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Modeling of coupling gasification and anaerobic digestion processes for maize bioenergy conversion

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Abstract

This work estimates the advantages of using maize as fuel in a power plant composed of an anaerobic digester, a gasifier and an Internal Combustion (IC) engine. The digester is fed with maize grains, while, the remaining part of the plant, the stover, is gasified. Then biogas and syngas streams are both used as fuel into the engine. The performance of this plant was evaluated coupling gasification and anaerobic digestion mathematical models. Results of the proposed solution are compared with the performance of a 100 kW biogas power plant fed with the whole crop silaged. Results show that the overall energy yield of the improved solution is 39% higher than the conventional one fed with maize silage. This method will lead to the design of small and cheap digesters as a result of the increased conversion rate. In fact, the solution proposed fully converts the high cellulose-fiber parts of the maize plant that were tough to degrade in anaerobic digesters.

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² 1. Introduction

Maize is the energy crop most widely used for biogas power production 3 [1, 2, 3]. Most of power plants runs on a combination of maize silage and 4 livestock wastes. Even if this combination assures high conversion rate, lit-5 erature suggests the possibility to run a power plant with almost 100% of 6 maize silage [4, 5], as modeled in this work for simplicity. During the years, 7 several attempts were made in the direction of increasing the overall power 8 plant efficiency, working on harvesting period as suggested by Bruni et al. 9 [6] or pre-treatments on the biomass used for reactors feed [7]. 10

The solution proposed in this work starts from the study of the maize 11 plant, defining how it can be considered composed of different parts: stalk, 12 cob, leaves, husk and grains. These components behave differently in the 13 digester [8, 9]. Among the different parts of maize plant, grains are the 14 most degradable and even the most productive in terms of specific biogas 15 production. In fact, grains are characterized by high starch and soluble 16 sugars content, as well as low lignin [9, 10, 11]. On the other hand, the corn 17 stover, composed of cobs, leaves, husk and stalks, has a lower productivity 18 in terms of biogas as result of its high percentage of cellulose and lignin [9]. 19

This suggests that a digester fed mainly with grains could theoretically be smaller than a digester fed with the whole plant, but also with corn silage. Furthermore, while the operation of silaging increases the productivity of each specific part of the maize plant, the same procedure reduces the amount
of organic matter of at least 10%. This phenomenon is due to an unavoidable
aerobic fermentations occurring in the silaging process [12].

As described above, the grains are the most important part of the maize plant in terms of anaerobic digestion process, while the stover is characterized by low conversion rates that drastically affect the bio-chemical processes.

This work describes two different scenarios: 1) the conventional power plant composed of an anaerobic digester fed 100% with corn silage and 2) a hybrid power plant composed of an anaerobic digester fed with grains and a downdraft gasifier fed with the stover.

Literature review revealed several studies about corn residues gasifica-33 tion. Zijp et al. in the 1980 published a technical report with the Twente 34 University where the use of corn stover in gasifier was discussed. It was found 35 that the particulate content in the gas was one of the major issues related to 36 this application. [13]. Zijp's results were cited in one of the most important 37 manuals for fixed bed gasifier design: Woodgas as engine fuel [14]. A more 38 recent work on gasification by Mavukwana et al. [15] models the stover gasi-39 fication process in Aspen plus finding optimal equivalence ratio and steam 40 to biomass values for this feedstock. Groeneveld and Van-Swaaij in the 1979 41 and Allesina et al. in 2015 discussed the possibility to use corn cobs gasifica-42 tion in micro power plants. Both the works are focused on energy shortage 43 problems in African villages [16, 17]. A fixed bed gasifier of 350 kW_{th} was 44 used by Biagini et al. in 2014, it was fed with corn cobs, reporting a gasi-45 fication efficiency of the system of 67% and a syngas heating value of 5.7 46 MJ/Nm³ [18]. Literature review shows how the stover can also be processed 47

⁴⁸ in fluidized bed gasifiers [19, 20].

The basic idea of this work is depicted in Figure 1. Aim of this study is 49 to demonstrate the advantages related to the separation of the grains from 50 the corn stover and the exploitation of the grains in the digester while the 51 stover is converted into syngas by a fixed bed gasifier. Literature does not 52 include many works on the possible effect of combined anaerobic digestion 53 and gasification. Li et al. [21] modeled a coupled system aimed at biomethane 54 production, while Chen at al. couples a fixed bed gasifier working with 55 corncob and cotton straws and a typical biogas plant for fueling household 56 furnaces in rural scenarios [22]. Other studies focus on the use of anaerobic 57 digestion and thermo-chemical conversion in cascade. Two possible work 58 groups can be found in literature. The first group attempts a further energy 59 conversion of the digestate disposed by the biogas power plant [23, 24, 25], the 60 other uses anaerobic digestion for the conversion of the unwanted products 61 of the gasification and pyrolysis processes (wastewater and tar) [26, 27, 28]. 62

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The system is modeled as a composition of two sub-models:

• The anaerobic digestion was simulated with the Anaerobic Plant Emulation (APE) model reported in [29].

The stover gasification process was simulated with a black-box equilibrium model in order to estimate the steady state behavior of the gasifier.
Different gasification conditions were tested with the final purpose of obtaining a syngas with higher heating value over 4 MJ/Nm³ and a tar content lower than 2 g/Nm³. This conditions are within the typical ranges reported by Milne for downdraft gasifiers [30]. However, the value obtained is too high for direct feeding of the gas to an engine.

Therefore a filtering process is mandatory to lower the tar content value
under 100 mg/Nm³ [31].

Due to the synergy of gasification and anaerobic digestion models, precise 75 data about the chemical and physical properties of the feedstocks used as 76 inputs are required. Literature reports several studies about composition 77 and behavior in anaerobic digesters of corn and corn silage but works about 78 the behavior of the separated parts of the maize plant in bio-digestion or 79 gasification are few. For example, Hutnan et al. [32] discusses the differences 80 between maize grains and maize silage for biogas production; Getachew et 81 al. analyzes differences between grains and silage, but no data are reported 82 about the corn stover in both these works. A complete characterization 83 of the stover can be found in Evans et al. about power production from 84 substitute fuels [20]. In the next section these sources, together with other 85 literature data, are used in order to define the characteristics of the grains 86 and the stover used in the improved solution, as well as the characteristic 87 of the hypothetical 'equivalent maize silage' that can be obtained from the 88 same maize. 89

Therefore, the total chemical energy content in the syngas-biogas stream was compared to the biogas chemical energy content in case of 100% silage digestion. Results shown the advantages related to this approach. A power production boost of about 26% was obtained not considering the advantage of avoiding silaging matter losses. Considering also this contribution, the advantage of the operation raises to 39%. On the other hand, the gasifier is characterized by a higher conversion rate but some issues related to corn stover processing have been outlined during preliminary experimental gasifi⁹⁸ cation tests reported in the results.

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¹⁰⁰ 2. Material and Methods

101 2.1. Case study

The anaerobic digestion of a 100 kW biogas plant fed only with maize silage was modeled in order to evaluate the biogas production rate and the biogas higher heating value HHV. Then, the corn stover was removed from the model 'recipe' and the system was simulated under this new condition using only grains as fuel. At the same time, the stover removed from the biogas model was used in the gasification one. These two cases are depicted in Figure 1 are summarized as:

(A) Ensilage of 100% of corn and its total exploitation in the biogas power plant.

(B) Separation of the stover from grains, gasification of the stover and exploitation of the grains in the biogas plant.

This work does not focus on the effect of different gases on the performance of the CHP engine, for this reason the two cases were compared on the basis of the chemical energy content in the gas streams.

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117 2.2. Definition of maize characteristics from literature

The methodology applied in this work is the following: the grains and the stover were characterized on the data reported by Evans [20] and by Getachew [33]. These data are summarized in Table 1 and Table 4. Once the data about the maize components are known, it is possible to define an "equivalent maize silage" obtained as a composition of the values previously cited. Its properties are defined here as the weighted mean of their respective components values. This approach is effective if the following assumptions are verified:

 The mass balance of the corn plant is equal to the sum of the components masses.

2. The final characteristics of "equivalent maize silage" are in line with
those found in actual practice (i.e. [34]).

Basically, the method proposed allowed to back calculate the characteristics of the silage (usually collected in milky-waxy stage) from the data reported for grains and stover all referring to the complete maturation of the feedstock. In results section the effectiveness this methodology is discussed

134 2.3. Gasification process modeling

Corn cobs gasification is not a newness. Literature reports ongoing researches focused on using maize cobs or stovers as fuel in fixed bed gasifier [16, 13] as well as in fluidized bed gasifiers [19, 20]. In this work the gasification process was simulated using a black-box model based on Barman's work [35]. The model generally works for downdraft gasifiers; it is based on the following generic gasification equation:

$$CH_x O_y N_z + w H_2 O + m (O_2 + 3.76N_2) \rightarrow$$

$$n_{H_2} H_2 + n_{CO} CO + n_{CO_2} CO_2 + n_{H_2O} H_2 O \qquad (1)$$

$$+ n_{CH_4} CH_4 + (z/2 + 3.76m) N_2 + n_{tar} CH_p O_q$$

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 [36]; w $[mol/mol_{bio}]$ in the specific molar amount of the biomass moisture calculated by Equation 2; $m [mol/mol_{bio}]$ is the specific molar amount of oxygen calculated by Equation 3; $n_{H_2}, n_{CO}, n_{CO_2}, n_{H_2O}, n_{CH_4}, n_{tar} [mol/mol_{bio}]$ are the specific molar amount of $H_2, CO, CO_2, H_2O, CH_4$ and tar which constitute the syngas.

$$w = \frac{MW_{bio,daf} * M}{MW_{H_2O} \left(1 - M/100 - ASH/100\right)}$$
(2)

$$m = ER * (1 + x/4 - y/2) \tag{3}$$

where M [%] is the total moisture; ER [ad] is the equivalence ratio as 148 defined by Reed and Das [37] and $MW_{bio,daf}$ [g/mol] is the molecular weight 149 of biomass in "daf" conditions. Equation 1 can be multiplied by the molar 150 biomass flow in "daf" conditions $\dot{n}_{bio,daf} [mol_{bio}/s]$ in order to assess the molar 151 flow of each component of the syngas as well as the syngas composition in 152 wet and dry conditions. The molar flow of tar is given by Equation 4, the 153 tar production versus the "daf" biomass input x_{tar} [% wt. "daf" biomass] is 154 calculated by Equation 5. Furthermore, Equation 6 can be used to evaluate 155 the volumetric tar amount $g_{tar,vol}$ [g/Nm³] in the syngas. 156

$$\dot{n}_{tar} = n_{tar} * \dot{n}_{bio,daf} \tag{4}$$

$$x_{tar} = \frac{n_{tar} * M W_{tar}}{M W_{bio}} \tag{5}$$

$$g_{tar,vol} = \frac{n_{tar} * MW_{tar}}{\frac{\dot{n}_{tot,dry}}{\dot{n}_{bio,daf}} * 0.022414}$$
(6)

The constant 0.022414 is the volume in m^3 of 1 mol of ideal gas at the normal conditions of 101325 Pa and 273 K [38]. Moreover, assuming the syngas components as ideal gases it is possible to calculate the normal volumetric flow of wet and dry syngas. Equations 7 and 8 allow us to estimate the "cold gas" efficiency of the gasifier and the *HHV* of the clean and dry syngas.

$$\eta_{g,cold} = \frac{\dot{V}_g H H V_{syngas,clean}}{\dot{m}_f H H V_{bio,ar}} \tag{7}$$

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$$HHV_{syngas,clean} = x_{H_2}HHV_{H_2} + x_{CO}HHV_{CO} + x_{CH_4}HHV_{CH_4}$$
(8)

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where x_{H_2} , x_{CO} , x_{CH_4} [% vol] are the volumetric fraction of H_2 , CO, CH_4 in the dry syngas and HHV_{H_2} , HHV_{CO} , HHV_{CH4} [MJ/Nm³] are the higher heating values of H_2 , CO and CH_4 .

However, the molar specific amount of the syngas components have to be estimated. An algorithm similar to the one suggested in [39] is adopted here. 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 $CH_4 + H_2O \leftrightarrow CO + 3H_2$

Equations 9 and 10 are reported in [40] and are used here to calculate K_1 and K_2 while Equation 11 is used to evaluate K_3 and it is taken from [41]:

$$K_1 = e^{\frac{4276}{T} - 3.961} \tag{9}$$

$$ln(K_2) = \frac{7082.842}{T} - 6.567 * ln(T) + \frac{7.467 * 10^{-3} * T}{2} - \frac{2.167 * 10^{-6} * T^2}{6} + \frac{0.702}{2 * T^2} + 32.541$$
(10)

$$K_3 = 1.198 * 10^{13} * e^{\frac{-26830}{T}}$$
(11)

The System 12 is composed of three chemical balances calculated from Equation 1 (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.

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

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

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

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

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

$$K_3 = \frac{n_{CO} * n_{H_2}^3}{\left(\frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}\right)^2 n_{H_2O} n_{CH_4}}$$
(12)

Once the molar specific amount of the syngas species are evaluated, it is possible to solve the thermodynamic energy balance of the system reported in Equation 13. In order to find the equilibrium temperature T_{new} , the system is considered adiabatic and the the Newton-Raphson method is applied to the equations.

As reported in [39], if $abs(T - T_{new}) < 0.1$ K then the calculated equilibrium temperature and molar specific gases amounts are the final results; instead, a new iteration is done in order to satisfy the previous condition.

$$\sum_{j=react} n_j * HF_j^0 = \sum_{i=prod} n_i * \left(HF_i^0 + \Delta H_{T,i} \right)$$
(13)

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 ¹⁹¹ [moles] and HF_i^0 [kJ/kmol] are the specific moles amount and the standard ¹⁹² heat of formation of the i-th product (H_2 , CO, CO_2 , H_2O , CH_4 and N_2) and ¹⁹³ $\Delta H_{T,i}$ is the enthalpy difference between any given state and the standard ¹⁹⁴ state for the i-th product. $\Delta H_{T,i}$ can be calculated starting from the specific ¹⁹⁵ heat of the product:

$$\Delta H_{T,i} = \int_{298.15}^{T} C_p(T) \, dT = \left| aT + b\frac{T^2}{2} + c\frac{T^3}{3} + d\frac{T^4}{4} \right|_{298.15}^{T} \tag{14}$$

where the coefficient a,b,c and d are defined for each gas in [39]. The model was implemented in Python. In this way once the biomass equivalent molecule is defined, the model works with the only definition of a ER and a temperature. The temperature input is used only as a starting point for the iterating system, after few cycles the temperature converges to the ones that satisfy both the chemical and thermal sub-systems.

202 2.4. Biogas modeling

The mathematical sub-model designed to simulate the anaerobic digestion was developed in a previous works [29]. This model is useful to design wet (or semi-wet) anaerobic digestion plants in steady state conditions. The input data are the characteristics of the feedstock and few basic parameters such as the CHP efficiency and the process temperature.

The APE model [29] consists of several different interlaced sections which can be grouped into two fundamental modules: the biological module used in this work and the heat module. The biological module goals are:

1. Estimating the degradation of selected biomasses.

- 2. Calculating the production of bio-methane and, consequently, the power
 output of the plant.
- 3. Selecting the optimal Hydraulic Retention Time (HRT) on the basis of
 the chosen degradation efficiency.
- 4. Designing the digester tanks as function of the selected layout and
 water content of the substrate.

Assuming hydrolysis as a limiting step of the anaerobic digestion reactions chain, the biomass degradation can be described as a first order kinetic model [42]. The mass balance of the substrate for a generic reactor with constant volume and flow rates can be written as follows [43]:

$$\frac{d\hat{S}_e}{dt} = \frac{Q_i}{V}\hat{S}_t - \frac{Q_e}{V}\hat{S}_e - k\hat{S}_e \tag{15}$$

where \hat{S}_e is the biomass concentration in the effluent flow rate $[kg_{ts}/m^3]$, 222 \hat{S}_t is the biomass concentration in the in-fluent flow rate $[kg_{ts}/m^3]$, Q_i is 223 the influent flow rate $[m^3/day]$, Q_e the effluent flow rate $[m^3/day]$, V is the 224 digestor active volume $[m^3]$ and k is the hydrolysis kinetic constant $[day^{-1}]$. 225 Starting from the concentration of the effluent flow, it is possible to cal-226 culate the efficiency of the degradation process and, from this, to trace back 227 the HRT. The amount of reacting mass in the digester is also calculated con-228 sidering the partial degradation of the substrate and the degree of dilution. 229 The latter is evualuated in terms of water flow rate or recirculation flow rate 230 as a function of the desired water content of the substrate and the desired 231 organic loading rate (ORL). 232

To increase the accuracy of the results, each biomass used as input is broken into its constituents: sugars, proteins, fats, cellulose, hemicellulose, lignin, ash, and non-degradables parts according to the Van Soest method
(also called NDF method) [9]. A specific hydrolysis kinetic constant is assigned to each constituent type, in order to take into account the different degradation rates which typically occurs in different substrates. Specific aspects of the modeling approach were inspired by several other works
[43, 9, 42, 44].

The methane productivity estimation of the matrices was carried out by 241 giving to each class of substance (carbohydrates, proteins, lipids) a particu-242 lar methane yield. This method is in accordance with the specific technical 243 standard [45] which recalls the initial works of Buswell [46] and Boyle [47]. 244 The share of non-degradable organic matter do not produces biogas due to 245 its tough-fibers structure which is abiotic for microorganisms (i.e. lignin and 246 the part of cellulose closely linked to it). In order to take into account this 247 issue, thus avoiding overestimation of biogas production, the model calcu-248 lates the proportion of non-degradable NDF (iNDF) starting from the known 240 iNDF/ADL ratio [48]. Every biomass has a specific iNDF/ADL ratio, that 250 can be deduced from the other parameters once the methane yield is know. 251 252

253 2.5. Evaluation collecting and processing costs

This work gives an overall energy balance evaluation of the proposed solution. Thanks to this evaluation is possible to demonstrate the advantages related to the improved utilization of maize. On the other hand it is important to assure that the proposed implementations an changes in the harvesting process do not affect the economy of system. The first step that needs to be taken is the evaluation of the cost differences between the two solutions ²⁶⁰ applied to a medium-size farm with more than 10 ha cultivated, fair degree ²⁶¹ of mechanization and corn productivity about 60 t/ha (wet) [49, 50, 51]. The ²⁶² next paragraph reports the results obtained from literature review about the ²⁶³ cost of the two solutions.

264 3. Results

²⁶⁵ 3.1. Evaluation collecting and processing costs

Literature suggests small cost differences between collecting or leaving the stover on the fields in these conditions [49, 50, 51, 52, 53] as explained below:

- Case A: the average cost for the cultivation of maize silage is about 30
 \$/t with 65% moisture [49]. If the humidity of the silage is theoretically
 reduced to the moisture content of the stover in field (12.5%), the silage
 cost rises to 75 \$/t. Silage harvesting does not leave enough organic
 substance on the field, therefore it is necessary to integrate nutrients
 in the soil for a cost of 6.50 \$/t [50].
- Case B: grains harvest is carried out by a combine harvester which separates grains from the plant and leaves the stover in the field. The average cost of the cultivation of the grain starts from 50 \$/t [51].
 The stover harvesting into bales is similar to the process done for the straw. It costs an average of 30 \$/t considering the cost of nutrient replacement. The mechanical operations required are: flail shredding and raking followed by baling without crop processor [50].

On the other hand, leaving the stover on the field is not sufficient for assuring that its organic substance is properly transferred to the soil. In fact, the process required extra mechanical operations such as straw chopping and soil plowing which contribute to costs raising.

The focus of this paper was kept on chemical and physical changes of 286 the matrices during their fate from the initial conditions. From this point of 287 view, corn silage is subject to significant energy losses during the lactic fer-288 mentation (and often exceeding what reported in this work); these losses do 289 not occur in the same entity during the drying of the grains. For this reason 290 some auxiliary sources of energy consumption or losses were not taken into 291 account. For example, while in Case A was neglected the energy consump-292 tion for the silaging process (stacking the trenches, pressing, covering), in 293 Case B was neglected the energy consumption for drying corn grains. These 294 losses will affect similarly both the solutions proposed. 295

296 3.2. Biogas modeling

The "equivalent maize silage" method is effective due to the good simi-297 larity with literature (i.e. [8, 54]) as reported in Table 4. With this approach 298 the two cases have all the inputs required for their modeling as reported in 299 Tables 4, 4 and 5. The higher differences are the ones related to the fiber 300 composition. In particular, the "equivalent maize silage" shows higher lignin 301 content compared to literature and real data [54, 9]. The main cause of this 302 deviation is that Evans's data presents high fiber content values in the first 303 place. 304

The results of the biogas simulations are reported in Table 5. The model gives the specific power output in terms of m_{biogas}^3/kg_{ts} . This value is used to calculate the equivalent silage specific primary energy $e_A [kWh_{PE}/kg_{ts}]$ that is the primary energy produced (biogas chemical energy) by one kg of dry maize silage. The model also calculates the methane productivity of the feedstocks which it is compared with literature data as reported in Table 5. The average error of 12% is considered acceptable for the model validation.

312 3.3. Gasification modeling

Table 4 resumes the major results obtained considering a wet flow of syn-313 gas. Due to the composition of the feedstock, a working point characterized 314 by low tar content was found with an ER slightly higher than the value 315 suggested for wood chips gasification [31, 37]. The stover resulted a suitable 316 biomass for gasification with a cold gas efficiency of 71.2 %. The tar content 317 resulted 1.32 g/Nm³, this value is slightly lower than literature review values, 318 for example Milne reports downdraft gasifiers that produce up to 5 g/Nm^3 319 of tars. It is important to consider that the gasification power plant imple-320 mented in this solution consists in the reactor only. The syngas produced in 321 the reactor can be directly sent into the biogas gasometer. In so doing there 322 is no need for filtering process. There are few studies about the behavior of 323 tars into the biogas gasometer, Torri and Fabbri, [28] suggested how some 324 oils and tars can be upgraded to hydrogen through anaerobic digestion, while 325 the work of Hübner [27] already integrates a biogas reactor for upgrading the 326 liquid phase of a pyrolysis power plant. Anyway, within the gasometer, the 327 syngas can be effectively cooled down and slowed. Under these conditions 328 tars are able to condense flowing into the reacting biomass in the digester. 329

The gasifier conversion rate is evaluated in terms in dry stover specific primary energy $e_{stover} [kWh_{PE}/kg_{ts}]$. In case of electrical power output eval³³² uation, it is important to take into account the generator and Internal Com-³³³ bustion (IC) engine efficiencies. This values lead to a new conversione rate ³³⁴ value $e_{plant,model}$ [$kWh_{el}/kg_{ts,stover}$].

335 3.4. Chemical power output

Table 6 contains the comparison between the conversion rates of the two 336 solutions. Table 6 points out the overboost of 39% that can be reached 337 in case B. This value derives from the higher efficiency of the gasification 338 process and from its capability to exploit ligno-cellulosic matter. While the 339 productivity of the maize silage is known ($e_A = 3.079 \ kWh_{PE}/kg_{ts}$), the 340 value for the combined effect of gasification of the stover (dried to M = 5%341 of moisture) and the anaerobic digestion of the grains is evaluated starting 342 from mass share value of grains $(f_{ts,grains})$ and its complementary value for 343 the stover $(1 - f_{ts,grains})$: 344

$$e_B = f_{ts,grains} * e_{grains} + (1 - f_{ts,grains}) * e_{stover} =$$

= 0.46 * 4.051 + (1 - 0.46) * 3.758 = 3.89 (16)

This value is 1.264 times higher than the silage one. Furthermore the 345 silage losses (about 10% [12]) increase the value to 1.39. This means that 346 a 100 kW, 100% silage power plant can be boosted to 139 kW through the 347 adoption of a gasifier. In case A the primary energy yielded for hectare 348 of soil is calculated considering a productivity of 26.41 tons per hectare of 349 dried maize plant (type FAO 500: milky-waxy) as suggested by [9, 34]. As 350 previously discussed, the value obtained needs to be resumed due to silaging 351 matter losses. The calculation of the primary energy in Case B considers a 352

productivity of 83.98 tons per hectare of as-received maize plant (type FAO 700: full ripeness stage) [5]. The moisture of Case B is 68.14 % as suggested by Blandino et al. [54]. The final remark need to be addressed to the cost of the two proposed plants. The costs of the refer 100 kW power plant can be found in literature, in particular in Italy, in 2013 the cost for maize power plants is almost $10 \in /W$ [55] considering:

• Approximatively 900 m³ digester tank with gasometer

- 100 kW combine heat and power system
- Auxiliary systems such as mixers, blowers, sensors and control systems
- Automatic feeding
- Storage tank

More difficult is the estimation of the cost of the improved solution. In fact, on one hand, the biogas part of the combined power plant is going to be cheaper due to smaller tanks and digesters, on the other hand the gasifier reactor and its auxiliary equipments are going to increase the price as well as the bigger engine required for this solution. As rough approximation it can be possible to assume the cost linear, fixing it at $10 \in /W$.

The cost of the final plant will be proportional (roughly 1.4 millions of euros) because on one hand the biogas part is reduced in terms of volumes of tanks, digesters and auxiliary equipments but, on the other hand the new system is provided whit a gasification reactor. No filtering system is required. Figure 2 resumes the energy conversion effectiveness in the two cases analyzed.

376 4. Conclusions

This work demonstrates the advantages related to the combination of 377 anaerobic digestion and gasification technologies. The costs of new solution 378 proposed are similar to the conventional solution. The model predicts the 379 behavior of the system with an error of 12% compensated by silaging matter 380 losses of the conventional solution. Therefore, the minimum performance 381 increase is 26% for the improved solution. Such a power increase justify 382 the higher complexity of the improved solution even more considering that 383 the dimension of the new digester would be almost half of the conventional 384 one. Future work will focus on collecting data for a model validation. Lastly 385 this work suggests the use of a singular engine instead of two. This solution 386 requires an engine with higher power and presumably higher efficiency. All 387 these features of the improved solution compete to increasing the overall 388 efficiency, assuring the effectiveness of this method to boost the performance 389 of existing facilities as well as new biogas power plants. Finally it is important 390 to outline that a secondary result derived from this study. The models used 391 here required coherent data about the maize feedstock under two different 392 ripening conditions: full ripeness and silage. The solution proposed for this 393 problema is the definition of a methodology able to give the properties of 394 the silage starting from the chemical characteristics of the plant parts. The 395 method chose was found to be effective and the results are in line with typical 396 literature data about maize silage. 397

398 Nomenclature

399	$\Delta H_{T,i}$	enthalpy difference[kJ/kmol	4 20	HF^0	standard heat of formation
400	\dot{m}	mass flow [kg/s]	421		[kJ/kmol]
401	\dot{n}	molar flow [mol/s]	422	HHV	higher heating value
402	\dot{V}	volumetric flow $[Nm^3/s]$	423		[MJ/Nm ³ or MJ/kg]
403	\hat{Se}	biomass concentration in t	424 he	HRT	hydraulic retention time
404		effluent flow rate $[kg_{ts}/m^3]$	425 426	INDF	tergent fiber
405	\hat{St}	biomass concentration in the function $f_{\rm eff}$	he 427	K	equilibrium constant [ad]
406	ADL	acid detergent fiber/lignin	428	k	hydrolysis kinetic constant
407	ASH	ash content of the biomass $[$	429 %]		$[day^{-1}]$
409	C	carbon	430	М	total moisture content of the
410	C_p	specific heat [J/(mol K)]	431	m	specific molar amount of ovv-
411	CHP	combined heat power	432	110	gen $[mol/mol_{bio}]$
412	E	energy [kWh]	434	MW	molecular weight $[g/mol]$
413	e	specific biomass energy pr	• <u>635</u>	N	nitrogen
414		ductivity $[kWh_{PE}/kg_{ts}]$	436	n	specific molar amount of gases
415	ER	equivalence ratio [ad]	437		and tar $[mol/mol_{bio}]$
416	f	mass fraction $[\%]$	438	NDF	neutral detergent fiber
417	g	volumetric tar amou	439 NU	0	oxygen
418		$[g/Nm^3]$	440	ORL	organic loading rate
419	Η	hydrogen	441	Q_e	effluent flow rate $[m^3/day]$

442	Q_i	influent flow rate $[m^3/day]_{454}$	p	hydrogen coefficient of tar
443	Т	temperature [K] 455	prod	product
444	V	digestor active volume $[\mathrm{m}^3]_{\ _{456}}$	q	oxygen coefficient of tar
445	w	${\rm specific \ molar \ amount \ } \mathfrak{g} \mathfrak{f}$	react	reactant
446		biomass moisture $[mol/mol_{458}]_{_{458}}$	ts	total solid
447	x_{tar}	tar production versus daf $_{459}$	vs	volatile solid
448		biomass input [% wt.]		
	C - 1 -	460	x	hydrogn coefficient of the
449	Subs	cripts 461		biomass
450	ar	as received	u	hydrogen coefficient of the
451	bio	biomass 463	0	biomass
452	daf	dry ash free 464	z	nitrogen coefficient of the
453	q	gas 465		biomass

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⁶⁸⁰ Figure and tables captions



Figure 1: System layout

Table 1: Corn plant total solid composition and soil productivity (full ripeness) [52]

	Grains	Straw	Leaves	Cobs	Husk
Total Solids distribution f_{ts}	46%	27%	12%	8%	7%
Dry matter/ha $[ton_{ts}/ha]$	6.13	3.58	1.58	1.08	0.93

Parameter	Symbol	Value
Stover parameters for gasification process [20, 52]		
Carbon amount [% wt.]	C_{ar}	44.18
Nitrogen amount [% wt.]	N_{ar}	0.53
Hydrogen amount [% wt.]	H_{ar}	5.52
Sulfur amount [% wt.]	S_{ar}	$\simeq 0.1$
Oxygen amount [% wt.]	O_{ar}	37.69
Moisture [% wt.]	M	5 (dried on field)
Ash amount [% wt.]	ASH	6.98
Higher heating value [MJ/kg]	HHV_{ar}	$19 \mathrm{~MJ/kg}$
Stover parameters for anaerobic digestion process [20, 52]		
Total solids [% wt.]	ts	21.8 (full ripeness)
Ash amount [% wt.]	ASH	6.98
Volatile solids [% of ts]	VS	91.8
Crude protein [% of ts]	XP	4.8
Crude fat $[\% \text{ of } ts]$	XL	4.27
Non fiber carbohydrate [% of ts]	NFC	3.60
Neutral detergent fiber $[\% \text{ of ts}]$	NDF	79.1
Acid detergent fiber [% of ts]	ADL	16.9
Non degradable fiber ratio	iNDF/ADL	2
Grains [33, 52]		
Total solids [% wt.]	ts	66 (full ripeness)
Ash amount [% wt.]	ASH	1.2
Volatile solids [% of ts]	VS	98.8
Crude protein $[\% \text{ of ts}]$	XP	8.2
Crude fat $[\% \text{ of ts}]$	XL	3.4
Non fiber carbohydrate [% of ts]	NFC	76.6
Neutral detergent fiber $[\% \text{ of ts}]$	NDF	10.7
Acid detergent fiber [% of ts]	ADL	0.5
Non degradable fiber ratio	iNDF/ADL	2

Table 2: Corn grains and stover properties



Figure 2: Energy conversion balances in the two cases

1		0 1 1		
Equivalent maize silage	Symbol	Value	Comparison with [54]	
Total solids [% wt.]	ts	31.86	31.89	
Ash amount [% wt.]	ASH	4.46	5.13	
Volatile solids [% of ts]	VS	95.09	94.87	
Crude protein [% of ts]	XP	6.42	7.44	
Crude fat $[\% \text{ of ts}]$	XL	3.86	3.86	
Non fiber carbohydrate [% of ts]	NFC	38.31	43.3	
Neutral detergent fiber $[\%~{\rm of}~{\rm ts}]$	NDF	46.95	40.24	
Acid detergent fiber [% of ts]	ADL	8.96	4.97	
Non degradable fiber ratio	iNDF/ADL	2	/	

Table 3: Equivalent maize silage properties

Variable	Symbol	Value
Equivalence ratio	ER	0.365
Biomass moisture	M	5%
H_2 molar fraction	H_2	17.29~%
H_2O molar fraction	H_2O	5.7 %
CO molar fraction	CO	17.19~%
CO_2 molar fraction	CO_2	11.74~%
CH_4 molar fraction	CH_4	0.78~%
N_2 molar fraction	N_2	47.31 %
Specific volumetric tar amount	$m_{tar,vol}$	1.32 g/Nm^3
Volumetric syngas flow	\dot{V}_{syngas}	$47.0 \ \mathrm{Nm^3/h}$
Wet syngas higher heating value	$HHV_{syngas,w}$	4.68 MJ/Nm^3
Dry syngas higher heating value	$HHV_{syngas,d}$	$4.93 \ \mathrm{MJ/Nm^3}$
Cold gas efficiency	η_{cold}	71.2~%
Dry stover specific primary energy	e_{stover}	$3.758 \ kWh_{PE}/kg_{ts}$
IC engine efficiency	η_{engine}	0.3
Electrical generator efficiency	η_{gen}	0.95
Power plant conversion rate	$e_{plant,model}$	$1.07 \ kWh_{el}/kg_{ts,stover}$

Table 4: Results of the stover gasification

Table 5: Results and validation of the anaerobic digestion model

Table	Table 5. Results and valuation of the anaerobic digestion model					
	Literature methane	Model methane	Absolut	Model biogas	Model specific	
	production $[32, 52]$	production	error	production	primary energy	
	$[m^3/kg_{vs}]$	$[m^3/kg_{vs}]$	[%]	$[m^3/kg_{ts}]$	$\left[kWh_{PE}/kg_{ts}\right]$	
Grains	0.360	0.410	14	0.771	4.051	
Corn Stover	0.274	0.234	14	0.403	2.148	
Equivalent Silage	0.350	0.326	7	0.582	3.079	
Mean value	-	-	12	-	-	

Table 6: Comparison between case (A) and case (B)

Variable	Symbol	Value
Case (A) specific primary energy	e_A	$3.079 \ kWh_{PE}/kg_{ts}$
Case (B) specific primary energy	e_B	$3.890 \ kWh_{PE}/kg_{ts}$
Increase	Δe	26%
Increase considering silaging matter losses	Δe_{sml}	39 %
Case (A) specific primary energy per hectare	E_A	73.173 MWh_{PE}/ha
Case (B) specific primary energy per hectare	E_B	$104.08 \ MWh_{PE}/ha$