This is the peer reviewd version of the followng article:

Effects of upgrading systems on energy conversion efficiency of a gasifier - fuel cell - gas turbine power plant / Pedrazzi, Simone; Allesina, Giulio; Tartarini, Paolo. - In: ENERGY CONVERSION AND MANAGEMENT. - ISSN 0196-8904. - 126:(2016), pp. 686-696. [10.1016/j.enconman.2016.08.048]

Terms of use:

The terms and conditions for the reuse of this version of the manuscript are specified in the publishing policy. For all terms of use and more information see the publisher's website.

28/04/2024 14:32

Elsevier Editorial System(tm) for Energy

Conversion and Management

Manuscript Draft

Manuscript Number: ECM-D-16-03320R1

Title: Effects of upgrading systems on energy conversion efficiency of a gasifier - fuel cell - gas turbine power plant

Article Type: Original research paper

Section/Category: 3. Clean Energy and Sustainability

Keywords: Biomass; Gasification; Modeling; Solide Oxide Fuel Cells; Zeolites; PPO membrane

Corresponding Author: Dr. Simone Pedrazzi, Ph.D.

Corresponding Author's Institution: University of Modena and Reggio Emilia

First Author: Simone Pedrazzi, Ph.D.

Order of Authors: Simone Pedrazzi, Ph.D.; Giulio Allesina, Ph.D.; Paolo Tartarini, Full professor

Abstract: This work focuses on a DG-SOFC-MGT (downdraft gasifier- solid oxide fuel cell - micro gas turbine) power plant for electrical energy production and investigates two possible performance-upgrading systems: polyphenylene oxide (PPO) membrane and zeolite filters. The first is used to produce oxygen-enriched air used in the reactor, while the latter separates the CO2 content from the syngas. In order to prevent power plant shutdowns during the gasifier reactor scheduled maintenance, the system is equipped with a gas storage tank. The generation unit consists of a SOFC-MGT system characterized by higher electrical efficiency when compared to conventional power production technology (IC engines, ORC and EFGT). Poplar wood chips with 10% of total moisture are used as feedstock. Four different combinations with and without PPO and zeolite filtrations are simulated and discussed. One-year energy and power simulation were used as basis for comparison between all the cases analyzed. The modeling of the gasification reactions gives results consistent with literature about oxygen-enriched processes. Results showed that the highest electrical efficiency obtained is 32.81%. This value is reached by the power plant equipped only with PPO membrane filtration. Contrary to the PPO filtering, zeolite filtration does not increase the SOFC-MGT unit performance while it affects the energy balance with high auxiliary electrical consumption. This solution can be considered valuable only for future work coupling a CO2 sequestration system to the power plant.



Department of Engineering "Enzo Ferrari" University of Modena and Reggio Emilia Via Vivarelli, 10/1 - 41125 Modena, Italy

Dr. Simone Pedrazzi



BEELab (Bio Energy Efficiency Laboratory) www.beelab.unimore.it

Department of Engineering "Enzo Ferrari" University of Modena and Reggio Emilia Via Vivarelli, 10/1 – 41125 Modena, Italy phone +39 059 205 6229, email: simone.pedrazzi@unimore.it

Professor Moh' Ahmad Al-Nimr Editor-in-Chief Energy Conversion and Management

Dear Professor Moh' Ahmad Al-Nimr:

I am pleased to submit an original research article entitled "**Effects of upgrading systems on energy conversion efficiency of a gasifier - fuel cell - gas turbine power plant**" by Simone Pedrazzi, Giulio Allesina and Paolo Tartarini for consideration for publication in the *Energy Conversion and Management* journal.

This work focuses on a downdraft gasifier- solid oxide fuel cell - micro gas turbine power plant for electrical energy production and investigates two possible performance-upgrading systems: polyphenylene oxide (PPO) membrane and zeolite filters. The first is used to produce oxygenenriched air before the reactor, while the latter separates the CO₂ content from the syngas. In order to prevent power plant shutdowns during the gasifier reactor scheduled maintenance, the system is equipped with a gas storage tank. The generation unit consists of a SOFC-MGT system characterized by higher electrical efficiency when compared to conventional power production technology (IC engines, ORC and EFGT). Poplar wood chips with 10 % of total moisture are used as feedstock. Four different combinations with and without PPO and zeolite filtrations are simulated and discussed. One-year energy and power simulation were used as basis for comparison between all the cases analyzed. The modeling of the gasification reactions gives results consistent with literature about oxygen-enriched processes. Results shown that the highest electrical efficiency obtained is 32.81 %. This value is reached by the power plant equipped only with PPO membrane filtration. Contrary to the PPO filtering, zeolite filtration does not increase the SOFC-MGT unit performance while it affects the energy balance with high auxiliary electrical consumption. This solution can be considered valuable only for future work coupling a CO₂ sequestration system to the power plant.

Furthermore, with this letter the authors certify that the content of the paper and its novelty results are the original work of the authors and it was not submitted before in this journal. With this letter all the authors mutually agree to submit their manuscript to Energy Conversion and Management. If you felt that the manuscript is appropriate for your journal, we suggest the following reviewers:

- Prof. S. Dasappa, Faculty, Center for Sustainable Technologies, Indian Institute of Science;
- Dr. Pierluigi Leone, Department of Energy, Polytechnic University of Turin;



Department of Engineering "Enzo Ferrari" University of Modena and Reggio Emilia Via Vivarelli, 10/1 - 41125 Modena, Italy



BEELab (Bio Energy Efficiency Laboratory) www.beelab.unimore.it

- Dr. Florian Monlau, APESA, Plateau Technique, France.

We look forward to hearing from you soon.

Best regards,

Dr. Simone Pedrazzi Dr. Giulio Allesina Pror. Paolo Tartarini

Highlights

- An advanced gasifier-SOFC-MGT system is modeled.
- An overall electrical efficiency of 32.81% is reached.
- Influence of all the sub-system modeled on the power plant efficiency is discussed.
- Compression storage of syngas is taken into account.

Effects of upgrading systems on energy conversion efficiency of a gasifier - fuel cell - gas turbine power plant

Simone Pedrazzi^{a,*}, Giulio Allesina^a, Paolo Tartarini^a

^aUniversity of Modena and Reggio Emilia, Department of Engineering 'Enzo Ferrari', Via Vivarelli 10/1, 41125 Modena, Italy

Abstract

THIS PAPER IS SUBMITTED WITH THE OPTION 'YOUR PAPER YOUR WAY'. FOR THIS REASON LAYOUT AND STYLE MAY DIFFER FROM THE JOURNAL ONE.

This work focuses on a DG-SOFC-MGT (downdraft gasifier- solid oxide fuel cell - micro gas turbine) power plant for electrical energy production and investigates two possible performance-upgrading systems: polyphenylene oxide (PPO) membrane and zeolite filters. The first is used to produce oxygenenriched air used in the reactor, while the latter separates the CO_2 content from the syngas. In order to prevent power plant shutdowns during the gasifier reactor scheduled maintenance, the system is equipped with a gas storage tank. The generation unit consists of a SOFC-MGT system characterized by higher electrical efficiency when compared to conventional power production technology (IC engines, ORC and EFGT). Poplar wood chips with 10% of total moisture are used as feedstock. Four different combinations with and without PPO and zeolite filtrations are simulated and discussed. One-year

Preprint submitted to Energy Conversion and Management

^{*}Corresponding author

Email address: simone.pedrazzi@unimore.it (Simone Pedrazzi)

energy and power simulation were used as basis for comparison between all the cases analyzed. The modeling of the gasification reactions gives results consistent with literature about oxygen-enriched processes. Results showed that the highest electrical efficiency obtained is 32.81%. This value is reached by the power plant equipped only with PPO membrane filtration. Contrary to the PPO filtering, zeolite filtration does not increase the SOFC-MGT unit performance while it affects the energy balance with high auxiliary electrical consumption. This solution can be considered valuable only for future work coupling a CO_2 sequestration system to the power plant.

Keywords:

Biomass, Gasification, Modeling, Solide Oxide Fuel Cells, Zeolites, PPO membrane

1 1. Introduction

Due to the abundant availability and distribution, biomasses hold keyroles in plans for renewable energy production. This trend is becoming even more relevant thanks to the good degree of reliability and efficiency of the biomass-based technologies together with the high subsidies granted by several government for sustainable electrical energy production [1].

Depending on the feedstock quality and availability, biomasses are converted into energy through different technologies. In the case of ligno-cellulosic biomasses, a technolgy of great validity is gasification. This thermo-chemical process turns solid biomass into a gaseous fuel known as syngas, which can be converted into electrical energy through all those systems used for power production from gaseous fuels [2]. Gasification is today one of the most effi¹³ cient technologies to convert wood into electricity and it is also sustainable ¹⁴ in terms of the environmental balance of CO_2 [3, 4].

Most of the gasification power plants use an IC engine-generator to con-15 vert the syngas chemical energy into electrical power. However, in some cases 16 other conversion machines are used, i.e. Organic Rankine Cycles (ORC), Ex-17 ternal Firing Gas Turbines (EFGT)[5] and Stirling engines are used with the 18 major advantage of having minor limitation about the syngas level of purifica-19 tion [2, 6, 7, 8, 9]. These systems are usually characterized by low conversion 20 efficiencies of about 10-12%. Major conversion rates can be obtained only 21 with electrochemical devices such as proton exchange membrane fuel cells 22 [10], Molten Carbonate Fuel Cells (MCFC) [11, 12], Solid Oxide Fuel Cells 23 (SOFC) [13, 14, 15], systems composed of SOFC and Micro Gas Turbines 24 (MGT) [16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26] and systems composed of 25 SOFC-MGT-ORC [27]. Despite the high rate of energy conversion, these 26 systems require perfectly clean syngas [28]. Downdraft gasifiers are the most 27 suitable architecture due to the low tar and particulate content in their gas 28 when compared to updraft, crossdraft or fluidized bed gasifiers [2, 6, 29]. 29 However, downdraft gasifiers commonly use air as gasification agent. This 30 solution generates a syngas with a low calorific value where the hydrogen, 31 methane and carbon monoxide are diluted in non-burnable gases: N_2 (about 32 50%) and CO_2 (from 10 to 20%). Otherwise, it is possible to choose oxygen 33 gasification that produces a syngas with negligible N_2 content. However, oxy-34 gen gasification is a complex and expensive technology due to the gasification 35 agent supply sub-systems and reactor material choice. Indeed, temperatures 36 inside the reactor can reach 1200-1300 K when oxygen is used instead of air 37

зв [30].

The basic system discussed in this study is composed of an air blowndowndraft-fixed bed gasifier fed with poplar wood chips. This work is aimed at investigating the effects of different power plant designs on the overall energy conversion efficiency.

The first power plant upgrading sub-system consists of a polyphenylene 43 oxide (PPO) membrane used to produce oxygen-enriched air. The gas sep-44 aration characterization of this membrane is reported in literature [31, 32]. 45 In practice, membrane gas separation is applied to increase the oxygen con-46 tent in the inlet air of biomass boilers [33]. Bisio et al. studied the ther-47 modynamics of combustion with enriched air and reviewed several types of 48 memebranes [34]. Coombe and Nieh developed a membrane-based device for 49 air enrichment in small scale burners [35]. Hao et al. applied an oxygen-50 permeable membrane to a reactor for the co-production of dimethyl ether 51 (DME)/methanol and electricity [36]. This paper uses PPO membrane in 52 order to obtain air with about 50% of oxygen then used as gasification agent. 53 This solution is a hybrid between air and pure oxygen gasification. Enriched 54 air reduces the reactor thermal stress compared to pure oxygen gasification, 55 while the syngas has a lower N_2 content than the one obtained in pure air 56 gasification. In addition, the syngas flow rate decreases because, for a fixed 57 power output, the enriched air flow required for gasification is lower than air 58 used in conventional gasification. This happens because the same amount of 59 oxygen is used in both cases and its concentration in enriched air is higher 60 than untreated air. Finally, the tar production is lower than air gasification 61 as consequence of the higher temperature that cracks more efficiently the 62

⁶³ primary tars from pyrolysis [37].

A second solution discussed in this work consists of a porous media used 64 to upgrade the syngas. In fact, syngas has a variable CO_2 content depending 65 on gasification process as well as several boundary conditions. This value 66 ranges from 10% to 30% and it reduces significantly the higher heating value 67 of the syngas [37]. A solution to overcome this issue is to adopt a pressure-68 swing selective synthetic zeolite filter. This system is placed before the gas 69 storage in order to separate carbon dioxide from syngas [38, 39]. The filter 70 can be constantly regenerated using a rotary valve packaged into modules 71 as described by Tagliabue et al. [40]. Literature investigation about zeolite 72 filtration outlines several works. Bacsik et al. studied the biogas CO_2 - CH_4 73 separation through zeolites [41]. Kacem et al. investigated the pressure swing 74 adsorption for CO_2/N_2 and CO_2/CH_4 separation using activated carbon and 75 several types of zeolites [42]. Dirar et al. investigated intrinsic adsorption 76 properties of CO2 on 5A and 13X zeolite [43]. 77

The syngas obtained from gasification is stored and then used in a SOFC 78 unit able to produce electrical and thermal energy. The number of stacks 79 within the cell is optimized taking into account the optimal electrical cur-80 rent density. The chosen number guarantees a good efficiency, however the 81 gas discharged from the cell still contains some chemical energy. For this 82 reason, this work suggests to convert this residual energy in a micro gas tur-83 bine (MGT). The syngas storage allows the generation unit to operate in its 84 optimal point, furthermore it prevent the power plant shoutting down dur-85 ing the maintenance operations of the gasifier. This management preserves 86 the SOFC and MGT reliability. However, it is difficult to design the stor-87

age capacity because an oversize storage rises the systems costs, while an undersized capacity reduces the time gained for the maintenance. For this reason, the storage was designed taking into account the tanks pressure, the electrical power production of the SOFC-MGT unit and the time required for scheduled stops of the gasifier for maintenance operations.

The mathematics of the whole system was developed starting from literature. The overall model has been implemented in Matlab SimulinkTM software environment in order to simulate the behavior of the system under different conditions over a year long simulation.

97 2. System modeling

The basic system layout is reported in Figure 1. The most relevant components are:

- Downdraft gasifier: The gasifier is equipped with a subsystem for the syngas filtering and cooling with water scrubber and electrostatic filters.
- Syngas storage: It consists of a tank of a total volume of 650 m³.
- SOFC unit: This subsystem consists of 10875 solid oxide cells and it is connected to the electrical grid by a power inverter.
- Micro gas turbine (MGT): this turbo-machinery is used to convert the last part of chemical energy content in the syngas purged by the SOFC.

This work investigates the effect of the implementation of the following sub-systems to the basic scheme:

• **PPO membrane filter module:** The PPO sub-system consists of the membrane filter and a compressor that increases the pressure of the air before the PPO membrane filter to about 1 MPa. The oxygen enriched air is sent to the gasifier at atmospheric pressure. A flow of nitrogen is purged from the PPO module.

• Zeolite (ZEO) filter module: the zeolite (ZEO) filter module is placed after the first syngas compressor. There is a further syngas compression stage ahead the storage tanks because the ZEO module works at 0.5 MPa of pressure as described in Section 2.3, while the pressure in the storage is often higher.

The syngas is used as fuel in the SOFC stack. In this device, the fuel reforming occurs at the anode and there is a recirculation of the 20% of the anode exhaust to increase the fuel reforming performance [18, 22]. The anode exhaust is used to preheat the syngas, then it is finally burned in the MGT burner together with the cathode exhaust. The air required for the electrochemical reaction is compressed and preheated in the recuperator of the MGT as well as in the air preheater of the SOFC.

The SOFC stack generates DC current which is converted into AC current by an inverter and it is sent to the electrical grid. The MGT drags the air compressor and the remaining mechanical energy is converted into electrical energy by an alternator.

132 2.1. PPO module modeling

Polymeric membranes allow to separate different gaseous components de-133 pending on the pore size and pressure applied to the filter [31]. In this work 134 a membrane is used to separate nitrogen from air. The membranes widely 135 used for this purpose are: Matrimid, Polyphenylenoxide (PPO) and Poly-136 dimethylsiloxan (PDMS) [31]. As showed in Figure 2, in membranes the 137 inlet air flow is divided in permeate and retentate molar flows. The inlet 138 flow $(Q_{air} \text{ [mol/s]})$ has a pressure p_{feed} [atm] and it is composed of x_{O_2Feed} 139 and x_{N_2Feed} molar fractions of oxygen and nitrogen. The permeate molar 140 flow $(Q_P \text{ [mol/s]})$ has a pressure $p_{permeate}$ [atm] and it is composed of y_{O_2} 141 and y_{N_2} molar fractions of oxygen and nitrogen. The retentate molar flow 142 $(Q_R \text{ [mol/s]})$ has a pressure p_{feed} [atm] and it is composed of $x_{O_2Retentate}$ and 143 $x_{N_2Retentate}$ molar fractions of oxygen and nitrogen. 144

Each membrane behavior is identified through two parameters: the selectivity (α) and the permeability to oxygen (γ). The first factor represents the attitude of the membrane to attract oxygen, the second quantifies the attitude of the membrane to be crossed by it. High selectivity and permeability ensure great filtering performance in terms of high value of y_{O2} and a small membrane surface area is required to filter a given amount of air. Table 1 presents the parameters of Matrimid, PPO and PDSM membranes.

The choice of a PPO membrane is a compromise in terms of acceptable values of selectivity and permeability. In order to simulate the behavior of the membranes, a mathematical model has been implemented from Melin and Rautenbach [31]. The model is based on the following assumptions:

156

• Air is considered a binary gas mixture with 21% oxygen and 79% ni-

157 trogen.

- Steady state conditions.
- Isotherm conditions.
- Isobaric conditions.
- Perfect gas law.
- Constant permeability.
- Perfect mixing conditions on upstream and downstream sides.
- Concentration polarization at the membrane is neglected.
- Pressure loss in the porous support layer is neglected.
- The permeate can drain off freely.

The calculation of the permeate composition is made with the following formula taken from the work of Melin and Rautenbach [31]:

$$y_{O_2} = \frac{1}{2} \left[1 + \phi * \left(x_{O_2Feed} + \frac{1}{\alpha - 1} \right) \right] - \sqrt{\left[\frac{1}{2} \left[1 + \phi * \left(x_{O_2Feed} + \frac{1}{\alpha - 1} \right) \right] \right]^2 - \frac{\alpha * \phi * x_{O_2Feed}}{\alpha - 1}}$$
(1)

$$y_{N_2} = 1 - y_{O_2} \tag{2}$$

where ϕ [-] is the feed-permeate pressure ratio given by the following equation:

$$\phi = p_{feed} / p_{permeate} \tag{3}$$

Figure 3 reports the permeate composition over pressure ratio for the 171 three membrane types considered. It can be seen that only a certain maxi-172 mum of oxygen ratio can be achieved because all the graphs are leveling off. 173 Therefore a pressure ratio of 1 MPa was chosen for further calculations as 174 suggested in Melin and Rautenbach [31] and the $p_{retentate}$ was fixed at 1 atm. 175 The Matrimid membrane is able to produce the highest oxygen ratio 176 of 0.58 % vol. in the permeate, however PPO membrane presents a good 177 value of oxygen ratio (0.49 % vol.) and an acceptable value of permeability, 178 therefore this membrane is adopted in the simulations. The active area of 179 the membrane can be assessed from the molar flow of oxygen required for 180 the gasification Q_{PO_2} [mol/s]: 181

$$A_{membrane} = \frac{Q_{PO_2}}{\gamma * (x_{O_2Feed} * p_{feed} + y_{O_2} * p_{permeate})}$$
(4)

The molar flow of nitrogen Q_{PN_2} [mol/s] and the total permeate molar flow Q_P [mol/s] is given by the following equations:

$$Q_{PN_2} = \frac{\gamma}{\alpha} * \left[p_{feed} * (1 - x_{O_2Feed}) + p_{permeate} * (1 - y_{O_2}) \right]$$
(5)

$$Q_P = Q_{PN_2} + Q_{PO_2} \tag{6}$$

The molar flow of the inlet air Q_{air} , the retentate molar flow Q_R and the retentate composition ($x_{O_2Retentate}$ and $x_{N_2Retentate}$) are calculated setting to zero the amount of oxygen in the retentate flow as suggested by Melin and ¹⁸⁷ Rautenbach[31]. Thus, a mass balance equation can be applied to estimate ¹⁸⁸ Q_{air} and Q_R :

$$Q_{air} * x_{O_2Feed} = Q_P * y_{O_2} \rightarrow Q_{air} = \frac{Q_P * y_{O_2}}{x_{O_2Feed}}$$
(7)

$$Q_{air} = Q_P + Q_R \to Q_R = Q_{air} - Q_P \tag{8}$$

Finally, the electrical power consumption to pressurize the inlet air flow is calculated as a polytropic compression by Equation 15 assuming $T_{in} = 20$ o C; m = 1.2 and $\eta_{comp} = 90$ %.

192 2.2. Gasifier modeling

In this work, the gasification process is simulated using a black-box model based on Barman's work [44]. The model is validated for downdraft gasifiers; it is based on the following gasification equation:

$$CH_{x}O_{y}N_{z} + wH_{2}O + m(O_{2} + 3.76N_{2}) \rightarrow$$

$$n_{H_{2}}H_{2} + n_{CO}CO + n_{CO_{2}}CO_{2} + n_{H_{2}O}H_{2}O$$

$$+ n_{CH_{4}}CH_{4} + (z/2 + 3.76m)N_{2} + n_{tar}CH_{p}O_{q}$$
(9)

where $CH_xO_yN_z$ is the equivalent chemical formula of "dry and ash free" (daf) biomass; CH_pO_q is the equivalent chemical formula of tar [45]; $w [mol/mol_{bio}]$ is the specific molar amount of the biomass moisture; m $[mol/mol_{bio}]$ is the specific molar amount of oxygen calculated starting from the equivalence ratio ER as suggested by Jarungthammachote and Dutta[46]; $n_{H_2}, n_{CO}, n_{CO_2}, n_{H_2O}, n_{CH_4}, n_{tar} [mol/mol_{bio}]$ are the specific molar amounts of $H_2, CO, CO_2, H_2O, CH_4$ and tar which constitute the syngas. This model is used and discussed in several other works [47, 48, 49]. It consists of a chemical and a thermal sub-models that converge to the final composition of the gas. The first step is to choose an initial temperature T[K] and calculate the equilibrium constant of the following reactions:

• **K1**: Water-gas shift
$$CO + H_2O \leftrightarrow CO_2 + H_2$$

- **K2:** Hydrogasification $C + 2H_2 \leftrightarrow CH_4$
- **K3:** Methane steam reforming $CH_4 + H_2O \leftrightarrow CO + 3H_2$

The system of equations 10 reported below is composed of three chemical balances calculated from Equation 9 (carbon, hydrogen and oxygen) and the three equilibrium constants for water-gas, hydrogasification and methane reforming reactions. The system is solved with the Newton-Raphson method.

$$n_{CO} + n_{CO_2} + n_{CH_4} + n_{tar} - 1 = 0$$

$$2n_{H_2} + 2n_{H_2O} + 4n_{CH_4} + pn_{tar} - x - 2w = 0$$

$$n_{CO} + 2n_{CO_2} + n_{H_2O} + qn_{tar} - w - 2m - y = 0$$

$$K_1 = \frac{n_{CO_2} * n_{H_2}}{n_{CO} * n_{H_2O}}$$

$$K_2 = \frac{n_{CH_4} * \frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}}{n_{H_2}^2}$$

$$K_3 = \frac{n_{CO} * n_{H_2}^3}{\left(\frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}\right)^2 n_{H_2O} n_{CH_4}}$$
(10)

Once the molar specific amounts of the syngas species are evaluated, it is possible to solve the thermodynamic energy balance of the system reported in the following equation:

$$\sum_{j=react} n_j * HF_j^0 = \sum_{i=prod} n_i * \left(HF_i^0 + \Delta H_{T,i} \right)$$
(11)

where n_j [moles] and HF_j^0 [kJ/kmol] are the specific moles amount and standard heat of formation of the j-th reagent (biomass, air and moisture); n_i [moles] and HF_i^0 [kJ/kmol] are the specific moles amount and the standard heat of formation of the i-th product (H_2 , CO, CO_2 , H_2O , CH_4 and N_2) and $\Delta H_{T,i}$ is the enthalpy difference between any given state and the standard state for the i-th product. $\Delta H_{T,i}$ can be calculated starting from the specific heat of the product:

$$\Delta H_{T,i} = \int_{298.15}^{T} C_p(T) \, dT = \left| aT + b\frac{T^2}{2} + c\frac{T^3}{3} + d\frac{T^4}{4} \right|_{298.15}^{T} \tag{12}$$

where the coefficient a,b,c and d are defined for each gas by Jarungtham-224 machote and Dutta[46]. In order to find the equilibrium temperature T_{new} , 225 the system is considered adiabatic and the the Newton-Raphson method is 226 applied to the equations. If $abs(T - T_{new}) < 0.1$ K then the calculated equi-227 librium temperature and molar specific gases amounts are the final results; 228 otherwise, a new iteration is done in order to satisfy the previous condi-229 tion. The model is implemented in Python and the input are the biomass 230 equivalent molecule, the equivalence ratio ER and the initial temperature. 231 The temperature input is used only as a starting point for the iterating 232 system; after few cycles the temperature converges to the ones that satisfy 233 both the chemical and thermal sub-systems. About the ER, a value of 0.335234

is assumed. This value is consistent with air blown gasification parameters
[50, 37] and it is confirmed by the low tar content in the syngas. Poplar wood
chips properties and gasifier model parameters are summarized in Table 1.

238 2.3. ZEO module modeling

The zeolite filter is able to reduce the total syngas molar flow of about 240 20% - 30 % by the adsorption of CO_2 . Zeolite 5A is chosen because it has 241 a great selectivity for carbon dioxide in comparison with the other gases 242 that constitute syngas [38]. The gas adsorption in porous solids has been 243 described by the Langmuir equation [38, 39]:

$$q_i = \frac{q_{mi} * B_i * p_i}{1 + \sum_{j=1}^n B_j * p_j}$$
(13)

where $q_i \, [\text{mmol/g}]$ is the adsorbed amount of the component $i; q_{mi} \, [\text{mmol/g}]$ 244 is the saturation adsorbed amount of the component i; B_i [1/kPa] is the Lang-245 muir constant of the component i; p_i [kPa] is the equilibrium partial pressure 246 of the component i; B_j [1/kPa] is the Langmuir constant of the component 247 j; p_j [kPa] is the equilibrium partial pressure of the component j; i and j are 248 the gas species of the syngas. Table 1 reports the Langmuir constants and 249 the saturation adsorbed amounts for Zeolite 5A, while Figure 4 depicts the 250 adsorption trends of the syngas gases as function of pressure. It can be noted 251 the high CO_2 selectivity of the zeolite in comparison with others gases. 252

The mass of zeolite required for adsorbing all the carbon dioxide of the syngas depends on the molar flow of the dry syngas, its CO_2 molar fraction and kinetic constant of adsorption. The ZEO filter module can be constantly regenerated using a rotary valve packaged into modules as described in [40]. The mass of zeolite that needs to be regenerated every cycle with duration of t_{cycle} can be calculated as follows:

$$m_{zeo} = t_{cycle} * \dot{n}_{DG} * \frac{1 + \sum_{j=1}^{n} B_j * p_j}{q_{m,CO_2} * B_{CO_2} * p_{ads} * x_{CO_2}}$$
(14)

where p_{ads} [kPa] is the total pressure of the syngas inside the ZEO filter. 259 A constant temperature of the zeolite filter and of the inlet syngas of 303 K is 260 assumed and the pressure of the inlet syngas is set to 500 kPa as suggested in 261 [38, 39]. The cycling time of regeneration depends on kinetic CO_2 adsorption 262 constant. In this study a plausible time of 60 seconds is assumed and future 263 work will investigate this aspect. Zeolites adsorption generates heat, Ranjani 264 et al. [51] suggests that 64-70 kJ are released for every mole of CO_2 adsorbed. 265 This heat needs to be discharge by the ZEO module in order to keep the 266 temperature constant at 303 K. In this preliminary study, no attention was 267 paid to the ZEO module heat balance. Furthermore, the gas filtering sub-268 system considered in this work is based on the power plant described by 269 Allesina et al. [50]. It was designed with the idea of coupling the gasifier with 270 an internal combustion engine. Since the minimum presence of tars could 271 negatively affect the performance of the zeolite adsorber, syngas purification 272 unit should be properly designed. A potential alternative to water scrubber 273 is oil scrubber with subsequent stripping of tars [52, 53]. 274

275 2.4. Compressor and storage system modeling

The modeling of the syngas compression is carried out considering it as a polytropic transformation. The electrical power required for compression is given by Pedrazzi et al. [54]:

$$P_{comp} = \frac{\dot{n}_{gas} L_{comp,is}}{\eta_{comp}} = \frac{\dot{n}_{gas}}{\eta_{comp}} \frac{zRT_{in}}{z-1} \left[1 - \left(\frac{p_{out}}{p_{in}}\right)^{\frac{z-1}{z}} \right]$$
(15)

where z is the polytrophic coefficient, R is the universal gas constant 279 equal to 8.314 J mol⁻¹ K⁻¹, T_{in} is the gas inlet temperature (25°C for syngas 280 compressor 1 and 30°C for syngas compressor 2), p_{in} and p_{out} [atm] are the gas 281 inlet and outlet pressures, \dot{n}_{gas} [mol/s] is the gas molar flow and η_{comp} is the 282 compressor efficiency available from manufacturer's data [54]. The maximum 283 pressure value inside the storage system is a fundamental parameter required 284 todesign the tanks and the compressor properly. Assuming ideal gas and a 285 constant syngas storage temperature $T_s = 25$ °C, the pressure inside the 286 tanks is calculated by the ideal gas law: 287

$$p_s = \frac{nRT_s}{V} \tag{16}$$

where n [mol] are the moles of syngas inside the tanks and $V \text{ [m^3]}$ is the storage total volume. Assuming a value of the initial syngas moles n_{in} inside the storage, the moles of syngas at the time τ [s] are given by:

$$n = n_{in} + \int_0^\tau \left(\dot{n}_{in,s}(t) - \dot{n}_{out,s}(s) \right) dt$$
 (17)

where $\dot{n}_{in,s}(t)$ and $\dot{n}_{out,s}(t)$ [mol/s] are the inlet and the outlet molar flow at the instantaneous time t [s]. Table 2 reports the model parameters of the storage and compressor sub-systems. The total volume of storage and the initial syngas amount in the tank are reduced of about 50% in comparison with the conventional system without PPO and ZEO modules as investigated in [55, 56]. This result is reached thanks to the PPO adoption that decreases the molar flow of dry syngas of about the 20% - 30%, while the filtration in the ZEO module further reduces the syngas molar flow of about another 20% - 30% as shown in the results.

300 2.5. SOFC modeling

The SOFC model used in this study is based on the work of Bang-Møller 301 and Rokni [18]. This model does not take into account the recirculation 302 of gas at the cell anode. This feature may strongly compromise the fuel 303 cell efficiency in case of the presence of gases that do not take part in the 304 electro-chemical, shift and reforming reactions. Unfortunately, syngas con-305 tains considerable amounts of CO_2 and N_2 . To overcome this issue, the 306 model previously cited is implemented with the reforming model presented 307 by Rami Salah El-Emam et al. [22]. As described by Rami Salah El-Emam, 308 the electro-chemical reactions take place in both the anode and the cathode 309 of the cell (Eq.22), while the reforming and the monoxide water shift occur 310 only near the anode (Eqs. 18, 19). Equation 22 presents the overall electro-311 chemical reaction that is divided into two sub-reactions: the hydrogen reacts 312 with the oxygen ions to form water and electrons according to Eq. 20 at 313 the anode, while, at the cathode, the oxygen from inlet air reacts with the 314 electrons from the anode (Eq. 21) to form oxygen ions that flow to the anode 315 through the solid oxide electrolyte. 316

$$CH_4 + H_2O \to CO + 3H_2 \tag{18}$$

$$CO + H_2O \to CO_2 + H_2 \tag{19}$$

$$H_2 + O^{2-} \to H_2O + 2e^-$$
 (20)

$$\frac{1}{2}O_2 + 2e^- \to O^{2-} \tag{21}$$

$$H_2 + \frac{1}{2}O_2 \to H_2O \tag{22}$$

The mathematical modeling of reforming and electrochemical reactions is explained in References [22] and [18]. Using these models, it is possible to calculate the electrical power production and the electrical conversion efficiency for a given syngas inlet flow with a specific composition. The SOFC model parameters adopted in the simulations are reported in Table 2.

322 2.6. MGT modeling

Mathematical description of gas turbines is well described in literature. Details and assumptions of the present model can be found in Bang-Møller and Rokni work [18]. Characteristics of the turbine and others components connected to the MGT are listed in Table 2.

327 3. Simulation results and discussion

In this work four different cases are simulated. First of all, the basic system composed of a downdraft gasifier, a storage tank and a SOFC-MGT is simulated. After this step, the two possible solutions consisting of N_2 purging from air or CO_2 separation from syngas are discussed separately. Finally, the complete system provided with PPO module and ZEO module is simulated.

334 3.1. Case I (Gasifier + SOFC + MGT)

The SOFC-MGT unit constantly produces energy all over the year of simulation in order to preserve the stability of the cells and their gaskets, which are very sensitive to thermal stresses [57]. The syngas molar flow consumed by the SOFC-MGT unit is calculated by Equation 23 that considers the cycling working of the DG:

$$\dot{n}_{syngas-SOFC} = \frac{h_{operation} \dot{n}_{DG}}{h_{operation} + h_{maintenance}}$$
(23)

340

Figure 5 depicts the overall model implemented in Matlab SimulinkTM soft-341 ware environment. Table 3 reports the simulation results. Gasifier cold 342 efficiency is about 79%, this value is confirmed by literature that suggests 343 an efficiency of 70% - 80% for air-blown downdraft gasifier [37, 4, 44]. Syn-344 gas composition consists of about 19% vol. of H_2 and 15% vol. of CO, the 345 higher heating value of 4.75 MJ/Nm^3 is similar to the results reported by 346 Basu [37] for this kind of gasifier. SOFC-MGT unit has a constant electrical 347 power production of 197.43 kW all over the simulated year. The auxiliary 348 consumption of the whole system strongly depends on tank pressurization 349 level. The average annual value is 34.24 kW. For this reason, the net average 350 power production is reduced to 163.19 kW and the electrical efficiency of the 351 system is 25.43%. 352

353 3.2. Case II (PPO + Gasifier + SOFC + MGT)

Table 4 shows the results of the simulation of the system previously described and now equipped with the PPO module. The oxygen enriched air

flow is about 1.27 mol/s. This value is lower than 2.94 mol/s obtained with-356 out PPO membrane (Case I). Syngas composition is consistent with the work 357 of Wang et al. [58], where oxygen-enriched air (50% oxygen and 50% nitro-358 gen) is used as gasifying agent in a double stage downdraft gasifier fueled 359 with pine sawdust pellets. Differences of 1-2% between the model outputs 360 and Wang's results about CO and H_2 contents are achieved (29% vs. 27%) 361 for H_2 and 26% Vs. 25% for CO). The gasification with oxygen-enriched air 362 assures high gasifier performance in terms of cold gas efficiency (92%), tar 363 production (0.27 g/Nm^3) and syngas higher heating value (7.55 MJ/Nm^3) . 364 The syngas outlet flow is 3.589 mol/s, consistently lower than Case I where 365 syngas flow is 5.01 mol/s. The average equilibrium temperature of gasifi-366 cation in this case is 931 K. This value is only 36 K higher than Case I. 367 Wang et al. [58] suggestes a peak temperature of about 1200 K with oxygen-368 enriched air. With this temperature, conventional material adopted in air 369 gasifier can be used (i.e. stainless steel and refractory brick [2, 6]). In Case 370 II, the overall net power production is 210.52 kW and the electrical efficiency 371 is boosted to 32.81%. The average auxiliary consumption is 42 kW, 8 kW 372 higher than Case I. This is due to the PPO module that uses air at 1 MPa 373 pressure generated by an air compressor. The electrical consumption of the 374 air compressor is 15.3 kW and it is fully compensated by the increasing of 375 the gasifier efficiency and the SOFC-MGT unit efficiency. 376

377 3.3. Case III (Gasifier + ZEO + SOFC + MGT)

Table 5 resumes the simulation results of the system with the ZEO filtering module instead of PPO membrane. The filtered syngas has a higher heating value of 5.9 MJ/Nm^3 . This value falls between Case I (4.75 MJ/Nm^3)

and Case II (7.55 MJ/Nm^3) . The pressure of the storage tank ranges between 381 0.267-0.488 MPa, similar to the values obtained in Case II (0.266-0.462 MPa) 382 and Case I (0.267-0.539 MPa). The zeolite mass required to perform con-383 tinuosly the filtration is 23.742 kg. The value obtained is consistent with 384 Tagliabue et al. work [40]. However, in future work, the CO_2 adsorbed by 385 the ZEO module can be stored in order to create a carbon sequestration sys-386 tem. The power production and the net electrical efficiency of the system is 387 low (148.74 kW of power production and 22.87% of electrical efficiency) as a 388 consequence of the energy absorbed by the syngas compressor 1 (see Figure 389 1) to increase the pressure of the syngas to 0.5 MPa before the ZEO module. 390 This electrical energy consumption is higher than Cases II and I, in addition 391 the efficiency of the SOFC-MGT module fueled with the filtered syngas is 392 lower. As shown in Table 5, the SOFC-MGT efficiency in Case III is about 393 34%, thus lower than Case II (42.08%) and Case I (37.46%). 394

395 3.4. Case IV (PPO + Gasifier + ZEO + SOFC + MGT)

The results about the fully equipped gasifier power system are reported 396 in Table 6. A high power production (194.53 kW) and electrical efficiency 397 (30.32%) is reached thanks to the high H_2 and CO amounts in the filtered 398 syngas. In fact, the H_2 volume percentange reaches 41.53% and the CO 399 volume percentange is boosted to 32.54%. As a consequence of this com-400 position, the higher heating value of the syngas is 10.19 MJ/Nm^3 , a value 401 typical for oxygen-blown gasifiers [37]. Therefore, the SOFC-MGT unit syn-402 gas consumption is 2.696 mol/s. This value is about 45% lower than Case I 403 (4.95 mol/s), 24% lower than Case II (3.545 mol/s) and 33% lower than Case 404 III (4.02 mol/s). A pressure range of 0.266-0.415 MPa is achieved. In this 405

case power production and efficiency is lower than Case II as result of higher average auxiliary consumption (51.80 kW) and lower SOFC-MGT unit efficiency (38.41%). However, the utilization of the ZEO module has several advantages: separates the CO_2 and reduces the storage peaks pressure.

410 3.5. Performance and energy considerations

Figure 6 shows the electrical efficiency and the average power production 411 in every scenario. Case II resulted the best in terms of energy conversion; the 412 overall electrical efficiency reaches 32% and the power production is about 413 210 kW. These values are higher than commercial gasification power systems 414 with internal combustion engines where the maximum electrical efficiency 415 hardly reaches 25% [37, 4, 2]. This result is given by the PPO module that 416 increases the gasifier efficiency to 92% (75% is the reference value for air 417 blown gasifier [37]) and the SOFC-MGT module which has a higher electri-418 cal conversion efficiency (about 42%) compared to common engine-alternator 419 generator units (about 27% [4, 59, 3]). Case II is the best in terms of energy 420 balance as shown in Figure 7. These graphs do not consider the thermal 421 energy that can be recovered from the gasifier or the SOFC-MGT unit. The 422 highest energy loss occurs at the SOFC-MGT unit (about 52%), while aux-423 iliary consumption of the blowers and the auxiliary equipment of the gasifier 424 are low (9%). In Cases I and III, the low efficiency of the gasifier reduces the 425 overall electrical performance of the system. In Cases III and IV the ZEO 426 module consumes energy to separate the CO_2 from the gas, however no effi-427 ciency increase occours in the SOFC-MGT unit with a CO_2 free syngas and 428 the final result is a lower power production. The system modeled in this work 429 is obtained starting from a reference power plant described by Allesina et. al 430

[50] where IC engines are used instead of the SOFC-MGT unit. The author 431 reports an experimental cold gasification efficiency of 67%. Considering an 432 electrical IC engine-alternator unit efficiency of about 27%, as suggested by 433 Puglia at al. [3], the total electrical efficiency is about 18%. This value is 434 30% lower than Case I and it is 45% lower than Case II. Another study made 435 by Patuzzi et al. [60] reports the values of the net electrical efficiency of three 436 different commercial biomass gasifier - IC engine power plants. The average 437 efficiency is about 20%, this value is consistent with the one obtained for 438 Allesina et al. [50]. 439

440 4. Conclusions

The biomass fueled system with PPO module (Case II) shows the higher 441 electrical efficiency of about 33%. The reasons behind this result are vari-442 ous. First of all, oxygen-enriched air boosts the gasifier cold efficiency from 443 79% (Case I with air) to 92% (Cases II and IV with oxygen-enriched air). 444 In addition, the SOFC-MGT unit presents a higher efficiency (about 42%) 445 compared to IC engine-alternator unit (about 27%), ORC cycle (about 20%) 446 or EFGT cycle (about 20%). In Cases III and IV, the zeolites adsoption 447 module consumes energy to increase the higher heating value of the syngas 448 but not the performance of the SOFC-MGT system, this reduces the overall 449 system efficiency. The energy balances of four cases investigated show that 450 the greater losses are in the SOFC-MGT unit. This unit has the difficult 451 task to convert the chemical energy of a gas fuel into electrical energy in 452 an efficient way. An efficiency of about 50% is reached with natural gas, in 453 this study the maximum electrical efficiency is about 42% using a syngas 454

produced with an oxygen-enriched air as gasifying agent. This difference is 455 given by the presence of several inert gases into the syngas that reduces the 456 electrochemical conversion of the SOFC. The removal or the conversion of 457 these gases into syntethic natural gases (SNG) is possible and, in this way, 458 the efficiency of the SOFC-MGT unit will be similar to the value reach for 459 natural gas one. But, cost and energy self-consumption of the upgrading 460 process are very high and not convenient for this kind of power plants. Cases 461 III and IV has a lower efficiency compared to Case II, however, with the 462 ZEO module, it is possible to separate the CO_2 content of the syngas with 463 environmental benefits in case the module is coupled with a CO_2 sequestra-464 tion system. Future work will consider exergy calculations and experimental 465 tests on a micro-scale power system (5-20 kW of electrical power) with PPO 466 module and SOFC module in order to validate modeling results and to assest 467 system durability. In addition, economical net present value analysis will be 468 done to estimate the economic sustainability of the power plant. 460

470 **References**

- [1] IEA. Technology roadmap bioenergy for heat and power. Technical
 report, IEA, 2012.
- [2] Thomas B. Reed and Agua Das. Handbook of Biomass Downdraft Gasi-*fier Engine Systems.* The biomass energy foundation press, 1988.
- [3] G. Allesina, S. Pedrazzi M.Puglia, and P. Tartarini. Upgrading or substituting the gasification process for electrical energy production: an
 energy-based comparison. In XXX UIT Conference, Bologna 2012, 2012.

- ⁴⁷⁸ [4] H.A.M. Knoef. Handbook of Biomass Gasification. BTG, 2005.
- [5] Amitava Datta, Ranjan Ganguly, and Luna Sarkar. Energy and exergy
 analyses of an externally fired gas turbine (efgt) cycle integrated with
 biomass gasifier for distributed power generation. *Energy*, 35(1):341 –
 350, 2010.
- [6] FAO Forestry Department Mechanical Wood Products Branch. Woodgas *as engine fuel*, volume ISBN 92-5-102436-7. F.A.O., 1986.
- [7] F. Martelli, G. Riccio, S. Maltagliati, and D. Chiaramonti. Technical
 study and environmental impact of an external fired gas turbine power
 plant fed by solid fuel. 1st world Conference of Biomass, Sevilla, 2000.
- 488 [8] V. Naso. La macchina di Stirling. CEA, 1991.
- [9] C. Souleymane. Motori a combustione interna e turbine a gas di piccola taglia per gas di sintesi. Master's thesis, Università degli Studi di
 Padova, Italy., 2012.
- ⁴⁹² [10] Farqad Al-Hadeethi, Moh'd Al-Nimr, and Mohammad Al-Safadi. Us⁴⁹³ ing the multiple regression analysis with respect to {ANOVA} and 3d
 ⁴⁹⁴ mapping to model the actual performance of {PEM} (proton exchange
 ⁴⁹⁵ membrane) fuel cell at various operating conditions. *Energy*, 90, Part
 ⁴⁹⁶ 1:475 482, 2015.
- [11] Guido Galeno. Modellizzazione di un micro cogeneratore basato sulla
 tecnologia mcfc accoppiata ad un gassificatore di biomassa. PhD thesis,
 University of Cassino, Italy, 2006-2007.

500	[12]	Giulio Donolo, Giulio De Simon, and Maurizio Fermeglia. Steady
501		state simulation of energy production from biomass by molten carbon-
502		ate fuel cells. Journal of Power Sources, 158(2):1282 – 1289, 2006.
503		;ce:title;Special issue including selected papers from the 6th Interna-
504		tional Conference on Lead-Acid Batteries (LABAT 2005, Varna, Bul-
505		garia) and the 11th Asian Battery Conference (11 ABC, Ho Chi Minh
506		City, Vietnam) together with regular papers;/ce:title;.

- ⁵⁰⁷ [13] Stefano Cordiner, Massimo Feola, Vincenzo Mulone, and Fabio Ro⁵⁰⁸ manelli. Analysis of a {SOFC} energy generation system fuelled with
 ⁵⁰⁹ biomass reformate. Applied Thermal Engineering, 27(4):738 747, 2007.
 ⁵¹⁰ jce:title; Energy: Production, Distribution and Conservation;/ce:title;.
- [14] E. Achenbach. Three-dimensional and time-dependent simulation of a
 planar solid oxide fuel cell stack. *Journal of Power Sources*, 49:333–348,
 1994.
- [15] M. Mortazaei and M. Rahimi. A comparison between two methods of
 generating power, heat and refrigeration via biomass based solid oxide
 fuel cell: A thermodynamic and environmental analysis. *Energy Con*-*version and Management*, 126:132 141, 2016.
- [16] F. Calise, M. Dentice dâTMAccadia, A. Palombo, and L. Vanoli. Simulation and exergy analysis of a hybrid solid oxide fuel cell (sofc)â"gas turbine system. *Energy*, 31(15):3278 3299, 2006. {ECOS} 2004 17th International Conference on Efficiency, Costs, Optimization, Simulation, and Environmental Impact of Energy on Process Systems 17th Interna-

- tional Conference on Efficiency, Costs, Optimization, Simulation, and
 Environmental Impact of Energy on Process Systems.
- [17] L. Fryda, K.D. Panopoulos, and E. Kakaras. Integrated {CHP} with
 autothermal biomass gasification and sofc-mgt. *Energy Conversion and Management*, 49(2):281 290, 2008.
- [18] C. Bang-Moller and M. Rokni. Thermodynamic performance study of
 biomass gasification, solid oxide fuel cell and micro gas turbine hybrid
 systems. *Energy Conversion and Management*, 51(11):2330 2339, 2010.
- [19] Made Sucipta, Shinji Kimijima, and Kenjiro Suzuki. Performance analysis of the sofc-mgt hybrid system with gasified biomass fuel. *Journal*of Power Sources, 174(1):124 135, 2007. ¡ce:title¿Hybrid Electric Vehicles¡/ce:title¿.
- ⁵³⁵ [20] Pegah Ghanbari Bavarsad. Energy and exergy analysis of internal re ⁵³⁶ forming solid oxide fuel cellâ"gas turbine hybrid system. International
 ⁵³⁷ Journal of Hydrogen Energy, 32(17):4591 4599, 2007. Fuel Cells.
- [21] C. Ozgur Colpan, Ibrahim Dincer, and Feridun Hamdullahpur. Thermodynamic modeling of direct internal reforming solid oxide fuel cells operating with syngas. *International Journal of Hydrogen Energy*, 32(7):787
 - 795, 2007. jce:title; Fuel Cellsj/ce:title;.
- [22] Rami Salah El-Emam, Ibrahim Dincer, and Greg F. Naterer. Energy
 and exergy analyses of an integrated {SOFC} and coal gasification system. International Journal of Hydrogen Energy, 37(2):1689 1697, 2012.
 jce:title; 10th International Conference on Clean Energy 2010;/ce:title;.

- ⁵⁴⁶ [23] Penyarat Chinda and Pascal Brault. The hybrid solid oxide fuel cell
 ⁵⁴⁷ (sofc) and gas turbine (gt) systems steady state modeling. International
 ⁵⁴⁸ Journal of Hydrogen Energy, 37(11):9237 9248, 2012.
- ⁵⁴⁹ [24] S.H. Chan, H.K. Ho, and Y. Tian. Modelling of simple hybrid solid
 ⁵⁵⁰ oxide fuel cell and gas turbine power plant. *Journal of Power Sources*,
 ⁵⁵¹ 109(1):111 120, 2002.
- ⁵⁵² [25] Tae Won Song, Jeong Lak Sohn, Jae Hwan Kim, Tong Seop Kim,
 ⁵⁵³ Sung Tack Ro, and Kenjiro Suzuki. Performance analysis of a tubu⁵⁵⁴ lar solid oxide fuel cell/micro gas turbine hybrid power system based on
 ⁵⁵⁵ a quasi-two dimensional model. *Journal of Power Sources*, 142(1â"2):30
 ⁵⁵⁶ 42, 2005.
- ⁵⁵⁷ [26] Mahsa Aghaie, Mehdi Mehrpooya, and Fathollah Pourfayaz. Introduc⁵⁵⁸ ing an integrated chemical looping hydrogen production, inherent carbon
 ⁵⁵⁹ capture and solid oxide fuel cell biomass fueled power plant process con⁵⁶⁰ figuration. *Energy Conversion and Management*, 124:141 154, 2016.
- ⁵⁶¹ [27] Masood Ebrahimi and Iraj Moradpoor. Combined solid oxide fuel cell,
 ⁵⁶² micro-gas turbine and organic rankine cycle for power generation (sofc ⁵⁶³ mgt-orc). Energy Conversion and Management, 116:120 133, 2016.
- ⁵⁶⁴ [28] Ph. Hofmann, K.D. Panopoulos, P.V. Aravind, M. Siedlecki,
 ⁵⁶⁵ A. Schweiger, J. Karl, J.P. Ouweltjes, and E. Kakaras. Operation of
 ⁵⁶⁶ solid oxide fuel cell on biomass product gas with tar levels >10 g
 ⁵⁶⁷ nmâ'3. International Journal of Hydrogen Energy, 34(22):9203 9212,
 ⁵⁶⁸ 2009.

- ⁵⁶⁹ [29] Lopamudra Devi, Krzysztof J Ptasinski, and Frans J.J.G Janssen. A re view of the primary measures for tar elimination in biomass gasification
 ⁵⁷¹ processes. *Biomass and Bioenergy*, 24(2):125–140, 2003.
- ⁵⁷² [30] Jinsong Zhou, Qing Chen, Hui Zhao, Xiaowei Cao, Qinfeng Mei,
 ⁵⁷³ Zhongyang Luo, and Kefa Cen. Biomass-oxygen gasification in a high⁵⁷⁴ temperature entrained-flow gasifier. *Biotechnology Advances*, 27(5):606
 ⁵⁷⁵ 611, 2009. Bioenergy Research & amp; Development in ChinaICBT
 ⁵⁷⁶ 2008.
- ⁵⁷⁷ [31] T. Melin and R. Rautenbach. Membranverfahren, Grundlagen der
 ⁵⁷⁸ Modul- und Anlagenauslegung. 2007.
- [32] K.C. Khulbe, T. Matsuura, G. Lamarche, and H.J. Kim. The morphology characterisation and performance of dense ppo membranes for gas
 separation. *Journal of Membrane Science*, 135(2):211 223, 1997.
- [33] Janusz Kotowicz and Adrian Balicki. Enhancing the overall efficiency
 of a lignite-fired oxyfuel power plant with {CFB} boiler and membranebased air separation unit. *Energy Conversion and Management*, 80:20
 31, 2014.
- [34] Giacomo Bisio, Alessandro Bosio, and Giuseppe Rubatto. Thermody namics applied to oxygen enrichment of combustion air. *Energy Con- version and Management*, 43(18):2589 2600, 2002.
- [35] H. Scott Coombe and Sen Nieh. Polymer membrane air separation per formance for portable oxygen enriched combustion applications. *Energy Conversion and Management*, 48(5):1499 1505, 2007.

- [36] Yanhong Hao, Yi Huang, Minhui Gong, Wenying Li, Jie Feng, and Qun
 Yi. A polygeneration from a dual-gas partial catalytic oxidation coupling
 with an oxygen-permeable membrane reactor. *Energy Conversion and Management*, 106:466 478, 2015.
- [37] Prabir Basu. Biomass Gasification and Pyrolysis: Practical Design and Theory. Academic Press, Elsevier, 2010.
- [38] Saeed Pakseresht, Mohammad Kazemeini, and Mohammad M. Akbarnejad. Equilibrium isotherms for co, co2, {CH4} and {C2H4} on the 5a
 molecular sieve by a simple volumetric apparatus. Separation and Pu-*rification Technology*, 28(1):53 60, 2002.
- [39] Gi-Moon Nam, Byung-Man Jeong, Seok-Hyun Kang, Byung-Kwon Lee,
 and Dae-Ki Choi. Equilibrium isotherms of ch4, c2h6, c2h4, n2, and h2
 on zeolite 5a using a static volumetric method. Journal of Chemical &
 Engineering Data, 50(1):72-76, 2005.
- [40] Marco Tagliabue, David Farrusseng, Susana Valencia, Sonia Aguado,
 Ugo Ravon, Caterina Rizzo, Avelino Corma, and Claude Mirodatos.
 Natural gas treating by selective adsorption: Material science and chemical engineering interplay. *Chemical Engineering Journal*, 155(3):553 –
 566, 2009.
- [41] Zoltán Bacsik, Ocean Cheung, Petr Vasiliev, and Niklas Hedin. Selective
 separation of {CO2} and {CH4} for biogas upgrading on zeolite naka
 and sapo-56. Applied Energy, 162:613 621, 2016.

- [42] Mariem Kacem, Mario Pellerano, and Arnaud Delebarre. Pressure swing
 adsorption for co2/n2 and co2/ch4 separation: Comparison between activated carbons and zeolites performances. *Fuel Processing Technology*,
 138:271 283, 2015.
- [43] Qassim Hassan Dirar and Kevin F. Loughlin. Intrinsic adsorption properties of co2 on 5a and 13x zeolite. Adsorption, 19(6):1149–1163, 2013.
- [44] Niladri Sekhar Barman, Sudip Ghosh, and Sudipta De. Gasification of
 biomass in a fixed bed downdraft gasifier a realistic model including
 tar. *Bioresource Technology*, 107:505–511, 2012.
- [45] Takashi Yamazaki, Hirokazu Kozu, Sadamu Yamagata, Naoto Murao,
 Sachio Ohta, Satoru Shiya, and Tatsuo Ohba. Effect of superficial velocity on tar from downdraft gasification of biomass. *Energy & Fuels*,
 19:1186–1191, 2005.
- [46] S. Jarungthammachote and A. Dutta. Thermodynamic equilibrium
 model and second law analysis of a downdraft waste gasifier. *Energy*,
 32(9):1660 1669, 2007.
- [47] Giulio Allesina, Simone Pedrazzi, Luca Guidetti, and Paolo Tartarini.
 Modeling of coupling gasification and anaerobic digestion processes for
 maize bioenergy conversion. *Biomass and Bioenergy*, 81:444 451, 2015.
- [48] Giulio Allesina, Simone Pedrazzi, Federico Sgarbi, Elisa Pompeo,
 Camilla Roberti, Vincenzo Cristiano, and Paolo Tartarini. Approaching
 sustainable development through energy management, the case of fongo
- tongo, cameroon. International Journal of Energy and Environmental
 Engineering, 6(2):121–127, 2014.
- [49] Giulio Allesina, Simone Pedrazzi, Emma La Cava, Michele Orlandi,
 Miriam Hanuskova, Caludio Fontanesi, and Paolo Tartarini. Energybased assessment of optimal operating parameters for coupled biochar
 and syngas production in stratified downdraft gasifiers. In *International Heat Transfer Conference 15, Kyoto, Japan*, 2014.
- ⁶⁴³ [50] Giulio Allesina, Simone Pedrazzi, and Paolo Tartarini. Modeling and
 ⁶⁴⁴ investigation of the channeling phenomenon in downdraft stratified
 ⁶⁴⁵ gasifers. *Bioresource Technology*, 146(0):704 712, 2013.
- [51] Ranjani V. Siriwardan, Ming-Shing Shen, and Edward P. Fisher. Adsorption of co2 on zeolites at moderate temperatures. *Energy and Fuels*, 19:1153–1159, 2005.
- [52] Thana Phuphuakrat, Tomoaki Namioka, and Kunio Yoshikawa. Absorptive removal of biomass tar using water and oily materials. *Bioresource Technology*, 102(2):543–549, 2011.
- [53] Shunsuke Nakamura, Shigeru Kitano, and Kunio Yoshikawa. Biomass
 gasification process with the tar removal technologies utilizing bio-oil
 scrubber and char bed. Applied Energy, 170:186 192, 2016.
- ⁶⁵⁵ [54] S. Pedrazzi, G. Zini, and P. Tartarini. Complete modeling and soft ⁶⁵⁶ ware implementation of a virtual solar hydrogen hybrid system. *Energy* ⁶⁵⁷ Conversion and Management, 51(1):122 129, 2010.

- [55] Simone Pedrazzi, Giulio Allesina, Alberto Muscio, and Paolo Tartarini.
 Modeling and simulation of a dg-sofc-mgt hybrid system. In 7° Congresso Nazionale AIGE, Rende (CZ) Italy, 2013.
- [56] Simone Pedrazzi. Modeling and optimization of advanced systems for
 electrical energy production from wood biomass. PhD thesis, HIGH ME CHANICS AND AUTOMOTIVE DESIGN & TECHNOLOGY, University of Modena and Reggio Emilia, Dep. of Engineering 'Enzo Ferrari',
 2013.
- ⁶⁶⁶ [57] DOE. Fuel Cell Handbook (Seventh Edition). 2004.
- [58] Zhiqi Wang, Tao He, Jianguang Qin, Jingli Wu, Jianqing Li, Zhongyue
 Zi, Guangbo Liu, Jinhu Wu, and Li Sun. Gasification of biomass with
 oxygen-enriched air in a pilot scale two-stage gasifier. *Fuel*, 150:386 –
 393, 2015.
- [59] H.A.M. Knoef. Handbook of Biomass Gasification, Second Edition.
 BTG, 2012.
- [60] Francesco Patuzzi, Dario Prando, Stergios Vakalis, Andrea Maria Rizzo,
 David Chiaramonti, Werner Tirler, Tanja Mimmo, Andrea Gasparella,
 and Marco Baratieri. Small-scale biomass gasification {CHP} systems:
 Comparative performance assessment and monitoring experiences in
 south tyrol (italy). *Energy*, 112:285 293, 2016.
- [61] S. Pedrazzi, G. Allesina, and P. Tartarini. Aige conference: A kinetic
 model for a stratified downdraft gasifier experimental assessment and

- ⁶⁸⁰ modeling of energy conversion effectiveness in a gasification power plant.
- International Journal of Heat and Technology, 30(1):41–44, 2012.
- 682 Figure captions and tables



Figure 1: DG-SOFC-MGT hybrid system with zeolite CO_2 adsorption and oxygenenriched air layout



Figure 2: Oxygen enriched air membrane separator principle



Figure 3: Characteristics of the separation membranes



Figure 4: Zeolite 5A adsorbing curve Vs. pressure



Figure 5: DG-SOFC-MGT hybrid system with zeolite CO_2 adsorption and oxygenenriched air implemented in Matlab SimulinkTM



Figure 6: Efficiencies and eletrical production values of the studied cases



Figure 7: Energy balances of the studied cases

Table 1: Mode	el parameters I	-	
Membranes p	roperties [31]		
Material	Selectivity	Oxigen Permeability	
	α [ad]	$\gamma \; [\mathrm{mol} \; \mathrm{m}^{-2} \; \mathrm{s}^{-1} \; \mathrm{bar}^{-1}]$	
Matrimid	6.7	$62.0 \ge 10^{-5}$	
PPO	4.7	$37.2 \ge 10^{-4}$	
PDMS	2.1	$39.6 \ge 10^{-2}$	
Poplar wood chip	s properties [6	51]	
Description	Symbol	Value	
Total moisture	M	10 %	
Carbon content (as received)	C_{ar}	41.62~%	
Hydrogen content (as received)	H_{ar}	5.30~%	
Nitrogen content (as received)	N_{ar}	0.52~%	
Oxygen content (as received)	O_{ar}	39.81~%	
Ash content	ASH	2.75~%	
Higher heating value (dry basis)	HHV_{db}	$15.7 \mathrm{~MJ/kg}$	
Gasifier model	parameters[48]		
Description	Symbol	Value	
As received biomass consumption	\dot{m}_{bio}	187 kg/h	
Nominal gasifier thermal power	$P_{th,gas}$	800 kW	
Initial calculation temperature	T_{in}	900 K	
Pressure	р	1 atm	
Equivalence ratio	\mathbf{ER}	0.335	
Gasifier and filters auxiliary consumption	$P_{DG,self}$	12.5 kW	
Cyclic operation hours	$h_{operation}$	360 h	
Cyclic maintenance hours	$h_{maintenance}$	4 h	
Zeolite 5A parameters of adsorption at 303 K [39]			
Component	$B \ [1/kPa]$	$q_m \; [\mathrm{mmol/g}]$	
CO_2	0.019500	3.91900	
H_2	0.000361	0.54464	
N_2	0.000837	2.62543	
CH_4	0.002535	2.75403	
CO	0.004350	2.75800	

Table 2: Model par	rameters II	
Storage and compressor model parameters		
Description	Symbol	Value
Politropic exponent of the syngas [54]	m	1.33
Syngas compressor efficiency [54]	η_{comp}	92~%
Storage tank temperature	T_s	$298.15~\mathrm{K}$
Initial syngas amount in the tanks	n_{in}	$7*10^4$ mol
Total tanks volume	V	650 m^3
SOFC model par	ameters	
Description	Symbol	Value
Fuel utilization factor	U_f	0.85
Recirculation factor	r	0.2
Operating temperature	T_{sofc}	$1073.15 { m K}$
Anode pressure loss	Δp_a	$500 \ Pa$
Cathode pressure loss	Δp_c	$1000 \ Pa$
Anode pressure loss	Δp_a	$500 \ Pa$
Current density	i	300 mA/cm
Active cell area	A_{cell}	$81~{\rm cm^2}$
Cells for each stack	$n_{cell,stack}$	75 cells
Number of stacks	n_{stack}	145 stacks
Cathode air excess	vent	1.15
Pressure ratio	PR	2.5
Steam to carbon coefficient	STC	1.4
Electrochemical parameters taken from [18]		
MGT model parameters [18]		

MGT model parameters [18]		
Description	Symbol	Value
Politropic exponent of the air	m	1.33
Turbine isoentropic efficiency	$\eta_{is,turb}$	84 %
Air compressor isoentropic efficiency	$\eta_{is,comp}$	75 %
Turbine mechanical efficiency	$\eta_{mec,turb}$	99~%
Air compressor mechanical efficiency	$\eta_{mec,comp}$	98 %
Recuperator effectiveness	η_{rec}	85 %
Burner efficiency	$\eta_{eff,burner}$	99 %
MGT generator efficiency	$\eta_{alt,MGT}$	95~%
Pressure ratio	PR	2.5

Gasifier		
Description	Symbol	Value
H_2 syngas fraction	x_{H_2}	19.03 %
H_2O syngas fraction	x_{H_2O}	7.78 %
CO syngas fraction	x_{CO}	15.10~%
CH_4 syngas fraction	x_{CH_4}	1.18~%
CO_2 syngas fraction	x_{CO_2}	13.99~%
N_2 syngas fraction	x_{N_2}	42.92~%
Air inlet flow	Q_{air}	2.94 mol/s
Syngas molar flow	\dot{n}_{syngas}	5.01 mol/s
Syngas higher heating value	$HHV_{syngas,db}$	$4.75 \ \mathrm{MJ/Nm^3}$
Specific volumetric tar production	m_{tar,Nm^3}	$23.63~{\rm g/Nm^3}$
Gasifier cold gas efficiency	η_{cold}	78.98~%
Average temperature of gasification	T	895 K
SOFC + MG	Т	
Description	Symbol	Value
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	4.95 mol/s
SOFC electrical power production	P_{SOFC}	$136.70~\mathrm{kW}$
MGT electrical power production	P_{MGT}	$60.73 \mathrm{~kW}$
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$197.43~\mathrm{kW}$
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	37.46~%
Overall system	m	
Description	Symbol	Value
Storage tank pressure range	p_{serb}	2.67-5.39 bar
Average electrical auxiliary consumption	P_{self}	$34.24 \ kW_{el}$
Average electrical total power production	P_{tot}	163.19 kW_{el}
Average total electrical efficiency	η_{tot}	25.43~%

Table 3: Case I (Gasifier + SOFC + MGT) simulation results

PPO module		
Air inlet flow	Q_{air}	2.97 mol/s
Permeate molar flow	Q_P	1.27 mol/s
Retentate molar flow	Q_R	1.69 mol/s
Molar fraction of O_2 in permeate	y_{O2}	48.9~%
Molar fraction of N_2 in permeate	y_{N2}	51.1 %
Electric power consumption	$P_{el,PPO}$	$15.3 \mathrm{~kW}$
Gasifier		
Description	Symbol	Value
H_2 syngas fraction	x_{H_2}	28.49 %
H_2O syngas fraction	x_{H_2O}	9.91~%
CO syngas fraction	x_{CO}	26.33~%
CH_4 syngas fraction	x_{CH_4}	1.66~%
CO_2 syngas fraction	x_{CO_2}	17.2~%
N_2 syngas fraction	x_{N_2}	16.41~%
Syngas molar flow	\dot{n}_{syngas}	3.589 mol/s
Syngas higher heating value	$HHV_{syngas,db}$	$7.55 \ \mathrm{MJ/Nm^3}$
Specific volumetric tar production	m_{tar,Nm^3}	$0.27~{ m g/Nm^3}$
Gasifier cold gas efficiency	η_{cold}	92.0~%
Average temperature of gasification	T	931 K
SOFC + MG'	T	
Description	Symbol	Value
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	3.545 mol/s
SOFC electrical power production	P_{SOFC}	$184.20~\mathrm{kW}$
MGT electrical power production	P_{MGT}	$68.33 \mathrm{~kW}$
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$252.53~\mathrm{kW}$
SOFC+MGT eletrical efficiency	$\eta_{SOFC+MGT}$	42.08~%
Overall system		
Description	Symbol	Value
Storage tank pressure range	p_{serb}	2.67-4.62 bar
Average electrical auxiliary consumption	P_{self}	$42.01 \ kW_{el}$
Average electrical total power production	P_{tot}	210.52 kW_{el}
Average total electrical efficiency	η_{tot}	32.81~%

Table 4: Case II (PPO + Gasifier + SOFC + MGT) simulation results

Gashiei (see Cas	e 1)	
ZEO		
Description	Symbol	Value
H_2 syngas fraction after adsoption	x_{H_2}	25.36~%
CO syngas fraction after adsoption	x_{CO}	17.08~%
CH_4 syngas fraction after adsoption	x_{CH_4}	1.43~%
CO_2 syngas fraction after adsoption	x_{CO_2}	0.39~%
N_2 syngas fraction after adsoption	x_{N_2}	55.73~%
Syngas molar flow after adsoption	\dot{n}_{syngas}	4.065 mol/s
Syngas higher heating value after adsoption	$HHV_{syngas,db}$	$5.9 \ \mathrm{MJ/Nm^3}$
Active zeolite mass for every regeration cycle	m_{zeo}	$23.742~\mathrm{kg}$
SOFC + MGT	Г	
Description	Symbol	Value
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	4.020 mol/s
SOFC electrical power production	P_{SOFC}	$138.50~\mathrm{kW}$
MGT electrical power production	P_{MGT}	$44.00~\mathrm{kW}$
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$182.5~\mathrm{kW}$
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	34.33~%
Overall system	n	
Description	Symbol	Value
Storage tank pressure range	p_{serb}	2.66-4.88 bar
Average electrical auxiliary consumption	P_{self}	33.76 kW_{el}
Average electrical total power production	P_{tot}	148.74 kW_{el}
Average total electrical efficiency	η_{tot}	22.87~%

 $\frac{\text{Table 5: Case III (Gasifier + ZEO + SOFC + MGT) simulation results}}{\text{Gasifier (see Case I)}}$

PPO module (see Case II)			
Gasifier (see Ca	Gasifier (see Case II)		
ZEO			
Description	Symbol	Value	
H_2 syngas fraction after adsoption	x_{H_2}	41.53 %	
CO syngas fraction after adsoption	x_{CO}	32.54~%	
CH_4 syngas fraction after adsoption	x_{CH_4}	2.21~%	
CO_2 syngas fraction after adsoption	x_{CO_2}	0.43~%	
N_2 syngas fraction after adsoption	x_{N_2}	23.30~%	
Syngas molar flow after adsoption	\dot{n}_{syngas}	$2.726~{\rm mol/s}$	
Syngas higher heating value after adsoption	$HHV_{syngas,db}$	$10.19~\mathrm{MJ}/\mathrm{Nm}^3$	
Active zeolite mass for every regeration cycle	m_{zeo}	$20.244~\rm kg$	
SOFC + MGT			
Description	Symbol	Value	
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	$2.696~{\rm mol/s}$	
SOFC electrical power production	P_{SOFC}	$184.70~\mathrm{kW}$	
MGT electrical power production	P_{MGT}	51.80 kW	
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$236.50~\mathrm{kW}$	
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	38.41 %	
Overall system			
Description	Symbol	Value	
Storage tank pressure range	p_{serb}	2.66-4.15 bar	
Average electrical auxiliary consumption	P_{self}	51.80 kW_{el}	
Average electrical total power production	P_{tot}	194.53 kW_{el}	
Average total electrical efficiency	η_{tot}	30.32~%	







Figure(s) Click here to download high resolution image









Response to Review Comments

Thanks to the reviewers for their work, I have revised my present research paper in the light of their useful suggestions and comments. The corrections in the manuscript were made in red color. I hope my revision has improved the paper to the level of their satisfaction:

(O) = reviewer observation

(A) = author answer

Editor

(O): Please update your literature survey by referring to the most recent and relevant references that have been published in highly ranked and prestigious journals including this journal. Please focus on relevant publications during the last few years.

(A): We add 5 recent references about fuel cells systems in the introduction section

Reviewer # 1

(O): The manuscript should be carefully checked again for typos, for example: line 18, 43, and 415 etc..

(A): Done

(O): A nomenclature section is needed to quickly refer to any symbol/acronym during the reading of the article.

(A): Done

(O): Fig. 5, Simulink model is not properly visible. An increase in font size or style maybe required.

(A): Done

(O): In introduction section, the authors have emphasized largely on the importance of PPO or ZEO module and their working principle. In one way, this is important to explain all details. But the reviewer suggest that trivial details about importance of PPO or ZEO module should be mentioned briefly with proper literature references (for further study) and more focus should be on literature review of these modules and their effect on the efficiency of the system. I could not find the authors mentioning any study related to this scope of work.

(A): Several works about PPO membrane air filtration and zeolites adsorption applications are now cited in the introduction section.

(O): The conclusion section maintains that including the PPO and ZEO might result in higher efficiencies, which is major focus of this paper. But, in results section, the discussion related to comparison of these cases with existing energy conversion systems (IC engine or ORC etc.) might be missing/less emphasized. It is suggested that a detailed comparison of energy conversion efficiencies of other technologies may be included in the discussion section.

(A): We add a comparison with a common gasifier-IC engine power plant in the "Performance and energy consideration" subsection.

(O): Table 1 (page 38), gasifier model parameters reference for data may be furnished.

(A): Reference added

(O): Table 3-6 show data obtained from different cases during simulation. It is suggested that a brief summary of all simulation results (in the form of a graph) might be helpful for the reader to get an overview of the results.

(A): Figure 6 reports efficiency and power production results reported in Table 3-6. In addition Figure 7 shows energy results of the fourth cases.

Thank you again for your review.

All the best,

Simone Pedrazzi

Effects of upgrading systems on energy conversion efficiency of a gasifier - fuel cell - gas turbine power plant

Simone Pedrazzi^{a,*}, Giulio Allesina^a, Paolo Tartarini^a

^aUniversity of Modena and Reggio Emilia, Department of Engineering 'Enzo Ferrari', Via Vivarelli 10/1, 41125 Modena, Italy

Abstract

THIS PAPER IS SUBMITTED WITH THE OPTION 'YOUR PAPER YOUR WAY'. FOR THIS REASON LAYOUT AND STYLE MAY DIFFER FROM THE JOURNAL ONE.

This work focuses on a DG-SOFC-MGT (downdraft gasifier- solid oxide fuel cell - micro gas turbine) power plant for electrical energy production and investigates two possible performance-upgrading systems: polyphenylene oxide (PPO) membrane and zeolite filters. The first is used to produce oxygenenriched air used in the reactor, while the latter separates the CO_2 content from the syngas. In order to prevent power plant shutdowns during the gasifier reactor scheduled maintenance, the system is equipped with a gas storage tank. The generation unit consists of a SOFC-MGT system characterized by higher electrical efficiency when compared to conventional power production technology (IC engines, ORC and EFGT). Poplar wood chips with 10% of total moisture are used as feedstock. Four different combinations with and without PPO and zeolite filtrations are simulated and discussed. One-year

Preprint submitted to Energy Conversion and Management

^{*}Corresponding author

Email address: simone.pedrazzi@unimore.it (Simone Pedrazzi)

energy and power simulation were used as basis for comparison between all the cases analyzed. The modeling of the gasification reactions gives results consistent with literature about oxygen-enriched processes. Results showed that the highest electrical efficiency obtained is 32.81%. This value is reached by the power plant equipped only with PPO membrane filtration. Contrary to the PPO filtering, zeolite filtration does not increase the SOFC-MGT unit performance while it affects the energy balance with high auxiliary electrical consumption. This solution can be considered valuable only for future work coupling a CO_2 sequestration system to the power plant.

Keywords:

Biomass, Gasification, Modeling, Solide Oxide Fuel Cells, Zeolites, PPO membrane

1 1. Introduction

Due to the abundant availability and distribution, biomasses hold keyroles in plans for renewable energy production. This trend is becoming even more relevant thanks to the good degree of reliability and efficiency of the biomass-based technologies together with the high subsidies granted by several government for sustainable electrical energy production [1].

Depending on the feedstock quality and availability, biomasses are converted into energy through different technologies. In the case of ligno-cellulosic biomasses, a technolgy of great validity is gasification. This thermo-chemical process turns solid biomass into a gaseous fuel known as syngas, which can be converted into electrical energy through all those systems used for power production from gaseous fuels [2]. Gasification is today one of the most effi¹³ cient technologies to convert wood into electricity and it is also sustainable ¹⁴ in terms of the environmental balance of CO_2 [3, 4].

Most of the gasification power plants use an IC engine-generator to con-15 vert the syngas chemical energy into electrical power. However, in some cases 16 other conversion machines are used, i.e. Organic Rankine Cycles (ORC), Ex-17 ternal Firing Gas Turbines (EFGT)[5] and Stirling engines are used with the 18 major advantage of having minor limitation about the syngas level of purifica-19 tion [2, 6, 7, 8, 9]. These systems are usually characterized by low conversion 20 efficiencies of about 10-12%. Major conversion rates can be obtained only 21 with electrochemical devices such as proton exchange membrane fuel cells 22 [10], Molten Carbonate Fuel Cells (MCFC) [11, 12], Solid Oxide Fuel Cells 23 (SOFC) [13, 14, 15], systems composed of SOFC and Micro Gas Turbines 24 (MGT) [16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26] and systems composed of 25 SOFC-MGT-ORC [27]. Despite the high rate of energy conversion, these 26 systems require perfectly clean syngas [28]. Downdraft gasifiers are the most 27 suitable architecture due to the low tar and particulate content in their gas 28 when compared to updraft, crossdraft or fluidized bed gasifiers [2, 6, 29]. 29 However, downdraft gasifiers commonly use air as gasification agent. This 30 solution generates a syngas with a low calorific value where the hydrogen, 31 methane and carbon monoxide are diluted in non-burnable gases: N_2 (about 32 50%) and CO_2 (from 10 to 20%). Otherwise, it is possible to choose oxygen 33 gasification that produces a syngas with negligible N_2 content. However, oxy-34 gen gasification is a complex and expensive technology due to the gasification 35 agent supply sub-systems and reactor material choice. Indeed, temperatures 36 inside the reactor can reach 1200-1300 K when oxygen is used instead of air 37

зв [30].

The basic system discussed in this study is composed of an air blowndowndraft-fixed bed gasifier fed with poplar wood chips. This work is aimed at investigating the effects of different power plant designs on the overall energy conversion efficiency.

The first power plant upgrading sub-system consists of a polyphenylene 43 oxide (PPO) membrane used to produce oxygen-enriched air. The gas sep-44 aration characterization of this membrane is reported in literature [31, 32]. 45 In practice, membrane gas separation is applied to increase the oxygen con-46 tent in the inlet air of biomass boilers [33]. Bisio et al. studied the ther-47 modynamics of combustion with enriched air and reviewed several types of 48 memebranes [34]. Coombe and Nieh developed a membrane-based device for 49 air enrichment in small scale burners [35]. Hao et al. applied an oxygen-50 permeable membrane to a reactor for the co-production of dimethyl ether 51 (DME)/methanol and electricity [36]. This paper uses PPO membrane in 52 order to obtain air with about 50% of oxygen then used as gasification agent. 53 This solution is a hybrid between air and pure oxygen gasification. Enriched 54 air reduces the reactor thermal stress compared to pure oxygen gasification, 55 while the syngas has a lower N_2 content than the one obtained in pure air 56 gasification. In addition, the syngas flow rate decreases because, for a fixed 57 power output, the enriched air flow required for gasification is lower than air 58 used in conventional gasification. This happens because the same amount of 59 oxygen is used in both cases and its concentration in enriched air is higher 60 than untreated air. Finally, the tar production is lower than air gasification 61 as consequence of the higher temperature that cracks more efficiently the 62

⁶³ primary tars from pyrolysis [37].

A second solution discussed in this work consists of a porous media used 64 to upgrade the syngas. In fact, syngas has a variable CO_2 content depending 65 on gasification process as well as several boundary conditions. This value 66 ranges from 10% to 30% and it reduces significantly the higher heating value 67 of the syngas [37]. A solution to overcome this issue is to adopt a pressure-68 swing selective synthetic zeolite filter. This system is placed before the gas 69 storage in order to separate carbon dioxide from syngas [38, 39]. The filter 70 can be constantly regenerated using a rotary valve packaged into modules 71 as described by Tagliabue et al. [40]. Literature investigation about zeolite 72 filtration outlines several works. Bacsik et al. studied the biogas CO_2 - CH_4 73 separation through zeolites [41]. Kacem et al. investigated the pressure swing 74 adsorption for CO_2/N_2 and CO_2/CH_4 separation using activated carbon and 75 several types of zeolites [42]. Dirar et al. investigated intrinsic adsorption 76 properties of CO2 on 5A and 13X zeolite [43]. 77

The syngas obtained from gasification is stored and then used in a SOFC 78 unit able to produce electrical and thermal energy. The number of stacks 79 within the cell is optimized taking into account the optimal electrical cur-80 rent density. The chosen number guarantees a good efficiency, however the 81 gas discharged from the cell still contains some chemical energy. For this 82 reason, this work suggests to convert this residual energy in a micro gas tur-83 bine (MGT). The syngas storage allows the generation unit to operate in its 84 optimal point, furthermore it prevent the power plant shoutting down dur-85 ing the maintenance operations of the gasifier. This management preserves 86 the SOFC and MGT reliability. However, it is difficult to design the stor-87

age capacity because an oversize storage rises the systems costs, while an undersized capacity reduces the time gained for the maintenance. For this reason, the storage was designed taking into account the tanks pressure, the electrical power production of the SOFC-MGT unit and the time required for scheduled stops of the gasifier for maintenance operations.

The mathematics of the whole system was developed starting from literature. The overall model has been implemented in Matlab SimulinkTM software environment in order to simulate the behavior of the system under different conditions over a year long simulation.

97 2. System modeling

The basic system layout is reported in Figure 1. The most relevant components are:

- Downdraft gasifier: The gasifier is equipped with a subsystem for the syngas filtering and cooling with water scrubber and electrostatic filters.
- Syngas storage: It consists of a tank of a total volume of 650 m³.
- SOFC unit: This subsystem consists of 10875 solid oxide cells and it is connected to the electrical grid by a power inverter.
- Micro gas turbine (MGT): this turbo-machinery is used to convert the last part of chemical energy content in the syngas purged by the SOFC.

This work investigates the effect of the implementation of the following sub-systems to the basic scheme:

• **PPO membrane filter module:** The PPO sub-system consists of the membrane filter and a compressor that increases the pressure of the air before the PPO membrane filter to about 1 MPa. The oxygen enriched air is sent to the gasifier at atmospheric pressure. A flow of nitrogen is purged from the PPO module.

• Zeolite (ZEO) filter module: the zeolite (ZEO) filter module is placed after the first syngas compressor. There is a further syngas compression stage ahead the storage tanks because the ZEO module works at 0.5 MPa of pressure as described in Section 2.3, while the pressure in the storage is often higher.

The syngas is used as fuel in the SOFC stack. In this device, the fuel reforming occurs at the anode and there is a recirculation of the 20% of the anode exhaust to increase the fuel reforming performance [18, 22]. The anode exhaust is used to preheat the syngas, then it is finally burned in the MGT burner together with the cathode exhaust. The air required for the electrochemical reaction is compressed and preheated in the recuperator of the MGT as well as in the air preheater of the SOFC.

The SOFC stack generates DC current which is converted into AC current by an inverter and it is sent to the electrical grid. The MGT drags the air compressor and the remaining mechanical energy is converted into electrical energy by an alternator.

132 2.1. PPO module modeling

Polymeric membranes allow to separate different gaseous components de-133 pending on the pore size and pressure applied to the filter [31]. In this work 134 a membrane is used to separate nitrogen from air. The membranes widely 135 used for this purpose are: Matrimid, Polyphenylenoxide (PPO) and Poly-136 dimethylsiloxan (PDMS) [31]. As showed in Figure 2, in membranes the 137 inlet air flow is divided in permeate and retentate molar flows. The inlet 138 flow $(Q_{air} \text{ [mol/s]})$ has a pressure p_{feed} [atm] and it is composed of x_{O_2Feed} 139 and x_{N_2Feed} molar fractions of oxygen and nitrogen. The permeate molar 140 flow $(Q_P \text{ [mol/s]})$ has a pressure $p_{permeate}$ [atm] and it is composed of y_{O_2} 141 and y_{N_2} molar fractions of oxygen and nitrogen. The retentate molar flow 142 $(Q_R \text{ [mol/s]})$ has a pressure p_{feed} [atm] and it is composed of $x_{O_2Retentate}$ and 143 $x_{N_2Retentate}$ molar fractions of oxygen and nitrogen. 144

Each membrane behavior is identified through two parameters: the selectivity (α) and the permeability to oxygen (γ). The first factor represents the attitude of the membrane to attract oxygen, the second quantifies the attitude of the membrane to be crossed by it. High selectivity and permeability ensure great filtering performance in terms of high value of y_{O2} and a small membrane surface area is required to filter a given amount of air. Table 1 presents the parameters of Matrimid, PPO and PDSM membranes.

The choice of a PPO membrane is a compromise in terms of acceptable values of selectivity and permeability. In order to simulate the behavior of the membranes, a mathematical model has been implemented from Melin and Rautenbach [31]. The model is based on the following assumptions:

156

• Air is considered a binary gas mixture with 21% oxygen and 79% ni-

157 trogen.

- Steady state conditions.
- Isotherm conditions.
- Isobaric conditions.
- Perfect gas law.
- Constant permeability.
- Perfect mixing conditions on upstream and downstream sides.
- Concentration polarization at the membrane is neglected.
- Pressure loss in the porous support layer is neglected.
- The permeate can drain off freely.

The calculation of the permeate composition is made with the following formula taken from the work of Melin and Rautenbach [31]:

$$y_{O_2} = \frac{1}{2} \left[1 + \phi * \left(x_{O_2Feed} + \frac{1}{\alpha - 1} \right) \right] - \sqrt{\left[\frac{1}{2} \left[1 + \phi * \left(x_{O_2Feed} + \frac{1}{\alpha - 1} \right) \right] \right]^2 - \frac{\alpha * \phi * x_{O_2Feed}}{\alpha - 1}}$$
(1)

$$y_{N_2} = 1 - y_{O_2} \tag{2}$$

where ϕ [-] is the feed-permeate pressure ratio given by the following equation:

$$\phi = p_{feed} / p_{permeate} \tag{3}$$

Figure 3 reports the permeate composition over pressure ratio for the 171 three membrane types considered. It can be seen that only a certain maxi-172 mum of oxygen ratio can be achieved because all the graphs are leveling off. 173 Therefore a pressure ratio of 1 MPa was chosen for further calculations as 174 suggested in Melin and Rautenbach [31] and the $p_{retentate}$ was fixed at 1 atm. 175 The Matrimid membrane is able to produce the highest oxygen ratio 176 of 0.58 % vol. in the permeate, however PPO membrane presents a good 177 value of oxygen ratio (0.49 % vol.) and an acceptable value of permeability, 178 therefore this membrane is adopted in the simulations. The active area of 179 the membrane can be assessed from the molar flow of oxygen required for 180 the gasification Q_{PO_2} [mol/s]: 181

$$A_{membrane} = \frac{Q_{PO_2}}{\gamma * (x_{O_2Feed} * p_{feed} + y_{O_2} * p_{permeate})}$$
(4)

The molar flow of nitrogen Q_{PN_2} [mol/s] and the total permeate molar flow Q_P [mol/s] is given by the following equations:

$$Q_{PN_2} = \frac{\gamma}{\alpha} * \left[p_{feed} * (1 - x_{O_2Feed}) + p_{permeate} * (1 - y_{O_2}) \right]$$
(5)

$$Q_P = Q_{PN_2} + Q_{PO_2} \tag{6}$$

The molar flow of the inlet air Q_{air} , the retentate molar flow Q_R and the retentate composition ($x_{O_2Retentate}$ and $x_{N_2Retentate}$) are calculated setting to zero the amount of oxygen in the retentate flow as suggested by Melin and ¹⁸⁷ Rautenbach[31]. Thus, a mass balance equation can be applied to estimate ¹⁸⁸ Q_{air} and Q_R :

$$Q_{air} * x_{O_2Feed} = Q_P * y_{O_2} \rightarrow Q_{air} = \frac{Q_P * y_{O_2}}{x_{O_2Feed}}$$
(7)

$$Q_{air} = Q_P + Q_R \to Q_R = Q_{air} - Q_P \tag{8}$$

Finally, the electrical power consumption to pressurize the inlet air flow is calculated as a polytropic compression by Equation 15 assuming $T_{in} = 20$ o C; m = 1.2 and $\eta_{comp} = 90$ %.

192 2.2. Gasifier modeling

In this work, the gasification process is simulated using a black-box model based on Barman's work [44]. The model is validated for downdraft gasifiers; it is based on the following gasification equation:

$$CH_{x}O_{y}N_{z} + wH_{2}O + m(O_{2} + 3.76N_{2}) \rightarrow$$

$$n_{H_{2}}H_{2} + n_{CO}CO + n_{CO_{2}}CO_{2} + n_{H_{2}O}H_{2}O$$

$$+ n_{CH_{4}}CH_{4} + (z/2 + 3.76m)N_{2} + n_{tar}CH_{p}O_{q}$$
(9)

where $CH_xO_yN_z$ is the equivalent chemical formula of "dry and ash free" (daf) biomass; CH_pO_q is the equivalent chemical formula of tar [45]; $w [mol/mol_{bio}]$ is the specific molar amount of the biomass moisture; m $[mol/mol_{bio}]$ is the specific molar amount of oxygen calculated starting from the equivalence ratio ER as suggested by Jarungthammachote and Dutta[46]; $n_{H_2}, n_{CO}, n_{CO_2}, n_{H_2O}, n_{CH_4}, n_{tar} [mol/mol_{bio}]$ are the specific molar amounts of $H_2, CO, CO_2, H_2O, CH_4$ and tar which constitute the syngas. This model is used and discussed in several other works [47, 48, 49]. It consists of a chemical and a thermal sub-models that converge to the final composition of the gas. The first step is to choose an initial temperature T[K] and calculate the equilibrium constant of the following reactions:

• **K1**: Water-gas shift
$$CO + H_2O \leftrightarrow CO_2 + H_2$$

- **K2:** Hydrogasification $C + 2H_2 \leftrightarrow CH_4$
- **K3:** Methane steam reforming $CH_4 + H_2O \leftrightarrow CO + 3H_2$

The system of equations 10 reported below is composed of three chemical balances calculated from Equation 9 (carbon, hydrogen and oxygen) and the three equilibrium constants for water-gas, hydrogasification and methane reforming reactions. The system is solved with the Newton-Raphson method.

$$n_{CO} + n_{CO_2} + n_{CH_4} + n_{tar} - 1 = 0$$

$$2n_{H_2} + 2n_{H_2O} + 4n_{CH_4} + pn_{tar} - x - 2w = 0$$

$$n_{CO} + 2n_{CO_2} + n_{H_2O} + qn_{tar} - w - 2m - y = 0$$

$$K_1 = \frac{n_{CO_2} * n_{H_2}}{n_{CO} * n_{H_2O}}$$

$$K_2 = \frac{n_{CH_4} * \frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}}{n_{H_2}^2}$$

$$K_3 = \frac{n_{CO} * n_{H_2}^3}{\left(\frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}\right)^2 n_{H_2O} n_{CH_4}}$$
(10)

Once the molar specific amounts of the syngas species are evaluated, it is possible to solve the thermodynamic energy balance of the system reported in the following equation:

$$\sum_{j=react} n_j * HF_j^0 = \sum_{i=prod} n_i * \left(HF_i^0 + \Delta H_{T,i} \right)$$
(11)

where n_j [moles] and HF_j^0 [kJ/kmol] are the specific moles amount and standard heat of formation of the j-th reagent (biomass, air and moisture); n_i [moles] and HF_i^0 [kJ/kmol] are the specific moles amount and the standard heat of formation of the i-th product (H_2 , CO, CO_2 , H_2O , CH_4 and N_2) and $\Delta H_{T,i}$ is the enthalpy difference between any given state and the standard state for the i-th product. $\Delta H_{T,i}$ can be calculated starting from the specific heat of the product:

$$\Delta H_{T,i} = \int_{298.15}^{T} C_p(T) \, dT = \left| aT + b\frac{T^2}{2} + c\frac{T^3}{3} + d\frac{T^4}{4} \right|_{298.15}^{T} \tag{12}$$

where the coefficient a,b,c and d are defined for each gas by Jarungtham-224 machote and Dutta[46]. In order to find the equilibrium temperature T_{new} , 225 the system is considered adiabatic and the the Newton-Raphson method is 226 applied to the equations. If $abs(T - T_{new}) < 0.1$ K then the calculated equi-227 librium temperature and molar specific gases amounts are the final results; 228 otherwise, a new iteration is done in order to satisfy the previous condi-229 tion. The model is implemented in Python and the input are the biomass 230 equivalent molecule, the equivalence ratio ER and the initial temperature. 231 The temperature input is used only as a starting point for the iterating 232 system; after few cycles the temperature converges to the ones that satisfy 233 both the chemical and thermal sub-systems. About the ER, a value of 0.335234
is assumed. This value is consistent with air blown gasification parameters
[50, 37] and it is confirmed by the low tar content in the syngas. Poplar wood
chips properties and gasifier model parameters are summarized in Table 1.

238 2.3. ZEO module modeling

The zeolite filter is able to reduce the total syngas molar flow of about 240 20% - 30 % by the adsorption of CO_2 . Zeolite 5A is chosen because it has 241 a great selectivity for carbon dioxide in comparison with the other gases 242 that constitute syngas [38]. The gas adsorption in porous solids has been 243 described by the Langmuir equation [38, 39]:

$$q_i = \frac{q_{mi} * B_i * p_i}{1 + \sum_{j=1}^n B_j * p_j}$$
(13)

where $q_i \, [\text{mmol/g}]$ is the adsorbed amount of the component $i; q_{mi} \, [\text{mmol/g}]$ 244 is the saturation adsorbed amount of the component i; B_i [1/kPa] is the Lang-245 muir constant of the component i; p_i [kPa] is the equilibrium partial pressure 246 of the component i; B_j [1/kPa] is the Langmuir constant of the component 247 j; p_j [kPa] is the equilibrium partial pressure of the component j; i and j are 248 the gas species of the syngas. Table 1 reports the Langmuir constants and 249 the saturation adsorbed amounts for Zeolite 5A, while Figure 4 depicts the 250 adsorption trends of the syngas gases as function of pressure. It can be noted 251 the high CO_2 selectivity of the zeolite in comparison with others gases. 252

The mass of zeolite required for adsorbing all the carbon dioxide of the syngas depends on the molar flow of the dry syngas, its CO_2 molar fraction and kinetic constant of adsorption. The ZEO filter module can be constantly regenerated using a rotary valve packaged into modules as described in [40]. The mass of zeolite that needs to be regenerated every cycle with duration of t_{cycle} can be calculated as follows:

$$m_{zeo} = t_{cycle} * \dot{n}_{DG} * \frac{1 + \sum_{j=1}^{n} B_j * p_j}{q_{m,CO_2} * B_{CO_2} * p_{ads} * x_{CO_2}}$$
(14)

where p_{ads} [kPa] is the total pressure of the syngas inside the ZEO filter. 259 A constant temperature of the zeolite filter and of the inlet syngas of 303 K is 260 assumed and the pressure of the inlet syngas is set to 500 kPa as suggested in 261 [38, 39]. The cycling time of regeneration depends on kinetic CO_2 adsorption 262 constant. In this study a plausible time of 60 seconds is assumed and future 263 work will investigate this aspect. Zeolites adsorption generates heat, Ranjani 264 et al. [51] suggests that 64-70 kJ are released for every mole of CO_2 adsorbed. 265 This heat needs to be discharge by the ZEO module in order to keep the 266 temperature constant at 303 K. In this preliminary study, no attention was 267 paid to the ZEO module heat balance. Furthermore, the gas filtering sub-268 system considered in this work is based on the power plant described by 269 Allesina et al. [50]. It was designed with the idea of coupling the gasifier with 270 an internal combustion engine. Since the minimum presence of tars could 271 negatively affect the performance of the zeolite adsorber, syngas purification 272 unit should be properly designed. A potential alternative to water scrubber 273 is oil scrubber with subsequent stripping of tars [52, 53]. 274

275 2.4. Compressor and storage system modeling

The modeling of the syngas compression is carried out considering it as a polytropic transformation. The electrical power required for compression is given by Pedrazzi et al. [54]:

$$P_{comp} = \frac{\dot{n}_{gas} L_{comp,is}}{\eta_{comp}} = \frac{\dot{n}_{gas}}{\eta_{comp}} \frac{zRT_{in}}{z-1} \left[1 - \left(\frac{p_{out}}{p_{in}}\right)^{\frac{z-1}{z}} \right]$$
(15)

where z is the polytrophic coefficient, R is the universal gas constant 279 equal to 8.314 J mol⁻¹ K⁻¹, T_{in} is the gas inlet temperature (25°C for syngas 280 compressor 1 and 30°C for syngas compressor 2), p_{in} and p_{out} [atm] are the gas 281 inlet and outlet pressures, \dot{n}_{gas} [mol/s] is the gas molar flow and η_{comp} is the 282 compressor efficiency available from manufacturer's data [54]. The maximum 283 pressure value inside the storage system is a fundamental parameter required 284 todesign the tanks and the compressor properly. Assuming ideal gas and a 285 constant syngas storage temperature $T_s = 25$ °C, the pressure inside the 286 tanks is calculated by the ideal gas law: 287

$$p_s = \frac{nRT_s}{V} \tag{16}$$

where n [mol] are the moles of syngas inside the tanks and $V \text{ [m^3]}$ is the storage total volume. Assuming a value of the initial syngas moles n_{in} inside the storage, the moles of syngas at the time τ [s] are given by:

$$n = n_{in} + \int_0^\tau \left(\dot{n}_{in,s}(t) - \dot{n}_{out,s}(s) \right) dt$$
 (17)

where $\dot{n}_{in,s}(t)$ and $\dot{n}_{out,s}(t)$ [mol/s] are the inlet and the outlet molar flow at the instantaneous time t [s]. Table 2 reports the model parameters of the storage and compressor sub-systems. The total volume of storage and the initial syngas amount in the tank are reduced of about 50% in comparison with the conventional system without PPO and ZEO modules as investigated in [55, 56]. This result is reached thanks to the PPO adoption that decreases the molar flow of dry syngas of about the 20% - 30%, while the filtration in the ZEO module further reduces the syngas molar flow of about another 20% - 30% as shown in the results.

300 2.5. SOFC modeling

The SOFC model used in this study is based on the work of Bang-Møller 301 and Rokni [18]. This model does not take into account the recirculation 302 of gas at the cell anode. This feature may strongly compromise the fuel 303 cell efficiency in case of the presence of gases that do not take part in the 304 electro-chemical, shift and reforming reactions. Unfortunately, syngas con-305 tains considerable amounts of CO_2 and N_2 . To overcome this issue, the 306 model previously cited is implemented with the reforming model presented 307 by Rami Salah El-Emam et al. [22]. As described by Rami Salah El-Emam, 308 the electro-chemical reactions take place in both the anode and the cathode 309 of the cell (Eq.22), while the reforming and the monoxide water shift occur 310 only near the anode (Eqs. 18, 19). Equation 22 presents the overall electro-311 chemical reaction that is divided into two sub-reactions: the hydrogen reacts 312 with the oxygen ions to form water and electrons according to Eq. 20 at 313 the anode, while, at the cathode, the oxygen from inlet air reacts with the 314 electrons from the anode (Eq. 21) to form oxygen ions that flow to the anode 315 through the solid oxide electrolyte. 316

$$CH_4 + H_2O \to CO + 3H_2 \tag{18}$$

$$CO + H_2O \to CO_2 + H_2 \tag{19}$$

$$H_2 + O^{2-} \to H_2O + 2e^-$$
 (20)

$$\frac{1}{2}O_2 + 2e^- \to O^{2-} \tag{21}$$

$$H_2 + \frac{1}{2}O_2 \to H_2O \tag{22}$$

The mathematical modeling of reforming and electrochemical reactions is explained in References [22] and [18]. Using these models, it is possible to calculate the electrical power production and the electrical conversion efficiency for a given syngas inlet flow with a specific composition. The SOFC model parameters adopted in the simulations are reported in Table 2.

322 2.6. MGT modeling

Mathematical description of gas turbines is well described in literature. Details and assumptions of the present model can be found in Bang-Møller and Rokni work [18]. Characteristics of the turbine and others components connected to the MGT are listed in Table 2.

327 3. Simulation results and discussion

In this work four different cases are simulated. First of all, the basic system composed of a downdraft gasifier, a storage tank and a SOFC-MGT is simulated. After this step, the two possible solutions consisting of N_2 purging from air or CO_2 separation from syngas are discussed separately. Finally, the complete system provided with PPO module and ZEO module is simulated.

334 3.1. Case I (Gasifier + SOFC + MGT)

The SOFC-MGT unit constantly produces energy all over the year of simulation in order to preserve the stability of the cells and their gaskets, which are very sensitive to thermal stresses [57]. The syngas molar flow consumed by the SOFC-MGT unit is calculated by Equation 23 that considers the cycling working of the DG:

$$\dot{n}_{syngas-SOFC} = \frac{h_{operation} \dot{n}_{DG}}{h_{operation} + h_{maintenance}}$$
(23)

340

Figure 5 depicts the overall model implemented in Matlab SimulinkTM soft-341 ware environment. Table 3 reports the simulation results. Gasifier cold 342 efficiency is about 79%, this value is confirmed by literature that suggests 343 an efficiency of 70% - 80% for air-blown downdraft gasifier [37, 4, 44]. Syn-344 gas composition consists of about 19% vol. of H_2 and 15% vol. of CO, the 345 higher heating value of 4.75 MJ/Nm^3 is similar to the results reported by 346 Basu [37] for this kind of gasifier. SOFC-MGT unit has a constant electrical 347 power production of 197.43 kW all over the simulated year. The auxiliary 348 consumption of the whole system strongly depends on tank pressurization 349 level. The average annual value is 34.24 kW. For this reason, the net average 350 power production is reduced to 163.19 kW and the electrical efficiency of the 351 system is 25.43%. 352

353 3.2. Case II (PPO + Gasifier + SOFC + MGT)

Table 4 shows the results of the simulation of the system previously described and now equipped with the PPO module. The oxygen enriched air

flow is about 1.27 mol/s. This value is lower than 2.94 mol/s obtained with-356 out PPO membrane (Case I). Syngas composition is consistent with the work 357 of Wang et al. [58], where oxygen-enriched air (50% oxygen and 50% nitro-358 gen) is used as gasifying agent in a double stage downdraft gasifier fueled 359 with pine sawdust pellets. Differences of 1-2% between the model outputs 360 and Wang's results about CO and H_2 contents are achieved (29% vs. 27%) 361 for H_2 and 26% Vs. 25% for CO). The gasification with oxygen-enriched air 362 assures high gasifier performance in terms of cold gas efficiency (92%), tar 363 production (0.27 g/Nm^3) and syngas higher heating value (7.55 MJ/Nm^3) . 364 The syngas outlet flow is 3.589 mol/s, consistently lower than Case I where 365 syngas flow is 5.01 mol/s. The average equilibrium temperature of gasifi-366 cation in this case is 931 K. This value is only 36 K higher than Case I. 367 Wang et al. [58] suggestes a peak temperature of about 1200 K with oxygen-368 enriched air. With this temperature, conventional material adopted in air 369 gasifier can be used (i.e. stainless steel and refractory brick [2, 6]). In Case 370 II, the overall net power production is 210.52 kW and the electrical efficiency 371 is boosted to 32.81%. The average auxiliary consumption is 42 kW, 8 kW 372 higher than Case I. This is due to the PPO module that uses air at 1 MPa 373 pressure generated by an air compressor. The electrical consumption of the 374 air compressor is 15.3 kW and it is fully compensated by the increasing of 375 the gasifier efficiency and the SOFC-MGT unit efficiency. 376

377 3.3. Case III (Gasifier + ZEO + SOFC + MGT)

Table 5 resumes the simulation results of the system with the ZEO filtering module instead of PPO membrane. The filtered syngas has a higher heating value of 5.9 MJ/Nm^3 . This value falls between Case I (4.75 MJ/Nm^3)

and Case II (7.55 MJ/Nm^3) . The pressure of the storage tank ranges between 381 0.267-0.488 MPa, similar to the values obtained in Case II (0.266-0.462 MPa) 382 and Case I (0.267-0.539 MPa). The zeolite mass required to perform con-383 tinuosly the filtration is 23.742 kg. The value obtained is consistent with 384 Tagliabue et al. work [40]. However, in future work, the CO_2 adsorbed by 385 the ZEO module can be stored in order to create a carbon sequestration sys-386 tem. The power production and the net electrical efficiency of the system is 387 low (148.74 kW of power production and 22.87% of electrical efficiency) as a 388 consequence of the energy absorbed by the syngas compressor 1 (see Figure 389 1) to increase the pressure of the syngas to 0.5 MPa before the ZEO module. 390 This electrical energy consumption is higher than Cases II and I, in addition 391 the efficiency of the SOFC-MGT module fueled with the filtered syngas is 392 lower. As shown in Table 5, the SOFC-MGT efficiency in Case III is about 393 34%, thus lower than Case II (42.08%) and Case I (37.46%). 394

395 3.4. Case IV (PPO + Gasifier + ZEO + SOFC + MGT)

The results about the fully equipped gasifier power system are reported 396 in Table 6. A high power production (194.53 kW) and electrical efficiency 397 (30.32%) is reached thanks to the high H_2 and CO amounts in the filtered 398 syngas. In fact, the H_2 volume percentange reaches 41.53% and the CO 399 volume percentange is boosted to 32.54%. As a consequence of this com-400 position, the higher heating value of the syngas is 10.19 MJ/Nm^3 , a value 401 typical for oxygen-blown gasifiers [37]. Therefore, the SOFC-MGT unit syn-402 gas consumption is 2.696 mol/s. This value is about 45% lower than Case I 403 (4.95 mol/s), 24% lower than Case II (3.545 mol/s) and 33% lower than Case 404 III (4.02 mol/s). A pressure range of 0.266-0.415 MPa is achieved. In this 405

case power production and efficiency is lower than Case II as result of higher average auxiliary consumption (51.80 kW) and lower SOFC-MGT unit efficiency (38.41%). However, the utilization of the ZEO module has several advantages: separates the CO_2 and reduces the storage peaks pressure.

410 3.5. Performance and energy considerations

Figure 6 shows the electrical efficiency and the average power production 411 in every scenario. Case II resulted the best in terms of energy conversion; the 412 overall electrical efficiency reaches 32% and the power production is about 413 210 kW. These values are higher than commercial gasification power systems 414 with internal combustion engines where the maximum electrical efficiency 415 hardly reaches 25% [37, 4, 2]. This result is given by the PPO module that 416 increases the gasifier efficiency to 92% (75% is the reference value for air 417 blown gasifier [37]) and the SOFC-MGT module which has a higher electri-418 cal conversion efficiency (about 42%) compared to common engine-alternator 419 generator units (about 27% [4, 59, 3]). Case II is the best in terms of energy 420 balance as shown in Figure 7. These graphs do not consider the thermal 421 energy that can be recovered from the gasifier or the SOFC-MGT unit. The 422 highest energy loss occurs at the SOFC-MGT unit (about 52%), while aux-423 iliary consumption of the blowers and the auxiliary equipment of the gasifier 424 are low (9%). In Cases I and III, the low efficiency of the gasifier reduces the 425 overall electrical performance of the system. In Cases III and IV the ZEO 426 module consumes energy to separate the CO_2 from the gas, however no effi-427 ciency increase occours in the SOFC-MGT unit with a CO_2 free syngas and 428 the final result is a lower production. The system modeled in this work 429 is obtained starting from a reference power plant described by Allesina et. al 430

[50] where IC engines are used instead of the SOFC-MGT unit. The author 431 reports an experimental cold gasification efficiency of 67%. Considering an 432 electrical IC engine-alternator unit efficiency of about 27%, as suggested by 433 Puglia at al. [3], the total electrical efficiency is about 18%. This value is 434 30% lower than Case I and it is 45% lower than Case II. Another study made 435 by Patuzzi et al. [60] reports the values of the net electrical efficiency of three 436 different commercial biomass gasifier - IC engine power plants. The average 437 efficiency is about 20%, this value is consistent with the one obtained for 438 Allesina et al. [50]. 439

440 4. Conclusions

The biomass fueled system with PPO module (Case II) shows the higher 441 electrical efficiency of about 33%. The reasons behind this result are vari-442 ous. First of all, oxygen-enriched air boosts the gasifier cold efficiency from 443 79% (Case I with air) to 92% (Cases II and IV with oxygen-enriched air). 444 In addition, the SOFC-MGT unit presents a higher efficiency (about 42%) 445 compared to IC engine-alternator unit (about 27%), ORC cycle (about 20%) 446 or EFGT cycle (about 20%). In Cases III and IV, the zeolites adsoption 447 module consumes energy to increase the higher heating value of the syngas 448 but not the performance of the SOFC-MGT system, this reduces the overall 449 system efficiency. The energy balances of four cases investigated show that 450 the greater losses are in the SOFC-MGT unit. This unit has the difficult 451 task to convert the chemical energy of a gas fuel into electrical energy in 452 an efficient way. An efficiency of about 50% is reached with natural gas, in 453 this study the maximum electrical efficiency is about 42% using a syngas 454

produced with an oxygen-enriched air as gasifying agent. This difference is 455 given by the presence of several inert gases into the syngas that reduces the 456 electrochemical conversion of the SOFC. The removal or the conversion of 457 these gases into syntethic natural gases (SNG) is possible and, in this way, 458 the efficiency of the SOFC-MGT unit will be similar to the value reach for 459 natural gas one. But, cost and energy self-consumption of the upgrading 460 process are very high and not convenient for this kind of power plants. Cases 461 III and IV has a lower efficiency compared to Case II, however, with the 462 ZEO module, it is possible to separate the CO_2 content of the syngas with 463 environmental benefits in case the module is coupled with a CO_2 sequestra-464 tion system. Future work will consider exergy calculations and experimental 465 tests on a micro-scale power system (5-20 kW of electrical power) with PPO 466 module and SOFC module in order to validate modeling results and to assest 467 system durability. In addition, economical net present value analysis will be 468 done to estimate the economic sustainability of the power plant. 460

470 Nomenclature

- 471 \dot{m} mass flow [kg/s]
- 472 \dot{n} molar flow [mol/s]
- 473 au time [s]
- ⁴⁷⁴ ASH ash content of the biomass [%]
- $_{475}$ B Langmuir constant [1/kPa]
- 476 C carbon

- 477 C_p specific heat [J/(mol K)]
- $_{478}$ DG downdraft gasifier
- 479 e electron
- $_{480}$ EFGT external firing gas turbine
- 481 ER equivalence ratio [ad]
- 482 H hydrogen
- 483 H_T enthalpy [kJ/kmol]
- $_{484}$ HF^0 standard heat of formation [kJ/kmol]
- $_{485}$ HHV higher heating value [MJ/Nm³ or MJ/kg]
- $_{486}$ IC internal combustion
- $_{487}$ K equilibrium constant [ad]
- 488 L work [kJ]
- 489 M total moisture content of the biomass [%]
- 490 m specific molar amount of oxygen $[mol/mol_{bio}]$
- 491 m_{tar,Nm^3} volumetric tar amount $[g/Nm^3]$
- $_{492}$ MCFC molten carbonate fuel cell
- $_{493}$ MGT micro gas turbine
- 494 MW molecular weight [g/mol]

- 495 N nitrogen
- 496 n specific molar amount of gases and tar $[mol/mol_{bio}]$
- 497 *O* oxygen
- ⁴⁹⁸ ORC organic rankine cycle
- 499 P power [kW]
- 500 p pressure [atm]
- ⁵⁰¹ *PDSM* polydimethylsiloxan
- 502 PPO polyphenylene oxide
- $_{503}$ Q molar flow [mol/s]
- $_{504}$ q adsorbed amount [mmol/g]
- $_{505}$ R universal gas constant [J/(mol K)]
- 506 SOFC solide oxide fuel cell
- 507 T temperature [K]
- 508 t time [s]
- 509 V volume [m³]
- 510 w specific molar amount of biomass moisture $[mol/mol_{bio}]$
- $_{511} x$ molar fraction
- $_{512}$ y molar fraction

513 z polytrophic coefficien

- 514 ZEO zeolite
- 515 α selectivity
- 516 Δ difference
- $_{517} \gamma$ permeability [mol m⁻2 s⁻1 bar⁻1
- 518 ϕ pressure ratio

519 Subscripts

- 520 ads adsorption
- 521 ar as received
- 522 *bio* biomass
- 523 *comp* compressor
- $_{524}$ daf dry ash free
- $_{525}$ db dry basis
- 526 *g* gas
- 527 in inlet
- 528 *m* saturation
- 529 *out* outlet
- 530 P permeate

- $_{531}$ p hydrogen coefficient of tar
- 532 prod product
- $_{533}$ q oxygen coefficient of tar
- $_{534}$ R retentate
- 535 *react* reactant
- 536 s storage
- $_{537}$ x hydrogn coefficient of the biomass
- $_{538}$ y hydrogen coefficient of the biomass
- $_{539}$ z nitrogen coefficient of the biomass

540 **References**

- [1] IEA. Technology roadmap bioenergy for heat and power. Technical
 report, IEA, 2012.
- [2] Thomas B. Reed and Agua Das. Handbook of Biomass Downdraft Gasi *fier Engine Systems*. The biomass energy foundation press, 1988.
- [3] G. Allesina, S. Pedrazzi M.Puglia, and P. Tartarini. Upgrading or substituting the gasification process for electrical energy production: an
 energy-based comparison. In XXX UIT Conference, Bologna 2012, 2012.
- ⁵⁴⁸ [4] H.A.M. Knoef. Handbook of Biomass Gasification. BTG, 2005.

- [5] Amitava Datta, Ranjan Ganguly, and Luna Sarkar. Energy and exergy analyses of an externally fired gas turbine (efgt) cycle integrated with biomass gasifier for distributed power generation. *Energy*, 35(1):341 – 350, 2010.
- ⁵⁵³ [6] FAO Forestry Department Mechanical Wood Products Branch. Woodgas
 ⁵⁵⁴ as engine fuel, volume ISBN 92-5-102436-7. F.A.O., 1986.
- F. Martelli, G. Riccio, S. Maltagliati, and D. Chiaramonti. Technical
 study and environmental impact of an external fired gas turbine power
 plant fed by solid fuel. 1st world Conference of Biomass, Sevilla, 2000.
- ⁵⁵⁸ [8] V. Naso. La macchina di Stirling. CEA, 1991.
- [9] C. Souleymane. Motori a combustione interna e turbine a gas di piccola taglia per gas di sintesi. Master's thesis, Università degli Studi di
 Padova, Italy., 2012.
- ⁵⁶² [10] Farqad Al-Hadeethi, Moh'd Al-Nimr, and Mohammad Al-Safadi. Us⁵⁶³ ing the multiple regression analysis with respect to {ANOVA} and 3d
 ⁵⁶⁴ mapping to model the actual performance of {PEM} (proton exchange
 ⁵⁶⁵ membrane) fuel cell at various operating conditions. *Energy*, 90, Part
 ⁵⁶⁶ 1:475 482, 2015.
- [11] Guido Galeno. Modellizzazione di un micro cogeneratore basato sulla
 tecnologia mcfc accoppiata ad un gassificatore di biomassa. PhD thesis,
 University of Cassino, Italy, 2006-2007.
- 570 [12] Giulio Donolo, Giulio De Simon, and Maurizio Fermeglia. Steady

state simulation of energy production from biomass by molten carbonate fuel cells. Journal of Power Sources, 158(2):1282 – 1289, 2006.
ice:title¿Special issue including selected papers from the 6th International Conference on Lead-Acid Batteries (LABAT 2005, Varna, Bulgaria) and the 11th Asian Battery Conference (11 ABC, Ho Chi Minh
City, Vietnam) together with regular papersi/ce:title¿.

[13] Stefano Cordiner, Massimo Feola, Vincenzo Mulone, and Fabio Romanelli. Analysis of a {SOFC} energy generation system fuelled with
biomass reformate. Applied Thermal Engineering, 27(4):738 – 747, 2007.
jce:title; Energy: Production, Distribution and Conservation;/ce:title;.

- [14] E. Achenbach. Three-dimensional and time-dependent simulation of a
 planar solid oxide fuel cell stack. *Journal of Power Sources*, 49:333–348,
 1994.
- [15] M. Mortazaei and M. Rahimi. A comparison between two methods of
 generating power, heat and refrigeration via biomass based solid oxide
 fuel cell: A thermodynamic and environmental analysis. *Energy Con*-*version and Management*, 126:132 141, 2016.
- [16] F. Calise, M. Dentice dâTMAccadia, A. Palombo, and L. Vanoli. Simulation and exergy analysis of a hybrid solid oxide fuel cell (sofc)â"gas turbine system. *Energy*, 31(15):3278 3299, 2006. {ECOS} 2004 17th
 International Conference on Efficiency, Costs, Optimization, Simulation, and Environmental Impact of Energy on Process Systems 17th International Conference on Efficiency, Costs, Optimization, Simulation, and Environmental Impact of Energy on Process Systems.

- [17] L. Fryda, K.D. Panopoulos, and E. Kakaras. Integrated {CHP} with
 autothermal biomass gasification and sofc-mgt. *Energy Conversion and Management*, 49(2):281 290, 2008.
- [18] C. Bang-Moller and M. Rokni. Thermodynamic performance study of
 biomass gasification, solid oxide fuel cell and micro gas turbine hybrid
 systems. *Energy Conversion and Management*, 51(11):2330 2339, 2010.
- [19] Made Sucipta, Shinji Kimijima, and Kenjiro Suzuki. Performance analysis of the sofc-mgt hybrid system with gasified biomass fuel. Journal
 of Power Sources, 174(1):124 135, 2007. jce:title¿Hybrid Electric Vehiclesj/ce:title¿.
- [20] Pegah Ghanbari Bavarsad. Energy and exergy analysis of internal re forming solid oxide fuel cellâ"gas turbine hybrid system. International
 Journal of Hydrogen Energy, 32(17):4591 4599, 2007. Fuel Cells.
- [21] C. Ozgur Colpan, Ibrahim Dincer, and Feridun Hamdullahpur. Thermodynamic modeling of direct internal reforming solid oxide fuel cells operating with syngas. *International Journal of Hydrogen Energy*, 32(7):787
 795, 2007. jce:title¿Fuel Cellsj/ce:title¿.
- [22] Rami Salah El-Emam, Ibrahim Dincer, and Greg F. Naterer. Energy
 and exergy analyses of an integrated {SOFC} and coal gasification system. *International Journal of Hydrogen Energy*, 37(2):1689 1697, 2012.
 ice:title; 10th International Conference on Clean Energy 2010i/ce:title;.
- ⁶¹⁶ [23] Penyarat Chinda and Pascal Brault. The hybrid solid oxide fuel cell

- (sofc) and gas turbine (gt) systems steady state modeling. International
 Journal of Hydrogen Energy, 37(11):9237 9248, 2012.
- [24] S.H. Chan, H.K. Ho, and Y. Tian. Modelling of simple hybrid solid
 oxide fuel cell and gas turbine power plant. *Journal of Power Sources*,
 109(1):111 120, 2002.
- [25] Tae Won Song, Jeong Lak Sohn, Jae Hwan Kim, Tong Seop Kim,
 Sung Tack Ro, and Kenjiro Suzuki. Performance analysis of a tubular solid oxide fuel cell/micro gas turbine hybrid power system based on
 a quasi-two dimensional model. *Journal of Power Sources*, 142(1â"2):30
 42, 2005.
- [26] Mahsa Aghaie, Mehdi Mehrpooya, and Fathollah Pourfayaz. Introducing an integrated chemical looping hydrogen production, inherent carbon
 capture and solid oxide fuel cell biomass fueled power plant process configuration. Energy Conversion and Management, 124:141 154, 2016.
- [27] Masood Ebrahimi and Iraj Moradpoor. Combined solid oxide fuel cell,
 micro-gas turbine and organic rankine cycle for power generation (sofc mgt-orc). Energy Conversion and Management, 116:120 133, 2016.
- [28] Ph. Hofmann, K.D. Panopoulos, P.V. Aravind, M. Siedlecki,
 A. Schweiger, J. Karl, J.P. Ouweltjes, and E. Kakaras. Operation of
 solid oxide fuel cell on biomass product gas with tar levels >10 g
 nmâ'3. International Journal of Hydrogen Energy, 34(22):9203 9212,
 2009.

- [29] Lopamudra Devi, Krzysztof J Ptasinski, and Frans J.J.G Janssen. A review of the primary measures for tar elimination in biomass gasification
 processes. *Biomass and Bioenergy*, 24(2):125–140, 2003.
- [30] Jinsong Zhou, Qing Chen, Hui Zhao, Xiaowei Cao, Qinfeng Mei,
 Zhongyang Luo, and Kefa Cen. Biomass-oxygen gasification in a hightemperature entrained-flow gasifier. *Biotechnology Advances*, 27(5):606
 611, 2009. Bioenergy Research & amp; Development in ChinaICBT
 2008.
- ⁶⁴⁷ [31] T. Melin and R. Rautenbach. Membranverfahren, Grundlagen der
 ⁶⁴⁸ Modul- und Anlagenauslegung. 2007.
- [32] K.C. Khulbe, T. Matsuura, G. Lamarche, and H.J. Kim. The morphology characterisation and performance of dense ppo membranes for gas
 separation. *Journal of Membrane Science*, 135(2):211 223, 1997.
- [33] Janusz Kotowicz and Adrian Balicki. Enhancing the overall efficiency
 of a lignite-fired oxyfuel power plant with {CFB} boiler and membranebased air separation unit. *Energy Conversion and Management*, 80:20
 31, 2014.
- [34] Giacomo Bisio, Alessandro Bosio, and Giuseppe Rubatto. Thermodynamics applied to oxygen enrichment of combustion air. *Energy Con- version and Management*, 43(18):2589 2600, 2002.
- [35] H. Scott Coombe and Sen Nieh. Polymer membrane air separation per formance for portable oxygen enriched combustion applications. *Energy Conversion and Management*, 48(5):1499 1505, 2007.

- [36] Yanhong Hao, Yi Huang, Minhui Gong, Wenying Li, Jie Feng, and Qun
 Yi. A polygeneration from a dual-gas partial catalytic oxidation coupling
 with an oxygen-permeable membrane reactor. *Energy Conversion and Management*, 106:466 478, 2015.
- [37] Prabir Basu. Biomass Gasification and Pyrolysis: Practical Design and
 Theory. Academic Press, Elsevier, 2010.
- [38] Saeed Pakseresht, Mohammad Kazemeini, and Mohammad M. Akbarnejad. Equilibrium isotherms for co, co2, {CH4} and {C2H4} on the 5a
 molecular sieve by a simple volumetric apparatus. Separation and Pu-*rification Technology*, 28(1):53 60, 2002.
- [39] Gi-Moon Nam, Byung-Man Jeong, Seok-Hyun Kang, Byung-Kwon Lee,
 and Dae-Ki Choi. Equilibrium isotherms of ch4, c2h6, c2h4, n2, and h2
 on zeolite 5a using a static volumetric method. *Journal of Chemical & Engineering Data*, 50(1):72–76, 2005.
- [40] Marco Tagliabue, David Farrusseng, Susana Valencia, Sonia Aguado,
 Ugo Ravon, Caterina Rizzo, Avelino Corma, and Claude Mirodatos.
 Natural gas treating by selective adsorption: Material science and chemical engineering interplay. *Chemical Engineering Journal*, 155(3):553 –
 566, 2009.
- [41] Zoltán Bacsik, Ocean Cheung, Petr Vasiliev, and Niklas Hedin. Selective
 separation of {CO2} and {CH4} for biogas upgrading on zeolite naka
 and sapo-56. Applied Energy, 162:613 621, 2016.

- [42] Mariem Kacem, Mario Pellerano, and Arnaud Delebarre. Pressure swing
 adsorption for co2/n2 and co2/ch4 separation: Comparison between activated carbons and zeolites performances. *Fuel Processing Technology*,
 138:271 283, 2015.
- [43] Qassim Hassan Dirar and Kevin F. Loughlin. Intrinsic adsorption properties of co2 on 5a and 13x zeolite. Adsorption, 19(6):1149–1163, 2013.
- [44] Niladri Sekhar Barman, Sudip Ghosh, and Sudipta De. Gasification of
 biomass in a fixed bed downdraft gasifier a realistic model including
 tar. *Bioresource Technology*, 107:505–511, 2012.
- [45] Takashi Yamazaki, Hirokazu Kozu, Sadamu Yamagata, Naoto Murao,
 Sachio Ohta, Satoru Shiya, and Tatsuo Ohba. Effect of superficial ve locity on tar from downdraft gasification of biomass. *Energy & Fuels*,
 19:1186–1191, 2005.
- [46] S. Jarungthammachote and A. Dutta. Thermodynamic equilibrium
 model and second law analysis of a downdraft waste gasifier. *Energy*,
 32(9):1660 1669, 2007.
- [47] Giulio Allesina, Simone Pedrazzi, Luca Guidetti, and Paolo Tartarini.
 Modeling of coupling gasification and anaerobic digestion processes for
 maize bioenergy conversion. *Biomass and Bioenergy*, 81:444 451, 2015.
- [48] Giulio Allesina, Simone Pedrazzi, Federico Sgarbi, Elisa Pompeo,
 Camilla Roberti, Vincenzo Cristiano, and Paolo Tartarini. Approaching
 sustainable development through energy management, the case of fongo

- tongo, cameroon. International Journal of Energy and Environmental
 Engineering, 6(2):121–127, 2014.
- [49] Giulio Allesina, Simone Pedrazzi, Emma La Cava, Michele Orlandi,
 Miriam Hanuskova, Caludio Fontanesi, and Paolo Tartarini. Energybased assessment of optimal operating parameters for coupled biochar
 and syngas production in stratified downdraft gasifiers. In *International Heat Transfer Conference 15, Kyoto, Japan*, 2014.
- ⁷¹³ [50] Giulio Allesina, Simone Pedrazzi, and Paolo Tartarini. Modeling and
 ⁷¹⁴ investigation of the channeling phenomenon in downdraft stratified
 ⁷¹⁵ gasifers. *Bioresource Technology*, 146(0):704 712, 2013.
- [51] Ranjani V. Siriwardan, Ming-Shing Shen, and Edward P. Fisher. Adsorption of co2 on zeolites at moderate temperatures. *Energy and Fuels*, 19:1153–1159, 2005.
- [52] Thana Phuphuakrat, Tomoaki Namioka, and Kunio Yoshikawa. Absorptive removal of biomass tar using water and oily materials. *Bioresource Technology*, 102(2):543–549, 2011.
- [53] Shunsuke Nakamura, Shigeru Kitano, and Kunio Yoshikawa. Biomass
 gasification process with the tar removal technologies utilizing bio-oil
 scrubber and char bed. *Applied Energy*, 170:186 192, 2016.
- ⁷²⁵ [54] S. Pedrazzi, G. Zini, and P. Tartarini. Complete modeling and soft⁷²⁶ ware implementation of a virtual solar hydrogen hybrid system. *Energy*⁷²⁷ *Conversion and Management*, 51(1):122 129, 2010.

- [55] Simone Pedrazzi, Giulio Allesina, Alberto Muscio, and Paolo Tartarini.
 Modeling and simulation of a dg-sofc-mgt hybrid system. In 7° Congresso Nazionale AIGE, Rende (CZ) Italy, 2013.
- [56] Simone Pedrazzi. Modeling and optimization of advanced systems for
 electrical energy production from wood biomass. PhD thesis, HIGH MECHANICS AND AUTOMOTIVE DESIGN & TECHNOLOGY, University of Modena and Reggio Emilia, Dep. of Engineering 'Enzo Ferrari',
 2013.
- ⁷³⁶ [57] DOE. Fuel Cell Handbook (Seventh Edition). 2004.
- ⁷³⁷ [58] Zhiqi Wang, Tao He, Jianguang Qin, Jingli Wu, Jianqing Li, Zhongyue
 ⁷³⁸ Zi, Guangbo Liu, Jinhu Wu, and Li Sun. Gasification of biomass with
 ⁷³⁹ oxygen-enriched air in a pilot scale two-stage gasifier. *Fuel*, 150:386 –
 ⁷⁴⁰ 393, 2015.
- [59] H.A.M. Knoef. Handbook of Biomass Gasification, Second Edition.
 BTG, 2012.
- [60] Francesco Patuzzi, Dario Prando, Stergios Vakalis, Andrea Maria Rizzo,
 David Chiaramonti, Werner Tirler, Tanja Mimmo, Andrea Gasparella,
 and Marco Baratieri. Small-scale biomass gasification {CHP} systems:
 Comparative performance assessment and monitoring experiences in
 south tyrol (italy). *Energy*, 112:285 293, 2016.
- [61] S. Pedrazzi, G. Allesina, and P. Tartarini. Aige conference: A kinetic
 model for a stratified downdraft gasifier experimental assessment and

- ⁷⁵⁰ modeling of energy conversion effectiveness in a gasification power plant.
- ⁷⁵¹ International Journal of Heat and Technology, 30(1):41–44, 2012.
- 752 Figure captions and tables



Figure 1: DG-SOFC-MGT hybrid system with zeolite CO_2 adsorption and oxygenenriched air layout



Figure 2: Oxygen enriched air membrane separator principle



Figure 3: Characteristics of the separation membranes



Figure 4: Zeolite 5A adsorbing curve Vs. pressure



Figure 5: DG-SOFC-MGT hybrid system with zeolite CO_2 adsorption and oxygenenriched air implemented in Matlab SimulinkTM



Figure 6: Efficiencies and eletrical production values of the studied cases



Figure 7: Energy balances of the studied cases

Table 1: Mode	Table 1: Model parameters I		
Membranes p	roperties [31]		
Material	Selectivity	Oxigen Permeability	
	α [ad]	$\gamma \; [\mathrm{mol} \; \mathrm{m}^{-2} \; \mathrm{s}^{-1} \; \mathrm{bar}^{-1}]$	
Matrimid	6.7	$62.0 \ge 10^{-5}$	
PPO	4.7	$37.2 \ge 10^{-4}$	
PDMS	2.1	$39.6 \ge 10^{-2}$	
Poplar wood chip	os properties [6	51]	
Description	Symbol	Value	
Total moisture	M	10 %	
Carbon content (as received)	C_{ar}	41.62~%	
Hydrogen content (as received)	H_{ar}	5.30~%	
Nitrogen content (as received)	N_{ar}	0.52~%	
Oxygen content (as received)	O_{ar}	39.81~%	
Ash content	ASH	2.75~%	
Higher heating value (dry basis)	HHV_{db}	$15.7 \mathrm{~MJ/kg}$	
Gasifier model parameters[48]			
Description	Symbol	Value	
As received biomass consumption	\dot{m}_{bio}	187 kg/h	
Nominal gasifier thermal power	$P_{th,gas}$	800 kW	
Initial calculation temperature	T_{in}	900 K	
Pressure	р	1 atm	
Equivalence ratio	\mathbf{ER}	0.335	
Gasifier and filters auxiliary consumption	$P_{DG,self}$	12.5 kW	
Cyclic operation hours	$h_{operation}$	360 h	
Cyclic maintenance hours	$h_{maintenance}$	4 h	
Zeolite 5A parameters of adsorption at 303 K [39]			
Component	$B \ [1/kPa]$	$q_m \; [\mathrm{mmol/g}]$	
CO_2	0.019500	3.91900	
H_2	0.000361	0.54464	
N_2	0.000837	2.62543	
CH_4	0.002535	2.75403	
CO	0.004350	2.75800	

Table 2: Model parameters II			
Storage and compressor model parameters			
Description	Symbol	Value	
Politropic exponent of the syngas [54]	m	1.33	
Syngas compressor efficiency [54]	η_{comp}	92~%	
Storage tank temperature	T_s	$298.15 { m K}$	
Initial syngas amount in the tanks	n_{in}	$7*10^4~{\rm mol}$	
Total tanks volume	V	650 m^3	
SOFC model parameters			
Description	Symbol	Value	
Fuel utilization factor	U_f	0.85	
Recirculation factor	r	0.2	
Operating temperature	T_{sofc}	$1073.15 { m K}$	
Anode pressure loss	Δp_a	500 Pa	
Cathode pressure loss	Δp_c	$1000 \ Pa$	
Anode pressure loss	Δp_a	500 Pa	
Current density	i	300 mA/cm^2	
Active cell area	A_{cell}	$81 \ \mathrm{cm}^2$	
Cells for each stack	$n_{cell,stack}$	75 cells	
Number of stacks	n_{stack}	145 stacks	
Cathode air excess	vent	1.15	
Pressure ratio	PR	2.5	
Steam to carbon coefficient	STC	1.4	
Electrochemical parameters taken from [18]			
MGT model parameters [18]			

MGT model parameters [18]			
Description	Symbol	Value	
Politropic exponenet of the air	m	1.33	
Turbine isoentropic efficiency	$\eta_{is,turb}$	84 %	
Air compressor isoentropic efficiency	$\eta_{is,comp}$	75 %	
Turbine mechanical efficiency	$\eta_{mec,turb}$	99~%	
Air compressor mechanical efficiency	$\eta_{mec,comp}$	98~%	
Recuperator effectiveness	η_{rec}	85 %	
Burner efficiency	$\eta_{eff,burner}$	99~%	
MGT generator efficiency	$\eta_{alt,MGT}$	95~%	
Pressure ratio	PR	2.5	

Gasifier		
Description	Symbol	Value
H_2 syngas fraction	x_{H_2}	19.03 %
H_2O syngas fraction	x_{H_2O}	7.78~%
CO syngas fraction	x_{CO}	15.10~%
CH_4 syngas fraction	x_{CH_4}	1.18~%
CO_2 syngas fraction	x_{CO_2}	13.99~%
N_2 syngas fraction	x_{N_2}	42.92~%
Air inlet flow	Q_{air}	2.94 mol/s
Syngas molar flow	\dot{n}_{syngas}	5.01 mol/s
Syngas higher heating value	$HHV_{syngas,db}$	$4.75 \ \mathrm{MJ/Nm^3}$
Specific volumetric tar production	m_{tar,Nm^3}	$23.63~{\rm g/Nm^3}$
Gasifier cold gas efficiency	η_{cold}	78.98~%
Average temperature of gasification	T	895 K
SOFC + MC	ЪТ	
Description	Symbol	Value
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	4.95 mol/s
SOFC electrical power production	P_{SOFC}	$136.70~\mathrm{kW}$
MGT electrical power production	P_{MGT}	$60.73 \mathrm{~kW}$
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$197.43~\mathrm{kW}$
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	37.46~%
Overall syste	em	
Description	Symbol	Value
Storage tank pressure range	p_{serb}	2.67-5.39 bar
Average electrical auxiliary consumption	P_{self}	$34.24 \ kW_{el}$
Average electrical total power production	P_{tot}	163.19 kW_{el}
Average total electrical efficiency	η_{tot}	25.43~%

Table 3: Case I (Gasifier + SOFC + MGT) simulation results

PPO module			
Air inlet flow	Q_{air}	2.97 mol/s	
Permeate molar flow	Q_P	1.27 mol/s	
Retentate molar flow	Q_R	1.69 mol/s	
Molar fraction of O_2 in permeate	y_{O2}	48.9~%	
Molar fraction of N_2 in permeate	y_{N2}	51.1~%	
Electric power consumption	$P_{el,PPO}$	$15.3 \mathrm{~kW}$	
Gasifier			
Description	Symbol	Value	
H_2 syngas fraction	x_{H_2}	28.49 %	
H_2O syngas fraction	x_{H_2O}	9.91~%	
CO syngas fraction	x_{CO}	26.33~%	
CH_4 syngas fraction	x_{CH_4}	1.66~%	
CO_2 syngas fraction	x_{CO_2}	17.2~%	
N_2 syngas fraction	x_{N_2}	16.41~%	
Syngas molar flow	\dot{n}_{syngas}	3.589 mol/s	
Syngas higher heating value	$HHV_{syngas,db}$	$7.55~{ m MJ/Nm^3}$	
Specific volumetric tar production	m_{tar,Nm^3}	$0.27~{\rm g/Nm^3}$	
Gasifier cold gas efficiency	η_{cold}	92.0~%	
Average temperature of gasification	T	931 K	
SOFC + MGT			
Description	Symbol	Value	
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	3.545 mol/s	
SOFC electrical power production	P_{SOFC}	$184.20~\mathrm{kW}$	
MGT electrical power production	P_{MGT}	$68.33 \mathrm{kW}$	
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$252.53~\mathrm{kW}$	
SOFC+MGT eletrical efficiency	$\eta_{SOFC+MGT}$	42.08 %	
Overall system			
Description	Symbol	Value	
Storage tank pressure range	p_{serb}	2.67-4.62 bar	
Average electrical auxiliary consumption	P_{self}	$42.01~kW_{el}$	
Average electrical total power production	P_{tot}	$210.52\ kW_{el}$	
Average total electrical efficiency	η_{tot}	32.81 %	

Table 4: Case II (PPO + Gasifier + SOFC + MGT) simulation results

Gasilier (see Cas	e I)	
ZEO		
Description	Symbol	Value
H_2 syngas fraction after adsoption	x_{H_2}	25.36~%
CO syngas fraction after adsoption	x_{CO}	17.08~%
CH_4 syngas fraction after adsoption	x_{CH_4}	1.43~%
CO_2 syngas fraction after adsoption	x_{CO_2}	0.39~%
N_2 syngas fraction after adsoption	x_{N_2}	55.73~%
Syngas molar flow after adsoption	\dot{n}_{syngas}	4.065 mol/s
Syngas higher heating value after adsoption	$HHV_{syngas,db}$	$5.9 \ \mathrm{MJ/Nm^3}$
Active zeolite mass for every regeration cycle	m_{zeo}	$23.742~\mathrm{kg}$
SOFC + MGT	Г	
Description	Symbol	Value
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	4.020 mol/s
SOFC electrical power production	P_{SOFC}	$138.50~\mathrm{kW}$
MGT electrical power production	P_{MGT}	$44.00~\mathrm{kW}$
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$182.5~\mathrm{kW}$
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	34.33~%
Overall system	n	
Description	Symbol	Value
Storage tank pressure range	p_{serb}	2.66-4.88 bar
Average electrical auxiliary consumption	P_{self}	33.76 kW_{el}
Average electrical total power production	P_{tot}	148.74 kW_{el}
Average total electrical efficiency	η_{tot}	22.87~%

 $\frac{\text{Table 5: Case III (Gasifier + ZEO + SOFC + MGT) simulation results}}{\text{Gasifier (see Case I)}}$

PPO module (see Case II)			
Gasifier (see Case II)			
ZEO			
Description	Symbol	Value	
H_2 syngas fraction after adsoption	x_{H_2}	41.53 %	
CO syngas fraction after adsoption	x_{CO}	32.54~%	
CH_4 syngas fraction after adsoption	x_{CH_4}	2.21~%	
CO_2 syngas fraction after adsoption	x_{CO_2}	0.43~%	
N_2 syngas fraction after adsoption	x_{N_2}	23.30~%	
Syngas molar flow after adsoption	\dot{n}_{syngas}	$2.726~{\rm mol/s}$	
Syngas higher heating value after adsoption	$HHV_{syngas,db}$	$10.19~\mathrm{MJ}/\mathrm{Nm}^3$	
Active zeolite mass for every regeration cycle	m_{zeo}	$20.244~\rm kg$	
SOFC + MG	T		
Description	Symbol	Value	
Syngas molar flow to SOFC-MGT unit	\dot{n}_{SOFC}	$2.696~{\rm mol/s}$	
SOFC electrical power production	P_{SOFC}	$184.70~\mathrm{kW}$	
MGT electrical power production	P_{MGT}	51.80 kW	
Total SOFC-MGT electrical power production	$P_{SOFC+MGT}$	$236.50~\mathrm{kW}$	
SOFC+MGT electrical efficiency	$\eta_{SOFC+MGT}$	38.41 %	
Overall system			
Description	Symbol	Value	
Storage tank pressure range	p_{serb}	2.66-4.15 bar	
Average electrical auxiliary consumption	P_{self}	51.80 kW_{el}	
Average electrical total power production	P_{tot}	194.53 kW_{el}	
Average total electrical efficiency	η_{tot}	30.32~%	

Table 6: Case IV (PPO + Gasifier + ZEO + SOFC + MGT) simulation results **PPO module (see Case II)**
LaTeX Source Files Click here to download LaTeX Source Files: Manuscript_ECM_review_august_2016_highlight.tex LaTeX Source Files Click here to download LaTeX Source Files: Manuscript_ECM_review_august_2016_not_highlight.tex LaTeX Source Files Click here to download LaTeX Source Files: bib.bib