Abundance, Activity and Community Structure of Denitrifiers in Drainage Ditches in Relation to Sediment Characteristics, Vegetation and Land-Use

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ABSTRACT

Drainage ditches are ubiquitous yet understudied features of the agricultural landscape. Nitrogen pollution disrupts the nutrient balance of drainage ditch ecosystems, as well as the waterbodies in which they drain. Denitrification can help ameliorate the impact of N-fertilization by converting reactive nitrogen into dinitrogen gas. However, factors affecting denitrification in drainage ditches are still poorly understood. In this study, we tested how within-ditch and regional environmental conditions affect denitrifier activity, abundance, and community structure, to understand controls on denitrification at multiple scales. To this end, we quantified in situ denitrification rates and denitrifier abundance in 13 drainage ditches characterized by different types of sediment, vegetation and land-use. We determined how denitrification rates relate to denitrifier abundance and community structure, using the presence of nirS, nirK and nosZ genes as a proxy. Denitrification rates varied widely between the ditches, ranging from 0.006 to 24 mmol N m−2 h−1. Ditches covered by duckweed, which contained high nitrate concentrations and had fine, sandy sediments, were denitrification hotspots. We found highest rates in ditches next to arable land, followed by those in grasslands; lowest rates were observed in peatlands and nature reserves. Denitrification correlated to nitrate concentrations, but not to nirK, nirS and nosZ abundance, whereas denitrifier-gene abundance correlated to organic matter content of the sediment, but not to nitrate concentrations. Our results show a mismatch in denitrification regulators at its different organizational scales. Denitrifier abundance is mostly regulated at within-ditch scales, whereas N-loads, regulated by landscape factors, are most important determinants of instantaneous denitrification rates.

Key words: agroecosystem; denitrification; DGGE; ecosystem functioning; nitrogen; macrophytes; qPCR; wetlands.
INTRODUCTION

Drainage ditches are man-made ecosystems, designed to regulate water levels in low-lying agricultural areas and reclaimed wetlands (Painter 1999; Herzon and Helenius 2008; Zhang and others 2016). In the Netherlands, they comprise a total length of 300,000 km (Peeters and others 2014). Besides their importance for drainage of agricultural fields and natural areas, they form a unique ecosystem type, consisting of numerous networks of shallow, low-gradient waterbodies, with low flow velocities. Typical features of the agricultural drainage ditch ecosystem are high nutrient loads and tight aquatic-terrestrial coupling, which makes them more similar to (constructed) wetlands than to streams, as which they are often classified. Ditches often receive runoff and nitrogen-rich groundwater from adjacent fields, leading to excessive production of macrophytes and macroalgae, floating plant dominance, and consequently hypoxia and biodiversity loss (Janse and Van Puijenbroek 1998; Scheffer and others 2003; van Gerven and others 2015). Nitrogen loads from ditches contribute to eutrophication in connected waters, such as canals, lakes and reservoirs (Needelman and others 2007). This can be partly counteracted by denitrification (de Klein and Koelmans 2011; Kröger and others 2014), a major N-removing process that reduces nitrate to gaseous nitrogen.

Denitrification is performed by microorganisms and requires an electron donor like easily degradable organic carbon, and nitrate as terminal electron acceptor (Knowles 1982). In freshwater sediments, denitrification usually takes place in anoxic–oxic boundary layers, where nitrate is supplied from the oxic zone (Seitzinger and others 2006). Similar to wetlands, agricultural ditches are potential denitrification hotspots, due to their tight terrestrial-aquatic coupling, high nitrate inputs, suitable redox conditions, ample anoxic–oxic interface and sediments rich in organic matter (McClain and others 2003; Veraart and others 2011a). However, denitrification rates in ditches have been found to vary widely (de Klein and Koelmans 2011), and it remains unclear which factors are most important in regulating denitrification in these ecosystems.

Factors affecting denitrification act at two different levels: they drive not only the abundance and diversity of the denitrifying microorganisms present, but also the amount of nitrate converted by the resultant denitrifying community (Wallenstein and others 2006). Although most denitrifiers present in the environment remain uncharacterized (Philippot and Hallin 2005), abundance of genes coding for denitrification enzymes can be used to probe denitrification potential (O’Connor and others 2006; Graham and others 2010), whereas variation within functional genes may be used as indicator of denitrifier diversity (Hallin and Lindgren 1999; Throßack 2006).

In this study, we determine how denitrification rates in drainage ditches relate to denitrifier abundance and community structure, using the presence of genes coding for nitrite reductase (nirS, nirK) and nitrous oxide reductase enzymes (nosZ clade I, hereafter: \(\text{nosZ}_i\)) as a proxy. Furthermore, we test how in turn denitrifier activity, abundance and community structure are affected by their environment, at both within-ditch and landscape scales. To this end, we quantify denitrification rates and denitrifier abundance in drainage ditches characterized by different types of sediments, vegetations and land-uses, using an in situ isotope labelling technique, and qPCR and DGGE of the denitrifying community. This way, we can capture controls on denitrification in drainage ditches at organizational scales from microbes to ecosystems.

METHODS

Study Sites and Sampling Design

We sampled 13 drainage ditches (Table 1; Figure 1); all ditches were straight (non-meandering), shallow (max depth 80 cm) and between 1.2 and 6 m wide. They were all situated in a flat landscape, resulting in standing water or negligible flow velocities, regulated by pumping stations. Seven of these ditches were located in peat areas, and were also used to quantify the ditches’ greenhouse gas emission (P1-P4 and SP1-SP3, (Schrier-Uijl and others 2011). We selected the other six ditches to include clay sediments (C1-C4) and fine, sandy sediments (FS1, FS2), the latter of which the top layer consisted of fine, decomposed, material. The ditches were situated in agricultural areas (crops or meadows) and nature reserves or protected areas (mostly peatlands), and therefore differed in yearly N-loads (Table 1). Each ditch was sampled once, in early summer of the same year. We measured denitrification rates, dissolved oxygen, temperature, dissolved organic carbon (DOC), \(\text{NO}_3^-\), \(\text{NH}_4^+\), electric conductivity (EC) and pH in the water column, and the organic matter content (OM%) and oxygen demand of the sediment (SOD). Additionally, the structure and abundance of the \(\text{nirK}, \text{nirS}\) and \(\text{nosZ}_i\) gene pools in the top layer of
Table 1. Locations, Sediment Types and Sampling Dates

| Ditch | Location   | Latitude      | Longitude     | Depth (m) | Width (m) | Soil type<sup>b</sup> | Sediment<sup>c</sup> | Vegetation | Land-use<sup>d</sup> | N-load<sup>e</sup> | Sampling date |
|-------|------------|---------------|---------------|-----------|-----------|------------------------|----------------------|-------------|----------------------|-----------------|--------------|
| C1    | Randwijk  | 51°57'15.35"N | 5°43'51.02"E | 0.35      | 1.2       | R7 Clay               | None/little Grassland | 95.5       | July 9, 2009          |
| C2    | Randwijk  | 51°57'15.55"N | 5°43'51.02"E | 0.45      | 3         | R7 Clay               | None/little Grassland | 95.5       | July 9, 2009          |
| C3    | Lienden   | 51°55'29.10"N | 5°31'47.55"E | 0.35      | 2         | R8 Clay               | Submerged Grassland   | 65.4       | July 10, 2009         |
| C4    | Lienden   | 51°55'29.22"N | 5°31'49.09"E | 0.25      | 2         | R8 Clay               | Submerged Crops       | 65.4       | July 10, 2009         |
| FS1   | Wageningen| 51°59'16.01"N | 5°39'02.54"E | 0.25      | 2.5       | Z20 Fine Sand         | Floating Crops        | 174.6      | July 7, 2009          |
| FS2   | Wageningen| 51°59'16.47"N | 5°39'00.28"E | 0.6       | 2.5       | Z20 Fine Sand         | Floating Crops        | 174.6      | July 7, 2009          |
| P1    | Nieuwkoop | 52°08'32"N    | 4°47'28"E     | 0.35      | 3         | V1 Peat               | None/little Peat/NR.  | 22.3       | June 24, 2009         |
| P2    | Nieuwkoop | 52°08'22"N    | 4°47'50"E     | 0.45      | 3         | V1 Peat               | None/little Peat/NR.  | 22.3       | July 2, 2009          |
| P3    | Oukoop    | 52°02'10.76"N | 4°46'48.42"E | 0.3       | 6         | V1 Peat               | Floating Grassland    | 23.3       | June 23, 2009         |
| P4    | Stein     | 52°01'8.13"N  | 4°46'43.67"E | 0.3       | 5         | V1 Peat               | None/little Grassland | 27.5       | July 1, 2009          |
| SP1   | Doosje    | 52°41'10.81"N | 6°07'57.88"E | 0.45      | 6         | V13 Sand/Peat         | Submerged Peat/NR.    | 23.6       | June 25, 2009         |
| SP2   | st Jansklooster | 52°40'48.53"N | 6°00'28.66"E | 0.3       | 2         | Z18x/A1 (V) Sand/Peat | None/little Grassland | 28         | June 29, 2009         |
| SP3   | Horstermeer | 52°14'25"N   | 5°04'17"E    | 0.8       | 3         | V6 Sand/Peat          | None/little Peat/NR.  | 27         | June 26, 2009         |

Vegetation: the dominant type of macrophytes in the ditch, characterized as either floating (water surface fully covered by Lemna sp.), submerged (sediment surface covered by more than 25% submerged macrophytes, mainly Elodea sp.) or no/little vegetation (less than 25% sediment surface covered by submerged macrophytes).

<sup>a</sup>Approximate coordinates.
<sup>b</sup>Soil classification of the ditch area according to the Dutch classification system (Steur and others 1985), classes are explained in Supplementary Information Table S1.
<sup>c</sup>Sediment type classified by the main soil type in the ditch area. In sand/peat ditches, both sand and peat occur in the first 120 cm of the soil.
<sup>d</sup>Dominant land-use type surrounding the ditch; Peat/NR. are peat ditches in nature reserves or protected areas with mostly swamp vegetation in its surroundings.
<sup>e</sup>Average annual N-load to the surface water (kg ha<sup>−1</sup> y<sup>−1</sup>) of the polder area surrounding the ditch (1985–2005), estimated by the STONE model (Wolf and others 2003).
the sediment were analysed. Furthermore, we determined dominant macrophyte structures by estimating the coverage of floating and submerged vegetation in each ditch.

**Denitrification Measurements**

Denitrification rates were measured in situ using the $^{15}$N isotope-pairing technique in split-box benthic measuring chambers (Figure 1). Use of benthic chambers allows study of denitrification rates under field conditions, and was found to give similar results as more labour intensive laboratory batch-mode assays (Nielsen and Glud 1996; Mengis and others 1997). Measuring under in situ conditions better reflects actual denitrification rates than laboratory incubations. The addition of $^{15}$N-labelled substrate enables denitrification measurements at relatively low NO$_3^-$ concentrations, which reduces overestimation of denitrification. The perspex split-box chambers consisted of a frame and three parallel chambers (12 l each). One split-box chamber was installed in each ditch. We placed the frames in the sediment at least 1 h before starting the measurements. After settling of the disturbed sediment, we placed the three parallel chambers on the frame, fully submersed in the ditch water, capturing submerged vegetation, when present. The top
of the chamber contained a screw cap opening with a septum, through which $^{15}$N could be injected and water samples could be taken. We placed a stirrer next to the septum opening in the chamber, to gently mix the $^{15}$N through the chamber water, and to optimize diffusion of $^{15}$N into the sediment. Measurements were performed in the morning, and each day one ditch was sampled. We injected 5 ml 0.16 M $^{15}$N[NaNO$_3$] through the septum of each chamber, to enrich the water in each chamber with 1 mg l$^{-1}$ $^{15}$N. Water samples for N$_2$ analysis were taken with an airtight glass syringe, whereupon 5 ml of sample was transferred into a helium flushed pre-evacuated 12-ml extainer (Labco Wycombe), which contained 100 µl (50% w:v) ZnCl$_2$, to stop microbial activity. Water samples (triplicates) were taken 0.25, 1, 2, and 3 h after injecting the $^{15}$N solution. Samples were stored at room temperature until the end of analysis. Before analysis, samples were vigorously shaken to transfer the dissolved N$_2$ into the helium headspace. Denitrification rates were calculated from the increase of $^{28}$N$_2$ and $^{30}$N$_2$ in the headspace (Nielsen 1992; Steingruber and others 2001), measured at a SerCon Cryoprep trace gas concentration system interfaced to a PDZ Europa 20-20 isotope ratio mass spectrometer (Sercon Ltd., Cheshire, UK). N$_2$O emission was measured in seven of the sampled ditches, and was found to be negligible (Schrier-Uijl and others 2011).

Conditions in the Water Column

Dissolved oxygen, temperature (T), electric conductivity (EC), and pH in the ditches were measured using an HQ multiprobe with a luminescent dissolved oxygen sensor (Hach Company, Loveland, Colorado, USA), at a location undisturbed by conditions in the water column. Dissolved oxygen, temperature (T), electric conductivity (EC), and pH in the ditches were measured using an HQ multiprobe with a luminescent dissolved oxygen sensor (Hach Company, Loveland, Colorado, USA), at a location undisturbed by placing the denitrification chambers. We measured depth profiles at 10-cm intervals for O$_2$ and T. EC and pH were measured at 20-cm depth in the water column.

For nutrient analysis, mixed water column samples were collected within 2 m from the denitrification chambers. Samples were filtered in the field using 0.45 µm cellulose nitrate filters (Whatman Ltd., Kent, UK), kept cool on ice during transportation and stored at $-20^\circ$C upon arrival in the laboratory. Nutrient concentrations (NO$_3^- + NO_2^-$, NH$_4^+$, ortho-PO$_4^{3-}$) were measured colorimetrically using a SANplus autoanalyzer (Skalar Analytical, Breda, the Netherlands) as described by Veraart and others (2011a) and references therein. DOC was measured using a Total Organic Carbon Analyser (Model 700, O.I. Inter-

national, College Station, TX, USA). Estimates of groundwater seepage in the ditch area were obtained from hydrological maps (van der Gaast and others 2006).

Sediment Characteristics

The top 3 cm of the sediment was sampled using a Kajak corer at 3 locations within 1 m of the denitrification chambers. Samples were mixed to create one mixed sediment sample per ditch, kept on ice during transportation and frozen at $-20^\circ$C until analysis. Organic matter content was determined from the loss on ignition at 550°C for 3 h. Sediment oxygen demand was measured in the field, by inserting a closed dark chamber (10 cm high, 10 cm wide) in bare sediment and monitoring oxygen decrease in the static water at 5-min intervals for at least 3 h using a luminescent dissolved oxygen sensor (Hach Company, Loveland, Colorado, USA).

nirK, nirS and nosZ$_I$ Community Structure

Diversity of the nirK, nirS and nosZ$_I$ genes encoding for nitrite-reductase and nitrous oxide reductase, key enzymes in denitrification, were used as a proxy of the denitrifier community structure. These genes were chosen because they code for those enzymes that produce the gaseous intermediates and products of denitrification, and thus mediate N-removal. Sediment samples from the top 3 cm of sediment were collected and stored as described above. Total DNA was extracted from each sediment sample using a FastDNA® SPIN Kit for Soil (MP Biomedicals, Irvine, CA, USA), according to the manufacturers’ protocol. After extraction, DNA templates were purified using a OneStep™ PCR Inhibitor Removal Kit (Zymo Research, Orange, CA, USA). DNA quality and quantity were checked using a Nanodrop ND-100 spectrophotometer (Thermo Scientific, San Jose, CA, USA) and 1% (w/v) agarose gel electrophoresis, after which the DNA templates were diluted to 20 ng DNA µl$^{-1}$.

Gene fragments were amplified with the primers F1aCu and R3Cu for nirK (Hallin and Lindgren 1999), Cd3aF and R3 cd for nirS (Michotey and others 2000; Throback and others 2006), and nosZF and nosZ622R for nosZ$_I$ (Kloos and others 2001; Enwall and others 2005), with reverse primers having a 33-bp GC-clamp attached to the 5’end. PCR amplification was performed in a total reaction volume of 50 µl, containing 10 µl of 5 × Green GoTaq Reaction Buffer (Promega), 200 µM of each
by Muyzer and Smalla (1998), using a Dcode analysis of amplicons was performed as described above. PCR protocols are described in Veraart and others (2014). PCR amplicons were analysed by 1% (w/v) agarose gel electrophoresis and visualized under UV light after SYBRsafe (Invitrogen) staining.

Denaturing Gradient Gel Electrophoresis (DGGE) analysis of amplicons was performed as described by Muyzer and Smalla (1998), using a Dcode Universal Mutation Detection System (Bio-Rad, Hercules, CA, USA). DGGE was performed on 8% polyacrylamide gels with a denaturant gradient from 40 to 70% (100% denaturing acrylamide was defined as 7 M urea and 40% (v/v) formamide). Aliquots of PCR products were loaded on the gel, and electrophoresis was carried out with 0.5% Tris acetic acid EDTA buffer at 60°C and at 85 V for 16 h, initiated by a pre-run of 10 min at 120 V. After electrophoresis, gels were silver-stained (Sanguinetti and others 1994) and scanned using a GS-800 Calibrated Densitometer (Bio-Rad Hercules, CA, USA).

Quantitative PCR

Abundance of the nirK, nirS and nosZI genes was determined using quantitative PCR. DNA was isolated and purified as described above. Fragments of the genes were amplified using primers nirK876 and nirK1040 for nirK (Henry and others 2004b), nirSCd3aF and nirSR3 cd for nirS (Kandeler and others 2006) and nosZ2F and nosZ2R for nosZI (Henry and others 2006). The 25 µl final volume reaction contained 12.5 µl IQ™ SYBR® Green supermix (BioRad Hercules, CA, USA), 1.4 µM of each primer, 0.25 µl BSA (final concentration 200 ng BSA µl−1), and 5 µl (corresponding to 10 ng) of sample DNA. Thermal cycling was performed using a BioRad CFX96 real-time thermal cycler (Bio-Rad, Hercules, CA, USA), as described in Veraart and others (2014). Specificity of fragment amplification was checked by observing a single band of expected size in a 1.5% agarose gel, and the presence of a single melting peak on the melting curve.

Data analysis and Bionumerics

DGGE band detection was performed using Bionumerics software (version 4.61 Applied Maths, Belgium), with an optimum of 0.5% and a 0.5% position tolerance, but with manual adjustment to avoid misplacing of bands. A reference marker, included on the gel in three different positions, was used as standard for normalization, ensuring sample-to-sample comparability. Similarity between DGGE profiles was determined by calculating similarity indices using the Dice similarity coefficient, which takes into account the presence or absence of specific bands. The unweighted pair-group method with arithmetic means (UPGMA) algorithm was used for dendrogram construction. Richness of the nirK, nirS and nosZI gene pools was obtained from the number of visible bands, which represent operational taxonomic units (OTUs). This method cannot capture the complete richness and diversity existing in these gene pools, but will only depict the most abundant species targeted by the primers used. Nonetheless, comparing DGGE-based richness values can point at potential richness effects at the community level.

We tested for differences in denitrification rates between sediment types and vegetation types using one-way ANOVA. Relations between denitrification and potential explaining variables were tested by linear regression analysis. If necessary, data were ln(x + 1) transformed to achieve a normal distribution. We used redundancy analysis (RDA) to test how environmental variables explained variation in nirK, nirS and nosZI OTUs. The absence/presence for each of the observed OTUs was entered as species data in the ordination. Nitrate, NH4–N, PO4–P, DOC, O2, T, EC and sediment organic matter % were ln(x + 1) transformed and entered as environmental data. Scaling was focused on inter-sample distances. Significance of the canonical axes was evaluated by Monte-Carlo permutation tests (499 permutations). Statistical analyses were performed in SPSS 23 (IBM SPSS statistics, Armonk, NY, USA) and CANOCO 5.03 (Ter Braak and Smilauer, Biometris, Wageningen, the Netherlands).

RESULTS

Conditions in the Ditches

The sampled ditches varied considerably in water column conditions (Table 2). Dissolved oxygen in the water column ranged from 1.0 to 7.3 mg l −1 (average 4.3 ± 2.2 sd mg l −1). Nitrate could only be detected in FS1, FS2 and C3. FS2 had a fivefold higher nitrate concentration than FS1 and a 15 fold higher nitrate concentration than C3. Sediment organic matter averaged 30.0 ± 27.5 sd%. Ditch temperatures were on average 19.7 ± 2.8 °C. SP3 had highest EC, due to high amounts of chloride-rich groundwater seepage.
Table 2. Water Column Conditions and Sediment Characteristics

| Ditch | Denitrification (umol N m\(^{-2}\) h\(^{-1}\)) | NO\(_3^-\) + NO\(_2^-\) (mg N l\(^{-1}\)) | NH\(_4^+\) (mg N l\(^{-1}\)) | PO\(_4^{3-}\) (µg P l\(^{-1}\)) | DOC (mg C l\(^{-1}\)) | DO (mg l\(^{-1}\)) | T (°C) | pH | EC (µS cm\(^{-1}\)) | SOD (g m\(^{-2}\) day\(^{-1}\)) | OM (%) | Seepage (mm d\(^{-1}\)) |
|-------|----------------------------------|-------------------------------|-----------------|-----------------|-----------------|-----------------|--------|----|-----------------|-----------------|--------|-----------------|
| C1    | 497 (93)                         | <0.01                         | 0.11            | 18              | 10.0 (0.2)      | 3.7 (0.6)       | 17.7 (0.1) | 7.2 | 470             | 1.10            | 14.8   | 0.23            |
| C2    | 31 (31)                          | <0.01                         | 0.04            | 32              | 9.7 (0.1)       | 6.4 (0.0)       | 18.2 (0.0) | 7.5 | 389             | n.a.            | 9.9    | 0.23            |
| C3    | 50 (4)                           | 0.32                          | 0.03            | 42              | 6.7 (0.4)       | 7.2 (0.0)       | 16.3 (0.0) | 7.3 | 491             | 2.86            | 12.6   | 0.06            |
| C4    | 143 (22)                         | <0.01                         | 0.03            | 42              | 7.8 (2.2)       | 7.3 (0.1)       | 16.3 (0.1) | 7.4 | 491             | n.a.            | 12.8   | 0.06            |
| FS1   | 3888 (983)                       | 0.88                          | 0.14            | 159             | 10.2 (0.1)      | 4.6 (0.3)       | 19.3 (0.2) | 7.0 | 294             | 2.76            | 2.9    | 0.70            |
| FS2   | 24344 (9955)                     | 4.86                          | 0.07            | 14              | 12.3 (0.4)      | 3.0 (1.5)       | 18.4 (0.3) | 7.1 | 430             | n.a.            | 20.3   | 0.66            |
| P1    | 13 (6)                           | <0.01                         | <0.01           | 21              | 14.6 (0.2)      | 5.9 (3.2)       | 21.0 (0.3) | n.a.| n.a             | 1.10            | 74.5   | –0.71           |
| P2    | 48 (7)                           | <0.01                         | <0.01           | 11              | 15.6 (0.1)      | 3.3 (1.6)       | 25.8 (0.9) | 7.2 | 408             | 0.21            | 74.8   | –0.75           |
| P3    | 137 (45)                         | <0.01                         | 0.02            | 91              | 37.9 (0.2)      | 2.2 (1.7)       | 18.7 (1.6) | 7.3 | 541             | 0.25            | 57.6   | –0.37           |
| P4    | 57 (15)                          | <0.01                         | <0.01           | 30              | 39.2 (1.7)      | 3.2 (2.0)       | 23.5 (0.3) | 6.7 | 231             | 1.73            | 62.9   | –0.38           |
| SP1   | 6 (8)                            | <0.01                         | <0.01           | 18              | 17.1 (0.0)      | 6.9 (2.4)       | 22.1 (1.2) | 7.8 | 420             | 1.52            | 1.0    | 0.40            |
| SP2   | 167 (6)                          | <0.01                         | 0.20            | 255             | 16.4 (0.8)      | 1.0 (0.5)       | 19.7 (2.1) | 7.1 | 325             | 0.74            | 11.7   | 0.00            |
| SP3   | 14 (5)                           | <0.01                         | <0.01           | 12              | 10.8 (1.7)      | 1.7 (1.3)       | 18.6 (1.7) | 6.8 | 832             | n.a.            | 35.2   | 2.00            |

Presented as single measurement or mean (sd)

Seepage estimated groundwater seepage for the ditch area (van der Gaast and others 2006).

DOC = dissolved organic carbon (n = 2); DO = average of depth profile of dissolved oxygen in the water column; T = average of depth profile of temperature in the water column; EC = electric conductivity (n = 1); SOD = sediment oxygen demand (n = 1); OM = organic matter percentage of the sediment (n = 1).

\(^a\)n = 3.

\(^b\)Before adding 1 mg l\(^{-1}\) \(^{15}\)N-NO\(_3^-\) to the denitrification chambers (n = 1).

\(^c\)n = 1.
Denitrification

Denitrification rates varied widely between the ditches, averaging 2261 ± 6718 μmol N m⁻² h⁻¹ (Table 2), with considerable within ditch variability (n = 3, Table 2). Denitrification rates differed among sediment types (One-way ANOVA, F₃ = 33.683, P < 0.001, Figure 2A), vegetation types (One-way ANOVA, F₂ = 19.307, P < 0.001, Figure 2B) and land-use types (One-way ANOVA, F₂ = 35.129, P < 0.001, Figure 2C). Denitrification rates were highest in the ditches with sediments consisting of fine sand (FS1 and FS2, Tukey post hoc, P < 0.01, Figure 2A), followed by clay and peat ditches, with significantly higher rates in clay ditches than in ditches containing mixtures of sand and peat. Rates were highest in ditches covered by floating vegetation (FS1, FS2, P3; Tukey post hoc, P < 0.01, Figure 2B), and in ditches in agricultural areas (Crops > Grasslands > Peatland/Nature reserves, Tukey post hoc, P < 0.01, Figure 2C). Denitrification rates significantly correlated to nitrate (R² = 0.667, P = 0.001, Figure 3A, Table S.2), although this correlation is heavily driven by three data points, whereas the other samples are at the detection limit. Denitrification also correlates to ammonium (R² = 0.317, P < 0.05), and total nitrogen (R² = 0.317, P < 0.05), but not to organic matter content, relative abundance, and DGGE-obtained richness of denitrifier gene pools (Table S.2, Figure S.2). When excluding peat ditches, which have more refractive organic matter, there was also no relation between organic matter content and denitrification rates (R² = 0.244, P = 0.213).

NirK, nirS and nosZ₁ Structure and Abundance

A total number of 94 nirK, 87 nirS and 67 nosZ₁ OTUs were observed in the ditches (Table 3). Highest similarity of OTU occurrence was observed for nosZ₁, where 57% of the DGGE-bands occurred in all ditches, whereas lowest similarity (40%) was observed for nirS. For all genes studied, several distinct clusters of ditches with similar denitrifier communities were observed based on the presence/absence of the different OTUs (Figure S.1). Band analysis clustered the ditches mostly in groups that related to their sediment characteristics or vegetation type: ditches with clay sediments, or fine, sandy sediments always showed distinct clusters, peat and sand/peat ditches showed more variation for nirK and nirS, but clustered according to sediment type for nosZ₁. For nirK all ditches with floating vegetation clustered together. Ditch SP3 formed the overall outgroup for nirS and nirK, with only 40 (nirS) or 50 (nirK) % similarity with the other ditches.

Redundancy analysis clustered the ditches by sediment and vegetation type (Figure 4). For nirK, the first RDA axis was mainly explained by conditions in the sediment and explained 19% of the variance in OTUs. The second RDA axis explained 15% of the variation and was mostly defined by concentrations of solutes, and primary production (Figure 4A), with vegetated ditches on the negative side of the axis, and most unvegetated ditches on the positive side of the axis. Variation in nirS OTUs was better explained than for nirK, the first axis explaining 26% of variation, mostly defined by
nutrient availability, the second 15%, mostly explained by conductivity and pH, showing the same general clustering pattern as nirK. For nosZII, the ditches clustered only by sediment type, regardless of vegetation, the first axis explaining 21% of the variation, the second 16%.

Copy numbers of nirK, nirS and nosZII per gram dry sediment were on average $9 \times 10^4 \pm 1.7 \times 10^5$ sd, $4.1 \times 10^4 \pm 5.4 \times 10^4$, 74 ± 76, respectively. NirK, nirS and nosZII copies per ng sample DNA were on average $1 \times 10^2 \pm 2 \times 10^2$, 55 ± 78, 0.1 ± 0.1, respectively (Table 3). Copy numbers of nirK were significantly correlated to organic matter percentage of the sediment ($\ln(nirK \text{ g dry sed}^{-1}) + 1)$ versus organic matter %, $R^2 = 0.421, P < 0.05$, Figure 2E), but negatively correlated to seepage ($R^2 = -0.367$, $P < 0.05$, Figure 2D). Copy numbers of nirS were only moderately related to seepage ($\ln(nirS \text{ g dry sed}^{-1}) + 1)$ versus organic matter %, $R^2 = 0.367$).

**Figure 3.** Relations between A denitrification and nitrate (linear regression $y = 4.02 + 3.73x$, $R^2 = 0.667$), B denitrification and dissolved oxygen, C denitrification and organic matter % of the sediment, D nirS gene copies per g. sediment and groundwater seepage. E nirK gene copies per g. sediment and organic matter % of the sediment (linear regression $y = 7.67 + 0.06x$, $R^2 = 0.421$), F nirK gene copies per g. sediment and groundwater seepage (linear regression: $y = 9.67 - 1.97x$, $R^2 = 0.367$).
sed$^{-1}$), $R^2 = 0.282, P < 0.1)$. Copy numbers of nosZ$_I$ only tended to be related to OM%, $\ln$(nosZ$_g$ dry sed$^{-1}$) $R^2 = 0.238, P < 0.1$, Table S.2, Figure S.2).

Ditches with high copy numbers of nirK or nosZ$_I$ also had more detected OTUs of the respective gene ($R^2 = 0.318, P < 0.05, R^2 = 0.766, P < 0.001$), but this did not apply to nirS.

**DISCUSSION**

Factors Affecting Drainage Ditch Denitrification at Local and Regional Scales

Denitrification rates of the studied ditches varied widely, and differed among sediment, vegetation and land-use types. Denitrification rates in the agricultural ditches in our study were high, similar to rates previously found in agricultural streams and rivers (García-Ruiz and others 1998; Pattinson and others 1998; Laursen and Seitzinger 2004; Schaller and others 2004). These are among the highest denitrification rates observed in aquatic systems (Piña-Ochoa and Álvarez-Cobelas 2006). However, as drainage ditch residence times are higher, due to low flow velocities, their overall N-removing capacity will be higher. Moreover, denitrification rates within ditches are higher than those within the meadows from which they receive N-rich runoff (comparing rates per m$^2$/h). For example, although in managed grasslands up to 25% of applied fertilizer can be lost due to denitrification, maximal rates are still orders of magnitude lower than average drainage ditch denitrification rates measured in this study (de Klein and Van Logtestijn 1994). These differences arise from the tight aquatic-terrestrial coupling and steep biogeochemical gradients in drainage ditches, leading to favourable heterogeneous oxygen conditions for coupled nitrification–denitrification.

We considered driving factors of denitrification at local scales (within-ditch) and regional (landscape) scales. At local scales, we considered sediment and vegetation types of the ditches as well as physical–chemical properties of sediment and water column and denitrifier’s presence. The regional scale is captured by considering regional N-loads and land-use (see Figure 5 for a schematic overview of effects at multiple scales). Ditches containing fine sand (FS) had higher denitrification rates than those with clay or peat sediments. Fine-textured sediments may support higher denitrification rates, because they have a larger proportion of anoxic microsites compared to coarser sediments (Valett and others 1996; García-Ruiz and others 1998; Martin and others 2001; Findlay and others 2011). Clay sediments have even smaller particle size, but may have had lower denitrification rates due to their lower porosity and therefore hampered diffusion of nitrate to the denitrification zone. However, these ditches were also situated in areas with lower nitrate loads and different vegetation types, which may be an alternative explanation for their lower denitrification rates. In this study, FS ditches had significantly higher nitrate concentrations than ditches of other sediment types, which can contribute to the extreme differences in denitrification rates. Additionally—and possibly as a consequence of high nutrient loads—these ditches were covered by duckweed. Complete duckweed coverage of the

**Table 3.** Abundance and Richness of the nirK, nirS and nosZ$_I$ Gene pools

| Ditch | # nirK OTUs | # nirS OTUs | # nosZ$_I$ OTUs | # copies nirK/g dr sed. | # copies nirS/g dr sed. | # copies nosZ$_I$/g dr sed. | # copies nirK/ng DNA | # copies nirS/ng DNA | # copies nosZ$_I$/ng DNA |
|-------|-------------|-------------|----------------|------------------------|------------------------|--------------------------|-------------------|-------------------|----------------------|
| C1    | 34          | 38          | 33             | 2.30E+03               | 4.38E+03               | 61                        | 4                 | 8                 | 0.11                 |
| C2    | 39          | 19          | 39             | 8.20E+03               | 1.71E+04               | 67                        | 21                | 44                | 0.17                 |
| C3    | 34          | 28          | 36             | 1.60E+04               | 4.37E+04               | 33                        | 24                | 46                | 0.05                 |
| C4    | 45          | 27          | 38             | 1.90E+04               | 6.40E+04               | 49                        | 26                | 87                | 0.07                 |
| FS1   | 45          | 25          | 29             | 5.70E+03               | 8.44E+03               | 35                        | 22                | 32                | 0.13                 |
| FS2   | 45          | 23          | 32             | 2.50E+04               | 3.57E+04               | 140                       | 28                | 41                | 0.16                 |
| P1    | 36          | 32          | 34             | 2.50E+05               | 6.93E+04               | 64                        | 217               | 60                | 0.05                 |
| P2    | 40          | 33          | 41             | 3.70E+05               | 8.69E+04               | 184                       | 199               | 46                | 0.10                 |
| P3    | 49          | 31          | 38             | 5.10E+05               | 1.92E+05               | 262                       | 791               | 298               | 0.41                 |
| P4    | 31          | 23          | 38             | 4.50E+03               | 2.20E+03               | 54                        | 8                 | 4                 | 0.09                 |
| SP1   | 39          | 27          | 21             | 1.80E+03               | 4.61E+03               | 8                         | 11                | 28                | 0.05                 |
| SP2   | 32          | 24          | 16             | 2.10E+02               | 4.82E+02               | 1                         | 1                 | 3                 | 0.00                 |
| SP3   | 32          | 30          | 21             | 8.30E+02               | 7.44E+02               | 10                        | 1                 | 1                 | 0.01                 |

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water column has been shown to significantly reduce water column oxygen concentrations, and as a result, stimulate denitrification rates (Veraart and others 2011a).

Low denitrification rates were observed in the peat ditches. This may be due to nitrogen limitation, because nitrate concentrations were below detection limit in all peat ditches. Also, due to the

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**Figure 4.** Redundancy analysis of *nirK*, *nirS* and *nosZ* OTUs and environmental variables. DOC dissolved organic carbon, DR denitrification rate, EC electric conductivity, OM organic matter content of the sediment, T temperature. Units are as in Table 2. Larger distance on the plot indicates greater dissimilarity between ditches based on OTUs. Arrows indicate the direction of the largest gradient in each environmental variable. *NirK* Eigenvalue of axis 1 ($x$) = 0.187; eigenvalue of axis 2 ($y$) = 0.149. Significance of all canonical axes: $P < 0.1$. *NirS* Eigenvalue of axis 1 ($x$) = 0.258; eigenvalue of axis 2 ($y$) = 0.150. Significance of all canonical axes: $P < 0.05$. *NosZ* Eigenvalue of axis 1 ($x$) = 0.207; eigenvalue of axis 2 ($y$) = 0.161. Significance of all canonical axes: $P = 0.132$. 

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high organic C to NO$_3^-$ ratios that generally prevail in peat sediments, dissimilatory nitrate reduction to ammonium (DNRA) may have been the dominant nitrogen reducing process (Tiedje and others 1982; Burgin and Hamilton 2007). However, the organic carbon may not have been available for nitrate reducing bacteria (those performing either DNRA or denitrification) due to the presence of phenolic compounds in peat. Phenolic substances are potent enzyme inhibitors that may not only inhibit nitrate reducing enzymes directly, but also slow down microbial decomposition under anaerobic conditions (Freeman and others 2001; Freeman and others 2004), resulting in lower nitrate reduction rates due to carbon limitation.

Ditches with floating plants had higher denitrification rates than those without plants or with submerged vegetation. However, distinguishing net vegetation effects on denitrification is complex, because the type of dominant vegetation present in aquatic ecosystems largely depends on nutrient loads, sediment conditions and maintenance strategies (Janse and Van Puijenbroek 1998; Scheffer and others 2003; Kosten and others 2009; van Gerven and others 2015) which all influence denitrification as well.

Overall, denitrification rates were best explained by nitrate and ammonium concentrations, although in the case of nitrate this was mainly caused by the presence of nitrate in three ditches. The relation between nitrate and aquatic denitrification rates is well known, and arises from denitrification reaction stoichiometry (Inwood and others 2005; Piña-Ochoa and Álvarez-Cobelas 2006; Mulholland and others 2008). Ammonium availability in the ditches potentially not only related to denitrification through coupled nitrification–denitrification (Eriksson and Weisner 1999), but may also reflect anoxia in the sediment, which inhibits conversion of ammonium to nitrate by nitrification.

Similar to the meta-analysis of Piña-Ochoa and Álvarez-Cobelas (2006), temperature was not a significant factor explaining denitrification rates in the ditches. This is opposed to studies of Veraart and others (2011b) and Bachand and Horne (1999) which both found a strong temperature effect on denitrification in experimental and single-wetland setups covering a similar temperature range. This may indicate that temperature plays a role within ecosystems, but when comparing rates in different ecosystems other factors limiting denitrification, such as nitrate availability, are more important. However, all measurements for the current study were done in summer, with relatively low variability in temperatures. Over the course of a year, temperature likely has an effect.
Relative Denitrifier Richness and Abundance

Denitrification rates in the ditches were not significantly related to DGGE-obtained nirK, nirS and nosZI richness estimates. The absence of a richness-function relationship in denitrifying communities may be explained by a high functional redundancy of denitrifiers (Wertz and others 2006). Also, richness estimates obtained by DGGE will not capture the true richness of denitrifiers within ditch sediments, because this method is generally assumed to visualize only those OTUs with relative abundances above 1%.

Abundances of the nirK gene observed in this study were in the same range as those reported for soils (Henry and others 2004a), but contrasted observations from North American drainage ditches (Baker and others 2015), in which nirK abundances were below detection limit. Abundance of nirS was lower than reported in North American drainage ditches (Baker and others 2015), and Dutch drainage ditch slurries, rich in organic matter (Kim and others 2015), which can be explained by the different sediment characteristics of the sampled ditches. Similar to other studies, nirK and nirS genes were far more abundant than nosZI genes (Hallin and others 2009; García-Lledo and others 2011), which may point at a large proportion of denitrifiers lacking the capacity to reduce $N_2O$ to $N_2$, and therefore a high risk of $N_2O$ emission. But, we cannot exclude that part of the nitrous oxide reducing community in the sampled ditches may belong to phylogenetic groups not covered by the primers used in this study. In freshwater sediments, abundance of sequences belonging to nosZ clade II can be similar or even higher than that of sequences belonging to clade I, which were measured in this study (Jones and others 2013).

Abundances of nirK, nirS and nosZI were not significantly related to denitrification rates, which is in agreement with results of Graham and others (2010) and Hallin and others (2009) for nirK and nirS but contrasts to the findings of O’Connor and others (2006) for nirK, and Hallin and others (2009) for nosZ. The absence of such a relation may be partly caused by the fact that the presence of a denitrification gene does not mean that it is expressed—and thus functional—in the environment. Denitrifiers containing the cytochrome cd1-nitrate reductase enzyme NirS are expected to dominate in systems with a thick layer of well-decomposed sludge and low oxygen concentrations (Knapp and others 2009). This is in line with our findings. Copy numbers of nirS exceeded those of nirK in ditches with a layer of decomposed sludge, being the fine-sand and clay ditches in our study.

Interestingly, nirK abundance was significantly related to organic matter in the sediment, as was also found by Kandeler and others (2006), but organic matter in the sediment was not related to denitrification. Also, abundance of denitrification genes tended to decline with increasing groundwater-seepage estimates, linking to long-term effects on pore-water chemistry, but possibly also because increased seepage leads to decreased water and solute residence time, leaving less time for denitrification and development of the denitrifier community. Unlike denitrification rates, denitrifier gene abundance in the ditches was not related to nitrate concentration. These results indicate that denitrifier abundance is mainly controlled by conditions in the sediment acting at longer time-scales, such as organic matter’s presence and micronutrients provided by seepage, whereas instantaneous denitrification rates are largely determined by nitrate availability (Graham 2010; Wallenstein 2006).

The absence of a strong denitrifier abundance–denitrification relationship can be explained by a combination of methodological and biological factors. Only one time point was sampled; thus, only capturing a snapshot of denitrification activity, the PCR primers used may not cover the entire diversity of denitrifiers present in the ditch, and lastly, denitrification genes present in the environment will only be expressed under the right conditions. Nitrate limitation may largely explain the absence of a strong relationship between denitrifier gene abundance and denitrification rates in the studied ditches.

Denitrifier Community Structure Related to Environmental Conditions

Community structures based on nirK, nirS and nosZI genes were similar for similar sediment and vegetation types. Part of the variation in community structure was explained by environmental variables, indicating that specific conditions favour some denitrifiers more than others. For all three tested genes, about two-thirds of the total variation in OTUs remained unexplained and may be due to specific conditions in the sediment or geographical factors. Interestingly, not all ditches that were geographically close to each other clustered together in the RDA (for example, P3-P4, SP1-SP2), indicating that local environmental conditions, and in relation to this the dominant vegetation present,
may play a more important role in structuring denitrifying communities. Another example of specific conditions creating different communities is ditch SP3, which was the overall outgroup in the DGGE clustering analysis and RDA of nirK and nirS. This ditch is situated in an area with high rates of groundwater seepage, which is rich in chloride and arsenic, very different hydrological and chemical conditions than the other ditches, which can have led to the different community structures in ditch SP3.

**Implications for Nitrogen Removal from Drainage Ditch Ecosystems**

When recalculating the measured denitrification rates to yearly values, the studied ditches would remove 0.7 to 2986 g m$^{-2}$ y$^{-1}$, corresponding to 4 to 1710% of their average summer nitrogen loads (as calculated by the STONE model and van Gerven and others (2016) (Table 1, Table S.3)). In most drainage ditches in agricultural areas (graslands and crops) more than half of the incoming N can be denitrified (Table S.3), suggesting that denitrification in ditches can indeed significantly contribute to purification of nitrogen polluted surface waters. Ditches in peat areas and nature reserves have lower denitrification efficiencies. In these systems, N-loads may have been overestimated, as their wide buffer zones protect from high N-influx. It is also important to note that these are estimates based on one time point, extrapolated to yearly values, and furthermore NO$_3^-$ addition to nitrate-poor ditches can lead to an overestimation of denitrification rates. More information on seasonal variation is needed to make more precise estimations. In practice there will be a seasonal mismatch between N-loads and denitrification potential. Highest loads occur in winter, when denitrification rates are low due to low temperatures. In summer, denitrification potential can be high, but nitrogen loads are lower than in winter, and incoming nitrogen is rapidly assimilated by the ditch vegetation (de Klein and Koelmans 2011). In some of the sampled ditches, the potential denitrification rate exceeds the estimated summer N-load (Table S.3). This implies that during our measurements all the incoming nitrate is denitrified or taken up by the macrophytes, which is reflected by the absence of nitrate in the water column. The high denitrification and nitrate concentration in fine-sand ditches can be explained by their high seepage rates, supplying the denitrifying community with additional nitrate from the groundwater. Such relation between groundwater seepage rates and locally high denitrification rates was also found in agricultural streams (Veraart and others 2014).

These results shed light on the interplay of landscape factors, hydrology, and biogeochemistry in determining denitrification rates. To some extent, denitrification potential can be estimated from land-use type and vegetation status, which can help to calculate regional nitrogen budgets. Moreover, whereas sediment conditions structure the denitrifier community, local N availability determines instantaneous denitrification rates. Denitrification 'hotspots'—zones with locally enhanced process rates relative to their surroundings—can occur when hydrological flow paths supply ‘missing’ reactants to zones of potential activity (McClain and others 2003). In the case of FS1 and FS2, upwelling nitrate-rich groundwater seepage likely supplied rate-limiting N to an already well-established denitrifying community, leading to excessively high denitrification rates.

**Conclusions**

Drainage ditches can sustain high denitrification rates, which can contribute to reducing N-loads to adjacent surface waters. Denitrification was mainly explained by availability of nitrogen, but not significantly related to abundance of nirK, nirS or nosZ (clade I), indicators of denitrifier presence. Drainage ditches in agricultural areas, with low oxygen concentrations due to a closed cover of floating plants, and high nitrate concentrations provided by groundwater seepage, were denitrification hotspots. Our results show that denitrifier abundance is mostly controlled by long-term conditions in the sediment, whereas nitrogen availability determines instantaneous denitrification rates. Our results show how both landscape factors and within-ditch conditions affect activity and abundance of denitrifier communities, driving denitrification rates.

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