Pinus sylvestris as a missing source of nitrous oxide and methane in boreal forest

Katerina Machacova1, Jaana Bäck3, Anni Vanhatalo3, Elisa Halmeenmäki2, Pasi Kolari2, Ivan Mammarella2, Jukka Pumpanen4, Manuel Acosta1, Otmar Urban1 & Mari Pihlatie2,5

Boreal forests comprise 73% of the world’s coniferous forests. Based on forest floor measurements, they have been considered a significant natural sink of methane (CH4) and a natural source of nitrous oxide (N2O), both of which are important greenhouse gases. However, the role of trees, especially conifers, in ecosystem N2O and CH4 exchange is only poorly understood. We show for the first time that mature Scots pine (Pinus sylvestris L.) trees consistently emit N2O and CH4 from both stems and shoots. The shoot fluxes of N2O and CH4 exceeded the stem flux rates by 16 and 41 times, respectively. Moreover, higher stem N2O and CH4 fluxes were observed from wet than from dry areas of the forest. The N2O release from boreal pine forests may thus be underestimated and the uptake of CH4 may be overestimated when ecosystem flux calculations are based solely on forest floor measurements. The contribution of pine trees to the N2O and CH4 exchange of the boreal pine forest seems to increase considerably under high soil water content, thus highlighting the urgent need to include tree-emissions in greenhouse gas emission inventories.

Methane (CH4) and nitrous oxide (N2O) are naturally produced in soils. The net CH4 and N2O flux at the soil–atmosphere interface is a balance of gas production, consumption and transport processes within soil (Supplementary Fig. S1). CH4 is produced by anaerobic methanogenesis1 in water saturated soils and oxidized by methanotrophic bacteria2. N2O is mainly formed during denitrification, anaerobic dissimilatory nitrate reduction to ammonium, and aerobic nitrification1. Denitrification is the only process consuming N2O by reduction to N2.

In addition to gas diffusion at the soil surface and ebullition1, it has been shown that plant-mediated transport3–13 can contribute significantly to CH4 and N2O exchange between the pedosphere and the atmosphere (Supplementary Fig. S1). CH4 and N2O produced in the soil can be taken up by roots, diffuse across root cortex3,6, and be transported into the above-ground plant tissues. This transport occurs via intercellular spaces and the aerenchyma system3,5,6,7,11 and/or in xylem via the transpiration stream4,5,8,13. Release of CH4 and N2O into the atmosphere takes place via lenticels or stomata3,6,11,13. Both gases may also be formed in plants, either by microorganisms living within the plant14–16 or by physiological and photochemical processes17–19.

In recent decades, N2O and CH4 fluxes from plants have predominantly been investigated in herbaceous plants from wetlands. Studies in trees are rather rare and restricted mostly to stem flux measurements on wetland species. Particularly, those upland tree species lacking an aerenchyma system have been poorly investigated8,9,13,20. This is despite the fact that upland soils seem to be an important natural source of N2O21 and a strong natural sink of CH422. Moreover, the current flux estimates of N2O and CH4 from forest ecosystems are based mostly on measurements from the forest floor, excluding the contribution of trees.

We quantified N2O and CH4 fluxes from stems, shoots (i.e. terminal branches of ca 15–20 cm length in upper canopy), and the forest floor of boreal forest dominated by Scots pine (Pinus sylvestris L.). We also investigated whether soil moisture level affects the N2O and CH4 exchange from trees and forest floor. This study is unique for its simultaneous determination of stem, shoot, and forest floor fluxes. Data were collected during May to July 2013 in a 50-year-old P. sylvestris stand23 in Southern Finland on two experimental plots (dimensions of...
20 × 15 m, a distance of 100 m apart) with naturally differing soil volumetric water content (VWC): dry plot with 0.33 ± 0.030 m³ m⁻³, wet plot with 0.75 ± 0.016 m³ m⁻³ (mean ± standard error).

Results and Discussion

N₂O fluxes. In dry field conditions typical for the studied boreal forest, we observed that P. sylvestris stems and shoots emitted N₂O at rates (medians) of 0.023 and 0.097 μg N₂O per m² of stem and projected leaf area, respectively, per hour (Supplementary Fig. S2a), accounting for 0.11 and 1.9 mg N₂O, respectively, after scaling up per hectare of ground area per hour (see Methods, Fig. 1a). To our knowledge, measurements of shoot fluxes of N₂O from mature trees have never been reported, and most studies assume negligible shoot emissions compared to stem fluxes⁵,⁹,¹⁰,¹²,¹³. Contrary to this current understanding, the shoot fluxes of N₂O from the studied pine trees exceeded the stem fluxes by more than 16 times. This underlines the important role of forest canopies in trace gas exchange. The N₂O fluxes from pine trees were accompanied by forest floor flux rates reaching 2.50 μg N₂O m⁻² h⁻¹ (24.9 mg N₂O ha⁻¹ h⁻¹; Supplementary Fig. S2a, Fig. 1a), which agrees with previous soil N₂O measurements in the same forest²⁴. In general, boreal forest soils are characterized by low availability of mineral N²³,²⁵ and low N deposition²³, resulting in low soil N₂O emissions, particularly when compared to 4 to 12 times higher emissions from temperate and tropical forests²⁵.

The up-scaled N₂O emission rates from trees, assuming the mean tree constitution and density of 1000 trees per hectare in the dry plot (see Methods), were equivalent to 8.0% of the forest floor emissions per hectare (Fig. 1a, comparison of medians). Thus, the N₂O emissions from trees constitute a significant part of the boreal pine forest N₂O flux. The N₂O flux from dry areas of the studied forest, including the contribution from forest floor and pine trees, reached approximately 26.9 mg N₂O ha⁻¹ h⁻¹ (8.0 g CO₂-e ha⁻¹ h⁻¹ using a global warming potential [GWP] of 298 [ref. 26]), which lies within the range of the global inventory estimates of N₂O flux rates for boreal forests²⁵. Based on the shoot-to-stem N₂O fluxes ratio of 16 at the dry area, the shoot fluxes at the wet plot could reach 3.3 mg N₂O ha⁻¹ h⁻¹ versus the measured stem fluxes of 0.20 mg N₂O ha⁻¹ h⁻¹. As follows, under high soil water content typical for studied wet areas of the forest with density of 1400 trees per hectare, the contribution of pine trees could be up to 18% (based on medians comparison) of the forest floor N₂O exchange.

The up-scaled N₂O emission rates from trees, assuming the mean tree constitution and density of 1000 trees per hectare in the dry plot (see Methods), were equivalent to 8.0% of the forest floor emissions per hectare (Fig. 1a, comparison of medians). Thus, the N₂O emissions from trees constitute a significant part of the boreal pine forest N₂O flux. The N₂O flux from dry areas of the studied forest, including the contribution from forest floor and pine trees, reached approximately 26.9 mg N₂O ha⁻¹ h⁻¹ (8.0 g CO₂-e ha⁻¹ h⁻¹ using a global warming potential [GWP] of 298 [ref. 26]), which lies within the range of the global inventory estimates of N₂O flux rates for boreal forests²⁵. Based on the shoot-to-stem N₂O fluxes ratio of 16 at the dry area, the shoot fluxes at the wet plot could reach 3.3 mg N₂O ha⁻¹ h⁻¹ versus the measured stem fluxes of 0.20 mg N₂O ha⁻¹ h⁻¹. As follows, under high soil water content typical for studied wet areas of the forest with density of 1400 trees per hectare, the contribution of pine trees could be up to 18% (based on medians comparison) of the forest floor N₂O exchange.

N₂O/CH₄ exchange between individual pine trees as well as between different tree species remain unknown due to lack of canopy flux measurements. We estimated that temporal variability in the shoot, stem and forest floor exchange.

N₂O and CH₄ exchange are expressed as percentage of the forest floor flux. Statistically significant differences at p < 0.017 (multiple comparison – Bonferroni correction) between flux components are indicated by different capital letters above bars.
Nitrous oxide (N₂O) and methane (CH₄) fluxes were higher than the spatial variability in the dry plot, whereas in the wet plot the spatial variability dominated the fluxes. This indicates that most of the variability in the fluxes in the dominating dry areas originates from day-to-day variation, whereas the fluxes in the wet areas, which form a minority of the forest, are dominated by high small-scale variation.

The pine stem N₂O fluxes correlated positively with forest floor fluxes (Spearman’s rank correlation coefficient: $p = 0.351$, $p < 0.05$), indicating that the tree-emitted N₂O could originate from soil. As N₂O is rather water soluble, and many plant species emit N₂O irrespective of the presence of a rhizosphere system, we hypothesize that N₂O is absorbed by roots from the soil, transported via xylem into the above-ground tree parts, and then emitted into the atmosphere.

CH₄ fluxes. Contrary to the CH₄ uptake by shoots (i.e. negative flux) found in Scots pine seedlings grown under field and laboratory conditions, we observed emissions of CH₄ from both shoots and stems of mature *P. sylvestris*. This difference in shoot CH₄ fluxes may result from (i) different soil water content and soil temperature (not reported for the seedlings experiment), (ii) known discrepancy in emission capacities of young and mature trees, and (iii) the fact that the seedlings were investigated in the absence of UV radiation, which is known to stimulate CH₄ formation. The CH₄ emission rates from pine stems and shoots were 0.005 and 0.050 μg CH₄ m⁻² h⁻¹ (medians), respectively (Supplementary Fig. S2b). Up-scaling emission rates at stand level were 0.03 and 1.1 mg CH₄ ha⁻¹ h⁻¹ (Fig. 1b) assuming mean tree constitution and density of 1000 trees per hectare (see Methods). As is the case of N₂O, pine shoots seem to be the primary tree surface emitting CH₄ into the atmosphere, given that shoot fluxes were 41 times higher than the stem fluxes. This contradicts the common assumption that basal regions of stems are the main source of CH₄ and N₂O from trees. Whereas trees were a source of CH₄, the forest floor was a sink (−14.4 μg CH₄ m⁻² h⁻¹, Supplementary Fig. S2b; −143 mg CH₄ ha⁻¹ h⁻¹, Fig. 1b). The estimated average pine tree CH₄ emission represented 0.8% of the forest floor uptake. The CH₄ uptake from the dry area of the studied forest (−4.9 g CO₂-e ha⁻¹ h⁻¹ using GWP of 34 [ref. 26]) is roughly 35% to 50% lower than are estimates of CH₄ uptake for boreal forests in global inventories.

The median stem CH₄ fluxes at the wet plot (0.100 μg CH₄ m⁻² h⁻¹) were one order of magnitude higher than those at the dry plot (0.013 μg CH₄ m⁻² h⁻¹) (Fig. 2b), while the soil remained a sink for CH₄ even under high soil water content (−7.09 μg CH₄ m⁻² h⁻¹, −70.7 mg CH₄ ha⁻¹ h⁻¹; Fig. 2d). Moreover, the stem-to-forest-floor CH₄ fluxes ratio increased with soil water content, underlining the importance of pine trees at wet areas in the balance of CH₄. Although direct measurement of shoot CH₄ flux at the wet plot was technically impossible, based on the shoot-to-stem CH₄ fluxes ratio of 41 at the dry plot, the shoot CH₄ flux at the wet plot were estimated to reach 24 mg CH₄ ha⁻¹ h⁻¹ in comparison to the stem CH₄ fluxes of 0.59 mg CH₄ ha⁻¹ h⁻¹. Under high soil water content, CH₄ emissions from pine trees could, therefore, account for up to 35% of the forest floor uptake. This estimate is rather higher than in a recent study by Pangala and colleagues, who found that CH₄ emissions mediated by *Alnus glutinosa* and *Betula pubescens* contribute up to 14% to the total CH₄ fluxes from a temperate forested wetland.

The partial soil origin of pine-emitted CH₄ is supported by strong positive correlation of stem CH₄ fluxes with forest floor CH₄ fluxes ($p = 0.716$, $p < 0.001$) and VWC in topsoil ($p = 0.802$, $p < 0.001$). In wet conditions, pine trees may therefore prevent CH₄ consumption in the upper soil layers by transporting CH₄, produced in deeper soil, into the atmosphere. We suggest that soil-produced CH₄ is transported into the above-ground parts of *P. sylvestris* mainly by the transpiration stream and then released into the atmosphere predominantly via stomata, thus explaining the higher CH₄ emissions from shoots as compared to stems. This assumption is supported by the positive correlation between the shoot CH₄ flux and transpiration ($p = 0.626$, $p < 0.05$), and stem CH₄ flux and sap flow ($p = 0.390$, $p < 0.01$). Therefore, alternative pathways, such as radial diffusion of CH₄ (and N₂O) in stems through intercellular spaces of the ray parenchyma and a release from the stem via lenticels, seem of a lesser importance.

Different mechanisms of CH₄ emissions from trees grown on dry plot as compared to those on wet plot are, however, likely. Limited shoot CH₄ production in deeper mineral soil layers and low mineral soil VWC (0.28 ± 0.02 m³ m⁻³) in the studied period give an assumption of negligible soil CH₄ production in the dry plot. Moreover, approximately half of the root system of *P. sylvestris* is located in the top soil organic layer with the rest of the roots equally distributed to mineral soil (0–40 cm). Therefore, it is probable that part of the CH₄ emitted from trees in the dry plot originated from anaerobic production processes within the wood and/or aerobic, non-microbial metabolic processes in the plant tissues.

*P. sylvestris* appears to be one of the missing sources for N₂O and CH₄ in boreal forests. N₂O emissions from boreal pine forests may previously have been underestimated and the uptake of CH₄ overestimated. Even though our measurements indicate only potential mechanisms, and more detailed measurements of spatio-temporal variability are necessary, the pine mediated N₂O and CH₄ emissions could account for up to 18% of forest floor N₂O emissions and 35% of forest floor CH₄ uptake, respectively, under high soil moisture conditions. This can be crucial for the future greenhouse gas budgets of boreal pine forests, especially if precipitation and evapotranspiration patterns will change due to climate change. Our findings highlight the important, but often neglected role of upland trees in N₂O and CH₄ exchange between the biosphere and the atmosphere and the importance of including tree emissions to the total forest ecosystem budgets of N₂O and CH₄.

**Methods**

**Site description and experimental design.** The measurements were performed in a 50-year-old stand of Scots pine (*Pinus sylvestris* L.) at the SMEAR II station (Station for Measuring Ecosystem–Atmosphere Relations) in Hyytiälä, Southern Finland (61°51′N, 24°17′E, 181 m a.s.l.) from 23 May to 19 July 2013. Established in 1962, the site is a boreal coniferous forest dominated by *P. sylvestris* with some additional Norway spruce (*Picea abies*)
and broadleaved trees in the understorey\textsuperscript{23,34}. The long-term annual mean temperature and precipitation are 3.5 °C and 711 mm, respectively\textsuperscript{35}. The soil is Haplic podzol on glacial till with irregularly distributed peat soil spots\textsuperscript{36}.

Naturally wet and dry plots (dimensions of 20 × 15 m, a distance of 100 m apart) with mean soil volumetric water content (VWC) 0.75 ± 0.016 m\(^3\) m\(^{-3}\) (mean ± standard error) and 0.33 ± 0.030 m\(^3\) m\(^{-3}\), respectively, were selected. During the measurement period, soil water content was measured using an HH2 Moisture Meter and Theta Probe (type ML2x, AT Delta-T Devices, Cambridge, UK) in A-horizon corresponding to depths 0–5 cm from the soil surface, and expressed as mean of three independent measurements close to each tree and soil chamber. Soil temperature was measured continuously by a DS1921G Maxim Thermochron iButtons (Maxim Integrated, San Jose, California, USA) in A-horizon next to each soil chamber.

On each plot, six representative trees were chosen for stem flux measurements (n = 6). Shoot fluxes were measured from the upper canopy of three trees used for the stem flux measurements at the dry plot. Shoot fluxes were not measured from the wet plot as installation of a scaffold tower was technically not possible. Forest floor CH\(_4\) and N\(_2\)O fluxes were measured at three representative positions in the dry plot and at three positions in the wet plot (n = 3). Four flux measurement campaigns, each taken over ca 2 weeks, were made for stem, shoot, and forest floor fluxes between 23 May and 19 July 2013 (for details concerning the number of replicates see the legends of Figs 1 and 2, and Supplementary Fig. S2 describing individual measuring campaigns). Simultaneous measurements of fluxes from each tree and the forest floor in its vicinity allowed a comparison of N\(_2\)O and CH\(_4\) fluxes between tree shoots, stems, and forest floor. The fluxes were determined on all measuring days at approximately same time to prevent possible variation in flux rates caused by flux diurnal cycle.

The mean tree height, length of living crown, and stem diameter at breast height (DBH) of the selected pine trees were 18.2 ± 0.4 m, 6.47 ± 0.32 m, and 0.162 ± 0.012 m, respectively, for the wet plot. For the dry plot, these were 17.7 ± 0.5 m, 7.22 ± 0.43 m, and 0.180 ± 0.004 m, respectively. These morphological parameters did not differ significantly when comparing wet and dry plots. The stand densities were estimated to be 1000 and 1400 trees

Figure 2. Stem and forest floor fluxes of N\(_2\)O (a,c) and CH\(_4\) (b,d) from dry and wet plots of boreal forest dominated by Scots pine (Pinus sylvestris). Mean volumetric water contents for dry and wet plots (± s.e.) were 0.33 ± 0.030 m\(^3\) m\(^{-3}\) and 0.75 ± 0.016 m\(^3\) m\(^{-3}\), respectively. Flux rates of all sampling points as follows are expressed as medians (solid lines) or means (broken lines) per m\(^2\) of surface area. Stem fluxes are determined from six trees per plot (n = 6; 3–6 measurement repetitions per tree), and forest floor fluxes from three soil chambers per plot (n = 3; 6–9 repetitions per chamber). The fluxes from the stems and from the forest floor were always measured simultaneously. Statistically significant differences at p < 0.05 are indicated by an asterisk.

For box plots description, see Fig. 1.
per hectare on the dry and wet plots, respectively. The stand basal area was measured directly on the plots using a rod relascope technique and was 19.5 and 26 m² ha⁻¹ on dry and wet plots, respectively.

**Chamber systems.** The stem fluxes were measured using 12 stem chambers (1 chamber per tree) enclosing the entire stem circumference and the skeleton of the stem chamber (volume between 0.0009 and 0.0015 m³ depending on stem diameter) was created by a flexible pipe from polyethylene-coated aluminium (Synflex, Eaton Hydraulics Group Europe, Morges, Switzerland), which was wrapped in a spiral around the stem. A tube-fitting brace was attached to this spiral and enabled fixation of inlet and outlet connectors. Teflon FEP film (0.05 mm thick, Fluoroplast, Maalahiti, Finland) impermeable for CH₄ and N₂O was wrapped 1.5 to 2 times around the tube spiral to create the chamber wall, and then sealed with adhesive FEP tape. Due to therequirement of mounting the stem chambers on the basal part of the rough pine bark (around 0.2 m above the forest floor), the surface of the dead outer bark was carefully removed from the upper and basal ends of the chamber. The upper and basal ends of the Teflon foil were sealed with elastic closed cell polyethylene foam and wide flexible ties to the carefully smoothed bark surface. The results of the stem flux measurements on twelve trees were used in the comparison of the stem and forest floor fluxes between dry and wet plots (Fig. 2).

Two different shoot chamber types were used to measure fluxes of CH₄ and N₂O: two cylindrical chambers with FEP foil walls (volume 0.0054 m³) and a methacrylic cylindrical shoot chamber (volume 0.005 m³). We did not observe any differences in flux rates obtained by these two types of chambers. The three chambers were installed in a follow-up study of the three trees on the dry plot. The air temperatures (DT 612 thermometer, CEM, Shenzhen, China) inside and outside of the chambers were regularly measured during chamber closures. To avoid overheating in the chambers, the shoot fluxes were measured only on cloudy days. The comparison of the shoot, stem and forest floor fluxes presented in Fig. 1 and Supplementary Fig. S2 is based on the measurements at the dry plot only. To compare the whole tree flux rates in dry and wet plots, we used the shoot-to-stem flux ratio from the dry plot where both shoot and stem flux measurements were performed.

In both stem and shoot chambers, the mixing of the air inside the chambers was provided by vacuum pumps (V 1500-GAS-12V standard vacuum pumps, Xavitech, Härnösand, Sweden; NMP 850.1.2, KNDC B, KNF Neuberger, Freiburg, Germany) gas-tightly connected to the chamber using Teflon tubes and stainless steel connectors (Swagelok, Ohio, USA). The chambers were non-steady-state flow-through chambers returning the air from the pump again into the chambers. Gas samples were taken with a syringe via a septum connected to the air circulation. Six gas samples (each 20 ml) were taken from the closed stem and shoot chambers at time intervals of 60 min over a period of 6 h. The possible under-pressure resulted from the gas sampling was compensated by the flexible foil wall. The stem chambers were flushed with ambient air for at least 30 min before sampling. Forest floor CH₄ and N₂O fluxes were measured using large opaque soil chambers made of aluminium. Three chambers were placed on the dry plot (volume of ca 0.091 m³ depending on vegetation inside the chamber), enclosed soil surface area of 0.298 m², and three on the wet plot (volume of ca 0.135 m³, soil area of 0.298 m²). The chambers were located in the vicinity of the measured trees. The ground vegetation in the soil chambers varied among chambers depending on the soil conditions and location, and consisted of *Sphagnum sp.*, *Polytrichum sp.*, *Dicranum polysetum*, *Pleurozium schreberi*, *Equisetum sylvaticum*, *Vaccinium myrtillus*, *Vaccinium vitis-idaea*, *Trientalis europaea*, and several representatives of *Poaceae*. The placement of chamber collars took place several days before the first sampling to allow the soil to settle and avoid soil disturbances. The soil chambers were closed for ca 40 min during which gas samples (each 20 ml) were taken at 2, 5, 10, 20, 30, and 40 minutes after the chamber closure. A fan was used to mix the headspace air during the closure. Chamber headspace temperature (DT 612 thermometer, CEM, China) was regularly monitored during the measurements.

**Gas analyses.** Gas samples from stem, shoot, and soil chambers were taken in 20 ml syringes (BD syringe, Franklin Lakes, New Jersey, USA) and immediately transferred to the evacuated 12 ml glass vials (Labco, Ceredigion, UK), then stored at 4 °C. The gas samples were analysed by an Agilent 7890A gas chromatograph (GC) (Agilent Technologies, Santa Clara, California, USA) equipped with a flame ionization detector (FID) and an electron capture detector (ECD) for CH₄ and N₂O analyses, respectively. Briefly, CH₄ was detected by FID (300 °C) supplied with synthetic air (450 ml min⁻¹) and hydrogen (H₂, 45 ml min⁻¹) and with nitrogen (N₂, 5 ml min⁻¹) as a make-up gas. Helium (He, 45 ml min⁻¹) was used in both cases as a carrier gas. Columns Porapak Q 80–100 Mesh and Hayesep Q 80–100 Mesh (Agilent Technologies, USA) were used for water vapour removal and gas separation. Oven temperature was kept at 60 °C. Retention times for CH₄ and N₂O were 3.6 and 4.3 min, respectively. The gas samples were automatically injected by an autosampler Gilson GX-271 Liquid Handler (Gilson, Middleton, Wisconsin, USA). An overpressure in vials was necessary for proper injection of gas samples and was an indicator of gas tightness of the vials. ChemStation B.03.02 software was used for the GC analyses.

The identification of CH₄ and N₂O peaks in gas samples and calculation of their molar fractions referred to dry air (hereinafter: “concentrations”) were performed using a four-point standard curve with the following concentrations: CH₄ (1.207, 1.810, 2.413, 3.017 ppm of CH₄ in synthetic air), N₂O (0.279, 0.330, 0.381, 0.457 ppm of N₂O in synthetic air). The four standards were analysed at the beginning of the analyses and after every ca 30 gas samples. A running standard (1.810 ppm CH₄, 0.330 ppm N₂O; in synthetic air) for detailed control was applied after every ca 15 gas samples.

**Calculation of N₂O and CH₄ flux rates.** The flux rates of N₂O and CH₄ from stems, shoots, and forest floor were calculated by linear least square fits of time series of N₂O and CH₄ concentrations as follows

\[ F = \frac{V}{A} \frac{M_{273.16}}{V_{m}} \frac{273.16}{3600} + \frac{1}{T} \]

(1)
where \( F \) is flux of \( \text{N}_2\text{O} \) or \( \text{CH}_4 \) from stem, shoot, or forest floor surface [\( \mu\text{g m}^{-2} \text{ (surface area)} \text{ h}^{-1} \)]; \( S \) is the slope of the linear fit to the \( \text{N}_2\text{O} \) or \( \text{CH}_4 \) concentrations over the chamber closure ([ppm s \(^{-1}\)]); \( V \) is volume of chamber [\( \text{m}^3 \)]; \( A \) is stem surface area, projected leaf area, or soil surface area enclosed in stem, shoot, or soil chamber, respectively [\( \text{m}^2 \)]; \( M \) is molecular mass of \( \text{N}_2\text{O} \) or \( \text{CH}_4 \) [44.01 and 16.04 g mol\(^{-1}\)], respectively; \( V_{\text{air}} \) is molar volume of an ideal gas at 1 atmosphere pressure and 25 \(^\circ\text{C} \) [0.0245 m\(^3\) mol\(^{-1}\)]; and \( T \) is the temperature [\(^\circ\text{C} \)] inside the chamber.

The flux estimates and upscaling to a stand level are saddled with uncertainties arising from sampling and gas analyses\(^{45,46} \), variables in Equation 1, and application of allometric relationships for estimation of total leaf area\(^{41} \), and from calculation of stem area per tree. At a stand level, uncertainties of flux rates are thus particularly given by spatio-temporal variability in the fluxes, heterogeneity of tree morphological parameters (height, length of living crown, stem diameter), and stand heterogeneity (tree density per hectare and species composition etc.). In addition, fast changes in transpiration and sap flow rates induced by dynamic light environment under variable sky conditions have also a potential to substantially influence the gas exchange over longer periods\(^{44} \). However, here such factors are of minor importance, as the measurements were predominantly conducted during overcast days.

Ancillary measurements. The following continuously measured variables at the SMEAR II experimental station were used for correlation analyses: a) soil water content (TDR-100, Campbell Scientific, North Logan, Utah, USA)\(^{45,46} \) and b) soil temperature (Philips KTY81, NXP, Eindhoven, Netherlands)\(^{46} \), both in four soil horizons (O-, A-, B- and C-corizon; corresponding to depths of − 4–0, 0–5, 5–23 and 23–60 cm from the mineral soil surface); c) air temperature at 4.2 m height within the forest stand (Pt100 sensors), d) photosynthetic photon flux density at 23 m height (Li-190SZ, Li-Cor, USA); on \( P. \text{sylvestris} \): e) stem sap flow using the Granier-type heat dissipation method\(^ {47,48} \) at a height of about 2 m; and f) shoot transpiration at the top canopy with dynamic enclosures\(^ {49} \).

Statistics. Datasets were tested for normal distribution (Shapiro–Wilk test) and homogeneity of variances in different subpopulations. The flux data were assumed independent. Because of non-normally distributed data and/or data with unequal variances, the non-parametric Mann–Whitney rank sum test was run at \( p < 0.05 \) to test the statistical significance a) among flux rates from stems, shoots, and forest floor; and b) between flux rates from dry and wet plots.

Correlation analyses (a) between stem, shoot, and forest floor flux rates of \( \text{N}_2\text{O} \) or \( \text{CH}_4 \), and (b) between the trace gases flux rates and micro-climatic and other tree parameters were performed using non-parametric correlation analyses (Spearman’s rank correlation). The statistical significance was defined at \( p < 0.05 \).

SigmaPlot 11.0 (Systat Software, San Jose, California, USA) was used for statistical analyses.

References

31. Hölttä, T. & Kolari, P. Interpreting the role of leaf to shoot flow data, and Marian Pavelka for providing us with part of the vacuum pumps used and for his valuable advice regarding methodology. This research was financially supported by the EU FP7 project ExpeER (Grant Agreement no. 262060), the ENVIMET project (CZ.1.07/2.3.00/20.0246), NPU I project (LO1415), the Emil Aaltonen Foundation, and Academy of Finland Research Fellow projects (263858, 288494), The Academy of Finland Centre of Excellence (project 1118615), ICOS-Finland (281255) and Helsinki University Centre for Environment, HENV1.

Acknowledgements

We would like to thank Hermanni Aaltonen for help with the gas chromatograph, Juho Aalto for help in installation of shoot chambers and calculation of stem and needle area, Teemu Paljakka for supplying us with sap flow data, and Marian Pavelka for providing us with part of the vacuum pumps used and for his valuable advice regarding methodology. This research was financially supported by the EU FP7 project ExpeER (Grant Agreement no. 262060), the ENVIMET project (CZ.1.07/2.3.00/20.0246), NPU I project (LO1415), the Emil Aaltonen Foundation, and Academy of Finland Research Fellow projects (263858, 288494), The Academy of Finland Centre of Excellence (project 1118615), ICOS-Finland (281255) and Helsinki University Centre for Environment, HENV1.

Author Contributions


Additional Information

Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: The authors declare no competing financial interests.
How to cite this article: Machacova, K. et al. *Pinus sylvestris* as a missing source of nitrous oxide and methane in boreal forest. *Sci. Rep.* 6, 23410; doi: 10.1038/srep23410 (2016).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/