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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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*Keywords:*

Stover, Corn grains, gasification, anaerobic digestion, modeling

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## 2 **1. Introduction**

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

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

20 This suggests that a digester fed mainly with grains could theoretically be  
21 smaller than a digester fed with the whole plant, but also with corn silage.  
22 Furthermore, while the operation of silaging increases the productivity of

23 each specific part of the maize plant, the same procedure reduces the amount  
24 of organic matter of at least 10%. This phenomenon is due to an unavoidable  
25 aerobic fermentations occurring in the silaging process [12].

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

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

33 Literature review revealed several studies about corn residues gasifica-  
34 tion. Zijp et al. in the 1980 published a technical report with the Twente  
35 University where the use of corn stover in gasifier was discussed. It was found  
36 that the particulate content in the gas was one of the major issues related to  
37 this application. [13]. Zijp's results were cited in one of the most important  
38 manuals for fixed bed gasifier design: Woodgas as engine fuel [14]. A more  
39 recent work on gasification by Mavukwana et al. [15] models the stover gasi-  
40 fication process in Aspen plus finding optimal equivalence ratio and steam  
41 to biomass values for this feedstock. Groeneveld and Van-Swaaij in the 1979  
42 and Allesina et al. in 2015 discussed the possibility to use corn cobs gasifica-  
43 tion in micro power plants. Both the works are focused on energy shortage  
44 problems in African villages [16, 17]. A fixed bed gasifier of 350 kW<sub>th</sub> was  
45 used by Biagini et al. in 2014, it was fed with corn cobs, reporting a gasi-  
46 fication efficiency of the system of 67% and a syngas heating value of 5.7  
47 MJ/Nm<sup>3</sup> [18]. Literature review shows how the stover can also be processed

48 in fluidized bed gasifiers [19, 20].

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

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

- 64 • The anaerobic digestion was simulated with the Anaerobic Plant Em-  
65 ulation (APE) model reported in [29].
  
- 66 • The stover gasification process was simulated with a black-box equilib-  
67 rium model in order to estimate the steady state behavior of the gasifier.  
68 Different gasification conditions were tested with the final purpose of  
69 obtaining a syngas with higher heating value over  $4 \text{ MJ/Nm}^3$  and a tar  
70 content lower than  $2 \text{ g/Nm}^3$ . This conditions are within the typical  
71 ranges reported by Milne for downdraft gasifiers [30]. However, the  
72 value obtained is too high for direct feeding of the gas to an engine.

73 Therefore a filtering process is mandatory to lower the tar content value  
74 under 100 mg/Nm<sup>3</sup> [31].

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

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

98 cation tests reported in the results.

99

## 100 **2. Material and Methods**

### 101 *2.1. Case study*

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

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

111 (B) Separation of the stover from grains, gasification of the stover and ex-  
112 ploitation of the grains in the biogas plant.

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

116

### 117 *2.2. Definition of maize characteristics from literature*

118 The methodology applied in this work is the following: the grains and  
119 the stover were characterized on the data reported by Evans [20] and by

120 Getachew [33]. These data are summarized in Table 1 and Table 4. Once  
121 the data about the maize components are known, it is possible to define an  
122 "equivalent maize silage" obtained as a composition of the values previously  
123 cited. Its properties are defined here as the weighted mean of their respective  
124 components values. This approach is effective if the following assumptions  
125 are verified:

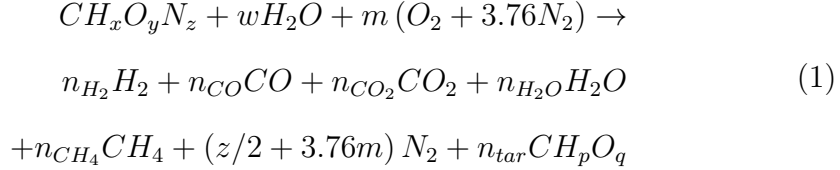
- 126 1. The mass balance of the corn plant is equal to the sum of the compo-  
127 nents masses.
- 128 2. The final characteristics of "equivalent maize silage" are in line with  
129 those found in actual practice (i.e. [34]).

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

### 134 *2.3. Gasification process modeling*

135 Corn cobs gasification is not a newness. Literature reports ongoing re-  
136 searches focused on using maize cobs or stovers as fuel in fixed bed gasifier  
137 [16, 13] as well as in fluidized bed gasifiers [19, 20]. In this work the gasi-  
138 fication process was simulated using a black-box model based on Barman's  
139 work [35]. The model generally works for downdraft gasifiers; it is based on  
140 the following generic gasification equation:





141 where  $CH_xO_yN_z$  is the equivalent chemical formula of "dry and ash  
142 free" (daf) biomass;  $CH_pO_q$  is the equivalent chemical formula of tar [36];  $w$   
143 [ $mol/mol_{bio}$ ] is the specific molar amount of the biomass moisture calculated  
144 by Equation 2;  $m$  [ $mol/mol_{bio}$ ] is the specific molar amount of oxygen cal-  
145 culated by Equation 3;  $n_{H_2}, n_{CO}, n_{CO_2}, n_{H_2O}, n_{CH_4}, n_{tar}$  [ $mol/mol_{bio}$ ] are the  
146 specific molar amount of  $H_2, CO, CO_2, H_2O, CH_4$  and  $tar$  which constitute  
147 the syngas.

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

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

148 where  $M$  [%] is the total moisture;  $ER$  [ad] is the equivalence ratio as  
149 defined by Reed and Das [37] and  $MW_{bio,daf}$  [g/mol] is the molecular weight  
150 of biomass in "daf" conditions. Equation 1 can be multiplied by the molar  
151 biomass flow in "daf" conditions  $\dot{n}_{bio,daf}$  [ $mol_{bio}/s$ ] in order to assess the molar  
152 flow of each component of the syngas as well as the syngas composition in  
153 wet and dry conditions. The molar flow of tar is given by Equation 4, the  
154 tar production versus the "daf" biomass input  $x_{tar}$  [% wt. "daf" biomass] is  
155 calculated by Equation 5. Furthermore, Equation 6 can be used to evaluate  
156 the volumetric tar amount  $g_{tar,vol}$  [g/Nm<sup>3</sup>] in the syngas.

$$\dot{n}_{tar} = n_{tar} * \dot{n}_{bio,daf} \quad (4)$$

$$x_{tar} = \frac{n_{tar} * MW_{tar}}{MW_{bio}} \quad (5)$$

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

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

$$\eta_{g,cold} = \frac{\dot{V}_g HHV_{syngas, clean}}{\dot{m}_f HHV_{bio, ar}} \quad (7)$$

163

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

164

165 where  $x_{H_2}$ ,  $x_{CO}$ ,  $x_{CH_4}$  [% vol] are the volumetric fraction of  $H_2$ ,  $CO$ ,  $CH_4$   
 166 in the dry syngas and  $HHV_{H_2}$ ,  $HHV_{CO}$ ,  $HHV_{CH_4}$  [MJ/Nm<sup>3</sup>] are the higher  
 167 heating values of  $H_2$ ,  $CO$  and  $CH_4$ .

168 However, the molar specific amount of the syngas components have to  
 169 be estimated. An algorithm similar to the one suggested in [39] is adopted  
 170 here. The first step is to choose an initial temperature  $T$  [K] and calculate  
 171 the equilibrium constant of the following reactions:

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

173 • **K2:** Hydrogasification  $C + 2H_2 \leftrightarrow CH_4$

174 • **K3:** Methane steam reforming  $CH_4 + H_2O \leftrightarrow CO + 3H_2$

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

$$K_1 = e^{\frac{4276}{T} - 3.961} \quad (9)$$

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

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

177 The System 12 is composed of three chemical balances calculated from  
178 Equation 1 (carbon, hydrogen and oxygen) and the three equilibrium con-  
179 stants for water-gas, hydrogasification and methane reforming reactions. The  
180 system is solved with the Newton-Raphson method.

$$\left\{ \begin{array}{l}
n_{CO} + n_{CO_2} + n_{CH_4} + n_{tar} - 1 = 0 \\
2n_{H_2} + 2n_{H_2O} + 4n_{CH_4} + pn_{tar} - x - 2w = 0 \\
n_{CO} + 2n_{CO_2} + n_{H_2O} + qn_{tar} - w - 2m - y = 0 \\
K_1 = \frac{n_{CO_2} * n_{H_2}}{n_{CO} * n_{H_2O}} \\
K_2 = \frac{n_{CH_4} * \frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}}{n_{H_2}^2} \\
K_3 = \frac{n_{CO} * n_{H_2}^3}{\left(\frac{\dot{n}_{tot,wet}}{\dot{n}_{bio,daf}}\right)^2 n_{H_2O} n_{CH_4}}
\end{array} \right. \quad (12)$$

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

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

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

189 where  $n_j$  [moles] and  $HF_j^0$  [kJ/kmol] are the specific moles amount and  
190 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:

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

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

#### 202 2.4. Biogas modeling

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

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

- 211 1. Estimating the degradation of selected biomasses.

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

218 Assuming hydrolysis as a limiting step of the anaerobic digestion reactions  
 219 chain, the biomass degradation can be described as a first order kinetic model  
 220 [42]. The mass balance of the substrate for a generic reactor with constant  
 221 volume and flow rates can be written as follows [43]:

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

222 where  $\hat{S}_e$  is the biomass concentration in the effluent flow rate [ $\text{kg}_{ts}/\text{m}^3$ ],  
 223  $\hat{S}_t$  is the biomass concentration in the in-fluent flow rate [ $\text{kg}_{ts}/\text{m}^3$ ],  $Q_i$  is  
 224 the influent flow rate [ $\text{m}^3/\text{day}$ ],  $Q_e$  the effluent flow rate [ $\text{m}^3/\text{day}$ ],  $V$  is the  
 225 digester active volume [ $\text{m}^3$ ] and  $k$  is the hydrolysis kinetic constant [ $\text{day}^{-1}$ ].

226 Starting from the concentration of the effluent flow, it is possible to cal-  
 227 culate the efficiency of the degradation process and, from this, to trace back  
 228 the HRT. The amount of reacting mass in the digester is also calculated con-  
 229 sidering the partial degradation of the substrate and the degree of dilution.  
 230 The latter is evaluated in terms of water flow rate or recirculation flow rate  
 231 as a function of the desired water content of the substrate and the desired  
 232 organic loading rate (ORL).

233 To increase the accuracy of the results, each biomass used as input is  
 234 broken into its constituents: sugars, proteins, fats, cellulose, hemicellulose,

235 lignin, ash, and non-degradables parts according to the Van Soest method  
236 (also called NDF method) [9]. A specific hydrolysis kinetic constant is as-  
237 signed to each constituent type, in order to take into account the differ-  
238 ent degradation rates which typically occurs in different substrates. Spe-  
239 cific aspects of the modeling approach were inspired by several other works  
240 [43, 9, 42, 44].

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

252

### 253 *2.5. Evaluation collecting and processing costs*

254 This work gives an overall energy balance evaluation of the proposed so-  
255 lution. Thanks to this evaluation is possible to demonstrate the advantages  
256 related to the improved utilization of maize. On the other hand it is impor-  
257 tant to assure that the proposed implementations an changes in the harvest-  
258 ing process do not affect the economy of system. The first step that needs to  
259 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*

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

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



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

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

### 296 *3.2. Biogas modeling*

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

305 The results of the biogas simulations are reported in Table 5. The model  
306 gives the specific power output in terms of  $m_{biogas}^3/kg_{ts}$ . This value is used

307 to calculate the equivalent silage specific primary energy  $e_A$  [ $kWh_{PE}/kg_{ts}$ ]  
308 that is the primary energy produced (biogas chemical energy) by one kg of  
309 dry maize silage. The model also calculates the methane productivity of the  
310 feedstocks which it is compared with literature data as reported in Table 5.  
311 The average error of 12% is considered acceptable for the model validation.

### 312 3.3. Gasification modeling

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

330 The gasifier conversion rate is evaluated in terms in dry stover specific  
331 primary energy  $e_{stover}$  [ $kWh_{PE}/kg_{ts}$ ]. In case of electrical power output eval-

332 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

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

$$\begin{aligned}
 e_B &= f_{ts,grains} * e_{grains} + (1 - f_{ts,grains}) * e_{stover} = \\
 &= 0.46 * 4.051 + (1 - 0.46) * 3.758 = 3.89
 \end{aligned}
 \tag{16}$$

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

353 productivity of 83.98 tons per hectare of as-received maize plant (type FAO  
354 700: full ripeness stage) [5]. The moisture of Case B is 68.14 % as suggested  
355 by Blandino et al. [54]. The final remark need to be addressed to the cost of  
356 the two proposed plants. The costs of the refer 100 kW power plant can be  
357 found in literature, in particular in Italy, in 2013 the cost for maize power  
358 plants is almost 10 €/W [55] considering:

- 359 • Approximately 900 m<sup>3</sup> digester tank with gasometer
- 360 • 100 kW combine heat and power system
- 361 • Auxiliary systems such as mixers, blowers, sensors and control systems
- 362 • Automatic feeding
- 363 • Storage tank

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

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

#### 376 **4. Conclusions**

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

#### 398 **Nomenclature**

399	$\Delta H_{T,i}$	enthalpy difference [kJ/kmol]	420	$HF^0$	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 <sup>3</sup> /s]	423		[MJ/Nm <sup>3</sup> or MJ/kg]
403	$\hat{S}_e$	biomass concentration in the	424	$HRT$	hydraulic retention time
404		effluent flow rate [kg <sub>ts</sub> /m <sup>3</sup> ]	425	$iNDF$	non-degradable neutral de-
405	$\hat{S}_t$	biomass concentration in the	426		tergent fiber
406		in-fluent flow rate [kg <sub>ts</sub> /m <sup>3</sup> ]	427	$K$	equilibrium constant [ad]
407	$ADL$	acid detergent fiber/lignin	428	$k$	hydrolysis kinetic constant
408	$ASH$	ash content of the biomass [%]	429		[day <sup>-1</sup> ]
409	$C$	carbon	430	$M$	total moisture content of the
410	$C_p$	specific heat [J/(mol K)]	431		biomass [%]
411	$CHP$	combined heat power	432	$m$	specific molar amount of oxy-
412	$E$	energy [kWh]	433		gen [ $mol/mol_{bio}$ ]
413	$e$	specific biomass energy pro-	434	$MW$	molecular weight [ $g/mol$ ]
414		ductivity [ $kWh_{PE}/kg_{ts}$ ]	435	$N$	nitrogen
415	$ER$	equivalence ratio [ad]	436	$n$	specific molar amount of gases
416	$f$	mass fraction [%]	437		and tar [ $mol/mol_{bio}$ ]
417	$g$	volumetric tar amount	438	$NDF$	neutral detergent fiber
418		[ $g/Nm^3$ ]	439	$O$	oxygen
419	$H$	hydrogen	440	$ORL$	organic loading rate
			441	$Q_e$	effluent flow rate [m <sup>3</sup> /day]

442	$Q_i$	influent flow rate [m <sup>3</sup> /day]	454	$p$	hydrogen coefficient of tar
443	$T$	temperature [K]	455	$prod$	product
444	$V$	digester active volume [m <sup>3</sup> ]	456	$q$	oxygen coefficient of tar
445	$w$	specific molar amount of	457	$react$	reactant
446		biomass moisture [ $mol/mol_{bio}$ ]	458	$ts$	total solid
447	$x_{tar}$	tar production versus daf	459	$vs$	volatile solid
448		biomass input [% wt.]	460	$x$	hydrogen coefficient of the
449	<b>Subscripts</b>		461		biomass
450	$ar$	as received	462	$y$	hydrogen coefficient of the
451	$bio$	biomass	463		biomass
452	$daf$	dry ash free	464	$z$	nitrogen coefficient of the
453	$g$	gas	465		biomass

466 **References**

467 **References**

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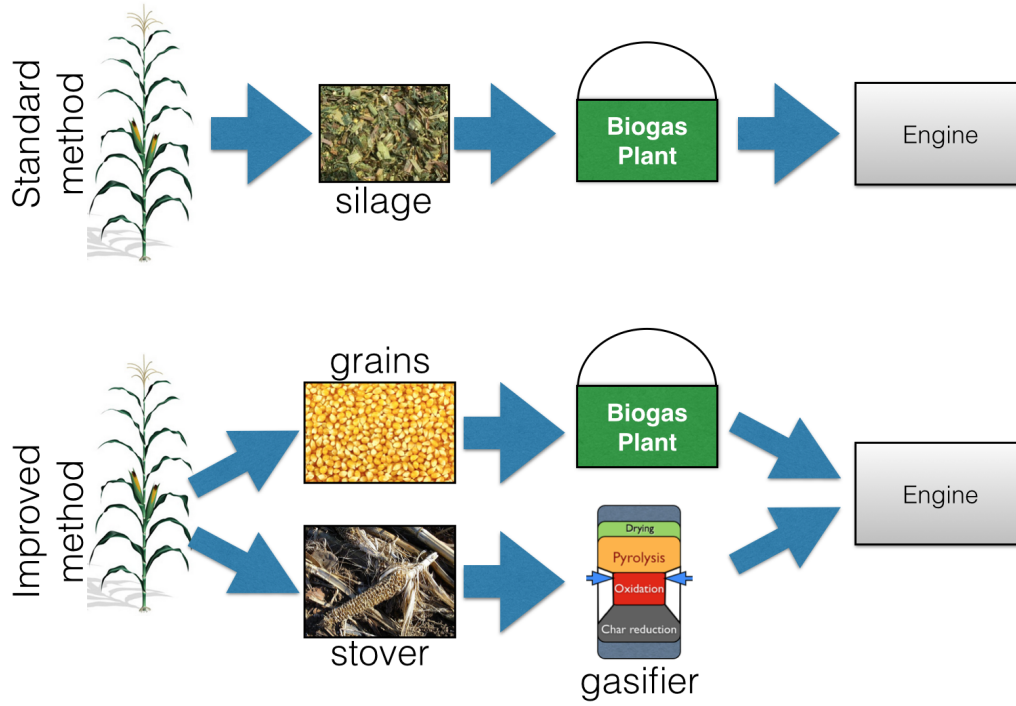


Figure 1: System layout

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 [ $\text{ton}_{ts}/\text{ha}$ ]	6.13	3.58	1.58	1.08	0.93

Table 2: Corn grains and stover properties

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 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 [% of ts]	XL	4.27
Non fiber carbohydrate [% of ts]	NFC	3.60
Neutral detergent fiber [% 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 [% of ts]	XP	8.2
Crude fat [% of ts]	XL	3.4
Non fiber carbohydrate [% of ts]	NFC	76.6
Neutral detergent fiber [% of ts]	NDF	10.7
Acid detergent fiber [% of ts]	ADL	0.5
Non degradable fiber ratio	iNDF/ADL	2

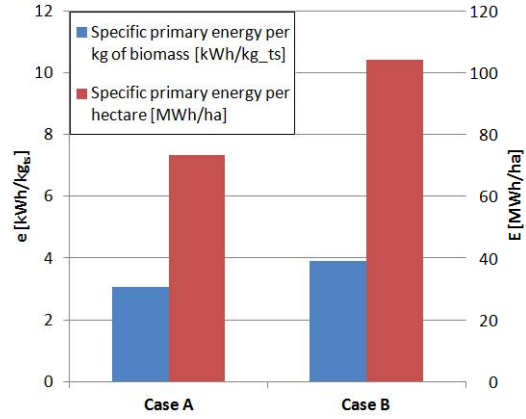


Figure 2: Energy conversion balances in the two cases

Table 3: Equivalent maize silage properties

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

Table 4: Results of the stover gasification

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 <sup>3</sup>
Volumetric syngas flow	$\dot{V}_{syngas}$	47.0 Nm <sup>3</sup> /h
Wet syngas higher heating value	$HHV_{syngas,w}$	4.68 MJ/Nm <sup>3</sup>
Dry syngas higher heating value	$HHV_{syngas,d}$	4.93 MJ/Nm <sup>3</sup>
Cold gas efficiency	$\eta_{cold}$	71.2 %
Dry stover specific primary energy	$e_{stover}$	3.758 kWh <sub>PE</sub> /kg <sub>ts</sub>
IC engine efficiency	$\eta_{engine}$	0.3
Electrical generator efficiency	$\eta_{gen}$	0.95
Power plant conversion rate	$e_{plant,model}$	1.07 kWh <sub>el</sub> /kg <sub>ts,stover</sub>

Table 5: Results and validation of the anaerobic digestion model

	Literature methane production [32, 52] [m <sup>3</sup> /kg <sub>vs</sub> ]	Model methane production [m <sup>3</sup> /kg <sub>vs</sub> ]	Absolut error [%]	Model biogas production [m <sup>3</sup> /kg <sub>ts</sub> ]	Model specific primary energy [kWh <sub>PE</sub> /kg <sub>ts</sub> ]
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 <sub>PE</sub> /kg <sub>ts</sub>
Case (B) specific primary energy	$e_B$	3.890 kWh <sub>PE</sub> /kg <sub>ts</sub>
Increase	$\Delta e$	26%
Increase considering silaging matter losses	$\Delta e_{sml}$	39 %
Case (A) specific primary energy per hectare	$E_A$	73.173 MWh <sub>PE</sub> /ha
Case (B) specific primary energy per hectare	$E_B$	104.08 MWh <sub>PE</sub> /ha