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1 **Prediction of organic matter accessibility and complexity in anaerobic** 2 **digestates**

3

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12

13 **Abstract**

14 Further characterization to properly assess the fate of organic matter quality during
15 anaerobic digestion and organic carbon mineralization in soils is required. Organic
16 matter quality based on its accessibility and complexity was employed to successfully
17 classify 28 substrate/digestate pairs through principal components and hierarchical
18 clustering analysis. The two first components explained 58.02% of the variability and
19 four main groups were separated according to the feedstock type. A decrease in the
20 accessibility (16-66%) and an increase in the complexity (34-98%) of the most
21 accessible fractions was noticed. Besides, an increase of non-biodegradable compounds
22 (17-66%) was globally observed after anaerobic digestion. The observed trends in the
23 conversion of organic matter during anaerobic digestion have allowed to fill the gap in
24 the modeling of the anaerobic digestion process chain. Indeed, partial least squares
25 regressions have accurately predicted the organic matter quality of digestates from their
26 inputs ($R^2 = 0.831$, $Q^2 = 0.593$) although the digester operational conditions

27 (temperature and hydraulic retention time) were non-explicative enough. As a novel
28 approach, the predicted digestate quality was used to feed a partial least squares
29 regression model previously developed to predict organic carbon mineralization in soil.
30 The combined models have predicted experimental organic carbon mineralization in soil
31 ($R^2 = 0.697$) with a model quality similar to the model for organic carbon mineralization
32 in soil ($R^2 = 0.894$). This is the first study that has successfully conceived an additional
33 step in the prediction of organic matter fate from raw substrate before anaerobic
34 digestion to soil carbon mineralization.

35

36 **Abbreviations**

37 3D, three dimension; AD, anaerobic digestion; BMP, biochemical methane potential; C, carbon; C_{bio},
38 biodegradable carbon; COD, chemical oxygen demand; DOM, Dissolved Organic Matter; EPS,
39 extracellular polymeric substances; FCI, fluorescence complexity index; HRT, Hydraulic Retention Time;
40 HCA, Hierarchical Clustering Analysis; NEOM, Non-Extractable Organic Matter; NMR, nuclear
41 magnetic resonance; OM, organic matter; PCA, Principal Components Analysis; PEOM, Poorly
42 Extractable Organic Matter; PLS, Partial least squares; Q^2 , Percent of variation of Y predicted by model
43 in cross-validation, R^2 , Correlation coefficient from PLS; REOM, Readily Extractable Organic Matter;
44 RMSE, Root Mean Square Error, RMSE_{CV}, Root Mean Square Error for Cross Validation; RMSEP,
45 RMSE calculated on validation dataset; PC1, Principal Component 1; PC2, Principal Component 2; Pf(i),
46 fluorescence proportion for a zone (i); SEOM, Slowly Extractable Organic Matter; SPOM, Extractable
47 Soluble from Particulate Organic Matter; T, Temperature; TS, total solids; Vf (i), fluorescence volume for
48 a zone (i); VS, volatile solids

49

50

51 **Keywords**

52 biogas effluent, stability, fluorescence, waste characterization, soil

53

54 **1. Introduction**

55 The current waste management model has started to evolve towards more sustainable
56 and resource recovery strategies (Fonoll et al., 2016; Vidal-Antich et al., 2021).

57 Anaerobic Digestion (AD) is a biological process widely used to convert the organic

58 matter (OM) present in different wastes into methane (Fernández-Domínguez et al.,
59 2020; Vinardell et al., 2021), along with the production of both OM and nutrient-rich
60 by-product called digestate (Fernandez-Bayo et al., 2018; Guo et al., 2018). Nowadays,
61 digestates represent alternative fertilizers used in agriculture either as organic
62 amendment or fertilizer depending on the process, post-treatment and substrate type
63 (Akhiar et al., 2017; Guilayn et al., 2020). However, digestate efficiency as organic
64 amendment mainly depends on their OM stability (Kögel-Knabner, 2002), which
65 remains a topic of ongoing research.

66 A need for accurate OM characterization added to the strict limitations by legislation on
67 contaminants, such as heavy metals for the agricultural reuse of specific feedstock, are
68 points of main consideration to enhance digestate management (Khakbaz et al., 2020).
69 Digestate stability has to be properly assessed before land application (Tambone et al.,
70 2013; Maynaud et al., 2017). Digestates often acquire higher biological stability and
71 nutrient availability (particularly nitrogen) than raw substrates, enhancing the interest
72 for land use (Provenzano et al., 2011). During AD, recalcitrant compounds are
73 concentrated since the labile organic structures are preferentially degraded (Insam et al.,
74 2015). Nonetheless, this stabilization could occur due to the interaction of different
75 factors, including (i) degradation and solubilization of simple compounds, (ii) molecular
76 complexification, (iii) complex microbial-related products release from biomass growth
77 and decay (Aemig et al., 2016).

78 For a suitable agronomic valorization of digestates, evaluating the OM fate during AD
79 and soil carbon (C) mineralization is a meaningful aspect to reduce associated
80 environmental impacts (e.g. C loss, greenhouse gases emission) or favor ecosystem
81 service like soil C sequestration that contributes to climate change mitigation (Minasny
82 et al., 2017). Thus, wider efforts are needed to better understand how OM from non-

83 digested substrate is transformed into digestate OM (Shakeri Yekta et al., 2019;
84 Tambone et al., 2015).

85 Standard methods to measure the biodegradability of organic wastes through
86 biodegradability assays are laborious and time-consuming. At least, 30 and 90 days are
87 required for anaerobic biodegradation potential tests and aerobic biodegradation during
88 soil incubations, respectively (Jimenez et al., 2017). Hence, biochemical fractionation is
89 a much more time-saving method to characterize the structural nature of organic wastes
90 and assess waste biodegradability using successive chemical extractions (Teglia et al.,
91 2011a). The Soest (1963) fractionation method was widely applied to associate OM
92 composition with biodegradability (Fernandes et al., 2009; Gunaseelan, 2007; Triolo et
93 al., 2012). Based on this fractionation method, the Iroc indicator (i.e. indicator of
94 residual organic C in soils) has been developed as a proxy of the potentially remaining
95 OM after application in soils for various organic amendments (Lashermes et al., 2009).
96 Moreover, it was successfully used to predict the long term evolution of OM in soils
97 after repeated applications of these organic amendments (Levavasseur et al., 2021).
98 However, biodegradable substrates and digestates were not included in the panel of OM
99 used for the development of the Iroc indicator. In addition, the Soest fractionation
100 method presented limitations in predicting the biodegradability for a wide range of
101 organic residues using biomethane potential tests (BMP) (Mottet et al., 2010).
102 Therefore, less aggressive methods were requested as a more representative way for
103 anaerobic biodegradability prediction (Bareha et al., 2018; Mottet et al., 2013).

104 With this purpose, Jimenez et al. (2015) suggested a less aggressive fractionation
105 method coupled with 3D fluorescence on the extracted fractions. These authors
106 successfully classified sixty organic residues based on their OM quality, which is
107 defined by the OM accessibility (i.e. compounds availability to microorganisms for intra

108 or extra-cellular degradation) and complexity (i.e. molecules structure) of organic
109 waste. This methodology is a promising approach compared with single spectroscopic
110 techniques. Indeed, it has allowed an accurate assessment of the OM quality (Muller et
111 al., 2014; Zhang et al., 2019) and has defined new inputs for AD modeling approaches
112 (Jimenez et al., 2020). Interestingly, in Jimenez et al. (2017), the OM quality of 82
113 samples, comprising organic waste of different origin, was used to accurately predict
114 both BMP and potential C mineralization in soil using Partial Least Square (PLS)
115 regression. Similarly, Bareha et al. (2018) have properly evaluated the correlation
116 between substrates organic nitrogen accessibility indicators with C biodegradability
117 using an extracellular polymeric substances (EPS) fractionation method modified from
118 Jimenez et al. (2015) in PLS regressions. The accessibility and complexity of a treated
119 substrate will probably vary during AD. However, no studies have been focused before
120 on the prediction of digestate OM quality from their corresponding inputs. This
121 approach would make it possible to further predict the potential C mineralization of
122 digestates in soil as described in Jimenez et al. (2017). Therefore, models could be an
123 interesting alternative to fill this gap, saving time and providing new tools to improve
124 AD performance.

125 Feedstock typology stands as a key factor to understand the final OM quality of
126 digestates (Rocamora et al., 2020). Indeed, Fourier-transform infrared spectroscopy
127 (FTIR) analysis showed that the main spectroscopic features in digestates depend on the
128 composition of the initial biomass (Provenzano et al., 2011). Nevertheless, structural
129 changes of labile and recalcitrant fractions of OM were reported comparing the nuclear
130 magnetic resonance (NMR) spectra of different substrates and their subsequent
131 digestates (Laera et al., 2019; Shakeri Yekta et al., 2019). However, the OM quality
132 evolution due to AD has not been evaluated for a broad range of feedstock. Nonetheless,

133 it could provide relevant information for different sectors implementing AD (e.g. farms,
134 wastewater treatment plants, solid waste treatment plants).

135 This study aims to assess the OM accessibility and complexity evolution during AD for
136 a wide variety of feedstocks. To this end, 28 substrate/digestate pairs including different
137 feedstock nature were classified using statistical analyses to evaluate how both
138 feedstock type and AD parameters influenced OM quality in the final digestate. Finally,
139 a model based on linear regression was proposed to predict digestate OM quality from
140 their inputs. The predicted quality was used to feed the PLS model for C
141 biodegradability in soil proposed by Jimenez et al. (2017) to determine the digestates
142 organic C mineralization in soil after land spreading.

143

144 **2. Material and methods**

145 **2.1. Substrate/digestate pairs**

146 Twenty-eight substrate/digestate pairs (a total of 56 samples) were collected from
147 different waste treatment plants in France. The selected organic substrates, including
148 diverse types of feedstocks and origins, were anaerobically digested in laboratory or
149 full-scale reactors under different digester operational conditions (Table 1). The studied
150 samples comprised 6 digestate types from the digestate fertilizing-value typology
151 reported in Guilayn et al. (2019). Each sample was characterized in terms of
152 accessibility and complexity as assessed by the biochemical fractionation method
153 coupled with 3D fluorescence developed in Jimenez et al. (2015) and described below.
154 Biochemical fractionation was conducted on the freeze-dried particular matter and
155 fluorescence on the aqueous phase.

156 **2.2. Fractionation method**

157 Following the reported methodology in Jimenez et al. (2017), all the fresh samples were
158 centrifuged (ca. 18,600 g for 30 min) at 4°C as a first step. The aqueous phase, namely
159 dissolved organic matter (DOM), was separated. Total Solids (TS) and Volatile Solids
160 (VS) on the raw sample as in the resulting particulate phase were determined (APHA,
161 2005). Then, the particulate phase was freeze-dried and ground (1 mm) and a quantity
162 of 0.5 g of freeze-dried sample was subjected to the successive chemical extractions (30
163 mL of each extractant). After every extraction stage, the sample was centrifuged
164 (18,600 g, 20 min, 4 °C) and the liquid phase was filtered (0.45 µm) and kept for
165 analyses. All tests were carried out in duplicate. The resulting fractions from the
166 fractionation method, ordered from more to less accessible, were defined as: (1)
167 Extractable Soluble from Particulate Organic Matter (SPOM), (2) Readily Extractable
168 Organic Matter (REOM), (3) Slowly Extractable Organic Matter (SEOM), (4) Poorly
169 Extractable Organic Matter (PEOM), (5) Non-Extractable Organic Matter (NEOM).
170 Chemical oxygen demand (COD) and 3D fluorescence spectroscopy were performed on
171 the four liquid extracted fractions (DOM, SPOM, REOM, PEOM). Regarding the
172 fractionation method reproducibility, the measurement error is below 5% (data not
173 shown).

174 COD was determined on the extracts in duplicate using Aqualytic® Vario COD kits (0-
175 1500 mg O₂/L) analyzed using an ultraviolet (UV) spectrophotometer MultiDirect from
176 Aqualytic®. Samples were diluted with deionized water if required. Units were in
177 mgO₂/L and mg O₂/g TS for the liquid and solid phases, respectively.

178 **2.3. Fluorescence spectroscopy analysis**

179 The 3D fluorescence spectroscopy analyses were conducted on the obtained liquid
180 fractions using a spectrofluorimeter Perkin Elmer LS55. The excitation wavelengths

181 varied from 200 to 600 nm and the values were recorded every 0.5 nm between 200 and
 182 600 nm, with increments of 10 nm. The resulting spectra were divided into seven
 183 different zones (I-VII) corresponding to a biochemical family with common complexity
 184 level for spectra interpretation, as described by Jimenez et al. (2014) and He et al.
 185 (2011): I, protein-like (Tyrosine); II, protein-like (Tryptophane), III, protein-like
 186 (Tyrosine, Tryptophane and microbial products); IV, fulvic acid-like; V, inner filter,
 187 glycolated protein-like; VI, melanoidin-like and lignocellulose-like; VII, humic acid-
 188 like. Afterward, the obtained data were processed to calculate the fluorescence volume
 189 of each zone $V_f(i)$ and then the fluorescence proportion of each zone $P_f(i)$ as described
 190 in Eqs. (1) and (2), respectively.

$$191 \quad V_f(i) (\text{U.A./mg COD} \cdot \text{L}^{-1}) = V_{f_raw}(i) / \text{COD}_{sample} \times 1 / \frac{S(i)}{\sum_{i=1}^7 S(i)} \quad (\text{Eq.1})$$

$$192 \quad P_f(i) (\%) = V_f(i) / \sum_{i=1}^7 V_f(i) \times 100 \quad (\text{Eq.2})$$

193

194 Where:

195 $V_f(i)$ is the normalized volume of a zone i ($\text{U.A./ mg O}_2 \cdot \text{L}^{-1}$)

196 V_{f_raw} is the raw fluorescence volume of a zone i ($\text{U.A./ mg O}_2 \cdot \text{L}^{-1}$)

197 COD_{sample} is the COD concentration of the sample ($\text{mg O}_2 \cdot \text{L}^{-1}$)

198 $S(i)$ is the area of a zone i (nm^2)

199 $P_f(i)$ is the fluorescence proportion of a zone i (%)

200 Moreover, a fluorescence complexity index (FCI) was calculated between the
 201 proportions of fluorescence volumes of the most complex molecules zones (IV-VII) and
 202 the less complex molecules zones (I-III), as stated in Eq. (3) (Aemig et al., 2016).

$$203 \quad FCI = \frac{\sum_{i=4}^7 V_f(i)}{\sum_{j=1}^3 V_f(i)} \quad (\text{Eq.3})$$

204

205 **2.4. Statistical analysis**

206 Principal component analysis (PCA), Hierarchical Clustering Analysis (HCA) and their
207 corresponding plots were carried out in R (4.0.3) (R Development Core Team, 2021).
208 The displayed groups for the PCA individuals were determined by HCA clustering.
209 Both analyses were conducted on a total of 56 samples (28 substrate/digestate pairs)
210 considered individually. A Euclidean distance matrix with center-scaled variables was
211 calculated before HCA analysis and the number of groups was determined heuristically.
212 PCA was performed on center-scaled variables. A total of 34 variables for each sample
213 was evaluated. Six variables were related to the accessibility of the sample and
214 corresponded to the biochemical fractions (DOM, SPOM, REOM, SEOM, PEOM,
215 NEOM) and 28 variables were linked to the complexity of the sample (seven
216 fluorescence variables for each extracted fraction). Tukey's methodology was followed
217 for the displayed boxplots.

218 In order to determine a linear prediction model of digestate quality from substrates
219 quality, PLS regressions were performed using SIMCA software from UMETRICS. In
220 its simplest form, a linear model specifies the linear relationship between a dependent
221 (response) variable Y, and a set of X predictor variables, the X's. Cross-validation was
222 then performed to test the model's quality. K-fold cross-validation was used. The
223 database was divided into 7 blocks and some samples were selected as validation
224 samples. This step was repeated. The mean and standard deviation of the scores were
225 calculated to estimate the bias and the variance of validation performance. The
226 validation step was performed on 5 samples not used in the calibration step among the
227 28 samples, chosen as representative of each type of digestate: Sludge_2_D, PS + Cow
228 Food_D, PS + Cow Manure_D, MW4_D and CM3_D.

229 The parameters from the PLS models used to assess model quality are the following:

230 Correlation coefficient: R^2 .
231 Root Mean Square Error (RMSE): used as an accuracy measurement of differences
232 between predicted values and measured model values.
233 RMSE_CV: which is the RMSE for the cross-validation and was applied as prediction
234 model error.
235 RMSEP: RMSE calculated on the validation dataset.
236 Q^2 : percentage of variation of Y predicted by the PLS model according to cross-
237 validation. This parameter indicates how well the model predicts the data. A large Q^2
238 (>0.5) indicates good predictivity.
239 The PLS model was based on 33 variables: the fractionation percentage of COD
240 (DOM+SPOM, REOM, SEOM, PEOM, NEOM) and the fluorescence percentage of the
241 seven zones coming from each fraction. The PLS model coefficients are presented in
242 Supplementary Material. The same X-Variables as Jimenez et al. (2017) model have
243 been used. Thus, DOM and SPOM fractions have been pooled and only SPOM
244 fluorescence percentage has been considered. To combine the PLS model for C
245 biodegradability in soil from Jimenez et al. (2017) and the digestate quality PLS model,
246 experimental data obtained from soil C mineralization tests as described in Jimenez et
247 al. (2017) have been used. Fourteen digestates have been tested among the 28 samples
248 previously defined. Table 2 presents the digestates considered and their respective
249 biodegradable C percentage.

250

251 **3. Results and Discussion**

252 **3.1. Substrates and digestates classification**

253 PCA and HCA analyses were conducted on 33 variables describing the accessibility and
254 complexity of the OM for the 56 samples considered individually. Scores and loadings
255 from PCA are presented in Figure 1a and 1b, respectively. HCA clusters are illustrated
256 in Figure 1c. The PCA analysis has showed that the first two components explained
257 58.02% of the total variance, meaning that the samples were rather well described by
258 the used characterization data. These results are in accord with previous studies
259 focusing on organic waste classification (Jimenez et al., 2015) or the characterization of
260 post-treated digestates (Maynaud et al., 2017). From the HCA, four main groups were
261 identified by different colors (A, B, C, D) in PCA and HCA plots (Figure 1a and 1c).

262 The Component 1 (PC1) was described by the complexity of the molecules, from the
263 simplest samples on the right part of the loadings plot (mainly fluorescences zones II-
264 III, related to simple sugars/proteins and microbial products with low complexity) to the
265 most complex samples on the left part (mainly fluorescences zones VI-VII, linked to
266 humic acids, lignocellulose and melanoidin). Hence, PC1 was significantly (p-value <
267 0.05) correlated to simple and intermediate fluorescence zones, such as REOM_I_II,
268 SEOM_I_II and PEOM_I_IV variables ($R^2 = 0.77, 0.67$ and 0.66 , respectively).

269 Meanwhile, it was anti-correlated (p-value < 0.05) to complex fluorescence zones (VI-
270 VII), particularly REOM_I_VII, REOM_I_VI and SEOM_I_VI variables ($R^2 = -0.94, -$
271 0.90 and -0.87 , respectively).

272 The second component (PC2) was explained by the complexity added to the
273 accessibility of the sample. A significant (p-value < 0.05) and positive correlation was
274 observed between REOM_C ($R^2 = 0.74$) and fluorescence zones from I to III
275 (SPOM_I_III, REOM_I_I and SEOM_I_I, with $R^2 = 0.78, 0.67$ and 0.65 , respectively).

276 Contrarily, a significant (p-value < 0.05) and negative correlation was defined between
277 PEOM_C ($R^2 = -0.79$) and fluorescence zones of intermediate complexity as

278 REOM_I_IV, SEOM_I_IV, SEOM_I_V ($R^2 = -0.95, -0.85, \text{ and } -0.85$, respectively).

279 Biochemical fractionation variables were among the main variance contributors for PC2

280 which allowed to identify the most accessible samples on the top (essentially due to

281 REOM) from the less accessible samples (mainly PEOM) on the bottom. DOM fraction

282 was not correlated with SPOM/REOM/SEOM nor to PC1 but to a lesser extent ($R^2 = -$

283 0.59 and $p\text{-value} < 0.05$). SEOM was positively correlated with the most accessible

284 fractions (SPOM and REOM) while PEOM was negatively correlated with REOM and

285 SEOM, as previously stated by Jimenez et al. (2015) and Aemig et al. (2016). Zhang et

286 al. (2019) also showed that SEOM fraction shared protein-like compounds fluorescent

287 peaks with SPOM and REOM fractions whereas PEOM presented a humic-like

288 compound peak as the main peak. Indeed, NMR spectroscopy on the solid fraction

289 showed that SEOM was mainly composed of proteins (Laera et al., 2019). NEOM

290 fraction was non-explicative enough in this study.

291 Figure 1c shows the four groups established from the HCA analysis based on the

292 extracted fractions and their complexity of each sample: (A) pig slurry and slurry

293 mixtures with primary sludge, agro-industrial waste or biowaste, (B) manure, fibers and

294 municipal solid waste, (C) pig slurry mixtures with fiber or food wastes, and (D) sludge.

295 All the groups were strictly related to the feedstock type, highlighting its influence on

296 organic waste classification, which is supported by other authors (Akhiar et al., 2021;

297 Bareha et al., 2018; Guilayn et al., 2019). Groups B, C, and D came from a different

298 main cluster than Group A, probably due to a higher DOM content related to complex

299 fluorescence zones (VI-VII) that could arise from the degradation of refractory

300 compounds of other fractions (Zhang et al., 2019). The high reported complexity in

301 animal slurries could be related to particular recalcitrant alkyl-C (e.g. sterols, lipids,

302 cutin) (Tambone et al., 2019). Group B was defined by PEOM (poorly accessible C)

303 and fluorescence zones IV and V. Indeed, these types of substrates are characterized to
304 present high C/N and TS besides complex proteins and humic/fulvic acids (Akhiar et
305 al., 2017). Finally, Groups C and D were related to low complexity zones (I-III) while
306 sludge showed higher fluorescence proportions on zones III than pig slurry mixtures. In
307 fact, zone III corresponds to protein content and microbial by-products, typically related
308 to activated sludge metabolism and growth/decay (Fang et al., 2015). These results have
309 been also reported by Zhang et al. (2019), who showed that the characteristic presence
310 of protein-like organics (zones II and III) was in the SPOM and REOM fractions of
311 sludge.

312 Interestingly, a broad variance on pig slurry mixtures clustering from fluorescence
313 zones I-III to VI-VII was found depending on the co-substrate added (e.g. pairs of pig
314 slurry + fiber co-substrate were clustered in Group C but spatially distributed close to
315 Group B, which is mainly composed of fiber-rich samples). Notwithstanding that the
316 percentage of co-substrate in raw mass was mainly below 20%, the addition of a co-
317 substrate seems to influence their classification and should be considered to properly
318 classify organic wastes (see Table 1).

319 Most of the digestates were clustered together with their substrates, confirming that the
320 main OM complexity and accessibility prevailed after AD (Provenzano et al., 2014).
321 Nevertheless, within the same cluster, variations in the classification between substrates
322 and digestates have been observed. Since the OM accessibility and complexity
323 conversion after AD remains unclear, it is discussed in the following section.

324 **3.2. OM accessibility and complexity: revealed groups characterization**

325 ***3.2.1. OM accessibility and fluorescence complexity index of substrates***

326 The proportion of total COD in the biochemical fractions and the FCI (defined in
327 Section 2.3) of substrates profiles have been evaluated to define each group and are
328 summarised in Figure 2.

329 The proportions of COD for each extracted fraction varied between groups according to
330 the different origins of the sample (Figure 2). The largest proportion of COD for Group
331 A was in the DOM fraction (30-55%), indicating a notable content of water-soluble
332 organic substances in slurries, such as simple sugars (e.g. sucrose, glucose and
333 fructose), proteins (mainly globular protein), volatile fatty acids or soluble recalcitrant
334 compounds. High DOM fraction content was also reported by Laera et al. (2019) (76%
335 of total COD) for a substrate from a household, slaughterhouse and industrial waste co-
336 digestion plant. For Groups B and C, PEOM fraction was the main COD fraction
337 extracted and ranged between 22 and 58% of total COD. Indeed, these results are
338 consistent since the PEOM fraction is rich in hemicellulose, cellulose-like, starch and
339 certain proteins (Laera et al., 2019), which characterizes the fibrous feedstocks of
340 Groups B and C. Finally, Group D extracted COD was dominated by the NEOM
341 fraction (23-33%). Similar values of NEOM (30-40%) have been reported by different
342 authors for sludge samples (Jimenez et al., 2015; Maynaud et al., 2017).

343 Comparing the accessibility between groups, Group D had the highest SEOM (27%)
344 and SPOM (22%) fractions probably due to high complex protein content (mainly
345 fibrous proteins) and simple sugars/proteins in sludge samples. Group C presented the
346 highest REOM (25%) and NEOM (39%) fractions probably because of the protein and
347 lipid content provided by food wastes. Group B presented the highest PEOM fraction
348 (58%), while the NEOM fraction was the second extracted fraction for all the groups
349 (except for Group D), indicating that all wastes had a considerable amount of non-

350 extractable OM. Whilst, SPOM and REOM fractions showed low percentages of the
351 total extracted COD for all the groups.

352 Regarding the FCI of the substrates, Groups A and B had higher FCI in all fractions
353 compared with Groups C and D (Figure 2). Indeed, Group C and D were mainly
354 characterized by fluorescence regions I to III likely they have less complex
355 proteins/lignified compounds than Groups A and B. Only the FCI of the SPOM fraction
356 of Group C (1.03-1.50) was higher than the one of the other Groups. The highest FCI
357 for all groups was for Groups A and B and corresponded to the less accessible fractions
358 (SEOM and PEOM), as reported by Muller et al. (2014). Concerning the FCI of each
359 group, the FCI of the SEOM fraction was the highest for the Group A (3.60). In fact,
360 this fraction targets recalcitrant compounds such as humic-like acids, fulvic-like acids
361 and complex proteins, which could explain that Group A was previously defined by
362 complex fluorescence zones (VI-VII) (see Figure 1). The FCI of the PEOM fraction
363 showed the highest values for Group B and D (2.94 and 1.97, respectively) whereas
364 Group C displayed slight differences in the FCI for all the extracted fractions.

365 Fluorescence spectroscopy was also performed on some DOM fractions samples (PS,
366 CM/S and HM/S + Slu, BW + CS + AI, Sludge_6 and Sludge_7). However, this data
367 was not considered in this study because (i) all the DOM fractions in the dataset were
368 not analyzed and (ii) Jimenez et al. (2017) did not consider it in their model.

369 Nonetheless, the fluorescence percentage for zone I to VII of SPOM and DOM of these
370 6 samples were similar for the evaluated substrates. Zhang et al. (2019) reported similar
371 results for sewage sludge. Therefore, DOM fluorescence was not relevant for an
372 accurate prediction of the final C mineralization in soil concerning the present study
373 (see Supplementary Material).

374 **3.2.2. OM accessibility and fluorescence complexity index conversion after AD**

375 The anaerobic biodegradability of OM also depends on the chemical nature of the
376 compounds, therefore, the more accessible fractions will not strictly be the more
377 biodegradable fractions (Bareha et al., 2019; Mottet et al., 2010). However, simpler
378 compounds (e.g. soluble sugars or proteins) will be extracted in the most easily
379 extractable fractions, while lignocellulose-like compounds are normally present in the
380 last extracted fractions. To evaluate these considerations, Table 3 shows the proportions
381 of COD for each extracted fraction and the FCI evolution between the substrates and
382 their corresponding digestates after AD. Meanwhile, Figure 3 displays the proportions
383 of COD for each extracted fraction (described in Section 2.2) and FCI profiles for the
384 digestates samples of each group.

385 From the four groups evaluated, general assumptions can be drawn regarding the
386 influence of AD in the OM quality of digestates. The SPOM and REOM fractions have
387 decreased after AD whereas the FCI increased for all feedstock groups (except SPOM
388 and REOM complexity of Group C which decreased). Aemig et al. (2019) have also
389 reported the highest biodegradation yields for the most accessible fractions (74% and
390 69% for SPOM and REOM, respectively) after sewage sludge AD. Thus, SPOM and
391 REOM have shown the lowest percentage of extracted COD (less than 10% of total
392 COD) for all digestates. Indeed, the sum of SPOM+REOM in digestates ranged between
393 3 and 27% of the total COD samples (data not shown). Low extracted COD proportions
394 for the most accessible fractions were also found in Bareha et al. (2018) and Laera et al.
395 (2019), where the sum of SPOM, REOM and SEOM was below the 10% of total COD.
396 Similarly, the sum of SPOM + REOM only accounted for the 2.7-10.6% of the total
397 COD of digestates OM by Maynaud et al. (2017). In accordance with the present study,
398 Aemig et al. (2019) and Zhang et al. (2019) have also reported an increase of the FCI

399 for SPOM and REOM fractions since simple fluorescence peaks remarkably decreased
400 after anaerobic digestion.

401 Regarding the DOM fraction, Group A kept the highest COD content in DOM (26-38%)
402 after AD. This could suggest that, DOM fraction could contain non-biodegradable
403 compounds or that AD could produce accessible but complex molecules (Jimenez et al.,
404 2015; Lashermes et al., 2009), which has also been shown during composting (Peltre et
405 al., 2011). Indeed, Zhang et al. (2019) showed that refractory organic compounds (i.e.
406 complex compounds) could be accumulated in DOM due to solubilization. In the
407 studied dataset, the DOM fraction has been displayed to be variable among the groups.
408 Actually, SPOM and DOM digestates spectra have been observed to present
409 differences, although without a significative general trend (see Supplementary
410 Material).

411 The SEOM fraction was kept similar or increased depending on the group. Laera et al.
412 (2019) have also noticed a SEOM fraction increase from 7 to 15% after AD. These
413 authors have associated it with the concentration of protein from the growth/decay of
414 microorganisms and to a preferable consumption of simple soluble compounds.
415 Similarly, the SEOM reduction was smaller compared with SPOM and REOM fractions
416 for sewage sludge and cow manure digestion, as they are expected to have more
417 complex and therefore less degradable compounds (Somers et al., 2021; Zhang et al.,
418 2019). The FCI of SEOM has tended to increase in the digestate of Groups A and C.
419 Meanwhile, the FCI of SEOM of Group B and D remained constant, whilst the SEOM
420 accessibility decreased, probably due to the biodegradation of non-fluorescent
421 compounds, as previously reported by Aemig et al. (2019).

422 PEOM fraction has also varied among feedstock types. The Groups with high initial
423 PEOM fraction (B and C) has shown a decrease while an increase was observed for the
424 Groups with low initial PEOM fraction (A and D). Similar findings were previously
425 reported in Bareha et al. (2018) and Aemig et al. (2016). These authors have reported a
426 PEOM fraction decrease for cow manure (i.e. rich in PEOM) and an increase for sewage
427 sludge (i.e. poor in PEOM), respectively. Therefore, higher degradation rates of PEOM
428 fraction (cellulose and hemicellulose) during AD could be expected when other simpler
429 molecules are missing (Tambone et al., 2013). Biodegradable cellulose/hemicellulose in
430 the PEOM fraction of Groups B and C have probably contributed to higher
431 biodegradability of the PEOM fraction. However, the PEOM fraction of Groups A and
432 D could contain non-identified complex compounds (e.g. complex proteins) that
433 increase the recalcitrance to AD. The FCI of PEOM has increased for all groups (except
434 for Group C). In contrast, Aemig et al. (2016) have reported no evolution on the FCI for
435 the less accessible fractions during sewage sludge digestion. Nevertheless, a slight
436 increase of the fluorescence intensity of humic-like compounds peaks in PEOM fraction
437 after AD (suggesting poor biodegradability and high complexity of this fraction), was
438 reported as a possible re-polymerization of humic acid (Tang et al., 2018; Zhang et al.,
439 2019).

440 The NEOM fraction has increased for all groups after AD and represented between 25
441 and 63% of the total COD in digestates. This lignocellulose-type fraction is
442 concentrated after AD due to its recalcitrance and poor biodegradability in anaerobic
443 conditions (Usman Khan and Kiaer Ahring, 2021). Thus, digestates are expected to be
444 enriched in recalcitrant compounds, enhancing their suitability as soil amendment
445 compared to the raw substrates (Jimenez et al., 2017; Pognani et al., 2010; Shakeri
446 Yekta et al., 2019; Teglia et al., 2011b).

447 The obtained results have shown that the OM conversions shared a similar pattern
448 regardless of the feedstock type: (i) an increase of complexity for the majority of the
449 fractions, (ii) a global decrease of accessibility of the most accessible fractions (SPOM
450 and REOM), and (iii) an increase of the non-extractable organic matter (NEOM)
451 fraction. Nonetheless, it should be noted that the discussed evolution of the OM quality
452 does not distinguish between the contribution due to the exogenous OM (i.e. substrate)
453 and the endogenous OM (i.e. microbial-related products). The contribution of each
454 compartment will be prioritized in forthcoming studies.

455 The accessibility conversion pattern is in agreement with Aemig et al. (2016), who have
456 also shown a decrease for the most accessible fractions (named DOM, S-EPS and RE-
457 EPS) during AD (22-65%, 49-50%, 23-35%, respectively), whereas NEOM relatively
458 increased. Similarly, cow manure AD was evaluated by Somers et al. (2021) and a
459 significant decrease in DOM, SPOM and REOM fractions of 24%, 62%, and 61%, was
460 associated with methane production and hydrolysis of organic matter. Laera et al.
461 (2019) have also noted a remarkable decrease from 76 to 28% for the DOM fraction
462 between raw and digested mixtures of household, slaughterhouse and industrial wastes.
463 Besides, an increase from 9% to 47% for PEOM + NEOM was also reported by these
464 authors. Moreover, Zhang et al. (2019) have also showed a decrease after AD from
465 46.74% to 39.42% for DOM, SPOM and REOM fractions and the FCI increased for all
466 the extracted fractions. Furthermore, the FCI of the REOM fraction was increased by
467 34% during cow manure AD (Somers et al., 2021) whereas a non-significant increase in
468 the FCI of SEOM and PEOM during sludge and cow manure AD was stated by Aemig
469 et al. (2019, 2016) and Somers et al. (2021), respectively. Therefore, general trends on
470 OM accessibility and complexity evolution during AD have been observed.

471 Additionally, the prediction of OM accessibility and complexity of digestates from their
472 inputs have been further assessed.

473

474 **3.3. Prediction of digestate OM quality from substrate OM quality**

475 ***3.3.1. PLS model for the prediction of digestate OM quality***

476 PLS regression was applied on 28 observations (substrate/digestate pair) split in two
477 datasets: a calibration dataset (23 samples) and a validation dataset (5 samples). Seven
478 models were tested (Table 4). Model n°1 used the substrate fractions DOM+SPOM,
479 REOM, SEOM and PEOM as X-variables to predict the digestate fractions (i.e.
480 DOM+SPOM, REOM, SEOM and PEOM). NEOM fraction was not included as this
481 fraction is calculated by difference. Model n°2 was similar to Model n°1 with the
482 addition of the reactor temperature (T) as X-Variable. Model n°3 was the Model n°1
483 with the addition of Hydraulic Retention Time (HRT) as X-Variables and Model n°4
484 merged Models n°2 and 3.

485 Model n°5 used both fractions and fluorescence percentage in each fraction measured in
486 the substrates (32 variables) as X-variables to predict the same 32 Y-variables in the
487 digestates. Models n°6, 7 and 8 were based on Model n°5 with the addition of the T,
488 HRT and both variables as X-variables, respectively. Table 4 presents the quality
489 parameters of each model. The errors of calibration, cross-validation and prediction
490 using cross-validation methods (RMSE, RMSE_CV, and RSMEP) of each Y-variable in
491 each model are specified in the Supplementary Material.

492 According to Table 4, all the models have shown good quality performances with $Q^2 >$
493 0.5 and correlation coefficients of prediction were between 0.691 and 0.832. Models n°5
494 to 8 had better correlation coefficients than Models n°1 to 4 and smaller calibration

495 errors (cf. RMSE in Supplementary Material). Indeed, the addition of fluorescence
496 variables from Model n°1 to n°5 improved the prediction of the 32 variables with 7
497 components. Overall, for Model n°5, R^2 was 0.831 and specific R^2 for each fraction
498 prediction was 0.869, 0.910, 0.736 and 0.882 for respectively DOM+SPOM, REOM,
499 SEOM and PEOM. Furthermore, the prediction error RMSEP was lower in Model n°5
500 for the fractions prediction than Model n°1 (see Supplementary Material).

501 The Model n°2 and 4 had the best performances among the models using only fractions
502 and operational parameters as X-Variables. Indeed, the addition of T (Model n°2) and
503 the addition of HRT (Model n°3) as X-variables increased R^2Y but needed a component
504 addition. However, HRT addition did not improve the quality parameters R^2Y and Q^2
505 (Model n°4 in Table 4). This trend was not the same in the case of Models n°6 to 8 in
506 comparison with Model n°5 without operational parameters. Indeed, despite a little
507 increase of R^2Y when T was added as X-variables (Model N°6), the prediction accuracy
508 (Q^2) was decreased when HRT and T were added. The results obtained in Model n°2
509 have shown that T was anti-correlated (p-value < 0.05) with SEOM ($R^2 = -0.593$) and,
510 less significantly with DOM+SPOM ($R^2 = -0.313$). However, reactors T were mainly
511 mesophilic except for MW digestates (55 °C). These digestates were associated with
512 lower DOM+SPOM and SEOM fractions than the others. Consequently, relying on our
513 dataset, the impact of T was related to the substrate type. To test the impact of T alone,
514 it would have been interesting to get substrate/digestate pairs from various feedstocks at
515 both mesophilic and thermophilic temperature.

516 Similarly, the obtained results in Model n°3 have shown that HRT was negatively
517 correlated with REOM ($R^2 = -0.628$) and, positively correlated with PEOM ($R^2 =$
518 0.397). However, low HRT values were mainly found for sludge digestion whereas
519 high HRT values were associated with cow manure digestates. Sludge digestate groups

520 were related to a high value of REOM and SEOM fractions whereas cow manure
521 digestates were characterized by high values of PEOM and low values of DOM+SPOM.
522 Again, it was not possible to distinguish the effect of HRT from substrate impact on
523 digestate quality prediction. Moreover, in the Model n°4, weight coefficients showed
524 that HRT and T were considered as the least important variables for quality prediction
525 (Supplementary Material).

526 As the operational conditions have been determined as non-explicative enough in this
527 study, and considering the high quality of the prediction, Model n°5 was selected.
528 Besides, the predicted variables of digestates quality are necessary for the PLS model
529 for C biodegradability in soil. To go further, the impact of the most significant X-
530 variables affecting the prediction of Y-variables derived from the calculation of the
531 weight of each variable for Model n°5 (without T and HRT) was analyzed (data not
532 shown).

533 The characteristics of input OM remarkably influenced the prediction of the digestate
534 OM characteristics, meaning that the main pattern of accessibility present in feedstock
535 input remained in digestate OM after AD. The variables of the most complex
536 fluorescence zones from substrate impacted significantly the digestate quality. The
537 recalcitrant compounds contained in the substrate were preserved in their subsequent
538 digestate, as possible humus precursors (Guilayn et al., 2020; Tambone et al., 2010).
539 Nonetheless, for the prediction of the simpler fluorescence zones, other factors such as:
540 (i) solubilisation/complexification of biodegradable/non-biodegradable compounds, (ii)
541 preferential compounds degradation, (iii) hydrolysis, (iv) prevalence of recalcitrant
542 compounds, (iv) compounds contribution from other fractions are possible hypothesis
543 that contribute to the explanation of the resulting prediction. Therefore, this is the first
544 approach to understand how OM quality varies with AD based on accessibility and

545 complexity for a wide range of feedstock. The addition of the T and the HRT as X-
546 variables to predict OM quality and accessibility confirmed that the operational
547 conditions were not informative enough for the studied dataset. To properly evaluate the
548 impact of the operational conditions on the prediction of digestate quality, the digestion
549 of the same substrate subjected to different T or HRT should be conceived in future
550 investigations.

551 ***3.3.2. Coupling digestate quality prediction with PLS model for carbon***

552 ***biodegradability in soil prediction***

553 To validate the digestate quality PLS model found, the PLS model for C
554 biodegradability in soil was applied on the digestate quality predicted by Model n°5.
555 Among all the samples that were used for the digestate quality model, 14 samples were
556 incubated in soil and biodegradable carbon (C_bio) was obtained after 91 days (Table
557 2).

558 First, the PLS model for C biodegradability in soil was tested. A comparison between
559 the 14 predicted values of proportion of biodegradable C in the digestates and
560 experimental data obtained through soil incubation was plotted in Figure 4a. Results
561 have shown that the PLS model for C biodegradability in soil was successfully able to
562 predict the biodegradable organic C of the 14 digestates ($R^2 = 0.739$) with low bias.
563 Then, Model n°5 was combined with the PLS model for C biodegradability in soil to
564 predict C_bio. Figure 4b shows that the C_bio prediction was not altered by the models'
565 combination. Indeed, the combined models were able to predict the experimental data
566 ($R^2 = 0.697$) with a similar bias as in Jimenez et al. (2017). The 5 validation samples
567 that were used for Model n°5 were plotted in black in Figure 4a and b. Prediction error
568 of C_bio obtained by the combined models ranged between 1% and 7%, with $R^2 =$

569 0.828, which represents a high quality of prediction. Therefore, the reproducibility of
570 the model was confirmed. Moreover, the model validation was performed with external
571 data not included in the dataset used for model calibration. Finally, the 28 pair samples
572 were used to compare both models' predictions as presented in Figure 4c. Results
573 showed that C_bio prediction by the combined models is quite similar to C_bio
574 prediction by Jimenez et al. (2017) ($R^2 = 0.894$).

575 Thus, Model n°5 provided an extra step on OM fate prediction from raw substrate to soil
576 C mineralization to better understand AD influence on OM accessibility and complexity
577 before land disposal. Additionally, the present study related multiple aspects of
578 scientific interest such as waste characterization, anaerobic transformation processes of
579 OM and soil C mineralization and supposes an innovative approach to enhance the
580 modeling of the AD process chain. Nonetheless, future actions could be addressed to
581 improve the models' application when specific substrates are subjected to different AD
582 operational conditions and soil typologies.

583

584 **4. Conclusions**

585 The prediction of digestate OM quality from their input was evaluated. PCA and HCA
586 analysis have allowed to classify 28 substrate/digestate pairs covering a wide diversity
587 of OM. This classification was based on the extracted fractions from the OM
588 (accessibility) and their complexity assessed by fluorescence. Substrates and their
589 respective digestates were clustered together according to the feedstock type.
590 Nonetheless, common trends on the conversions of OM quality were observed,
591 indicating potential for the prediction of digestate quality of the entire dataset regardless
592 of the feedstock type. Thus, this study proposed a digestate quality PLS model that

593 accurately predicted ($Q^2 = 0.593$) the digestates OM quality from the substrate OM
594 characteristics. However, future investigations should be focused on subjecting the same
595 substrate to different T or HRT to properly evaluate the impact of operational conditions
596 on the prediction of digestate quality. The predicted digestate OM characteristics
597 validated the prediction of their biodegradability in soils using the PLS model for C
598 biodegradability in soil previously developed. This work performed the combination of
599 both models. OM conversion during AD and soil C mineralization was precisely
600 predicted using a rapid analysis indicator (biochemical fractionation and 3D
601 fluorescence). Such combined models brought a major contribution in the modeling of
602 the AD process chain favoring the development of decision-making tools to properly
603 manage the digestates.

604

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612

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816 **Table 1.** Summary of the raw substrate/digestate pairs type, origin and digester
 817 operational conditions

Type of feedstock ^a	Digestate type ^b	Number of substrate/digestate pairs	Scale	Origin	T (°C)	HRT (d)
Cow manure	4	5	Industrial	Farms	35-41	56-75
Cow manure mixtures:		2				
CM + Straw		1	Industrial	Farms	35-37	56
50% CM/S + 23% HM/S + 27% Slu		1	Industrial	Farms	35-37	56
Straw	1	1	Lab scale	Farms	-	-
Biowaste	2	1	Industrial	Municipal solid waste plant	55	21
Biowaste mixtures:		1				
50% BW + 20% CS + 30% AIW	5	1	Industrial	Municipal solid waste plant	37	90
Municipal Waste	5	4	Industrial	Municipal solid waste plant	55	20-28
Pig Slurry	3	1	Industrial	Farms	38	60
Pig Slurry mixtures:		6				
45% PS + 40% PsluAI + 15% others	3	1	Lab scale	Farms	38	60
93% PS + 7% Cow Food	3	1	Lab scale	Farms	38	24
93% PS + 7% Horse Food	3	1	Lab scale	Farms	38	24
80% PS + 20% Maize Silage	1	1	Lab scale	Farms	38	25
62% PS + 38% CM	1	1	Lab scale	Farms	38	24
80% PS + 20% BW	3	1	Lab scale	Farms	38	21
Sludge	2	7	Industrial	Wastewater treatment plant	37	15-25

818 ^aType of feedstock: AIW: agroindustrial waste; BW: biowaste; CM: cow manure; CS: cow slurry; HM: horse manure; PS: pig
 819 slurry; PsluAI: primary sludge from agroindustry; S: slurry; Slu: Sludge. ^bDigestate type based on Guilayn et al. (2019): (1) Fibrous
 820 feedstock (2) Sewage sludge, Biowaste, food agroindustrial residues (FAI) mono/co-digestion; (3) Organic fraction of municipal
 821 solid waste (OFMSW), Food waste (FW), FAI, PS mono/co-digestion; (4) Manure/other co-digestion; (5) OFMSW and BW
 822 mono/co-digestion; (6) Fibrous feedstock: Cattle manure, green waste, silage.

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839 **Table 2.** Biodegradable carbon percentage after soil incubation of several studied
 840 digestates coming from Jimenez et al. (2017) data

Digestate name	C_bio (%C)
Sludge2_D	39%
Sludge7_D	31%
CM3_D	15%
MW1_D	25%
MW3_D	32%
MW4_D	19%
MW5_D	27%
PS +BW_D	46%
PS + Cow Food_D	43%
PS + Cow Manure_D	24%
PS + Horse Food_D	35%
PS + Maize Silage_D	44%
BW_D	24%
PS + PrimSluAI_D	31%

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859 **Table 3.** Proportions of COD for each extracted fraction and fluorescence complexity
 860 index evolution after AD in the different groups. The results display the relative
 861 percentage of the increase or decrease as expressed in Tambone et al. (2013).

Group	Accessibility (% in COD)						Fluorescence complexity index (-)			
	DOM	SPOM	REOM	SEOM	PEOM	NEOM	SPOM	REOM	SEOM	PEOM
A	--	--	--	+	++	++	++++	++	+	++
B	N.A.	--	-	-	-	++	++	++	0	+
C	++++	---	--	++	--	+	--	-	+++	-
D	--	--	--	-	+	++	++	++	0	+++

862 Relative conversion ranges: +: 0 to 25%; ++: 25 to 50%; +++: 50 to 75%; ++++: 75 to 100%; -: 0 to -25%; --: -25 to -50%; ---: -50
 863 to -75%; ----: -75 to -100%. Relative percentage = (final value in the digestates - initial value in the substrates)/initial value in the
 864 substrates) × 100. N.A. = no presence of DOM for Group B (solid digestates)

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887 **Table 4.** Quality parameters of the PLS models

Model	Variables		Model quality			
	X	Y	Components number	R ² X	R ² Y	Q ²
Model n°1	DOM+SPOM; REOM; SEOM; PEOM	DOM+SPOM; REOM; SEOM; PEOM	2	0.892	0.691	0.621
Model n°2	DOM+SPOM; REOM; SEOM; PEOM;T	DOM+SPOM; REOM; SEOM; PEOM	3	0.919	0.763	0.681
Model n°3	DOM+SPOM; REOM; SEOM; PEOM;HRT	DOM+SPOM; REOM; SEOM; PEOM	3	0.922	0.757	0.639
Model n°4	DOM+SPOM; REOM; SEOM; PEOM;HRT;T	DOM+SPOM; REOM; SEOM; PEOM	4	0.938	0.775	0.587
Model n°5	DOM+SPOM; REOM; SEOM; PEOM;Pf_i	DOM+SPOM; REOM; SEOM; PEOM;Pf_i	7	0.953	0.831	0.593
Model n°6	DOM+SPOM; REOM; SEOM; PEOM;Pf_i;T	DOM+SPOM; REOM; SEOM; PEOM;Pf_i	7	0.950	0.832	0.588
Model n°7	DOM+SPOM; REOM; SEOM; PEOM;Pf_i; HRT	DOM+SPOM; REOM; SEOM; PEOM;Pf_i	7	0.951	0.829	0.58
Model n°8	DOM+SPOM; REOM; SEOM; PEOM;Pf_i;T;HRT	DOM+SPOM; REOM; SEOM; PEOM;Pf_i	7	0.949	0.830	0.576

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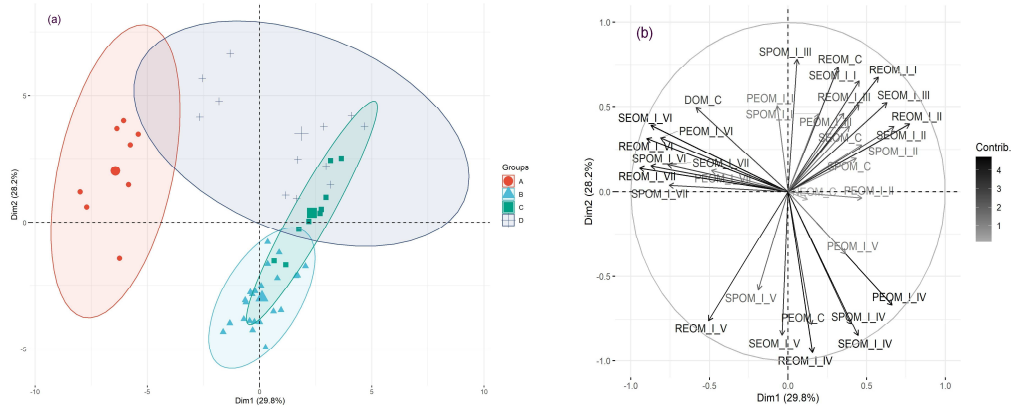
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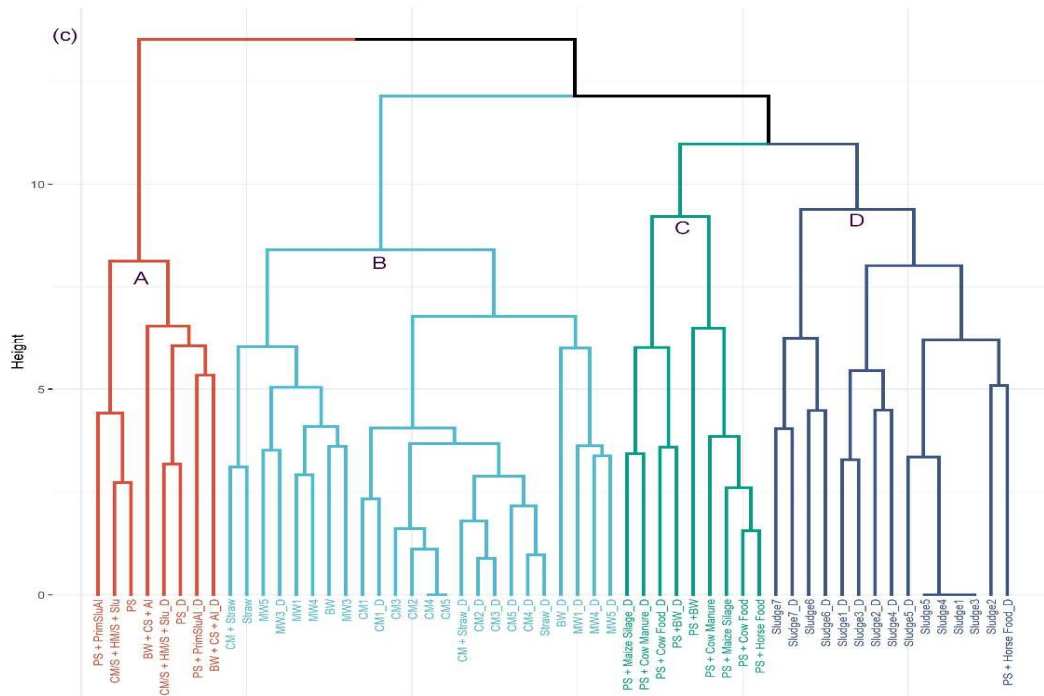
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909 **Figure 1.** Scores plot (a) and loadings (b) obtained from the PCA analysis, and (c) HCA
 910 analysis for the 56 samples studied. PCA individuals are distinguished by shape and
 911 color according to the HCA revealed groups (A-D). Ellipses show 95% confidence
 912 intervals. Loadings intensity color (plot b) is related to the variables contribution,
 913 from low (1) to high (4). Groups: (A) pig slurry and slurry mixtures with primary
 914 agro-industrial waste or biowaste, (B) manure, fibers and municipal solid waste, (C) pig
 915 slurry mixtures with fiber or food wastes, and (D) sludge

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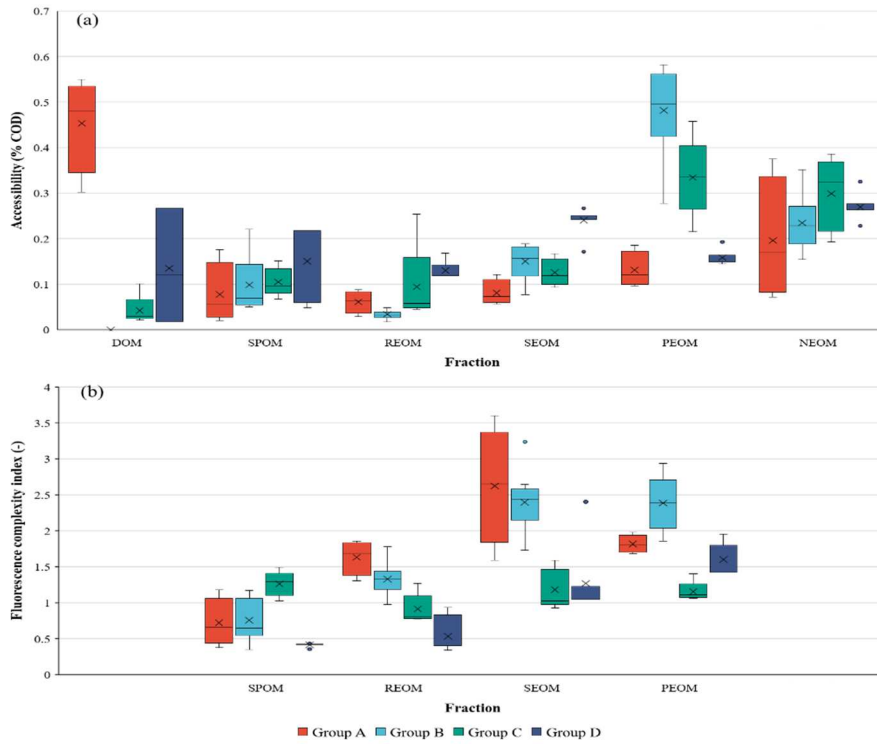
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925 **Figure 2.** Accessibility (a) and fluorescence complexity index (b) boxplots of substrates
926 for the HCA revealed groups. Groups: (A) pig slurry and slurry mixtures with primary
927 sludge, agro-industrial waste or biowaste, (B) manure, fibers and municipal solid waste,
928 (C) pig slurry mixtures with fiber or food wastes, and (D) sludge

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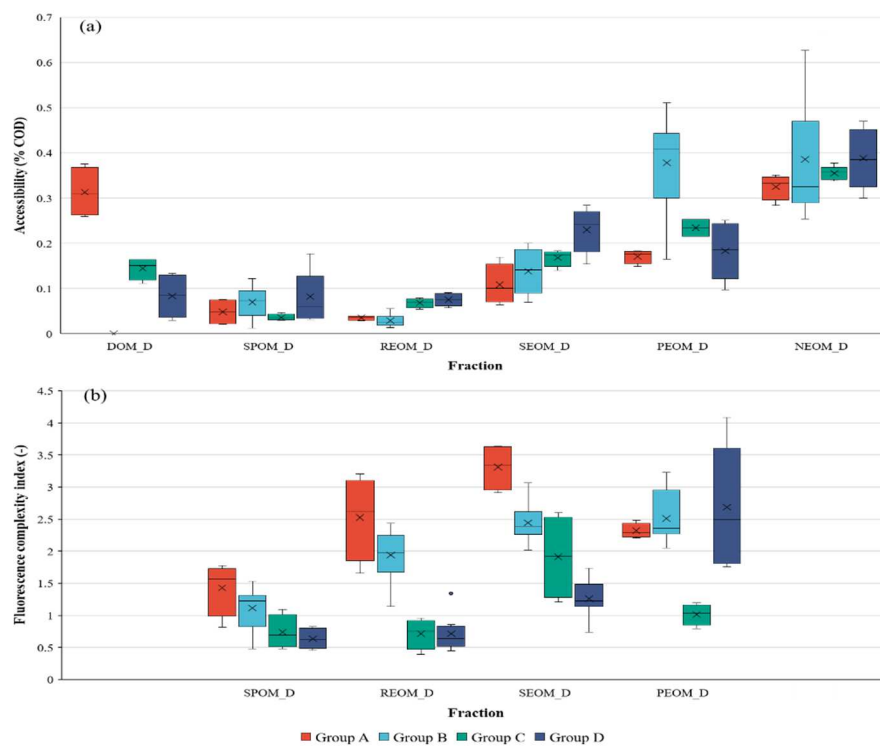
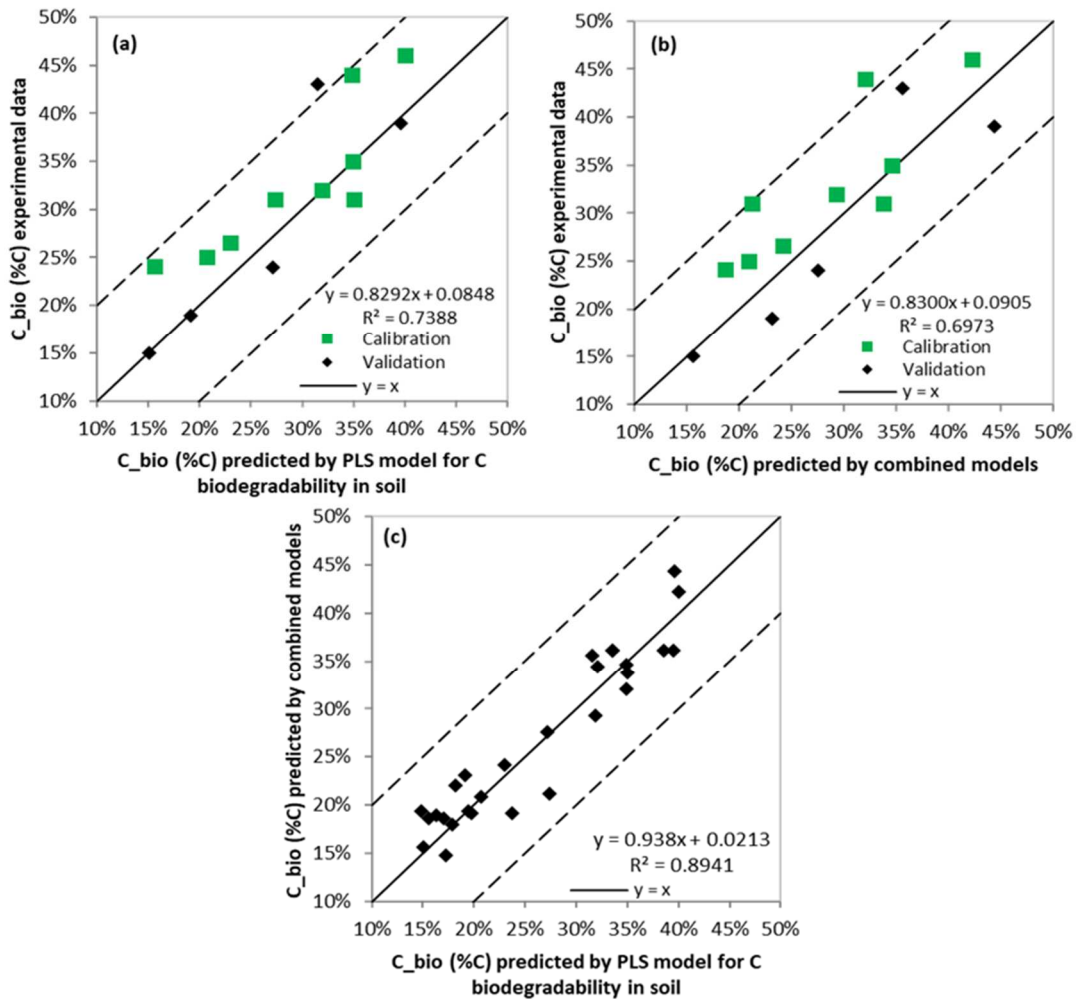


Figure 3. Accessibility (a) and fluorescence complexity index (b) boxplots of digestates for the HCA revealed groups. Groups: (A) pig slurry and slurry mixtures with primary sludge, agro-industrial waste or biowaste, (B) manure, fibers and municipal solid waste, (C) pig slurry mixtures with fiber or food wastes, and (D) sludge



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973 **Figure 4.** Validation of model PLS: comparison between experimental data from
 974 biodegradable organic carbon tests on soil with PLS model for C biodegradability in
 975 soil (a) and combined models (b); and comparison between combined models and PLS
 976 model for C biodegradability in soil (c)

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