{S}$	<i>Molar entropy (kJ/kmol.K)</i>
$T, \bar{V}$	<i>Temperature (K), Molar volume (m<sup>3</sup>/kmol)</i>
$\dot{W}$	<i>Shaft-Power (MW)</i>
$Y, Z$	<i>Molar fraction, Compressibility factor</i>

## Greek Symbols

$\mu_k$	<i>Chemical potential of k<sup>th</sup> species</i>
$\eta, \eta_P$	<i>Exergy and polytropic efficiencies</i>
$\dot{\Omega}_S$	<i>Entropy creation rate (kW/K)</i>

## Superscripts

<i>exported, imported</i>	<i>Exported, Imported</i>
$W$	<i>Mechanical power</i>

## Subscripts

<i>Burn, Emitted</i>	<i>Combustion, Emitted</i>
<i>in, in.total</i>	<i>Inlet, Total Inlet</i>
<i>inj</i>	<i>Full-Injection</i>
<i>out, out,total</i>	<i>Outlet, Total Outlet</i>
<i>Produced, Exp</i>	<i>Produced, Exported</i>

## 4.1. Introduction

According to the International Energy Agency (IEA, 2018), the global energy demand will increase by ~25% until 2040, making oil and natural gas (NG) responsible for ~50% of the global energy consumption in the same period. Additionally, NG partially replaces coal in power generation until 2030, becoming the second source in the global energy matrix (Copenhagen Economics, 2017). In 2018 oil and NG were responsible for 5.2 gigatonnes of CO<sub>2</sub> equivalent (CO<sub>2eq</sub>) liberated into the atmosphere, where ~2/3 of such emissions resulted from in-place power/heat productions (IOGP, 2016). Meanwhile, the Paris Agreement forced the oil-and-gas industry to limit operational emissions, resulting that major oil companies worldwide are now considering climate-change policies on planning and investment decisions; i.e., environmental and economic objectives must be balanced because regulatory policies, such as carbon-taxation and efficiency standards, can affect supply/demand and prices of fossil fuels (Chevron, 2018).

Given the growing oil-gas demands, deepwater oil-gas fields attracted investments worldwide in which Floating Production, Storage and Offloading (FPSO) vessels are the preferred production concept for large-scale projects at remote offshore locations without pipeline systems, primarily due to lower installation-decommissioning costs, flexibility and large storage capacity (Araújo et al., 2017). Currently, 186 FPSOs operate worldwide, with 24 FPSO awards expected by 2020, 8 of them in Pre-Salt Basin, Brazil (Rystad Energy, 2019).

A typical FPSO operates with low-efficiency power-producing, power-consuming and heat-producing systems. On one hand, gas-turbines (GT) are the main power-producing systems and operate without heat-recovery steam-generators; i.e., only waste-heat recovery units transfer some heat from flue-gas to pressurized-hot-water (PHW) and combined-cycles are vetoed due to footprint/weight/safety restrictions. On the other hand, major power-consuming systems are single-shaft large centrifugal compressors (SSLC) designed for high gas-loads, which waste power in anti-surge gas recycles at low gas-loads as shown in Cruz et al. (2018) who investigated benefits of deep-seawater to cool cooling-water (CW) for SSLC trains that respond for 52% of the power demand of a Pre-Salt FPSO processing CO<sub>2</sub>-rich NG and dispatching CO<sub>2</sub> to Enhanced Oil Recovery (EOR).

New oil-and-gas sector regulations have demanded technology improvements to mitigate environmental impacts of offshore units. Rational and optimized energy utilization leads to less

power/emission intensive FPSOs. Pierobon et al. (2014) investigated waste-heat recovery in offshore rigs. Allahyarzadeh-Bidgoli et al. (2018) performed energy optimization of Pre-Salt FPSOs. Barrera, Bazzo and Kami (2015), Reis and Gallo (2018) and Veloso et al. (2018) studied optimization of FPSO power production with organic Rankine-Cycles, while Roussanaly et al. (2018) techno-economically analyzed offshore power generation with carbon capture and storage.

#### 4.1.1. Exergy Analysis of Offshore Rigs

Nguyen et al. (2013) performed exergy analyses of North Sea oil-gas rigs, while Voldsund et al. (2014) investigated the respective exergy destructions. Nguyen et al. (2014) exergetically analyzed upstream oil plants on mature fields and Gallo et al. (2017) performed energy/exergy analyses of Pre-Salt FPSOs, concluding that SSLCs stand as major power sinks whose anti-surge recycles increasingly devour exergy as gas-load decreases.

The exergy concept for open systems combines the 1<sup>st</sup> and 2<sup>nd</sup> Laws of Thermodynamics. The exergy flow rate of a stream is the maximum obtainable power when reaching equilibrium with a reference environment reservoir (RER) (Teixeira et al., 2016). Exergy analysis assesses exergy flows, discriminating wasted exergy to the environment and exergy destruction due to irreversibilities. Both can be minimized, though the latter invoke different approaches for reduction. Exergy destruction is a reflex of systemic irreversibilities requiring interventions for better exergy efficiency (Soundararajan et al., 2014). However, system re-design is problematic. FPSO design is constrained by equipment technology (e.g., efficiencies), field characteristics and export conditions (oil/gas temperature, pressure and specifications). Additionally, economic and operational risks must be considered in new designs. There are works on exergy-based design optimization and on minimizing energy degradation, normally disregarding economy implications. Panton et al. (2014) compared conventional and electrified oil-gas platforms via exergy analysis, while da Silva and de Oliveira (2018) determined exergy cost of oil-gas offshore production and Silva et al. (2019) exergetically analyzed gas-turbines on offshore rigs.

As seen above, many works have applied exergy analysis to offshore rigs and FPSOs, but none has used exergy assessments to compare SSLC with new compressor schemes.

### 4.1.2. The Present Work

Due to rough working conditions and long distance to coast, deep-water offshore oil production with associated gas has a past of extreme exergy inefficiency. About 25 years ago it was a common practice of some oil companies to simply burn all produced gas – exceeding power/heating/gas-lift utilizations – through bizarrely giant flares requiring huge radiation shields for crew protection. With the advent of climate-change such absurd practices were banned, and a growing interest in better FPSO exergy efficiency appeared in the literature. But even nowadays oil is still the supreme goal in deep-water enterprises and gas is, not rarely, considered worthless in the sense that power-consuming systems are designed for less efficient operation privileging low investment; e.g., SSLC schemes with inefficient anti-surge recycles at low gas-load. Gallo et al. (2017) suggested replacing SSLC by multiple-paralleled smaller compressors (MPSC), which are gradually turned-off as gas production falls, but they did not develop MPSC. Replacing SSLC by MPSC to rule out anti-surge recycles can increase FPSO exergy efficiency at the expense of rising fixed capital investment (FCI); hence a trade-off appears between FCI and exergy efficiency, requiring further examination. This study fills this gap comparing SSLC and MPSC in offshore oil-gas rigs on following grounds: (i) exergy efficiency; (ii) power demand, fuel consumption and CO<sub>2</sub> emissions per equivalent oil barrel (BOE) produced; and (iii) FCI/footprint. Calculations adopted rigorous thermodynamic simulation of processes, exergy analysis with two RER definitions and 2<sup>nd</sup> Law consistency checks.

## 4.2. Methods

For deep-water oil-gas FPSOs handling CO<sub>2</sub>-rich NG, MPSC is proved to have benefits over SSLC. To do this, SSLC and MPSC comparisons on exergy conservation, power consumption and capital/footprint grounds were conducted at three FPSO gas-loads (~100%, ~50%, ~25%) representing phases of typical Pre-Salt FPSO campaigns. Two RER approaches (RER-1, RER-2) were considered for exergy analysis, entailing 6 process simulations and 12 exergy analyses performed. FCI/footprint analyses contemplate only 100% gas-load, upon which MPSC and SSLC gas-plants were sized; i.e., the design condition. Fig. 4.1 is a flowchart of steps in this study: (i) red-hexagon consolidates literature information and data from environmental impact assessments of the real FPSO (Cruz et al., 2018) selected for this study; (ii) blue-boxes

represent computational tasks for process simulation, exergy analysis, equipment sizing and FCI/footprint estimation; and (iii) green-flags represent task inputs and/or task results.

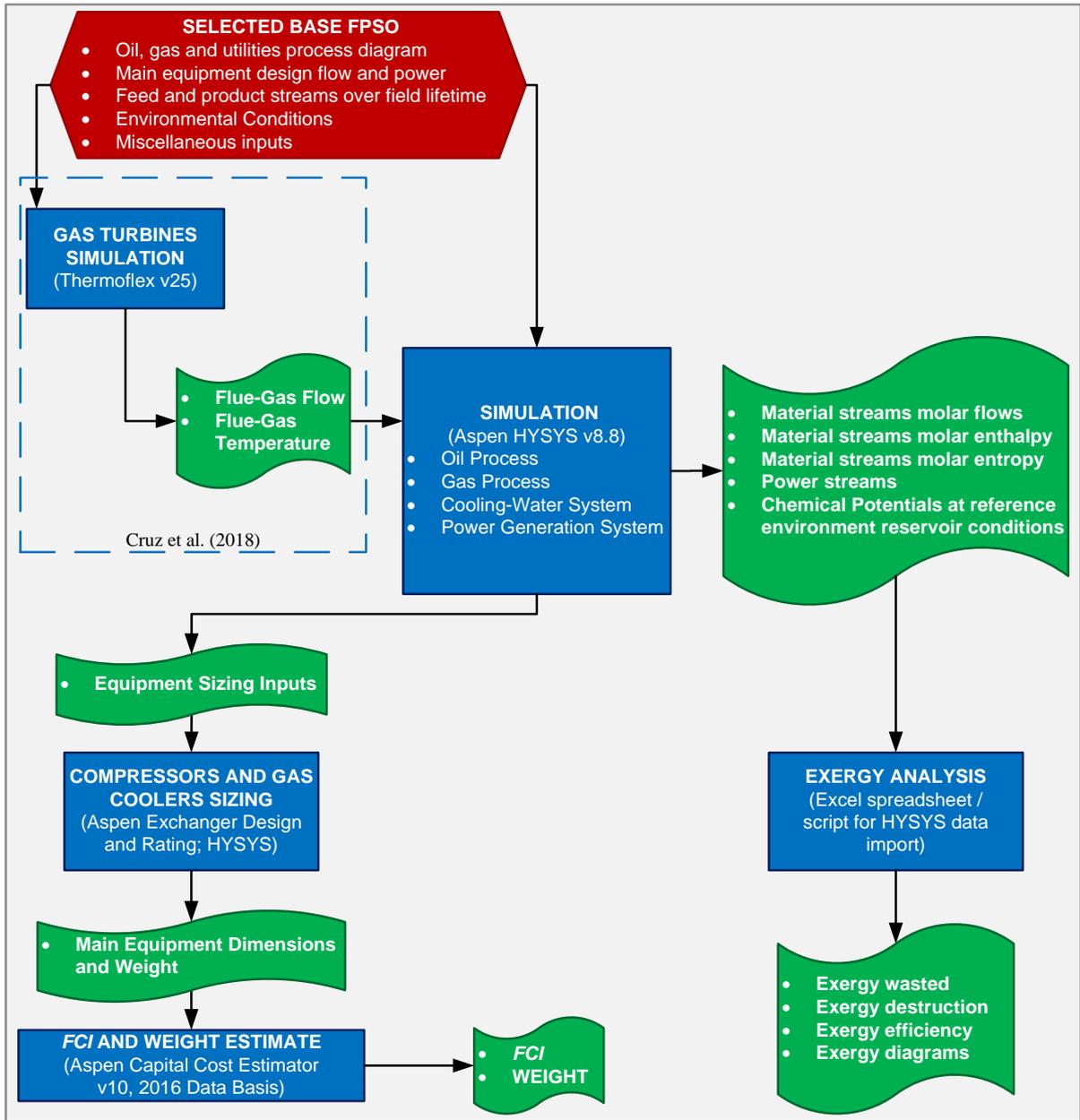
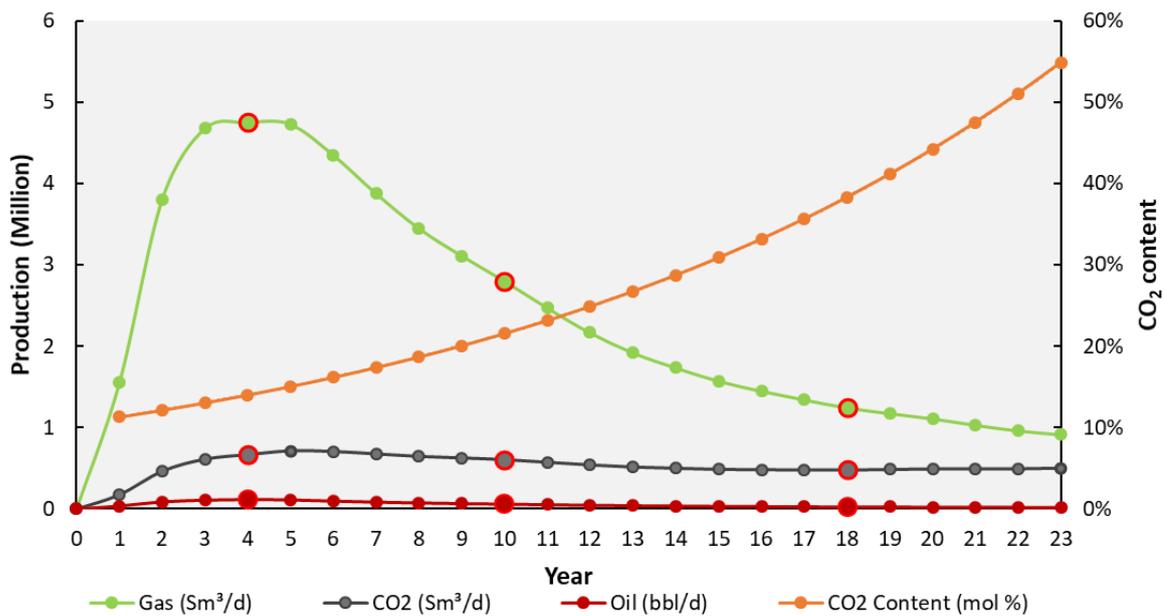


Figure 4.1. Study Flowchart.

#### 4.2.1. Process Simulation

The chosen FPSO (Cruz et al., 2018) operates on Pre-Salt (Santos) Basin with gas and oil respective capacities of 5 MMSm<sup>3</sup>/d and 100,000 bbl/d. On Pre-Salt fields, CO<sub>2</sub> separation, EOR injection and NG exportation require high pressures, entailing highly power-intensive gas processing due to massive use of SSLC centrifugal compressors. FPSO and oil-gas field data

come from Santos Basin Environmental Impacts Assessment (Petrobras, 2013) as the profiles (Fig. 4.2) of oil/gas/CO<sub>2</sub> productions and %mol CO<sub>2</sub> of raw NG along field lifetime, where the large circles represent three phases of field campaign: maximum (year 4), medium (year 10) and minimum (year 18) gas-loads at which the FPSO respectively operates at ~100%/~50%/~25% of design gas capacity. Steady-state FPSO configurations, SSLC-Case and MPSC-Case, were simulated at ~100%/~50%/~25% gas-loads. Fig. 4.3 depicts SSLC-Case flowsheet with 319 material-streams, 40 power-streams and 197 units. The legends auto-explain Fig. 4.3, where red-box and green-box respectively envelope oil-plant and gas-plant; blue-box encloses CW system; grey-box envelopes power generation gas-turbines; and black-box represents FPSO topside with streams 1-10 as feeds/products. All envelopes were simulated, except the production-water plant (Fig. 4.3, bottom) because it is not affected by gas-plants nor generate feeds to them. The oil-plant is absent in exergy/economic analyses since it is not affected by SSLC/MPSC gas-plants.



**Figure 4.2. Oil/gas/CO<sub>2</sub> productions along field lifetime. (Petrobras, 2013).**

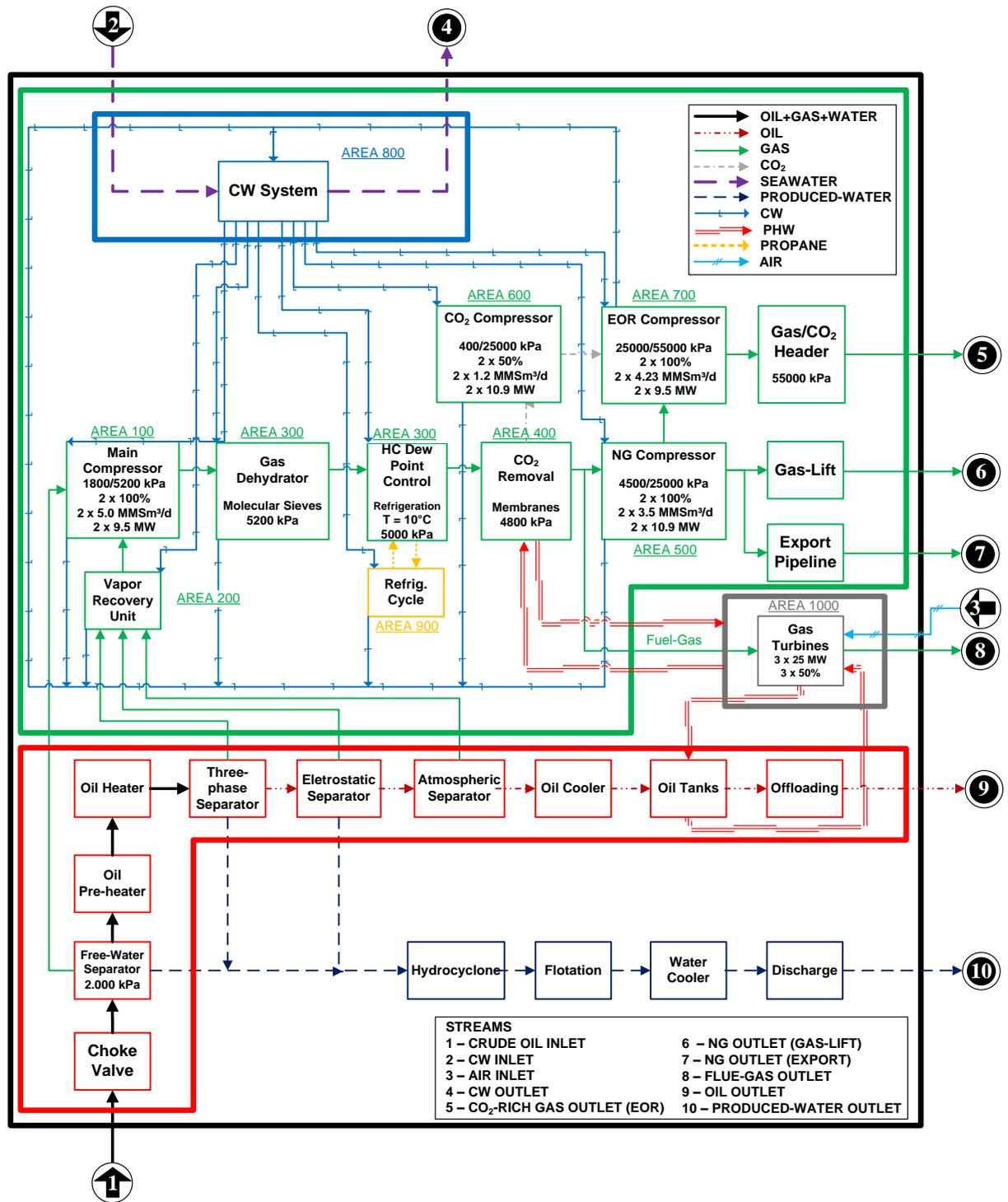


Figure 4.3. SSLC-Case: oil-plant (red), gas-plant (green) and water-plant (bottom-right).

#### 4.2.1.1. Oil-Plant

Fig. 4.4 depicts the oil-plant flowsheet with oil and produced-water inputs in Table 4.1 at ~100%/~50%/~25% gas-loads. Oil-plant feeds the gas-plant with raw gas streams that vary according to gas-load (Table 4.2) and is the same for SSLC-Case and MPSC-Case.

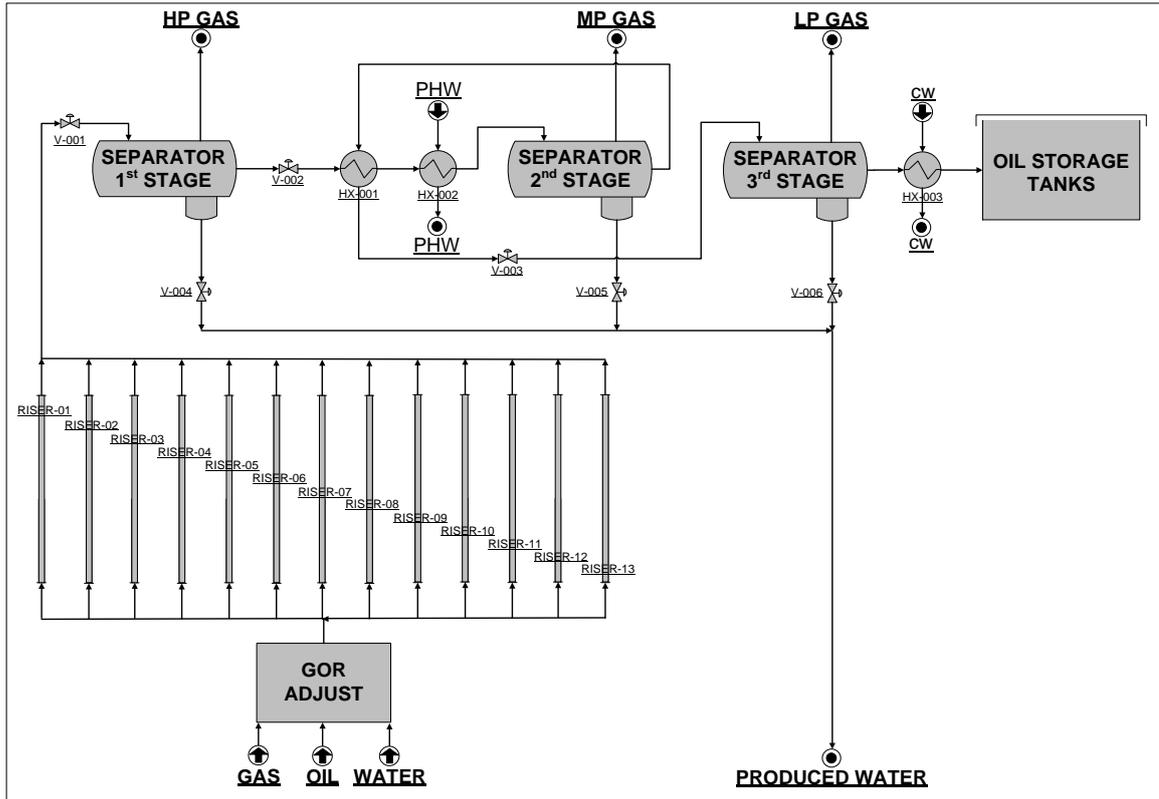


Figure 4.4. Oil-Plant (HP=high-pressure, MP=medium-pressure, LP=low-pressure)

Table 4.1. Oil-Plant: simulation inputs.

<i>FPSO Gas-Load</i>	<i>100%</i>	<i>50%</i>	<i>25%</i>
<i>Operational Year</i>	<i>4</i>	<i>10</i>	<i>18</i>
<i>Oil (bbl/d)</i>	<i>111467</i>	<i>56771</i>	<i>24037</i>
<i>Production-Water (bbl/d)</i>	<i>2907</i>	<i>41748</i>	<i>79153</i>
<i>Raw NG (Sm<sup>3</sup>/d)</i>	<i>4750184</i>	<i>2801137</i>	<i>1244432</i>
<i>CO<sub>2</sub> (Sm<sup>3</sup>/d)</i>	<i>664971</i>	<i>603644</i>	<i>476677</i>
<i>Gas-Oil Separators</i>	<i>3</i>	<i>3</i>	<i>3</i>
<i>Gas-Oil Ratio (Sm<sup>3</sup>/m<sup>3</sup>)</i>	<i>257</i>	<i>294</i>	<i>303</i>
<i>Water-Oil Ratio (m<sup>3</sup>/m<sup>3</sup>)</i>	<i>0.025</i>	<i>0.691</i>	<i>3.09</i>
<i>Reservoir Temperature (°C)</i>	<i>40</i>	<i>40</i>	<i>40</i>
<i>Reservoir Pressure (kPa)</i>	<i>41368</i>	<i>41368</i>	<i>41368</i>

**Table 4.2. Oil-Plant: raw NG streams.**

<i>FPSO</i>	<i>100%</i>			<i>50%</i>			<i>25%</i>		
<i>Gas-Load</i>									
<i>Gas-Oil Separator</i>	<i>1<sup>st</sup></i>	<i>2<sup>nd</sup></i>	<i>3<sup>rd</sup></i>	<i>1<sup>st</sup></i>	<i>2<sup>nd</sup></i>	<i>3<sup>rd</sup></i>	<i>1<sup>st</sup></i>	<i>2<sup>nd</sup></i>	<i>3<sup>rd</sup></i>
<i>Gas Stream</i>	101	208	201	270	320	370	420	470	520
<i>MMSm<sup>3</sup>/d</i>	4.44	0.25	0.06	2.65	0.12	0.03	1.19	0.05	0.01
<i>T(°C)</i>	20	60	44	29	60	48	36	60	44
<i>P(kPa)</i>	1850	655	250	1850	655	250	1850	655	250
<i>Mol Fractions</i>									
<i>CO<sub>2</sub></i>	0.1410	0.1774	0.2471	0.2126	0.2451	0.3417	0.3610	0.4114	0.5111
<i>H<sub>2</sub></i>	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
<i>N<sub>2</sub></i>	0.0057	0.0003	0.0011	0.0051	0.0003	0.0010	0.0041	0.0003	0.0007
<i>Methane</i>	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
<i>Ethane</i>	0.6993	0.1839	0.1705	0.6309	0.1585	0.1519	0.5053	0.1379	0.1133
<i>Propane</i>	0.0898	0.1849	0.1426	0.0834	0.1576	0.1198	0.0678	0.1230	0.0850
<i>i-Butane</i>	0.0444	0.2408	0.2056	0.0442	0.2152	0.1722	0.0374	0.1571	0.1212
<i>n-Butane</i>	0.0050	0.0411	0.0436	0.0055	0.0395	0.0380	0.0049	0.0285	0.0274
<i>i-Pentane</i>	0.0092	0.0830	0.0879	0.0103	0.0819	0.0780	0.0096	0.0596	0.0572
<i>n-Pentane</i>	0.0014	0.0155	0.0209	0.0018	0.0165	0.0195	0.0018	0.0125	0.0153
<i>n-Hexane</i>	0.0017	0.0200	0.0275	0.0022	0.0216	0.0259	0.0023	0.0166	0.0208
<i>n-Heptane</i>	0.0008	0.0103	0.0185	0.0011	0.0119	0.0180	0.0013	0.0097	0.0157
<i>n-Octane</i>	0.0003	0.0041	0.0093	0.0004	0.0048	0.0091	0.0006	0.0042	0.0087
<i>n-Nonane</i>	0.0001	0.0020	0.0059	0.0002	0.0025	0.0058	0.0003	0.0021	0.0055
<i>n-Decane</i>	0.0000	0.0006	0.0021	0.0001	0.0007	0.0021	0.0001	0.0006	0.0020
<i>n-C11</i>	0.0000	0.0002	0.0008	0.0000	0.0002	0.0008	0.0000	0.0002	0.0008
<i>n-C12</i>	0.0000	0.0000	0.0002	0.0000	0.0001	0.0002	0.0000	0.0000	0.0002
<i>H<sub>2</sub>O</i>	0.0000	0.0000	0.0002	0.0000	0.0000	0.0002	0.0000	0.0000	0.0002
<i>Ar</i>	0.0014	0.0357	0.0161	0.0022	0.0435	0.0157	0.0033	0.0362	0.0148

#### 4.2.1.2. Gas-Plant, Cooling-Water System and Power Generation

Gas-plant, GT power generation and CW system of SSLC-Case and MPSC-Case were simulated in HYSYS using Peng-Robinson Equation-of-State (PR-EOS) with Free-Water for thermodynamic modeling of oil/gas/CO<sub>2</sub>/water process streams and HYSYS-Steam-Table for CW, PHW and seawater (SW) streams. Air temperature (23°C), relative humidity (87%) and SW temperature (23°C near sea surface) represent average values at FPSO location (Petrobras, 2013), which are necessary for GT simulations (validated in Cruz et al., 2018) and CW calculations. Fig. 4.5 depicts SSLC-Case flowsheet comprising gas-plant, CW system and GT area, where VRU, MP, WDPA, HCDPA, C3, C3+, C6+ respectively stand for Vapor-Recovery Unit, Membrane-Permeation, Water Dew-Point Adjustment, Hydrocarbon Dew-Point Adjustment, propane, propane-and-heavier-alkanes, and hexane-and-heavier-alkanes. HCDPA is set at 3°C to guarantee C6+ below 0.1%mol, avoiding MP condensation issues. MP CO<sub>2</sub> separation was simulated using HYSYS MP Extension (Arinelli et al., 2019).

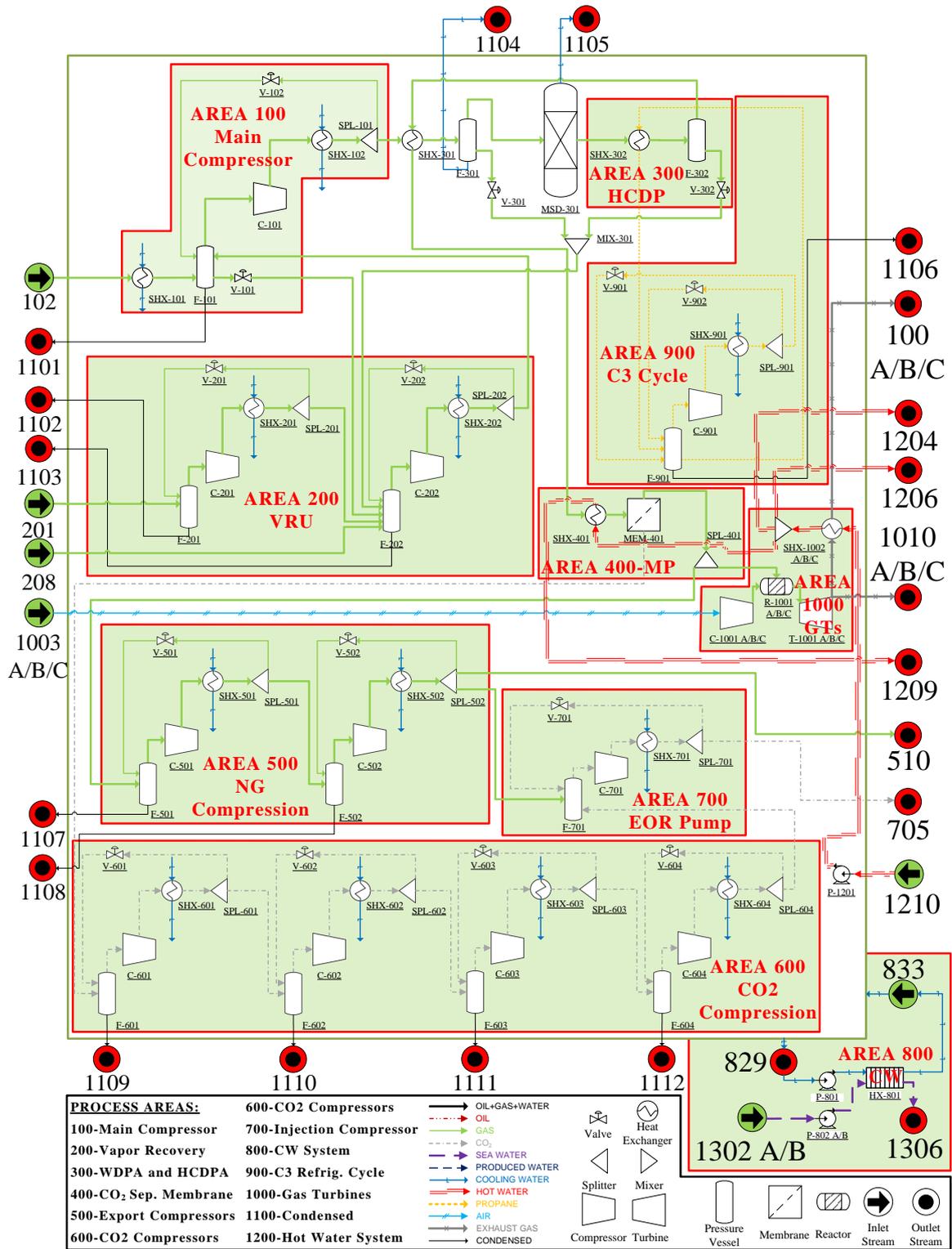


Figure 4.5. SSLC-Case: gas-plant, CW system and GT area.

#### 4.2.1.3. MPSC-Case Simulation

MPSC-Case is modeled as the SSLC-Case (Figs. 4.3-4.5) just replacing some SSLC compressors with anti-surge recycles by sets of MPSCs deprived of anti-surge control. This

eliminates exergy destruction by anti-surge recycles in MPSC-Case. The number of equal MPSCs replacing a given large SSLC in MPSC-Case is defined using the minimum achievable flow rate via variable-speed drivers (VSD) which decrease RPM of SSLC to avoid surge at low inlet flow rates. VSD can also move the operating point to the optimal efficiency region via RPM manipulation, regardless inlet flow variations (Albusaidi and Pilidis, 2015). Since VSD flows are limited to 50%–60% of design flows, 55% of design inlet flow is assumed as VSD lower limit. Therefore, if a SSLC experiences less than 55% flow rate at ~25% FPSO gas-load, it is replaced in MPSC-Case by a set of MPSCs each one designed somewhat above such minimum SSLC flow and totaling little above the integral SSLC flow at 100% FPSO gas-load. Thus, at 100% FPSO gas-load all MPSCs are active and are turned off, one-at-a-time, as FPSO gas-load falls. Since the polytropic efficiency of centrifugal compressors varies with flow rate, MPSC efficiencies were estimated via a correlation of Cruz et al. (2018).

However, the minimum VSD flow is limited to 50%–60% of the compressor design flow. In this work, 55% of the design inlet volumetric flow is adopted as limit. Below this limit, the inlet flow is divided and smaller paralleled compressors are considered. When the gas-load becomes lesser than 55% of design capacity one of the smaller compressors is turned off and the operation proceeds with the other one. Table 4.3 shows the compressors scheme for SSLC-Case and MPSC-Case. The polytropic efficiency of centrifugal compressors varies with inlet flow rate. This effect is considered in this study, as observed in Table 4.3. The polytropic efficiency is estimated using a correlation of Cruz et al. (2018).

#### 4.2.2. Exergy Analysis of Processes

Exergy analysis follows Teixeira et al. (2016), with main steps described in the following.

##### 4.2.2.1. Process division into subsystems

Fig. 4.5 shows such division for SSLC-Case, which is the same for MPSC-Case. Envelopes for exergy analyses of subsystems expose feed/product material-streams, CW/PHW/SW streams and power-streams.

#### 4.2.2.2. Energy and mass balances

Energy-Mass balances are solved through process simulation (Sec. 4.2.1) obtaining process parameters, material-streams properties and power-streams for determination of inlet/outlet exergy flows.

#### 4.2.2.3. RER Configuration

RER is an infinite ground-level equilibrium system constituted of process species at temperature  $T_0$ , pressure  $P_0$  and  $i^{th}$  chemical potential  $\mu_i^0$  ( $i=1..nc$ ). Exergy flow rates (kW) of material-streams are relative to the chosen RER; i.e., exergy flows depend on the associated material-streams and also on RER definition, while for power-streams the exergy flows are the respective power values. Adequate RER definition is crucial for a useful exergy analysis (Dinçer and Rosen, 2013); i.e., inappropriate RER choices would entail enormous values of inlet/outlet exergy flows of material-streams, making exergy destruction rates – which do not depend on RER choice – relatively insignificant. In such cases, exergy efficiencies would have artificially high values close to 100%, making the assessment useless.

Two RER definitions are adopted: RER-1 and RER-2. Both initially consist of standard dry air ( $N_2=78.08\%mol$ ,  $O_2=20.95\%mol$ ,  $Ar=0.93\%mol$  and  $CO_2=0.04\%mol$ ) at  $T_0=298.15$  K,  $P_0=1.013$  bar, which was put in equilibrium with an infinite body of liquid water becoming water-saturated at  $T_0=298.15$  K,  $P_0=1.013$  bar. The existence of liquid water in RER is convenient to lower exergy flows of numerous CW/PHW/SW streams. At this point, the calculation of  $\mu_i^0$  for  $i \in \{N_2, O_2, Ar, CO_2, H_2O\}$  is immediate as an atmospheric ideal gas mixture.

The differences between RER-1 and RER-2 emerge from the different  $\mu_i^0$  states chosen for the hydrocarbons of raw NG. RER-1 is adequate for combustion or chemically reactive processes (e.g., GTs), while RER-2 is appropriate for non-reactive processes (e.g., compressors, valves, exchangers and physical separation operations). In RER-1, the  $\mu_i^0$  of hydrocarbons is obtained from  $\mu_i^0$   $i \in \{N_2, O_2, Ar, CO_2, H_2O\}$  via chemically equilibrated gas-phase combustion reactions (e.g.,  $\mu_{CH_4}^0 = \mu_{CO_2}^0 + 2\mu_{H_2O}^0 - 2\mu_{O_2}^0$ ,  $\mu_{C_2H_6}^0 = 2\mu_{CO_2}^0 + 3\mu_{H_2O}^0 - 3.5\mu_{O_2}^0$ , etc), entailing very low  $\mu_i^0$  for hydrocarbons and giving high exergy flows for NG streams. On the other hand, in RER-

2 hydrocarbons are not chemically equilibrated with air species; i.e., they are physically added to the water-saturated air at low contents compatible with typical atmospheres surrounding oil-gas facilities (e.g., *44.1 ppm-mol* for CH<sub>4</sub>, etc) based on microseepage studies (Chang et al., 2014). The gas phase composition of RER-2 is updated due to the small hydrocarbon contents, but the ideal gas behavior is still valid. RER-2 is critical for exergy analysis of non-reactive processes that do not change the number of molecules; i.e., in RER-2 the  $\mu_i^0$  of hydrocarbons are not too low. This reduces the “chemical part” of exergy flows of feed/product streams of physical operations, enhancing the visibility of their exergy destructions and producing meaningful exergy efficiencies. Regardless of RER-1 or RER-2 choices, the rate of exergy destruction is always the same for a given steady-state operation, but the exergy efficiency is calculated with inlet exergy flows and, as such, is very dependent of RER choice.

$\mu_i^0$  for  $i \in \{N_2, O_2, Ar, CO_2, H_2O\}$  are calculated respecting HYSYS enthalpy/entropy reference-states for the chosen thermodynamic packages. PR-EOS for process gas/oil/water streams and HYSYS-Steam-Table for CW/PHW/SW streams have the same water reference-states; i.e., have datum compatibility. Since HYSYS does not export chemical potentials,  $\mu_i^0$  for  $i \in \{N_2, O_2, Ar, CO_2, H_2O\}$  is calculated with ideal gas formula in the left term of Eq. (1), where  $P^*$  is a sufficiently low pressure for pure  $i$  ideal gas at  $T_0$ . For light gases  $\{N_2, O_2, Ar, CO_2\}$   $P^*=P_0$  is chosen, while for water  $P^*=0.01$  bar is adequate.  $\mu_i^{Pure}(T_0, P^*)$  is obtained for a pure  $i$  stream at  $(T_0, P^*)$  via the right term in Eq. (4.1) with PR-EOS. For trace hydrocarbon  $i$  in RER-2, Eq. (1) is used with pure  $i$  at  $P^*=0.01$  bar. Table 4.3 presents molar fractions ( $y_i^0$ ) and  $\mu_i^0$  for NG species in RER-1 and RER-2.

$$\mu_i^0 = \mu_i^{Pure}(T_0, P^*) + RT_0 \cdot \ln\left(\frac{P_0 y_i^0}{P^*}\right), \mu_i^{Pure}(T_0, P^*) = \bar{H}_i^{Pure}(T_0, P^*) - T_0 \cdot \bar{S}_i^{Pure}(T_0, P^*) \quad (4.1)$$

**Table 4.3. RER-1 and RER-2: gas molar fractions ( $y_i^0$ ) and chemical potentials ( $\mu_i^0$ ) for NG operations.**

	<i>RER-1</i>		<i>RER-2</i>	
<i>T(K)</i>	$T_0=298.15$		$T_0=298.15$	
<i>P(bar)</i>	$P_0=1.0133$		$P_0=1.0133$	
<i>Species</i>	$y_i^0$	$\mu_i^0$ (kJ/mol)	$y_i^0$	$\mu_i^0$ (kJ/mol)
$N_2$	0.7593	-47.4	0.7593	-47.4
$O_2$	0.2037	-52.5	0.2037	-52.5
$H_2O$	0.0276	-301.3	0.0276	-301.3
<i>Argon</i>	0.0090	-47.1	0.0090	-47.1
$CO_2$	0.0004	-464.2	0.0004	-464.2
$CH_4$	N/A	-961.7	4.41E-05	-154
$C_2H_6$	N/A	-1648	4.12E-11	-201
$C_3H_8$	N/A	-2335	2.31E-11	-212
<i>i-C<sub>4</sub>H<sub>10</sub></i>	N/A	-3022	4.48E-12	-247
$C_4H_{10}$	N/A	-3022	1.17E-11	-228
<i>i-C<sub>5</sub>H<sub>12</sub></i>	N/A	-3709	6.33E-12	-237
$C_5H_{12}$	N/A	-3709	6.58E-12	-242
$C_6H_{14}$	N/A	-4395	3.29E-12	-268
$C_7H_{16}$	N/A	-5082	1.65E-12	-303
$C_8H_{18}$	N/A	-5769	8.23E-13	-313
$C_9H_{20}$	N/A	-6455	4.11E-13	-379
$C_{10}H_{22}$	N/A	-7142	2.06E-13	-416
$C_{11}H_{24}$	N/A	-7829	1.03E-13	-452
$C_{12}H_{26}$	N/A	-8516	5.14E-14	-487

#### 4.2.2.4. Exergy Flow of Inlet/Outlet Energy/Material Streams

Eqs. (4.2a)-(4.2b) calculate inlet/outlet exergy flows of a system regarding the chosen RER (Teixeira et al., 2016), where  $\dot{B}_{in}$ ,  $\dot{B}_{out}$  are inlet/outlet exergy flow rates (kW) of material-streams, while  $\dot{B}_{in}^W$ ,  $\dot{B}_{out}^W$  respectively comprehend sums of positive power-streams  $\dot{W}_j^{imported}$ ,  $\dot{W}_j^{exported}$  on the right-hand side of Eqs. (4.2a)-(4.2b) that are imported/exported by the system (e.g., electricity to compressors/pumps drivers). All terms of Eqs. (4.2a)-(4.2b) were extracted from HYSYS simulations and exported to spreadsheets via an automatized procedure to avoid mistakes.

$$\dot{B}_{in,total} = \dot{B}_{in} + \dot{B}_{in}^W = \sum_{j=1}^{nfs} F_j \left( \bar{H}_{Fj} + P_0 \bar{V}_{Fj} - T_0 \bar{S}_{Fj} - \sum_{k=1}^{nc} \mu_k^0 y_{k,Fj} \right) + \sum_{j=1}^{nwi} |\dot{W}_j^{imported}| \quad (4.2a)$$

$$\dot{B}_{out,total} = \dot{B}_{out} + \dot{B}_{out}^W = \sum_{j=1}^{nps} K_j \left( \bar{H}_{Kj} + P_0 \bar{V}_{Kj} - T_0 \bar{S}_{Kj} - \sum_{k=1}^{nc} \mu_k^0 y_{k,Kj} \right) + \sum_{j=1}^{nwe} |\dot{W}_j^{exported}| \quad (4.2b)$$

#### 4.2.2.5. Exergy Balances and Exergy Destructions

Eq. (4.3) calculates the exergy destruction rate (kW) of a system ( $\Delta \dot{B}$ ) by subtracting Eqs (4.2a) and (4.2b). It can be proved that  $\Delta \dot{B}$  is also  $\dot{W}^{LOST}$ , the rate of lost work given in Eq. (4.4) via the rate of entropy creation in the Universe ( $\dot{\Omega}_S$ ) due to the system steady-state operation (Teixeira et al., 2016).

$$\Delta \dot{B} = (\dot{B}_{out} + \dot{B}_{out}^W) - (\dot{B}_{in} + \dot{B}_{in}^W) \quad (4.3)$$

$$\dot{W}^{LOST} = T_0 \cdot \dot{\Omega}_S \quad (4.4)$$

A total of six exergy balances – for SSLC-Case and MPSC-Case with three gas-loads each – were conducted for RER-1 and RER-2. Material-streams and power-streams are classified as inlet or outlet and are attributed to a system or subsystem. Fig. 4.5 depicts the subsystems for SSLC-Case and MPSC-Case, including the respective overall gas-plants. RER-1 is suitable only for exergy analysis of GT area, while all other subsystems and gas-plants use RER-2.

#### 4.2.2.6. Exergy Efficiencies

Exergy efficiencies of oil/gas processing with RER-1 leads to useless high values due to high chemical exergy of hydrocarbons; i.e., RER-1 exergy efficiencies via Eq. (4.5a) tend to be numerically high and similar for corresponding systems/subsystems of SSLC-Case and MPSC-Case at analogous gas-loads. Moreover, they also present low sensitivity to process changes and are inefficient for exploring potential FPSO improvements and trade-offs.

On the other hand, RER-2 exergy analysis of non-reactive systems allows using simple exergy efficiency formulas, Eq (4.5a), to compare SSLC-Case and MPSC-Case, excluding the respective GT areas which are appropriate for RER-1 efficiencies. To validate the present methods, exergy efficiencies of GTs with RER-1 were calculated via Eq. (4.5b) and compared with counterparts of Gallo et al. (2017) in Appendix 4.A.

$$\eta = \frac{\dot{B}_{out,total}}{\dot{B}_{in,total}} \quad (4.5a)$$

$$\eta_{GT} = \frac{\dot{B}_{out,electricity}}{\dot{B}_{in,fuel-gas}} \quad (4.5b)$$

### 4.2.3. Energy Metrics

SSLC-Case and MPSC-Case are also compared via energy metrics such as Fuel Intensity ( $\text{BOE}_{\text{Burn}}/10^3 \cdot \text{BOE}_{\text{Produced}}$ ),  $\text{CO}_2$  Intensity ( $\text{tCO}_2^{\text{Emitted}}/10^3 \cdot \text{BOE}_{\text{Produced}}$ ) and FPSO Power Consumption – considering  $\approx 23\text{MW}$  of background power consumption (Cruz et al., 2018).

### 4.2.4. Economic and Footprint Assessments

Equipment *FCI* and weight were estimated for SSLC-Case and MPSC-Case via Aspen Capital-Cost Estimator (2016-database). The inputs for turbo-generators are power (kVA) and GT-driver type. MPSC-Case heat exchangers were sized to smaller duties in Table 4.4 from corresponding exchangers of SSLC-Case downsized proportionally to the ratios of heat duty ( $Q$ ) per log-mean temperature difference (*LMTD*) at 100% gas-load (data in Table B1.2, Supplementary Materials) Table 4.5 presents compressors data extracted from simulation at 100% gas-load for *FCI* and weight estimation. *FCI* of heat exchangers of SSLC-Case were estimated with data in Table B2.1.1 (Supplementary Materials).

**Table 4.4. Heat exchanger areas: SSLC-Case and MPSC-Case.**

<i>TAG</i>	<i>Area SSLC (m<sup>2</sup>)*</i>	<i>Δ(Q/LMTD)</i>	<i>Area MPSC (m<sup>2</sup>)</i>	<i>Paralleled Exchangers MPSC</i>	<i>Total Design-Area MPSC (m<sup>2</sup>)</i>	<i>Spares</i>	<i>Identical Exchangers SSLC</i>	<i>Identical Exchangers MPSC</i>
<i>HX-102</i>	1093.0	52.8%	516.2	2	1032.5	1	2	3
<i>HX-202</i>	41.0	65.0%	14.3	3	43.0	1	2	4
<i>HX-204</i>	108.5	71.5%	30.9	3	92.7	1	2	4
<i>HX-501</i>	505.5	67.7%	163.4	3	490.2	1	2	4
<i>HX-502</i>	563.6	67.7%	182.3	3	546.8	1	2	4
<i>HX-601</i>	254.4	19.5%	204.7	1	204.7	1	2	2
<i>HX-602</i>	198.7	18.9%	161.2	1	161.2	1	2	2
<i>HX-603</i>	203.3	18.6%	165.6	1	165.6	1	2	2
<i>HX-604</i>	239.4	18.8%	194.4	1	194.4	1	2	2
<i>HX-701_exp</i>	683.2	76.7%	159.0	1	867.9	1	2	2
<i>HX-702_inj</i>		48.1%	354.4	2		1	2	3
<i>HX-901</i>	8.6	46.6%	4.6	2	9.2	1	2	3

\*Cruz et al. (2018)

**Table 4.5. Compressors: data for FCI and weight estimation.**

<i>SSLC-Case</i>															
<i>TAG</i>	<i>Casing Material</i>	<i>Flow<sup>Inlet</sup> (m<sup>3</sup>/h)</i>	<i>P<sup>Inlet</sup> (kPag)</i>	<i>T<sup>Inlet</sup> (°C)</i>	<i>P<sup>Discharge</sup> (kPag)</i>	<i>MW (kg/kmol)</i>	<i>Cp/Cv</i>	<i>Z<sup>Inlet</sup></i>	<i>Z<sup>Outlet</sup></i>	<i>Tubes Material</i>	<i>Driver</i>	<i>Driver Power (kW)</i>	<i>Reduced Gear Driver</i>	<i>Paralleled Machines</i>	<i>Identical Items</i>
<i>C-101</i>	CS <sup>#</sup>	10477	1699	20.3	5149	24.33	1.335	0.921	0.921	SS316L	Motor	8223	No	1	2
<i>C-201</i>	CS	1100	124	38.55	556	39.12	1.163	0.964	0.964	SS316L	Motor	139	No	1	2
<i>C-202</i>	CS	1878	531	20.45	1749	38.22	1.211	0.901	0.901	SS316L	Motor	605	No	1	2
<i>C-501</i>	CS	2808	4399	32.3	10541	22.27	1.467	0.868	0.868	CS	Motor	4577	No	1	2
<i>C-502</i>	CS	1029	10491	38.39	24949	22.27	1.872	0.88	0.88	SS316L	Motor	4635	No	1	2
<i>C-601</i>	CS	10635	299	31.63	1034	30.62	1.302	0.986	0.986	CS	Motor	2123	No	1	2
<i>C-602</i>	CS	3867	1009	37.31	3051	30.62	1.329	0.967	0.967	CS	Motor	2241	No	1	2
<i>C-603</i>	CS	1288	3026	32.38	8775	30.62	1.443	0.923	0.923	SS316L	Motor	2238	No	1	2
<i>C-604</i>	CS	377	8725	29.71	24949	30.62	2.050	0.886	0.886	SS316L	Motor	2032	No	1	2
<i>C-701</i>	CS	584	24899	31.08	34470*	24.16	1.88	1.137	1.137	SS316L	Motor	7767	No	1	2
<i>C-901</i>	CS	1929	279	0.17	1619	44.10	1.179	0.791	0.791	CS	Motor	494	No	1	2
<i>MPSC-Case</i>															
<i>TAG</i>	<i>Casing Material</i>	<i>Flow<sup>Inlet</sup> (m<sup>3</sup>/h)</i>	<i>P<sup>Inlet</sup> (kPag)</i>	<i>T<sup>Inlet</sup> (°C)</i>	<i>P<sup>Discharge</sup> (kPag)</i>	<i>MW (kg/kmol)</i>	<i>Cp/Cv</i>	<i>Z<sup>Inlet</sup></i>	<i>Z<sup>Outlet</sup></i>	<i>Tubes Material</i>	<i>Driver</i>	<i>Driver Power (kW)</i>	<i>Reduced Gear Driver</i>	<i>Paralleled Machines</i>	<i>Identical Items</i>
<i>C-101 A/B</i>	CS	5240	1699	20.4	5149	24.34	1.335	0.921	0.921	SS316L	Motor	4059	Yes	2	3
<i>C-201 A/B/C</i>	CS	402	124	40	556	39.07	1.162	0.965	0.965	SS316L	Motor	43	Yes	3	4
<i>C-202 A/B/C</i>	CS	619	531	20.68	1749	38.25	1.211	0.901	0.901	SS316L	Motor	180	Yes	3	4
<i>C-501 A/B/C</i>	CS	952	4399	33.78	10541	22.27	1.461	0.870	0.870	CS	Motor	1619	Yes	3	4
<i>C-502 A/B/C</i>	CS	349	10491	40	24949	22.27	1.851	0.879	0.879	SS316L	Motor	1584	Yes	3	4
<i>C-601</i>	CS	10635	299	30.78	1034	30.62	1.276	0.986	0.986	CS	Motor	1707	Yes	1	2
<i>C-602</i>	CS	3867	1009	40	3051	30.62	1.327	0.968	0.968	CS	Motor	1824	Yes	1	2
<i>C-603</i>	CS	1287	3026	40	8775	30.62	1.426	0.931	0.931	SS316L	Motor	1864	Yes	1	2
<i>C-604</i>	CS	377	8725	40	24949	30.62	1.887	0.908	0.908	SS316L	Motor	1762	Yes	1	2
<i>C-701 A</i>	CS	121	24899	40	34470*	30.62	2.192	1.048	1.048	SS316L	Motor	1552	Yes	1	2
<i>C-701 B/C</i>	CS	296	24899	40	34470*	24.13	1.879	1.136	1.136	SS316L	Motor	3685	Yes	2	3
<i>C-901 A/B</i>	CS	970	279	-6	1619	44.10	1.186	0.777	0.777	CS	Motor	229	Yes	2	3

<sup>#</sup>Carbon-Steel. \*Aspen Capital-Cost Estimator limit ( $P^{Discharge,C-701}=54949$  kPag).

### 4.3. Results and Discussion

#### 4.3.1. Energy and Environmental Analyses

Six simulation cases – SSLC-Case and MPSC-Case at  $\approx 100\%/ \approx 50\%/ \approx 25\%$  gas-loads – comprehend a large amount of data, hence only relevant information is presented. Table 4.6 shows compressor flow rates of SSLC-Case and the counterparts of MPSC-Case.

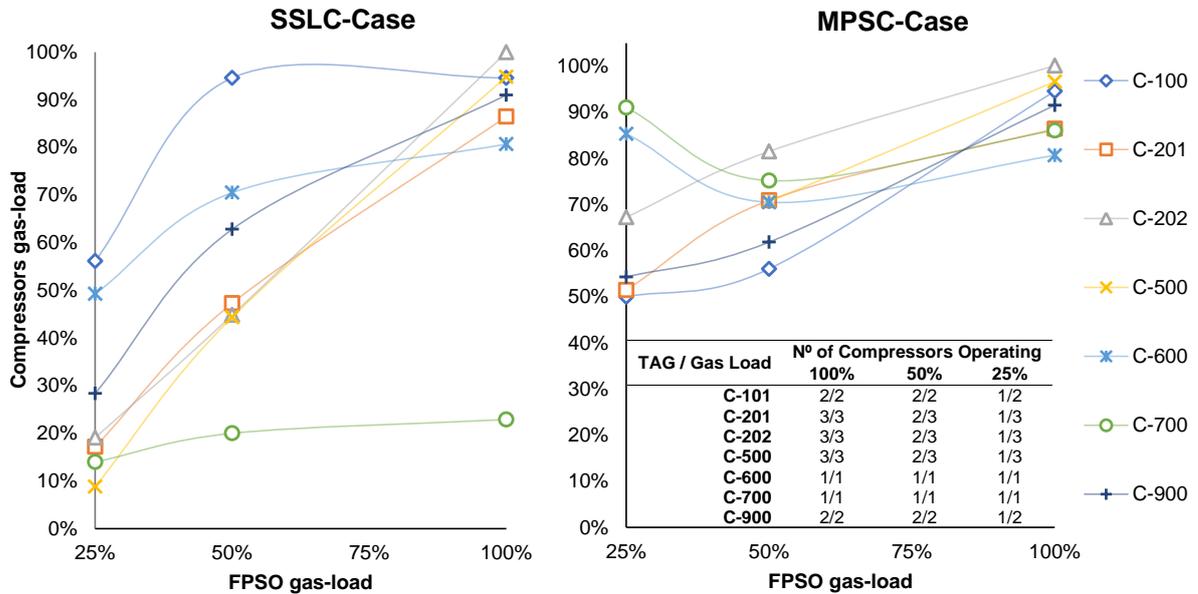
**Table 4.6. Compressor schemes: SSLC-Case and MPSC-Case.**

TAG	SSLC-Case					MPSC-Case							
	Inlet Flow ( $m^3/h$ ) at FPSO %Gas- Load			$\eta_p$	VSD Min. Flow ( $m^3/h$ )	Parallel Comps.	Design Flow ( $m^3/h$ )	VSD		Min. Flow ( $m^3/h$ )	Total Flow ( $m^3/h$ )	Min. Flow ( $m^3/h$ )	$\eta_p$
	100%	50%	25%					Min.	Flow				
C-101	10477	6418	2943	82%	5239	2	5239	2881	10477	2881	79%		
C-201	1100	610	219	74%	550	3	367	202	1100	202	69%		
C-202	1878	974	415	75%	939	3	626	344	1878	344	71%		
C-501	2802	1286	243	77%	1401	3	934	514	2802	514	72%		
C-502	1027	473	91	73%	514	3	342	188	1027	188	69%		
C-601	10635	9198	6289	83%	5318	1	10635	5850	10635	5850	82%		
C-602	3867	3368	2343	79%	1933	1	3867	2127	3867	2127	78%		
C-603	1288	1114	767	74%	644	1	1288	708	1288	708	73%		
C-603*	1288		1001		644								
C-604	377	316	204	70%	189	1	377	207	377	207	69%		
C-604*	377		324		189								
C-701	121	100	64	68%	61	1	121	67	576	67	65%		
C-701*	576		108		288	2	227	125			67%		
C-901	1929	1331	601	75%	964	2	964	530	1929	530	72%		

\*Full-injection mode.

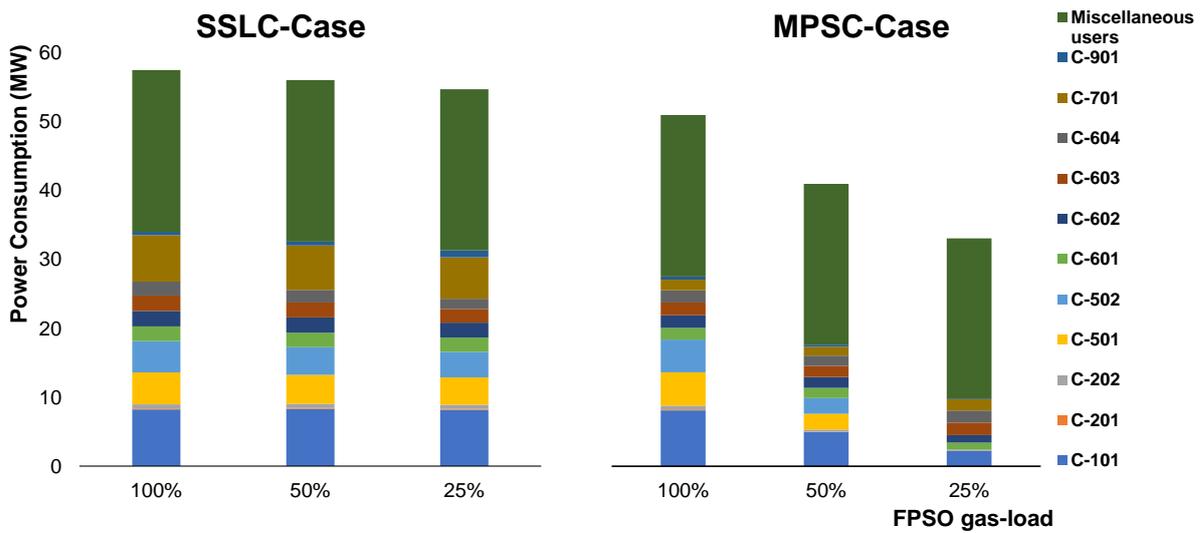
Only the CO<sub>2</sub> compressors (C-600) were not divided into MPSCs because as the gas-load reduces, the %CO<sub>2</sub> of raw gas increases, giving CO<sub>2</sub> flow rates always above 55% of design capacity for such SSLCs (Table 4.6, VSD Min. Flow); i.e., VSD alone can prevent surge. In all other cases, two or more compressors in parallel were necessary to achieve minimum flow with VSD. As compressor efficiencies slightly reduce as flow rate decreases, the advantage of dismissing anti-surge recycles is shadowed a little. Fig. 4.6 shows compressor gas-loads versus FPSO gas-load for SSLC-Case and MPSC-Case. In SSLC-Case compressors generally operate above 80% of design capacities for 100% FPSO gas-load, while at 25% gas-load all compressors operate below 60% of design capacities, being C-700 – EOR compressor – the

only exception since it must inject all processed gas if gas exportation is not possible (i.e., full-injection mode). Consequently, C-700 operates at 15%-20% of its design capacity when NG is being exported; i.e., in SSLC-Case all compressors operate at high recycle ratios, especially the C-700.

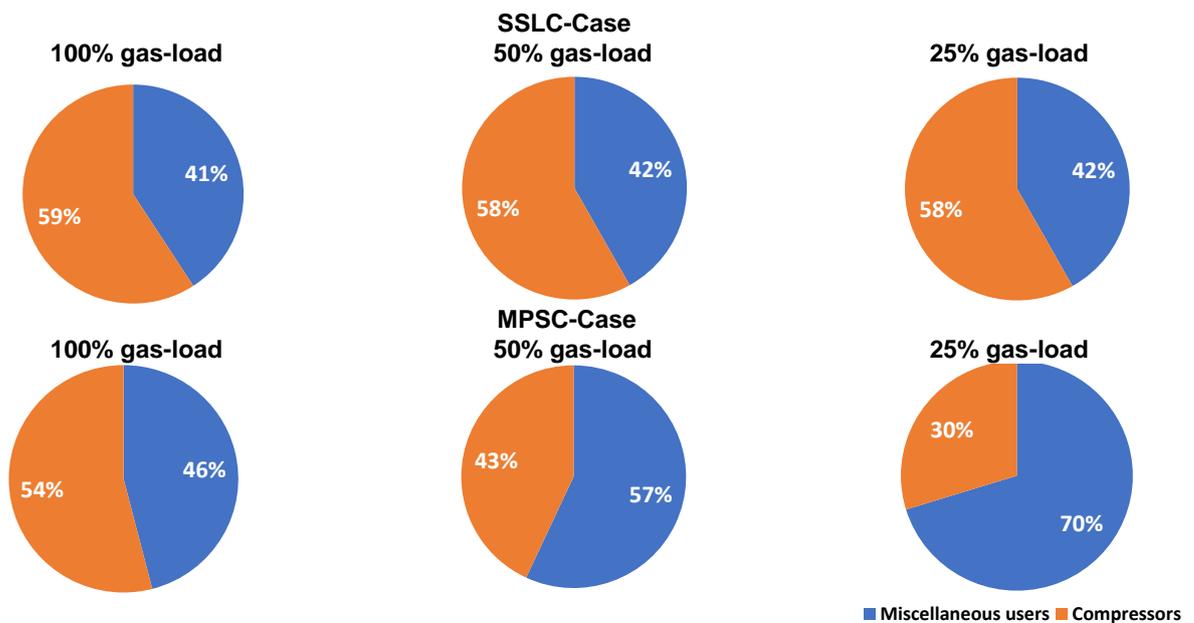


**Figure 4.6. Compressors gas-loads versus FPSO gas-load.**

As FPSO gas-load decreases in MPSC-Case, compressors operate above 55% of design capacities along the entire FPSO lifetime. It is only necessary to shut down part of the paralleled compressors (as in the table inside Fig. 4.6) and using VSD to make anti-surge recycles unnecessary. Fig. 4.7 demonstrates MPSC power savings, showing power consumption of SSLC-Case almost constant along FPSO lifespan due to anti-surge recycles that keep compressor design flow at partial gas-loads. Fig. 4.8 shows total power consumption of SSLC-Case modestly decreasing as gas-load reduces, while MPSC-Case shows power savings of 11% and 39% respectively at 100% and 25% gas-loads due to absence of anti-surge recycles.



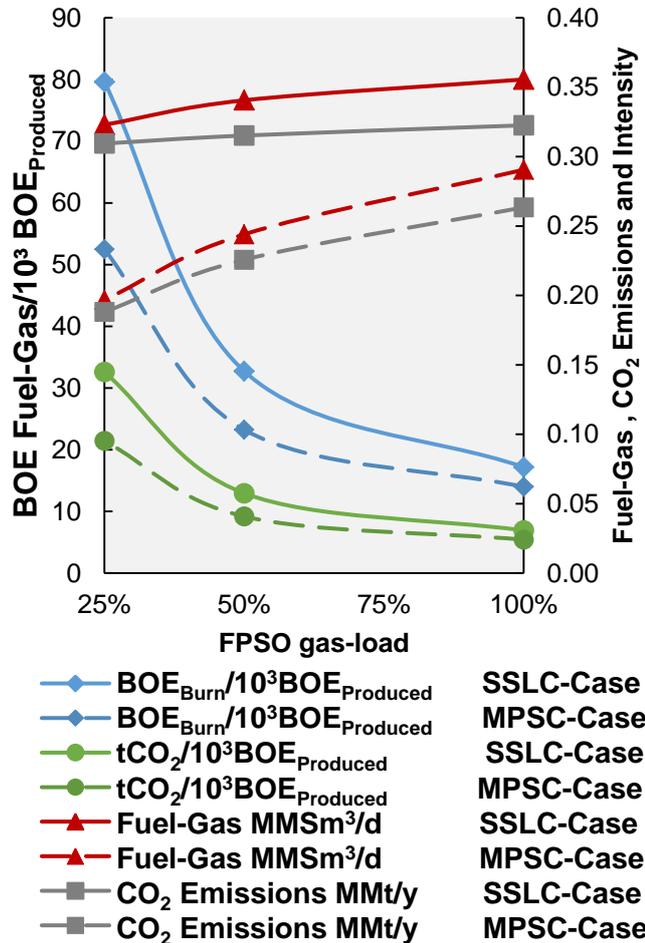
**Figure 4.7. Power consumptions versus FPSO gas-load: compressors and miscellaneous units.**



**Figure 4.8. Compressor share of total power consumption versus FPSO gas-load.**

In MPSC-Case at 25% gas-load, the power consumption of compressors is not anymore the greatest share, while in SSLC-Case compressors continue to be the greatest power sink due to anti-surge gas recycles, as depicted in Fig. 4.8. The SSLC-Case gas-plant consumes a considerable fraction of its carbon input, especially at low gas-loads. Given the current climate-change scenario, it is important to increase the usable energy ratio of oil-gas production. Consequently, the carbon intensity of FPSO products should be reduced to meet sustainability commitments. There are many energy return ratios reported in the literature; e.g., energy return on investment (EROI) and net energy ratio (NER). In this work, net energy

ratio and carbon intensity of oil-gas are respectively reported in terms of fuel-gas consumption and tCO<sub>2</sub> emitted both per BOE produced. These ratios are reported in Fig 4.9 to unveil the different energy productivities of SSLC-Case and MPSC-Case gas-plants at 25%/50%/100% gas-loads.



**Figure 4.9. SSLC-Case and MPSC-Case: Fuel-Gas and CO<sub>2</sub> intensities versus FPSO gas-load.**

The power consumption of compressors has a significant impact on FPSO fuel-gas consumption and CO<sub>2</sub> emissions. Therefore, SSLC-Case energy and environmental performance are drastically affected by its anti-surge recycles, as demonstrated in Fig. 4.9, which also shows that MPSC-Case is more environmentally friendly in terms of net energy ratio and CO<sub>2</sub> intensity, especially at low gas-loads. At 25% gas-load the fuel-gas and CO<sub>2</sub> intensities of MPSC-Case are 34% lower than the counterparts of SSLC-Case. More results are available in Supplementary Materials, Sec. B2.1 (SSLC-Case) and Sec. B2.2 (MPSC-Case).

### 4.3.2. Exergy Analysis

To shed more light on the thermodynamic advantages of MPSC-Case over SSLC-Case, 12 exergy analyses were performed for MPSC-Case and SSLC-Case, with RER-1 and RER-2, at three FPSO gas-loads. Exergy flows and exergy balances for each gas-load of MPSC-Case and SSLC-Case, are available in Supplementary Materials B, Section B3.

Fig 4.10 reports overall flow rates of exergy destruction and wasted for SSLC-Case and MPSC-Case considering RER-1. RER-1 is satisfactory only for exergy analysis of GTs, because it makes other exergy flows quite invariant compared to changes observed in GTs exergy flows; that is, RER-1 makes the exergy efficiency of physical units falsely high (~99%) in all gas-loads and cases. This results from the enormous values of inlet/outlet chemical part of the exergy flows that are invariant in physical units, making changes of physical exergy insignificant. Fig 4.10 shows that at 100% gas-load the rate of exergy destruction and wasted is similar for SSLC-Case and MPSC-Case, where GTs are major exergy destructors and exergy wasters comparatively to other systems, because GTs operate with highest spontaneity and irreversibility on FPSO topside thanks to highly irreversible combustion chemical reactions. Considering flue-gas contributing to the outlet exergy flow, the exergy efficiency of GTs via Eq. (4.5a) is  $\approx 61\%$  for both SSLC-Case and MPSC-Case at any FPSO gas-loads. On the other hand, operating at 25% gas-load, there is a reduction of 46.7% on the flows of destroyed plus wasted (flue-gas+ SW discharge) exergy of the overall MPSC-Case vis-à-vis SSLC-Case, confirming the thermodynamic superiority of MPSC-Case over SSLC-Case.

Fig 4.11 depicts SSLC-Case and MPSC-Case exergy Sankey diagrams for GTs with RER-1 at all gas-loads, showing that inlet and outlet GT exergy flows are almost the same in SSLC-Case (Fig. 4.11, left), including the exergy destruction rate, no matter the FPSO gas-load. The SSLC anti-surge recycles keep constant compressor inlet gas flows, consequently FPSO power and fuel-gas consumptions are almost constant as gas-load decreases (Fig. 4.8). Therefore, GTs operate at almost constant throughput along FPSO lifetime. Counterpointing this, in MPSC-Case (Fig. 4.11, right) as FPSO gas-load decreases, inlet fuel-gas exergy flow decreases, as well as the flows of destroyed exergy, wasted exergy (flue-gas and SW discharge) and electricity, all consequences of inexistent anti-surge recycles.

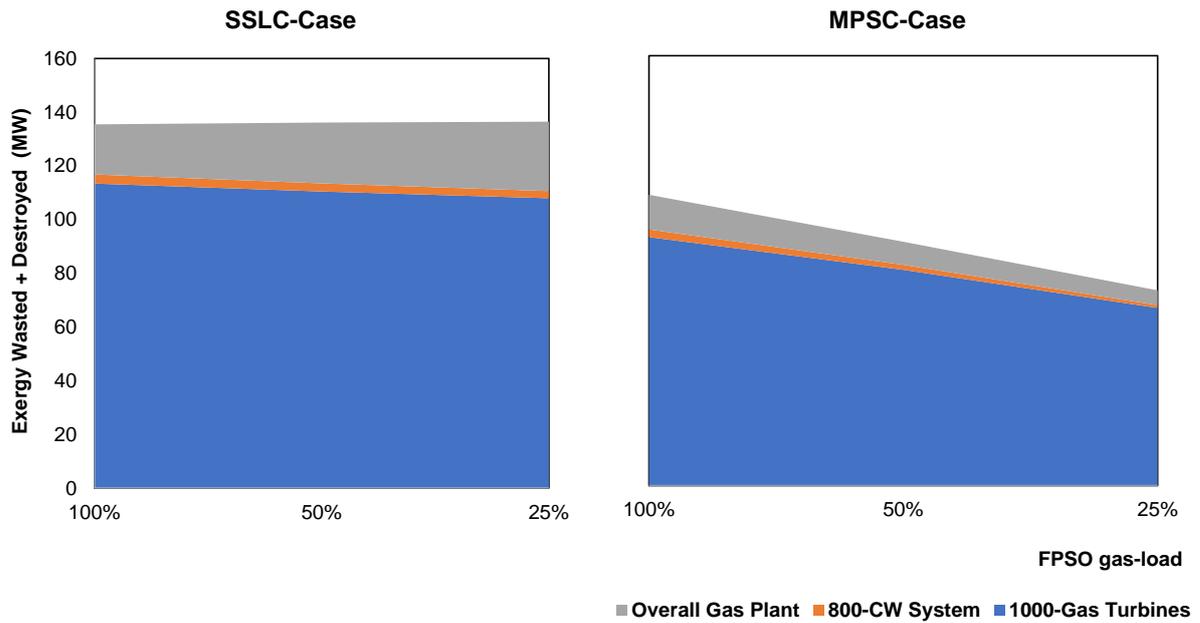


Figure 4.10. Destroyed and wasted exergy versus FPSO gas-load with RER-1.

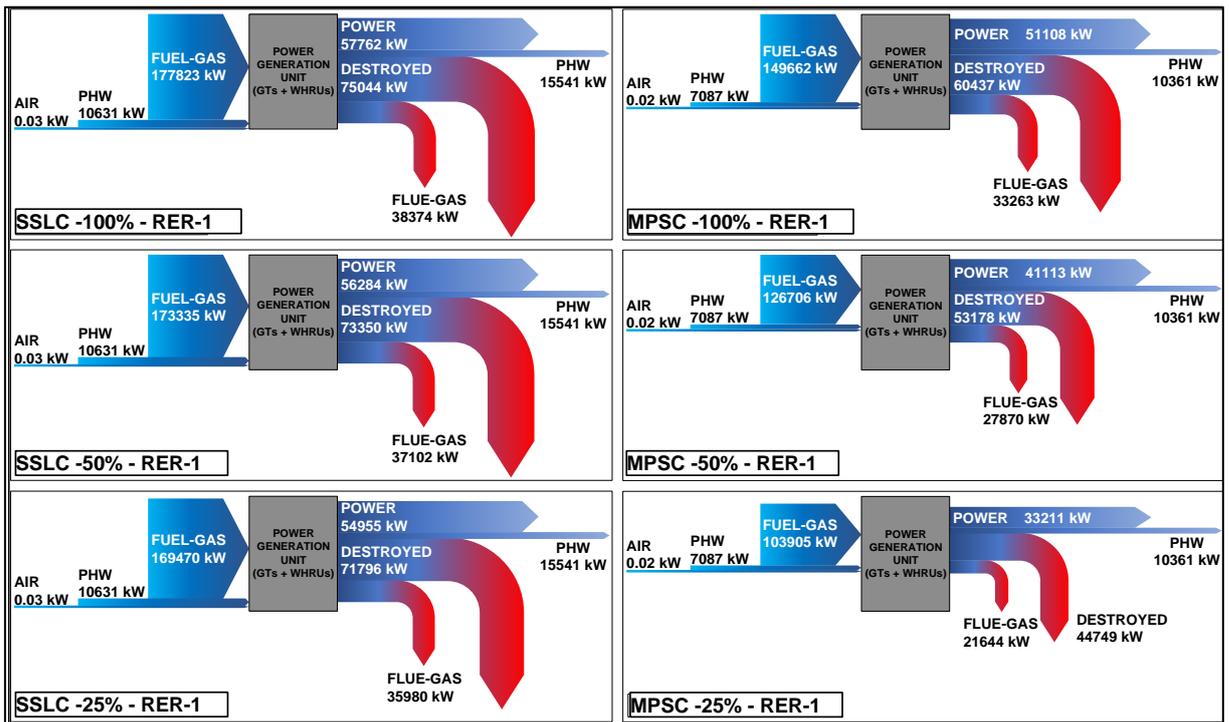
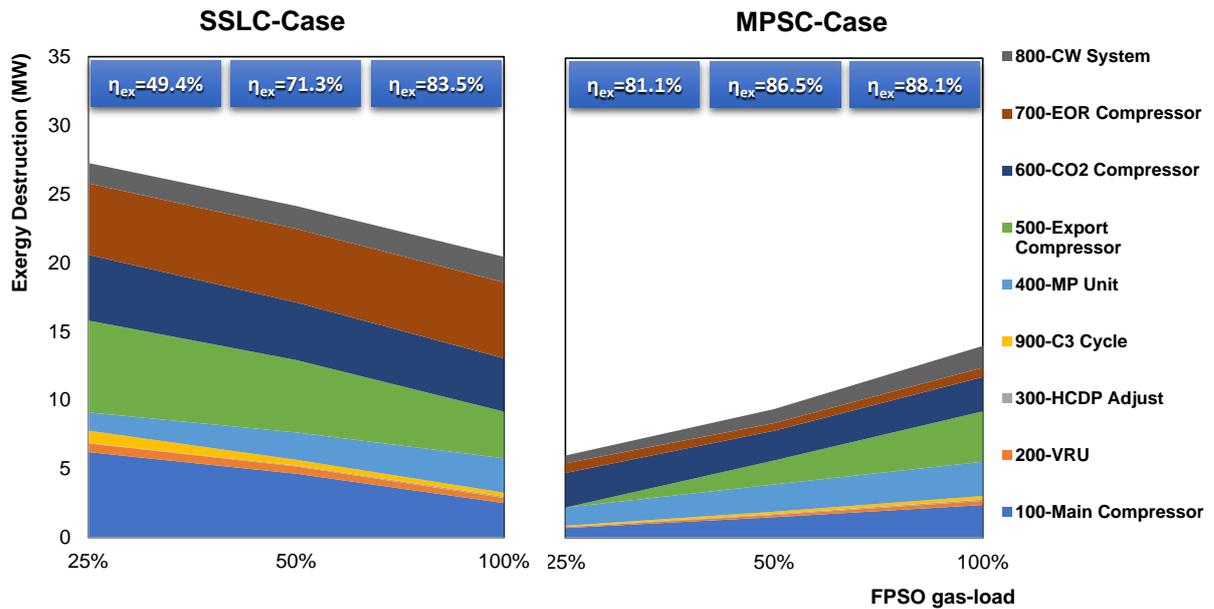


Figure 4.11. Exergy Sankey diagrams for GT power generators with RER-1: SSLC-Case (left) and MPSC-Case (right) at various FPSO gas-loads.

Fig 4.12 depicts flow rates of exergy destruction with RER-2 for subsystems of SSLC-Case and MPSC-Case, including overall exergy efficiencies via Eq. (4.5a) for the respective gas-plants (green-box, Fig. 4.3). The exergy destruction rates are differences between inlet and outlet exergy flow rates across the boundaries of subsystems in Fig. 4.5. Fig. 4.12 shows that RER-2 exergy analysis is much more sensitive to irreversibilities in physical subsystems

(compressors, exchangers, separators) in comparison with RER-1 exergy analysis, unveiling that total exergy destruction rates of MPSC-Case gas-plant are 32.5%, 62.2% and 78.6% lower than the SSLC-Case gas-plant counterparts at 100%/50%/25% FPSO gas-loads, respectively. It is also interesting to observe that while exergy destruction rate increase with gas-load in MPSC-Case the opposite is seen in SSLC-Case, proving that the increased activity of anti-surge recycles at decreasing FPSO gas-loads increase exergy destruction rates even at low FPSO gas-loads.



**Figure 4.12. Exergy destruction rate versus FPSO gas-load with RER-2: SSLC-Case, MPSC-Case.**

Fig. 4.13 depicts exergy Sankey diagrams with RER-2 for SSLC-Case and MPSC-Case at 25%/50%/100% FPSO gas-loads, showing increasing exergy destruction rate of SSLC-Case as FPSO gas-load decreases. At 25% FPSO gas-load, 37% of the inlet exergy flow is destroyed in SSLC-Case gas-plant (green-box, Fig. 4.3). As a result, the decreases of outlet NG exergy (53% at 50% gas-load and 91% at 25% gas-load) are more pronounced than the respective decreases in the feed gas (respectively, 43% and 76%). In MPSC-Case, on the other hand, the decreases of outlet NG exergy flow rate are proportional to the gas feed reductions. The exception is for FPSO gas-load below 25%, because the entire gas is necessary as fuel-gas for power production. Operating at such extremely low gas-load would demand several small paralleled compressors (in MPSC-Case) or full gas recycles (in SSLC-Case), becoming both SSLC-Case and MPSC-Case very inefficient in terms of *FCI* and/or energy use, respectively. Adding such low NG load to the CO<sub>2</sub>-rich EOR stream seems to be a

more reasonable solution, which was adopted in MPSC-Case at 25% FPSO gas-load in Fig. 4.13.

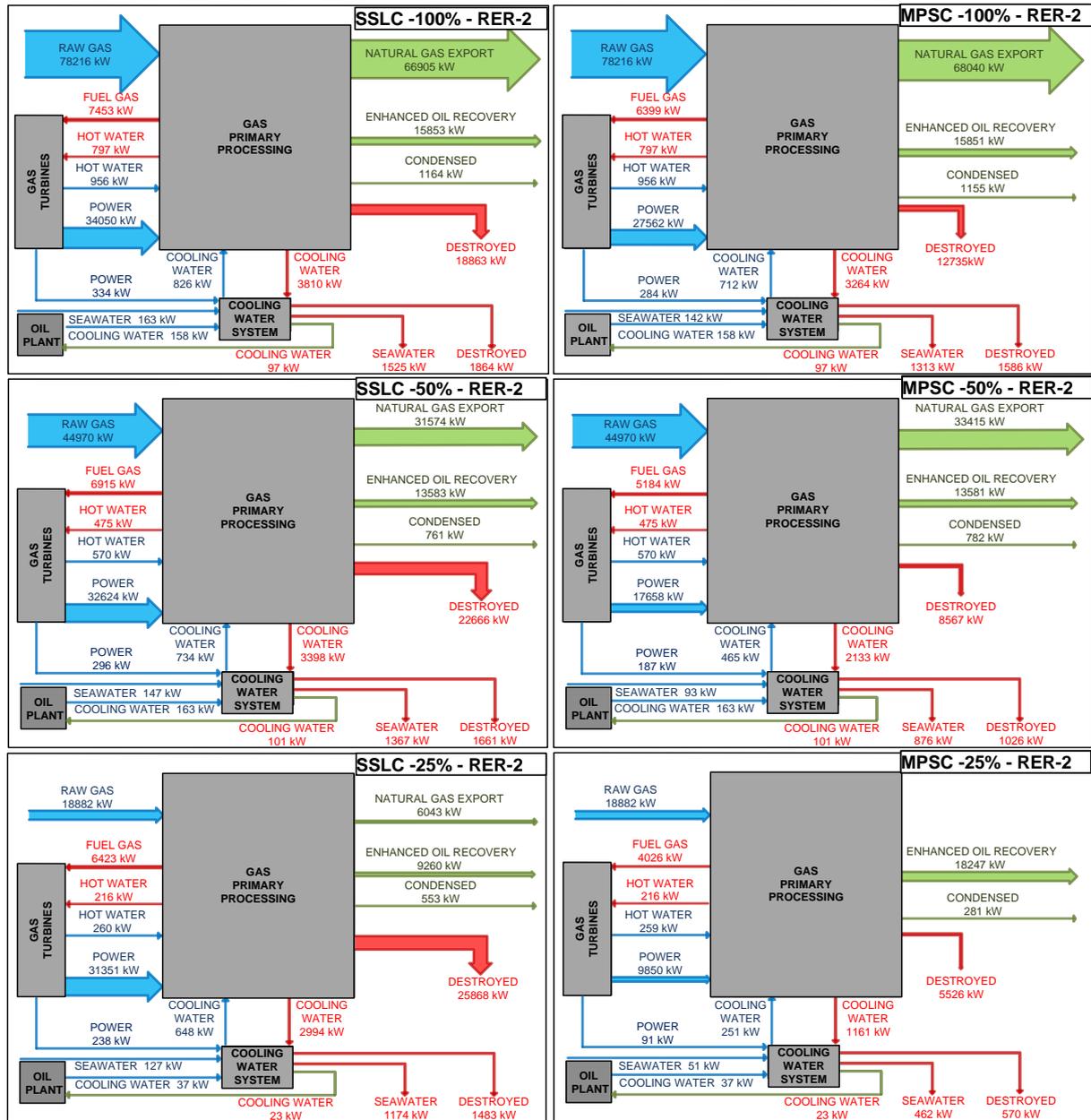
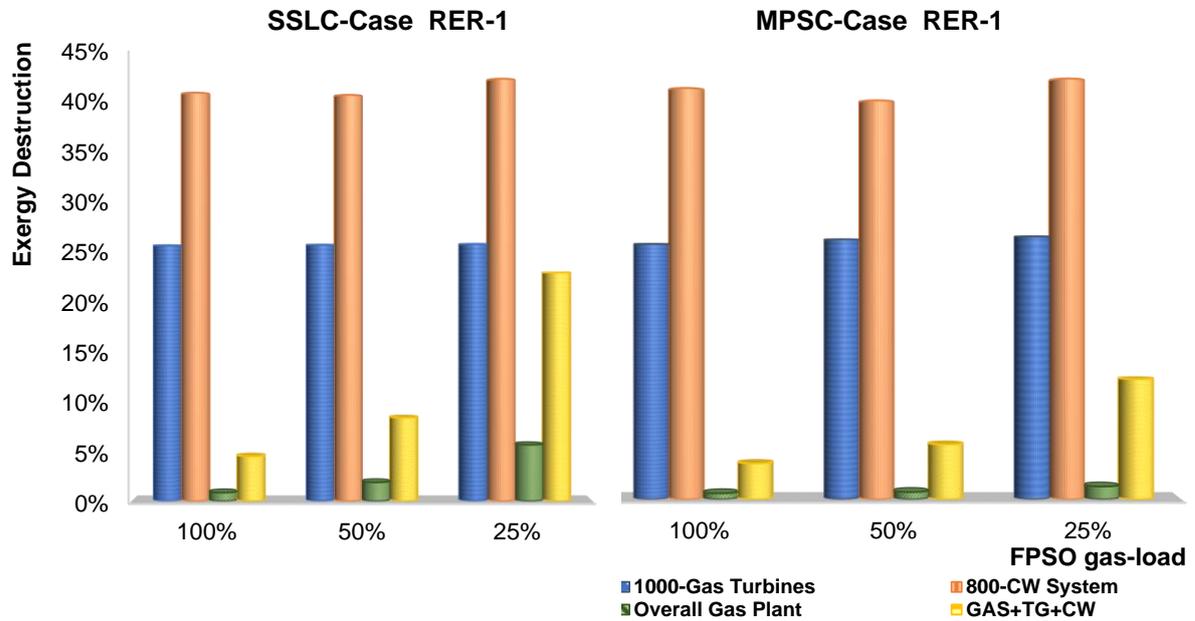


Figure 4.13. Exergy Sankey diagrams versus FPSO gas-load with RER-2: SSLC-Case, MPSC-Case.

Figs 4.14 and 4.15 show percent exergy destructions of gas-plant, GTs, CW system and subsystems, respectively considering RER-1 and RER-2. Percent exergy destructions in Fig. 4.14 are obtained with absolute destructions (Fig. 4.10) and inlet exergy flow rates of envelopes in Fig. 4.3. Percent exergy destructions are similar for GT and CW envelopes of SSLC-Case and MPSC-Case, but, the high percent exergy destruction of CW system is illusionary and consequence of the very low inlet exergy flow rates of CW streams in RER-1

which create high relative values. Thanks to the anti-surge recycles of SSLC-Case, the percent exergy destruction of gas-plant envelope increases as gas-load reduces. Without such anti-surge recycles in MPSC-Case gas-plant, such increase is almost unnoticeable.



**Figure 4.14. Exergy destructions (%) versus gas-load with RER-1: SSLC-Case, MPSC-Case.**

Fig. 4.15 provides segmented portraits of percent exergy destructions of SSLC-Case and MPSC-Case gas-plants using RER-2. Percent exergy destructions are calculated with absolute destructions (Fig. 4.12) and respective inlet exergy flow rates of subsystems. Gas-plant percent exergy destructions in Fig. 4.15 are much higher than Fig. 4.14 counterparts. The underlying reason has to do with the considered RER: Despite RER invariance of absolute exergy destructions, RER-1 embodies chemical exergy in inlet exergy flows, dramatically inflating them and appreciably reducing percent destructions, while RER-2 put inlet exergy flows into appropriate scales entailing better discrimination of gas-plant irreversibilities. Fig. 4.15 also evinces gas-export compressors (C-500) as the SSLC-Case subsystem with highest percent exergy destruction at 25% gas-load, but at 100% gas-load – i.e., design flow – percent destruction goes down thanks to no utilization of anti-surge recycle. EOR compressor (C-700) and CO<sub>2</sub> compressors (C-600) behave differently, keeping high percent exergy destruction even at 100% gas-load, since they always operate with high anti-surge recycle ratios due to large sizing suited for full-injection operation mode. In MPSC-Case most compressors operate with low percent exergy destruction comparatively to SSLC-Case.

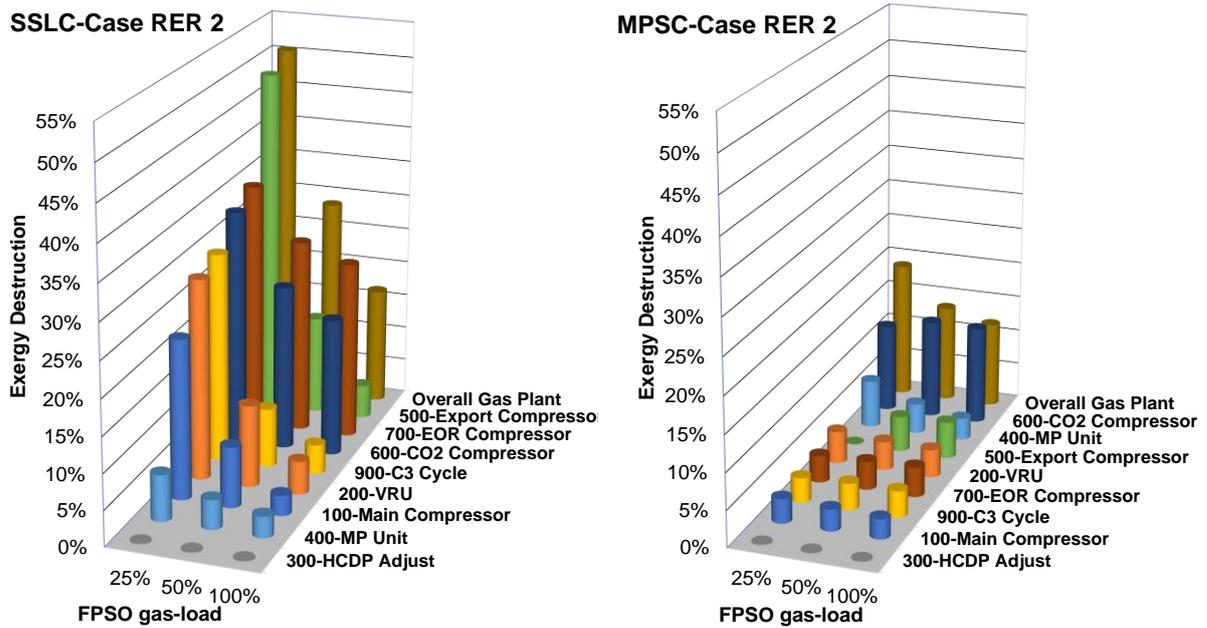


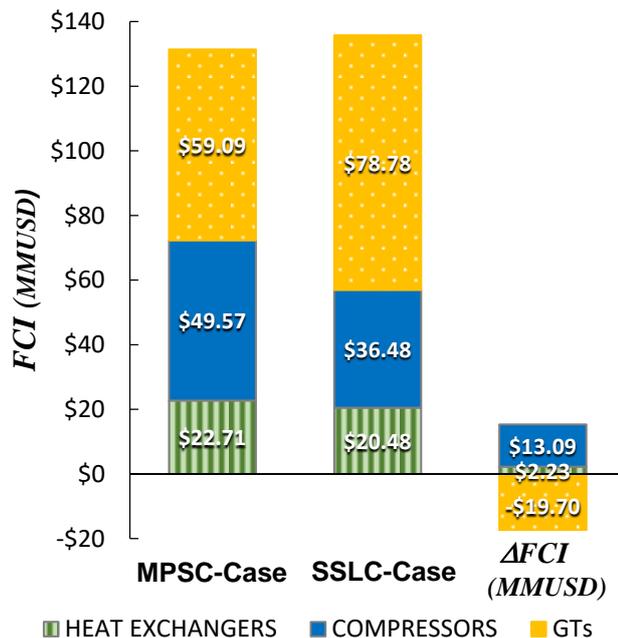
Figure 4.15. Exergy destructions (%) with RER-2: SSLC-Case, MPSC-Case.

#### 4.3.2.1. Exergy Analysis: Consistency Check

Following Teixeira et al. (2016), the consistency of exergy analysis can be checked by calculating the lost power (work)  $\dot{W}^{LOST}$  using two thermodynamically independent routes: via the rate of exergy destruction  $\dot{\Delta B}$  in Eq. (4.3) and via the 2<sup>nd</sup> Law  $\dot{W}^{LOST}$  formula in Eq. (4.4). Since all calculations of exergy flows rely on steady-state simulation of heavy flowsheets, with hundreds of streams and dozens numerically iterated recycles, followed by a numerically-intensive treatment with thermodynamic properties extracted from the flowsheet already incorporating some round-off errors, it is natural that  $\dot{\Delta B}$  and  $\dot{W}^{LOST}$  present some unavoidable discrepancy. In the present study, very low discrepancies smaller than 0.1% were encountered for the majority of the 12 exergy analyses. The exception corresponds to SSLC-Case at 100% gas-load with RER-1, which reached 1% of discrepancy between  $\dot{\Delta B}$  and  $\dot{W}^{LOST}$ . Results of consistency checks are in Tables B3.2.1 to B3.2.4, Supplementary Materials B, Section B3.2.

### 4.3.3. Investment and Footprint Assessments

MPSC-Case frankly outperformed SSLC-Case on energy-efficiency and exergy-efficiency grounds. But it is also important to compare *FCI* and costs of SSLC-Case and MPSC-Case to establish, or not, the economic feasibility of MPSC-Case and of MPSC design in FPSOs. Fig. 4.16 summarizes *FCI* comparison of SSLC-Case and MPSC-Case. Considering only compressors and intercoolers/aftercoolers, MPSC-Case has a 27% greater *FCI*, or 15.3MMUSD. However, since one less GT is necessary in MPSC-Case, there is also a *FCI* reduction of 19.7MMUSD, resulting a final MPSC-Case *FCI* (compressors, intercoolers/aftercoolers and GTs) 4.4MMUSD lower than the SSLC-Case counterpart. Therefore, in addition to being more energy/exergy efficient, MPSC-Case design also entailed 3% of *FCI* savings. However, the risk of using only two GTs in MPSC-Case is evident: at 100% gas-load GTs would operate at  $\approx 99\%$  of design capacity; i.e., with zero clearance for extra demand. Operation with 3 GTs – now at 67% of design load at 100% gas-load – is safer.

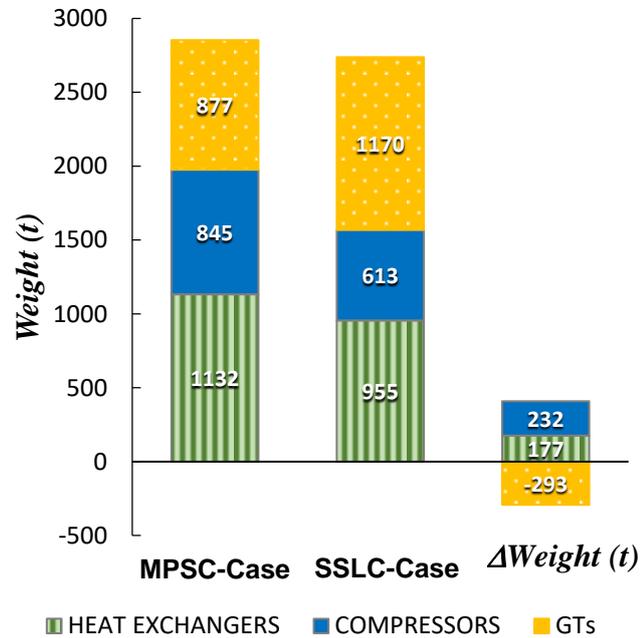


**Figure 4.16. *FCI* comparison: MPSC-Case versus SSLC-Case.**

### 4.3.4. Weight Comparison

Fig. 4.17 compares equipment weights showing MPSC-Case with  $\approx 115$  t more weight than SSLC-Case. Despite the criticality of equipment weight and footprint on FPSO topside, the

weight difference of only  $\approx 4\%$  against MPSC-Case is acceptable vis-à-vis its sound exergy, energy and environmental benefits.



**Figure 4.17. Equipment weight comparison: MPSC-Case versus SSLC-Case.**

#### 4.4. Conclusions

Process simulations, exergy analyses and *FCI*/footprint estimation of FPSO gas-plants adopting different compressor schemes – SSLC-Case (single-shaft larger compressors with anti-surge recycles) and MPSC-Case (multiple-paralleled smaller compressors without anti-surge recycles) – were performed unveiling an outstanding superiority of MPSC-Case over SSLC-Case on exergy-efficiency, energy-efficiency and environment grounds, contrasted by a small superiority of SSLC-Case in terms of *FCI*/footprint.

SSLC-Case and MPSC-Case were compared in terms of exergy-efficiency, *FCI*/footprint and CO<sub>2</sub> emissions. Simulations revealed that oversized compressors with anti-surge recycles (SSLC-Case) lead to almost constant power consumption along the field lifespan, regardless the gas-load processed. Consequently, fuel-gas and CO<sub>2</sub> intensities increase in SSLC-Case as gas-load decreases. It was shown that the efficiency of compressors can be kept high using VSD and smaller paralleled compressors. Moreover, the proposed MPSC-Case eliminates anti-surge recycles, turning the power demand proportional to gas-load, improving FPSO fuel-gas and CO<sub>2</sub> intensities.

Exergy analyses were conducted using two Reference Environmental Reservoirs, RER-1 and RER-2. RER-1 inflates exergy flows with the high chemical exergy of hydrocarbons, producing too high exergy efficiencies of physical operations (e.g., compressors, exchangers and separators); i.e., the modest irreversibilities of physical operations and of the entire gas-plant are masked giving exergy efficiencies for SSLC-Case and MPSC-Case always close to ~99%, regardless the gas-load. However, RER-1 is useful for chemically reactive operations with high spontaneity such as gas-turbines and combustors. For such operations, RER-1 produces reliable exergy efficiencies because there is huge rate of exergy destruction in gas-fired systems corresponding to high fractions of the inlet exergy flow. Counterpointing this, RER-2 deflates exergy flows by excluding the high chemical exergy of hydrocarbons, entailing that the exergy assessments of physical operations and of typical gas-plants (i.e., gas-turbines excluded) are now meaningful. Exergy analyses corroborate the simulation achievements, unveiling that MPSC-Case entails a much lower FPSO exergy destruction rate according to both RER-1 and RER-2. For FPSO gas-load ranging from 25% to 100%, the RER-2 exergy efficiency of SSLC-Case lies between 49% and 83%, whereas the counterpart of MPSC-Case is always from 81% to 88%.

Investment and footprint assessments indicate that MPSC-Case, besides being more exergy-efficient, energy-efficient and environmentally adequate, also entails 3% of *FCI* savings, despite increasing only 4% the equipment weight. The lower power demand of MPSC-Case allowed one less GT in the FPSO, compensating the *FCI*/footprint increases of compressors and exchangers.

In summary, MPSC-Case has much superior exergy/energy and environmental performances relatively to SSLC-Case. Surprisingly, the superiority of MPSC-Case extends to economic grounds, thanks to the elimination of a GT power generator. In a carbon taxation scenario, MPSC-Case would be even more profitable.

#### **4.5. References of Chapter 4**

- Albusaidi, W., Pilidis, P., 2015. An iterative method to derive the equivalent centrifugal compressor performance at various operating conditions: Part I: Modelling of suction parameters impact. *Energies* 8, 8497–8515. <https://doi.org/10.3390/en8088497>
- Allahyarzadeh-Bidgoli, A., Salviano, L.O., Dezan, D.J., de Oliveira Junior, S., Yanagihara, J.I., 2018. Energy optimization of an FPSO operating in the Brazilian Pre-salt region.

Energy 164, 390–399. <https://doi.org/10.1016/J.ENERGY.2018.08.203>

- Araújo, O.Q.F., Reis, A.C., de Medeiros, J.L., Nascimento, J.F., Grava, W.M., Musse, A.P.S., 2017. Comparative analysis of separation technologies for processing carbon dioxide rich natural gas in ultra-deepwater oil fields. *J. Clean. Prod.* 155, 12–22. <https://doi.org/10.1016/j.jclepro.2016.06.073>
- Arinelli, L.O., de Medeiros, J.L., Teixeira, A.M., Araújo, O.Q.F., 2019. Modeling of Supersonic Separators and Membrane Permeation Units for Processing of CO<sub>2</sub>-Rich Natural Gas with HYSYS Implementation, in: *Offshore Processing of CO<sub>2</sub>-Rich Natural Gas with Supersonic Separator: Multiphase Sound Speed, CO<sub>2</sub> Freeze-Out and HYSYS Implementation*. Springer International Publishing, Cham, pp. 163–213. [https://doi.org/10.1007/978-3-030-04006-2\\_6](https://doi.org/10.1007/978-3-030-04006-2_6)
- Barrera, J.E., Bazzo, E., Kami, E., 2015. Exergy analysis and energy improvement of a Brazilian floating oil platform using Organic Rankine Cycles. *Energy* 88, 67–79. <https://doi.org/10.1016/J.ENERGY.2015.03.091>
- Chang, K., Baril, R., Hull, M., Pepe, N.T.M., Nwachuku, I.B., Doezema, L.A., 2014. Microseepage of C<sub>2</sub>-C<sub>5</sub> alkanes over the Baldwin Hills in Los Angeles. *Atmos. Environ.* 87, 170–174. <https://doi.org/10.1016/j.atmosenv.2014.01.047>
- Chevron, 2018. *Climate Change Resilience: A Framework for Decision Making*. <https://www.chevron.com/-/media/shared-media/documents/climate-change-resilience.pdf> (Accessed 21-October-2019).
- Copenhagen Economics, 2017. The future of fossil fuels: How to steer fossil fuels use in a transition to a low-carbon energy system. An analysis of fossil fuels trajectories in low-carbon scenarios. [http://www.energy-transitions.org/sites/default/files/ETC\\_Copenhagen\\_Economics\\_-\\_The\\_future\\_of\\_fossil\\_fuels\\_-\\_Summary\\_Paper.pdf](http://www.energy-transitions.org/sites/default/files/ETC_Copenhagen_Economics_-_The_future_of_fossil_fuels_-_Summary_Paper.pdf) (Accessed 2-July-2018).
- Cruz, M.A., Araújo, O.Q.F., de Medeiros, J.L., 2018. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. *J. Nat. Gas Sci. Eng.* 56, 193–211. <https://doi.org/10.1016/j.jngse.2018.06.011>
- da Silva, J.A.M., de Oliveira Junior, S., 2018. Unit exergy cost and CO<sub>2</sub> emissions of offshore petroleum production. *Energy* 147, 757–766. <https://doi.org/10.1016/j.energy.2018.01.100>
- Dinçer, I., Rosen, M. *EXERGY: Energy, Environment and Sustainable Development*. Elsevier Science. <https://doi.org/10.1016/C2010-0-68369-6>
- Gallo, W.L.R., Gallego, A.G., Acevedo, V.L., Dias, R., Ortiz, H.Y., Valente, B.A., 2017. Exergy analysis of the compression systems and its prime movers for a FPSO unit. *J. Nat. Gas Sci. Eng.* 44, 287–298. <https://doi.org/10.1016/j.jngse.2017.04.023>
- Veloso, T.G.C., Sotomonte, C.A.R., Coronado, C.J.R., Nascimento, M.A.R., 2018. Multi-objective optimization and exergetic analysis of a low-grade waste heat recovery ORC application on a Brazilian FPSO. *Energy Convers. Manag.* 174, 537–551.

<https://doi.org/10.1016/j.enconman.2018.08.042>

- IEA, 2018. 2018 World Energy Outlook: Executive Summary. Oecd/Iea. <https://webstore.iea.org/download/summary/190?fileName=English-WEO-2018-ES.pdf> (Accessed 21-October-2019).
- IOGP, 2016. Environmental Performance Indicators - 2015 data, IOGP. London. [www.iogp.org](http://www.iogp.org) (Accessed 21-October-2019).
- Silva, F.C.N., Flórez-Orrego, D., de Oliveira Junior, S., 2019. Exergy assessment and energy integration of advanced gas turbine cycles on an offshore petroleum production platform. *Energy Convers. Manag.* 197, 111846. <https://doi.org/10.1016/j.enconman.2019.111846>
- Nguyen, T., Pierobon, L., Elmegaard, B., Haglind, F., Breuhaus, P., Voldsund, M., 2013. Exergetic assessment of energy systems on North Sea oil and gas platforms. *Energy* 62, 23–36. <https://doi.org/10.1016/j.energy.2013.03.011>
- Nguyen, T.V., Jacyno, T., Breuhaus, P., Voldsund, M., Elmegaard, B., 2014. Thermodynamic analysis of an upstream petroleum plant operated on a mature field. *Energy*. <https://doi.org/10.1016/j.energy.2014.02.040>
- Panton, L.K., Eide, B., Solutions, A., Førde, T., 2014. Exergy analysis of conventional and electrified oil and gas platforms. Norwegian University of Science and Technology. [https://ntnuopen.ntnu.no/ntnu-xmlui/bitstream/handle/11250/235802/757579\\_FULLTEXT01.pdf?sequence=2&isAllowed=y](https://ntnuopen.ntnu.no/ntnu-xmlui/bitstream/handle/11250/235802/757579_FULLTEXT01.pdf?sequence=2&isAllowed=y) (Accessed 23-October-2019).
- Petrobras, 2013. Polo Pre-Sal - Bacia de Santos - Etapa 1. IBAMA. [http://licenciamento.ibama.gov.br/Petroleo/Producao/Producao - Polo Pre-Sal - Bacia de Santos - Etapa 1 - Petrobras/](http://licenciamento.ibama.gov.br/Petroleo/Producao/Producao-Polo-Pre-Sal-Bacia-de-Santos-Etapa-1-Petrobras/) (Accessed 4-July-2017).
- Pierobon, L., Benato, A., Scolari, E., Haglind, F., Stoppato, A., 2014. Waste heat recovery technologies for offshore platforms. *Appl. Energy* 136. <https://doi.org/10.1016/j.apenergy.2014.08.109>
- Reis, M.M.L., Gallo, W.L.R., 2018. Study of waste heat recovery potential and optimization of the power production by an organic Rankine cycle in an FPSO unit. *Energy Convers. Manag.* 157, 409–422. <https://doi.org/10.1016/J.ENCONMAN.2017.12.015>
- Roussanaly, S., Aasen, A., Anantharaman, R., Danielsen, B., Jakobsen, J., Heme-De-Lacotte, L., Neji, G., Sødal, A., Wahl, P.E., Vrana, T.K., Dreux, R., 2019. Offshore power generation with carbon capture and storage to decarbonise mainland electricity and offshore oil and gas installations: A techno-economic analysis. *Appl. Energy* 233–234, 478–494. <https://doi.org/10.1016/j.apenergy.2018.10.020>
- Rystad Energy, 2019. FPSO market is booming with Brazil fueling demand. <https://www.rystadenergy.com/newsevents/news/press-releases/FPSO-market-is-booming-with-Brazil-fueling-demand/> (Accessed 21-October-2019).
- Soundararajan, K., Ho, H.K., Su, B., 2014. Sankey diagram framework for energy and exergy flows. *Appl. Energy* 136, 1035–1042. <https://doi.org/10.1016/j.apenergy.2014.08.070>

- Teixeira, A.M., Arinelli, L.O., de Medeiros, J.L., Araújo, O.Q.F., 2016. Exergy Analysis of Monoethylene glycol recovery processes for hydrate inhibition in offshore natural gas fields. *J. Nat. Gas Sci. Eng.* 35, 798–813. <https://doi.org/10.1016/J.JNGSE.2016.09.017>
- Voldsund, M., Nguyen, T.V., Elmegaard, B., Ertesvåg, I.S., Røsjorde, A., Jøssang, K., Kjelstrup, S., 2014. Exergy destruction and losses on four North Sea offshore platforms: A comparative study of the oil and gas processing plants. *Energy* 74, 45–58. <https://doi.org/10.1016/j.energy.2014.02.080>

## Appendix 4A: Comparison of Exergy Efficiencies of Gas-Turbines

Exergy flows of FPSO gas-plants were calculated with a novel methodology (Teixeira et al., 2016) mostly using RER-2. On the other hand, GTs exergy efficiency of SSLC-Case and MPSC-Case were determined with exergy flows according to RER-1 via Eq. (4.5a). However, the exergy efficiency of GTs according to Gallo et al. (2017) in Eq. (4.5b) offers an opportunity to compare and validate the present approach. This comparison is available in Table 4A.1 which shows that, despite the completely different methodologies for exergy flows, the present exergy efficiency of GTs using RER-1 and Eq. (4.5b) gave values very close to the value of Gallo et al. (2017).

**Table 4A.1. Exergy Efficiency Comparison for GTs (Eq. (4.5b) with RER-1).**

	<i>This work</i>		<i>Gallo et al.</i>
	<i>100% Gas-Load SSLC-Case</i>	<i>100% Gas-Load MPSC-Case</i>	<i>(2017)</i>
<i>FPSO Gas-Load [MMSm<sup>3</sup>/d]</i>	4.75	4.75	6.00
<i>Active GTs</i>	3	3	3
<i>Net Power [kW]</i>	57762	51108	69134
<i><math>\eta_{GT}</math></i>	32.5%	34.1%	33.8%

## 5. IMPACT OF SOLID WASTE TREATMENT FROM SPRAY DRYER ABSORBER ON THE LEVELIZED COST OF ENERGY OF A COAL-FIRED POWER PLANT

*This chapter is published a full-length original article in the Journal of Cleaner Production.*

CRUZ, M. DE A. et al. Impact of solid waste treatment from spray dryer absorber on the levelized cost of energy of a coal-fired power plant. **Journal of Cleaner Production**, v. 164, 2017.

### Abstract

Coal-fired power plants with semi-dry flue-gas desulfurization (semi-dry FGD) system produce daily tones of ashes contaminated with calcium sulfite. To turn this solid waste useful, e.g. to the cement industry, and avoid landfill disposal, the present study suggests a semi-dry FGD solid waste treatment unit, that promotes the dry oxidation of the calcium sulfite to calcium sulfate. Sizing of main equipment using pilot-plant data and patents allows economic evaluation of capital expenditure, operational and maintenance costs, and sale of the treated residue, allowing estimation of levelized cost of energy to assess the impact of the technology on the electricity price of a power plant using the proposed solid waste treatment unit. As base case, a Brazilian coal-fired power plant facing decision making process on semi-dry FGD waste destination is selected. Results demonstrate that the semi-dry FGD, without the solid treatment unit, has total levelized cost of energy increased in 0.56% (from 94.44 to 94.97 \$/MWh) resulting from solids waste disposal. If the treated semi-dry FGD waste was transferred (at zero revenue) as additive to a cement industry, the levelized cost of energy of the power plant would remain approximately unchanged. This is because the increase of 0.51\$/MWh resulting from the investment and operation and maintenance cost of the treatment unit is compensated by the decrease of 0.53\$/MWh, in virtue of the avoided waste disposal costs. However, if the commercialization as raw material of the treated semi-dry FGD waste is considered, a reduction of 2.83 \$/MWh (~3%) on the levelized cost of energy (to 92.14 \$/MWh) would occur. In both cases, the proposed treatment unit shows small impact on the total power plant levelized cost of energy, besides solving the solid management problems of landfill saturation, land use and costs related to landfill maintenance. Thus, it is adequate to implement the semi-dry FGD waste treatment unit on the power plant in question. The conclusion can be extended to plants with similar design and economic parameters.

**Keywords:** Coal-Fired Power Plant; Flue-Gas Desulfurization; Spray Dryer Absorbers; Solid Waste Treatment; Calcium Sulphite Oxidation; Levelized Cost of Energy.

Supplementary Materials for this chapter are found in Appendix H, Section C.

## Nomenclature

### Abbreviations

<i>CAPEX</i>	Capital Expenditure
<i>FBR</i>	Fluidized Bed Reactor
<i>FGD</i>	Flue-Gas Desulfurization
<i>IECM</i>	Integrated Environmental Control Model software
<i>OPEX</i>	Operational Expenditure
<i>PCC</i>	Pulverized Coal Combustion
<i>PFD</i>	Process Flow Diagram
<i>SDA</i>	Spray Dryer Absorber

### Roman letters

<i>AE</i>	Annual energy output (MWh/yr)
<i>D</i>	Diameter (m)
<i>f</i>	Annuity factor (%)
<i>HHV</i>	Higher Heating Value (kJ/kg)
<i>L</i>	Length (m)
<i>LCOE</i>	Levelized cost of energy (\$/MWh)
<i>MM</i>	Molecular mass (kmol/kg)
<i>n</i>	Polytropic exponent
<i>P</i>	Pressure (kPa)
<i>P<sub>w</sub></i>	Power (kW)
<i>R<sub>i</sub></i>	Reaction number <i>i</i> , where <i>i</i> is a counter
<i>q</i>	Flow rate (kg/h)
<i>R</i>	Gas constant (J/mol.K)
<i>r<sub>p</sub></i>	Pressure ratio
<i>t</i>	Lifetime of the plant (yr)
<i>T</i>	Temperature (K)
<i>TAC</i>	Total annualized capital costs (\$/yr)
<i>Z</i>	Average compressibility factor
<i>z</i>	Discount Rate (%)

### Greek Letters

$\Delta H_r$	Enthalpy of reaction (kcal/mol)
$\eta_p$	Polytropic efficiency

## 5.1. Introduction

Brazilian electricity matrix is dominated by hydropower generation. However, because of the recent water scarcity crisis in Brazil, hydroelectricity has been supplemented with electrical power generated by thermal power plants, resulting in 18% increase during 2013-2014 and presently represents 28.2% of the total Brazilian electricity source. In this same period, electricity produced by coal power plants has increased 24.2%, with mineral coal representing 9.6% of the thermopower source in Brazil (EPE, 2015a). The Brazilian energy demand will increase in an average of 3.6% yearly until 2019, thus it is expected that the use of coal-fired power plants continues to increase in short to medium term (EPE, 2015b). Furthermore, an average power plant technical lifetime of about 40 years for coal compared to 34 years for gas and 34 years for oil-fired power plants is estimated (Farfan and Breyer, 2017), indicating that the next decade will sustain supply of fossil energy accompanied by growing environmental legislation and policies. Consequently, technologies for reducing post-combustion emissions will assume a protagonist role, notably for carbon dioxide and sulfur oxides, the latter – flue-gas desulfurization (FGD) being a fingerprint of coal-fired power plants.

The commercial application of FGD is a challenge in terms of removal efficiency, price and availability of sorbent (Ma et al., 2000). Wet limestone FGD system is the most widely used process for flue-gas desulfurization because of its high performance and reduced operating cost (Cordoba, 2015). Less capital-intensive alternatives to wet FGD are sought, but are characterized by modest SO<sub>2</sub> removal despite reduced capital costs (Sage and Ford, 1996). semi-dry flue-gas desulfurization (semi-dry FGD) is an intermediate between dry and wet flue-gas scrubbing, with slightly lower costs than wet FGD, but presents residue disposal problems (Sage and Ford, 1996).

Nevertheless, about 12% of USA power plants were using semi-dry FGD systems in 2007. (EPRI, 2009) According to Alston Power, the market was running about 40% - 60% in favour of semi-dry FGD. The semi-dry technology has typically been employed on small to moderate size plants. Because of size limitations on the absorber tower, the maximum power served by a spray dryer is about 250 – 350 MW. Power plants with semi-dry FGD usually burns low-sulfur coal, in virtue of a limit of 95% on SO<sub>2</sub> removal efficiency. The semi-dry technology is interesting in regions where the water supply is limited (e.g., northeast of Brazil and western of the United States), because it consumes 30-40 less water than Wet-FGD. In terms of capital

expenditure (CAPEX), the cost of a semi-dry FGD is about 60% lower than the wet technology.(Blankinship, 2005) Environmental impact analyses of FGD alternatives are available (Wu et al., 2017), they mostly treat energy consumption - resource consumption and pollutant emissions, despite solid residues being recognized as the most critical for FGD technologies (Feng et al., 2014).

The semi-dry FGD solid waste is basically composed of calcium sulfite ( $\text{CaSO}_3$ ), varying amounts of unreacted fly ashes and lime. Most units do not present fly ash pre-collectors, often resulting in semi-dry FGD waste with high ash percentage (EPRI, 2009). Fly ashes from combustion of mineral coal show very low agglomerating property, but, in the presence of water, react with calcium hydroxide ( $\text{Ca}(\text{OH})_2$ ), at ambient temperature, to form aggregating (pozzolanic) compounds (Thomas, 2007; ABNT, 2015). In USA only 22% is used in commercial applications, with mining applications representing 83% of its use. Other uses include oil/gas field services, cementitious products, cement replacement in concrete (pozzolanic material), engineering applications, agriculture, soil stabilization and as wet FGD sorbent (ACAA, 2014).

Most coal-fired power plants that use semi-dry FGD systems currently dispose its solid residue on landfills. The solids transportation and landfill maintenance is expensive, unhealthy to local workers (EPRI, 1998) and unsustainable in long term, because the landfill becomes saturated, as could be noticed in Fig.5.1. The power plant complex of Pecém, shown in Fig. 5.1, has three 360 MW coal-fired power plants units, with semi-dry FGD for  $\text{SO}_2$  control. The start-up occurred in 2012 and after 4 years of operation 2 ash landfills becomes saturated and a third one is being built. In Brazil and USA, among other countries, the use of semi-dry FGD systems is expected to grow, or at least be maintained, creating the need for increased alternatives for utilization for the semi-dry FGD waste.



Fig. 5.1. Ash landfills of a Brazilian coal-fired power plant in December of 2016.  
Source: Google Earth (satellite images) and personal archive (landfill picture).

Among possible uses, FGD gypsum is recognized as a substitute for natural gypsum in the cement industry (Galos et al., 2002). In fact, as stated by Mikulčić et al. (2016), the challenge for the cement industry is to use alternative raw materials especially wastes originated from other industries, highlighting, among others, fly ashes and gypsum from coal power plants.

To explore this synergy, i.e., the use of the semi-dry FGD waste as pozzolanic material, solids treatment to comply with standards are required. Besides other parameters, standards (e.g., ABNT, 2015; ASTM, 2015) define threshold limit for  $\text{CaSO}_3$  content in the semi-dry FGD waste. For instance, in Brazil, when  $\text{CaSO}_3$  exceeds 5% (mass) it is considered inadequate to be sold as pozzolanic or cementitious material. The Brazilian ABNT standard (ABNT, 2015) is similar to the American ASTM Standard C618 (ASTM, 2015).

In this context of expanding energy demand, continued share of coal among energy supply sources, intensification of legislation to enforce  $\text{SO}_2$  emissions and the social, economic and environmental impacts of landfill as destination to semi-dry FGD waste, this study evaluates the effect on the levelized cost of electricity (LCOE) posed by using a treatment unit. Specifically, the work approaches a new technology which employs fluidized bed reactor

(FBR) to convert the  $\text{CaSO}_3$  present in the semi-dry FGD waste by high temperature dry oxidation, resulting in residue compliance with the class C pozzolanic material standard. Additionally, a comparison is presented of the LCOE with FGD solids management and the LCOE of the original project, i.e., landfill destination.

The LCOE is recognized worldwide as one of the most adequate methodology to compare and evaluate the economic competitiveness of different electricity generation technologies. (Tolmasquim, 2016) The LCOE gives the cost of the energy that is generated over the lifetime of a given power plant per unit of energy produced. In a simple manner, it is calculated by dividing the total annualized cost of the power plant by the total annual energy generated in the same period (Santoyo-Castelazo and Azapagic, 2014; Short et al., 1995). The total annualized cost is based on a levelized average lifetime cost approach, using the discounted cash flow (DCF) methodology. Several general, local and technology specific assumptions must be considered for various technical and economic parameters. Costs are calculated at the plant level (busbar), and do not include transmission and distribution costs, nor considers other systemic costs or externalities beyond  $\text{CO}_2$  emissions (IEA and NEA, 2015). A detailed procedure describing how to calculate the LCOE is supplied by NREL (Short et al., 1995) and IEA (2016). To standardize and simplify LCOE calculation, several computational tools are available. For coal-fired power plant, the IECM (Berkenpas and Grol, 2009) by Rochedo et al. (2016).

Although the semi-dry FGD waste treatment solves the solid waste problem, it demands CAPEX for building the solid waste treatment unit (FBR and auxiliary equipment), Operation and maintenance (O&M) fixed and variable costs, and consumes energy (the air used to oxidize  $\text{CaSO}_3$  must be slightly compressed (by a blower) and sometimes heated above 500 °C). Although this costs increases the LCOE, it allows revenues (pozzolanic/cementitious material and reduction of landfill related costs), which could render it profitable.

Hence, the main objective of this study is determining the impact of the semi-dry FGD waste treatment unit on the LCOE of a coal-fired power plant. The work is organized in four sections. In Section 2, process premises and methods of calculating LCOE are presented. Results and discussion follows in Section 3 and main conclusions are given in Section 4. Supplementary material is available on line with results from Aspen Process Economic Analyzer (Aspentech Inc) and details for FBR sizing.

## 5.2. Process Premises and Methods

The block diagram of Fig. 5.2 presents the calculation procedure used in this study.

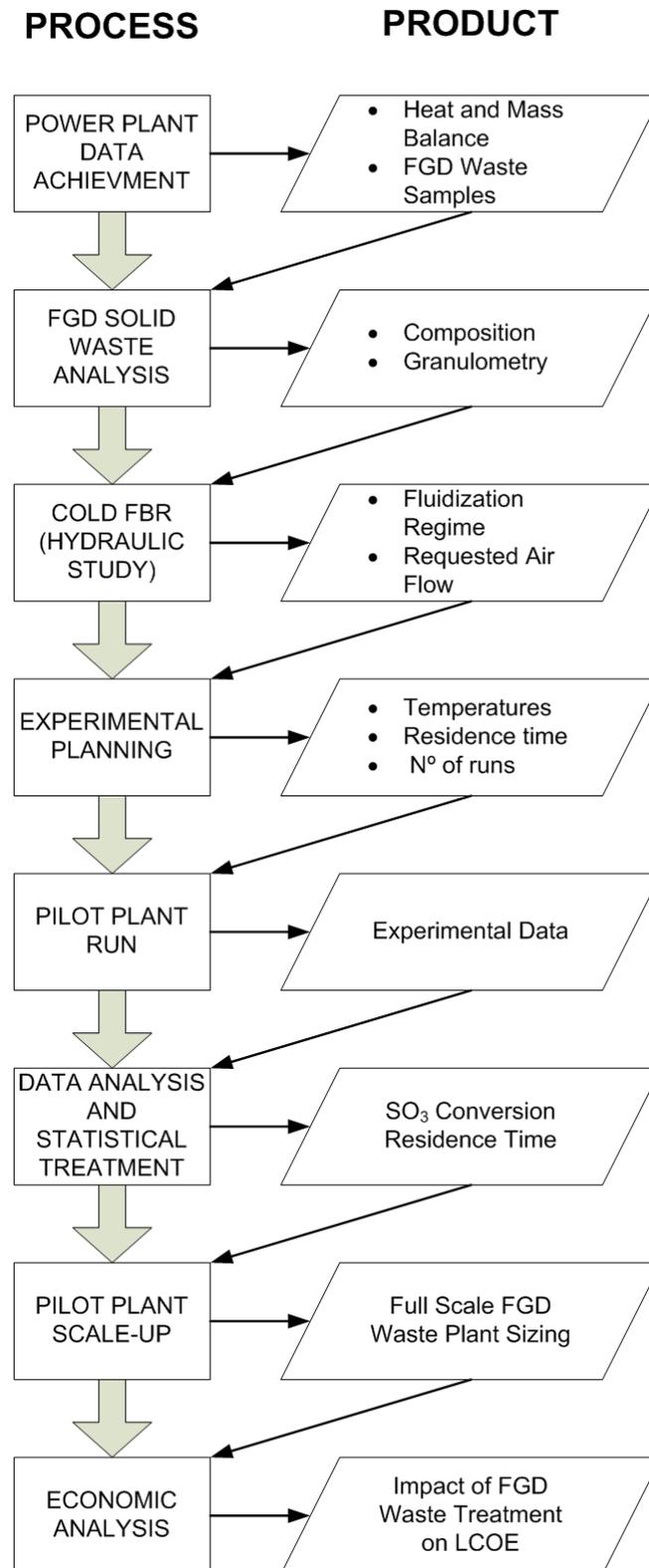


Fig. 5.2 block diagram of calculation procedure

The work is based on a 360 MW coal-fired power plant. This is a conventional pulverized coal combustion (PCC) plant, located in the Brazilian northeast. Coal from Colombia is supplied to the power plant. It is transported by ship and sent from the port terminal to the coal stock by a mechanical belt.

The proposed semi-dry FGD waste treatment unit is based on a pilot-plant, designed and constructed by the authors. It is composed of a fluidized bed reactor (FBR), where occurs oxidation of  $\text{CaSO}_3$ , an air compressor, a heater, a cyclone, an economizer and an air filter. The cyclone collect particles above  $10\ \mu\text{m}$  back to the FBR. The economizer recovers part of the heat of the exhaust air from the FBR. The air filter avoids emission of small particles, below  $10\ \mu\text{m}$ , to the atmosphere. The pilot-plant FBR has a diameter of 200 mm and 1100 mm of height. Depending on temperature and residence time, conversions of  $\text{CaSO}_3$  in  $\text{CaSO}_4$  up to 90% is reached. Fig. 5.3 presents a diagram of the pilot-plant.

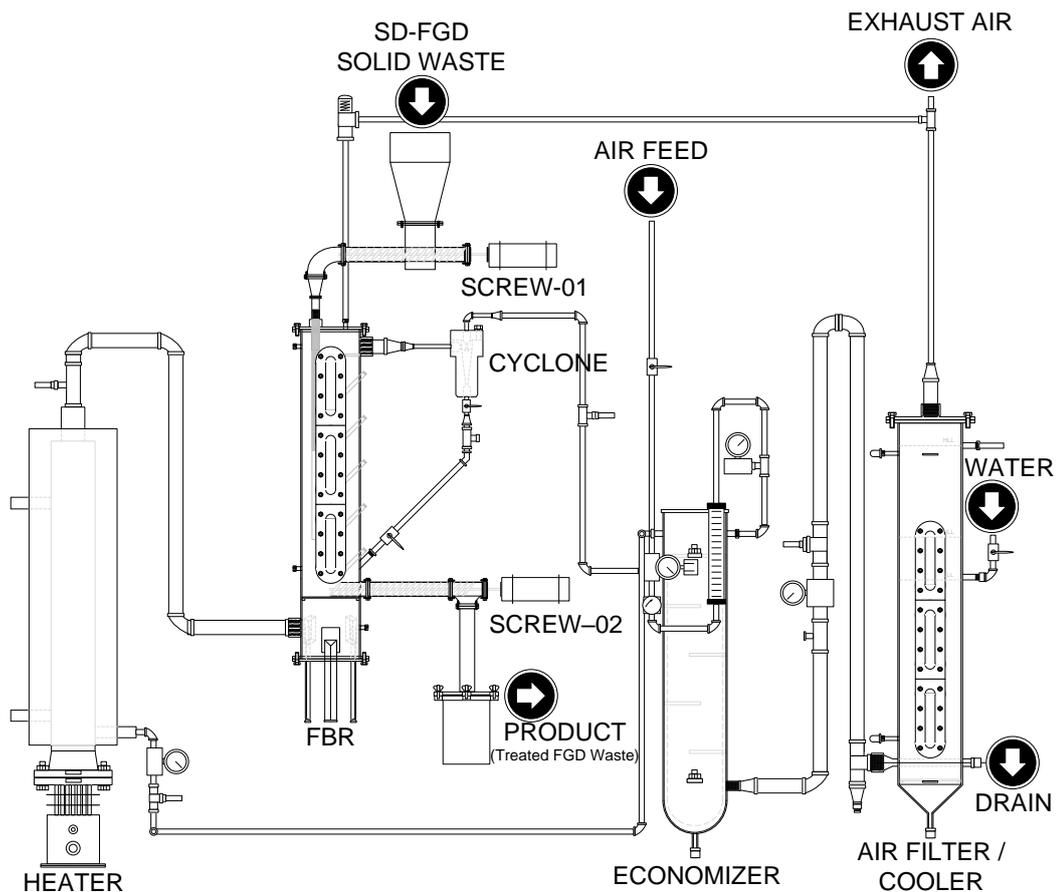


Fig. 5.3. Diagram of the semi-dry FGD waste treatment pilot-plant

A simplified process flow diagram of the power plant and the semi-dry FGD waste treatment unit was developed and is presented in Fig. 5.4.

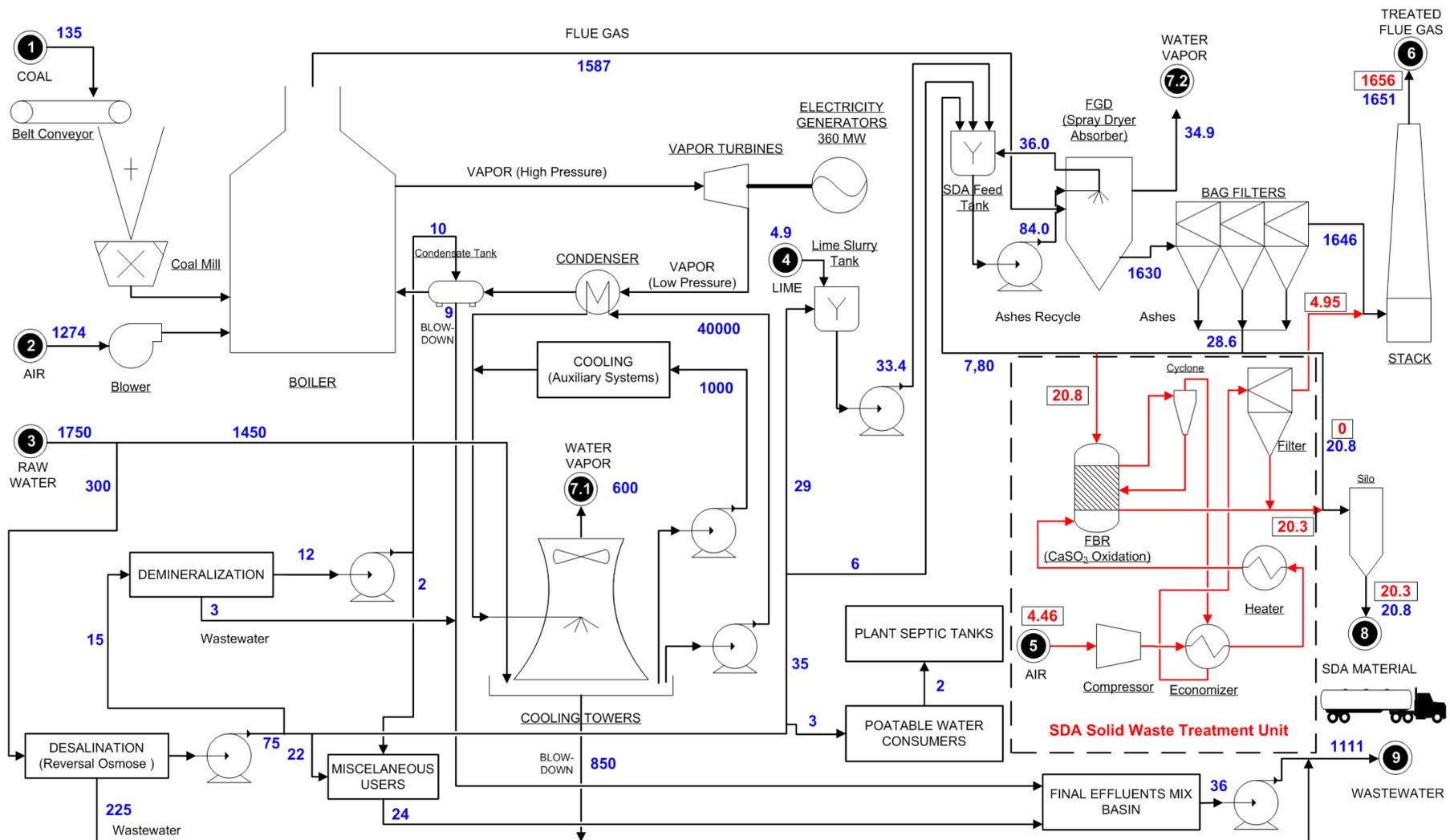


Fig. 5.4. Process flow diagram of the coal-fired power plant and the semi-dry FGD waste treatment unit (indicated with a dashed box). Numbers along stream arrows indicate original mass balance, while numbers within boxes correspond to mass balance of modified power plant (with solid waste treatment).

### 5.2.1. Power Plant

The global mass balance of the power plant follows the original layout (without treatment of the semi-dry FGD waste) of the reference power plant. The main process streams are represented in the simplified process flow diagram presented on Fig. 5.4, where mass flows are expressed in tons per hour. Original power plant streams mass flow (t/h) represented by numbers along stream arrows, and the semi-dry FGD waste treatment unit with numbers inside boxes.

To perform the heat balance and validate the mass balance of the reference PCC plant, the software IECM v 9.2.1 (Berkenpas; Grol, 2009) is adopted. Main set parameters are based on the power plant's data book and environmental impacts assessment (EIA) study (DEEPL, 2008). Fig. 5.5 presents the plant representation on IECM interface.

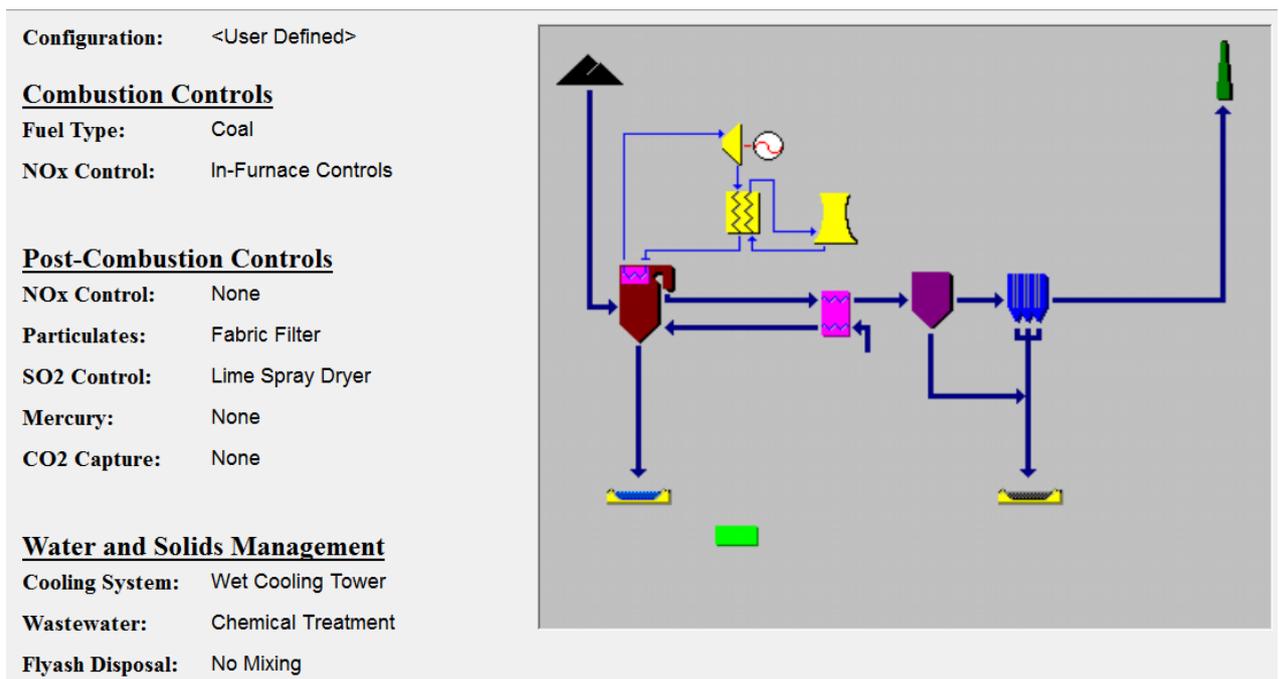


Fig. 5.5. PCC Base Plant Design. Source: IECM v 9.2.1 Interface.

### 5.2.2. Semi-dry FGD waste Treatment Unit

Basically, all information to design auxiliary equipment of the semi-dry FGD waste treatment unit depends on the determination of the inlet and outlet FBR air flow (Section C1 of Supplementary Materials C). The FBR heat and mass balances follow information presented

by Jons et al. (1987) and in data acquired from the pilot-plant experiments (not reported). FBR inlet and outlet air streams (standard and actual volumetric flows) of the full-scale plant are determined using the process simulator PRO-II v9.3 (Schneider Electric SimSci), as presented in Fig. 5.6. Compressor power and heaters duty are also determined with the simulator. The economizer is designed using the Aspen Exchanger Design and Rating v8.8 (Aspentech Inc).

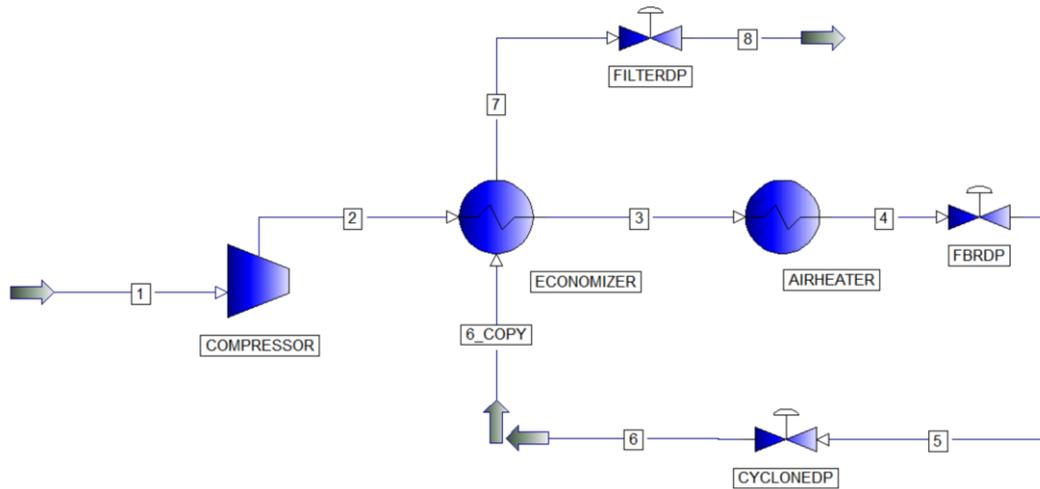


Fig. 5.6. Simulation diagram of the pilot fluidization plant (PRO-II). Valves with DP after equipment names represents the pressure drop of the referred equipment.

### 5.2.3. Levelized Cost of Energy

The LCOE of the coal-fired power plant is calculated using the software IECM v 9.2.1 (Berkenpas; Grol, 2009), while the LCOE for the semi-dry FGD waste treatment unit is based on main equipment economic analysis, performed in Aspen Process Economic Analyzer v8.8 (APEA, Aspentech Inc) (results are presented in Section C2 of the Supplementary Materials C. APEA can estimate CAPEX and O&M of a given industrial process, based on main equipment specification. Tables C2.1 and C2.2, from Supplementary Materials C, present the equipment specifications and main economic parameters assumed in APEA for CAPEX, O&M and LCOE evaluation of the semi-dry FGD waste treatment unit.

### 5.2.4. Semi-dry FGD waste Treatment Unit Heat and Mass Balance

The set of reactions expected to occur inside the FBR is presented in Table 5.1 (Jons et al., 1987).

Table 5.1. FBR set of reactions.

Reaction	$\Delta H_r$ (kcal/mol)	Temperature Range (°C)
R <sub>1</sub> - $\text{CaSO}_3 \cdot \frac{1}{2} \text{H}_2\text{O} \rightarrow \text{CaSO}_3 + \frac{1}{2} \text{H}_2\text{O}$	6.87	370-390
R <sub>2</sub> - $\text{CaSO}_4 \cdot 2 \text{H}_2\text{O} \rightarrow \text{CaSO}_4 + 2 \text{H}_2\text{O}$	26.07	130-150
R <sub>3</sub> - $\text{Ca}(\text{OH})_2 \rightarrow \text{CaO} + \text{H}_2\text{O}$	25.7	> 470
R <sub>4</sub> - $\text{CaSO}_3 + \frac{1}{2} \text{O}_2 \rightarrow \text{CaSO}_4$	-65.8	> 550

The semi-dry FGD waste composition and mass flow, shown in Table 5.2, is obtained from the heat and mass balance of the reference power plant. This is the worst expected case, as it considers coal with 1.5% of sulphur, on a weight basis. When the coal sulphur content is lower, SO<sub>2</sub> content in the fly-ash decreases.

Table 5.2. Composition of the semi-dry FGD waste – worst case, coal with 1.5% of sulfur.

<b>Mass Flow (t/h)</b>	20.78
<b>Composition (mass %)</b>	
CaSO <sub>3</sub> ·½H <sub>2</sub> O	38.40%
CaSO <sub>4</sub> ·2H <sub>2</sub> O	0.00%
Ca(OH) <sub>2</sub>	8.30%
CaSO <sub>3</sub>	0.00%
CaSO <sub>4</sub>	0.00%
CaO	0.00%
H <sub>2</sub> O	2.00%
Fly-ashes (inert)	51.30%

Based on Angevine et al. (1985), considering a temperature of 550 °C and 5% of O<sub>2</sub> in excess, it is possible to achieve a SO<sub>3</sub> mass composition of 3.2% on the treated residue, complying with ASTM Standard C618 (ASTM, 2015) for class C or F fly ash, that establishes a limit of 5% (mass) of SO<sub>3</sub> for this type of cement additive. A conversion of 91.3% is considered for reaction R<sub>4</sub> and 100% for reactions R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> (Table 5.1). These results are considered in performing

the heat and mass balance of the semi-dry FGD waste treatment unit. The semi-dry FGD waste heat capacity used to estimate the energy consumption due to the heating of the residue from 80 °C to 550 °C is considered 730 J/kg.K, the same value of a class C fly ash (Bentz et al., 2011). The inlet and outlet air flow and composition is calculated in function of the FBR demand (reactions R<sub>1</sub> to R<sub>4</sub> of Table 5.1).

The detailed calculation is presented in Table C1.2, Section C1 of Supplementary Materials C. The air is pre-heated to 350 °C by the economizer. The heater is designed to complete the service, rising the air temperature to the set-point (550 °C). It is worth noting that the energy supplied by the oxidation of CaSO<sub>3</sub> and integration with steam purges of the power plant could bring the air heating energy input to zero. Thus, the air heater could be used only for the start-up of the system.

## 5.2.5. Equipment Scale-up and Sizing

### 5.2.5.1. Fluidized Bed Reactor

As the FBR is not available in industrial scale, determination of the fluidization air flow needs to be estimated. In the pilot scale, where FBR deals with a 10% w/w CaSO<sub>3</sub>.½H<sub>2</sub>O feed, the ratio of fluidization air flow to stoichiometric air flow is 3.75. Since in this study the CaSO<sub>3</sub> mass fraction is 38.4%, the stoichiometry air flow with 5% of oxygen (O<sub>2</sub>) excess is considered sufficient to promote bed fluidization. The length/diameter (*L/D*) ratio of the full scale FBR is considered the same of the pilot scale equipment (*L/D*=5), as well as the maximum level of solids inside the reactor (30% of *L*). The maximum residence time of the semi-dry FGD waste is considered 500 seconds, based in Jons et al. (1987).

### 5.2.5.2. Air Compressor

A fan, air blower or compressor is necessary to make the air stream passes through the economizer, air heater, FBR, cyclones and air filter, reaching the end of the process with a pressure slightly higher than atmospheric, with estimated pressure drop of 1.5 bar. As the air inlet is nearly at atmospheric pressure, the necessary absolute outlet pressure of the compressor is 250 kPa, and the pressure ratio is 2.47. This range of pressure ratio turns the use of fans and common blowers inappropriate, being a compressor more adequate for the service.

The power of centrifugal compressor is calculated using Eq. 5.1 (GPSA, 2004).

$$Pw = \frac{q \cdot Z \cdot R \cdot T \cdot \left[ (r_p)^{\frac{n-1}{n}} - 1 \right]}{3.600 \cdot \eta_p \cdot MM \cdot \frac{n-1}{n}} \quad (5.1)$$

where  $Pw$  is the brake horsepower (kW),  $q$  is the gas flow rate (kg/h),  $Z$  is the average compressibility factor,  $R$  is the gas constant (8.314 kJ/kmol.K),  $T$  is the gas inlet temperature (K),  $MM$  is the molecular mass (kmol/kg),  $r_p$  is the pressure ratio,  $n$  is the polytrophic exponent and  $\eta_p$  is the polytrophic efficiency. The pressure ratio is calculated dividing the outlet pressure,  $P_2$  (kPa), by the inlet pressure  $P_1$  (kPa).  $P_2$  is considered 250 kPa (gauge).  $P_1$  is the atmospheric pressure (0 kPa gauge).  $\eta_p$  is considered 80%,  $q$  of inlet air stream (stream 1 of Fig. 5.5), 4460 kg/h, is calculated in function of the FBR demand (see Table C1.2 of Section C1 of Supplementary Materials C).

### 5.2.5.3. Air Heater

Although it is expected that the energy supplied by the oxidation of  $CaSO_3$  and integration with steam purges from the closed loop steam cycle could heat the air inside the reactor to the required level (550 °C), as by the reaction  $R_4$  (see Table 5.1), the air heater is necessary to start-up the system and to heat the air when the  $SO_3$  content in the semi-dry FGD waste is lower than expected, in function of the use of a coal with less sulphur. In this case, the oxidation reaction could not be sufficient to supply the necessary energy input and the air heater must have capacity to reach the reaction  $R_4$  temperature set-point. PRO-II is used to calculate the heater duty, to heat the air from the compressor discharge temperature (~130 °C) to 550 °C. It was assumed that during the start-up no heat is recovered by the economizer. A maximum pressure drop of 30 kPa is considered in the process side of the heater.

### 5.2.5.4. Economizer

The economizer recovers part of the heat of the exhaust air from the FBR to pre-heat the air coming from the compressor to 350 °C. Inlet and outlet streams temperatures and pressures are obtained from the simulation. A vertical BEU TEMA type shell-and-tube heat exchanger is considered. The design of the equipment is developed on Aspen Heat Exchanger Design and

Rating software (Aspentech Inc). A maximum pressure drop of 30 kPa is considered on shell and tubes sides.

#### 5.2.5.5. Cyclone and Air Filter

A preliminary sizing of both equipment is not required, as the economic analysis tool (APEA) demands only the air flow to estimate costs, obtained from process simulation.

#### 5.2.6. Semi-dry FGD Waste Treatment Unit Levelized Cost of Energy

Traditionally, the LCOE can be calculated by Eq. 5.2 (Santoyo-Castelazo and Azapagic, 2014).

$$LCOE = TAC/AE \quad (5.2)$$

where, *LCOE* is the levelized or unit cost of energy (\$/MWh), *TAC* is the total annualised cost of generating electricity (\$/yr) and *AE* is the annual energy generation (MWh/yr). *TAC* is the sum of annualized capital costs (\$/yr), fixed costs (\$/yr), variable costs (\$/yr) and fuel costs (\$/yr). However, sale of treated semi-dry FGD waste could be deducted from the annual variable cost parcel of *TAC*. The annualised capital costs are calculated multiplying the total capital cost by an annuity factor (*f*), that is calculated according to Eq. 5.3 (Santoyo-Castelazo and Azapagic, 2014).

$$f = z(1 + z) e^t / [(1 + z)e^t - 1] \quad (5.3)$$

where, *z* is the discount rate (%) and *t* is the lifetime of the plant (yr). The discount rate measures the time value of the money, it is used to calculate the present value of a cash flow and, often, to account the inherent risk of an investment.

The proper selection of the discount rate is very important to any economic analysis and depends on several factors, as rate of return, risk premium, planning horizon, interest rates and taxes. Therefore, discount rates vary from place to place, industry to industry and company to company (Short et al., 1995). In the present study, a nominal discount rate of 8% is assumed, based on Tomalsquim (2016). A lifetime of 30 years is considered for the power plant to calculate the LCOE of the semi-dry FGD waste treatment unit. Furthermore, this study considers that the treated residue can be used as cement kiln raw material, assumed as 47 \$/t (an approximate value in Brazil).

Tables C2.1 and C2.2, Section C2 of Supplementary Materials C, present the equipment specifications and main economic parameters assumed on APEA to perform CAPEX, O&M and LCOE evaluation of the semi-dry FGD waste treatment unit.

### 5.2.7. Levelized Cost of Energy of the Coal-Fired Power Plant

Tables 5.3 and 5.4 summarize the main parameters input for IECM tool, adapted to reproduce the reference power plant heat and mass balances and performance.

Table 5.3. IECM input parameters.

<b>Set Parameters</b>	<b>Value</b>	<b>Reference</b>
Capacity Factor (%)	85	(Tolmasquim, 2016)
Ambient Air Dry Bulb Temperature (°C)	27	ENEVA
Ambient Air Pressure (MPa)	0.1013	Plant at sea level
Relative Humidity (%)	80.84	ENEVA
Discount Rate (%)	8.0	(Tolmasquim, 2016)
Plant or Project Book Life (yr)	30	
Federal Tax Rate (%)	15	Minimum IECM value
State Tax Rate (%)	2	(DEEPL, 2008)
Property Tax rate (%)	0	Government incentive program
Internal Cost of Electricity (COE)	Total Plant COE	
O&M Escalation Rate (%/yr)	3.5	APEA
Coal Cost (\$/ton)	90	(Tolmasquim, 2016)
Gross Electrical Output (MW)	373.8	ENEVA
Steam cycle type	Sub-Critical	(DEEPL, 2008)
Boiler Efficiency (%)	90	(DEEPL, 2008)
Gas Temperature Exiting Economizer (°C)	307	ENEVA
Gas Temperature Exiting Air Preheater (°C)	118	ENEVA
Miscellaneous (%MW)	5	Set to reach reference heat rate (10097 kJ/kWh)
Construction time (yr)	4	(Tolmasquim, 2016)
Royalties Fees (%)	0	Imported Coal, royalty free.
Number of Operating Jobs	80	(DEEPL, 2008)
Maximum SO <sub>2</sub> removal (%)	95	(DEEPL, 2008)
Reagent Stoichiometry (molCa/molS in)	1.27	Set to reach the reference CaO consumption (4.9 t/h)
Ca content of lime (%)	90	(DEEPL, 2008)
SDA Power Requirement (% gross MW)	0.38	Set to reach the reference value (1.4 MW)
Cooling Water Inlet Temperature (°C)	39	ENEVA
Tower overdesign factor (% total load)	20	Set to reach the reference cooling water flow (~ 40000 t/h)
Tower Power Requirement (% gross MW)	0.719	Set to reach reference (2.688 MW)

Table 5.4. IECM Customized Fuel and Ash Properties.

Fuel		Ash	
Higher Heating Value (kJ/kg)	25289	Composition (wt%)	
Composition (wt%)		SiO <sub>2</sub>	59.00
Carbon	60.66	Al <sub>2</sub> O <sub>3</sub>	22.00
Hydrogen	4.40	Fe <sub>2</sub> O <sub>3</sub>	8.00
Oxygen	9.38	CaO	2.50
Chlorine	0.03	MgO	1.50
Sulfur	1.50	Na <sub>2</sub> O	0.70
Nitrogen	1.23	K <sub>2</sub> O	1.80
Ash	10.80	TiO <sub>2</sub>	1.10
Moisture	12.00	P <sub>2</sub> O <sub>5</sub>	0.30
Total	100.00	SO <sub>3</sub>	2.50
Default Cost (\$/t)	90.00	MnO <sub>2</sub>	0.00
		Other	0.60
		Total	100.00

Source: (DEEPL, 2008)

### 5.3. Results and Discussion

#### 5.3.1. Semi-dry FGD waste Treatment Unit Heat and Mass Balance

Based on information from Tables 5.1, 5.2 and fly ash heat capacity (730 J/kg.K), the heat balance of the FBR is performed.

The inlet air and semi-dry FGD waste heating demands are 248 kW and 1980 kW, respectively. Reaction R<sub>1</sub> demands 494 kW, reaction R<sub>2</sub> does not occur with the considered semi-dry FGD waste composition, reaction R<sub>3</sub> demands 697 kW and reaction R<sub>4</sub> (exothermic) liberates 4260 kW. The final heat balance gives -841 kW. Hence, considering all stated premises, the reaction could be auto sufficient in terms of energy. Additional energy is necessary only to start up the FBR (diesel) and to supply the compressor (electricity).

The semi-dry FGD waste production is estimated in 20.3 t/h. The treated residue has final composition of CaSO<sub>3</sub> of 3.66%, complying with the ASTM standard (< 5.0%). The FBR air demand is estimated in 4,460.0 kg/h and the calculated outlet air flow as 4,491.0 kg/h (6.9 Sm<sup>3</sup>/h). Detailed information on the calculation of mass flow and composition of the treated

semi-dry FGD waste is presented in Supplement C (Table C2.1) as well as the inlet and outlet air mass flows, which promote the oxidation of the CaSO<sub>3</sub> inside the FBR. (Table C2.2).

### 5.3.2. Semi-dry FGD Waste Treatment Unit Economic Analysis

Considering the main equipment specification of Table C2.1 (Section C2 of Supplementary Materials C), CAPEX and OPEX of the semi-dry FGD waste treatment unit are estimated. The main equipment cost and weight are individually presented in Table C2.3, Section C2 of Supplementary Materials C. The final economic analysis (CAPEX and OPEX) is shown in Table 5.5.

Table 5.5. Semi-dry FGD waste Treatment Unit CAPEX and OPEX Analysis Summary.

<u>Item</u>	<u>Value</u>
Total Project Capital Cost (\$)	6,804,026.36
Total Products Sales (\$/yr)	7,108,999.10
Total Operating Labor and Maintenance Cost (\$/yr)	573,261.31
Total Utilities Cost (\$/yr)	112,257.85
Total Operating Cost (\$/yr)	1,049,948.80
Operating Labor Cost (\$/yr)	558,825.00
Maintenance Cost (\$/yr)	14,436.31
Operating Charges (\$/yr)	25
Plant Overhead (\$/yr)	286,630.66
Subtotal Operating Cost (\$/yr)	972,174.82
General and Administration Cost (\$/yr)	77,773.99

From Table 5.5, it can be noticed that, in 1 year of operation, the semi-dry FGD waste revenue could pay all the capital investment on its treatment unit. Details on the treated semi-dry FGD waste sales are presented in Table 5.6.

Table 5.6. SDA Solids Commercialization.

Products Sales per Hour (\$/h)	954.1
Product Name	Ash + SDA Residue
Product Rate (t/h)	20.3
Product Unit Cost (\$/t)*	47
Product Rate per Period (t/yr)	151,255.30
Product Sales (\$/yr)	7,108,999.10

\*Considering sale as cement kiln raw material in Brazil.

With data provided by APEA, it is possible to calculate LCOE of the semi-dry FGD waste treatment unit, as shown in Table 5.7. It is worth noting that the impact of the treatment unit on the power plant LCOE depends on the revenue price of the treated residue.

Table 5.7. SDA Solids Treatment LCOE.

Discount Rate	8%
<i>f</i>	8.88%
Annualized Capital Cost (\$/yr)	604,384.20
O&M Fixed Cost (\$/yr)	573,261.31
O&M Variable Cost (\$/yr)	-6,996,741.25
Fuel Cost (\$/yr)	0.00
TAC (\$/yr)	-5,819,095.74
TAC w/o product sales (\$/yr)	1,289,903.36
Capacity Factor	85%
AE (MWh/yr)	2,535,000
LCOE without product sales (\$/MWh)	0.51
LCOE with product sales (\$/MWh)	-2.30

### 5.3.3. IECM Results for the Coal Fired Power Plant

Considering the input parameters of Tables 5.3 and 5.4, IECM produces a comprehensive set of results. For the intended analysis, a realistic coal and calcium oxide (CaO) consumption is fundamental, to produce the informed flow of semi-dry FGD waste (20.8 t/h). The mass balance results are presented in Figs. 5.7 and 5.8. Other important parameter is the net plant heat rate, that reflects the final net efficiency of the plant.

Results demonstrate that the SO<sub>2</sub> control (semi-dry FGD), without the solid treatment unit, has total LCOE of 19.56 \$/MWh, where 0.53 \$/MWh (2.7%) corresponds to the semi-dry FGD waste disposal. Thus, the total plant LCOE is increased in 0.56% (from 94.44 to 94.97 \$/MWh) resulting from disposal of the semi-dry FGD waste.

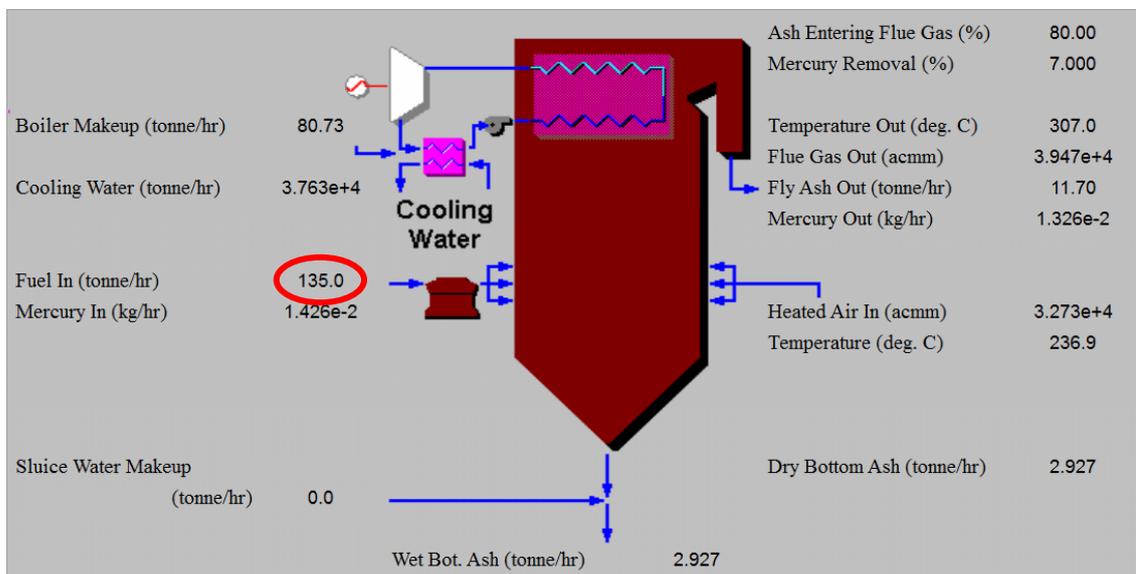


Fig. 5.7. Boiler Diagram. Source: IECM v 9.2.1 Interface.

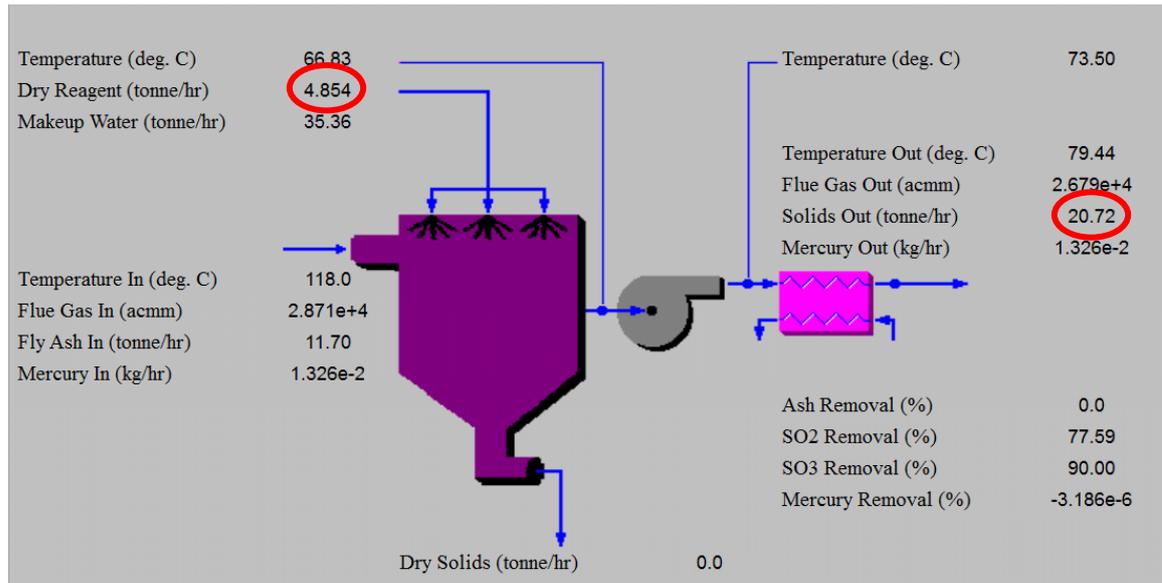


Fig. 5.8. Semi-dry FGD Diagram. Source: IECM v 9.2.1 Interface.

To validate the consistence and accuracy of the IECM results, key performance parameters are compared with values reported in the literature and reference power plant information, presented in Table 5.8, which shows that all key performance parameters are in strong agreement with the literature and reference power plant data. This accuracy is relevant to reach a realistic LCOE result. IECM calculates LCOE, or revenue required from electricity price, based on the total O&M and annualized capital costs, presented in Table 5.9.

Table 5.8. Overall PCC Plant Performance Validation.

Performance Parameter	Result	Reference Value*	Deviation (%)
Coal Consumption (ton/h)	135	135	0.00%
Primary Fuel Input (MW)	948.06	942.35	0.61%
Gross Electrical Output (MW)	373.8	373.8	0.00%
Plant Electricity Requirements			
Base Plant Use (MW)	28.6		
In-Furnace NO <sub>x</sub> Use (MW)	0		
Fabric Filter Use (MW)	1.152		
Spray Dryer Use (MW)	1.140	1.14	0.00%
Cooling Tower Use (MW)	2.688	2.688	0.00%
Wastewater Plant Use (MW)	0.016		
Net Electrical Output (MW)	340.2	336	1.25%
Annual Power Generation (kWh/yr)	2.535x10 <sup>9</sup>		
Gross Plant Heat Rate, HHV (kJ/kWh)	9133		
Net Plant Heat Rate, HHV (kJ/kWh)	10030	10097	-0.66%
Net Plant Efficiency, HHV (%)	35.82	35.8	0.06%
CaO Consumption (t/h)	4.85	4.9	-0.94%
Ash Production	20.72	20.8	-0.38%

\*Reference values from the power plant data (ENEVA) and Environmental Impacts Assessment Study (DEEPL, 2008).

Table 5.9. PCC Plant Total Cost.

<i>Technology</i>	<i>Fixed O&amp;M (M\$/yr)</i>	<i>Variable O&amp;M (M\$/yr)</i>	<i>Total O&amp;M (M\$/yr)</i>	<i>Annualized Capital (M\$/yr)</i>	<i>Levelized Annual Cost (M\$/yr)</i>
Combustion NOx Control	0.1181	0	0.1181	0.4622	0.5803
TSP Control	1.552	3.21	4.762	2.584	7.347
SO <sub>2</sub> Control	7.452	6.862	14.31	5.248	19.56
Subtotal	9.122	10.07	19.19	8.295	27.49
Cooling Tower	2.032	5.061	7.092	3.909	11
Wastewater Control	0.776	0.3172	1.093	0.96	2.053
Base Plant	63.63	87.65	151.3	48.86	200.1
Land	0	0	0	5.02E-02	5.02E-02
<b>Total</b>	<b>75.56</b>	<b>103.1</b>	<b>178.7</b>	<b>62.08</b>	<b>240.7</b>

Based on Table 5.9, information, IECM calculates total capital required, capital required per net kW, annual revenue required and, the objective of this study, LCOE (\$/MWh), as shown in Table 5.10.

Table 5.10. PCC plant cost summary.

<i>Technology</i>	<i>Capital Required (M\$)</i>	<i>Capital Required (\$/kW-net)</i>	<i>Revenue Required (M\$/yr)</i>	<i>Revenue Required (\$/MWh)</i>
Combustion NOx Control	5.344	15.71	0.5803	0.2289
Post-Combustion NOx Control	0	0	0	0
Mercury Control	0	0	0	0
TSP Control	29.88	87.82	7.347	2.898
SO <sub>2</sub> Control	60.67	178.3	19.56	7.717
Combined SOx/NOx Control	0	0	0	0
CO <sub>2</sub> Control	0	0	0	0
Subtotal	95.89	281.9	27.49	10.84
Cooling Tower	45.19	132.8	11	4.34
Wastewater Control	11.1	32.62	2.053	0.8099
Base Plant	564.9	1660	200.1	78.95
Land	0.5798	1.704	5.02E-02	1.98E-02
Emission Taxes	0	0	0	0
<b>Total</b>	<b>717.7</b>	<b>2109</b>	<b>240.7</b>	<b>94.97</b>

The final LCOE is 94.97 \$/MWh. This value strongly depends on the considered discount rate. For a discount rate of 7%, closer to the one assumed in this study (8%), the International Energy Agency (IEA and NEA, 2015) presents a range of 75 – 110 \$/MWh for the LCOE of

coal-fired power plant. Thus, it can be considered that the LCOE calculated using IECM is in good agreement with the international values, allowing to determine the impact of the semi-dry FGD waste treatment unit on LCOE, as presented in Table 5.11.

Table 5.11. Final LCOE Comparison for Base PCC and PCC + SDA Solid Treatment Plants

<i>Case</i>	<i>PCC</i>	<i>PCC + Solids Treatment</i>
Air Compressor Power (MW)	-	0.135
Net Electrical Power Output (MW)	340.200	340.065
Plant Efficiency	35.88%	35.87%
Reference Plant LCOE (\$/MWh)	94.97	94.97
FGD waste disposal cost (\$/MWh)	0.53	0.00
LCOE with FGD waste sales (\$/MWh)		92.14
LCOE reduction with FGD waste sales		3.0%
LCOE without FGD waste sales (\$/MWh)		94.95
LCOE reduction without FGD waste sales (\$/MWh)		0.02%

As shown in Table 5.11, there is no need of a heater for operating the semi-dry FGD waste treatment unit, because the exothermic oxidation of  $\text{CaSO}_3$  ( $R_4$ ) can supply all the energy required to heat air and solids flow and promote the endothermic reactions ( $R_1$ ,  $R_2$ , and  $R_3$ ). The only energy requirement is from the air compressor power demand (0.135 MW), merely 0.04% of the PCC plant net power output (340 MW). Thus, the only factors that considerably impact LCOE are CAPEX and O&M costs of the semi-dry FGD waste treatment unit, and the revenue from treated residue price.

If the treated semi-dry FGD waste was transferred (at zero cost and null revenue) as additive to a cement industry, LCOE of the power plant would remain approximately the same, because the increase of 0.51\$/MW resulting from CAPEX and O&M costs of the treatment unit is compensated by the decrease of 0.53\$/MWh, in virtue of the avoided semi-dry FGD waste disposal costs. However, if commercialization as raw material of the treated semi-dry FGD waste (for cement kiln) is considered, there is a reduction of 3% (2.83 \$/MWh) on the power plant LCOE (to 92.14 \$/MWh).

It is worth noting that that regional economic scenarios may change the impact of semi-dry FGD waste treatment on LCOE. For instance, Liu et al. (2016) surveyed 7 coal fired-power plants in China to collect detailed field data to examine the costs and benefits of flue-gas desulfurization. They state that a PCC plant in China which installs and properly operate FGD equipment can receive 15 Yuan/MWh (~2.89 \$/MWh) premium tariff on top of their on-grid

tariffs. However, their study shows that this incentive is insufficient to cover FGD costs of most of the sample plants surveyed. Liu et al. (2016) propose that, to cope with the penalty on LCOE, in China's scenario of dispatch regulation, allocating generation hour based on the pollutant emission would provide strong incentive for sulfur dioxide mitigation.

Lastly, coal fired generation with CCS has estimated LCOE of 139.5 \$/MWh (EIA, 2017), a penalty of 44.5 \$/MWh, considering the reference power plant (without solid waste treatment) of the present work (~95 \$/MWh). Comparatively, semi-dry-FGD has positive impact in LCOE, and the proposed semi-dry FGD waste treatment would turn a penalty of 0.53\$/MWh (because of semi-dry FGD waste disposal cost) in an extra revenue (as LCOE is reduced in 2.83 \$/MWh).

Aiming to show the impact of variations on treated semi-dry FGD waste revenue price, a sensitivity analysis of LCOE, based on variation of final product price between 0 and 100 \$/ton is presented in Fig. 5.9. Remembering that, the base price adopted in the present study was 47 \$/ton (points with black board on Fig. 5.9). The detailed calculation is presented on supplementary material Tables C3.1 e C3.2, Section C3 of Supplementary Materials C.

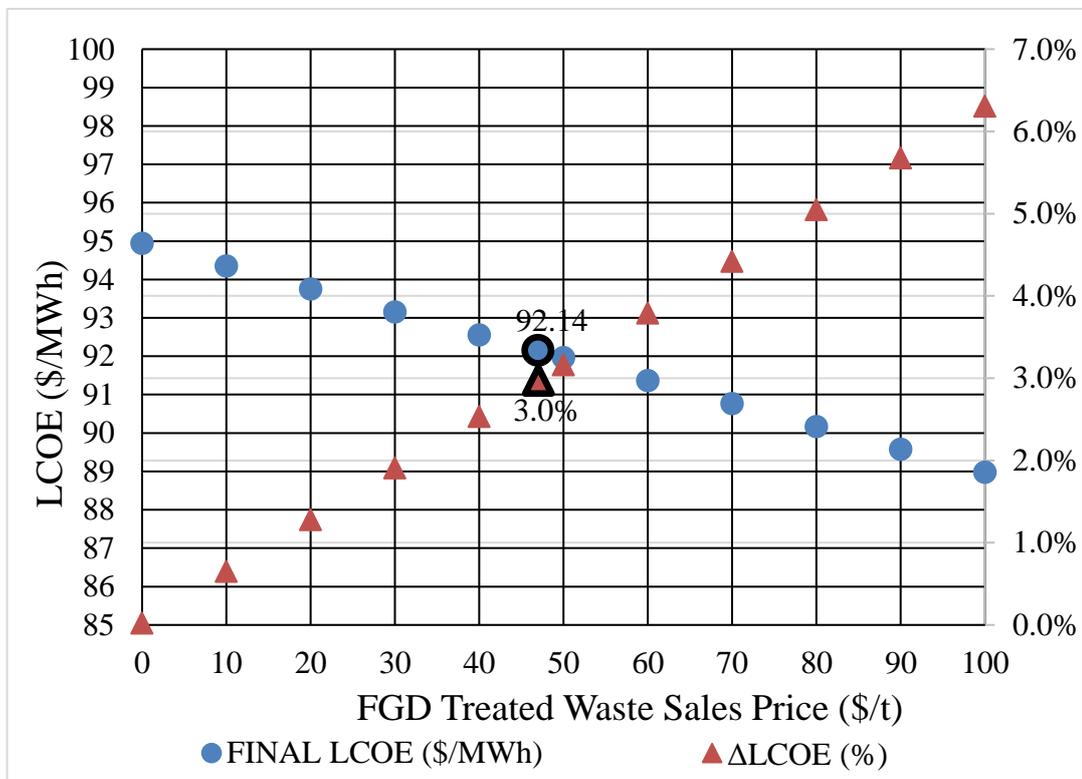


Fig. 5.9. Sensitivity Analysis of LCOE versus Treated Semi-dry FGD Residue Revenue Price

The sensitivity analysis shown a linear variation on the LCOE according to the treated semi-dry FGD residue revenue price. Even if the residue were donated for free there is no impact on LCOE, because of the saving with landfill disposal (0.53 \$/MWh). If the revenue price increases about 100% behind the base value, the impact on LCOE would be more 3.3% of decrease, achieving a final value of 89 \$/MWh, or 6.3% of decrease (based on the reference plant, that is 95 \$/MWh).

## 5.4. Conclusions

This study analyzes the impact on LCOE from installing a semi-dry FGD waste treatment unit in a real PCC power plant (the reference power plant) that is currently facing decision making process on semi-dry FGD waste destination. The calculated LCOE, 94.97 \$/MWh, is consistent with world values for this type of power plant. The analysis shows that the extra energy demanded by the novel semi-dry FGD waste treatment unit is nearly negligible due to the exothermicity of the  $\text{CaSO}_3$  oxidation reaction and the proposed economizer for heating air and solids inlet streams. The power necessary to air compressor operation is only 0.04% of the PCC plant net power output (340 MW). Thus, the only factors that impact LCOE are cost (CAPEX and O&M) of the semi-dry FGD waste treatment unit and the residue revenue price.

Furthermore, this study demonstrates that, if the residue revenue is not considered, the achieved reduction in LCOE is only 0.02% (0.02 \$/MWh). If the residue is commercialized as a raw material for cement kiln, the LCOE could be reduced in ~3% (2.83 \$/MWh) to 92.14 \$/MWh. Therefore, with or without treated semi-dry FGD waste revenue, the proposed treatment unit has a negative, but small, impact on the total power plant LCOE, besides solving the problem of landfill saturation, land use and costs related to landfill maintenance. Thus, it is adequate to implement the semi-dry FGD waste treatment unit on PCC power plants with similar design and financial parameters compared to the reference coal-fired power plant studied in this work.

## 5.5. References of Chapter 5

ACAA. American Coal Ash Association. Beneficial Use of Coal Combustion Products. An American Recycling Success Story, 2014 Available at: <https://www.acaa-usa.org/Portals/9/Files/PDFs/Production-and-Use-Brochure.pdf>. Last access date: 4/7/2016.

Angevine, P.A., Bengtsson, S., Koudijs, G.P. Method for oxidation of flue gas desulfurization absorbent and the product produced thereby. US Patent No. 4544542, 195. ABNT. Associação Brasileira de Normas Técnicas. NBR 12653 - Materiais pozolânicos, 2015. Available at: <http://www.abntcatalogo.com.br/>, Last access date: 4/7/2016.

ASTM. C618-15, Standard Specification for Coal Fly Ash and Raw or Calcined Natural Pozzolan for Use. Annu. B. ASTM Stand, 2015. <http://dx.doi.org/10.1520/C0618-15>.

Bentz, D., Peltz, M., Duran-Herrera, A., Valdez, P., Juarez, C. Thermal properties of high-volume fly ash mortars and concretes. J. Build. Phys. 34, 263–275, 2011. <http://dx.doi.org/10.1177/1744259110376613>.

Berkenpas, M.B. (Carnegie M.U., Grol, E.P.E. (National E.T.L., 2009. Integrated Environmental Control Model User Manual. U.S. Department of Energy, Pittsburgh, PA. Cordoba, P. Status of Flue Gas Desulphurisation (FGD) systems from coal-fired power plants: Overview of the physic-chemical control processes of wet limestone FGDs. Fuel, 144, 274-286, 2015. <http://dx.doi.org/10.1016/j.fuel.2014.12.065>.

Blankinship, S. Looking for a Good Scrubbing: Today's FGD Technology. Power Eng. 109, 9; 19-24, 2005. Available at: <http://www.power-eng.com/articles/print/volume-109/issue-9/features/looking-for-a-good-scrubbing-todayrsquos-fgd-technology.html> Last access date: 03/15/2017.

DEEPL. Diferencial Energia Empreendimentos e Participação Ltda. Estudo de Impacto Ambiental (EIA). Empreendimento da Usina Termoeletrica Porto do Itaqui (UTE Porto do Itaqui), 2008.

EIA. U.S. Energy Information Agency. Levelized Cost and Levelized Avoided Cost of New Generation Resources in the Annual Energy Outlook 2016, August 2016. Available at: [http://www.eia.gov/outlooks/aeo/pdf/electricity\\_generation.pdf](http://www.eia.gov/outlooks/aeo/pdf/electricity_generation.pdf). Last access date: 01/23/2017.

EPE. Empresa de Pesquisa Energética. Balanço Energético Nacional 2015, 2015a. Available at: [https://ben.epe.gov.br/downloads/Relatorio\\_Final\\_BEN\\_2015.pdf](https://ben.epe.gov.br/downloads/Relatorio_Final_BEN_2015.pdf) Last access date: 4/7/2016.

EPE. Empresa Brasileira de Planejamento Energético. Previsões de carga para o Planejamento Anual da Operação Energética 2015 – 2019 1a Revisão Quadrimestral, 2015b. Rio de Janeiro.

EPRI. Electric Power Research Institute. Coal Ash: Its origin, disposal, use and potential health issues. Report from EPRI's Environment Division. Environmental Focus. Palo Alto, CA, 1998. Available at: <http://ecosmartconcrete.com/docs/trepri98.pdf>. Last access date: 01/23/2017.

EPRI. Electric Power Research Institute. Characterization of Spray Dryer Absorber Products for Use in Cement and Concrete Applications 1017580. Palo Alto, CA, 2009.

Farfen, J., Breyer, C. Structural changes of global power generation capacity towards sustainability and the risk of stranded investments supported by a sustainability indicator. Journal of Cleaner Production, 141, 370–384, 2017. <http://dx.doi.org/10.1016/j.jclepro.2016.09.068>.

Feng, C., Gao, X., Tang, Y., Zhang, T. Comparative life cycle environmental assessment of flue gas desulphurization technologies in China. *Journal of Cleaner Production*, 68, 81-92, 2014. <http://dx.doi.org/10.1016/j.jclepro.2013.10.023>.

Galos, K. A., Smakowski, T.S., Szlugaj, J. Flue-gas desulphurisation products from Polish coal-fired power-plants. *Applied Energy*, 75(3-4), 257–265, 2003. [http://dx.doi.org/10.1016/S0306-2619\(03\)00039-4](http://dx.doi.org/10.1016/S0306-2619(03)00039-4).

GPSA. Gas Processors Suppliers Association. *Engineering Data Book*, 20th ed., 2004. Gas Processors Association, Tulsa, Oklahoma.

IEA, NEA. *Projected Costs of Generating Electricity*. Organisation for Economic Co-operation and Development/International Energy Agency, and Organisation for Economic Co-operation and Development/Nuclear Energy Agency, Paris, 2015 Edition. Available at: <http://www.iea.org/Textbase/npsum/ElecCost2015SUM.pdf>. Last access date: 1/23/2017.

Jons, E.S., Liborius, E., Veltman, P.L., Vernenkar, K.N., 1987. Method for treating by-products from flue gas. US Patent 4666694.

Liu, X., Lin, B., Zhang, Y. Sulfur dioxide emission reduction of power plants in China: current policies and implications. *Journal of Cleaner Production*, 113, 133–143, 2016. <http://dx.doi.org/10.1016/j.jclepro.2015.12.046>.

Ma, X., Kaneko, T., Tashimo, T., Yoshida, T., Kato, K. Use of limestone for SO<sub>2</sub> removal from flue gas in the semidry FGD process with a powder-particle spouted bed. *Chemical Engineering Science*, 55(20), 4643-4652, 2000. [http://dx.doi.org/10.1016/S0009-2509\(00\)00090-7](http://dx.doi.org/10.1016/S0009-2509(00)00090-7).

Mikulčić, H., Klemeš, J. J., Vujanović, M., Urbaniec, K., Duić, N. Reducing greenhouse gasses emissions by fostering the deployment of alternative raw materials and energy sources in the cleaner cement manufacturing process. *Journal of Cleaner Production*, 136, Part B, 119–132, 2016. <http://dx.doi.org/10.1016/j.jclepro.2016.04.145>.

Rochedo, P.R.R., Costa, I.V.L., Império, M., Hoffmann, B.S., Merschmann, P.R. de C., Oliveira, C.C.N., Szklo, A., Schaeffer, R., 2016. Carbon capture potential and costs in Brazil. *J. Clean. Prod.* 131, 280–295. <http://dx.doi.org/10.1016/j.jclepro.2016.05.033>.

Sage, P. W., Ford, N. W. J. Review of Sorbent Injection Processes for Low-Cost Sulphur Dioxide Control. *Proceedings of the Institution of Mechanical Engineers, Part A: Journal of Power and Energy*, 210 (3), 183-190, 1996. Available at: [http://journals.sagepub.com/doi/pdf/10.1243/PIME\\_PROC\\_1996\\_210\\_031\\_02](http://journals.sagepub.com/doi/pdf/10.1243/PIME_PROC_1996_210_031_02). Last access date: 01/23/2017.

Santoyo-Castelazo, E., Azapagic, A., 2014. Sustainability assessment of energy systems: integrating environmental, economic and social aspects. *J. Clean. Prod.* 80, 119–138. <http://dx.doi.org/10.1016/j.jclepro.2014.05.061>

Short, W., Packey, D.J., Holt, T., 1995. *A Manual for the Economic Evaluation of Energy Efficiency and Renewable Energy Technologies*. National Renewable Energy Laboratory (NREL), Colorado.

Thomas, M. Optimizing the Use of Fly Ash in Concrete. Portland Cement Association. Illinois, USA, 2007. Available at: [http://www.cement.org/docs/default-source/fc\\_concrete\\_technology/is548-optimizing-the-use-of-fly-ash-concrete.pdf](http://www.cement.org/docs/default-source/fc_concrete_technology/is548-optimizing-the-use-of-fly-ash-concrete.pdf). Last access date: 01/23/2017.

Tolmasquim, M.T., 2016. Energia Termelétrica: Gás Natural, Biomassa, Carvão, Nuclear, 1st ed. Empresa de Pesquisa Energética (EPE), Rio de Janeiro.

Wu, X., Wu, K., Zhang, Y., Hong, Q., Zheng, C., Gao, X., Cen, K., 2017. Comparative life cycle assessment and economic analysis of typical flue-gas cleaning processes of coal-fired power plants in China. *J. Clean. Prod.* 142, 3236–3242. doi:10.1016/j.jclepro.2016.10.146



## 6. ENVIRONMENTAL PERFORMANCE OF A SOLID WASTE MONETIZATION PROCESS APPLIED TO A COAL-FIRED POWER PLANT WITH SEMI-DRY FLUE-GAS DESULFURIZATION

*This chapter is published a full-length original article in the Journal of Journal of Sustainable Development of Energy, Water and Environment Systems*

### Abstract

Mixing of semi-dry flue-gas desulfurization solids and fly-ash from coal-fired power plants results in a solid waste contaminated by calcium sulfite. Therefore, it becomes useless for industry and is often landfilled. To support decision-making on process configurations to monetize this solid residue a gate-to-gate life cycle assessment was performed, considering three scenarios: BASE case – standard 360 MW power plant, CASE I – base plant adopting dry thermal oxidation treatment of spray dryer solids, CASE II – bypass of desulfurization system. Cases I and II allow commercialization of the solid residue as class C fly-ash. Evaluated alternatives were compared based on quantitative potential environmental impacts, using United States Environmental Protection Agency waste reduction algorithm. Based on the results, the BASE case was more aggressive to the environment, due to solid waste production. CASE II increased photochemical oxidation and acidification potentials. CASE I was the more environmentally friendly but demands additional capital and operational expenditure.

**Keywords:** Calcium sulfite dry oxidation, Coal fired power plant, Life cycle assessment, Semi-dry flue-gas desulfurization, Solid waste treatment, Spray dryer absorbers

### Nomenclature

$M_{j,out}$	output mass flows of $j$ streams	[-]
$MW$	molecular weight	[kg mol/kg]
$n$	polytropic exponent	[-]
$P_W$	power	[kW]
$P_i$	Pressure at $i$ , where $i$ is a counter	[kPa]
$q$	gas flow rate	[kg/h]
$R$	universal gas constant (8.314 J/molK)	[-]
$r_p$	pressure ratio	[-]
$T$	temperature	[K]
$x_{kj}$	$k$ component composition on $j$ output stream	[-]
$Z$	compressibility factor	[-]

### Greek letters

$\alpha$	impacts categories
$\eta_p$	polytropic efficiency
$\Psi_{ki}$	normalized score of $i$ category and $k$ component

## Abbreviations

AP	Acidification Potential
ATP	Aquatic Toxicity Potential
CCP	Coal Combustion Products
EPRI	Electric Power Research Institute
FBR	Fluidized Bed Reactor
FGD	Flue-Gas Desulfurization
GDP	Gross Domestic Product
GHG	Greenhouse Gas
GWP	Global Warming Potential
HTPE	Human Toxicity Potential by Exposure
HTPI	Human Toxicity Potential by Ingestion
LCA	Life Cycle Assessment
ODP	Ozone Depletion Potential
PCC	Pulverized Coal Combustion
PCOP	Photochemical Oxidation Potential
PEI	Potential of Environmental Impacts
PFD	Process Flow Diagram
SDA	Spray Dryer Absorber
SD-FGD	Semi-dry Flue-Gas Desulfurization
SD-FGD-R	Semi-dry Flue-Gas Desulfurization Solid Residue
TTP	Terrestrial Toxicity Potential
WAR	Waste Reduction Algorithm

## 6.1. Introduction

Coal-fired power plants are responsible to fuel 41% of global electricity demand [1]. In some countries, this share is much higher. In China for instance, the world's largest coal producer and consumer, the use of coal for power generation is not expected to decrease in the short to medium term [2]. Despite the on-going transition to a low carbon economy driving a move to renewable sources of energy, the supply of base-load remains dependent on fossil fuel to face their intermittent supply. In this scenario, coal is the most plentiful, and one of the cheapest, among fossil alternatives. As an example, the water scarcity crisis that occurred in Brazil, during 2013-2015, limited the hydropower generation. It is known that the majority of Brazilian electricity is supplied by hydro sources [3], this source is responsible to supply 65% of the total electricity demand [4]. As result, electricity from coal-fired power plants has increased 24.2%, presently, mineral coal represents 9.6% of the thermoelectric power source in Brazil [5].

In the face of the huge amount of solid waste produced by coal-fired power plants, many initiatives were raised in the last decades, aiming to improve waste management of such processes. Common associated solid wastes are: fly and bottom ash, flue-gas desulfurization sludges, boiler blowdown and coal pile runoff, chemicals and other materials related to power plants operation. Within all named solid wastes, fly-ash, bottom ash, slag and scrubber sludge are the ones produced in higher volume [6].

Coal combines organic and mineral components in varying proportions, with ash yields ranging from 3 to 49%. Consequently, coal power generation produces significant amounts of solid wastes, Coal Combustion Products (CCP), consisting of fly-ash, bottom ash, boiler slag, and material from Flue-Gas-Desulfurization (FGD, process applied to flue-gas stream to chemically trap sulfur) [7]. The term coal ash has been used to refer to all the different ash types [8]. CCP is composed basically of non-combustible minerals and a small fraction of unreacted carbon [1]. Depending on burner and pollution control technologies (e.g., FGD), the solid wastes composition varies significantly. Wet CCP is disposed in large surface impoundments while dry CCP is disposed in landfills. To reduce landfill occupation, there is a need for utilization of CCP into valuable materials.

Semi-Dry FGD (SD-FGD) is a technology that uses Spray Dryer Absorption (SDA) to control Sulfur dioxide ( $\text{SO}_2$ ) emissions by flue-gases, by chemical reaction with lime. According to Electric Power Research Institute [9], in 2007, about 12% of USA power plants were using SD-FGD systems, whose water use is 30 to 40% lower than the Wet-FGD technology, being attractive in regions where water supply is limited. However, while Wet-FGD CCP has commercial value for gypsum production, SD-FGD solid is almost useless, having landfills as usual destination. In general, CCP produced by SD-FGD systems is composed of Calcium sulfite ( $\text{CaSO}_3$ ), fly-ash and unreacted lime. Most power plants with SD-FGD do not have fly-ash pre-collectors resulting in solid waste with high ash content (> 50%).

CCP plays an important role in the cement industry. Besides reducing the need for landfill space, the use of fly-ash as substitute for traditional cement brings environmental benefits: Greenhouse Gas (GHG) emissions and primary raw material reduction. In fact, CCP has been used for decades, as a substitute for mined or manufactured materials, lowering construction costs [10]. Fly-ash is not required to pass through the clinker kiln, an energy-intensive step of Portland cement production. Furthermore, concrete from fly-ash is durable, strong and corrosion resistant [11]. There are patented processes for dry oxidation of  $\text{CaSO}_3$  from SD-

FGD waste into Calcium sulfate ( $\text{CaSO}_4$ ). In general they claim technologies to transform CCP into cementitious material or suitable for other applications. Patent 4,478,810, authored by Bloss *et al.* [12], claims a method of treating final products from FGD. Patent 4,544,542, authored by Angevine *et al.* [13], claims a method for oxidation of FGD absorbent and the product produced thereby. Patent 4,666,694, authored by Jons *et al.* [14], claims a method for treating by-products from flue-gas.

Alternative methods aiming to improve CCP properties and applications have also been highlighted in the literature. Li *et al.* [15] reported improving the pozzolanic degree of fly-ash using chemical activators solutions of Sodium hydroxide ( $\text{NaOH}$ ), Sodium sulfate ( $\text{Na}_2\text{SO}_4$ ) and Sodium chloride ( $\text{NaCl}$ ) injected into the fluidized fly-ash through a side spray device, in a Fluidized Bed Reactor (FBR). Ren-ping *et al.* [16] studied the oxidation characteristics of ashes containing  $\text{CaSO}_3$ . SDA material has been used commercially to manufacture cement in Germany after treatment in a fluidized bed process [17]. In fact, post-treatment is necessary since the use of SD-FGD solid residue as cementitious (pozzolanic) material must comply with the ASTM C618 standard or similar country-specific standards [18]. According to ASTM C618, when the  $\text{CaSO}_3$  content of fly-ash exceeds 5% by mass, it is considered inadequate for commercialization as cement additive or replacement material for concrete.

Despite the economic advantage of using SD-FGD waste as cement, the commercial application of this residue remains a challenge. In USA only 22% of SD-FGD residue is used, with mining applications representing 83% of this use. In general, coal fired power plants with SD-FGD dispose its solid waste on landfills, with massive land use. In USA, the production of SD-FGD waste was about  $3.5 \times 10^6$  tonnes in 2009 and is expected to double by 2019 [9].

Clearly, increased utilization of SD-FGD solid residue is needed [19]. The SD-FGD waste landfill is a potential source of contaminants. Besides landfill soil and nearby vegetation ash contamination, leaching of CCP landfills could carry toxic substances, like mercury [2], hexavalent chromium [20] and other contaminants [21], posing potential impact to groundwater.

Additionally, landfill construction and maintenance present economic penalty to electricity generation. Furthermore, the air inside and around the landfill is unhealthy to local workers, because of the high concentration of particulate matter.

Animal tests revealed that SD-FGD waste is not a skin sensitizer but, is irritating to eyes. If ingested, it is an irritant to the digestive tract, causing gastro-intestinal disturbances, erosion or hemorrhage. A moderately acute oral and injection toxicity was indicated in animals. Sulfites are recognized as a food allergen. Breathing difficulty, sneezing, throat swelling and hives could be observed after minutes of ingestion. The inhalation of sulfite aerosol caused mild lung changes in rats and effects on respiratory tract of dogs [22].

Attempts to add use and commercial value to fly-ash appear in the literature since decades. Mulder [23] investigated mechanical properties of coal fly-ash for road base construction material application. Camilleri *et al.* [24] studied the viability of use of fly-ash from coal-fired power plant as a cement replacement in concrete mixes. Today this topic is still being explored by many researchers. Use as Geopolymer is proposed by Chindapasirt and Rattanasak [25] and Xu *et al.* [26]. Doudart de la Grée *et al.* [27] investigated the use of fly-ashes as building materials. Ding *et al.* [28] proposed the recovery of alumina from fly-ash.

A Brazilian coal-fired power plant complex, located in the Northeast region, is considering an alternative destination for its SDA solid waste. This complex has 3 identical 360 MW Pulverized Coal Combustion (PCC) power plants, equipped with SD-FGD for reduction of SO<sub>2</sub> emissions. After 4 years of operation, 2 landfills, with total area of ~79,500 m<sup>2</sup> of area, became almost full with CCP and a third one is being built for operation guarantee (see Figure 6.1).

Aiming to solve the environmental challenge related to CCP landfilling, a SD-FGD waste treatment pilot-plant was designed and constructed at the Federal University of Rio de Janeiro [29]. It is based on the above-mentioned patents information with a modified layout and innovative equipment design. The main equipment is a FBR, to oxidize CaSO<sub>3</sub>, reducing the sulfite (SO<sub>3</sub><sup>-2</sup>) content of the FGD waste, allowing the treated residue to be used as pozzolanic material. The FBR of mini-pilot plant has diameter of 200 mm and 1,100 mm of height. The pilot-plant has a heater, a cyclone (to collect and return particles above 10 μm back to the FBR), an economizer (to partially recover the heat of the hot outlet air stream leaving the cyclone) and an air filter (to avoid emission to the atmosphere of small particles, not captured by the cyclone). de Castro *et al.* [29] reported SO<sub>3</sub><sup>-2</sup> content reduction to below 5% w/w under dry oxidation on FBR at temperatures above 500 °C.

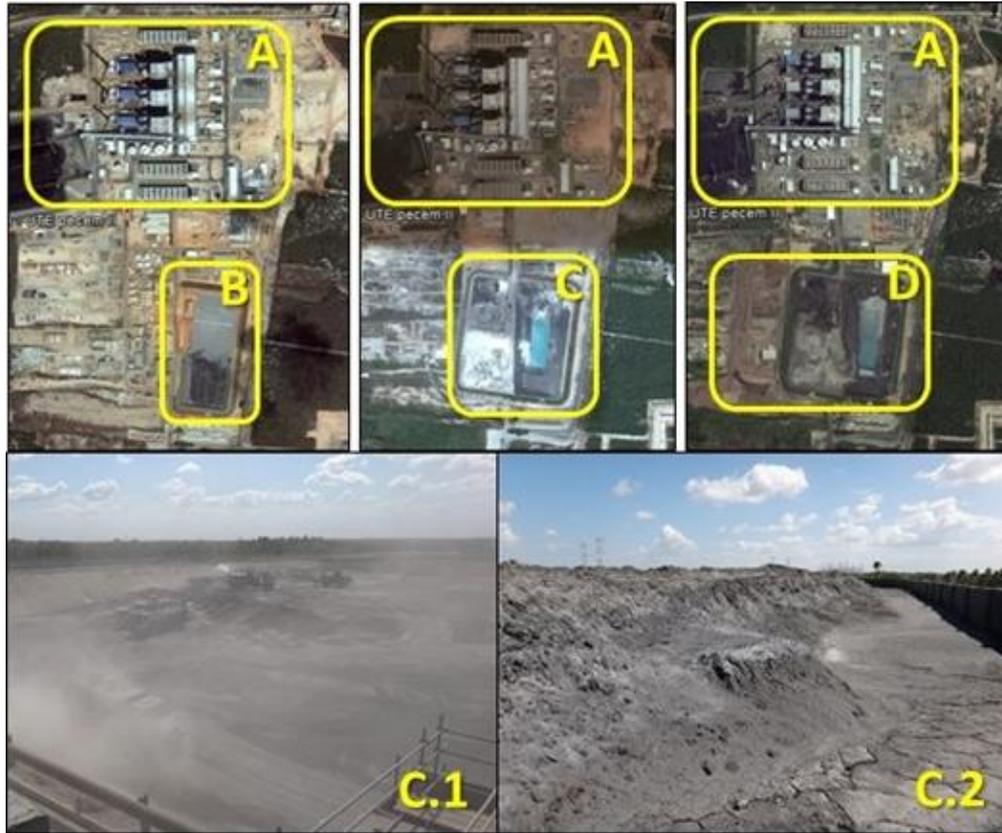


Figure 6.1. Time evolution of ash landfills at a Brazilian coal fired power plant [source: Google Earth (satellite images) and site pictures (landfills 1 and 2):  $3 \times 360$  MW power plants (a); year of 2012 – power plant operations start-up – Landfill I (b); year of 2015 – Landfill I is full, Landfill II in use (c); Landfill II in 2015 (c.1); Landfill I in 2015 (c.2) and year of 2017 – Landfill II is almost full, and Landfill III is under construction (d)]

Based on experimental results of de Castro *et al.* [29] and patent information [13], this work assesses the potential environmental impacts avoided if a full-scale SD-FGD waste treatment unit were put in operation. For a full-scale plant, an air compressor is required to supply air at the FBR pressure. Pressure losses through the economizer, air heater, FBR, cyclone and filter are estimated in 150 kPa, and the compressor pressure ratio is 2.47.

Environmental impacts of waste management are assessed using Waste Reduction Algorithm (WAR) [30] for three alternative destinations of CCP: BASE case, CASE I and CASE II. BASE case is the coal-fired power plant (Figure 6.1) operating with the FGD process and the resulting CCP destined to landfills, considered as waste on WAR. CASE I adapts the power plant to operate with the proposed full-scale FGD waste treatment unit, converting the SDA residue into a class C pozzolanic material. Although CASE I manages CCP without increasing  $\text{SO}_2$  emissions, it demands capital investments (CAPEX) for building the solid waste treatment unit. Although the air used to oxidize  $\text{CaSO}_3$  must be heated above 400-600

°C, the oxidation reaction is exothermic and, depending on the residue composition, could be autothermic. However, extra energy is necessary (e.g., for plant start-up or compensation of heat losses). Integration with hot gases, vapor purge or combustion air from the power plant process would avoid fuel consumption. CASE II consists of turning-off the SD-FGD, making possible to commercialize the residue directly as Class C pozzolanic material, because ashes are not contaminated by desulfurization products.

It is worth noting that CASE I is an environmentally friendly approach for CCP management, while CASE II prioritizes economic performance at the expense of environmental impacts. That alternative is legally possible only if the SO<sub>2</sub> concentration in exhausted gas complies with local environmental regulation (in Brazil, 400 mg/Nm<sup>3</sup>, according to CONAMA 03/1990 [31]<sup>2</sup>). Adjusting the FGD operation and using low sulfur coal, SO<sub>2</sub> emissions will probably be very close to the regulation limit. In the event of surpassing emission limit, increased atmospheric pollution would result, comparatively to CASE I and BASE case.

The main objective of the study is to evaluate, based on a gate-to-gate Life Cycle Assessment (LCA) methodology, the environmental performance of the three CCP management alternatives, considering a set of environmental impact metrics (i.e., not restricted to solely evaluating SO<sub>2</sub> emissions). The results aim to quantify how much CASE I is less polluting than CASE II and BASE case, proving the relevance of SDA waste treatment unit for coal-fired power plants operating with SD-FGD system.

The present results and the proposed methodology contribute to the decision-making process of CCP managing of coal-fired power plants using SD-FGD. No similar work was found in the scientific literature, proving the originality of this study.

## 6.2. Materials and Methods

The assessment of environmental impacts of a process or product systems is useful as a decision-making tool and can be achieved using LCA [4]. ISO 14040 [32] establish four basic steps to perform a LCA:

- Goal and scope definition;
- Inventory analysis;
- Impact assessment;
- Interpretation of results.

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<sup>2</sup> See Erratum, item 1.

### 6.2.1. Goal and scope

The main goal is support decision-making on process configurations to monetize mixed coal combustion products from a 360 MW pulverized coal power plant with semi-dry FGD. A gate-to-gate life cycle assessment is performed, considering three scenarios: BASE – standard power plant [33], CASE I – base plant adopting dry thermal oxidation treatment of spray dryer solids, CASE II – bypass of desulfurization system. Cases I and II allow commercialization of the solid residue as class C fly-ash.

### 6.2.2. Heat and mass balances, and streams inventory

A global mass balance of each process was performed, classifying the streams as: inlet, waste outlet and product outlet. These streams are based on the Process Flow Diagram (PFD) of the Brazilian Coal-Fired power plant pictured in Figure 6.2, used as case study of the proposed methodology. The missing information was calculated from mass balance.

The power plant is supplied with Colombian Coal, with composition assumed as similar to Colombian field IGM 1238 [34]. The considered set of reactions expected to occur inside the FBR and the SDA solid residue composition and mass flow is presented by Cruz *et al.* [33]. The last was obtained from the heat and mass balances of the power plant used as case study and considers coal with 1.5%w/w of sulfur.

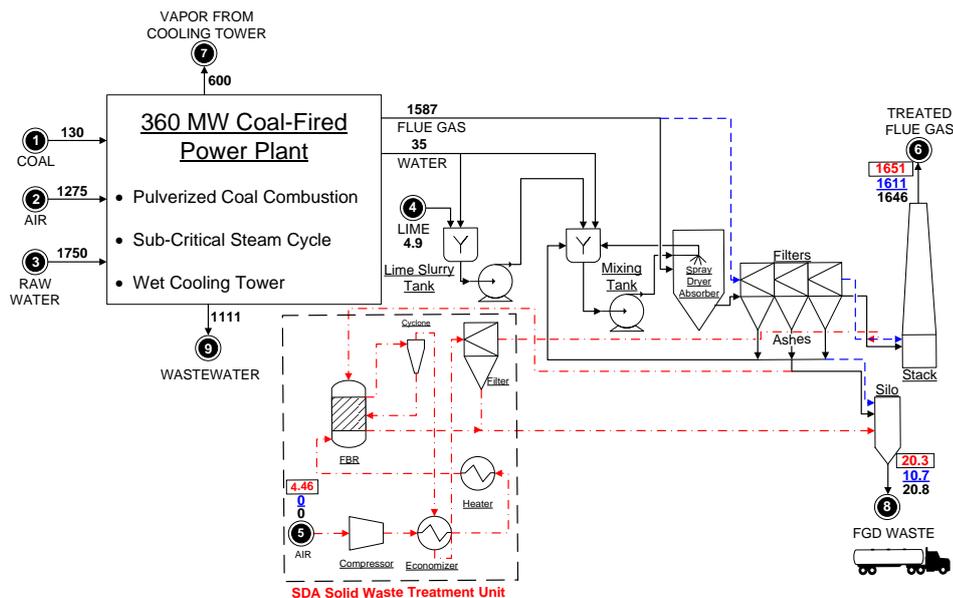


Figure 6.2. Flow diagram of the coal-fired power plant with SD-FGD waste treatment unit (dashed box), numbers in black indicate mass flow of original power plant [t/h], while red numbers within boxes correspond to CASE I and underlined blue numbers correspond to CASE II

Based on Angevine *et al.* [13], considering a temperature of 550 °C and 5% of excess O<sub>2</sub>, it is possible to achieve a SO<sub>3</sub> mass composition of 3.2% on the treated SDA waste, complying with ASTM Standard C-618 [18] Sulfur trioxide (SO<sub>3</sub>) limit for class C or F fly-ash. Based on experiments of de Castro *et al.* [29], a conversion of 91.3% was considered for SO<sub>3</sub> oxidation reaction. These results were considered for mass and energy balances of SDA treatment unit of CASE I. The specific heat of SDA residue, used to estimate energy consumption for heating the SDA residue from 80 °C to 550 °C was considered 730 J/kgK, the same value of a class C fly-ash [35].

As the FBR does not exist in industrial scale, the fluidization air flow was estimated. In a pilot scale FBR dealing with a 10%w/w CaSO<sub>3</sub>×½H<sub>2</sub>O feed, the flow ratio of fluidization air to stoichiometric air was taken as 3.75. Considering the calcium sulfite mass fraction as 38.4%, the stoichiometry air flow with 5% of O<sub>2</sub> excess was considered enough to promote bed fluidization. The air compressor power was calculated by eq. (6.1) [36]:

$$P_w = \frac{q \times Z \times R \times T \left[ (r_p)^{\frac{n-1}{n}} - 1 \right]}{3,600 \times \eta_p \times MW \frac{n-1}{n}} \quad (6.1)$$

where  $P_w$  is the brake horsepower [kW],  $q$  is the gas flow rate [kg/h],  $Z$  is the average compressibility factor,  $R$  is the gas constant (8.314 kJ/kmolK),  $T$  is the gas inlet temperature [K],  $MW$  is the molecular mass [kg mol/kg],  $r_p$  is the pressure ratio,  $n$  is the polytropic exponent and  $\eta_p$  is the polytropic efficiency. The pressure ratio is calculated dividing the outlet pressure,  $P_2$  [kPa], by the inlet pressure  $P_1$  (kPa).  $P_2$  is considered 250 kPa (gauge).  $P_1$  is the atmospheric pressure (0 kPa gauge).  $\eta_p$  is considered 80%,  $q$  of inlet air stream (stream 1 of Figure 6.3), calculated as a function of the FBR demand.

The air is pre-heated to 250 °C by the economizer. The heater service is to heat the air to the reactor temperature (550 °C). However, the energy supplied by the oxidation of CaSO<sub>3</sub> and integration with high temperature steam purges from the closed loop steam cycle could bring the air heating energy input to zero (the air heater can be used only for the start-up of the system). The SDA power consumption was obtained from the Environmental Impacts Assessment Study of the power plant used as case study [37].

### 6.2.3. Waste Reduction Algorithm Methodology

Evaluated alternatives are compared based on quantitative Potential Environmental Impacts (PEI), using United States Environmental Protection Agency (USEPA) Waste Reduction Algorithm [38]. To compare the environmental friendliness of chemical processes, WAR algorithm uses the concept of PEI balance. It is based on the idea that the PEI of a certain amount of material and energy can be defined as the effect that they would have on the environment if they were emitted [30]. As PEI is a conceptual quantity, it cannot be directly measured, but can be calculated from measurable parameters, using functional relations [38]. The balance considers the flow of PEI (mass + energy) across the process boundary [PEI/h]. From the balance, PEI indexes are calculated, providing the degree of environmental friendliness of the process.

WAR algorithm describes the Potential of Environmental Impacts Rate [PEI/h] for each category using the eq. (6.2) [38]:

$$\hat{I}_{\text{out}} = \sum_i^{\text{category}} (\alpha_i) \sum_j^{\text{stream}} (M_{j,\text{out}}) \sum_k^{\text{component}} (x_{kj} \psi_{ki}) \quad (6.2)$$

where  $\hat{I}_{\text{out}}$  is the output PEI rate,  $\alpha_i$  is the user defined weight factor for the  $i^{\text{th}}$  impact category,  $M_{j,\text{out}}$  is the output mass flow of the  $j^{\text{th}}$  stream,  $x_{kj}$  is the composition of the  $k^{\text{th}}$  component in the  $j^{\text{th}}$  output stream,  $\psi_{ki}$  is the normalized score of  $i^{\text{th}}$  impact category for the  $k^{\text{th}}$  component ( $\text{score}_{ki} / \langle \text{score}_{ki} \rangle$ ).  $\langle \text{score}_{ki} \rangle$  is the average score of all components in a same category. According to Young and Cabezas [30], WAR classifies PEI in impact categories, with the global PEI resulting from their weighted sum (with user defined weights). Table 6.1 shows the impact categories and weights adopted for the current evaluation. The objective of the study is comparing scenarios. Therefore, the weight and absolute value of each category individually does not matter in the proposed analysis.

We are interested in the difference between cases. Using different weights for some categories might be considered an attempt to manipulating the conclusions. Thus, it was decided to keep all weights equal to 1, for all the 3 cases.

The inventory streams of cases BASE, I and II were used as input of WAR algorithm, through the software WAR GUI Version 1.0.17 (2008), namely chemical composition and flow rates of

mass streams entering and leaving the process. Energy input were ignored (considered zero) since all the alternative cases present similar energy use.

Table 6.1. WAR environmental impacts categories and adopted weights

Impact	Description	Weight
HTPI	Human Toxicity Potential by Ingestion	1
HTPE	Human Toxicity Potential by Exposure	1
ATP	Aquatic Toxicity Potential	1
TTP	Terrestrial Toxicity Potential	1
GWP	Global Warming Potential	1
ODP	Ozone Depletion Potential	1
PCOP	Photochemical Oxidation Potential	1
AP	Acidification Potential	1

### 6.3. Results and Discussion

The methodology stated on the last section was successfully applied and the main results are presented below.

#### 6.3.1. Fluidized Bed Reactor heat and mass balance

According to Cruz *et al.* [33], the mass flow of SDA solid residue is 20.8 tons/h, fly ash specific heat is 730 J/kgK. The initial  $\text{CaSO}_3 \times \frac{1}{2}\text{H}_2\text{O}$  content on SDA residue is 38.4% w/w, with 2% of water (humidity), 8.3% of Calcium hydroxide  $[\text{Ca}(\text{OH})_2]$  and 51.3% of inert minerals (fly-ash). Therefore, it is possible to calculate the heat balance around the FBR, product mass flow and composition as well as the air mass flows (in and out) to promote CCP oxidation inside the FBR, as shown in Tables 6.2-6.4.

Table 6.2. FBR heat balance

Item	Energy [kW]
Air heating	372
Solids heating	1,980
Reaction 1 ( $\text{CaSO}_3 \times \frac{1}{2}\text{H}_2\text{O} \rightarrow \text{CaSO}_3 + \text{H}_2\text{O}$ )	494
Reaction 2 ( $\text{CaSO}_4 \times \frac{1}{2}\text{H}_2\text{O} \rightarrow \text{CaSO}_4 + \text{H}_2\text{O}$ )	0.0
Reaction 3 [ $\text{Ca}(\text{OH})_2 \rightarrow \text{CaO} + \text{H}_2\text{O}$ ]	697
Reaction 4 ( $\text{CaSO}_3 + \frac{1}{2}\text{O}_2 \rightarrow \text{CaSO}_4$ )	-4,260
Balance	-717

Table 6.3. FBR product stream

Component	MW [g/mol]	Flow [mol/h]	Flow [kg/h]	Composition [% weight]
CaSO <sub>3</sub> ×½H <sub>2</sub> O	129	-	-	0.00
CaSO <sub>4</sub> ×2H <sub>2</sub> O	172	-	-	0.00
Ca(OH) <sub>2</sub>	74	-	-	0.00
CaSO <sub>3</sub>	120	6,186	742	3.7
CaSO <sub>4</sub>	136	55,671	7,571	37.3
CaO	56	23,307	1,305	6.5
H <sub>2</sub> O	18	-	-	0.00
Inert	-	-	10,660	52.5
Total			20,279	100.0

Table 6.4. FBR air inlet and outlet streams

FBR air inlet stream				
Component	MW [g/mol]	Flow [mol/h]	Flow [kg/h]	% Molar [mole %]
O <sub>2</sub>	32	32,475	1,039	21.0
N <sub>2</sub>	28	122,167	3,421	79.0
Air	29	154,642	4,460	100.0
FBR air outlet stream				
Component	MW [g/mol]	Flow [mol/h]	Flow [kg/h]	% Molar [mole %]
O <sub>2</sub>	32	4,639	148	2.3
N <sub>2</sub>	28	122,167	3,421	59.8
H <sub>2</sub> O	18	77,325	1,392	37.9
Air	24	204,131	4,961	100.0

The FBR heat balance shows that, considering all stated premises, the reaction could be self-sufficient in terms of energy, and energy input is necessary only to start up the FBR and to supply the compressor. As shown in Table 6.5, the extra energy is 1,275 kW. This is only 0.35% of the plant turbine power output (360 MW) and was not considered in the WAR algorithm analysis.

Table 6.5. Overall power plant heat balance for CASE I and CASE II

Case	Unit	I	II
Coal consumption	[tons/h]	135	135
Boiler duty	[kW]	987,368	987,368
Turbine output	[kW]	360,000	360,000
Compressor power	[kW]	135	0.00
SDA consumption	[kW]	1,140	0.00
Net electrical power output	[kW]	358,725	360,000
SDA + Ash treatment energy penalty	[kW]	1,275	0.00
SDA + Ash treatment energy penalty	[%]	0.35	0.00
Plant efficiency (LHV)	[%]	36.33	36.46

### 6.3.2. Waste Reduction Algorithm Results

Based on streams inventory, the PEI generation rate of each case (BASE, I and II) were calculated using the software WAR. Tables 6.6-6.8 show the streams inventory of each case.

The results for each environmental impact category and the total PEI rate are summarized in Figure 6.3.

Table 6.6. BASE case streams inventory

Stream	1	2	3	4	5	6	7	8	9
Type	Inlet	Inlet	Inlet	Inlet	Inlet	Waste outlet	Waste outlet	Waste outlet	Waste outlet
Name	Coal	Air inlet 1	Raw water	Lime	Air inlet 2	Flue-gas	Water vapor	FGD waste	Wastewater
Flow [tons/h]	129	1,275	1,750	4.9	-	1,646	600	20.4	857
Coal	1.000	-	-	-	-	-	-	-	-
N <sub>2</sub>	-	0.8113	-	-	0.8113	0.6845	-	-	-
O <sub>2</sub>	-	0.1887	-	-	0.1887	0.0508	-	-	-
H <sub>2</sub> O	-	-	1.0000	-	-	0.0761	1.0000	0.0200	1.000
SO <sub>2</sub>	-	-	-	-	-	0.0002	-	-	-
CO <sub>2</sub>	-	-	-	-	-	0.1884	-	-	-
SiO <sub>2</sub>	-	-	-	-	-	-	-	0.1319	-
Al <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.1391	-
CaO	-	-	-	0.9500	-	-	-	0.1368	-
MgO	-	-	-	0.0500	-	-	-	0.0144	-
Fe <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.0649	-
TiO <sub>2</sub>	-	-	-	-	-	-	-	0.0094	-
P <sub>2</sub> O <sub>5</sub>	-	-	-	-	-	-	-	0.0030	-
CaSO <sub>3</sub>	-	-	-	-	-	-	-	0.3840	-
SO <sub>4</sub> <sup>*</sup>	-	-	-	-	-	-	-	0.0964	-

Table 6.7. CASE I streams inventory

Stream	1	2	3	4	5	6	7	8	9
Type	Inlet	Inlet	Inlet	Inlet	Inlet	Waste outlet	Waste outlet	Product	Waste outlet
Name	Coal	Air inlet 1	Raw water	Lime	Air inlet 2	Flue-gas	Water vapor	Fly-ash	Wastewater
Flow [tons/h]	129	1,275	1,750	4.9	4.9	1,651	600	20.4	857
Coal	1.000	-	-	-	-	-	-	-	-
N <sub>2</sub>	-	0.8113	-	-	0.8113	0.6845	-	-	-
O <sub>2</sub>	-	0.1887	-	-	0.1887	0.0508	-	-	-
H <sub>2</sub> O	-	-	1.0000	-	-	0.0761	1.0000	-	1.000
SO <sub>2</sub>	-	-	-	-	-	0.0002	-	-	-
CO <sub>2</sub>	-	-	-	-	-	0.1884	-	-	-
SiO <sub>2</sub>	-	-	-	-	-	-	-	0.1352	-
Al <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.1426	-
CaO	-	-	-	0.9500	-	-	-	0.1195	-
MgO	-	-	-	0.0500	-	-	-	0.0148	-
Fe <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.0665	-
TiO <sub>2</sub>	-	-	-	-	-	-	-	0.0097	-
P <sub>2</sub> O <sub>5</sub>	-	-	-	-	-	-	-	0.0031	-
CaSO <sub>3</sub>	-	-	-	-	-	-	-	0.0365	-
SO <sub>4</sub> <sup>*</sup>	-	-	-	-	-	-	-	0.4722	-

\*See Erratum, item 2.

Table 6.8. CASE II streams inventory

Stream	1	2	3	4	5	6	7	8	9
Type	Inlet	Inlet	Inlet	Inlet	Inlet	Waste outlet	Waste outlet	Product	Waste outlet
Name	Coal	Air inlet 1	Raw water	Lime	Air inlet 2	Flue-gas	Water vapor	Fly-ash	Wastewater
Flow [tons/h]	129	1,275	1,715	0	0	1,611	600	10.7	921
Coal	1.0000	-	-	-	-	-	-	-	-
N <sub>2</sub>	-	0.8113	-	-	0.8113	0.6942	-	-	-
O <sub>2</sub>	-	0.1887	-	-	0.1887	0.0487	-	-	-
H <sub>2</sub> O	-	-	1.0000	-	-	0.0582	1.0000	-	1.000
SO <sub>2</sub>	-	-	-	-	-	0.0028	-	-	-
CO <sub>2</sub>	-	-	-	-	-	0.1961	-	-	-
SiO <sub>2</sub>	-	-	-	-	-	-	-	0.2572	-
Al <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.2712	-
CaO	-	-	-	0.9500	-	-	-	0.1048	-
MgO	-	-	-	0.0500	-	-	-	0.0281	-
Fe <sub>2</sub> O <sub>3</sub>	-	-	-	-	-	-	-	0.1264	-
TiO <sub>2</sub>	-	-	-	-	-	-	-	0.0184	-
P <sub>2</sub> O <sub>5</sub>	-	-	-	-	-	-	-	0.0059	-
CaSO <sub>3</sub>	-	-	-	-	-	-	-	-	-
SO <sub>4</sub> <sup>-</sup>	-	-	-	-	-	-	-	0.1880	-

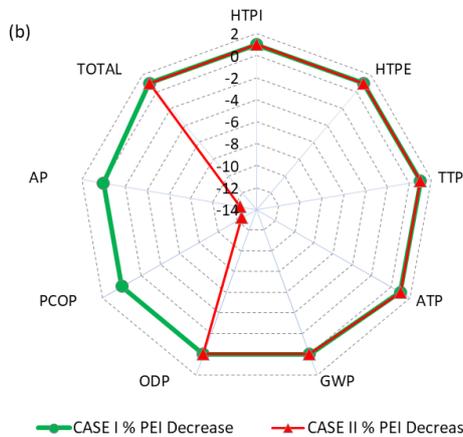
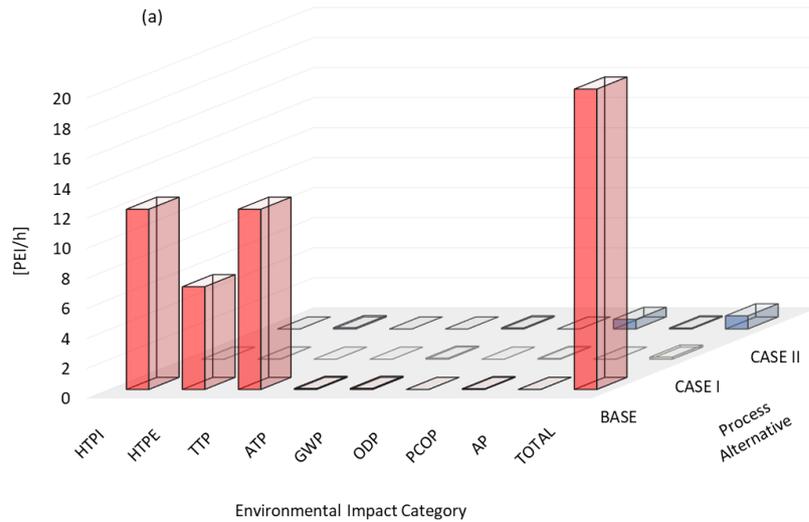


Figure 6.3. Environmental impact assessment: PEI/h for each impact categories (a) and decrease of PEI generation rates for CASE I and CASE II with respect to BASE case (b)

\*See Erratum, item 2.

Table 6.6 refers to BASE case streams inventory. On BASE case it can be noticed that the stream 8 is considered as waste, because the 20.4 tons/h of SD-FGD solid is landfilled. Table 6.7 represents CASE I, this is the only one when the stream 5 (air) is not zero. This air is used on the FBR reactor, to oxidize the  $\text{CaSO}_3$  to  $\text{CaSO}_4$ . On Tables 6.7 and 6.8 the stream 8 is considered a product, and not a waste. In this way, environmental impacts of those streams are not considered by the software on PEI generation rates. On Table 6.8 (CASE II), it is noticed that the solids production (stream 8) is lower. It happens because FGD is out of operation. There is no lime consumption (stream 4 flow is zero), the only solid waste source is coal combustion. CASE II presents a 2% decrease on water consumption (stream 3). The reason is that FGD uses 35 tons/h of water, that evaporates on the SDA. It is shown by the difference on waste flue-gas mass flow (stream 6) of CASE II, compared to CASE I and BASE case. Stream 6 of CASE I presents a higher flow because the air used by the FBR is mixed with the flue-gas from boiler. CASE II presents a lower flow because, as the flue-gases do not pass through SDA, no water vapor is mixed with this stream.

Figure 6.3a shows clearly that BASE case scores are higher in categories related to human health and terrestrial toxicity (HTPI, HTPe and TTP), proving that FGD waste is indeed an environmental problem. As the PEI rate of these categories were an order of magnitude higher related to the other ones, results are presented in Figure 6.4, for PEI generation rates [PEI/h] in categories ATP, GWP, ODP, PCOP, AP, and decrease in PEI generation of CASES I and II with respect to BASE case.

The absence of  $\text{SO}_2$  recovery system resulted in a photochemical oxidation and acidification potential PEI generation rate 1,245% higher for CASE II. That happens because these categories are directly affected by  $\text{SO}_2$  emissions. The total PEI generation reduction of CASE I was approximately 500% related to CASE II, showing definite inferiority of CASE II with respect to CASE I. It is worth noting that CASES I and II have very lower Total PEI generation rates since both CCP (solid wastes from FGD) comply with specifications for commercial use, hence being considered products and, as such, are not computed as waste (reducing PEI generation) by WAR. Clearly, the more environmentally friendly alternative to FGD solid waste problem is CASE I.

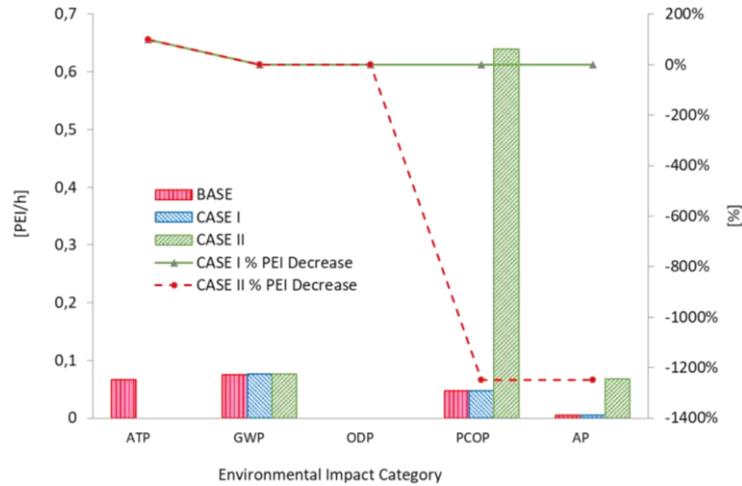


Figure 6.4. PEI generation rates [PEI/h] for categories ATP, GWP, ODP, PCOP, AP and decrease in PEI generation of Cases I and II with respect to BASE case

## 6.4. Conclusions

Heat and mass balances were performed for three modes of operation of the Semi-Dry FGD section of a Coal Fired power plant in the northeast of Brazil. WAR results demonstrated that BASE case is much more aggressive to the environment, due to the large amount of useless FGD waste produced. The treatment of FGD waste (CASE I) or the bypass of the SDA system (CASE II) were compared separately, as alternatives to transform the solid waste into a class C fly-ash. Because of the SDA system bypassing,  $\text{SO}_2$  emission was responsible for increasing PCOP and AP by 1,245%. CASE I was demonstrated to be the more environmentally friendly alternative, although resulting in capital expenditure to install an FBR and auxiliary equipment, to oxidize  $\text{CaSO}_3$  and solve the problem of landfill use. The SDA and FBR operation also entail an increment of operational expenditures, like energy (1,275 kW or 0.35% of the total turbine power output), water (35 t/h) and lime consumption (4.9 tons/h).

Both Cases I and II allow the commercialization of the solids coming from the SDA as class C fly-ash. Thus, considering only the economic point of view, CASE II is better, but this study proves that the environmental impacts related to  $\text{SO}_2$  emissions increases dramatically, and could be prohibitive in countries where the environmental legislation is more restrictive, like in Western Europe and USA. CASE I is more sustainable, because it solves  $\text{SO}_2$  emissions, while reducing environmental impacts in other impact categories, contrarily to

CASE II, which favors economics, increasing air pollution to mitigate landfill related environmental impacts. In the long term, depending on the ash and cement market, CASE I could become profitable, resulting from commercialization of treated CCP. Future work must include new data from the recently improved pilot plant and ash analytical methodology, aiming to generate a more accurate streams inventory. Results from this work could be validated, using other LCA software and data basis, like SimaPro and Ecoinvent. Use of low-grade heat from power plant could favor the economic and environmental performance of the full-scale SD-FGD treatment system. This effect must be investigated, and results included in future LCA studies.

## 6.5. References of Chapter 6

1. World Coal Association, Coal & Electricity, 2017, <https://www.worldcoal.org/coal/uses-coal/coal-electricity>, [Accessed: 14-March-2017]
2. Wang, Y., Duan, Y., Yang, L., Huang, Z., Meng, S. and Zhao, C., Mercury Speciation and Emission from the Coal-fired Power Plant Filled with Flue Gas Desulfurization Equipment, *Can. J. Chem. Eng.*, Vol. 88, No. 5, pp 867-873, 2010, <https://doi.org/10.1002/cjce.20331>
3. de Almeida Prado, F. A., da Silva, A. L. R., Avila, E. M. and Matsuyama, G., Clean Energy Certification in Brazil: A Proposal, *J. Sustain. Dev. Energy, Water Environ. Syst.*, Vol. 3, No. 1, pp 95-105, 2015, <https://doi.org/10.13044/j.sdewes.2015.03.0007>
4. Geller, M. T. B. and Meneses, A. A. M., Life Cycle Assessment of a Small Hydropower Plant in the Brazilian Amazon, *J. Sustain. Dev. Energy, Water Environ. Syst.*, Vol. 4, No. 4, pp 379-391, 2016, <https://doi.org/10.13044/j.sdewes.2016.04.0029>
5. Empresa de Pesquisa Energética (EPE), National Energetic Balance 2015 (in Portuguese), 2015, [https://ben.epe.gov.br/downloads/S%25c3%25adntese do Relat%25c3%25b3rio Final\\_2015\\_Web.pdf](https://ben.epe.gov.br/downloads/S%25c3%25adntese%20do%20Relat%25c3%25b3rio%20Final_2015_Web.pdf), [Accessed: 07-April-2016]
6. Row, R. W., Developments in the Management of Wastes from Coal-fired Power Plants, *Waste Manag.*, Vol. 14, No. 3, pp 299-308, 1994, [https://doi.org/10.1016/0956-053X\(94\)90076-0](https://doi.org/10.1016/0956-053X(94)90076-0)
7. Pierce, B. S. and Dennen, K. O., The National Coal Resource Assessment Overview, *U.S. Geol. Surv. Prof. Pap.*, Vol. 1625, No. F, p 402, 2009.
8. Heidrich, C., Feuerborn, H.-J. and Weir, A., Coal Combustion Products: A Global Perspective, *Proceedings of the World of Coal Ash Conference*, Lexington, Kentucky, USA, 2013.
9. Electric Power Research Institute (EPRI), Characterization of Spray Dryer Absorber Products for Use in Cement and Concrete Applications 1017580, Palo Alto, California,

- USA, 2009.
10. American Coal Ash Association (ACAA), Key Findings 2015, Coal Combustion Products Utilization, U.S. Historical Perspective and Forecast, 2015.
  11. World Coal Association (WCA), Coal Combustion Products, 2018, <https://www.worldcoal.org/coal/uses-coal/coal-combustion-products>, [Accessed: 14-March-2017]
  12. Bloss, W. and Mohn, U., Method of Treating Final Products from Flue Gas Desulfurization, Patent Number 4,478,810, 1984.
  13. Angevine, P. A., Bengtsson, S. and Koudijs, G. P., Method for Oxidation of Flue Gas Desulfurization Absorbent and the Product Produced Thereby, Patent Number 4,544,542, 1985.
  14. Jons, E. S., Liborius, E., Veltman, P. L. and Vernenkar, K. N., Method for Treating By-Products from Flue Gas, Patent Number 4,666,694, 1987.
  15. Li, C., Zhu, H., Wu, M., Wu, K. and Jiang, Z., Pozzolanic Reaction of Fly Ash Modified by Fluidized Bed Reactor-vapor Deposition, *Cem. Concr. Res.*, Vol. 92, pp 98-109, 2017, <https://doi.org/10.1016/j.cemconres.2016.11.016>
  16. Liu, R., Guo, B., Ren, A. and Bian, J., The Chemical and Oxidation Characteristics of Semi-dry Flue Gas Desulfurization Ash from a Steel Factory, *Waste Manag. Res.*, Vol. 28, No. 10, pp 865-871, 2010, <https://doi.org/10.1177/0734242X09339952>
  17. Electric Power Research Institute (EPRI), A Review of Literature Related to the Use of Spray Dryer Absorber Material Production, Characterization, Utilization Applications, Barriers, and Recommendations, 1014915, Technical Report, Palo Alto, California and Grand Forks, North Dakota, USA, 2007.
  18. ASTM, C618-15, Standard Specification for Coal Fly Ash and Raw or Calcined Natural Pozzolan for Use, *Annual Book of ASTM Standards*, p 5, ASTM International, West Conshohocken, Pennsylvania, USA, 2015.
  19. American Coal Ash Association (ACAA), Beneficial Use of Coal Combustion Products, An American Recycling Success Story, 2014, <https://www.aaa-usa.org/Portals/9/Files/PDFs/Production-and-Use-Brochure.pdf>, [Accessed: 07-April-2016]
  20. Huggins, F. E., Rezaee, M., Honaker, R. Q. and Hower, J. C., On the Removal of Hexavalent Chromium from a Class F Fly Ash, *Waste Manag.*, Vol. 51, pp 105-110, 2016, <https://doi.org/10.1016/j.wasman.2016.02.038>
  21. Hjelmar, O., Leachate from Land Disposal of Coal Fly Ash, *Waste Manag. Res.*, Vol. 8, No. 6, pp 429-449, 1990, [https://doi.org/10.1016/0734-242X\(90\)90019-J](https://doi.org/10.1016/0734-242X(90)90019-J)
  22. European Chemicals Agency (ECHA), Product of Semi-Dry Absorption method of Flue Gas Desulfurization – Registration Dossier, <https://echa.europa.eu/registration-dossier/-/registered-dossier/15224/7/2/2#>, [Accessed: 30-March-2017]

23. Mulder, E., A Mixture of Fly Ashes as Road Base Construction Material, *Waste Manag.*, Vol. 16, No. 1, pp 15-20, 1996, [https://doi.org/10.1016/S0956-053X\(96\)00026-8](https://doi.org/10.1016/S0956-053X(96)00026-8)
24. Camilleri, J., Sammut, M. and Montesin, F. E., Utilization of Pulverized Fuel Ash in Malta, *Waste Manag.*, Vol. 26, No. 8, pp 853-860, 2006, <https://doi.org/10.1016/j.wasman.2005.11.022>
25. Chindapasirt, P. and Rattanasak, U., Utilization of Blended Fluidized Bed Combustion (FBC) Ash and Pulverized Coal Combustion (PCC) Fly Ash in Geopolymer, *Waste Manag.*, Vol. 30, No. 4, pp 667-672, 2010, <https://doi.org/10.1016/j.wasman.2009.09.040>
26. Xu, H., Li, Q., Shen, L., Zhang, M. and Zhai, J., Low-reactive Circulating Fluidized Bed Combustion (CFBC) Fly Ashes as Source Material for Geopolymer Synthesis, *Waste Manag.*, Vol. 30, No. 1, pp 57-62, 2010, <https://doi.org/10.1016/j.wasman.2009.09.014>
27. Doudart de la Grée, G. C. H., Florea, M. V. A., Keulen, A. and Brouwers, H. J. H., Contaminated Biomass Fly Ashes – Characterization and Treatment Optimization for Reuse as Building Materials, *Waste Manag.*, Vol. 49, pp 96-109, 2016, <https://doi.org/10.1016/j.wasman.2015.12.023>
28. Ding, J., Ma, S., Shen, S., Xie, Z., Zheng, S. and Zhang, Y., Research and Industrialization Progress of Recovering Alumina From Fly Ash: A Concise Review, *Waste Manag.*, Vol. 60, pp 375-387, 2017, <https://doi.org/10.1016/j.wasman.2016.06.009>
29. de Castro, R. de P. V., de Medeiros, J. L., Araújo, O. de Q. F., Cruz, M. de A., Ribeiro, G. T. and de Oliveira, V. R., Fluidized Bed Treatment of Residues of Semi-dry Flue Gas Desulfurization Units of Coal-fired Power Plants for Conversion of Sulfites to Sulfates, *Energy Convers. Manag.*, Vol. 143, pp 173-187, 2017, <https://doi.org/10.1016/j.enconman.2017.03.078>
30. Young, D. M. and Cabezas, H., Designing Sustainable Processes with Simulation: The Waste Reduction (WAR) Algorithm, *Comput. Chem. Eng.*, Vol. 23, No. 10, pp 1477-1491, 1999, [https://doi.org/10.1016/S0098-1354\(99\)00306-3](https://doi.org/10.1016/S0098-1354(99)00306-3)
31. National Environment Council (CONAMA), CONAMA Resolution No. 003 from June 28<sup>th</sup> 1990 (in Portuguese), 1990.
32. The International Standards Organisation, Environmental Management — Life Cycle Assessment — Principles and Framework, ISO 14040, 2006.
33. Cruz, M. de A., Araújo, O. de Q. F., de Medeiros, J. L., de Castro, R. de P. V., Ribeiro, G. T. and de Oliveira, V. R., Impact of Solid Waste Treatment from Spray Dryer Absorber on the Levelized Cost of Energy of a Coal-fired Power Plant, *J. Clean. Prod.*, Vol. 164, pp 1623-1634, 2017, <https://doi.org/10.1016/j.jclepro.2017.07.061>
34. Tewalt, S. J., Finkelman, R. B., Torres, I. E. and Simoni, F., Colombia, World Coal Quality Inventory: South America, 2006, [http://pubs.usgs.gov/of/2006/1241/Chapter 5-Colombia.pdf](http://pubs.usgs.gov/of/2006/1241/Chapter%205-Colombia.pdf), [Accessed: 28-May-2015]
35. Bentz, D., Peltz, M., Duran-Herrera, A., Valdez, P. and Juarez, C., Thermal Properties of

High-volume Fly Ash Mortars and Concretes, *J. Build. Phys.*, Vol. 34, No. 3, pp 263-275, 2011, <https://doi.org/10.1177/1744259110376613>

36. GPSA, *Engineering Data Book* (20<sup>th</sup> ed.), Gas Processors Suppliers Association, Tulsa, Oklahoma, USA, 2004.
37. Diferencial Energia Empreendimentos e Participação Ltda (DEEPL), Environmental Impact Assessment, Port of Itaquí Power Plant Enterprise (in Portuguese), p 1395, Rio de Janeiro, Brasil, 2008.
38. Cabezas, H., Bare, J. C. and Mallick, S. K., Pollution Prevention with Chemical Process Simulators: The Generalized Waste Reduction (WAR) Algorithm — Full Version, *Comput. Chem. Eng.*, Vol. 23, No. 4-5, pp 623-634, 1999, [https://doi.org/10.1016/S0098-1354\(98\)00298-1](https://doi.org/10.1016/S0098-1354(98)00298-1)

## 7. CO<sub>2</sub> CAPTURE FROM FLUE-GASES BY PHASE-CHANGING ABSORPTION SOLVENTS

*This first part of this chapter is based on the conference paper SDEWES2019.0276, presented at the 14<sup>th</sup> Sustainable Development on Environment Water and Energy Systems Conference – Dubrovnik – 2019. Additional content is included on the second part, to update and complement this topic.*

### 7.1. Chemical Absorption of CO<sub>2</sub> from Flue-Gases: Experiments with Phase-Changing Solvents in a Bench-Scale Plant

Cruz, M. de A. et al. (2019) **SDEWES2019.0276 Chemical Absorption of CO<sub>2</sub> from Flue-Gases: Experiments with Phase Changing Solvents in a Bench Scale Plant**, 14th Conference on Sustainable Development of Energy, Water and Environment Systems. Edited by A. Mudrovčić and M. Ban. Zagreb: Faculty of Mechanical Engineering and Naval Architecture. Available at: <https://www.dubrovnik2019.sdewes.org/>.

#### Abstract

The energy penalty of solvent regeneration is a barrier for the deployment of chemical absorption post-combustion carbon capture. Phase-changing absorption solvents have been proposed to overcome this issue. CO<sub>2</sub> absorption triggers phase separation with only the CO<sub>2</sub>-rich phase requiring regeneration, potentially reducing energy demand. A systematic review supports the choice of a set of solvents for experimental investigation, based on economic, process and energy-related criteria. Until now the selected absorbents were only investigated in laboratory scale. They need to move from lab to industrial scale to contribute to the global warming mitigation. Three selected biphasic solvents are selected and tested, to confirm the results reported by its developers. Solvent A, based on monoethanolamine/1-propanol, was considered the more suitable one. This blend presented 26% of reduction on the lower (CO<sub>2</sub>-rich) liquid phase, compared to the initial volume of fresh solvent. monoethanolamine is the more traditional chemical absorbent for CO<sub>2</sub> capture applications and both components can be considered standard chemicals (low-cost), which is a remarkable advantage. The viscosity of the CO<sub>2</sub>-rich phase of solvent A is 10 mPa.s at 25°C, which is considered low compared to other candidates. Solvent B, based on diethylene-triamine and N,N,N',N'',N'''-pentamethyldiethylenetriamine, was disregarded for further evaluations. It presented only 10% of volume reduction on CO<sub>2</sub> rich phase and prohibitively high viscosity (360 mPa.s at 40°C). Furthermore, chemical components of this solvent are considered specialty (high cost).

Solvent C, based on N-methylcyclohexylamine and N,N-dimethylcyclohexylamine, was also disregarded for further tests. Although the costs of its chemical components are in the same range of monoethanolamine and the CO<sub>2</sub>-rich liquid phase presents an acceptable viscosity (58 mPa.s at 25°C), this blend had an issue. It presented solids precipitation on the CO<sub>2</sub> rich phase, what is a potential source of operational problems on industrial application. Solvent A was the only one tested on a bench scale screening plant, designed for absorption and desorption of chemical absorption solvents. This blend presented a CO<sub>2</sub> loading of 2.8 mol/kg on the lower phase, 76% higher than MEA 30%. It is an opportunity to reduce the energy penalty of carbon capture of CO<sub>2</sub> by chemical absorption. The solvent A was selected to further evaluation on a continuous mode pilot-plant (under construction).

**Keywords:** Post-combustion carbon capture, phase change solvents, biphasic solvents, chemical absorption pilot plant, combustion exhaust gases.

## Nomenclature

<i>AMP</i>	2-amino-2-methyl-1-propanol
<i>CCS</i>	Carbon Capture and Storage
<i>DEEA</i>	Diethylaminoethanol
<i>DETA</i>	Diethylene-triamine
<i>DMCA</i>	N,N-dimethylcyclohexylamine
<i>LLPS</i>	Liquid-liquid phase separation
<i>LPST</i>	Liquid phase separation temperature
<i>MAPA</i>	N-Methyl-1,3-Propanediamine
<i>MCA</i>	N-methylcyclohexylamine
<i>MEA</i>	Monoethanolamine
<i>PCAS</i>	Phase-changing absorption solvents
<i>PCASP</i>	Phase-change absorption screening plant
<i>PMDETA</i>	N,N,N',N'',N'''-pentamethyldiethylenetriamine
<i>TBS</i>	Termomorphic biphasic solvent

### 7.1.1. Introduction

Paris Agreement established a compromise to keep the global warming below 2°C to the end of this century. It imposes at least 70% emissions reductions through 2050, compared to 2010

levels. (IPCC, 2014b) Carbon Capture and Storage (CCS) is expected to account for 14% (140Gt) of this target, considering the power and industrial sectors. (Global CCS Institute, 2017) Nevertheless, the pace of CCS development is not enough to achieve this goal. (IEA, 2016) Cost is a remarked hindrance to CCS deployment on industries where CO<sub>2</sub> separation step is not inherent to the process. CCS could bring up to 70% addition on lifecycle cost of production for power generation, 68% for cement production and 41% for steel manufacturing. (Global CCS Institute, 2017)

Major CCS costs come from the energy penalty of capture and compression processes. Usually, CO<sub>2</sub> capture corresponds to 65% - 80% of this penalty on power plants. (Goto; Yogo; Higashii, 2013) Chemical absorption with alkanolamines is a mature post-combustion capture process for exhaust gases with low CO<sub>2</sub> partial pressure (4 – 30 kPa). Gas scrubbing with an aqueous solution of monoethanolamine (MEA) is the benchmark technology. (Rochelle, 2009; Rubin; Chen; Rao, 2007) The first process was proposed by Bottoms (1930) and imposes an energy consumption of 3.7 GJ/ton of CO<sub>2</sub> captured. (Knudsen et al., 2009) Major energy demand comes from desorption of CO<sub>2</sub> from the solvent. Usually, heat is supplied by low-pressure steam, from boilers or steam cycles. As a result, the net energy efficiency of those processes is impaired. For each GJ/ton CO<sub>2</sub> reduced, 2% net efficiency improvement is achieved on coal-fired power plants. (Goto; Yogo; Higashii, 2013). Additionally, low CO<sub>2</sub> absorption capacity per mol of solvent (loading), thermal and oxidative degradation and corrosivity are frequently reported as weaknesses of MEA. (Zhang et al., 2013)

Process improvements and optimization on conventional amine scrubbing have been exhaustively studied. The energy penalty of Bottom's process cannot be considered a reference nowadays. (Boot-Handford et al., 2014) Frailie et al. (2013) evaluated inter-stage cooling and heating on absorber and stripper columns, respectively. Li et al. (2011) investigated heating integration and exhaust gas recycle. Park et al. (2016) performed an optimization study concerning stripper pressure. Rochelle and collaborators deeply investigated absorber performance (Zhang; Rochelle, 2014) and intercooling (Rezazadeh et al., 2014, 2017). Commercial processes applying state-of-the-art technology combines process optimization and advanced solvents. KM-CDR process reached an energy penalty of 2.11 GJ/ton of CO<sub>2</sub> captured, about 43% lower than early MEA scrubbing processes. (Miyamoto et al., 2017) Nevertheless, the energy penalty of carbon capture by chemical absorption still

being a barrier for its deployment. However, there are few opportunities for energy saving and lifecycle cost reduction of CCS by amine scrubbing.

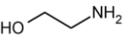
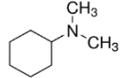
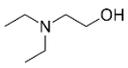
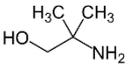
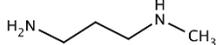
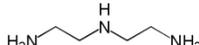
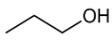
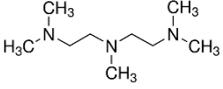
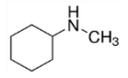
#### 7.1.1.1. Phase-changing absorption solvents

Phase-changing absorption solvents (PCAS) emerged as a real opportunity to cut the energy penalty of carbon capture. Most PCAS are called thermomorphic, which means that they present a lower critical CO<sub>2</sub> loading that is a function of temperature. After saturated in CO<sub>2</sub> and above a certain temperature, liquid-liquid phase separation (LLPS) occurs. (Budzianowski, 2016) Preferably, one phase must be rich and other lean on CO<sub>2</sub> (Raynal et al., 2014) and liquid phase separation temperature (LPST) must be higher than absorption temperature. Thereby, the CO<sub>2</sub> lean phase is recycled to the absorber and only the rich phase is sent to desorption. It entails reduced circulation rate and reboiler heat duty. (Coulier et al., 2017; Liebenthal et al., 2013) Some PCAS presents lower desorption temperature, enabling the use of low-grade heat. Higher desorption pressure is another possibility, reducing CO<sub>2</sub> compression penalty.

Early PCAS was already demonstrated in pilot scale. DMX process (Raynal et al., 2014) reported a specific reboiler heat consumption of 2.5 GJ/t of CO<sub>2</sub>, 19% reduction on energy penalty and 20% lower cost of CO<sub>2</sub> avoided, compared to MEA runs. BiCAP process (Lu, 2017) reported 34% of energy penalty reduction and 50% reduction on the cost of CO<sub>2</sub> avoided. Both DMX and BiCAP used undisclosed proprietary blends of PCAS. DEEA (Diethylaminoethanol) and MAPA (N-Methyl-1,3-Propanediamine) aqueous solutions was tested in Gløshaugen (NTNU/SINTEF) pilot plant. The average reboiler duty was 2.3 GJ/t of CO<sub>2</sub> (Pinto et al., 2014), 30% lower than the benchmark (MEA).

Recently, new PCAS (Barzagli; Mani; Peruzzini, 2017; Wang et al., 2017; Zhang et al., 2013, 2017a, 2017b; Zhou et al., 2017) emerged, aiming to overtake weaknesses of early ones, notably: proprietary blends, low net CO<sub>2</sub> loading, LPST below absorption temperature, high cost of employed unconventional amines and high viscosity of loaded solution. This work selected three PCAS in the early stage of development from literature. Table 7.1 lists retail prices of all components of selected PCAS, for comparison.

Table 7.1. Retail price of PCAS chemicals (Sigma-Aldrich, 2018)

Component	Structural Formula	Price (US\$/liter)	Component	Structural Formula	Price (US\$/liter)
MEA		51.00	DMCA		74.60
DEEA		54.50	AMP		56.50
MAPA		297.50	DETA		63.40
1-propanol		79.50	PMDETA		270.00
MCA		383.65			

The first blend is MEA/1-propanol/H<sub>2</sub>O, claimed by Zhang et al. (2017a, 2017b) as a promising PCAS, because it is based on MEA, the most common CO<sub>2</sub> absorbent and 1-propanol, an ordinary alcohol. 1-propanol presents affordable prices, in the same order of magnitude of MEA (see Table 7.1). The author performed tests in a simple lab absorption apparatus. Basically, pure CO<sub>2</sub> was bubbled in an Erlenmeyer flask, filled with the absorption solvent. Preliminary results confirmed LPST at around 30°C. The volume of rich phase could be reduced up to 67% of the initial volume of solvent with loading up to 2.6 molCO<sub>2</sub>/kg of solvent, 30% higher than MEA 30% (w/w). The author suggested that more tests, in pilot scale, are required to demonstrate the suitability and potential for energy penalty reduction of this PCAS.

The second was developed by Zhou et al. (2017), being composed of diethylene-triamine (DETA) and N,N,N',N'',N'''-pentamethyldiethylenetriamine (PMDETA). That blend is not cost attractive as the first one, especially because PMDETA price (see Table 7.1). But bulk orders are available for both components, with much more affordable prices. The advantage is that, usually, none of them needs special permits of army or police to be purchased in large quantities. Another advantage is the improved loading (0.613 mol CO<sub>2</sub>/mol amine), 21% higher than MEA and 26% higher than DEEA/MAPA. LPST is 50°C, within the temperature range of flue-gases from power plants. The volume of rich phase is 57% of total solvent volume. The author reported 2.3 GJ/t of CO<sub>2</sub> (reduction of 38% compared to MEA). Once

again, tests were performed in small glass apparatus on the lab. Pilot scale experiments are required, to prove the advantage of using this PCA in full-scale CCS applications.

Lastly, Zhang et al. (2013) studied a so-called TBS (termomorphic biphasic solvent), that mixes N-methylcyclohexylamine (MCA) as an absorption activator, N,N-dimethylcyclohexylamine (DMCA), as regeneration promoter and amine 2-amino-2-methyl-1-propanol (AMP), as a solubilizer to increase LPST. This PCAS do not use expensive amines, as shown in Table 7.1, and performs well in terms of CO<sub>2</sub> loading (3.5 molCO<sub>2</sub>/kg – 75% higher than MEA). LPST is between 30°C - 40 °C. Desorption occurs at 80 °C, enabling the use of low-grade heat. Experiments were performed in bench scale plant but used glass columns of only 0.04 mm inner diameter at atmospheric pressure. The author reported 2.0 GJ/t of CO<sub>2</sub> (reduction of 46% compared to MEA). However, this result could be unrealistic, because of wall effects and other consequences of such small-scale plant.

PCAS with high viscosity could be a challenge for pumping systems. Zhang et al. (2013) reported viscosities of 16 mPa.s for loaded TBS-3, what is higher than MEA (3.3 mPa.s) but lower than DETA/PMDETA (250 mPa.s (Zhou et al., 2017)) and MEA/1-propanol (60 mPa.s (Zhang et al., 2017b)). The effect of viscosity on the full-scale system must be considered on choosing the more suitable PCAS.

#### 7.1.1.2. Solvent Testing Plant (Batch)

Technical, economic and environmental performance of large-scale post-combustion processes with selected PCAS are necessary to prove the advantage of such solvents over early ones. Move from lab to pilot scale plants is the first step to achieve this goal. In this work, a bench scale screening plant is constructed to the preliminary evaluation of selected PCAS. Based on lab preliminary tests and techno-economic criteria, suitable PCAS are tested on the screening plant. Based on the results of this work and previous pilot plant studies, a continuous mode pilot plant will be designed. Data from experiments in both, bench and pilot-scale plants, will support mathematical modeling and computational simulation of full-scale CCS using PCAS.

The absorption capacity of solvents is affected by testing methods, solvent composition, temperature (of gas and solvent), pressure, and gas composition. (Huertas et al., 2015) To reproduce the results of previous studies, the bench scale plant is designed to run experiments

at the same conditions used by authors. In future tests, a more realistic standard condition must be set, to enable comparison between different PCAS. Therefore, the bench plant must be able to work with a wide range of CO<sub>2</sub>, O<sub>2</sub> and N<sub>2</sub> flows (gas composition), temperatures and pressures. It is also important to keep the solvent temperature constant along with the experiment, using a process thermostat.

Although showing promising results, most previous studies were performed at lab conditions (ambient temperature and pressure, small glass apparatus and bubbling pure CO<sub>2</sub> into a few milliliters of solvent). However, higher temperature (40 - 70°C) low CO<sub>2</sub> partial pressures (5 – 30 kPa) and oxygen (1% - 15%) are found in flue-gases from power stations, gas-turbines, furnaces, refinery, cement and steel plants (Ulrich, 2010). As a result, the loading reported at lab conditions, by previous studies, could be overestimated compared to real conditions. Consequently, the overall energy penalty of the process is underestimated.

Oxygen irreversibly reacts with amines along the time, reducing its CO<sub>2</sub> absorption capacity. (Huertas et al., 2015) Considering aqueous solutions concentrations between 12% and 42% (w/w) the process is controlled by kinetics and O<sub>2</sub> mass transfer. However, MEA loss due to oxidative degradation is in the range of 0.29 – 0.73 kg per ton of CO<sub>2</sub> captured in large scale processes. (Thong et al., 2012) Therefore, this effect is expected to be negligible for batch experiments with virgin PCAS, but relevant for the continuous mode pilot plant.

Low CO<sub>2</sub> partial pressures and higher temperature reduces the capture efficiency of the absorption process. Lower efficiency means increased saturation time for batch experiments and higher absorbent flux (reboiler duty) and column size for continuous process. Once again, CO<sub>2</sub> partial pressure is not so relevant parameter for batch experiments, but very important to a continuous process.

The main objective of this work is evaluating selected PCAS, in terms of CO<sub>2</sub> absorption capacity and rate, volume ratio and viscosity of upper and lower liquid phases. Besides process and energetic matters, economic, environmental and safety inherent characteristics of PCAS components will be taken into account on solvent choice. Results support continuous pilot-plant design and operation. The more suitable PCAS is selected to a trial test and further studied on a batch pilot plant and, in the future, tested at a continuous mode pilot-plant (under construction).

## 7.1.2. Methodology

### 7.1.2.1. Preliminary experiments

Preliminary absorption experiments (Figure 7.1) are performed with the three selected PCAS, at the lab, to observe the phase-change behavior and properties (density and viscosity) of liquid phases. Tests were executed at ambient temperature (around 25 °C), based on previous studies, according to Table 7.2. These preliminary tests consisted of bubbling a pure CO<sub>2</sub> gas stream into a recipient containing 50 ml of solvent, as shown in Figure 7.2. The gas flow is kept constant until there is no more difference between the absorption liquid and the ambient temperatures. It means that the solvent is already saturated because the absorption reaction is exothermic. The required chemical components to formulate the PCAS are gathered in Table 7.1. MEA was purchased from Oxiteno S.A, 1-propanol (99.5%) from Isofar, MCA, DMCA, DETA, PMDETA, and AMP were obtained from Merck (Sigma-Aldrich, Inc).



Figure 7.1. Absorption Preliminary Tests Apparatus

Table 7.2. Selected phase-change absorption solvents - composition

PCAS	Components	Total Amine			Reference
		Mole Concentration	Components Mole Concentration		
A	MEA/1-propanol	5.00	5.00	: 5.35	(Zhang et al., 2017a)
B	DETA/PMDETA	5.00	4.00	: 1.00	(Zhou et al., 2017)
C	MCA/DMCA/AMP	5.50	1.00	: 3.00 : 1.50	(Zhang et al., 2013)

Feed gas flow and composition used on absorption experiments are shown in Table 7.3.

Table 7.3. Absorption tests – feed gas parameters

T	P	Gas Vol. Flow (NLPM)		CO <sub>2</sub> Partial Pressure
		Air	CO <sub>2</sub>	(kPa)
(°C)	(kPa)			
25	101.3	-	0.50	101

Preliminary absorption experiments are performed to enable comparison between results obtained by this work and reference ones. It employs a standard gas composition, without O<sub>2</sub>, SO<sub>x</sub>, and NO<sub>x</sub>. Such contaminants could react with amines and modify some results.

### 7.1.2.2. Phase-Change Absorption Screening Plant (PCASP)

The PCASP is shown in Figures 7.2 and 7.3. It is a bench unit, mainly comprised of two bubble columns (V-02 and V-03, with 0.1 m diameter and 0.675 m of height), where absorption and desorption PCAS tests take place. A supervisory control and data acquisition (SCADA) system (Elipse E3) is used to control and record temperatures, pressures, flow rates, and gas CO<sub>2</sub> concentrations at strategic points of the system. Gas cylinders (from Linde Gases LTDA.) supplies CO<sub>2</sub> (99.8% at 5800 kPa) and Nitrogen (N<sub>2</sub>) (99.8% at 20000 kPa). Dry compressed air can be used to adjust Oxygen (O<sub>2</sub>) composition of the feed gas. The flows of all inlet gases are measured by thermal mass flow controllers (Brooks SLA 5800 series). Standard flow is double checked by local rotameters. Maximum standard flows of CO<sub>2</sub>, air, and N<sub>2</sub> are 3 normal liters per minute (NLPM), 10 NLPM and 15 NLPM, respectively. The CO<sub>2</sub> pressure regulator and the downstream pipe has an electrical trace, enabling gas heating and avoiding frozen. Streams of CO<sub>2</sub>, air, and N<sub>2</sub> are mixed and follows to vessel V-01. If a humid and warm gas is desired (to simulate a flue-gas from a real power plant), V-01 must be partially filled with water and its immersed electrical resistance turned on. The feed gas temperature can be set in the range of 30°C – 50°C. Experiments can be realized between 101.3 and 500 kPa, resulting in feed gas CO<sub>2</sub> partial pressures in the range of 5 – 500 kPa.

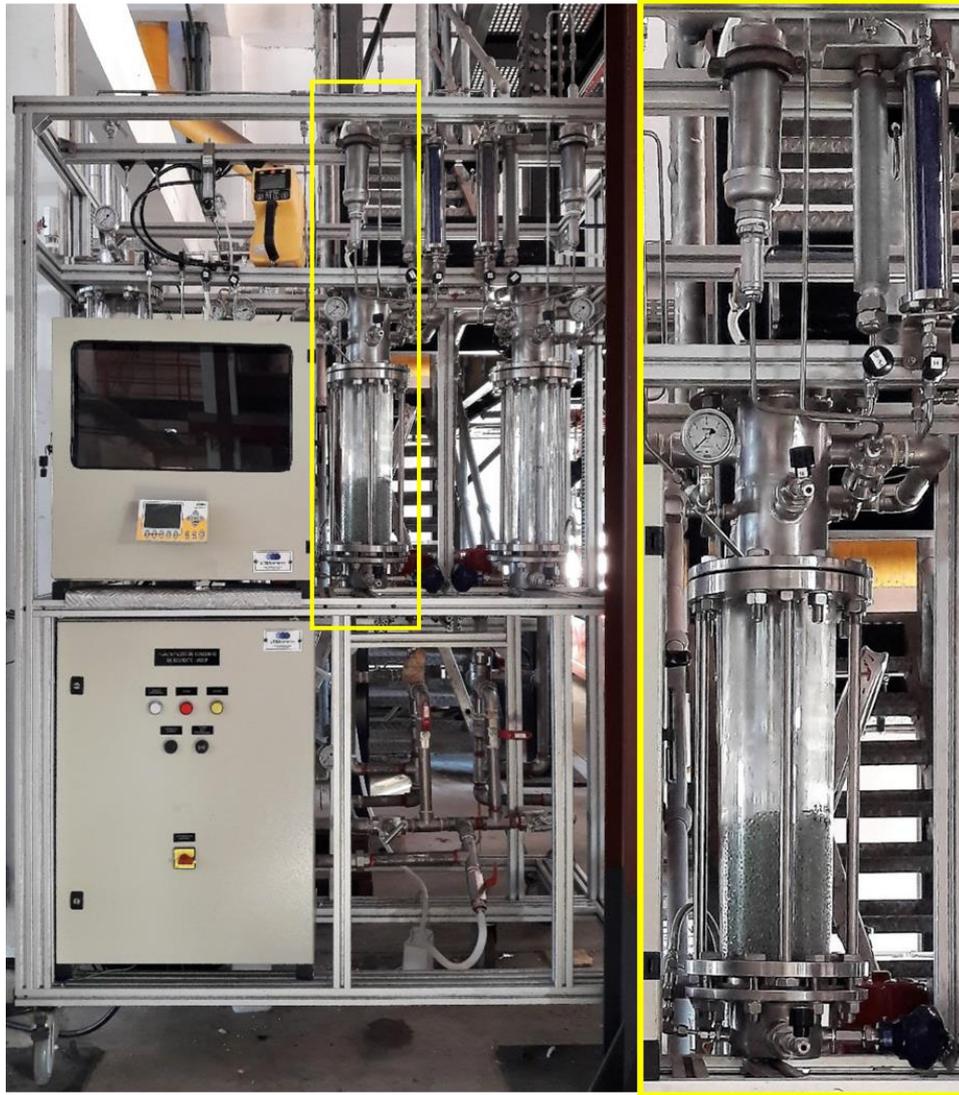


Figure 7.2. Absorption & Desorption Screening Plant

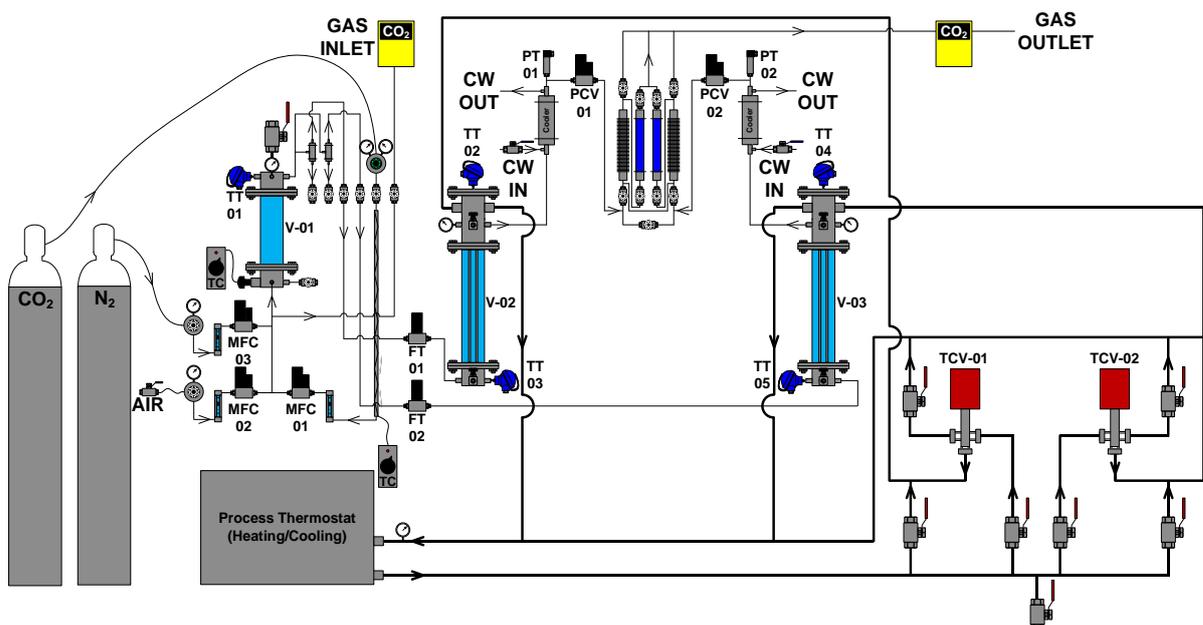


Figure 7.3. Absorption & Desorption Screening Plant Flowsheet

Two different solvents can be tested simultaneously. Each vessel (V-02 and V-03) can receive up to 4 liters of solvent, considering the maximum level as half of the vessel height). Filled with 5 kg of 5 mm glass beads, the inventory of solvent is reduced to 1.5 liters of solvent. These vessels have internal coils. Inside these coils a thermal fluid (Kryo 65) flows, coming from a process thermostat (Lauda Integral XT-150). This equipment enables to keep the temperature constant during absorption (cooling mode) and desorption (heating mode). TCV-01 and TCV-02 are 3-way temperature control valves (Badger Meter RCV type 1118 with electronic actuator model EVA-1). They adjust the thermal fluid flow in each coil, enabling individually set the temperature inside V-01 and V-02. Each vessel has a water condenser and a silica gel filter after gas outlet, to reduce solvent loss and avoid liquids and humidity into CO<sub>2</sub> sensors.

### 7.1.2.3. Absorption Experiment Using the PCASP

Procedure and Setup:

Solvent temperature set-point: 30°C during absorption; 40 °C after solvent saturation

- Absorption vessel pressure (V-02): 101.3 kPa
- Gas composition: 16.7% CO<sub>2</sub> + 83.3% dry air (9 min); 66.7% CO<sub>2</sub> + 33.3% dry air (10 min); 100% CO<sub>2</sub> (28 min)
- CO<sub>2</sub> flow: 2.0 l/min
- Airflow: 10.0 l/min (9 min); 1.0 l/min (10 min). Air was used to increase the feed gas flow, promoting heat and composition homogenization inside absorption vessel (V-02)
- Solvent: 1.5 liters of solvent A (5M MEA/6M 1-propanol)

The CO<sub>2</sub> absorption rate was calculated by the difference between the inlet and outlet gas concentration as proposed by Wang et al. (2013), according to Equation (7.1):

$$r_{CO_2} = \frac{273.15 \times P Q_g (C_{in} - C_{out})}{101.3 \times 22.4 T} \quad (7.1)$$

Where  $r_{CO_2}$  is the CO<sub>2</sub> instantaneous absorption rate (mol/min), P is the absorption pressure (kPa),  $Q_g$  is the CO<sub>2</sub> standard flow rate (l/min),  $C_{in}$  and  $C_{out}$  are the CO<sub>2</sub> concentrations on inlet and outlet gas streams (% vol.) and T is the absorption temperature (K).

The CO<sub>2</sub> concentration of the feed and lean gas can be continuously monitored by in-line infrared sensors (Witt model PA 7.0). CO<sub>2</sub> loading in liquid phases is analyzed by the Chittick method. (Zhang et al., 2018) The viscosity of liquid phases is measured using a viscometer (Brookfield DV1). Density is determined by Mettler Toledo DM40 density meter.

### 7.1.3. Results and Discussion

#### 7.1.3.1. Preliminary tests

The three tested PCAS candidates (A, B and C) formed two phases after being loaded with CO<sub>2</sub>. This is the first criterion that must be regarded before considering a test on the PCASP.

Although it was confirmed the formation of two phases after CO<sub>2</sub> absorption by solvents A, B, and C, there was a 26% reduction in the volume of the lower (supposedly CO<sub>2</sub> rich) phase of solvent A, 10% in the lower phase of the solvent B and 20% of solvent C. The modest volume reduction of solvent B would lead to negligible reduction in the energy penalty of regeneration. The viscosity of each phase of the solvents was also analyzed, as shown in Figure 7.5. High viscosity could be a challenge for pumping systems and must be considered when aiming a full-scale application.

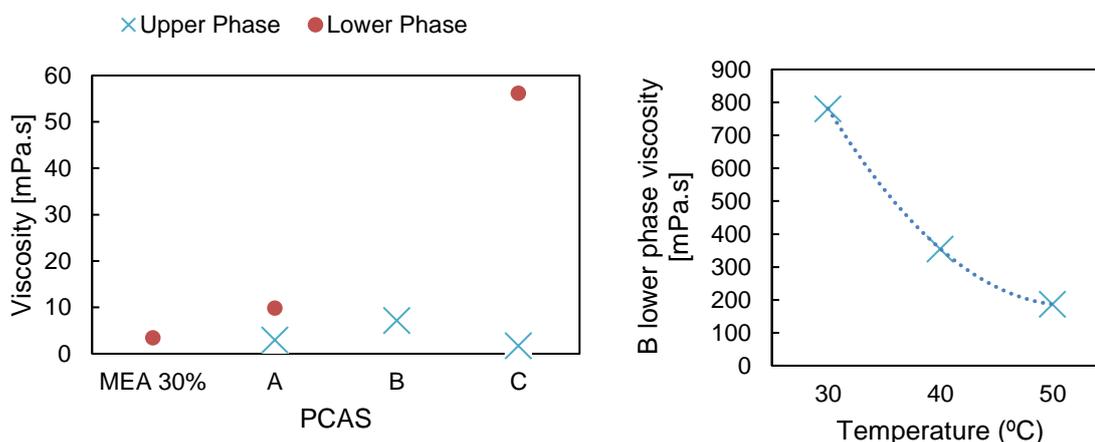


Figure 7.5. Viscosity of liquid phases after CO<sub>2</sub> absorption at 25°C – (A) 5M MEA/4M 1-propanol; (B) 4M DETA/1 M PMDETA; (C) 1M MCA/3M DMCA/1.5M AMP. Viscosity versus temperature of solvent B.

Solvent B presented high viscosity for the lower phase, exceeding the maximum value supported by the viscometer (1200 mPa.s) at 25 °C. Heating was required to enable a measure of lower phase viscosity, as shown in Figure 7.6. The viscosity only reaches the values reported by Zhou et al. (2017) (250 mPa.s) above 50°C. This viscosity is prohibitive because

brings operation and maintenance issues on industrial application. This characteristic together with the low reduction of volume of the CO<sub>2</sub> rich phase resulted in the elimination of solvent B from further experiments on PCASP.

The viscosity of the lower phase of solvents A and C was around 183% and 1500% greater than the reference (30% MEA), respectively. The slightly higher viscosity of A could imply marginal impacts on the overall energy penalty of the process, but this effect could be more representative for solvent C. Besides presented a moderate viscosity, crystals precipitation was noted in solvent C after standing at ambient temperature. The precipitate can bring operational problems – e.g. clogging of liquid dispersers and process pipes. For this reason, the use of solvent C may not be convenient on plants designed for liquid-liquid PCAS. Another negative point of C was difficult on visualizing the liquid-liquid interface, due to the color similarity of the two phases. For these reasons, solvent C was not tested on the PCASUS.

#### 7.1.3.2. Absorption Experiments on the PCASP

The solvent started to become saturated in CO<sub>2</sub> after 40 minutes of absorption. This event was visually observed by the increased turbidity of the solvent inventory, by the gas temperature drop (Figure 7.6) and by the increase in CO<sub>2</sub> concentration at the gas outlet (Figure 7.7). The solvent was considered fully saturated after 46 minutes (Figure 7.7) when the liquid became totally turbid and the CO<sub>2</sub> concentration at the outlet reached 100%. At this stage, the set-point of the temperature control valve (TV-01) was modified to 40 °C. After 70 minutes, when the liquid temperature reached 34 °C, the phase split began, and the gas feed was interrupted to stop the stirring and enable the total liquid-liquid separation. The turbid interface gradually narrowed until becoming a line dividing the two liquid phases (Figure 7.6). At the end of the experiment, samples of the upper and lower phases were collected. The CO<sub>2</sub> loading, density, and viscosity of each phase were analyzed, and results are shown in Table 7.4. The instant CO<sub>2</sub> absorption rate was calculated according to Equation 7.1 and is shown in Figure 7.7.

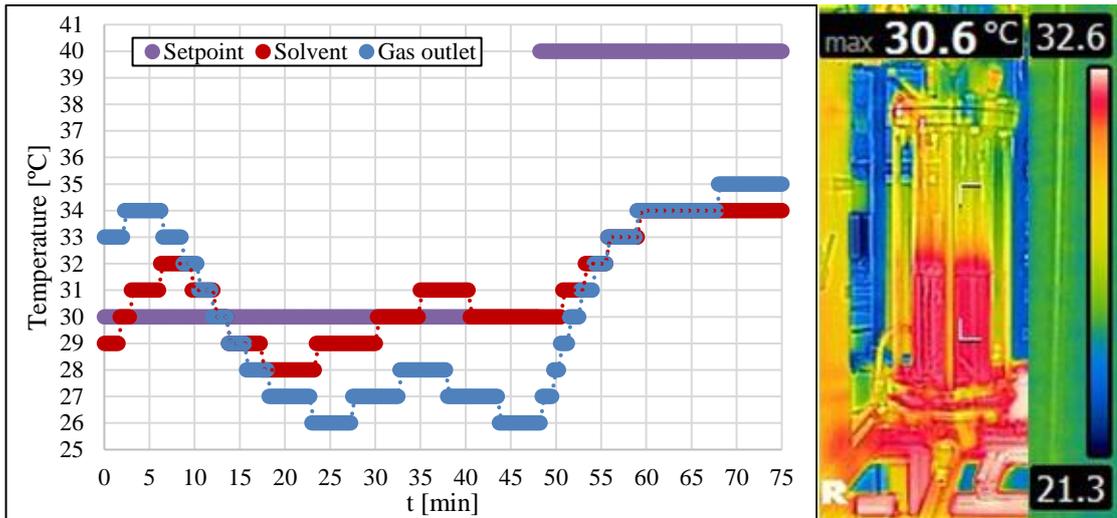


Figure 7.6. Average solvent A (5M MEA/6M 1-propanol) temperature along with the test at PCASP (left side). Thermal image of the vessel during the experiment (right side).

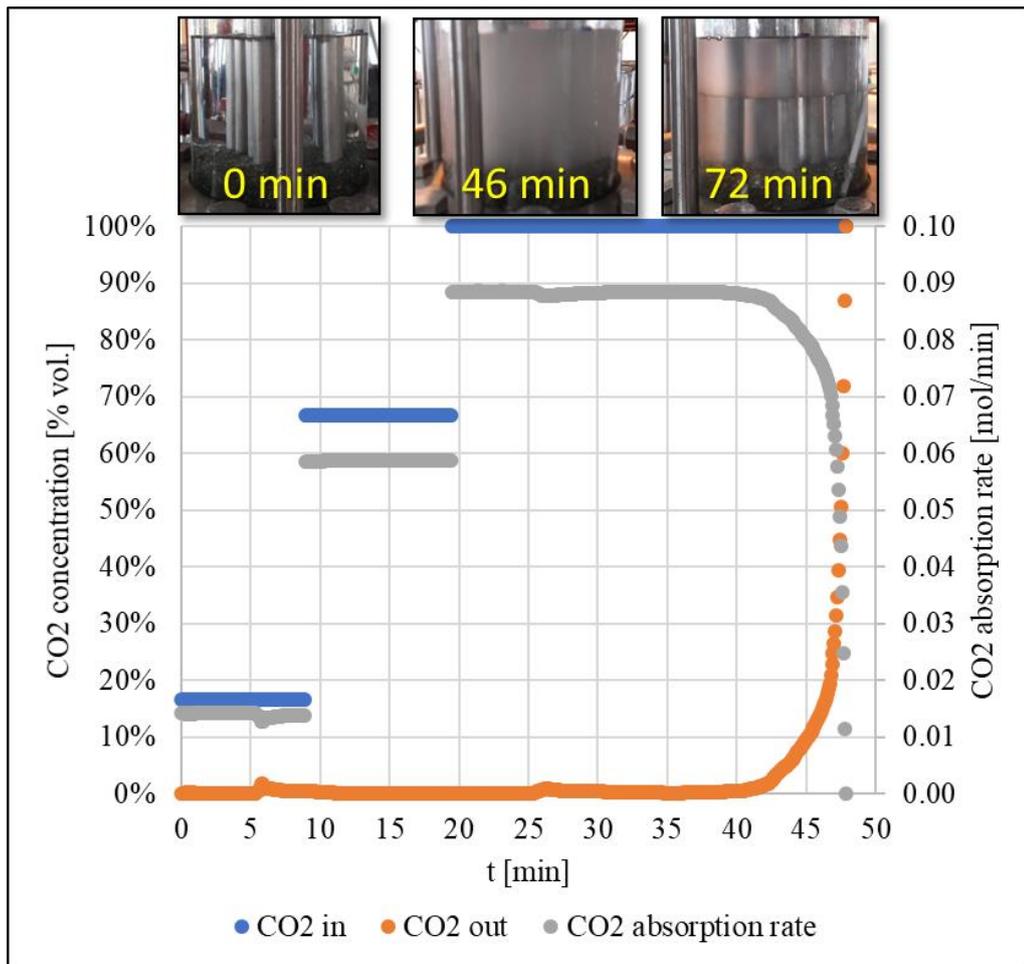


Figure 7.7. Gas CO<sub>2</sub> concentration and absorption rate of solvent A (5M MEA/6M 1-propanol) along with the test at PCASP

Table 7.4. Main results of the test of solvent A (5M MEA/6M 1-propanol) on the PCASP compared with results of ZHANG et al. (2017a).

Property	Lower Phase			Upper Phase		
	This work	Ref.*	Diff.	This work	Ref.*	Diff.
<b>Density (g/cm<sup>3</sup>)</b>	1.11	1.06	5.1%	0.86	0.79	8.9%
<b>Viscosity (cP)</b>	10	12	-16.7%	3	N/A	N/A
<b>CO<sub>2</sub> loading (mol/kg)</b>	2.80	3.00	-6.5%	0.47	0.49	-4.6%
<b>Volume (ml)</b>	1050	-	-	450	-	-

\*(Zhang et al., 2017a)

The CO<sub>2</sub> loading, densities, and viscosities are in agreement with the results reported by Zhang et al. (2017a). According to the present study, the loading of solvent A is 76% higher than MEA 30%. Measure the energy penalty reduction of the carbon capture process depends on further desorption experiments, that enables to determine the cyclic capacity of the solvent. Nevertheless, it is possible to infer that solvent A would bring energy savings and reduced stripper footprint when applied to a carbon capture process. The energy penalty benefit comes from the increased loading of CO<sub>2</sub> per mol of solvent that is sent to the stripper. If the CO<sub>2</sub> loading of the lower phase were the same as MEA 30% solutions, the reduction on the initial volume of solvent would have no consequence on the regeneration energy. The reduction of the lower phase (30%) compared to the initial volume of solvent was almost the same volume of 1-propanol added to the MEA 30% solution. According to Wang et al., (2019), the only advantage off add 1-propanol to the solvent is that it works as a physical solvent, increasing the diffusivity of the CO<sub>2</sub> and consequently its solubility and absorption rate. After reach a critical loading, carbamates encircled by water expels 1-propanol molecules, forming another liquid phase. This effect is named salting-out.

### 7.1.3.3. Potential Energy Savings from the use of PCAS (solvent A)

The presence of 1-propanol (even in low concentration) on the lower phase is an advantage. Alcohols have a low dielectric constant, promoting solvent regeneration. (Zhang et al., 2017a).

Simulating a capture process using the same PCAS considered in this study (MEA 30%/1-propanol 40% w/w) Wang et al. (2019) reported an energy penalty of 2.87 GJ/t, setting a pressure of 200 kPa at the stripper, which resulted in a temperature of 127 °C on the reboiler. The energy saving estimated by Wang et al. (2019) is 28% lower than the baseline (MEA 30% - 3.99 GJ/tCO<sub>2</sub>). Disregarding simulation constrains and accuracy, the major source of

energy penalty is the volume reduction coming from the split of the rich liquid phase considered by Wang et al. (2019) and also reported in the present study. The author reported a 43.6% volume fraction of the upper phase against the total income solution while in the experiments performed in the present study this fraction is 30%. This factor influence MEA concentration, CO<sub>2</sub> loading on rich phase and the flow rate of the solvent sent to the stripper, which directly influences the reboiler duty and temperature and stripper CO<sub>2</sub> recovery.

The feed gas of a NGCC flue-gas has a CO<sub>2</sub> partial pressure ranging around 5 to 10 kPa. Wang considered a partial pressure of 16 kPa (compatible with PCC flue-gas), which makes the chemical absorption process less intensive on energy. Using a NGCC flue-gas entails a higher solvent per CO<sub>2</sub> flow ratio (capture ratio) increasing the energy penalty.

#### 7.1.4. Conclusion

This study investigated the state-of-the-art on PCAS. Three blends of solvents were selected and submitted to preliminary tests. One of them was elected to be tested on a batch screening plant, named PCASP, designed to evaluate biphasic solvents properties and phase change behavior during absorption and desorption of CO<sub>2</sub> coming from the synthetic exhaust gas. The test confirmed that using PCAS based on MEA and 1-propanol is an opportunity to reduce the energy penalty of carbon capture of CO<sub>2</sub> by chemical absorption.

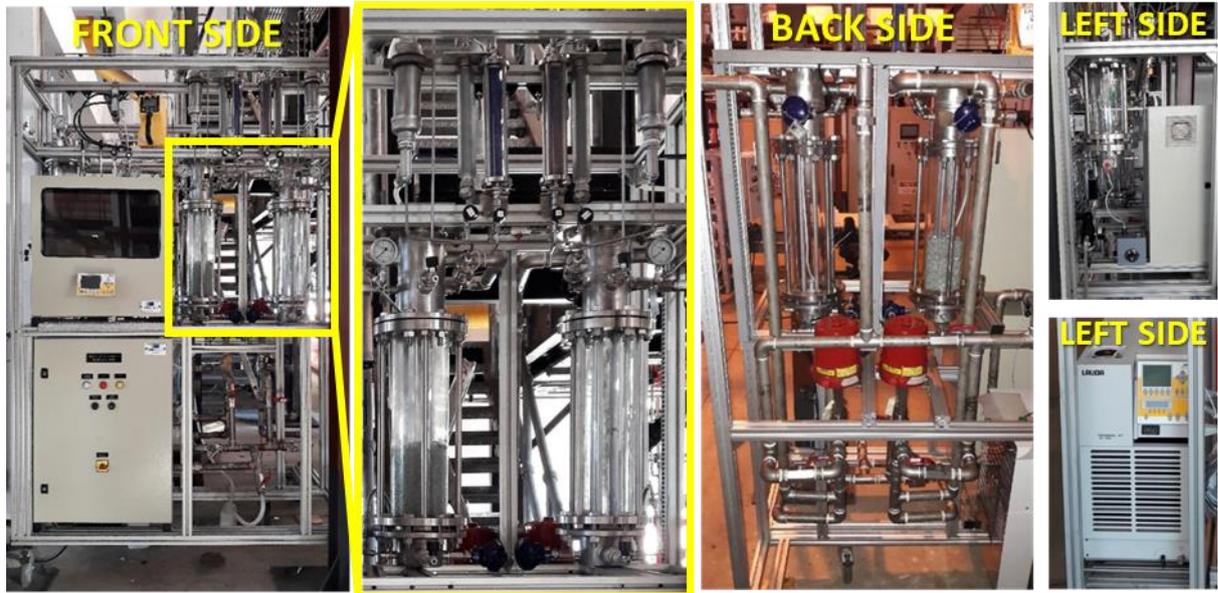
The next step is testing the best PCAS under different compositions and absorption conditions, simulating a more realistic flue-gas (higher temperature, pressure, presence of contaminants and lower CO<sub>2</sub> partial pressure). Desorption experiments also must be executed to confirm the cyclic capacity and comparison with results reported by other authors. Results will serve as a reference to the design of a pilot plant that will expose the solvent to a continuous process and long runs, in order to evaluate more precisely the energy penalty, degradation of the solvent and other process parameters. This information is necessary to validate the simulation and scale-up of a full-scale application of a PCAS.

## **7.2. Pilot Plants Developed for Testing CO<sub>2</sub> Capture with Phase-Changing Absorption Solvents**

A pilot plant was designed to perform long-run experiments with PCAS, to up scaling the first plant, PCASP, presented in section 7.1 – the Phase-Changing Absorption Pilot Plant (PCAPP). This section describes the two plants.

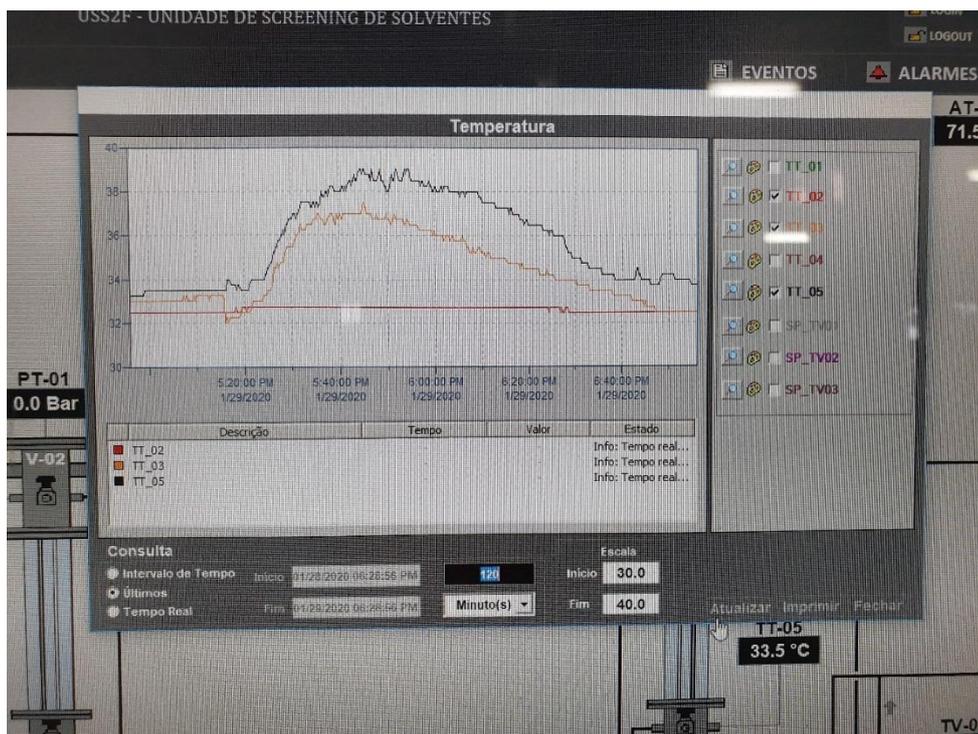
### 7.2.1. Phase-Changing Absorption Solvents Screening Unit - PCASP

The PCASP was presented in section 7.1 and is detailed in this section. The construction of the plant lasted six months and was finished in January 2018. Fig.7.8 shows the unit.



**Figure 7.8. PCASP Overview**

On the front side of the plant, the control and automation panels and the two main vessels are visualized. On the opposite side, are installed the temperature control valves and thermal fluid piping. On the left side, the process thermostat and the gas heating and water saturation vessel are placed. The commissioning and startup of the plant were performed in May 2018 and experiments were performed along 2018 and 2019. A 5M MEA solution was used to perform an absorption test with a CO<sub>2</sub>-rich feed gas. The plant provided a capture efficiency >90%. The skid has three glass absorption vessels. Each vessel has a temperature sensor (PT-100) and a mechanic pressure gauge. The inlet and outlet gas flows are connected to the CO<sub>2</sub> analyzers. Temperature, gas flows and CO<sub>2</sub> concentration trends were registered at the SCADA of the PCASP, as shown in Fig. 7.9.



**Figure 7.9. Temperature trends in typical PCAS experiment.**

### 7.2.2. Phase-Changing Absorption Pilot-Plant - PCAPP

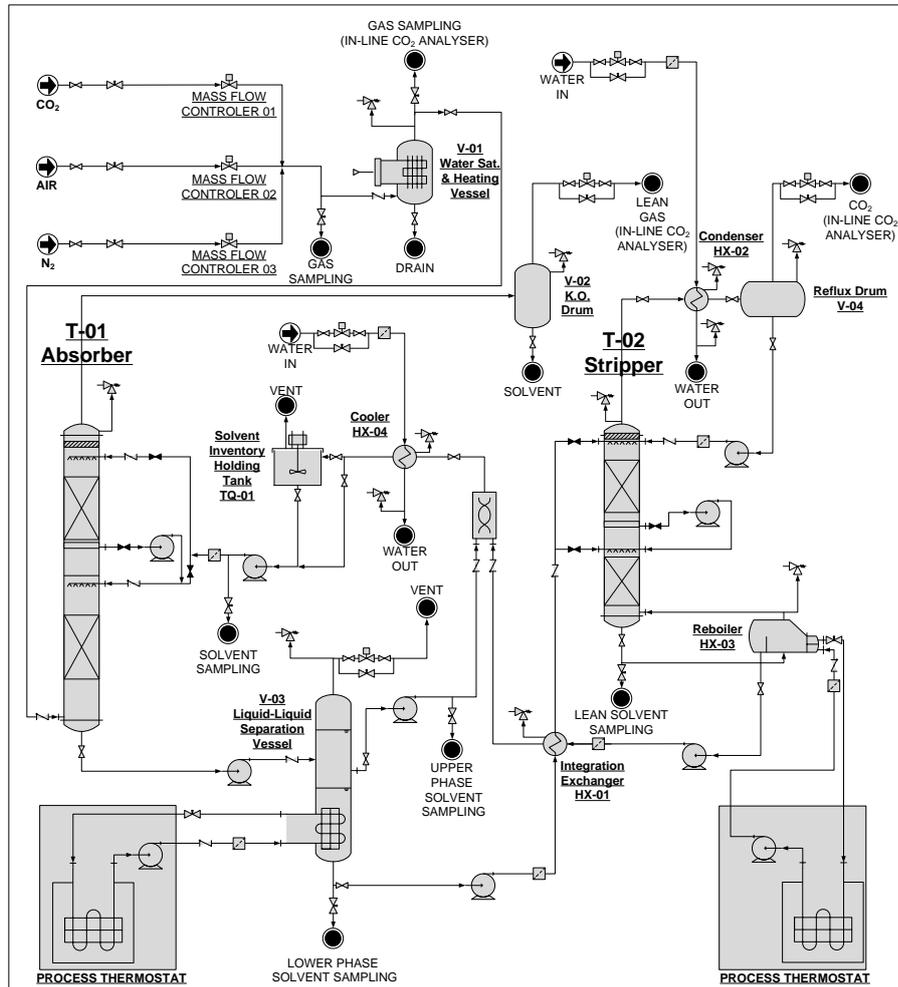
This section presents the process description and basic engineering design data related to the PCAPP, a pilot-plant intended to testing PCAS in continuous mode. The PCAPP will be located in the *Natural Gas Center of Excellence (CE-GN)*. The CE-GN has the process utilities described in Table 7.5.

**Table 7.5 CE-GN Available Utilities**

UTILITY	PROCESS PARAMETERS
Chilled Cooling-Water	5°C
Tower Cooling-Water	28°C
Compressed air	800 kPag
N <sub>2</sub> gas	400 kPag (generator)/10000 kPag (cylinder)
CO <sub>2</sub> gas	5700 kPag (cylinder)
Mixed Gas (30% CO <sub>2</sub> /70% N <sub>2</sub> )	15000 kPag (tank)
Electricity	380V triphasic 60Hz (AC) 220V monophasic 60Hz (AC)
Heating (thermal fluid)	Up to 150°C (11,2 kW)

The PCAPP is designed to capture CO<sub>2</sub> from low-pressure flue-gases with PCAS. The main difference with relation to conventional chemical absorption units is the liquid-liquid

separation vessel placed downstream the absorber bottom liquid outlet. The PCAPP process flow diagram is depicted in Figure 7.10. The P&ID of the plant is available in Fig. 7.11.



**Figure 7.10. PCAPP Flowsheet**

The plant is designed to absorb a maximum flow rate of 5kg/h of CO<sub>2</sub>. Table 7.6 shows the range of CO<sub>2</sub> concentration and partial pressure considered – the higher the CO<sub>2</sub> partial pressure the more favored is the absorption. The NGCC flue-gases are well known to present low CO<sub>2</sub> partial pressures (4 – 8 kPa) making the chemical absorption less efficient. The considered partial pressures are expected on flue-gases from NGCC power plants and some refinery units.

**Table 7.6. Design CO<sub>2</sub> partial pressures and dry flue-gas mass flows**

CO <sub>2</sub> Composition		SFG Mass Flow	SFG Inlet Pressure	CO <sub>2</sub> Partial Pressure
% mass	% mol	kg/h	kPag.	kPag
5,0	~3,5	94,7	100 – 200	3.5 - 7.0
10,0	~7,0	50,0	50 – 200	3.5 - 14
15,0	~10	33,4	50 – 200	5.0 - 20
20,0	~14	25,0	50 – 200	7.0 - 28



- **Synthetic Flue-Gas Production**

The synthetic flue-gas (SFG) is produced by mixing N<sub>2</sub>, Air and CO<sub>2</sub>. The SFG composition is manipulated using mass-flow controllers (MFCs 1 to 3, Fig 7.11). Before reaching the MFC, the pressure of each gas stream is broken by a pressure regulator. For safety reasons, the inlet gas pressures must be set to a maximum of 500 kPa. The dry SFG is sent to the heating and saturation vessel (V-01), where it could be bubbled in hot water if a more realistic flue-gas composition is desired. The SFG from V-01 proceed to the absorption column (T-01).

- **CO<sub>2</sub> Absorption**

The CO<sub>2</sub>-rich SFG (SFG<sub>rich</sub>) is fed at the bottom of the (T-01), where it passes through two sections of a 2.2m height packed beds, countercurrent to the lean solvent (SOLV<sub>lean</sub>). CO<sub>2</sub> is selectively removed from the SFG<sub>rich</sub> by chemical absorption, and the lean gas leaves the T-01 (SFG<sub>lean</sub>). SOLV<sub>lean</sub> is fed at the top of each packed bed. A knock-out drum (V-02) is placed downstream of the T-01 top gas outlet. This vessel is designed to hold eventual entrained liquid. The minimum CO<sub>2</sub> capture efficiency of the T-01 must be 90% (weight basis). The T-01 is designed to operate between 50 and 200 kPag. This parameter is controlled by a pressure control valve (PCV-01), which acts adjusting the outlet flow of SFG<sub>lean</sub>. The pump P-02 sends the CO<sub>2</sub>-rich solvent to the liquid-liquid separation vessel (V-03).

- **Liquid Phases Split**

Most of the investigated solvents present spontaneous phase-split under absorption temperature after loaded in CO<sub>2</sub>. However, some of them (TBS) need to be heated to 50°C - 80°C (depending on the PCAS) to start the liquid phase separation. Hence, the V-03 was provided with an internal heater. The heat is delivered by thermal fluid circulation guided by a process thermostat. Inside the V-03, total liquid-liquid phase split occurs, forming a CO<sub>2</sub>-rich liquid phase (SOLV<sub>richp</sub>) and a CO<sub>2</sub>-lean liquid phase (SOLV<sub>leanp</sub>). The SOLV<sub>leanp</sub> is sent back to the tank of solvent (TQ-01) by pump P-03. SOLV<sub>richp</sub> is pumped by P-04 to the top of the regeneration column (T-02). Before reaching the T-02, SOLV<sub>richp</sub> is pre-heated at the integration heat exchanger HX-01. Manipulation of V-03 pressure is possible and optionally releases CO<sub>2</sub>. In this case, V-03 would work as a batch solvent regenerator.

- **Solvent Regeneration**

Pre-heated  $SOLV_{richp}$  follows a downward path through the packed beds of T-02, in countercurrent with vapor coming from the reboiler (HX-03), when  $CO_2$  is stripped from  $SOLV_{richp}$ . The  $CO_2$ -rich hot gas ( $HOTGAS_{rich}$ ) leaves T-02 from the top and is cooled to nearly  $40^\circ C$  at the condenser (HX-02). From HX-02, the cooled stream is sent to the reflux drum (V-04) where the condensed liquid is separated from the saturated  $CO_2$ -rich gas ( $SATGAS_{rich}$ ). The  $SATGAS_{rich}$  passes through the  $CO_2$  analyzer AI-04 and leaves the system. The condensate returns to the top of T-02, through pump P-05. HX-02 uses chilled water ( $5^\circ C$ ) as a cold utility.

The stripper temperature and pressure have a remarkable effect on energy consumption and solvent flow rate. The pressure of the T-02 is controlled between 150 - 400 kPa by PCV-02. The reboiler uses thermal fluid as a heat source to promote the partial vaporization of the solvent. For MEA-based solvents, the maximum reboiler pressure is  $\cong 300$  kPa. Higher pressures imply temperatures above  $130^\circ C$  and consequent MEA degradation and heat-stable salts formation. The lean solvent ( $SOLV_{lean}$ ) is pumped from the bottom of T-02 to the reboiler by P-12. This pump is necessary because of the low height of the liquid column between the T-02 bottom outlet nozzle and the reboiler inlet nozzle. This liquid column could not be enough to surpass the reboiler internal pressure flooding the tower. The hot lean solvent ( $HOTSOLV_{lean}$ ) from the bottom of the reboiler is sent to the TQ-01 by pump P-07. Before reaching the tank,  $HOTSOLV_{lean}$  is partially cooled at the HX-01 and is mixed with the  $SOLV_{leanp}$ , forming the lean solvent stream ( $SOLV_{lean}$ ).  $SOLV_{lean}$  is cooled to  $40^\circ C$  at the HX-04 and finally returns to TQ-01. HX-04 uses chilled water as a cold utility. The regenerated solvent is recirculated to T-01 top by pump P-08.

The detailed procedure of designing the pilot plant is available in Appendix I.

### 7.3. References of Chapter 7

Aspentech. Aspen HYSYS. Disponível em: <<https://www.aspentech.com/en/products/engineering/aspens-hysys>>. Acesso em: 3 mar. 2020.

Barzagli, F., Mani, F., Peruzzini, M., 2017. Novel water-free biphasic absorbents for efficient  $CO_2$  capture. Int. J. Greenh. Gas Control. <https://doi.org/10.1016/j.ijggc.2017.03.010>

- Boot-Handford, M.E., Abanades, J.C., Anthony, E.J., Blunt, M.J., Brandani, S., Mac Dowell, N., Fernández, J.R., Ferrari, M.-C., Gross, R., Hallett, J.P., Haszeldine, R.S., Heptonstall, P., Lyngfelt, A., Makuch, Z., Mangano, E., Porter, R.T.J., Pourkashanian, M., Rochelle, G.T., Shah, N., Yao, J.G., Fennell, P.S., 2014. Carbon capture and storage update. *Energy Environ. Sci.* 7, 130–189. <https://doi.org/10.1039/C3EE42350F>
- Bottoms, R.R., 1930. Process for separating acidic gases. US17883901A.
- Budzianowski, W.M., 2016. Explorative analysis of advanced solvent processes for energy efficient carbon dioxide capture by gas–liquid absorption. *Int. J. Greenh. Gas Control* 49, 108–120. <https://doi.org/10.1016/J.IJGGC.2016.02.028>
- Coulier, Y., Lowe, A.R., Coxam, J.-Y., Ballerat-Busserolles, K., 2017. Thermodynamic Modeling and Experimental Study of CO<sub>2</sub> Dissolution in New Absorbents for Post-Combustion CO<sub>2</sub> Capture Processes. *ACS Sustain. Chem. Eng.* *acssuschemeng.7b03280*. <https://doi.org/10.1021/acssuschemeng.7b03280>
- Frailie, P.T., Madan, T., Sherman, B.J., Rochelle, G.T., 2013. Energy performance of advanced stripper configurations. *Energy Procedia* 37, 1696–1705. <https://doi.org/10.1016/j.egypro.2013.06.045>
- Global CCS Institute, 2017. The Global Status of CCS: 2017. Melbourne, Australia.
- Goto, K., Yogo, K., Higashii, T., 2013. A review of efficiency penalty in a coal-fired power plant with post-combustion CO<sub>2</sub> capture. *Appl. Energy* 111, 710–720. <https://doi.org/10.1016/J.APENERGY.2013.05.020>
- Huertas, J.I., Gomez, M.D., Giraldo, N., Garzón, J., 2015. CO<sub>2</sub> Absorbing Capacity of MEA. *J. Chem.* 2015, 1–7. <https://doi.org/10.1155/2015/965015>
- IEA, 2016. 20 Years of Carbon Capture and Storage - Accelerating Future Deployment. Paris, France.
- IPCC, 2014. Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change. Geneva, Switzerland.
- Knudsen, J.N., Jensen, J.N., Vilhelmsen, P.J., Biede, O., 2009. Experience with CO<sub>2</sub> capture from coal flue gas in pilot-scale: Testing of different amine solvents. *Energy Procedia* 1, 783–790. <https://doi.org/10.1016/j.egypro.2009.01.104>
- Koch-Glitsch. KG-TOWER® Software. Disponível em: <<https://koch-glitsch.com/kg-tower-software>>. Acesso em: 2 mar. 2020.
- Li, H., Haugen, G., Ditaranto, M., Berstad, D., Jordal, K., 2011. Impacts of exhaust gas recirculation (EGR) on the natural gas combined cycle integrated with chemical absorption CO<sub>2</sub> capture technology. *Energy Procedia* 4, 1411–1418. <https://doi.org/10.1016/j.egypro.2011.02.006>
- Liebenthal, U., Di D. Pinto, D., Monteiro, J.G.M.S., Svendsen, H.F., Kather, A., 2013. Overall process analysis and optimisation for CO<sub>2</sub> Capture from coal fired power plants

- based on phase change solvents forming two liquid phases. *Energy Procedia* 37, 1844–1854. <https://doi.org/10.1016/j.egypro.2013.06.064>
- Lu, Y., 2017. Development of a Novel Biphasic CO<sub>2</sub> Absorption Process with Multiple Stages of Liquid–Liquid Phase Separation for Post-Combustion Carbon Capture. DOE/NETL, Pittsburgh, PA.
- Micropump. Pumps - Micropump. Disponível em: <[http://www.micropump.com/product\\_list.aspx?ProductFamilyID=5](http://www.micropump.com/product_list.aspx?ProductFamilyID=5)>. Acesso em: 3 mar. 2020.
- Miyamoto, O., Maas, C., Tsujiuchi, T., Inui, M., Hirata, T., Tanaka, H., Yonekawa, T., Kamijo, T., 2017. KM CDR Process™ Project Update and the New Novel Solvent Development. *Energy Procedia* 114, 5616–5623. <https://doi.org/10.1016/J.EGYPRO.2017.03.1700>
- Park, T., Bae, J., Lee, C.J., Lee, J.M., 2016. A Sequential Method for Determining Optimal Stripper Pressure and Terminal Pressure in CO<sub>2</sub> Capture and Liquefaction Process Using MEA. *IFAC-PapersOnLine* 49, 657–662. <https://doi.org/10.1016/j.ifacol.2016.07.250>
- Pinto, D.D.D., Knuutila, H., Fytianos, G., Haugen, G., Mejdell, T., Svendsen, H.F., 2014. CO<sub>2</sub> post combustion capture with a phase change solvent. Pilot plant campaign. *Int. J. Greenh. Gas Control* 31, 153–164. <https://doi.org/10.1016/j.ijggc.2014.10.007>
- Raynal, L., Briot, P., Dreillard, M., Broutin, P., Mangiaracina, A., Drioli, B.S., Politi, M., La Marca, C., Mertens, J., Thielens, M.L., Laborie, G., Normand, L., 2014. Evaluation of the DMX process for industrial pilot demonstration - methodology and results. *Energy Procedia* 63, 6298–6309. <https://doi.org/10.1016/j.egypro.2014.11.662>
- Rezazadeh, F., Gale, W.F., Rochelle, G.T., Sachde, D., 2017. Effectiveness of absorber intercooling for CO<sub>2</sub> absorption from natural gas fired flue gases using monoethanolamine solvent. *Int. J. Greenh. Gas Control* 58, 246–255. <https://doi.org/10.1016/j.ijggc.2017.01.016>
- Rezazadeh, F., Gale, W.F., Sachde, D., Rochelle, G.T., 2014. Absorber intercooling configurations using aqueous piperazine for capture from sources with 4 to 27% CO<sub>2</sub>. *Energy Procedia* 63, 1637–1656. <https://doi.org/10.1016/j.egypro.2014.11.174>
- Rochelle, G.T., 2009. Amine Scrubbing for CO<sub>2</sub> Capture. *Science* (80-. ). 325, 1652 LP – 1654.
- Rubin, E.S., Chen, C., Rao, A.B., 2007. Cost and performance of fossil fuel power plants with CO<sub>2</sub> capture and storage. *Energy Policy* 35, 4444–4454. <https://doi.org/10.1016/j.enpol.2007.03.009>
- Sigma-Aldrich, 2018. Order Preview.
- Thong, D., Dave, N., Feron, P., Azzi, M., 2012. Process Modelling for Amine-based PostCombustion Capture Plant.
- Ulrich, J., 2010. Flue Gas Analysis in Industry: Practical Guide for Emission and Process

Measurements, 2nd ed. TESTO. <https://doi.org/0981.2773/hd/R/08.2004>

- Wang, L., An, S., Li, Q., Yu, S., Wu, S., 2017. Phase change behavior and kinetics of CO<sub>2</sub> absorption into DMBA/DEEA solution in a wetted-wall column. *Chem. Eng. J.* <https://doi.org/10.1016/j.cej.2016.12.033>
- Wang, R., Liu, S., Wang, L., Li, Q., Zhang, S., Chen, B., Jiang, L., Zhang, Y., 2019. Superior energy-saving splitter in monoethanolamine-based biphasic solvents for CO<sub>2</sub> capture from coal-fired flue gas. *Appl. Energy* 242, 302–310. <https://doi.org/10.1016/J.APENERGY.2019.03.138>
- Wang, Z., Fang, M., Pan, Y., Yan, S., Luo, Z., 2013. Amine-based absorbents selection for CO<sub>2</sub> membrane vacuum regeneration technology by combined absorption–desorption analysis. *Chem. Eng. Sci.* 93, 238–249. <https://doi.org/10.1016/J.CES.2013.01.057>
- Zhang, J., Qiao, Y., Wang, W., Misch, R., Hussain, K., Agar, D.W., 2013. Development of an energy-efficient CO<sub>2</sub> capture process using thermomorphic biphasic solvents. *Energy Procedia* 37, 1254–1261. <https://doi.org/10.1016/j.egypro.2013.05.224>
- Zhang, S., Shen, Y., Shao, P., Chen, J., Wang, L., 2018. Kinetics, Thermodynamics, and Mechanism of a Novel Biphasic Solvent for CO<sub>2</sub> Capture from Flue Gas. *Environ. Sci. Technol.* 52, 3660–3668. <https://doi.org/10.1021/acs.est.7b05936>
- Zhang, W., Jin, X., Tu, W., Ma, Q., Mao, M., Cui, C., 2017a. A Novel CO<sub>2</sub> Phase Change Absorbent: MEA/1-propanol/H<sub>2</sub>O. *Energy & Fuels* [acs.energyfuels.7b00090](https://doi.org/10.1021/acs.energyfuels.7b00090). <https://doi.org/10.1021/acs.energyfuels.7b00090>
- Zhang, W., Jin, X., Tu, W., Ma, Q., Mao, M., Cui, C., 2017b. Development of MEA-based CO<sub>2</sub> phase change absorbent. *Appl. Energy* 195, 316–323. <https://doi.org/10.1016/j.apenergy.2017.03.050>
- Zhang, Y., Rochelle, G.T., 2014. Absorber Performance with High CO<sub>2</sub>. *Energy Procedia* 63, 1329–1338. <https://doi.org/10.1016/j.egypro.2014.11.142>
- Zhou, X., Liu, F., Lv, B., Zhou, Z., Jing, G., 2017. Evaluation of the novel biphasic solvents for CO<sub>2</sub> capture: Performance and mechanism. *Int. J. Greenh. Gas Control.* <https://doi.org/10.1016/j.ijggc.2017.03.013>



## **8. CONCLUSIONS AND SUGGESTIONS**

The causal nexus of economic growth, energy consumption and ecological footprint is reviewed in this thesis. A challenging scenario of increasing energy demand constrained by global warming is presented. The persistence and high share of fossil sources in the global TPES for the next decades are also evidenced. Aiming to mitigate some environmental impacts of carbon-based power, this thesis identifies technological gaps within the fossil energy value chain. Four technologies, related to three research lines (R1, R2 and R3), are proposed and evaluated through technical, economic and/or environmental assessments.

In general, the technologies and alternative process designs developed in this thesis proved to be beneficial in at least one of the three considered dimensions (technical, economic or environmental). The technologies have the potential to be applied in full-scale and achieve commercial maturity. However, to reach this stage more R&D effort to scale-up and troubleshooting technical issues is necessary. The utilization of the proposed technologies would raise the competitiveness and support decision-making on some chains of the fossil energy production process. It makes clear that energy efficiency is a key point in future developments. However, the economic impacts of new technologies are the main barrier to its development, especially in developing and poor countries. Meet the triple bottom line of sustainability on the fossil-energy sector is a challenging task. The introduction of a carbon tax would be the way towards a sustainable future. The taxation would favor environmental-friendly technologies, like the ones proposed in this thesis.

More specific findings, contributions and answers to the questions that motivated the development of the present thesis are addressed below, separated by research line.

### **8.1. R1 - Offshore Processing of CO<sub>2</sub>-Rich Natural Gas**

#### **R1.1. Deep Seawater Intake for Primary Cooling in Tropical Offshore Processing of Natural Gas with High Carbon Dioxide Content: Energy, Emissions and Economic Assessments**

The use of DSW intake at 900 m of depth instead of conventional seawater intake as the primary cooling utility of a FPSO is investigated. The resulting variations of gas processing layout, electricity generation capacity, energy usage efficiency and CO<sub>2</sub> emissions were assessed.

The overall energy usage efficiency increase with DSW intake is in the range of 2.7% to 5.0%, depending on the gas processing flow. The highest value corresponds to full capacity. The CO<sub>2</sub> emissions had decreased in the same proportion, thanks to the reduced fuel gas consumption for power generation. The use of DSW promotes modest cost and weight savings when the entire FPSO is considered, but the results are more expressive considering only the gas processing plant and power generation units. Furthermore, DSW intake also leads to other indirect advantages like the elimination of the refrigeration cycle for HCDPA and 6% reduction of water content in the gas feed to dehydration TSA units for WDPA. The last could result in further cost and energy savings, not assessed in this investigation. The study reached its main goals, but some simplification was considered to limit the scope of the investigation. It was necessary because of the complexity and size of the FPSO flowsheet and the massive calculation workforce demanded to perform the necessary assessments. Some limitations of the cost estimation software, like equipment pressure and size, could lead to underestimated cost and weight.

As a recommendation for future work, a lifecycle cost evaluation of the DSW intake pipelines must be performed to determine the ultimate feasibility of the investigated technology. Additionally, the feasibility of DSW intake should be investigated within a continuous or mixed-integer non-linear optimization framework so that certain features that were assumed pre-defined and constant in this study could vary to seek optimum performance. For instance, the present analysis showed that certain FPSO units become problematic if cooled with cold CW at 7°C. Therefore, a possible optimization formulation would consider two independent CW circuits, working at two temperature ranges – 35°C-55°C and 7°C-27°C – whose service heat loads, allocation points, exchanger/pump sizing and circulation flow rates are continuous or mixed-integer decision variables to be sought. In this case, it is conceivable to let the outlet temperature of DSW in the plate exchanger free, up to the regulated outlet limit of 40°C. The DSW flow rate and the cost of intake piping, insulations and pumps would be minimized. Relaxing the exiting temperature of DSW seems a reasonable point to be questioned by optimizations as it has some thermodynamic support in the context of exergy analysis. The flow of wasted exergy could be reduced by returning a lower flow rate of hotter DSW to the sea at the expense of using larger plate heat exchangers in the process.

**R1.2.** Exergy, Energy and Emissions Analysis of Compressors Schemes in Offshore Rigs: CO<sub>2</sub>-Rich Natural Gas Processing

An alternative process layout applied to the offshore processing of CO<sub>2</sub>-Rich NG is developed and evaluated. MPSC (multiple paralleled smaller compressors) are proposed instead of conventional SSLC (single-shaft larger compressors with anti-surge recycles). SSLC-Case and MPSC-Case were compared in terms of exergy efficiency, FCI, footprint, CO<sub>2</sub> and energy intensities. This integrated holistic approach is unusual in conventional studies. Simulations revealed that oversized compressors with anti-surge recycle lead to almost constant power consumption along the field lifespan, even with a falling gas flow being processed. Consequently, fuel-gas and CO<sub>2</sub> intensities increase as gas-load decreases. It is shown that the efficiency of compressors is kept higher using VSD and smaller paralleled compressors. Moreover, eliminating anti-surge recycles the FPSO power demand becomes proportional to gas-load, reducing fuel-gas and CO<sub>2</sub> intensities.

Regarding the exergy analysis, two Reference Environmental Reservoirs are considered, RER-1 and RER-2. RER-1 inflates exergy flows with the high chemical exergy of hydrocarbons, producing too high exergy efficiencies for physical operations, e.g. compressors, exchangers and separators. Therefore, the exergy efficiencies for SSLC-Case and MPSC-Case are always ~99%, no matter the gas-load. RER-1 is considered useful for chemically reactive operations with high spontaneity, such as gas turbines and combustors in general. For such operations, RER-1 produces reliable exergy efficiencies. RER-2 deflates exergy flows by excluding the high chemical exergy of hydrocarbons, making the exergy assessments of physical operations and the overall gas plant meaningful. Exergy analyses corroborate the simulation achievements, unveiling that MPSC-Case entails a much lower FPSO exergy destruction rate. For FPSO gas-load ranging from 25% to 100%, the RER-2 exergy efficiency of SSLC-Case lies between 49% and 83%, whereas the MPSC-Case is always from 81% to 88%.

Besides being more exergy and energy-efficient, MPSC-Case leads to 3% of *FCI* savings, despite increasing only 4% the overall equipment weight. The lower power demand of MPSC-Case allowed removal of one GT in the FPSO, compensating the *FCI* and footprint increases of compressors and exchangers. In a carbon taxation scenario, MPSC-Case would be even more profitable.

Aiming to prove the technical feasibility potential of innovation of using MPSC, future work would address dynamic simulation of the compression systems. It would validate the proposed process design under several operation modes. The developed methodology of

exergy analysis would also be applied to further evaluation of DSW intake. Additionally, the DSW intake and MPSC designs would be mixed and evaluated simultaneously.

## **8.2. R2 - Desulfurization Residues from Coal-Fired Power Plants**

The environmental burden of SD-FGDR landfills is introduced. A treatment of SD-FGDR through dry-oxidation is proposed to make this residue useful. In Chapter 5 the impact of the treatment in the LCOE of a real PCC power plant facing decision-making process on SD-FGDR destination is assessed. A LCOE of 94.97 \$/MWh is calculated for a conventional PCC, in agreement with reported values of similar power plants. The energy demanded by the novel SD-FGDR treatment unit is negligible due to the exothermal nature of the  $\text{CaSO}_3$  oxidation reaction and the pre-heater of inlet air and solids. The air compressor power is only 0.04% of the net power plant output (340 MW). The LCOE is impacted by the investment, operation and maintenance costs of the SD-FGDR treatment unit and the residue revenue price. Disregarding the commercialization of residue the LCOE increases only 0.02%. If the residue is sold as a raw material for cement kiln, the LCOE decreases by ~3%, to 92.14 \$/MWh. Therefore, the SD-FGDR revenue has a small beneficial impact on the LCOE, besides solving the landfill environmental issues and maintenance cost. In conclusion, this study demonstrates that it is technically and economically feasible to implement the dry-oxidation SD-FGDR treatment on the investigated PCC power plants and others with similar design.

In Chapter 6, an assessment of the environmental impacts of three different SD-FGDR management scenarios is performed for the same PCC power plant considered in Chapter 5. Heat and mass balances were calculated for each alternative. It is shown that the Base Case has a higher rate of potential environmental impacts due to the massive production of SD-FGDR. The Base Case is compared to the PCC operating with the treatment of the residue (CASE I) or the bypass of the SDA system (CASE II). CASE I and CASE II enable selling the SD-FGDR mixed with coal ashes as class C fly-ash. The SDA bypassing lead to higher  $\text{SO}_2$  emissions and an increase of 1245% in the photochemical oxidation and acidification potentials. It is concluded that using de SD-FGDR treatment is a more environmental-friendly alternative.

Considering only the economic point of view, bypass the FGD is better, but this study proves that the environmental impacts related to SO<sub>2</sub> emissions increase dramatically and could be prohibitive in countries where environmental legislation is restrictive, like in Western Europe and the USA. CASE I is more sustainable because it solves SO<sub>2</sub> emissions while reducing other environmental impact categories. In the long term, depending on the ash and cement market, CASE I could become profitable, because of the revenue of fly-ash.

Future work must include updated data from pilot-plant runs and improvements on the ash composition analytical methodology. These data would enable a more accurate streams inventory. This inventory could serve as input for extended LCA. The use of low-grade heat coming from the power plant could favor the economic and environmental performance of a full-scale SD-FGDR treatment system. This effect must be investigated, and results included in future studies.

### **8.3. R3 - CO<sub>2</sub> Capture from Flue-Gases by Phase-Changing Absorption Solvents**

The state-of-the-art in PCAS is investigated. Solvents were selected and preliminary tests performed to confirm phase-changing behavior, CO<sub>2</sub> loading and other relevant process parameters. One solvent was selected for further evaluation in a batch screening plant, named PCASP. The plant is designed to perform absorption and desorption tests using a synthetic exhaust gas. The test confirmed that the PCAS based on MEA and 1-propanol has the potential to reduce the energy penalty of CO<sub>2</sub> carbon capture by chemical absorption using conventional and affordable chemical components.

The next step is testing the best PCAS under different compositions and absorption conditions, simulating a more realistic flue-gas (higher temperature, pressure, presence of contaminants and lower CO<sub>2</sub> partial pressure). Desorption experiments also must be executed to confirm the cyclic capacity and support comparison with results reported by other authors.

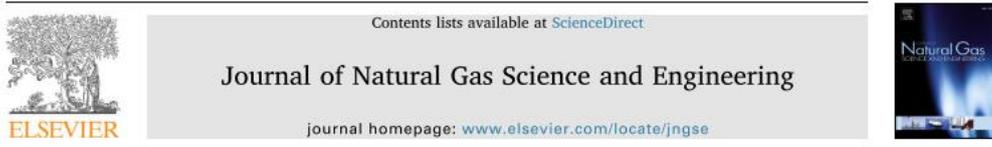
A pilot plant is designed to operate in continuous mode. It will enable submit solvents to long run experiments. The pilot experiments will support a more precise estimation of the potential energy penalty, solvent loss and degradation rate, and other useful process parameters. Pilot data is necessary to validate process simulations and to scale-up the PCAS technology.



## APPENDIX A. PUBLISHED PAPER OF CHAPTER 3 - JNGSE

CRUZ, M. DE A.; ARAÚJO, O. DE Q. F.; DE MEDEIROS, J. L. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. **Journal of Natural Gas Science and Engineering**, v. 56, n. June, p. 193–211, 2018.

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### Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments



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#### ABSTRACT

In deepwaters offshore oil-gas rigs, centrifugal compressor trains are major power consumers, requiring intercoolers conventionally designed assuming surface seawater for primary cooling, limiting compressor inlet gas temperatures to 40 °C at tropical sites. On the other hand, at tropical deepwaters the available deep seawater at 4 °C can be exploited to reduce compression power – nearly proportional to inlet gas absolute temperature – entailing energy, economic and environmental benefits. This work considers a new primary cooling for deepwaters offshore platforms based on deep seawater (DSW) intake at 4 °C from depths around 900 m, reducing the outlet temperature of intercoolers to 12 °C. DSW intake alternative is assessed in terms of power consumption, CO<sub>2</sub> emissions and economy employing detailed equipment sizing and cost estimation. Depending on gas flow rate, it is shown that DSW intake lowers compressors power up to 9.2%, besides several indirect benefits: elimination of one CO<sub>2</sub> compressor; 30% less heat transfer areas; 4.5% less fuel gas consumption; 4% less gas turbines power; 9.5% (15 MMUS\$) less investment; 14.4% (226 t) less topside weight, while making refrigeration unnecessary for dew point adjustment. DSW intake also entails 5% more efficient energy usage and 9327 tCO<sub>2</sub>/y less emissions, boosting economic performance under carbon taxation.

#### 1. Introduction

Since 2000, offshore fields respond for ≈30% of the world production of oil and natural gas (Rui et al., 2017). Due to still modest competitiveness of renewable energy sources, fossil sources will continue to play significant role in global energy matrix in the short to mid-term, especially natural gas (NG). However, oil price decline and new climate change mitigation policies – e.g. carbon taxation – have been a challenge to this industry. CO<sub>2</sub> emission taxation is already a reality in many countries, with Sweden imposing the highest tax of 140 US\$/t (IEA, 2016). According to the International Association of Oil and Gas Producers (IOGP, 2016) main oil and gas companies emitted 280 Mt CO<sub>2e</sub> of greenhouse gases (GHG) in 2015, of which 68% is related to fuel combustion for in-place energy production. Thus, energy usage efficiency and CO<sub>2</sub> emissions are becoming not solely an environmental, but also, an economic issue for oil and gas producers. As energy usage efficiency and CO<sub>2</sub> emissions are inversely interrelated concepts, using processing strategies with higher energy usage efficiency implies lowering CO<sub>2</sub> emissions, alleviating the environmental burden of oil and gas industries. In other words, there is no option to the carbon fossil

industry but questing for better energy usage efficiency.

##### 1.1. Offshore oil and gas processing: improving efficiency of energy usage

New developments on offshore oil and gas primary processing represent opportunities to, cumulatively, improve efficiency of energy usage, reduce GHG emissions, lower topsides footprint and, consequently, reduce costs. All these effects contribute to increase the economic and environmental feasibility of offshore oil and gas production. The literature presents several recent works on energy assessment and optimization of primary processing on offshore oil and gas rigs, comprising measures to improve energy usage efficiency (Nguyen et al., 2016a), better power generation schemes with organic cycles (Pierobon et al., 2013), air-bottoming cycles (Pierobon and Haglind, 2014), steam-bottoming cycles (Nguyen et al., 2014a), heat-exchanger network optimization for minimum energy consumption (Pierobon et al., 2013; Pierobon and Haglind, 2014; Nguyen et al., 2014a) and offshore power production via combined cycles (Rivera-Alvarez et al., 2015). To improve energy usage efficiency of the process as a whole, including carbon capture units such as post-combustion amine plants, Nguyen

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## APPENDIX B. PAPER SUBMITTED OF CHAPTER 4 – JNGSE

*Title: Exergy, Energy and Emissions Analysis of Compressors Schemes in Offshore Rigs: CO<sub>2</sub>-Rich Natural Gas Processing*

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Keywords: Offshore gas processing; CO <sub>2</sub> -rich natural gas; Centrifugal compressors; Anti-surge recycle; Exergy analysis; CO <sub>2</sub> emissions
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Abstract: Deepwater oil and associated gas productions resort to floating rigs operating at continuously decreasing gas-loads during the last three quarters of the field campaign. As centrifugal compressors are sized at maximum loads, anti-surge recycles are used making operation inefficient in terms of power consumption and emissions per oil barrel produced. Smaller paralleled compressors and variable-speed drivers are investigated at peak and partial gas-loads and compared to traditional anti-surge recycle designs in terms of exergy efficiency, investment, footprint and emissions. Oversized compressors with anti-surge recycles result in almost constant power consumption along process lifespan, regardless the gas-load, increasing fuel and CO <sub>2</sub> intensities as gas-load decreases and attaining exergy efficiencies of 49% and 83% at 25% and 100% gas-loads, respectively. On the other hand, with variable-speed drivers and smaller paralleled compressors, power consumption becomes proportional to gas-load with exergy efficiencies always between 80% and 88%, and attaining 11% and 39% less power consumptions at 100% and 25% gas-loads. Moreover, CO <sub>2</sub> intensity and investment are, respectively 34% and 3% less than in traditional layouts with oversized compressors. These savings resulted from eliminating a gas turbine thanks to lower power demand when no anti-surge recycles are used.
Research Data Related to this Submission ----- There are no linked research data sets for this submission. The following reason is given: Data will be made available on request

## APPENDIX C. PUBLISHED PAPER OF CHAPTER 5 – JCLEPRO

CRUZ, M. DE A. et al. Impact of solid waste treatment from spray dryer absorber on the levelized cost of energy of a coal-fired power plant. *Journal of Cleaner Production*, v. 164, 2017a.

Journal of Cleaner Production 164 (2017) 1623–1634



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Journal of Cleaner Production

journal homepage: [www.elsevier.com/locate/jclepro](http://www.elsevier.com/locate/jclepro)



### Impact of solid waste treatment from spray dryer absorber on the levelized cost of energy of a coal-fired power plant



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#### ABSTRACT

Coal-fired power plants with semi-dry flue gas desulfurization (semi-dry FGD) system produce daily tones of ashes contaminated with calcium sulphite. To turn this solid waste useful (e.g. to the cement industry) and avoid landfill disposal, the present study suggests a semi-dry FGD solid waste treatment unit, that promotes the dry oxidation of calcium sulfite to calcium sulfate. Sizing of main equipment using pilot-plant and patents data allows economic evaluation of capital expenditure, operational and maintenance costs, and sale of the treated residue, which permits estimation of levelized cost of energy to assess the impact of the technology on the electricity price of a power plant using the proposed solid waste treatment unit. As base case, a Brazilian coal-fired power plant facing decision making process on semi-dry FGD waste destination is selected. Results demonstrate that the semi-dry FGD, without the solid treatment unit, has total levelized cost of energy increased in 0.56% (from 94.44 to 94.97 \$/MWh) resulting from solids waste disposal. If the treated semi-dry FGD waste was transferred (at zero revenue) as additive to a cement industry, the levelized cost of energy of the power plant would remain approximately unchanged. This is because the increase of 0.51\$/MWh resulting from the investment and operation and maintenance cost of the treatment unit is compensated by the decrease of 0.53\$/MWh, in virtue of the avoided waste disposal costs. However, if the commercialization as raw material of the treated semi-dry FGD waste is considered, a reduction of 2.83 \$/MWh (~3%) on the levelized cost of energy (to 92.14 \$/MWh) would occur. In both cases, the proposed treatment unit shows small impact on the total power plant levelized cost of energy, besides solving the solid management problems of landfill saturation, land use and costs related to landfill maintenance. Thus, it is adequate to implement the semi-dry FGD waste treatment unit on the power plant in question. The conclusion can be extended to plants with similar design and economic parameters.

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#### 1. Introduction

The Brazilian electricity matrix is dominated by hydropower generation. However, because of the recent water scarcity crisis in Brazil, hydroelectricity has been supplemented with electrical power generated by thermal power plants, resulting in 18% increase

during 2013–2014 and presently represents 28.2% of the total Brazilian electricity sources. In this same period, electricity produced by coal power plants has increased 24.2%, with mineral coal representing 9.6% of the thermopower sources in Brazil (EPE, 2015a). The Brazilian energy demand will increase in an average of 3.6% yearly until 2019, thus it is expected that the use of coal-fired power plants will continue to increase in short to medium term (EPE, 2015b). Furthermore, an average power plant technical lifetime of about 40 years for coal compared to 34 years for gas and 34 years for oil-fired power plants is estimated (Farfen and Breyer, 2017), indicating that the next decade will sustain supply of fossil energy accompanied by growing environmental legislation and

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## APPENDIX D – PUBLISHED PAPER OF CHAPTER 6 - JSDEWES

CRUZ, M. D. A. et al. Environmental Performance of a Solid Waste Monetization Process Applied to a Coal-Fired Power Plant with Semi-Dry Flue Gas Desulfurization. **Journal of Sustainable Development of Energy, Water and Environment Systems**, v. 7, n. 3, p. 506–520, 2018.



*Journal of Sustainable Development of Energy, Water  
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<http://www.sdewes.org/jsdewes>

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### **Environmental Performance of a Solid Waste Monetization Process Applied to a Coal-Fired Power Plant with Semi-Dry Flue Gas Desulfurization**

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#### **ABSTRACT**

Mixing of semi-dry flue gas desulfurization solids and fly-ash from coal-fired power plants results in a solid waste contaminated by calcium sulfite. Therefore, it becomes useless for industry and is often landfilled. To support decision-making on process configurations to monetize this solid residue a gate-to-gate life cycle assessment was performed, considering three scenarios: BASE case – standard 360 MW power plant, CASE I – base plant adopting dry thermal oxidation treatment of spray dryer solids, CASE II – bypass of desulfurization system. Cases I and II allow commercialization of the solid residue as class C fly-ash. Evaluated alternatives were compared based on quantitative potential environmental impacts, using United States Environmental Protection Agency waste reduction algorithm. Based on the results, the BASE case was more aggressive to the environment, due to solid waste production. CASE II increased photochemical oxidation and acidification potentials. CASE I was the more environmentally friendly but demands additional capital and operational expenditure.

#### **KEYWORDS**

*Calcium sulfite dry oxidation, Coal fired power plant, Life cycle assessment, Semi-dry flue gas desulfurization, Solid waste treatment, Spray dryer absorbers.*

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## APPENDIX E – CONFERENCE PAPER – SDEWES 2019

Cruz, M. de A. et al. (2019) **SDEWES2019.0276 Chemical Absorption of CO<sub>2</sub> from Flue Gases: Experiments with Phase Changing Solvents in a Bench Scale Plant**, 14th Conference on Sustainable Development of Energy, Water and Environment Systems. Edited by A. Mudrovčić and M. Ban. Zagreb: Faculty of Mechanical Engineering and Naval Architecture. Available at: <https://www.dubrovnik2019.sdwes.org/>.

Conference on Sustainable Development of Energy, Water and Environment Systems, Dubrovnik, 1.-6.10.2019

### **SDEWES2019.0276**

#### **Post-Combustion Carbon Capture: Investigation of Chemical Absorption with Phase-Changing Solvents in a Bench-Scale Plant Operating in Batch Mode**

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#### **Abstract**

The energy penalty of solvent regeneration is a barrier for the deployment of chemical absorption post-combustion carbon capture. Phase-changing absorption solvents have been proposed to overcome this issue. CO<sub>2</sub> absorption triggers phase separation with only the CO<sub>2</sub>-rich phase requiring regeneration, potentially reducing energy demand. A systematic review supports the choice of a set of solvents for experimental investigation, based on economic, process and energy-related criteria. Until now the selected absorbents were only investigated in laboratory scale. They need to move from lab to industrial scale to contribute to the global warming mitigation. Three selected biphasic solvents are selected and tested, to confirm the results reported by its developers. Solvent A, based on monoethanolamine/1-propanol, was considered the more suitable one. This blend presented 26% of reduction on the lower (CO<sub>2</sub>-rich) liquid phase, compared to the initial volume of fresh solvent. monoethanolamine is the more traditional chemical absorbent for CO<sub>2</sub> capture applications and both components can be considered standard chemicals (low-cost), which is a remarkable advantage. The viscosity of the CO<sub>2</sub>-rich phase of solvent A is 10 mPa.s at 25°C, which is considered low compared to other candidates. Solvent B, based on diethylene-triamine and N,N,N',N',N''-pentamethyldiethylenetriamine, was disregarded for further evaluations. It presented only 10% of volume reduction on CO<sub>2</sub> rich phase and prohibitively high viscosity (360 mPa.s at 40°C). Furthermore, chemical components of this solvent are considered specialty (high cost). Solvent C, based on N-methylcyclohexylamine and N,N-dimethylcyclohexylamine, was also disregarded for further tests. Although the costs of its chemical components are in the same range of monoethanolamine and the CO<sub>2</sub>-rich liquid phase presents an acceptable viscosity (58 mPa.s at 25°C), this blend had an issue. It presented solids precipitation on the CO<sub>2</sub> rich phase, what is a potential source of operational problems on industrial application. Solvent A was the only one tested on a bench scale screening plant, designed for absorption and desorption of chemical absorption solvents. This blend presented a CO<sub>2</sub> loading of 2.8 mol/kg on the lower phase, 76% higher than MEA 30%. It is an opportunity to reduce the energy penalty of carbon capture of CO<sub>2</sub> by chemical absorption. The solvent A was selected to further evaluation on a continuous mode pilot-plant (under construction). Based on thermodynamic process simulation, the use of the solvent A in replacement of MEA 30% could result in a decrease of 7.9% on the regeneration energy penalty.

## APPENDIX F – PUBLISHED PAPER - ECM

DE CASTRO, R. DE P. V. et al. Fluidized bed treatment of residues of semi-dry flue gas desulfurization units of coal-fired power plants for conversion of sulfites to sulfates. **Energy Conversion and Management**, v. 143, 2017.

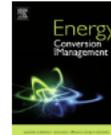
Energy Conversion and Management 143 (2017) 173–187



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### Fluidized bed treatment of residues of semi-dry flue gas desulfurization units of coal-fired power plants for conversion of sulfites to sulfates



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#### ABSTRACT

Coal-fired power plants with semi-dry flue gas desulfurization produce several metric tons per day of residues with 10–18%w/w of hemi-hydrated calcium sulfite. The rest of the 90–80%w/w of such residues contains silica, aluminium-silicates and calcium/magnesium carbonates, sulfates and hydroxides. This material could be added to cement, but sulfites degrade the cement quality and lead to costs of landfill disposal. To test upgrading desulfurization residues to turn it into an acceptable cement feedstock, a pilot plant was built to oxidize residues with hot air converting sulfites to sulfates. This pilot comprehends a fluidized bed reactor, an air heater, a cyclone and a heat recovery exchanger. Its operation showed that residues react favorably under fluidization. The effect of independent variables, residence time and temperature, was investigated and sulfite conversions up to 89.4% were observed. With statistical treatment of pilot experimental data, response surface of conversion of sulfites to sulfates was developed, allowing to estimate the effect of independent coordinates on conversion and the optimal oxidation conditions. Experimental data and model predictions showed agreement leading to low estimated variance.

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#### 1. Introduction

Coal-fired power plants play a vital role in electricity generation worldwide. According to World Coal Association, coal-fired power plants currently are responsible by 41% of global electricity production and the global coal demand is projected to increase 15% until 2040, mainly due to the growing energy demand of Asian economies such as China and India [1]. New plant efficiencies, tight emissions controls and cleaner technologies are expected to comply with this increasing demand of coal energy via-à-vis the increasing climate concerns.

The primary flue gas (FG) of a typical 360 MW coal-fired power plant comprehends impressive flow rates of problematic gases, e.g.  $\approx 452$  t/h of carbon dioxide ( $\text{CO}_2$ ) and  $\approx 4.2$  t/h of sulfur dioxide ( $\text{SO}_2$ ). In order to abate the atmospheric emissions of  $\text{SO}_2$  the flue gas (FG) is treated in the operation known as Flue Gas Desulphurization (FGD), which is located just prior the liberation of FG into the atmosphere.

##### 1.1. FGD systems of coal-fired power plants

There are a few FGD technologies for coal-fired power plants. An important aspect in connection with FGD systems in such plants is the presence of very fine fly-ash, which can be deposited with the solid FGD residues. FGD systems of coal-fired plants are generally classified as Dry or Semi-Dry FGD and Wet FGD.

In Wet FGD the FG at the dew point is contacted in large vessels with a fine-grain aqueous slurry of grinded limestone containing calcium carbonate ( $\text{CaCO}_3$ ) and magnesium carbonate ( $\text{MgCO}_3$ ) with air injected in the bottom. In the Wet FGD  $\text{SO}_2$  is the only component of FG which reacts with the slurry of limestone via Eqs. (1a)–(1d) producing basically a slurry of hydrated calcium sulfate or gypsum ( $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ ), magnesium sulfate ( $\text{MgSO}_4 \cdot 6\text{H}_2\text{O}$ ), calcium sulfite ( $\text{CaSO}_3 \cdot 0.5\text{H}_2\text{O}$ ), magnesium sulfite ( $\text{MgSO}_3 \cdot 6\text{H}_2\text{O}$ ) and  $\text{CO}_2$  gas. As the weight ratio sulfates:sulfites is  $\approx 5$  (w/w), an important economic factor of Wet FGD is the exportation of gypsum as a masonry feedstock. For the 360 MW coal power plant, the flow rate of 4.2 t/h of  $\text{SO}_2$  in the FG will entail a production of  $\approx 9$  t/h of gypsum + sulfites or  $\approx 28$  t/h of solids (gypsum + sulfites + fly-ashes + limestone) with a consumption of  $\approx 7$  t/h of limestone and  $\approx 39$  t/h of water.

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0196-8904/© 2017 Elsevier Ltd. All rights reserved.

## APPENDIX G – PUBLISHED PAPER – Petróleo & Gás

CRUZ, M. DE A. et al. Estudo de mercados de GNL, GLP, propano e butano liquefeitos, visando aproveitamento comercial do gás natural do pré-sal. *Revista Petróleo & Gás*, v. 369, p. 16–22, 2017b.

Artigo Técnico

# Estudo dos mercados de GNL, GLP, propano e butano liquefeitos, visando aproveitamento comercial do gás natural do Pré-sal

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### Resumo

Investimentos de grande soma têm sido feitos no polo Pré-sal da Bacia de Santos, visando o desenvolvimento dos seus campos e aumento da produção de óleo. Com isso, há expectativa de que a produção de gás natural (GN) dobre até 2030. Esse GN pode ser reinjetado, com o objetivo de aumentar a recuperação de óleo dos campos, ou escoado para terra, por gasodutos. Essa última alternativa depende da expansão da malha de gasodutos, ou seja, mais investimento, mas abriria oportunidades para redução do déficit nacional do mercado de GNL e exploração dos mercados mundiais de  $C_3$ ,  $C_4$  e GLP, contribuindo para o desenvolvimento da economia brasileira. O principal objetivo do estudo é fornecer subsídios para definir-se a melhor estratégia de utilização do GN produzido no Pré-sal. São estudadas as cadeias produtivas desses produtos e apresentadas unidades de processamento e seus custos típicos. A seguir é apresentado o mercado nacional e internacional de GNL, GLP,  $C_3$  e  $C_4$ . O comércio exterior brasileiro foi estudado através das ferramentas Radar Comercial e Alice Web. Evidenciaram-se a complexidade e o dinamismo do mercado de GNL,  $C_3$ ,  $C_4$  e GLP. A análise dos dados de comércio exterior indica que o mercado de GNL é muito maior (em volume e valores, a nível nacional e internacional) que o de GLP,  $C_3$  e  $C_4$  e que os maiores consumidores desses produtos pertencem à região Ásia-Pacífico, com destaque para o Japão, Coreia e China. O Radar Comercial indica que a prioridade deve ser dada a projetos de  $C_3$  e  $C_4$ . Projetos de GLP não seriam tão oportunos para o Brasil. Com isso, investir em UPLs e UFLs pode ser mais interessante do que em UPGNs. Além disso, haveria maior disponibilização de gás natural no Brasil, reduzindo o déficit da balança comercial de GNL, hoje importado para complementar o suprimento de gás natural ao mercado nacional.

### 1. Introdução

A partir do gás natural (GN) bruto são obtidos diversos produtos, tais como: o Gás Natural Especificado, Gás Natural liquefeito (GNL), Gás Liquefeito de Petróleo (GLP), Propano Liquefeito ( $C_3$ ), Butano Liquefeito ( $C_4$ ) e Gasolina Natural ( $C_4$ ). Como será mostrado, o Brasil sempre foi importador destes produtos.

A figura 1 mostra que a oferta doméstica de GN, a partir dos campos da Petrobras e de seus parceiros, deve mais que dobrar até 2030, refletindo os investimentos que têm sido feitos para aumentar a produção e o escoamento do GN produzido no polo Pré-sal. Tal previsão assume a premissa de escoamento do GN, dos campos *offshore* para terra. Inicialmente são previstas três rotas, com capacidade total de 40,8 MM Sm<sup>3</sup>/d, como ilustrado pela figura 2. Porém, há campos do Pré-sal com previsão de gerar até 100 MM Sm<sup>3</sup>/d de GN no pico da produção, o que demonstra a necessidade de mais gasodutos de exportação.



Figura 1. Oferta de GN da Petrobras 2013-2030.

Fonte: Petrobras

## **APPENDIX H – SUPPLEMENTARY MATERIALS**

### **SUPPLEMENTARY MATERIALS A**

CRUZ, M. DE A.; ARAÚJO, O. DE Q. F.; DE MEDEIROS, J. L. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. **Journal of Natural Gas Science and Engineering**, v. 56, n. June, p. 193–211, 2018.

**Supplement A1. Simulation Methodology**

**Supplement A2. Equipment Sizing**

**Supplement A3. Cost and Weight Estimation**

**Supplement A4. Simulation, Sizing and CAPEX Results**

**References of Supplementary Materials A**

## Supplement A1 – Simulation Methodology

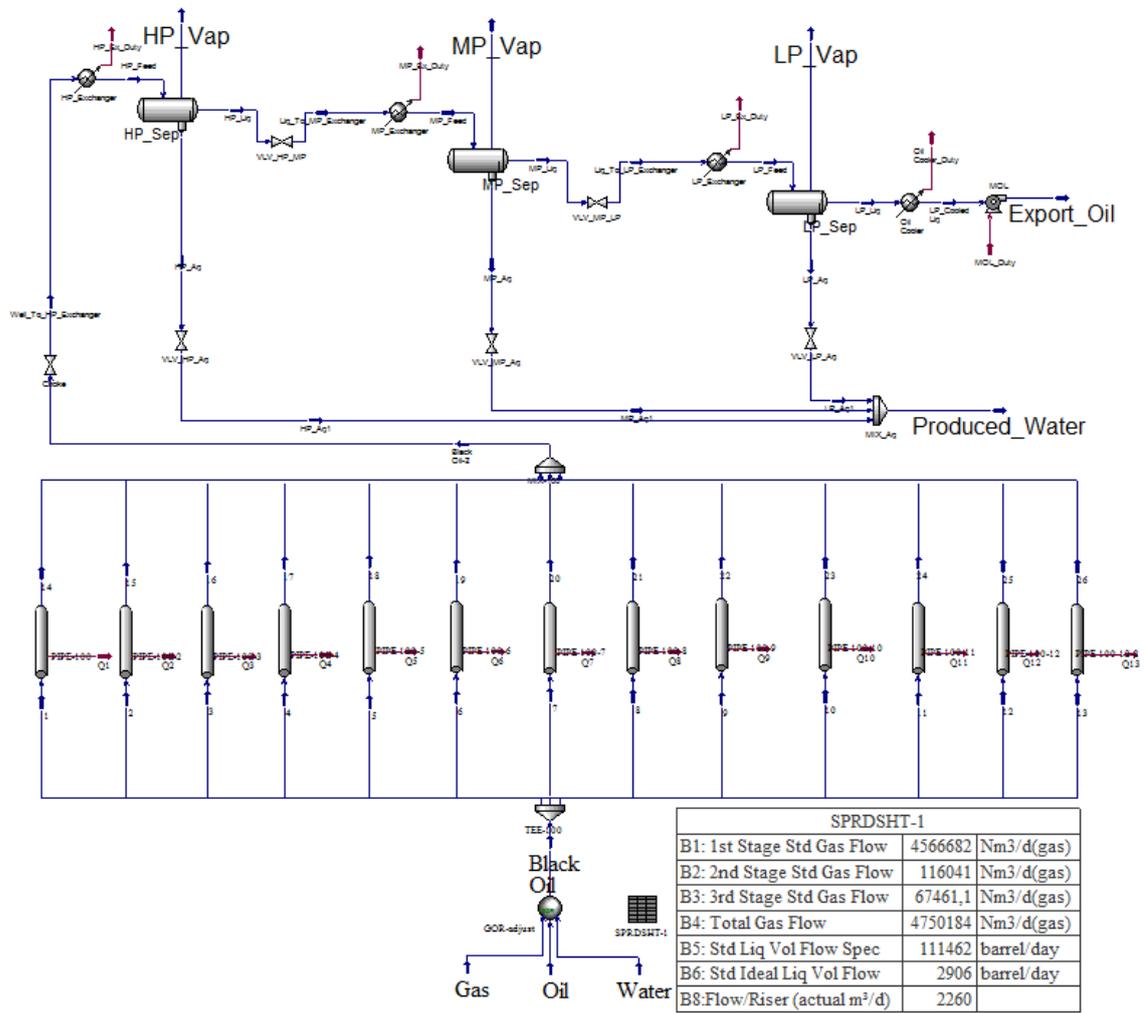


Figure A1.1. Oil Plant Simulation Flowsheet

BASE CASE		DSW CASE	
SEQ	DESCRIPTION	SEQ	DESCRIPTION
	BEGIN CONTROL LOOP FOR 'SVC-101'		BEGIN CONTROL LOOP FOR 'SVC-101'
1	HX 'SHX-101'	1	HX 'SHX-101'
2	CONTROLLER 'SVC-101'	2	CONTROLLER 'SVC-101'
	END CONTROL LOOP FOR 'SVC-101'		END CONTROL LOOP FOR 'SVC-101'
	BEGIN RECYCLE LOOP 'LOOP1'		BEGIN RECYCLE LOOP 'LOOP2'
3	FLASH 'F-101'	3	FLASH 'F-101'
4	VALVE 'V-101'	4	VALVE 'V-101'
5	COMPRESSOR 'C-101'	5	COMPRESSOR 'C-101'
	BEGIN CONTROL LOOP FOR 'SVC-102'	7	HX 'SHX-102'
6	HX 'SHX-102'		BEGIN CONTROL LOOP FOR 'SVC-102'
7	CONTROLLER 'SVC-102'	8	HX 'SHX-103'
	END CONTROL LOOP FOR 'SVC-102'	9	CONTROLLER 'SVC-102'
8	SPLITTER 'SPL-101'		END CONTROL LOOP FOR 'SVC-102'
9	VALVE 'V-102'	10	SPLITTER 'SPL-102'
10	FLASH 'F-301'	11	MIXER 'MIX-101'
11	VALVE 'V-301'	12	VALVE 'V-102'
12	STRM CALC 'MSD-301'	13	FLASH 'F-301'
13	HX 'SHX-301'	14	VALVE 'V-301'
14	HX 'SHX-302'	15	STRM CALC 'MSD-301'
	BEGIN CONTROL LOOP FOR 'SVC-901'	16	HX 'SHX-301'
15	HX 'SHX-303'		BEGIN CONTROL LOOP FOR 'SVC-301'
16	CONTROLLER 'SVC-901'	17	HX 'SHX-302'
	END CONTROL LOOP FOR 'SVC-901'	18	CONTROLLER 'SVC-301'
17	HX 'SHX-901'		END CONTROL LOOP FOR 'SVC-301'
18	FLASH 'F-901'	19	FLASH 'F-302'
19	COMPRESSOR 'C-901'	20	VALVE 'V-302'
20	SPLITTER 'SPL-901'	21	MIXER 'MIX-301'
21	VALVE 'V-902'		BEGIN RECYCLE LOOP 'LOOP1'
	BEGIN CONTROL LOOP FOR 'SVC-902'	22	FLASH 'F-201'
22	HX 'SHX-902'	23	COMPRESSOR 'C-201'
23	VALVE 'V-901'	24	SPLITTER 'SPL-201'
24	CONTROLLER 'SVC-902'		BEGIN CONTROL LOOP FOR 'SVC-201'
	END CONTROL LOOP FOR 'SVC-902'	25	HX 'SHX-201'
25	FLASH 'F-302'	26	CONTROLLER 'SVC-201'
26	VALVE 'V-302'		END CONTROL LOOP FOR 'SVC-201'
27	MIXER 'MIX-301'	27	SPLITTER 'SPL-202'
28	FLASH 'F-201'	28	MIXER 'MIX-201'
29	COMPRESSOR 'C-201'	29	VALVE 'V-201'
	BEGIN CONTROL LOOP FOR 'SVC-201'		END RECYCLE LOOP 'LOOP1'
30	HX 'SHX-201'	30	COMPRESSOR 'C-202'
31	CONTROLLER 'SVC-201'	31	SPLITTER 'SPL-203'
	END CONTROL LOOP FOR 'SVC-201'		BEGIN CONTROL LOOP FOR 'SVC-202'
32	SPLITTER 'SPL-201'	32	HX 'SHX-202'
33	VALVE 'V-201'	33	CONTROLLER 'SVC-202'
34	COMPRESSOR 'C-202'		END CONTROL LOOP FOR 'SVC-202'
	BEGIN CONTROL LOOP FOR 'SVC-202'	34	SPLITTER 'SPL-204'
35	HX 'SHX-202'	35	MIXER 'MIX-202'
36	CONTROLLER 'SVC-202'	36	VALVE 'V-202'
	END CONTROL LOOP FOR 'SVC-202'	37	FLASH 'F-202'
37	SPLITTER 'SPL-202'		BEGIN MVC LOOP FOR 'MVC-401'
38	VALVE 'V-202'	38	HX 'SHX-401'
39	FLASH 'F-202'	39	STRM CALC 'MEM-401'
	END RECYCLE LOOP 'LOOP1'	40	MVC 'MVC-401'
	BEGIN MVC LOOP FOR 'MVC-401'		END MVC LOOP FOR 'MVC-401'
40	HX 'SHX-401'		BEGIN MVC LOOP FOR 'MVC-1001A'
41	STRM CALC 'MEM-401'		ENTER RECYCLE LOOP 'LOOP2'
42	FLASH 'F-002'	41	SPLITTER 'SPL-401'
43	MVC 'MVC-401'	42	VALVE 'V-1001'
	END MVC LOOP FOR 'MVC-401'	43	SPLITTER 'SPL-1001'
	BEGIN MVC LOOP FOR 'MVC-1001A'	44	VALVE 'V-1002A'
44	SPLITTER 'SPL-401'	45	COMPRESSOR 'C-1001A'
45	FLASH 'F-003'	46	CONV REAC 'R-1001A'
46	VALVE 'V-1001'		END RECYCLE LOOP 'LOOP2'
47	SPLITTER 'SPL-1001'		BEGIN RECYCLE LOOP 'LOOP3'
48	VALVE 'V-1002A'	47	EXPANDER 'T-1001A'
49	COMPRESSOR 'C-1001A'	48	SPLITTER 'SPL-1002A'
50	CONV REAC 'R-1001A'	49	HX 'SHX-1001A'
	BEGIN RECYCLE LOOP 'LOOP2'		END RECYCLE LOOP 'LOOP3'
	BEGIN RECYCLE LOOP 'LOOP3'	50	MVC 'MVC-1001A'
51	EXPANDER 'T-1001A'		END MVC LOOP FOR 'MVC-1001A'
52	SPLITTER 'SPL-1002A'		BEGIN RECYCLE LOOP 'LOOP4'
53	HX 'SHX-1001A'	51	FLASH 'F-501'
	END RECYCLE LOOP 'LOOP2'	52	COMPRESSOR 'C-501'
	BEGIN RECYCLE LOOP 'LOOP3'	53	SPLITTER 'SPL-501'
54	MVC 'MVC-1001A'		BEGIN MVC LOOP FOR 'MVC-501'
	END MVC LOOP FOR 'MVC-1001A'		ENTER RECYCLE LOOP 'LOOP4'
	BEGIN RECYCLE LOOP 'LOOP3'	54	HX 'SHX-501'
55	FLASH 'F-501'	55	SPLITTER 'SPL-502'
56	COMPRESSOR 'C-501'	56	MIXER 'MIX-501'
	BEGIN CONTROL LOOP FOR 'SVC-501'	57	VALVE 'V-501'
57	HX 'SHX-501'		END RECYCLE LOOP 'LOOP4'
58	CONTROLLER 'SVC-501'	58	MVC 'MVC-501'
	END CONTROL LOOP FOR 'SVC-501'		END MVC LOOP FOR 'MVC-501'
59	SPLITTER 'SPL-501'		BEGIN RECYCLE LOOP 'LOOP5'
60	VALVE 'V-501'	59	FLASH 'F-502'
	END RECYCLE LOOP 'LOOP3'	60	COMPRESSOR 'C-502'
			BEGIN CONTROL LOOP FOR 'SVC-502'
		61	HX 'SHX-502'
		62	CONTROLLER 'SVC-502'
			END CONTROL LOOP FOR 'SVC-502'
		63	SPLITTER 'SPL-503'
		64	VALVE 'V-502'
			END RECYCLE LOOP 'LOOP5'

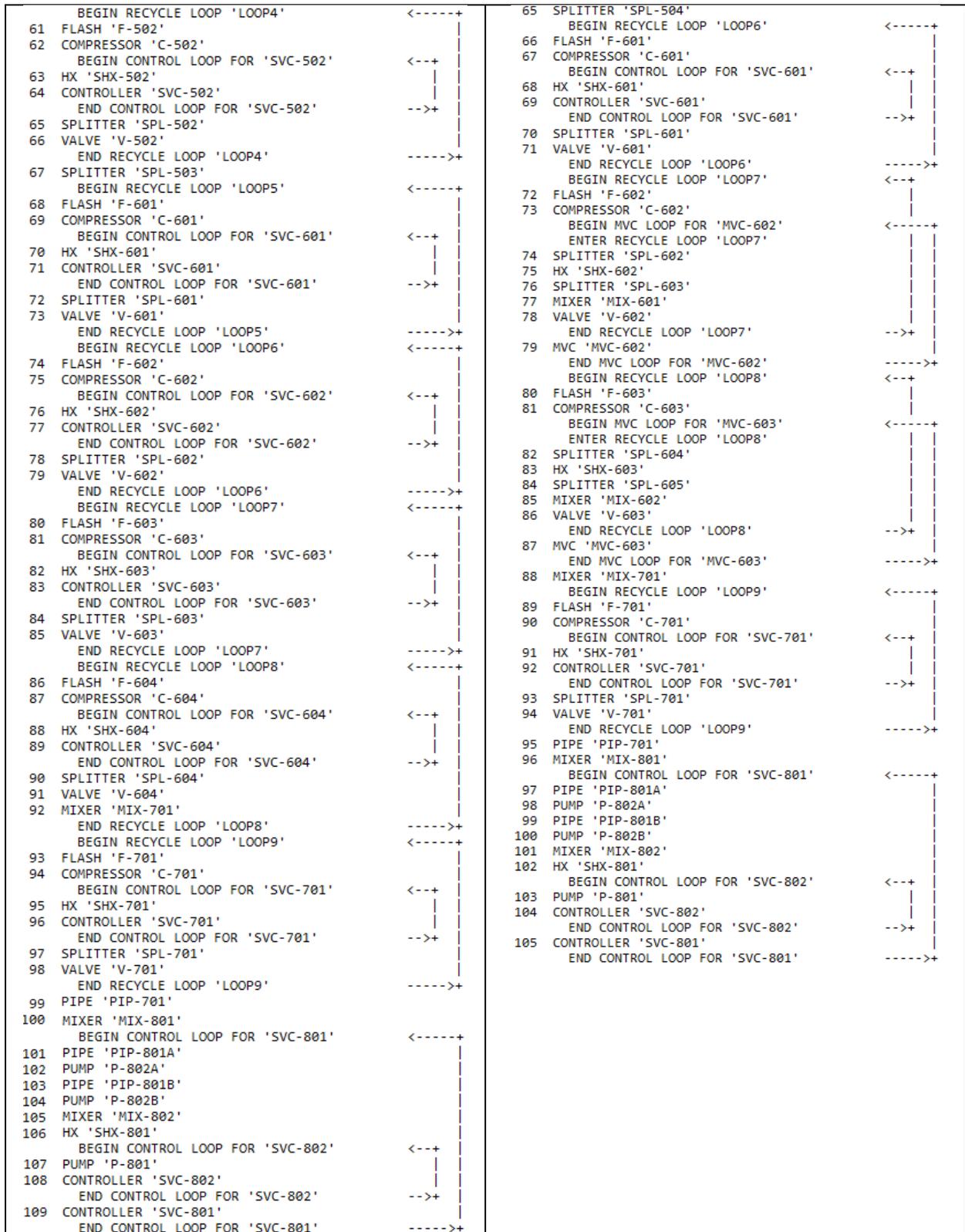


Figure A1.2. Simulation Calculation Sequence

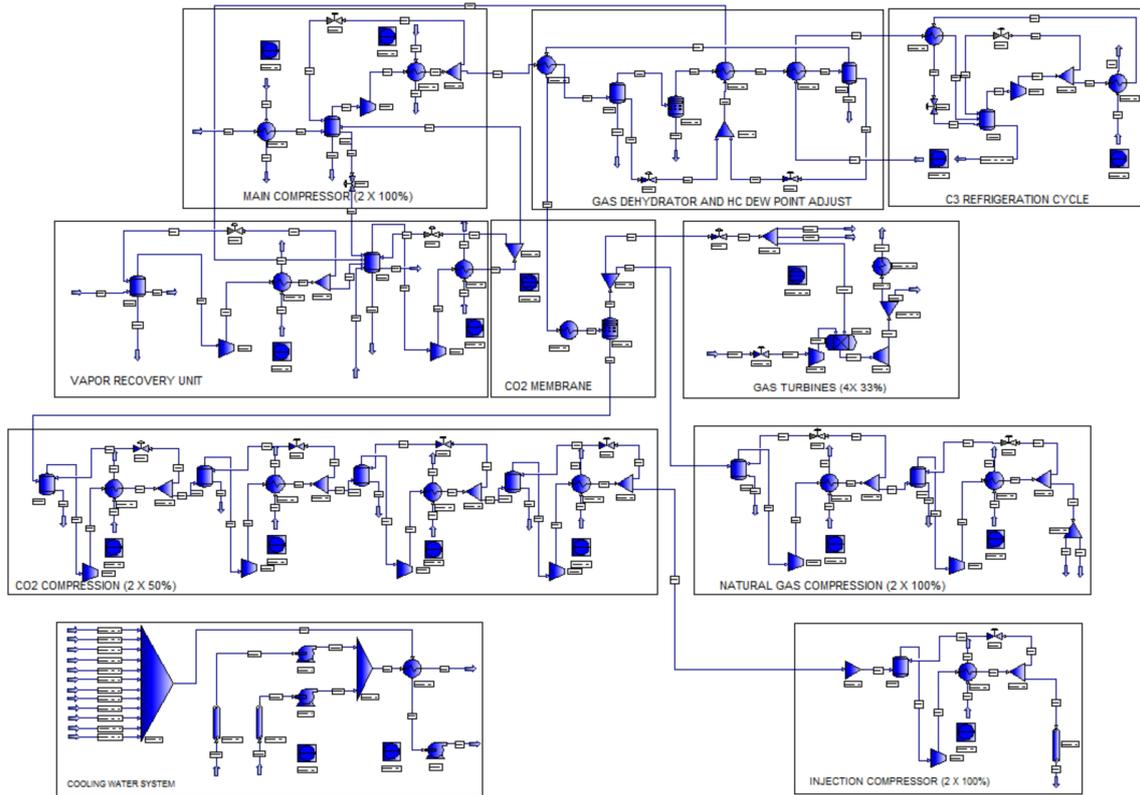


Figure A1.3. Simulation Flowsheet: Base-Case

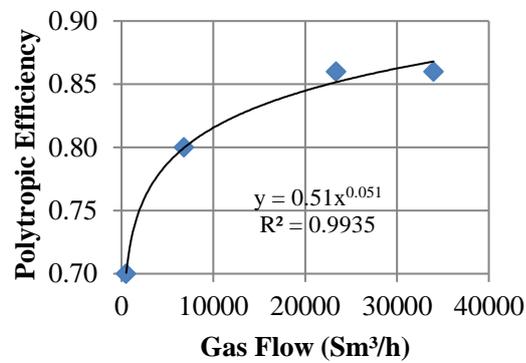


Figure A1.4 – Polytopic efficiency versus gas flow rate (Fitted with data fromn Fig. 13-23 of GPSA [2])

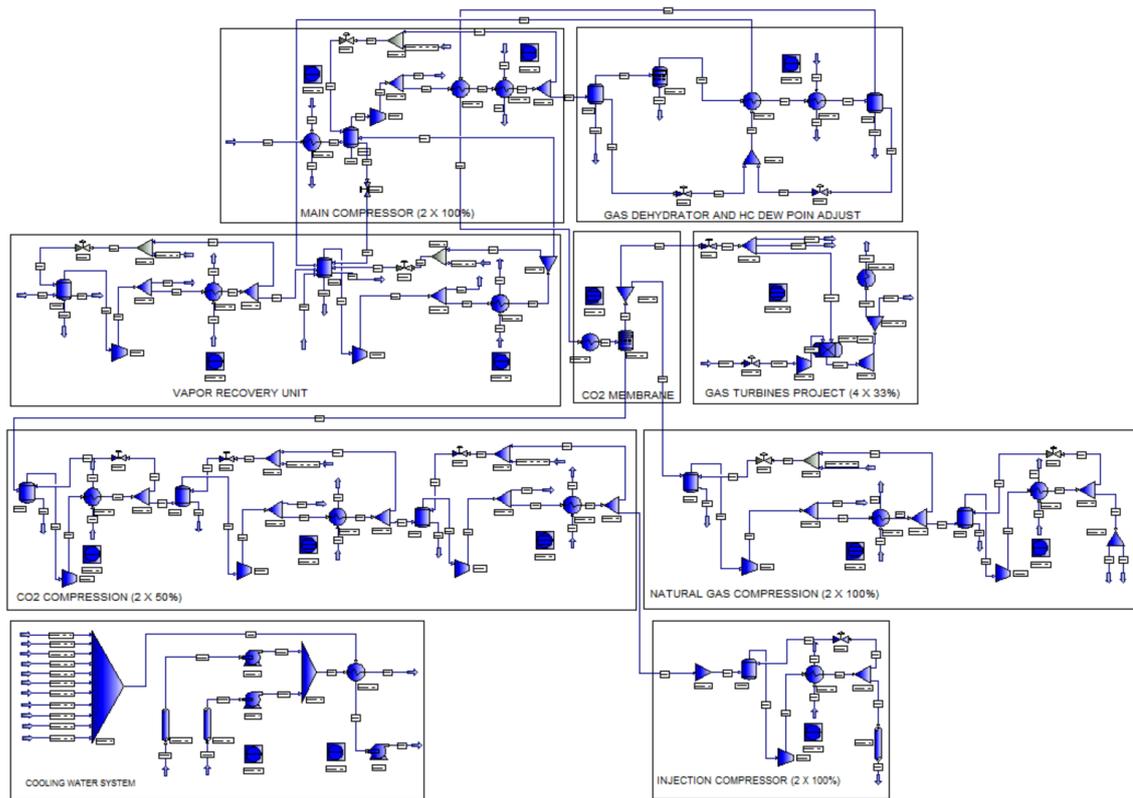


Figure A1.5. Simulation Flowsheet: DSW-Case

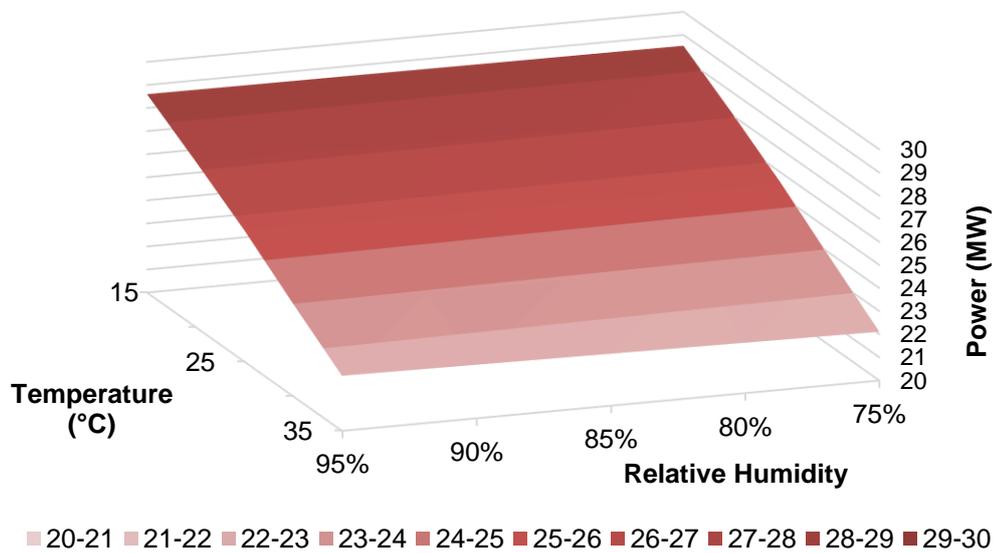
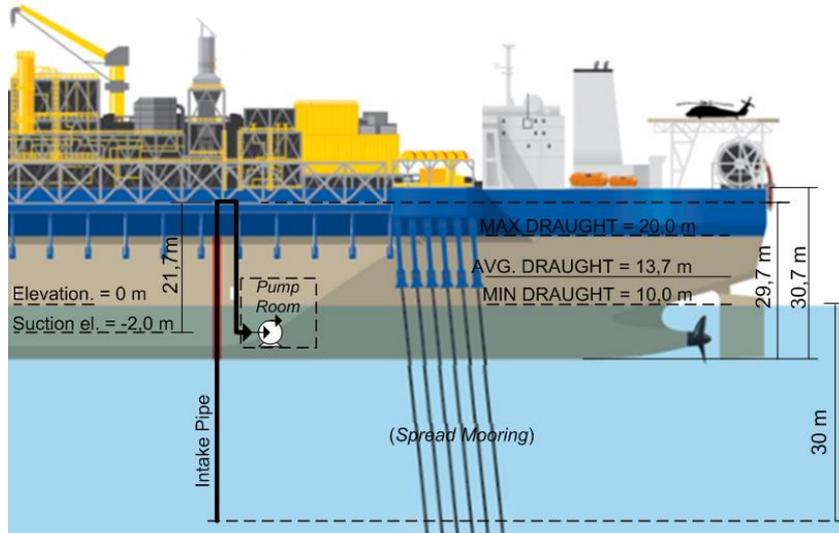


Figure A1.6. GE LM2500 GT Power vs Air Temperature and Relative Humidity



**Figure A1.7. Base-Case Seawater Intake System  
(based on EIA [1])**

**Table A1.1. Simulation of Plant Inlet Streams (EIA (Petrobras, 2013))**

	OIL	GAS	WATER
Temperature (°C)	40	40	40
Pressure (kPa)	41368	41368	41368
Composition (mol fraction)			
CO <sub>2</sub>	0.0000	0.1513	0.0000
H <sub>2</sub> S	0.0000	0.0000	0.0000
N <sub>2</sub>	0.0000	0.0051	0.0000
Methane	0.0000	0.6396	0.0000
Ethane	0.0000	0.0924	0.0000
Propane	0.0050	0.0591	0.0000
i-Butane	0.0019	0.0099	0.0000
n-Butane	0.0081	0.0207	0.0000
i-Pentane	0.0065	0.0052	0.0000
n-Pentane	0.0122	0.0075	0.0000
C <sub>6</sub> *	0.0295	0.0063	0.0000
C <sub>7</sub> *	0.0559	0.0009	0.0000
C <sub>8</sub> *	0.0777	0.0016	0.0000
C <sub>9</sub> *	0.0658	0.0005	0.0000
C <sub>10</sub> *	0.0604	0.0000	0.0000
C <sub>11</sub> *	0.0405	0.0000	0.0000
C <sub>12</sub> *	0.0537	0.0000	0.0000
C <sub>13</sub> *	0.0488	0.0000	0.0000
C <sub>14</sub> *	0.0436	0.0000	0.0000
C <sub>15</sub> *	0.0308	0.0000	0.0000
C <sub>16</sub> *	0.0339	0.0000	0.0000
C <sub>17</sub> *	0.0224	0.0000	0.0000
C <sub>18</sub> *	0.0236	0.0000	0.0000
C <sub>19</sub> *	0.0224	0.0000	0.0000
C <sub>20</sub> +	0.3573	0.0000	0.0000
H <sub>2</sub> O	0.0000	0.0000	1.0000
MM C <sub>20</sub> +		500	
Density C <sub>20</sub> +		0.9496	

**Table A1.2. Compressors Flows and Anti-Surge Recycle – Base-Case**

Compressor	Design Flow (MMSm <sup>3</sup> /d)	Base-Case 100% Operation 5.0 MM Sm <sup>3</sup> /d		Base-Case 75% Operation 3.75 MM Sm <sup>3</sup> /d		Base-Case 50% Operation 2.5 MM Sm <sup>3</sup> /d		Base-Case 25% Operation 1.25 MM Sm <sup>3</sup> /d	
		Flow	Recycle	Flow	Recycle	Flow	Recycle	Flow	Recycle
		C-101	5.082	5.082	0.0%	5.082	25.2%	5.081	50.3%
C-201	0.071	0.071	0.0%	0.071	26.0%	0.071	52.0%	0.071	76.0%
C-202	0.302	0.302	0.0%	0.301	26.0%	0.301	52.0%	0.301	77.0%
C-501	3.500	3.506	0.0%	3.506	27.5%	3.508	50.1%	3.504	82.8%
C-502	3.500	3.506	0.0%	3.506	27.5%	3.508	50.1%	3.504	82.8%
C-601	1.200	1.200	16.6%	1.200	37.5%	1.200	58.3%	1.200	79.1%
C-602	1.200	1.200	16.6%	1.200	37.5%	1.200	58.3%	1.200	79.1%
C-603	1.200	1.200	16.6%	1.200	37.5%	1.200	58.3%	1.200	79.1%
C-604	1.200	1.200	16.6%	1.200	37.5%	1.200	58.3%	1.200	79.1%
C-701	4.230	4.231	76.4%	4.230	82.3%	4.230	88.2%	4.230	94.1%
C-901	0.254	0.254	0.0%	0.254	18.5%	0.254	37.8%	0.254	57.0%

**Table A1.3. Heat Exchangers Specification – Base-Case**

ID in Fig. 10	TAG	Utility Fluid	T <sub>in</sub> (°C)	T <sub>out</sub> (°C)	ΔP (kPa)	Specification	Value (°C)	ΔP gas (kPa)
1	SHX-101	CW	35	45	100	Hot Fluid T	40	50
2	SHX-102	CW	35	55	50	Hot Fluid T	40	50
3	SHX-201	CW	35	50	50	Hot Fluid T	40	25
4	SHX-202	CW	35	55	50	Hot Fluid T	40	50
5	SHX-301	-				Hot Fluid T	5	50
6	SHX-302	-				Hot T <sub>in</sub> – Cold T <sub>out</sub>	3	50
7	SHX-303	Propane	0	0	13.8	Hot Fluid T	10	50
8	SHX-501	CW	35	55	100	Hot Fluid T	40	50
9	SHX-502	CW	35	55	100	Hot Fluid T	40	50
10	SHX-601	CW	35	55	50	Hot Fluid T	40	25
11	SHX-602	CW	35	55	50	Hot Fluid T	40	25
12	SHX-603	CW	35	55	100	Hot Fluid T	40	50
13	SHX-604	CW	35	55	50	Hot Fluid T	40	50
14	SHX-701	CW	35	55	50	Hot Fluid T	40	50
15	SHX-801	SW	32	40	60	Hot Fluid T	35	60
16	SHX-901	-				°C Above Dew Point	10	0
17	SHX-902	CW	35	55	50	Hot Fluid Vapor Fraction	0	0

**Table A1.4. Anti-Surge Control Loop, Hot Bypass Flow and Minimum Allowable Temperature – DSW-Case (100% gas capacity)**

Compressor	Design Mass Flow (MMSm <sup>3</sup> /d)	Simulated Flow (MM Sm <sup>3</sup> /d)	% Anti-Surge Recycle	Hot Bypass (Sm <sup>3</sup> /d)	Hydrate T (°C)	Min. T (°C)
C-101	4.956	4.956	0.0%	0	8.7/0	5
C-201	0.071	0.071	0.0%	0	-7.8/0	5
C-202	0.238	0.238	0.0%	0	3.6/0	5
C-501	3.500	3.520	0.0%	0	-	5
C-502	3.500	3.520	0.0%	0	-	5
C-601	1.200	1.201	16.5%	0	-	5
C-602	1.200	1.201	16.5%	4228	-	5
C-603	1.200	1.201	16.5%	51291	-	5
C-701	4.200	4.230	76.3%	0	-	5

**Table A1.5. Surge Control Loop, Hot Bypass Flow and Minimum Allowable Temperature – DSW-Case (75% gas capacity)**

Compressor	Design Mass Flow (MMSm <sup>3</sup> /d)	Simulated Flow (MM Sm <sup>3</sup> /d)	% Anti-Surge Recycle	Hot Bypass (Sm <sup>3</sup> /d)	Hydrate T (°C)	Min. T (°C)
C-101	4.956	4.956	25.0%	300000	8.7/0	5
C-201	0.071	0.071	27.0%	1500	-7.8/0	5
C-202	0.238	0.239	26.0%	12500	3.6/0	5
C-501	3.500	3.503	27.1%	180000	N/P	5
C-502	3.500	3.503	27.1%	0	N/P	5
C-601	1.200	1.200	37.5%	0	N/P	5
C-602	1.200	1.200	37.5%	75222	N/P	5
C-603	1.200	1.200	37.5%	191159	N/P	5
C-701	4.200	4.230	82.3%	0	N/P	5

**Table A1.6. Anti-Surge Control Loop, Hot Bypass Flow and Minimum Allowable Temperature – DSW-Case (50% gas capacity)**

Compressor	Design Mass Flow (MMSm <sup>3</sup> /d)	Simulated Flow (MM Sm <sup>3</sup> /d)	% Anti-Surge Recycle	Hot Bypass (Sm <sup>3</sup> /d)	Hydrate T (°C)	Min. T (°C)
C-101	4.956	4.956	50.0%	525000	8.7/0	5
C-201	0.071	0.070	52.5%	3750	-7.8/0	5
C-202	0.238	0.239	51.0%	39000	3.5/0	5
C-501	3.500	3.502	54.5%	820000	N/P	5
C-502	3.500	3.502	54.5%	0	N/P	5
C-601	1.200	1.201	58.4%	0	N/P	5
C-602	1.200	1.201	58.4%	146977	N/P	5
C-603	1.200	1.201	58.4%	332054	N/P	5
C-701	4.200	4.230	88.2%	0	N/P	5

**Table A1.7. Anti-Surge Control Loop, Hot Bypass Flow and Minimum Allowable Temperature – DSW-Case (25% gas capacity)**

Compressor	Design Mass Flow (MMSm <sup>3</sup> /d)	Simulated Flow (MM Sm <sup>3</sup> /d)	% Anti-Surge Recycle	Hot Bypass (Sm <sup>3</sup> /d)	Hydrate T (°C)	Min. T (°C)
C-101	4.956	4.956	75.0%	750000	8.7/0	5
C-201	0.071	0.071	77.0%	6750	-7.8/0	5
C-202	0.238	0.239	76.1%	58000	3.5/0	5
C-501	3.500	3.503	82.3%	1440000	N/P	5
C-502	3.500	3.503	82.3%	0	N/P	5
C-601	1.200	1.200	79.2%	0	N/P	5
C-602	1.200	1.200	79.2%	217204	N/P	5
C-603	1.200	1.200	79.2%	470246	N/P	5
C-701	4.200	4.230	94.1%	0	N/P	5

**Table A1.8. Heat Exchangers Specification – DSW-Case**

ID Fig. 10	TAG	Utility Fluid	T <sub>in</sub> (°C)	T <sub>out</sub> (°C)	ΔP (kPa)	Specification	Value (°C)	ΔP gas (kPa)
1	SHX-101	CW	7	17	100	Hot Fluid T	12	50
2	SHX-103	CW	7	27	50	Hot Fluid T	20	50
3	SHX-201	CW	7	27	50	Hot Fluid T	12	25
4	SHX-202	CW	7	27	50	Hot Fluid T	12	50
5	SHX-102	-	-	-	50	Cold Fluid T	35	50
6	SHX-301	-	-	-	50	Hot T <sub>in</sub> – Cold T <sub>out</sub>	3	50
7	SHX-302	CW	7	13	50	Hot Fluid T	10	50
8	SHX-501	CW	7	27	100	Hot Fluid T	12	50
9	SHX-502	CW	7	27	100	Hot Fluid T	40	50
10	SHX-601	CW	7	27	50	Hot Fluid T	12	25
11	SHX-602	CW	7	27	50	Hot Fluid T	12	25
12	SHX-603	CW	7	27	100	Hot Fluid T	12	50
14	SHX-701	CW	7	27	50	Hot Fluid T	40	50
15	SHX-801	SW	4	11	60	Hot Fluid T	7	60

**Table A1.9. Fuel Gas Properties**

	CASE	BASE	DSW
Pressure (kPa)		3500	3500
Temperature (°C)		22.5	22.5
LHV (kJ/kg)		43487	43486
HHV (kJ/kg)		47848	47847
Molar Mass (kg/kgmol)		22.91	22.92
COMPOSITION (MOLAR FRACTION)			
N <sub>2</sub>		0.00672	0.00673
CO <sub>2</sub>		0.05000	0.05000
Methane		0.69910	0.70100
Ethane		0.14380	0.14000
Propane		0.06802	0.06966
n-Butane		0.01684	0.01687
n-Pentane		0.00567	0.00583
Hexane		0.00093	0.00099
Isobutane		0.00892	0.00892

**Table A1.10. FPSO Typical Electricity Consumers**

ENGINE ROOM AND VESSEL	POWER (kW)
Compressors	1000
Oil Treatment Panel	300
Engine Command Panel	2190
Cranes	290
Offloading Hose Reel	150
Hot Water Pumps	90
Inert Gas Generator	125
Deaerator Pump	132
Naval Lighting	150
Hypochlorite Generator	330
Main Generation/ Auxiliary Equipment	112
Ballast System	110
Emergency Cooling Pump	95
Emergency Hot Water	90
Inert Gas Generator Cooling Pump	110
<b>ACCOMMODATIONS -100 people</b>	
Miscellaneous Equipment	157
Gym	0.59
Laundry	7.14
Hospital	2.89
Kitchen	36.34
Others	39.14
Electric Devices	96.50
<b>PROCESS (Oil, Gas, Water)</b>	
Flare System	2300
CO <sub>2</sub> removal	1500
Gas Dehydration, Fuel Gas and HCDP	560
Injection Manifolds	155
Oil Processing and Treatment	5000
Sulfates Removal and Water Injection System	300
Chemical Storage	3500
Utilities	515
Laydown Area	65
Automation	1200
Pipe Rack	1450
Flare Tower	655
Laboratory	350
Cooling Water Pumps	2000
<b>TOTAL</b>	<b>23.16 MW</b>

(adapted from Martins et. al (2014) (Martins; Delfino; Fachini, 2014))

## Supplement A2 – Equipment Sizing

**Table A2.1. Heat Exchangers Sizing – Base-Case**

	RHX-101	RHX-102	RHX-201	RHX-202
Heat Duty	SHX-101	SHX-102	SHX-201	SHX-202
Hot Fluid Side	Tubes	Tubes	Tubes	Tubes
Cold Fluid Side	Shell	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	50	50	25	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	25	50	50	50
Number of Shells in Serie	2	1	1	1
Number of Shells in Parallel	1	1	1	1
Number of Tube Passes /Shell	1	4	2	2
TEMA Type	AEL	NFU	NFU	NFU
Shell Diameter (mm)	900	1800	350	500
Shell Material	A 516	A 516	A 516	A 516
Tubes Material	Inconel	Inconel	Inconel	Inconel
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	25.4	19.05	19.05	19.05
Tubes BWG	18	18	18	18
Tubes Length (m)	5.7	6.096	6.096	6.096
Pitch (mm)	31.7	25.4	25.4	25.4
Tubes Pattern	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Baffles Type	Single	Double	Single	Single
Baffles Cut (%)	25	30	25	25
Baffles Spacing (mm)	250	350	150	170
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.1. Heat Exchangers Sizing – Base-Case (Continued)**

	RHX-301	RHX-302	RHX-303	RHX-501
Heat Duty	SHX-301	SHX-302	SHX-303	SHX-501
Hot Fluid Side	Shell	Tubes	Tubes	Tubes
Cold Fluid Side	Tubes	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	25	50	50	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	25	50	0	50
Number of Shells in Series	2	1	1	2
Number of Shells in Parallel	1	1	1	1
Number of Tube Passes /Shell	2	2	2	2
TEMA Type	BHM	NFU	NFU	DFU
Shell Diameter (mm)	1150	750	800	800
Shell Material	A 516	SS316	A 516	A 516
Tubes Material	Inconel	316LS	316LS	316LS
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	19.05	25.4	25.4	19.05
Tubes BWG	18	18	18	18
Tubes Length (m)	11.6	4.5	6.6	10.6
Pitch (mm)	25.4	31.7	31.7	25.4
Tubes Pattern	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Baffles Type	Single	Single	Single	Double
Baffles Cut (%)	25	25	25	33
Baffles Spacing (mm)	350	120	150	600
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.1. Heat Exchangers Sizing – Base-Case (Continued)**

	RHX-502	RHX-601	RHX-602	RHX-603
Heat Duty	SHX-502	SHX-601	SHX-602	SHX-603
Hot Fluid Side	Tubes	Tubes	Tubes	Tubes
Cold Fluid Side	Shell	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	50	50	50	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	50	50	50	50
Number of Shells in Serie	2	1	2	-
Number of Shells in Parallel	1	2	1	-
Number of Tube Passes /Shell	2	2	2	-
TEMA Type	DFU	NFU	NFU	DFU
Shell Diameter (mm)	800	900	700	550
Shell Material	A 516	A 516	A 516	A 516
Tubes Material	316LS	A179	A179	316LS
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	19.05	19.05	19.05	19.05
Tubes BWG	16	18	18	18
Tubes Length (m)	11.8	8.4	11	9.4
Pitch (mm)	25.4	25.4	25.4	25.4
Tubes Pattern	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Baffles Type	Double	Single	Double	-
Baffles Cut (%)	33	30	30	-
Baffles Spacing (mm)	750	300	500	-
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.1. Heat Exchangers Sizing – Base-Case (Continued)**

	RHX-604	RHX-701	RHX-901	RHX-902
Heat Duty	SHX-604	SHX-701	SHX-901	SHX-902
Hot Fluid Side	Tubes	Shell	Shell	Tubes
Cold Fluid Side	Shell	Tubes	Tubes	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	50	50	0	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	50	50	0	50
Number of Shells in Serie	-	-	1	1
Number of Shells in Parallel	-	-	1	1
Number of Tube Passes /Shell	-	-	1	6
TEMA Type	DFU	DFU	BEM	AJL
Shell Diameter (mm)	-	-	325	850
Shell Material	A516	A516	A 516	A 516
Tubes Material	316LS	316LS	A179	A179
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	19.05	19.05	25.4	19.05
Tubes BWG	-	-	18	18
Tubes Length (m)	-	-	1.5	5.2
Pitch (mm)	-	-	31.7	25.4
Tubes Pattern	-	-	Triangular 30°	Triangular 30°
Baffles Type	-	-	-	Single
Baffles Cut (%)	-	-	-	25
Baffles Spacing (mm)	-	-	-	80
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.2. Plate Heat Exchangers HX-801 Sizing – Base-Case**

	Cold Side	Hot Side
Fluid	SW	CW
Mass Flow (kg/s)	calculated	162.9 x 4
Inlet Temperature (°C)	32	53.6
Outlet Temperature (°C)	40	35
Outlet Pressure (kPa abs.)	800	500
$\Delta P$ max (kPa)	60	60
Fouling Resistance (m <sup>2</sup> .K/W)	0.3	0.2

**Table A2.3. Heat Exchangers Sizing – DSW-Case**

	RHX-101	RHX-102	RHX-103	RHX-201	RHX-202
Heat Duty	SHX-101	SHX-102	SHX-103	SHX-201	SHX-202
Hot Fluid Side	Tubes	Shell	Tubes	Tubes	Tubes
Cold Fluid Side	Shell	Tubes	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}$ /Shell (kPa)	50	50	50	50	50
$\Delta P_{\max \text{ tubes}}$ /Shell (kPa)	25	50	50	25	25
Number of Shells in Serie	2	1	1	1	2
Number of Shells in Parallel	1	2	1	1	1
Number of Tube Passes /Shell	2	1	2	2	2
TEMA Type	AEL	BHM	NFU	NFU	NFU
Shell Diameter (mm)	1280	450	1000	350	400
Shell Material	A 516				
Tubes Material	Inconel	Inconel	Inconel	Inconel	Inconel
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	25.4	19.05	19.05	19.05	19.05
Tubes BWG	18	18	18	18	18
Tubes Length (m)	4.3	3.1	11.6	12	8.4
Pitch (mm)	31.7	25.4	25.4	25.4	25.4
Tubes Pattern	Triangular 30°				
Baffles Type	Single	Single	Double	Single	Single
Baffles Cut (%)	25	25	30	25	25
Baffles Spacing (mm)	400	250	500	200	250
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611	0.17611

**Table A2.3. Heat Exchangers Sizing – DSW-Case (continued)**

	RHX-301	RHX-302	RHX-501	RHX-502
Heat Duty	SHX-301	SHX-302	SHX-501	SHX-502
Hot Fluid Side	Shell	Tubes	Tubes	Tubes
Cold Fluid Side	Tubes	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	50	50	50	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	50	50	50	50
Number of Shells in Serie	1	2	2	1
Number of Shells in Parallel	1	1	1	1
Number of Tube Passes /Shell	6	2	2	2
TEMA Type	BHM	NFU	DFU	DFU
Shell Diameter (mm)	500	1000	1050	620
Shell Material	A 516	A 516	A 516	A 516
Tubes Material	SS 316	SS 316	SS 316	SS 316
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	25.4	25.4	19.05	19.05
Tubes BWG	18	18	18	16
Tubes Length (m)	4.7	10.4	9	9
Pitch (mm)	31.7	31.7	25.4	25.4
Tubes Pattern	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Baffles Type	Single	Double	Double	Double
Baffles Cut (%)	25	30	33	33
Baffles Spacing (mm)	500	280	850	400
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.3. Heat Exchangers Sizing – DSW-Case (continued)**

	RHX-601	RHX-602	RHX-603	RHX-701
Heat Duty	SHX-601	SHX-602	SHX-603	SHX-701
Hot Fluid Side	Tubes	Tubes	Tubes	Tubes
Cold Fluid Side	Shell	Shell	Shell	Shell
$\Delta P_{\max \text{ shell}}/\text{Shell}$ (kPa)	50	50	50	50
$\Delta P_{\max \text{ tubes}}/\text{Shell}$ (kPa)	50	50	50	50
Number of Shells in Serie	1	1	-	-
Number of Shells in Parallel	1	1	-	-
Number of Tube Passes /Shell	2	2	-	-
TEMA Type	NFU	NFU	DFU	DFU
Shell Diameter (mm)	900	780	-	-
Shell Material	A 516	A 516	A516	A516
Tubes Material	A179	A179	SS316	SS316
Tubes Abs. Roughness (mm)	0.04572	0.04572	0.04572	0.04572
Tubes Outer Diameter (mm)	19.05	19.05	19.05	19.05
Tubes BWG	18	18	-	-
Tubes Length (m)	9.2	12	-	-
Pitch (mm)	25.4	25.4	-	-
Tubes Pattern	Triangular 30°	Triangular 30°	-	-
Baffles Type	Single	Single	-	-
Baffles Cut (%)	25	30	-	-
Baffles Spacing (mm)	500	650	-	-
Fouling Resistance (m <sup>2</sup> .K/kW)	0.17611	0.17611	0.17611	0.17611

**Table A2.4. Plate Heat Exchangers HX-801 Sizing – DSW-Case**

	Cold Side	Hot Side
Fluid	SW	CW
Mass Flow (kg/s)	818.5 x 2	133.1 x 5
Inlet Temperature (°C)	4	23.8
Outlet Temperature (°C)	11	7
Outlet Pressure (kPa abs.)	800	500
ΔP max (kPa)	60	60
Fouling Resistance (m <sup>2</sup> .K/W)	0.3	0.2

**Supplement A3 – Cost and Weight Estimation****Table A3.1. Inputs for Compressors CAPEX Estimation – Base-Case**

	C-101	C-201	C-202	C-501	C-502	C-601
Compressor Material	CS	CS	CS	CS	CS	CS
Inlet Flow (m <sup>3</sup> /h)	11975.6	1238.5	1838.7	2894.0	1110.4	12939.3
Inlet Pressure (kPag)	1699	149	566	4399	10491	299
Inlet Temperature (°C)	39.48	55	25.01	28.86	40	30
Discharge Pressure (kPag)	5149	591	1749	10541	24949	1034
Gas Mol. Mass	25.35	39.83	34.50	22.56	22.56	31.87
Cp/Cv	1.3049	1.1579	1.2179	1.4858	1.8807	1.308
Z <sub>inlet</sub>	0.9462	0.9837	0.9517	0.8555	0.7451	0.9879
Z <sub>outlet</sub>	0.9484	0.9744	0.9317	0.8923	0.9366	0.9887
Tubes Material	SS316L	SS316L	SS316L	CS	SS316L	CS
Driver	Motor	Motor	Motor	Motor	Motor	Motor

**Table A3.1. Inputs for Compressors CAPEX Estimation – Base-Case (continued)**

	C-602	C-603	C-604	C-701	C-901
Compressor Material	CS	CS	CS	CS	CS
Inlet Flow (m <sup>3</sup> /h)	605	609	613	702	904
Inlet Pressure (kPag)	606	610	614	703	905
Inlet Temperature (°C)	4727.7	1583.2	466.5	312.8	1884.9
Discharge Pressure (kPag)	1009	3025	8724	24899	375
Gas Mol. Mass	40	40	40	37.42	0
Cp/Cv	3050	8774	24949	54949	1637
Z <sub>inlet</sub>	31.87	31.87	31.87	25.22	44.10
Z <sub>outlet</sub>	1.3316	1.436	1.9648	n/a	1.1992
Tubes Material	0.97	0.9147	0.7607	n/a	0.898
Driver	0.976	0.949	0.9419	n/a	0.7835
Compressor Material	CS	SS316L	SS316L	SS316L	CS
Inlet Flow (m <sup>3</sup> /h)	Motor	Motor	Motor	Motor	Motor

**Table A3.2. Inputs for Heat Exchangers CAPEX Estimation – Base-Case**

	SHX-101	SHX-102	SHX-201	SHX-202	SHX-301
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	587.50	1093.00	41.00	108.50	1153.00
Number of Shells <sup>[2]</sup>	2	1	1	1	2
TEMA Type <sup>[3]</sup>	AEL	NFU	NFU	NFU	BHM
Tubes Material <sup>[2]</sup>	Inconel	Inconel	Inconel	Inconel	Inconel
Tubes Operation P (kPag) <sup>[1]</sup>	1749	5149	591	1749	4799
Tubes Design P (kPag) <sup>[4]</sup>	2094	5494	761	2094	5144
Tubes Operation T (°C) <sup>[1]</sup>	55	126	116	92	35
Tubes Design T (°C) <sup>[5]</sup>	125	156	146	125	125
Tubes Ext. Diameter (mm) <sup>[2]</sup>	25.4	25.4	19.1	19.1	25.4
Shell Material <sup>[2]</sup>	A 516				
Shell Operation P (kPag) <sup>[1]</sup>	438	438	438	438	5099
Shell Design P (kPag) <sup>[4]</sup>	608	608	608	608	5444
Shell Operation T (°C) <sup>[1]</sup>	45	55	50	55	40
Shell Design T (°C) <sup>[5]</sup>	125	125	125	125	125
Tubes Length (m) <sup>[2]</sup>	5.70	9.00	10.00	12.20	11.60
Tubes BWG <sup>[2]</sup>	18	18	18	18	18
Pitch (mm) <sup>[2]</sup>	31.7	25.4	25.4	25.4	25.4
Tubes Pattern <sup>[2]</sup>	Triangular 30°				
Shell Diameter (mm) <sup>[1]</sup>	900	1800	350	500	1150
Channel Material	A 516				
Head Material	A 516				
Clad Material	Inconel	Inconel	Inconel	Inconel	Inconel
Clad Side	Head	Head	Head	Head	Head
Number of Tube Passes/Shell <sup>[2]</sup>	1	4	2	2	2
Number of /Shell <sup>[2]</sup>	1	2	2	2	4

**Table A3.2. Inputs for Heat Exchangers CAPEX Estimation – Base-Case (continued)**

	SHX-302	SHX-303	SHX-501	SHX-502	SHX-601
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	1153	76,8	128,2	505,5	563,6
Number of Shells <sup>[2]</sup>	2	1	1	2	2
TEMA Type <sup>[3]</sup>	BHM	NFU	NFU	DFU	DFU
Tubes Material <sup>[2]</sup>	Inconel	316LS	316LS	316LS	316LS
Tubes Operation P (kPag) <sup>[1]</sup>	4799	4899	4849	10541	24949
Tubes Design P (kPag) <sup>[4]</sup>	5144	5244	5194	11068	26196
Tubes Operation T (°C) <sup>[1]</sup>	35	21,6	18	102	110
Tubes Design T (°C) <sup>[5]</sup>	125	22	22	132	140
Tubes Ext. Diameter (mm) <sup>[2]</sup>	19.05	25.4	25.4	19.05	19.05
Shell Material <sup>[2]</sup>	A 516	SS316	A 516	A 516	A 516
Shell Operation P (kPag) <sup>[1]</sup>	5099	616	375	438	438
Shell Design P (kPag) <sup>[4]</sup>	5444	786	545	608	608
Shell Operation T (°C) <sup>[1]</sup>	40	-11	-1	55	55
Shell Design T (°C) <sup>[5]</sup>	125	-41	-31	125	125
Tubes Length (m) <sup>[2]</sup>	11.6	4.5	6.6	10.6	11.8
Tubes BWG <sup>[2]</sup>	18	18	18	18	16
Pitch (mm) <sup>[2]</sup>	25.4	31.7	31.7	25.4	25.4
Tubes Pattern <sup>[2]</sup>	Triangular	Triangular	Triangular	Triangular	Triangular
	30°	30°	30°	30°	30°
Shell Diameter (mm) <sup>[1]</sup>	1150	750	800	800	800
Channel Material	A 516	316L	A 516	316L	316L
Head Material	A 516	316L	A 516	316L	316L
Clad Material	Inconel	-	-	-	-
Clad Side	Head	-	-	-	-
Number of Tube Passes/Shell <sup>[2]</sup>	2	2	2	2	2
Number of Passes /Shell <sup>[2]</sup>	4	2	2	2	2

**Table A3.2. Inputs for Heat Exchangers CAPEX Estimation – Base-Case (continued)**

	SHX-602	SHX-603	SHX-901	SHX-902
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	198.7	203.3	8.64	266.1
Number of Shells <sup>[2]</sup>	1	2	1	1
TEMA Type <sup>[3]</sup>	NFU	DFU	BEM	AJL
Tubes Material <sup>[2]</sup>	A179	316LS	A179	A179
Tubes Operation P (kPag) <sup>[1]</sup>	3050	8774	375	438
Tubes Design P (kPag) <sup>[4]</sup>	3395	9213	545	608
Tubes Operation T (°C) <sup>[1]</sup>	142	150	10	45
Tubes Design T (°C) <sup>[5]</sup>	172	180	22	125
Tubes Ext. Diameter (mm) <sup>[2]</sup>	19.05	19.05	19.05	19.05
Shell Material <sup>[2]</sup>	A 516	A 516	A 516	A 516
Shell Operation P (kPag) <sup>[1]</sup>	438	438	1637	1637
Shell Design P (kPag) <sup>[4]</sup>	608	608	1807	1807
Shell Operation T (°C) <sup>[1]</sup>	55	55	50	66
Shell Design T (°C) <sup>[5]</sup>	125	125	125	125
Tubes Length (m) <sup>[2]</sup>	11	9.4	1.5	5.2
Tubes BWG <sup>[2]</sup>	18	18	18	18
Pitch (mm) <sup>[2]</sup>	25.4	25.4	31.7	25.4
Tubes Pattern <sup>[2]</sup>	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Shell Diameter (mm) <sup>[1]</sup>	700	550	325	850
Channel Material	A 516	316L	A 516	A 516
Head Material	A 516	316L	A 516	A 516
Clad Material	-	-	-	-
Clad Side	-	-	-	-
N° Tube Passes/Shell <sup>[2]</sup>	2	2	1	6
N° of Passes /Shell <sup>[2]</sup>	2	2	1	1

Notes:

[1] From simulation

[2] From Table B-1

[3] From Table B-1

[4] Calculated by Aspen Capital cost Estimator

[5] Calculated by Aspen Capital cost Estimator

**Table A3.3. Inputs for Heat Exchangers SHX-604 and SHX-701 CAPEX Estimation – Base-Case**

	SHX-604	SHX-701
U (W/m <sup>2</sup> .K)	591	629
UA (W/K)	141477	429733
Heat Exchange Area (m <sup>2</sup> )	239.4	683.2
Number of Shells	2	2
TEMA Type	DFU	DFU
Tubes Material	316LS	316LS
Tubes Operation P (kPag)	24949	54949
Tubes Design P (kPag)	26196	57696
Tubes Operation T (°C)	147	103
Tubes Design T (°C)	177	133
Tubes Ext. Diameter (mm)	19.05	19.05
Shell Material	A516	A516
Shell Operation P (kPag)	438	438
Shell Design P (kPag)	608	608
Shell Operation T (°C)	55	55
Shell Design T (°C)	125	125
Tubes Length (m)	0	0
Tubes BWG	-	-
Pitch (mm)	-	-
Tubes Pattern	-	-
Shell Diameter (mm)	-	-
Channel Material	316L	316L
Head Material	316L	316L
Clad Material	-	-
Clad Side	-	-
N° Tube Passes/Shell	2	2
N° of Passes /Shell	2	2

**Table A3.4. Inputs for Plate Heat Exchanger SHX-801 CAPEX Estimation – Base-Case**

	SHX-801
Number of Heat Exchangers	3 + 1 spare
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	2415.5
Number of Plates <sup>[1]</sup>	669
Operation P (kPag)	792.85
Design P (kPag)	963
Operation T (°C)	53.6
Design T (°C)	125
Material	titanium

[1] Heat exchange area and number of plates from Aspen Exchanger Design and Rating.

**Table A3.5. Inputs for CW System Pumps CAPEX Estimation – Base-Case**

	P1	P2
Model	DCP API 610	DCP API 610
Pumps in Operation	1x100%	2x50%
Spares	1	1
Material	Carbon Steel	Steel/Copper
Flow (L/s) <sup>[1]</sup>	647.7	564.4
Head (m) <sup>[1]</sup>	36.6	75.4

[1] From Simulation

**Table A3.6. Inputs for Gas Turbines CAPEX Estimation – Base-Case**

Gas Turbines in Operation <sup>[1]</sup>	3
Spares <sup>[1]</sup>	1
Design Power (kW) <sup>[1]</sup>	28491
Power Factor	0.9
Power (kVA) <sup>[2]</sup>	31657

[1] From EIA(Petrobras, 2013); [2] Power (kVA) = Power (kW) / Power Factor

**Table A3.7 – Inputs for Compressors CAPEX Estimation – DSW-Case**

	C-101	C-201	C-202	C-501	C-502	C-601	C-602	C-603	C-701
Compressor Material	CS	CS	CS	CS	CS	CS	CS	CS	CS
Inlet Flow (m <sup>3</sup> /h)	10428	1239	1426	2894	835	12888	2943	581	313
Inlet Pressure (kPag)	1699	149	566	4399	10491	299	1473	6165	24899
Inlet Temperature (°C)	10.7	55.0	19.3	28.9	12.0	28.9	12.0	12.0	28.7
Discharge Pressure (kPag)	5149	591	1749	10541	24949	1498	6190	24949	54949
Gas Mol. Mass	24.78	39.83	33.41	22.56	22.56	31.84	31.84	31.84	25.21
Cp/Cv	1.353	1.158	1.230	1.486	2.474	1.309	1.392	2.036	n/a
Z <sub>inlet</sub>	0.928	0.984	0.953	0.855	0.616	0.988	0.940	0.739	n/a
Z <sub>outlet</sub>	0.929	0.974	0.937	0.892	0.873	0.990	0.960	0.947	n/a
Tubes Material	SS316L	SS316L	SS316L	CS	SS316L	CS	CS	SS316L	SS316L
Driver	Motor	Motor	Motor	Motor	Motor	Motor	Motor	Motor	Motor

**Table A3.8. Inputs for Heat Exchangers CAPEX Estimation – DSW-Case**

	SHX-101	SHX-102	SHX-103	SHX-201
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	883.20	89.30	439.00	49.20
Number of Shells <sup>[2]</sup>	2	2	1	1
TEMA Type <sup>[3]</sup>	AEL	BHM	NFU	NFU
Tubes Material <sup>[2]</sup>	Inconel	Inconel	Inconel	Inconel
Tubes Operation P (kPag) <sup>[1]</sup>	1749	4799	5099	591
Tubes Design P (kPag) <sup>[4]</sup>	2094	5144	5444	761
Tubes Operation T (°C) <sup>[1]</sup>	55	35	70	116
Tubes Design T (°C) <sup>[5]</sup>	125	125	125	146
Tubes Ext. Diameter (mm) <sup>[2]</sup>	25.4	19.1	19.1	19.1
Shell Material <sup>[2]</sup>	A 516	A 516	A 516	A 516
Shell Operation P (kPag) <sup>[1]</sup>	438	5149	438	438
Shell Design P (kPag) <sup>[4]</sup>	608	5494	608	608
Shell Operation T (°C) <sup>[1]</sup>	7	96	27	27
Shell DesignT (°C) <sup>[5]</sup>	22	126	125	125
Tubes Length (m) <sup>[2]</sup>	4.30	3.10	11.60	12.00
Tubes BWG <sup>[2]</sup>	18	18	18	18
Pitch (mm) <sup>[2]</sup>	31.7	25.4	25.4	25.4
Tubes Pattern <sup>[2]</sup>	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Shell Diameter (mm) <sup>[1]</sup>	1280	450	1000	350
Channel Material	A 516	A 516	A 516	A 516
Head Material	A 516	A 516	A 516	A 516
Clad Material	Inconel	-	Inconel	Inconel
Clad Side	Head	-	Head	Head
N° Tube Passes/Shell <sup>[2]</sup>	1	1	2	2
N° of Passes /Shell <sup>[2]</sup>	1	2	2	2

**Table A3.8. Inputs for Heat Exchangers CAPEX Estimation – DSW-Case (continued)**

	SHX-202	SHX-301	SHX-302	SHX-501
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	91.60	91.6	62.6	660
Number of Shells <sup>[2]</sup>	2.00	2	1	2
TEMA Type <sup>[3]</sup>	NFU	NFU	BHM	NFU
Tubes Material <sup>[2]</sup>	Inconel	Inconel	SS 316	SS 316
Tubes Operation P (kPag) <sup>[1]</sup>	1749	1749	616	4849
Tubes Design P (kPag) <sup>[4]</sup>	2094	2094	786	5194
Tubes Operation T (°C) <sup>[1]</sup>	89	89	-19	18
Tubes Design T (°C) <sup>[5]</sup>	125	125	-49	22
Tubes Ext. Diameter (mm) <sup>[2]</sup>	19.1	19.05	25.4	25.4
Shell Material <sup>[2]</sup>	A 516	A 516	A 516	A 516
Shell Operation P (kPag) <sup>[1]</sup>	438	438	4899	438
Shell Design P (kPag) <sup>[4]</sup>	608	608	5244	608
Shell Operation T (°C) <sup>[1]</sup>	27	27	20	7
Shell Design T (°C) <sup>[5]</sup>	125	125	22	22
Tubes Length (m) <sup>[2]</sup>	8.40	8.4	4.7	10.4
Tubes BWG <sup>[2]</sup>	18	18	18	18
Pitch (mm) <sup>[2]</sup>	25.4	25.4	31.7	31.7
Tubes Pattern <sup>[2]</sup>	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Shell Diameter (mm) <sup>[1]</sup>	400	400	500	1000
Channel Material	A 516	A 516	A 516	A 516
Head Material	A 516	A 516	A 516	A 516
Clad Material	Inconel	Inconel	-	-
Clad Side	Head	Head	-	-
Number of Tube Passes/Shell <sup>[2]</sup>	2,00	2	6	2
Number of Passes /Shell <sup>[2]</sup>	2,00	2	4	2

**Table A3.8. Inputs for Heat Exchangers CAPEX Estimation – DSW-Case (continued)**

	SHX-502	SHX-601	SHX-602
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	748.7	125.8	272
Number of Shells <sup>[2]</sup>	2	1	1
TEMA Type <sup>[3]</sup>	DFU	DFU	NFU
Tubes Material <sup>[2]</sup>	SS 316	SS 316	A179
Tubes Operation P (kPag) <sup>[1]</sup>	10541	24949	6190
Tubes Design P (kPag) <sup>[4]</sup>	11068	26196	6535
Tubes Operation T (°C) <sup>[1]</sup>	102	70	149
Tubes Design T (°C) <sup>[5]</sup>	132	125	179
Tubes Ext. Diameter (mm) <sup>[2]</sup>	19.05	19.05	19.05
Shell Material <sup>[2]</sup>	A 516	A 516	A 516
Shell Operation P (kPag) <sup>[1]</sup>	438	438	438
Shell Design P (kPag) <sup>[4]</sup>	608	608	608
Shell Operation T (°C) <sup>[1]</sup>	27	27	27
Shell Design T (°C) <sup>[5]</sup>	125	125	125
Tubes Length (m) <sup>[2]</sup>	9	9	12
Tubes BWG <sup>[2]</sup>	18	16	18
Pitch (mm) <sup>[2]</sup>	25.4	25.4	25.4
Tubes Pattern <sup>[2]</sup>	Triangular 30°	Triangular 30°	Triangular 30°
Shell Diameter (mm) <sup>[1]</sup>	1050	620	780
Channel Material	316L	316L	A 516
Head Material	316L	316L	A 516
Clad Material	-	-	-
Clad Side	-	-	-
Number of Tube Passes/Shell <sup>[2]</sup>	2	2	2
Number of Passes /Shell <sup>[2]</sup>	2	2	2

[1] From simulation

[2] From Table B-3

[3] From Table B-3

[4] Calculated by Aspen Capital cost Estimator

[5] Calculated by Aspen Capital cost Estimator

**Table A3.9. Inputs for Heat Exchangers SHX-603, SHX-701 CAPEX Estimation – DSW-Case**

	SHX-603	SHX-701
U (W/m <sup>2</sup> .K)	433	463
UA (W/K)	152561	140588
Heat Exchange Area (m <sup>2</sup> )	352.3	303.6
Number of Shells	1	1
TEMA Type	DFU	DFU
Tubes Material	SS316	SS316
Tubes Operation P (kPag)	24949	54949
Tubes Design P (kPag)	26196	57696
Tubes Operation T (°C)	150	90
Tubes Design T (°C)	180	125
Tubes Ext. Diameter (mm)	19.05	19.05
Shell Material	A516	A516
Shell Operation P (kPag)	438	438
Shell Design P (kPag)	608	608
Shell Operation T (°C)	27	27
Shell Design T (°C)	125	125
Tubes Length (m)	0	0
Tubes BWG	-	-
Pitch (mm)	-	-
Tubes Pattern	-	-
Shell Diameter (mm)	-	-
Channel Material	316L	316L
Head Material	316L	316L
Clad Material	-	-
Clad Side	-	-
Number Tube Passes/Shell	2	2
Number of Passes /Shell	2	2

**Table A3.10. Inputs for Plate Heat Exchanger SHX-801 CAPEX Estimation – DSW-Case**

	SHX-801
Numberof Heat Exchangers	4 + 1 spare
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	1908.2
Number of Plates <sup>[1]</sup>	687
Operation P (kPag)	792.85
Design P (kPag)	963
Operation T (°C)	23.8
Design T (°C)	125
Material	titanium

[1] Heat exchange area and number of plates from Aspen Exchanger Design and Rating.

**Table A3.11. Inputs for CW System Pumps CAPEX Estimation – DSW-Case**

	P1	P2
Model	DCP API 610	DCP API 610
Pumps in Operation	1x100%	2x50%
Spares	1	1
Material	Carbon Steel	Steel/Copper
Flow (L/s) <sup>[1]</sup>	664.7	818.5
Head (m) <sup>[1]</sup>	36.6	75.1

[1] From Simulation

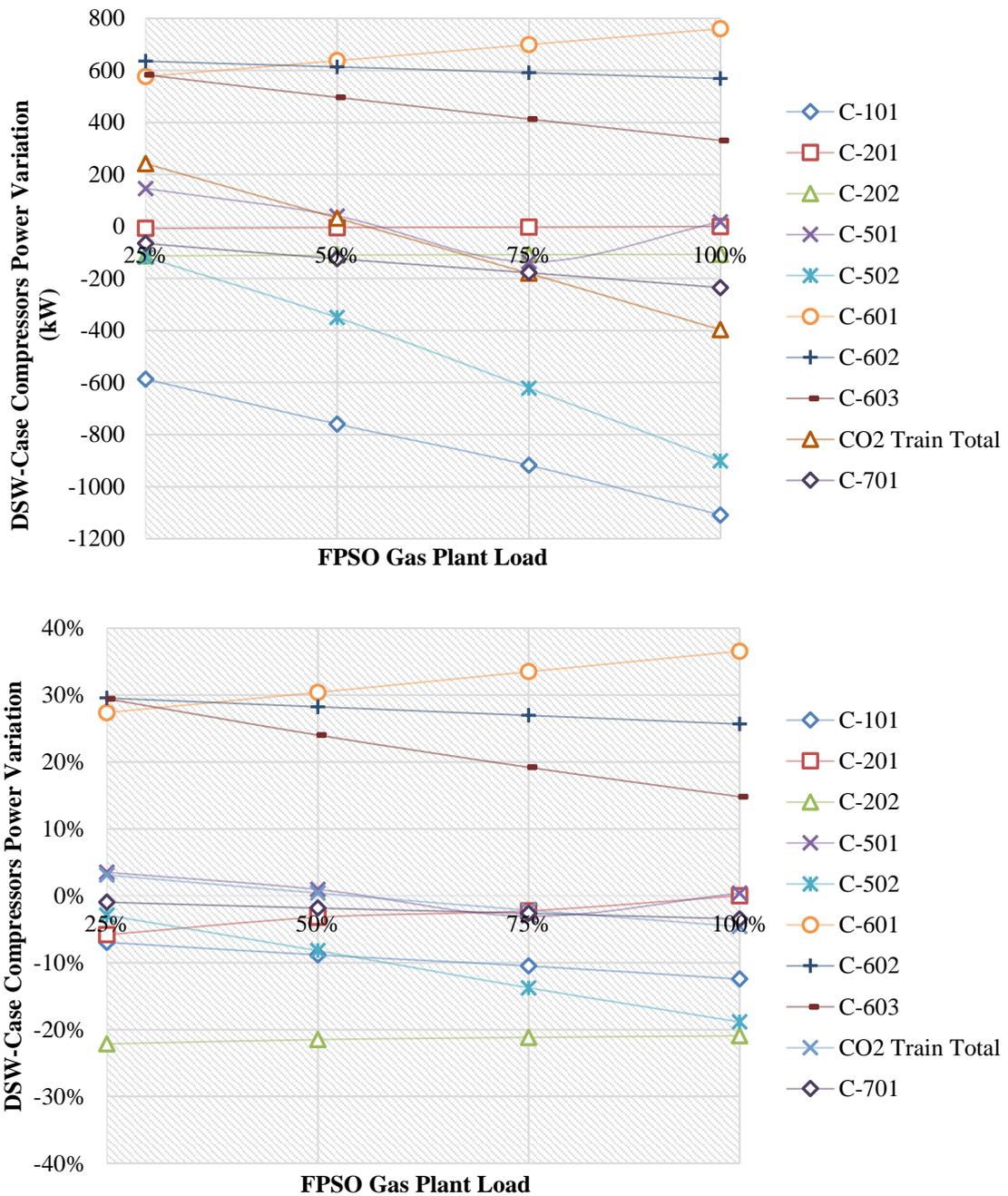
**Table A3.12. Inputs for Gas Turbines CAPEX Estimation – DSW-Case**

GTs in Operation <sup>[1]</sup>	3
Spares <sup>[1]</sup>	1
Design Power (kW) <sup>[1]</sup>	27017
Power Factor	0.9
Power (kVA) <sup>[2]</sup>	30019

[1] From EIA (Petrobras, 2013)

[2] Power (kVA) = Power (kW) / Power Factor

**Supplement A4 – Simulation, Sizing and CAPEX Results**



**Figure A4.1. DSW-CASE compressors power reduction at partial load**

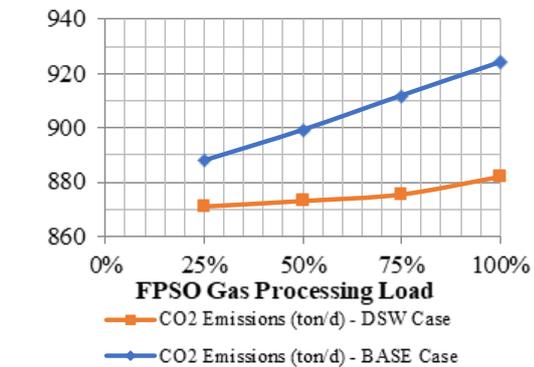
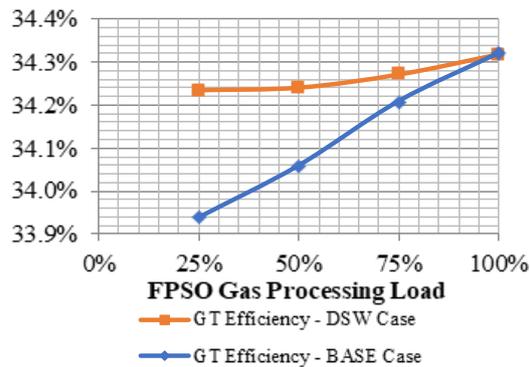
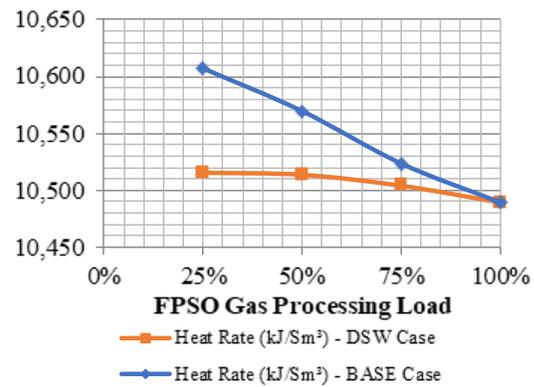
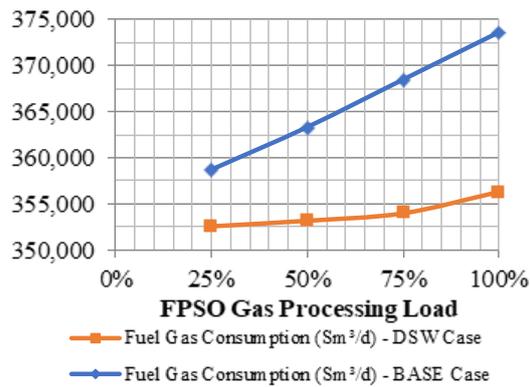
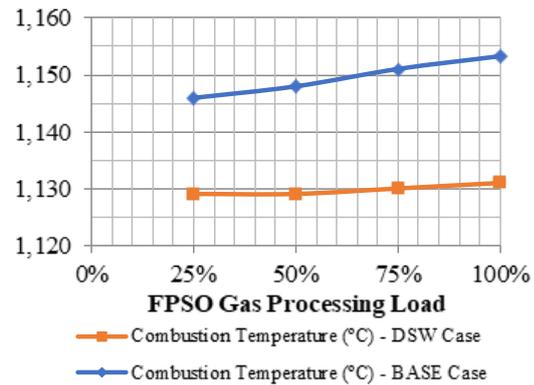
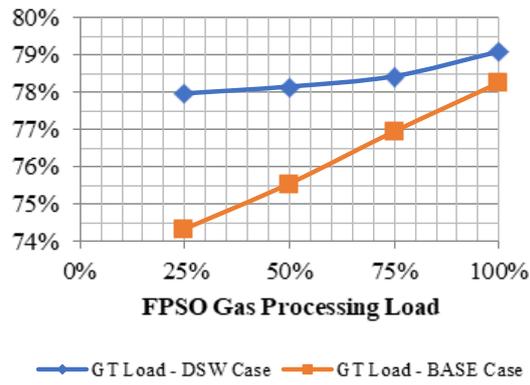


Figure A4.2. Gas turbines (GTs) performance comparison at partial load

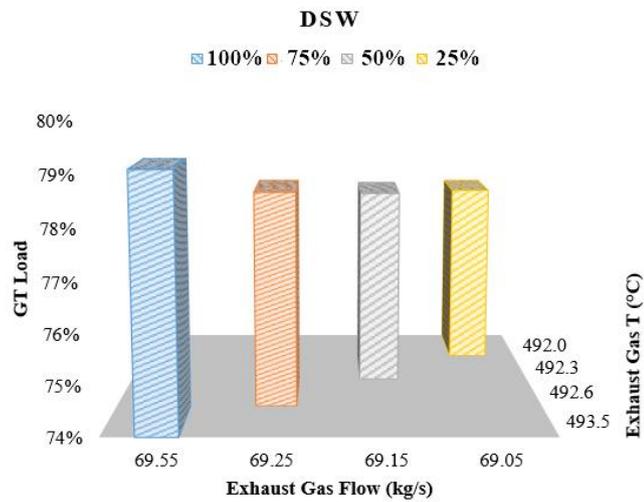
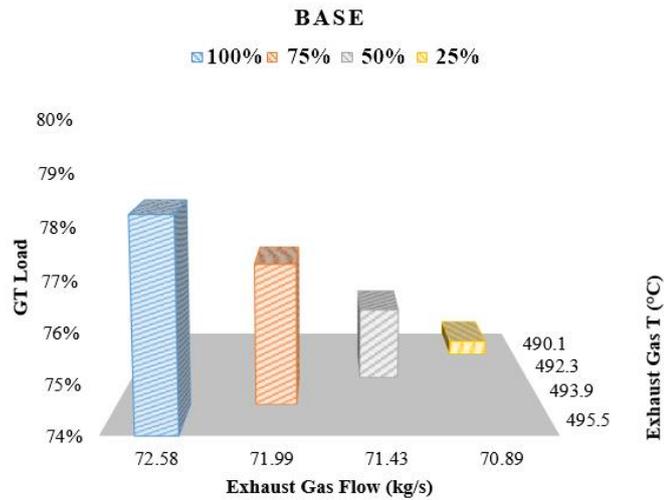


Figure A4.3. Gas turbines (GTs) load versus exhaust gas flow rate and temperature

Table A4.1. Compressors Results

CASE TAG	BASE			DSW			
	Inlet Flow (m <sup>3</sup> /h)	Inlet T (°C)	Power (kW)	Inlet Flow (m <sup>3</sup> /h)	Inlet T (°C)	Power (kW)	Power Cut
C-101	11,976	39	8,901	10,428	11	7,792	12.5%
C-201	1,239	55	137	1,239	55	137	0.0%
C-202	1,839	25	511	1,426	19	404	20.9%
C-501	2,894	29	4,596	2,894	29	4,595	0.0%
C-502	1,110	40	4,768	835	12	3,856	19.1%
C-601	12,939	30	2,080	12,888	29	2,879	-38.4%
C-602	4,728	40	2,234	2,943	12	2,866	-28.3%
C-603	1,583	40	2,301	581	12	2,718	-18.1%
C-604	466	40	2,196	-	-	-	-
C-701	313	37	7,672	313	29	7,385	3.7%
C-901	1,885	0	455	-	-	-	-
<b>TOTAL</b>			<b>35,850</b>			<b>32,633</b>	<b>9.0%</b>

**Table A4.2. Heat Exchangers Results and Comparison**

CASE		Q (MJ/h)			LMTD			A <sub>o</sub> (m <sup>2</sup> )		
BASE	DSW	BASE	DSW	ΔQ	BASE	DSW	ΔLMTD	BASE	DSW	ΔA <sub>o</sub>
SHX-101	SHX-101	7.43	20.7	179%	7.30	16.6	127%	588	883	50%
SHX-102	SHX-103	42.3	23.1	-45%	23.5	25.0	6%	1,093	439	-60%
SHX-201	SHX-201	1.02	1.53	51%	20.2	25.0	24%	41.0	49.2	20%
SHX-202	SHX-202	1.98	2.87	45%	12.7	16.2	27%	109	86.1	-21%
SHX-301	SHX-102	11.4	11.4	0%	7.50	60.5	712%	1,153	89.3	-92%
SHX-302	SHX-301	1.84	0.92	-50%	10.0	11.3	13%	85.7	62.6	-27%
SHX-303	SHX-302	4.57	4.87	6%	14.5	3.90	-73%	128	660	415%
SHX-501	SHX-501	24.1	37.7	57%	18.8	26.1	39%	506	749	48%
SHX-502	SHX-502	30.1	13.2	-56%	20.9	37.8	81%	564	126	-78%
SHX-601	SHX-601	7.08	12.4	75%	24.5	38.6	57%	254	277	9%
SHX-602	SHX-602	9.17	14.0	53%	28.8	36.9	28%	199	272	37%
SHX-603	SHX-603	12.0	20.2	69%	30.6	36.8	20%	203	352	73%
SHX-604	-	15.2	-	-	29.8	-	-	239	-	-
SHX-701	SHX-701	29.3	23.6	-19%	19.0	46.7	146%	683	304	-56%
SHX-801	SHX-801	131	168	28%	10.4	6.50	-37%	6,649	8,232	24%
SHX-901	-	0.20	-	-	43.5	-	-	8.60	-	-
SHX-902	-	6.22	-	-	10.0	-	-	288	-	-
TOTAL		335	355	6%				12,791	12,581	1.6%

**Table A4.3. GTs Simulation Validation against Thermoflex Results**

Gas Turbines Parameters (air at 23°C and 87% RH)	Base-Case			DSW-Case		
	HYSYS	Thermoflex	Error	HYSYS	Thermoflex	Error
Compressor Efficiency	84.2%	N/A	-	85.7%	-	-
Turbine Efficiency	88.0%	N/A	-	87.5%	-	-
Net Power (kW) (generator eff. = 97%)	25754	25780	-0.10%	24300	24337	-0.15%
Gas Inlet Flow (kg/s)	1.650	1.648	0.09%	1.571	1.571	0.03%
Air Inlet Flow(kg/s)	80.85	80.85	0.00%	77.06	77.06	0.00%
Fuel LHV (kJ/Sm <sup>3</sup> ) (from simulation)	40737	40737	0.00%	40751	40750	0.00%
Heat Rate (kJ/kWh) <sup>[1]</sup>	10017	9998	0.19%	10114	10096	0.18%
Efficiency <sup>[2]</sup>	35.9%	36.0%	0.19%	35.6%	35.7%	0.18%
Exhaust Gas Temperature (°C)	519.5	519.5	0.00%	521.9	521.9	0.00%
Fire Temperature (°C)	1196	N/A	-	1179	-	-
Exhaust Gas Flow (kg/s)	81.67	81.61	0.07%	77.85	77.83	0.02%

[1] Heat Rate = (Fuel Flow\*Fuel LHV\*3600) / Net Power

[2] Efficiency = 3600 / Heat Rate

**Table A4.4. CW, SW and DSW Pumps: Summary of Design Results**

Pump	CW P-801 (1x 100%)		$\Delta$ (%)
	BASE-CASE	DSW-CASE	
Flow (L/s)	647.7	664.7	2.6
Head (m)	36.6	36.4	-0.5
Power (kW)	307.5	315.6	2.6
Pump	SW/DSW P-802 A/B (2 x 50%)		$\Delta$ (%)
	BASE-CASE	DSW-CASE	
Flow (L/s)	564.4	818.5	31.0%
Head (m)	75.4	75.1	-0.4%
Power (kW)	553.5	802.7	31.0%
NPSH (kPa)	45	32	-28.0%

**Table A4.5. Compressors: Summary of Weight Results**

Compressor	Weight (t)			
	BASE-CASE	DSW-CASE	$\Delta$	$\Delta$ (%)
C-101	74	62.6	-11.4	-15%
C-201	9.4	9.4	0.0	0%
C-202	14.4	13.2	-1.2	-8%
C-501	47.8	47.8	0.0	0%
C-502	43.4	38.6	-4.8	-11%
C-601	34.6	45.2	10.6	31%
C-602	29.2	35.0	5.8	20%
C-603	33	33.0	0.0	0%
C-604	28.6	-	-28.6	-
C-701	20.8	20.8	0.0	0%
C-901	13.8	-	-13.8	-
<b>TOTAL</b>	<b>349</b>	<b>305.6</b>	<b>-43.4</b>	<b>-12%</b>

**Table A4.6. Heat Exchangers: Summary of Weight Results**

Case		Weight (t)			
BASE	DSW	BASE-CASE	DSW-CASE	$\Delta$	$\Delta$ (%)
SHX-101	SHX-101	34.8	60.4	25.6	74%
SHX-102	SHX-103	57.2	8.8	-48.4	-64%
SHX-201	SHX-201	2.4	20.4	18.0	33%
SHX-202	SHX-202	6.0	3.2	-2.8	-7%
SHX-301	SHX-102	120	5.6	-114.4	-93%
SHX-302	SHX-301	5.8	6.6	0.8	14%
SHX-303	SHX-302	8.2	34.0	25.8	315%
SHX-501	SHX-501	30.8	47.6	16.8	55%
SHX-502	SHX-502	48.8	13.2	-35.6	-73%
SHX-601	SHX-601	11.0	12.0	1.0	9%
SHX-602	SHX-602	9.4	13.2	3.8	40%
SHX-603	SHX-603	14.0	33.0	19.0	136%
SHX-604	-	23.6	-	-23.6	-
SHX-701	SHX-701	57.6	26.8	-30.8	-53%
SHX-801	SHX-801	86.0	102.6	16.6	14%
SHX-901	-	1.04	-	-1.0	-
SHX-902	-	12.8	-	-12.8	-
TOTAL		529,4	387.4	-142.0	-27%

**Table A4.7. CW System: Summary of Weight Results**

Pumps	Weight (t)		$\Delta$	$\Delta$ (%)
	BASE-CASE	DSW-CASE		
P-801	7.60	7.80	0.20	2.6%
P-802 A/B	10.40	13.60	3.20	30.8%
TOTAL	18.00	21.40	3.40	18.9%

**Table A4.8. Compressors CAPEX Comparison**

Compressor	CAPEX US\$ <sup>[1]</sup>		$\Delta$ (%)
	BASE-CASE	DSW-CASE	
C-101	6,741,600	5,650,400	-16.2%
C-201	1,839,700	1,839,700	0.0%
C-202	2,267,600	2,187,900	-3.5%
C-501	4,539,800	4,539,800	0.0%
C-502	4,970,100	4,608,200	-7.3%
C-601	3,458,700	4,728,600	36.7%
C-602	2,804,500	3,674,100	31.0%
C-603	3,625,000	3,964,700	9.4%
C-604	3,677,200	-	-
C-701	3,155,200	3,155,500	0.0%
C-901	2,077,400	-	-
TOTAL	39,156,800	34,348,900	-12.3%

[1] For two pieces of equipment (one spare)

**Table A4.9. Heat Exchangers CAPEX Comparison**

Heat Exchanger		CAPEX US\$ <sup>[1]</sup>		$\Delta$ (%)
		BASE-CASE	DSW-CASE	
SHX-101	SHX-101	2,568,200	3,526,400	37.3%
SHX-102	SHX-103	4,503,900	2,259,000	-49.8%
SHX-201	SHX-201	404,500	428,700	6.0%
SHX-202	SHX-202	832,400	859,800	3.3%
SHX-301	SHX-102	4,945,300	946,400	-80.9%
SHX-302	SHX-301	504,800	380,100	-24.7%
SHX-303	SHX-302	570,900	1,468,500	157.2%
SHX-501	SHX-501	2,138,300	3,047,000	42.5%
SHX-502	SHX-502	4,199,700	1,796,000	-57.2%
SHX-601	SHX-601	307,700	327,400	6.4%
SHX-602	SHX-602	255,700	384,000	50.2%
SHX-603	SHX-603	1,247,800	2,860,700	129.3%
SHX-604	-	2,799,100	-	-
SHX-701	SHX-701	4,549,500	2,694,100	-40.8%
SHX-801	SHX-801	6,594,500	8,213,300	24.5%
SHX-901	-	123,800	-	-
SHX-902	-	336,000	-	-
TOTAL		36,882,100	29,191,400	-20.9%

[1] For two pieces of equipment (one spare)

**Table A4.10. CW Pumps CAPEX Comparison**

Pumps	CAPEX US\$ <sup>[1]</sup>		$\Delta$ (%)
	BASE-CASE	DSW-CASE	
P-801	740,700	759,500	2.5%
P-802	1,228,500	1,633,900	24.8%
TOTAL	\$1,969,200	\$2,393,400	17.7%

[1] For two pieces of equipment (one spare)

## References of Supplement A

Boyce MP. GasTurbine Engineering Handbook. 2<sup>nd</sup> Ed. Houston: Gulf Professional Publishing; 2002.

TEMA. Standards of the Tubular Exchanger Manufacturers Association. 9th ed. Tarrytown, NY: Tubular Exchanger Manufacturers Association, INC.; 2007.

## **SUPPLEMENTARY MATERIALS B**

*Title: Exergy, Energy and Emissions Analysis of Compressors Schemes in Offshore Rigs: CO<sub>2</sub>-Rich Natural Gas Processing*

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*Manuscript Code: JNGSE-D-20-00340*

**Supplement B1. Heat Exchangers**

**Supplement B2. Process Simulation**

**Supplement B3. Exergy Analysis**

**References of Supplementary Materials B**

## Supplement B1 – Heat Exchangers

**Table B1.1. Heat Exchangers Design and Materials**

	SHX-101	SHX-202	SHX-204	SHX-501	SHX-502	SHX-601	SHX-602	SHX-603	SHX-604	SHX-701	SHX-901
Heat Exchange Area (m <sup>2</sup> ) <sup>[1]</sup>	1093	41.0	108.5	505,5	563,6	254.4	198.7	203.3	239.4	683.2	266.1
Number of Shells <sup>[1]</sup>	1	1	1	2	2	1	1	2	2	2	1
TEMA Type <sup>[1]</sup>	NFU	NFU	NFU	DFU	NFU	NFU	NFU	DFU	DFU	DFU	AJL
Tubes Material <sup>[1]</sup>	Inconel	Inconel	Inconel	316LS	316LS	A179	A179	316LS	316LS	316LS	A179
Tubes Design P (kPag) <sup>[1]</sup>	5494	761	2094	11068	20500 <sup>[2]</sup>	1204.05	3395	9213	20500 <sup>[3]</sup>	20500 <sup>[4]</sup>	608
Tubes Ext. Diameter (mm) <sup>[1]</sup>	25.4	19.05	19.05	19.05	19.05	19.05	19.05	19.05	19.05	19.05	19.05
Shell Material <sup>[1]</sup>	A 516	A 516	A 516	A 516	A516	A516	A 516				
Shell Design P (kPag) <sup>[1]</sup>	608	608	608	608	608	608.0	608	608	608	608	1789
Tubes Pattern <sup>[1]</sup>	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°	Triangular 30°
Shell Diameter (mm) <sup>[1]</sup>	1800	350	500	800	800	900	700	550	-	-	850
Channel Material	316L	316L	316L	316L	316L	A 516	A 516	316L	316L	316L	A 516
Number of Tube Passes/Shell <sup>[1]</sup>	4	2	2	2	2	2	2	2	2	2	6
Number of Passes /Shell <sup>[1]</sup>	2	2	2	2	2	2	2	2	2	2	1

Notes:

[1] From Cruz et al. (2018)

[2] Pressure limited to 20500 kPa by ACCEv10. The real design pressure of the HX-502 is 26196 kPag

[3] Pressure limited to 20500 kPa by ACCEv10. The real design pressure of the HX-604 is 26196 kPag

[4] Pressure limited to 20500 kPa by ACCEv10. The real design pressure of the HX-701 is 57696 kPag

**Table B1.2. Heat Exchangers Area Estimation for Multiple Smaller Compressors in Parallel Case**

<b>SSLC</b>	<b>HX-102</b>	<b>HX-202</b>	<b>HX-204</b>	<b>HX-501</b>	<b>HX-502</b>	<b>HX-601</b>	<b>HX-602</b>	<b>HX-603</b>	<b>HX-604</b>	<b>HX-701</b>	<b>HX-901</b>	
Q (MJ/h)	30569	645	2405	25301	28914	7358	8997	10937	12892	25797	5679	
UA (MJ/h.°C)	1846	35	165	1273	1414	295	320	383	489	1530	314	
LMTD (°C)	16.6	18.4	14.6	19.9	20.4	25.0	28.1	28.5	26.3	16.9	18.1	
Min. Approach (°C)	5	5	5	5	5	5	5	5	5	5	10	
<b>MSCP</b>	<b>HX-102</b>	<b>HX-202</b>	<b>HX-204</b>	<b>HX-501</b>	<b>HX-502</b>	<b>HX-601</b>	<b>HX-602</b>	<b>HX-603</b>	<b>HX-604</b>	<b>HX-701 exp</b>	<b>HX-702 inj</b>	<b>HX-901</b>
Q (MJ/h)	15089	202	720	8748	9918	5862	7496	9594	11921	7103	14853	2788
UA (MJ/h.°C)	872	12	47	411	457	237	260	312	397	356	794	168
LMTD (°C)	17.3	16.5	15.4	21.3	21.7	24.7	28.9	30.7	30.0	19.9	18.7	16.6
Min. Approach (°C)	5	5	5	5	5	5	5	5	5	5	5	10
<b>DIFFERENCE (SSLC-MSCP)</b>												
Q <sub>reduction</sub> (MJ/h)	15480	443	1685	16554	18996	1496	1500	1343	970	18694	10944	2891
Q <sub>reduction</sub> (%)	50.6%	68.6%	70.1%	65.4%	65.7%	20.3%	16.7%	12.3%	7.5%	72.5%	42.4%	50.9%
LMTD (°C)	-0.75	1.90	-0.75	-1.38	-1.24	0.25	-0.76	-2.20	-3.66	-3.08	-1.85	1.45
LMTD (%)	-4.5%	10.3%	-5.2%	-7.0%	-6.1%	1.0%	-2.7%	-7.7%	-13.9%	-18.3%	-11.0%	8.0%
Q/LMTD (MJ/h.°C)	974	23	118	861	957	58	60	71	92	1174	736	146
Q/LMTD (%)	52.8%	65.0%	71.5%	67.7%	67.7%	19.5%	18.9%	18.6%	18.8%	76.7%	48.1%	46.6%
<b>AREA</b>												
Design Area of SSLC (m <sup>2</sup> )*	1093	41	108.5	505.5	563.6	254.4	198.7	203.3	239.4	683.2		8.64
Design Area of MSCP (m <sup>2</sup> )	516.2	14.3	30.9	163.4	182.3	204.7	161.2	165.6	194.4	159.0	354.4	4.8
Exchangers in Parallel MSCP	2	3	3	3	3	1	1	1	1	1	2	2
Total Required Area MSCP (m <sup>2</sup> )	1032	43	93	490	547	205	161	166	194	868		9
Spares (SSLC AND MSCP)	1	1	1	1	1	1	1	1	1	1	1	1
N° of identical exchangers SSLC	2	2	2	2	2	2	2	2	2	2	2	2
N° of identical exchangers MSCP	3	4	4	4	4	2	2	2	2	2	3	3

\*from Cruz et al. (2018)

## Supplement B2 – Process Simulation

### B2.1. SSLC-Case

**Table B2.1.1. Compressors and Pumps Simulation Power - SSLC**

<b>COMPRESSORS AND PUMPS</b>	
C-101 Std Gas Flow (Sm <sup>3</sup> /d)	5000318
C-201 Std Gas Flow (Sm <sup>3</sup> /d)	70793
C-202 Std Gas Flow (Sm <sup>3</sup> /d)	341542
C-501 Std Gas Flow (Sm <sup>3</sup> /d)	3499879
C-502 Std Gas Flow (Sm <sup>3</sup> /d)	3499879
C-601 Std Gas Flow (Sm <sup>3</sup> /d)	1200505
C-602 Std Gas Flow (Sm <sup>3</sup> /d)	1200505
C-603 Std Gas Flow (Sm <sup>3</sup> /d)	1200505
C-604 Std Gas Flow (Sm <sup>3</sup> /d)	1200505
C-701 Std Gas Flow (Sm <sup>3</sup> /d)	4229609
C-901 Std Gas Flow (Sm <sup>3</sup> /d)	226185
C-101 Power (kW)	8223
C-201 Power (kW)	139
C-202 Power (kW)	605
C-501 Power (kW)	4635
C-502 Power (kW)	4577
C-601 Power (kW)	2123
C-602 Power (kW)	2241
C-603 Power (kW)	2238
C-604 Power (kW)	2032
C-701 Power (kW)	6744
C-901 Power (kW)	494
C-200 Total power (kW)	744
C-500 Total power (kW)	9212
C-600 Total power (kW)	8634
Compressors Total power (kW)	34050
Pumps power (kW)	481
Other users power (kW)	23164
Total drivers power (kW)	34531
Total power demand (kW)	57695
Power per GT (kW)	19232

**Table B2.1.2. Gas Turbines Simulation: SSLC-Case**

<b>1000 - GAS TURBINES</b>	
NG Mass Flow/GT (kg/s)	1.29
NG Mass Flow/GT (kg/h)	4652
NG Std Gas Flow (Sm <sup>3</sup> /d)	118533
NG Vapour Fraction	1
NG Higher Heating Value (kJ/kg)	43500
NG Lower Heating Value (kJ/kg)	47945
NG Pressure (kPa)	3500
NG Temperature (°C)	27.5
NG Molecular Weight	22.27
AIR Mass Flow (kg/s)	65.57
AIR Mass Flow (kg/h)	236055
AIR Std Gas Flow (Sm <sup>3</sup> /d)	4667006
AIR INLET Pressure (kPa)	101.3
AIR INLET Temperature (°C)	23.0
C-1001 Power (kW)	34319
T-1001 Power (kW)	54150
NET Power (kW)	19831
Net Electricity (kW -97% gen. eff.)	19237
Heat Rate (BTU/kWh)	11246
LHV Efficiency (%)	32.01
Exhaust Gas Mass Flow (kg/s)	66.19
Exhaust Gas Mass Flow (kg/h)	238299
Exhaust Gas Temperature (°C)	518.5
Fire Temperature (°C)	1177.5
Exhaust CO <sub>2</sub> Comp (mole %)	3.3%
Exhaust CO <sub>2</sub> Emissions (kg/h)	12275
Total NG Consumption (kg/h - 3 TGs)	13955
Total NG Consumption (Sm <sup>3</sup> /h - 3 TGs)	355598
Total CO <sub>2</sub> Emissions (kg/h - 3 TGs)	36825
GT used capacity (%)	74.62%
Power demand/GT (kW)	19232
Check Power (generated - demand) (kW)	4.8
Error ((generated - demand)/demand) (kW)	0.025%
Loss due to Gen. Efficiency (3%) (kW)	594.9

**Table B2.1.3. Propane Refrigeration Cycle Simulation: SSLC-Case**

<b>900 - C3 REFRIGERATION CYCLE</b>	
Liquid T sat (°C)	-7
Liquid P sat (kPa)	380
Vapor T sat (°C)	50
Vapor P sat (kPa)	1720
Stream 901 T (°C)	-6.0
Stream 905 T (°C)	45
Superheating (°C)	1.0
Subcooling (°C)	5
HX-303 ΔH (kJ/kg)	244.2
C-901 ΔH (kJ/kg)	101.1
Pot / ton refriger.	1.45
COP	2.42
Stream 901 Mass Flow (kg/h)	15977
Stream 902 std gas flow (Sm <sup>3</sup> /d)	226185
Stream 902 mass flow (kg/h)	17576
NG Inlet T (°C)	10
NG Outlet T (°C)	3
HX-303 Duty (kJ/h)	3902082
HX-303 BTU/h	3698393
T.R.	308
Condenser CW flow (kg/h)	131744
CW inlet T (°C)	35
CW outlet T (°C)	45
Condenser Duty (kJ/h)	5679369
Outlet Gas C6+ content (ppmv)	483

**Table B2.1.4. Cooling-Water System Simulation: SSLC-Case**

<b>800 - COOLING WATER SYSTEM</b>	
HX-003 Mass Flow (kg/h)	228442
HX-102 Mass Flow (kg/h)	354331
HX-201 Mass Flow (kg/h)	1843
HX-202 Mass Flow (kg/h)	7474
HX-203 Mass Flow (kg/h)	19748
HX-204 Mass Flow (kg/h)	27880
HX-501 Mass Flow (kg/h)	293281
HX-502 Mass Flow (kg/h)	335161
HX-601 Mass Flow (kg/h)	85288
HX-602 Mass Flow (kg/h)	104282
HX-603 Mass Flow (kg/h)	126772
HX-604 Mass Flow (kg/h)	149429
HX-701 Mass Flow (kg/h)	299029
HX-901 Mass Flow (kg/h)	131744
TOTAL CW INLET (kg/h)	2164705
CW OUTLET STREAM T (°C)	35.0
SW OUTLET STREAM T (°C)	40.0
CW INLET STREAM T (°C)	52.7
SW INLET STREAM T (°C)	23.0

## B2.2. MPSC-Case

**Table B2.2.1. Compressors and Pumps Simulation Power: MPSC-Case**

<b>COMPRESSORS AND PUMPS</b>		
	Flow (Sm <sup>3</sup> /d)	Power (kW)
C-101 A	2364552	4059
C-101 B	2364552	4059
C-201 A	20109	43
C-201 B	20109	43
C-201 C	20109	43
C-202 A	97030	180
C-202 B	97030	180
C-202 C	97030	180
C-501 A	1124749	1619
C-501 B	1124749	1619
C-501 C	1124749	1619
C-502 A	1124749	1584
C-502 B	1124749	1584
C-502 C	1124749	1584
C-601	968160	1707
C-602	968160	1824
C-603	968160	1864
C-604	968160	1762
C-701 A	968160	1552
C-901 A	103459	229
C-901 B	103459	229
C-100 Total	4729104	8118
C-200 Total	291090	669
C-500 Total	3374248	9607
C-600 Total	968160	7158
C-900 Total	206919	459
Compressors Total power (kW)	27562	
Pumps power (kW)	382	
Other users power (kW)	23164	
Total drivers power (kW)	27944	
Total power demand (kW)	51108	
Power per GT (kW)	25554	

**Table B2.2.2. Gas Turbines Simulation: MPSC-Case**

<b>1000 - GAS TURBINES</b>	
NG Mass Flow/GT (kg/s)	1.63
NG Mass Flow/GT (kg/h)	5873
NG Std Gas Flow (Sm <sup>3</sup> /d)	149637
NG Vapour Fraction	1
NG Higher Heating Value (kJ/kg)	43500
NG Lower Heating Value (kJ/kg)	47945
NG Pressure (kPa)	3500
NG Temperature (°C)	27.5
NG Molecular Weight	22.27
AIR Mass Flow (kg/s)	80.10
AIR Mass Flow (kg/h)	288371
AIR Std Gas Flow (Sm <sup>3</sup> /d)	5701330
AIR INLET Pressure (kPa)	101.3
AIR INLET Temperature (°C)	23.0
C-1001 Power (kW)	41338
T-1001 Power (kW)	67682
NET Power (kW)	26344
Net Electricity (kW -97% gen. eff.)	25554
Heat Rate (BTU/kWh)	10688
LHV Efficiency (%)	33.68
Exhaust Gas Mass Flow (kg/s)	80.92
Exhaust Gas Mass Flow (kg/h)	291300
Exhaust Gas Temperature (°C)	519.5
Fire Temperature (°C)	1191.8
Exhaust CO <sub>2</sub> Comp (mole %)	3.4%
Exhaust CO <sub>2</sub> Emissions (kg/h)	15491
Total NG Consumption (kg/h - 3 TGs)	11745
Total NG Consumption (Sm <sup>3</sup> /h - 3 TGs)	299274
Total CO <sub>2</sub> Emissions (kg/h - 3 TGs)	30982
GT used capacity (%)	99.1%
Power demand/GT (kW)	25554
Check Power (generated - demand) (kW)	0.3
Error ((generated - demand)/demand) (kW)	0.001%
Loss due to Gen. Efficiency (3%) (kW)	790.3

**Table B2.2.3. Propane Refrigeration Cycle Simulation: MPSC-Case**

<b>900 - C3 REFRIGERATION CYCLE</b>	
Liquid T sat (°C)	-7
Liquid P sat (kPa)	380
Vapor T sat (°C)	50
Vapor P sat (kPa)	1720
Stream 901 T (°C)	-6.1
Stream 905 T (°C)	45
Superheating (°C)	0.9
Subcooling (°C)	5
HX-303 ΔH (kJ/kg)	244.1
C-901 ΔH (kJ/kg)	102.7
Pot / ton refrig.	1.47
COP	2.38
Stream 901 Mass Flow (kg/h)	16079
Stream 902 std gas flow (Sm <sup>3</sup> /d)	206919
Stream 902 mass flow (kg/h)	16079
NG Inlet T (°C)	10
NG Outlet T (°C)	3
HX-303 Duty (kJ/h)	3925034
HX-303 BTU/h	3720148
T.R.	310
Condenser CW flow (kg/h)	64681
CW inlet T (°C)	35
CW outlet T (°C)	45
Condenser Duty (kJ/h)	2788247
Outlet Gas C6+ content (ppmv)	482

**Table B2.2.4. Cooling-Water System Simulation: MPSC-Case**

<b>800 - COOLING WATER SYSTEM</b>	
HX-003 Mass Flow (kg/h)	228442
HX-102 A/B Mass Flow (kg/h)	354331
HX-201 Mass Flow (kg/h)	1843
HX-202 Mass A/B/C Flow (kg/h)	7474
HX-203 Mass Flow (kg/h)	19748
HX-204 A/B/C Mass Flow (kg/h)	27880
HX-501 A/B/C Mass Flow (kg/h)	293281
HX-502 A/B/C Mass Flow (kg/h)	335161
HX-601 Mass Flow (kg/h)	85288
HX-602 Mass Flow (kg/h)	104282
HX-603 Mass Flow (kg/h)	126772
HX-604 Mass Flow (kg/h)	149429
HX-701 A Mass Flow (kg/h)	299029
HX-901 A/B Mass Flow (kg/h)	131744
TOTAL CW INLET (kg/h)	2164705
CW OUTLET STREAM T (°C)	35.0
SW OUTLET STREAM T (°C)	40.0
CW INLET STREAM T (°C)	52.7
SW INLET STREAM T (°C)	23.0

## Supplement B3 – Exergy Analysis

### B3.1. Determination of exergy flows of streams

- RER-1

**Table B3.1.1 Streams exergy flow- Single-shaft compressors – 100% FPSO gas -load – RER-1**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>(kW)</i>	$H - T_0 S + P_0 V - \sum N_k \cdot \mu_k^0$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt; 0</i>
102	S	IN	Overall Gas Plant	-123689	163.76	2.173	-2281951.7	1907850.8	-105369	OK
201	S	IN	Overall Gas Plant	-156362	183.40	0.030	-51158.7	44918.0	-1609	OK
208	S	IN	Overall Gas Plant	-176963	176.71	0.098	-173097.0	150557.8	-5144	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.463	-1646213.1	151.2	-91031	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.115	-34726.0	3.2	-1920	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.430	-129531.9	11.9	-7163	OK
813	S	IN	Overall Gas Plant	-284622	56.26	4.522	-1362576.3	125.2	-75347	OK
815	S	IN	Overall Gas Plant	-284622	56.26	5.168	-1557150.0	143.0	-86106	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.315	-396246.1	36.4	-21911	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.608	-484492.7	44.5	-26791	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.955	-588977.4	54.1	-32569	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.304	-694244.5	63.8	-38390	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.611	-1389279.6	127.6	-76823	OK
827	S	IN	Overall Gas Plant	-284622	56.26	2.031	-612079.3	56.2	-33846	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.463	-1646213.1	724.9	98948	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.115	-34726.0	15.4	2087	OK

810	S	OUT	Overall Gas Plant	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.430	-129531.9	57.0	7786	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	4.522	-1362576.3	600.0	81900	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	5.168	-1557150.0	685.7	93595	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.315	-396246.1	174.5	23817	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.608	-484492.7	213.4	29121	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.955	-588977.4	259.4	35401	OK
824	S	OUT	Overall Gas Plant	-283068	61.15	2.304	-694244.5	305.7	41729	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	4.611	-1389279.6	611.8	83505	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	2.031	-612079.3	139.7	35340	OK
510	S	OUT	Overall Gas Plant	-99659	134.20	1.624	-1891287.2	1664873.1	64552	OK
402	S	OUT	Overall Gas Plant	-96058	156.34	0.174	-202695.8	177915.9	8059	OK
705	S	OUT	Overall Gas Plant	-245894	118.13	0.474	-332775.6	199653.4	16581	OK
1101	S	OUT	Overall Gas Plant	-285888	52.58	0.000	-67.6	0.0	3	OK
1102	S	OUT	Overall Gas Plant	-284367	57.17	0.000	-50.2	0.0	3	OK
1103	S	OUT	Overall Gas Plant	-180135	87.06	0.025	-78540.9	73445.3	638	OK
1104	S	OUT	Overall Gas Plant	-286955	49.90	0.003	-783.2	0.6	38	OK
1105	S	OUT	Overall Gas Plant	-286501	49.58	0.001	-234.8	0.1	11	OK
1106	S	OUT	Overall Gas Plant	-122967	80.40	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-127699	118.00	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-97645	145.67	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-239637	172.98	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-239613	164.78	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-240373	154.22	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-242403	140.59	0.000	0.0	0.0	0	OK
C-101_P	P	IN	Overall Gas Plant					8222.8	0	OK
C-201_P	P	IN	Overall Gas Plant					139.0	0	OK
C-202_P	P	IN	Overall Gas Plant					604.8	0	OK
C-501_P	P	IN	Overall Gas Plant					4635.3	0	OK
C-502_P	P	IN	Overall Gas Plant					4576.7	0	OK
C-601_P	P	IN	Overall Gas Plant					2122.9	0	OK

C-602_P	P	IN	Overall Gas Plant					2241.1	0	OK
C-603_P	P	IN	Overall Gas Plant					2237.8	0	OK
C-604_P	P	IN	Overall Gas Plant					2031.7	0	OK
C-701_P	P	IN	Overall Gas Plant					6744.0	0	OK
C-901_P	P	IN	Overall Gas Plant					493.8	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.767	-231168.6	955.8	-17590	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.767	-231168.6	796.7	17131	OK
102	S	IN	GAS+TG+CW	-123689	163.76	2.173	-2281951.7	1907850.8	-105369	OK
201	S	IN	GAS+TG+CW	-156362	183.40	0.030	-51158.7	44918.0	-1609	OK
208	S	IN	GAS+TG+CW	-176963	176.71	0.098	-173097.0	150557.8	-5144	OK
510	S	OUT	GAS+TG+CW	-99659	134.20	1.624	-1891287.2	1664873.1	64552	OK
705	S	OUT	GAS+TG+CW	-245894	118.13	0.474	-332775.6	199653.4	16581	OK
1101	S	OUT	GAS+TG+CW	-285888	52.58	0.000	-67.6	0.0	3	OK
1102	S	OUT	GAS+TG+CW	-284367	57.17	0.000	-50.2	0.0	3	OK
1103	S	OUT	GAS+TG+CW	-180135	87.06	0.025	-78540.9	73445.3	638	OK
1104	S	OUT	GAS+TG+CW	-286955	49.90	0.003	-783.2	0.6	38	OK
1105	S	OUT	GAS+TG+CW	-286501	49.58	0.001	-234.8	0.1	11	OK
1106	S	OUT	GAS+TG+CW	-122967	80.40	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-127699	118.00	0.000	0.0	0.0	0	OK
1108	S	OUT	GAS+TG+CW	-97645	145.67	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-239637	172.98	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-239613	164.78	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-240373	154.22	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-242403	140.59	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1009-A	S	OUT	GAS+TG+CW	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1009-A	S	OUT	GAS+TG+CW	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1009-A	S	OUT	GAS+TG+CW	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1010-A	S	OUT	GAS+TG+CW	-16646	196.12	0.024	-1926.1	168.2	1366	OK

1010-A	S	OUT	GAS+TG+CW	-16646	196.12	0.024	-1926.1	168.2	1366	OK
1010-A	S	OUT	GAS+TG+CW	-16646	196.12	0.024	-1926.1	168.2	1366	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	6.193	-1866058.4	7715.2	141991	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	5.515	-1661717.3	6870.4	126442	OK
1209	S	OUT	GAS+TG+CW	-277945	75.40	0.767	-231168.6	796.7	17131	OK
1303-A	S	IN	GAS+TG+CW	-285558	53.18	17.466	-5262603.5	81.4	-275068	OK
1303-B	S	IN	GAS+TG+CW	-285558	53.18	17.466	-5262603.5	81.4	-275068	OK
1306	S	OUT	GAS+TG+CW	-284236	57.51	34.931	-10525207.1	1525.4	594950	OK
835	S	OUT	GAS+TG+CW	-284620	56.27	3.522	-1061333.6	98.1	58695	OK
802	S	IN	GAS+TG+CW	-284234	57.51	3.522	-1061333.6	157.6	-59995	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					594.9	0	OK
Gen Loss	P	OUT	GAS+TG+CW					594.9	0	OK
Gen Loss	P	OUT	GAS+TG+CW					594.9	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.285	-125039.0	0.0	-111286	OK
1002-A	S	IN	1000-GT	-96058	158.13	0.058	-67565.3	59274.5	-2717	OK
1002-A	S	IN	1000-GT	-96058	158.13	0.058	-67565.3	59274.5	-2717	OK
1002-A	S	IN	1000-GT	-96058	158.13	0.058	-67565.3	59274.5	-2717	OK
1009-A	S	OUT	1000-GT	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1009-A	S	OUT	1000-GT	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1009-A	S	OUT	1000-GT	-19509	192.30	2.329	-190682.2	12623.1	132626	OK
1010-A	S	OUT	1000-GT	-16646	196.12	0.024	-1926.1	168.2	1366	OK
1010-A	S	OUT	1000-GT	-16646	196.12	0.024	-1926.1	168.2	1366	OK
1010-A	S	OUT	1000-GT	-16646	196.12	0.024	-1926.1	168.2	1366	OK
C1001A	P	IN	1000-GT					34318.9	0	OK

C1001A	P	IN	1000-GT						34318.9	0	OK
C1001A	P	IN	1000-GT						34318.9	0	OK
T1001A	P	OUT	1000-GT						54150.4	0	OK
T1001A	P	OUT	1000-GT						54150.4	0	OK
T1001A	P	OUT	1000-GT						54150.4	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4		3543.6	-90311	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4		3543.6	-90311	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4		3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4		5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4		5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4		5180.5	95341	OK
1303-A	S	IN	800-CW System	-285558	53.18	17.466	-5262603.5		81.4	-275068	OK
1303-B	S	IN	800-CW System	-285558	53.18	17.466	-5262603.5		81.4	-275068	OK
830	S	IN	800-CW System	-283242	60.61	33.378	-10057161.8		4037.5	-599089	OK
1306	S	OUT	800-CW System	-284236	57.51	34.931	-10525207.1		1525.4	594950	OK
832	S	OUT	800-CW System	-284615	56.26	33.378	-10057161.8		1145.6	556141	OK
P-802A	P	IN	800-CW System						74.3	0	OK
P-802B	P	IN	800-CW System						74.3	0	OK
P-801	P	IN	800-CW System						185.7	0	OK

**Table B3.1.2. Streams exergy flow- Single-shaft compressors – 50% FPSO gas load – RER-1**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>(kW)</i>	$H-T_0S + P_0V - \sum N_k \cdot \mu_k^0$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-146519	164.81	1.298	-1321446.9	1067969.5	-63338	OK
201	S	IN	Overall Gas Plant	-178368	183.66	0.016	-26695.9	22925.0	-881	OK
208	S	IN	Overall Gas Plant	-204760	176.13	0.046	-74560.7	62658.7	-2416	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.843	-1760425.3	161.7	-97347	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.040	-12180.4	1.1	-674	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.103	-31057.8	2.9	-1717	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.132	-39764.9	3.7	-2199	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.340	-102381.6	9.4	-5661	OK
813	S	IN	Overall Gas Plant	-284622	56.26	3.392	-1022120.6	93.9	-56520	OK
815	S	IN	Overall Gas Plant	-284622	56.26	3.648	-1099152.0	101.0	-60780	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.290	-388600.7	35.7	-21489	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.586	-477876.2	43.9	-26425	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.899	-572142.2	52.6	-31638	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.167	-652824.6	60.0	-36099	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.376	-1318543.4	121.1	-72912	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.703	-513218.8	47.1	-28380	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.843	-1760425.3	775.2	105813	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.040	-12180.4	1.8	689	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.103	-31057.8	13.8	1867	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.132	-39764.9	9.1	2296	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.340	-102381.6	45.1	6154	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	3.392	-1022120.6	450.1	61436	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	3.648	-1099152.0	484.0	66066	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.290	-388600.7	171.1	23357	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.586	-477876.2	210.4	28724	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.899	-572142.2	252.0	34390	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	2.167	-652824.6	287.5	39239	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	4.376	-1318543.4	580.6	79253	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.703	-513218.8	117.1	29632	OK
510	S	OUT	Overall Gas Plant	-100232	134.01	0.760	-899225.2	792870.0	30167	OK
402	S	OUT	Overall Gas Plant	-96657	155.92	0.167	-197235.7	173422.3	7698	OK
705	S	OUT	Overall Gas Plant	-273140	115.63	0.414	-273791.9	146509.9	14180	OK
1101	S	OUT	Overall Gas Plant	-285676	53.53	0.001	-175.4	0.0	9	OK
1102	S	OUT	Overall Gas Plant	-284732	56.03	0.000	-99.7	0.0	5	OK
1103	S	OUT	Overall Gas Plant	-180790	86.07	0.016	-51431.5	48151.2	405	OK
1104	S	OUT	Overall Gas Plant	-287076	50.34	0.002	-599.4	0.5	30	OK
1105	S	OUT	Overall Gas Plant	-286416	49.88	0.001	-158.6	0.1	8	OK
1106	S	OUT	Overall Gas Plant	-118741	94.93	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-132940	108.54	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-99269	141.93	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-266265	171.66	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-266266	163.39	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-267244	152.16	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-269677	137.26	0.000	0.0	0.0	0	OK
C-101_P	P	IN	Overall Gas Plant					8313.4	0	OK
C-201_P	P	IN	Overall Gas Plant					136.5	0	OK
C-202_P	P	IN	Overall Gas Plant					603.1	0	OK
C-501_P	P	IN	Overall Gas Plant					4196.2	0	OK
C-502_P	P	IN	Overall Gas Plant					4027.6	0	OK
C-601_P	P	IN	Overall Gas Plant					2109.2	0	OK
C-602_P	P	IN	Overall Gas Plant					2219.4	0	OK
C-603_P	P	IN	Overall Gas Plant					2159.1	0	OK
C-604_P	P	IN	Overall Gas Plant					1846.1	0	OK
C-701_P	P	IN	Overall Gas Plant					6438.8	0	OK
C-901_P	P	IN	Overall Gas Plant					574.5	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.457	-137799.6	569.7	-10485	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.457	-137799.6	474.9	10212	OK

102	S	IN	GAS+TG+CW	-146519	164.81	1.298	-1321446.9	1067969.5	-63338	OK
201	S	IN	GAS+TG+CW	-178368	183.66	0.016	-26695.9	22925.0	-881	OK
208	S	IN	GAS+TG+CW	-204760	176.13	0.046	-74560.7	62658.7	-2416	OK
510	S	OUT	GAS+TG+CW	-100232	134.01	0.760	-899225.2	792870.0	30167	OK
705	S	OUT	GAS+TG+CW	-273140	115.63	0.414	-273791.9	146509.9	14180	OK
1101	S	OUT	GAS+TG+CW	-285676	53.53	0.001	-175.4	0.0	9	OK
1102	S	OUT	GAS+TG+CW	-284732	56.03	0.000	-99.7	0.0	5	OK
1103	S	OUT	GAS+TG+CW	-180790	86.07	0.016	-51431.5	48151.2	405	OK
1104	S	OUT	GAS+TG+CW	-287076	50.34	0.002	-599.4	0.5	30	OK
1105	S	OUT	GAS+TG+CW	-286416	49.88	0.001	-158.6	0.1	8	OK
1106	S	OUT	GAS+TG+CW	-118741	94.93	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-132940	108.54	0.000	0.0	0.0	0	OK
1108	S	OUT	GAS+TG+CW	-99269	141.93	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-266265	171.66	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-266266	163.39	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-267244	152.16	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-269677	137.26	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1009-A	S	OUT	GAS+TG+CW	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1009-A	S	OUT	GAS+TG+CW	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1009-A	S	OUT	GAS+TG+CW	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1010-A	S	OUT	GAS+TG+CW	-16636	196.14	0.023	-1874.3	163.9	1330	OK
1010-A	S	OUT	GAS+TG+CW	-16636	196.14	0.023	-1874.3	163.9	1330	OK
1010-A	S	OUT	GAS+TG+CW	-16636	196.14	0.023	-1874.3	163.9	1330	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	2.623	-790324.3	3267.6	60137	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	9.395	-2830820.5	11704.0	215401	OK

1209	S	OUT	GAS+TG+CW	-277945	75.40	0.457	-137799.6	474.9	10212	OK
1303-A	S	IN	GAS+TG+CW	-285558	53.18	15.652	-4716189.7	73.5	-246509	OK
1303-B	S	IN	GAS+TG+CW	-285558	53.18	15.652	-4716189.7	73.5	-246509	OK
1306	S	OUT	GAS+TG+CW	-284236	57.51	31.304	-9432379.3	1367.2	533179	OK
835	S	OUT	GAS+TG+CW	-284620	56.27	3.650	-1099717.2	102.0	60817	OK
802	S	IN	GAS+TG+CW	-284234	57.51	3.650	-1099717.2	163.3	-62164	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					579.7	0	OK
Gen Loss	P	OUT	GAS+TG+CW					579.7	0	OK
Gen Loss	P	OUT	GAS+TG+CW					579.7	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.223	-121685.1	0.0	-108301	OK
1002-A	S	IN	1000-GT	-96657	157.69	0.056	-65745.2	57778.2	-2595	OK
1002-A	S	IN	1000-GT	-96657	157.69	0.056	-65745.2	57778.2	-2595	OK
1002-A	S	IN	1000-GT	-96657	157.69	0.056	-65745.2	57778.2	-2595	OK
1009-A	S	OUT	1000-GT	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1009-A	S	OUT	1000-GT	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1009-A	S	OUT	1000-GT	-19578	192.21	2.266	-185559.9	12203.3	128991	OK
1010-A	S	OUT	1000-GT	-16636	196.14	0.023	-1874.3	163.9	1330	OK
1010-A	S	OUT	1000-GT	-16636	196.14	0.023	-1874.3	163.9	1330	OK
1010-A	S	OUT	1000-GT	-16636	196.14	0.023	-1874.3	163.9	1330	OK
C1001A	P	IN	1000-GT					33398.4	0	OK
C1001A	P	IN	1000-GT					33398.4	0	OK
C1001A	P	IN	1000-GT					33398.4	0	OK
T1001A	P	OUT	1000-GT					52722.4	0	OK
T1001A	P	OUT	1000-GT					52722.4	0	OK
T1001A	P	OUT	1000-GT					52722.4	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK

1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1303-A	S	IN	800-CW System	-285558	53.18	15.652	-4716189.7	73.5	-246509	OK
1303-B	S	IN	800-CW System	-285558	53.18	15.652	-4716189.7	73.5	-246509	OK
830	S	IN	800-CW System	-283253	60.57	30.168	-9090005.8	3621.1	-541158	OK
1306	S	OUT	800-CW System	-284236	57.51	31.304	-9432379.3	1367.2	533179	OK
832	S	OUT	800-CW System	-284615	56.26	30.168	-9090005.8	1035.4	502658	OK
P-802A	P	IN	800-CW System					66.0	0	OK
P-802B	P	IN	800-CW System					66.0	0	OK
P-801	P	IN	800-CW System					163.7	0	OK

**Table B3.1.3. Streams exergy flow- Single-shaft compressors – 25% FPSO gas load – RER-1**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>(kW)</i>	$\frac{H-T_0S + P_0V - \sum N_k \cdot \mu_k^0}{N_k \cdot \mu_k^0}$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt; 0</i>
102	S	IN	Overall Gas Plant	-193477	164.61	0.581	-536568.9	395934.5	-28303	OK
201	S	IN	Overall Gas Plant	-225649	181.98	0.006	-7996.2	6352.3	-317	OK
208	S	IN	Overall Gas Plant	-254181	174.38	0.019	-25648.8	19752.7	-996	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.658	-1704929.6	156.6	-94278	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.006	-1714.2	0.2	-95	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.094	-28201.7	2.6	-1559	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.046	-13809.6	1.3	-764	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.338	-101853.6	9.4	-5632	OK
813	S	IN	Overall Gas Plant	-284622	56.26	2.702	-814069.2	74.8	-45016	OK
815	S	IN	Overall Gas Plant	-284622	56.26	2.584	-778458.4	71.5	-43047	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.261	-380016.5	34.9	-21014	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.523	-458840.6	42.1	-25373	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.702	-512874.3	47.1	-28360	OK
823	S	IN	Overall Gas Plant	-284622	56.26	1.749	-526925.7	48.4	-29137	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.031	-1214669.5	111.6	-67168	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.730	-521244.5	47.9	-28823	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.658	-1704929.6	750.8	102478	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.006	-1714.2	0.2	97	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.094	-28201.7	12.5	1695	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.046	-13809.6	3.2	797	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.338	-101853.6	44.9	6122	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	2.702	-814069.2	358.5	48931	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	2.584	-778458.4	342.8	46791	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.261	-380016.5	167.3	22842	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.523	-458840.6	202.1	27579	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.702	-512874.3	225.9	30827	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	1.749	-526925.7	232.0	31672	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	4.031	-1214669.5	534.9	73010	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.730	-521244.5	119.0	30096	OK
510	S	OUT	Overall Gas Plant	-100257	134.26	0.145	-170754.8	150458.6	5763	OK
402	S	OUT	Overall Gas Plant	-97821	154.85	0.158	-192265.0	169550.5	7250	OK
705	S	OUT	Overall Gas Plant	-309848	111.68	0.289	-175364.1	76147.5	9570	OK
1101	S	OUT	Overall Gas Plant	-286104	52.78	0.001	-326.1	0.1	17	OK
1102	S	OUT	Overall Gas Plant	-284945	55.42	0.000	-28.3	0.0	2	OK
1103	S	OUT	Overall Gas Plant	-192339	87.03	0.005	-18581.5	17396.6	140	OK
1104	S	OUT	Overall Gas Plant	-287549	50.63	0.001	-207.0	0.2	10	OK
1105	S	OUT	Overall Gas Plant	-286423	49.85	0.000	-80.9	0.0	4	OK
1106	S	OUT	Overall Gas Plant	-83662	201.20	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-130193	106.31	0.006	-12628.2	11685.2	184	OK
1108	S	OUT	Overall Gas Plant	-100086	139.52	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-302035	169.53	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-302152	160.91	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-303646	147.99	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-306947	130.49	0.000	0.0	0.0	0	OK
C-101_P	P	IN	Overall Gas Plant					8161.0	0	OK
C-201_P	P	IN	Overall Gas Plant					138.2	0	OK
C-202_P	P	IN	Overall Gas Plant					612.2	0	OK
C-501_P	P	IN	Overall Gas Plant					3970.1	0	OK
C-502_P	P	IN	Overall Gas Plant					3702.3	0	OK
C-601_P	P	IN	Overall Gas Plant					2091.4	0	OK
C-602_P	P	IN	Overall Gas Plant					2174.0	0	OK
C-603_P	P	IN	Overall Gas Plant					1979.2	0	OK
C-604_P	P	IN	Overall Gas Plant					1484.6	0	OK
C-701_P	P	IN	Overall Gas Plant					6032.8	0	OK
C-901_P	P	IN	Overall Gas Plant					1005.6	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.208	-62806.1	259.7	-4779	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.208	-62806.1	216.5	4654	OK

102	S	IN	GAS+TG+CW	-193477	164.61	0.581	-536568.9	395934.5	-28303	OK
201	S	IN	GAS+TG+CW	-225649	181.98	0.006	-7996.2	6352.3	-317	OK
208	S	IN	GAS+TG+CW	-254181	174.38	0.019	-25648.8	19752.7	-996	OK
510	S	OUT	GAS+TG+CW	-100257	134.26	0.145	-170754.8	150458.6	5763	OK
705	S	OUT	GAS+TG+CW	-309848	111.68	0.289	-175364.1	76147.5	9570	OK
1101	S	OUT	GAS+TG+CW	-286104	52.78	0.001	-326.1	0.1	17	OK
1102	S	OUT	GAS+TG+CW	-284945	55.42	0.000	-28.3	0.0	2	OK
1103	S	OUT	GAS+TG+CW	-192339	87.03	0.005	-18581.5	17396.6	140	OK
1104	S	OUT	GAS+TG+CW	-287549	50.63	0.001	-207.0	0.2	10	OK
1105	S	OUT	GAS+TG+CW	-286423	49.85	0.000	-80.9	0.0	4	OK
1106	S	OUT	GAS+TG+CW	-83662	201.20	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-130193	106.31	0.006	-12628.2	11685.2	184	OK
1108	S	OUT	GAS+TG+CW	-100086	139.52	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-302035	169.53	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-302152	160.91	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-303646	147.99	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-306947	130.49	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1009-A	S	OUT	GAS+TG+CW	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1009-A	S	OUT	GAS+TG+CW	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1009-A	S	OUT	GAS+TG+CW	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1010-A	S	OUT	GAS+TG+CW	-16593	196.13	0.022	-1829.9	160.2	1299	OK
1010-A	S	OUT	GAS+TG+CW	-16593	196.13	0.022	-1829.9	160.2	1299	OK
1010-A	S	OUT	GAS+TG+CW	-16593	196.13	0.022	-1829.9	160.2	1299	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	0.558	-168025.7	694.7	12785	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	11.709	-3528112.6	14587.0	268459	OK

1209	S	OUT	GAS+TG+CW	-277945	75.40	0.208	-62806.1	216.5	4654	OK
1303-A	S	IN	GAS+TG+CW	-285558	53.18	13.431	-4046874.6	63.6	-211525	OK
1303-B	S	IN	GAS+TG+CW	-285558	53.18	13.431	-4046874.6	63.6	-211525	OK
1306	S	OUT	GAS+TG+CW	-284235	57.51	26.862	-8093749.1	1174.0	457526	OK
835	S	OUT	GAS+TG+CW	-284620	56.27	0.825	-248690.8	23.2	13753	OK
802	S	IN	GAS+TG+CW	-284234	57.51	0.825	-248690.8	36.9	-14058	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					566.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					566.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					566.0	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.172	-118900.2	0.0	-105822	OK
1002-A	S	IN	1000-GT	-97821	156.57	0.053	-64088.3	56489.9	-2444	OK
1002-A	S	IN	1000-GT	-97821	156.57	0.053	-64088.3	56489.9	-2444	OK
1002-A	S	IN	1000-GT	-97821	156.57	0.053	-64088.3	56489.9	-2444	OK
1009-A	S	OUT	1000-GT	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1009-A	S	OUT	1000-GT	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1009-A	S	OUT	1000-GT	-19604	192.10	2.214	-181162.4	11833.1	125933	OK
1010-A	S	OUT	1000-GT	-16593	196.13	0.022	-1829.9	160.2	1299	OK
1010-A	S	OUT	1000-GT	-16593	196.13	0.022	-1829.9	160.2	1299	OK
1010-A	S	OUT	1000-GT	-16593	196.13	0.022	-1829.9	160.2	1299	OK
C1001A	P	IN	1000-GT					32634.0	0	OK
C1001A	P	IN	1000-GT					32634.0	0	OK
C1001A	P	IN	1000-GT					32634.0	0	OK
T1001A	P	OUT	1000-GT					51501.8	0	OK
T1001A	P	OUT	1000-GT					51501.8	0	OK
T1001A	P	OUT	1000-GT					51501.8	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK

1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1303-A	S	IN	800-CW System	-285558	53.18	13.431	-4046874.6	63.6	-211525	OK
1303-B	S	IN	800-CW System	-285558	53.18	13.431	-4046874.6	63.6	-211525	OK
830	S	IN	800-CW System	-283160	60.86	24.248	-7306298.2	3123.6	-437015	OK
1306	S	OUT	800-CW System	-284235	57.51	26.862	-8093749.1	1174.0	457526	OK
832	S	OUT	800-CW System	-284615	56.26	24.248	-7306298.2	832.2	404021	OK
P-802A	P	IN	800-CW System					56.0	0	OK
P-802B	P	IN	800-CW System					56.0	0	OK
P-801	P	IN	800-CW System					126.5	0	OK

**Table B3.1.4. Streams exergy flow- Multiple paralleled compressors – 100% FPSO gas load – RER-1**

<i>Stream</i>	<i>Stream Type</i>	<i>Direction</i>	<i>System</i>	<i>H</i> (kJ/kgmol)	<i>S</i> (kJ/kgmol.K)	<i>Flow</i> (kgmol/s)	$\sum N_k \cdot \mu_k^0$ (kW)	$H-T_0S + P_0V - \sum N_k \cdot \mu_k^0$ (kW)	<i>S<sub>K</sub></i> kW	<i>Check</i> <i>Ex &gt; 0</i>
102	S	IN	Overall Gas Plant	-123689	163.76	2.173	-2281951.7	1907850.8	-105369	OK
201	S	IN	Overall Gas Plant	-156362	183.40	0.030	-51158.7	44918.0	-1609	OK
208	S	IN	Overall Gas Plant	-176963	176.71	0.098	-173097.0	150557.8	-5144	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.394	-1625184.2	149.3	-89868	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.109	-32694.6	3.0	-1808	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.386	-116320.6	10.7	-6432	OK
813	S	IN	Overall Gas Plant	-284622	56.26	4.690	-1413300.6	129.8	-78152	OK
815	S	IN	Overall Gas Plant	-284622	56.26	5.318	-1602345.2	147.2	-88605	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.048	-315688.3	29.0	-17457	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.340	-403707.8	37.1	-22324	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.715	-516649.3	47.5	-28569	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.131	-642009.6	59.0	-35501	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.270	-382547.8	35.1	-21154	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.995	-601016.6	55.2	-33235	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.394	-1625184.2	715.7	97684	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.109	-32694.6	14.5	1965	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.386	-116320.6	51.2	6992	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	4.690	-1413300.6	622.4	84949	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	5.318	-1602345.2	705.6	96312	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.048	-315688.3	139.0	18975	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.340	-403707.8	177.8	24266	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.715	-516649.3	227.5	31054	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	2.131	-642009.6	282.7	38589	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.270	-382547.8	168.5	22994	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.995	-601016.6	137.2	34702	OK
510	S	OUT	Overall Gas Plant	-99660	134.20	1.652	-1923425.2	1693169.5	65645	OK
402	S	OUT	Overall Gas Plant	-96059	156.34	0.146	-170595.3	149740.2	6783	OK
705	S	OUT	Overall Gas Plant	-245893	118.14	0.474	-332755.7	199640.8	16581	OK
1101	S	OUT	Overall Gas Plant	-285880	52.60	0.000	-64.0	0.0	3	OK
1102	S	OUT	Overall Gas Plant	-284254	57.53	0.000	-27.6	0.0	2	OK
1103	S	OUT	Overall Gas Plant	-180564	87.06	0.025	-78132.1	73044.5	636	OK
1104	S	OUT	Overall Gas Plant	-286952	49.90	0.003	-789.2	0.6	39	OK
1105	S	OUT	Overall Gas Plant	-286498	49.59	0.001	-235.2	0.1	11	OK
1106	S	OUT	Overall Gas Plant	-123654	77.86	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-127302	118.83	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-97534	146.03	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-239668	172.88	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-239507	165.12	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-240046	155.27	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-241741	142.64	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					4058.8	0	OK
C-101-B_P	P	IN	Overall Gas Plant					4058.8	0	OK
C-201-A_P	P	IN	Overall Gas Plant					42.9	0	OK
C-201-B_P	P	IN	Overall Gas Plant					42.9	0	OK
C-201-C_P	P	IN	Overall Gas Plant					42.9	0	OK
C-202-A_P	P	IN	Overall Gas Plant					180.0	0	OK
C-202-B_P	P	IN	Overall Gas Plant					180.0	0	OK
C-202-C_P	P	IN	Overall Gas Plant					180.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-501-B_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-501-C_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-502-A_P	P	IN	Overall Gas Plant					1583.6	0	OK
C-502-B_P	P	IN	Overall Gas Plant					1583.6	0	OK

C-502-C_P	P	IN	Overall Gas Plant					1583.6	0	OK
C-601_P	P	IN	Overall Gas Plant					1707.3	0	OK
C-602_P	P	IN	Overall Gas Plant					1824.1	0	OK
C-603_P	P	IN	Overall Gas Plant					1864.4	0	OK
C-604_P	P	IN	Overall Gas Plant					1762.3	0	OK
C-701-A_P	P	IN	Overall Gas Plant					1551.8	0	OK
C-701-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant					229.4	0	OK
C-901-B_P	P	IN	Overall Gas Plant					229.4	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.767	-231176.8	955.8	-17591	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.767	-231176.8	796.8	17131	OK
102	S	IN	GAS+TG+CW	-123689	163.76	2.173	-2281951.7	1907850.8	-105369	OK
201	S	IN	GAS+TG+CW	-156362	183.40	0.030	-51158.7	44918.0	-1609	OK
208	S	IN	GAS+TG+CW	-176963	176.71	0.098	-173097.0	150557.8	-5144	OK
510	S	OUT	GAS+TG+CW	-99660	134.20	1.652	-1923425.2	1693169.5	65645	OK
705	S	OUT	GAS+TG+CW	-245893	118.14	0.474	-332755.7	199640.8	16581	OK
1101	S	OUT	GAS+TG+CW	-285880	52.60	0.000	-64.0	0.0	3	OK
1102	S	OUT	GAS+TG+CW	-284254	57.53	0.000	-27.6	0.0	2	OK
1103	S	OUT	GAS+TG+CW	-180564	87.06	0.025	-78132.1	73044.5	636	OK
1104	S	OUT	GAS+TG+CW	-286952	49.90	0.003	-789.2	0.6	39	OK
1105	S	OUT	GAS+TG+CW	-286498	49.59	0.001	-235.2	0.1	11	OK
1106	S	OUT	GAS+TG+CW	-123654	77.86	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-127302	118.83	0.000	0.0	0.0	0	OK
1108	S	OUT	GAS+TG+CW	-97534	146.03	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-239668	172.88	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-239507	165.12	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-240046	155.27	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-241741	142.64	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.791	-152750.7	0.0	-135949	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.791	-152750.7	0.0	-135949	OK
1009-A	S	OUT	GAS+TG+CW	-19787	193.16	2.848	-235672.9	16424.5	162902	OK

1009-A	S	OUT	GAS+TG+CW	-19787	193.16	2.848	-235672.9	16424.5	162902	OK
1010-A	S	OUT	GAS+TG+CW	-17445	196.24	0.029	-2380.5	207.0	1672	OK
1010-A	S	OUT	GAS+TG+CW	-17445	196.24	0.029	-2380.5	207.0	1672	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	6.193	-1866070.3	7715.3	141992	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	1.356	-408715.7	1689.8	31100	OK
1209	S	OUT	GAS+TG+CW	-277945	75.40	0.767	-231176.8	796.8	17131	OK
1303-A	S	IN	GAS+TG+CW	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
1303-B	S	IN	GAS+TG+CW	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
1306	S	OUT	GAS+TG+CW	-284236	57.51	30.061	-9057714.1	1313.3	512008	OK
835	S	OUT	GAS+TG+CW	-284620	56.27	3.522	-1061333.6	98.6	58694	OK
802	S	IN	GAS+TG+CW	-284234	57.51	3.522	-1061333.6	157.6	-59995	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					790.3	0	OK
Gen Loss	P	OUT	GAS+TG+CW					790.3	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.791	-152750.7	0.0	-135949	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.791	-152750.7	0.0	-135949	OK
1002-A	S	IN	1000-GT	-96059	158.13	0.073	-85297.6	74831.2	-3430	OK
1002-A	S	IN	1000-GT	-96059	158.13	0.073	-85297.6	74831.2	-3430	OK
1009-A	S	OUT	1000-GT	-19787	193.16	2.848	-235672.9	16424.5	162902	OK
1009-A	S	OUT	1000-GT	-19787	193.16	2.848	-235672.9	16424.5	162902	OK
1010-A	S	OUT	1000-GT	-17445	196.24	0.029	-2380.5	207.0	1672	OK
1010-A	S	OUT	1000-GT	-17445	196.24	0.029	-2380.5	207.0	1672	OK
C1001A	P	IN	1000-GT					41337.8	0	OK
C1001A	P	IN	1000-GT					41337.8	0	OK
T1001A	P	OUT	1000-GT					67682.1	0	OK
T1001A	P	OUT	1000-GT					67682.1	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK

1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1303-A	S	IN	800-CW System	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
1303-B	S	IN	800-CW System	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
830	S	IN	800-CW System	-283266	60.53	29.249	-8813109.3	3477.9	-524333	OK
1306	S	OUT	800-CW System	-284236	57.51	30.061	-9057714.1	1313.3	512008	OK
832	S	OUT	800-CW System	-284615	56.26	29.249	-8813109.3	1003.9	487346	OK
P-802A	P	IN	800-CW System					63.1	0	OK
P-802B	P	IN	800-CW System					63.1	0	OK
P-801	P	IN	800-CW System					157.6	0	OK

**Table B3.1.5. Streams exergy flow- Multiple paralleled compressors – 50% FPSO gas load – RER-1**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>		<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>(kW)</i>	$H-T_0S + P_0V - \sum N_k \cdot \mu_k^0$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>kW</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-146519	164.81	1.298	-1321446.9	1067969.5	-63339	OK
201	S	IN	Overall Gas Plant	-178368	183.66	0.016	-26695.9	22925.0	-881	OK
208	S	IN	Overall Gas Plant	-204760	176.13	0.046	-74560.7	62658.7	-2416	OK
803	S	IN	Overall Gas Plant	-284622	56.26	3.683	-1109831.4	101.9	-61371	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.040	-12180.4	1.1	-674	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.060	-18115.0	1.7	-1002	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.132	-39764.9	3.7	-2199	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.149	-44804.8	4.1	-2478	OK
813	S	IN	Overall Gas Plant	-284622	56.26	2.219	-668756.7	61.4	-36980	OK
815	S	IN	Overall Gas Plant	-284622	56.26	2.585	-778940.0	71.6	-43073	OK
817	S	IN	Overall Gas Plant	-284622	56.26	0.882	-265819.6	24.4	-14699	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.180	-355461.3	32.7	-19656	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.547	-466125.2	42.8	-25775	OK
823	S	IN	Overall Gas Plant	-284622	56.26	1.934	-582608.8	53.5	-32217	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.057	-318397.9	29.2	-17606	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.349	-406417.2	37.3	-22474	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	3.683	-1109831.4	488.7	66708	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.040	-12180.4	1.8	689	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.060	-18115.0	8.0	1089	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.132	-39764.9	9.1	2296	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.149	-44804.8	19.7	2693	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	2.219	-668756.7	294.5	40197	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	2.585	-778940.0	343.0	46819	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	0.882	-265819.6	117.1	15978	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.180	-355461.3	156.5	21366	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.547	-466125.2	205.3	28017	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	1.934	-582608.8	256.6	35019	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.057	-318397.9	140.2	19138	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.349	-406417.2	92.7	23466	OK
510	S	OUT	Overall Gas Plant	-100221	134.02	0.805	-951393.1	838826.0	31933	OK
402	S	OUT	Overall Gas Plant	-96648	155.92	0.122	-144185.2	126770.2	5630	OK
705	S	OUT	Overall Gas Plant	-273135	115.64	0.414	-273771.2	146500.7	14179	OK
1101	S	OUT	Overall Gas Plant	-285273	54.86	0.000	0.0	0.0	0	OK
1102	S	OUT	Overall Gas Plant	-284261	57.54	0.000	-54.2	0.0	3	OK
1103	S	OUT	Overall Gas Plant	-181522	85.12	0.016	-52360.4	48960.9	415	OK
1104	S	OUT	Overall Gas Plant	-287081	50.33	0.003	-768.6	0.6	38	OK
1105	S	OUT	Overall Gas Plant	-286420	49.86	0.001	-158.0	0.1	8	OK
1106	S	OUT	Overall Gas Plant	-123639	77.92	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-129428	116.12	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-98082	145.88	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-266336	171.41	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-266086	163.95	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-266675	154.00	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-268594	140.81	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					2482.2	0	OK
C-101-B_P	P	IN	Overall Gas Plant					2482.2	0	OK
C-201-A_P	P	IN	Overall Gas Plant					34.9	0	OK
C-201-B_P	P	IN	Overall Gas Plant					34.9	0	OK
C-201-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-A_P	P	IN	Overall Gas Plant					139.7	0	OK
C-202-B_P	P	IN	Overall Gas Plant					139.7	0	OK
C-202-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					1156.2	0	OK
C-501-B_P	P	IN	Overall Gas Plant					1156.2	0	OK
C-501-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-A_P	P	IN	Overall Gas Plant					1140.3	0	OK
C-502-B_P	P	IN	Overall Gas Plant					1140.3	0	OK

C-502-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-601_P	P	IN	Overall Gas Plant					1476.1	0	OK
C-602_P	P	IN	Overall Gas Plant					1587.8	0	OK
C-603_P	P	IN	Overall Gas Plant					1612.1	0	OK
C-604_P	P	IN	Overall Gas Plant					1484.2	0	OK
C-701-A_P	P	IN	Overall Gas Plant					1280.5	0	OK
C-701-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant					155.1	0	OK
C-901-B_P	P	IN	Overall Gas Plant					155.1	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.457	-137710.3	569.4	-10479	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.457	-137710.3	474.6	10205	OK
102	S	IN	GAS+TG+CW	-146519	164.81	1.298	-1321446.9	1067969.5	-63339	OK
201	S	IN	GAS+TG+CW	-178368	183.66	0.016	-26695.9	22925.0	-881	OK
208	S	IN	GAS+TG+CW	-204760	176.13	0.046	-74560.7	62658.7	-2416	OK
510	S	OUT	GAS+TG+CW	-100221	134.02	0.805	-951393.1	838826.0	31933	OK
705	S	OUT	GAS+TG+CW	-273135	115.64	0.414	-273771.2	146500.7	14179	OK
1101	S	OUT	GAS+TG+CW	-285273	54.86	0.000	0.0	0.0	0	OK
1102	S	OUT	GAS+TG+CW	-284261	57.54	0.000	-54.2	0.0	3	OK
1103	S	OUT	GAS+TG+CW	-181522	85.12	0.016	-52360.4	48960.9	415	OK
1104	S	OUT	GAS+TG+CW	-287081	50.33	0.003	-768.6	0.6	38	OK
1105	S	OUT	GAS+TG+CW	-286420	49.86	0.001	-158.0	0.1	8	OK
1106	S	OUT	GAS+TG+CW	-123639	77.92	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-129428	116.12	0.000	0.0	0.0	0	OK
1108	S	OUT	GAS+TG+CW	-98082	145.88	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-266336	171.41	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-266086	163.95	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-266675	154.00	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-268594	140.81	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.436	-133352.4	0.0	-118685	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.436	-133352.4	0.0	-118685	OK
1009-A	S	OUT	GAS+TG+CW	-19329	192.58	2.483	-203394.8	13755.4	141638	OK

1009-A	S	OUT	GAS+TG+CW	-19329	192.58	2.483	-203394.8	13755.4	141638	OK
1010-A	S	OUT	GAS+TG+CW	-16644	196.15	0.025	-2054.5	179.8	1457	OK
1010-A	S	OUT	GAS+TG+CW	-16644	196.15	0.025	-2054.5	179.8	1457	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	2.623	-790324.3	3267.6	60137	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	5.237	-1577928.3	6523.9	120067	OK
1209	S	OUT	GAS+TG+CW	-277945	75.40	0.457	-137710.3	474.6	10205	OK
1303-A	S	IN	GAS+TG+CW	-285557	53.18	10.021	-3019416.3	48.0	-157824	OK
1303-B	S	IN	GAS+TG+CW	-285557	53.18	10.021	-3019416.3	48.0	-157824	OK
1306	S	OUT	GAS+TG+CW	-284236	57.51	20.042	-6038832.6	875.7	341360	OK
835	S	OUT	GAS+TG+CW	-284620	56.27	3.650	-1099717.2	103.1	60816	OK
802	S	IN	GAS+TG+CW	-284234	57.51	3.650	-1099717.2	163.3	-62164	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					635.8	0	OK
Gen Loss	P	OUT	GAS+TG+CW					635.8	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.436	-133352.4	0.0	-118685	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.436	-133352.4	0.0	-118685	OK
1002-A	S	IN	1000-GT	-96648	157.69	0.061	-72092.6	63353.1	-2847	OK
1002-A	S	IN	1000-GT	-96648	157.69	0.061	-72092.6	63353.1	-2847	OK
1009-A	S	OUT	1000-GT	-19329	192.58	2.483	-203394.8	13755.4	141638	OK
1009-A	S	OUT	1000-GT	-19329	192.58	2.483	-203394.8	13755.4	141638	OK
1010-A	S	OUT	1000-GT	-16644	196.15	0.025	-2054.5	179.8	1457	OK
1010-A	S	OUT	1000-GT	-16644	196.15	0.025	-2054.5	179.8	1457	OK
C1001A	P	IN	1000-GT					36600.7	0	OK
C1001A	P	IN	1000-GT					36600.7	0	OK
T1001A	P	OUT	1000-GT					57792.9	0	OK
T1001A	P	OUT	1000-GT					57792.9	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK

1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1303-A	S	IN	800-CW System	-285557	53.18	10.021	-3019416.3	48.0	-157824	OK
1303-B	S	IN	800-CW System	-285557	53.18	10.021	-3019416.3	48.0	-157824	OK
830	S	IN	800-CW System	-283330	60.33	20.467	-6166940.4	2321.9	-365703	OK
1306	S	OUT	800-CW System	-284236	57.51	20.042	-6038832.6	875.7	341360	OK
832	S	OUT	800-CW System	-284615	56.26	20.467	-6166940.4	702.4	341017	OK
P-802A	P	IN	800-CW System					41.2	0	OK
P-802B	P	IN	800-CW System					41.2	0	OK
P-801	P	IN	800-CW System					104.2	0	OK

**Table B3.1.6. Streams exergy flow- Multiple paralleled compressors – 25% FPSO gas load – RER-1**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>		<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>(kW)</i>	$\frac{H-T_0S + P_0V - \sum N_k \cdot \mu_k^0}{N_k \cdot \mu_k^0}$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>kW</i>	<i>Check</i> <i>Ex &gt; 0</i>
102	S	IN	Overall Gas Plant	-193477	164.61	0.581	-536568.9	395934.5	-28303	OK
201	S	IN	Overall Gas Plant	-225649	181.98	0.006	-7996.2	6352.3	-317	OK
208	S	IN	Overall Gas Plant	-254182	174.37	0.019	-25648.8	19752.7	-996	OK
803	S	IN	Overall Gas Plant	-284622	56.26	1.851	-557702.2	51.2	-30839	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.006	-1714.2	0.2	-95	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.021	-6443.1	0.6	-356	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.046	-13802.5	1.3	-763	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.056	-16854.6	1.5	-932	OK
813	S	IN	Overall Gas Plant	-284622	56.26	0.000	0.0	0.0	0	OK
815	S	IN	Overall Gas Plant	-284622	56.26	0.000	0.0	0.0	0	OK
817	S	IN	Overall Gas Plant	-284622	56.26	0.564	-170056.2	15.6	-9404	OK
819	S	IN	Overall Gas Plant	-284622	56.26	0.834	-251322.7	23.1	-13897	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.620	-488056.4	44.8	-26988	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.208	-665227.8	61.1	-36785	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.264	-380897.8	35.0	-21063	OK
827	S	IN	Overall Gas Plant	-284622	56.26	0.584	-175892.0	16.2	-9726	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	1.851	-557702.2	245.6	33522	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.006	-1714.2	0.2	97	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.021	-6443.1	2.9	387	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.046	-13802.5	3.1	797	OK
812	S	OUT	Overall Gas Plant	-283068	61.16	0.056	-16854.6	7.4	1013	OK
814	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
816	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	0.564	-170056.2	74.9	10222	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	0.834	-251322.7	110.7	15106	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.620	-488056.4	214.9	29335	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	2.208	-665227.8	292.9	39985	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.264	-380897.8	167.7	22894	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	0.584	-175892.0	40.1	10156	OK
402	S	OUT	Overall Gas Plant	-97760	154.88	0.097	-117905.5	103955.1	4455	OK
405	S	OUT	Overall Gas Plant	-97760	154.88	0.000	0.0	0.0	0	OK
705	S	OUT	Overall Gas Plant	-221506	121.21	0.500	-431717.8	302892.1	17966	OK
1101	S	OUT	Overall Gas Plant	-284838	56.72	0.000	0.0	0.0	0	OK
1102	S	OUT	Overall Gas Plant	-284275	57.55	0.000	-6.0	0.0	0	OK
1103	S	OUT	Overall Gas Plant	-192973	82.99	0.006	-19978.1	18644.5	151	OK
1104	S	OUT	Overall Gas Plant	-287563	50.60	0.002	-522.5	0.6	26	OK
1105	S	OUT	Overall Gas Plant	-286432	49.82	0.000	-80.1	0.0	4	OK
1106	S	OUT	Overall Gas Plant	-123645	77.89	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-134540	111.38	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-302216	168.76	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-301763	162.00	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-219786	118.38	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-217511	145.10	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					2269.6	0	OK
C-101-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-201-A_P	P	IN	Overall Gas Plant					26.0	0	OK
C-201-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-201-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-A_P	P	IN	Overall Gas Plant					117.8	0	OK
C-202-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-A_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-B_P	P	IN	Overall Gas Plant					0.0	0	OK

C-502-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-601_P	P	IN	Overall Gas Plant					1008.9	0	OK
C-602_P	P	IN	Overall Gas Plant					1103.6	0	OK
C-603_P	P	IN	Overall Gas Plant					1804.1	0	OK
C-604_P	P	IN	Overall Gas Plant					1755.3	0	OK
C-701-A_P	P	IN	Overall Gas Plant					1634.9	0	OK
C-701-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant					129.9	0	OK
C-901-B_P	P	IN	Overall Gas Plant					0.0	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.208	-62651.6	259.0	-4767	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.208	-62651.6	215.9	4643	OK
102	S	IN	GAS+TG+CW	-193477	164.61	0.581	-536568.9	395934.5	-28303	OK
201	S	IN	GAS+TG+CW	-225649	181.98	0.006	-7996.2	6352.3	-317	OK
208	S	IN	GAS+TG+CW	-254182	174.37	0.019	-25648.8	19752.7	-996	OK
405	S	OUT	GAS+TG+CW	-97760	154.88	0.000	0.0	0.0	0	OK
705	S	OUT	GAS+TG+CW	-221506	121.21	0.500	-431717.8	302892.1	17966	OK
1101	S	OUT	GAS+TG+CW	-284838	56.72	0.000	0.0	0.0	0	OK
1102	S	OUT	GAS+TG+CW	-284275	57.55	0.000	-6.0	0.0	0	OK
1103	S	OUT	GAS+TG+CW	-192973	82.99	0.006	-19978.1	18644.5	151	OK
1104	S	OUT	GAS+TG+CW	-287563	50.60	0.002	-522.5	0.6	26	OK
1105	S	OUT	GAS+TG+CW	-286432	49.82	0.000	-80.1	0.0	4	OK
1106	S	OUT	GAS+TG+CW	-123645	77.89	0.000	0.0	0.0	0	OK
1107	S	OUT	GAS+TG+CW	-134540	111.38	0.000	0.0	0.0	0	OK
1108	S	OUT	GAS+TG+CW	0	0.00	0.000	0.0	0.0	0	OK
1109	S	OUT	GAS+TG+CW	-302216	168.76	0.000	0.0	0.0	0	OK
1110	S	OUT	GAS+TG+CW	-301763	162.00	0.000	0.0	0.0	0	OK
1111	S	OUT	GAS+TG+CW	-219786	118.38	0.000	0.0	0.0	0	OK
1112	S	OUT	GAS+TG+CW	-217511	145.10	0.000	0.0	0.0	0	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.014	-110206.6	0.0	-98085	OK
1003-A	S	IN	GAS+TG+CW	-6020	164.49	2.014	-110206.6	0.0	-98085	OK
1009-A	S	OUT	GAS+TG+CW	-19653	191.74	2.051	-167471.3	10673.9	116482	OK

1009-A	S	OUT	GAS+TG+CW	-19653	191.74	2.051	-167471.3	10673.9	116482	OK
1010-A	S	OUT	GAS+TG+CW	-16403	196.11	0.021	-1691.6	148.3	1203	OK
1010-A	S	OUT	GAS+TG+CW	-16403	196.11	0.021	-1691.6	148.3	1203	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1201-A	S	IN	GAS+TG+CW	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1204	S	OUT	GAS+TG+CW	-277139	77.42	0.558	-168025.7	694.7	12785	OK
1206	S	OUT	GAS+TG+CW	-277139	77.42	7.551	-2275285.6	9407.2	173129	OK
1209	S	OUT	GAS+TG+CW	-277945	75.40	0.208	-62651.6	215.9	4643	OK
1303-A	S	IN	GAS+TG+CW	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
1303-B	S	IN	GAS+TG+CW	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
1306	S	OUT	GAS+TG+CW	-284235	57.51	10.569	-3184558.2	462.0	180018	OK
835	S	OUT	GAS+TG+CW	-284621	56.26	0.825	-248690.8	23.5	13753	OK
802	S	IN	GAS+TG+CW	-284234	57.51	0.825	-248690.8	36.9	-14058	OK
Other users	P	OUT	GAS+TG+CW					23164.0	0	OK
Gen Loss	P	OUT	GAS+TG+CW					513.6	0	OK
Gen Loss	P	OUT	GAS+TG+CW					513.6	0	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.014	-110206.6	0.0	-98085	OK
1003-A	S	IN	1000-GT	-6020	164.49	2.014	-110206.6	0.0	-98085	OK
1002-A	S	IN	1000-GT	-97760	156.61	0.049	-58952.7	51952.7	-2252	OK
1002-A	S	IN	1000-GT	-97760	156.61	0.049	-58952.7	51952.7	-2252	OK
1009-A	S	OUT	1000-GT	-19653	191.74	2.051	-167471.3	10673.9	116482	OK
1009-A	S	OUT	1000-GT	-19653	191.74	2.051	-167471.3	10673.9	116482	OK
1010-A	S	OUT	1000-GT	-16403	196.11	0.021	-1691.6	148.3	1203	OK
1010-A	S	OUT	1000-GT	-16403	196.11	0.021	-1691.6	148.3	1203	OK
C1001A	P	IN	1000-GT					30610.2	0	OK
C1001A	P	IN	1000-GT					30610.2	0	OK
T1001A	P	OUT	1000-GT					47729.1	0	OK
T1001A	P	OUT	1000-GT					47729.1	0	OK
1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK

1201-A	S	IN	1000-GT	-278742	73.33	4.158	-1252981.4	3543.6	-90311	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1202-A	S	OUT	1000-GT	-277139	77.42	4.158	-1252981.4	5180.5	95341	OK
1303-A	S	IN	800-CW System	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
1303-B	S	IN	800-CW System	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
830	S	IN	800-CW System	-283211	60.70	9.879	-2976660.3	1228.4	-177585	OK
1306	S	OUT	800-CW System	-284235	57.51	10.569	-3184558.2	462.0	180018	OK
832	S	OUT	800-CW System	-284616	56.26	9.879	-2976660.3	339.0	164601	OK
P-802A	P	IN	800-CW System					21.4	0	OK
P-802B	P	IN	800-CW System					21.4	0	OK
P-801	P	IN	800-CW System					48.2	0	OK

- RER-2

**Table B3.1.7. Streams exergy flow- Single-shaft compressors – 100% FPSO gas load – RER-2**

Stream	Stream Type	Direction	System	H	S	Flow	$\sum N_k \cdot \mu_k^0$	$\frac{H-T_0S + P_0V - \sum N_k \cdot \mu_k^0}{N_k \cdot \mu_k^0}$	$S_K$	Check
	S/P	In/Out		(kJ/kgmol)	(kJ/kgmol.K)	(kgmol/s)	Flow (kW)	Flow (kW)	(kW)	Ex >0
102	S	IN	Overall Gas Plant	-123689	163.76	2.173	-446929.1	72828.2	-105369	OK
201	S	IN	Overall Gas Plant	-156362	183.40	0.030	-7474.2	1233.6	-1609	OK
208	S	IN	Overall Gas Plant	-176963	176.71	0.098	-26693.3	4154.1	-5144	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.463	-1646213.1	151.2	-91031	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.115	-34726.0	3.2	-1920	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.430	-129531.9	11.9	-7163	OK
813	S	IN	Overall Gas Plant	-284622	56.26	4.522	-1362576.3	125.2	-75347	OK
815	S	IN	Overall Gas Plant	-284622	56.26	5.168	-1557150.0	143.0	-86106	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.315	-396246.1	36.4	-21911	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.608	-484492.7	44.5	-26791	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.955	-588977.4	54.1	-32569	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.304	-694244.5	63.8	-38390	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.611	-1389279.6	127.6	-76823	OK
827	S	IN	Overall Gas Plant	-284622	56.26	2.031	-612079.3	56.2	-33846	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.463	-1646213.1	724.9	98948	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.115	-34726.0	15.4	2087	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.430	-129531.9	57.0	7786	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	4.522	-1362576.3	600.0	81900	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	5.168	-1557150.0	685.7	93595	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.315	-396246.1	174.5	23817	OK

820	S	OUT	Overall Gas Plant	-283068	61.15	1.608	-484492.7	213.4	29121	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.955	-588977.4	259.4	35401	OK
824	S	OUT	Overall Gas Plant	-283068	61.15	2.304	-694244.5	305.7	41729	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	4.611	-1389279.6	611.8	83505	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	2.031	-612079.3	139.7	35340	OK
510	S	OUT	Overall Gas Plant	-99659	134.20	1.624	-293318.9	66904.8	64552	OK
402	S	OUT	Overall Gas Plant	-96058	156.34	0.174	-31436.0	6656.1	8059	OK
705	S	OUT	Overall Gas Plant	-245894	118.13	0.474	-148975.3	15853.1	16581	OK
1101	S	OUT	Overall Gas Plant	-285888	52.58	0.000	-67.6	0.0	3	OK
1102	S	OUT	Overall Gas Plant	-284367	57.17	0.000	-50.2	0.0	3	OK
1103	S	OUT	Overall Gas Plant	-180135	87.06	0.025	-6259.1	1163.5	638	OK
1104	S	OUT	Overall Gas Plant	-286955	49.90	0.003	-783.2	0.6	38	OK
1105	S	OUT	Overall Gas Plant	-286501	49.58	0.001	-234.8	0.1	11	OK
1106	S	OUT	Overall Gas Plant	-122967	80.40	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-127699	118.00	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-97645	145.67	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-239637	172.98	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-239613	164.78	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-240373	154.22	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-242403	140.59	0.000	0.0	0.0	0	OK
C-101_P	P	IN	Overall Gas Plant					8222.8	0	OK
C-201_P	P	IN	Overall Gas Plant					139.0	0	OK
C-202_P	P	IN	Overall Gas Plant					604.8	0	OK
C-501_P	P	IN	Overall Gas Plant					4635.3	0	OK
C-502_P	P	IN	Overall Gas Plant					4576.7	0	OK
C-601_P	P	IN	Overall Gas Plant					2122.9	0	OK
C-602_P	P	IN	Overall Gas Plant					2241.1	0	OK
C-603_P	P	IN	Overall Gas Plant					2237.8	0	OK
C-604_P	P	IN	Overall Gas Plant					2031.7	0	OK
C-701_P	P	IN	Overall Gas Plant					6744.0	0	OK
C-901_P	P	IN	Overall Gas Plant					493.8	0	OK

1208	S	IN	Overall Gas Plant	-277139	77.42	0.767	-231168.6	955.8	-17590	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.767	-231168.6	796.7	17131	OK
1303-A	S	IN	800-CW System	-285558	53.18	17.466	-5262603.5	81.4	-275068	OK
1303-B	S	IN	800-CW System	-285558	53.18	17.466	-5262603.5	81.4	-275068	OK
830	S	IN	800-CW System	-283242	60.61	33.378	-10057161.8	4037.5	-599089	OK
1306	S	OUT	800-CW System	-284236	57.51	34.931	-10525207.1	1525.4	594950	OK
832	S	OUT	800-CW System	-284615	56.26	33.378	-10057161.8	1145.6	556141	OK
P-802A	P	IN	800-CW System					74.3	0	OK
P-802B	P	IN	800-CW System					74.3	0	OK
P-801	P	IN	800-CW System					185.7	0	OK
102	S	IN	100-Main Comp.	-123689	163.76	2.173	-446929.1	72828.2	-105369	OK
803	S	IN	100-Main Comp.	-284622	56.26	5.463	-1646213.1	151.2	-91031	OK
107	S	OUT	100-Main Comp.	-126529	155.67	2.315	-483882.1	84325.0	106702	OK
804	S	OUT	100-Main Comp.	-283068	61.15	5.463	-1646213.1	724.9	98948	OK
1101	S	OUT	100-Main Comp.	-285888	52.58	0.000	-67.6	0.0	3	OK
214	S	IN	100-Main Comp.	-168313	160.53	0.142	-37020.6	6345.6	-6756	OK
C-101_P	P	IN	100-Main Comp.					8222.8	0	OK
404	S	IN	500-Export Comp.	-96058	156.34	1.624	-293318.9	62105.9	-75199	OK
813	S	IN	500-Export Comp.	-284622	56.26	4.522	-1362576.3	125.2	-75347	OK
815	S	IN	500-Export Comp.	-284622	56.26	5.168	-1557150.0	143.0	-86106	OK
510	S	OUT	500-Export Comp.	-99659	134.20	1.624	-293318.9	66904.8	64552	OK
814	S	OUT	500-Export Comp.	-283068	61.15	4.522	-1362576.3	600.0	81900	OK
816	S	OUT	500-Export Comp.	-283068	61.15	5.168	-1557150.0	685.7	93595	OK
1107	S	OUT	500-Export Comp.	-127699	118.00	0.000	0.0	0.0	0	OK
1108	S	OUT	500-Export Comp.	-97645	145.67	0.000	0.0	0.0	0	OK
C-501_P	P	IN	500-Export Comp.					4635.3	0	OK
C-502_P	P	IN	500-Export Comp.					4576.7	0	OK
403	S	IN	600-CO2 Comp.	-239668	172.88	0.474	-148975.3	11119.9	-24265	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	1.315	-396246.1	36.4	-21911	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	1.608	-484492.7	44.5	-26791	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.955	-588977.4	54.1	-32569	OK

823	S	IN	600-CO2 Comp.	-284622	56.26	2.304	-694244.5	63.8	-38390	OK
620	S	OUT	600-CO2 Comp.	-245003	126.37	0.474	-148975.3	15119.2	17737	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	1.315	-396246.1	174.5	23817	OK
820	S	OUT	600-CO2 Comp.	-283068	61.15	1.608	-484492.7	213.4	29121	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.955	-588977.4	259.4	35401	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	2.304	-694244.5	305.7	41729	OK
1109	S	OUT	600-CO2 Comp.	-239637	172.98	0.000	0.0	0.0	0	OK
1110	S	OUT	600-CO2 Comp.	-239613	164.78	0.000	0.0	0.0	0	OK
1111	S	OUT	600-CO2 Comp.	-240373	154.22	0.000	0.0	0.0	0	OK
1112	S	OUT	600-CO2 Comp.	-242403	140.59	0.000	0.0	0.0	0	OK
C-601_P	P	IN	600-CO2 Comp.					2122.9	0	OK
C-602_P	P	IN	600-CO2 Comp.					2241.1	0	OK
C-603_P	P	IN	600-CO2 Comp.					2237.8	0	OK
C-604_P	P	IN	600-CO2 Comp.					2031.7	0	OK
620	S	IN	700-EOR Comp.	-245003	126.37	0.474	-148975.3	15119.2	-17737	OK
825	S	IN	700-EOR Comp.	-284622	56.26	4.611	-1389279.6	127.6	-76823	OK
705	S	OUT	700-EOR Comp.	-245894	118.13	0.474	-148975.3	15853.1	16581	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	4.611	-1389279.6	611.8	83505	OK
C-701_P	P	IN	700-EOR Comp.					6744.0	0	OK
307	S	IN	400-MP Unit	-126286	155.39	2.272	-473730.2	82223.5	-104563	OK
401	S	OUT	400-MP Unit	-96058	156.34	1.798	-324754.9	68762.0	83258	OK
403	S	OUT	400-MP Unit	-239668	172.88	0.474	-148975.3	11119.9	24265	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.767	-231168.6	955.8	-17590	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.767	-231168.6	796.7	17131	OK
901	S	IN	900-C3 Cycle	-106646	141.76	0.101	-21354.6	6396.1	-4225	OK
827	S	IN	900-C3 Cycle	-284622	56.26	2.031	-612079.3	56.2	-33846	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.101	-21354.6	6518.3	3019	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	2.031	-612079.3	139.7	35340	OK
C-901_P	P	IN	900-C3 Cycle					493.8	0	OK

201	S	IN	200-VRU	-156362	183.40	0.030	-7474.2	1233.6	-1609	OK
208	S	IN	200-VRU	-176963	176.71	0.098	-26693.3	4154.1	-5144	OK
317	S	IN	200-VRU	-154621	111.94	0.039	-9162.3	1820.2	-1296	OK
805	S	IN	200-VRU	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	200-VRU	-284622	56.26	0.115	-34726.0	3.2	-1920	OK
809	S	IN	200-VRU	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	200-VRU	-284622	56.26	0.430	-129531.9	11.9	-7163	OK
806	S	OUT	200-VRU	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	200-VRU	-283067	61.15	0.115	-34726.0	15.4	2087	OK
810	S	OUT	200-VRU	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	200-VRU	-283068	61.15	0.430	-129531.9	57.0	7786	OK
214	S	OUT	200-VRU	-168313	160.53	0.142	-37020.6	6345.6	6756	OK
1102	S	OUT	200-VRU	-284367	57.17	0.000	-50.2	0.0	3	OK
1103	S	OUT	200-VRU	-180135	87.06	0.025	-6259.1	1163.5	638	OK
C-201_P	P	IN	200-VRU					139.0	0	OK
C-202_P	P	IN	200-VRU					604.8	0	OK
303	S	IN	300-HCDP Adj.	-127848	150.50	2.299	-479959.6	83592.0	-102463	OK
305	S	OUT	300-HCDP Adj.	-128021	149.37	2.272	-473730.2	82330.1	100514	OK
312	S	OUT	300-HCDP Adj.	-153765	106.80	0.027	-6229.4	1283.9	844	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.101	-21354.6	6518.3	-3019	OK
901	S	OUT	300-HCDP Adj.	-106646	141.76	0.101	-21354.6	6396.1	4225	OK
801	S	IN	Oil Plant	-284622	56.26	3.522	-1061333.6	97.5	-58689	OK
802	S	OUT	Oil Plant	-284234	57.51	3.522	-1061333.6	157.6	59995	OK

**Table B3.1.8. Streams exergy flow - Single-shaft compressors – 50% FPSO gas load – RER approach II**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \cdot \mu_k^0$ <i>Flow (kW)</i>	$H - T_0 S + P_0 V - \sum N_k \cdot \mu_k^0$ <i>Flow (kW)</i>	<i>Check</i> <i>Ex &gt; 0</i>
102	S	IN	Overall Gas Plant	-146519	164.81	1.298	-295989.8	42512.4	OK
201	S	IN	Overall Gas Plant	-178368	183.66	0.016	-4408.1	637.2	OK
208	S	IN	Overall Gas Plant	-204760	176.13	0.046	-13722.7	1820.7	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.843	-1760425.3	161.7	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.040	-12180.4	1.1	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.103	-31057.8	2.9	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.132	-39764.9	3.7	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.340	-102381.6	9.4	OK
813	S	IN	Overall Gas Plant	-284622	56.26	3.392	-1022120.6	93.9	OK
815	S	IN	Overall Gas Plant	-284622	56.26	3.648	-1099152.0	101.0	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.290	-388600.7	35.7	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.586	-477876.2	43.9	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.899	-572142.2	52.6	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.167	-652824.6	60.0	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.376	-1318543.4	121.1	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.703	-513218.8	47.1	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.843	-1760425.3	775.2	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.040	-12180.4	1.8	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.103	-31057.8	13.8	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.132	-39764.9	9.1	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.340	-102381.6	45.1	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	3.392	-1022120.6	450.1	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	3.648	-1099152.0	484.0	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.290	-388600.7	171.1	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.586	-477876.2	210.4	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.899	-572142.2	252.0	OK
824	S	OUT	Overall Gas Plant	-283068	61.15	2.167	-652824.6	287.5	OK

826	S	OUT	Overall Gas Plant	-283068	61.15	4.376	-1318543.4	580.6	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.703	-513218.8	117.1	OK
510	S	OUT	Overall Gas Plant	-100232	134.01	0.760	-137929.7	31574.5	OK
402	S	OUT	Overall Gas Plant	-96657	155.92	0.167	-30253.4	6440.0	OK
705	S	OUT	Overall Gas Plant	-273140	115.63	0.414	-140865.2	13583.2	OK
1101	S	OUT	Overall Gas Plant	-285676	53.53	0.001	-175.4	0.0	OK
1102	S	OUT	Overall Gas Plant	-284732	56.03	0.000	-99.7	0.0	OK
1103	S	OUT	Overall Gas Plant	-180790	86.07	0.016	-4041.0	760.8	OK
1104	S	OUT	Overall Gas Plant	-287076	50.34	0.002	-599.4	0.5	OK
1105	S	OUT	Overall Gas Plant	-286416	49.88	0.001	-158.6	0.1	OK
1106	S	OUT	Overall Gas Plant	-118741	94.93	0.000	0.0	0.0	OK
1107	S	OUT	Overall Gas Plant	-132940	108.54	0.000	0.0	0.0	OK
1108	S	OUT	Overall Gas Plant	-99269	141.93	0.000	0.0	0.0	OK
1109	S	OUT	Overall Gas Plant	-266265	171.66	0.000	0.0	0.0	OK
1110	S	OUT	Overall Gas Plant	-266266	163.39	0.000	0.0	0.0	OK
1111	S	OUT	Overall Gas Plant	-267244	152.16	0.000	0.0	0.0	OK
1112	S	OUT	Overall Gas Plant	-269677	137.26	0.000	0.0	0.0	OK
C-101_P	P	IN	Overall Gas Plant					8313.4	OK
C-201_P	P	IN	Overall Gas Plant					136.5	OK
C-202_P	P	IN	Overall Gas Plant					603.1	OK
C-501_P	P	IN	Overall Gas Plant					4196.2	OK
C-502_P	P	IN	Overall Gas Plant					4027.6	OK
C-601_P	P	IN	Overall Gas Plant					2109.2	OK
C-602_P	P	IN	Overall Gas Plant					2219.4	OK
C-603_P	P	IN	Overall Gas Plant					2159.1	OK
C-604_P	P	IN	Overall Gas Plant					1846.1	OK
C-701_P	P	IN	Overall Gas Plant					6438.8	OK
C-901_P	P	IN	Overall Gas Plant					574.5	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.457	-137799.6	569.7	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.457	-137799.6	474.9	OK
1303-A	S	IN	800-CW System	-285558	53.18	15.652	-4716189.7	73.5	OK

1303-B	S	IN	800-CW System	-285558	53.18	15.652	-4716189.7	73.5	OK
830	S	IN	800-CW System	-283253	60.57	30.168	-9090005.8	3621.1	OK
1306	S	OUT	800-CW System	-284236	57.51	31.304	-9432379.3	1367.2	OK
832	S	OUT	800-CW System	-284615	56.26	30.168	-9090005.8	1035.4	OK
P-802A	P	IN	800-CW System					66.0	OK
P-802B	P	IN	800-CW System					66.0	OK
P-801	P	IN	800-CW System					163.7	OK
102	S	IN	100-Main Comp.	-146519	164.81	1.298	-295989.8	42512.4	OK
803	S	IN	100-Main Comp.	-284622	56.26	5.843	-1760425.3	161.7	OK
107	S	OUT	100-Main Comp.	-149531	155.29	1.373	-317311.2	48780.4	OK
804	S	OUT	100-Main Comp.	-283068	61.15	5.843	-1760425.3	775.2	OK
1101	S	OUT	100-Main Comp.	-285676	53.53	0.001	-175.4	0.0	OK
214	S	IN	100-Main Comp.	-191775	162.13	0.076	-21496.0	3204.2	OK
C-101_P	P	IN	100-Main Comp.					8313.4	OK
404	S	IN	500-Export Comp.	-96657	155.92	0.760	-137929.7	29361.0	OK
813	S	IN	500-Export Comp.	-284622	56.26	3.392	-1022120.6	93.9	OK
815	S	IN	500-Export Comp.	-284622	56.26	3.648	-1099152.0	101.0	OK
510	S	OUT	500-Export Comp.	-100232	134.01	0.760	-137929.7	31574.5	OK
814	S	OUT	500-Export Comp.	-283068	61.15	3.392	-1022120.6	450.1	OK
816	S	OUT	500-Export Comp.	-283068	61.15	3.648	-1099152.0	484.0	OK
1107	S	OUT	500-Export Comp.	-132940	108.54	0.000	0.0	0.0	OK
1108	S	OUT	500-Export Comp.	-99269	141.93	0.000	0.0	0.0	OK
C-501_P	P	IN	500-Export Comp.					4196.2	OK
C-502_P	P	IN	500-Export Comp.					4027.6	OK
403	S	IN	600-CO2 Comp.	-266339	171.41	0.414	-140865.2	9558.5	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	1.290	-388600.7	35.7	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	1.586	-477876.2	43.9	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.899	-572142.2	52.6	OK
823	S	IN	600-CO2 Comp.	-284622	56.26	2.167	-652824.6	60.0	OK
620	S	OUT	600-CO2 Comp.	-272265	123.61	0.414	-140865.2	12966.2	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	1.290	-388600.7	171.1	OK

820	S	OUT	600-CO2 Comp.	-283068	61.15	1.586	-477876.2	210.4	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.899	-572142.2	252.0	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	2.167	-652824.6	287.5	OK
1109	S	OUT	600-CO2 Comp.	-266265	171.66	0.000	0.0	0.0	OK
1110	S	OUT	600-CO2 Comp.	-266266	163.39	0.000	0.0	0.0	OK
1111	S	OUT	600-CO2 Comp.	-267244	152.16	0.000	0.0	0.0	OK
1112	S	OUT	600-CO2 Comp.	-269677	137.26	0.000	0.0	0.0	OK
C-601_P	P	IN	600-CO2 Comp.					2109.2	OK
C-602_P	P	IN	600-CO2 Comp.					2219.4	OK
C-603_P	P	IN	600-CO2 Comp.					2159.1	OK
C-604_P	P	IN	600-CO2 Comp.					1846.1	OK
620	S	IN	700-EOR Comp.	-272265	123.61	0.414	-140865.2	12966.2	OK
825	S	IN	700-EOR Comp.	-284622	56.26	4.376	-1318543.4	121.1	OK
705	S	OUT	700-EOR Comp.	-273140	115.63	0.414	-140865.2	13583.2	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	4.376	-1318543.4	580.6	OK
C-701_P	P	IN	700-EOR Comp.					6438.8	OK
307	S	IN	400-MP Unit	-149330	155.00	1.341	-309048.3	47254.1	OK
401	S	OUT	400-MP Unit	-96657	155.92	0.927	-168183.1	35801.0	OK
403	S	OUT	400-MP Unit	-266339	171.41	0.414	-140865.2	9558.5	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.457	-137799.6	569.7	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.457	-137799.6	474.9	OK
901	S	IN	900-C3 Cycle	-106648	141.75	0.069	-14744.1	4416.2	OK
827	S	IN	900-C3 Cycle	-284622	56.26	1.703	-513218.8	47.1	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.069	-14744.1	4500.5	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	1.703	-513218.8	117.1	OK
C-901_P	P	IN	900-C3 Cycle					574.5	OK
201	S	IN	200-VRU	-178368	183.66	0.016	-4408.1	637.2	OK
208	S	IN	200-VRU	-204760	176.13	0.046	-13722.7	1820.7	OK
317	S	IN	200-VRU	-171930	112.91	0.030	-7506.0	1347.1	OK

805	S	IN	200-VRU	-284622	56.26	0.040	-12180.4	1.1	OK
807	S	IN	200-VRU	-284622	56.26	0.103	-31057.8	2.9	OK
809	S	IN	200-VRU	-284622	56.26	0.132	-39764.9	3.7	OK
811	S	IN	200-VRU	-284622	56.26	0.340	-102381.6	9.4	OK
806	S	OUT	200-VRU	-284235	57.51	0.040	-12180.4	1.8	OK
808	S	OUT	200-VRU	-283067	61.15	0.103	-31057.8	13.8	OK
810	S	OUT	200-VRU	-283846	58.74	0.132	-39764.9	9.1	OK
812	S	OUT	200-VRU	-283068	61.15	0.340	-102381.6	45.1	OK
214	S	OUT	200-VRU	-191775	162.13	0.076	-21496.0	3204.2	OK
1102	S	OUT	200-VRU	-284732	56.03	0.000	-99.7	0.0	OK
1103	S	OUT	200-VRU	-180790	86.07	0.016	-4041.0	760.8	OK
C-201_P	P	IN	200-VRU					136.5	OK
C-202_P	P	IN	200-VRU					603.1	OK
303	S	IN	300-HCDP Adj.	-150800	150.26	1.359	-313432.5	48117.3	OK
305	S	OUT	300-HCDP Adj.	-151092	148.89	1.341	-309048.3	47317.6	OK
312	S	OUT	300-HCDP Adj.	-171059	107.28	0.018	-4384.1	818.0	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.069	-14744.1	4500.5	OK
901	S	OUT	300-HCDP Adj.	-106648	141.75	0.069	-14744.1	4416.2	OK
801	S	OUT	Oil Plant	-284622	56.26	3.650	-1099717.2	101.0	OK
802	S	IN	Oil Plant	-284234	57.51	3.650	-1099717.2	163.3	OK

**Table B3.1.9. Streams exergy flow - Single-shaft compressors – 25% FPSO gas load – RER-2**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \mu_k^0$ <i>Flow (kW)</i>	$H-T_0S + P_0V - \sum N_k \mu_k^0$ <i>Flow (kW)</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-193477	164.61	0.581	-158649.3	18014.9	OK
201	S	IN	Overall Gas Plant	-225649	181.98	0.006	-1844.7	200.8	OK
208	S	IN	Overall Gas Plant	-254181	174.38	0.019	-6562.0	665.8	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.658	-1704929.6	156.6	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.006	-1714.2	0.2	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.094	-28201.7	2.6	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.046	-13809.6	1.3	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.338	-101853.6	9.4	OK
813	S	IN	Overall Gas Plant	-284622	56.26	2.702	-814069.2	74.8	OK
815	S	IN	Overall Gas Plant	-284622	56.26	2.584	-778458.4	71.5	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.261	-380016.5	34.9	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.523	-458840.6	42.1	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.702	-512874.3	47.1	OK
823	S	IN	Overall Gas Plant	-284622	56.26	1.749	-526925.7	48.4	OK
825	S	IN	Overall Gas Plant	-284622	56.26	4.031	-1214669.5	111.6	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.730	-521244.5	47.9	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.658	-1704929.6	750.8	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.006	-1714.2	0.2	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.094	-28201.7	12.5	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.046	-13809.6	3.2	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.338	-101853.6	44.9	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	2.702	-814069.2	358.5	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	2.584	-778458.4	342.8	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.261	-380016.5	167.3	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.523	-458840.6	202.1	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.702	-512874.3	225.9	OK
824	S	OUT	Overall Gas Plant	-283068	61.15	1.749	-526925.7	232.0	OK

826	S	OUT	Overall Gas Plant	-283068	61.15	4.031	-1214669.5	534.9	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.730	-521244.5	119.0	OK
510	S	OUT	Overall Gas Plant	-100257	134.26	0.145	-26339.0	6042.8	OK
402	S	OUT	Overall Gas Plant	-97821	154.85	0.158	-28921.2	6206.7	OK
705	S	OUT	Overall Gas Plant	-309848	111.68	0.289	-108476.8	9260.3	OK
1101	S	OUT	Overall Gas Plant	-286104	52.78	0.001	-326.1	0.1	OK
1102	S	OUT	Overall Gas Plant	-284945	55.42	0.000	-28.3	0.0	OK
1103	S	OUT	Overall Gas Plant	-192339	87.03	0.005	-1434.0	249.0	OK
1104	S	OUT	Overall Gas Plant	-287549	50.63	0.001	-207.0	0.2	OK
1105	S	OUT	Overall Gas Plant	-286423	49.85	0.000	-80.9	0.0	OK
1106	S	OUT	Overall Gas Plant	-83662	201.20	0.000	0.0	0.0	OK
1107	S	OUT	Overall Gas Plant	-130193	106.31	0.006	-1246.0	303.1	OK
1108	S	OUT	Overall Gas Plant	-100086	139.52	0.000	0.0	0.0	OK
1109	S	OUT	Overall Gas Plant	-302035	169.53	0.000	0.0	0.0	OK
1110	S	OUT	Overall Gas Plant	-302152	160.91	0.000	0.0	0.0	OK
1111	S	OUT	Overall Gas Plant	-303646	147.99	0.000	0.0	0.0	OK
1112	S	OUT	Overall Gas Plant	-306947	130.49	0.000	0.0	0.0	OK
C-101_P	P	IN	Overall Gas Plant					8161.0	OK
C-201_P	P	IN	Overall Gas Plant					138.2	OK
C-202_P	P	IN	Overall Gas Plant					612.2	OK
C-501_P	P	IN	Overall Gas Plant					3970.1	OK
C-502_P	P	IN	Overall Gas Plant					3702.3	OK
C-601_P	P	IN	Overall Gas Plant					2091.4	OK
C-602_P	P	IN	Overall Gas Plant					2174.0	OK
C-603_P	P	IN	Overall Gas Plant					1979.2	OK
C-604_P	P	IN	Overall Gas Plant					1484.6	OK
C-701_P	P	IN	Overall Gas Plant					6032.8	OK
C-901_P	P	IN	Overall Gas Plant					1005.6	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.208	-62806.1	259.7	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.208	-62806.1	216.5	OK
1303-A	S	IN	800-CW System	-285558	53.18	13.431	-4046874.6	63.6	OK

1303-B	S	IN	800-CW System	-285558	53.18	13.431	-4046874.6	63.6	OK
830	S	IN	800-CW System	-283160	60.86	24.248	-7306298.2	3123.6	OK
1306	S	OUT	800-CW System	-284235	57.51	26.862	-8093749.1	1174.0	OK
832	S	OUT	800-CW System	-284615	56.26	24.248	-7306298.2	832.2	OK
P-802A	P	IN	800-CW System					56.0	OK
P-802B	P	IN	800-CW System					56.0	OK
P-801	P	IN	800-CW System					126.5	OK
102	S	IN	100-Main Comp.	-193477	164.61	0.581	-158649.3	18014.9	OK
803	S	IN	100-Main Comp.	-284622	56.26	5.658	-1704929.6	156.6	OK
107	S	OUT	100-Main Comp.	-196887	153.66	0.614	-169510.5	20672.1	OK
804	S	OUT	100-Main Comp.	-283068	61.15	5.658	-1704929.6	750.8	OK
1101	S	OUT	100-Main Comp.	-286104	52.78	0.001	-326.1	0.1	OK
214	S	IN	100-Main Comp.	-238694	160.88	0.035	-11187.4	1302.1	OK
C-101_P	P	IN	100-Main Comp.					8161.0	OK
404	S	IN	500-Export Comp.	-97821	154.85	0.151	-27585.0	5919.9	OK
813	S	IN	500-Export Comp.	-284622	56.26	2.702	-814069.2	74.8	OK
815	S	IN	500-Export Comp.	-284622	56.26	2.584	-778458.4	71.5	OK
510	S	OUT	500-Export Comp.	-100257	134.26	0.145	-26339.0	6042.8	OK
814	S	OUT	500-Export Comp.	-283068	61.15	2.702	-814069.2	358.5	OK
816	S	OUT	500-Export Comp.	-283068	61.15	2.584	-778458.4	342.8	OK
1107	S	OUT	500-Export Comp.	-130193	106.31	0.006	-1246.0	303.1	OK
1108	S	OUT	500-Export Comp.	-100086	139.52	0.000	0.0	0.0	OK
C-501_P	P	IN	500-Export Comp.					3970.1	OK
C-502_P	P	IN	500-Export Comp.					3702.3	OK
403	S	IN	600-CO2 Comp.	-302265	168.76	0.289	-108477.0	6563.7	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	1.261	-380016.5	34.9	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	1.523	-458840.6	42.1	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.702	-512874.3	47.1	OK
823	S	IN	600-CO2 Comp.	-284622	56.26	1.749	-526925.7	48.4	OK
620	S	OUT	600-CO2 Comp.	-309043	119.18	0.289	-108477.0	8851.2	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	1.261	-380016.5	167.3	OK

820	S	OUT	600-CO2 Comp.	-283068	61.15	1.523	-458840.6	202.1	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.702	-512874.3	225.9	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	1.749	-526925.7	232.0	OK
1109	S	OUT	600-CO2 Comp.	-302035	169.53	0.000	0.0	0.0	OK
1110	S	OUT	600-CO2 Comp.	-302152	160.91	0.000	0.0	0.0	OK
1111	S	OUT	600-CO2 Comp.	-303646	147.99	0.000	0.0	0.0	OK
1112	S	OUT	600-CO2 Comp.	-306947	130.49	0.000	0.0	0.0	OK
C-601_P	P	IN	600-CO2 Comp.					2091.4	OK
C-602_P	P	IN	600-CO2 Comp.					2174.0	OK
C-603_P	P	IN	600-CO2 Comp.					1979.2	OK
C-604_P	P	IN	600-CO2 Comp.					1484.6	OK
620	S	IN	700-EOR Comp.	-309043	119.18	0.289	-108477.0	8851.2	OK
825	S	IN	700-EOR Comp.	-284622	56.26	4.031	-1214669.5	111.6	OK
705	S	OUT	700-EOR Comp.	-309848	111.68	0.289	-108476.8	9260.3	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	4.031	-1214669.5	534.9	OK
C-701_P	P	IN	700-EOR Comp.					6032.8	OK
307	S	IN	400-MP Unit	-196983	153.38	0.598	-164983.2	19974.9	OK
401	S	OUT	400-MP Unit	-97821	154.85	0.309	-56506.1	12126.6	OK
403	S	OUT	400-MP Unit	-302265	168.76	0.289	-108477.0	6563.7	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.208	-62806.1	259.7	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.208	-62806.1	216.5	OK
901	S	IN	900-C3 Cycle	-106645	141.76	0.031	-6656.2	1993.7	OK
827	S	IN	900-C3 Cycle	-284622	56.26	1.730	-521244.5	47.9	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.031	-6656.2	2031.8	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	1.730	-521244.5	119.0	OK
C-901_P	P	IN	900-C3 Cycle					1005.6	OK
201	S	IN	200-VRU	-225649	181.98	0.006	-1844.7	200.8	OK
208	S	IN	200-VRU	-254181	174.38	0.019	-6562.0	665.8	OK
317	S	IN	200-VRU	-209972	113.61	0.015	-4243.0	615.7	OK

805	S	IN	200-VRU	-284622	56.26	0.006	-1714.2	0.2	OK
807	S	IN	200-VRU	-284622	56.26	0.094	-28201.7	2.6	OK
809	S	IN	200-VRU	-284622	56.26	0.046	-13809.6	1.3	OK
811	S	IN	200-VRU	-284622	56.26	0.338	-101853.6	9.4	OK
806	S	OUT	200-VRU	-284235	57.51	0.006	-1714.2	0.2	OK
808	S	OUT	200-VRU	-283067	61.15	0.094	-28201.7	12.5	OK
810	S	OUT	200-VRU	-283846	58.74	0.046	-13809.6	3.2	OK
812	S	OUT	200-VRU	-283068	61.15	0.338	-101853.6	44.9	OK
214	S	OUT	200-VRU	-238694	160.88	0.035	-11187.4	1302.1	OK
1102	S	OUT	200-VRU	-284945	55.42	0.000	-28.3	0.0	OK
1103	S	OUT	200-VRU	-192339	87.03	0.005	-1434.0	249.0	OK
C-201_P	P	IN	200-VRU					138.2	OK
C-202_P	P	IN	200-VRU					612.2	OK
303	S	IN	300-HCDP Adj.	-198411	148.49	0.606	-167190.5	20325.2	OK
305	S	OUT	300-HCDP Adj.	-198808	147.05	0.598	-164983.2	20004.0	OK
312	S	OUT	300-HCDP Adj.	-211405	107.91	0.008	-2207.3	330.1	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.031	-6656.2	2031.8	OK
901	S	OUT	300-HCDP Adj.	-106645	141.76	0.031	-6656.2	1993.7	OK
801	S	IN	Oil Plant	-284622	56.26	0.825	-248690.8	22.8	OK
802	S	OUT	Oil Plant	-284234	57.51	0.825	-248690.8	36.9	OK

**Table B3.1.10. Streams exergy flow - Multiple paralleled compressors – 100% FPSO gas load – RER-2**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \mu_k^0$ <i>(kW)</i>	$\frac{H-T_0S + P_0V - \sum N_k \mu_k^0}{N_k \mu_k^0}$ <i>(kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-123689	163.76	2.173	-446929.1	72828.2	-105369	OK
201	S	IN	Overall Gas Plant	-156362	183.40	0.030	-7474.2	1233.6	-1609	OK
208	S	IN	Overall Gas Plant	-176963	176.71	0.098	-26693.3	4154.1	-5144	OK
803	S	IN	Overall Gas Plant	-284622	56.26	5.394	-1625184.2	149.3	-89868	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.109	-32694.6	3.0	-1808	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.386	-116320.6	10.7	-6432	OK
813	S	IN	Overall Gas Plant	-284622	56.26	4.690	-1413300.6	129.8	-78152	OK
815	S	IN	Overall Gas Plant	-284622	56.26	5.318	-1602345.2	147.2	-88605	OK
817	S	IN	Overall Gas Plant	-284622	56.26	1.048	-315688.3	29.0	-17457	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.340	-403707.8	37.1	-22324	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.715	-516649.3	47.5	-28569	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.131	-642009.6	59.0	-35501	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.270	-382547.8	35.1	-21154	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.995	-601016.6	55.2	-33235	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	5.394	-1625184.2	715.7	97684	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.109	-32694.6	14.5	1965	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.386	-116320.6	51.2	6992	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	4.690	-1413300.6	622.4	84949	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	5.318	-1602345.2	705.6	96312	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	1.048	-315688.3	139.0	18975	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.340	-403707.8	177.8	24266	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.715	-516649.3	227.5	31054	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	2.131	-642009.6	282.7	38589	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.270	-382547.8	168.5	22994	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.995	-601016.6	137.2	34702	OK
510	S	OUT	Overall Gas Plant	-99660	134.20	1.652	-298296.2	68040.5	65645	OK
402	S	OUT	Overall Gas Plant	-96059	156.34	0.146	-26456.9	5601.9	6783	OK
705	S	OUT	Overall Gas Plant	-245893	118.14	0.474	-148965.9	15851.1	16581	OK
1101	S	OUT	Overall Gas Plant	-285880	52.60	0.000	-64.0	0.0	3	OK
1102	S	OUT	Overall Gas Plant	-284254	57.53	0.000	-27.6	0.0	2	OK
1103	S	OUT	Overall Gas Plant	-180564	87.06	0.025	-6242.4	1154.8	636	OK
1104	S	OUT	Overall Gas Plant	-286952	49.90	0.003	-789.2	0.6	39	OK
1105	S	OUT	Overall Gas Plant	-286498	49.59	0.001	-235.2	0.1	11	OK
1106	S	OUT	Overall Gas Plant	-123654	77.86	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-127302	118.83	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-97534	146.03	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-239668	172.88	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-239507	165.12	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-240046	155.27	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-241741	142.64	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					4058.8	0	OK
C-101-B_P	P	IN	Overall Gas Plant					4058.8	0	OK
C-201-A_P	P	IN	Overall Gas Plant					42.9	0	OK
C-201-B_P	P	IN	Overall Gas Plant					42.9	0	OK
C-201-C_P	P	IN	Overall Gas Plant					42.9	0	OK
C-202-A_P	P	IN	Overall Gas Plant					180.0	0	OK
C-202-B_P	P	IN	Overall Gas Plant					180.0	0	OK
C-202-C_P	P	IN	Overall Gas Plant					180.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-501-B_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-501-C_P	P	IN	Overall Gas Plant					1618.7	0	OK
C-502-A_P	P	IN	Overall Gas Plant					1583.6	0	OK
C-502-B_P	P	IN	Overall Gas Plant					1583.6	0	OK

C-502-C_P	P	IN	Overall Gas Plant					1583.6	0	OK
C-601_P	P	IN	Overall Gas Plant					1707.3	0	OK
C-602_P	P	IN	Overall Gas Plant					1824.1	0	OK
C-603_P	P	IN	Overall Gas Plant					1864.4	0	OK
C-604_P	P	IN	Overall Gas Plant					1762.3	0	OK
C-701-A_P	P	IN	Overall Gas Plant					1551.8	0	OK
C-701-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant					229.4	0	OK
C-901-B_P	P	IN	Overall Gas Plant					229.4	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.767	-231176.8	955.8	-17591	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.767	-231176.8	796.8	17131	OK
1303-A	S	IN	800-CW System	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
1303-B	S	IN	800-CW System	-285558	53.18	15.030	-4528857.0	70.8	-236717	OK
830	S	IN	800-CW System	-283266	60.53	29.249	-8813109.3	3477.9	-524333	OK
1306	S	OUT	800-CW System	-284236	57.51	30.061	-9057714.1	1313.3	512008	OK
832	S	OUT	800-CW System	-284615	56.26	29.249	-8813109.3	1003.9	487346	OK
P-802A	P	IN	800-CW System					63.1	0	OK
P-802B	P	IN	800-CW System					63.1	0	OK
P-801	P	IN	800-CW System					157.6	0	OK
102	S	IN	100-Main Comp.	-123689	163.76	2.173	-446929.1	72828.2	-105369	OK
803	S	IN	100-Main Comp.	-284622	56.26	5.394	-1625184.2	149.3	-89868	OK
107	S	OUT	100-Main Comp.	-126532	155.67	2.315	-483975.5	84343.9	106720	OK
804	S	OUT	100-Main Comp.	-283068	61.15	5.394	-1625184.2	715.7	97684	OK
1101	S	OUT	100-Main Comp.	-285880	52.60	0.000	-64.0	0.0	3	OK
214	S	IN	100-Main Comp.	-168264	160.43	0.142	-37110.4	6364.6	-6770	OK
C-101-A_P	P	IN	100-Main Comp.					4058.8	0	OK
C-101-B_P	P	IN	100-Main Comp.					4058.8	0	OK
404	S	IN	500-Export Comp.	-96059	156.34	1.652	-298296.2	63159.8	-76474	OK
813	S	IN	500-Export Comp.	-284622	56.26	4.690	-1413300.6	129.8	-78152	OK
815	S	IN	500-Export Comp.	-284622	56.26	5.318	-1602345.2	147.2	-88605	OK
510	S	OUT	500-Export Comp.	-99660	134.20	1.652	-298296.2	68040.5	65645	OK

814	S	OUT	500-Export Comp.	-283068	61.15	4.690	-1413300.6	622.4	84949	OK
816	S	OUT	500-Export Comp.	-283068	61.15	5.318	-1602345.2	705.6	96312	OK
1107	S	OUT	500-Export Comp.	-127302	118.83	0.000	0.0	0.0	0	OK
1108	S	OUT	500-Export Comp.	-97534	146.03	0.000	0.0	0.0	0	OK
C-501-A_P	P	IN	500-Export Comp.					1618.7	0	OK
C-501-B_P	P	IN	500-Export Comp.					1618.7	0	OK
C-501-C_P	P	IN	500-Export Comp.					1618.7	0	OK
C-502-A_P	P	IN	500-Export Comp.					1583.6	0	OK
C-502-B_P	P	IN	500-Export Comp.					1583.6	0	OK
C-502-C_P	P	IN	500-Export Comp.					1583.6	0	OK
403	S	IN	600-CO2 Comp.	-239667	172.88	0.474	-148965.9	11119.3	-24264	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	1.048	-315688.3	29.0	-17457	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	1.340	-403707.8	37.1	-22324	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.715	-516649.3	47.5	-28569	OK
823	S	IN	600-CO2 Comp.	-284622	56.26	2.131	-642009.6	59.0	-35501	OK
620	S	OUT	600-CO2 Comp.	-245004	126.36	0.474	-148965.9	15118.5	17735	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	1.048	-315688.3	139.0	18975	OK
820	S	OUT	600-CO2 Comp.	-283068	61.15	1.340	-403707.8	177.8	24266	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.715	-516649.3	227.5	31054	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	2.131	-642009.6	282.7	38589	OK
1109	S	OUT	600-CO2 Comp.	-239668	172.88	0.000	0.0	0.0	0	OK
1110	S	OUT	600-CO2 Comp.	-239507	165.12	0.000	0.0	0.0	0	OK
1111	S	OUT	600-CO2 Comp.	-240046	155.27	0.000	0.0	0.0	0	OK
1112	S	OUT	600-CO2 Comp.	-241741	142.64	0.000	0.0	0.0	0	OK
C-601_P	P	IN	600-CO2 Comp.					1707.3	0	OK
C-602_P	P	IN	600-CO2 Comp.					1824.1	0	OK
C-603_P	P	IN	600-CO2 Comp.					1864.4	0	OK
C-604_P	P	IN	600-CO2 Comp.					1762.3	0	OK
620	S	IN	700-EOR Comp.	-245004	126.36	0.474	-148965.9	15118.5	-17735	OK

825	S	IN	700-EOR Comp.	-284622	56.26	1.270	-382547.8	35.1	-21154	OK
705	S	OUT	700-EOR Comp.	-245893	118.14	0.474	-148965.9	15851.1	16581	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	1.270	-382547.8	168.5	22994	OK
C-701-A_P	P	IN	700-EOR Comp.					1551.8	0	OK
C-701-B_P	P	IN	700-EOR Comp.					0.0	0	OK
307	S	IN	400-MP Unit	-126285	155.39	2.272	-473719.1	82222.3	-104561	OK
401	S	OUT	400-MP Unit	-96059	156.34	1.798	-324753.2	68761.6	83257	OK
403	S	OUT	400-MP Unit	-239667	172.88	0.474	-148965.9	11119.3	24264	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.767	-231176.8	955.8	-17591	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.767	-231176.8	796.8	17131	OK
901	S	IN	900-C3 Cycle	-106651	141.74	0.101	-21491.4	6437.2	-4252	OK
827	S	IN	900-C3 Cycle	-284622	56.26	1.995	-601016.6	55.2	-33235	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.101	-21491.4	6560.1	3039	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	1.995	-601016.6	137.2	34702	OK
C-901-A_P	P	IN	900-C3 Cycle					229.4	0	OK
C-901-B_P	P	IN	900-C3 Cycle					229.4	0	OK
201	S	IN	200-VRU	-156362	183.40	0.030	-7474.2	1233.6	-1609	OK
208	S	IN	200-VRU	-176963	176.71	0.098	-26693.3	4154.1	-5144	OK
317	S	IN	200-VRU	-154625	111.92	0.039	-9212.9	1830.2	-1303	OK
805	S	IN	200-VRU	-284622	56.26	0.028	-8563.0	0.8	-474	OK
807	S	IN	200-VRU	-284622	56.26	0.109	-32694.6	3.0	-1808	OK
809	S	IN	200-VRU	-284622	56.26	0.304	-91748.2	8.4	-5073	OK
811	S	IN	200-VRU	-284622	56.26	0.386	-116320.6	10.7	-6432	OK
806	S	OUT	200-VRU	-284235	57.51	0.028	-8563.0	1.2	484	OK
808	S	OUT	200-VRU	-283067	61.15	0.109	-32694.6	14.5	1965	OK
810	S	OUT	200-VRU	-283846	58.74	0.304	-91748.2	20.9	5297	OK
812	S	OUT	200-VRU	-283068	61.15	0.386	-116320.6	51.2	6992	OK
214	S	OUT	200-VRU	-168264	160.43	0.142	-37110.4	6364.6	6770	OK
1102	S	OUT	200-VRU	-284254	57.53	0.000	-27.6	0.0	2	OK
1103	S	OUT	200-VRU	-180564	87.06	0.025	-6242.4	1154.8	636	OK

C-201-A_P	P	IN	200-VRU					42.9	0	OK
C-201-B_P	P	IN	200-VRU					42.9	0	OK
C-201-C_P	P	IN	200-VRU					42.9	0	OK
C-202-A_P	P	IN	200-VRU					180.0	0	OK
C-202-B_P	P	IN	200-VRU					180.0	0	OK
C-202-C_P	P	IN	200-VRU					180.0	0	OK
303	S	IN	300-HCDP Adj.	-127848	150.51	2.299	-480004.8	83602.2	-102475	OK
305	S	OUT	300-HCDP Adj.	-128021	149.37	2.272	-473719.1	82328.9	100511	OK
312	S	OUT	300-HCDP Adj.	-153751	106.79	0.027	-6285.7	1295.6	851	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.101	-21491.4	6560.1	-3039	OK
901	S	OUT	300-HCDP Adj.	-106651	141.74	0.101	-21491.4	6437.2	4252	OK
801	S	OUT	Oil Plant	-284622	56.26	3.522	-1061333.6	97.5	58689	OK
802	S	IN	Oil Plant	-284234	57.51	3.522	-1061333.6	157.6	-59995	OK

**Table B3.1.11. Streams exergy flow - Multiple paralleled compressors – 50% FPSO gas load – RER-2**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \mu_k^0$ <i>Flow</i> <i>(kW)</i>	$H-T_0S + P_0V - \sum N_k \mu_k^0$ <i>Flow (kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-146519	164.81	1.298	-295989.8	42512.4	-63339	OK
201	S	IN	Overall Gas Plant	-178368	183.66	0.016	-4408.1	637.2	-881	OK
208	S	IN	Overall Gas Plant	-204760	176.13	0.046	-13722.7	1820.7	-2416	OK
803	S	IN	Overall Gas Plant	-284622	56.26	3.683	-1109831.4	101.9	-61371	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.040	-12180.4	1.1	-674	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.060	-18115.0	1.7	-1002	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.132	-39764.9	3.7	-2199	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.149	-44804.8	4.1	-2478	OK
813	S	IN	Overall Gas Plant	-284622	56.26	2.219	-668756.7	61.4	-36980	OK
815	S	IN	Overall Gas Plant	-284622	56.26	2.585	-778940.0	71.6	-43073	OK
817	S	IN	Overall Gas Plant	-284622	56.26	0.882	-265819.6	24.4	-14699	OK
819	S	IN	Overall Gas Plant	-284622	56.26	1.180	-355461.3	32.7	-19656	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.547	-466125.2	42.8	-25775	OK
823	S	IN	Overall Gas Plant	-284622	56.26	1.934	-582608.8	53.5	-32217	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.057	-318397.9	29.2	-17606	OK
827	S	IN	Overall Gas Plant	-284622	56.26	1.349	-406417.2	37.3	-22474	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	3.683	-1109831.4	488.7	66708	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.040	-12180.4	1.8	689	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.060	-18115.0	8.0	1089	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.132	-39764.9	9.1	2296	OK
812	S	OUT	Overall Gas Plant	-283068	61.15	0.149	-44804.8	19.7	2693	OK
814	S	OUT	Overall Gas Plant	-283068	61.15	2.219	-668756.7	294.5	40197	OK
816	S	OUT	Overall Gas Plant	-283068	61.15	2.585	-778940.0	343.0	46819	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	0.882	-265819.6	117.1	15978	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	1.180	-355461.3	156.5	21366	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.547	-466125.2	205.3	28017	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	1.934	-582608.8	256.6	35019	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.057	-318397.9	140.2	19138	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	1.349	-406417.2	92.7	23466	OK
510	S	OUT	Overall Gas Plant	-100221	134.02	0.805	-145982.4	33415.3	31933	OK
402	S	OUT	Overall Gas Plant	-96648	155.92	0.122	-22123.9	4708.9	5630	OK
705	S	OUT	Overall Gas Plant	-273135	115.64	0.414	-140851.6	13581.2	14179	OK
1101	S	OUT	Overall Gas Plant	-285273	54.86	0.000	0.0	0.0	0	OK
1102	S	OUT	Overall Gas Plant	-284261	57.54	0.000	-54.2	0.0	3	OK
1103	S	OUT	Overall Gas Plant	-181522	85.12	0.016	-4181.3	781.7	415	OK
1104	S	OUT	Overall Gas Plant	-287081	50.33	0.003	-768.6	0.6	38	OK
1105	S	OUT	Overall Gas Plant	-286420	49.86	0.001	-158.0	0.1	8	OK
1106	S	OUT	Overall Gas Plant	-123639	77.92	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-129428	116.12	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	-98082	145.88	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-266336	171.41	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-266086	163.95	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-266675	154.00	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-268594	140.81	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					2482.2	0	OK
C-101-B_P	P	IN	Overall Gas Plant					2482.2	0	OK
C-201-A_P	P	IN	Overall Gas Plant					34.9	0	OK
C-201-B_P	P	IN	Overall Gas Plant					34.9	0	OK
C-201-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-A_P	P	IN	Overall Gas Plant					139.7	0	OK
C-202-B_P	P	IN	Overall Gas Plant					139.7	0	OK
C-202-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					1156.2	0	OK
C-501-B_P	P	IN	Overall Gas Plant					1156.2	0	OK
C-501-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-A_P	P	IN	Overall Gas Plant					1140.3	0	OK
C-502-B_P	P	IN	Overall Gas Plant					1140.3	0	OK

C-502-C_P	P	IN	Overall Gas Plant						0.0	0	OK
C-601_P	P	IN	Overall Gas Plant						1476.1	0	OK
C-602_P	P	IN	Overall Gas Plant						1587.8	0	OK
C-603_P	P	IN	Overall Gas Plant						1612.1	0	OK
C-604_P	P	IN	Overall Gas Plant						1484.2	0	OK
C-701-A_P	P	IN	Overall Gas Plant						1280.5	0	OK
C-701-B_P	P	IN	Overall Gas Plant						0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant						155.1	0	OK
C-901-B_P	P	IN	Overall Gas Plant						155.1	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.457	-137710.3		569.4	-10479	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.457	-137710.3		474.6	10205	OK
1303-A	S	IN	800-CW System	-285557	53.18	10.021	-3019416.3		48.0	-157824	OK
1303-B	S	IN	800-CW System	-285557	53.18	10.021	-3019416.3		48.0	-157824	OK
830	S	IN	800-CW System	-283330	60.33	20.467	-6166940.4		2321.9	-365703	OK
1306	S	OUT	800-CW System	-284236	57.51	20.042	-6038832.6		875.7	341360	OK
832	S	OUT	800-CW System	-284615	56.26	20.467	-6166940.4		702.4	341017	OK
P-802A	P	IN	800-CW System						41.2	0	OK
P-802B	P	IN	800-CW System						41.2	0	OK
P-801	P	IN	800-CW System						104.2	0	OK
102	S	IN	100-Main Comp.	-146519	164.81	1.298	-295989.8		42512.4	-63339	OK
803	S	IN	100-Main Comp.	-284622	56.26	3.683	-1109831.4		101.9	-61371	OK
107	S	OUT	100-Main Comp.	-149576	155.28	1.372	-316995.0		48685.0	63092	OK
804	S	OUT	100-Main Comp.	-283068	61.15	3.683	-1109831.4		488.7	66708	OK
1101	S	OUT	100-Main Comp.	-285273	54.86	0.000	0.0		0.0	0	OK
214	S	IN	100-Main Comp.	-192756	162.62	0.074	-21005.3		3108.8	-3578	OK
C-101-A_P	P	IN	100-Main Comp.						2482.2	0	OK
C-101-B_P	P	IN	100-Main Comp.						2482.2	0	OK
404	S	IN	500-Export Comp.	-96648	155.92	0.805	-145982.4		31071.1	-37152	OK
813	S	IN	500-Export Comp.	-284622	56.26	2.219	-668756.7		61.4	-36980	OK
815	S	IN	500-Export Comp.	-284622	56.26	2.585	-778940.0		71.6	-43073	OK
510	S	OUT	500-Export Comp.	-100221	134.02	0.805	-145982.4		33415.3	31933	OK

814	S	OUT	500-Export Comp.	-283068	61.15	2.219	-668756.7	294.5	40197	OK
816	S	OUT	500-Export Comp.	-283068	61.15	2.585	-778940.0	343.0	46819	OK
1107	S	OUT	500-Export Comp.	-129428	116.12	0.000	0.0	0.0	0	OK
1108	S	OUT	500-Export Comp.	-98082	145.88	0.000	0.0	0.0	0	OK
C-501-A_P	P	IN	500-Export Comp.					1156.2	0	OK
C-501-B_P	P	IN	500-Export Comp.					1156.2	0	OK
C-501-C_P	P	IN	500-Export Comp.					0.0	0	OK
C-502-A_P	P	IN	500-Export Comp.					1140.3	0	OK
C-502-B_P	P	IN	500-Export Comp.					1140.3	0	OK
C-502-C_P	P	IN	500-Export Comp.					0.0	0	OK
403	S	IN	600-CO2 Comp.	-266335	171.41	0.414	-140851.6	9557.7	-21018	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	0.882	-265819.6	24.4	-14699	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	1.180	-355461.3	32.7	-19656	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.547	-466125.2	42.8	-25775	OK
823	S	IN	600-CO2 Comp.	-284622	56.26	1.934	-582608.8	53.5	-32217	OK
620	S	OUT	600-CO2 Comp.	-272261	123.61	0.414	-140851.6	12965.4	15157	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	0.882	-265819.6	117.1	15978	OK
820	S	OUT	600-CO2 Comp.	-283068	61.15	1.180	-355461.3	156.5	21366	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.547	-466125.2	205.3	28017	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	1.934	-582608.8	256.6	35019	OK
1109	S	OUT	600-CO2 Comp.	-266336	171.41	0.000	0.0	0.0	0	OK
1110	S	OUT	600-CO2 Comp.	-266086	163.95	0.000	0.0	0.0	0	OK
1111	S	OUT	600-CO2 Comp.	-266675	154.00	0.000	0.0	0.0	0	OK
1112	S	OUT	600-CO2 Comp.	-268594	140.81	0.000	0.0	0.0	0	OK
C-601_P	P	IN	600-CO2 Comp.					1476.1	0	OK
C-602_P	P	IN	600-CO2 Comp.					1587.8	0	OK
C-603_P	P	IN	600-CO2 Comp.					1612.1	0	OK
C-604_P	P	IN	600-CO2 Comp.					1484.2	0	OK
620	S	IN	700-EOR Comp.	-272261	123.61	0.414	-140851.6	12965.4	-15157	OK

825	S	IN	700-EOR Comp.	-284622	56.26	1.057	-318397.9	29.2	-17606	OK
705	S	OUT	700-EOR Comp.	-273135	115.64	0.414	-140851.6	13581.2	14179	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	1.057	-318397.9	140.2	19138	OK
C-701-A_P	P	IN	700-EOR Comp.					1280.5	0	OK
C-701-B_P	P	IN	700-EOR Comp.					0.0	0	OK
307	S	IN	400-MP Unit	-149333	155.01	1.341	-308957.9	47232.4	-61537	OK
401	S	OUT	400-MP Unit	-96648	155.92	0.926	-168106.3	35780.0	42782	OK
403	S	OUT	400-MP Unit	-266335	171.41	0.414	-140851.6	9557.7	21018	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.457	-137710.3	569.4	-10479	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.457	-137710.3	474.6	10205	OK
901	S	IN	900-C3 Cycle	-106641	141.78	0.068	-14519.8	4348.9	-2873	OK
827	S	IN	900-C3 Cycle	-284622	56.26	1.349	-406417.2	37.3	-22474	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.068	-14519.8	4432.0	2053	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	1.349	-406417.2	92.7	23466	OK
C-901-A_P	P	IN	900-C3 Cycle					155.1	0	OK
C-901-B_P	P	IN	900-C3 Cycle					155.1	0	OK
201	S	IN	200-VRU	-178368	183.66	0.016	-4408.1	637.2	-881	OK
208	S	IN	200-VRU	-204760	176.13	0.046	-13722.7	1820.7	-2416	OK
317	S	IN	200-VRU	-172156	112.94	0.028	-7109.9	1274.2	-949	OK
805	S	IN	200-VRU	-284622	56.26	0.040	-12180.4	1.1	-674	OK
807	S	IN	200-VRU	-284622	56.26	0.060	-18115.0	1.7	-1002	OK
809	S	IN	200-VRU	-284622	56.26	0.132	-39764.9	3.7	-2199	OK
811	S	IN	200-VRU	-284622	56.26	0.149	-44804.8	4.1	-2478	OK
806	S	OUT	200-VRU	-284235	57.51	0.040	-12180.4	1.8	689	OK
808	S	OUT	200-VRU	-283067	61.15	0.060	-18115.0	8.0	1089	OK
810	S	OUT	200-VRU	-283846	58.74	0.132	-39764.9	9.1	2296	OK
812	S	OUT	200-VRU	-283068	61.15	0.149	-44804.8	19.7	2693	OK
214	S	OUT	200-VRU	-192756	162.62	0.074	-21005.3	3108.8	3578	OK
1102	S	OUT	200-VRU	-284261	57.54	0.000	-54.2	0.0	3	OK
1103	S	OUT	200-VRU	-181522	85.12	0.016	-4181.3	781.7	415	OK

C-201-A_P	P	IN	200-VRU					34.9	0	OK
C-201-B_P	P	IN	200-VRU					34.9	0	OK
C-201-C_P	P	IN	200-VRU					0.0	0	OK
C-202-A_P	P	IN	200-VRU					139.7	0	OK
C-202-B_P	P	IN	200-VRU					139.7	0	OK
C-202-C_P	P	IN	200-VRU					0.0	0	OK
303	S	IN	300-HCDP Adj.	-150801	150.26	1.357	-313165.1	48062.3	-60404	OK
305	S	OUT	300-HCDP Adj.	-151094	148.90	1.341	-308957.9	47295.9	59113	OK
312	S	OUT	300-HCDP Adj.	-171194	107.31	0.017	-4207.2	784.2	536	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.068	-14519.8	4432.0	-2053	OK
901	S	OUT	300-HCDP Adj.	-106641	141.78	0.068	-14519.8	4348.9	2873	OK
801	S	OUT	Oil Plant	-284622	56.26	3.650	-1099717.2	101.0	60811	OK
802	S	IN	Oil Plant	-284234	57.51	3.650	-1099717.2	163.3	-62164	OK

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**Table B3.1.12. Streams exergy flow- Multiple paralleled compressors – 25% FPSO gas load – RER-2**

<i>Stream</i>	<i>Stream Type</i> <i>S/P</i>	<i>Direction</i> <i>In/Out</i>	<i>System</i>	<i>H</i> <i>(kJ/kgmol)</i>	<i>S</i> <i>(kJ/kgmol.K)</i>	<i>Flow</i> <i>(kgmol/s)</i>	$\sum N_k \mu_k^0$ <i>Flow</i> <i>(kW)</i>	$\frac{H-T_0S + P_0V - \sum N_k \mu_k^0}{N_k \mu_k^0}$ <i>Flow (kW)</i>	<i>S<sub>K</sub></i> <i>(kW)</i>	<i>Check</i> <i>Ex &gt;0</i>
102	S	IN	Overall Gas Plant	-193477	164.61	0.581	-158649.3	18014.9	-28303	OK
201	S	IN	Overall Gas Plant	-225649	181.98	0.006	-1844.7	200.8	-317	OK
208	S	IN	Overall Gas Plant	-254182	174.37	0.019	-6562.0	665.8	-996	OK
803	S	IN	Overall Gas Plant	-284622	56.26	1.851	-557702.2	51.2	-30839	OK
805	S	IN	Overall Gas Plant	-284622	56.26	0.006	-1714.2	0.2	-95	OK
807	S	IN	Overall Gas Plant	-284622	56.26	0.021	-6443.1	0.6	-356	OK
809	S	IN	Overall Gas Plant	-284622	56.26	0.046	-13802.5	1.3	-763	OK
811	S	IN	Overall Gas Plant	-284622	56.26	0.056	-16854.6	1.5	-932	OK
813	S	IN	Overall Gas Plant	-284622	56.26	0.000	0.0	0.0	0	OK
815	S	IN	Overall Gas Plant	-284622	56.26	0.000	0.0	0.0	0	OK
817	S	IN	Overall Gas Plant	-284622	56.26	0.564	-170056.2	15.6	-9404	OK
819	S	IN	Overall Gas Plant	-284622	56.26	0.834	-251322.7	23.1	-13897	OK
821	S	IN	Overall Gas Plant	-284622	56.26	1.620	-488056.4	44.8	-26988	OK
823	S	IN	Overall Gas Plant	-284622	56.26	2.208	-665227.8	61.1	-36785	OK
825	S	IN	Overall Gas Plant	-284622	56.26	1.264	-380897.8	35.0	-21063	OK
827	S	IN	Overall Gas Plant	-284622	56.26	0.584	-175892.0	16.2	-9726	OK
804	S	OUT	Overall Gas Plant	-283068	61.15	1.851	-557702.2	245.6	33522	OK
806	S	OUT	Overall Gas Plant	-284235	57.51	0.006	-1714.2	0.2	97	OK
808	S	OUT	Overall Gas Plant	-283067	61.15	0.021	-6443.1	2.9	387	OK
810	S	OUT	Overall Gas Plant	-283846	58.74	0.046	-13802.5	3.1	797	OK
812	S	OUT	Overall Gas Plant	-283068	61.16	0.056	-16854.6	7.4	1013	OK
814	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
816	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
818	S	OUT	Overall Gas Plant	-283068	61.15	0.564	-170056.2	74.9	10222	OK
820	S	OUT	Overall Gas Plant	-283068	61.15	0.834	-251322.7	110.7	15106	OK
822	S	OUT	Overall Gas Plant	-283068	61.15	1.620	-488056.4	214.9	29335	OK

824	S	OUT	Overall Gas Plant	-283068	61.15	2.208	-665227.8	292.9	39985	OK
826	S	OUT	Overall Gas Plant	-283068	61.15	1.264	-380897.8	167.7	22894	OK
828	S	OUT	Overall Gas Plant	-283846	58.74	0.584	-175892.0	40.1	10156	OK
510	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
402	S	OUT	Overall Gas Plant	-97760	154.88	0.097	-17760.7	3810.3	4455	OK
705	S	OUT	Overall Gas Plant	-221506	121.21	0.500	-147073.0	18247.3	17966	OK
1101	S	OUT	Overall Gas Plant	-284838	56.72	0.000	0.0	0.0	0	OK
1102	S	OUT	Overall Gas Plant	-284275	57.55	0.000	-6.0	0.0	0	OK
1103	S	OUT	Overall Gas Plant	-192973	82.99	0.006	-1614.3	280.7	151	OK
1104	S	OUT	Overall Gas Plant	-287563	50.60	0.002	-522.5	0.6	26	OK
1105	S	OUT	Overall Gas Plant	-286432	49.82	0.000	-80.1	0.0	4	OK
1106	S	OUT	Overall Gas Plant	-123645	77.89	0.000	0.0	0.0	0	OK
1107	S	OUT	Overall Gas Plant	-134540	111.38	0.000	0.0	0.0	0	OK
1108	S	OUT	Overall Gas Plant	0	0.00	0.000	0.0	0.0	0	OK
1109	S	OUT	Overall Gas Plant	-302216	168.76	0.000	0.0	0.0	0	OK
1110	S	OUT	Overall Gas Plant	-301763	162.00	0.000	0.0	0.0	0	OK
1111	S	OUT	Overall Gas Plant	-219786	118.38	0.000	0.0	0.0	0	OK
1112	S	OUT	Overall Gas Plant	-217511	145.10	0.000	0.0	0.0	0	OK
C-101-A_P	P	IN	Overall Gas Plant					2269.6	0	OK
C-101-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-201-A_P	P	IN	Overall Gas Plant					26.0	0	OK
C-201-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-201-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-A_P	P	IN	Overall Gas Plant					117.8	0	OK
C-202-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-202-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-A_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-501-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-A_P	P	IN	Overall Gas Plant					0.0	0	OK
C-502-B_P	P	IN	Overall Gas Plant					0.0	0	OK

C-502-C_P	P	IN	Overall Gas Plant					0.0	0	OK
C-601_P	P	IN	Overall Gas Plant					1008.9	0	OK
C-602_P	P	IN	Overall Gas Plant					1103.6	0	OK
C-603_P	P	IN	Overall Gas Plant					1804.1	0	OK
C-604_P	P	IN	Overall Gas Plant					1755.3	0	OK
C-701-A_P	P	IN	Overall Gas Plant					1634.9	0	OK
C-701-B_P	P	IN	Overall Gas Plant					0.0	0	OK
C-901-A_P	P	IN	Overall Gas Plant					129.9	0	OK
C-901-B_P	P	IN	Overall Gas Plant					0.0	0	OK
1208	S	IN	Overall Gas Plant	-277139	77.42	0.208	-62651.6	259.0	-4767	OK
1209	S	OUT	Overall Gas Plant	-277945	75.40	0.208	-62651.6	215.9	4643	OK
1303-A	S	IN	800-CW System	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
1303-B	S	IN	800-CW System	-285556	53.18	5.284	-1592279.1	25.6	-83232	OK
830	S	IN	800-CW System	-283211	60.70	9.879	-2976660.3	1228.4	-177585	OK
1306	S	OUT	800-CW System	-284235	57.51	10.569	-3184558.2	462.0	180018	OK
832	S	OUT	800-CW System	-284616	56.26	9.879	-2976660.3	339.0	164601	OK
P-802A	P	IN	800-CW System					21.4	0	OK
P-802B	P	IN	800-CW System					21.4	0	OK
P-801	P	IN	800-CW System					48.2	0	OK
102	S	IN	100-Main Comp.	-193477	164.61	0.581	-158649.3	18014.9	-28303	OK
803	S	IN	100-Main Comp.	-284622	56.26	1.851	-557702.2	51.2	-30839	OK
107	S	OUT	100-Main Comp.	-197062	153.61	0.612	-168944.7	20529.7	27835	OK
804	S	OUT	100-Main Comp.	-283068	61.15	1.851	-557702.2	245.6	33522	OK
1101	S	OUT	100-Main Comp.	-284838	56.72	0.000	0.0	0.0	0	OK
214	S	IN	100-Main Comp.	-244174	161.29	0.031	-10295.5	1159.3	-1495	OK
C-101-A_P	P	IN	100-Main Comp.					2269.6	0	OK
C-101-B_P	P	IN	100-Main Comp.					0.0	0	OK
405	S	IN	500-Export Comp.	-97760	154.88	0.000	0.0	0.0	0	OK
813	S	IN	500-Export Comp.	-284622	56.26	0.000	0.0	0.0	0	OK
815	S	IN	500-Export Comp.	-284622	56.26	0.000	0.0	0.0	0	OK
510	S	OUT	500-Export Comp.	0	0.00	0.000	0.0	0.0	0	OK

814	S	OUT	500-Export Comp.	0	0.00	0.000	0.0	0.0	0	OK
816	S	OUT	500-Export Comp.	0	0.00	0.000	0.0	0.0	0	OK
1107	S	OUT	500-Export Comp.	-134540	111.38	0.000	0.0	0.0	0	OK
1108	S	OUT	500-Export Comp.	0	0.00	0.000	0.0	0.0	0	OK
C-501-A_P	P	IN	500-Export Comp.					0.0	0	OK
C-501-B_P	P	IN	500-Export Comp.					0.0	0	OK
C-501-C_P	P	IN	500-Export Comp.					0.0	0	OK
C-502-A_P	P	IN	500-Export Comp.					0.0	0	OK
C-502-B_P	P	IN	500-Export Comp.					0.0	0	OK
C-502-C_P	P	IN	500-Export Comp.					0.0	0	OK
403	S	IN	600-CO2 Comp.	-302215	168.76	0.289	-108459.9	6563.7	-14460	OK
406	S	IN	600-CO2 Comp.	-97760	154.88	0.211	-38613.1	8283.8	-9686	OK
817	S	IN	600-CO2 Comp.	-284622	56.26	0.564	-170056.2	15.6	-9404	OK
819	S	IN	600-CO2 Comp.	-284622	56.26	0.834	-251322.7	23.1	-13897	OK
821	S	IN	600-CO2 Comp.	-284622	56.26	1.620	-488056.4	44.8	-26988	OK
823	S	IN	600-CO2 Comp.	-284622	56.26	2.208	-665227.8	61.1	-36785	OK
620	S	OUT	600-CO2 Comp.	-220847	128.84	0.500	-147073.0	17446.0	19097	OK
818	S	OUT	600-CO2 Comp.	-283068	61.15	0.564	-170056.2	74.9	10222	OK
820	S	OUT	600-CO2 Comp.	-283068	61.15	0.834	-251322.7	110.7	15106	OK
822	S	OUT	600-CO2 Comp.	-283068	61.15	1.620	-488056.4	214.9	29335	OK
824	S	OUT	600-CO2 Comp.	-283068	61.15	2.208	-665227.8	292.9	39985	OK
1109	S	OUT	600-CO2 Comp.	-302216	168.76	0.000	0.0	0.0	0	OK
1110	S	OUT	600-CO2 Comp.	-301763	162.00	0.000	0.0	0.0	0	OK
1111	S	OUT	600-CO2 Comp.	-219786	118.38	0.000	0.0	0.0	0	OK
1112	S	OUT	600-CO2 Comp.	-217511	145.10	0.000	0.0	0.0	0	OK
C-601_P	P	IN	600-CO2 Comp.					1008.9	0	OK
C-602_P	P	IN	600-CO2 Comp.					1103.6	0	OK
C-603_P	P	IN	600-CO2 Comp.					1804.1	0	OK
C-604_P	P	IN	600-CO2 Comp.					1755.3	0	OK

620	S	IN	700-EOR Comp.	-220847	128.84	0.500	-147073.0	17446.0	-19097	OK
825	S	IN	700-EOR Comp.	-284622	56.26	1.264	-380897.8	35.0	-21063	OK
705	S	OUT	700-EOR Comp.	-221506	121.21	0.500	-147073.0	18247.3	17966	OK
826	S	OUT	700-EOR Comp.	-283068	61.15	1.264	-380897.8	167.7	22894	OK
C-701-A_P	P	IN	700-EOR Comp.					1634.9	0	OK
C-701-B_P	P	IN	700-EOR Comp.					0.0	0	OK
307	S	IN	400-MP Unit	-197022	153.39	0.598	-164833.6	19943.1	-27147	OK
401	S	OUT	400-MP Unit	-97760	154.88	0.308	-56373.7	12094.0	14141	OK
403	S	OUT	400-MP Unit	-302215	168.76	0.289	-108459.9	6563.7	14460	OK
1208	S	IN	400-MP Unit	-277139	77.42	0.208	-62651.6	259.0	-4767	OK
1209	S	OUT	400-MP Unit	-277945	75.40	0.208	-62651.6	215.9	4643	OK
901	S	IN	900-C3 Cycle	-106645	141.77	0.030	-6372.9	1908.8	-1261	OK
827	S	IN	900-C3 Cycle	-284622	56.26	0.584	-175892.0	16.2	-9726	OK
907	S	OUT	900-C3 Cycle	-117416	101.30	0.030	-6372.9	1945.3	901	OK
828	S	OUT	900-C3 Cycle	-283846	58.74	0.584	-175892.0	40.1	10156	OK
C-901-A_P	P	IN	900-C3 Cycle					129.9	0	OK
C-901-B_P	P	IN	900-C3 Cycle					0.0	0	OK
201	S	IN	200-VRU	-225649	181.98	0.006	-1844.7	200.8	-317	OK
208	S	IN	200-VRU	-254182	174.37	0.019	-6562.0	665.8	-996	OK
317	S	IN	200-VRU	-210693	113.69	0.012	-3509.1	507.2	-414	OK
805	S	IN	200-VRU	-284622	56.26	0.006	-1714.2	0.2	-95	OK
807	S	IN	200-VRU	-284622	56.26	0.021	-6443.1	0.6	-356	OK
809	S	IN	200-VRU	-284622	56.26	0.046	-13802.5	1.3	-763	OK
811	S	IN	200-VRU	-284622	56.26	0.056	-16854.6	1.5	-932	OK
806	S	OUT	200-VRU	-284235	57.51	0.006	-1714.2	0.2	97	OK
808	S	OUT	200-VRU	-283067	61.15	0.021	-6443.1	2.9	387	OK
810	S	OUT	200-VRU	-283846	58.74	0.046	-13802.5	3.1	797	OK
812	S	OUT	200-VRU	-283068	61.16	0.056	-16854.6	7.4	1013	OK
214	S	OUT	200-VRU	-244174	161.29	0.031	-10295.5	1159.3	1495	OK
1102	S	OUT	200-VRU	-284275	57.55	0.000	-6.0	0.0	0	OK

1103	S	OUT	200-VRU	-192973	82.99	0.006	-1614.3	280.7	151	OK
C-201-A_P	P	IN	200-VRU					26.0	0	OK
C-201-B_P	P	IN	200-VRU					0.0	0	OK
C-201-C_P	P	IN	200-VRU					0.0	0	OK
C-202-A_P	P	IN	200-VRU					117.8	0	OK
C-202-B_P	P	IN	200-VRU					0.0	0	OK
C-202-C_P	P	IN	200-VRU					0.0	0	OK
303	S	IN	300-HCDP Adj.	-198452	148.50	0.604	-166741.9	20248.6	-26574	OK
305	S	OUT	300-HCDP Adj.	-198844	147.08	0.598	-164833.6	19972.1	26030	OK
312	S	OUT	300-HCDP Adj.	-211834	107.98	0.007	-1908.3	284.6	213	OK
907	S	IN	300-HCDP Adj.	-117416	101.30	0.030	-6372.9	1945.3	-901	OK
901	S	OUT	300-HCDP Adj.	-106645	141.77	0.030	-6372.9	1908.8	1261	OK
801	S	OUT	Oil Plant	-284622	56.26	0.825	-248690.8	22.8	13752	OK
802	S	IN	Oil Plant	-284234	57.51	0.825	-248690.8	36.9	-14058	OK

### B3.2. Exergy balances

**Table B3.2.1. Exergy balance for single-shaft compressors (base case) – RER-1**

System	Ex <sub>in</sub> (kW)	Ex <sub>out</sub> (kW)	Ex <sub>destroyed</sub> (kW)	S <sub>generated</sub> (kW)	Error	Ex <sub>destroyed</sub>	Ex <sub>waste</sub> (kW)	Ex <sub>waste+destroyed</sub>	η <sub>ex</sub>
<b>Case SSLC - 100% Gas Load - RER I</b>									
1000-GT	291411	216366	75044	75057	-0.016%	25.8%	38374	38.9%	61.1%
800-CW System	4535	2671	1864	1865	-0.080%	41.1%	1525	74.7%	25.3%
Overall Gas Plant	2139159	2120495	18664	18873	-1.118%	0.9%		0.9%	99.1%
GAS+TG+CW	2114278	2018301	95977	95968	0.009%	4.5%	39899	6.4%	93.6%
<b>Case SSLC - 50% Gas Load - RER I</b>									
1000-GT	284161	210810	73350	73362	-0.016%	25.8%	37102	38.9%	61.1%
800-CW System	4064	2403	1661	1661	-0.011%	40.9%	1367	74.5%	25.5%
Overall Gas Plant	1187481	1164827	22654	22669	-0.064%	1.9%		1.9%	98.1%
GAS+TG+CW	1164494	1066452	98042	97857	0.189%	8.4%	38469	11.7%	88.3%
<b>Case SSLC - 25% Gas Load - RER I</b>									
1000-GT	278003	206026	71976	71988	-0.016%	25.9%	35980	38.8%	61.2%
800-CW System	3489	2006	1483	1481	0.115%	42.5%	1174	76.2%	23.8%
Overall Gas Plant	454299	428449	25849	25869	-0.076%	5.7%		5.7%	94.3%
GAS+TG+CW	432834	333226	99609	99437	0.172%	23.0%	37154	31.6%	68.4%

**Table B3.2.2 Exergy balance for Multiple paralleled compressors – RER-1**

System	Ex <sub>in</sub> (kW)	Ex <sub>out</sub> (kW)	Ex <sub>destroyed</sub> (kW)	S <sub>generated</sub> (kW)	Error	Ex <sub>destroyed</sub>	Ex <sub>waste</sub> (kW)	Ex <sub>waste+destroyed</sub>	η <sub>ex</sub>
<b>Case MSCP - 100% Gas Load - RER I</b>									
1000-GT	239425	178988	60437	60447	-0.017%	25.2%	32065	38.6%	61.4%
800-CW System	3903	2317	1586	1586	0.000%	40.6%	1265	73.0%	27.0%
Overall Gas Plant	2132556	2119657	12899	12729	1.320%	0.6%		0.6%	99.4%
GAS+TG+CW	2110713	2035477	75236	74917	0.424%	3.6%	33330	5.1%	94.9%
<b>Case MSCP - 50% Gas Load - RER I</b>									
1000-GT	206995	153817	53178	53186	-0.016%	25.7%	27148	38.8%	61.2%
800-CW System	2604	1578	1026	1026	0.000%	39.4%	849	72.0%	28.0%
Overall Gas Plant	1172246	1163666	8579	8567	0.145%	0.7%			100.0%
GAS+TG+CW	1160900	1097839	63061	62911	0.237%	5.4%	27997	7.8%	92.2%
<b>Case MSCP - 25% Gas Load - RER I</b>									
1000-GT	172213	127463	44749	44757	-0.016%	26.0%	21393	38.4%	61.6%
800-CW System	1371	801	570	570	0.000%	41.6%	452	74.5%	25.5%
Overall Gas Plant	432399	426869	5530	5526	0.077%	1.3%		1.3%	98.7%
GAS+TG+CW	429215	378176	51039	50916	0.241%	11.9%	21845	17.0%	83.0%

**Table B3.2.3. Exergy balance for single-shaft compressors (base case) – RER-2**

System	Ex <sub>in</sub> (kW)	Ex <sub>out</sub> (kW)	Ex <sub>destroyed</sub> (kW)	S <sub>generated</sub> (kW)	Error	Ex <sub>destroyed</sub>	η <sub>ex</sub>
<b>Case SSLC - 100% Gas Load - RER II</b>							
100-Main Comp.	87548	85050	2498	2498	0.01%	2.9%	97.1%
200-VRU	7976	7604	372	372	0.00%	4.7%	95.3%
300-HCDP Adj.	90110	90010	100	100	0.00%	0.1%	99.9%
900-C3 Cycle	6946	6658	288	288	0.04%	4.1%	95.9%
400-MP Unit	83179	80679	2501	2501	0.00%	3.0%	97.0%
500-Export Comp.	71586	68191	3395	3396	0.00%	4.7%	95.3%
600-CO2 Comp.	19952	16072	3880	3880	0.01%	19.4%	80.6%
700-EOR Comp.	21991	16465	5526	5526	0.00%	25.1%	74.9%
800-CW System	4535	2671	1864	1865	-0.08%	41.1%	58.9%
Overall Gas Plant	114048	95185	18863	18873	-0.05%	16.5%	83.5%
<b>Case SSLC - 50% Gas Load - RER II</b>							
100-Main Comp.	54192	49556	4636	4637	-0.02%	8.6%	91.4%
200-VRU	4562	4035	527	527	0.00%	11.6%	88.4%
300-HCDP Adj.	52618	52552	66	66	-0.05%	0.1%	99.9%
900-C3 Cycle	5038	4618	420	420	-0.01%	8.3%	91.7%
400-MP Unit	47824	45834	1989	1989	0.00%	4.2%	95.8%
500-Export Comp.	37780	32509	5271	5271	-0.01%	14.0%	86.0%
600-CO2 Comp.	18084	13887	4197	4197	0.00%	23.2%	76.8%
700-EOR Comp.	19526	14164	5362	5362	0.00%	27.5%	72.5%
800-CW System	4064	2403	1661	1661	-0.01%	40.9%	59.1%
Overall Gas Plant	78898	56232	22666	22669	-0.01%	28.7%	71.3%
<b>Case SSLC - 25% Gas Load - RER II</b>							
100-Main Comp.	27635	21423	6211	6212	0.00%	22.5%	77.5%
200-VRU	2246	1612	634	634	0.00%	28.2%	71.8%
300-HCDP Adj.	22357	22328	29	29	-0.03%	0.1%	99.9%
900-C3 Cycle	3047	2151	896	896	0.00%	29.4%	70.6%
400-MP Unit	20235	18907	1328	1328	0.00%	6.6%	93.4%
500-Export Comp.	13739	7047	6691	6691	0.00%	48.7%	51.3%
600-CO2 Comp.	14465	9678	4787	4787	0.00%	33.1%	66.9%
700-EOR Comp.	14996	9795	5200	5200	0.01%	34.7%	65.3%
800-CW System	3489	2006	1483	1481	0.11%	42.5%	57.5%
Overall Gas Plant	51141	25273	25868	25869	0.00%	50.6%	49.4%

**Table B3.2.4. Exergy balance for Multiple paralleled compressors – RER-2**

System	Ex <sub>in</sub> (kW)	Ex <sub>out</sub> (kW)	Ex <sub>destroyed</sub> (kW)	S <sub>generated</sub> (kW)	Error	Ex <sub>destroyed</sub>	η <sub>ex</sub>
<b>Case MSCP - 100% Gas Load - RER II</b>							
100-Main Comp.	87460	85060	2400	2400	0.00%	2.7%	97.3%
200-VRU	7909	7607	302	302	0.00%	3.8%	96.2%
300-HCDP Adj.	90162	90062	101	101	0.00%	0.1%	99.9%
900-C3 Cycle	6951	6697	254	254	0.00%	3.7%	96.3%
400-MP Unit	83178	80678	2500	2500	0.00%	3.0%	97.0%
500-Export Comp.	73044	69368	3675	3675	0.00%	5.0%	95.0%
600-CO2 Comp.	18450	15946	2504	2504	0.00%	13.6%	86.4%
700-EOR Comp.	16706	16020	686	686	0.00%	4.1%	95.9%
800-CW System	3903	2317	1586	1586	0.00%	40.6%	59.4%
Overall Gas Plant	107445	94710	12735	12729	0.05%	11.9%	88.1%
<b>Case MSCP - 50% Gas Load - RER II</b>							
100-Main Comp.	50688	49174	1514	1514	0.00%	3.0%	97.0%
200-VRU	4092	3929	163	163	0.00%	4.0%	96.0%
300-HCDP Adj.	52494	52429	65	65	0.00%	0.1%	99.9%
900-C3 Cycle	4696	4525	172	172	0.00%	3.7%	96.3%
400-MP Unit	47802	45812	1989	1989	0.00%	4.2%	95.8%
500-Export Comp.	35797	34053	1744	1744	0.00%	4.9%	95.1%
600-CO2 Comp.	15871	13701	2170	2170	0.00%	13.7%	86.3%
700-EOR Comp.	14275	13721	554	554	0.00%	3.9%	96.1%
800-CW System	2604	1578	1026	1026	0.00%	39.4%	60.6%
Overall Gas Plant	63663	55096	8567	8567	0.00%	13.5%	86.5%
<b>Case MSCP - 25% Gas Load - RER II</b>							
100-Main Comp.	21495	20775	720	720	0.00%	3.3%	96.7%
200-VRU	1521	1454	68	68	0.00%	4.4%	95.6%
300-HCDP Adj.	22194	22165	28	28	0.00%	0.1%	99.9%
900-C3 Cycle	2055	1985	69	69	0.00%	3.4%	96.6%
400-MP Unit	20202	18874	1329	1329	0.00%	6.6%	93.4%
500-Export Comp.	-	-	-	-	-	-	-
600-CO2 Comp.	20664	18139	2525	2525	0.00%	12.2%	87.8%
700-EOR Comp.	19116	18415	701	701	0.00%	3.7%	96.3%
800-CW System	1371	801	570	570	0.00%	41.6%	58.4%
Overall Gas Plant	29241	23715	5526	5526	0.00%	18.9%	81.1%

## References of Appendix B

Cruz, M. de A.; Araújo, O. de Q. F.; De Medeiros, J. L. Deep seawater intake for primary cooling in tropical offshore processing of natural gas with high carbon dioxide content: Energy, emissions and economic assessments. **Journal of Natural Gas Science and Engineering**, v. 56, n. June, p. 193–211, 2018.

## SUPPLEMENTARY MATERIALS C

CRUZ, M. DE A. et al. Impact of solid waste treatment from spray dryer absorber on the levelized cost of energy of a coal-fired power plant. **Journal of Cleaner Production**, v. 164, 2017a.

**Supplement C1. FBR solid product and air flows determination**

**Supplement C2. Aspen process economic analysis**

**Supplement C3. Sensitivity analysis (LCOE versus treated residue revenue price)**

### References of Supplementary Materials C

#### Supplement C1. FBR solid product and air flows determination

Table C1.1. FBR Solid Product Stream.

Component	MW (kg/mol)	Flow (mol/h)	Flow (kg/h)	% Weight
CaSO <sub>3</sub> .½H <sub>2</sub> O	1.29E-01	-	-	0.00%
CaSO <sub>4</sub> .2H <sub>2</sub> O	1.72E-01	-	-	0.00%
Ca(OH) <sub>2</sub>	7.40E-02	-	-	0.00%
CaSO <sub>3</sub>	1.20E-01	6186	742	3.66%
CaSO <sub>4</sub>	1.36E-01	55671	7571	37.34%
CaO	5.60E-02	23307	1305	6.44%
H <sub>2</sub> O*	1.80E-02	-	-	0.00%
Inert	-	-	10660	52.57%
Total			20279	100.00%

Table C1.2. FBR Air Inlet and Outlet Streams.

<b>FBR Air Inlet Stream</b>				
	MW (kg/mol)	Flow (mol/h)	Flow (kg/h)	% Molar
O <sub>2</sub>	0.032	32475	1039	21.00%
N <sub>2</sub>	0.028	122167	3421	79.00%
Air	0.029	154642	4460	100.00%
<b>FBR Air Outlet Stream</b>				
O <sub>2</sub>	0.032	4639	148	2.27%
N <sub>2</sub>	0.028	122167	3421	59.85%
H <sub>2</sub> O	0.018	77325	1392	37.88%
Air	0.024	204131	4961	100.00%

## Supplement C2. Aspen process economic analysis

Table C2.1. Aspen Process Economic Analysis Equipment Inputs.

<b>Reactor (FBR)</b>	
Vessel Diameter (mm)	101.6
Vessel Height (m)	5.08
Design Gauge Pressure (kPa)	100
Design Temperature (°C)	800
Operating Temperature (°C)	550
Design Temperature (°C)	750
Solids Volume (% of vessel's volume)	30
<b>Compressor</b>	
Actual Gas Flow Rate (m <sup>3</sup> /h)	3717.5
Design Gauge Pressure (kPa)	148.67
Design Temperature Inlet (°C)	20
<b>Air Heater</b>	
Material	SS304
Duty (MW)	0.5572
Gas Flow Rate (Sm <sup>3</sup> /h)	3466
Process Type	GAS
Design Gauge Pressure (kPa)	250
Design Temperature (°C)	550
<b>Economizer</b>	
Heat Transfer Area (m <sup>2</sup> )	17.1
Number of Shells	1
TEMA symbol	BEU
Tube Material	SS304
Tube Design Gauge Pressure (kPa)	200
Tube Operating Temperature (°C)	350
Tube Outside Diameter (mm)	19.05
Shell Material	SS304
Shell Design Gauge Pressure (kPa)	100
Shell Operating Temperature (°C)	550
Shell Diameter (mm)	406
<b>Air Filter</b>	
Gas Flow Rate (Sm <sup>3</sup> /h)	4575
Air Temperature (°C)	400
<b>Cyclones</b>	
Gas Flow Rate (Sm <sup>3</sup> /h)	4575

Table C2.2. Aspen Process Economic Analysis Financial Set Parameters.

<b>Time Period</b>		
Period Description		Year
Operating Hours per Period	Hours/period	7451
Number of Weeks per Period	Weeks/period	52
Number of Periods for Analysis	Period	30
<b>Capital Costs Parameters</b>		
Working Capital Percentage	%/period	5
<b>Operating Costs Parameters</b>		
Operating Supplies (lump-sum)	Cost/period	25
Laboratory Charges (lump-sum)	Cost/period	0
User Entered Operating Charge (%)	%/period	25
Operating Charges (% of Operating Labor Costs)	%/period	0
Plant Overhead (% of O&M Costs)	%/period	50
G and A Expenses (% of Subtotal Operating Costs)	%/period	8
<b>Operating Labor and Maintenance Costs</b>		
Operating Labour		
Operators per Shift		2
Unit Cost	Cost/Operator/H	20
Supervision		
Supervisors per Shift		1
Unit Cost	Cost/Supervisor/H	35
<b>Utilities Costs</b>		
Electricity Unit Cost	Cost/kWh	0.095
<b>General Investment Parameters</b>		
Project Type		Grass Roots
Tax Rate	%/period	40
Interest Rate (or discount rate)	%/period	8
Economic Life of Project	Period	30
Salvage Value (Fraction of Initial Capital Cost)	%	20
Depreciation Method		Straight Line
<b>Escalation</b>		
Project Capital Escalation	%/period	5
Products Escalation	%/period	5
Raw Material Escalation	%/period	3.5
Operating and Maintenance Labor Escalation	%/period	3
Utilities Escalation	%/period	3

Table C2.3. Semi-Dry FGD Treatment Unit CAPEX.

<b>Component Name</b>	<b>Total Direct Cost (USD)*</b>	<b>Equipment Cost (USD)*</b>	<b>Equipment Weight (kg)</b>	<b>Installed Weight (kg)</b>
Compressor	400,100.00	250,800.00	6,500.00	13,671.00
Reactor (FBR)	223,400.00	49,500.00	1,700.00	9,748.00
Heater	324,700.00	194,400.00	13,500.00	19,636.00
Economizer	120,800.00	19,900.00	760	6,149.00
Air Filter	29,500.00	13,800.00	1,900.00	3,294.00
Cyclones	25,300.00	14,200.00	260	735
<b>Total</b>	<b>1,123,800.00</b>	<b>542,600.00</b>	<b>24,620</b>	<b>53,233</b>

\*1<sup>st</sup> quarter 2014 price basis.

### Supplement C3. Sensitivity analysis.

Table C3.1. Sensitivity Analysis Fixed Parameters

<b>Fixed Parameters</b>	<b>Value</b>
Sales (ton/h)	20.3
Operating Hours (h/yr)	7,451
Sales (ton/yr)	151,255
Project Lifetime (yr)	30
Interest Rate (%/yr)	8
Total Utilities Cost (\$/yr)	112,257.85
Annualised Capital Cost (\$/yr)	604,384.20
O&M Fixed Cost (\$/yr)	573,261.31
AE (MWh/yr)	2,535,000
Base Plant LCOE (\$/MWh)	94.97
FGD waste disposal cost (\$/MWh)	0.53

Table C3.2. Sensitivity Analysis of LCOE versus Treated Semi-dry FGD Revenue Price

<b>Residue Price (\$/ton)</b>	<b>Annual Revenue (\$/yr)</b>	<b>Sales Profit (\$/yr)</b>	<b>TAC * (\$/yr)</b>	<b>ΔLCOE (\$/MWh)</b>	<b>FINAL LCOE (\$/MWh)</b>	<b>ΔLCOE (%)</b>
0	0.00	-112257.85	-1289903.36	0.02	94.95	0.0%
10	1512553.00	1400295.15	222649.64	0.62	94.35	0.7%
20	3025106.00	2912848.15	1735202.64	1.21	93.76	1.3%
30	4537659.00	4425401.15	3247755.64	1.81	93.16	1.9%
40	6050212.00	5937954.15	4760308.64	2.41	92.56	2.5%
47	7108999.10	6996741.25	5819095.74	2.83	92.14	3.0%
50	7562765.00	7450507.15	6272861.64	3.00	91.97	3.2%
60	9075318.00	8963060.15	7785414.64	3.60	91.37	3.8%
70	10587871.00	10475613.15	9297967.64	4.20	90.77	4.4%
80	12100424.00	11988166.15	10810520.64	4.79	90.18	5.0%
90	13612977.00	13500719.15	12323073.64	5.39	89.58	5.7%
100	15125530.00	15013272.15	13835626.64	5.99	88.98	6.3%

\* TAC = Total Annualized Cost of semi-dry FGD solid waste treatment unit = sales profit - annualized capital cost - O&M fixed cost.

# APPENDIX I – PILOT-PLANT DETAILED DESIGN

## I.1. Process Simulation

Preliminary process simulation was developed in Aspen HYSYS v8.8 (Aspentech, 2020), according to the process design. A blend of MEA/1-propanol/Water was considered as the PCAS. The Acid Gas fluid package was chosen to perform the simulation. This thermodynamic package does not predict phase split when the solvent absorbs CO<sub>2</sub>. A stream splitter module was used to force the phase-split, based on the volume ratios determined experimentally at the PCASP. The information generated in the simulation was used as a design basis for sizing and specification of the main equipment, instruments, valves and pipes. Figure I.1 shows the simulation flowsheet and Table I.1 summarizes the main simulation inputs, constrains, targets and other assumptions.

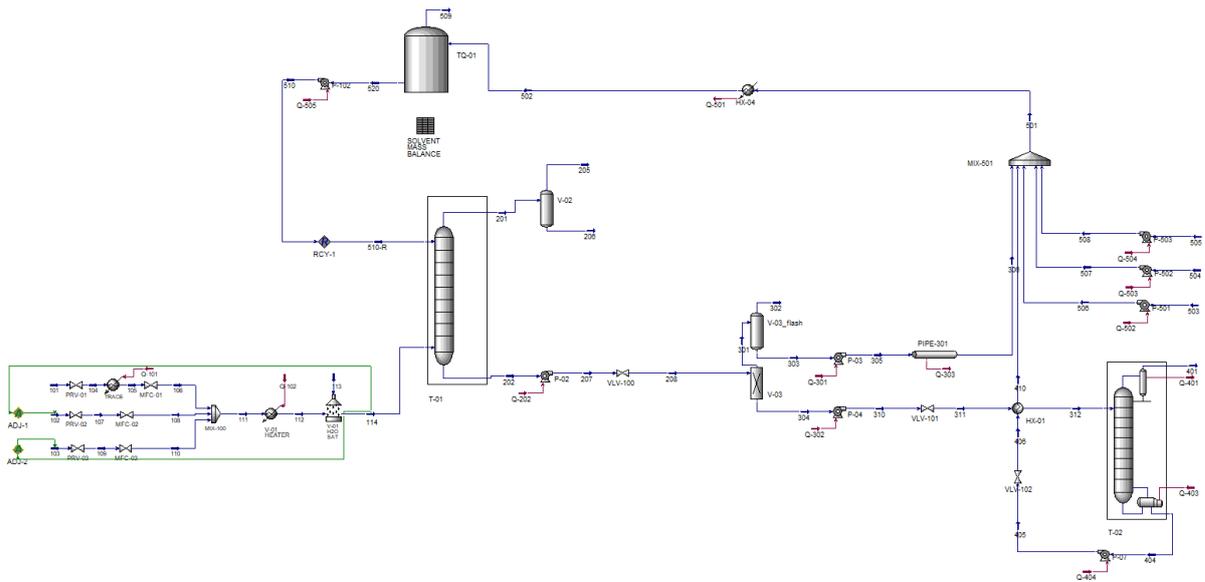


Figure I.1. PCAPP Simulation Flowsheet

**Table I.1. PCAPP Simulation assumptions**

Parameter	Value
Flue-Gas CO <sub>2</sub> flow rate	2.5 – 5.0 kg/h
Flue-Gas CO <sub>2</sub> concentration	4% - 15% (mol/mol)
Lean solvent composition	30% MEA, 40% 1-propanol (w/w)
Solvent flow rate	50 – 200 kg/h
Absorption column pressure	101-200 kPa
Absorption column internals	MELLAPAK 500Y metal, 135mm diameter, 4400mm section height
Minimum CO <sub>2</sub> capture efficiency	90%
Integration heat exchanger minimum approach	10°C
Regeneration pressure	150 – 300 kPa
Reboiler maximum temperature	130°C
Condenser temperature	40°C
Regeneration column internals	MELLAPAK 500Y metal, 108mm diameter, 4400mm section height

## I.2. Equipment Design and Specification

- **Towers**

To design the columns, it was considered a hypothetical wet flue-gas from a natural gas combined cycle (NGCC) power plant. The flue-gas CO<sub>2</sub> molar concentration was considered 4% at 200 kPa and 40°C. The flue-gas inlet flow rate is manipulated to reach a CO<sub>2</sub> flow rate of 5kg/h. The solvent flow rate was considered 100 kg/h (30% MEA, 40% 1-propanol).

The diameters of the absorption and desorption towers were defined in 5” (141mm internal diameter) and 4” (114mm internal diameter), respectively. Both columns have 2 sections of packed beds with 2.2m of height each. The column's height is limited in around 8 meters by the ceiling of the shed. The hydraulics of the towers were checked using the software KG-Tower (Koch-Glitsch, 2020). This software requires the properties summarized in Table I.2 to calculate the pressure drop through the packed beds, % of flooding, and other operational parameters. All inputs were imported from the process simulations.

**Table I.2. KG-Tower Inputs**

<b>Inputs</b>	
<b>Liquid</b> <i>(for each theoretical stage or the plate with maximum internal flow rate)</i>	Nominal, Minimum and Maximum Flow Rates (kg/h); Density (kg/m <sup>3</sup> ); Viscosity (cP); Surface Tension (dyne/cm)
<b>Vapor</b> <i>(for each theoretical stage or the plate with maximum internal flow rate)</i>	Nominal, Minimum and Maximum Flow Rates (kg/h); Density (kg/m <sup>3</sup> ); Viscosity (cP)
<b>Tower</b>	System Factor; Packing Type; Height (m), Diameter (m)

The KG-Tower reports generated for the T-01 and T-02 are depicted in Figs. I.2 and I.3.

**KOCH-GLITSCH®**  
 KG-TOWER® Software v 5.4  
 Registered To: Matheus Cruz, UFRJ

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**PACKED TOWER RATING DATA**

Project Name	UPAS2F	Date : 02-Mar-2020	Page : 1
Tower Name	T-01 Absorption Tower	File : T01-10_REVD.kgt	
Case Name	100kg/h 30MEA40PROP 5kg/h CO2	By : Matheus	Revision : 0

ZONE DESCRIPTION	Top Max Liq	Top Max Liq	Top Max Liq
BED NUMBER	3	MIN	MAX
<b>% OF LOADING</b> (%V/%L)	100	50/ 75	105/ 200
<b>LOADINGS</b>			
Vapor Rate	kg/hr	88	44
Vapor Density	kg/m3	1.829	1.829
Vapor Volume	m3/s	0.01	0.01
Vapor Viscosity	cP	0.0187	0.0187
Liquid Rate	kg/hr	103	77
Liquid Density	kg/m3	948.17	948.17
Liquid Volume	m3/hr	0.11	0.08
Surface Tension	mN/m	26.26	26.26
Liquid Viscosity	cP	1.789	1.789
<b>System Factor</b>		0.70	0.70
<b>Packing Type</b>	FLEXIPAC® 500Y structured packing METAL	FLEXIPAC® 500Y structured packing METAL	FLEXIPAC® 500Y structured packing METAL
Tower Diameter	mm	135	135
Tower Area	m2	0.01	0.01
Packing Height	mm	4400	4400
Fs	m/s*(kg/m3) <sup>0.5</sup>	1.26	0.63
Cv	m/s	0.04	0.02
Liquid Loading	m3/hr/m2	7.58	5.68
Calculated Capacity	%	92	49
Constant L/V			109
Pressure Drop	mbar/m	1.69	<0.5
Total Packing Pressure Drop	mm Hg		5.57 ( Min. 1.42 Max. N/A )
<i>Note: The total packing pressure drop is the sum of the calculated pressure drop for each loading.</i>			
<b>WARNINGS:</b>		1,	2,3,

**Figure I.2. KG-Tower Report for the T-01 Hydraulic Analysis**

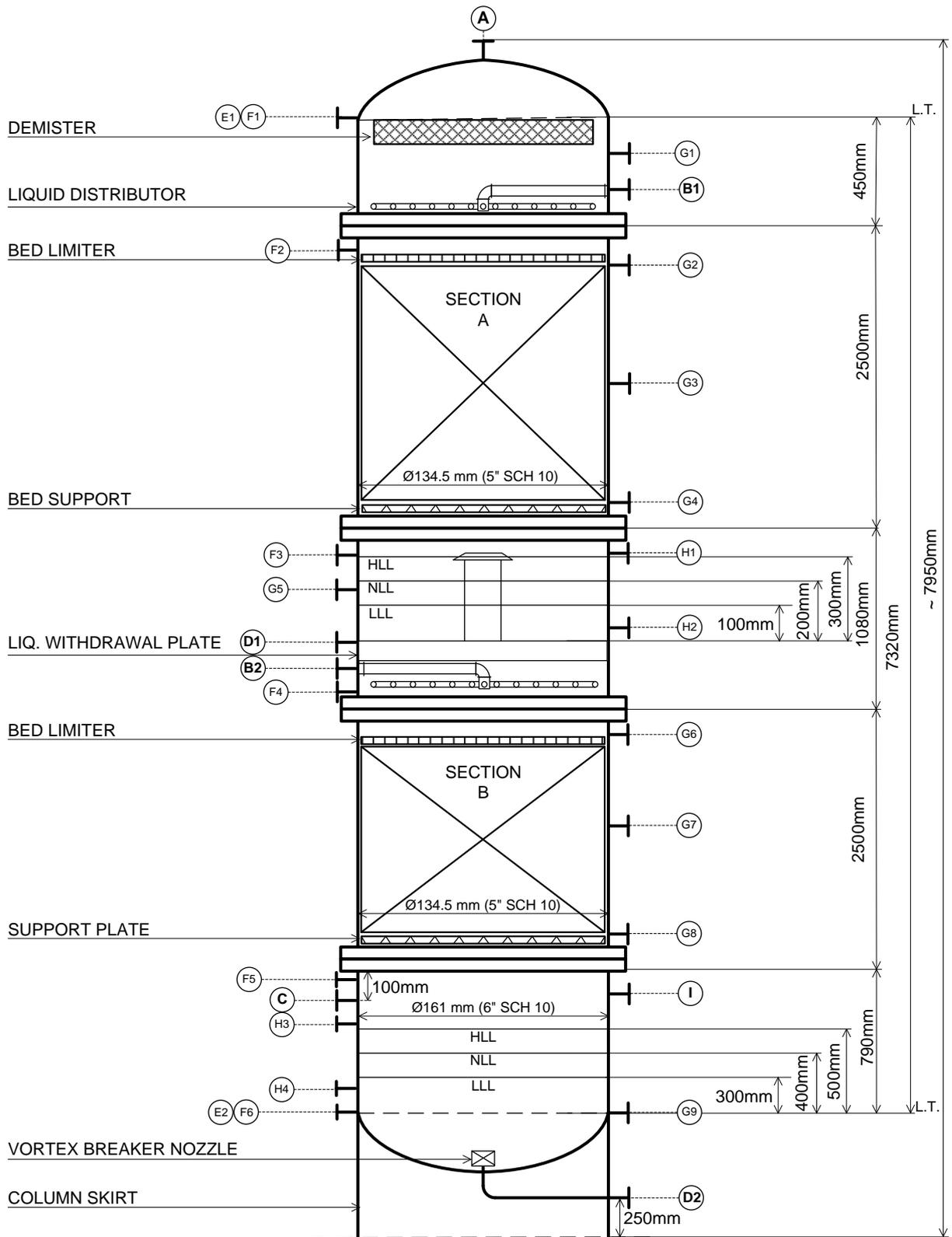
Project Name	UPAS2F	Date	02-Mar-2020	Page	1
Tower Name	T-02 Stripper	File	T02-5_REV0.kgt	By	Matheus
Case Name	100kg/h 30MEA40PROP 5kg/h CO2	Revision	0		
ZONE		Bottom	Bottom	Bottom	
DESCRIPTION		Max L&G Rates	Max L&G Rates	Max L&G Rates	
BED NUMBER		20	MIN	MAX	
% OF LOADING	(%V/%L)	100	50/ 75	105/ 200	
<b>LOADINGS</b>					
Vapor Rate	kg/hr	11	5	11	
Vapor Density	kg/m3	2.854	2.854	2.854	
Vapor Volume	m3/s	0.00	0.00	0.00	
Vapor Viscosity	cP	0.0140	0.0140	0.0140	
Liquid Rate	kg/hr	81	61	162	
Liquid Density	kg/m3	959.75	959.75	959.75	
Liquid Volume	m3/hr	0.08	0.06	0.17	
Surface Tension	mN/m	23.65	23.65	23.65	
Liquid Viscosity	cP	0.711	0.711	0.711	
System Factor		0.70	0.70	0.70	
Packing Type		FLEXIPAC® 500Y structured packing METAL	FLEXIPAC® 500Y structured packing METAL	FLEXIPAC® 500Y structured packing METAL	
Tower Diameter	mm	102	102	102	
Tower Area	m2	0.01	0.01	0.01	
Packing Height	mm	4400	4400	4400	
Fs	m/s*(kg/m3) <sup>0.5</sup>	0.22	0.11	0.23	
Cv	m/s	0.01	0.00	0.01	
Liquid Loading	m3/hr/m2	10.26	7.69	20.51	
Calculated Capacity	%	26	15	35	
Constant L/V					
Pressure Drop	mbar/m	<0.5	<0.5	<0.5	
Total Packing Pressure Drop	mm Hg		0.19	( Min. 0.05 Max. 0.23 )	

Note: The total packing pressure drop is the sum of the calculated pressure drop for each loading.

**Figure I.3. KG-Tower Report for the T-02 Hydraulic Analysis**

The % of flooding is above 90% for the T-01 at 100% CO<sub>2</sub> capacity (5kg/h). This result is above 80%, the maximum recommended by the manufacturers of tower internals. Thus, if the flooding occurs the plant should operate with a lower CO<sub>2</sub> inlet flow rate. The hydraulic analysis of T-02 unveiled that it will operate far from the flooding point because of the small internal vapor flow rates.

Figs. I.4 and I.5 present the internals and nozzles of the towers T-01 and T-02, respectively. The description of each nozzle is presented in Table I.3.



**Figure I.4. Absorber (T-01) Internals and Nozzles**



**Table I.3. Nozzles list of the absorption and desorption columns**

NOZZLE	ABSORBER (T-01)			STRIPPER (T-02)		
	QUANT.	DIAM. (in)	DESCRIPTION	QUANT.	DIAM. (in)	DESCRIPTION
<b>A</b>	1	1	Top gas outlet	1	1	Top gas outlet
<b>B</b>	2	1/2	Liquid inlet	2	1	Liquid inlet
<b>C</b>	1	1	Gas inlet	1	1/2	Liquid inlet (reflux)
<b>D</b>	2	1	Liquid outlet	2	1	Liquid outlet
<b>E</b>	2	1/4	Pressure gauge	1	1	Gas inlet
<b>F</b>	6	1/4	Pressure sensor	2	1/4	Pressure gauge
<b>G</b>	9	1/4	Temperature sensor	6	1/4	Pressure sensor
<b>H</b>	4	1/2	Level sensor	9	1/4	Temperature sensor
<b>I</b>	1	1/2	Liquid inlet	4	1/2	Level sensor
<b>J</b>	-	-	-	1	1	Liquid inlet

Fig. I.6 shows pictures of the columns and their internals.



**Figure I.6. PCAPP Columns and their internals**

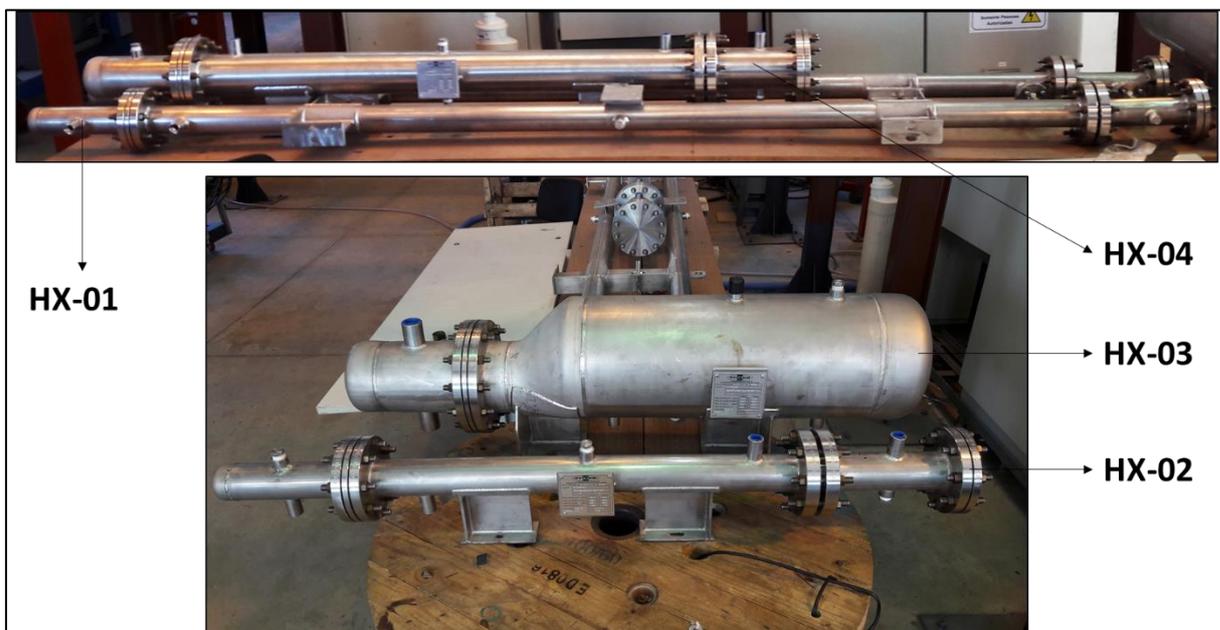
- **Heat Exchangers**

Table I.4 summarize the heat exchangers design.

**Table I.4. PCAPP Heat Exchangers**

TAG	TEMA Type	Duty (kW)	DN Shell (mm)	Tubes Length (mm)	Tubes Design T (°C)	Shell Design T (°C)	Tubes Design P (kPa)	Shell Design P (kPa)	Heat Exchange Area (m <sup>2</sup> )
HX-01	BEW	4,5	67	2500	155	155	400	500	1
HX-02	BEW	0,6	67	900	70	125	400	500	0,2
HX-03	BKU	7,0	161	600	215	155	400	500	0,8
HX-04	BEW	5,0	95	1400	105	105	400	500	0,7

Fig. I.7 shows pictures of the heat exchangers.



**Figure I.7. PCAPP Heat Exchangers**

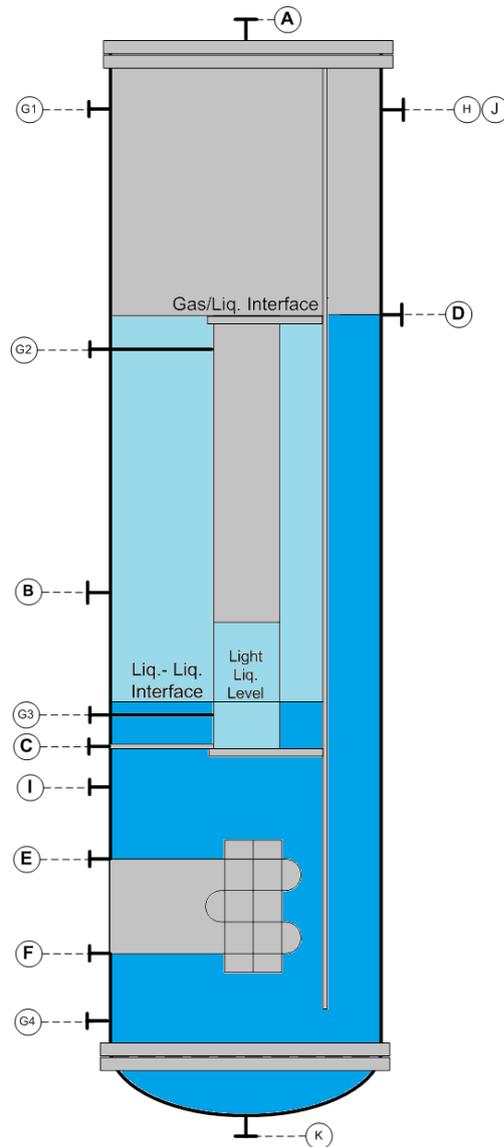
- **Pressure Vessels**

Table I.5 summarize the pressure vessels design.

**Table I.5. PCAPP Pressure Vessels**

TAG	Volume (l)	Design P (kPag)	Design T (°C)	DN (in)	Height (m)	Service
V-01	15,0	500	70	4	1,70	Flue-Gas heating and saturation with H <sub>2</sub> O
V-02	10,0	500	80	6	0,53	Knock-Out
V-03	15,0	500	90	8	0,90	Liquid-liquid phase separation
V-04	1,50	500	140	4	0,58	Reflux condensate separation

The vessel V-03 is illustrated in Fig. I.8 and its nozzles list is shown in Table I.6.

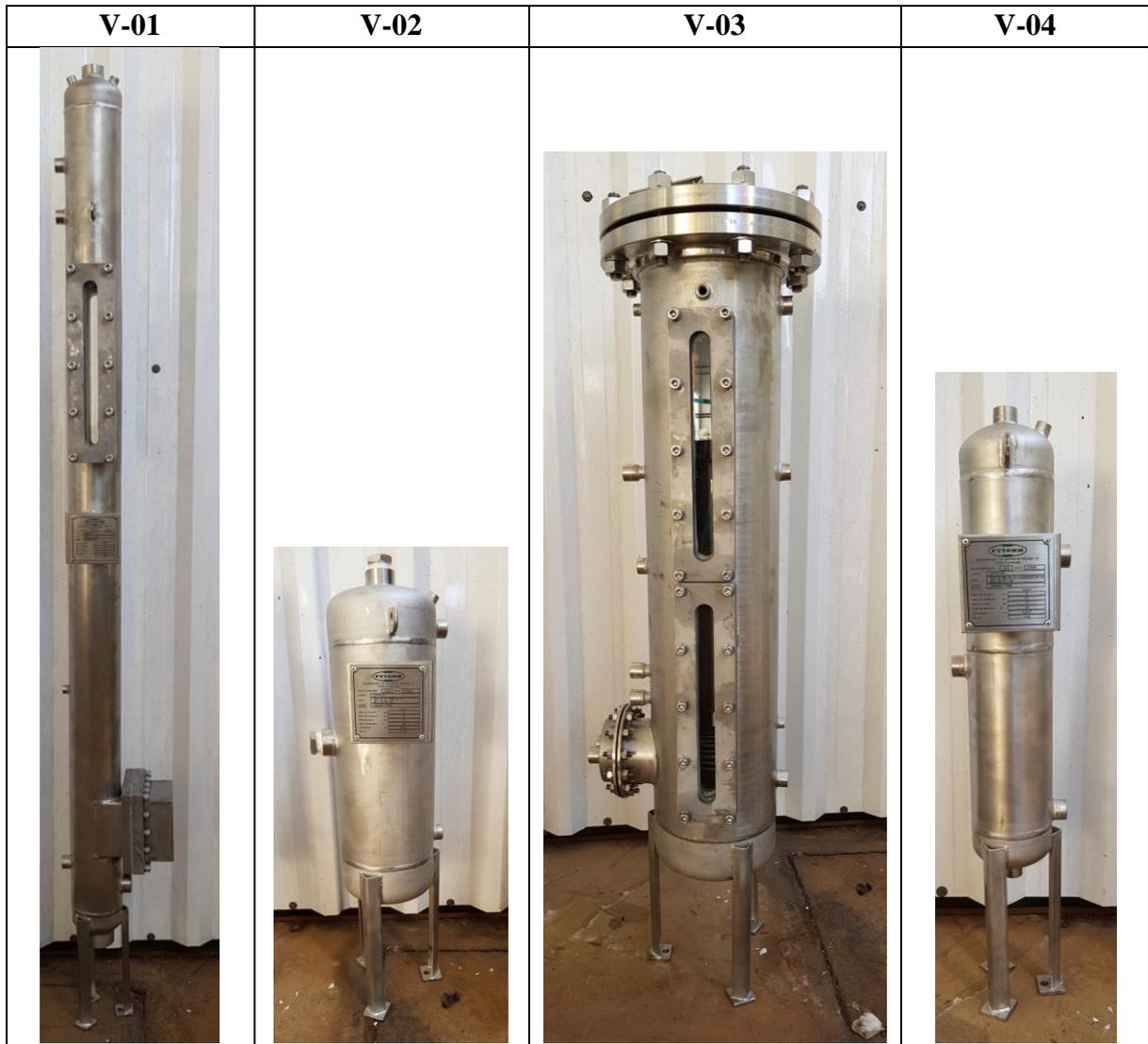


**Figure I.8. V-03 Internals and Nozzles**

**Table I.6. Nozzles list of the V-03**

NOZZLE	QUANT.	DIAM. (in)	DESCRIPTION
A	1	1/2	Top gas outlet
B	1	1/2	Liquid inlet
C	1	1/2	Light liquid outlet
D	1	1/2	Heavy liquid outlet
E	1	1/2	Thermal fluid inlet
F	1	1/2	Thermal fluid outlet
G	4	1/2	Level sensor
H	1	1/4	Pressure gauge
I	1	1/4	Temperature sensor
J	1	1/4	Pressure sensor
K	1	1	Drain

Fig. I.9 shows pictures of the four vessels of the PCAPP.



**Figure I.9. PCAPP Pressure Vessels**

- **Pumps**

The pump selection was challenging due to the very low flow rates, high temperatures and chemical aggressivity of the fluids. The Micropump external gear pumps (Micropump, 2020) GA and GB series were considered suitable for the service. Table I.7 summarize the specification of the pumps.

However, the PCAPP is intended to deal with multiple PCAS and CO<sub>2</sub> flow rates and partial pressures. As a result, a great flow rate flexibility is necessary. To meet this demand VSD and gearboxes were adopted. Table I.8 shows the range of flow rates and revolutions per minute (rpm) necessary to achieve the range of flow rates shown in Table I.7.

**Table I.7. PCAPP Pumps**

TAG	Volumetric Flow Rates			P <sub>inlet</sub> (barg)	P <sub>outlet</sub> (barg)	T <sub>operation</sub> (°C)	T <sub>design</sub> (°C)	Max viscosity (cP)	Design density (kg/m <sup>3</sup> )
	Min (l/h)	Nominal (l/h)	Max (l/h)						
<b>P-01</b>	36	90	210	0.5	3.0	45	80	250	1200
<b>P-02</b>	36	90	210	0.5	1.5	40	80	250	1200
<b>P-03</b>	15	30	120	0.1	3.0	40	90	250	1200
<b>P-04</b>	24	66	180	0.1	3.0	40	90	250	1200
<b>P-05A</b>	9.0	24.0	36.0	1.0	3.0	40	90	1.1	1000
<b>P-05B</b>	1.0	3.6	6.0	1.0	3.0	40	90	1.1	1000
<b>P-06</b>	36	75	210	2.0	3.0	130	150	20	1000
<b>P-07</b>	36	75	210	2.0	3.0	130	150	20	1000
<b>P-08</b>	36	105	210	0,0	3.0	40	50	250	1200
<b>P-12</b>	36	75	210	2.0	3.0	130	150	20	1000

**Table I.8. PCAPP pumps: drives, variable speed drives, gearboxes and rpm range**

TAG	Pump Model	Drive	VSD	Gearbox Ratio	RPM RANGE					
					Min	Nom	Max	Min*	Nom*	Max
<b>P-01</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1425	3325	33%	83%	97%
<b>P-02</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1425	3325	33%	83%	97%
<b>P-03</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	4:1	238	475	1900	28%	55%	55%
<b>P-04</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	3:1	380	1045	2850	33%	91%	83%
<b>P-05A</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	599	1596	2394	35%	93%	70%
<b>P-05B</b>	GA	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	4:1	204	733	1221	24%	85%	71%
<b>P-06</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1188	3325	33%	69%	97%
<b>P-07</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1188	3325	33%	69%	97%
<b>P-08</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1663	3325	33%	97%	97%
<b>P-12</b>	GB	WEG - 220/380V 3PH 60Hz 0,75CV, IEC 71-B14	WEG CFW 500	2:1	570	1188	3325	33%	69%	97%

\*considering gearbox

- **Instruments, Control and Automation**

Table I.9 lists all the instruments specified to the PCAPP

**Table I.9. PCAPP List of Instruments**

<b>Type</b>	<b>Qty.</b>	<b>Sign</b>
Pressure Sensor/Transmitter (PT)	15	4 – 20 mA
Temperature Sensor/Transmitter (TT)	35	4 – 20 mA
Differential Pressure Sensor/Transmitter (PDT)	4	4 – 20 mA
Flow Meter/Transmitter (FT)	12	4 – 20 mA
Level Sensor/Transmitter (PT)	7	4 – 20 mA
Analyzers (AT)	8	4 – 20 mA

Table I.10 lists all the control loops of the PCAPP.

**Table I.10. PCAPP List of Control Loops**

<b>Type</b>	<b>Qty.</b>	<b>Control Element</b>
Pressure Control	3	Valve (PCV)
	2	Valve (TCV)
Temperature Control	1	Solid State Relay (SSR)
	2	Process Thermostat
Mass Flow Control	3	Mass Flow Controller (MFC)
Level Control	9	VSD of pumps
Tank Mixer rpm	1	VSD

Table I.11 lists the control panel components and Fig. I.10 shows a picture of the panel.

**Table I.11. PCAPP List of components of Programmable Logic Control Panel**

<b>Description</b>	<b>Qty.</b>
Programmable Logic Control Wago model PFC200	1
CPU - 24 Vdc, Modbus TCP/IP, PFC200, Slot Micro SD	1
Analog Card 8 inputs, 4..20mA	11
Digital Card 16 inputs, 24Vdc, 3ms	2
Digital Card 16 outputs, 24Vdc	2
Analog Card 4 outputs, 4..20mA	4
Analog Card 4 outputs, 0..20mA	2
Module	1
Power source: In:100..240Vac / Out:24Vdc, 10A	1
Relay, 24Vdc	64
Jumper (for relay) 10 ways	10

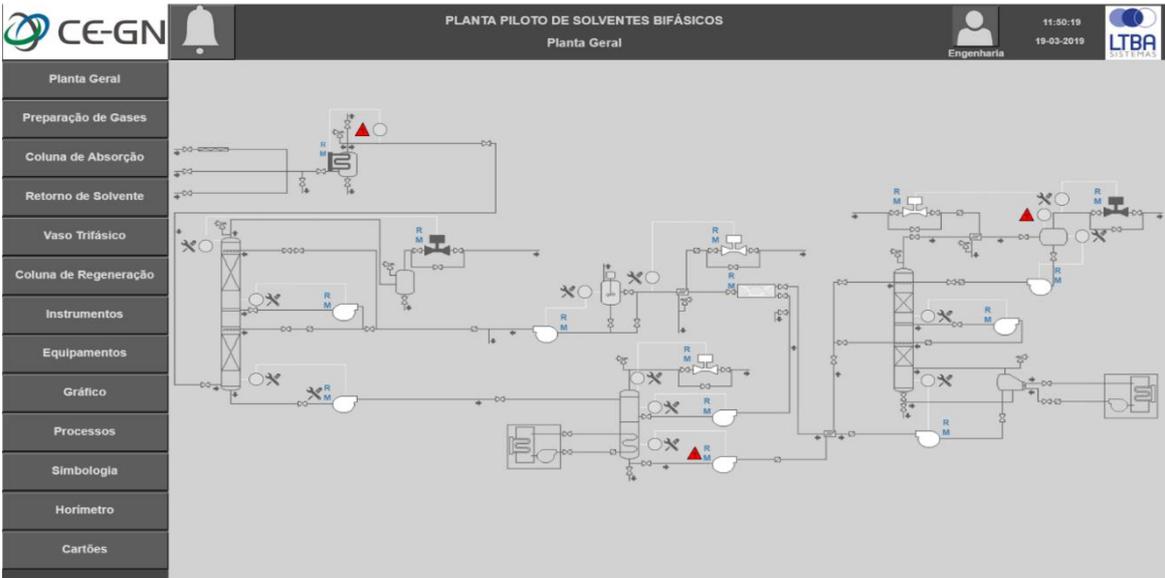


Figure I.10. PCAPP Control Panel

## ERRATUM

CRUZ, M. D. A. et al. Environmental Performance of a Solid Waste Monetization Process Applied to a Coal-Fired Power Plant with Semi-Dry Flue Gas Desulfurization. **Journal of Sustainable Development of Energy, Water and Environment Systems**, v. 7, n. 3, p. 506–520, 2018.

1 – The reference [31] (CONAMA 03/1990), cited in the page 510, line 13 of the original article (reproduced in the Chapter 6 of this thesis) is a national regulation for air quality in Brazil. However, the limit of 400 mg/Nm<sup>3</sup> for SO<sub>2</sub> emissions from coal-fired Power Plants is established by the BNDES, through the following reference:

BNDES, 2020. Environmental criteria to support the power generation segment. [WWW Document]. URL [http://www.https://www.bndes.gov.br/SiteBNDES/bndes/bndes\\_en/Institucional/Social\\_and\\_Environmental\\_Responsibility/socioenvironmental\\_policy/environmental\\_criteria\\_power\\_generation.html](http://www.https://www.bndes.gov.br/SiteBNDES/bndes/bndes_en/Institucional/Social_and_Environmental_Responsibility/socioenvironmental_policy/environmental_criteria_power_generation.html) (accessed 5.15.20).

2 – In the last line of Tables 6, 7, and 8 (Tables 6.6, 6.7, and 6.8 of this thesis): where it reads “SO<sub>4</sub>”, it should read “CaSO<sub>4</sub>”.