

1 **Potential short-term losses of N<sub>2</sub>O and N<sub>2</sub> 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 NH<sub>3</sub> losses.  
11 Its proposed injection into soils as a counter-measure has been suggested to promote the generation of N<sub>2</sub>O, leading  
12 to a potential trade-off. Furthermore, the effect of high nutrient concentrations on N<sub>2</sub> losses as they may appear  
13 after injection of BD into soil has not yet been evaluated. Hence, we performed an incubation experiment with soil  
14 cores in a helium-oxygen atmosphere to examine the influence of soil substrate (loamy sand, clayey silt), water-  
15 filled pore space (WFPS; 35, 55, 75%) and application rate [0, 17.6 and 35.2 mL BD per soil core (250 cm<sup>3</sup>)] on  
16 the emission of N<sub>2</sub>O, N<sub>2</sub> and CO<sub>2</sub> after the usage of high loads of BD. To determine the potential capacity for  
17 gaseous losses, we applied anaerobic conditions by purging with helium for the last 24 h of incubation. Immediate  
18 N<sub>2</sub>O and N<sub>2</sub> emissions as well as the N<sub>2</sub>/(N<sub>2</sub>O+N<sub>2</sub>) product ratio depended on soil type and increased with WFPS  
19 indicating a crucial role of soil gas diffusivity for the formation and emission of nitrogenous gases in agricultural  
20 soils. However, emissions did not increase with the application rate of BD. This is probably due to an inhibitory  
21 effect of the high NH<sub>4</sub><sup>+</sup> content in BD on nitrification. Our results suggest a larger potential for N<sub>2</sub>O formation  
22 immediately following BD injection in the fine-textured clayey silt compared to coarse loamy sand. By contrast,  
23 the loamy sand showed a higher potential for N<sub>2</sub> production under anaerobic conditions. Our results suggest that  
24 short-term N losses of N<sub>2</sub>O and N<sub>2</sub> after injection may be higher than probable losses of NH<sub>3</sub> following surface  
25 application of BD.

## 26 **1 Introduction**

27 Nitrous oxide (N<sub>2</sub>O) is a potent greenhouse gas (Myhre et al., 2013), with agriculture being its largest single  
28 anthropogenic source, contributing about 4.1 Tg N<sub>2</sub>O-N yr<sup>-1</sup> or 66% of total gross anthropogenic emissions, mainly  
29 as a result of mineral nitrogen (N) fertiliser and manure application (Davidson and Kanter, 2014). The generation  
30 of nitrogen gas (N<sub>2</sub>) is of agronomic interest in terms of nutrient management, since such gaseous losses may imply  
31 a significant loss of N from the soil/plant system (Cameron et al., 2013; Friedl et al., 2016). However, from an  
32 environmental stance, N<sub>2</sub> is innocuous and, thus, the preferred type of gaseous N-loss from soil (Davidson et al.,  
33 2015). In general, the improvement of N use efficiency and thus the decrease of N losses in crop production are  
34 paramount in the presence of challenges like food security, environmental degradation and climate change (Zhang  
35 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<sub>4</sub><sup>+</sup>) and a narrowed carbon (C)/N ratio of BD (Möller and Müller, 2012). These altered chemical properties  
39 may promote biochemical reactions in the soil that are responsible for the formation of gaseous N species like  
40 N<sub>2</sub>O, nitric oxide (NO), N<sub>2</sub> and ammonia (NH<sub>3</sub>) (Nkoa, 2013).

41 Significant losses of N as NH<sub>3</sub> may occur within the first hours after manure application (Quakernack et al., 2012).  
42 To reduce NH<sub>3</sub> losses, the application of BD by injection is recommended, but this measure can simultaneously  
43 increase the potential for N<sub>2</sub>O losses compared to surface-application (Velthof and Mosquera, 2011; Wulf et al.,  
44 2002). On the one hand, high NH<sub>4</sub><sup>+</sup> concentrations in the injection band promote nitrification, consuming O<sub>2</sub> and  
45 releasing N<sub>2</sub>O (Christensen and Rowe, 1984). On the other hand, increased amounts of C in the injection band also  
46 promote respiration, additionally depleting O<sub>2</sub> supply (Dell et al., 2011). Altogether, the conditions during the

47 initial phase after injection of BD foster microsites favourable for microbial denitrification, which promote also  
48 the formation of N<sub>2</sub> due to anaerobic conditions (Köster et al., 2015; Webb et al., 2010).  
49 There is a wealth of biotic and abiotic processes in soils that produce N<sub>2</sub>O and N<sub>2</sub>, depending on mineral N content,  
50 C availability as well as on temperature, most of which are enhanced by anoxic or at least suboxic conditions  
51 (Butterbach-Bahl et al., 2013). The amounts and the relative share of N<sub>2</sub> and N<sub>2</sub>O in the overall gaseous N  
52 emissions depend – among other factors – on the degree of O<sub>2</sub> restriction (Firestone and Davidson, 1989). Soil  
53 physical and biotic factors [i.e. diffusion permitted by soil porosity in conjunction with water-filled pore space  
54 (WFPS) as well as consumption of O<sub>2</sub> by heterotrophic respiration and nitrification] control the aerobic status of  
55 a soil (Ball, 2013; Maag and Vinther, 1999; Uchida et al., 2008). In general, clayey soils exhibit a lower gas  
56 diffusivity compared to coarse textured soils. This regularly results in higher denitrification in the former with  
57 higher N<sub>2</sub>O emission rates, but also a higher probability for the consecutive reduction to N<sub>2</sub> (Ball, 2013; Gu et al.,  
58 2013; Senbayram et al., 2014).  
59 There is a general lack of knowledge about the effects of high BD concentration on gaseous N-losses as they might  
60 appear after injection into soils and their interactions with O<sub>2</sub> limiting factors like soil texture and WFPS, as well  
61 as temperature and heterotrophic respiration. Thus, we applied the helium-oxygen (He-O<sub>2</sub>) incubation technique  
62 (Butterbach-Bahl et al., 2002) in a laboratory experiment to evaluate the effect of the factors suggested above on  
63 the emission of N<sub>2</sub>O and N<sub>2</sub> from different soils. Simultaneously, CO<sub>2</sub> flux was determined as an indicator for  
64 microbial O<sub>2</sub> consumption, O<sub>2</sub> diffusion and also for the degradability of organic C applied with BD (Blagodatsky  
65 and Smith, 2012). We hypothesised that (1) N<sub>2</sub>O and N<sub>2</sub> emissions will increase with WFPS, (2) gaseous N losses  
66 will also be affected by BD application rate, i.e. the hypothetical concentration of C and N resulting from injection,  
67 and (3) the clayey silt will induce higher gaseous N losses than the coarse loamy sand.

## 68 **2 Material and Methods**

### 69 **2.1 Selected soils, sampling of soil cores and biogas digestate**

70 Two soils were selected and both were adjusted to three levels of WFPS and three quantities of BD (Table 1),  
71 resulting in 18 factor combinations with three replicates. Temperature was increased from 2 °C during the first  
72 two days to 15 °C for the last three days of the incubation. Intact soil cores (diameter 7.2 cm, height 6.1 cm, volume  
73 250 cm<sup>3</sup>) were taken with sample rings in the range from 0–0.10 m depth from two sites with different textures,  
74 i.e. sandy loam and clayey silt. The sandy loam samples were collected from a stagnic luvisol (IUSS Working  
75 Group WRB, 2006) located in Gülzow (North-East Germany) in the ground moraine of the Weichselian glacial  
76 period at 53° 48' 35" N and 12° 4' 20" E. The clayey silt samples were collected from a haplic luvisol located in  
77 Dornburg between the foothills and the lowlands of Central Germany at 51° 0' 8" N and 11° 39' 25" E (see Table  
78 2 for more details on soil characteristics). After field sampling, the soil cores were dried for 48 h at 40 °C to  
79 facilitate adjustment of WFPS.

80 Both sites have been cultivated with similar crop rotations used as feedstock for biogas production and have been  
81 amended with biogas digestate for the past nine years. The crop rotation on the sandy loam consisted of maize  
82 (*Zea mays* L.), rye (*Secale cereale* L.), sorghum (*Sorghum bicolor* (L.) MOENCH), winter triticale (×  
83 *Triticosecale* Wittmack), ryegrass (*Lolium perenne* L.) and winter wheat (*Triticum aestivum* L.). The only

84 difference in the crop rotation on the clayey silt was the cultivation of sudan grass (*Sorghum × drummondii*)  
85 instead of sorghum.

86 The biogas digestate used for the incubation was obtained from a biogas plant at 'Gut Dalwitz', an organic farm  
87 in northeast Germany. The feedstock for the anaerobic fermentation in the plant consisted of 60 % maize, 20 %  
88 solid cattle manure, 10 % dry chicken manure and 10 % rye. The digestate was analysed by LUFA Rostock,  
89 Germany and had a pH of 8.3, 2.91% organic C, 0.16% dissolved organic C (DOC), 0.54% N and 0.27% NH<sub>4</sub>-N  
90 in undried material with a dry matter content of 9.4%.

## 91 **2.2 Adjustment of WFPS and addition of N**

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

## 101 **2.3 Determination of gas fluxes**

102 The measurements of N<sub>2</sub>, N<sub>2</sub>O and CO<sub>2</sub> fluxes were applied following the He-O<sub>2</sub> method (Butterbach-Bahl et al.,  
103 2002; Scholefield et al., 1997). Six soil cores (i.e. the repetitions of two factor combinations at a time, Table 3)  
104 were placed simultaneously in special gas-tight incubation vessels inside a climate chamber. Analyses were  
105 conducted in the laboratory of the Institute for Landscape Biogeochemistry, Leibniz Centre for Agricultural  
106 Landscape Research (ZALF), Müncheberg, Germany. Before flux measurements, the vessels were evacuated to  
107 0.047 bar and flushed with an artificial He/O<sub>2</sub> gas mixture (20.49 % O<sub>2</sub>, 345.5 ppm CO<sub>2</sub>, 359 ppb N<sub>2</sub>O, 1863 ppb  
108 CH<sub>4</sub>, 2.46 ppm N<sub>2</sub>, rest He) four times consecutively to remove ambient N<sub>2</sub>. Subsequently, the air temperature of  
109 the climate chamber was set to 2 °C and a continuous He/O<sub>2</sub> gas flow rate of 15 ml min<sup>-1</sup> was applied to the vessel  
110 headspace for 72 h to remove residues of N<sub>2</sub> from soil cores by diffusion, including a restricted N<sub>2</sub> production by  
111 decreased microbial activity. After this pre-incubation in the following two days the headspace concentration of  
112 N<sub>2</sub>O and CO<sub>2</sub> was measured once daily in the morning. To compensate for the lower precision of the detector for  
113 N<sub>2</sub> in relation to the detector for N<sub>2</sub>O and CO<sub>2</sub> (Eickenscheidt et al., 2014), N<sub>2</sub> concentrations were measured  
114 consecutively three times daily in the morning. Immediately after the last measurement on the second day, the  
115 temperature was set to 15 °C and the measurements were continued for another two days. Finally, the He/O<sub>2</sub> gas  
116 mixture was substituted by pure He and, following 24 h of acclimatisation, gas measurements were carried out  
117 once again (Figure 1) to determine the generation of N<sub>2</sub>O and N<sub>2</sub> in a completely anaerobic soil matrix. The latter  
118 step is important to get a clue about the actual potential for gaseous N losses after highly concentrated BD  
119 application. The settings of the chromatographs for gas analyses are described in Eickenscheidt et al. (2014). Gas  
120 fluxes were calculated according to Eq. (1):

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

where  $f$  is the flux ( $\text{N}_2$  and  $\text{CO}_2$ :  $\text{mg m}^{-2} \text{h}^{-1}$ ,  $\text{N}_2\text{O}$ :  $\mu\text{g m}^{-2} \text{h}^{-1}$ ),  $M$  the molar mass in  $\text{g mol}^{-1}$  ( $\text{N}_2$ : 28,  $\text{CO}_2$ : 44,  $\text{N}_2\text{O}$ : 44),  $p$  the air pressure (Pa),  $v$  the air flow ( $\text{L h}^{-1}$ ),  $R$  the gas constant ( $8.31 \text{ J mol}^{-1} \text{ K}^{-1}$ ),  $T$  the temperature inside the chamber (K),  $A$  the area of the incubation vessel ( $\text{m}^2$ ), and  $dc$  the difference of gas concentrations ( $\text{N}_2$  and  $\text{CO}_2$ : ppm,  $\text{N}_2\text{O}$ : ppb) between inlet and outlet of a vessel.

To enhance the tightness against atmospheric  $\text{N}_2$  contamination, the lids of the incubation vessels were permanently purged with helium. We obtained blank values by inserting aluminium blocks into the vessels before each measurement cycle. Since this blank values were usually steady with means of 1.9 ( $1\sigma = 0.9$ ) ppm  $\text{N}_2$ , 349.6 ( $1\sigma = 11.4$ ) ppb  $\text{N}_2\text{O}$  and 353.9 ( $1\sigma = 13.5$ ) ppm  $\text{CO}_2$ , we suggest that the vessels were tight. Derived from the blank 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)  $\mu\text{g N}_2\text{O-N m}^{-2} \text{h}^{-1}$  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 measured at the respective outlet. Estimated fluxes from the soil cores smaller than the respective blank fluxes of each day were set to zero.

134

## 135 2.4 Soil analyses after incubation

136 After incubation, the soil cores were stored at 2 °C until they were extracted with 0.1 M KCl solution (soil to extract  
137 ratio 1:4, standardised extraction method of the commissioned laboratory at Leibniz Centre for Agricultural  
138 Landscape Research e. V.) and analysed for  $\text{NH}_4^+$  and nitrate ( $\text{NO}_3^-$ ) by spectrophotometry according to DIN ISO  
139 14256 with a continuous flow analyser ‘CFA-SAN’, Skalar Analytical B.V., the Netherlands and for DOC by  
140 combustion according to DIN ISO 10694 with an analyser ‘RC 612’, Leco Instruments GmbH, Germany.

## 141 2.5 Statistical analysis

142 All statistical analyses were done using R version 3.2.3 (R Core Team, 2015) with the data of the measuring days  
143 under He- $\text{O}_2$  atmosphere. Data from the vessels with the factor combination of 35% WFPS and LOBD with clayey  
144 silt were omitted due to technical reasons during sample preparation. For the final period of pure He headspace,  
145 some gas concentration data could not be documented. For loamy sand, this affects all WFPS levels with LOBD  
146 ( $\text{N}_2$  and  $\text{N}_2\text{O}$ ), the treatment 75% WFPS with 320  $\text{kg N h}^{-1}$  ( $\text{N}_2\text{O}$  and  $\text{CO}_2$ ) and for the clayey silt the treatment  
147 35% WFPS without amendment ( $\text{N}_2\text{O}$  and  $\text{CO}_2$ ).

148 To account for repeated measurement of vessels, linear mixed effect models were applied with package ‘lmerTest’  
149 version 2.0-33 (Kuznetsova et al., 2016) for fluxes of each gas type. The three pseudo-replicated fluxes from the  
150  $\text{N}_2$  measurements of each vessel were averaged for each day to obtain the same number of observations as for  $\text{N}_2\text{O}$   
151 and  $\text{CO}_2$  fluxes. The fixed structure of models included soil type, WFPS, amount of digestate, temperature,  $\text{NO}_3^-$   
152 and DOC contents after incubation as well as the fluxes of  $\text{N}_2\text{O}$  (in the model for  $\text{N}_2$ ) and  $\text{CO}_2$  (in the models for  
153  $\text{N}_2$ ,  $\text{N}_2\text{O}$  and  $\text{N}_2/[\text{N}_2+\text{N}_2\text{O}]$  product ratio). Soil  $\text{NH}_4^+$  was omitted since it showed high autocorrelation with the  
154 amount of BD applied. The individual soil cores in the vessels were set as random effect (nested within the week  
155 of incubation and with allowance for a variable slope of the effect each day) with regard to lack of independence  
156 of consecutive measurements. The model responses for  $\text{N}_2$ ,  $\text{N}_2\text{O}$  and  $\text{CO}_2$  were log transformed ( $\ln[\text{value} + 1]$ )

157 since gas fluxes from soils usually show lognormal distributions (Kaiser et al., 1998). The function 'step' was used  
158 for automatic backward selection of models based on AIC (Akaike's 'An Information Criterion'). The skewness  
159 ( $\gamma$ ) was calculated with R package 'moments' version 0.14 (Komsta and Novomestky, 2015) to check residuals for  
160 normal distribution and  $|\gamma| \leq 2$  was assumed as appropriate (West et al., 1995). For mixed effects models,  $p$ -values  
161 of the ANOVA (type 2) were calculated based on Satterthwaite's approximation.

162 Cumulated gas fluxes were estimated with a bootstrap method using function 'auc.mc' of R package 'flux' version  
163 0.3-0 (Jurasinski et al., 2014) for the R statistical software version 3.2.3 (R Core Team, 2015). In short, the fluxes  
164 for the period of aerobic headspace were cumulated in 100 iterations, while for each run two fluxes were omitted  
165 randomly. Then, the resulting data were used to calculate means and standard deviations.

## 166 **3 Results**

### 167 **3.1 Soil NH<sub>4</sub><sup>+</sup>, NO<sub>3</sub><sup>-</sup> and DOC contents**

168 The calculated application of NH<sub>4</sub><sup>+</sup>-N from BD per kg soil approximated for the sandy loam 135.8 mg (LOBD)  
169 and 271.5 mg (HIBD), and for the clayey silt 126.7 mg (LOBD) and 253.4 mg (HIBD). The NO<sub>3</sub><sup>-</sup> content of BD  
170 was negligible. In general, the NH<sub>4</sub><sup>+</sup> content of the soils after incubation increased with digestate application with  
171 lower amounts detected in the clayey silt. Nitrate was found almost exclusively in the latter soil. (Fig. 2).

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

### 174 **3.2 CO<sub>2</sub> fluxes**

175 CO<sub>2</sub> fluxes showed clear differences between the soils: under all combinations of temperature and oxygen, the  
176 fluxes were always larger from loamy sand compared to clayey silt (Table A1). In general, mean fluxes from  
177 loamy sand increased with the amount of digestate during each of the different periods regarding temperature and  
178 headspace aerobicity, but showed no obvious pattern with WFPS. There was no clear trend of fluxes with the  
179 amount of amendment, but a slight trend of decreasing fluxes with increasing WFPS for the clayey silt. However,  
180 the predictive power of WFPS on CO<sub>2</sub>-C fluxes was minor since it was eliminated during stepwise regression  
181 fitting. By contrast, soil type, amount of digestate, temperature as well as the DOC content after the incubation  
182 had significant ( $p < 0.01$ ) effects (Table 5).

### 183 **3.3 N<sub>2</sub>O fluxes**

184 The mean N<sub>2</sub>O fluxes from the loamy sand in the He-O<sub>2</sub> headspace were virtually zero, independent of temperature  
185 and WFPS as well as the amount of BD application (Fig. 3, Table A2). In contrast, the emissions of the clayey silt  
186 increased with temperature and were highest at 15 °C with intermediate WFPS and amount of BD, i.e. 6.2 mg  
187 N<sub>2</sub>O-N m<sup>-2</sup> h<sup>-1</sup> at 55% with LOBD, respectively. Surprisingly, at 15 °C, increasing the amount of BD up to HIBD  
188 did not increase the observed N<sub>2</sub>O efflux; rather it decreased the efflux significantly ( $p < 0.05$ , Tuckey's HSD) at  
189 55% and also, but not significantly, at 75% WFPS (Fig. 3, Table A2). According to the linear mixed model for

190 N<sub>2</sub>O fluxes in aerobic conditions, WFPS, amount of digestate, temperature, DOC content of soil after incubation  
191 and CO<sub>2</sub> fluxes had significant ( $p < 0.001$ ) effects on N<sub>2</sub>O flux (Table 5).

192 Under anaerobic headspace conditions, the overall highest mean N<sub>2</sub>O flux was observed from the clayey silt at  
193 35% WFPS with HIBD (11.7 mg N<sub>2</sub>O-N m<sup>-2</sup> h<sup>-1</sup>). The same soil showed a tendency of decreasing N<sub>2</sub>O fluxes with  
194 increasing WFPS and amendment. In the loamy sand, the pure He-atmosphere induced increasing mean N<sub>2</sub>O fluxes  
195 (up to 1.3 mg N<sub>2</sub>O-N m<sup>-2</sup> h<sup>-1</sup>) with increasing WFPS (Fig. 3, Table A2). Thus, the anaerobic headspace induced a  
196 change only in the loamy sand by increasing emissions.

### 197 **3.4 N<sub>2</sub> fluxes**

198 From the loamy sand, no or only small rates of N<sub>2</sub> were detected at both temperatures under He-O<sub>2</sub> atmosphere  
199 (Fig. 4, Table A3). The clayey silt showed mean fluxes of up to 1.4 mg N<sub>2</sub> m<sup>-2</sup> h<sup>-1</sup> at 2 °C (all incubations with  
200 75% WFPS) and up to 3.8 mg N<sub>2</sub> m<sup>-2</sup> h<sup>-1</sup> at 15 °C (75% WFPS with LOBD), but no fluxes in all BD treatments  
201 with 35% WFPS. Put simply, temperature had a small effect on N<sub>2</sub> emissions from the sandy loam with no  
202 consistent influence of WFPS and the amount of BD. In contrast, the clayey silt emitted increasing fluxes with  
203 increasing temperature and WFPS. However, the application rise from LOBD up to HIBD at 15 °C resulted in  
204 slightly, but not significantly ( $p > 0.05$ , Tuckey's HSD) decreased fluxes (Fig. 4, Table A3). The summary of the  
205 linear mixed model for N<sub>2</sub> fluxes under aerobic conditions revealed significant effects ( $p < 0.05$ ) of soil type,  
206 WFPS, the amount of digestate, temperature, DOC content after incubation and N<sub>2</sub>O flux (Table 5).

207 After switching the atmosphere to pure He, the N<sub>2</sub> fluxes from the sandy loam increased more than 60-fold. In  
208 contrast to aerobic conditions, all measured factor combinations showed mean fluxes up to 35.1 mg N<sub>2</sub> m<sup>-2</sup> h<sup>-1</sup>  
209 (55% with 320 kg N ha<sup>-1</sup>) (Fig. 2, Day 5 in Table A3). Mean fluxes from clayey silt increased only up to 9.3 mg  
210 N<sub>2</sub> m<sup>-2</sup> h<sup>-1</sup> in amended treatments. Thus, the loamy sand exhibited a much more intense reaction under anaerobic  
211 headspace conditions.

### 212 **3.5 N<sub>2</sub>/(N<sub>2</sub> + N<sub>2</sub>O-N) product ratio**

213 No clear trend of the product ratio of N<sub>2</sub>/(N<sub>2</sub> + N<sub>2</sub>O-N) was found for incubations of the loamy sand. However,  
214 there was a clear distinction of the ratios for this soil under aerobic and anaerobic atmospheres: while the ratios  
215 were close to zero in the former, they were close to one in the latter (Fig. 5). In contrast, in the clayey silt the ratios  
216 increased with WFPS and were affected by digestate amendment under both the aerobic and the anaerobic  
217 atmospheres, where the highest ratios (up to 0.8) were found in treatments without digestate and at least 55%  
218 WFPS. The digestate-amended treatments showed ratios around or above 0.5, with exception of the 35% WFPS  
219 treatments, which had ratios close to zero. According to the linear mixed model, the product ratio under aerobic  
220 conditions was affected significantly ( $p < 0.01$ ) by soil type and the amount of digestate (Table 5).

## 221 4 Discussion

### 222 4.1 Increased BD application rate did not increase N<sub>2</sub>O and N<sub>2</sub> losses probably due to inhibitory effect 223 of high NH<sub>4</sub><sup>+</sup> concentrations

224 In the loamy sand, the higher NH<sub>4</sub><sup>+</sup> content measured after the incubation cycle compared to the calculated NH<sub>4</sub><sup>+</sup>  
225 application rates may result from heterogeneity of BD itself (Andruschkewitsch et al., 2013). By contrast, the  
226 considerable lower values after incubation in the clayey silt could be attributed to a higher fixation of NH<sub>4</sub><sup>+</sup> as NH<sub>3</sub>  
227 by clay minerals, enhanced by the increased pH of BD (Kissel et al., 2008).

228 The overall N<sub>2</sub>O fluxes corresponded well with those from other studies with similar incubation conditions and  
229 application rates of BD in terms of N ha<sup>-1</sup> (Köster et al., 2015; Senbayram et al., 2012; Severin 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 than we actually applied. Although we observed differences in N<sub>2</sub>O emissions between soils,  
232 soil type was not confirmed as a significant effect. Nevertheless, WFPS and temperature, which are well known  
233 controls of N<sub>2</sub>O generation (Maag and Vinther, 1999), showed significant influences. Both are physical (by gas  
234 diffusion) and biological (by increased metabolic activity and consequently increased O<sub>2</sub> consumption by  
235 respiration) drivers for O<sub>2</sub> availability (Ball, 2013; Maag and Vinther, 1999). Accordingly, CO<sub>2</sub> flux (resulting  
236 from respiration of O<sub>2</sub>) generally increased with temperature and was also identified as significant by regression  
237 selection.

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

247 N<sub>2</sub>/(N<sub>2</sub>+N<sub>2</sub>O) product 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 loamy sand to generally adverse conditions for the formation of N<sub>2</sub>O and N<sub>2</sub>, i.e. a sufficient supply of O<sub>2</sub>  
250 (see section 4.2). Contrary, the influence of WFPS apparently mirrored favourable conditions in the clayey silt  
251 (Table 5). Simultaneously, with increasing WFPS, the reduction of N<sub>2</sub>O as an alternative electron acceptor under  
252 reduced O<sub>2</sub> supply accelerates (Tiedje, 1988). Accordingly, no or rather small fluxes of the investigated gaseous  
253 N species were generally found in our presumably well aerated treatments with 35% WFPS.

254 In our study, one treatment (clayey silt, 55% WFPS, LOBD) showed exceptionally large mean N<sub>2</sub>O fluxes of up  
255 to 7.1 mg N m<sup>-2</sup> h<sup>-1</sup> (Fig. 3, Table A2). This could be evidence that the injection of such commonly applied amounts  
256 of BD-N (i.e., 160 kg N ha<sup>-1</sup>) may favour much larger losses of N<sub>2</sub>O compared to an even distribution of BD in a  
257 soil surface due to larger substrate concentration in injection slits. However, with higher amendments (i.e. HIBD),  
258 we observed partially significant ( $p < 0.05$ , Tuckey's HSD) reductions of N<sub>2</sub>O and a decreasing tendency of N<sub>2</sub>



259 emissions (Table A2, Table A3). In line with this, the amount of BD showed a significant effect during the  
260 regression selection on N<sub>2</sub>O, but not on N<sub>2</sub> fluxes (Table 5). A coherent reasoning for the rather smaller emissions  
261 of highly amended HIBD treatments might lie in an inhibitory effect of NH<sub>3</sub> on nitrification. Accordingly, Kim et  
262 al. (2006) found a selective inhibition of NO<sub>2</sub><sup>-</sup> oxidation in the presence of 14 to 17 mg NH<sub>3</sub>-N L<sup>-1</sup>. Our calculated  
263 application rates in the treatments with HIBD amounted to at least 253.4 mg NH<sub>4</sub><sup>+</sup>-N (kg soil)<sup>-1</sup> for the clayey silt  
264 (Fig. 3) which corresponds to 13.0 mg NH<sub>3</sub>-N (kg soil)<sup>-1</sup> at 15 °C when applying the pH of the BD and assuming  
265 all extractable NH<sub>4</sub><sup>+</sup>-N to be in solution (Emerson et al., 1975). Hence, we consider this inhibitory effect as the  
266 reason for the missing increase of N<sub>2</sub>O and N<sub>2</sub>. Nevertheless, because we mixed the BD with the soil, one would  
267 expect a lower NH<sub>3</sub> fixation by clay minerals in tubular injection slits *in situ* (Kissel et al., 2008), resulting in  
268 probably lower N<sub>2</sub>O and N<sub>2</sub> fluxes from clayey soils due to an more marked inhibitory effect.  
269 High NH<sub>4</sub><sup>+</sup> loads in conjunction with an increased pH favour NO<sub>2</sub><sup>-</sup> accumulation because NO<sub>2</sub><sup>-</sup> oxidising bacteria  
270 are less resilient against high concentrations of NH<sub>3</sub> than NH<sub>3</sub> oxidising bacteria (Anthonisen et al., 1976). This  
271 NO<sub>2</sub><sup>-</sup> should have protonated then partly to toxic and unstable HNO<sub>2</sub>, which drives biological and chemical  
272 production of NO and N<sub>2</sub>O for detoxification (Venterea et al., 2015). Although we did not determine NO<sub>2</sub><sup>-</sup>, we  
273 suggest a dominant role of nitrifier denitrification, i.e., NO<sub>2</sub><sup>-</sup> reduction, in the generation of N<sub>2</sub>O during our  
274 experiment, especially during the anaerobic headspace conditions at the end of the incubation, resulting in the  
275 relatively small NO<sub>3</sub><sup>-</sup> recovery in both soils. Accordingly, coupled nitrification-denitrification and bacterial  
276 denitrification have been found to dominate the production of N<sub>2</sub>O directly after application of BD (Köster et al.,  
277 2011; Senbayram et al., 2009). However, N<sub>2</sub>O-N losses were clearly larger than N<sub>2</sub> losses under aerobic headspace  
278 in the clayey silt. This indicates that much of the N gas loss was driven by processes other than canonical  
279 denitrification. Under the above-mentioned conditions, NO-N losses may exceed N<sub>2</sub>O losses (Venterea et al.,  
280 2015), calling for taking account of NO measurements in future studies.  
281 Supposed that 15 % of NH<sub>4</sub><sup>+</sup>-N is volatilised as NH<sub>3</sub> within the first ten hours after surface application of BD  
282 (Quakernack et al., 2012), the losses from the NH<sub>4</sub><sup>+</sup> amounts we applied would averaged to 80 mg NH<sub>3</sub>-N m<sup>-2</sup> h<sup>-1</sup>  
283 (LOBD) and 160 mg NH<sub>3</sub>-N m<sup>-2</sup> h<sup>-1</sup> (HIBD). The actual losses of up to 11.7 mg N<sub>2</sub>O-N m<sup>-2</sup> h<sup>-1</sup> at 30 % WFPS in  
284 the clayey soil (Table A2) or of up to 35.1 mg N<sub>2</sub> m<sup>-2</sup> h<sup>-1</sup> at 55 % WFPS in the sandy loam (Table A3) from our  
285 HIBD treatments add up to 117 mg N<sub>2</sub>O-N and 351 mg N<sub>2</sub>, respectively, for the same period. Hence, increased  
286 N<sub>2</sub>O and N<sub>2</sub> emissions following injection of BD might effectively cause higher N losses compared to a surface  
287 application and deserve closer attention in future.

## 288 4.2 Different effects of soil diffusivity on N<sub>2</sub>O and N<sub>2</sub> fluxes

289 Apparently, the tested factors affected the N<sub>2</sub>O and N<sub>2</sub> fluxes from both soils in a different way. A specific soil  
290 characteristic that exhibits such a fundamental control on biogeochemical processes such as denitrification is the  
291 diffusivity for O<sub>2</sub> (Ball, 2013; Letey et al., 1980; Parkin and Tiedje, 1984), which is a main soil characteristic  
292 responsible for the appearance of anaerobic microsites. In general, diffusivity integrates the soil porosity, i.e., pore  
293 continuity and size as well as WFPS, which control both soil N<sub>2</sub>O and N<sub>2</sub> emissions (Balaine et al., 2016; Ball,  
294 2013; Letey et al., 1980). Soils with a coarser texture like the loamy sand have a higher proportion of macropores  
295 and thus a higher gas diffusion compared with fine textured soils like the clayey silt we used (Groffman and Tiedje,  
296 1991). This lets us expect conditions that are more favourable for N<sub>2</sub>O and N<sub>2</sub> generation in the latter due to

297 relatively poor diffusion characteristics and, thus, a smaller O<sub>2</sub> supply. Actually, although we incubated the soils  
298 at comparable levels of WFPS and BD amendments, the apparent lower diffusivity led to larger N<sub>2</sub>O and N<sub>2</sub>  
299 production in the treatments with the clayey silt in relation to the loamy sand.

300 The role of the distinct diffusivities of both soils is corroborated by our observations of the gas fluxes in anaerobic  
301 headspace. With switching the He-O<sub>2</sub> atmosphere in the headspace to pure He, the denitrification potential can be  
302 tested because anaerobicity eliminates respiration processes that use O<sub>2</sub> as electron acceptor (Parkin and Tiedje,  
303 1984). We acknowledge e.g. DNRA and anammox as possible additional sources of N<sub>2</sub>O and N<sub>2</sub> under such  
304 conditions but we were not able to quantify their contribution. The anaerobic headspace induced a considerable  
305 increase of N<sub>2</sub>O fluxes in the loamy sand, but not in the clayey silt. Concurrently, the N<sub>2</sub> fluxes increased in both  
306 soils, but pronounced, i.e. more than 60-fold, in the sandy loam. These observed changes resulting from oxygen  
307 deprivation imply that, during the previous aerobic conditions, the diffusivity of the sandy loam was too high to  
308 allow for a sufficient establishment of anaerobic microsites while the clayey silt ensured a moderate diffusional  
309 constraint to maintain suboxic conditions. In general, only N<sub>2</sub>O fluxes from treatments with negligible fluxes  
310 during the previous aerobic period increased under anaerobic conditions, including all treatments with loamy sand  
311 (Fig. 3, Table A2). At the same time, there was a reduction of N<sub>2</sub>O fluxes in most clayey silt treatments. However,  
312 a closer look reveals that virtually all of the latter treatments showed increased N<sub>2</sub> flux rates. Hence, there was an  
313 enhanced reduction of N<sub>2</sub>O to N<sub>2</sub>, which is reflected in the increased N<sub>2</sub>/(N<sub>2</sub> + N<sub>2</sub>O) product ratio (Fig. 5) and  
314 points to intensified reduction of N<sub>2</sub>O due to the lack of oxygen (Parkin and Tiedje, 1984). The much larger N<sub>2</sub>  
315 fluxes from the loamy sand compared to the clayey silt might have been caused as well by poor NO<sub>3</sub><sup>-</sup> availability  
316 (Fig. 2) and a high availability of C (Table 4), which promoted the reduction of N<sub>2</sub>O to N<sub>2</sub> (Senbayram et al.,  
317 2012). Further, we found no evidence for any shortage of substrate in the clayey silt during the subsequent  
318 anaerobic headspace conditions. However, the cumulated fluxes of both N<sub>2</sub> and N<sub>2</sub>O amounted to a maximum  
319 absolute loss of 9.4 (1σ = 0.3) mg N per kg soil in the clayey silt with LOBD and 55% WFPS, which was 7.4 %  
320 of the calculated NH<sub>4</sub><sup>+</sup>-N applied with BD (Fig. 2). On the other hand, the N<sub>2</sub>/(N<sub>2</sub>+N<sub>2</sub>O) product ratios increased  
321 only slightly (Fig. 5) and, in contrast to the loamy sand, there were still significant N<sub>2</sub>O fluxes in the clayey silt  
322 (Fig. 3). This points to still sufficient stocks of NO<sub>3</sub><sup>-</sup> in the latter (Senbayram et al., 2012). In fact, the NO<sub>3</sub><sup>-</sup> stock  
323 was greater in the clayey silt than in loamy sand after incubation (Fig. 2). Thus, we suggest that the gas fluxes  
324 were unaffected by the change to anaerobic headspace in the clayey silt due to already low O<sub>2</sub> concentrations as  
325 a result of poor diffusivity. In conclusion, distinct gas diffusivities of both soils can be proposed as the main reason  
326 for the differing N<sub>2</sub>O and N<sub>2</sub> fluxes.

327 In interaction with soil diffusivity, respiration affects the aerobicity of a soil matrix by concurrent consumption  
328 and formation of O<sub>2</sub> and CO<sub>2</sub> as well. Depending on microbial C availability, respiration could be indicated by  
329 DOC, though not all DOC might be readily degradable (Cook and Allan, 1992). Generally, DOC content after our  
330 incubation increased with application rate of BD (Table 4), but DOC content was always smaller in clayey silt.  
331 This might reflect a stronger sorption of C and thus a lower availability for respiration in the clayey silt compared  
332 to loamy sand (Kaiser and Guggenberger, 2000). If we compare DOC concentrations with cumulated flux rates of  
333 CO<sub>2</sub> over the period of aerobic headspace, we find a good regression fit ( $R^2 = 0.91$ ,  $p < 0.001$ ) for both soils (Fig.  
334 6) indicating a sufficient availability of C from BD for respiration and, thus, implicitly also for denitrification  
335 (Reddy et al., 1982). Moreover, as increased DOC enhanced respiration (Table A1), it consequently affected O<sub>2</sub>

336 consumption and, thus, also the emergence of anaerobic microsites (Azam et al., 2002). Accordingly, there is also  
337 a good correlation between cumulated CO<sub>2</sub> and N<sub>2</sub>O + N<sub>2</sub> fluxes for the same period from the clayey silt ( $R^2 =$   
338  $0.93$ ,  $p = 0.001$ ), when the treatments with 35 % WFPS (which showed virtually no N emissions) are omitted (Fig.  
339 7). 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<sub>2</sub>O and N<sub>2</sub>, which, nevertheless,  
341 interacted with the inhibitory effect of high NH<sub>4</sub><sup>+</sup> 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<sub>2</sub> and N<sub>2</sub>O 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. We assume also the measurements after  
348 only 24 hours of anaerobicity in the headspace to be representative for the emission potential since Wang et al.  
349 (2011; 2013) showed in similar studies that the emission of N<sub>2</sub> and N<sub>2</sub>O peaked within less than 24 hours after  
350 switching their headspace from aerobic to anaerobic conditions.

351 In summary, as hypothesised, N<sub>2</sub>O and N<sub>2</sub> emissions as well as the N<sub>2</sub>/(N<sub>2</sub>O+N<sub>2</sub>) product ratio increased with  
352 WFPS, most probably due to restricted supply of O<sub>2</sub>. Contrary to our second hypothesis, the gaseous losses of N<sub>2</sub>O  
353 and N<sub>2</sub> did not increase with the application rate of BD. This indicates an inhibitory effect of high NH<sub>3</sub> and NH<sub>4</sub><sup>+</sup>  
354 concentrations on nitrification, which are found typically in BD. At the same time, the N<sub>2</sub>/(N<sub>2</sub>O+N<sub>2</sub>) product ratio  
355 tended to decrease with application rate as supposed, probably due to a copious supply with NO<sub>2</sub><sup>-</sup> and NO<sub>3</sub><sup>-</sup> from  
356 oxidised BD-NH<sub>4</sub><sup>+</sup>. Confirming our third hypothesis, the fine textured clayey silt induced larger gaseous N losses  
357 and a higher N<sub>2</sub>/(N<sub>2</sub>O+N<sub>2</sub>) ratio than the coarse loamy sand by the apparent distinct diffusivities of both soils.  
358 Overall, there was a larger potential for formation of N<sub>2</sub>O in the fine-textured clayey silt compared to the coarse  
359 loamy sand after the application of high concentrations of BD as they may appear after injection. However, the  
360 loamy sand showed a large potential for N<sub>2</sub> formation under anaerobic headspace conditions.

361 Since coupled nitrification-denitrification N losses from injected BD seem to be massive in your study, the short-  
362 term emissions of N<sub>2</sub>O and N<sub>2</sub> after injection appear to offset the reduced NH<sub>3</sub>-N losses that would have arose  
363 hypothetically from surface application. Further investigations are needed in regarding the dynamics and the  
364 duration of the observed effects and their reliability for field conditions.

## 365 **6 Data availability**

366 The data underlying the figures are accessible publicly via the Supplement.

367

368 **Acknowledgements** We thank the editor Karsten Kalbitz and three anonymous referees for their careful reading,  
369 critical comments and valuable suggestions. We are very grateful to Heinrich Graf von Bassewitz and Matthias  
370 Haß from Gut Dalwitz for their straightforward support with substrate from their anaerobic digester. We thank  
371 Madlen Pohl from the ZALF, Institute for Landscape Biogeochemistry, Müncheberg, Germany, most sincerely

372 for managing the laboratory analyses of the soil samples. The joint research project underlying this report was  
373 funded by the German Federal Ministry of Food and Agriculture under the funding identifier 22007910.

374

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519

520

521 **Table 1: The examined factors soil texture, water-filled pore space (WFPS), and amount (i.e., concentration) of nitrogen**  
522 **(N) applied with biogas digestate (BD) with their respective levels applied in the present study, resulting in 18**  
523 **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 <sup>-1</sup> ) [3]	0	160	320
Temperature (°C) [2]	2	15	

524



525 **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,  $n = 3$ )**  
 526 **and mineral N ( $\text{NO}_3^-$  and  $\text{NH}_4^+$ ,  $n = 3$ ) of both soils in 0–10 cm depth after field sampling.**

Texture	C ( $\text{mg g}^{-1}$ ) <sup>a</sup>	N ( $\text{mg g}^{-1}$ ) <sup>a</sup>	pH <sup>b</sup>	Bulk density ( $\text{g cm}^{-3}$ ) <sup>c</sup>	$\text{NO}_3^-$ ( $\text{mg kg}^{-1}$ ) <sup>d</sup>	$\text{NH}_4^+$ ( $\text{mg kg}^{-1}$ ) <sup>d</sup>
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)

527 <sup>a</sup> measured with analyser “Truspec CNS”, Leco Instruments GmbH, Germany, performed according to ISO 10694 (“elemental analysis”) for C and according to ISO 13878  
 528 (“elemental analysis”) for N

529 <sup>b</sup> measured in  $\text{H}_2\text{O}$  with TitraMaster85, Radiometer Analytical SAS, France, performed according to VDLUFA Methodenbuch, Vol. 1, chap. 5.1.1

530 <sup>c</sup> measured on 250  $\text{cm}^3$  soil cores

531 <sup>d</sup> measured with analyser “CFA-SAN”, Skalar Analytical B.V., the Netherlands, performed according to ISO 14256

532 **Table 3: Chronological order of the incubated factor combinations. Two different factor combinations with their**  
 533 **respective repetitions ( $n = 3$ ) were placed together for each weekly incubation course (cf. Fig. 1). The factors were**  
 534 **combined by (1) soil (loamy sand: LS, clayey silt: CS), (2) amount (kg) of applied N from digestate per ha and (3) WFPS**  
 535 **(%).**

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%

536

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

kg digestate-N ha <sup>-1</sup>	WFPS (%)	mg DOC (kg soil) <sup>-1</sup>	
		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)

539 n.a.: data not available

540 **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<sub>2</sub>, N<sub>2</sub>O, N<sub>2</sub>/(N<sub>2</sub>+N<sub>2</sub>O) product**  
 541 **ratio and CO<sub>2</sub> in aerobic He-O<sub>2</sub> atmosphere. Soil type, water-filled pore space (WFPS), amount of digestate, temperature, NO<sub>3</sub><sup>-</sup> and DOC content of soil after incubation as well as**  
 542 **fluxes of N<sub>2</sub>O and CO<sub>2</sub> were set as possible independent variables. The individual soil rings were set as random effect (nested within the respective week and with the allowance for**  
 543 **varying slopes for each day of measurements). The random effect was always significant.**

544

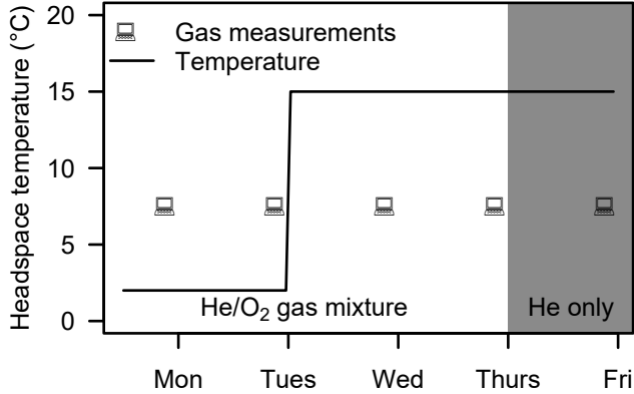
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Response	Soil type	WFPS	Digestate amount	Temperature	NO <sub>3</sub> <sup>-</sup> post	DOC post	N <sub>2</sub> O flux	CO <sub>2</sub> flux
N <sub>2</sub>	0.026	< 0.001	0.008	0.037	†	0.001	< 0.001	†
N <sub>2</sub> O	†	< 0.001	< 0.001	< 0.001	†	< 0.001	*	< 0.001
N <sub>2</sub> /(N <sub>2</sub> +N <sub>2</sub> O)	0.005	0.004	†	†	†	†	*	†
CO <sub>2</sub>	<0.001	†	<0.001	<0.001	†	0.007	†	*

545 † Variable eliminated during stepwise regression selection

546 \* Variable was not included into original regression

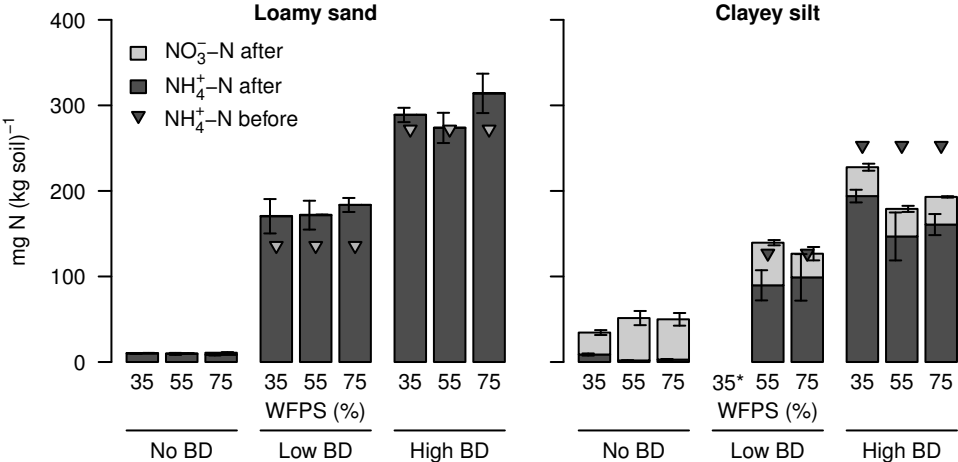
547 **Figure 1: Course of incubation and gas measurements with respect to atmosphere and temperature of the headspace**  
548 **after two days of pre-incubation at 2 °C in He/O<sub>2</sub> gas mixture. Gas concentrations of the headspace were determined**  
549 **on five consecutive days, i.e. Monday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

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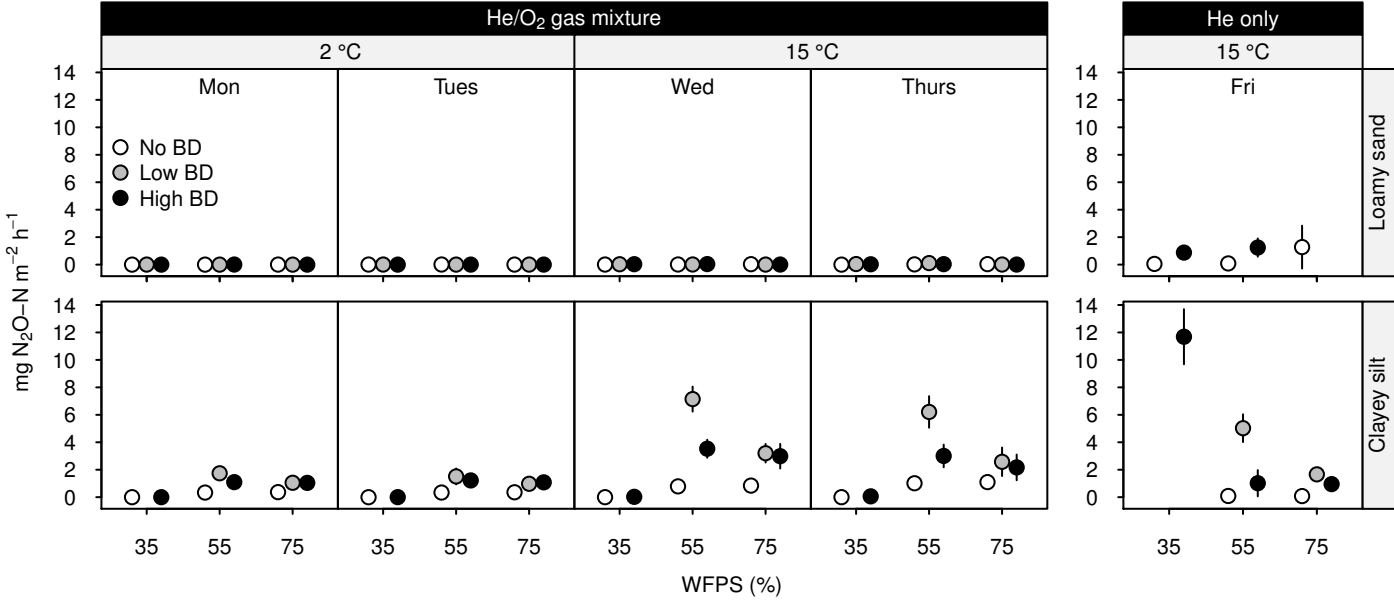
553 **Fig. 2: Ammonium and nitrate contents from loamy sand and clayey silt after incubation with different water-filled**  
554 **pore spaces (WFPS, %) and amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and 35.2 mL:**  
555 **'High BD'). Error bars denote standard deviations. In general, the ammonium content increased with digestate**  
556 **application with lower amounts detected in the clayey silt. Nitrate was found almost exclusively in the latter soil. For**  
557 **comparison, inverted triangles show calculated amounts of applied ammonium, which may differ from actual rates due**  
558 **to heterogeneity of biogas digestate. One treatment (\*) was omitted from all analyses due to technical reasons.**

559

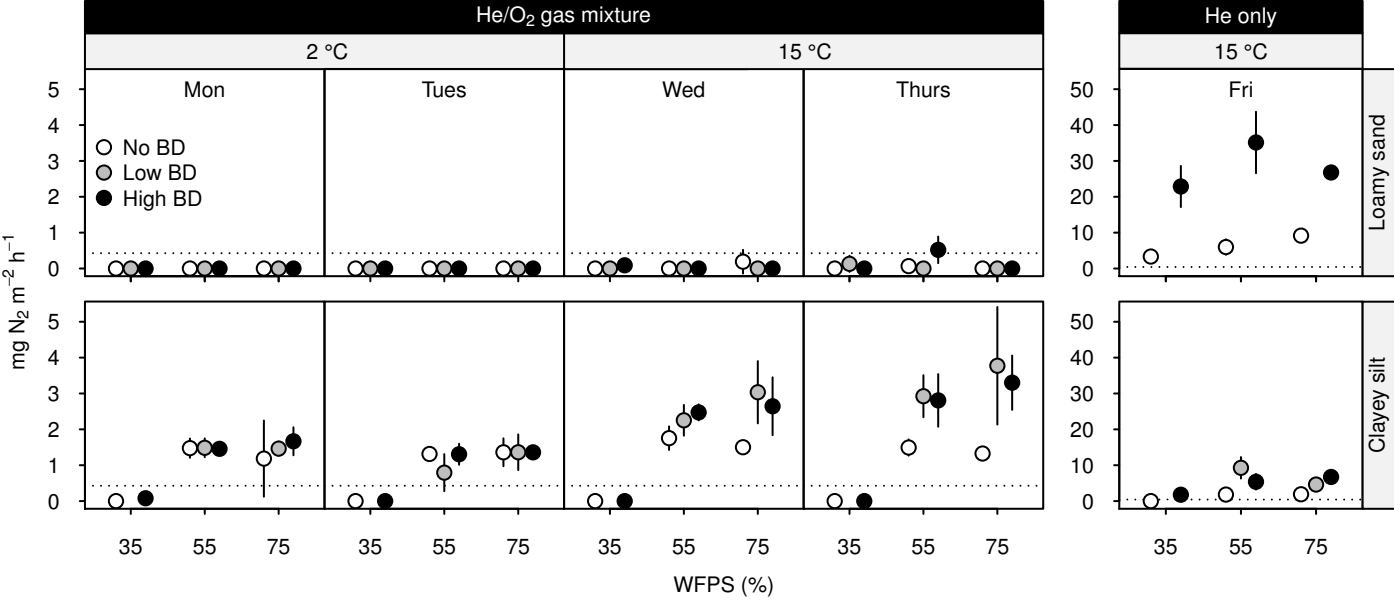




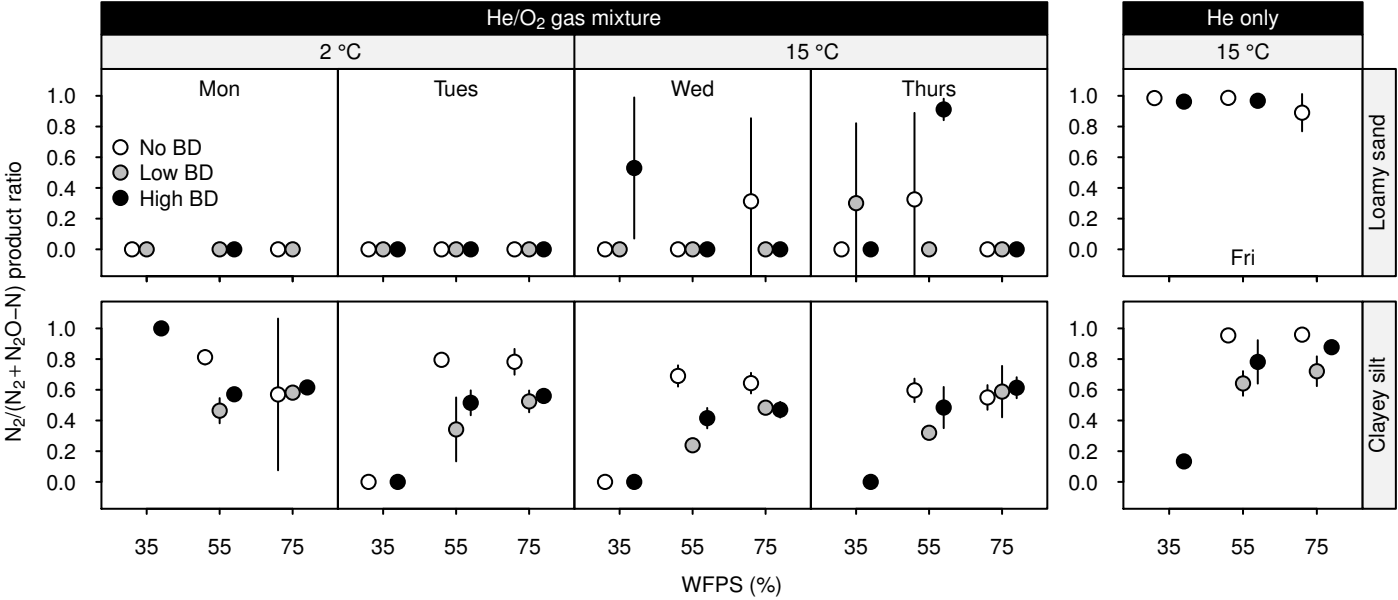
560 Fig. 3: Mean N<sub>2</sub>O fluxes (mg N m<sup>-2</sup> h<sup>-1</sup>) from a loamy sand and a clayey silt incubated under different water-filled pore  
561 spaces (WFPS, %) with different amounts of digestate (0 mL per sample ring: 'No BD', 17.6 mL: 'Low BD' and 35.2  
562 mL: 'High BD'). The first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



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

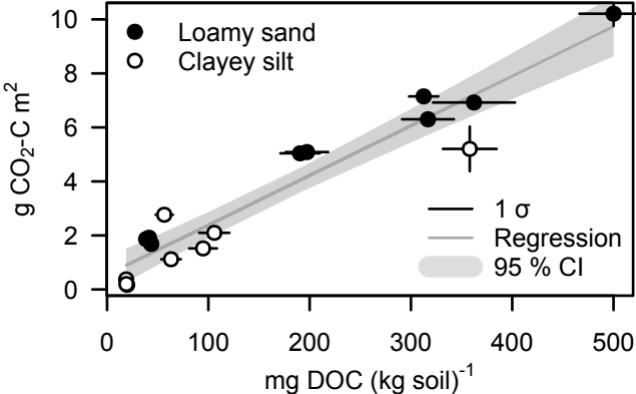


581 **Fig. 5: Mean  $N_2/(N_2 + N_2O-N)$  product ratio from a loamy sand and a clayey silt incubated under different water-filled**  
582 **pore spaces (WFPS, %) with different amounts of digestate (0 mL per sample ring: ‘No BD’, 17.6 mL: ‘Low BD’ and**  
583 **35.2 mL: ‘High BD’). The first till the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



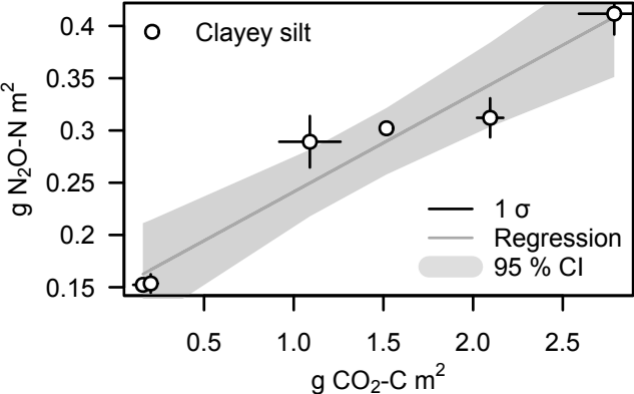
590 **Fig. 6: Regression between DOC (mg per 100 g soil) measured after the incubation and the respective cumulated CO<sub>2</sub>**  
591 **emissions (g C m<sup>-2</sup>) during the period of aerobic headspace with their standard deviations and confidence interval**  
592 **(95%). If error bars are not visible, they are smaller than the symbols of the means. Both soils showed increasing**  
593 **emissions with increasing soil DOC contents as well a good regression fit ( $R^2 = 0.91, p < 0.001$ ).**

594





595 **Fig. 7: Regression between cumulated CO<sub>2</sub> emissions (g C m<sup>-2</sup>) and the respective cumulated N<sub>2</sub>O + N<sub>2</sub> emissions (g N**  
596 **m<sup>-2</sup>) from the clayey silt with WFPS > 35 % during the period of aerobic headspace with their standard deviations and**  
597 **confidence interval (95%). If error bars are not visible, they are smaller than the symbols of the means. The proportional**  
598 **increase of CO<sub>2</sub> and the N gas species shows a good regression fit of  $R^2 = 0.93$ , ( $p = 0.001$ ).**



599 **Table A1: Mean CO<sub>2</sub>-C fluxes with standard deviations in mg m<sup>-2</sup> h<sup>-1</sup> from the loamy sand and the clayey silt, treated**  
600 **with different water-filled pore spaces (WFPS, %), amounts of digestate (kg N ha<sup>-1</sup>) as well as different temperature**  
601 **regimes (°C) under aerobic (He-O<sub>2</sub>) 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 <sup>-1</sup>	mg CO <sub>2</sub> -C m <sup>-2</sup> h <sup>-1</sup>	
					Loamy sand	Clayey silt
1	He-O <sub>2</sub>	2	35	0	6.8 ± 2.4 cd	0 ± 0 c
1	He-O <sub>2</sub>	2	35	160	22 ± 3.5 bcd	NA
1	He-O <sub>2</sub>	2	35	320	23.3 ± 9.3 bc	22.8 ± 2.8 ab
1	He-O <sub>2</sub>	2	55	0	6 ± 0.7 d	4.6 ± 7.9 bc
1	He-O <sub>2</sub>	2	55	160	34.4 ± 3.1 b	34.5 ± 11.6 a
1	He-O <sub>2</sub>	2	55	320	28 ± 3.2 b	15.9 ± 3.4 abc
1	He-O <sub>2</sub>	2	75	0	9.4 ± 1.4 cd	0 ± 0 c
1	He-O <sub>2</sub>	2	75	160	37.5 ± 6 b	15.5 ± 12.1 abc
1	He-O <sub>2</sub>	2	75	320	68.3 ± 12.1 a	24.5 ± 2.7 a
2	He-O <sub>2</sub>	2	35	0	9.8 ± 3.5 c	1.3 ± 1.4 b
2	He-O <sub>2</sub>	2	35	160	23 ± 3.9 bc	NA
2	He-O <sub>2</sub>	2	35	320	30.9 ± 2.2 b	22.2 ± 2.4 a
2	He-O <sub>2</sub>	2	55	0	8.7 ± 1.5 c	0.6 ± 1 b
2	He-O <sub>2</sub>	2	55	160	33.4 ± 0.9 b	27.6 ± 12.3 a
2	He-O <sub>2</sub>	2	55	320	35.9 ± 2.7 b	14.4 ± 1.9 ab
2	He-O <sub>2</sub>	2	75	0	8.3 ± 1.5 c	0 ± 0 b
2	He-O <sub>2</sub>	2	75	160	31.9 ± 3 b	13 ± 9.3 ab
2	He-O <sub>2</sub>	2	75	320	57.6 ± 14.8 a	18.3 ± 4 a
3	He-O <sub>2</sub>	15	35	0	42.5 ± 4.5 c	6.7 ± 0.7 b
3	He-O <sub>2</sub>	15	35	160	114.3 ± 12.2 b	NA
3	He-O <sub>2</sub>	15	35	320	149.5 ± 9.4 b	130.9 ± 105 a
3	He-O <sub>2</sub>	15	55	0	41.3 ± 3.5 c	3.2 ± 0.4 b
3	He-O <sub>2</sub>	15	55	160	108.7 ± 10.1 b	57.8 ± 12.2 bc
3	He-O <sub>2</sub>	15	55	320	162.1 ± 9.6 b	26.8 ± 0.7 bc
3	He-O <sub>2</sub>	15	75	0	44.1 ± 9.8 c	3.2 ± 0.7 b
3	He-O <sub>2</sub>	15	75	160	150.4 ± 19 b	26.4 ± 11.8 bc
3	He-O <sub>2</sub>	15	75	320	249.7 ± 53.5 a	35.3 ± 6 bc
4	He-O <sub>2</sub>	15	35	0	48.7 ± 6 c	15.1 ± 4.9 cd
4	He-O <sub>2</sub>	15	35	160	114.3 ± 6.4 b	NA
4	He-O <sub>2</sub>	15	35	320	156.9 ± 15.4 a	65.7 ± 2.2 a
4	He-O <sub>2</sub>	15	55	0	48 ± 3.4 c	4.2 ± 0.2 d
4	He-O <sub>2</sub>	15	55	160	109 ± 14.4 b	51.2 ± 15.1 ab

4	He-O <sub>2</sub>	15	55	320	177.7 ± 7.5 a	26.6 ± 2.3 cd
4	He-O <sub>2</sub>	15	75	0	34 ± 7.8 c	6.7 ± 4 d
4	He-O <sub>2</sub>	15	75	160	168.7 ± 0.4 a	22.1 ± 14.8 cd
4	He-O <sub>2</sub>	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

604

605 **Table A2: Mean N<sub>2</sub>O-N fluxes with standard deviations in mg m<sup>-2</sup> h<sup>-1</sup> from the loamy sand and the clayey silt, treated**  
606 **with different water-filled pore spaces (WFPS, %), amounts of digestate (kg N ha<sup>-1</sup>) as well as different temperature**  
607 **regimes (°C) under aerobic (He-O<sub>2</sub>) 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.**

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

4	He-O <sub>2</sub>	15	55	320	0 ± 0 ab	3 ± 0.8 b
4	He-O <sub>2</sub>	15	75	0	0 ± 0 ab	1.1 ± 0.3 bc
4	He-O <sub>2</sub>	15	75	160	0 ± 0 b	2.6 ± 1 b
4	He-O <sub>2</sub>	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

610

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

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

4	He-O <sub>2</sub>	15	55	320	0.5 ± 0.4 a	2.8 ± 0.7 a
4	He-O <sub>2</sub>	15	75	0	0 ± 0 b	1.3 ± 0.2 bc
4	He-O <sub>2</sub>	15	75	160	0 ± 0 b	3.8 ± 1.6 a
4	He-O <sub>2</sub>	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

616