

Anaerobic digestates from sewage sludge used as fertilizer on a poor alkaline sandy soil and on a peat substrate: Effects on tomato plants growth and on soil properties

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Abstract: Anaerobic digestates from sewage sludge (SSADs) are a by-product of the wastewater treatment process that still preserves a certain agronomic interest for its richness in plant nutrients and organic matter. Fertilizing properties of two liquid and two dewatered SSADs were tested on tomato plants (*Solanum lycopersicum* L.). Pot experiments were performed on sandy soil and peat substrate under greenhouse conditions with a SSADs application rate of 170 kg N/ha over a period of three months. Beneficial effects of SSADs were reported on different growth parameters, revealing an increase in biomass and height up to 37.5 and 6-folds over untreated control. No phytotoxic effect occurred on SSAD-exposed plants. Chemical analysis of soils treated with SSADs showed enrichment of macro- and micro-nutrients as well as organic matter. In some cases, the chemical characterization of leaves revealed an enhancement of uptaken macronutrients. This study contributed in general to deepen the knowledge on the short-term growing season fertilizing effects of SSAD. Despite the treatment dosage was calculated only on nitrogen requirements, the study highlighted the importance of the other nutrients and organic matter on plant growth.

1 Title

2 **Anaerobic digestates from sewage sludge used as fertilizer on a poor alkaline sandy**
3 **soil and on a peat substrate: effects on tomato plants growth and on soil properties**

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26 **Declaration of Interest:**

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28 **Abstract**

29 Anaerobic digestates from sewage sludge (SSADs) are a by-product of the wastewater treatment
30 process that still preserves a certain agronomic interest for its richness in plant nutrients and organic
31 matter. Fertilizing properties of two liquid and two dewatered SSADs were tested on tomato plants
32 (*Solanum lycopersicum* L.). Pot experiments were performed on sandy soil and peat substrate under
33 greenhouse conditions with a SSADs application rate of 170 kg N/ha over a period of three months.
34 Beneficial effects of SSADs were reported on different growth parameters, revealing an increase in
35 biomass and height up to 37.5 and 6-folds over untreated control. No phytotoxic effect occurred on
36 SSAD-exposed plants. Chemical analysis of soils treated with SSADs showed enrichment of macro-
37 and micro-nutrients as well as organic matter. In some cases, the chemical characterization of leaves
38 revealed an enhancement of uptaken macronutrients. This study contributed in general to deepen the
39 knowledge on the short-term growing season fertilizing effects of SSAD. Despite the treatment
40 dosage was calculated only on nitrogen requirements, the study highlighted the importance of the
41 other nutrients and organic matter on plant growth.

42

43 **Keywords:** waste management; nitrogen; soil organic matter; nutrient recycling; sewage sludge;
44 tomato plants.

45

46 1. Introduction

47 Globally, the demand of the three primary plant nutrients used for soil fertilization (N, P₂O₅ and K₂O) is
48 increasing (Vanotti et al., 2019). In 2015, the total fertilizer nutrient demand was around 184 Mt and,
49 by the end of 2020, it is expected to overcome 200 Mt (FAO, 2017). The production processes of
50 these fertilizers are very expensive in terms of energy (ammonia) and non-renewable resources
51 (phosphorus and potassium), with heavy environmental costs (Li et al., 2009). Ammonia production is
52 mainly performed via the Haber-Bosch process which requires a large amount of fossil fuel (Basosi et
53 al., 2014). Phosphate rock is the principal raw material exploited in the production of nearly all
54 phosphate fertilizers (Fixen and Johnston, 2012; Reijnders, 2014). This non-renewable resource may
55 contain many toxic heavy metals such as As, Hg, Ni, V (Mortvedt, 1995), Cd, Cr, Cu, Pb, Zn (Sabiha-
56 Javied et al., 2009), fluorine (Mirlean and Roisenberg, 2007) and uranium (Schnug and Lottermoser,
57 2013). The P₂O₅ extraction can cause environmental pollution by contaminants accumulating in air,
58 soil, and water bodies around the manufacturing place (Mirlean et al., 2008; Sabiha-Javied et al.,
59 2009). It has been observed that these impurities can persist into phosphate fertilizers, provoking a
60 subsequent accumulation in agricultural soils (De López Camelo et al., 1997). Potassium derives from
61 non-renewable resources like minerals such as sylvite, sylvinit, hartsalz and langbeinite (Fixen and
62 Johnston, 2012). Furthermore, world distribution of phosphorous and potassium mines is not uniform:
63 45% of global phosphate rock is concentrated in Morocco and the Western Sahara (Fixen and
64 Johnston, 2012).

65 Within a circular economy perspective, the reuse of sewage sludge (SS) as fertilizer is an interesting
66 scenario. SS can be defined as “the residue generated from the treatment of wastewater” (Smith et
67 al., 2009). This matrix is a valuable source in terms of plant nutrients: a study conducted on 240 dried
68 samples from Pennsylvania revealed an average N, P and K content of 4.74%, 2.27%, and 0.31%,
69 respectively (Stehouwer et al., 2000). Furthermore, SS can contain many micronutrients (e.g. Ca, Mg,

Abbreviations: A_N: assimilation; ANRE: Apparent Nitrogen Recovery Efficiency; ANUE: Agronomic Nitrogen Use Efficiency; C: centrifuged SSAD; CEC: cation exchange capacity; CCI: Chlorophyll Content Index; C_i: CO₂ concentration in substomatal cavity; CRF: controlled release fertilizer; D: dried SSAD; D.M.: dry matter; EC: electrical conductivity; EDC: endocrine disrupting compounds; EmC: Emerging Contaminants; EU: European Union; g_s: stomatal conductance; IRGA: infra-red gas analyzer; M: mineral fertilizer; OM: organic matter; P: primary SSAD; QL: quantification limits; S: secondary SSAD; SS: sewage sludge; SSAD: anaerobic digestate from sewage sludge; T: non-treated, control thesis; WWTP: wastewater treatment plant.

70 S, Fe, Mn, Cu, Zn and B) which are important for plant growth, but usually not included in most
71 commercial fertilizers (Warman and Termeer, 2005). The percentage of the nutrients appears low, but
72 it is important to underline that every year a huge amount of wastewater is produced. An empirical
73 study revealed that approximately 330 km³ of municipal wastewater are produced worldwide yearly
74 (Mateo-Sagasta et al., 2015). Therefore, also the SS production has dramatically risen thanks to
75 policies dealing with the improvement of wastewater treatment and of standard quality of effluents,
76 such as the E.U. directive 91/271/EEC (Council of the European Communities, 1991a).

77 The considerable presence of organic carbon and organic matter in SS is another strength of its reuse
78 (Alvarenga et al., 2015; Mateo-Sagasta et al., 2015). In fact, land application of organic matter (OM)
79 improves soil physical properties such as cation exchange capacity (CEC), soil structure, soil
80 moisture content and retention (Epstein, 2002). Furthermore, the addition of SS can enhance the
81 amount of organic carbon in soils (Kladivko and Nelson, 1979; Perez-Espinosa et al., 1999) and thus
82 reverse the current reduction of organic matter in soils (known as *SOM decline*) (Schulze and
83 Freibauer, 2005).

84 Today, SS is classified as waste and its safe disposal represents a very important issue in waste
85 management (Epstein, 2002; Singh and Agrawal, 2008). The four main destinations of SS are
86 incineration, landfilling, composting and agricultural use. In Italy, according to Eurostat data (Eurostat,
87 2019), the majority of SS is sent to landfill (50.8%), while 34.7% is reused in agriculture, 4% is
88 incinerated and 10.4% is sent to other destinations. The Council Directive 86/278 (Council of the
89 European Communities, 1986) regulates the agricultural SS reuse in Europe to prevent soil
90 contamination. In fact, this practice has three principal problems that limit its unconditioned use:
91 biological risk, heavy metal contamination and contamination by organic pollutants. The biological risk
92 is principally represented by pathogens such as *Salmonella* spp., *Escherichia coli* (enterotoxigenic
93 and enteropathogenic variants), *Campylobacter* spp., *Clostridium* spp., and *Yersinia* spp. (Arthurson,
94 2008); stabilization treatments can reduce significantly their presence in SS and are mandatory before
95 subsequent SS applications (Dumontet et al., 1999). For instance, one of the most diffused
96 stabilization techniques is anaerobic digestion (Liu et al., 2012), in which the reduction of pathogens,
97 putrescence and odor is coupled with biogas production, allowing energy recovery (Epstein, 2002).

98 Heavy metal content (normally represented by Cd, Cr, Cu, Hg, Ni, Pb, Zn) can be abated by means of
99 chemical (e.g. chelating addition), physical (e.g. electroremediation) or biological (e.g.
100 vermicomposting) treatments (Camargo et al., 2016). Finally, some organic pollutants (e.g. pesticides,
101 antibiotics and hormones) can be volatilized or degraded through biotic or abiotic processes (Harrison
102 et al., 2006). Concerning organic pollutants, their abatement is trickier. Indeed, class of emerging
103 contaminants (EmC) in wastewater is increasingly gaining more interest within the organic
104 compounds. EmC include molecules such as endocrine disrupting compounds (EDC, e.g. hormones),
105 pharmaceutically active compounds (e.g. antibiotics), illicit drugs and pesticides (Fijalkowski, 2019).
106 EmC abatement is becoming even more required both on the effluent of WWTPs with advanced
107 treatments (e.g. activated carbon absorption, advanced oxidation processes, reverse osmosis) and on
108 sewage sludge (Gadupudi et al., 2019). Some studies affirmed that anaerobic digestion is the
109 stabilization strategy ensuring the best EmC removal, especially when the sludge is pretreated (e.g.
110 via ozonation) (Neumann et al., 2016). However, further studies are still required to improve the
111 performances and to reduce the costs of these techniques, which nowadays are rarely applied at
112 WWTP level since they are money and/or time consuming (Camargo et al., 2016). The
113 abovementioned EU directive regulates the SS soil application in the EU and establishes threshold
114 values of some of these pathogens and pollutants in SS.

115 On the basis of these opportunities and threats related to SS, this work aims to deepen the
116 knowledge about SS fertilizing effects over time in terms of nutrients and OM on a poor alkaline sandy
117 soil. This kind of soil was selected because: i) nutrient depletion constrains plant growth to depend on
118 treatment application; ii) a high pH both hinders the nutrient adsorption and reduce the metal
119 bioavailability (Alvarenga et al., 2016); iii) sandy-textured soil lacks nutrients and has low water-
120 holding capacity. These results were compared to the one obtained with a richer peat substrate. Pot
121 experiments were performed on tomato plants (*Solanum lycopersicum* L.) in a greenhouse to
122 evaluate nutrient provision of anaerobic digestates from sewage sludge (SSADs). Tomato plant was
123 chosen because: i) it is one of the most exploited vegetables crop (Jones Jr, 2008); ii) there is an
124 increasing interest on alternative nutrient sources for this crop (Zucco et al., 2015); iii) it has a high
125 fertilizer requirements (Zucco et al., 2015); iv) plenty of scientific literature is available for this crop

126 (Jones Jr, 2008). In this work, no analysis on pathogens was carried out since anaerobic digestion is
127 considered one of the safest technologies for pathogen reduction in SS (Epstein, 2002).
128 Nevertheless, this aspect may be taken into consideration in future researches. Concerning the use of
129 SS in agronomic experiments, a lack in details about SS typology is provided. Indeed, in many work
130 no detail on stabilization strategy is provided (Bakshi et al., 2019), or the kind of SS digestion is not
131 specified (Hossain et al., 2015). In the present work, the digestates used derived from the same
132 WWTP and were obtained with consequent treatments (Cristina et al., 2019). As far as we know, this
133 is the first example of use of four different and consequent SSADs to fertilize tomato plants. The
134 paper examines agronomic parameters of tomato plants, the nutrient distribution in soil and nutrient
135 absorption by plants after the application of SSADs. Furthermore, numerous plants physiological
136 parameters were evaluated over a span of three months in order to better understand the effects in a
137 time course approach.

138 2. Materials and methods

139 2.1. Characterizations

140 2.1.1. Anaerobic digestates from sewage sludge

141 Four SSADs were used in the experiment: two liquid (primary (P) and secondary (S)) and two
142 dewatered (centrifuged (C) and dried (D)). Physical and chemical characterization of the SSADs is
143 described in a previous work (Cristina et al., 2019); characterization of the four SSADs is reported in
144 **Supplementary Material -Section I (Table S1)**.

145 2.1.2. Cultivation substrates

146 Two types of substrates were used: a sandy soil and a commercial peat substrate (**Table 1**). The
147 sandy soil was sampled within 20 and 100 cm depth in Grugliasco (TO), Italy (45°03'58.4"N,
148 7°35'32.9"E). Analytical methods used for characterization of the sandy soil and the peat substrate
149 are specified in **Supplementary material - Section II**. Based on the distribution of the particle size
150 (sand: 94% ± 2; silt: 3% ± 1; clay: 3% ± 1), the selected soil was classified as sandy (Buol et al.,
151 2011). Based on ARPAV soil analysis (Arpa Veneto, 2007), the soil was considered alkaline (8.2 ±

152 0.16), very poor in OM ($0.38 \pm 0.12\% < 0.8\%$) and very poor in macronutrients such as nitrogen (0.29
153 ± 0.09 g/kg < 0.5 g/kg), phosphorous (1.8 ± 1.3 mg/kg < 7 mg/kg), potassium (18 ± 1 mg/kg < 40
154 mg/kg) and magnesium (15 ± 5 mg/kg < 50 mg/kg). On the other hand, content of calcium (675 ± 27
155 mg/kg < 1000 mg/kg) and some microelements such as iron (2.5 mg/kg $< 6.7 \pm 1.1$ mg/kg < 20 mg/kg)
156 and manganese (2 mg/kg $< 6.5 \pm 3.0$ mg/kg < 10 mg/kg) resulted normal.

157 The peat substrate used consisted of a commercial blend of blond and black peat (15:85, Turco
158 Silvestro, Italy), mixed with perlite (80:20 v/v). The substrate was steamed at 90°C for 30 minutes
159 before use. The substrate had the following characteristics as indicated by the manufacturer: pH 6.1;
160 E.C.: 0.56 dS/m; bulk density 250 kg/m³; C total 175 g kg⁻¹; N total 7 g kg⁻¹; organic matter 32% d.m.;
161 P₂O₅ 10 g kg⁻¹ ; K₂O 11 g kg⁻¹. Hence, the peat substrate could be reasonably considered a good
162 cultivation substrate, satisfying the requirements as a benchmark to be compared with the poor sandy
163 soil.

164 2.2. Experimental set-up

165 A greenhouse experiment was performed over three months during the summer season in a
166 greenhouse of the Centre of Competence AGROINNOVA – University of Torino, located in Grugliasco
167 (TO), Italy. The experimental campaign was carried out with commercial plastic pots of 2.5 L (\varnothing 17
168 cm, height 20 cm, surface area 0.227 m²). Four types of SSADs (P, S, C, D) were applied as
169 treatments, and compared to a commercial fertilizer (M) (NPK 22-5-6 + 2MgO, “Osmocote Topdress”,
170 ICL, Israel) and an untreated control (T). The experiment was designed in a completely randomized
171 block, with 15 replications per each thesis. The same experimental set-up was adopted on the two
172 cultivation substrates (sandy soil and peat substrate). Each treatment was applied at the dosage of
173 170 kg N/ha, in line with the European Nitrates Directive (Council of the European Communities,
174 1991b). Moreover, this application rate was chosen as it showed the best results in a preliminary
175 study (Cristina et al., 2019). Three untreated seeds of tomato (*Solanum lycopersicum* L. cv. Beefsteak,
176 “Furia sementi”, Parma, Italy) were sown in each pot. Automatic sprinkler irrigation was set three
177 times a day for 2-3 minutes in order to keep 40-50% WHC. Ten days after sowing a thinning was

178 conducted and the best plant from each pot was kept. At the end of each month, five pre-selected
179 replicates of each treatment were removed to carry out all the measurements.

180 2.3. Measurement of plant parameters

181 At the end of every month, the five removed replicates were examined. Firstly, height was measured,
182 then, leaves, inflorescences and fruits were counted, if present. After that, the Chlorophyll Content
183 Index (CCI) was evaluated with a SPAD 502 chlorophyll meter (CCM-200, Opti Sciences, Inc.,
184 Hudson, NH, USA) using the method described in the previous work (Cristina et al., 2019). One
185 month after sowing, it was not possible to measure CCI on the sand specimen because the minimum
186 leaves size was not satisfied. At the end of the second month, assimilation (A_N), stomatal
187 conductance (g_s) and CO_2 concentration in substomatal cavity (C_i) were measured by the means of
188 an Infrared Gas Analyzer (IRGA, ADC, Hoddesdon, UK). These measurements were performed on
189 three fully formed leaves in each replicate. The selected leaves had to be non-senescing, at the same
190 physiological age (in the middle part of the plant, considering the third to fourth leaf from the shoot
191 apex) and directly exposed to sunlight. After all the measurements were taken, each plant was
192 subsequently cut and immediately weighed to record the fresh biomass value. In order to evaluate the
193 mean dry biomass, each plant was dried at 105°C for at least 72 hours. Subsequently, agronomic
194 nitrogen use efficiency (ANUE) was calculated as:

$$ANUE = \frac{(Dry\ biomass\ treated\ samples - Dry\ biomass\ control\ samples)}{Amount\ of\ nitrogen\ applied\ in\ treated\ samples}$$

195 2.4. Chemical analysis

196 Substrates were chemically characterized at the end of the second month, once the aerial plant part
197 had been cut. Chemical analyses were performed on samples from the treatment with SSADs (P, S,
198 C, D) as well as on minerally fertilized ones (M) and untreated control (T). The samples were
199 collected excluding the upper 3 cm of topsoil and the rhizosphere area. The analyses were performed
200 with the same methods exploited for the chemical characterization of substrates prior to the
201 experiment (see **Supplementary material - Section II**). Chemical analyses of the leaves were

202 conducted at the end of the second month, after the biomass measurement, in order to assess the
203 content of nitrogen, phosphorus and potassium in the leaves. In the case of the samples from the
204 sandy soil, the measurements were performed on samples treated with one liquid digestate (P), one
205 solid digestate (D) and the mineral fertilizer (M). It was not possible to analyze samples from the
206 negative control (T) due to the low biomass production. On the peat substrate, it was possible to
207 evaluate N-P-K content not only in the P, D, and M samples, but also in the negative control ones (T).
208 The plant samples were firstly processed with a humid digestion protocol (Mills and Jones Jr, 1996).
209 Then, nitrogen was measured through the Kjeldahl method, phosphorus was evaluated through
210 colorimetry (molybdovanadate method) and potassium was quantified by Atomic Absorption
211 Spectroscopy (AAS). Finally, the N, P and K percentages were used to calculate the mean total
212 element present in the epigeal part of the plant using the following formula:

$$\frac{\text{dry sample biomass (g)}}{N, P, K \text{ in sample (\%)}} * \frac{100}{1000} = \text{total N, P, K presence in single sample (mg)}$$

213 Using data of soil and leaves chemical analyses, nitrogen apparent balance was calculated as
214 reported in **Supplementary material – Section III**.

215 2.5. Statistical analysis

216 The experimental data were subjected to statistical analyses. Two-way ANOVA was used to compare
217 the average results of different treatments on plant measurements. Differently, one-way ANOVA was
218 used to compare the mean results of different treatments on the chemical analyses of soils and leaf
219 nutrient content. After the ANOVA, Tukey's post-hoc test ($P < 0.05$) was performed. The statistical
220 software R (version 3.5.1 - Feather Spray - 2018) was used for all statistical analysis.

221 3. Results

222 3.1. Plant measurements

223 3.1.1. Dry biomass

224 On the sandy soil at the end of the first month, the dry biomass of the tomato plants grown with
225 digestates did not show any difference between each other. Despite the absence of significant
226 differences, it must be pointed out that biomass of S was 28.7 and 12.7-folds higher than control (T)
227 and mineral fertilizer (M), respectively. At the end of second month, all digestates (P, S, C, D) showed
228 a dry biomass production significantly higher (26.7, 33, 35.3 and 37.5-folds, respectively) than control.
229 At the same time, S, C and D showed a higher biomass than mineral fertilizer (2.9, 3.1 and 3.3-folds,
230 respectively). At the end of third month, dewatered SSADs proved to be the most productive
231 treatments, with C and D displaying the highest yields (10.23 g and 10.97 g). Their biomasses
232 doubled mineral fertilizer one (5.13 g), which was only comparable to the biomass produced by plants
233 treated with SSADs after two months. Furthermore, C and D yields were 16 and 17-folds higher than
234 T (0.64 g), respectively (**Figure 1.A**).

235 On the peat substrate, no significant differences between treatments were appreciable within the
236 same month. The only significant differences emerged between biomass values between three
237 different months (**Figure 1.B**).

238 Results of ANUE showed significant differences only on sandy soil (**Table S3**). Moreover, it must be
239 pointed out that ANUE values of SSADs in sandy soil were up to 23, 3.5 and 2.4-folds higher than
240 mineral ones after one, two and three months after sowing, respectively.

241 3.1.2.Height

242 On the sandy soil, no differences in plant height were present at the end of the first month.
243 Nevertheless, S treatment revealed the tallest tomato plants, up to 2.6 and 2.5-folds higher than T
244 and M. In the second month, all SSADs-treated plants were significantly taller than control and
245 mineral fertilizer, with D treatment displaying a height 6 and 2.1-folds higher than T and M,
246 respectively. After three months, the mean height of T was still the lowest. The mean height of the
247 plants grown on P and D was comparable to plants grown on mineral fertilizer. Plants grown with S
248 and C treatments had a statistically higher height than mineral fertilizer (M). It is worth highlighting that
249 the mean heights of the plants grown on all digestates was at least 3.5-folds higher than the control
250 ones (**Figure 2.A**).

251 On the peat substrate, no significant differences were observed between the different treatments
252 within the same month. The only significant differences emerged between the height of the samples
253 between three different months (data not shown).

254 3.1.3. Leaves and inflorescences

255 After the first month, the plants grown on sandy soil in presence of D and C treatments showed a
256 number of leaves comparable to control and minerally fertilized plants. On the other hand, samples
257 from liquid SSADs (P and S) revealed a higher mean leaf number than control. After two months, the
258 leaves number on plants grown with digestates was significantly higher only than negative control
259 plants. At the end of the experiment, samples from S and D treatments showed the highest number of
260 leaves, which were not statistically different from samples from C treatment. Plants grown with P had
261 similar number of leaves than C and mineral fertilizer, while leaves number in negative control was
262 still the lowest one (**Figure 2.B**).

263 With regards to the number of inflorescences, no plant on sandy soil showed flowers one month after
264 sowing. At the end of the second month, plants in T and M were still not revealing any flower.
265 Differently, P, S, C and D had some inflorescences, but no significant difference between treatments
266 was present. At the end of the experiment, negative control plants still did not show any flower. Plants
267 treated with P and S had a number of inflorescences statistically similar to mineral fertilizer. The
268 highest number of inflorescences was found on C and D treatments (**Figure 2.C**).

269 As regards the number of leaves and inflorescences of plants grown on peat substrate, no differences
270 between treatments at the same month were highlighted by statistical analysis (data not shown).

271 3.1.4. Chlorophyll Content Index (CCI)

272 On sandy soil, leaves dimension after one month was too small to measure CCI. At the end of second
273 month leaves of plants treated with P, S and C showed a CCI higher than control and comparable to
274 mineral fertilizer. The mean CCI value of plants grown with D digestate was statistically higher than
275 mineral fertilizer (M) but comparable to the others SSADs. CCI measures performed at the end of
276 third month revealed a substantial decrease in CCI values registered in all SSADs and in mineral

277 fertilizer, whose values were not significantly different from the control. The only significant difference
278 emerging at the endpoint was between P and mineral fertilizer measure. (**Figure 2.D**).

279 On peat substrate, the only differences were recorded between the CCI measure of D and P at the
280 end of second month, and S and control at the end of the third month (data not shown).

281 3.1.5. Infra-red gas analyzer (IRGA)

282 As regards IRGA measurements, on sandy soil the lowest A_N value was found in control, where
283 significantly higher values were recorded on C and S. Detailed results are reported in **Supplementary**
284 **material - Section V (Table S4)**.

285 3.2. Chemical analysis

286 3.2.1. Substrates analyses

287 Results of chemical analyses performed on the sandy soil after two months from treatments
288 application are summarized in **Table 2A**. SSADs showed all an intermediate mean pH included
289 between control (8.3) and mineral fertilizer samples (8.0). OM was significantly higher in P, C and D
290 treatments than in S, mineral fertilizer and control. As expected, values of organic carbon showed a
291 trend similar to OM. Total nitrogen (Kjeldahl) was lower in control, mineral fertilizer and S than P, C
292 and D treatments. All results of nitrite analysis were below quantification limits (QL). Nitrates were
293 detectable only in S, C and D treatments, showing very low concentrations (between 1 and 4 mg/kg)
294 with respect to M sample (60 mg/kg). Organic nitrogen values were roughly similar to total Kjeldahl
295 nitrogen ones. Regarding C/N ratio, the lowest value was calculated in control and mineral fertilizer,
296 while all SSADs revealed higher values. Olsen phosphorus was below QL in T and M samples;
297 differently, phosphorous content in samples treated with SSADs was higher. The lowest value of
298 exchangeable calcium was observed in S samples followed by negative control, D, P, C and mineral
299 fertilizer. A great difference in exchangeable sodium content was found between negative control
300 samples and all the treatments. Available zinc ranged between 0.21 mg/kg in control samples, and
301 1.00 mg/kg in D ones, with samples treated with liquid SSADs and mineral fertilizer showing an
302 intermediate behavior. Digestates showed intermediate values of CEC, included between control

303 (2.81 cmol/kg) and mineral fertilizer samples (3.58 cmol/kg). Values of electrical conductivity,
304 ammonia nitrogen (NH_4^+), exchangeable K, exchangeable Mg, available Mn and available Cu did not
305 show any significant difference between treatments on sandy soil.

306 Results of chemical analyses performed on peat substrate two months after treatments application
307 are summarized in **Table 2B**. pH values ranged from a minimum of 6.6 (M) to a maximum of 7.4 (D).
308 Total Kjeldahl nitrogen was lower in control samples and gradually increased along with the dry
309 matter of SSADs; the highest value was displayed by mineral fertilizer samples. Organic nitrogen
310 values were analogous to total Kjeldahl nitrogen in terms of values, trend and differences between
311 treatments. Nitrites, nitrates, extractable Mn, Cu and Zn were below detection limits. All other
312 parameters did not show any significant difference.

313 3.2.2. Leaf analysis

314 On sandy soil, chemical characterization of leaves showed a concentration of nitrogen and potassium
315 in P and D significantly lower than mineral fertilizer samples. As regards phosphorous, no significant
316 difference emerged. The total nitrogen accumulated in leaves in D plants was significantly higher than
317 in P ones. The mean phosphorous uptake by plants was significantly different across D, P and M
318 samples. Finally, the potassium uptaken in leaves did not show significant differences between thesis
319 (**Table 3A**).

320 On plants grown on peat substrate, concentrations and total uptake of both nitrogen and potassium
321 on control, P and D were statistically similar to each other, but they resulted lower in comparison with
322 mineral fertilizer ones. Concentration and total uptake of phosphorous in leaves, control showed the
323 lowest values while D samples the highest ones (**Table 3B**).

324 4. Discussion

325 4.1. Agronomic and physiological evaluations

326 For many years extensive studies and reviews have shown that soil and plant benefit from SS.
327 Indeed, SS is a good source of macro and micro nutrients as well as of OM; this enhances soil fertility
328 and, as a consequence, crop production even in a more effective way than commercial fertilizers

329 (Singh and Agrawal, 2008). The results of the present work were in agreement with literature and the
330 better performances of SS compared to inorganic fertilizers have been confirmed. **Table 4** shows
331 technical details and results of other works dealing with SS treatment of tomato plant with pot
332 experiments. It is important to notice that not only SSAD application rate was considerably lower in
333 the present work, but also that the results obtained were remarkably higher. For instance, biomass
334 and height of treated tomato plants at two months after sowing were up to 37.5 and 6-folds higher,
335 respectively, than control plants (corresponding to an increase of 3652% and 500%), results never
336 reached before in other works on tomato plants. Interestingly, fertilizing performances of SSAD also
337 overcame the ones of mineral fertilizer, especially one month after sowing, when S treatment revealed
338 biomass and height of tomato plants up to 12.7 and 2.5-folds higher than M. From here on out,
339 differences between SSAD treatments and M samples were less accentuated, probably because
340 nutrients release of the mineral fertilizer was faster after an initial “lag” phase. As a corollary, biomass
341 values were reflected by ANUE ones, which were higher than the ones reported in literature for
342 tomato plants grown in pot under greenhouse conditions treated with a 10-folds higher nitrogen
343 application (Wang et al., 2013). Improvement in terms of leaves number and chlorophyll content were
344 less intense, but still higher than the examples reported in literature (Bakshi et al., 2019; Elloumi et al.,
345 2016; He et al., 2016; Hossain et al., 2015).

346 To a broader extent, results of the present study in terms of biomass and plant height can be
347 compared to other works conducted with a similar experimental setup but exploiting different model
348 species. In order to biomass, the general trend was an increase in dry matter ranging usually between
349 4 (*Capsicum annuum* L.; Pascual et al., 2008) and 16-folds (*Triticum aestivum* L.; Eid et al., 2019)
350 more than untreated control. The findings of the present work confirmed and went beyond these
351 results, considering also that the most used SS application rates ranged between the dosage used in
352 this work and a 35-folds higher one (Eid et al., 2019). On the other hand, the improvements in plant
353 height were in line with the results obtained by Eid and colleagues on cucumber (*Cucumis sativus* L.)
354 (2017) and wheat (*Triticum aestivum* L.) (2019), reporting a stem length improvement up to 3 and 6-
355 folds, respectively, over untreated control. The only case with a striking higher biomass production
356 was described for the sunflower (*Heliantus annuus* L.), whose production increased up to 125-folds

357 more than the untreated control. However, the SS dosage was up to 35-folds higher than the present
358 study. Moreover improvement in terms of height was comparable to the present work (Bourioug et al.,
359 2018). Taking into account the works using SS dosages comparable to 170 kg N/ha, the majority
360 were open field experiments. For instance, triticale (*X Triticosecale Wittmack*) (Kchaou et al., 2018)
361 revealed a biomass increase of 2-folds. Furthermore, results of the present work corroborate positive
362 effects on biomass of SS application on soils poor in nutrients (Walter et al., 2000) and strongly
363 alkaline (Zuo et al., 2019).

364 SSAD application on tomato crops resulted also in an augmented number of leaves and
365 inflorescences with respect to control and mineral fertilizer. Moreover, inflorescences number of
366 SSAD-treated plants increased from 2 to 3-folds over the last month. These findings were in general
367 agreement with other results reported on tomato grown in presence of SS (Bakshi et al., 2019),
368 despite the higher treatment dosages.

369 Number of leaves and inflorescences are developmental parameters considered also with other plant
370 species when testing the fertilizing effects of SSAD. For instance, Eid and colleagues (2017)
371 registered on cucumber a boost in the number of leaves of more than 2-folds, which is in line with the
372 results of the present work. Similar outcomes have been reported in terms of number of flowers in
373 common bean (*Phaseolus vulgaris* L.)(Fernández-Luqueño et al., 2010) and marigold (*Tagetes erecta*
374 L.)(Solanki et al., 2017) grown in SS dosages lower and higher, respectively, than the present work.
375 In contrast with these results, Tariq and co-workers (2012) described a decrease up to 60% in flowers
376 number in *Dahlia x hortensis*, whose growth had probably been compromised by an excessive SS
377 dosage.

378 Results of the present work confirmed the positive effects of SS application on net photosynthesis
379 (Bourioug et al., 2018; Pascual et al., 2008) and chlorophyll content. Leaf chlorophyll content was
380 directly correlated with indirect chlorophyll measurements such as readings through SPAD and CCI-
381 meters (Xiong et al., 2015), whose value can be compared to each other with the equations proposed
382 by Parry and colleagues (2014). Application of SSAD improved chlorophyll content values of tomato
383 plants grown on sandy soil at the end of second month, as well as dry biomass and net
384 photosynthesis (A_N). This beneficial effect has been already observed also in sunflower (Bourioug et

385 al., 2018), sorghum (Alvarenga et al., 2016) and triticale (Kchaou et al., 2018). On the other hand,
386 literature provides examples of reduction of leaf chlorophyll content in tomato after treatment with SS
387 (Elloumi et al., 2016), which is probably due to the excessive heavy metals presence in the used SS
388 (Singh and Agrawal, 2007). However, this aspect was likely not linked with the reduction of chlorophyll
389 content over time observed in the present study. Indeed, this phenomenon has been already
390 observed in other SS-treated plant species, such as common bean (Fernández-Luqueño et al., 2010).
391 A possible explanation of this reduction in CCI at the end of the third month could be the deficiency of
392 nutrients in soil. A second hypothesis for CCI decrease has been proposed by de Oliveira and co-
393 workers (2017): after the initial blooming of the plant, gradual degradation of chlorophyll occurs due to
394 the beginning of the fruit development phase, which induces a metabolic change in the plant, with a
395 more sustained nutrients accumulation in the fruit. Taking into account the relationship between leaf
396 nitrogen and chlorophyll content (Xiong et al., 2015), a third justification for CCI decrease can be
397 provided by the so-called nitrogen dilution curve. In fact, biomass increase in tomato plant was
398 accompanied by a reduction in nitrogen concentration (and, consequently in chlorophyll content)
399 because the structural compartment (lower in N%) becomes proportionally more massive than
400 metabolic active one (higher in N%) (Tei et al., 2002).

401 4.2. Chemical analysis

402 4.2.1. Substrates analyses

403 The application of SS on soil can affect different physical and chemical soil characteristics (Epstein,
404 2002). Likewise, many changes were documented in this experiment (both on sandy soil and on peat
405 substrate) two months after treatments application. Although peat substrate was low in nutrient
406 content, it showed a consistently higher amount of microelement than sandy soil. Moreover, peat
407 substrate has many other advantages such as lightweight, high water holding capacity and high air
408 space (Gruda et al., 2016). All these peculiarities most probably contributed to the minor differences
409 registered on peat substrate.

410 Soil analysis results revealed a change in soil pH after the treatments application. Many works
411 reported an increase (Bayoumi Hamuda et al., 2009; Ferreiro-Domínguez et al., 2011) or a decrease

412 (Mosquera-Losada et al., 2016; Singh and Agrawal, 2007) in soil pH. In the present work, acidification
413 occurred in treated sandy soil samples, probably due to both the lower pH of SSADs and the nitrogen
414 mineralization (Rasouli-Sadaghiani and Moradi, 2014). In particular, the nitrification process ($\text{NH}_4^+ \rightarrow$
415 NO_3^-) (Stamatiadis et al., 1999) induces the release of H^+ in soil solution media and the leaching of
416 NO_3^- by water (Whitehead, 1995). Another conceivable theory for soil acidification in SSAD-treated
417 samples could be the generation of organic acids during sewage sludge mineralization (Angin et al.,
418 2012; Bouriouq et al., 2018). Additionally, the low buffering capacity might be yet another plausible
419 effect occurring in the sandy soil case.

420 Electrical conductivity values (both on sandy soil and on peat substrate) did not statistically change
421 after treatments application unlike many other works (Bouriouq et al., 2018; Singh and Agrawal,
422 2007), likely due to the consistently lower SSAD application rates. Nevertheless, it must be pointed
423 out that, concerning sandy soil, EC values in M were approximatively doubled compared to SSAD
424 ones, which in turns were somewhat higher than control. High EC of M might be due to the
425 particularly higher concentration of nitrates, likely released as bioavailable form nitrogen by the
426 commercial fertilizer. However, these relatively elevate nitrate amounts were likely not necessary, as
427 confirmed by the better growth parameters and ANUE values of tomato plants growing on SSAD
428 amended soil. On the contrary, excess of nitrates may result in undesired drawbacks such as
429 leaching and hyperaccumulation in plant tissues, feature in agreement with the foliar analyses.
430 Moreover, sodium might have affected EC values both in mineral fertilizer and in SSAD treatments
431 (probably influenced by sodium presence in the digestates). However, Na did not affect the
432 physiological parameters of tomato plants as confirmed by IRGA measurements.

433 The thesis of a possible increasing of soil OM in soils treated with SSADs (Kladivko and Nelson,
434 1979; Perez-Espinosa et al., 1999) was confirmed by the present work. Despite the OM percentage
435 was very low in all samples, the value in SSADs treated theses was higher than control and mineral
436 fertilizer. This may partially justify the better performances of treated samples in term of biomass and
437 height, according to the well-known soil OM benefits on plants growth (Bot and Benites, 2005).

438 CEC significantly increased in SSADs-treated soil, which was probably caused by the OM increment.
439 This effect is even more pronounced on alkaline soils (Bohn et al., 2001) and similar results were
440 found in other works (Angin et al., 2012; Ferreiro-Domínguez et al., 2011).

441 Total N, available P, exchangeable Ca and Na and available Fe and Zn concentrations increased in
442 the sandy soil amended with SSADs due to their higher concentration in SS (Singh and Agrawal,
443 2007).

444 Two months after treatments application, N_{Tot} (Kjeldahl) was higher in C and D than liquid SSADs (P
445 and S), probably due to their solid form that plausibly induced a slower release, both on sandy soil
446 and peat substrate. Other studies revealed that total soil nitrogen can persist in higher concentrations
447 also for longer periods after SSAD treatment application (Bourioug et al., 2015). Anyway, all samples
448 showed a total N content lower than before digestates application. It meant that a remarkable part of
449 nitrogen both already present in sandy soil and added with digestates was absorbed, transformed or
450 lost after two months, as suggested by the apparent nitrogen balance (**Table S2**). Concerning this
451 balance, it worth specifying that no significant difference was found between P, D and M treatments.

452 The significant variation in N and OM content in treated sandy soils changed C/N ratio. The results
453 obtained with SSADs were still low (< 9; Arpa Veneto, 2007), but higher than in control and mineral
454 fertilizer. The small changes in C/N and the relatively low values across treatments likely indicated
455 that nitrogen mineralization could have prevailed over microbial immobilization. Therefore, nitrogen in
456 SSAD treated samples was surely bioavailable and used efficiently by plants, as also confirmed by
457 ANUE values. However, it should be also noticed that mineralisation was likely a slow nitrogen
458 release process, as evidenced by soil nitrate and leaf nitrogen analyses. Indeed, these evidenced that
459 nitrogen was much more bioavailable in M treatments, but less efficiently utilizable, according to
460 ANUE values.

461 In all SSADs treated soils, the available P was higher than control and mineral fertilizer. Considering
462 that the different dosages were normalized on N dosage per each thesis, the difference in P content
463 between the samples treated with SSADs can be explained by the different percentages of P in the
464 four SSADs. This diversity could also explain the differences among different treatments on
465 physiological parameters of tomato. Moreover, the addition of OM probably enhanced the availability

466 of P in soil treated with SSADs (Fekri et al., 2011). In fact, this can increase the abundance and the
467 activity of microorganisms, favoring P capture (Nobile et al., 2019). Similar results in increase of soil P
468 were obtained by Singh and Agrawal (2007) and Walter and colleagues (2000).

469 For what it concerns K, no differences were registered in soil after digestates application, probably
470 due to their low concentration in this macronutrient. These results agree with other works (Bourioug et
471 al., 2015; Walter et al., 2000).

472 Many SS are rich of Ca due to the stabilization by means of liming (Epstein, 2002). Although the
473 SSADs exploited in this work did not undergo Ca addition at WWTP level, its content was pretty high
474 (> 4.64% D.M.). Considering the medium content in the initial soil, exchangeable Ca increased in
475 some cases in treated soils, confirming the results of Ferreiro Dominguez and Singh (Ferreiro-
476 Domínguez et al., 2011; Singh and Agrawal, 2007).

477 A significant increase of exchangeable Na was measured in all treated soils due to the sodium
478 percentage in SSAD and confirmed by two abovementioned works (Ferreiro-Domínguez et al., 2011;
479 Singh and Agrawal, 2007). The excess of Na is a well-known limiting factor for plants growing (Jones
480 Jr., 2012) but Na has been recently defined as a “new beneficial element” (Morgan, 2000) that, in
481 small quantities, can increase tomato yields (Jones Jr., 2012).

482 The consistent presence of Fe and Zn in SSADs likely provoked the increase in their concentration in
483 sandy soil, confirming the results of Angin and colleagues (2012).

484 4.2.2. Leaf analysis

485 In some cases, in literature the use of SS enhanced the percentage of macronutrients in leaves
486 (Angin et al., 2012; Zuo et al., 2019), in other ones no change took place (Kotecki et al., 2014; Pinna
487 et al., 2009) and still in other ones concentration increased only for some nutrients (Bakshi et al.,
488 2019; De Andres et al., 2010). This work belongs to the third category, since only foliar P% and total
489 uptaken P of control plants grown on peat substrate were significantly lower than SSADs ones. On
490 sandy soil, content of uptaken P was significantly higher in D and P treatments, which was likely
491 influenced by the phosphorous amount in the initial application. Nevertheless, no significant
492 differences emerged in foliar P% despite the difference in uptaken P content between SSADs and

493 mineral fertilizer: probably, the controlled nutrient release of the mineral fertilizer compensated the
494 higher quantity of P in the SSADs. Moreover, it could be inferred that differences in foliar
495 macronutrient content could have been appreciated between control and treated samples on sandy
496 soil. However, the too low biomass of untreated samples made impossible this investigation.

497 The total amount of N and K uptaken in leaves had varied results. On sandy soil, D samples revealed
498 a significantly higher N content than P ones due to the different biomass production. Concerning
499 plants grown on peat substrate, P and D showed a nitrogen plant uptake similar to negative control,
500 but lower than mineral fertilizer, likely due to the characteristics of the fertilizer, such as the controlled
501 nutrient release and the presence of readily bioavailable nitrogen forms. As regards K, despite its
502 higher amount in mineral fertilizer, total K uptaken in leaves did not result significantly different
503 between the treatments applied on sandy soil, due to the different aboveground biomass production.
504 On the other hand, on peat substrate, the $K_{\text{Extractable}}$ content of plant with mineral fertilizer was the
505 highest considering the similar biomass production.

506 5. Conclusions

507 In the present work, pot experiments under greenhouse conditions on two different substrates were
508 performed to evaluate fertilizing effects of four different SSADs over a time span of three months. The
509 application of these digestates clearly highlighted beneficial effects on different growth parameters of
510 tomato plants, especially when cultivated on a sandy, alkaline and poor (in nutrient and OM) soil. For
511 instance, it is important to point out that plant biomass and height reached values up to 37.5 and 6-
512 folds, respectively, higher than untreated control; additionally, SSAD-treated plants showed values of
513 biomass and height up to 12.7 and 2.5-folds, respectively, higher than mineral treatment, indicating
514 that SSAD could be a valuable alternative to mineral fertilizers to boost fertility in poor and sandy soils
515 . Moreover, the present work confirmed the thesis of the enhancement of soil OM with the use of
516 SSAD. Furthermore, it is important to notice the increments of some macro- (nitrogen, phosphorous
517 and calcium) and micro-nutrients (iron and zinc) in sandy soil, showing significant differences with
518 respect to untreated control. Nevertheless, some of the registered values were low and it can be
519 reasonably assumed that most of nutrients had already been assimilated to let the plant grow. This

520 aspect was confirmed by leaves analysis, which showed a remarkable uptake in N, P and K by
521 tomato plants. With respect to these macronutrients, it is worth emphasizing that the experiment was
522 designed to administer plants, across the different treatments, the same nitrogen dosage as sludge
523 application rate is usually based on plants nitrogen requirements. However, the differences in SSADs
524 composition implied a remarkable imbalance in terms of other nutrients and OM. Hence, we can
525 assume that these differences likely influenced plant growth, providing consistent differences between
526 different theses.

527 Future work should include on one side a deeper analysis of the issues tackled in the present paper,
528 and on the other hand it should consider also related aspects. Concerning the formers, chemical
529 characterization of the treated substrates and plants should be carried out in a time-course fashion,
530 allowing to properly describe the mass balance of the elements (including the study of leaching
531 effects) and their release dynamics over time. Consequently, it should allow a more detailed
532 evaluation of the fertilizing indexes (e.g. ANRE, ANUE). As regards the new related aspects to be
533 addressed, soil application of SSAD should be explored both analyzing the presence of organic
534 pollutants (e.g. antibiotics, EDC) as well as considering microbiological aspects, such as the effects
535 on microbial communities and the study of metagenomics and metatranscriptomics traits (e.g.
536 antibiotics resistance genes).

537 Despite reserves and resources for N, P and K appear adequate for the near future, it is necessary to
538 find less impactful solutions to produce fertilizers in the short term. In this way, the reuse of SS can
539 reduce the negative effects connected by the extraction, manufacturing and the use of mineral
540 fertilizers derived from non-renewable resources. Furthermore, this experiment showed how the
541 positive effects of SSADs are emphasized if applied on a poor alkaline soil.

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768 **Table 1.** Physiscal and chemical anlalysis of soil and peat used in the present work. CEC: Cation-Exchange Capacity; AAS: Atomic Absorption
 769 Spectroscopy.

Sandy soil			Peat substrate		
Parameter	Unit	Value	Parameter	Unit	Value
Stones	-	absent	Stones	-	-
Sand (2.0 - 0.020 mm)	%	94 ± 2	Sand (2.0 - 0.020 mm)	-	-
Silt (0.020 - 0.002 mm)	%	3 ± 1	Silt (0.020 - 0.002 mm)	-	-
Clay (< 0.002 mm)	%	3 ± 1	Clay (< 0.002 mm)	-	-
Texture	-	sandy	Texture	-	-
pH	-	8.2 ± 0.16	pH	-	6.2 ± 0.1
Electrical conductivity	dS/m	0.131 ± 0.018	Electrical conductivity	dS/m	0.722 ± 0.146
Organic matter	%	0.38 ± 0.12	Organic matter	-	-
Organic carbon	%	0.22 ± 0.07	Organic carbon	-	-
N - Tot (Kjeldahl)	g/kg	0.29 ± 0.09	N - Tot (Kjeldahl)	%	0.42 ± 0.06
N - NO₂⁻	mg/kg	< 0,2	N - NO₂⁻	mg/l	< QL
N - NO₃⁻	mg/kg	6.33 ± 1.53	N - NO₃⁻	mg/l	30.4 ± 7.2
N - NH₄⁺	mg/kg	3 ± 1	N - NH₄⁺	mg/l	1.3 ± 0.3
N - Org	g/kg	0.29 ± 0.09	N - Org	%	0.4 ± 0.40
C/N		7.6 ± 0.2	C/N		-
P_{Olsen}	mg/kg	1.8 ± 1.3	P_{extractable}	mg/l	8.1 ± 2.3
K_{exchangeable}	mg/kg	18 ± 1	K_{extractable}	mg/l	41.1 ± 6.8
Mg_{exchangeable}	mg/kg	15 ± 5	Mg_{extractable}	mg/l	28 ± 7
Ca_{exchangeable}	mg/kg	675 ± 27	Ca_{extractable}	mg/l	36 ± 8
Na_{exchangeable}	mg/kg	6 ± 3	Na_{extractable}	mg/l	16 ± 11
Fe_{available}	mg/kg	6.7 ± 1.1	Fe_{extractable}	mg/l	0.79 ± 0.21
Mn_{available}	mg/kg	6.5 ± 3.0	Mn_{extractable}	mg/l	0.15 ± 0.04
Cu_{available}	mg/kg	0.69 ± 0.29	Cu_{extractable}	mg/l	< QL
Zn_{available}	mg/kg	0.47 ± 0.29	Zn_{extractable}	mg/l	0.02 ± 0.00
CEC	cmol/kg	3.65 ± 0.35	CEC		-

770 **Table 2.** Chemical characterization performed two months after treatments application on sandy soil (A) and on peat substrate (B). Data are
 771 expressed as mean \pm standard deviation. Asterisks mean significant differences according to ANOVA test (*, **, *** differences between means
 772 significant at $P \leq 0.05$, 0.01 and 0.001, respectively). CEC, cation exchange capacity; QL, quantification limit.

773 **A**

Parameter	Unit of measure	Control (T)			Primary (P)			Secondary (S)			Centrifuged (C)			Dried (D)			Mineral fertiliser (M)		
pH	-	8.3	\pm 0.1	***	8.2	\pm 0.1	***	8.2	\pm 0.1	***	8.1	\pm 0.1	***	8.1	\pm 0.1	***	8.0	\pm 0.1	***
Electrical conductivity	dS/m	0.155	\pm 0.020		0.219	\pm 0.032		0.201	\pm 0.010		0.197	\pm 0.023		0.198	\pm 0.025		0.399	\pm 0.146	
Organic matter	%	0.16	\pm 0.01	***	0.24	\pm 0.01	***	0.18	\pm 0.02	***	0.25	\pm 0.02	***	0.26	\pm 0.01	***	0.16	\pm 0.02	***
Organic carbon	%	0.09	\pm 0.00	***	0.14	\pm 0.00	***	0.11	\pm 0.01	***	0.14	\pm 0.01	***	0.15	\pm 0.00	***	0.10	\pm 0.01	***
N - Tot (Kjeldahl)	g/kg	0.17	\pm 0.01	***	0.19	\pm 0.01	***	0.15	\pm 0.01	***	0.20	\pm 0.00	***	0.22	\pm 0.01	***	0.17	\pm 0.01	***
N - NO ₂ ⁻	mg/kg	< QL			< QL			< QL			< QL			< QL			< QL		
N - NO ₃ ⁻	mg/kg	< QL			< QL			1	\pm 1	*	4	\pm 4	*	2	\pm 1	*	60	\pm 40	*
N - NH ₄ ⁺	mg/kg	< QL			< QL			< QL			1	\pm 1		2	\pm 0		1	\pm 0	
N - Org	g/kg	0.17	\pm 0.01	***	0.19	\pm 0.01	***	0.15	\pm 0.01	***	0.20	\pm 0.00	***	0.22	\pm 0.01	***	0.17	\pm 0.01	***
C/N	-	5.5	\pm 0.2	**	7.3	\pm 0.6	**	6.9	\pm 0.5	**	7.3	\pm 0.5	**	7.1	\pm 0.3	**	5.8	\pm 1.0	**
P _{Olsen}	mg/kg	< QL			4.2	\pm 0.5	**	10.3	\pm 0.8	**	15.4	\pm 4.6	**	36.2	\pm 11.4	**	< QL		
K _{exchangeable}	mg/kg	14	\pm 3		11	\pm 3		12	\pm 1		12	\pm 2		9	\pm 1		13	\pm 2	
Mg _{exchangeable}	mg/kg	11	\pm 2		21	\pm 7		26	\pm 3		25	\pm 4		22	\pm 4		25	\pm 2	
Ca _{exchangeable}	mg/kg	524	\pm 26	*	594	\pm 25	*	491	\pm 62	*	626	\pm 94	*	579	\pm 48	*	646	\pm 62	*
Na _{exchangeable}	mg/kg	16	\pm 1	***	35	\pm 2	***	33	\pm 3	***	26	\pm 3	***	32	\pm 3	***	27	\pm 5	***
Fe _{available}	mg/kg	5.7	\pm 0.1	***	7.6	\pm 0.5	***	9.3	\pm 0.2	***	11.2	\pm 0.7	***	12.5	\pm 0.4	***	6.1	\pm 0.21	***
Mn _{available}	mg/kg	5.1	\pm 0.4		14.8	\pm 16.6		33.7	\pm 2.5		35.8	\pm 27.1		20.5	\pm 28.1		5.2	\pm 0.21	
Cu _{available}	mg/kg	0.40	\pm 0.08		0.47	\pm 0.13		0.60	\pm 0.06		0.85	\pm 0.12		0.96	\pm 0.29		0.40	\pm 0.01	
Zn _{available}	mg/kg	0.21	\pm 0.03	***	0.36	\pm 0.04	***	0.37	\pm 0.02	***	0.73	\pm 0.08	***	1.00	\pm 0.07	***	0.43	\pm 0.05	***
CEC	cmol/kg	2.81	\pm 0.13	*	3.32	\pm 0.17	*	2.83	\pm 0.33	*	3.47	\pm 0.43	*	3.24	\pm 0.20	*	3.58	\pm 0.32	*

QL: N - NO₂⁻ = 0.2 mg/kg; N - NO₃⁻ = 1 mg/kg; N - NH₄⁺ = 1 mg/kg; P = 1 mg/kg.

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775 **B**

Parameter	Unit of measure	Control (T)	Primary SSAD (P)	Secondary SSAD (S)	Centrifuged SSAD (C)	Dried SSAD (D)	Mineral (M)
pH	-	7.0 ± 0.4 *	6.7 ± 0.3 *	7.2 ± 0.2 *	6.9 ± 0.2 *	7.4 ± 0.3 *	6.6 ± 0.2 *
Electrical conductivity	dS/m	0.235 ± 0.040	0.436 ± 0.220	0.183 ± 0.038	0.495 ± 0.134	0.225 ± 0.074	0.523 ± 0.202
N - Tot (Kjeldahl)	% D.M.	0.23 ± 0.03 *	0.24 ± 0.03 *	0.25 ± 0.03 *	0.28 ± 0.04 *	0.31 ± 0.04 *	0.32 ± 0.04 *
N - NO₂⁻	mg/l	< QL	< QL	< QL	< QL	< QL	< QL
N - NO₃⁻	mg/l	1.0 ± 0.9	1.5 ± 0.5	1.0 ± 0.2	2.6 ± 0.4	1.2 ± 0.3	2.6 ± 1.7
N - NH₄⁺	mg/l	< QL	< QL	< QL	< QL	< QL	< QL
N - Org	% D.M.	0.22 ± 0.03 *	0.24 ± 0.03 *	0.25 ± 0.03 *	0.28 ± 0.04 *	0.31 ± 0.04 *	0.32 ± 0.04 *
P extractable	mg/l	0.4 ± 0.1	1.2 ± 0.4	1.2 ± 0.5	0.6 ± 0.3	1.1 ± 0.1	< QL
K extractable	mg/l	2.9 ± 0.3	2.0 ± 0.3	2.4 ± 0.8	2.4 ± 0.4	5.0 ± 2.2	2.6 ± 0.2
Mg extractable	mg/l	8 ± 2	22 ± 16	5 ± 2	24 ± 12	6 ± 3	26 ± 17
Ca extractable	mg/l	13 ± 4	26 ± 16	12 ± 3	33 ± 14	14 ± 3	32 ± 17
Na extractable	mg/l	24 ± 2	29 ± 7	17 ± 3	32 ± 4	23 ± 7	31 ± 4
Fe extractable	mg/l	1.17 ± 0.26	0.52 ± 0.46	0.80 ± 0.13	0.52 ± 0.30	0.73 ± 0.06	0.28 ± 0.11
Mn extractable	mg/l	< QL	< QL	< QL	< QL	< QL	0.03 ± 0.01
Cu extractable	mg/l	< QL	< QL	< QL	< QL	< QL	< QL
Zn extractable	mg/l	< QL	< QL	< QL	< QL	< QL	0.02 ± 0

QL: N - NO₂⁻ = 0.05 mg/l; N - NH₄⁺ = 0.06 mg/l; P = 0.3 mg/l; Mn = 0.03 mg/l; Cu = 0.03 mg/l; Zn = 0.02 mg/l.

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781 **Table 3.** Results of leaves analyses performed after two months after treatments application on sandy soil (A) and on peat substrate (B). Different
 782 letters indicate differences between treatments that are significant at $P < 0.05$ (Tukey HSD). Data are expressed as mean \pm standard deviation.

783 **A**

Parameter	Unit of measure	Primary SSAD (P)			Dried SSAD (D)			Mineral fertilizer (M)		
N	%	1.10	\pm 0.05	b	1.35	\pm 0.28	b	2.95	\pm 0.36	a
	Total (mg)	46.64	\pm 7.45	b	81.08	\pm 15.63	a	60.83	\pm 9.26	ab
P	%	0.14	\pm 0.01	a	0.16	\pm 0.01	a	0.13	\pm 0.02	a
	Total (mg)	5.97	\pm 1.25	b	9.61	\pm 1.26	a	2.56	\pm 0.33	c
K	%	1.46	\pm 0.38	b	1.40	\pm 0.18	b	3.63	\pm 0.57	a
	Total (mg)	61.61	\pm 16.41		84.54	\pm 10.95		74.24	\pm 7.52	

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785 **B**

Parameter	Unit of measure	Control (T)			Primary SSAD (P)			Dried SSAD (D)			Mineral fertilizer (M)		
N	%	1.26	\pm 0.08	b	1.32	\pm 0.04	b	1.29	\pm 0.19	b	2.05	\pm 0.36	a
	Total (mg)	319.01	\pm 11.24	b	323.86	\pm 20.48	b	360.41	\pm 53.17	b	550.22	\pm 102.49	a
P	%	0.23	\pm 0.01	b	0.29	\pm 0.02	ab	0.31	\pm 0.04	a	0.27	\pm 0.02	ab
	Total (mg)	58.02	\pm 7.29	b	69.82	\pm 2.92	ab	85.44	\pm 9.77	a	72.95	\pm 8.78	ab
K	%	1.83	\pm 0.04	b	1.92	\pm 0.04	b	1.74	\pm 0.21	b	2.53	\pm 0.27	a
	Total (mg)	465.11	\pm 37.08	b	470.69	\pm 24.49	b	486.97	\pm 57.85	b	677.03	\pm 53.15	a

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788 **Table 4.** Comparison of the results from other works in literature on the effects of treatment with sewage sludge on tomato plants. Application
789 dosages are shown as reported in the original works; values in brackets indicate how many folds more is the SS application rate with respect to the
790 present study. n.a., not available.

Cultivar	SS typology	Dosage	Kind of experiment	Cultivation substrate	Differences with respect to untreated control				Reference
					Biomass increase	Plant height	Leaves and inflorescences	Chlorophyll content	
Cherry	Digested ^a	10 t/ha (2X)	Pot experiment Greenhouse 16 weeks	Chromosol	<i>Dry biomass</i> : + 20%	10 weeks: + 50% 13 weeks: + 20% 15 weeks: + 7%	n.a.	n.a.	Hossain et al., 2015
Red Robin	n.a.	SS:soil 1:10 (65X)	Pot experiment Growth chamber 120 days	Loamy soil	<i>Fresh biomass</i> Stem: + 70% Leaves: + 142%	+ 43%	<i>Leaves</i> : + 33% <i>Flowers</i> : +130%	<i>Chlorophyll</i> ^b : <i>a</i> : + 18.3% <i>b</i> : + 34.8%	Bakshi et al., 2019
Rio Grande	Aerobically digested	2.5%; 5.0%; 7.5% (11X; 22X; 33X)	Pot experiment Greenhouse 30 days	Sandy soil	<i>Dry biomass</i> : + 180%; + 280%; +140%	n.a	n.a.	<i>Chlorophyll a+b</i> ^b : + 17.5%; - 40%; - 68.5%	Elloumi et al., 2016
n.a.	Aerobically digested	400 - 800 kg N/ha (2.35X; 4.7X)	Pot experiment Greenhouse 90 days	Clay soil	<i>Dry biomass</i> : + 18.6% + 29.6%	+ 19.2%; + 24.5%	n.a.	n.a.	He et al. 2016
Beefsteak	Anaerobically digested (4 typologies: P, S, C, D)	170 kg N/ha	Pot experiment Greenhouse 120 days	Sandy soil	<i>Dry biomass</i> up to + 3652% (D treatment, II month)	up to + 500% (D treatment, II month)	<i>Leaves</i> : up to + 180% (S treatments, I month) <i>Flowers</i> : not observed in untreated control	<i>CCI</i> : up to + 172% (D treatment, II month)	This work
				Peat substrate	<i>Dry biomass</i> : up to + 70% (C treatment, I month)	up to + 24% (P treatment, I month)	n.a.	<i>CCI</i> : up to + 64% (D treatment, III month)	

^aIn this work, no details about the typology of digestion are provided.

^bIn these works, leaf chlorophyll content was evaluated with methods based on extraction with organic solvents followed by spectrophotometrical quantification.

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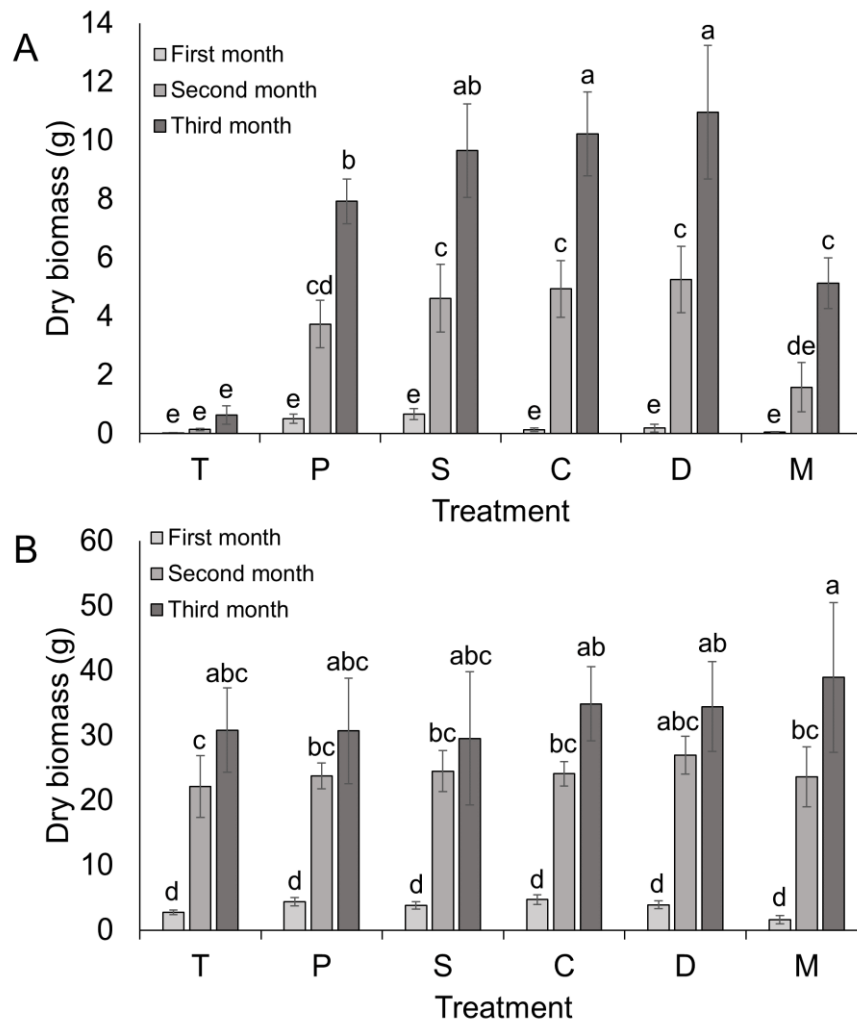
1 **Figure 1.** Mean dry biomasses of *Solanum lycopersicum* L. grown on sandy soil (A) and peat

2 substrate (B) with different treatments among three months. Different letters indicate differences

3 between treatments that are significant at $P < 0.05$ (Tukey HSD). Each error bar represents one

4 standard deviation. T: non-treated, control thesis; P: primary digestate; S: secondary digestate; C:

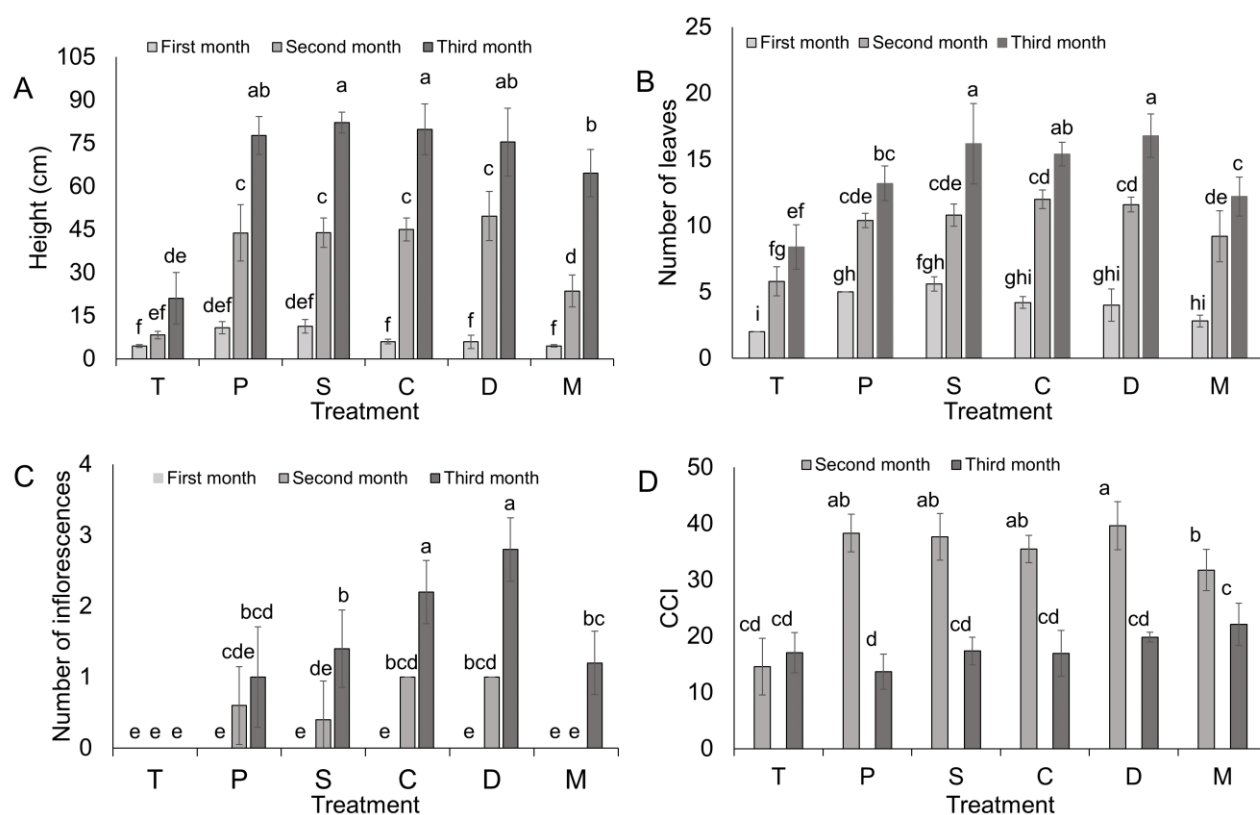
5 centrifuged digestate; D: dried digestate; M: mineral fertilizer.



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1 **Figure 2.** (A) Mean height of *Solanum lycopersicum* L. grown on sandy soil with different treatments
 2 among three months; (B) Mean number of leaves of *Solanum lycopersicum* L. grown on sandy soil
 3 with different treatments among three months; (C) Mean number of flower of *Solanum lycopersicum* L.
 4 grown on sandy soil with different treatments among three months. (D) Mean Chlorophyll Content Index
 5 (CCI) of leaves of *Solanum lycopersicum* L. grown on sandy soil with different treatments among 3
 6 months. Different letters indicate differences between treatments that are significant at $P < 0.05$ (Tukey
 7 HSD). Each error bar represents one standard deviation. T: non-treated, control thesis; P: primary
 8 digestate; S: secondary digestate; C: centrifuged digestate; D: dried digestate; M: mineral fertilizer.



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Supplementary material

of

“Anaerobic digestates from sewage sludge used as fertilizer on a poor alkaline sandy soil and on a peat substrate: effects on tomato plants growth and on soil properties”

Giulio Cristina, Enrico Camelin, Tonia Tommasi, Debora Fino and Massimo Pugliese.

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Section I: Characterization of anaerobic digestates from sewage sludge (SSADs) used

Primary, secondary, centrifuged and dried SSADs were sampled directly at WWTP and stored at 4°C until chemical characterisation and further uses. The chemical analyses were performed according to “Analytical Methods for Fertilizers” by the Italian Ministry of Agriculture and Forestry (M.P.A.A.F., 2006) and “Methods for Analysis of Sewage Sludge by Water Research Institute of National Council of Researches (IRSA-CNR, 1985), unless specified differently. Results are reported in **Table S1**. pH and electrical conductivity were measured on distilled water extracts (1:10 m/v) by potentiometry and conductometry, respectively. Dry matter content and humidity were measured by gravimetry, drying the samples at 105°C until constant weight. Ashes were determined with calcination at 550°C for 5 hours.

Total organic carbon was evaluated as reported on “Official methods of soil analysis” by the Italian Ministry of Agriculture and Forestry (Italian Ministerial Decree, 1999), exploiting the Walkley-Black method: sample digestion with potassium dichromate and sulphuric acid is followed by titration with iron(II) sulphate heptahydrate. Organic matter content was calculated with the Van Bemmelen conversion factor (1.724) (Pribyl, 2010).

Total nitrogen (N_{Tot}) was measured with the Kjeldahl method, which allows to titrate both organic and inorganic forms of nitrogen. Ammonium nitrogen (NH_4^+) was evaluated through distillation with magnesium oxide followed by titration with sulphuric acid, while nitrates (N-NO_3^-) were determined by the means of ionic chromatography. Organic nitrogen (N_{Org}) was then calculated by subtraction: $N_{\text{Org}} = N_{\text{Tot}} - (\text{N-NH}_4^+)$. Other macronutrients (K and P), micronutrients (Ca, Mg, Na, Fe, Mn, B, Zn) and heavy metals (Pb, Cr, Ni, Cu) were extracted with mineral acid digestion and then analysed by the means of inductively coupled plasma optical emission spectrometry (ICP-OES). Other contaminants such as Cd and As were extracted with the same digestion protocol, but analysed with graphite furnace atomic absorption spectroscopy (GF-AAS). Hg was evaluated with hydride generation atomic absorption spectroscopy (HGAAS) after microwave mineralisation, while Cr^{6+} was determined by colorimetry after complexation with diphenylcarbazide.

Table S1. Physicochemical properties of the four anaerobic digestates from sewage sludge used. Last three columns on right specify analysis methods for sewage sludge, Italian law limits for Land application of sewage sludges (Italian Decree Law 99/1992, n.d.), and law limits for heavy metals in fertilizers (Italian Decree Law 75/2010). d.m.b., Dry matter basis.

Parameter	Unit of measure	Anaerobic digestates				Method of analysis	Technique	Italian Law Land application of sewage sludge (D. Lgs 99/92)	Italian Law Discipline on fertilizers (D.Lgs 75/2010)
		Primary (P)	Secondary (S)	Centrifuged (C)	Dried (D)				
Dry matter	%	4.4	4.8	25.8	88.8	Calculation	Calculation		
Humidity	%	95.6	95.2	74.2	11.2	M.P.A.A.F., 2006 Method III.1	Gravimetry		
Ashes	% d.m.b.	35.3	31.5	36.1	35.6	Calculation	Calculation		
pH (1:10)		7.7	7.5	7.3	6.8	M.P.A.A.F., 2006 Method III.3	Potentiometry		
E.C.	mS/cm	0.378	0.36	1.069	1.575	M.P.A.A.F., 2006 Method III.4	Conductometry		
Organic matter	% d.m.b.	64.7	68.5	63.9	64.4	Calculation	Calculation		
TOC	% d.m.b.	37.5	39.7	37.1	37.3	D.M. 13/09/99 GU 248 21/10/199 met. VII.3	Walkley & Black method	>20	
N - Tot	% d.m.b.	7.4	7.5	6.3	5	CNR IRSA 6 Q64 vol.3, 1985	Kjeldahl method	>1.5	
N - Org	% d.m.b.	5.84	6.16	5.33	4.75	M.P.A.A.F., 2006 Method IV.12	Calculation		
N - NO ₃ ⁻	% d.m.b.	<0.01	<0.01	<0.01	<0.01	M.P.A.A.F., 2006 Method IV.12	Ionic chromatography		
N - NH ₄ ⁺	% d.m.b.	1.56	1.34	0.97	0.25	M.P.A.A.F., 2006 Method IV.12	Distillation and titration		
N - org / N - Tot	%	79	82	84	94	Calculation	Calculation		
C/N		5.1	5.3	5.9	7.4	Calculation	Calculation		
P	% d.m.b.	4.16	5.75	6.74	6.26	M.P.A.A.F., 2006 Method VIII	Acid digestion + ICP-OES	>0.4	
K	% d.m.b.	0.55	0.69	0.39	0.18	M.P.A.A.F., 2006 Method VIII	Acid digestion + ICP-OES		
Ca	% d.m.b.	6.46	4.69	5.02	4.64	M.P.A.A.F., 2006 Method VIII	Acid digestion + ICP-OES		
Mg	% d.m.b.	1.78	1.53	1.45	1.16	M.P.A.A.F., 2006 Method VIII	Acid digestion + ICP-OES		
Na	% d.m.b.	1.05	1.03	0.34	0.19	M.P.A.A.F., 2006 Method VIII	Acid digestion + ICP-OES		
B	mg/kg d.m.b.	51	60	52	41	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES		
Zn	mg/kg d.m.b.	918	650	849	719	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES	2500	500
Fe	% d.m.b.	2.43	3.32	3.99	3.48	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES		
Mn	mg/kg d.m.b.	255	190	268	228	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES		
Cu	mg/kg d.m.b.	357	340	406	396	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES	1000	230
Pb	mg/kg d.m.b.	92	70	92	79	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES	750	140
Cr	mg/kg d.m.b.	245	210	245	217	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES	<200*	
Cd	mg/kg d.m.b.	1	0.6	0.8	<0.1	M.P.A.A.F., 2006 Method IX	Acid digestion + GF-AAS	20	1.5
Ni	mg/kg d.m.b.	163	120	155	137	M.P.A.A.F., 2006 Method IX	Acid digestion + ICP-OES	300	100
As	mg/kg d.m.b.	2.8	2.1	0.9	<0.1	M.P.A.A.F., 2006 Method IX	Acid digestion + GF-AAS	<20*	
Hg	mg/kg d.m.b.	<0.1	<0.1	<0.1	<0.1	Internal method	HGAAS	10	1.5
Cr ⁶⁺	mg/kg d.m.b.	<0.1	<0.1	<0.1	<0.1	CNR IRSA 16 Q64 vol.3, 1986	Colorimetry	<2*	0.5

* Values introduced with Italian Law 130/2018

Section II: Method of analysis of sandy soil and peat substrate

The soil used in this study was sampled in Grugliasco (TO), Italy (45°03'58.4"N, 7°35'32.9"E). It was collected within 20 and 100 cm depth, sieved at 2 mm and not previously sterilized.

Physical and chemical analysis were performed according to the official methods of soil analysis of Italian Ministry of Agriculture and Forestry (Italian Ministerial Decree, 1999), except for available Fe, Mn, Cu and Zn. Stones were evaluated by sieving (2 mm) (Method II.1) while soil texture was determined by granulometry (wet sieve analysis; Method II.6). Measure of pH and electrical conductivity, organic matter, nitrogen forms and phosphorous was conducted on an aqueous extract obtained following the Sonneveld method (Sonneveld and Voogt, 2009). pH and electrical conductivity were measured by potentiometry (Method III.1) and conductometry (Method IV.1). Organic carbon was measured with the Walkley-Black method (Method VII.3). Organic matter content was calculated with the Van Bemmelen conversion factor (1.724) (Pribyl, 2010). Total nitrogen was measured with the Kjeldahl method (Method XIV.3). Mineral forms of nitrogen were extracted with an aqueous solution of KCl 2M (Method XIV.4); ammonium was measured through distillation (Method XIV.6), while nitrate and nitrite were quantified through continuous flux colorimetry (Griess-Ilosvay reaction; Method XIV.12 and XIV.13). Organic nitrogen and C/N ratio were obtained by calculation. Available phosphorous was determined by Olsen method (Method XV.3).

Measure of cation exchange capacity (C.E.C.) and exchangeable bases (Na, K, Mg, Ca) were performed on an extract obtained with an aqueous solution of BaCl₂ – triethanolamine (pH 8.2); C.E.C. was determined through complexometric titration (Method XIII.2) while exchangeable bases were measured by the means of flame atomic absorption spectroscopy (FAAS) (Method XIII.5).

Analysis of available Fe, Mn, Cu and Zn was performed according to Italian Ministerial Decree (Italian Ministerial Decree, 1992) Method 37, which exploits the Lindsay-Norwell method, that is an extraction through an aqueous solution of DTPA, CaCl₂ and triethanolamine (pH 7.3) followed by quantification with flame atomic absorption spectroscopy (FAAS).

Peat substrate was mixed with perlite and then sterilised before each application. Chemical characterization of peat substrate was performed on an aqueous extract 1:2 (v/v water/peat

substrate) according to Sonneveld method (Sonneveld and van den Ende, 1971). The analytical methods for peat analysis were all internal methods. pH and electrical conductivity were measured by potentiometry and conductometry. Total Nitrogen was evaluated with Kjeldahl method while organic nitrogen was calculated. Inorganic forms of nitrogen (ammonium, nitrite, nitrate) were measured by colorimetry (indophenol-blue method, diazotization method and dimethylphenol method respectively). Phosphorous were measured by colorimetry (molybdovanadate method). K, Mg, Ca, Na, Fe, Mn, Cu and Zn were measured through flame atomic absorption spectroscopy (FAAS).

Section III: Apparent balance of nitrogen in sandy soil

Apparent N balance in sandy soil (**Table S2**) was calculated according to Yang and co-workers (2020), with some modifications. Total nitrogen was considered as the sum of Kjeldahl nitrogen, nitrites and nitrates per pot. Total initial nitrogen was calculated as the sum of N present in nude sandy soil and N added with treatments. Stored nitrogen in soil was the N still present after two months; stored nitrogen in plants was the N measured in plants (epigeal part) two months after sowing. N loss was calculated as difference between initial and stored nitrogen. Statistical analysis of N loss highlighted no significant differences between treatments. With regards to peat substrate, nitrogen storage was not calculated since no significant differences were observed in biomass and ANUE results.

Table S2. Results of apparent balance of nitrogen in sandy soil. Nitrogen amounts are reported as mean values \pm standard deviation.

Treatment	Initial nitrogen			Nitrogen storage		Loss of nitrogen	
	Soil nitrogen storage (g)	Nitrogen added (g)	Total nitrogen (g)	Soil (g)	Plant (mg)	(g)	%
Primary SSAD	1.16 \pm 0.32	0.39	1.55 \pm 0.32	0.76 \pm 0.03	46.64 \pm 7.45	0.75 \pm 0.04	48.14 \pm 2.51
Dried SSAD	1.16 \pm 0.32	0.39	1.55 \pm 0.32	0.87 \pm 0.04	81.08 \pm 15.63	0.60 \pm 0.03	38.46 \pm 2.24
Mineral fertilizer	1.16 \pm 0.32	0.39	1.55 \pm 0.32	0.91 \pm 0.20	60.83 \pm 9.26	0.58 \pm 0.17	37.45 \pm 11.17

Section IV: Agronomic nitrogen use efficiency (ANUE) calculation

Table S3. Agronomic nitrogen use efficiency (ANUE) on sandy soil samples. Data are reported as mean value \pm standard deviation and are expressed in g gN^{-1} . Different letters indicate differences between treatments that are significant at $P < 0.05$ (Tukey HSD).

Month	SSAD												Mineral fertilizer		
	Primary			Secondary			Centrifuged			Dried					
1	1.26	\pm 0.39	d	1.66	\pm 0.50	d	0.28	\pm 0.16	d	0.43	\pm 0.35	d	0.07	\pm 0.05	d
2	9.32	\pm 2.09	c	11.60	\pm 2.99	c	12.43	\pm 2.51	c	13.25	\pm 2.93	c	3.74	\pm 2.17	d
3	18.89	\pm 1.99	b	25.09	\pm 1.76	a	24.85	\pm 3.70	a	27.99	\pm 3.45	a	11.65	\pm 2.26	c

Section V: Infra-red gas analyzer (IRGA)

Net photosynthesis (A_N) of tomato plants leaves grown on sandy soil showed significant differences between treatments (**Table S4**). The lowest A_N value was found in control ($4.08 \mu\text{mol CO}_2 \text{ m}^{-2}\text{s}^{-1}$), while the significantly highest values were recorded on C ($10.56 \mu\text{mol CO}_2 \text{ m}^{-2}\text{s}^{-1}$) and S ($10.21 \mu\text{mol CO}_2 \text{ m}^{-2}\text{s}^{-1}$). P, D and mineral fertilizer (M) displayed values comparable with both the lowest and highest ones. Moving to the stomatal conductance (g_s) and CO_2 concentration in substomatal cavity (C_i), no statistically significant difference was registered. The overall mean g_s value was $0.25 \text{ mmol H}_2\text{O m}^{-2}\text{s}^{-1}$ while the overall mean C_i value was 276.22 ppm.

As regards IRGA measurements on peat substrate, A_N did not show any significative difference between the treatments and overall mean calculated value was $8.64 \mu\text{mol CO}_2 \text{ m}^{-2}\text{s}^{-1}$. The stomatal conductance of the digestates and of the control was higher than on mineral fertilizer. Concerning C_i , no significant difference between the treatments was found; the overall mean of C_i value was 280.89 ppm.

Table S4. Results of infra-red gas analyzer (IRGA) on plants grown on sandy soil. Different letters indicate differences between treatments that are significant at $P < 0.05$ (Tukey HSD). Data are expressed as mean \pm standard deviation. A_N : net assimilation ($\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$); g_s : stomatal conductance ($\text{mmol H}_2\text{O m}^{-2} \text{ s}^{-1}$); C_i : CO_2 concentration in substomatal cavity (ppm).

Treatment	Parameter	A_N ($\mu\text{mol CO}_2 \text{ m}^{-2}\text{s}^{-1}$)	g_s ($\text{mmol H}_2\text{O m}^{-2}\text{s}^{-1}$)	C_i (ppm)
Control (T)		4.08 \pm 1.87 b	0.17 \pm 0.09	300.11 \pm 10.19
Primary digestate (P)		9.96 \pm 1.12 ab	0.27 \pm 0.06	264.00 \pm 11.35
Secondary digestate (S)		10.21 \pm 1.48 a	0.28 \pm 0.02	269.56 \pm 8.18
Centrifuged digestate (C)		10.56 \pm 2.37 a	0.28 \pm 0.04	260.22 \pm 20.72
Dry digestate (D)		8.29 \pm 3.37 ab	0.26 \pm 0.07	281.56 \pm 27.84
Mineral fertilizer (M)		7.30 \pm 1.97 ab	0.24 \pm 0.07	281.89 \pm 12.36

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Declaration of interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Author contributions

Giulio Cristina: Conceptualization, Methodology, Writing - Original Draft

Enrico Camelin: Writing - Review & Editing, Visualization, Formal analysis, Software

Tonia Tommasi: Data Curation, Project administration

Massimo Pugliese: Conceptualization, Methodology, Validation, Resources, Supervision

Debora Fino: Funding acquisition, Investigation