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Review of biochar role as additive in anaerobic digestion processes

- 2 Marco Chiappero^a, Omid Norouzi^b, Mingyu Hu^c, Francesca Demichelis^a, Franco
- 3 Berruti^d, Francesco Di Maria^b, Ondřej Mašek^c, Silvia Fiore^a *

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- 5 a DIATI (Department of Engineering for Environment, Land and Infrastructures),
- 6 Politecnico di Torino, Corso Duca degli Abruzzi 24, 10129, Torino, Italy
- 7 b LAR Laboratory, Department of Engineering, University of Perugia, Perugia, Italy
- 8° UK Biochar Research Centre, School of Geoscience, University of Edinburgh,
- 9 Edinburgh, United Kingdom
- 10 d Institute for Chemicals and Fuels from Alternative Resources (ICFAR), Chemical and
- 11 Biochemical Engineering Department, Western University, London, Ontario, Canada

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- 13 corresponding author: Prof Silvia Fiore, DIATI, Politecnico di Torino, Corso Duca
- 14 degli Abruzzi, 24; 10129, Torino, Italy; Email: silvia.fiore@polito.it; Phone: +39
- 15 0110907613

16

17 Abstract

- 18 Anaerobic digestion (AD) could be considered as a mature technology and nowadays it
- 19 can still play a pivot role because of the urgent need to provide renewable energy sources
- 20 and efficiently manage the continuously growing amount of organic waste. Biochar (BC)
- 21 is an extremely versatile material, which could be produced by carbonization of organic
- 22 materials, including biomass and wastes, consistently with Circular Economy principles,
- 23 and "tailor-made" for specific applications. The potential BC role as additive in the
- 24 control of the many well-known critical issues of AD processes has been increasingly

25 explored over the past few years. However, a clear and comprehensive understanding of 26 the connections between BC and AD is still missing. This review paper analyses and 27 discusses significant references (review articles, research papers and international 28 databases and reports), mostly published in the last 10 years. This review is aimed at 29 addressing three key issues related to the better understanding of the BC role in AD 30 processes: 1. Investigation of the influence of BC properties on AD performances and of 31 their ability to counteract its main challenges; 2. Assessment of the optimal BC 32 production chain (i.e. feedstock-pyrolysis-activation) to achieve the desired features; 3. 33 Evaluation of the economic and environmental advantages connected to BC use in AD 34 processes, compared to conventional solutions applied to address AD challenges.

35

36 Highlights

- 37 Biochar key properties were specific surface, pores and surface functional groups
- 38 Lignocellulosic biomass, slow pyrolysis and physical activation made the best biochar
- 39 Biochar addition exhibited economic and environmental advantages
- 40 Biochar improved methane production (in average 25 %) and digestate quality

41

42 Keywords: anaerobic digestion; biochar; bioenergy; biomass; circular economy; waste

43

44 Abbreviations

45 AD: anaerobic digestion; BC: biochar; BET: Brunauer–Emmett–Teller; CE: circular 46 economy; CEC: cationic exchange capacity; CIC: controlling internal circulation; CLSM: 47 confocal laser scanning microscopy; COD: chemical oxygen demand; DIET: direct 48 interspecies electron transfer; EAC: electron accepting capacity; EDC: electron donating 49 capacity; EU: European union; FAN: free ammonia nitrogen; FISH: Fluorescence in situ

50 hybridization; FW: food waste; GAC: granular activated carbon; GHG: green-house gas; 51 HRT: hydraulic retention time; IET: interspecies electron transfer; LCA: life cycle 52 analysis; MSW: municipal solid waste; OFMSW: organic fraction of municipal solid 53 waste; OLR: organic loading rate; PAH: polyaromatic hydrocarbons; PCBs: 54 polychlorinated biphenyls; S/I: substrate to inoculum ratio; SMP: specific methane 55 potential; SS: suspended solids; SSA: specific surface area; TAN: total ammonia 56 nitrogen; TPAD: two phased anaerobic digestion; TSS: total suspended solids; UASB: 57 up-flow anaerobic sludge blanket; VFA: volatile fatty acids; WAS: waste activated 58 sludge; WWTP: wastewater treatment plant; WV: working volume; V: volume.

59

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82 1. Introduction

83 The global energy consumption increased from around 8,800 million tonnes of oil 84 equivalents (Mtoe, including coal, gas, oil, electricity, heat and biomass) in 1990 to 85 14,400 Mtoe in 2018 [1]. In 2016 renewable energy sources met this demand for about 86 14 % of primary energy supply, while fossil fuels accounted for 81% of greenhouse gas 87 (GHG) emissions [2]. The need to decrease fossil fuel dependency and GHG emissions 88 and to generate economic benefits is stimulating initiatives to produce energy as well as 89 high-value chemicals and products from sustainable non-food biomass, residues, co-90 products and wastes.

91 Anaerobic digestion (AD) is a well-established technology for the treatment of 92 wastewater and organic waste, e.g. the organic fraction of the municipal solid waste 93 (OFMSW), waste activated sludge (WAS), animal manure, etc. Biogas can directly 94 replace fossil fuels in heat and power generation or be upgraded to bio-methane as vehicle 95 fuel or injected in the gas grid [3], while digestate can be employed as soil amendment 96 [4,5]. Other benefits of AD, compared to alternative treatment options applied to organic 97 waste, include control of odor and GHG emissions, removal of pathogens and adaptability 98 to many substrates [6]. AD has been extensively implemented, from small-scale 99 applications, particularly in rural areas and developing countries, to large-scale industrial 100 plants [7]. Despite the rapidly growing number of biogas installations in Europe (from 101 about 6,200 to 18,202 between 2009 and 2018) [8], many challenges regarding AD 102 processes still persist. Different options have been explored to address these challenges: 103 optimization of working parameters and bio-reactor configuration, co-digestion and 104 nutrients control [9]; adoption of pre-treatments to enhance the degradability of various 105 substrates [10,11]; use of inorganic and biological additives to support biomass

106 immobilization, supplement nutrients, mitigate inhibitors and improve process stability 107 [12–14].

108 Among the additives, conductive carbonaceous materials, as granular activated carbon 109 (GAC) and biochar (BC), gained interest because of their ability of enhancing methane 110 production [15,16]. BC may present several advantages over other additives; it can be 111 produced with a wide variety of physico-chemical properties by controlling feedstock, 112 pyrolysis operating conditions and the activation process [17,18], resulting "tailor-made" 113 for specific applications. In recent years, several authors confirmed the possibility of 114 increasing methane production through BC addition (Table 1), suggesting different 115 potential mechanisms: (1) increase of the buffering capacity of the AD system; (2) 116 mitigation of inhibition phenomena or agents; (3) support media for biomass 117 immobilization; (4) promotion of syntrophic metabolisms; (5) enhancement of digestate 118 quality; (6) biogas cleaning and upgrading. The most significant and updated scientific 119 literature investigating the BC role within AD processes can be classified as follows 120 (Table 1):

- 121 Studies considering the advantages of coupling AD with thermal processes
- 122 (gasification, pyrolysis and hydrothermal carbonization) from a general point of view;
- 123 Studies exploring the positive influence of BC on AD, based on methane and/or biogas
- production enhancement and/or improvement of digestate quality;
- 125 Studies analysing in detail the BC features that may influence one or more of the
- above-mentioned mechanisms.
- 127 Considering the digestate, BC has been reported to enhance its quality in terms of nutrient
- 128 retention, increase of the carbon-to-nitrogen ratio and reduction of nutrient leaching [19].

129 Consequently, BC doesn't need to be separated from digestate at the end of its utilization 130 in AD.

131

132 Table 1. Overview of literature studies exploring the connections between BC and AD

G-1: 11 1 41	C(1' 1 ' 41	C(1' 1 ' 41	C(1' 1'
Studies considering the	Studies exploring the positive influence of BC	Studies exploring the positive influence of BC	Studies analysing in details the biochar features
general advantages of coupling AD with thermal	on AD, based on a general	on AD, based on a general	that may influence one or
processes	analysis of methane and/or	analysis of the	more of the six above-
processes	biogas production	improvement of digestate	mentioned mechanisms
	enhancement	quality	memorica memamonio
Pecchi and Baratieri,	Linville et al., 2017 [24]	Bruun et al., 2011 [32]	Fagbohungbe et al., 2017
2019 [20]	Torri and Fabbri, 2014	Shen et al., 2016 [33]	[19]
Fabbri and Torri, 2016	[25]	Shen et al., 2017 [34]	Ye et al., 2018 [13]
[21]	Sunyoto et al., 2016 [26]	5.1011 ev a, 2017 [5 1]	González et al., 2018 [15]
Hübner and Mumme,	Jang et al., 2018 [27]		Zhang et al., 2018 [16]
2015 [22]	Sun et al., 2019 [28]		Codignole Luz et al.,
Salman et al., 2017 [23]	Zhao et al., 2015 [29]		2018 [35]
	Inthapanya et al., 2012		Masebinu et al., 2019 [36]
	[30]		Pelaez-Samaniego et al.,
	Inthapanya and Preston,		2018 [37]
	2013 [31]		G. Wang et al., 2018b
			[38]
			D. Wang et al., 2017 [39]
			Fagbohungbe et al., 2016
			[40]
			Mumme et al., 2014 [41]
			Shanmugam et al., 2018
			[42]
			Su et al., 2019 [43]
			Lü et al., 2016 [44]
			Cruz Viggi et al., 2017
			[45]
			Li et al., 2018 [46]
			Martínez et al., 2018 [47]
			Luo et al., 2015 [48]
			Wang et al., 2019 [49]
			C. Wang et al., 2018 [50] Zhao et al., 2016 [51]
			Yu et al., 1999 [52]
			Cooney et al., 2016 [53]
			Dang et al., 2017 [54] Sasaki et al., 2010 [55]
			Martins et al., 2018 [56]
			S. Chen et al., 2014 [57]
			Pan et al., 2019b [58]
			Qiu et al., 2019 [59]
			Z.a o. a, 2017 [57]

135 To the best of our knowledge, despite the growing number of studies investigating the 136 connections between BC and AD, three main issues still need to be explored: a clear 137 comprehension of the above-mentioned six mechanisms; a correlation between BC 138 features and their effects on AD processes; environmental and economic outcomes related 139 to BC use. The present review is therefore aimed at addressing the following issues: 1) 140 investigation of the influence of BC properties on AD performance and of their ability to 141 counteract its main challenges; 2) assessment of the optimal BC production chain (i.e. 142 feedstock-pyrolysis-activation) to achieve the desired features; 3) evaluation of the 143 economic and environmental advantages connected to BC use in AD processes, compared 144 to conventional solutions applied to address AD challenges.

145 2. Review methodology

146 The selection of scientific literature was made considering the following criteria:

- 147 a) Relevant international databases and information sources. Bibliometric sources as 148 Web of Science, Science Direct and Scopus were used to retrieve articles, book-chapters 149 and proceedings of indexed conferences. Official international databases (Eurostat, 150 International Energy Agency, European Biogas Association, etc.) were searched for the 151 selection of technical reports and statistics;
- 152 b) Chronological order. The majority of the references in this review are from 2015 to 153 2019 (67 %) and from 2009 to 2014 (26 %), with the balance before 2009 (7 %);
- 154 c) Relevant keywords for the topics of interest. A first survey adopted the following 155 keywords in different combinations: "biochar, anaerobic digestion, methane, pyrolysis, 156 economic assessment, environmental assessment, life cycle analysis, LCA". Based on

157 this preliminary selection, the review questions have been identified. Thereby, additional 158 keywords have been adopted to face specific issues, as: "biochar, buffer capacity, 159 anaerobic digestion, volatile fatty acids", "biochar, ammonia inhibition, ammonia 160 removal, anaerobic digestion", "syntrophic metabolisms, biomass immobilization, 161 biochar, anaerobic digestion, DIET, electron transfer". 162 d) Selection of the references based on content analysis. After a first screening 163 considering the abstracts of all identified reference sources, 259 relevant references 164 were analyzed as full content and included in this review. The relevant scientific 165 content was sourced from scientific articles (240 research papers and reviews – 93 %), book chapters and indexed conference proceedings (11 documents - 4 %), and 166

167 international databases and reports (8 documents - 3 %). Considering only the 168 scientific research papers and the reviews, they were related to (Figure 1): AD (31 %) and pyrolysis (10 %); coupling AD and thermal processes (10 %); BC features (25 %); BC and AD interaction mechanisms (14 %); economic and environmental aspects related to BC use (10 %). The distribution of the relative contributes of the scientific literature to the different topics concerning the investigation of BC role in AD in 2009 - 2019 exhibits a clear increasing trend due to a growing interest, particularly in the 174 last five years, for the topics covered by the present work and consequently strengthens the need to address the above mentioned review questions.

176 This review is structured according to the contents outline. At the end of each section, the 177 last paragraph summarizes the main findings and research gaps of the specific topic 178 explored, which are finally condensed in the conclusions.

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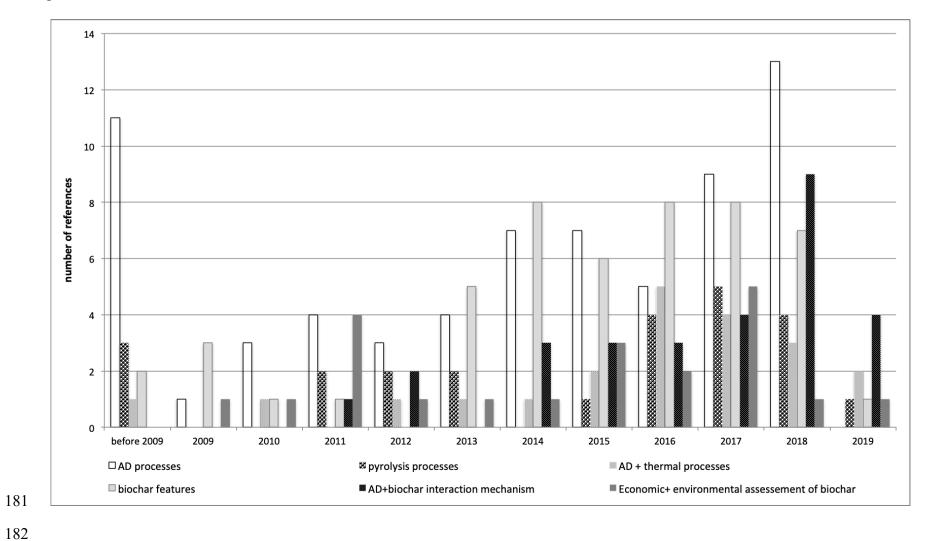
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180 Figure 1. Relative contributions of the scientific literature selected for this work



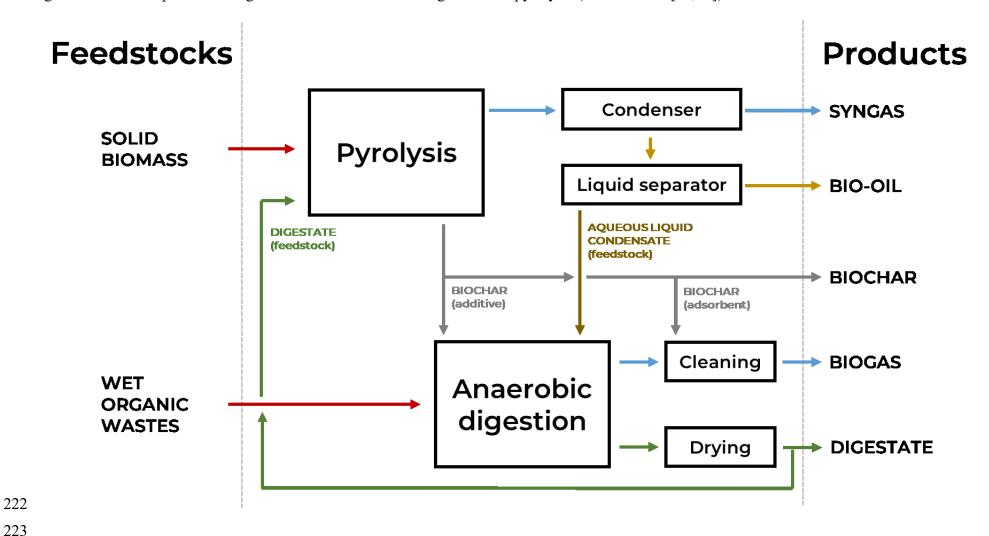
183 3. Biochar

184 BC is the porous carbonaceous solid residue of the complex physical and chemical 185 phenomena simultaneously occurring during the thermochemical treatment of raw 186 biomass [60–63]. While AD is particularly suitable to produce biogas and digestate from 187 wet organic waste (Figure 2), pyrolysis is a thermochemical process able to convert solid 188 dry biomass in an oxygen-limited environment into valuable liquid chemicals (bio-oil), 189 as well as gaseous biofuels and charcoal (biochar, BC) [64–67]. Although pyrolysis was 190 conventionally mostly focusing on the production of liquid biofuels from woods and 191 purpose-grown energy crops, the most recent studies found that it is not cost effective 192 unless the feedstock is inexpensive, dry and readily available, and there are valuable 193 chemical co-products (liquid bio-oil and solid BC) or favorable government policies [68]. 194 Both AD and pyrolysis can contribute to fulfill Circular Economy targets, but research 195 on biological and thermochemical processes traditionally progressed along parallel and 196 separate pathways, often in competition. 197 In recent years, improved understanding of the characteristics of bio-oil and BC has 198 helped to develop key quality requirements to expand their potential value [69]. High 199 value liquid bio-oil products can be obtained from the pyrolysis of biomass residues, such 200 as phenolic substitutes for adhesives or antioxidants [70,71]. However, there are 201 limitations: the most valuable chemicals are usually mixed with impurities that impair 202 their properties. Similarly, there are many promising applications for BC, such as soil 203 improver [18], composite materials [72], activated carbon [73], electrodes for batteries or 204 electrolysis cells [74], metallurgical coke substitute [75], catalyst for tars [76], and could 205 be eligible for carbon credits [77]. The efficiency of BC in most applications significantly

206 depends on its physico-chemical properties and elemental composition, which are subject 207 to variations depending on feedstock, pyrolysis conditions, pre- and post-processing 208 treatments.

209 The characteristics of AD and pyrolysis processes and products may offer interesting 210 opportunities for the integration of such technologies (Figure 2), with the objective of 211 contributing to the Circular Economy. For example, pyrolysis could be utilized to convert 212 dewatered digestate into BC, which could then be used for a variety of applications 213 [17,78,79], including its potential use to improve the quality of biogas and digestate [37], 214 or to reduce instabilities and inhibition in digestors [16,24]. On the other hand, the acetic 215 acid-rich aqueous pyrolysis condensate could be utilized as a feedstock for AD, possibly 216 with the addition of BC for the selective adsorption of toxic inhibitors [21–23,25,80–88]. 217 In conclusion, BC represents the key link between pyrolysis and AD and, consequently, 218 in this section we wish to review its characteristics, including suitable feedstocks, 219 production technologies, activation processes and characterization techniques.

221 Figure 2. Outline of potential integrations between anaerobic digestion and pyrolysis (derived from [20,21]).



224 3.1. Feedstocks

225 During thermal processing under oxygen free or oxygen-limited conditions (as pyrolysis), 226 as the temperature increases, the polymeric components of lignocellulosic biomass crack 227 and liquid intermediate phases are formed with different visco-elastic behaviors [89,90]; 228 under slow progressive heating, cellulose remains hard and elastic up to 400 °C, whereas 229 hemicellulose softens within a relatively narrow temperature range (230 to 280 °C) and 230 remains viscous, while lignin softens and bubbles over a broader temperature range (150 231 to 350 °C) and remains mainly elastic. The overall process cannot be simply designed 232 based on the linear combination of the behaviors of the individual components, but it is 233 the result of their interactions at different temperatures and it is catalyzed by the mineral 234 matter present in the ashes. As the temperature increases over these ranges, the 235 depolymerized liquid undergoes two competitive processes, oligomer evaporation and 236 crosslinking reactions, with the former favored at high heating rates, and the latter under 237 slow pyrolysis. These findings explain why, under slow pyrolysis, the char yield is 238 greater, and the particles globally keep the same macro-structure of the original biomass, 239 although forming an intermediate soft material. Montoya et al. (2017) [91] confirmed the 240 presence of molten liquid intermediates formed from cellulose, xylan and lignin and 241 showed that the presence of minerals prevents the complete liquefaction of the xylan, 242 therefore contributing to the preservation of the general shape and structure of the solid 243 material during pyrolysis. 244 Therefore, BC can be produced from a wide range of lignocellulosic materials, containing 245 cellulose, hemicellulose (xylan), lignin, small quantities of other organic compounds 246 (phenols, phytosterols and fats) and inorganics (P, N, S, Si, alkali and alkaline earth 247 metals and traces of other minerals). Such materials include wood, agricultural and forest

248 residues, food waste, sugars, industrial organics, sewage sludge and manure. BC can be 249 mesoporous or microporous depending on the operating conditions employed for its 250 production and on the feedstock.

251 3.2. Thermochemical production

252 Recent reviews illustrated the different technologies and processes used experimentally 253 and industrially to produce BC [17,92]. These include torrefaction, pyrolysis, 254 gasification, combustion and hydrothermal carbonization [93]. Scale and potential 255 mobility are important considerations in relation to the feedstock supply, logistics, 256 seasonality, further refining, of products quantities, characteristics and value, and 257 potential markets [17,60].

258 *Torrefaction* involves biomass treatment at atmospheric pressure and within a 259 temperature range of 200–300 °C, without oxygen or with limited oxygen supplies [94]. 260 As discussed earlier, partial depolymerization and liquefaction of the biomass 261 components take place reducing the biomass tenacity and improving grindability. 262 Torrefaction produces char with less moisture, higher energy density, lower weight, lower 263 O/C and H/C ratio, increased hydrophobic nature and resistance to biological degradation 264 with respect to the original feedstock. Typical yields of torrefied biomass range between 265 50 and 80 % [95].

266 Gasification is a thermochemical process carried out at temperatures higher than 750 °C 267 in the presence of a gasifying agent (air, oxygen, or steam) at atmospheric or at high 268 pressures. Under these conditions, biochar yields are not sufficiently significant to 269 consider gasification an appropriate BC production process. Similarly, combustion is not 270 a suitable BC production process, since, under good combustion conditions, BC yield 271 should be negligible.

272 The various modes of *pyrolysis* include slow, intermediate, fast, flash, and ultra-pyrolysis, 273 carried out under vacuum, atmospheric pressure or under pressure [65,69]. As a result of 274 the balance between primary, secondary cracking and recombination reactions, bio-oil 275 yields are typically maximized (up to 70 %) at intermediate temperatures (450-550 °C), 276 faster heating rates (100-500 °C/s) and short vapor residence times (< 1~2 s), which are 277 characteristic of fast and flash pyrolysis. Under these conditions, BC yields are typically 278 of the order of 15~20 %. On the other hand, higher BC yields (25~40 %) are achieved at 279 moderate temperatures (300-450 °C), slower heating rates (~ 1 °C/s) and longer vapor 280 residence times ($> 5 \sim 10$ s), representative of slow or intermediate pyrolysis, when bio-oil 281 yields vary between 40 and 50 % with the balance being gas. 282 Hydrothermal carbonization (HTC) is performed on wet biomass (moisture > 10 %) as 283 feedstock, making the process highly energy-intensive [96]. HTC happens under water in 284 a sealed confined system and heated at the temperature range of 175–300 °C for up to 16 285 h under saturated pressure under subcritical conditions producing tar-free BC (hydrochar) 286 with large number of functional groups [97–99]. Hydrochars are usually obtained at 287 yields of 20-25 % [96], and they contain mainly aliphatic compounds and more oxygen 288 functional groups and higher cation exchange capacity than conventional BC. On the 289 other hand, they have lower surface area, microporosity and carbon stability [96,100]. 290 Among the various thermal technologies, pyrolysis has been the most investigated 291 technique and it is considered the best technology for the production of BC [101] from 292 relatively dry feedstock (moisture content below 10 %). However, also HTC is gaining 293 interest to produce hydrochar, particularly from wet biomass [96]. In a comparison 294 between the chars, it should be noticed that BC yields are higher, as well as porous 295 structure and aromatics and ash contents, while hydrochar exhibits a non-porous

296 structure, lower fixed and total carbon contents and higher amounts of alkyl functional 297 groups [96].

298 3.3. Activation

299 BC with different properties can be produced from a wide range of feedstock and by 300 adjusting processing conditions. Further modification of BC chemical and physical 301 properties can be achieved through a process defined "activation", aimed at increasing 302 BC porosity and modifying its pore size distribution (Figure 3), as well as to some extent 303 surface chemistry. Activation can be carried out in a number of ways, depending on type 304 of activation agents (e.g. physical and chemical activation) or mode of operation into 305 single-stage or multi-stage activation. 306 Chemical activation utilizes chemical agents, e.g., H₃PO₄, HNO₃, KOH, NaOH, H₂SO₄, 307 and ZnCl₂ [102,103]. It typically involves two steps; in the first the feedstock is 308 impregnated with a selected chemical agent, and then thermally treated in the second step. 309 Another option is the activation of already produced BC by soaking it in a chemical agent, 310 followed by a thermal treatment. Depending on the agent selected and thermal treatment 311 conditions used, different degrees of activation can be achieved. The activated BC needs 312 to be thoroughly washed with deionized water to neutralize its pH and to remove any 313 remaining chemicals [104], and this procedure can contribute to a negative environmental 314 impact of the technology. Oxidative activation that uses acidic or alkaline agents is among 315 the most common activation methods. Besides enhancing porosity and surface area, it 316 also creates oxygen-containing functional groups on the surface of BC (e.g. carboxyl, 317 hydroxyl, lactone, phenol, carbonyl, and peroxide groups). These functional groups play 318 an essential role in different applications of BC, such as for contaminant removal or 319 nutrient sorption.

320 *Physical activation* does not utilize chemicals, thus avoiding some of the negative 321 environmental impacts that can be associated with chemical activation. Instead, BC is 322 exposed to physical oxidizing agents, such as steam, CO₂, ozone or air/O₂, typically at 323 temperatures above 700 °C. These gaseous agents penetrate the internal structure of BC 324 gasifying the carbon atoms, which results in opening and widening of pores [105]. 325 Similarly to chemical activation, physical activation not only enhances BC porosity, but 326 also modifies its surface functional groups, increasing the abundance of oxygen 327 functional groups [105]. 328 In conclusion, both chemical and physical agents can be used effectively to activate BC

329 and obtain high porosity, surface areas over 1000 m²/g, and modified surface functional 330 groups, tuned to specific applications. The key challenges of BC activation are without 331 any doubt related to their potential negative environmental impacts, mainly for chemical 332 activations, and high-energy intensity need.

333 3.4. Physico-chemical characteristics

334 The variability of BC physico-chemical properties resulting from the proper selection of 335 feedstock, pyrolysis and activation can be adapted to a wide range of applications.

336 Specific surface area (SSA), expressed as m²g⁻¹, is the ratio between the total surface area

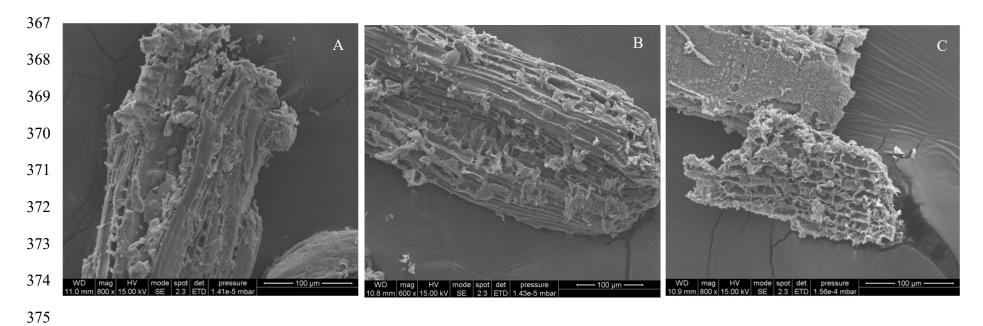
337 and the total particle mass of BC [106], commonly determined by Brunauer–Emmett–338 Teller (BET) analysis. The reported SSA values of BC can vary significantly, in the range 339 10^o - 10² m²g⁻¹ [107,108] depending on pyrolysis temperature and feedstock, and up to 340 over 10³ m²g⁻¹ after activation. SSA is a key parameter enhancing the adsorption capacity 341 of BC, even if the chemical composition of its surface is determinant in the interactions 342 with other species [109]. SSA is related to BC pore size distribution [18].

343 Porosity can be described by the pore volume (m³g¹¹) and by the pore size distribution, 344 including micro-, meso-, and macro-pores (respectively < 2nm, 2-50 nm, >50 nm) [110], 345 based on their internal diameters. The relative abundance of these pores produces 346 different BC structures (Figure 3), since micro-pores have the major contribute to BC 347 surface area, micro- and meso-pores are essential for its adsorption capacity, while macro-348 pores can provide an appropriate habitat for microorganisms [18,111]. Yin et al. (2017) 349 [112] stated that within a proper range, a high temperature results in a large surface area 350 and pore formations. Trigo et al. (2016) [113] found that SSA increased between 350°C 351 and 700 °C for different hardwood biochars, while Chen et al. (2014) [114] revealed an 352 enhanced porous structure of BC from sewer sludge varying the temperature from 500 °C 353 to 900 °C.

354 Specifically considering the effect of the feedstock on BC characteristics [101], yields 355 and aromatic functional groups are much higher from lignin (48.8 %) than from 356 hemicellulose (21.1 %), while no differences were found in the elemental composition of 357 resulting BCs. High lignin and mineral content in the feedstock result in higher yields of 358 BC [115,116]. Woody and grassy biomasses generate a more carbon-rich BC when 359 compared to sewage sludge and manure [32,117,118]. Specifically considering the effect 360 of pyrolysis process on BC characteristics [119], increasing pyrolysis temperature 361 decreased concentrations of aliphatic carbons, oxygenated functional groups, nitrogen, 362 oxygen and hydrogen. However, carbon mass fraction, micropore volume, ash content, 363 fixed carbon and carbon stability increase with higher pyrolysis temperatures.

364

365 Figure 3. SEM micrographs of (A) raw biomass (eucalyptus) and of derived (B) biochar and (C) physically activated biochar (Courtesy of 366 the authors. The samples were produced at ICFAR and analysed at Politecnico di Torino).



377 BC composition is mainly of C, H, O, N, and S, along with mineral trace components, 378 such as K, Ca and Mg, varying according to pyrolysis conditions and feedstock [110]. 379 The mineral content can be enhanced by increasing the temperature of pyrolysis [107]. 380 The cationic exchange capacity (CEC) of BC can be defined as its ability of absorbing 381 and attracting cations (nutrients and heavy metals) decreasing with higher pyrolysis 382 temperature [18]. A high CEC value is due to a negative surface charge of BC. The 383 presence of surface functional groups such as hydroxyl, carboxylic, and amino, expressed 384 respectively as atomic ratio of H/C, O/C, and N/C (generally studied by Fourier transform 385 infrared (FTIR) spectroscopy), mostly depends on the feedstock and it is reduced with a 386 higher temperature of pyrolysis [107]. Further, the H/C ratio is considered as a measure 387 of the degree of aromatization of BC and the O/C ratio as an indication of more 388 oxygenated functional groups, contributing to high CEC values and hydrophilicity of BC 389 [120,121]. An increase in pyrolysis temperature has been reported to negatively affect the 390 amount of acidic functional groups and the CEC [112,122]. 391 pH values are generally basic and increase as the pyrolysis temperature grows, due to the 392 volatilization of acidic functional groups and to the higher ash content [18]. Moreover, 393 BC can present redox characteristics, being able to donate and accept electrons, and these 394 tendencies are quantified respectively by the *electron donating* and *accepting capacities* 395 [110]. Phenolic groups may be the electron donating species, while quinones and 396 polycondensed aromatic structures the electron accepting compounds [110,123]. In 397 addition, the *electrical conductivity*, expressed as S m⁻¹, decrease as consequence of the 398 increase of the pyrolysis temperature [17]. 399 A full characterization of BC involves several analytical methods [124]. Scanning

400 electron microscopy (SEM) is employed to investigate the superficial structure of BC, as

401 well as its chemical composition if coupled with energy dispersive X-ray spectroscopy 402 (SEM-EDX); transmission electron microscopy (TEM) allows to observe BC bulk 403 structure; X-ray diffraction (XRD) analysis detects the crystalline phases; nuclear 404 magnetic resonance (NMR) spectroscopy investigates BC composition, specifically the 405 aromatic structure; X-ray photoelectron spectroscopy (XPS) quantitatively analyses the 406 superficial composition.

407 4. Application of biochar as additive in anaerobic digestion

408 4.1. Buffer capacity and alkalinity

409 The volatile fatty acids (VFAs) produced as intermediates during AD tend to lower the 410 pH [125]. This effect is generally counteracted by syntrophic acetogens and methanogens 411 microorganisms, which convert VFAs to methane and carbon dioxide [126]. However, in 412 case of high organic loads of easily biodegradable wastes (i.e. when the VFAs production 413 rate exceeds the consumption rate), the accumulation of VFAs can occur, resulting in pH 414 drop and even in the failure of AD [126,127]. The buffer capacity in a digestor is linked 415 to the ability to neutralize VFAs, and it is considered a rate-limiting step in AD processes 416 [38,39,128]. The buffer capacity is determined by the alkalinity of the AD system, mainly 417 in the form of carbon dioxide and bicarbonate [6,125]. So it is technologically challenging 418 to build a bioreactor aimed at digesting easily biodegradable feedstock with high pH 419 buffer capacity [16]. Even if several methods have been proposed (Table 2) so far, the 420 most diffused solution to improve buffer capacity implemented both at lab [129] and full 421 scale [130] is co-digestion. In particular, manure or sewage sludge showed high buffer 422 capacity, mainly due to the quite high ammonia content, which is able to counteract the

423 pH drop due to VFAs production. Therefore, manure or sewage sludge are often co-424 digested with OFMSW and/or agricultural waste biomass.

425

426 Table 2. Advantages and disadvantages of conventional methods for creating buffering 427 capacity in bioreactors

Methods	Advantages	Disadvantages
Alkaline pretreatment [131]	Cellulose more available to the enzymes	Expensive and considered as the technological bottleneck
Commercial alkaline materials (e.g. NaOH and CaCO ₃) [132,133]	Easy pH adjustment	pH continually should be analyzed Temporary Lower biogas output at higher organic loads
Co-digestion [16]	Increasing methane yield Low cost and green	Hard to achieve an appropriate mixture of feedstocks to limit VFAs production
Geopolymer (Composite and monolith) [132,133]	Controllable alkali leaching Prevent the need for continuous pH adjustment	25 g L ⁻¹ costs around 0.10 \$

428

429 However, it still remains a great challenge to develop simple, permanent and cost-430 effective methods to improve buffer capacity of AD systems. Nowadays, BC is 431 considered as an attractive alternative to the aforementioned methods for two important 432 reasons. Firstly, it can be produced via cost-effective and environmentally friendly 433 approaches. Secondly, its physical and chemical properties can be matched with the 434 operational conditions [19,40,41].

435 BC's buffer capacity mainly depends on two factors:

436 I. Functional groups: Rapid accumulation of VFAs during AD process results in a 437 medium with low pH value in which some functional groups of BC like amine adsorbs H⁺ and accepts electron. This phenomenon could mitigate the sudden pH 438 439 drop.

Inorganic materials: Ash portion of biochar contains inorganic materials such as Ca,
K, Mg, Na, Al, Fe, Si and S. Among them, alkali and alkaline earth metals (AAEMs)
are responsible for alkalinity of biochar via reaction 1 (Ca and CxHxCOOH are
selected as representative of AAEMs and VFAs, respectively) [16]:

444
$$CaCO_3 + 2C_xH_xCOOH \rightarrow [C_xH_xCOO]2Ca + H_2O + CO_2$$
 (1)

445 Table 3 shows the literature data related to BC buffering capacity and its effects on bio-446 methane production, showing total alkalinity, pH values and the increase in CH₄ 447 production (%), ranging from 8 % up to about 26 %. Some studies suggest that the 448 alkalinity of BC due to AAEMs in ash fraction could effectively contribute to the 449 buffering capacity of AD against VFAs inhibition. Jang et al. (2018) [27] investigated the 450 effects of BC from dairy manure on AD of dry dairy manure at 20°C, 35°C, and 55°C. In 451 each case they observed a lower total VFAs concentration and increased methane 452 production. They suggested the role of high nutrients (9.1% Ca, 3.6% Mg, 1.3% N, 0.14% 453 P) and alkalinity potential of BC in enhancing methane production. Wang et al. (2017) 454 [39] investigated the role of BC from vermicompost on the buffering capacity of AD of 455 high organic loads of kitchen waste and chicken manure. The authors proved the high 456 buffer capacity of vermicompost BC to different short-chain VFAs (700-3800 mg L⁻¹), 457 probably due to the presence of AAEMs and superficial functional groups. Linville et al. 458 (2017) [24] studied the effects of BC from walnut shell during the AD of FW in 459 mesophilic and thermophilic conditions. They observed that BC improved process 460 stability by enhancing the total alkalinity from 2800 to 4800-6800 mg L⁻¹ CaCO₃ and pH 461 (initial pH >8) in BC amended digesters. Wei et al. (2020) [134] found enhanced methane 462 production and solids removal by adding BC from corn stover (rich of alkaline earth 463 metals) to AD of primary sludge. The authors observed higher total alkalinity (3500-4700

464 mg L⁻¹ CaCO₃) and pH in BC amended reactors, suggesting that BC provided a strong 465 buffering capacity. Ambaye et al. (2020) [135] investigated the role of BC from sewage 466 sludge on the AD of fruit waste. They observed that the BC addition enhanced methane 467 production and VFAs degradation. Further, the deficiency of some nutrients or trace 468 elements in the substrate may cause an increase of VFAs inducing inhibition of microbial 469 activity during AD [33,135]. Extra doses of trace metals may be required for an effective 470 activity of methanogens without VFAs accumulation [136]. Thereby, it could be 471 speculated that BC could provide adequate concentrations of trace metals for the stability 472 of the AD system [135]. 473 In summary, BC addition can counteract VFAs inhibition in case of high loads of easy 474 degradable wastes such as FW, OFMSW, primary sludge. The alkaline nature of BC 475 determining its pH buffering capacity may contribute to prevent VFAs inhibition. The 476 ash fraction of BC contains AAEMs possibly contributing to its acid-buffering capacity 477 and important trace elements for microorganisms. However, further investigations are 478 required to confirm the effectiveness of these mechanisms and to identify the optimal 479 dose of BC related to these properties. Higher doses can be toxic for AD [24,134], perhaps 480 due to inhibitory concentrations of alkali metals [137]. Besides, other mechanisms could 481 be responsible of BC role in alleviating VFAs inhibition. Porous BC could provide 482 support for biofilm growth and protection to selectively enriched functional microbes 483 closely attached to it under acid stress [48]. Secondly, BC could promote the activity of 484 microbial partners enhancing the syntrophic VFAs degradation and methane production 485 under high organic loads [38,138]. The latter microbial mechanisms will be fully addressed

486 in section 4.3.

488 Table 3. Results of pH drop and CH₄ production/yield of AD processes in the presence 489 of various types of BCs

Pyr	olysis O _l	perational co	ondition		AD Ope	rational con	ditions		Results of buffering effect					
Temp	Time	Reactor	Feedstock	Temp	HRTor	Substrate	inoculum	Total alk	Total alkalinity pH			Increase		
(C ₀)	(min)			(C°)	duration (d)			(g L ⁻¹ Ca	aCO ₃)			in CH ₄ (%)		
								Control	BC	Control	BC			
350	180	Muffle furnace	dairy manure	20	35	dairy manure	Anaerobic sludge	1.8-4.8	5.3- 7.3	7.18	7.71	26.47ª	[27]	
350	180	Muffle furnace	dairy manure	35	35	dairy manure	Anaerobic sludge	1.8-4.8	5.3- 7.3	7.09	7.68	24.90a	[27]	
350	180	Muffle furnace	dairy manure	55	35	dairy manure	Anaerobic sludge	1.8-4.8	5.3- 7.3	7.12	7.55	24.69a	[27]	
500	120	Tube furnace	vermi- compost	35	50	chicken manure	Anaerobic sludge	6.0	7.3	7.0-7.1	7.8- 8.0	N.P.	[39]	
500	120	Tube furnace	vermi- compost	35	50	kitchen waste	Anaerobic sludge	0.0	2.3	3.5-3.7	4.7- 4.9	N.P.	[39]	
600	120	Kiln	corn stover	55	15	primary sludge	Anaerobic sludge	2.8	3.5- 4.7	7.2	7.6- 8.0	8.6- 17.8 ^a	[134]	
350	15		sewage sludge digestate	37		fruit waste	Anerobic sludge	N.P.	N.P.	4.5-5	7-8	13-27ª	[135]	
550	15		sewage sludge digestate	37		fruit waste	Anerobic sludge	N.P.	N.P.	4.5-5.5	7- 8.6	12-22ª	[135]	
500	120	Slow pyrolysis	rice straw	35	25	Municipal solid waste	Anaerobic sludge			7.8	7.5	11.69ª	[139]	

a) CH4 Yield; N.P: Not provided.

490

491

492 4.2. Adsorption of inhibitors

493 Inhibition is often defined as the predominant cause of reduction of bio-methane yields 494 and instability of AD processes. A substance can be identified as "inhibitor" when it 495 determines an adverse shift in the microbial population or arrests of the bacterial growth 496 [137]. There are direct inhibitors, such as metals (Cu²⁺, Zn²⁺, Cr³⁺, Cd, Ni, Pb⁴⁺, Hg²⁺, 497 Na⁺, K⁺, Mg²⁺, Ca²⁺, Al³⁺), organic compounds (chlorophenols, halogenated aliphatics, 498 pesticides, antibiotics, lignocellulose hydrolysate); and indirect inhibitors, such as VFAs, 499 long-chain fatty acids, hydrogen, ammonium and sulphides [19]. Among the inhibitors, 500 ammonia (NH₃ or free ammonia nitrogen, FAN) and ammonium (NH₄⁺) concentrations 501 are considered crucial (ammonia and ammonium are together considered as total 502 ammonia nitrogen, TAN). The optimal control of ammonia and ammonium may ensure

504 system. However, excess FAN/TAN concentrations are major causes of AD failure 505 [140,141]. Chen et al. (2008) [137] reported a wide range of TAN concentrations 506 (between 1.7 and 14 g L⁻¹) causing a 50 % reduction in methane production. FAN 507 concentration is considered the main cause of inhibitions for methanogens [140,142] and 508 it was reported to have a toxic effect to anaerobes ranging from 150 to 1200 mg L⁻¹ 509 [143,144]. Various mitigation strategies have been investigated to effectively counteract 510 TAN/FAN inhibitory effects on AD. Some were based on the removal and 511 immobilization of the inhibitor, as: dilution, co-digestion and microbial adaptation [145– 512 147]; struvite precipitation [148]; use of a microbial desalinisation cell [149]; ammonia 513 stripping [150]. An alternative approach is based on the use of adsorbents, inorganic as 514 zeolites [142] and organic as GAC [143] and BC [41]. 515 Based on several studies focusing on BC impacts on AD and ammonia inhibition (Table 516 4), it seems that BC could effectively mitigate ammonia inhibition, resulting in reduced 517 lag phase and enhanced methane production respect to control reactors. Some authors 518 [33,34,151] promoted BC use in AD of sewer sludge. Mumme et al. (2014) [41] revealed 519 that BC from pyrolysis of paper sludge and wheat husks could limit mild ammonia 520 inhibition (2.1 g TAN kg-1). Su et al. (2019) [43] stated that the addition of BC may 521 alleviate inhibition in case of up to 1500 mg L⁻¹ ammonia-N in food waste AD [152]. Lü 522 et al. (2016) [44] revealed that BC can support AD under high ammonium stress (up to 7 523 g-N L⁻¹). The above-mentioned studies suggest a positive effect of BC towards ammonia 524 inhibition, however, there isn't full agreement on the mitigation mechanisms 525 hypothesized: cation exchange capacity [34,151]; chemical and/or physical adsorption 526 capacity and surface functional groups [33,34,151]; promotion of direct interspecies 527 electron transfer (DIET) [44,153]; immobilization of microorganisms [43,44].

528 Therefore, BC may contribute to ammonia mitigation by direct (cation exchange capacity, 529 adsorption, surface functionality) and/or indirect factors (DIET and immobilization of 530 microorganisms), depending on the characteristics of both BC and digested substrate, 531 along with the operating conditions of the AD process (e.g. pH and temperature). 532 Focusing on direct factors, a better understanding of the interactions between BC and 533 FAN/TAN is a key step to identify its physico-chemical properties able to maximise the 534 ammonia removal. The mechanisms for ammonia adsorption on BC from wastewater and 535 digestate have been studied in literature (Table 5), resulting in promising values of 536 ammonium adsorption capacity (up to hundreds of mg NH₄-N g⁻¹ BC). Yin et al. (2017) 537 [112] reported that physical sorption could be supported by high SSA and large porous 538 structure. However, various studies suggested that porosity and SSA may not be the 539 predominant factors in ammonium adsorption [154,155]. For instance, ion exchange may 540 occur between acidic functional groups on BC surface and ammonium [112,156,157], 541 and CEC could have a major role in enhancing the ammonium adsorption capacity of BC 542 [157]. Zhang et al. (2014) [158] found that BC derived from pyrolysis of corn cob at 400 543 °C exhibited higher ammonium adsorption capacity than those produced at 600 °C due to 544 the presence of functional groups with higher acidity. Thus, the definition of the proper 545 pyrolysis temperature and the other control parameters is crucial for enhancing the 546 adsorption capacity of BC. 547 In conclusion, various studies reported the potential of BC in alleviating ammonia 548 inhibition (Tables 4 and 5) through different mechanisms. The direct adsorption of 549 ammonia on BC may contribute to the mitigation of ammonia inhibition. However, 550 further studies are required to clarify the effective role of direct mechanisms in complex 551 AD systems in presence of other phenomena, i.e. the close bond between ammonia and

552 VFAs inhibition. In some cases the direct removal of ammonium may not be the main 553 contribution to the mitigation of ammonia inhibition by BC [44,82,153]. Instead, other 554 indirect mechanisms (immobilization and faster acclimation of biomass, DIET) have been 555 suggested in literature and will be discussed in the next sections.

557 Table 4. A summary of selected works focusing on ammonia/ammonium inhibition mitigation by BC in AD processes (FAN: free ammonia 558 nitrogen; TAN: total ammonia nitrogen).

Biochar								Anaerobic d	igestion			Reference
Feedstoc k	Process: temperat ure [°C]	Dose of biochar	Substrate	Inoculum	Inoculum to substrate ratio (VS basis)	Temperatur e [°C]	рН	Critical TAN cone. or as specified [mg L ⁻¹]	Critical FAN conc. or as specifi ed [mg L-	Ammonia removal by biochar	Results and comments	
Fruitwo od	Pyrolysi s: 800- 900°C	10 g L ⁻¹	Glucose solution (6 g L ⁻¹)	Granular sludge from AD of paper mill wastewater	1:6	35	7	260 3500 7000	- - -		• ↑ Specified CH ₄ prod. rates increased by 18.6%, 10.1% and 23.5% respectively • ↓ Lag phase by 30.4%, 12.7% and 23.8% respectively	[44]
1:2 (v/v) mix: - paper sludge - wheat husks	Pyrolysi s: 500°C	6.67% w/w (2 g BC per 30 g inoculum)	-	Meso. AD (TS = 4.05%) + Ammoniu m carbonate	Only 30 g of inoculum	42	7.66 7.73 7.93 7.84 8.07	1626 2126 3126 4126 6626	- - - -	2.4 mg TAN g ⁻¹ BC 2.0 mg TAN g ⁻¹ BC 4.2 mg TAN g ⁻¹ BC 4.5 mg TAN g ⁻¹ BC 6.8 mg TAN g ⁻¹	No clear effect on biogas production was observed	[41]
Macada nia nut hells	Pyrolysi s: 350°C	33.3 g L ⁻¹	Food waste (+ water + NaHCO ₃)	AD from UASB (SS =34 g	-	Room temperatu re		1500	-	BC	• ↑ COD removal from 78% without BC to 90% with BC	[43]
Corn stover	Gasificat ion	1.82, 2.55, 3.06, 3.64 g BC g ⁻¹ TS sludge	WWTP sludge	Thermo WWTP AD	2:1	55	7.5	1100 (total N)	600 (NH ₃ - N)		CH ₄ yield, biomethanation rate constant and ↑ max CH ₄ prod. rate up to 7.0%, 8.1% and 27.6% respectively ↑ NH ₃ -N concentration by 41.5% after AD without BC, while by 0.2-18.1% with BC	[151]
Pine pellets	Gasificat ion	2.49, 4.97 g BC g ⁻¹ TS sludge	WWTP primary sludge	Meso. and thermo. WWTP AD	2:1	37, 55	7.2- 7.3	750 (total N)	400- 450 (NH ₃ - N)		• ↑ CH ₄ yield by 3.9-9.5% • ↑ NH ₃ -N concentration by 67.0% after AD without BC, while by -7.2 to 4.7% with BC	[33]

White oak pellets		2.20, 4.40 g BC g ⁻¹ TS sludge	TS = 1.25% (inoc +subs)	TS = 1.25% (inoc +subs)							• ↑ CH ₄ yield by 5.7-9.6% • ↑ NH ₃ -N concentration by 67.0% after AD without BC, while by -7.2 to 4.7% with BC	
Corn stover	Gasificat ion: 850°C	0.25, 0.375, 0.5, 1.0 g d ⁻¹	Primary sludge + WAS	WWTP AD from TPAD:	2:1	55	7.4 (sta ge	-	-	6.2-13.2%	• ↑ CH ₄ content by 13.7-25.3% • ↑ CH ₄ prod. rate by 5.5-36.9%	[34]
Pine			(TS = 7.01%)	1 stage (TS = 6.14%) 2 stage (TS = 3.77%)			1); 5.3- 6.0 (sta ge II)	-	-	3.6-11.2%	• \uparrow CH ₄ content by 0.7-9.1% • \uparrow CH ₄ prod. rate by -2.3-16.6%	
Wheat bran pellets	Pyrolysi s: 800°C	25 g L ⁻¹	Food waste fermentate	Anaerobic methanoge nic culture (from WAS	-	20	7.5		200- 250 (NH ₃ - N)		After acclimation: ↑ rate of VFAs degradation and ↓ lag-phase; No adsorption of ammonia by BC	[45]
Coppice d woodlan ds	Pyrolysi s: 500°C			AD)					,			
Orchard pruning	Pyrolysi s: 500°C											
Wheat straw	Pyrolysi s: 350, 450, 550°C	5% w/w (chicken manure)	Chicken manure (+ water)	Chicken manure meso. AD	Inoculum: 20% of WV of AD	35	Initi al: 6.7- 7.6	3450-3540 (average)	472- 600 (max)		↑ Cum. CH ₄ yield ↓ TAN compared to control	[82]
Fruit	Pyrolysi						Initi	3440-3650	421-	up to 25% TAN	• ↑ Cum. CH4 yield up to 69%	
wood	s: 350, 450, 550°C						al: 6.7- 7.0	(average)	634 (max)		 ↓ TAN compared to control ↓ FAN up to 58% compared to control 	
Air-	Pyrolysi						Initi	3880-3960	496-		• ↑ Cum. CH4 yield	
dried chicken	s: 350, 450,						al: 6.9-	(average)	701 (max)		• \ TAN compared to control	
manure	550°C						7.3		(111111)			

Table 5. Ammonium adsorption capacity of different BCs (NA: not available)

Biochar production			Biochar p	roperties	3		Adsorption			Reference
Feedstock	Pyrolysis	temperature	BET-	pН	CEC	Total pore	Source of NH ₄ ⁺ -N	Initial	NH ₄ ⁺ adsorption	_
	[°C]		SA		[cmol _c kg ⁻¹]	volume		NH4 ⁺ -N	capacity	
			$[m^2g^{-1}]$			[mL g ⁻¹]		[mg L ⁻¹]	[mg NH ₄ ⁺ -N g ⁻¹ biochar]	
Oak wood	400-450		NA	9.9	59.4 ± 8.1	NA	Ammonium solution	1000	100.9 ± 3.4	
Oak wood	-		NA	9.7	105.8 ±	NA		-	129.4 ± 34.8	
					12.1					
Greenhouse waste	-		NA	10.6	109.5 ±	NA	-	-	118.2 ± 26.9	
					21.8					
Municipal waste	-		NA	9.5	65.7 ± 16.2	NA	-	-	137.3 ± 0.6	
Presscake from	-		NA	10.3	51.0 ± 5.5	NA	-	-	105.8 ± 11.5	
AD										[154]
Oak wood	600-650		NA	10.3	76.6 ± 0.7	NA	-	-	114.4 ± 3.4	
Oak wood	-		NA	8.6	65.2 ± 20.2	NA	-	-	123.5 ± 28.7	
Greenhouse waste	-		NA	11.0	146.2 ±	NA	-	-	99.3 ± 28.5	
					32.3					
Municipal waste	-		NA	10.2	67.9 ± 12.5	NA	-	-	128.3 ± 6.7	
Presscake from	-		NA	10.1	52.6 ± 11.5	NA	-	-	136.2 ± 18.1	
AD										
Hardwood	600		147.0	9.80	NA	0.176	AD slurry	500-580	114.2	
Corncobs	-		23.0	8.92	NA	0.098	-	-	108.9	[159]
Mixed sawdust	-		6.80	8.60	NA	0.038	-	-	24.7	

Mixed wood	600	273.623	9.80	NA	0.176	Swine manure AD slurry	$1400~\mathrm{mg}~\mathrm{N}~\mathrm{L}^{1}$	44.64 ± 0.602	
Rice husk	/	10.995	7.80	NA	0.038	-	-	39.8 ± 0.54	[155]
1:2 (v/v) mix:	500	NA	NA	NA	NA	During anaerobic	1626 mg TAN kg ⁻	2.4 mg TAN g ⁻¹	
- paper sludge						digestion	1		
- wheat husks									
		-	-	-	-	-	2126 mg TAN kg ⁻	2.0 mg TAN g ⁻¹	
							1		
		-	-	-	-	-	3126 mg TAN kg ⁻	4.2 mg TAN g ⁻¹	[41]
							1		
		-	-	-	-	-	4126 mg TAN kg ⁻	4.5 mg TAN g ⁻¹	
							1		
		-	-	-	-	-	6626 mg TAN kg-	6.8 mg TAN g ⁻¹	
							1		
Fruitwood	800-900	NA	NA	NA	NA	NH ₄ Cl solution	NA	2-3 mg N g ⁻¹	[44]

563 4.3. Effects on microbial populations

564 Many authors reported the positive effects of BC on microbial populations in AD (Table 565 6). In semi-continuous and continuous AD experiments, BC addition improved methane 566 production [34] keeping it stable even with rising OLRs [29,49] and resulted in higher 567 COD removal rates [43,50,51].

568 Although there is a good agreement among the studies in terms of enhancement of AD 569 process performances (11-30 % reduction of lag phases and 11-50 % increments of 570 methane production rates) (Table 6), there is not a common hypothesis to explain BC 571 influence on microbial populations, and different possible mechanisms are proposed as 572 follows.

573 4.3.1. Microbial attachment and acclimation

574 Immobilization and acclimation of balanced microbial consortia on various support media 575 are commonly adopted for counteracting various drawbacks of AD processes [160], 576 thanks to the intensification of syntrophic conversion relationships, along with a major 577 resistance to inhibition effects [161]. Wang et al. (2018) [50] found that BC acted as an 578 inert core for microbial aggregation, resulting in a higher microbial growth rate and 579 accelerating sludge granulation. Cooney et al. 2016 [53] investigated the possibility of 580 accelerating biofilm formation by adding BC during the start-up of a packed bed 581 anaerobic digester at pilot scale. In a relatively short time, the AD system reached stable 582 and good performances, suggesting the rapid development of biofilms rich in active 583 methanogens.

584 High SSA, proper porosity structure and particle size, superficial hydrophobicity seemed 585 to be important properties favouring microbial immobilization [12,52]. Further, 586 conductive materials may act as ideal support media due to their surface hydrophobicity

587 and porous structure [54,55]. In particular, macropores can help the attachments of 588 bacterial cells [19,162,163]. Lü et al. (2016) [44] explored the influence of different 589 particle sizes of BC (2-5 mm, 0.5-1 mm, 75-150 µm) on the microbial distribution during 590 the AD of glucose under ammonium stress. They stated that bacteria could access more 591 easily fine particles than coarse particles. As a consequence, the attachment and 592 colonization of microbial populations on BC can limit the risks of wash-out, accelerate 593 the acclimation of microbes during substrate-induced inhibition, reduce the distance 594 between syntrophic bacteria and methanogens, facilitate interspecies electron transfer and 595 exchanges of VFAs or other metabolites [16,36,137]. Li et al. (2018) [46] found that 596 methanogens survived under acidic stress in presence of BC during co-digestion of FW 597 and WAS. Further, the immobilization of microbes could significantly attenuate ammonia 598 inhibition [43,44]. The colonization of porous materials by microbes can alter the 599 dominant species, making them more resistant and more rapidly acclimatized to substrate-600 induced inhibition [29,36,164]. Magnetic BC favoured the enrichment of acido/acetogens 601 and methanogens absorbed on its surface shortening the microbial contact distance, thus 602 VFAs produced by acido/acetogen bacteria could be more quickly transported to 603 methanogens than in control digesters during AD of OFMSW in batch tests [16,139]. The 604 distance of less than 1 µm has been reported to be essential for the oxidation of VFAs and 605 hydrogen production [19,165].

606 4.3.2. Selective colonization of functional microbes

607 The effect of BC addition on microbial communities was investigated with the aim of 608 identifying the most abundant bacterial and archaeal populations and, indirectly, the main 609 interspecies interactions (Table 6). In detail, most studies focused on: biofilm formation 610 [26,48,53]; shifts of microbial populations [28,34,46,47]; selective enrichment of

611 microbial -DIET partners [38,44,49,50]; promotion of DIET [38,44–46,49,51,139]. The 612 relevant anaerobic bacteria and archaea enriched in BC amended digesters are reported 613 in Table 7, together with BC properties, substrate and inoculum used in AD tests. Many 614 bacterial species were found in reactors supplemented with BCs, none of them identified 615 as more recurrent. Among archaeae, most studies identified *methanosaeta*, 616 *methanosarcina*, *methanobacterium*, and *methanolinea* species in BC amended reactors. 617 Different studies [44,48,51] investigated the spatial distribution of bacteria and archaea 618 by dividing sludge samples into different fractions, from suspended to attached to BC. Lü 619 et al. (2016) [44] postulated an explanation for the spatial distribution of methanogens 620 into BC pores by their cell morphology and dimension. The short fibrous form of 621 Methanosaeta (0.8-7 μm in size) could explain its attachment into internal and external 622 pores, while the long fibrous form of Methanobacterium (1.2-120 μm in length) could 623 limit its penetration into BC pores [44,166].

624 4.3.3. Promotion of syntrophic metabolisms

- 625 Many studies (Table 7) suggested that BC addition may improve electron transfer 626 mechanisms between anaerobic bacteria and archaea closely attached to BC surface. The 627 overall AD efficiency depends on effective syntrophic interactions between bacteria and 628 methanogens exchanging electrons to satisfy their energy requirements [56], happening 629 through various routes:
- 630 Indirect interspecies electron transfer (IIET) via soluble (i.e. hydrogen, formate,
- acetate) [167,168] and insoluble (humic substances) [169,170] compounds;
- 632 Direct interspecies electron transfer (DIET) via electrical conductive pili, membrane-
- bound electron transport proteins, and conductive materials (i.e. magnetite, biochar,
- 634 granular activated carbon, carbon cloth) [171–173].

635 In IIET hydrogen and formate operate as electron shuttles between syntrophic-producing 636 bacteria and consuming-methanogens [56]. Diffusion regulates the transfer of a 637 metabolite between microorganisms, as defined by Fick's Law [174]: the shorter the 638 distance, the higher the flux of metabolites between microbes. Thereby, when cells 639 aggregate the rate of interspecies hydrogen transfer is enhanced by the moment anaerobic 640 bacteria and methanogenic archaea form compact structures acting as an organ [174,175]. 641 However, the diffusion of soluble metabolites is considered a relatively slow mechanism 642 of energy and information transfer [173] and hydrogen IET is regarded as a bottleneck in 643 methane production [171]. On the contrary, DIET consists in the formation of an electric 644 current between electron-donating and electron-acceptor microorganisms without the 645 mediation of electron shuttles [56]. DIET could be faster and more specific compared to 646 IIET [173,176]. Park et al. (2018) [171] reported that direct exchange of electrons via 647 conductive pili was observed in co-cultures between geobacter metalliriducens and 648 geobacter sulfurreducens [176], methanosaeta harundinacea [177], or methanosarcina 649 barkeri [178]. Conductive materials such as magnetite [179], GAC [180], and BC [57] 650 were shown to effectively mediate DIET between syntrophic partners. 651 Based on changes in the microbial community composition, many studies justified the 652 enhancement of AD activity by means of the improvement of hydrogen and formate 653 interspecies transfer mechanisms or, more frequently, by DIET via conductive biochar 654 (Table 7). These findings are usually based on indirect observations, i.e. the enrichment 655 of bacterial and archaeal species able to participate to DIET function as potential partners. 656 Martinez et al. (2018) [47] found an enrichment of homoacetogenic bacteria, as 657 Clostridium, Eubacterium and Syntrophomonas, and H₂ using methanogens through the 658 analysis of microbial communities in digesting WAS and orange peels with BC,

659 suggesting the formation of co-cultures enhancing methane production. Zhao et al. (2016) 660 [51] observed the selective enrichment on BC of Geobacter and Methanosaeta during 661 AD of synthetic wastewater with butyrate and propionate in UASB reactors. They 662 suggested that butyrate and propionate could be degraded via DIET in the presence of a 663 conductive material, and they found abundance of Syntrophomonas and Smithella, 664 concluding that the metabolism via interspecies H₂ transfer for butyrate and propionate 665 degradation was probably present. Wang et al. (2018) [50] showed that the microbial 666 community analysis in a BC amended reactor during AD of synthetic wastewater resulted 667 in the selective enrichment of potential DIET-partners, as Geobacter and Bacteroidetes, 668 as well as archaea Methanosaeta and Methanosarcina. They suggested that BC could 669 enhance DIET among electrogenic microbes and archaea, improving the electron transfer 670 characteristics of granular sludge, as well as COD removal and methane yield. Wang et 671 al. (2018) [38] found that the addition of different BC doses increased methane production 672 rate and shortened the lag phase during mesophilic AD of dewatered WAS and FW. 673 According to the authors, BC counteracted the pH decrease due to VFAs accumulation 674 through its buffering capacity, and it seemed to promote DIET. 675 Aside from physical properties as SSA and porosity, favouring microbial colonization, 676 other chemical and conductive characteristics of BC may be crucial in the promotion of 677 electrons transfer. The role of electrical conductivity (EC) of BC in AD has been 678 investigated in literature [59]. EC of digestate has been reported to increase in presence of 679 BC [50,151]. However, the EC of digestate seemed unrelated to the conductivity of BC, 680 which varies depending on the metabolism and composition of microbial species [56]. 681 The capability of BC in promoting DIET appeared to be comparable to that of GAC, even 682 if the EC of BC was roughly 1000 times lower [57,171]. Barua and Dhar (2017) [181]

683 reported that multi-species aggregates from anaerobic digesters exhibited conductivity 684 ranging 0.2-36.7 μS cm⁻¹, suggesting its relation with DIET via conductive pili. Martins 685 et al. (2018) [56] stated that conductive materials could have a similar role of humic 686 substances in DIET, acting as electron shuttles by receiving and donating electrons. Wang 687 et al. (2019) [49] suggested that BC from sawdust may act as a temporary electron 688 acceptor for VFAs oxidation during thermophilic AD. They found a significantly higher 689 and more stable methane yield at higher OLR values during anaerobic co-digestion in 690 semi-continuous mode with the addition of BC from sawdust, while they did not show 691 any enhancement of AD with the addition of BC from sewage sludge. The main effects 692 of sawdust BC seemed to be the enhancements of microbial activities and syntrophic 693 oxidation of VFAs. The EC of both BCs was similar, suggesting it was not a determinant 694 factor in the promotion of syntrophic oxidation of VFAs. Instead, the authors speculated 695 that the presence of redox-active phenazine structures in the BC from sawdust could 696 promote VFAs degradation via DIET. Thereby, for better investigating whether BC from 697 sawdust could substitute hydrogen as electron acceptor in syntrophic oxidation of VFAs, 698 they conducted a series of batch experiments with butyrate or propionate as substrates in 699 which methanogenesis was inhibited. The control reactors did not show degradation of 700 butyrate and propionate, while the addition of BC stimulated their oxidation as well as 701 the production of acetate, supporting the hypothesis of the electron-accepting capacity of 702 BC in the syntrophic process. 703 The adoption of a further control in AD tests amended with a non-conductive material

703 The adoption of a further control in AD tests amended with a non-conductive material 704 can be crucial to investigate whether the stimulatory effects of BC on methane production 705 can be more closely linked to its physical properties (i.e. SSA and porosity) rather than 706 its electrical properties [56]. This was evident in the study of Cruz Viggi et al. (2017)

707 [45], where they introduced two controls without BC and with non-conductive silica sand 708 for AD of FW. They found that VFAs degradation and methane production were faster 709 in the case of BC amended reactors than both the control reactors, suggesting the 710 predominant influence of the electrical properties of BCs.

711 Table 6. A summary of selected works focusing on positive effects of BC on microbial populations in AD processes

Biochar		Anaerobic di	igestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	
k		biochar			re	mode			
					[°C]				
Sawdust	Pyrolysis:	10 g L ⁻¹	Food waste	WAS	55°C	Batch	By ↑ OLR:	• ↑ buffer capacity	[46]
	• 500°C		(TS = 94.2 g)	$(TS = 89.1 \text{ g L}^{-})$		V = 120 mL	• ↓ lag time	Microbial attachment and acclimation on BC	
	• 1 h		L-1)	1)		S/I (VS) = 0.25	• ↑ CH ₄ prod. rate	• Promotion of DIET	
	• 20°C min ⁻					- 3		Selective succession of microbes	
	1								
Pine	Pyrolysis:	8.3 - 33.3 g	Food waste	WWTP-sludge	35°C	Batch	1st phase:	1st phase:	[26]
sawdust	• 650°C	L-1	(bread)	(heated at		V = 100 mL	• ↓ lag phase	• Promotion biofilm formation	
	• 20 min		(TS = 61.2%)	95°C)		TPAD	• Faster VFAs generation	Providing temporary nutrients	
							• ↑ H ₂ prod. rate	Buffering pH by BC	
							• ↑ Cum. H ₂ prod.		
							2 nd phase:	2 nd phase:	
							• ↓ lag phase (41-45%)	Promotion methanogenic biofilm formation	
							Faster VFA degradation	• No buffering pH by BC	
							• ↑ CH4 prod. rate		
Macadam	Pyrolysis:	33.3 g L ⁻¹	Food waste +	AD from	Room	CIC reactor	• ↑ COD removal by 15%	• ↓ NH ₃ inhibition by BC	[43]
ia nut	• 350°C		water + (↑ N-		temperatur	V = 4.6 L		• ↑ Alkalinity by BC	
shells	• 2 h		NH4 conc.)		e	HRT=16 h		• ↓ VFAs accumulation	

Biochar		Anaero	bic di	gestion						References
Feedstoc	Production	Dose	of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	-
k		biochar				re	mode			
						[°C]				
	• 10°C min ⁻								• ↑ Selective enrichment of functional microbes	
	1								Microbial attachment and acclimation on BC	
					$(SS = 34 \text{ g L}^{-1})$					
Sawdust	Pyrolysis:	2 - 15 g	g L-1	Dewatered	Meso. AD	35°C	Batch	• ↓ lag phase by 27.5–64.4%	Buffer capacity by BC (alleviate ↓ pH due to ↑ VFAs)	[38]
	• 500°C			WAS	from brewery		Serum bottles	• ↑ Max CH4 prod. rate	$\bullet\uparrow$ DIET by BC (temporary electron acceptor, due to the	
	• 1.5 h			+ Food waste	factory		WV = 90 mL	by 22.4%–40.3%	richness of surface functional groups)	
	• 10°C min-			(TS = 8.8%)	(TS = 6.8%)		S/I = 0.75 - 3		• Selective enrichment of microbial DIET-partners by BC	
	1			+ water			(VS)			
Sawdust	Pyrolysis:	15 g L	1	Food waste	Brewery Meso.	55°C	Semi-	• ↑ CH ₄ yield by	• ↑ VFAs syntrophic oxidation by BC	[49]
	• 500°C			+ Sewage	AD		continuous	16.0%-55.2%	• High SA of BC \rightarrow microbial attachment (?), closer	
				sludge	(TS = 5.2%)		Serum bottles	• Stable CH4 prod. at ↑	association of syntrophic partners	
				(TS = 9.2%)			WV = 150 mL	OLRs	\bullet Electroactive functional groups of BC \Rightarrow stimulation of	
				FW/Sludge =			$\downarrow \text{HRT}, \uparrow \text{OLR}$		DIET (?)	
				4:1 (TS)						
Corn	Gasificatio	0.25 - 1	.0 g	Primary	WWTP AD	55°C	Semi-	• ↑ % CH ₄ by 13.7-25.3%	Adsorption and precipitation of CO ₂ by BC	[34,182]
stover	n:	d-1		sludge +			continuous	• ↑ CH ₄ prod. rate by 5.5-	• ↓ TAN	
	• 850°C	(2 nd pha	ase)	WAS			V = 500 mL	36.9%	• ↑ Alkalinity	
				(TS = 7.01%)			TPAD:		• ↑ Macro-/micro-nutrients in digestate	
									Shift in bacterial community	

Biochar		Anaerobic	digestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	
k		biochar			re	mode			
					[°C]				
				TPAD:		1) HRT = 5, 15		• Inhibition at high BC doses	
				1 ph. (TS =		d			
				6.14%)		2) HRT = 13 -			
				2 ph. (TS =		30 d			
				3.77%)					
Pine	-	-	-	-	-	-	• ↑ % CH ₄ by 0.7-9.1%		
							• ↑ CH ₄ prod. rate by 2.3-		
							16.6%		
Corn	Gasificatio	1.82 -3.64	WWTP	Thermo.	55°C	Batch	• ↑ % CH ₄ (88.5-96.7%)	• ↑ buffer capacity	[151]
stover	n	g BC g ⁻¹	sludge	WWTP AD		V = 600 mL	• ↑ CH ₄ yield	• ↑ electrical conductivity in digester (+37%)	
		TS sludge				26 days		→ ? ↑ extracellular electron transfer	
								• ↓ NH ₃ inhibition	
								• CO ₂ sequestration by BC	
								• ↑ macro-/micro-nutrients in digestate	
Rice	Gasificatio	1%, 3%	Cattle	-	35°C	Batch	• ↑ Biogas prod. by 31% by	• ? Microbial biofilm formation on BC	[30]
husks	n:	w/w	manure +			V = 1400 mL	1% BC		
	• 900-		water				• ↓ % CH4 by 7% by 1% BC		
	1000°C		(TS = 5%)				• No benefits by 3% BC		

Biochar		Anaerobic d	igestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	
k		biochar			re	mode			
					[°C]				
Rice	Gasificatio	1%, 3%	Cattle	-	25-30°C	Semi-	• ↑ Biogas prod. by 4-5% by	• ? Not enough time for biofilm formation	[31]
husks	n:	w/w	manure +			continuous	BC		
	• 900-		water			WV = 12 L	• ↓ % CH ₄		
	1000°C		(Mix: TS =			HRT = 20 days			
			5%)						
Dry dairy	Pyrolysis:	0, 1, 10 g	Dry dairy	Inoculum from	20°C	Batch:	• ↑ CH ₄ yield up to 26.5%	• No biofilm formation on BC (?)	[27]
manure	• 350°C	L-1	manure	lagoon, fed		Serum bottles	• ↓ lag phase	• No effects on DIET by BC (?)	
	• 3 h			with dried		V = 280 mL	• ↓ total VFAs	• ↑ Alkalinity and ↑ pH (buffer capacity)	
	• 10°C min-			manure		S/I = 1 (VS)			
	1			(TS = 115.85 g)					
				L-1)					
-	-	-	-	-	35°C	-	• ↑ CH ₄ yield up to 24.9%	• ↑ Alkalinity and ↑ pH (buffer capacity)	
							• ↓ lag phase		
							• ↓ total VFAs		
-	-	-	-	-	55°C	-	• ↑ CH ₄ yield up to 24.7%	• ↑ Alkalinity and ↑ pH (buffer capacity)	
							• ↓ lag phase		
							• ↓ total VFAs		
Vineyard	Pyrolysis:	10, 30 g L ⁻¹	Co-digestion:	WWTP AD	37°C	Batch:	Batch:	Promotion of synthrophic metabolism by BC	[47,183]
prunings	• 550°C					• V = 250 mL	• \ lag phase	Adsorption of inhibitors	

	Anaerobic	digestion						References
Production	Dose o	of Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	
	biochar			re	mode			
				[°C]				
		• Orange	$(TS = 35.5 \text{ g kg}^{-})$			• ↑ CH4 prod.	Adhesion and growth of microorganisms by BC SA	
		peels	1)		Semi-	Semi-continuous:		
		(TS = 311g)			continuous	• ↑ CH ₄ prod.		
		kg-1)			• V = 3 L			
		• Sewage			• HRT = 10-30			
		sludge			days			
		(TS = 28.7g						
		kg ⁻¹)						
Pyrolysis:	1:1 (TS)	Citrus peel	WWTP AD	35°C	Batch	• ↓ lag phase	Limonene adsorption by BC	[40]
• 450°C		(TS = 16.6%)	(TS = 11.0 %)		V=500 mL	• ↑ Cum. CH ₄ prod.		
					S/I = 0.31 - 0.33			
					(VS)			
-	1:1 (TS)	-	-	-	-	• ↓ lag phase	• Limonene adsorption by BC	
						• ↑ Cum. CH ₄ prod.	Adhesion and growth of microbes on BC	
-	1:1; 1:2	2; -	-	-	-	• ↓ lag phase	• Limonene adsorption by BC	
	1:3; 2:	1				• ↑ Cum. CH ₄ prod.	Adhesion and growth of microbes on BC	
	,	•				1 F	B · · · · ·	
	Pyrolysis: • 450°C	Pyrolysis: 1:1 (TS) • 450°C - 1:1 (TS)	• Orange peels (TS = 311g kg ⁻¹) • Sewage sludge (TS = 28.7g kg ⁻¹) Pyrolysis: 1:1 (TS) Citrus peel • 450°C (TS = 16.6%)	Production Dose of Substrate Inoculum biochar • Orange (TS = 35.5 g kg peels 1) (TS = 311g kg 1) • Sewage sludge (TS = 28.7g kg 1) Pyrolysis: 1:1 (TS) Citrus peel WWTP AD • 450 °C (TS = 16.6%) (TS = 11.0 %)	Production Dose of Substrate Inoculum Temperatu biochar re [°C] • Orange (TS = 35.5 g kg peels 1) (TS = 311g kg 1) • Sewage sludge (TS = 28.7g kg 1) • Yerolysis: 1:1 (TS) Citrus peel WWTP AD 35°C • 450°C (TS = 16.6%) (TS = 11.0 %)	Production Dose of biochar Substrate biochar Inoculum Temperatu re mode Experimental mode • Orange (TS = 35.5 g kg²) • Semicontinuous • V = 3 L • HRT = 10-30 • Kg²) • Sewage sludge • HRT = 10-30 • HRT = 10-30 • Kg²¹) • WWTP AD 35°C Batch • 450°C (TS = 16.6%) (TS = 11.0 %) V=500 mL • S/I = 0.31 - 0.33 (VS) - 1:1 (TS) - - - 1:1; 1:2; - - -	Production Dose of biochar Substrate biochar Inoculum Temperatu re mode Experimental mode Results • Orange (TS = 35.5 g kg² pcels ¹) • Orange (TS = 35.5 g kg² pcels ¹) • ↑ ↑ ℂH₄ prod. • ↑ ↑ ℂH₄ prod. • (TS = 311g kg²) • ○ Semi-continuous • ↑ ℂH₄ prod. • ∨ 3 L • → ↑ ℂH₄ prod. • Sewage sludge (TS = 28.7g kg²) • HRT = 10-30 days • → ↑ ℂum. ℂH₄ prod. • 450°C (TS = 16.6%) (TS = 11.0 %) V=500 mL v↑ ↑ ℂum. ℂH₄ prod. • √ Cum. ℂH₄ prod. • ↑ ↑ ℂum. ℂH₄ prod. • ↑ ↑ ℂum. ℂH₄ prod. • ↑ ↑ ℂum. ℂH₄ prod. • ↑ ↑ ℂum. ℂH₄ prod. • ↑ ↑ ℂum. ℂH₄ prod.	Production Dose of Substrate Inoculum Temperatu Experimental Results Possible mechanisms

Biochar		Anaerobic di	gestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	=
k		biochar			re	mode			
					[°C]				
Rice	In FeCl ₃	0.5% w/w	OFMSW +	OFMSW	35°C	Batch	• ↓ lag phase	• No effects on NH ₃ by BC	[139]
straw +	solution:		water	thermo. AD +		Serum bottles	• ↑ CH ₄ prod. by 11.69%	• No effects on pH by BC	
FeCl ₃	• 2h		(TS = 1.64%)	water		V = 500 mL,		• ↑ syntrophic associations of bacteria on BC	
(3.2 g	Carbonizati			(TS = 2.19%)		S/I = 1 (VS)		• (?) DIET by high conductivity of BC	
FeCl ₃ :10	on:								
0 g rice-	• 500°C								
straw)	• 2h								
Rice	Carbonizati	-	-	-	-	-	• ↓ lag phase	• No effects on NH ₃ by BC	
straw	on:						• ↓ CH ₄ prod.	• No effects on pH by BC	
	• 500°C								
	• 2h								
Cow	Pyrolysis:	2 – 14 g L ⁻¹	Dry beer lees	Meso. WWTP	35°C	Batch	• ↓ lag phase	• Promotion of DIET by BC conductive properties (?)	[28]
manure	• 500°C		(TS = 62.5%)	AD		WV = 150 mL	• ↑ Max cum. CH4 prod. up	• Microbial colonisation and biofilm formation on BC (?)	
	• 4 h			(TS = 36.7%)		TS = 25%	to 82.9%	• ↑ Alkalinity and ↑ pH (buffer capacity)	
	• 100°C h ⁻¹					S/I = 3 (TS)		• Selective enrichment of methanogens by BC	
-	-	-	-	Thermo.	55°C	-	• ↓ lag phase	-	
				WWTP AD			• ↑ Max cum. CH4 prod. up		
				(TS = 38.5%)			to 47.2%		
Rice	Pyrolysis:	4 g L ⁻¹	Synthetic	WWTP sludge	35°C	UASB	UASB:	No effects on pH by BC	[50]
straw	• 500°C		wastewater				• ↑ COD removal rate	BC inert core for microbial aggregation	

Biochar		Anaero	obic di	gestion						References
Feedstoc	Production	Dose	of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	-
k		biocha	r			re	mode			
						[°C]				
					$(SS = 25.2 \text{ g L}^{-})$		• Continuous	• ↑ VFAs degradation	Selective enrichment of microbial DIET-partners by BC	
							mode	• ↑ gran. sludge		
							• V = 5500 mL	conductivity and quality		
							• HRT = 12 h,			
							12-6 h	Batch:		
								• ↓ lag phase by 28.6%		
							Batch	• ↑ biogas yield		
							• Serum bottles	• ↑ %CH4		
							• V = 550 mL			
-	Biochar	-		Grease trap	-	37°C	Biochar packed	• COD removal:68%	• ↑ Methanogenic biofilm communities on BC	[53]
				waste			bed anaerobic	• Total VFAs: from 4.7	• BS as packing material for growth and retention of biofilm	
				wastewater			digester	(feed) to 1.46 g L ⁻¹		
				(TSS = 1.04 g)			• V: 900 L	(effluent)		
				L-1)			+1500 L+ 1500	• %CH ₄ > 60%		
							L	• Start-up: 59 days		
							• ↓ HRT (3.1 –			
							1.8 days)			

Biochar		Anaerobic d	igestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	
k		biochar			re	mode			
					[°C]				
Wheat	Pyrolysis:	25 g L ⁻¹	Food waste	Methanogenic	20°C	Batch	After acclimation:	• No effects on pH via BC	[45]
bran	• 800°C		fermentate	culture (from		V = 120 mL	• ↑ rate of VFAs	• No effects on NH3 via BC	
pellets	• 3h			WAS digestate)			degradation	• Promotion of IET by BC	
							• ↓ lag-phase		
Coppiced	Pyrolysis:	-	-	-	-	-		-	
woodland	• 500°C								
s									
Orchard	Pyrolysis:	-	-	-	-	-		-	
pruning	• 500°C								
Fruitwoo	Pyrolysis:	10 g L ⁻¹	Glucose	AD gran.	35°C	Batch	• ↓ lag phase by 5.9-23.9%	• NOT NH ₄ adsorption	[44]
d	• 800-		(6 g L ⁻¹)			Serum bottles	• ↑max CH4 prod. rate by	• NOT ↑ buffer capacity	
	900°C						23.5-47.1%	DIET promotion via BC	
							Faster VFAs degradation	Affinity of methanogens with BC	

Biochar		Anaero	bic di	igestion						References
Feedstoc	Production	Dose	of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	=
k		biochai	r			re	mode			
						[°C]				
					Ammonium:					
					0.26, 3.5, 7 g-N					
					L-1					
Fruit	Pyrolysis:	10 g L	-1	Nutrient	Meso. pulp	35°C	Batch	• ↓ lag phase by 11.4-30.3%	Selective colonization of functional microbes by BC	[48]
wood	• 800°C			solution	sewage		Serum bottles	• ↑ Max CH ₄ prod. rate by	• Not ↑ buffer capacity	
				+ glucose	digestate		V = 500 mL	5.2-86.6%	• ? Biofilm growth on BC	
				(2 - 8 g L ⁻¹)	(1 g VS L ⁻¹)			• ↑ VFAs production and	• ? Promotion of DIET or Hydrogen IET by BC	
								degradation		
Pine	Pyrolysis:	5 g L ⁻¹		Synthetic	WAS digestate	37°C	UASB	• ↑ CH ₄ prod. rate by 25%	• ↑ Butyrate degradation via DIET in UASB via BC	[51,57]
wood	• 600°C			wastewater	(TSS = 13.1 g)		WV = 1000 mL	• ↑ COD removal	• Selective enrichment of microbial DIET-partners by BC	
	• 2h			with butyrate						
				Synthetic				• ↑ CH ₄ prod. rate by 16%	• ↑ Propionate degradation via DIET in UASB via BC	
				wastewater				• ↑ COD removal	• Selective enrichment of microbial DIET-partners by BC	
				with						
				propionate						

Biochar		Anaerobic d	igestion						References
Feedstoc	Production	Dose of	Substrate	Inoculum	Temperatu	Experimental	Results	Possible mechanisms	-
k		biochar			re	mode			
					[°C]				
-	Biochar	2.5 g L ⁻¹	Synthetic	WAS digestate	37°C	UASB	• ↑ COD removal: ≥ 93%	• (?) Promotion syntrophic metabolism via DIET with BC	[29]
			wastewater	(TSS = 13.1 g)		WV = 1000 mL	(control: 75-83%)	in UASB reactors	
			with ethanol			↓ HRT (24 - 8	• ↑ CH ₄ prod. rate		
						h)			

CIC: controlling internal circulation; COD: chemical oxygen demand; DIET: direct interspecies electron transfer; FW: food waste; HRT: hydraulic retention time; IET: interspecies electron transfer; OLR: organic loading rate; OFMSW: organic fraction of municipal solid waste; SA: surface area; S/I: substrate to inoculum ratio; SMP: specific methane potential; SS: suspended solids; TPAD: two phased anaerobic digestion; TSS: total suspended solids; UASB: up-flow anaerobic sludge blanket; VFA: volatile fatty acid; WAS: waste activated sludge; WWTP: wastewater treatment plant; WV: working volume; V: volume.

714 Table 7. A summary of studies reporting the selective enrichment of bacteria and archaea by BC addition during AD processes

Biochar production		Biochar prope	rties		Anaerobic diges	tion				Reference
Feedstock	Pyrolysis temperature	BET-SSA	Electri	Total pore	Temperature	Substrate	Identification technique	Enriched Bacteria	Enriched Archaea	•
	[°C]	$[m^2g^{-1}]$	cal	volume	[°C]					
			condu	[cm ³ g ⁻¹]						
			ctivity							
			[S m							
			1]							
Wheat bran pellets	800	55 ± 1	49.9	0.0445	20	Food waste fermentate	FISH- CLSM		More Methanosarcina-like	[45]
									Archaea rather than	
									Methanosaeta like Archaea	
Coppiced	500	61 ± 1	1.6	0.0483						
woodlands										
Orchard pruning	500	13.7 ± 0.5	0.5	0.0165						
Macadam	350	12.7	-	-	Room	Food waste + water	16S rRNA sequencing	Bacteroidales	Methanoregulaceae	[43]
ia nut shells	330	12.7			temperature	rood waste - water	105 He W sequencing	Anaerolineales	Methanotrichaceae	[45]
ia nut snens					temperature			Syntrophobacterales	Methanobacteriaceae	
	000 000					a.	460 PM			5443
Fruitwoods	800-900	-	-	-	35	Glucose	16S rRNA sequencing	Enterobacteriaceae	Methanobacterium	[44]
									Methanosaeta	
									Methanosarcina	
Fruit wood	800	-	-	-	35	Glucose	16S rRNA sequencing	Syntrophomonas	Methanobacterium	[48]
								Clostridium	Methanosaeta	
								Clostridiaceae	Methanosarcina	
Rice straw + FeCl ₃	500	51.14	-	0.0328	35	OFMSW + water	16S rRNA sequencing	Bacteroides	Methanosarcina	[139]
(3.2 g FeCl ₃ :100 g								Clostridiaceae	Methanobacterium	
rice-straw)								Porphyromonadaceae	OTU in Methanomicrobiales	
								Moraxellaceae	Methanosaeta	
Rice straw	500	111.5	-	-	35	Synthetic wastewater	16S rRNA sequencing	Bacteroidetes unclassified (23.65%)	Methanosaeta (77.18%)	[50]
								Bacteroidales unclassified (9.19%)	Methanosarcina (11.65%)	

												T (C 420/)		
												Treponema (6.43%)		
												Smithella (5.56%)		
												Brooklawnia (5.45%)		
												Geobacter (5.42%)		
Sawdust	500		248.6		-	-	35		Dewatered WAS +	16S rRNA sequence	ing	Anaerolineaceae	Methanosaeta	[38]
									food waste + water			Porphyromonadaceae	Methanobacterium	
													Methanolinea	
Cow manure	500		112.6		-	0.0156	35		Dry beer lees	16S rRNA sequence	ing	Chloroflexi	Methanospirillum	[28]
												Bacteroides	Methanosarcina	
												Proteobacteria	Methanolinea	
Vineyard prunings	550		240 ± 4.8	8	-	-	37		Orange peels	16S rRNA sequence	ing	Bellilinea	Methanosaeta	[47]
												Trepomena	Thermogymnomonas	
												Clostridium	Methanolinea	
												Petrimonas	Methanofollis	
												Proteiniphilum		
												Bacteroides		
Vineyard prunings		550		240				37	Sewage sludge	16S	rRNA	Clostridium	Methanosaeta	
			± 4.8							sequencing		Longilinea	Methanolinea	
												Curvibacter	Thermogymnomonas	
												Eubacterium	Methanobacterium	
												Syntrophomonas		
Pinewood		600		209				37	Synthetic wastewater	16S	rRNA	Attached sludge:	Attached sludge:	[51]
					.33 μS				with butyrate	sequencing		Geobacter	Methanosaeta	
					cm ⁻¹							Thermanaerovibrio	Methanosarcina	
					cm ⁻¹							Thermanaerovibrio Syntrophomonas	Methanosarcina Methanospirillum	
Pinewood	600		209		cm ⁻¹	-	37		Synthetic wastewater	16S rRNA sequenc	ing			
Dinewood	600		209			-	37		Synthetic wastewater with propionate	16S rRNA sequenc	ing	Syntrophomonas	Methanospirillum	

								Syntrophus	Methanosphaerula	
Biochar	-	-	-	-	37	Grease trap waste	16S rRNA sequencing	Attached to BC:	Attached to BC:	[53]
						wastewater		Aminobacterium	Methanobacterium	
								Syntrophomonas	Methanosarcina	
								Sporanaerobacter	Methanobrevibacter	
								Escherichia		
Sawdust	500	248 ± 34	-	-	55	Food waste	16S rRNA sequencing	Coprothermobacter (36.3%)	Methanosaeta (43.9%)	[46]
								Fervidobacterium (20.0%)	Methanosarcina (15.8%)	
								Syntrophothermus		
								Treponema		
Corn stover	850	315.2	-	0.09	55	Primary sludge +	16S rRNA sequencing	Firmicutes (Clostridia)	Methanothermobacter (>90%)	[34]
						WAS		Bacteroidetes	Methanosarcina	
								Proteobacteria		
Pine		353.1	-	0.23						
Sawdust	500	248.6 ± 9.4	0.11	-	55	Food waste	16S rRNA sequencing	Defluviitoga	Methanothermobacter	[49]
			μS			+ Sewage sludge		Tepidimicrobium	Methanosarcina	
			cm ⁻¹							

NA: not available; CLSM: confocal laser scanning microscopy; FISH: Fluorescence in situ hybridization; OFMSW: organic fraction of municipal solid waste; SSA: surface area; WAS: waste activated sludge.

717 4.4. Effect of biochar on digestate quality

Anaerobic digestate has been considered as soil improver because it is rich in nutrients [184–186]. However, challenges related to digestate management have recently grown in association with EU regulations on ammonia, volatile organic acids, phenolic compounds, heavy metals, PAHs and PCBs [4,187]. So far, most technologies available to exploit digestate as soil improver are based on mechanical/physical (e.g. mechanical dewatering, semipermeable membranes and evaporation) or chemical processes (e.g. ammonia stripping and nutrients adjustment) (Table 8).

Table 8. Advantages and disadvantages of conventional technologies for digestate processing [188–191].

Methods	Advantages	Disadvantages
Solid-liquid separation	Rich in phosphorus and suitable for	Liquid fraction has been considered
	fertilizer application	as a surface and ground water
		pollutions
Belt and drum dryers	treatment and upgrading of digestate	Rapid volatilization of ammonium
	to a solid or concentrated product	causes severe ammonia emission
High-Tech technologies (e.g.	Production of several streams with	Expensive
ammonia stripping, membrane	different physical and chemical	
process and vacuum evaporation	properties.	
exist)		
Mixing solid digestate with desired	Increasing value per ton	-
nutrient		

Dehydrated digestate could be used in non-agricultural markets as a heating fuel, however, this use implies nutrient and metal losses, which in turn have negative impacts

on the environment and crops. Consequently, three approaches have been conventionally applied to reduce diffuse pollution resulting from digestate application to land [19,188]:

- 733 Nutrient recovery from digestate;
- 734 Carbon to Nitrogen ratio adjustment;
- 735 Increasing of nutrient retention capacity using an additive.

One of the key aspects that should be considered when supplying additives to AD process is their effects on the quality of digestate for subsequent uses, especially as soil improver. Despite literature has not explored enough the fate and properties of digestate with BC to land applications, some potential benefits of BC amendment can be identified as follows. BC remaining in digestate after AD acts as a nutrient retention improver and catalyst, mitigates leaching of heavy metals and pollutants via physical and chemical absorption of organics, phosphate, ammonium, nitrate, nitrite, metals and CO₂ [19,192]. The improvements on digestate quality can be related to BC features as SSA, surface functional groups, ash content and presence of metals (Table 9).

Table 9. Biochar properties able to improve digestate quality

Factors	Function	Literature	Reference
	Higher content of functional	HCl, NH ₃ ·H ₂ O and KMnO ₄ modification	[193]
	group (e.g. Si-O-Si, O-H and	were performed to obtain functional biochar	
Textural	C=O)	from Swine manure digestate.	
properties	Increase of the pH and cation	H ₂ O ₂ and KOH modification were	[104]
(surface area, pore	exchange capacity	performed to obtain functional biochar from	
size)		domestic sewage sludge digestate	
Ash content	Alkali and alkaline earth metals	Pinewood and white oak biochars made	[33]
	increase the alkalinity	digestate a great alternative to agricultural	
		lime fertilizer.	

Metals on the	The cation form of the metal can	Ex:	[151]
surface	bind with soluble phosphorus	$3\text{Fe}(\text{OH})_2 + 2\text{H}_3\text{PO}_4^{3-} \rightarrow \text{Fe}_3(\text{PO}_4)_2 + 6\text{H}_2\text{O}$	
Functional group	Negatively charged functional	Manganese oxide-modified biochar	[194]
	groups forms complexes with	composite derived from corn straws	
	heavy metals	improved heavy metal stabilization in the	
		digestate	

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BC supplement in digesters may be beneficial to the fertilizer value of digestates. Shen et al. [34,151] have found that BC addition can cause a substantial increase of the macroand micro-nutrients as N, P, K, Ca, Mg, and Fe in digestate. Zhang et al. (2020) [195] reported a similar rise of nutrient content in digestate amended with BC, even if concentrations of certain nutrients did not fully meet limits of EU regulation on fertilizing products (Regulation EU 2019/1009). Research has shown that the joint amendment of BC and anaerobic digestate can reduce atmospheric greenhouses emissions from fields, such as N₂O [196] and CO₂ [197–199]. A mixture of dried anaerobic digestate and BC may be an alternative to standard formulations in horticultural potting media [200]. In addition, BC may contribute to improve the composting of digestate [153,201,202], particularly in terms of process performance, compost quality and its benefits on plants [203–205]. Further research is needed to understand interactions between BC, digestate and soil for the potential use of the mixture as soil improver after AD. Future studies about the use of BC as additive in AD should also consider its effects on the agronomic value of anaerobic sludge (i.e. content in macro- and micro-nutrients, germination and phytotoxicity tests, and others [206]).

765 4.5. Effects on biogas upgrading

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766 Raw biogas from AD consists mainly of CH₄ (50-70 % v/v) and CO₂ (30-50 % v/v), along with minor compounds as water vapour, H₂S, NH₃, O₂ and N₂ [207]. Upgrading and 768 cleaning biogas are required to meet the requirements and standards for engines and 769 pipelines, although they represent energetic and economic costs up to 55 % of the total 770 biomethane production cost [33,208,209]. To date, conventional technologies involve water scrubbing, cryogenic separation, physico-chemical absorption, and membranes, 772 among the others [210,211]. Recently, BC has been investigated as adsorbent of CO₂ and H₂S for in-situ and ex-situ applications, as addressed in the following paragraphs.

774 4.5.1. In-situ biogas upgrading

A series of studies [24,33,34,151] investigated the feasibility of in-situ biogas upgrading by the addition of BC, obtaining methane contents up to pipeline quality (Table 10). Shen et al. (2015) [151] investigated the possibility of sequestering CO₂ with BC during thermophilic AD of WAS for in-situ biogas cleaning and upgrading. They reported average methane contents of 88.5-96.7 % in BC amended reactors, compared to 67.9 % in control reactor, reaching CO₂ removals of 54.9-86.3 % and residual H₂S content below 5 ppb. They suggested that CO₂ removal could be promoted by the high porosity of BC, by the large SSA rich of basic sites and of hydrophobic sites. Shen et al. (2016) [33] assessed the AD of WAS with the addition of two woody BCs. They observed average methane content up to 92.3 % and 79.0 % in biogas from BCs amended reactors in mesophilic and thermophilic conditions compared to control reactors, corresponding to CO₂ removals by up to 66.2 % and 32.4 %, respectively. They stated that both BCs owned desirable properties for CO₂ sequestration, and in particular high values of: SSA, porosity,

chemical stability, degree of carbonization and alkaline nature. Linville et al. (2017) [24] investigated the influence of particle size and dose of BC from walnut shell on AD of FW in mesophilic and thermophilic conditions. They found higher CO₂ removals compared to control reactors in the case of smaller particle size of BC (61.0 %) than coarse one (51.0 %), due to the larger SSA and ash content. However, according to other studies [33,151], they observed a reduction of methane production with higher BC doses, concluding that this could lead to inhibition caused by higher concentrations of cations released by the BC. Shen et al. (2017) [34] studied the effects of two BCs from corn stover and pine wood on AD of WAS in two-stage digesters. They reported an average methane content of 81.0-88.6 % in the reactor with BC from corn stover and 72.1-76.6 % with BC from pine wood, compared to around 70.0 % for the control. They stated that BCs would release base cations sequestrating CO₂ by chemical sorption and forming bicarbonate/carbonate salts, and that its surface structure would help CO₂ adsorption. Apart from CO₂ adsorption on BC, the major formation of CH₄ depends on the stronger syntrophic cooperation between organic acid-oxidising bacteria and CO₂ reducing methanogens [36,58], underlying the key role of efficient interspecies electron transfers. Further confirmation by other authors to the attractive findings about biogas in-situ upgrading by BC would be beneficial.

806 4.5.2. Ex-situ biogas cleaning and upgrading

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The use of BC and other carbonaceous adsorbents for CO₂ capture from various gaseous streams recently attracted a growing attention [212–214]. Considering ex-situ applications of BC for biogas upgrading and cleaning, Table 11 shows CO₂ and H₂S adsorption capacity of different BCs. Clearly, the adsorption capacity for both CO₂ and H₂S is in a wide range, 0.4-2.3 mmol g⁻¹ and 0.2-19.1 mmol g⁻¹ respectively. Most of the

studies regarding CO₂ capture do not specifically focus on biogas, investigating different BCs eventually subjected to activation. Sethupathi et al. (2017) [215] assessed the adsorption of CH₄, CO₂ and H₂S in a synthetic biogas stream by four BCs in fixed bed adsorbers during continuous experiments. They reported that just CO₂ and H₂S were captured by BCs, which exhibited adsorption capacities up to 0.208 mmol g⁻¹ for H₂S and $0.126 \text{ mmol g}^{-1}$ for CO_2 . Creamer et al. (2014) [216] investigated the adsorption of CO_2 into BCs from bagasse and hickory wood. They found that BC could effectively capture CO₂ (adsorption capacity up to 73.55 mg g⁻¹ or 1.67 mmol g⁻¹), suggesting the importance of surface area and superficial nitrogen groups in CO₂ sequestration, mainly through physical adsorption. Creamer and Gao (2016) [217] reported that the main mechanism for CO₂ sequestration by BC is physical adsorption, suggesting the importance of high SSA [100], adequate pore size (0.5-0.8 nm) [218] and pore volume, thanks to Van der Waals and electrostatic forces. However, the adsorption of CO₂ can also be influenced by BC chemical properties, such as the presence of basic surface functional groups or alkali and alkaline earth metals, hydrophobicity and non-polarity [214]. For instance, Xu et al. (2016) [219] found that the adsorption of CO₂ by three BCs during batch equilibrium tests was due to the presence of alkali and alkaline earth metals (Ca, Fe, K, Mg) by CO₂ mineralogical reactions together with physical sorption. Activation and surface treatments of BCs can provide high surface area and micropores for physical sorption and enrich surface functional groups and metal oxides for chemical sorption, leading to remarkable CO₂ adsorption capacities (5.0-7.4 mmol g⁻¹) [220]. Other studies applied BC for removing H₂S from biogas (Table 11). Sahota et al. 2018 [221] found 84.2 % removal of H₂S from biogas with BC from leaf waste. Kanjanarong et al. (2017) [222] obtained a removal of 98 % of H₂S (8.02 mmol g⁻¹) from biogas with

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BC, suggesting carboxylic and hydroxide radical groups as responsible of H₂S adsorption. Finally, Pelaez-Samaniego et al. (2018) [37] found that BC from AD digestate could effectively remove H₂S from a synthetic biogas, possibly facilitated by the presence of ash, porosity, or aromatics in BC. In contrast with CO₂ for which adsorption onto BC seemed to be mainly physical, absorption of H₂S seemed to involve many chemical mechanisms with BC surface [223].

Overall, biochar seems to be a promising adsorbent for ex-situ biogas cleaning and upgrading applications. However, additional studies [215] should focus on the adsorption of CO₂ and H₂S along with NH₃ from real or synthetic biogas, considering their competitive adsorptions, along with the influence of water vapour and the eventual removal of CH₄.

Table 10. Physic-chemical properties of biochars used in biogas in-situ upgrading

Feedstock	Production	BET-SA	Total pore	Average pore	H/C	O/C	Ash	Reference
	temperature	$[m^2g^{-1}]$	volume	diameter	[molar ratio]	[molar ratio]	[% wt]	
	[°C]		$[cm^3g^{-1}]$	[nm]				
Corn stover	Gasification	315.30	0.09	6.50	0.075 ± 0.007	0.004 ± 0.001	45.18 ± 0.40	[34,151]
Pine pellets	Gasification	310.19	0.19	5.07	0.078 ± 0.009	0.249 ± 0.014	18.69 ± 0.44	[33,34]
White oak	Gasification	296.81	0.15	4.92	0.109 ± 0.026	0.051 ± 0.010	34.90 ± 0.65	
Walnut shell	Gasification: 900 °C	86.5	0.16	7.06	0.20 ± 0.01	0.06 ± 0.01	43.2 ± 0.2	[24]

Table 11. Biochar adsorption capacity (mmol g⁻¹) of H₂S and CO₂ during ex-situ applications.

Feedstock	Pyrolysis/activati	Surfac	Total	Gas	Inlet H ₂ S or	H ₂ S	CO ₂	Referen
	on	e area	pore		CO_2	Adsorpti	Adsorpti	ce
		$[m^2g^-]$	volum		concentrati	on	on	
		¹]	e		on	capacity	capacity	
			[cm ³ g ⁻		[ppm]	[mmol g ⁻	[mmol g ⁻	
			1]			1]	1]	

AD	PY/500°C/60	134	0.037	Syntheti	2000	3.96 a		[37]
digestate fibres	min	134	0.037	c biogas	2000	3.90		[37]
110140	PY/600°C/60min	142	0.035	Syntheti c biogas	2000	15.90 a		
	PY/600°C/60min +Na ₂ CO ₃	-	-	Syntheti c biogas	2000	19.13 a		
Biomass	C/450°C/6min	60		N ₂ +H ₂ S	1000	2.14 a		[224]
from black liquor	C/ 150 C/OHIII	00		1121125	1000	2.11		[221]
Pig manure	PY/500°C/4hour s	47.4		Air + H ₂ S	1% (v/v)	1.75 a		[225]
Sewage sludge	PY/500°C/4hour s	71.6		Air + H ₂ S	1% (v/v)	1.29 a		
Potato peel waste	C/500°C/5min	63		N ₂ + H ₂ S	1000	1.56 a		[226]
Camphor	PY/400°C/5hour	20		1125		3.21 a		[227]
Rice hull	s PY/400°C/5hour	115				11.23 a		
Bamboo	s PY/400°C/5hour	58				9.88 a		
80% wood chips 20% AD	PY/600°C			Biogas	1020	8.02 a		[222]
digestate Perilla	PY/700°C	473.4	0.1	C414:		0.537	2.312	[215]
leaf				Syntheti c biogas				[215]
Korean oak	PY/400°C	270.8	0.1	Syntheti c biogas		0.178	0.597	
Japanese oak	PY/500°C	475.6	0.2	Syntheti c biogas		0.167	0.379	
Soybean stover	PY/700°C	420.3	0.2	Syntheti c biogas		0.308	0.707	
Sawdust	G/850°C	182.0 4	0.003	N ₂ /CO ₂			1.08 b	[228]
	G/850°C + monoethanolami	3.17	0.007	N ₂ /CO ₂			1.02 b	
Sugarcane	ne PY/600°C	401.0					1.67 ^b	[216]
bagasse Whitewoo	PY/500°C	840	0.55	He/CO ₂	30 % mol		1.34 b	[229]
d	+ steam activation	0.10	0.55	115, 502	20.31101			[==/]
Whitewoo d	PY/500°C + CO ₂ activation	820	0.45	He/CO ₂	30 % mol		1.43 b	
Whitewoo d	PY/500°C + KOH	1400	0.62	He/CO ₂	30 % mol		1.77 ^b	
Walnut	activation C900°C/1.5hours	397	0.198				1.65 b	[230]
shell Pig	PY/500°C/4hour	31.57	0.044	N ₂ /CO ₂			0.53 b	[219]
manure Wheat	s PY/500°C/4hour	20.20	0.041	N ₂ /CO ₂			0.78 b	
straw Sewage	s PY/500°C/4hour	10.12	0.022	N ₂ /CO ₂			0.41 b	
sludge	S							

5. Economic and environmental assessments

An economic and environmental evaluation of BC production and application in AD was performed, considering specifically four perspectives: 1. BC production according to feedstock composition and 2. to pyrolysis process conditions; 3. BC application in AD compared to current state of the art technologies addressing AD challenges; 4. integration of AD and pyrolysis processes (to our knowledge, specific studies related to the economic and environmental assessment of BC use in AD processes are not available).

5.1. Economic assessment

Considering BC production, the key parameters to evaluate the economic benefits of feedstocks (perspective 1) are: ash and lignin contents and O/C ratio, which affect pyrolysis yield, molecular weight of bio-oil and BC amount production [231]. Li et al. (2017) [232], based on regression model applied to 346 lignocellulosic feedstocks, stated that higher ash content increases BC production in a range of 12.5-15.5 %, reducing bio-oil production, and consequently the incomes coming out from bio-oil trade, which makes pyrolysis economically profitable. With lower ash content and higher O/C ratio of feedstocks, higher biofuel yields are produced, which leads to better economic performance, in fact minimum fuel selling prices for lignocellulosic feedstocks ranged from 0.53-1.1 Euro/L.

Considering the pyrolysis process (perspective 2), the investment costs for BC production from lignocellulosic feedstocks are: 43 % for pre-treatment and pyrolysis, 35 % for H2 generation and 22 % for cooling and fractionations [232]. The average operating cost varies from 0.68 Euro/L for woody biomass to 0.86 Euro/L for straw biomass, due to the higher costs of disposal and pre-treatment of straw biomass, respectively 32 % and 34 %

of total operational costs [233]. Based on the economic analysis performed by Harsono et al. (2013) [234] and Sahoo et al. (2019) [235], the investment and operational costs related to BC production from lignocellulosic feedstock can only be balanced by a BC trade price of 470 Euro/t.

Considering perspective 3, the need to enhance AD feasibility and applicability to unconventional substrates implied higher costs, which should be exceeded by the increased methane production and therefore by the additional electric energy potentially associated. Inorganic and biological additives as iron, micronutrients and ash are conventionally employed to reduce inhibition and facilitate organic matter solubilisation, thus improving methane production. Nevertheless, the application of additives in AD accounts as 3.60-4.10 euro/L of enzyme and as 13-16 euro/L of nutrients [236]. The overall costs of BC, depending on feedstock, pyrolysis process and activating agent (Table 12), could range from 0.2 to 0.5 USD/kg, making BC cheaper than granular activated carbon (GAC), which has production costs ranging between 0.6 and 20 USD/kg.

Table 12. Comparison of granular activated carbon (GAC) and biochar (BC) production costs depending on feedstock, production process and activating agent

Additive	Feedstock	Production process/ Activating agent		Cost of production (USD kg ⁻¹)	Reference
GAC	Poultry litter derived carbon	Steam	USA	1.44	[237]
GAC	Rice bran	NaOH	China	3.58- 3.77	[237]
GAC	Red oak wood	Steam	USA	0.62- 1.27	[237]
GAC - commercial	-	-	USA	1.19- 16.34	[237]
GAC	Fruit processing waste	Steam	Malaysia	1.67	[237]
GAC	Rice bran	CO_2	Brazil	3.54	[237]

GAC	Acid treated rice	CO_2	Brazil	20.45	[237]
GAC	Wood	КОН	-	2.49	[237]
GAC	Charcoal	КОН	-	1.25	[237]
GAC	Lignite	КОН	-	2.18	[237]
GAC - commercial				1.93	[237]
ВС	Empty fruit bunches	Slow PY	Malaysia	0.533	[234]
BC	Straw	Slow PY (large scale)	UK	0.203	[18]
BC	Straw	Slow PY (medium scale)	UK	0.447	[18]
BC	Straw	Slow PY (small scale)	UK	0.351	[18]
ВС	Short rotation coppicing, forestry residues, short rotation forestry	Slow PY (large scale)	UK	0.266	[18]
ВС	Short rotation coppicing, forestry residues, short rotation forestry	Slow PY (medium scale)	UK	0.500	[18]
ВС	Short rotation coppicing	Slow PY (small scale)	UK	0.434	[18]
BC	Forestry residue	Slow PY (large scale)	UK	0.345	[18]
BC	Forestry residue	Slow PY (medium scale)	UK	0.584	[18]
ВС	Arboricultural arisings	Slow PY (small scale)	UK	0.213	[18]
BC	Pine wood	Slow PY	USA	0.220- 0.280	[238]
ВС	-	-	European market (2014)	0.662- 0.811 ^a	[239]
BC	-	-	-	0.207	[239]
BC	-	-	-	0.600	[68]
ВС	-	-	Global, market price (mean)	2.06	[240]
ВС	-	-	USA, market price (mean)	2.48 (0.08-13.48)	[241]

GAC: granular activated carbon; PY: pyrolysis.
(a) exchange rate USD/euro (January 2014): 1.3516

893 Still considering perspective 3 and moving from the additive's cost to the improvement 894 of AD performances, 895 The economic benefits of the integration of AD and pyrolysis technologies (perspective 896 4) has been explored by literature. The integrated technologies of AD of waste biomass 897 and pyrolysis of the digestate could increase the net electricity production respect to AD 898 alone [80] and enhance its quality as soil amendment [81] with economic and 899 environmental benefits [242]. However, up to date, there is uncertainty regarding the 900 balance between input costs of BC supplementation and output of energy production from 901 AD. Qiu et al. (2019) [59] proposed the energy input-output LCA method to estimate the 902 total energy input for biogas production, by calculating the energy associated with each 903 component used in AD, and the actual energy return of investment. Zhang et al. (2020) 904 [243] investigated the addition of woody BC to improve the thermophilic AD of FW. 905 They concluded that BC supplementation could be economically feasible to enhance 906 thermophilic AD of FW.

5.2. Environmental assessment

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Considering feedstocks (perspective 1), life cycle analysis (LCA) from cradle to cradle of the pyrolysis of lignocellulosic feedstocks stated that GHG emissions for husk/shell/pit ranged from 120-250 g CO_2 eq/MJ, while for other organic waste, wood and straw they ranged between 20-50 g CO_2 eq/MJ [232]. This difference was due to the dominant contribution of indirect land use change from food production. Feedstocks having higher O/C ratio and 0-2 % ash content increased the GHG emissions [244]. Based on GREET database [245], GHG emissions reduction for lignocellulosic feedstocks was 85 - 98 % compared to the GHG emissions for petroleum fuels refining, which was equal to 93

916 gCO₂eq/MJ. GHG emission reductions for lignocellulosic feedstocks satisfy the 50 % 917 share of renewable fuel standard for GHG emission reduction requirement for advanced 918 fuels [246]. 919 Considering BC production process (perspective 2), LCA from cradle to gate showed that 920 BC produced from palm oil empty fruit bunches through slow pyrolysis had an energy 921 content higher than the energy required for producing BC [234]. Furthermore, LCA 922 cradle-to grave proved the positive energy balance of BC produced from different 923 lignocellulosic materials [247]. Hence, the positive energy balance of BC production and 924 application, due to the high-energy content of lignocellulosic feedstocks, represents a 925 crucial benefit both from economic and environmental perspectives. One of the main 926 issue of BC produced from renewable feedstocks as lignocellulose wastes (i.e. WAS, 927 wood) and not from purpose grown feedstocks is the higher risk of having contaminants 928 as heavy metals or organic compounds like dioxins, PAHs and PCBs [248]. 929 Still considering BC process production (perspective 2), pyrolysis and gasification were 930 compared from the perspective of carbon equivalent abatement (CA): pyrolysis achieved 931 the highest CA, ranging from 0.07 to 1.25 t CO₂ eq/t feedstock, as cardboard and wood 932 waste, while gasification reached the highest electricity generation outputs with 0.9 933 MWhe/t of feedstocks [249]. Activation was proven to represent a high item cost both in 934 terms of economic and environmental perspectives [250]. BC chemical activation costs, 935 in line with non-renewable GAC activation, range between 1.38 and 1.48 Euro/kg, 936 respectively with acidic and basic treatments [251]. 937 However, perspective 2 could also be explored considering the existing literature related 938 to the conventional applications of BC as soil improver, adsorbent for water and air 939 pollutants, catalyst for syngas upgrading and biodiesel production (Table 13). LCA

studies from cradle-to-grave of conventional BC applications measured in all cases positive environmental benefits compared to conventional perspectives. BC produced from waste forestry feedstock and applied as soil improver contributed to GHGs emission reduction up to 2.74 kg CO₂ eq/ kg BC for the impact categories climate change (CC), natural gas avoided for fossil depletion (FD) and urea avoided for freshwater eutrophication (FE) and human toxicity (HT) [252]. A LCA from cradle to cradle [247] showed that BC production and application as adsorbent presented, compared to GAC deriving from virgin non-renewable feedstock, lower environmental impacts in terms of climate change (CC), fossil depletion (FD), freshwater eutrophication (FE) and terrestrial acidification (TA) impact categories, and that wood and wood chips achieved the highest environmental credits among the others feedstocks. In details, considering CC (expressed as kg CO₂/kg adsorption material), GAC produced 1.44, while wood chips and corn stover had credits ranging between -3.42 to -3.57, whereas oil palm produced 11.1 [253]. Regarding FE (evaluated as kg P eq/ kg adsorption material), BC from lignocellulosic feedstock exhibited higher values than GAC, respectively ranging between 6.2 to 10.9 % [247], due to the organic matter of renewable feedstock [254]. Considering both TA and FD (respectively estimated as kg SO₂ eq/ kg adsorption material and kg oil eq/ kg adsorption material), BC from lignocellulosic feedstock achieved higher credits than GAC, between 9.5 % and 32.0 %. Considering BC application in AD processes (perspective 3), to our knowledge the available scientific literature only focused on sequential processes, as pyrolysis followed by AD. A LCA from cradle to grave [255] considered BC produced from corn stover applied as carburant and soil amendment with sequential AD, achieving respectively: -

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2.47 kg CO₂eq/t and energy saving of -6.53 MJ/t for the first scenario and - 4.67 kg CO₂eq/t and energy saving of -9.73 MJ/t for the second scenario.

Table 13. Environmental assessment of biochar (BC) production and uses

	Positive effects	Negative effects
Feedstock (perspective 1)	Lignocellulosic biomass	Feedstock provision (transport) [93]
	exhibited more positive effects	Potential presence of contaminants
	because of higher energy	(heavy metals, persistent organic
	potential [247]	pollutants) in waste feedstocks
	Waste biomasses presented	(sewage sludge) [256]
	more advantages (e.g. avoided	
	waste management) [256]	
Production process	Renewable energy from syngas	Large centralized pyrolysis units
(perspective 2)	and bio-oil [93]	higher transportation distances
	Stabilization of C in biomass	compared to smaller decentralized
	feedstock [18]	units [256]
Activation process		Activation was proven to represent a
(perspective 2)		high item cost both in terms of
		economic and environmental
		perspectives [250].
		Use of chemicals and electricity is
		associated to higher impacts [257].
Applications		
Soil improver	Biochar (C) sequestration in the	Binding /deactivation of pesticides,
	soil [93]	herbicides and nutrients in soil [258]
	Reduced fertiliser requirements	BC as source of potential toxicants
	[93]	(heavy metals, PAHs, organics) [258]
	Reduced N ₂ O emissions from	
	soil [93]	
	Enhanced plants growth [18]	
	Reduced fossil fuel use in	
	irrigation and cultivation [18]	
	Enhanced nutrient availability	
	[258]	
	Increased H ₂ O retention [258]	
	Reduced leaching and run-off of	
	nutrients [258]	
Adsorbent	BC lower impacts than activated	
	carbon [247]	

In conclusion, focusing the environmental assessment on perspective 4, the integration of

AD of waste biomass and pyrolysis of digestate, as shown in Figure 2, could increase the

net electricity production respect to AD alone [80] and enhance its quality as soil amendment [81] with economic and environmental benefits [242]. In conclusion, the integrated approach has been investigated by life cycle analysis and exhibited positive environmental outcomes if compared with non-integrated processes [255,259].

6. Conclusions

976 This review addressed three key issues related to the comprehension of BC role in AD

1. Investigation of the influence of BC properties on AD performances and of their ability

977 processes:

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recovery as soil improver.

- 979 to counteract its main challenges. It is understood that BC properties are determined 980 by the feedstock and by pyrolysis and activation processes. The key features were SSA, porous structure and distribution, nature of surface functional groups (related to 982 CEC and adsorption capacity, buffer capacity, ability to immobilize microbial 983 communities), elemental composition and ash content. However, some mechanisms 984 (e.g. BC role in ammonium adsorption and BC influence on microbial mechanisms) 985 still need to be fully understood and explained. Other challenges for future research 986 are related to digestate management; in detail, the influence of BC relatively high doses on the rheological properties of the digestate should be explored, as well as the
 - 2. Assessment of the optimal BC production chain (i.e. feedstock-pyrolysis-activation) to achieve the desired features. Lignocellulosic biomasses, slow pyrolysis and physical activation seemed to be a good combination in general, while other feedstocks and/or chemical activation should be evaluated for specific needs and tailor-made applications.

eventual leaching of pollutants in the environment as a consequence of digestate

- However, a systematic investigation of the correlations linking BC physico-chemical characteristics and AD performances, carefully exploring one by one the BC effects mentioned in this review, is highly needed for a deep understanding of BC role as additive in AD processes.
- 3. Evaluation of the economic and environmental advantages connected to BC use in AD processes, compared to conventional solutions applied to address AD challenges. The main research gap related to this issue is the absence of specific literature related to BC use in AD processes, therefore only general statements could be formulated. The use of BC as additive could be cheaper and has less environmental impacts than of conventional AD improvers (e.g. physico-chemical-biological pre-treatments) and of GAC. The integration of AD and pyrolysis achieved economic feasibility and positive environmental performances if compared with non-integrated processes. Future research could investigate the optimization of technical, economic and environmental performances of BC production chain and its integration in AD processes.

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