Soil biochar amendment affects the diversity of nosZ transcripts: Implications for N₂O formation

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Microbial nitrogen transformation processes such as denitrification represent major sources of the potent greenhouse gas nitrous oxide (N₂O). Soil biochar amendment has been shown to significantly decrease N₂O emissions in various soils. However, the effect of biochar on the structure and function of microbial communities that actively perform nitrogen redox transformations has not been studied in detail yet. To analyse the community composition of actively denitrifying and N₂O-reducing microbial communities, we collected RNA samples at different time points from a soil microcosm experiment conducted under denitrifying conditions and performed Illumina amplicon sequencing targeting nirK, typical nosZ and atypical nosZ mRNA transcripts. Within 10 days, biochar significantly increased the diversity of nirK and typical nosZ transcripts and resulted in taxonomic shifts among the typical nosZ-expressing microbial community. Furthermore, biochar addition led to a significant increase in transcript production among microbial species that are specialized on direct N₂O reduction from the environment. Our results point towards a potential coupling of biochar-induced N₂O emission reduction and an increase in microbial N₂O reduction activity among specific groups of typical and atypical N₂O reducers. However, experiments with other soils and biochars will be required to verify the transferability of these findings to other soil-biochar systems.

Nitrous oxide (N₂O) is a strong greenhouse gas that has, based on its radiative capacity, an almost 300-fold greater global warming potential than CO₂. Since 1750 its atmospheric concentration increased from approximately (estimates from ice cores) 270 to 324 ppb in 2011. Microbial nitrogen transformation processes in soils, such as nitrification and denitrification, represent the world’s largest sources of atmospheric N₂O. N₂O emissions have been increasing mainly as a result of excessive application of nitrogen fertilization in agricultural systems. Due to the increasing demand for food and animal feed, N₂O emissions will likely further increase in the future.

Biochar, a carbonaceous solid manufactured for the deliberate purpose of applying it to soil, has recently been suggested as a promising N₂O mitigation tool. Biochar is produced by thermal decomposition of organic biomass under low oxygen conditions and has gained a lot of attention among agronomists and soil scientists because of its soil quality-enhancing properties and its recalcitrance to biodegradation. Biochar’s physico-chemical properties vary strongly with the type of organic material used as feedstock and the charring conditions during production. However, many biochars are produced by slow pyrolysis of woody plant material at temperatures between 400 and 700 °C and these biochars often share common physicochemical properties such as a high carbon content, a neutral to alkaline pH, a large surface area, and a highly aromatic carbon structure.

For wood-derived but also for other biochars, it has been shown that soil biochar amendment can considerably improve soil quality (e.g. by impacting soil pH, water holding capacity, and nutrient retention), while simultaneously sequestering carbon from the atmosphere. In addition, a recent meta-analysis which summarized the results of several laboratory and field studies demonstrated that the addition of wood-derived biochars to arable soils results in a significant decrease of N₂O emissions.

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In the field, highest N$_2$O emissions are frequently observed under denitrifying conditions (oxygen limitation, high nitrate and organic carbon concentrations), which occur for example after heavy rain fall on fertilized soil$^{15,16}$. Denitrification is defined as the stepwise enzymatic reduction of nitrate (NO$_3^-$) to dinitrogen (N$_2$), with nitrite (NO$_2^-$), nitric oxide (NO), and N$_2$O occurring as obligate intermediates. Denitrification is a common functional trait in many facultative and strict anaerobic chemoorganotrophic bacteria and has been also detected in some archaea and fungi$^{19,20}$. The enzymes catalysing the reactions of the different reduction steps in denitrification are encoded by the functional genes narG and napA (nitrate reductases), nirK and nirS (nitrite reductases), norB (nitric oxide reductase), and nosZ (nitrous oxide reductase)$^{19,21}$. Because some denitrifiers lack a functional nosZ gene and nitrous oxide reductases are highly oxygen- and pH-sensitive, the last step of denitrification (N$_2$O reduction to N$_2$) is often impaired$^{22-24}$. Since the microbial reduction of N$_2$O to N$_2$ via nosZ-encoded nitrous oxide reductases represents the only known sink of N$_2$O$^{1}$, this process directly controls the N$_2$O/(N$_2$O + N$_2$) emission ratio and thus the quantity of nitrogen that is released as N$_2$O. Until recently N$_2$O reduction was thought to be a functional trait exclusively associated with ‘classical’ denitrifiers. These microorganisms usually contain, in addition to the nosZ gene, also the functional genes that encode the enzymes catalysing the other steps towards complete denitrification (narG/napA, nirK/nirS, and norB). Most of the ‘classical’ denitrifiers are affiliated with the phylum Proteobacteria$^{15,25}$. However, recent studies revealed that the ability to perform N$_2$O reduction is not restricted to these functional groups of microorganisms, but instead is also very common among microbes that belong to other phyla such as Actinobacteria, Bacteroidetes, Firmicutes and Chloroflexi$^{26-27}$. The majority of these microorganisms carry a fully functional but atypical form of the nosZ gene and about half of them do not contain a nitrite reductase gene$^{27-29}$.

Although it has been confirmed in many studies that soil biochar amendment holds the potential to act as promising N$_2$O mitigation strategy the underlying processes causing a decrease in N$_2$O emissions are still not fully understood. Because N$_2$O is formed during microbial nitrogen transformation reactions such as nitrification and especially denitrification, several studies aimed to link N$_2$O emission suppression to changes in the structure and activity of microbial communities involved in nitrogen cycling$^{30-35}$. It is known from 16S rRNA gene-based molecular fingerprinting and sequencing studies that the addition of biochar can significantly affect the microbial diversity in soil. Several studies reported a biochar-induced increase in relative abundance of microbial taxa that are known to be involved in nitrogen transformation processes such as nitrogen fixation, nitrification, and denitrification$^{36,35,36}$. Other studies revealed that soil biochar amendment can significantly increase typical and atypical nosZ gene and transcript copies suggesting a higher abundance and activity of N$_2$O reducers in response to soil biochar amendment$^{31,32,34,37,38}$. Furthermore, it has been reported in a recent sequencing study, targeting typical and atypical nosZ genes, that biochar incorporation into soil can cause taxonomic shifts among microbes that carry the genetic potential for N$_2$O reduct$^{39}$. However, the impact of soil biochar amendment on the diversity and taxonomic composition of active denitrifiers and N$_2$O reducing microbial taxa has not yet been systematically investigated on the transcript level.

In order to investigate biochar-induced changes in diversity and taxonomic composition of active denitrifiers and N$_2$O reducers, we amplified and sequenced nirK, typical nosZ, and atypical nosZ transcripts in samples collected at different time points of sampling (0, 4, and 10 days) from a soil microcosm experiment that has been performed under denitrifying conditions$^{35}$.

**Material and Methods**

The RNA extracts analysed in this study were obtained from soil samples collected in a microcosm experiment previously described by Harter, et al.$^{34}$. In the following we briefly describe the soil and biochar used for this soil microcosm experiment and its experimental design. For a detailed description of the experiment please refer to Harter, et al.$^{34}$.

**Soil sampling and biochar production.** The soil for the soil microcosm experiment was collected from the top 10 cm of an Anthrosol (World Reference Base for Soil Resources, 2014) located at an urban gardening site of the University of Tuebingen, Germany (48° 32′ N, 9° 4′ E). The soil is classified as sandy clay loam (USDA) with a particle size distribution of 49.8% sand, 25.8% silt and 24.4% clay. Biochar was produced by Swiss Biochar Sàrl (Belmont-sur-Lausanne, Switzerland) from green waste by slow pyrolysis (620°C) according to the standards of the European Biochar Certificate (EBC) (http://www.european-biochar.org). Important physical and chemical properties of soil and biochar are summarized in Table 1. Detailed descriptions of the methods used to determine the physical and chemical properties of soil and biochar can be found in Harter, et al.$^{34}$.

**Soil microcosm experiment.** Prior to microcosm setup the field moist (water content: 29% w/w) soil and the biochar were passed through a 2 mm sieve and thoroughly homogenized. Soil microcosms were set up in 250 ml glass beakers. Microcosms contained either the field moist equivalent of 100 g dry soil (control microcosms) or the field moist equivalent of 95 g dry soil and 5 g dry biochar resulting in a final biochar content of 5% (biochar microcosms). Control and biochar microcosms were set up in triplicates. After preparation, soil microcosms were homogenized and compacted (10 repetitions of dropping a 125 g hammer from a height of 20 cm onto the soil surface). In order to simulate conventional fertilization, all soil microcosms were fertilized with a NH$_4$NO$_3$ solution at a rate of 332 mg N kg$^{-1}$ (equivalent to 100 kg N ha$^{-1}$ estimated based on the soil surface area in the microcosms). The amount of water added with the NH$_4$NO$_3$ solution was calculated to adjust the water filled pore space (WFPS) to 90%. While the experiment was running the soil microcosms were covered with a perforated aluminium foil to allow gas exchange with the ambient atmosphere and to decrease evaporation. Soil microcosms were incubated at a constant temperature of 20°C and the WFPS was held constant by periodically replenishing the evaporated water. Soil microcosms were incubated for 10 days with sampling taking place after 0, 1, 2, 4, 7 and 10 days of incubation. To allow destructive soil sampling of a set of 6 microcosms (3 control and 3...
biochar microcosms) at each time point of sampling, we set up 36 soil microcosms in total. During soil sampling the entire soil of the soil microcosm was transferred into a sterile container, homogenized using a spatula and aliquoted for parallel RNA extraction. All samples for RNA extraction contained the equivalent of 2 g dry soil and were directly frozen at −80 °C. For the present sequencing study only RNA extracts obtained from soil samples collected at day 0, 4, and 10 were analysed.

**RNA extraction and reverse transcription.** Total RNA was extracted using the RNA PowerSoil Total RNA Isolation Kit according to the manufacturer protocol (MO BIO Laboratories, Carlsbad, CA, USA). RNA concentration and quality of the resulting extracts were determined spectrophotometrically (NanoDrop 1000, Thermo Scientific, Waltham, MA, USA), fluorometrically (Qubit 2.0 Fluorometer, Life Technologies, Carlsbad, CA, USA), and by gel electrophoresis (Experion Automated Electrophoresis Station, Bio-Rad Laboratories, Hercules, CA, USA). Residual DNA in RNA extracts was digested using the Ambion TURBO DNA-free Kit (Life Technologies, Carlsbad, CA, USA) according to the instructions given by the manufacturer. Successful removal of DNA was confirmed by PCR using primers 27F (5′-AGAGTTTGATCMTGGCTCAG-3′) and PC5 (5′-TACCTTGTTACGACTT-3′) with the following conditions: hot start at 70 °C, 5 min at 95 °C, 35 cycles with 1 min at 95 °C, 1 min at 44 °C and 3 min at 72 °C followed by a final elongation step of 10 min at 72 °C. If the PCR revealed no PCR products, RNA extracts were used for cDNA synthesis. Reverse transcription of pure RNA extracts was performed with the SuperScript III Reverse Transcriptase using random primers according to the manufacturer’s protocol (Life Technologies, Carlsbad, CA, USA). cDNA samples were quality checked and quantified using agarose gel electrophoresis and Nanodrop (NanoDrop 1000, Thermo Scientific, Waltham, MA, USA).

**Illumina amplicon sequencing.** In order to sequence nirK, typical nosZ and atypical nosZ mRNA transcripts we amplified the corresponding fragments from the cDNA samples from the control and biochar microcosms at day 0, 4 and 10 (n = 18) using PCR. PCRs were performed using target-specific primers fused with overhang adapter sequences at the 5′-end to allow the addition of index sequences and sequencing adapters in a second
index PCR. Transcripts of nirK genes were amplified using primer F1aCu (5′-ATCATGGTSCGCGCG-3′) and R3Cu (5′-GCCTCGATCAGRTTGTGGTT-3′)41. For typical nosZ gene transcripts primer nosZ2F (5′-CGCRACCGCAAAAGTMSMTTGT-3′) and nosZ2R (5′-CAKRGCAKGCRTGCGAA-3′)37 were used. Transcripts of atypical nosZ genes were amplified with the primers nosZ-II-F (5′-CTNGGNCNNVTKCAYAC-3′) and nosZ-II-R (5′-GCNGARARAAANTCGBRC-3′) developed by Jones, et al.46. PCRs were performed with the FastStart High Fidelity PCR system (Roche Diagnostics, Rotkreuz, Switzerland) using the thermal protocols and reaction mixtures described in Table S1 in the supplementary information. The produced amplicons were purified using AMPure XP beads (Beckman Coulter, Brea, CA, USA) at a ratio of 1:1 (v/v). Quality and quantity of the purified amplicons were determined using agarose gel electrophoresis and Nanodrop (NanoDrop 1000, Thermo Scientific, Waltham, MA, USA). Subsequent library preparation steps and sequencing were performed by IMGM Laboratories GmbH ( Martinsried, Germany) according to the amplicon sequencing guidelines given by Illumina (San Diego, CA, USA). Sequencing was performed on an Illumina MiSeq sequencing system (Illumina, San Diego, CA, USA) using the 2 × 300 bp MiSeq Reagent Kit v3 (600 cycle) (Illumina, San Diego, CA, USA). The MiSeq Reporter Software v2.5.1.3 (Illumina, San Diego, CA, USA) was used for signal processing, de-multiplexing and trimming of adapter sequences.

Sequence analysis. Quality control of raw nirK, typical nosZ and atypical nosZ read pairs was performed using Cutadapt v1.943 and USEARCH v8.1.181244,45. At first target-specific primers were detected and trimmed using Cutadapt. Afterwards, paired-end nirK and typical nosZ reads were merged using the fastq_mergepairs algorithm implemented in USEARCH. Due to the large size of atypical nosZ amplicons (>600 bp) paired-end reads could not be merged. Instead, the forward read was trimmed to 200 bp to ensure the removal of bases with low phred scores and to obtain sequences of equal length. Trimming was performed with fastx_truncate using USEARCH. Merged (nirK and typical nosZ) and trimmed (atypical nosZ) sequences were quality filtered using the USEARCH fastq_filter function with a maximum error rate threshold of 1%. Chimeric sequences were de novo detected and removed using the uchime_denoovo algorithm implemented in USEARCH.

All quality filtered non-chimeric nirK, typical nosZ and atypical nosZ sequences were mapped against the NCBI Reference Sequence (RefSeq) protein database (release 66). Mapping was performed using DIAMOND (v0.7.9.58)47 in blastx mode with a minimum protein sequence identity cut-off of 70% and an e-value cut-off of 10−10. The top 50 database hits of each sequence were used for further analysis with MEGAN6 Ultimate Edition48,49. For taxonomic placement the Lowest Common Ancestor (LCA) algorithm of MEGAN was applied50. The LCA analysis parameters ‘Top Percent’ and ‘Min Support’ were set to 0.5% (all hits within the top 0.5% of the best bit score are considered) and 10 (taxa need to obtain a minimum of 10 reads to be considered), respectively. Shannon-Weaver and Simpson-Reciprocal diversity indices were computed on the species level in MEGAN.

Absolute sequence counts and relative sequence abundances of all assigned taxa were exported from MEGAN for further analyses. Relative sequence abundances were calculated based on all sequences that matched database entries in the RefSeq protein database.

Typical and atypical nosZ transcript copy numbers of individual species at day 4 were calculated by multiplying absolute transcript copy numbers determined by qPCR in a previous study conducted by Harter, et al.34 with the corresponding relative sequence abundances of individual nosZ-expressing species obtained in this study. This was possible because sequencing and qPCRs were performed on the same RNA samples with the same sets of primers in the present and the previous study by Harter, et al.34. Individual species with an average relative sequence abundance below 1% were not considered.

Figures 1, 2 and 3 were generated with GraPhAAn11 using the LCA-based taxonomy and the corresponding relative sequence abundance information from MEGAN. According to the basic principle of the LCA algorithm, sequences that are conserved among different species (e.g. as consequence of horizontal gene transfer) will only be assigned to taxa of higher rank48. Nonetheless, it is very difficult to directly prove that a given nirK or nosZ gene (typical and atypical) appears in a specific microbial taxon. Thus, whenever we mention specific taxa in the results and discussion we refer to a taxonomic group of microorganisms that contains a nirK or nosZ gene closely related to the nirK or nosZ gene of the respective taxa.

Statistical analysis. The effects of biochar addition on diversity indices (Tables 2 and 3) and the relative sequence abundance of each assigned microbial species (Figs 1,2,3) were statistically evaluated at each time point of sampling (0, 4, and 10 days) using t-tests. In addition, we used t-tests to determine significant biochar effects on the absolute nosZ transcript copy numbers of individual species at day 4 (Fig. 4). Differences in diversity indices among the average (all control and biochar microcosms at all time points of sampling, n = 18) nirK, typical nosZ and atypical nosZ-expressing microbial communities (Tables 2 and 3) were determined using one-way ANOVA with Tukey’s HSD post-hoc test. T-tests and one-way ANOVAs were carried out in SAS (SAS 9.2, SAS Institute, Cary, NC, USA) using PROC TTEST and PROC GLM, respectively. Dissimilarities among nirK, typical nosZ and atypical nosZ-expressing microbial communities in control and biochar microcosms and over time were statistically evaluated using permutational multivariate analysis of variance (PERMANOVA) with biochar addition and time as main factors and 105 permutations (Table 4). PERMANOVAs were performed based on Bray-Curtis dissimilarity matrices using the adonis function implemented in the R package vegan53.

Data availability. Illumina sequencing reads have been deposited in the ENA Sequence Read Archive (SRA) under accession number PRJEB14348.
Results

Sequencing statistics. Transcript sequencing resulted in 3,404,618 raw pairs of reads in total with \( \text{nirK} \), typical \( \text{nosZ} \), and atypical \( \text{nosZ} \) accounting for 733,518 (40,751 ± 13,287 per sample), 2,064,131 (114,674 ± 35,377 per sample), and 606,969 (33,721 ± 9,523 per sample) raw read pairs, respectively. Quality processing resulted in 375,665 merged high quality \( \text{nirK} \) sequences of which 374,783 (20,821 ± 8,672 per sample) matched entries in the RefSeq database (protein sequence identity cut-off: 70%, e-value cut-off: \( 10^{-10} \)). Merged high quality \( \text{nirK} \) sequences had an average sequence length of 436 bp.

For typical \( \text{nosZ} \), 1,456,124 merged high quality sequences remained after quality processing and 1,420,025 (78,890 ± 24,006 per sample) matched database entries (protein sequence identity cut-off: 70%, e-value cut-off: \( 10^{-10} \)). The average sequence length was 223 bp.

Quality processing of atypical \( \text{nosZ} \) sequences led to 288,167 high quality sequences with a sequence length of 200 bp. 166,786 sequences (9266 ± 2243 per sample) matched database entries (protein sequence identity cut-off: 70%, e-value cut-off: \( 10^{-10} \)).

Diversity of the microbial communities expressing \( \text{nirK} \), typical \( \text{nosZ} \), and atypical \( \text{nosZ} \). In order to investigate the impact of soil biochar amendment on the diversity of expressed \( \text{nirK} \) and \( \text{nosZ} \) genes, we computed Shannon–Weaver and Simpson–Reciprocal diversity indices for all samples. Shannon–Weaver indices ranged from 0.99 to 3.49 and were generally highest for atypical \( \text{nosZ} \) and lowest for \( \text{nirK} \) (Table 2). On average (all control and biochar microcosms at all sampling time points, \( n = 18 \)), the \( \text{nirK} \)-expressing community had a value of 1.56. In comparison with \( \text{nirK} \), typical and atypical \( \text{nosZ} \)-expressing communities showed significantly higher values of 2.74 (\( p < 0.001 \)) and 3.20 (\( p < 0.001 \)), respectively. Atypical
nosZ had a significantly higher average Shannon-Weaver index than typical nosZ ($p < 0.001$). Biochar addition altered the Shannon-Weaver index at all time points of sampling. For nirK and typical nosZ a biochar-induced increase in diversity was observed (Table 2). Atypical nosZ diversity slightly decreased in the presence of biochar. While this slight decrease was not statistically significant at any time point of sampling, the values for nirK at day 10 ($p = 0.002$) and typical nosZ at day 4 ($p = 0.014$) were significantly higher in biochar compared to control microcosms.

The Simpson-Reciprocal diversity index showed a very similar pattern and ranged from 1.53 to 6.97 (Table 3). With an average value of 5.21, the highest diversity was determined for the atypical nosZ-expressing microbial community. The average value for typical nosZ was slightly, but not significantly, lower (4.72). However, with a value of 2.27 the nirK-expressing community had a significantly lower average diversity compared to the microbial communities producing typical ($p < 0.001$) and atypical ($p < 0.001$) nosZ transcript. Biochar addition significantly increased the Simpson-Reciprocal diversity index of typical nosZ transcripts at day 4 ($p = 0.016$) and 10 ($p = 0.049$) (Table 3). In addition, significantly higher values in biochar compared to control microcosms were also determined for nirK at day 10 ($p = 0.003$). For atypical nosZ soil biochar amendment resulted in significantly lower values at day 4 ($p = 0.012$).

In order to determine if biochar addition or time caused significant dissimilarities among nirK, typical nosZ and atypical nosZ-expressing microbial communities, we performed two-way PERMANOVA. As shown in Table 4, PERMANOVAs revealed a significant time effect for nirK ($p < 0.001$), typical nosZ ($p < 0.001$) and atypical nosZ-expressing microbial communities ($p < 0.001$) and a significant biochar effect for the typical nosZ-expressing microbial community ($p = 0.017$).

Figure 2. Circular taxonomic tree of typical nosZ-expressing taxa. Microbial taxa are indicated by circles. Different classes are shown with specific branch colours. Large, filled, circles depict taxa with average relative sequence abundances above 1%. Light colour shaded areas indicate orders with a relative sequence abundance above 1%. Dark colour shaded boxes with numbers (1–8) represent species with a relative sequence abundance above 1%. External rings show the relative sequence abundance of the corresponding species. Ring 1 shows the average relative sequence abundance in percent (all microcosms at all time points of sampling, $n = 18$). Rings 2–7 show the relative sequence abundances in control and biochar microcosms at day 0, 4, and 10 as circular heatmaps ($n = 3$).
Figure 3. Circular taxonomic tree of the atypical nosZ-expressing taxa. Microbial taxa are indicated by circles. Different phyla are shown with specific branch colours. Large, filled circles depict taxa with average relative sequence abundances above 1%. Light colour shaded areas indicate orders with a relative sequence abundance above 1%. Dark colour shaded boxes with numbers (1–15) represent species with a relative sequence abundance above 1%. External rings show the relative sequence abundance of the corresponding species. Ring 1 shows the average relative sequence abundance in percent (all microcosms at all time points of sampling, n = 18). Rings 2–7 show the relative sequence abundances in control and biochar microcosms at day 0, 4, and 10 as circular heatmaps (n = 3).

Table 2. Shannon-Weaver diversity index for nirK, typical nosZ, and atypical nosZ, in control and biochar microcosms after 0, 4, and 10 days. “average” shows average values of all microcosms at all time points of sampling. Values represent means ± standard deviation (individual samples: n = 3, average: n = 18). Significant differences between individual control and biochar samples are shown in the column “sig.” (ns = not significant, *p < 0.05, **p < 0.01). Significant differences (p < 0.05) between the average values of nirK, typical nosZ and atypical nosZ are indicated by superscripted letters next to the corresponding values.
Classification of nirK transcripts. Regardless of the time point of sampling and the type of microcosm (biochar or control) all microbial taxa that produced copper nitrite reductase gene (nirK) transcripts were affiliated with the phylum Proteobacteria (Fig. 1). 99.8% of all nirK sequences belonged to the order Rhizobiales. 74.4% could be classified down to the species level with Mesorhizobium ciceri (47.0%), Rhodopseudomonas palustris (13.3%), Mesorhizobium opportunistum (6.12%), Mesorhizobium australicum (4.02%), and Chelativornas

Table 3. Simpson-Reciprocal diversity index for nirK, typical nosZ, and atypical nosZ, in control and biochar microcosms after 0, 4, and 10 days. “average” shows average values of all microcosms at all time points of sampling. Values represent means ± standard deviation (individual samples: n = 3, average: n = 18). Significant differences between individual control and biochar samples are shown in the column “sig.” (ns = not significant, *p < 0.05, **p < 0.01). Significant differences (p < 0.05) between the average values of nirK, typical nosZ and atypical nosZ are indicated by superscripted letters next to the corresponding values.

Table 4. Results from Bray-Curtis dissimilarity based two-way PERMANOVAs. The table shows pseudo F-statistics (F), explained variances (R²), and the level of significance (p) for the two main effects “biochar” and “time” and their interaction “biochar*time”. Significant effects indicated by p-values below 0.05 are shown in bold font.

Figure 4. Calculated absolute typical (A) and atypical (B) nosZ transcript copy numbers originating from specific species in control (white bars) and biochar (black bars) microcosms at day 4. The figure shows only species where biochar addition had a statistically significant effect on transcript production (p < 0.05). Bars and error indicators represent means and standard errors (n = 3), respectively.
sp. BNC1 (2.05%) showing a relative sequence abundance above 1% (Fig. 1). A complete list with the relative sequence abundances of all nirK-expressing microbial species identified in this study can be found in Table S2 in the supplementary information.

The taxonomic composition and distribution of nirK transcripts slightly differed in control and biochar microcosms at all time points of sampling (0, 4, and 10 days) (Fig. 1). However, biochar related changes in relative sequence abundance of specific species did not follow a clear trend. Accordingly, significantly higher and lower relative sequence abundances in biochar compared to control microcosms were only detected for Mesorhizobium australicum at day 0 (control: 3.0%, biochar: 6.3%, p = 0.048) and Mesorhizobium ciceri at day 10 (control: 44.0%, biochar: 33.6%, p = 0.011), respectively (Fig. 1).

Classification of typical nosZ transcripts. Similar to nirK, all nosZ transcripts were affiliated with the phylum Proteobacteria (Fig. 2). With a relative sequence abundance of 77.1%, most sequences belonged to the order Rhizobiales. Rhodocyclales, Burkholderiales, and Rhodobacterales had relative sequence abundances of 7.8%, 6.9%, and 1.2%, respectively. In total, 90.2% of all typical nosZ sequences could be classified down to the species level. The most abundant species were: Oligotropha carboxidovorans (31.8%), Bradyrhizobium diazoefficiens (15.1%), Hyphomicrobiurn nitrativorans (12.4%), Ensifer meliloti (10.6%), Azorarcus sp. KH32C (6.57%), Ensifer fredii (4.68%), Polymorphum gilvum (2.45%), and Azorarcus sp. BH72 (1.24%). All of these species had a relative sequence abundance above 1% (Fig. 2). A complete list with the relative sequence abundances of all typical nosZ-expressing microbial species identified in this study can be found in Table S3 in the supplementary information.

In agreement with the PERMANOVA results, the taxonomic composition and distribution of typical nosZ transcripts was significantly affected by biochar addition (Fig. 2). Although the contribution of certain species to total typical nosZ transcripts was different in control and biochar microcosms throughout all time points of sampling (0, 4 and 10 days), statistically significant differences in relative sequence abundance were only determined after 4 and 10 days of incubation. At day 4, soil biochar amendment significantly increased the relative sequence abundance of Ensifer meliloti (control: 9.3%, biochar: 12.2%, p = 0.019) and Azorarcus sp. BH72 (control: 0.1%, biochar: 2.9%, p = 0.010). The relative sequence abundances of Hyphomicrobiurn nitrativorans (control: 20.6%, biochar: 16.1%, p = 0.002) and Azorarcus sp. KH32C (control: 5.4%, biochar: 4.1%, p = 0.048) were significantly lower in the presence of biochar (Fig. 2). After 10 days of incubation, significantly higher relative sequence abundances in biochar compared to control microcosms were determined for Bradyrhizobium diazoefficiens (control: 16.9%, biochar: 30.3%, p = 0.025), Ensifer fredii (control: 2.6%, biochar: 3.3%, p = 0.021), and Azorarcus sp. BH72 (control: 0.4%, biochar: 2.7%, p = 0.021). Significantly lower values were found for Oligotropha carboxidovorans (control: 36.8%, biochar: 22.2%, p = 0.006) (Fig. 2).

Classification of atypical nosZ transcripts. Atypical nosZ-expressing microorganisms belonged to a broad diversity of different phyla (Fig. 3). About 36.9% of all sequences were affiliated with the phylum Chlorobii. Other important phyla were: Proteobacteria (19.5%), Bacteroidetes (15.1%), Chloroflexi (14.2%), and Firmicutes (13.8%), respectively. Although differences in relative sequence abundance between control and biochar microcosms, atypical nosZ transcripts were affiliated with the phylum Proteobacteria (Fig. 2). With a relative sequence abundance of 77.1%, most sequences belonged to the order Rhizobiales. Rhodocyclales, Burkholderiales, and Rhodobacterales had relative sequence abundances of 7.8%, 6.9%, and 1.2%, respectively. In total, 90.2% of all typical nosZ sequences could be classified down to the species level. The most abundant species were: Oligotropha carboxidovorans (31.8%), Bradyrhizobium diazoefficiens (15.1%), Hyphomicrobiurn nitrativorans (12.4%), Ensifer meliloti (10.6%), Azorarcus sp. KH32C (6.57%), Ensifer fredii (4.68%), Polymorphum gilvum (2.45%), and Azorarcus sp. BH72 (1.24%). All of these species had a relative sequence abundance above 1% (Fig. 2). A complete list with the relative sequence abundances of all typical nosZ-expressing microbial species identified in this study can be found in Table S3 in the supplementary information.

In agreement with the PERMANOVA results, the taxonomic composition and distribution of typical nosZ transcripts was significantly affected by biochar addition (Fig. 2). Although the contribution of certain species to total typical nosZ transcripts was different in control and biochar microcosms throughout all time points of sampling (0, 4 and 10 days), statistically significant differences in relative sequence abundance were only determined after 4 and 10 days of incubation. At day 4, soil biochar amendment significantly increased the relative sequence abundance of Ensifer meliloti (control: 9.3%, biochar: 12.2%, p = 0.019) and Azorarcus sp. BH72 (control: 0.1%, biochar: 2.9%, p = 0.010). The relative sequence abundances of Hyphomicrobiurn nitrativorans (control: 20.6%, biochar: 16.1%, p = 0.002) and Azorarcus sp. KH32C (control: 5.4%, biochar: 4.1%, p = 0.048) were significantly lower in the presence of biochar (Fig. 2). After 10 days of incubation, significantly higher relative sequence abundances in biochar compared to control microcosms were determined for Bradyrhizobium diazoefficiens (control: 16.9%, biochar: 30.3%, p = 0.025), Ensifer fredii (control: 2.6%, biochar: 3.3%, p = 0.021), and Azorarcus sp. BH72 (control: 0.4%, biochar: 2.7%, p = 0.021). Significantly lower values were found for Oligotropha carboxidovorans (control: 36.8%, biochar: 22.2%, p = 0.006) (Fig. 2).

Typical and atypical nosZ transcript copies of individual species at day 4. At day 4, soil biochar amendment significantly altered the calculated abundance of typical nosZ transcripts in 5 relevant (relative sequence abundance >1%) species. The calculated number of typical nosZ transcript copies of the following species was significantly increased in the presence of biochar (Fig. 4A): Oligotropha carboxidovorans (p = 0.042), Bradyrhizobium diazoefficiens (p = 0.028), Ensifer meliloti (p = 0.028), Ensifer fredii (p = 0.021), and Azorarcus sp. BH72 (p = 0.016). Transcript copy numbers originating from other species were not significantly affected by biochar addition.

Similar to typical nosZ, significant biochar-induced changes in the production of atypical nosZ transcripts at day 4 were determined for only 5 (relative sequence abundance >1%) species (Fig. 4B). Calculated atypical nosZ transcript copy numbers of Melioribacter roseus (p = 0.005), Pedobacter saltans (p = 0.024), B. baltica
(p = 0.041), Runella slithyformis (p = 0.029), and Flavobacteriaceae bacterium 3519-10 (p = 0.019) were significantly higher in biochar compared to control microcosms. Absolute transcript production of other atypical nosZ-expressing species was not significantly altered by soil biochar amendment.

Discussion

Highest N₂O emission rates usually occur after heavy rain fall when high amounts of organic carbon and mineral nitrogen are mobilized and oxygen availability decreases due to high water filled pore spaces (WFPS) [17, 18]. These conditions promote the growth and activity of microbial strains capable of performing anaerobic nitrogen transformation processes such as denitrification [19, 20]. In order to simulate such conditions, we set up a soil microcosm experiment at a constant WFPS of 90% to which we added a NH₄NO₃ solution at concentrations that reflect common agricultural application rates. Although the amount of N₂O emitted under these conditions can account for large fractions of total annual emissions, a high WFPS is usually occurring only a few days after rain events in most soils [17, 18]. Accordingly, our experiment was designed to mimic the frequently observed high N₂O emissions occurring in response to soil water saturation over a short time period (10 days of incubation, 6 time points of sampling).

Harter, et al. [24] described and discussed the impact of biochar addition on nitrogen transformation rates, absolute gene and transcript copy numbers of denitrification marker genes, and the formation and release of N₂O and N₂. According to Harter, et al. [24], biochar addition significantly decreased N₂O emissions and increased transcript copy numbers of specific functional denitrification genes. While transcripts of the denitrification genes napA, narG and nirS did not show significant biochar effects or were subject to a significant interaction effect (biochar*time), nirK, typical nosZ and atypical nosZ transcript copy numbers were significantly increased by biochar addition. Thus, the current Illumina amplicon sequencing study described here focuses on biochar-induced changes in diversity and relative abundance among the microbial communities actively expressing nirK, typical nosZ and atypical nosZ transcripts.

According to both diversity indices (Shannon-Weaver and Simpson-Reciprocal) the nirK-expressing microbial community had a significantly lower diversity than the microbial communities producing typical nosZ and atypical nosZ transcripts. A lower diversity of the nirK-harbouring community compared to the microbial community carrying typical nosZ has also been reported by Palmer and Horn [25]. The microbial community that actively expressed atypical nosZ genes was significantly more diverse (Shannon-Weaver diversity index) than the community expressing typical nosZ genes. These findings confirm the results of Jones, et al. [26] who determined an up to 3-fold higher Faith's phylogenetic diversity (PD) for atypical nosZ compared to typical nosZ genes. In accordance with our results, Jones and colleagues also demonstrated a significant relationship between the relative abundance and the phylogenetic diversity of the nosZ community and the ability of the soil microbial community to reduce N₂O. Using network and co-occurrence analyses, Jones, et al. [26] found that functional groups that were identified as significant indicators of reduced soil N₂O emissions were dominated by atypical nosZ communities. Notably, Jones and colleagues found specific atypical nosZ groups in high abundance in soils for which the lowest N₂O emissions were observed. These atypical nosZ lineages lacked either nir or genes but seemed to be critical for the N₂O sink capacity of the respective soil. Interestingly, the observation made by Jones, et al. [26] with respect to nirK and nosZ gene abundance and diversity are very similar to what we report here for a biochar-amended soil and nirK and nosZ transcript abundance and diversity under conditions of reduced N₂O emissions.

The taxonomic compositions of the microbial communities expressing nirK, typical nosZ, and atypical nosZ described here, were in good agreement with other sequencing studies performed on soil. In accordance with the findings of Bremer, et al. [27] and Henry, et al. [28], the nirK-expressing community was largely dominated by the order Rhizobiales and the genus Mesorhizobium. The typical nosZ-expressing soil community was dominated by Alpha- and Betaproteobacteria and species affiliated with the genera Oligotropha, Bradyrhizobium, Hyphomicrobiurn, Ensifer, and Azorarcus. The importance of these genera within the typical nosZ gene-carrying community, as well as the dominance of microbial taxa that belong to the classes Alpha- and Betaproteobacteria has been reported in several previous soil studies [28, 29, 33, 35, 36]. While all nirK and typical nosZ transcripts were produced by microbial taxa affiliated with the phylum Proteobacteria, the atypical nosZ-expressing community comprised a board variety of different microbial species affiliated with several phyla including Bacteroidetes, Chlororobi, Chloroflexi, and Firmicutes. These results support the findings of Jones, et al. [26] and Sanford, et al. [27] who showed that N₂O reduction is a widespread functional trait among soil microbes. Furthermore, the determined species reflected the taxonomic composition of atypical nosZ-carrying microbial species identified in similar sequencing studies and a recent soil metagenome analysis [28].

 Soil biochar amendment significantly increased the diversity indices computed for nirK (day 10) and typical nosZ (day 4 and 10). Higher diversity indices in biochar-amended compared to biochar-free soils have been reported frequently in studies targeting 16S rRNA genes [31, 35, 37]. The reason for the biochar-induced increase in microbial diversity is mostly unknown. However, as mentioned in other studies, it seems likely that organic compounds attached to the biochar particles, the high pH of biochar-amended soils, or other biochar properties (surface area, pore space) promoted the formation of specific niches, which supported the growth and activity of a diverse range of taxa [33, 35, 57, 60].

PERMANOVA analyses indicated that the composition of nirK, typical nosZ and atypical nosZ-expressing microbial communities were significantly altered over the whole duration of the experiment (p < 0.001) and that the composition of typical nosZ transcripts was significantly affected by biochar addition (p < 0.017). Among the typical nosZ-expressing community biochar addition resulted in a significant increase in relative abundance of Ensifer meliloti (day 4), Azorarcus sp. BH72 (day 4 and 10), Bradyrhizobium diazoefficiens (day 10), and Ensifer fredii (day 10). Interestingly all of these microbial species are primarily known for their ability to perform nitrogen fixation and usually form a symbiotic relationship with legumes or grasses [61–64]. Except for Azorarcus sp. BH72, which does not contain nirK or nir genes, all species carry the full set of denitrification genes (napA/narG,
In order to further our understanding of the effects biochar has on microbial N$_2$O reduction, we calculated the absolute numbers of typical and atypical transcript copies produced by individual species based on the qPCR data from Harter, et al.\textsuperscript{34} and the relative transcript abundances determined in the present sequencing study. Species-specific transcript expression was specifically determined at day 4, because biochar most significantly affected total transcript copy numbers and N$_2$O emission rates at this day\textsuperscript{34}.

Biochar addition to soil significantly increased the calculated production of nosZ transcripts in 5 typical and 5 atypical nosZ-carrying species. Typical nosZ transcript production was enhanced in Oligotropha carboxidovorans, Bradyrhizobium diaezaeuliens, Ensifer mellitii, Ensifer fredii, and Azoarcus sp. BH72. An increased transcription of atypical nosZ genes in biochar microcosms was determined for Melioribacter roseus, Pedobacter saltans, Belliella balitica, Flavobacteriaceae bacterium 3519-10, and Runella slithyformis. The calculated production of typical and atypical nosZ transcripts in all other relevant microbial species we identified based on the sequencing of typical and atypical nosZ transcripts was not significantly affected by biochar addition. Recent studies revealed that well-studied strains of some of these species (Ensifer mellitii 1021 and Bradyrhizobium diaezaeuliens USDA 110) are able to grow with externally supplied N$_2$O as sole electron acceptor\textsuperscript{66, 67}. Thus, these strains take up N$_2$O from the environment and further reduce it to N$_2$. This is in contrast to other denitrifying strains such as Pseudomonas aeruginosa PAO1 which cannot grow on exogenous N$_2$O as sole electron acceptor\textsuperscript{79}. In addition to these species that are presumably able to physiologically specialize on N$_2$O reduction many other species that had a significantly higher nosZ transcript production in biochar microcosms such as Azoarcus sp. BH72, Flavobacteriaceae bacterium 3519-10, Belliella balitica, Melioribacter roseus, Pedobacter saltans, and Runella slithyformis lack at least one of the functional genes encoding nitrate, nitrite, and nitric oxide reductases\textsuperscript{79}. Thus these species are forced to perform a truncated version of the denitrification pathway by taking up denitrification intermediates (e.g. N$_2$O) from the environment that have been produced by other denitrifiers\textsuperscript{71}. Taken together these findings suggest that soil biochar amendment significantly increased nosZ transcript expression in microbial species capable of reducing exogenous N$_2$O from the environment.

Our results are in good agreement with the findings from a recent DNA-based sequencing study in which biochar addition significantly increased relative sequence abundances of microbial species capable to use exogenous N$_2$O as electron acceptor directly from the environment\textsuperscript{35}. The cause for the increase in nosZ expression in these specific species might be due to the higher quantities of entrapped N$_2$O in biochar microcosms\textsuperscript{35} that can serve as electron acceptor and thus lead to an increase in their transcriptional activity. Based on the high availability of N$_2$O, species specialized on N$_2$O reduction many other species that had a significantly higher nosZ transcript production in biochar microcosms might gain a competitive advantage over species performing the full denitrification pathway considering the thermodynamic advantages associated with directly reducing N$_2$O to N$_2$. However, although N$_2$O entrapment in biochar-amended soils might be an important stimulant for increased nosZ gene expression in exogenous N$_2$O reducers, we cannot exclude that other factors that are directly related to the physicochemical properties of biochar might have also contributed to the observed increase in nosZ transcription among the identified taxa. As reported in other studies several properties of biochar have the potential to significantly alter microbial communities by directly affecting microbial growth and activity of microorganisms (e.g. pH, biochar’s redox activity, nutrient sorption)\textsuperscript{65, 73–78}. In our study biochar’s high pH might have stimulated alkaliphilic Ensifer mellitii and Ensifer fredii strains\textsuperscript{61, 67}. Furthermore, it is possible that the high aromaticity of biochar or organic compounds that were attached to the biochar particles promoted Bradyrhizobium diaezaeuliens strains that are able to degrade aromatic substances\textsuperscript{78}.

Independent of the exact mechanism causing higher transcript production in these species, a higher N$_2$O reduction activity of microbes that are potentially capable to reduce exogenous N$_2$O from the environment will result in a decrease of the N$_2$O/(N$_2$O + N$_2$) ratio. Thus, together with the findings from our previous study\textsuperscript{34}, the decreased N$_2$O emissions from biochar-amended soils are most likely caused by N$_2$O entrapment resulting in retention and increased availability of N$_2$O for microbial reduction in the water-filled pore space. The lower N$_2$O/(N$_2$O + N$_2$) ratio is then caused by a higher microbial N$_2$O reduction activity of nosZ-containing microorganisms able to directly reduce exogenous N$_2$O from the environment.

In conclusion, this soil microcosm study showed that soil biochar amendment can lead to dynamic changes in the diversity of active denitrifier populations over time. Sequencing of nirK, typical nosZ and atypical nosZ transcripts revealed a significant biochar-induced increase in nirK and typical nosZ diversity and a higher relative sequence abundance of typical nosZ-expressing Rhizobia species. Furthermore, biochar addition led to a significantly higher calculated typical and atypical nosZ transcript expression among microbial species that are capable of directly reducing exogenous N$_2$O from the environment. Our results suggest that the increased activity of these specific microbial species might be responsible for the observed dynamic changes in N$_2$O/(N$_2$O + N$_2$) ratios in biochar vs control microcosms. We relate the observed increase in soil N$_2$O sink capacity to the higher activity of specific nosZ gene taxa in response to N$_2$O entrapment in the water-filled pore spaces of the biochar-amended soil. These findings further improve our understanding of the mechanisms responsible for biochar-induced N$_2$O emission mitigation. However, it has to be taken into account that species-specific transcript expression was calculated based on qPCR and sequencing data collected from a short-term experiment performed in plant-free, fertilized soil microcosms with a high water content. Hence, generalizations and extrapolation to field scales and extended time periods should be done cautiously until the findings from the present study have been evaluated using different biochars and soils at spatial and temporal scales that more closely translate to field conditions.
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Author Contributions
The study was designed by J.H., S.B., A.K. All microcosm experiments and sample preparation steps were done by J.H. Data analysis was done by M.E.H., D.H.H., and J.H. Table and figure preparation was performed by J.H. The manuscript was written by J.H. and S.B. All authors discussed the results and helped to improve the manuscript.

Additional Information
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