# Gaseous nitrogen and carbon fluxes in riparian alder stands

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Riparian buffer zones are considered to be important elements of agricultural watersheds, in that they control nutrient and carbon fluxes. Although the water purification effect of riparian ecosystems has been well studied, there is little knowledge of their internal cycling, especially in the area of gaseous emissions. We measured fluxes of nitrous oxide (N<sub>2</sub>O), dinitrogen (N<sub>2</sub>), methane (CH<sub>4</sub>), and carbon dioxide (CO<sub>2</sub>) in riparian grey alder stands in southern Estonia and black alder stands in Brandenburg, Germany. Dinitrogen emission was the most important component in N retention in the studied riparian grey alder forests. In 2001–2003, the median values of N, emission in the grey alder sites significantly exceeded the N<sub>2</sub> emission rates in the black alder sites, varying from 700 to 1200 and from 360 to 400 kg N<sub>2</sub>-N ha<sup>-1</sup> year<sup>-1</sup>, respectively. In contrast, the median values of N<sub>2</sub>O flux were higher in the black alder sites than in the grey alder sites, i.e., 0.9–2.6 and 0.2-0.7 kg N<sub>2</sub>O-N ha<sup>-1</sup> year<sup>-1</sup>, respectively. The black alder sites acted as a sink for CH<sub>4</sub>, whereas the grey alder sites emitted a small amount of CH<sub>4</sub>. The CO<sub>2</sub>-C flux was higher in the black alder stands. The estimated N<sub>2</sub>-N emission in the grey alder stands for 1994–1995 was 51.2, whereas  $N_2$ O-N emission was 0.5 kg N ha<sup>-1</sup> year<sup>-1</sup>. The significant increase in  $N_a$  emission from 1994–1995 to 2001–2003 can be related to changes in microbial activity during the succession of the pioneer grey alder stand into a more stable mixed forest community. Due to CO, fluxes and N,O fluxes from sites with altered water regime, the estimated GWP of the studied riparian alder stands was relatively high. Further investigations should concentrate on the factors that regulate rates of N<sub>2</sub>O and N<sub>2</sub> emission from riparian buffer zones.

## Introduction

In their role as the interface between terrestrial and aquatic components of the landscape, riparian buffer zones are important ecotechnological measures to control water quality in agricultural catchments (Kuusemets and Mander 1999). Although the water purification effect of riparian ecosystems has been thoroughly studied (Haycock and Pinay 1993, Vought *et al.* 1994, Mander *et al.* 1995, 1997a), little is known about their internal cycling (Lowrance *et al.* 1983, Peterjohn and Correll 1984), especially concerning gaseous emissions (Groffman *et al.* 1991,

Teiter and Mander 2005). Moreover, some studies have shown that water purification efficiency can be less favorable in riparian zones, which function as hotspots of greenhouse gas emissions with high global warming potential (GWP; Groffman *et al.* 2000).

Alders are known as a host species of symbiotic dinitrogen (N<sub>2</sub>) fixing bacteria (actinobacteria) from the *Frankia* group (Rytter *et al.* 1989). Due to high rates of N<sub>2</sub> fixation, some authors have seen alder forests as sources of water body pollution with excess nitrogen (N) (Binkley et al. 1992). Several other studies consider riparian alder stands to be effective N removal ecosystems (Mander et al. 1995, 1997a, 1997b). This contradiction is mainly due to the positioning of alder stands in the landscape: in riparian zones the excess N is mainly denitrified, whereas in more aerated conditions of higher altitude locations (see Binkley et al. 1992) leaching takes place. Denitrification, which is generally referred to as the microbial reduction of nitrate-N (NO<sub>3</sub>-N) to nitrite-N (NO<sub>2</sub>-N) and further to the gaseous forms nitric oxide (NO), (nitrous oxide) N<sub>2</sub>O and N<sub>2</sub> (Knowles 1982), has been found in numerous studies to be a significant process in N removal in riparian buffer zones (Groffman et al. 1991, Ambus and Christensen, 1993, Hanson et al. 1994, Weller et al. 1994, Gold et al. 1998, Hefting and Klein 1998, Groffman et al. 2000). In the majority of these studies, N<sub>2</sub>O fluxes have been measured, while only a few studies pay attention to N<sub>2</sub> emission (Watts and Seitzinger 2000, Butterbach-Bahl *et al.* 2002).

Nitrous oxide, as one of the greenhouse gases, is increasing in the atmosphere at a rate of about 0.3% year<sup>-1</sup> (Mosier 1998). It has an atmospheric lifetime of about 120 years, a GWP of 296 relative to carbon dioxide (CO<sub>2</sub>), over a 100year time horizon, and is responsible for about 5% of the anticipated warming (IPCC 2001). Riparian zones have the potential to be hotspots of N<sub>2</sub>O production in the landscape (Groffman et al. 2000). Likewise, riparian wetlands and wet riparian forests can be sources of methane (CH<sub>4</sub>) (Jones and Mulholland 1998, Rush and Rennenberg 1998), which is another greenhouse gas that is increasing in the atmosphere at a rate of about 0.8% year<sup>-1</sup> (Mosier 1998). Methane in the atmosphere has a lifetime of 8.4 years. On a 100-year time horizon,  $CH_4$  has a global warming potential of 23 relative to  $CO_2$ , and is responsible for about 20% of anticipated warming (IPCC 2001).

Both denitrification and CH<sub>4</sub> formation depend on the oxygen status of the soil or sediment. As a result, the spatial and temporal variability of fluxes of both N<sub>2</sub>O (Robertson and Tiedje 1984, Ambus and Christensen 1993, Augustin *et al.* 1998, Gold *et al.* 1998, Jacinthe *et al.* 1998) and CH<sub>4</sub> (Saarnio *et al.* 1997, Willison *et al.* 1998) are high. Denitrification rates in soils are mainly influenced by carbon availability, NO<sub>3</sub><sup>-</sup> availability, temperature and pH (Nõmmik 1956, Knowles 1982). Methane is produced in anoxic soils and sediments, while well-drained soils act as a sink for atmospheric CH<sub>4</sub> due to CH<sub>4</sub> oxidation, through either ammonia oxidizers or methanotrophs (Hanson *et al.* 1993).

Several studies consider CO<sub>2</sub> emissions and sequestration in riparian wetlands (Mitsch and Gosselink 1993) and buffer zones (Brumme *et al.* 1999, Gulledge and Schimel 2000, Tufekcioglu *et al.* 2001, Larmola *et al.* 2003, Scott *et al.* 2004, Teiter and Mander 2005, von Arnold *et al.* 2005) Depending on meteorological and hydrological conditions, riparian ecosystems, especially wetlands, can be either sources or sinks of carbon (C) (Gulledge and Schimel 2000).

The main objectives of this research were to quantify and compare  $N_2O$ ,  $N_2$ ,  $CH_4$  and  $CO_2$  emission rates in two different riparian alder forests: grey alder, *Alnus incana*, and black alder, *A. glutinosa*, and to estimate the global warming potential (GWP) of the analyzed greenhouse gases. Based on our measurements and data from an earlier study, we further estimate the role of  $N_2O$  and  $N_2$  emission in the nitrogen budget of the grey alder stand.

### Material and methods

#### Study sites

The Porijõgi study area represented a grey alder stand. It is situated in the moraine plain of south-eastern Estonia (Tartu County, Sirvaku; 58°13′N, 26°47′E), in the riparian zone of a small river, the Porijõgi, which flows in a primeval valley where

agricultural activities ceased in 1992. The landscape study transect in this valley crosses several plant communities: an abandoned field (last cultivated in 1992) on Planosols and Podzoluvisols; an abandoned cultivated grassland (last mown in 1993) on Colluvial Podzoluvisol (dominated by Dactylis glomerata and Alopecurus pratensis); an 11 m wide wet grassland on Gleysol (two parallel communities, one dominated by Filipendula ulmaria, another by Aegopodium podagraria) and a 20-m wide grey alder stand on Mollic Gleysol. In the grey alder stand, 3 sites: Edge, Wet and Dry were chosen for gas and soil analyses. The main soil characteristics of the Porijõgi study area are presented in Table 1. For a more detailed description see Kuusemets et al. (2001).

The Gumnitz study area is located in Müncheberg (52°50′N, 14°14′E) in Brandenburg State in northeastern Germany. Vegetation cover at this site represented a black alder forest on Gleysols and Histosols, more specifically a *Carici acutiformis–Alnetum* community, in which *Carex acutiformis* and *Carex canescens* dominated. The Gumnitz A site represented a riparian alder stand with lowered water table due to drainage, whereas the Gumnitz B has an unchanged water regime. The main soil characteristics of the two riparian study sites Gumnitz A and Gumnitz B are presented in Table 1.

#### Gas sampling and analyses

For the measurement of N<sub>2</sub>O, N<sub>2</sub>, CH<sub>4</sub> and CO<sub>2</sub> emissions, the closed chamber method (Hutchinson and Livingston 1993) and the He-O method (Butterbach-Bahl *et al.* 1997, Scholefield *et al.* 1997, Mander *et al.* 2003) were used. The latter

allows measurement of  $N_2$  fluxes. Five gas samplers were installed: one at each of the 3 different sites (Edge, Wet and Dry) in the Porijõgi riparian buffer zone, and on the stream bank (A) and the alder stand edge (B) in Gumnitz. The samplers were closed chambers with a cover made of PVC, height 50 cm,  $\varnothing$  50 cm, volume 65 l, sealed with a water-filled ring on the soil surface, painted white to avoid heating during application.

Gas sampling was carried out according to the following schedule: (1) in Porijõgi, once a month in May, June and September 2000, October and November 2001, March, May to December 2002, and January to March, July and October 2003; (2) in Gumnitz, 3-4 times a month from January to September 2000 and once a month in January and May 2001 and in May 2002. At the end of the 1-hr measuring period, gas samples were taken from the enclosures of samplers with previously evacuated gas bottles (100 ml; see Augustin et al. 1998). The soil temperature and redox potential, and water depth in the sampling wells was measured simultaneously, and the NH<sub>4</sub>-N and NO<sub>3</sub>-N concentration in soil samples was analysed using the Kjeldahl method (APHA 1989).

Intact soil cores (diameter 6.8 cm, height 6 cm) for use with the He-O method were taken from the topsoil (0–10 cm) at the gas sampler sites each time after gas sampling was completed. Soil samples were weighed, kept at low temperature (4 °C) and transported to the laboratory of the Institute of Primary Production and Microbial Ecology of the Centre for Agricultural Landscape and Land Use Research (ZALF) in Germany. At the lab, the soil cores were introduced into special gas-tight incubation vessels. In these vessels, N, was removed using 3 sub-

**Table 1**. Main soil characteristics of the study sites.

Study site	Average water table (m)	рН	Dry matter (%)	N (%)	NH <sub>4</sub> +-N (mg 100 g <sup>-1</sup> )	NO <sub>3</sub> -N (mg 100 g <sup>-1</sup> )	C (%)
Gumnitz A	0.60	5.2	39.8	1.6	0.43	20.90	20.1
Gumnitz B	0.02-0.04	5.7	25.7	1.9	0.46	14.30	39.5
Porijõgi Wet	0-0.05	6.5	55.0	0.4	1.88	0.09	4.0
Porijõgi Dry	0.45-0.95	6.3	44.4	0.8	0.33	0.14	4.5
Porijõgi Edge	0.45-0.95	6.3	65.5	0.3	0.61	0.11	5.3

sequent slight evacuation/flushing cycles with an artificial gas mixture (21.3% O<sub>2</sub>, 78.6% He, 337 ppm CO<sub>2</sub>, 374 ppb N<sub>2</sub>O, 1882 ppb CH<sub>4</sub> and approximately 5 ppm  $N_2$ ). This was followed by the establishment of a new flow equilibrium by continuously flushing the vessel headspace with the gas mixture at 10 ml per minute for 12 hours. For the start value, N, and the greenhouse gas concentration in the gas mixture was measured. The gas concentrations in the incubation headspace were measured (final value) after closing the incubation headspace for one hour to accumulate the emission of N, and the greenhouse gases. The final accumulation value minus the start continuous flow value served as the basis for the calculation of the emission rates. During the flushing the redox potential of the soil cores was regularly measured and regulated so that it was comparable with the field conditions. The gas concentration in the collected air was determined by using gas chromatograpy (electron capture detector and flame ionization detector; Loftfield et al. 1997) in the lab of the Institute of Primary Production and Microbial Ecology, Centre for Agricultural Landscape and Land Use Research (ZALF), Germany. The procedures used for the determination of the emission rates of gases are described by Mander et al. (2003).

### Water sampling and analyses

In Porijõgi, shallow groundwater samples from the upper aquifer were collected once to twice a month from 6 piezometers, 3 of them installed on the upper border and 3 within the grey alder stand. The depth of ground water varied from 10-80 cm. Filtered water samples were analysed for NH<sub>4</sub>-N, NO<sub>2</sub>-N, NO<sub>3</sub>-N, total Kjeldahl-N, phosphate (PO<sub>4</sub><sup>3</sup>-P), total phosphorus (total-P), sulphate (SO<sub>4</sub><sup>2-</sup>), iron (Fe) and calcium (Ca<sup>2+</sup>) in the Laboratory of Plant Biochemistry of the Estonian Agricultural University following standard methods for the examination of water and wastewater quality (APHA 1989). Groundwater discharge was estimated on the basis of both Darcy's law and through gauging with weirs installed in groundwater seeping sites. Average annual input discharge was estimated to be  $8.3 \pm 3.5 \text{ m}^3 \text{ ha}^{-1} \text{ d}^{-1}$ . Meteorological analyses

are based on precipitation, air temperature, wind velocity and humidity data, measured six times a day, as well as daily, monthly and annual averages from the Ülenurme Meteorology Station of the Estonian Meteorology and Hydrology Institute (EMHI). The station is located near the weir of the Porijõgi transect. Mean annual precipitation for the study period varied from 555 to 711 mm.

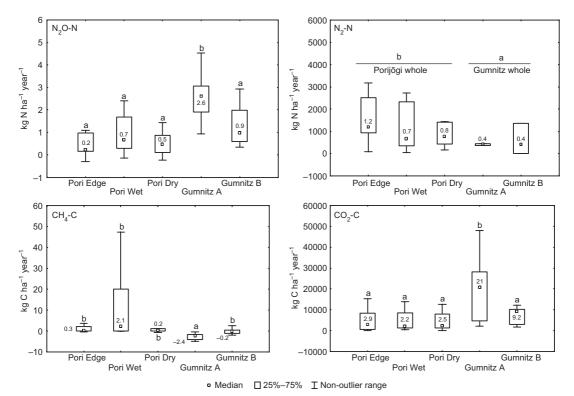
# Soil sampling and analyses in the Estonian study area

In Porijõgi, three soil samples of 50 cm³ were taken at two depths (0–10 cm and 10–20 cm) from all plant communities through the riparian buffer zone. Sampling was carried out twice a year: in spring (May) and autumn (October). Soil pH value, organic matter (loss of ignition), Kjeldahl-N, and lactate-soluble P concentrations were analyzed in all soil samples using the standard methods (APHA 1989).

# Calculations and statistical analyses

In order to estimate the final  $N_2$ -N flux values, we used the  $N_2$ : $N_2$ O ratio calculated using the He-O method. The GWP of the studied systems was calculated by converting the fluxes of  $N_2$ O and  $CH_4$  into  $CO_2$  equivalents (IPCC 2001).

The normality of variable distributions was checked using the Kolmogorov-Smirnov, Lilliefors, and Shapiro-Wilk's tests. In most cases for gas analyses, the distribution differed from the normal distribution, and hence non-parametric tests were performed. Medians, 25% and 75% percentiles and non-outlier range values of variables are presented. We used the Kruskal-Wallis ANOVA and multiple comparison of mean ranks to check the significance of differences between the gas fluxes at different sites. For CH<sub>4</sub>-C, the Duncan test was used. The Mann-Whitney Utest was used to check the difference between the gas fluxes in the study areas in Estonia and Germany. The statistical analysis was carried out using Statistica ver. 7.1 (StatSoft Inc.). The level of significance of  $\alpha = 0.05$  was accepted in all cases.



**Fig. 1.** Median, 25% and 75% percentiles and non-outlier ranges of  $N_2O$ ,  $N_2$ ,  $CH_4$ , and  $CO_2$  fluxes from three sites (Edge, Wet, Dry) of the Porijōgi grey alder stand and two sites (A, B) of the Gumnitz black alder stand. Nitrous oxide and  $CH_4$  fluxes are given in kg ha<sup>-1</sup> year<sup>-1</sup>, and  $N_2$  and  $CO_2$  are given in the ha<sup>-1</sup> year<sup>-1</sup>. Numbers indicate median values, and letters show significantly (p < 0.05) differing values according to the multiple comparison of test sites: for  $CO_2$ ,  $N_2O$  and  $N_2$  using the Kruskal-Wallis test, and for  $CH_4$  using the Duncan test.

# Results

#### Gaseous emissions

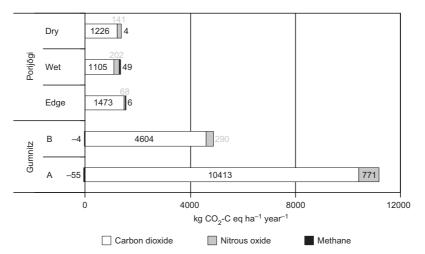
We found remarkable differences in the median values of gaseous emissions between different study sites (Fig. 1). Riparian grey alder sites emitted significantly less N<sub>2</sub>O-N than black alder sites: the median values for N2O flux varied from 0.2 to 0.7 and from 0.9 to 2.6 kg N ha<sup>-1</sup> year-1 respectively, and for N<sub>2</sub> flux from 700 to 1200 and from 360 to 400 kg N ha<sup>-1</sup> year<sup>-1</sup> correspondingly (Fig. 1). No significant difference was found between different sites in the grey alder forest, while the N<sub>2</sub>O-N flux in black alder stand with altered water regime (Gumnitz A: 2.6 kg N ha<sup>-1</sup> year<sup>-1</sup>) was significantly higher than in the black alder stand with unchanged water regime (Gumnitz B: 0.9 kg N ha<sup>-1</sup> year<sup>-1</sup>; Fig. 1). No significant differences between N<sub>2</sub>-N:N<sub>2</sub>O-N ratios in Estonian grey alder stands were found.

The CH<sub>4</sub> emission showed elevated values in the grey alder sites, and was highest in wetter conditions (Porijõgi Wet: 2.1 kg C ha<sup>-1</sup> year<sup>-1</sup>). At the same time, the black alder forests acted as a methane sink (–0.2 and –2.4 kg C ha<sup>-1</sup> year<sup>-1</sup> in Gumnitz B and A, respectively; Fig. 1).

The emission of CO<sub>2</sub>-C was highest in the black alder forest with lowered water table (Gumnitz A: 21 t C ha<sup>-1</sup> year<sup>-1</sup>), whereas no significant differences were found between the study sites in the grey alder forest (2.2–2.9 t C ha<sup>-1</sup> year<sup>-1</sup>; Fig. 1).

# Global warming potential of studied systems

The studied riparian buffer zones did show relatively high GWP values: from 1356–1547 kg CO<sub>2</sub>-C equivalents (eq) ha<sup>-1</sup> year<sup>-1</sup> in Porijõgi to 4870–11 130 CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup> in Gumnitz



**Fig. 2**. The share of major greenhouse gases from the Porijõgi and Gumnitz riparian study sites from October 2001 to November 2003, presented as median CO<sub>2</sub> equivalent values (kg CO<sub>2</sub>-C ha<sup>-1</sup> year<sup>-1</sup>). The conversion of the flux rates into CO<sub>2</sub> equivalents is given with 296 for N<sub>2</sub>O and 23 for CH<sub>4</sub> in a time horizon of 100 years (IPCC 2001). Carbon dioxide values are reduced by 50% assuming that this proportion can be assimilated in photosynthesis (*see* Butnor *et al.* 2003).

(Fig. 2). Carbon dioxide constituted the largest part of the total GWP of the riparian alder forests, showing average values of 1473, 1105 and 1226 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup> in Porijõgi Edge, Wet and Dry sites, respectively, and 10 413 and 4604 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup> in Gumnitz A and B sites, correspondingly (Fig. 2). The share of N<sub>2</sub>O in GWP was significantly higher in the black alder stands in Gumnitz (290–791 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup>) than in the grey alder stand in Porijõgi (68–202 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup>; Fig. 2). The share of methane fluxes in GWP was very low, while it was negative in the black alder stands at both Gumnitz sites (–4 to –55 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup>).

# **Discussion**

Several investigations into N<sub>2</sub>O and CH<sub>4</sub> emissions from riparian zones show significant variability in emission rates, as well as in the percentage share of N<sub>2</sub>O-N in the N load (Table 2). The N<sub>2</sub>O and CH<sub>4</sub> fluxes have ranged from -0.22 to 20 kg N<sub>2</sub>O-N ha<sup>-1</sup> year<sup>-1</sup> and from -4.4 to 36 kg CH<sub>4</sub>-C ha<sup>-1</sup> year<sup>-1</sup>, while the share of N<sub>2</sub>O-N emission in the total N input has varied from 0.02% to 4.2%. Likewise, our earlier studies showed a remarkable variability of greenhouse

gas emissions from the Porijõgi alder forest (Mander *et al.* 2005, Teiter and Mander 2005)

Dinitrogen emission was the most important component of N retention within the Porijõgi riparian grey alder forest. The estimated N<sub>2</sub>-N emission for 1994–1995 was 51.2 kg N ha<sup>-1</sup> year<sup>-1</sup> (Lõhmus *et al.* 2002), while the latest measurements in 2001–2003 gave median values as high as 700-1200 kg N ha-1 year-1. The intensive N<sub>2</sub> emission from the Porijõgi test area can be related to the high microbial activity in alder forests (Hart et al. 1997, Dilly et al. 2000, Lõhmus et al. 2002), which could be assumed to lead to higher denitrification activity. One of the N<sub>2</sub> sources could be denitrification (degassing) in the groundwater, although this has been measured in riparian wetlands (Blicher-Mathiesen et al. 1998, Mookherji et al. 2003). Mogge et al. (1998) found that denitrification in a black alder forest was approximately 15 times greater than in beech forest. The comparison of nitrogen-fixing red alder and Douglas fir forests has shown that denitrification enzyme activity was greater in the alder forest. Also, in alder forest denitrification was limited only by organic matter and not by NO<sub>3</sub>-, whereas in Douglas-fir soils it was frequently limited by both NO<sub>3</sub><sup>-</sup> and energy (Griffiths et al. 1998). On the other hand, the CH<sub>4</sub> oxidation capacity in alder forests has been found to be significantly lower than in other forests (Reay *et al.* 2001, 2005). This explains the higher CH<sub>4</sub> emission values in alder stands (*see* the Wet site in Porijõgi; Fig. 1).

To determine the possible impact of stand age and site conditions on the forms of N emission, we measured gaseous emissions from additional grey alder stands in southern Estonia in July and October 2003. In a Viiratsi riparian grey alder stand (Kuusemets et al. 2001, Lõhmus et al. 2002, Mander et al. 1997) two sites in the 50year-old part and one in the < 10-year-old buffer strip were involved in the study. In Holvandi, emissions were measured in a 9-year-old grey alder plantation on an abandoned field (Uri et al. 2003a) where the soil N pool was 3-4 times lower than in riparian grey alder stands. In all of the grey alder stands studied (Porijõgi, Holvandi, and Viiratsi) the N<sub>2</sub>-N:N<sub>2</sub>O-N ratio was very high.

In riparian grey alder stands  $N_2$  emissions were significantly lower in 50-year-old sites than in younger ones, which indicates that  $N_2$  emission in alder stands could be higher in earlier successional phases. This is most likely related to changes in microbial activity during the succession of a pioneer grey alder stand into a more stable mixed forest community.

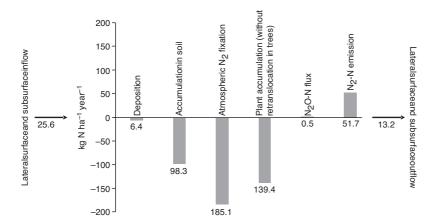
Based on the total inputs and outputs as well as accumulation and transformation rates of nitrogen estimated in our earlier publications (Lõhmus *et al.* 2002, Mander *et al.* 2005) the N budget of the grey alder stand in Porijõgi was established (Fig. 3). Total inputs (kg N ha<sup>-1</sup> year<sup>-1</sup>), which consist of symbiotic  $N_2$  fixation (184.6), subsurface and overland flow (25.6), deposition of mineral nitrogen (6.4), and non-symbiotic  $N_2$  fixation (0.5), were 217.1 kg N ha<sup>-1</sup> year<sup>-1</sup>. This relatively high value is due to high symbiotic fixation, whereas the non-symbiotic  $N_2$  fixation was estimated to be negligible.

Nitrogen accumulation in plants after litter fall was 139.4 kg N ha<sup>-1</sup> year<sup>-1</sup> (Fig. 3). The annual N accumulation in soil was remarkably high (98.3 kg N ha<sup>-1</sup> year<sup>-1</sup>), about half of the fixed N<sub>2</sub>-N. The total N demand in the Porijõgi grey alder stand in 1994 was 285.3 kg N ha<sup>-1</sup> year<sup>-1</sup>; 81.9 kg N ha<sup>-1</sup> year<sup>-1</sup> was transformed and assimilated by plants, whereas the litter amounted to 132 kg N ha<sup>-1</sup> year<sup>-1</sup> (*see* Lõhmus *et al.* 2002).

Table 2. Emission of nitrous oxide and methane in riparian buffer zones.

Reference	Country	Ecosystem type	N <sub>2</sub> O-N emission (kg N ha <sup>-1</sup> year <sup>-1</sup> )	CH₄-C emission³ Initial load³ (kg C ha⁻¹ year⁻¹) (kg N ha² year⁻¹)	Initial load <sup>a</sup> (kg N ha <sup>-2</sup> year <sup>-1</sup> )	N <sub>2</sub> O % of N input	Comment
Weller <i>et al.</i> 1994	USA	Riparian	0.16-0.88ª	I	24.1–102ª	0.65-0.87ª	range of
Jones <i>et al.</i> 1995	NSA	Riparian (hyporheic)	1	0.32-14.9ª	I	I	range
Jacinthe et al. 1998	NSA	Riparian forested wetland	0.66-11.0 <sup>a</sup>	I	65.7–296ª	0.02-3.7a	range
Groffman <i>et al.</i> 2000	NSA	Riparian forested	I	I	I	0.09-0.45a	range
Hefting et al. 2003	The Netherlands	wettarius Riparian black alder foroct	20.0ª	I	475ª	4.2ª	average value
Teiter & Mander 2005	Estonia	Riparian gray alder	-0.22-3.6	-0.44-36	11.0–292	1.2–2	range of mean
This paper	Germany	Riparian black alder forest	0.44–7.8	-4.4-3.1	I	I	values range of mean values

original data given in µg m-2 h-1 or in mg m-2 d-1.



**Fig. 3.** Nitrogen fluxes (kg ha<sup>-1</sup> year<sup>-1</sup>) in the Porijõgi grey alder forest in 1994–1995. Compiled after Lõhmus *et al.* 2002.

Nitrogen output into groundwater and streams was  $13.2 \text{ kg N ha}^{-1} \text{ year}^{-1} \text{ (Fig. 3)}$ . The estimated N<sub>2</sub>-N emission for 1994–1995 was 51.2, whereas emission of N<sub>2</sub>O-N, which contributes to global warming and damages the ozone layer, was 0.5 kg N ha<sup>-1</sup> year<sup>-1</sup> (Fig. 3). Thus the nitrous oxide flux was 3 orders lower than the dinitrogen emission.

Considering all inputs and outputs, the N removal efficiency in grey alder stands decelerates with increasing age (see also Mander et al. 2005). Thus grey alder buffer communities should be managed by regeneration cutting and tending so as to keep their N removal rate high.

Due to intensive N and C retention, the P retention in alder forests is also remarkable (Giardina *et al.* 1995, Zou *et al.* 1995, Uri *et al.* 2003b, Mander *et al.* 2005). Thus alder forests with intensive internal nutrient cycling work as effective soil producers without significant leaching of nutrients. In riparian zones denitrification is the key factor in balancing N fluxes.

The relatively high  $\mathrm{CO}_2$  emission from the Gumnitz study area is related to the significantly higher total C concentration of soils in Gumnitz as compared with that in the Porijõgi study area (20%–40% and 4.0%–5.3%, respectively). In our study, the  $\mathrm{CO}_2$  emission is not connected with fluxes related to plant photosynthesis. Therefore only data for cold periods can be considered as losses to the atmosphere. In the calculation of net ecosystem  $\mathrm{CO}_2$  exchange, a more advanced measurement technique is required.

However, some studies on C sequestration in wetlands and forest ecosystems (Butnor *et al.*)

2003) allow us to estimate that about 50% of the  $\rm CO_2$  released during soil respiration, will in the vegetation period be assimilated by trees through the photosynthesis. Therefore this 50% reduction has been taken into account in our results on  $\rm CO_2$  emissions from forest ecosystems (Fig. 3).

In comparing the greenhouse potential of CH<sub>4</sub> and N<sub>2</sub>O over a long time scale (100–500) years), one can speculate that due to the short adjustment time for CH<sub>4</sub> in the atmosphere (8.4) years; IPCC 2001), the radiative forcing of CH<sub>4</sub> will fall relative to CO<sub>2</sub> (Whiting and Chanton 2001). Nitrous oxide, with its atmospheric lifespan of about 120 years and GWP value of 296, however, has a more significant impact. Our data show a minor CH<sub>4</sub> emission from riparian alder forests. Likewise, the N<sub>2</sub>O emission was relatively low as compared with emissions from constructed wetlands for wastewater treatment (Teiter and Mander 2005). However, the high radiative forcing value of N<sub>2</sub>O makes its share in total GWP remarkable, being highest in the riparian black alder forest with altered water regime (771 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup> in Gumnitz A; Fig. 3). The lowering of the water table in wetlands is a well-known reason for N<sub>2</sub>O emission (Martikainen et al. 1993). The planned increase in fertilization intensity and reconstruction of abandoned drainage systems in several Eastern European countries after the political and socioeconomic changes of the 1990s (see Mander et al. 2000, Stålnacke et al. 2003) may change the N balance of riparian ecosystems. However, its dynamics and especially the change in gaseous N fluxes are relatively unpredictable. Therefore further investigations should concentrate on the factors that regulate  $N_2O$  and  $N_2$  emission rates from riparian buffer zones.

# **Conclusions**

Gaseous emissions from the studied riparian alder stands showed significant spatial variation. The median values of  $N_2O-N$ ,  $N_2-N$  and  $CH_4-C$  ranged from 0.2 to 2.6, from 360 to 1200 and from –2.4 to 2.1 kg ha<sup>-1</sup> year<sup>-1</sup>, respectively. The  $CO_2-C$  fluxes varied from 2.2 to 21 t ha<sup>-1</sup> year<sup>-1</sup>. In the grey riparian alder forest sites the  $CH_4$  and  $N_2$  fluxes were higher and the  $CO_2$  and  $N_2O$  fluxes were lower than in the black alder sites.

Dinitrogen emission was found to be the most important component in N retention from the studied riparian grey alder forests (up to 1200 kg N<sub>2</sub>-N ha<sup>-1</sup> year<sup>-1</sup>). On the other hand, we found a significantly higher N<sub>2</sub>-N emission and lower N<sub>2</sub>O-N flux from the grey alder stand on mineral soils than from the black alder stands on organic soils. Nitrous oxide emission was significantly higher in a study site with a lowered groundwater table, while the methane emission showed higher values in wetter conditions at Porijõgi Wet Site.

Carbon dioxide constituted the largest part of the total GWP of riparian alder forests, being significantly higher in black alder sites (from 4604 to 10413 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup>) as compared with that in the grey alder sites (from 1105 to 1473 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup>). The share of N<sub>2</sub>O in the GWP was also significantly higher in the black alder stands: 290–771 and 68–202 kg CO<sub>2</sub>-C eq ha<sup>-1</sup> year<sup>-1</sup> for black alder and grey alder stands, respectively.

Riparian grey alder forests as pioneer communities are optimal buffer zones, and their N removal capacity is high despite the N accumulation in soil due to symbiotic  $N_2$  fixation. Instead of leaching, in these ecosystems denitrification works down the soil N storage accumulated during succession.

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### References

- APHA 1989. Standard methods for the examination of water and waste water, 17th ed. American Public Health Organisation, Washington, DC.
- Ambus P. & Christensen S. 1993. Denitrification variability and control in a riparian fen irrigated with agricultural drainage water. Soil Biol. Biochem. 25: 915–923.
- Augustin J., Merbach W. & Rogasik J. 1998. Factors influencing nitrous oxide and methane emissions from minerotrophic fens in northeast Germany. *Biol. Fert. Soils* 28: 1–4.
- Binkley D., Sollins S., Bell R., Sachs D. & Myrold D. 1992. Biogeochemistry of adjacent conifer and alder-conifer stands. *Ecology* 73: 2022–2033.
- Blicher-Mathiesen G., McCarty G.W. & Nielsen L.P. 1998. Denitrification and degassing in groundwater estimated from dissolved dinitrogen and argon. *J. Hydrol*. 208: 16–24.
- Brumme R., Borken W. & Finke S. 1999. Hierarchical control on nitrous oxide emission in forest ecosystems. *Glob. Biogeochem. Cy.* 13: 1137–1148.
- Butnor J.R., Johnsen K.H., Oren R. & Katul G.G. 2003. Reduction of forest floor respiration by fertilization on both carbon dioxide-enriched and reference 17-year-old loblolly pine stands. *Glob. Change Biol.* 9: 849–861.
- Butterbach-Bahl K., Willibald G. & Papen H. 1997. A new method for simultaneous measurements of N<sub>2</sub> and N<sub>2</sub>Oemissions from intact soil cores. In: van Cleemput O., Haneklaus S., Hofman G., Schnug E., Vermoesen A. (eds.), Fertilization for sustainable plant production and soil fertility, Proceedings of 11th World Fertilizer Congress of CIEC, vol. 2, Ghent University Press, Ghent, Belgium, pp. 618–624.
- Butterbach-Bahl K., Willibald G. & Papen H. 2002. Soil core method for direct simultaneous determination of N<sub>2</sub> and N<sub>2</sub>O emissions from forest soils. *Plant Soil* 240: 105–116.
- Dilly O., Bach H.-J., Buscot F., Eschenbach C., Kutsch W., Middelhoff U., Pritsch K. & Munch J.C. 2000. Characteristics and energetic strategies of the rhizosphere in ecosystems of the Bornhöved Lake district. *Appl. Soil Ecol.* 15: 201–210.
- Giardina C.P., Huffman S., Binkley D. & Caldwell B.A. 1995. Alders increase soil phosphorus availability in a Douglas fir plantation. Can J. Forest Res. 25: 1652-1657.
- Gold A.J., Jacinthe P.A., Groffman P.M., Wright W.R. & Puffer R.H. 1998 Patchiness in groundwater nitrate removal in a riparian forest. J. Environ. Qual. 27: 146–155.
- Griffiths R.P., Homann P.S. & Riley R. 1998. Denitrification enzyme activity of Douglas-fir and red alder forest

- soils of the Pacific Northwest. *Soil Biol. Biochem.* 30: 1147–1157.
- Groffman P.M., Axelrod E.A., Lemunyon J.L. & Sullivan W.M. 1991. Denitrification in grass and forest vegetated filter strips. J. Environ. Qual. 20: 671–674.
- Groffman P.M., Gold A. & Addy K. 2000. Nitrous oxide production in riparian zones and its importance to national emission inventories. Chemosphere Global Change Science 2: 291–299.
- Gulledge J. & Schimel J.P. 2000. Controls on soil carbon dioxide and methane fluxes in a variety of taiga forest stands in interior Alaska. *Ecosystems* 3: 269–282.
- Hanson G.C., Groffman P.M. & Gold A.J. 1994. Denitrification in riparian wetlands receiving high and low groundwater nitrate inputs. J. Environ. Qual. 23: 917–922.
- Hart C.H., Binkley D. & Perry D.A. 1997. Influence of red alder on soil nitrogen transformations in two conifer forests of contrasting productivity. *Soil Biol. Biochem*. 29: 1111–1123.
- Haycock N.E. & Pinay G. 1993. Groundwater nitrate dynamics in grass and poplar vegetated riparian buffers trips during the winter. J. Environ. Qual. 22: 273–278.
- Hefting M.M. & de Klein J.J.M. 1998. Nitrogen removal in buffer strips along a lowland stream in the Netherlands: a pilot study. *Environ. Pollut*. 102(S1): 521–526.
- Hefting M.M., Bobbink R. & de Caluwe H. 2003. Nitrous oxide emission and denitrification in chronically nitrateloaded riparian buffer zones. *J. Environ. Qual.* 32: 1194–1203.
- Hutchinson G.L. & Livingston G.P. 1993. Use of chamber systems to measure trace gas fluxes. In: Harper L.E., Mosier A.R., Duxbury J.M. & Rolston D.E. (eds.), Agricultural ecosystems effects on trace gases and global climate change, ASA Special Publication No. 55. American Society of Agronomy, Madison, MI, USA, pp. 1–55.
- IPCC 2001. Atmospheric chemistry and greenhouse gases. In: Houghton J.T., Ding Y., Griggs D.J., Noguer M., van der Linden P.J., Dai X., Maskell K. & Johnson C.A. (eds.), Climate change 2001. The scientific basis, Cambridge University Press, Cambridge and New York, pp. 239–287.
- Jacinthe P.A., Groffman P.M., Gold A.J. & Mosier A. 1998.
  Patchiness in microbial nitrogen transformations in groundwater in a riparian forest. *J. Environ. Qual.* 27: 156–164.
- Jones J.B., Holmes R.M., Fisher S.G., Grimm N.B. & Greene D.M. 1995. Methanogenesis in Arizona, USA dryland streams. *Biogeochemistry* 31: 155–173.
- Jones J.B. & Mulholland P.J. 1998. Methane input and evasion in a hardwood forest stream: effects of subsurface flow from shallow and deep pathways. *Limnol. Ocea*nogr. 43: 1243–1250.
- Knowles R. 1982. Denitrification. *Microbiol. Rev.* 46: 43–70.
  Kuusemets V. & Mander Ü. 1999. Ecotechnological measures to control nutrient losses from catchments. *Water Sci. Technol.* 40: 195–202.
- Kuusemets V., Mander Ü., Lõhmus K. & Ivask M. 2001. Nitrogen and phosphorus variation in shallow groundwater and assimilation in plants in complex riparian buffer zones. Water Sci. Technol. 44: 615–622.

- Larmola T., Alm J., Juutinen S., Martikainen P.J. & Silvola J. 2003. Ecosystem CO<sub>2</sub> exchange and plant biomass in the littoral zone of a boreal eutrophic lake. *Freshwater Biol*. 48: 1295–1310.
- Lawrance R.R., Todd R.L. & Asmussen L.E. 1983. Waterborne nutrient budgets for the riparian zone of an agricultural watershed. *Agric. Ecosyst. Environ.* 10: 371–384.
- Loftfield N., Flessa H., Augustin J. & Beese F. 1997. Automated gas chromatographic system for rapid analysis of the atmospheric trace gases methane, carbon dioxide, and nitrous oxide. *J. Environ. Qual.* 26: 560–564.
- Lõhmus K., Kuusemets V., Ivask M., Teiter S., Augustin J. & Mander Ü. 2002. Budgets of nitrogen fluxes in riparian grey alder forests. Arch. Hydrobiol. 141 Suppl. Large Rivers 13: 321–332.
- Mander Ü., Kull A., Kuusemets V. & Tamm T. 2000. Nutrient runoff dynamics in a rural catchment: Influence of landuse changes, climatic fluctuations and ecotechnological measures. *Ecol. Eng.* 14: 405–417.
- Mander Ü., Kuusemets V. & Ivask M. 1995. Nutrient dynamics of riparian ecotones: a case study from the Porijõgi River catchment, Estonia. *Landscape Urban Plan*. 31: 333–348.
- Mander Ü., Kuusemets V., Lõhmus K. & Mauring T. 1997a.Efficiency and dimensioning of riparian buffer zones in agricultural catchments. *Ecol. Eng.* 8: 299–324.
- Mander Ü., Lõhmus K., Kuusemets V. & Ivask M. 1997b.
  The potential role of wet meadows and grey alder forests as buffer zones. In: Haycock N.E., Burt T.P., Goulding K.W.T. & Pinay G. (eds.), Buffer zones: their processes and potential in water protection, Quest Environmental, Foundation for Water Research, Oxford, UK, pp. 35–46.
- Mander Ü., Kuusemets V., Lõhmus K., Mauring T., Teiter S. & Augustin J. 2003. Nitrous oxide, dinitrogen and methane emission in a subsurface flow constructed wetland. Water Sci. Technol. 48: 135–142.
- Mander Ü., Löhmus K., Kuusemets V., Ivask M., Teiter S. & Augustin J. 2005. Budgets of nitrogen and phosphorus fluxes in riparian grey alder forest. In: Vymazal J. (ed.), Natural and constructed wetlands: nutrients, metals and management. Backhuys Publishers, Leiden, pp. 1–19.
- Martikainen P.J., Nykänen H., Crill P. & Silvola J. 1993.
  Effect of a lowered water table on nitrous oxide fluxes from northern peatlands. *Nature* 366: 51–53.
- Mitsch W.J. & Gosselink J.G. 1993. *Wetlands*. Van Nostrand Reinhold, New York.
- Mogge B., Kaiser E.-A. & Munch J.-C. 1998. Nitrous oxide emissions and denitrification N-losses from forest soils in the Bornhöved lake region (northern Germany). *Soil Biol. Biochem.* 30: 703–710.
- Mookherji S., McCarty G.W. & Angier J.T. 2003. Dissolved gas analysis for assessing the fate of nitrate in wetlands. *J. Am. Water Resour. As.* 39: 381–387.
- Mosier A.R. 1998. Soil processes and global changes. *Biol. Fertil. Soils* 27: 221–229.
- Nõmmik H. 1956. Investigations on denitrification in soil. *Acta Agric. Scand.* 6: 195–228.
- Peterjohn W.T. & Correll D.L. 1984. Nutrient dynamics in an agricultural watershed: observations on the role of a riparian forest. *Ecology* 65: 1466–1475.

- Reay D.S., Nedwell D.B., McNamara N. & Ineson P. 2005. Effect of tree species on methane and ammonium oxidation in forest soils. *Soil Biol. Biochem.* 37: 719–730.
- Reay D.S., Radajewski J.C., Murrell J.C., McNamara N. & Nedwell D.B. 2001. Effects of land-use on the activity and diversity of methane oxidizing bacteria in forest soils. Soil Biol. Biochem. 33: 1613–1623.
- Robertson G.P. & Tiedje J.M. 1984. Denitrification and nitrous oxide production in successional and old-growth Michigan forest. Soil Sci. Soc. Am. J. 48: 383–389.
- Rusch H. & Rennenberg H. 1998. Black alder (Alnus glutinosa (L.) Gaertn.) trees mediate methane and nitrous oxide emission from the soil to the atmosphere. Plant Soil 201: 1–7.
- Rytter L., Slapokas T. & Granhall U. 1989. Woody biomass and litter production of fertilized grey alder plantations on a low-humified peatbog. *Forest Ecol. Manag.* 28: 161–176.
- Saarnio S., Alm J., Silvola J., Lohila A., Nykänen H. & Martikainen P.J. 1997. Seasonal variation in CH<sub>4</sub> emissions and production and oxidation potentials at microsites on an oligotrophic pine fen. *Oecologia* 110: 414–422.
- Scholefield D., Hawkins J.M.B. & Jackson S.M. 1997. Development of a helium atmosphere soil incubation technique for direct measurement of nitrous oxide and dinitrogen fluxes during denitrification. Soil Biol. Biochem. 29: 1345–1352.
- Scott R.L., Edwards E.A., Shuttleworth W.J., Huxman T.E., Watts C. & Goodrich D.C. 2004. Interannual and seasonal variation in fluxes of water and carbon dioxide from a riparian woodland ecosystem. Agr. Forest Meteorol. 122: 65–84.
- Stålnacke P., Grimvall A., Libiseller C., Laznik A. & Kokorite I. 2003. Trends in nutrient concentrations in Latvian rivers and the response to the dramatic change in agriculture. *J. Hydrol.* 283: 184–205.
- Teiter S. & Mander Ü. 2005. Emission of N<sub>2</sub>O, N<sub>2</sub>, CH<sub>4</sub> and

- CO<sub>2</sub> from constructed wetlands for wastewater treatment and from riparian buffer zones. *Ecol. Eng.* 25: 528–541.
- Tufekcioglu A., Raich J.W., Isenhart T.M. & Schultz R.C. 2001. Soil respiration within riparian buffers and adjacent crop fields. *Plant Soil* 229: 117–124.
- Uri V., Lõhmus K. & Tullus H. 2003a. Annual net nitrogen mineralization in a grey alder (*Alnus incana* (L.) Moench) plantation on abandoned agricultural land. Forest Ecol. Manage. 184: 167–176.
- Uri V., Tullus H. & Lõhmus K. 2003b. Nutrient allocation, allocation and above-ground biomass in grey alder and hybrid alder plantations. Silva Fennica 37: 301–311.
- von Arnold K., Nilsson M., Hånell, B., Weslien P. & Klemedtsson L. 2005. Fluxes of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O from drained organic soils in deciduous forests. *Soil Biol. Biochem.* 37: 105–1071.
- Vought L.B.M., Dahl J., Pedersen C.L. & Lacoursiere J.O. 1994. Nutrient retention in riparian ecotones. Ambio 23: 342–348.
- Watts S. & Seitzinger S.P. 2000. Denitrification rates in organic and mineral soils from riparian sites: a comparison of N<sub>2</sub> flux and acetylene inhibition methods. Soil Biol. Biochem. 32: 1383–1392.
- Weller D.E., Correll D.L. & Jordan T.E. 1994. Denitrification in riparian forests receiving agricultural discharges. In: Mitsch W.J. (ed.), Global wetlands: old world and new, Elsevier, New York, USA, pp. 117–131.
- Whiting G.J. & Chanton J.P. 2001. Greenhouse carbon balance of wetlands: methane emission versus carbon sequestration. *Tellus B* 53: 521–528.
- Willison T.W., Baker J.C. & Murphy D.V. 1998. Methane fluxes and nitrogen dynamics from a drained fenland peat. *Biol. Fertil. Soils* 27: 279–283.
- Zou X.M., Binkley D. & Caldwell B.A. 1995. Effects of dinitrogen-fixing trees on phosphorus biogeochemical cycling in contrasting forests. Soil Sci. Soc. Am. J. 59: 1452–1458.