

# 1 **Process Analysis of Pressurized Oxy-Coal Power Cycle for Carbon Capture Application** 2 **Integrated with Liquid Air Power Generation and Binary Cycle Engines**

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## 8 **Abstract**

9 In this paper, the thermodynamic advantage of integrating liquid air power generation (LAPG)  
10 process and binary cycle waste heat recovery technology to a standalone pressurized oxy-coal  
11 combustion supercritical steam power generation cycle is investigated through modeling and  
12 simulation using Aspen Plus<sup>®</sup> simulation software version 8.4. The study shows that the  
13 integration of LAPG process and the use of binary cycle heat engine which convert waste  
14 heat from compressor exhaust to electricity, in a standalone pressurized oxy-coal combustion  
15 supercritical steam power generation cycle improves the thermodynamic efficiency of the  
16 pressurized oxy-coal process. The analysis indicates that such integration can give about 12 –  
17 15% increase in thermodynamic efficiency when compared with a standalone pressurized  
18 oxy-coal process with or without CO<sub>2</sub> capture. It was also found that in a pressurised oxy-  
19 coal process, it is better to pump the liquid oxygen from the cryogenic ASU to a very high  
20 pressure prior to vapourization in the cryogenic ASU main heat exchanger and subsequently  
21 expand the gaseous oxygen to the required combustor pressure than either compressing the  
22 atmospheric gaseous oxygen produced from the cryogenic ASU directly to the combustor  
23 pressure or pumping the liquid oxygen to the combustor pressure prior to vapourization in the  
24 cryogenic ASU main heat exchanger. The power generated from the compressor heat in the  
25 flue gas purification, carbon capture and compression unit using binary cycle heat engine was  
26 also found to offset about 65% of the power consumed in the flue gas cleaning and  
27 compression process.

28 The work presented here shows that there is a synergistic and thermodynamic advantage of  
29 utilizing the nitrogen-rich stream from the cryogenic ASU of an oxy-fuel power generation  
30 process for power generation instead of discarding it as a waste stream.

31 **Keywords:** Liquid air energy storage; Pressurised Oxy-coal combustion; Air separation unit;  
32 Process integration; Process simulation

## 33 **1. Introduction**

### 34 **1.1 Coal-fired power generation and Post-combustion Carbon Capture**

35 The global climate change resulting from CO<sub>2</sub> and other greenhouse gas emissions has  
36 become one of the greatest environmental threats of our time. Reduction in CO<sub>2</sub> emission  
37 especially from coal-fired power plants has been the mainstay of many researches on climate  
38 change mitigation. Studies show that a one percent point improvement in the efficiency of a  
39 conventional pulverised coal combustion power plant can result in a 2 – 3% reduction in CO<sub>2</sub>  
40 emission (WCA, 2014).

41 Despite the historical tightening of emission constraints from coal-fired power plants, its use  
42 for power generation has been on the increase mainly due to its availability, cost and the ever  
43 increasing global energy demand ((IEA, 2013, IEA, 2012, IEA, 2011). This shows that coal  
44 will continuously play a major role in meeting the global energy need. However, the success  
45 will depend on the development of technologies to control pollution and CO<sub>2</sub> emissions from  
46 such plants (Ciferno et al., 2000) especially now that CO<sub>2</sub> is becoming regulated in the US  
47 and Europe.

48 Carbon capture and sequestration (CCS) technologies have been in development for over a  
49 decade (Hagi et al., 2013) and is required to provide a long term solution by virtually  
50 eliminating CO<sub>2</sub> emission from coal-fired power plants. One of the CCS technologies which  
51 has been under investigation for decades now is the post-combustion process. Post-  
52 combustion technology involves capturing the CO<sub>2</sub> contained in the flue gas after the  
53 combustion process. Unlike the pre-combustion technology, this technology can easily be  
54 added to existing fossil fuel power plants for CO<sub>2</sub> capture. The efficiency of this technology  
55 largely depends on the concentration of the CO<sub>2</sub> in the flue gas. The major barriers opposing  
56 the commercialisation of this technology include: (a) high capital and operation cost; (b)  
57 steam consumption for solvent regeneration (Ciferno et al., 2000).

## 58 **1.2 Oxy-fuel combustion and its recent development**

59 Conventional coal based power plants produces flue gas with up to 10 - 15 vol% CO<sub>2</sub>  
60 (Ciferno et al., 2000, Hong et al., 2009). The low CO<sub>2</sub> content of the flue gas makes the  
61 capture operation energy intensive. One way of improving the CO<sub>2</sub> concentration in the flue  
62 gas is by using pure oxygen in the combustion process instead of air. The replacement of air  
63 with oxygen otherwise known as oxy-fuel combustion helps to produce a flue gas that  
64 contains mainly CO<sub>2</sub> and water vapour (Hong et al., 2009, Hu and Yan, 2011, Roy and  
65 Bhattacharya, 2013), which can easily be separated.

66 Studies had shown that there are more potential in pressurized oxy-fuel process compared to  
67 atmospheric oxy-fuel combustion power cycles. ENEL work in the area of pressurized oxy-  
68 fuel combustion using a novel pressurized oxy-combustion technology, known as the  
69 Isotherm Pwr® technology shows an increased heat transfer rates on the heat recovery steam  
70 generator (HRSG) compared to atmospheric oxy-combustion process (Zheng, 2011, Barbucci,  
71 2008). In the Isotherm Pwr® process, combustion takes place at elevated pressures and at  
72 1400 – 1600 °C.

73 However, one big challenge facing oxy-fuel process is the high energy requirement of the  
74 cryogenic air separation unit (ASU) which is currently the only mature air separation  
75 technology that can produce high purity, high tonnage oxygen required by the power plant.  
76 Thus improving the energy efficiency of the ASU is paramount to the success of oxy-fuel  
77 combustion process.

78 Several studies have been carried out on how to reduce the energy demand of cryogenic ASU  
79 for oxy-fuel combustion application. Some of the studies include:(1) improving the energy  
80 efficiency of heat exchangers and compressors and (2) the use of control system with real  
81 time optimization capability (Castle, 2002, Rübberdt, 2009), (3) use of self-heat recuperation  
82 process (Kansha et al., 2011), (4) pumping the liquid oxygen produced to a very high  
83 pressure prior to vaporization and expansion to the required process pressure (Manenti et al.,

84 2013), (5) recovery of the compressor heat using organic Rankine cycle system (Aneke and  
85 Wang, 2015b, Aneke and Wang, 2015a).

### 86 **1.3 Liquid air power generation and Motivation for process integration**

87 The recent breakthrough in liquid air power generation (LAPG) provides a new synergistic  
88 advantage for minimizing the high energy concern associated with oxy-fuel combustion of  
89 fossil fuels through integration with other processes. In a typical oxy-fuel combustion process,  
90 high tonnage gaseous nitrogen rich stream which contains about 80 mol% - 95 mol% nitrogen  
91 is produced together with the 95 mol% oxygen required for the oxy-fuel process. Presently,  
92 this gaseous nitrogen rich stream is discarded as waste since there is usually no need for  
93 nitrogen on site. However, demonstration plant studies had shown that this waste gas stream  
94 is an ideal working fluid for LAPG (Strahan, 2013). Thus, LAPG could be annexed to an  
95 oxy-fuel combustion process in order to improve both the overall power output and energy  
96 efficiency of the process. More heat integration opportunities could be possible depending on  
97 the plant configuration.

### 98 **1.4 Novel contributions of this study and outline of this paper**

99 In this study, the thermodynamic advantage of integrating LAPG and binary cycle waste heat  
100 recovery heat engines to pressurized oxy-coal combustion with supercritical steam power  
101 cycle will be investigated through modeling and simulation using Aspen Plus® version 8.4  
102 simulation software. The entire process will be analysed to evaluate the impact of integrating  
103 the aforementioned process to the thermodynamic efficiency of a pressurized oxy-coal  
104 supercritical steam power cycle process which is used as the base case scenario. Different  
105 process scenarios will be investigated with/without carbon capture consideration. To the  
106 knowledge of the authors, this is the only work that has introduced the concept of utilizing  
107 the nitrogen rich stream from the ASU of an oxy-fuel combustion process for power  
108 generation instead of the current practice where it is discarded as a waste stream.

109 This paper is presented in five sections. The first section is the introduction and covers the  
110 carbon capture technologies, the liquid air power generation technology and the synergistic  
111 advantage of integrating oxy-fuel technology with liquid air power generation technology  
112 which is the underlining novelty of the paper. In the second section, the description of the  
113 different processes is presented while in section three, the Aspen simulation of the overall  
114 process is presented together with the modeling parameters and process efficiency definitions  
115 and equations. In section four, the results from the process simulation were presented and  
116 discussed while in section five, the conclusions drawn from the study carried out is presented.

## 117 **2. Process Description**

118 Figure 1 shows the process flow diagram of the pressurized oxy-coal combustion  
119 supercritical steam power cycle integrated with liquid air power generation and binary cycle  
120 heat engines proposed in this study. The process shown in Figure 1 consists of eight primary  
121 units: 1) Cryogenic ASU; 2) Pressurized oxy-coal combustor unit; 3) Steam generation unit;  
122 4) Supercritical steam power generation unit; 5) Air liquefaction unit; 6) LAPG unit; 7) Flue  
123 gas purification, Carbon dioxide capture and compression unit; 8) Binary cycle waste heat  
124 recovery units.

125 The pressurized oxy-coal combustor, steam generation unit and supercritical steam power  
126 generation unit were adapted from Hong et al., (2009) and Zheng (2011). The cryogenic ASU

127 and heat recovery using binary cycles were based on the previous work of the authors (Aneke  
128 et al., 2012, Aneke and Wang, 2015a, Aneke et al., 2011b), flue gas purification, carbon  
129 dioxide capture and compression unit were adapted from White et. al., (2009) while the air  
130 liquefaction and LAPG unit were adapted from a demonstration plant study carried out by the  
131 authors.

132 An overview of the process flow diagram shown in Figure 1 is as follows: the 10 bar  
133 pulverised coal slurry (stream 5) is first dried at the entrance to the combustor. This is  
134 followed by the decomposition of the coal into its constituents based on the properties of coal  
135 as shown in Table 1.

136 **Figure 1: Overall Process Flow Diagram of the Pressurized Oxy-coal Power Cycle for**  
137 **Carbon Capture Application Integrated with LAPG and Binary Cycle Engine**

138 **Table 1: Properties of coal used in the simulation (adapted from (Hu and Yan, 2011))**

139 The pressurized coal is burnt with 95.68 mol% oxygen from the cryogenic ASU (stream 4)  
140 and a portion of the recycled flue gas, (stream 10) which helps to maintain the combustion  
141 temperature at 1550 °C. A second portion of the flue gas recycle (stream 9) is used to control  
142 the temperature at the heat recovery steam generator (HRSG) inlet, which is kept close to 800  
143 °C to avoid slagging (Zheng, 2011) and also minimize hot corrosion and oxidation (Hong et  
144 al., 2009). The hot temperature-controlled flue gas (stream 6) is passed through the HRSG  
145 where it is used to generate supercritical steam used in the supercritical steam power  
146 generation unit.

147 The HRSG consists of two superheaters, a once-through boiler and an economizer. The steam  
148 reaches the supercritical state of 600 °C at 250 bar before being delivered to high pressure  
149 turbine (HPT) of the supercritical steam power generation unit. The superheater also acts as a  
150 re-heater to generate two reheat subcritical steam of 620 °C at 50 bar and 10 bar for the  
151 intermediate pressure turbine (IPT) and the low pressure turbine (LPT) respectively. Due to  
152 the presence of SO<sub>x</sub> and NO<sub>x</sub> in the flue gas, flue gas from the exit of the HRSG must be  
153 maintained at a temperature higher than the acid dew point. This is maintained at 239 °C in  
154 this present work. The remaining flue gas after the recycle is compressed to 15 bar and sent to  
155 the flue gas purification unit where water, SO<sub>x</sub>, NO<sub>x</sub> and inert gases are removed from the  
156 flue gas to achieve about 95.35 mol% CO<sub>2</sub> which is compressed to 110 bar (stream 52) and  
157 sent to the storage facility.

158 The nitrogen rich gas stream gas (stream 3) from the ASU unit is compressed and liquefied to  
159 produce the working fluid for the LAPG unit. The liquid nitrogen-rich stream otherwise  
160 known as the liquid air is pumped to a pressure of 120 bar (stream 33) vaporized using heat  
161 extracted from the supercritical steam power generation unit and expanded in a 4 stage  
162 turbine and reheat arrangement in the LAPG unit to produce power.

163 The compressor heat from the intercooler and after-cooler heat exchangers in the cryogenic  
164 ASU unit, LAPG unit, the flue gas purification and the CO<sub>2</sub> compression unit were utilized  
165 for power generation using binary cycle heat engine which uses R134a as the working fluid.  
166 The detailed description of each of the process unit is as given below.

167 **2.1 Cryogenic Air Separation Unit**

168 As aforementioned, oxy-coal combustion process uses oxygen at >95mol% purity instead of  
169 air for combustion operation. For the pressurized oxy-coal combustion process developed in  
170 this study, the combustion pressure is assumed to be 10 bar while the combustion chamber is  
171 maintained at 1550 °C using a mixture of oxygen and flue gas recycle. The oxygen used in  
172 the combustion chamber is produced from a cryogenic ASU and should be supplied at a  
173 pressure of 10 bar.

174 The process flow diagram of the cryogenic ASU for the specialist application proposed in this  
175 work is shown in Figure 2. Unlike the conventional ASU, the ASU proposed in this work  
176 makes use of only 2 columns to produce nitrogen-rich gaseous stream and 95.68 mol% liquid  
177 oxygen stream since there is no need for pure nitrogen and argon stream. The nitrogen-rich  
178 gaseous stream and the liquid oxygen stream exchanges heat with the in-coming air stream in  
179 the main heat exchanger in order to obtain an all gaseous product.

180 In the process, atmospheric air is filtered, cleaned and compressed in a 3 stage compressor  
181 with inter-cooling to a pressure of 6.35 bar. The compressed air is split into two, cooled and  
182 partially liquefied against leaving product streams (gaseous nitrogen-rich stream and liquid  
183 oxygen-rich stream (95.68 mol% purity)). One of the streams is sent to the high pressure  
184 column (HPC) where nitrogen is separated at a pressure of about 6 bar. The other stream is  
185 expanded in an air expander to a pressure of about 1.2 bar and sent to the low pressure  
186 column (LPC). The top nitrogen product from the HPC is condensed against the boiling  
187 oxygen in the reboiler of the LPC, before being depressurized and sent to the top of the LPC.  
188 The bottom liquid product from the HPC is also sent to the LPC after been depressurized in  
189 the JT valve. In the LPC, gaseous nitrogen-rich stream which contains about 94.54 mol%  
190 nitrogen leaves through the top of the column while the liquid oxygen rich stream (95.68  
191 mol% oxygen) which is required for the oxy-coal combustion leaves through the bottom of  
192 the column.

193 It is proposed in this work that the liquid oxygen produced in the cryogenic ASU be pumped  
194 to about 200 bar prior to vapourization in the main heat exchanger against in-coming air feed  
195 to the system. The high pressure gaseous oxygen from the main heat exchanger will then be  
196 expanded in the turbine to 10 bar (required combustion pressure), thus generating extra  
197 energy.

## 198 **Figure 2: Process Flow Diagram of Cryogenic ASU**

### 199 **2.2 Pressurized Coal Combustion, Steam Generation Unit and Supercritical Steam** 200 **Power Generation Unit**

201 The pressurized coal combustor process is based on the novel Isotherm Pwr® technology  
202 developed by ENEL (Zheng, 2011, Hong, 2009). The combustion is assumed to take place at  
203 a pressure of 10 bar and temperature of 1550 °C. In the process, wet coal is first dried and  
204 burnt with a mixture of 95.68 mol% oxygen and 23.4% (by mass) recycled flue gas in order  
205 to maintain the temperature of the combustor at 1550 °C. The exit flue gas from the  
206 combustion chamber is mixed with about 66.6% of the recycled flue gas in order to maintain  
207 the temperature to the HRSG at about 730 °C.

208 The supercritical steam generation unit otherwise known as the HRSG comprises of two  
209 superheaters, a once-through boiler and an economizer. The heating in the steam generation  
210 unit is provided by the temperature controlled flue gas at 730 °C in a counter-current flow

211 arrangement. The flue gas enters the first superheat of the HRSG at about 730 °C and is used  
212 to generate a supercritical steam at 600 °C and 250 bar and a subcritical steam (reheat stream)  
213 at 620 °C and 50 bar. The effluent flue gas from the first superheater is sent to the second  
214 superheater where it is used to further generate a subcritical steam (reheat stream) at 620 °C  
215 and 10 bar. From the second superheater, the flue gas enters the once-through boiler and the  
216 economiser where it is used to preheat and generate steam respectively before exiting the  
217 HRSG at about 239 °C.

218 The steam power generation unit comprises of a supercritical Rankine cycle. The  
219 supercritical steam at 600 °C and 250 bar generated in the first superheater of the HRSG  
220 enters into the high pressure turbine (HPT). The steam is expanded in the HPT to produce  
221 power. Part of the steam from the HPT is also injected into the pressurized combustor. Part of  
222 the exit steam from the HPT is reheated in the HRSG and used to provide steam for the  
223 intermediate pressure turbine (IPT) while the remaining is sent to the deaerator via a heat  
224 exchanger arrangement which is used to preheat the feed water system. The reheated steam at  
225 620 °C and 50 bar is expanded in the IPT to produce more power. Steam bleeding from the  
226 IPT is also used to preheat the feed water from the deaerator. Part of the exit steam from the  
227 IPT is returned to the HRSG and reheated to 620 °C before being sent to the low pressure  
228 turbine (LPT) while the remaining steam is sent to the deaerator. In the LPT, the steam is  
229 expanded for power generation. The exit steam from the LPT is condensed using cooling  
230 water at 25 °C. The condensed stream is preheated using heat from the water used to cool the  
231 combustion chamber wall. The combustor is assumed to lose 2% of the lower heating value  
232 of the coal to the water-cooled wall of the combustor (Hong et al., 2009). The preheated  
233 stream is sent to the deaerator where the whole liquid stream together with the makeup water  
234 is collected and pumped back into the HRSG to complete the steam cycle. The process flow  
235 diagram for these units is shown in Figure 3.

236 **Figure: 3 Process Flow Diagram of Pressurized Combustor Unit, Steam Generation**  
237 **Unit and Supercritical Steam Power Generation Unit**

### 238 **2.3 Air Liquefaction Unit**

239 Figure 4 shows the liquefaction unit where the nitrogen-rich (94.54 mol %) gaseous stream  
240 from the cryogenic ASU is liquefied to produce liquid nitrogen-rich stream otherwise known  
241 as liquid air. The process is based on the principle of Claude liquefaction cycle. The nitrogen-  
242 rich gas stream from the ASU is compressed in a 3 stage compressor with intercooler and  
243 after-cooler arrangement. The compressed gaseous stream is further compressed using a  
244 compressor joined to the shaft of an expander. The compressed gas is cooled with water and  
245 further cooled in the cold box with cryogenic gaseous stream from the expander outlet. The  
246 cold gaseous stream is expanded in the expander and the pressure dropped to produce a two  
247 phase stream (liquid and gas). The gaseous stream is used to cool incoming stream in the cold  
248 box while the liquid phase is sent to the cryogenic tank.

249 **Figure 4: Process Flow Diagram of Air Liquefaction Unit**

### 250 **2.4 Liquid Air Power Generation (LAPG) Unit**

251 In this unit, the liquid air in the cryogenic tank is pumped to about 120 bar and vapourized  
252 indirectly using a hot water loop which collects heat from the steam power generation unit.

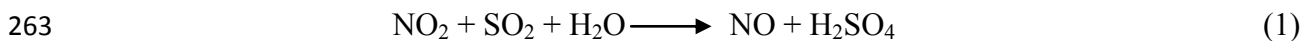
253 The vapourized air is expanded in a series of 4 stage expanders with inter-heaters to produce  
254 power (Figure 5).

## 255 **Figure 5: Liquid Air Power Generation Unit**

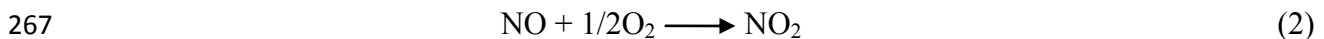
### 256 **2.5 Flue gas purification, carbon dioxide capture and compression unit**

257 After recycling of about 90% of the flue gas leaving the HRSG, the remaining flue gas is  
258 cleaned to remove the NO<sub>x</sub>, SO<sub>x</sub> and the inert gases. The cleaning of the flue gas starts with  
259 an increase in the pressure of the flue gas to 15 bar. After the compression operation, direct  
260 contact scrubbing was used to cool the flue gas product as well as condense water vapour  
261 present in the flue gas and remove residual ash particles and highly soluble SO<sub>3</sub>.

262 The removal of SO<sub>2</sub> involves the reaction of NO<sub>2</sub> with SO<sub>2</sub> to form sulphuric acid.



264 This reaction is fast and so is considered to be equilibrium limited. Since most of the NO<sub>x</sub> in  
265 the flue gas occurs as NO, the NO would have to be converted to NO<sub>2</sub> to allow equation 1 to  
266 proceed. The conversion of NO to NO<sub>2</sub> occurs as



268 Reaction 2 is a third order reaction with reaction rate given as (White et al., 2009)

$$269 \quad d[\text{NO}_2]/dt = 2k[\text{NO}]^2 \cdot [\text{O}_2] \quad (3)$$

270 where k, in l<sup>2</sup> mol<sup>-2</sup> s<sup>-1</sup> is 1200 x 10<sup>230/T</sup> where T is in Kelvin (White et al., 2009, Tsukahara et  
271 al., 1999).

272 Since the rate is proportional to pressure to the 3<sup>rd</sup> power, it is assumed that the reaction rate  
273 will become significant at the pressure of 15 bars and low temperature obtained after  
274 scrubbing.

275 After the removal of SO<sub>2</sub>, the flue gas is compressed to about 30 bar, dried and then purified  
276 further to remove the inert gases such as nitrogen and argon using low temperature  
277 processing. In the process, the impure flue gas is cooled in HX1 and HX2 against evaporating  
278 lower pressure liquid CO<sub>2</sub> streams to a temperature of -55 °C, which is close to its triple point.  
279 The inert stream leaving the cold equipment at about 30 bars is further heated and expanded  
280 to produce power while the 95.35 mol% pure CO<sub>2</sub> streams leaving the cold equipment are  
281 compressed adiabatically in a second stage of CO<sub>2</sub> compression to a pressure of 110 bars. The  
282 process flow diagram is shown in Figure 6.

## 283 **Figure 6: Process Flow Diagram of flue gas Cleaning, CO<sub>2</sub> purification and** 284 **compression unit**

### 285 **2.6 Binary Cycle/Organic Rankine Cycle (ORC) Waste Heat Recovery Unit**

286 Because there is no need for further heat integration in the process, the ORC unit is used to  
287 convert the waste heat from the compressor exhausts to electricity by acting as the  
288 compressor intercooler and after cooler system (Aneke and Wang, 2015b). In the ORC unit,  
289 the waste heat from the compressor exhaust is used to preheat and vapourise an organic  
290 working fluid in the evaporator. The vapourised organic fluid from the evaporator passes  
291 through the turbine where it expands to produce work which turns the shaft connected to

292 the generator to produce electricity (Aneke et al., 2012). The process flow diagram of a  
293 typical ORC system is shown in Figure 7. The system is used in all the compressors as shown  
294 in Figures 2 to 6 to compressor heat to electricity. The process uses R134a as the working  
295 fluid (Aneke et al., 2012, Aneke et al., 2011a).

### 296 **Figure 7: Process Flow Diagram of Organic Rankine Cycle**

297 The individual process flow diagrams presented in this work are linked to each other using  
298 alphabets from A to J.

### 299 **3. Aspen Plus<sup>®</sup> Simulation of the Overall Process**

300 The overall model of the process was developed using Aspen Plus<sup>®</sup> v8.4. Aspen is modular  
301 mode steady state simulation software. It contains modules of unit operations (like heat  
302 exchangers, reactors, turbine, flash, pumps etc.) used in the development of the model of the  
303 entire process described in this paper. The model equations used to model the individual unit  
304 operations and the physical property calculator used to model the process stream property in  
305 Aspen were standard equations. Due to the complexity of the flowsheet and in order to  
306 improve the convergence and the simulation time, the entire flowsheet of the process as  
307 represented in Figure 1 is split into five main sub-processes: 1) pressurised coal combustion,  
308 steam generation and supercritical steam power generation unit, 2) cryogenic air separation  
309 unit, 3) nitrogen-rich stream air liquefaction and liquid air power generation unit, 4) flue gas  
310 purification, carbon capture and CO<sub>2</sub> compression unit and 5) binary cycle heat engine units.  
311 The sub-processes were simulated individually and the results transferred to the appropriate  
312 sub-process as inputs.

313 Coal is modeled as a non-conventional component using the ultimate and proximate analysis  
314 (Table 1). The coal combustor in this work is modeled using the RGibbs reactor while the  
315 coal decomposition is modeled using RYield reactor together with the proximate and ultimate  
316 analysis of the coal. The fluid property of the overall process is modeled using Peng  
317 Robinson while the steam properties is modeled using STEAM-TA (ASME 1967 steam table  
318 correlations). The SO<sub>2</sub> removal is modeled using RadFrac column with chemical reaction.  
319 The pressure drop in the heat exchangers is assumed to be negligible. The simulation  
320 parameter for the overall process is given in Table 2.

### 321 **Table 2: Process Simulation Parameters**

322 In order to investigate the contribution of each of the process unit to the efficiency of a  
323 standalone pressurised oxy-coal process with/without carbon capture, different kinds of  
324 process efficiencies were evaluated and analysed in this study. The first three process  
325 scenarios show the impact of how gaseous oxygen is supplied from the cryogenic ASU on the  
326 efficiency of the pressurized oxy-coal process. Three different methods of supplying  
327 pressurized gaseous oxygen to the combustion chamber were investigated in the paper. The  
328 first is by pressurizing the gaseous oxygen produced in the cryogenic ASU to combustion  
329 chamber pressure of 10 bar (Case 1), the second is pumping the liquid oxygen produced in  
330 the cryogenic ASU to 10 bar before vapourizing it in the ASU main heat exchanger (Case 2)  
331 while the third is by pumping the liquid oxygen to 200 bar before vapourizing it in the ASU  
332 main heat exchanger and then expanding the high pressure gaseous oxygen to the required  
333 combustion chamber pressure of 10 bar (Case 3) thus producing extra power from the  
334 expander.



335 The baseline process scenario also known as the standalone pressurized oxy-coal process  
 336 (Case 1) is regarded as the configuration where there is no heat recovery from compressors of  
 337 the cryogenic ASU unit, no integration with liquid air power generation unit and where  
 338 gaseous oxygen product from the cryogenic ASU is compressed to the combustor pressure of  
 339 10 bars before being mixed with the recycle flue gas stream going into the combustion  
 340 chamber. As aforementioned, other alternatives include: pumping the liquid oxygen produced  
 341 in the LPC of the cryogenic ASU to 10 bars prior to vapourization in the main ASU heat  
 342 exchanger and subsequent mixing with the recycled flue gas into the combustor (Case 2) and  
 343 pumping to 200 bar prior to vapourization and expanding the gaseous oxygen to 10 bars  
 344 before been sent to the combustor with the recycled flue gas (Case 3). All the process  
 345 scenarios were evaluated with/without carbon capture.

346 For processes without carbon capture, the efficiencies are defined as follows:

347 For Case 1, the efficiency of the standalone pressurised oxy-coal combustion process with  
 348 supercritical steam power cycle and 10 bar gaseous oxygen from the cryogenic ASU  
 349 (baseline process) is defined as:

$$350 \quad \mu_{ox} = \frac{(\sum G_p - \sum D_{asu})}{Q_{in}} \quad (4)$$

351 where,

352  $\sum G_p$  is the sum of the power generated from the supercritical steam power generation unit  
 353 and the power from the cryogenic ASU air expander;  $\sum D_{asu}$  is the sum of the power  
 354 consumption by the air and gaseous oxygen compressors in the ASU as well as the pumps in  
 355 the supercritical power generation unit and  $Q_{in}$  the thermal energy input into the system from  
 356 the coal.

357 For Case 2, the efficiency of the standalone pressurized oxy-coal combustion process with  
 358 pumped liquid oxygen to 10 bars is defined as:

$$359 \quad \mu_{ox1} = \frac{(\sum G_p - \sum D_{asup})}{Q_{in}} \quad (5)$$

360 where,

361  $\sum G_p$  is the sum of the power generated from the supercritical steam power generation unit  
 362 and the power from the cryogenic ASU air expander;  $\sum D_{asup}$  is the sum of the power  
 363 consumption by the air compressors of ASU, liquid oxygen pump and the pumps in the  
 364 supercritical power generation unit and  $Q_{in}$  is the thermal energy input into the system from  
 365 the coal.

366 For case 3 where oxygen is pumped to 200 bars prior to vapourization in the main ASU heat  
 367 exchanger followed by expanding to 10 bars, the efficiency is defined as:

$$368 \quad \mu_{ox2} = \frac{(\sum G_{p2} - \sum D_{asup})}{Q_{in}} \quad (6)$$

369 where,

370  $\sum G_{p2}$  is the sum of the power generated from the supercritical steam power generation unit,  
 371 the power from the ASU air expander and the power from the oxygen expander;  $\sum D_{asupp}$  is  
 372 the sum of the power consumption by the air compressors of ASU, liquid oxygen pump and  
 373 the pumps in the supercritical power generation unit and  $Q_{in}$  is the heat input into the system  
 374 from the fuel.

375 In Case 4, the efficiency improvement of Case 3 as a result of the recovery of the compressor  
 376 waste heat using binary cycle heat engine is evaluated. In this case, the efficiency of the  
 377 standalone pressurized oxy-coal combustion process with pumped oxygen to 200 bars prior to  
 378 vapourization in the main ASU heat exchanger followed by expanding to 10 bars (i.e. Case 3)  
 379 together with binary waste heat recovery from the ASU compressors is defined as:

$$380 \quad \mu_{ox3} = (\sum G_{p3} - \sum D_{asupp}) / Q_{in} \quad (7)$$

381 where,

382  $\sum G_{p3}$  is the sum of the power generated from the supercritical steam power generation unit,  
 383 the power from the ASU air expander, the power from the oxygen expander and the power  
 384 from the binary cycle for heat recovery from ASU compressors;  $\sum D_{asupp}$  is the sum of the  
 385 power consumption by the air compressors of ASU, liquid oxygen pump, pumps in the  
 386 supercritical power generation unit and the binary cycle pump and  $Q_{in}$  is the thermal energy  
 387 input into the system from the fuel.

388 Case 5 focuses on the liquid air power generation unit of the entire process. The efficiency is  
 389 defined as:

$$390 \quad \mu_{la} = (G_{la} - D_{aux}) / \sum Q_{scp} \quad (8)$$

391 where,

392  $G_{la}$  is the power generated from the liquid air power generation unit;  $D_{aux}$  is the discharging  
 393 pump power consumption and  $\sum Q_{scp}$  is the sum of the heat input into the system by the  
 394 steam from the steam generation unit and the power consumption by the liquefaction  
 395 compressor (i.e. the charging power).

396 In Case 6, the impact of the binary heat recovery from the charging compressor exhaust heat  
 397 was investigated (i.e. Case 5 with heat recovery from the charging compressor exhaust heat  
 398 using binary cycle heat engine). The efficiency of the liquid air power generation unit with  
 399 binary cycle waste heat recovery from the compressors is given as:

$$400 \quad \mu_{la1} = (\sum G_{la1} - \sum D_{aux1}) / \sum Q_{scp} \quad (9)$$

401 where,

402  $\sum G_{la1}$  is the sum of the power generated from the liquid air power generation unit and the  
 403 binary cycle waste heat recovery from compressor;  $\sum D_{aux1}$  is the sum of the pump power  
 404 consumption by the liquid air power generation during the discharging operation and by the

405 binary cycle engine pump and  $\sum Q_{scp}$  is the sum of the heat input into the system by the  
406 steam from the steam power generation unit and the liquefaction process compression power.

407 In Case 7, the efficiency of the entire process is considered. The efficiency of the pressurized  
408 oxy-coal combustion process with pumped oxygen to 200 bars prior to vapourization in the  
409 main ASU heat exchanger followed by expanding to 10 bars integrated with liquid air power  
410 generation unit and binary cycle engine (i.e. Case 4 + Case 6) is defined as

$$411 \quad \eta_{RT} = \frac{(\sum G_{ov} - \sum D_{ov})}{Q_{in}} \quad (10)$$

412 where,

413  $\sum G_{ov}$  is the sum of the power generated from the overall process except the inert expander  
414 This comprises of the power from the supercritical steam power generation unit, the liquid air  
415 power generation unit, the binary cycle waste heat recovery units (except those in the flue gas  
416 cleaning, CO<sub>2</sub> purification and compression unit), the oxygen expander, the air expander of  
417 the cryogenic ASU,  $\sum D_{ov}$  is the sum of the power consumption by the auxiliary equipment  
418 such as pumps and ASU compressors except those in the flue gas cleaning, CO<sub>2</sub> purification  
419 and compression unit;  $\sum Q_{in}$  is the sum of the thermal energy input from the coal and power  
420 consumption by the compressors of the liquefaction process (charging power consumption).

421 The corresponding efficiencies for the process with carbon capture is obtained by including  
422 the power generation and consumption in the flue gas cleaning, CO<sub>2</sub> purification and  
423 compression unit. This include: power generation in the inert expander, power generation in  
424 the binary cycle heat engines attached to the CO<sub>2</sub> compressor train exhaust and power  
425 consumption by the pumps and compressors.

#### 426 **4. Results and Discussions**

427 The summary of the simulation results, stream conditions (mass flowrate, temperature and  
428 pressure) and the efficiency results for the different cases investigated in this work are shown  
429 in Tables 3, 4 and 5 respectively.

430 From Table 3, it can be seen that the oxygen purity generated from the ASU is about 95.68  
431 mol% which met the requirement for a typical oxy-coal combustion process. The specific  
432 power consumption of the ASU is 0.309 kWh/kg-O<sub>2</sub>, which is within the range of values  
433 quoted in the literature (Lombardi et al., 2011, Hong et al., 2009, Kansha et al., 2011). For a  
434 standalone pressurised oxy-coal process with gaseous oxygen compression to 10 bar (Case 1),  
435 the gross power generation from the supercritical steam power generation system was  
436 572595.28 kW while the gross power consumption by the supercritical steam power  
437 generation unit pumps, gaseous oxygen compressor and cryogenic ASU compressors were  
438 22821.52 kW, 20526.91 kW and 89094.10 kW respectively. This translates to an efficiency  
439 of 43.75% without carbon capture. The inclusion of carbon capture reduces the efficiency to  
440 39.99% giving an energy penalty of 3.76%. Replacing the gaseous oxygen compressor with  
441 liquid oxygen pump (Case 2) increases the process efficiency to 45.79% and 42.03% for  
442 scenarios without and with carbon capture respectively. Case 3 shows that pumping the liquid  
443 oxygen to a pressure of 200 bars and expanding the gaseous oxygen to the required  
444 combustion pressure of 10 bars improved the efficiency by 0.66% and 2.7% respectively for

445 processes with and without carbon capture when compared with processes with only liquid  
446 oxygen pump and gaseous oxygen compression.

447 **Table 3: Summary of Simulation Results**

448 **Table 4: Stream Temperature, Pressure and Mass Flowrate**

449 **Table 5: Process Efficiencies**

450 This is because more power is generated during oxygen expansion than is consumed during  
451 the pumping operation. For example, about 1603.21 kW of power was consumed in pumping  
452 the liquid oxygen to 200 bars while about 8182.53 kW of power is generated by expanding  
453 the high pressure gaseous oxygen to the required combustor pressure of 10 bars. The  
454 temperature of the oxygen stream changes from 20 °C before the expansion process to -  
455 139.50 °C after the expansion process. The exit oxygen from the expander is mixed with part  
456 of the recycled flue gas to obtain a stream at a temperature of 210 °C which is sent back to the  
457 combustion chamber. Although there is an improvement in process efficiency when Case 3 is  
458 compared with Case 2 and 1, doing this might not be economically attractive especially for  
459 Case 3 vs Case 2 since the improvement in efficiency of 0.66% might not be able to justify  
460 the cost of installing an additional oxygen expander. Case 4 shows the importance of  
461 recovering the waste heat from the compressor exhaust using binary cycle heat engine instead  
462 of the conventional water based inter-cooling/ after-cooling arrangement. The recovery of the  
463 compressor heat in the ASU adds a net power output of 9416.91 kW. The increase in the  
464 power translates to an increase in the efficiency to 47.38 % and 43.62% for cases without and  
465 with carbon capture.

466 **Figure 8: P-h Diagram of R134a working fluid used in the ORC System attached to the**  
467 **Cryogenic ASU**

468 In the same vein, cases 5 and 6 shows the efficiency of the liquid air power generation unit  
469 with and without waste heat recovery from the compressor exhaust stream using binary cycle  
470 heat engines. Without waste heat recovery, the efficiency of the liquid air power generation  
471 unit is 40.17%. This is improved to 42.69% when the waste heats from the compressors are  
472 converted to power using binary cycle heat engines. Figure 8 shows the Pressure-enthalpy  
473 diagram of the binary cycle heat engine (ORC) attached to the compressors in the cryogenic  
474 ASU. The flue gas cleaning, CO<sub>2</sub> purification and compression unit modeled in this work  
475 consumes about 38009.75 kW of power to generate 89.30 kg/s of CO<sub>2</sub> with purity of 95.35  
476 mol.%. This gives a specific power consumption of 0.118 kWh/kg-CO<sub>2</sub>. The round trip  
477 efficiency of the overall process is calculated as 56.01% and 54.74% for cases without and  
478 with carbon capture respectively. It is interesting to see that the round trip efficiency for the  
479 case with carbon capture is slightly lower than that without carbon capture. The reason is  
480 because, the sum of power obtained from the binary cycle waste heat recovery heat engines  
481 attached to the compressors in the flue gas cleaning, CO<sub>2</sub> purification and compression unit  
482 as well as the power from the inert expander is lower than the sum of the powers consumed  
483 by the compressors and pumps in the purification and compressor unit (39979.04 kW vs  
484 26112.06 kW). Thus, with the use of binary waste heat recovery heat engines to convert the  
485 waste heat from the compressors used in the flue gas purification unit, it is possible to  
486 generate up to 65% of the power consumed in the flue gas cleaning, CO<sub>2</sub> purification and  
487 compression unit.

488 **5. Conclusions**

489 The study carried out in this paper shows that the integration of liquid air power generation  
490 process to a standalone pressurized oxy-coal supercritical steam power generation process  
491 brings an enormous thermodynamic advantage as seen by the increase in the thermodynamic  
492 efficiency of the integrated process when compared to the standalone process. The results  
493 also show that the recovery of the waste heat from the compressors using binary cycle heat  
494 engines also helps to improve the overall power generated from the process. It was also found  
495 that the power generated by the binary cycle heat engine which recovers waste heat from the  
496 entire compressor train of the flue gas cleaning, carbon dioxide purification and compressor  
497 unit is capable of offsetting up to 65% of the power required for the flue gas cleaning, carbon  
498 capture and compression process. The results also show the significance of incorporating  
499 waste heat recovery technology to recover compressor waste heat from processes which make  
500 use of compressors.

501 **Acknowledgement**

502 The authors would like to thank UK EPSRC (through EPSRC Capital Investment on Energy  
503 Storage) and University of Hull for their financial support.

504 **References**

- 505 ANEKE, M., AGNEW, B. & UNDERWOOD, C. (2011a) Approximate Analysis of the Economic  
506 Advantage of a Dual Source ORC System Over two Single ORC Systems in the Conversion  
507 of Dual low and Mid Grade Heat Sources. *Journal of EUEC*, 5, 1-17.
- 508 ANEKE, M., AGNEW, B. & UNDERWOOD, C. (2011b) Performance analysis of the Chena binary  
509 geothermal power plant. *Applied Thermal Engineering*, 31, 1825-1832.
- 510 ANEKE, M., AGNEW, B., UNDERWOOD, C., WU, H. & MASHEITI, S. (2012) Power Generation  
511 from Waste Heat in a Food Processing Application. *Applied Thermal Engineering*, 36, 171-  
512 180.
- 513 ANEKE, M. & WANG, M. (2015a) Improving the Energy Efficiency of Cryogenic Air Separation  
514 Unit (ASU) through Compressor Waste Heat Recovery using Direct Binary Heat Engine  
515 Cycle. IN GERNAEY, V. K., HUUSOM, K. J. & GANI, R. (Eds.) *12th International  
516 Symposium on Process Systems Engineering and 25th European Symposium on Computer  
517 Aided Process Engineering*. Copenhagen, Denmark, Elsevier.
- 518 ANEKE, M. & WANG, M. (2015b) Potential for improving the energy efficiency of cryogenic air  
519 separation unit (ASU) using binary heat recovery cycles. *Applied Thermal Engineering*, 81,  
520 223 - 231.
- 521 BARBUCCI, P. (2008) The Enel's CCS Project. *Enel - Engineering and Innovation Division- Italy*.
- 522 CASTLE, W. F. (2002) Air separation and liquefaction: recent developments and prospects for the  
523 beginning of the new millennium. *International Journal of Refrigeration*, 25, 158 - 172.
- 524 CIFERNO, J. P., FOUT, T. E., JONES, A. P. & MURPHY, J. T. (2000) Capturing Carbon from  
525 Existing Coal-Fired Power Plants. *American Institute of Chemical Engineers*, April, 33 - 41.
- 526 HAGI, H., NEMER, M., LE MOULLEC, Y. & BOUALLOU, C. (2013) Assessment of the Flue Gas  
527 Recycle Strategies on Oxy-Coal Power Plants using an Exergy-based Methodology. *Chemical  
528 engineering Transactions*, 35, 343 - 346.
- 529 HONG, J., CHAUDHRY, G., BRISSON, J. G., FIELD, R. & GAZZINO, M. (2009) Anaysis of oxy-  
530 fuel combustion power cycle utilizing a pressurized coal combustor. *Energy*, 34, 1332 - 1340.
- 531 HU, Y. & YAN, J. (2011) Characterization of flue gas in oxy-coal combustion processes for CO<sub>2</sub>  
532 capture. *Applied Energy*, 90, 113 - 121.
- 533 IEA (2011) CO<sub>2</sub> Emissions from fuel combustion: Highlights. *International Energy Agency*.
- 534 IEA (2012) Key World Energy Statistics. *International Energy Agency*, [www.iea.org](http://www.iea.org).
- 535 IEA (2013) Key World Energy Statistics. *International Energy Agency*.

- 536 KANSHA, Y., KISHIMOTO, A., NAKAGAWA, T. & TSUTSUMI, A. (2011) A novel cryogenic air  
537 separation process based on self-heat recuperation. *Separation and Purification Technology*,  
538 77, 389 - 396.
- 539 LOMBARDI, L., CARNEVALE, E. & CORTI, A. (2011) Analysis of energy recovery potential using  
540 innovative technologies of waste gasification. *Waste Management*, 32, 640 - 652.
- 541 MANENTI, F., ROSSI, F., CROCE, G., GROTTOLI, M. G. & ALTAVILLA, M. (2013) Intensifying  
542 Air Separation Units. *Chemical engineering Transactions*, 35.
- 543 ROY, B. & BHATTACHARYA, S. (2013) Oxy-fuel Combustion Performance of Victorian Brown  
544 Coals in a 10 kW<sub>th</sub> Fluidized Bed. *3rd Oxyfuel Combustion Conference*. Ponferrada, Spain.
- 545 RÜBBERDT, K. (2009) Producing oxygen and nitrogen: Air separation techniques continue to  
546 advance. *ACHEMA 2009, 29th International Exhibition Congress in Chemical Engineering*,  
547 *Environmental Protection and Biotechnology*. Frankfurt am Main, Germany.
- 548 STRAHAN, D. (2013) Liquid Air Technologies - a guide to potential. *Centre for Cryogenic Energy*  
549 *Storage, University of Birmingham, UK*.
- 550 TSUKAHARA, H., ISHIDA, T. & MAYUMI, M. (1999) Gas-Phase Oxidation of Nitric Oxide:  
551 Chemical Kinetics and Rate Constant. *Ideal*, 3, 191 - 198.
- 552 WCA (2014) Coal & Electricity. World Coal Association, Available at: [www.worldcoal.org](http://www.worldcoal.org),  
553 Accessed 29th October: 2014.
- 554 WHITE, V., TORRENTE-MURCIANO, L., STURGEON, D. & CHADWICK, D. (2009) Purification  
555 of Oxyfuel-Derived CO<sub>2</sub>. *Energy Procedia*, 1, 399 - 406.
- 556 ZHENG, L. (Ed.) (2011) *Oxy-Fuel Combustion for Power Generation and Carbon Dioxide (CO<sub>2</sub>)*  
557 *Capture*, Woodhead, UK.
- 558
- 559

Figure

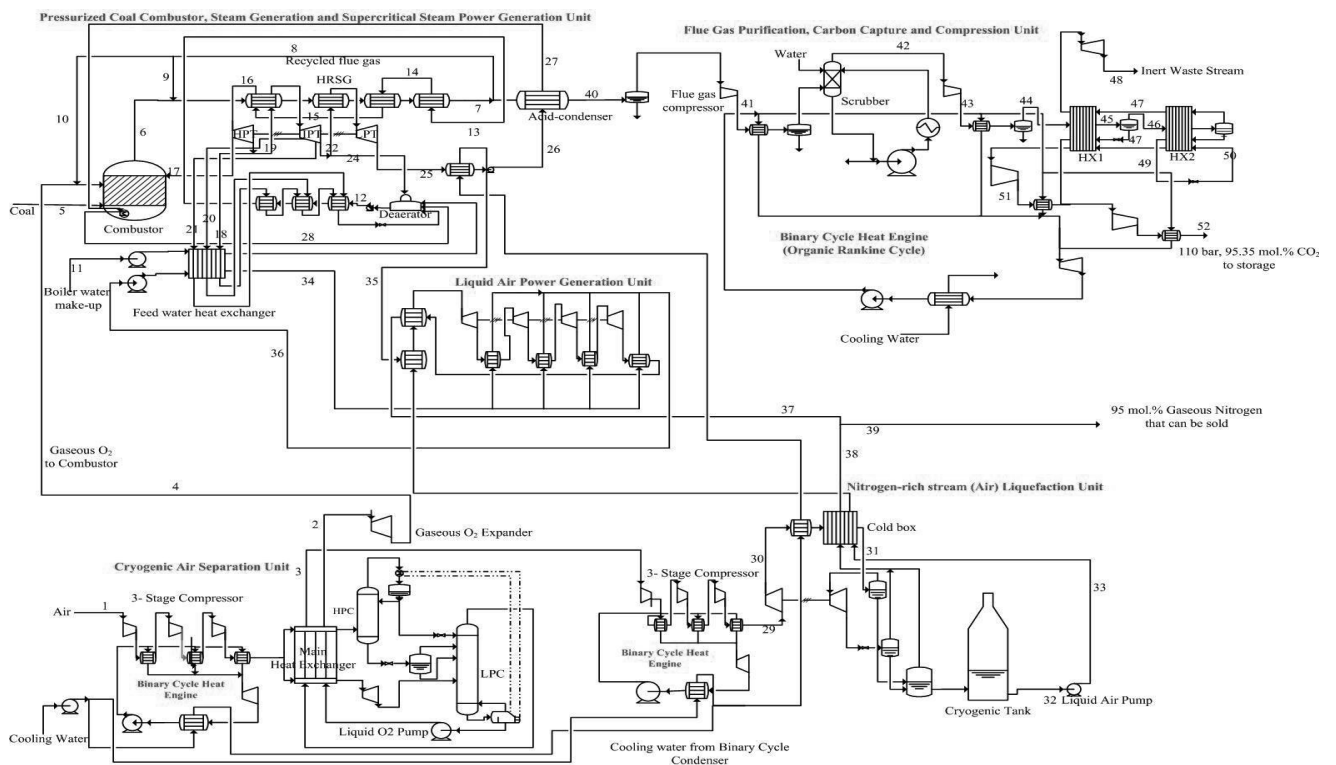
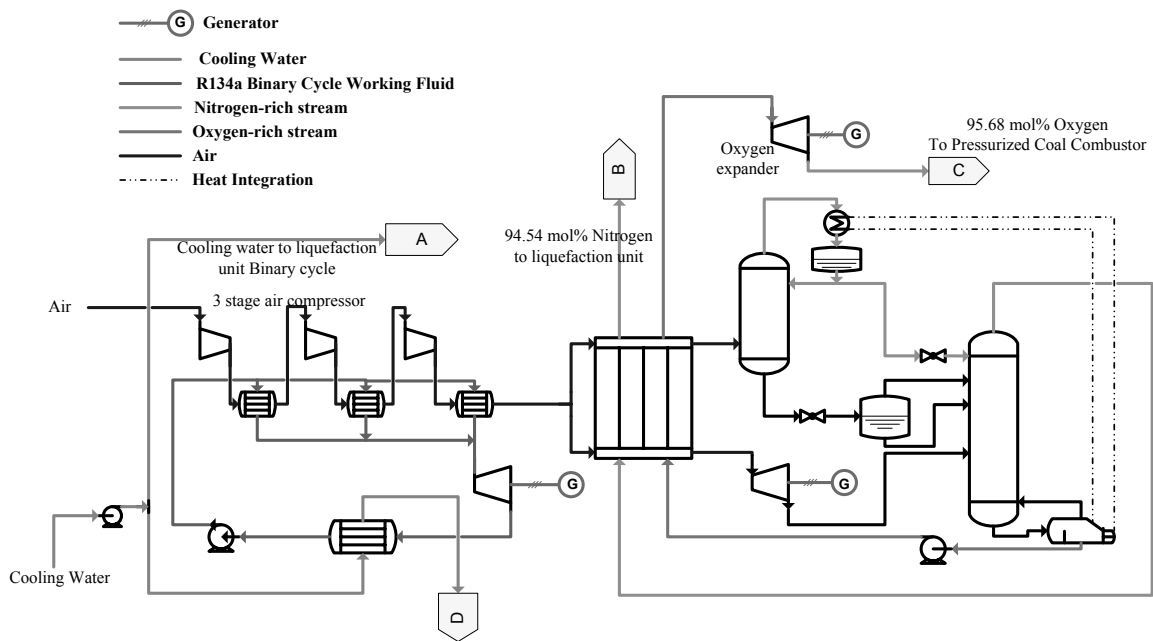
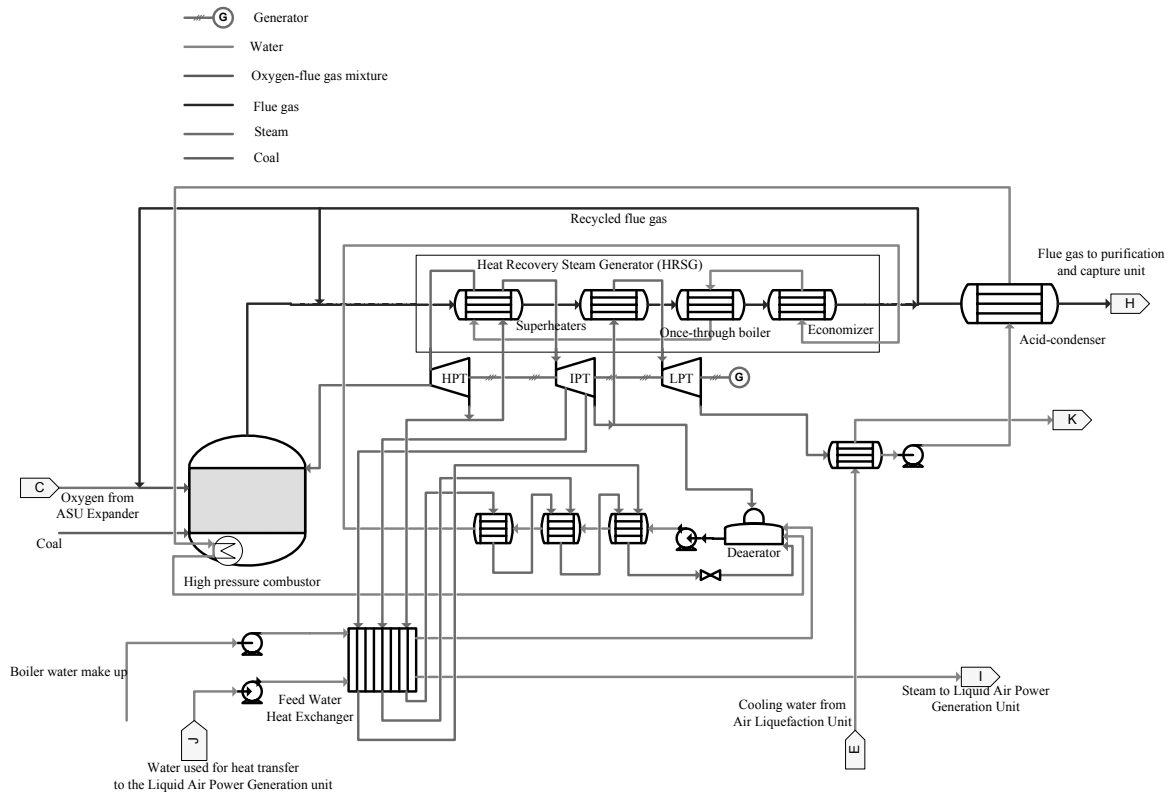


Figure 1: Overall Process Flow Diagram of the Pressurized Oxy-coal Power Cycle for Carbon Capture Application Integrated with LAPG and Binary Cycle Engine

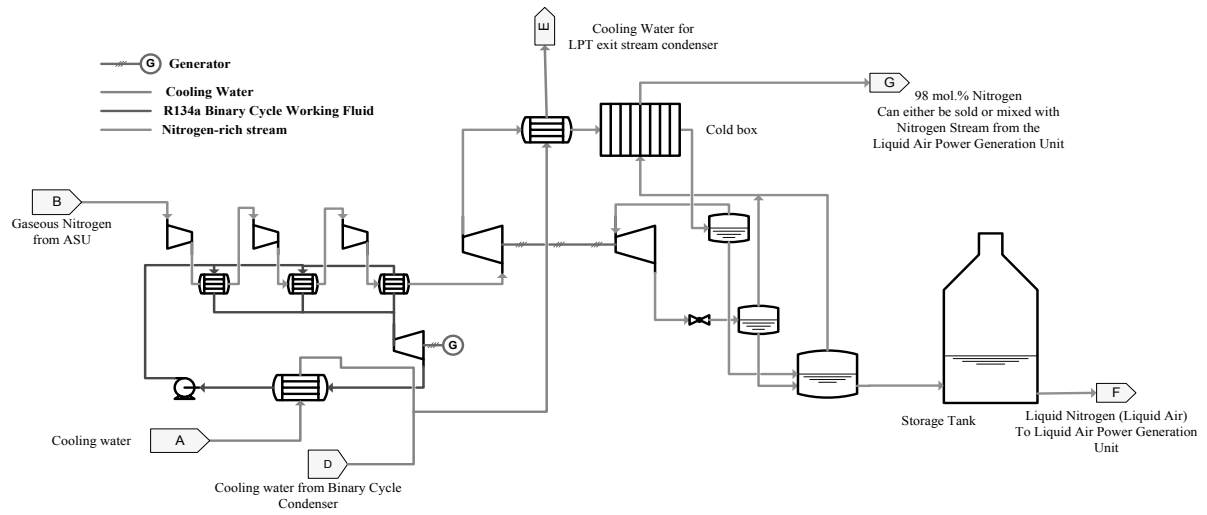


**Figure 2: Process Flow Diagram of Cryogenic ASU**

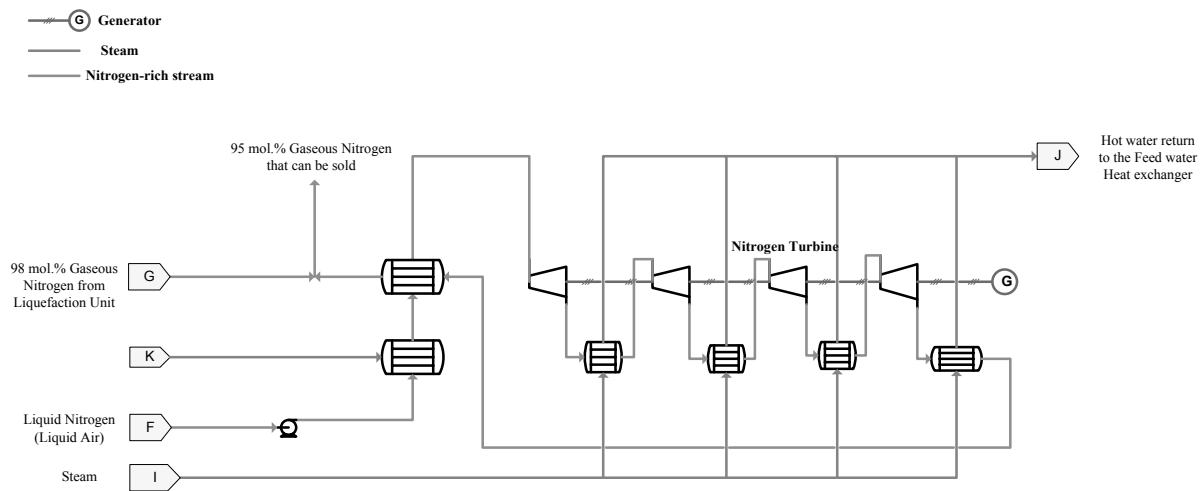




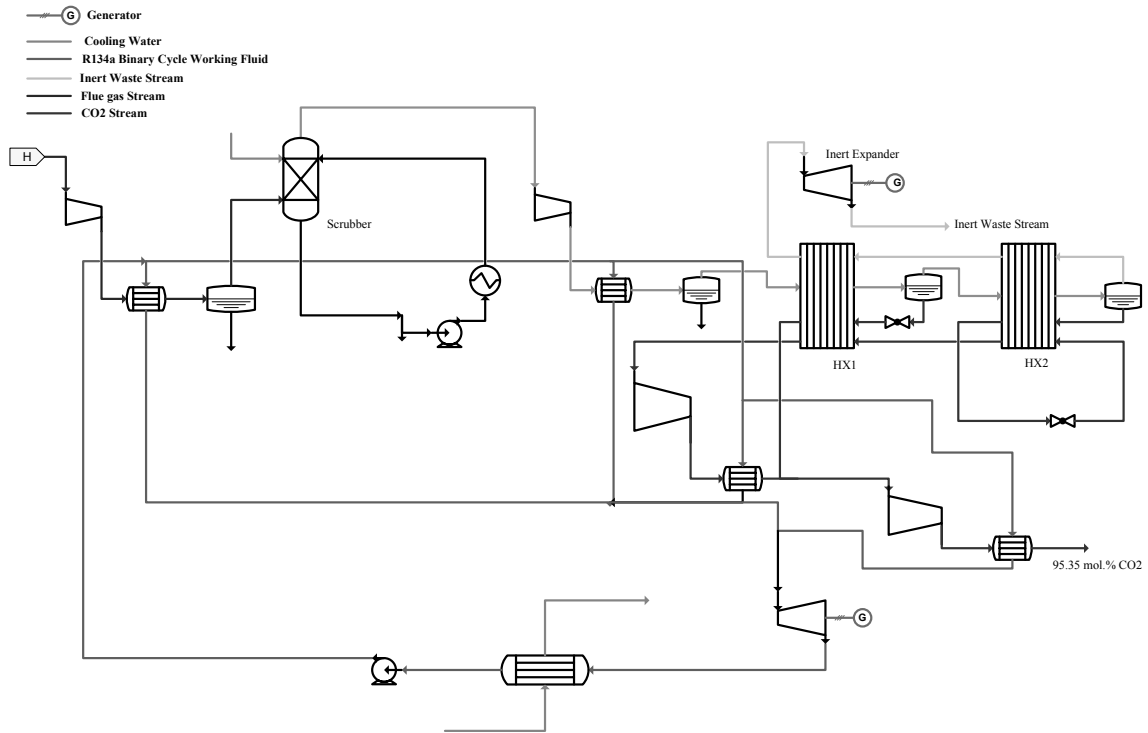
**Figure: 3 Process Flow Diagram of Pressurized Combustor Unit, Steam Generation Unit and Supercritical Steam Power Generation Unit**








**Figure 4: Process Flow Diagram of Air Liquefaction Unit**

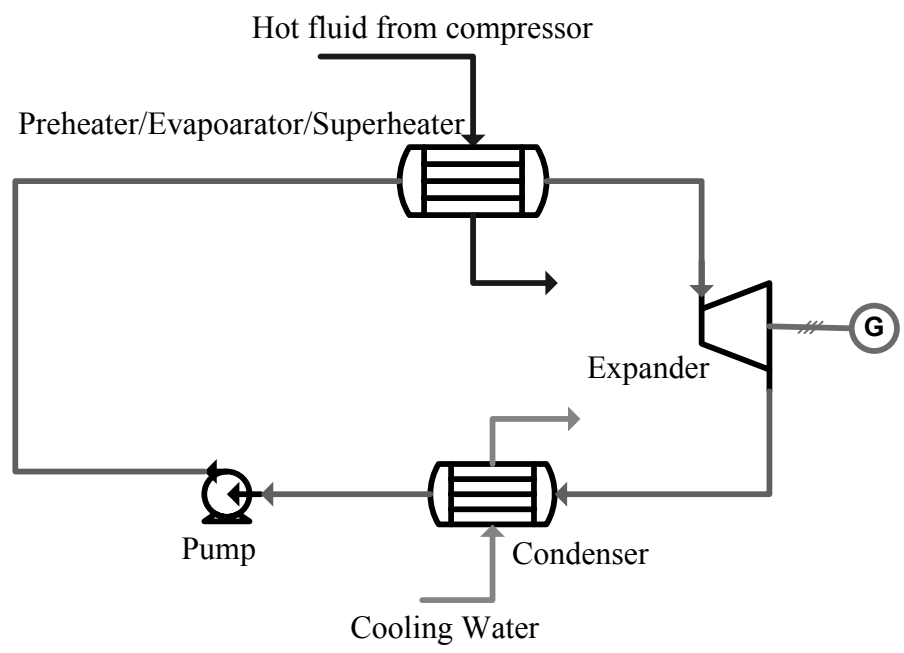


**Figure 5: Liquid Air Power Generation Unit**

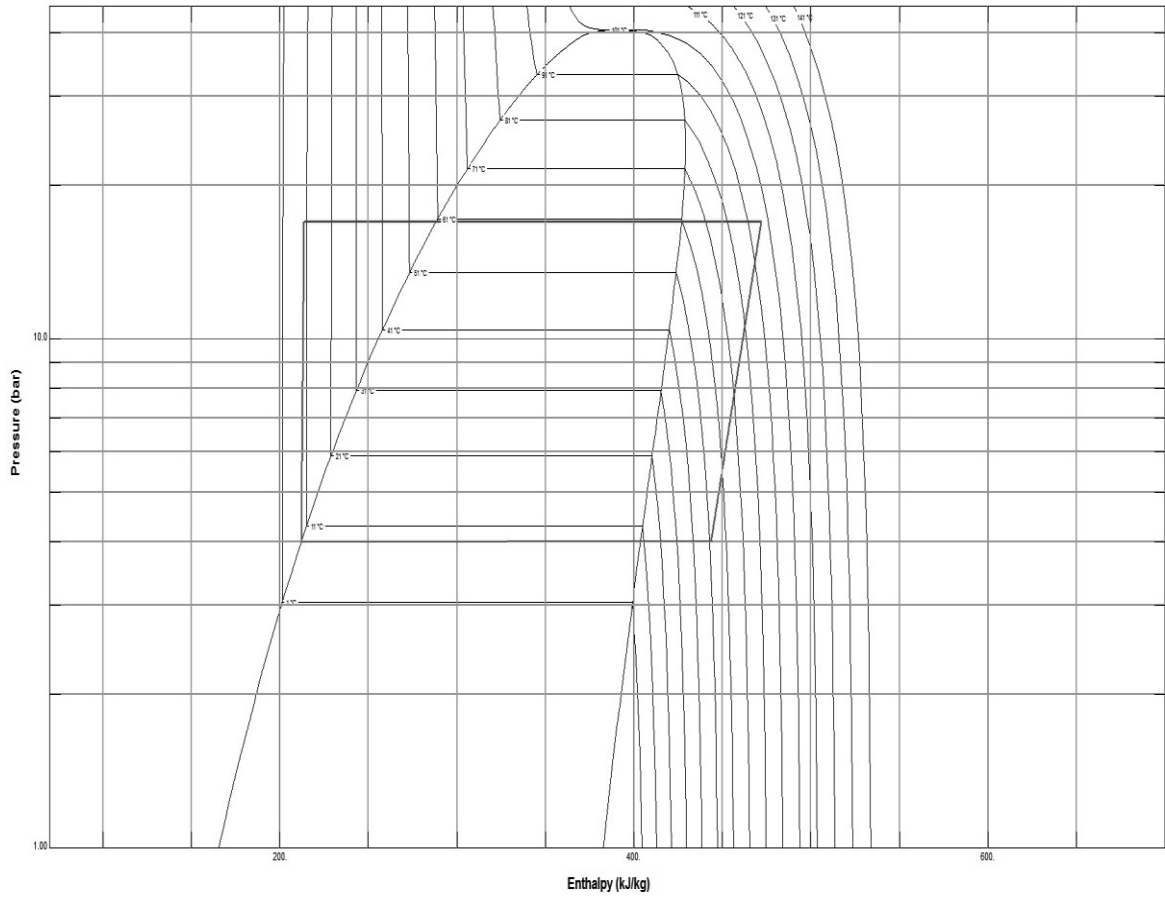


**Figure 6: Process Flow Diagram of flue gas Cleaning, CO<sub>2</sub> purification and compression unit**

-  **Generator**
-  **Cooling Water**
-  **R134a Binary Cycle Working Fluid**
-  **Nitrogen-rich stream**
-  **Hot fluid from compressor**



**Figure 7: Process Flow Diagram of Organic Rankine Cycle**



**Figure 8: P-h diagram of R134a used in the ORC system attached to the Cryogenic ASU**



Tables

**Table 1: Properties of coal used in the simulation (adapted from (Hu and Yan, 2011))**

<b>Coal Type</b>	<b>Bituminous</b>
<i>Moisture (wt.%) as analysed</i>	1.07
<i>Proximate (wt.%) (dry)</i>	
Ash	8.75
Volatile matter	35.33
Fixed carbon	54.85
<i>Ultimate (wt.%) (dry)</i>	
Carbon	77.65
Hydrogen	5.04
Nitrogen	1.49
Sulfur	0.95
Ash	8.84
Oxygen	6.03
Heating value (MJ/kg)	32.51



**Table 2: Process Simulation Parameters**

<b>Parameters</b>	<b>Value</b>
<b><i>Cryogenic ASU</i></b>	
Air mass flowrate	400 kg/s
Air composition (mol. %)	
Nitrogen	78.12
Oxygen	20.95
Argon	0.93
Compressor discharge pressure	6.35 bar
LPC Pressure	1.2 bar
HPC Pressure	5.7 bar
No of stages in LPC	69
No of stages in HPC	45
Liquid oxygen pump discharge pressure	200 bar
Gaseous oxygen expander inlet pressure	200 bar
<b><i>Pressurized Oxy-Coal Process</i></b>	
Coal mass flowrate	31.11 kg/s
Combustor pressure	10 bar
Combustor temperature	1550 °C
<b><i>Steam Generation Unit</i></b>	
Flue gas inlet temperature to HRSG	730 °C
Flue gas outlet temperature from HRSG	239 °C
<b><i>Supercritical Power Generation Unit</i></b>	
Turbine inlet pressure	250 bar
Turbine inlet temperature	600 °C
Reheat Temperature	620 °C
Deaerator Pressure	10 bars
<b><i>Air Liquefaction Unit</i></b>	
Compressor discharge pressure	8 bars
Liquid air temperature	-189 °C
Liquid air Pressure	1.99 bars
<b><i>Liquid Air Power Generation Unit</i></b>	
Turbine inlet pressure	120 bars
Reheat temperature	70 °C
<b><i>Flue Gas Cleaning, CO<sub>2</sub> purification and compression unit</i></b>	
Inert expander inlet pressure	30 bars
<b><i>Overall Process</i></b>	
Pump efficiency	0.80
Compression efficiency	0.80
Turbine Efficiency	0.80

**Table 3: Summary of Simulation Results**

<b>Performance Parameters</b>	<b>Value</b>
<b><i>Cryogenic ASU</i></b>	
<b><i>Materials</i></b>	
Oxygen stream mass flowrate (kg/s)	78.01
Oxygen stream composition (mol. %)	
Nitrogen	0.80
Oxygen	95.68
Argon	3.52
Nitrogen-rich stream mass flowrate (kg/s)	321.99
Nitrogen-rich stream composition (mol. %)	
Nitrogen	94.54
Oxygen	5.08
Argon	0.38
<b><i>Power Consumption &amp; Generation (kW)</i></b>	
Compressor Power consumption	89094.10
Liquid Oxygen Pump power consumption	1603.21
Gaseous Oxygen Expander Power output	8182.53
Air Expander Power output	2340.59
<b><i>Supercritical Power Generation Unit (kW)</i></b>	
Gross Turbine Power output	572595.28
Gross Pump Power consumption	22821.52
<b><i>Air (Nitrogen-rich stream) Liquefaction Unit Power Consumption (kW)</i></b>	
Compressor Power Consumption (Charging Power)	78219.00
<b><i>Liquid Air Power Generation Unit</i></b>	
Liquid Air Pump Power consumption (kW)	5048.40
Liquid Air Turbine Power output (kW)	125908.31
Thermal Energy Input from steam for Liquid Air Vapourization (kW <sub>th</sub> )	222639.45
<b><i>Flue Gas Cleaning, Carbon dioxide purification and Compression Unit</i></b>	
Flue gas purification and CO <sub>2</sub> compression train power consumption (kW)	37989.27
Scrubber recirculating stream pump power consumption (kW)	204.77
Inert Expander Power generation (kW)	184.29
CO <sub>2</sub> stream purity (mol. %)	95.35
CO <sub>2</sub> stream mass flowrate (kg/s)	89.30
<b><i>Binary Cycle Heat Engine Power (kW)</i></b>	
ASU compressor heat recovery binary cycle power output	9860.37
ASU binary cycle pump power consumption	443.46
Liquefaction compressor heat recovery binary cycle power output	7978.31
Liquefaction binary cycle pump power consumption	406.62
Flue gas purification and CO <sub>2</sub> compression binary cycle power output	25927.77
Flue gas purification and CO <sub>2</sub> compression binary cycle pump power consumption	1785.00

**Table 4: Stream Temperature, Pressure and Mass Flowrate**

<b>Stream no</b>	<b>Mass flowrate (kg/s)</b>	<b>Pressure (bar)</b>	<b>Temperature (°C)</b>
1	400.00	1.01	30.00
2	78.01	200.00	20.00
3	321.99	1.10	20.00
4	78.01	10.00	-139.50
5	31.11	10.00	25.00
6	803.85	10.00	1550.00
7	2106.73	10.00	239.07
9	1602.88	10.00	239.07
10	563.18	10.00	239.07
12	657.78	250.00	184.46
13	657.78	250.00	215.00
16	657.78	250.00	600.00
18	105.24	50.00	340.59
19	420.98	50.00	340.59
22	188.60	10.00	384.85
25	188.60	0.41	232.18
26	188.60	11.20	32.96
30	321.99	11.77	64.40
31	321.99	11.77	-163.00
32	289.15	1.99	-189.00
33	289.15	120.00	-184.08
40	240.67	10.00	167.19
41	199.69	15.00	217.91
42	92.03	15.00	26.05
43	92.03	30.00	86.93
44	91.95	30.00	40.00
45	91.95	30.00	-10.00
46	55.88	30.00	-10.00
47	2.70	30.00	-20.00
48	2.70	7.00	-61.94
49	53.17	30.00	-20.00
50	53.17	5.00	-61.52
51	53.17	20.00	161.41
52	89.24	110.00	37.78

**Table 5: Process Efficiencies**

Performance Parameters	Case 1	Case 2	Case 3	Case 4	Case 5	Case 6	Case 7
$Q_{in}$	1011386.10 <sup>a,b</sup>	1011386.10 <sup>a,b</sup>	1011386.10 <sup>a,b</sup>	1011386.10 <sup>a,b</sup>			1089605.10 <sup>a,b</sup>
$\Sigma Q_{scp}$					300858.45	300858.45	
$\Sigma G_P$	574935.87 <sup>a</sup> 575120.16 <sup>b</sup>	574935.87 <sup>a</sup> 575120.16 <sup>b</sup>					
$\Sigma G_{P2}$			583118.40 <sup>a</sup> 583302.69 <sup>b</sup>				
$\Sigma G_{P3}$				592978.77 <sup>a</sup> 593163.06 <sup>b</sup>			
$G_{la}$					125908.31		
$\Sigma G_{ov}$							724524.80 <sup>a</sup> 750636.86 <sup>b</sup>
$\Sigma G_{la1}$						133886.62	
$\Sigma D_{asu}$	132442.53 <sup>a</sup> 170636.57 <sup>b</sup>						
$\Sigma D_{asup}$		111847.11 <sup>a</sup> 150041.15 <sup>b</sup>	113379.74 <sup>b</sup> 151573.78 <sup>b</sup>				
$\Sigma D_{asupp}$				113823.20 <sup>a</sup> 152017.24 <sup>b</sup>			
$D_{aux}$					5048.40		
$\Sigma D_{aux1}$						5455.02	
$\Sigma D_{ov}$							114229.83 <sup>a</sup> 154208.87 <sup>b</sup>
<b>Efficiency</b>	$\mu_{ox}$	$\mu_{ox1}$	$\mu_{ox2}$	$\mu_{ox3}$	$\mu_{la}$	$\mu_{la1}$	$\eta_{RT}$
	43.75% <sup>a</sup> 39.99% <sup>b</sup>	45.79% <sup>a</sup> 42.03% <sup>b</sup>	46.45% <sup>a</sup> 42.69% <sup>b</sup>	47.38% <sup>a</sup> 43.62% <sup>b</sup>	40.17%	42.69%	56.01% <sup>a</sup> 54.74% <sup>b</sup>

All parameters are in kW; (a) means without carbon capture; (b) means with carbon capture