

1 **Potential short-term losses of N₂O and N₂ from high**
2 **concentrations of biogas digestate in arable soils**

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10 **Abstract.** Biogas digestate (BD) is increasingly used as organic fertiliser, but has a high potential for NH₃
11 losses. Its proposed injection into soils as a counter-measure has been suggested to promote the generation of
12 N₂O, leading to a potential trade-off. Furthermore, the effect of high nutrient concentrations on N₂ losses as they
13 may appear after injection of BD into soil has not yet been evaluated. Hence, we performed an incubation
14 experiment with soil cores in a helium-oxygen atmosphere to examine the influence of soil substrate (loamy
15 sand, clayey silt), water-filled pore space (WFPS; 35, 55, 75%) and application rate (0, 17.6 and 35.2 mL BD per
16 soil core [250 cm³]) on the emissions of N₂O, N₂ and CO₂ after the usage of high loads of BD. To determine the
17 potential capacity for gaseous losses, we applied anaerobic conditions by purging with helium for the last 24 h of
18 incubation. Immediate N₂O and N₂ emissions as well as the N₂/(N₂O+N₂) ratio depended on soil type and
19 increased with WFPS indicating a crucial role of soil gas diffusivity for the formation of nitrogenous gases in
20 agricultural soils. However, the emissions did not increase with the application rate of BD probably due to an
21 inhibitory effect of the high NH₄⁺ content in BD. Our results suggest a larger potential for N₂O formation in the
22 fine-textured clayey silt compared to the coarse loamy sand after applying high concentrations of BD as
23 appearing after injection. However, the loamy sand showed basically a large potential for N₂ formation under
24 anaerobic headspace conditions. Nevertheless, our results show the need for further investigations on the
25 dynamics and the duration of the observed effects and their significance for field conditions.

26 1 Introduction

27 Nitrous oxide (N₂O) is a potent greenhouse gas (Myhre et al., 2013), with agriculture being the largest single
28 source of anthropogenic N₂O emissions, contributing about 4.1 Tg N₂O-N yr⁻¹ or 66% of total gross
29 anthropogenic emissions mainly as a result of mineral nitrogen (N) fertiliser and manure application (Davidson
30 and Kanter, 2014). The generation of nitrogen gas (N₂) is of agronomic interest in terms of nutrient management,
31 since such gaseous losses may imply a significant loss of N from the soil/plant system (Friedl et al., 2016, 2016;
32 Cameron et al., 2013). However, from an environmental stance, N₂ is innocuous and, thus, the preferred type of
33 gaseous N-loss from soil (Davidson et al., 2015). In general, the improvement of N use efficiency and thus the
34 decrease of N losses in crop production are paramount in the presence of challenges like food security,
35 environmental degradation and climate change (Zhang et al., 2015).

36 Digestion residues (biogas digestate, BD) from biogas plants are used as organic amendment in agriculture. But,
37 compared to undigested amendments, digestion results in an increased pH, a higher proportion of ammonium

38 (NH_4^+) and a narrowed C to N of the BD (Möller and Müller, 2012). This altered chemical properties may
39 promote biochemical reactions in the soil that are responsible for the formation of gaseous N species like N_2O ,
40 nitric oxide (NO), N_2 and ammonia (NH_3) (Nkoa, 2013).

41 Significant losses of N as NH_3 can occur within the first hours after manure application (Quakernack et al.,
42 2012). To reduce NH_3 losses, the application of BD by injection is recommended, but this measure can
43 simultaneously increase the potential for N_2O losses compared to surface-application (Wulf et al., 2002; Velthof
44 and Mosquera, 2011). On the one hand, high NH_4^+ concentrations in the injection band promote nitrification,
45 which is a significantly O_2 consuming process releasing N_2O (Christensen and Rowe, 1984). On the other hand,
46 increased amounts of C in the injection band also promote respiration and, thus, additionally deplete the O_2
47 supply (Dell et al., 2011). Altogether, the conditions during the initial phase after injection of BD foster
48 microsites favourable for microbial denitrification, which may promote also the formation of N_2 due to anaerobic
49 conditions (Köster et al., 2015; Webb et al., 2010).

50 There is a wealth of biotic and abiotic processes in soils that produce N_2O and N_2 , depending on mineral N
51 content, carbon (C) availability as well as on temperature, most of which are enhanced by anaerobic or at least
52 suboxic conditions (Butterbach-Bahl et al., 2013). The amounts and the relative share of N_2 and N_2O in the
53 overall gaseous N emissions depend – among other factors – on the degree of O_2 restriction (Firestone and
54 Davidson, 1989). Soil physical and biotic factors (i.e. diffusion permitted by soil porosity in conjunction with
55 water-filled pore space [WFPS] as well as consumption of O_2 by heterotrophic respiration and nitrification)
56 control the aerobic status of a soil (Ball, 2013; Uchida et al., 2008; Maag and Vinther, 1999). In general, fine
57 textured soils with higher clay contents exhibit a lower gas diffusivity compared to coarse textured soils, which
58 result regularly in higher denitrification activity in the former with higher N_2O emission rates, but also a higher
59 probability for the consecutive reduction to N_2 (Senbayram et al., 2014; Gu et al., 2013; Ball, 2013).

60 There is a general lack of knowledge about the effects of high BD concentrations on gaseous N-losses as they
61 might appear after injection into soils and their interactions with O_2 limiting factors like soil texture and WFPS,
62 as well as temperature and heterotrophic respiration. Thus, we applied the helium-oxygen (He-O_2) incubation
63 technique (Butterbach-Bahl et al., 2002) in a laboratory experiment to evaluate the effect of above suggested
64 factors on the emission of N_2O and N_2 from different soils. Simultaneously, CO_2 flux was determined as an
65 indicator for microbial O_2 consumption, O_2 diffusion and also for the degradability of organic C applied with BD
66 (Blagodatsky and Smith, 2012), but with the restriction that inorganic sources could not be differentiated. We
67 hypothesised that (1) N_2O and N_2 emissions will increase with WFPS, (2) this gaseous N losses will also be

68 affected by BD application rate, i.e. the hypothetical concentration resulting from injection, and (3) the fine
69 textured clayey silt will induce higher gaseous N losses than the coarse loamy sand.

70 2 Material and Methods

71 2.1 Selected soils, sampling of soil cores and biogas digestate

72 Two soils were selected and both were adjusted to three levels of WFPS and three quantities of BD (Table 1),
73 resulting in 18 factor combinations with three repetitions each. Temperature was increased from 2 °C during the
74 first two days to 15 °C for the last three days of the incubation. Intact soil cores (diameter 7.2 cm, height 6.1 cm,
75 volume 250 cm³) were taken with sample rings in the range from 0–0.10 m depth from two sites with different
76 textures, i.e. sandy loam and clayey silt. The sandy loam samples were collected from a stagnic luvisol (IUSS
77 Working Group WRB, 2006) located in Gülzow (North-East Germany) in the ground moraine of the
78 Weichselian glacial period at 53° 48' 35" N and 12° 4' 20" E. The clayey silt samples were collected from a
79 haplic luvisol located in Dornburg between the foothills and the lowlands of Central Germany at 51° 0' 8" N and
80 11° 39' 25" E (see Table 2 for more details on soil characteristics). After field sampling, the soil cores were dried
81 for 48 h at 40 °C to facilitate adjustment of WFPS.

82 Both sites have been cultivated with similar crop rotations used as feedstock for biogas production and have been
83 amended with biogas digestate for the past nine years. The crop rotation on the sandy loam consisted of maize
84 (*Zea mays* L.), rye (*Secale cereale* L.), sorghum (*Sorghum bicolor* (L.) MOENCH), winter triticale (×
85 *Triticosecale* Wittmack), ryegrass (*Lolium perenne* L.) and winter wheat (*Triticum aestivum* L.). The only
86 difference in the crop rotation on the clayey silt was the cultivation of sudangrass (*Sorghum × drummondii*)
87 instead of sorghum.

88 The biogas digestate used for the incubation was obtained from a biogas plant at 'Gut Dalwitz', an organic farm
89 in northeast Germany. The feedstock for the anaerobic fermentation in the plant consisted of 60 % maize, 20 %
90 solid cattle manure, 10 % dry chicken manure and 10 % rye. The digestate was analysed by 'LUFÄ', Rostock,
91 Germany and had a pH of 8.3, 2.91% organic C, 0.16% dissolved organic C (DOC), 0.54% N and 0.27% NH₄-N
92 in undried material with a dry matter content of 9.4%.

93 2.2 Adjustment of WFPS and addition of N

94 For adjustment of WFPS, the dry and undisturbed soil cores were moistened dropwise. The respective quantities
95 of water were calculated based on the bulk density, an assumed particle density of 2.65 g cm^{-1} and reduced by
96 the expected moisture input from subsequent addition of BD. The soil cores were then mixed with BD and
97 finally repacked to reach nutrient concentrations comparable to that in injection bands. The amounts of added
98 BD were calculated with an assumed injection of 160 kg N ha^{-1} into soil with row spaces of 0.15 m (narrow
99 injection bands with low BD concentration, LOBD) and 0.30 m (wide injection bands with high BD
100 concentration, HIBD), which are common ranges used by injection machinery and which correspond to 17.6 and
101 25.3 mL BD , respectively, per sample ring. After this procedure, the soil cores were sealed with plastic lids and
102 stored immediately at $2 \text{ }^\circ\text{C}$ until the beginning of the incubation within a week.

103 2.3 Determination of gas fluxes

104 The measurements of N_2 , N_2O and CO_2 fluxes were applied following the He- O_2 method (Scholefield et al.,
105 1997; Butterbach-Bahl et al., 2002). Six soil cores (i.e. the repetitions of two factor combinations at a time, Table
106 3) were placed simultaneously in special gas-tight incubation vessels inside a climate chamber. Analyses were
107 conducted in the laboratory of the Institute for Landscape Biogeochemistry, Leibniz Centre for Agricultural
108 Landscape Research (ZALF), Müncheberg, Germany. Before flux measurements, the vessels were evacuated
109 moderately (0.047 bar) and flushed with an artificial He/ O_2 gas mixture ($20.49 \text{ } \%$ O_2 , 345.5 ppm CO_2 , 359 ppb
110 N_2O , 1863 ppb CH_4 , 2.46 ppm N_2 , rest He) four times consecutively to remove ambient N_2 . Subsequently, the air
111 temperature of the climate chamber was set to $2 \text{ }^\circ\text{C}$ and a continuous He/ O_2 gas flow rate of 15 ml min^{-1} was
112 applied to the vessel headspaces for 72 h to remove residues of N_2 from soil cores by diffusion, including a
113 restricted N_2 production by decreased microbial activity. After this pre-incubation, during the following two
114 days, the headspace concentration of N_2O and CO_2 was measured once daily in the morning. To compensate for
115 the lower precision of the detector for N_2 in relation to the detector for N_2O and CO_2 (Eickenscheidt et al.,
116 2014), N_2 concentrations were measured consecutively three times daily in the morning. Immediately after the
117 last measurement on the second day, the temperature was set to $15 \text{ }^\circ\text{C}$ and the measurements were continued for
118 another two days. Finally, the He/ O_2 gas mixture was substituted by pure He and, following 24 h of
119 acclimatisation, gas measurements were carried out once again (Figure 1) to determine the generation of N_2O
120 and N_2 in a completely anaerobic soil matrix. The latter step is important to get a clue about the actual potential

121 for gaseous N losses after highly concentrated BD application. The settings of the chromatographs for gas
122 analyses are described in Eickenscheidt et al. (2014). Gas fluxes were calculated according to Eq. (1):

$$123 \quad f = \frac{M \times p \times v \times dc}{R \times T \times A}, \quad (1)$$

124 where f is the flux (N_2 and CO_2 : $\text{mg m}^{-2} \text{h}^{-1}$, N_2O : $\mu\text{g m}^{-2} \text{h}^{-1}$), M the molar mass in g mol^{-1} (N_2 : 28, CO_2 : 44,
125 N_2O : 44), p the air pressure (Pa), v the air flow (L h^{-1}), R the gas constant ($8.31 \text{ J mol}^{-1} \text{ K}^{-1}$), T the temperature
126 inside the chamber (K), A the area of the incubation vessel (m^2), and dc the difference of gas concentrations (N_2
127 and CO_2 : ppm, N_2O : ppb) between inlet and outlet of a vessel.

128 To enhance the tightness against atmospheric N_2 contamination, the lids of the incubation vessels were purged
129 permanently with helium. We obtained blank values by inserting aluminium blocks into the vessels before each
130 measurement cycle. Since this blank values were usually steady with means of 1.9 ($1\sigma = 0.9$) ppm N_2 , 349.6 (1σ
131 = 11.4) ppb N_2O and 353.9 ($1\sigma = 13.5$) ppm CO_2 , we suggest that the vessels were tight. Derived from the blank
132 values, lowest detectable fluxes were on average 0.427 ($1\sigma = 0.271$) $\text{mg N}_2\text{-N m}^{-2} \text{h}^{-1}$, 3.6 (3.1) $\mu\text{g N}_2\text{O-N m}^{-2} \text{h}^{-1}$
133 and 0.918 (0.693) $\text{mg CO}_2\text{-C m}^{-2} \text{h}^{-1}$. For flux estimation, the blank values were subtracted from the values
134 measured at the respective outlet. Estimated fluxes from the soil cores smaller than the respective blank fluxes of
135 each day were set to zero.

136

137 2.4 Soil analyses after incubation

138 After incubation, the soil cores were stored at 2 °C until they were extracted with 0.1 M KCl solution (soil to
139 extract ratio 1:4, standardised extraction method of the commissioned laboratory at Leibniz Centre for
140 Agricultural Landscape Research e. V.) and analysed for NH_4^+ and nitrate (NO_3^-) by spectrophotometry
141 according to DIN ISO 14256 with a continuous flow analyser 'CFA-SAN', Skalar Analytical B.V., the
142 Netherlands and for DOC by combustion according to DIN ISO 10694 with an analyser 'RC 612', Leco
143 Instruments GmbH, Germany.

144 2.5 Statistical analysis

145 All statistical analyses were done using R version 3.2.3 (R Core Team, 2015) with the data of the measuring
146 days under He- O_2 atmosphere. Data from the vessels with the factor combination of 35% WFPS and LOBD with
147 clayey silt were omitted due to technical reasons during sample preparation. For the final period of pure He

148 headspace, some gas concentration data are missing due to logistical reasons. For the loamy sand, this affects all
149 WFPS levels with LOBD (N_2 and N_2O), the treatment 75% WFPS with 320 kg N h^{-1} (N_2O and CO_2) and for the
150 clayey silt the treatment 35% WFPS without amendment (N_2O and CO_2).

151 To account for repeated measurement of vessels, linear mixed effect models were applied with package
152 'lmerTest' version 2.0-33 (Kuznetsova et al., 2016) for fluxes of each gas type. The three pseudo-replicated
153 fluxes from the N_2 measurements of each vessel were averaged for each day to obtain the same number of
154 observations as for N_2O and CO_2 fluxes. The fixed structure of models included soil type, WFPS, amount of
155 digestate, temperature, NO_3^- and DOC contents after incubation as well as the fluxes of N_2O (in the model for
156 N_2) and CO_2 (in the models for N_2 , N_2O and $N_2/[N_2+N_2O]$ product ratio). Soil NH_4^+ was omitted since it showed
157 high autocorrelation with the amount of BD applied. The individual soil cores in the vessels were set as random
158 effect (nested within the week of incubation and with allowance for a variable slope of the effect each day) with
159 regard to lack of independence of consecutive measurements. The model responses for N_2 , N_2O and CO_2 were
160 log transformed ($\ln[\text{value} + 1]$) since gas fluxes from soils usually show lognormal distributions (Kaiser et al.,
161 1998). The function 'step' was used for automatic backward selection of models based on AIC (Akaike's 'An
162 Information Criterion'). The skewness (γ) was calculated with R package 'moments' version 0.14 (Komsta and
163 Novomestky, 2015) to check residuals for normal distribution and $|\gamma| \leq 2$ was assumed as appropriate (West et
164 al., 1995). For mixed effects models, p -values of the ANOVA (type 2) were calculated based on Satterthwaite's
165 approximation)
166 Cumulated gas fluxes were estimated with a bootstrap method using function 'auc.mc' of R package 'flux'
167 version 0.3-0 (Jurasinski et al., 2014) for the R statistical software version 3.2.3 (R Core Team, 2015). In short,
168 the fluxes for the period of aerobic headspace were cumulated in 100 iterations, while for each run 2 fluxes were
169 omitted randomly. Then, the resulting data were used to calculate means and standard deviations.

170 **3 Results**

171 **3.1 Soil NH_4^+ , NO_3^- and DOC contents**

172 The calculated application of NH_4^+ -N from BD per kg soil approximated for the sandy loam 247.0 mg (LOBD)
173 and 494.0 mg (HIBD), and for the clayey silt 266.0 mg (LOBD) and 532.0 mg (HIBD). The NO_3^- content of BD
174 was negligible. In general, the NH_4^+ content of the soils after incubation increased with digestate application
175 with lower amounts detected in the clayey silt. Nitrate was found almost exclusively in the latter soil. (Fig. 2).

176 The amounts of measured DOC increased with the application rate of BD, but with higher magnitudes for the
177 loamy sand than for the clayey silt (Table 4).

178 3.2 CO₂ fluxes

179 CO₂ fluxes showed clear differences between the soils: under all combinations of temperature and oxygen, the
180 fluxes were always larger from the loamy sand compared with the clayey silt (Table A1). In general, the mean
181 fluxes from the loamy sand increased with the amount of digestate during each of the different periods regarding
182 temperature and headspace aerobicity, but showed no obvious pattern with WFPS. There was no clear trend of
183 fluxes with the amount of amendment, but a slight trend of decreasing fluxes with increasing WFPS for the
184 clayey silt. However, the predictive power of WFPS on CO₂-C fluxes was minor since it was eliminated during
185 stepwise regression fitting. By contrast, soil type, amount of digestate, temperature as well as the DOC content
186 after the incubation had significant ($p < 0.01$) effects (Table 5).

187 3.3 N₂O fluxes

188 The mean N₂O fluxes from the loamy sand in the He-O₂ headspace were virtually zero, independent of
189 temperature and WFPS as well as the amount of BD application (Fig. 3, Table A2). In contrast, the emissions of
190 the clayey silt increased with temperature and were highest at 15 °C with intermediate WFPS and amount of BD,
191 i.e. 6.2 mg N₂O-N m⁻² h⁻¹ at 55% with LOBD, respectively. Surprisingly, at 15 °C, increasing the amount of BD
192 up to HIBD did not increase the observed N₂O efflux; rather it decreased the efflux significantly ($p < 0.05$,
193 Tuckey's HSD) at 55% and also, but not significantly, at 75% WFPS (Fig. 3, Table A2). According to the linear
194 mixed model for N₂O fluxes in aerobic conditions, WFPS, amount of digestate, temperature, DOC content of
195 soil after incubation and CO₂ fluxes had significant ($p < 0.001$) effects on N₂O flux (Table 5).

196 Under anaerobic headspace conditions, the overall highest mean N₂O flux was observed from the clayey silt at
197 35% WFPS with HIBD (11.7 mg N₂O-N m⁻² h⁻¹). The same soil showed a tendency of decreasing N₂O fluxes
198 with increasing WFPS and amendment. In the loamy sand, the pure He-atmosphere induced increasing mean
199 N₂O fluxes (up to 1.3 mg N₂O-N m⁻² h⁻¹) with increasing WFPS (Fig. 3, Table A2). Thus, the anaerobic
200 headspace induced a change only in the loamy sand by increasing emissions.

201 3.4 N₂ fluxes

202 From the loamy sand, no or negligible N₂ fluxes were detected at both temperatures under He-O₂ atmosphere
203 (Fig. 4, Table A3). The clayey silt showed mean fluxes of up to 1.4 mg N₂ m⁻² h⁻¹ at 2 °C (all incubations with
204 75% WFPS) and up to 3.8 mg N₂ m⁻² h⁻¹ at 15 °C (75% WFPS with LOBD), but no fluxes in all BD treatments
205 with 35% WFPS. Put simply, temperature had a small effect on N₂ emissions from the sandy loam with no
206 consistent influence of WFPS and the amount of BD. In contrast, the clayey silt emitted clearly increasing fluxes
207 with increasing temperature and WFPS. However, the application raise from LOBD up to HIBD at 15 °C,
208 resulted in slightly, but not significantly ($p > 0.05$, Tuckey's HSD), decreased fluxes (Fig. 4, Table A3). The
209 summary of the linear mixed model for N₂ fluxes under aerobic conditions revealed significant effects ($p < 0.05$)
210 of soil type, WFPS, the amount of digestate, temperature, DOC content after incubation and N₂O flux (Table 5).
211 After switching the atmosphere to pure He, the N₂ fluxes from the sandy loam increased more than 60-fold. In
212 contrast to aerobic conditions, all measured factor combinations showed mean fluxes up to 35.1 mg N₂ m⁻² h⁻¹
213 (55% with 320 kg N ha⁻¹) (Fig. 2, Day 5 in Table A3). The mean fluxes from the clayey silt increased only up to
214 9.3 mg N₂ m⁻² h⁻¹ in amended treatments. Thus, the loamy sand exhibited a much more intense reaction under
215 anaerobic headspace conditions.

216 3.5 N₂/(N₂ + N₂O-N) product ratio

217 No clear trend of the product ratio of N₂/(N₂ + N₂O-N) was found for incubations of the loamy sand. However,
218 there was a clear distinction of the ratios for this soil under aerobic and anaerobic atmospheres: while the ratios
219 were close to zero in the former, they were close to 1 in the latter (Fig. 5). In contrast, in the clayey silt the ratios
220 increased with WFPS and were affected by digestate amendment under both the aerobic and the anaerobic
221 atmospheres, where the highest ratios (up to 0.8) were found in treatments without digestate and at least 55%
222 WFPS. The digestate-amended treatments showed mostly ratios around or above 0.5, with exception of the 35%
223 WFPS treatments, which had ratios close to zero. According to the linear mixed model, the product ratio under
224 aerobic conditions was affected significantly ($p < 0.01$) by soil type and the amount of digestate (Table 5).

225 **4 Discussion**

226 **4.1 Increased BD application rate did not increase N₂O and N₂ losses probably due to inhibitory effect**
227 **of high NH₄⁺ concentrations**

228 The overall N₂O fluxes corresponded well with those from other studies with similar incubation conditions and
229 application rates of BD in terms of N ha⁻¹ (Severin et al., 2015; Senbayram et al., 2012; Köster et al., 2015).
230 However, the latter studies assumed a distribution of BD into soil by a cultivator, which implies a smaller
231 concentration of BD compared to its occurrence in injection slits. Although we observed differences in N₂O
232 emissions between soils, soil type was not confirmed as a significant effect. Nevertheless, WFPS and
233 temperature, which are well known controllers of N₂O generation (Maag and Vinther, 1999), showed significant
234 influences. Both are physical (by gas diffusion) and biological (by increased metabolic activity and consequently
235 increased O₂ consumption by respiration) drivers for O₂ availability, respectively (Maag and Vinther, 1999; Ball,
236 2013). Accordingly, the CO₂ flux (resulting from respiration of O₂) generally increased with temperature and
237 was also identified as significant by regression selection.

238 The mean N₂ fluxes of up to 0.5 (loamy sand) and 3.8 mg N m⁻² h⁻¹ (clayey silt) at 15° C (Fig. 5, Table A3) were
239 considerably smaller than the mean fluxes of up to 13.0 mg m⁻² h⁻¹ observed by Köster et al. (2015) during the
240 first five days of their incubation. Although the amount of BD in terms of applied N (250 kg ha⁻¹) was
241 comparable, Köster et al. (2015) used a higher WFPS of 90%, which may have increased the generation of N₂. In
242 contrast to N₂O emission rates, the observed N₂ fluxes depended not only on WFPS, but also on soil type (Table
243 5), most likely due to the direct influence of soil structure on diffusivity and, thus, the supply with O₂ (Balaine et
244 al. 2016; Butterbach-Bahl et al. 2013). N₂O flux showed also a significant effect during regression selection for
245 N₂. N₂O is the direct precursor of N₂ in denitrification and, hence, the flux of the latter depends on the
246 availability of the former. However, temperature showed no significant effect.

247 The N₂/(N₂+N₂O) ratios were significantly determined only by soil type and WFPS: while no clear trend was
248 observable for the loamy sand, there was a pronounced effect in the clayey silt (Fig 4). We attribute the lack of a
249 trend in the loamy sand to generally adverse conditions for the formation of N₂O and N₂. Contrary, the influence
250 of WFPS apparently mirrored favourable conditions in the clayey silt (Table 5). Simultaneously, with increasing
251 WFPS, the reduction of N₂O accelerates as an alternative electron acceptor under reduced O₂ supply (Tiedje,
252 1988). Accordingly, no or rather small fluxes of the investigated gaseous N species were generally found in our
253 presumably well aerated treatments with 35% WFPS.

254 In our study, one treatment (clayey silt, 55% WFPS, LOBD) showed exceptionally large mean N₂O fluxes of up
255 to 7.1 mg N m⁻² h⁻¹ (Fig. 3, Table A2). This could be evidence that injection of such commonly applied amounts
256 of BD-N (i.e., 160 kg N ha⁻¹) may favour much larger losses of N₂O compared to an even distribution of BD in
257 soils due to larger substrate concentration in injection slits. However, with higher amendments (i.e. HIBD), we
258 observed surprisingly partially significant ($p < 0.05$, Tuckey's HSD) reductions of N₂O and a decreasing
259 tendency of N₂ emissions (Table A2, Table A3). In line with this, the amount of BD showed a significant effect
260 during the regression selection on N₂O, but not on N₂ fluxes (Table 5). A coherent reason for the rather smaller
261 emissions of highly amended HIBD treatments might be the inhibitory effect of NH₃ on nitrification.
262 Accordingly, Anthonisen et al. (1976) found an inhibition by concentrations from 0.1 to 150 mg NH₃ L⁻¹. The
263 application rate in the treatments with HIBD amounted to approximately 500 mg NH₄⁺-N (kg soil)⁻¹ (Fig. 3)
264 which correspond to 25.8 mg NH₃-N (kg soil)⁻¹ at 15 °C if we use the pH of the BD and assume that all
265 extractable NH₄⁺-N was in solution (Emerson et al., 1975). Hence, we consider this inhibitory effect as the
266 reason for the missing increase of N₂O and N₂. Additionally, due to the increased pH of BD (Möller and Müller,
267 2012), the amount of NH₄⁺ fixed as NH₃ by soil organic matter increases and, moreover, this fixed NH₃ is not
268 readily extractable by the KCl method we have applied (Kissel et al., 2008). This is consistent with the
269 observation of generally higher N₂O and N₂ fluxes from the clayey silt since clay increases the sorption capacity
270 of soils for NH₄⁺ and may, thus, reduce the inhibitory effect on nitrification (Kissel et al., 2008). However,
271 because we mixed the BD with the soil, we would expect a lower NH₃ fixation in tubular injection slits *in situ*,
272 resulting in probably lower N₂O and N₂ fluxes from clayey soils.

273 Actually, high NH₄⁺ loads in conjunction with an increased pH favour NO₂⁻ accumulation, because NO₂⁻
274 oxidising bacteria are less resilient against high concentrations of NH₃ than NH₃ oxidising bacteria (Anthonisen
275 et al., 1976). This NO₂⁻ should have protonated then partly to the toxic and unstable HNO₂, which drives
276 biological and chemical production of NO and N₂O for detoxification (Venterea et al., 2015). Although we have
277 not determined NO₂⁻, we suggest a dominant role of nitrifier denitrification, i.e., NO₂⁻ reduction, in the
278 generation of N₂O during our experiment, especially during the anaerobic headspace conditions at the end of the
279 incubation, resulting in the relatively small NO₃⁻ recovery in both soils. Accordingly, coupled nitrification-
280 denitrification and bacterial denitrification have been found to dominate the production of N₂O directly after
281 application of BD (Köster et al., 2011; Senbayram et al., 2009). However, N₂O-N losses were clearly larger than
282 N₂ losses under aerobic headspace in the clayey silt, indicating that much of the N gas loss was driven by

283 processes other than canonical denitrification. Under the above mentioned conditions, NO-N losses may exceed
284 N₂O losses (Venterea et al., 2015), calling for taking account of NO measurements in future studies.

285 4.2 Different effects of soil diffusivity on N₂O and N₂ fluxes

286 Apparently, the tested factors affected the N₂O and N₂ fluxes from both soils in a different way. A specific soil
287 characteristic that exhibits such a fundamental control on biogeochemical processes such as denitrification is the
288 diffusivity for O₂ (Ball, 2013, 2013; Letey et al., 1980; Parkin and Tiedje, 1984), which is a main soil
289 characteristic responsible for the appearance of anaerobic microsites. In general, diffusivity integrates the soil
290 porosity, i.e., pore continuity and size as well as WFPS, which control both soil N₂O and N₂ emissions (Balaine
291 et al., 2016; Letey et al., 1980; Ball, 2013). Soils with a coarser texture like the loamy sand have a higher
292 proportion of macro-pores and thus a higher gas diffusion compared with fine textured soils like the clayey silt
293 we used (Groffman and Tiedje, 1991). This lets us expect conditions that are more favourable for N₂O and N₂
294 generation in the latter due to relatively poor diffusion characteristics and, thus, a smaller O₂ supply. Actually,
295 although we incubated the soils at comparable levels of WFPS and BD amendments, the apparent lower
296 diffusivity led to larger N₂O and N₂ production in the treatments with the clayey silt in relation to the loamy
297 sand.

298 The role of the distinct diffusivities of both soils is corroborated by our observations of the gas fluxes in
299 anaerobic headspace. With switching the He-O₂ atmosphere in the headspace to pure He, the denitrification
300 potential can be tested because anaerobicity eliminates respiration processes that use O₂ as electron acceptor
301 (Parkin and Tiedje, 1984). We acknowledge e.g. DNRA and anammox as possible additional sources of N₂O and
302 N₂, respectively, under such conditions, but we were not able to quantify their contribution. The anaerobic
303 headspace induced a considerable increase of N₂O fluxes in the loamy sand, but not in the clayey silt.
304 Concurrently, the N₂ fluxes increased in both soils, but pronounced, i.e. more than 60-fold, in the sandy loam.
305 These observed changes resulting from oxygen deprivation imply that, during the previous aerobic conditions,
306 the diffusivity of the sandy loam was too high to allow for a sufficient establishment of anaerobic microsites,
307 while the clayey silt ensured a moderate diffusional constraint to maintain suboxic conditions. In general, only
308 N₂O fluxes from treatments with negligible fluxes during the previous aerobic period increased under anaerobic
309 conditions, including all treatments with loamy sand (Fig. 3, Table A2). At the same time, there was a reduction
310 of N₂O fluxes in most clayey silt treatments. However, when we take a closer look at the simultaneous changes
311 of N₂ fluxes after atmosphere change, virtually all of the respective treatments showed increased rates. Hence,

312 there was an enhanced reduction of N_2O to N_2 , which is reflected in the increased $N_2/(N_2 + N_2O)$ ratio (Fig. 5)
313 and points to intensified reduction of N_2O due to the lack of oxygen (Parkin and Tiedje, 1984). The much larger
314 N_2 fluxes from the loamy sand compared to the clayey silt might have been caused additionally by small NO_3^-
315 availability (Fig. 2) and a high availability of C (Table 4), which promoted the reduction of N_2O to N_2
316 (Senbayram et al., 2012). Further, we found no evidence for any shortage of substrate in the clayey silt during
317 the subsequent anaerobic headspace conditions. However, the cumulated fluxes of both N_2 and N_2O amounted to
318 a maximum absolute loss of 9.4 ($1\sigma = 0.3$) mg N per kg soil in the clayey silt with LOBD and 55% WFPS, which
319 was roughly 3.5% of the calculated NH_4^+ -N applied with BD (Fig. 2). On the other hand, the $N_2/(N_2+N_2O)$ ratios
320 increased only slightly (Fig. 5) and, in contrast to the loamy sand, there were still significant N_2O fluxes in the
321 clayey silt (Fig. 3), which point to still sufficient stocks of NO_3^- in the latter (Senbayram et al., 2012). In fact, the
322 NO_3^- stock was greater in the clayey silt than in loamy sand after incubation (Fig. 2). Thus, we suggest that the
323 gas fluxes were unaffected by the change to anaerobic headspace in the clayey silt due to already low O_2
324 concentrations as a result of poor diffusivity. In conclusion, distinct gas diffusivities of both soils can be
325 proposed as the main reason for the differing N_2O and N_2 fluxes.

326 In interaction with soil diffusivity, also respiration affects the aerobicity of a soil matrix by concurrent
327 consumption and formation of O_2 and CO_2 , respectively. Depending on microbial availability of carbon,
328 respiration could be indicated approximately by DOC, though not all DOC might be readily degradable (Cook
329 and Allan, 1992). Generally, the DOC contents after our incubation increased with application rate of BD (Table
330 4), but the DOC contents were always smaller in the clayey silt. This might reflect a stronger sorption of C and
331 thus a lower availability for respiration in the clayey silt compared to the loamy sand (Kaiser and Guggenberger,
332 2000). If we compare the DOC values with the cumulated flux rates of CO_2 over the period of aerobic
333 headspace, we find a good regression fit ($R^2 = 0.91$, $p < 0.001$) for both soils (Fig. 6) indicating a sufficient
334 availability of C from BD for respiration and, thus, implicitly also for denitrification (Reddy et al., 1982).
335 Moreover, as increased DOC enhanced respiration (Table A1), it consequently affected O_2 consumption and,
336 thus, also the emergence of anaerobic microsites (Azam et al., 2002). Accordingly, there is also a good
337 correlation between cumulated CO_2 and $N_2O + N_2$ fluxes for the same period from the clayey silt ($R^2 = 0.93$, $p =$
338 0.001), when the treatments with 35 % WFPS (which showed virtually no N emissions) are omitted (Fig. 7).
339 However, there was no such a correlation for the loamy sand. This confirms the interactive effect of diffusivity
340 (induced by both the soils and WFPS) and C availability on the emissions of N_2O and N_2 , which, nevertheless,
341 interacted with the inhibitory effect of high NH_4^+ loads on nitrification (see chapter 4.1).

342 5 Relevance and implications

343 Our aim was to estimate the effect of differing soil environmental conditions on gaseous N losses – and not to
344 draw conclusions about the long-term dynamics of N_2 and N_2O emissions after BD application in concentrations
345 similar to injection. In another laboratory study at a WFPS of 65%, Senbayram et al. (2009) measured only one
346 peak within two days without a repeated increase later, regardless the amount of applied BD. Thus, we assume a
347 single peak shortly after application holds also true for our incubation as well. We assume also the measurements
348 after only 24 hours of anaerobicity in the headspace as representative for the emission potential since Wang et al.
349 (2011; 2013) showed in similar studies to ours that the emission of N_2 and N_2O peaked within less than 24 hours
350 after switching their headspace from aerobic to anaerobic conditions.

351 However, as hypothesised, N_2O and N_2 emissions as well as the $N_2/(N_2O+N_2)$ ratio increased with WFPS, most
352 probably due to restricted supply of O_2 . Contrary to our second hypothesis, the gaseous losses of N_2O and N_2 did
353 not increase with the application rate of BD. This indicates an inhibitory effect of high NH_3 and NH_4^+
354 concentrations, respectively, on nitrification, which are found typically in biogas digestates (BD). Nevertheless,
355 the $N_2/(N_2O+N_2)$ ratio tended to decrease with application rate as supposed, probably due to a copious supply
356 with NO_2^- and NO_3^- from oxidised $BD-NH_4^+$. Confirming our third hypothesis, the fine textured clayey silt
357 induced **larger** gaseous N losses and a higher $N_2/(N_2O+N_2)$ ratio than the coarse loamy sand by the apparent
358 distinct diffusivities of both soils. Overall, there was a larger potential for formation of N_2O in the fine-textured
359 clayey silt compared to the coarse loamy sand **after the application of high concentrations of BD as they may**
360 **appear after** injection. However, the loamy sand showed a large potential for N_2 formation under anaerobic
361 headspace conditions. Nevertheless, further investigations are needed in regarding **the dynamics and** the duration
362 of the observed effects and their reliability for field conditions.

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514

515 **Table 1: The examined factors soil texture, water-filled pore space (WFPS), and amount (i.e., concentration) of**
516 **nitrogen (N) applied with biogas digestate (BD) with their respective levels applied in the present study, resulting in**
517 **18 treatments with three replicates each. The temperature was manipulated consecutively during the incubation.**

Factor [<i>n</i>]	Levels		
Soil texture [2]	loamy sand	clayey silt	
WFPS (%) [3]	35	55	75
BD-N (kg ha ⁻¹) [3]	0	160	320
Temperature (°C) [2]	2	15	

518

519 **Table 2: Characteristics of both soils. Texture and mean values with standard deviations (in brackets) for carbon (C, $n = 9$), nitrogen (N, $n = 9$), pH ($n = 3$), bulk density (BD, $n =$**
 520 **3) and mineral N (NO_3^- and NH_4^+ , $n = 3$) of both soils in 0–10 cm depth after field sampling.**

Texture	C (mg g^{-1}) ^a	N (mg g^{-1}) ^a	pH ^b	Bulk density (g cm^{-3}) ^c	NO_3^- (mg kg^{-1}) ^d	NH_4^+ (mg kg^{-1}) ^d
Loamy sand	6.99 (0.29)	0.67 (0.05)	7.2 (0.1)	1.4 (0.0)	1.0 (0.2)	0.6 (0.3)
Clayey silt	10.77 (0.28)	1.19 (0.06)	7.2 (0.0)	1.5 (0.0)	1.8 (0.2)	0.3 (0.2)

521 ^a measured with analyser “Truspec CNS”, Leco Instruments GmbH, Germany, performed according to ISO 10694 (“elemental analysis”) for C and according to ISO

522 13878 (“elemental analysis”) for N

523 ^b measured in H_2O with TitraMaster85, Radiometer Analytical SAS, France, performed according to VDLUFA Methodenbuch, Vol. 1, chap. 5.1.1

524 ^c measured on 250 cm^3 soil cores

525 ^d measured with analyser “CFA-SAN”, Skalar Analytical B.V., the Netherlands, performed according to ISO 14256

526 **Table 3: Chronological order of the incubated factor combinations. Two different factor combinations with their**
 527 **respective repetitions ($n = 3$) were placed together for each weekly incubation course (cf. Fig. 1). The factors were**
 528 **combined by (1) soil (loamy sand: LS, clayey silt: CS), (2) amount (kg) of applied N from digestate per ha and (3)**
 529 **WFPS (%).**

Week	Factor combination 1	Factor combination 2
1	LS - 0 N - 35%	LS - 0 N - 55%
2	LS - 0 N - 75%	LS - 160 N - 35%
3	LS - 160 N - 55%	LS - 160 N - 75%
4	LS - 320 N - 35%	LS - 320 N - 55%
5	LS - 320 N - 75%	CS - 0 N - 35%
6	CS - 0 N - 55%	CS - 0 N - 75%
7	CS - 160 N - 35%	CS - 160 N - 55%
8	CS - 160 N - 75%	CS - 320 N - 35%
9	CS - 320 N - 55%	CS - 320 N - 75%

530

531 **Table 4: Mean DOC values from soils, measured after incubation with standard deviations in brackets for the**
 532 **respective treatments differing in amount of applied biogas digestate (BD) and water-filled pore space (WFPS).**

kg digestate-N ha ⁻¹	WFPS (%)	mg DOC (kg soil) ⁻¹	
		Loamy sand	Clayey silt
0	35	41.4 (2.7)	18.9 (1.1)
	55	38.6 (3.1)	19.8 (1.4)
	75	43.7 (1.4)	19.0 (1.8)
160	35	197.4 (20.7)	n.a.
	55	190.5 (19.3)	68.3 (12.7)
	75	362.2 (40.0)	63.2 (9.6)
320	35	316.8 (25.3)	358.1 (26.3)
	55	312.5 (14.3)	94.8 (13.6)
	75	500.1 (33.4)	105.9 (14.8)

533 n.a.: data not available

534 **Table 5: ANOVA table (type 2, p -values calculated based on Satterthwaite's approximation) of the linear mixed effects models for estimated fluxes of N_2 , N_2O , $N_2/(N_2+N_2O)$**
 535 **product ratio and CO_2 in aerobic He- O_2 atmosphere. Soil type, water-filled pore space (WFPS), amount of digestate, temperature, NO_3^- and DOC content of soil after incubation**
 536 **as well as fluxes of N_2O and CO_2 were set as possible independent variables. The individual soil rings were set as random effect (nested within the respective week and with the**
 537 **allowance for varying slopes for each day of measurements). The random effect was always significant.**

538

Response	Fixed effects							
	Soil type	WFPS	Digestate amount	Temperature	NO_3^- post	DOC post	N_2O flux	CO_2 flux
N_2	0.026	< 0.001	0.008	0.037	†	0.001	< 0.001	†
N_2O	†	< 0.001	< 0.001	< 0.001	†	< 0.001	*	< 0.001
$N_2/(N_2+N_2O)$	0.005	0.004	†	†	†	†	*	†
CO_2	<0.001	†	<0.001	<0.001	†	0.007	†	*

539 † Variable eliminated during stepwise regression selection

540 * Variable was not included into original regression

541 **Figure 1: Course of incubation and gas measurements with respect to atmosphere and temperature of the headspace**
542 **after two days of pre-incubation at 2 °C in He/O₂ gas mixture. Gas concentrations of the headspace were determined**
543 **on five consecutive days, i.e. Monday to Friday in the morning. After the first two measurement days, the headspace**
544 **temperature was increased from 2 to 15 °C. Additionally, after the fourth measurement day, the aerobic**
545 **Helium/oxygen gas mixture in the headspace was replaced by a pure Helium atmosphere.**

546

547 **Fig. 2: Ammonium and nitrate contents from loamy sand and clayey silt after incubation with different water-filled**
548 **pore spaces (WFPS, %) and amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and 35.2 mL:**
549 **'High BD'). Error bars denote standard deviations. In general, the ammonium content increased with digestate**
550 **application with lower amounts detected in the clayey silt. Nitrate was found almost exclusively in the latter soil. For**
551 **comparison, calculated amounts of ammonium applied with biogas digestate are shown by triangles. One treatment**
552 **(*) was omitted from all analyses due to technical reasons.**

553

554 Fig. 3: Mean N₂O fluxes (mg N m⁻² h⁻¹) from a loamy sand and a clayey silt incubated under different water-filled pore
555 spaces (WFPS, %) with different amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and 35.2
556 mL: 'High BD'). The first till the fourth day of the incubation were measured in an aerobic He-O₂ headspace (with
557 two days at 2 °C followed by another two days at 15 °C) while on the fifth day measurements were conducted in an
558 anaerobic headspace with pure He (at 15 °C). Error bars show standard deviations; if bars are not visible, they are
559 smaller than the symbols of the means. Under aerobic atmosphere, N₂O fluxes from loamy sand were negligible, while
560 fluxes from clayey silt showed an increase with temperature, especially with higher WFPS and intermediate amounts
561 of digestate. Under anaerobic atmosphere, mean fluxes from loamy sand increased slightly, but significantly (Tukey's
562 HSD, *p* < 0.05). The fluxes from clayey silt showed no significant differences (Tukey's HSD, *p* < 0.05) compared to the
563 day before, with the exception of 35% WFPS, where mean flux increased strongly in the treatment with 320 kg
564 digestate-N ha⁻¹.

565 Fig. 4: Mean N₂ fluxes (mg m⁻² h⁻¹) from a loamy sand and a clayey silt incubated under different water-filled pore
566 spaces (WFPS, %) with different amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and 35.2
567 mL: 'High BD'). The first till the fourth day of the incubation were measured in an aerobic He-O₂ headspace (with
568 two days at 2 °C followed by another two days at 15 °C) while on the fifth day measurements were conducted in an
569 anaerobic headspace with pure He (at 15 °C). Error bars show standard deviations; if bars are not visible, they are
570 smaller than the symbols of the means. The dotted horizontal lines depict the average blank value; single flux rates
571 lower than the respective blank value were set zero. Under aerobic atmosphere, N₂ fluxes from loamy sand were zero
572 or rather negligible, while fluxes from clayey silt show a distinct increase with WFPS and higher fluxes at 15 °C.
573 Under anaerobic atmosphere, mean fluxes from loamy sand increased by orders of magnitude, while the fluxes from
574 clayey silt increased as well, but more gently compared to the sand.

575 **Fig. 5: Mean $N_2/(N_2 + N_2O-N)$ product ratio from a loamy sand and a clayey silt incubated under different water-filled**
576 **pore spaces (WFPS, %) with different amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and**
577 **35.2 mL: 'High BD'). The first till the fourth day of the incubation were measured in an aerobic He-O₂ headspace**
578 **(with two days at 2 °C followed by another two days at 15 °C) while on the fifth day measurements were conducted**
579 **in an anaerobic headspace with pure He (at 15 °C). Error bars show standard deviations; if bars are not visible, they**
580 **are smaller than the symbols of the means. For the loamy sand, there was a clear distinction of the ratios between**
581 **aerobic and anaerobic atmospheres: while the ratios tended to 0 in the former, they tended to 1 in the latter,**
582 **irrespectively of temperature or amount of digestate. For the clayey silt, ratios increased with WFPS and were highest**
583 **from the unamended treatments under both the aerobic and the anaerobic atmospheres.**

584 **Fig. 6: Regression between DOC (mg per 100 g soil) measured after the incubation and the respective cumulated CO₂**
585 **emissions (g C m⁻²) during the period of aerobic headspace with their standard deviations and confidence interval**
586 **(95%). If error bars are not visible, they are smaller than the symbols of the means. Both soils showed increasing**
587 **emissions with increasing soil DOC contents as well a good regression fit ($R^2 = 0.91, p < 0.001$).**

588

589 **Fig. 7: Regression between cumulated CO₂ emissions (g C m⁻²) and the respective cumulated N₂O + N₂ emissions (g N**
590 **m⁻²) from the clayey silt with WFPS > 35 % during the period of aerobic headspace with their standard deviations**
591 **and confidence interval (95%). If error bars are not visible, they are smaller than the symbols of the means. The**
592 **proportional increase of CO₂ and the N gas species shows a good regression fit of $R^2 = 0.93$, ($p = 0.001$).**

593 **Table A1: Mean CO₂-C fluxes with standard deviations in mg m⁻² h⁻¹ from the loamy sand and the clayey silt, treated**
594 **with different water-filled pore spaces (WFPS, %), amounts of digestate (kg N ha⁻¹) as well as different temperature**
595 **regimes (°C) under aerobic (He-O₂) and anaerobic (He) atmosphere. Column 'Day' denotes the consecutive measuring**
596 **days of the respective incubation cycle. Different letters after fluxes indicate significant differences (Tukey's HSD, *p* <**
597 **0.05) within each soil and measuring day. Zeros as last digits were omitted.**

Day	Atmosphere	Temperature (°C)	WFPS (%)	kg N ha ⁻¹	mg CO ₂ -C m ⁻² h ⁻¹	
					Loamy sand	Clayey silt
1	He-O ₂	2	35	0	6.8 ± 2.4 cd	0 ± 0 c
1	He-O ₂	2	35	160	22 ± 3.5 bcd	NA
1	He-O ₂	2	35	320	23.3 ± 9.3 bc	22.8 ± 2.8 ab
1	He-O ₂	2	55	0	6 ± 0.7 d	4.6 ± 7.9 bc
1	He-O ₂	2	55	160	34.4 ± 3.1 b	34.5 ± 11.6 a
1	He-O ₂	2	55	320	28 ± 3.2 b	15.9 ± 3.4 abc
1	He-O ₂	2	75	0	9.4 ± 1.4 cd	0 ± 0 c
1	He-O ₂	2	75	160	37.5 ± 6 b	15.5 ± 12.1 abc
1	He-O ₂	2	75	320	68.3 ± 12.1 a	24.5 ± 2.7 a
2	He-O ₂	2	35	0	9.8 ± 3.5 c	1.3 ± 1.4 b
2	He-O ₂	2	35	160	23 ± 3.9 bc	NA
2	He-O ₂	2	35	320	30.9 ± 2.2 b	22.2 ± 2.4 a
2	He-O ₂	2	55	0	8.7 ± 1.5 c	0.6 ± 1 b
2	He-O ₂	2	55	160	33.4 ± 0.9 b	27.6 ± 12.3 a
2	He-O ₂	2	55	320	35.9 ± 2.7 b	14.4 ± 1.9 ab
2	He-O ₂	2	75	0	8.3 ± 1.5 c	0 ± 0 b
2	He-O ₂	2	75	160	31.9 ± 3 b	13 ± 9.3 ab
2	He-O ₂	2	75	320	57.6 ± 14.8 a	18.3 ± 4 a
3	He-O ₂	15	35	0	42.5 ± 4.5 c	6.7 ± 0.7 b
3	He-O ₂	15	35	160	114.3 ± 12.2 b	NA
3	He-O ₂	15	35	320	149.5 ± 9.4 b	130.9 ± 105 a
3	He-O ₂	15	55	0	41.3 ± 3.5 c	3.2 ± 0.4 b
3	He-O ₂	15	55	160	108.7 ± 10.1 b	57.8 ± 12.2 bc
3	He-O ₂	15	55	320	162.1 ± 9.6 b	26.8 ± 0.7 bc
3	He-O ₂	15	75	0	44.1 ± 9.8 c	3.2 ± 0.7 b
3	He-O ₂	15	75	160	150.4 ± 19 b	26.4 ± 11.8 bc
3	He-O ₂	15	75	320	249.7 ± 53.5 a	35.3 ± 6 bc
4	He-O ₂	15	35	0	48.7 ± 6 c	15.1 ± 4.9 cd
4	He-O ₂	15	35	160	114.3 ± 6.4 b	NA
4	He-O ₂	15	35	320	156.9 ± 15.4 a	65.7 ± 2.2 a
4	He-O ₂	15	55	0	48 ± 3.4 c	4.2 ± 0.2 d
4	He-O ₂	15	55	160	109 ± 14.4 b	51.2 ± 15.1 ab
4	He-O ₂	15	55	320	177.7 ± 7.5 a	26.6 ± 2.3 cd

4	He-O ₂	15	75	0	34 ± 7.8 c	6.7 ± 4 d
4	He-O ₂	15	75	160	168.7 ± 0.4 a	22.1 ± 14.8 cd
4	He-O ₂	15	75	320	166.3 ± 23.1 a	34.1 ± 5.7 bc
5	He	15	35	0	11.2 ± 0.6 d	NA
5	He	15	35	160	54.8 ± 9.3 c	NA
5	He	15	35	320	149.3 ± 3.9 a	45.8 ± 2.1 a
5	He	15	55	0	13.6 ± 1.9 d	3.4 ± 0.6 c
5	He	15	55	160	55.2 ± 4.4 bc	32 ± 11.4 ab
5	He	15	55	320	164.5 ± 3.5 a	15.2 ± 10.7 bc
5	He	15	75	0	20.9 ± 2.3 d	3.6 ± 0.1 c
5	He	15	75	160	75 ± 7.3 b	20.6 ± 8.5 bc
5	He	15	75	320	NA	26.1 ± 2.6 ab

598

599 **Table A2: Mean N₂O-N fluxes with standard deviations in mg m⁻² h⁻¹ from the loamy sand and the clayey silt, treated**
600 **with different water-filled pore spaces (WFPS, %), amounts of digestate (kg N ha⁻¹) as well as different temperature**
601 **regimes (°C) under aerobic (He-O₂) and anaerobic (He) atmosphere. Column 'Day' denotes the consecutive measuring**
602 **days of the respective incubation cycle. Different letters after fluxes indicate significant differences (Tukey's HSD, *p* <**
603 **0.05) within each soil and measuring day. Zeros as last digits were omitted.**

Day	Atmosphere	Temperature (°C)	WFPS (%)	kg N ha ⁻¹	mg N ₂ O-N m ⁻² h ⁻¹	
					Loamy sand	Clayey silt
1	He-O ₂	2	35	0	0 ± 0	0 ± 0 c
1	He-O ₂	2	35	160	0 ± 0	NA
1	He-O ₂	2	35	320	0 ± 0	0 ± 0 c
1	He-O ₂	2	55	0	0 ± 0	0.3 ± 0.1 c
1	He-O ₂	2	55	160	0 ± 0	1.7 ± 0.4 a
1	He-O ₂	2	55	320	0 ± 0	1.1 ± 0.1 b
1	He-O ₂	2	75	0	0 ± 0	0.4 ± 0.1 c
1	He-O ₂	2	75	160	0 ± 0	1 ± 0.1 b
1	He-O ₂	2	75	320	0 ± 0	1 ± 0.2 b
2	He-O ₂	2	35	0	0 ± 0	0 ± 0 d
2	He-O ₂	2	35	160	0 ± 0	NA
2	He-O ₂	2	35	320	0 ± 0	0 ± 0 cd
2	He-O ₂	2	55	0	0 ± 0	0.3 ± 0.1 bc
2	He-O ₂	2	55	160	0 ± 0	1.5 ± 0.6 a
2	He-O ₂	2	55	320	0 ± 0	1.2 ± 0.2 a
2	He-O ₂	2	75	0	0 ± 0	0.4 ± 0.1 bc
2	He-O ₂	2	75	160	0 ± 0	1 ± 0.1 ab
2	He-O ₂	2	75	320	0 ± 0	1.1 ± 0.2 a
3	He-O ₂	15	35	0	0 ± 0 cd	0 ± 0 c
3	He-O ₂	15	35	160	0 ± 0 abc	NA
3	He-O ₂	15	35	320	0 ± 0 ab	0 ± 0 c
3	He-O ₂	15	55	0	0 ± 0 bcd	0.8 ± 0.2 c
3	He-O ₂	15	55	160	0 ± 0 bcd	7.1 ± 0.9 a
3	He-O ₂	15	55	320	0 ± 0 a	3.5 ± 0.7 b
3	He-O ₂	15	75	0	0 ± 0 ab	0.8 ± 0.2 c
3	He-O ₂	15	75	160	0 ± 0 d	3.2 ± 0.7 b
3	He-O ₂	15	75	320	0 ± 0 cd	3 ± 0.9 b
4	He-O ₂	15	35	0	0 ± 0 b	0 ± 0 c
4	He-O ₂	15	35	160	0 ± 0 ab	NA
4	He-O ₂	15	35	320	0 ± 0 ab	0.1 ± 0.1 c
4	He-O ₂	15	55	0	0 ± 0 b	1 ± 0.2 bc
4	He-O ₂	15	55	160	0.1 ± 0.1 a	6.2 ± 1.1 a
4	He-O ₂	15	55	320	0 ± 0 ab	3 ± 0.8 b

4	He-O ₂	15	75	0	0 ± 0 ab	1.1 ± 0.3 bc
4	He-O ₂	15	75	160	0 ± 0 b	2.6 ± 1 b
4	He-O ₂	15	75	320	0 ± 0 b	2.2 ± 0.9 b
5	He	15	35	0	0.1 ± 0	NA
5	He	15	35	160	NA	NA
5	He	15	35	320	0.9 ± 0.1	11.7 ± 2 a
5	He	15	55	0	0.1 ± 0	0.1 ± 0 c
5	He	15	55	160	NA	5 ± 1 b
5	He	15	55	320	1.2 ± 0.7	1.4 ± 0.8 c
5	He	15	75	0	1.3 ± 1.6	0.1 ± 0 c
5	He	15	75	160	NA	1.7 ± 0.3 c
5	He	15	75	320	NA	1 ± 0.3 c

604

605 **Table A3: Mean N₂ fluxes with standard deviations in mg m⁻² h⁻¹ from the loamy sand and the clayey silt, treated with**
606 **different water-filled pore spaces (WFPS, %), amounts of digestate (kg N ha⁻¹) as well as different temperature**
607 **regimes (°C) under aerobic (He-O₂) and anaerobic (He) atmosphere. Column 'Day' denotes the consecutive measuring**
608 **days of the respective incubation cycle. Different letters after fluxes indicate significant differences (Tukey's HSD, *p* <**
609 **0.05) within each soil and measuring day. Zeros as last digits were omitted.**

	Atmosphere	Temperature (°C)	WFPS (%)	kg N ha ⁻¹	mg N ₂ m ⁻² h ⁻¹	
					Loamy sand	Clayey silt
1	He-O ₂	2	35	0	0 ± 0	0 ± 0 bc
1	He-O ₂	2	35	160	0 ± 0	NA
1	He-O ₂	2	35	320	0 ± 0	0.1 ± 0.1 bc
1	He-O ₂	2	55	0	0 ± 0	1.5 ± 0.3 a
1	He-O ₂	2	55	160	0 ± 0	1.5 ± 0.3 a
1	He-O ₂	2	55	320	0 ± 0	1.5 ± 0 a
1	He-O ₂	2	75	0	0 ± 0	1.2 ± 1.1 a
1	He-O ₂	2	75	160	0 ± 0	1.5 ± 0.2 a
1	He-O ₂	2	75	320	0 ± 0	1.7 ± 0.4 a
2	He-O ₂	2	35	0	0 ± 0	0 ± 0 c
2	He-O ₂	2	35	160	0 ± 0	NA
2	He-O ₂	2	35	320	0 ± 0	0 ± 0 c
2	He-O ₂	2	55	0	0 ± 0	1.3 ± 0.1 a
2	He-O ₂	2	55	160	0 ± 0	0.8 ± 0.5 b
2	He-O ₂	2	55	320	0 ± 0	1.3 ± 0.3 a
2	He-O ₂	2	75	0	0 ± 0	1.4 ± 0.4 a
2	He-O ₂	2	75	160	0 ± 0	1.4 ± 0.5 a
2	He-O ₂	2	75	320	0 ± 0	1.4 ± 0.1 a
3	He-O ₂	15	35	0	0 ± 0 b	0 ± 0 e
3	He-O ₂	15	35	160	0 ± 0 b	NA
3	He-O ₂	15	35	320	0.1 ± 0.1 ab	0 ± 0 e
3	He-O ₂	15	55	0	0 ± 0 b	1.8 ± 0.3 cd
3	He-O ₂	15	55	160	0 ± 0 b	2.3 ± 0.4 bc
3	He-O ₂	15	55	320	0 ± 0 b	2.5 ± 0.2 ab
3	He-O ₂	15	75	0	0.2 ± 0.3 a	1.5 ± 0.2 d
3	He-O ₂	15	75	160	0 ± 0 b	3 ± 0.9 a
3	He-O ₂	15	75	320	0 ± 0 b	2.6 ± 0.8 ab
4	He-O ₂	15	35	0	0 ± 0 b	0 ± 0 c
4	He-O ₂	15	35	160	0.1 ± 0.2 b	NA
4	He-O ₂	15	35	320	0 ± 0 b	0 ± 0 c
4	He-O ₂	15	55	0	0.1 ± 0.1 b	1.5 ± 0.2 b
4	He-O ₂	15	55	160	0 ± 0 b	2.9 ± 0.6 a
4	He-O ₂	15	55	320	0.5 ± 0.4 a	2.8 ± 0.7 a

4	He-O ₂	15	75	0	0 ± 0 b	1.3 ± 0.2 bc
4	He-O ₂	15	75	160	0 ± 0 b	3.8 ± 1.6 a
4	He-O ₂	15	75	320	0 ± 0 b	3.3 ± 0.8 a
5	He	15	35	0	3.3 ± 0.4 d	0 ± 0 c
5	He	15	35	160	NA	NA
5	He	15	35	320	22.9 ± 5.7 b	1.8 ± 0.1 c
5	He	15	55	0	6 ± 2.2 cd	1.8 ± 0.2
5	He	15	55	160	NA	9.5 ± 2.7 a
5	He	15	55	320	35.1 ± 8.6 a	5.1 ± 1.8 bc
5	He	15	75	0	9.2 ± 0.4 c	1.9 ± 0.1 c
5	He	15	75	160	NA	4.8 ± 1.6 bc
5	He	15	75	320	26.8 ± 1.1 b	6.7 ± 0.8 b

611