

온대 낙엽 활엽수림에서의 강수량에 따른 메탄 흡수 감소

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Precipitation Decreases Methane Uptake in a Temperate Deciduous Forest

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ABSTRACT

Soil moisture regulates the fate of methane (CH₄) in forest soil via biological and chemical processes. The instant effect of variable precipitation on CH₄ uptake is, however, unclear in the forest ecosystems. Here, we measured CH₄ flux in a temperate forest soil immediately after variable volume of water applications equivalent to 10, 20 40, and 80 mm m⁻² day⁻¹ precipitation. CH₄ uptake was significantly higher when the water was not applied. The CH₄ uptake decreased significantly with increasing water application. CH₄ uptake was linked with air filled porosity and water filled porosity. CH₄ uptake response to actual precipitation intensity was in agreement with CH₄ uptake results in this study. CH₄ uptake decreased 55% at highest precipitation intensity. Since annual CH₄ flux is calculated with interpolation of weekly or biweekly field observations, instant effect of precipitation can mislead the interpolated annual results.

Key words : Methane; precipitation; intensity; temperate forest; uptake

1. Introduction

Methane (CH₄) with the atmospheric concentration of approximately 1.8 mg/L is the second most abundant greenhouse gas in the atmosphere after carbon dioxide (CO₂) (IPCC 2014). CH₄ contributes to 32% of the current radiative forcing and its global warming potential is 25 times higher than CO₂ (IPCC 2014). It explains approximately 18% of the recent increase in global temperature. Forest soils are recognized as an important sink for CH₄ (Reeburgh 2003). Approximately 9 to 42 Tg of CH₄ is oxidized in unsaturated soils worldwide per year (Kirschke et al., 2013). Temperate forest's ecosystems contribute 30-50% of total soil-based CH₄ sink worldwide (Dutaur and Verchot., 2007; Ojima et al., 1993).

CH₄ is produced by methanogens under anaerobic condition in subsoil and oxidized by methanotrophs under

aerobic condition in topsoil (Le Mer and Roger 2001). CH₄ emission from the temperate forests is linked with biological, chemical, and physical changes in soil. It is mainly controlled by organic carbon substrate, soil temperature, soil water content, and so on (Smith et al., 2003; Von Fischer and Hedin., 2007). CH₄ production depends on the availability of organic carbon for methanogens, which is produced under anaerobic decomposition of organic matter such as plant biomass, leaf litter, and fine roots in soil (Dalal et al., 2008). Soil temperature controls microbial growth rate and subsequently CH₄ emission. Thirty degrees in Celcius is reported optimal temperature for microbial activity in soil (Gütlein et al., 2017; Moore et al., 2018). Soil water content in the forest soil controls diffusive transport of CH₄ in soil (Wei et al., 2018).

Soil submersion in water triggers methanogenic activity

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due to the formation of an anaerobic condition and it decreases methanotrophic activity by reducing the oxidized zone. CH_4 oxidation occurs at 20 to 60% of soil water content in dry season and it decreases at higher than 60% of soil water content in rainy season (Castro et al., 1995). Temperate forest regions with lower precipitation are known for CH_4 oxidation (Castro et al., 1995). CH_4 oxidation is suppressed in wet summer due to the inhibition of oxygen diffusion and CH_4 production in anoxic microsites (Itoh et al., 2009). Forest soils can emit CH_4 in wet summer (e.g. Keller and Reiners., 1994; Weitz et al., 1999; Davidson et al., 2004; Vasconcelos et al., 2004; Teh et al., 2005). CH_4 dynamics in forest soils may differ in regions with heavy summer precipitation. Itoh et al. (2009) reported CH_4 oxidation $-0.45 \text{ kg ha}^{-1} \text{ y}^{-1}$ in a dry season and CH_4 emission $1.80 \text{ kg ha}^{-1} \text{ y}^{-1}$ in a rainy season. Wetting of dry soils generally increases the microbial activity within minutes (Borken et al., 2003; Lee et al., 2004; Sponseller 2007) or hours (Pulleman & Tietema., 1999; Prieme & Christensen., 2001). Diffusion of CH_4 from the atmosphere into soil usually explained with Fick's first law (Ishizuka et al., 2000; Nakano et al., 2004; Wang et al., 2014). Soil water content controls CH_4 uptake by regulating CH_4 diffusion from the atmosphere into mineral soils (Castro et al., 1994; Czepiel et al., 1995; Whalen and Reeburgh., 1996). Soil wetting and drying experiments revealed significant reduction in CH_4 uptake with wetting (Kim et al., 2012). Kessavalou et al., (1998) reported that CH_4 uptake declined by about 60% after rewetting of dry soil. To best of our knowledge instant effect of variable intensity of precipitation on CH_4 uptake has not been reported. Moreover, CH_4 fluxes in forest soils are monitored weekly or biweekly using manually closed chamber method and then results are interpolated to estimate annual fluxes. Instant change in CH_4 flux due to precipitation may mislead the total annual CH_4 flux. Precipitation varies throughout a year and this variation affects soil moisture, which thereby affects CH_4 uptake or emission. The objectives of this study were to investigate the instant effect of variable precipitation on CH_4 uptake and to estimate the contribution of precipitation in reducing net CH_4 uptake in temperate forest. We hypothesized that CH_4 emission will occur when it starts to rain because rain water will replace CH_4 present in

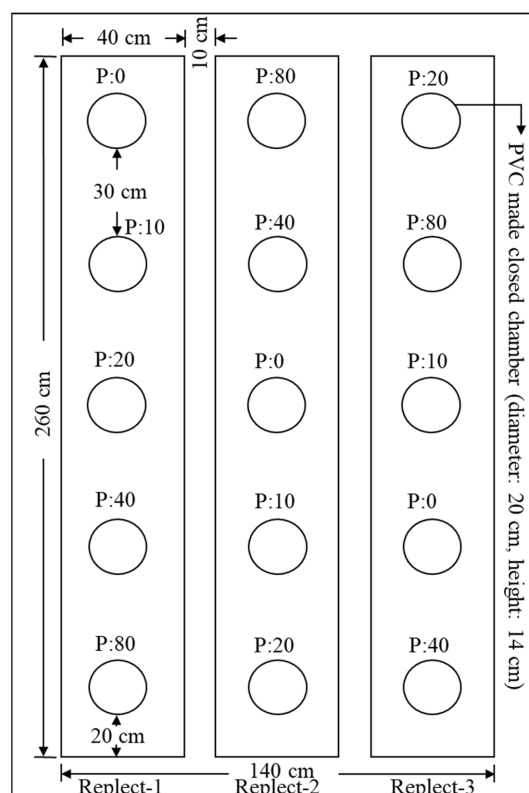


Fig. 1. On field experimental treatments to determine the effect of variable precipitation on CH_4 uptake.

subsoil. CH_4 uptake may decrease after precipitation due to water-filled pore space and in result limited space for atmospheric CH_4 uptake.

2. Materials and Methods

The experiment was conducted in a mature *Platanus occidentalis* forest on Hanyang University campus, Seoul, Republic of Korea ($37^{\circ}33'33''\text{N}$, $127^{\circ}02'47''\text{E}$). The soil texture was sandy loam with sand, silt, and clay proportions of 55.1, 33.8, and 11.1%, respectively. Daily temperature and precipitation varied between -18.6 to 36.7°C and 0.1 to 260 mm , respectively (Korea Metrological Administration 2010-2017).

Three experimental plots were located as shown in Fig. 1. Each plot was $40 \times 260 \text{ cm}$ and distance between two adjacent plots was 10 cm . Each plot comprised with five treatments such as P-0, P-10, P-20, P-40, and P-80, where (P) is precipitation and the number followed by P is amount of the water equivalent to precipitation (mm day^{-1}). The

water 0.34, 0.67, 1.35, and 2.69 L was sprayed in P-10, P-20, P-40, and P-80, respectively. Water was sprayed inside the chamber bases on alternate gas sampling days. When water was not sprayed we assume no precipitation (NP), henceforth mentioned as (NP-0, NP-10, NP-20, NP-40, and NP-80). To minimize soil disturbance, one plot was exclusively dedicated for soil sampling and remaining two plots were used for gas sampling. P-0 was used as control treatment and water was not sprayed in this treatment. The volume of water for corresponding precipitation that was calculated by using guidelines of food and agriculture organization (Dastane 1978). The volume of water used in this experiment was within the range of average daily precipitation 0.1 to 260 mm in 2010-2017 (Korea Metrological Administration 2010-2017). The volume of water corresponded to precipitation below 10 mm was too low to spray on given surface area of chamber. Precipitation above 80 mm was much higher than the volume of closed chamber above ground. Therefore, treatments for precipitation below 10 mm and above 80 mm were not installed were not installed.

Five polyvinyl chloride (PVC) chamber bases of (20 cm diameter and 20 cm height) were randomly inserted 5 cm into the ground in each plot. An air-tight lid made of PVC was kept on the chamber base for one hour and a 30 mL gas sample was collected from the chamber at 0, 15, 30, 45 and 60 min after chamber closure. All gas samples were stored in 25 mL glass vials sealed with aluminum caps and gray butyl septa. Samplings were conducted between 09:00 to 10:00 between 14th September to 15th October in 2018 every third day. Gas samples were analyzed using a gas chromatograph (YL 6100, Young Lin Instrument Co., Korea) equipped with a flame ionization detector and GS-Alumina Agilent column (length, 50 m; inner diameter, 0.53 mm). The temperatures of the column, injector, and detector were 120, 250, and 250°C, respectively. Helium was used as the carrier gas at a flow rate of 30 ml min⁻¹.

Hourly CH₄ flux was calculated from the change in gas concentration over 60 min chamber closure for first experiment and 30 min closure for second experiment (Rolston 1986):

$$F = \frac{V}{A} \times \frac{dc}{dt} \times \left(\frac{273}{273 + T} \right) \quad (1)$$

where F is the hourly CH₄ flux (μg m⁻² h⁻¹), V is the gas volume (m³), A is the area of the chamber base (m²), and $\frac{dc}{dt}$ is the rate of CH₄ concentration change over a 60 min period in the chamber (μg m⁻³ h⁻¹).

Temperatures of ambient air, the air inside the chamber, and the soil were recorded at the time of CH₄ sampling. Soil temperature and water content were monitored at 10, 20, and 30 cm depth of one plot on each sampling day. Soil samples were collected inside the chambers using a sampling tube with 2.5 cm internal diameter and 100 cm height. Soil gravimetric water content (θ_g) was determined using the oven drying method at the controlled temperature of 105°C for 24 h. Bulk density (ρ_b) of soil was measured before and after the experiment at 10, 20 and 30 cm depth using core sampler. Soil samples for bulk density were collected outside and inside of each chamber before and after experiment, respectively. The volumetric water content (θ_v) was calculated as:

$$\theta_v = \rho_b \times \theta_g \quad (2)$$

Volumetric water content was converted into absolute air-filled porosity (AFP, cm³ cm⁻³) knowing the bulk density (ρ_b) and the particle density of soil (ρ_s) with the equation (Epron et al., 2016):

$$AFP = (1 - \rho_b / \rho_s) - \theta_v \quad (3)$$

Soil particle density (ρ_s) was assumed 2.65 g cm⁻³ of rock, sand grains and other soil mineral particles (Gao et al., 2018; Zhu et al., 2013). The water-filled pore space (WFPS) was calculated with the equation (Gao et al., 2018):

$$WFPS = \theta_v / (1 - \rho_b / 2.65) \quad (4)$$

Both AFP and WFPS were then converted in percent by multiplying with 100.

2.1. Statistical analysis

The SPSS 20 statistical software package was used for statistical analysis. Independent-sample t-test was used to test the significant difference between control and litter P-(0-80) and NP-(0-80) treatments. One-way ANOVA was used to test the significant difference between the results of CH₄ emission in all treatments of P-(0-80) and NP-(0-80). The difference level was set at p<0.05. linear regression

analysis was performed to establish correlation between CH_4 uptake and (soil moisture content, soil temperature, AFP, and WFPS).

3. Results and discussion

Average CH_4 uptake in the entire experimental period was 30.6, 8.3, 5.6, 5.5, and 4.4 $\mu\text{g m}^{-2} \text{h}^{-1}$ in P-0, P-10, P-20, P-40, and P-80, respectively. Average CH_4 uptake 29.3, 42.5, 44.4, 26.2, and 21.7 was observed in NP-0, NP-10, NP-20, NP-40, and NP-80, respectively (Fig. 2a). Average CH_4 uptake in P-0, P-10, P-20, P-40, and P-80 was 5, 80, 87, 79, and 80%, respectively, lower than NP-0, NP-10, NP-20, NP-40, and NP-80, respectively. Average CH_4 uptake in P-(10-80) was significantly lower than NP-(10-80) treatments ($p=0.05$). No significant difference was observed in control treatments P-0 and NP-0 ($p=0.05$). In all treatments, soil temperature decreased consistently throughout the experimental period (Fig. 2b). Maximal and minimal soil temperature was observed on September and October, respectively. Average soil temperature 17.9, 18.2, 18.5, 19.7, and 20.7°C was observed in P-0, P-10, P-20, P-40, and P-80, respectively. Relatively low temperature 16.4, 16.2, 16.1, 16.3, and 16.6 °C was observed in NP-0, NP-10, NP-20, NP-40, and NP-80, respectively. Soil temperature was not significantly different among the treatments in both P (0-80) and NP (0-80) ($p=0.05$). Soil temperature was positively correlated with CH_4 uptake in P-0, P-10, P-20, P-40, and P-80 ($R^2=0.14, 0.49, 0.17, 0.44$, and 0.12 , respectively). Soil temperature was also positively correlated with CH_4 uptake in NP-0, NP-10, NP-20, NP-40, and NP-80 ($R^2=0.74, 0.65, 0.57, 0.03$, and 0.29 , respectively). Average soil water content in entire experimental period was 20.8, 26.2, 24.4, 24.4, and 27.2% in P-0, P-10, P-20, P-40, and P-80, respectively. Average soil water content in NP-0, NP-10, NP-20, NP-40, and NP-80 was 22.3, 21.7, 20.5, 20.7, and 24.5, respectively. Statistically there was no significant difference in P (0-80) and NP (0-80) ($p=0.05$). Average soil water content was positively correlated with average CH_4 uptake in P-0, P-10, P-20, P-40, and P-80 ($R^2=0.84, 0.61, 0.40, 0.64$, and 0.13 , respectively). Average soil water content was also positively correlated with average CH_4 uptake in NP-0, NP-10, NP-20, NP-40, and NP-80 ($R^2=$

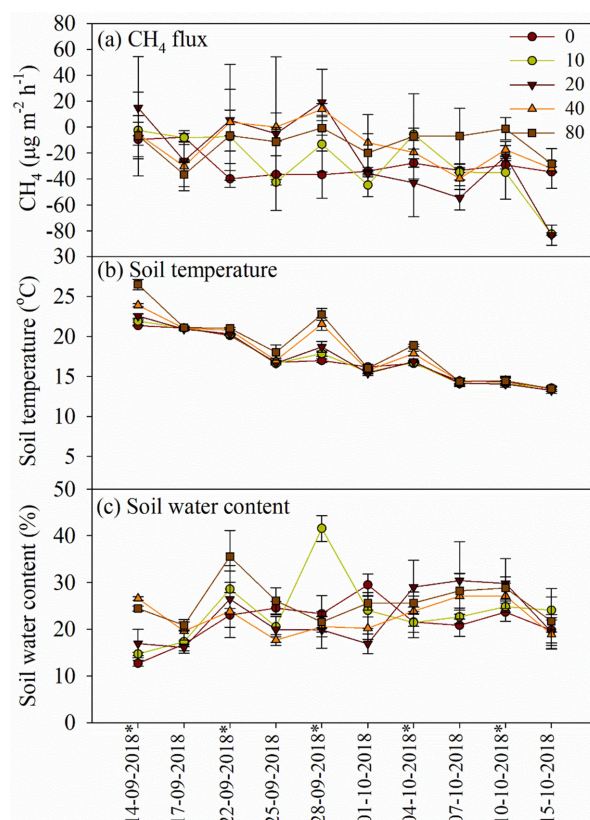


Fig. 2. (a), CH_4 flux; (b), soil temperature; and (c), soil water content with variable precipitation 0, 10, 20, 40, and 80 mm per day. Error bars represent ± 1 standard error of mean. *water was applied on these dates.

0.40, 0.64, 0.10, 0.40, and 0.96, respectively).

Average soil air filled porosity in 0-10 cm soil depth was 48.1, 43.0, 36.2, 37.9, and 33.9%, in P-0, P-10, P-20, P-40, and P-80, respectively. AFP in 10-20 cm soil depth was 39.0, 23.3, 31.2, 25.9, and 21.2%, in P-0, P-10, P-20, P-40, and P-80, respectively. AFP in 20-30 cm soil depth was 40.7, 31.5, 33.6, 25.8, and 23.4%, in P-0, P-10, P-20, P-40, and P-80, respectively. Average AFP in all soil depths (0-30 cm) of P-10 treatment was not significantly different from P-0 ($p=0.05$). Average AFP in all soil depths (0-30 cm) of P-20, P-40, and P-80 treatment was significantly different from P-0 ($p=0.05$). CH_4 uptake was positively correlated with AFP in all soil depths (0-30 cm) and all treatments (Fig. 3). CH_4 uptake decrease significantly in P-80 due to lowest AFP. CH_4 uptake was weakly correlated with AFP (P-80) in all soil depths (0-30 cm). Average AFP in soil depths 0-10, 10-20, and 20-30 cm in NP-0, NP-10, NP-20, NP-40, and NP-80 treatments was (50.2, 46.6, 41.78, 39.5,

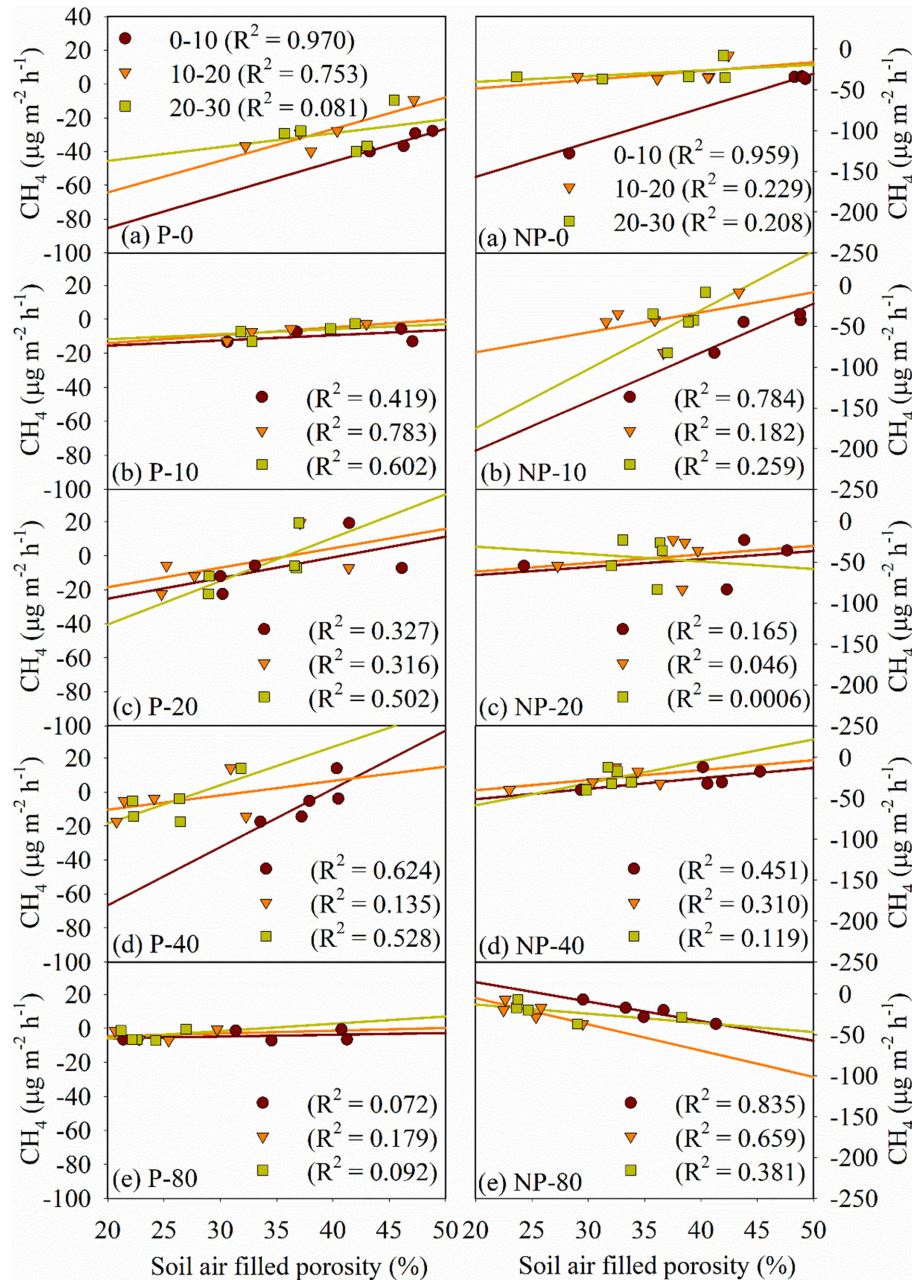


Fig. 3. Relationships (a-e, P (0-80) and a-e, NP (0-80)) between CH₄ emission and soil air filled porosity in different soil depths (0-10, 10-20, and 20-30 cm).

and 35.2%), (37.8, 36.0, 36.3, 31.3, 25.2%), and (35.6, 38.3, 34.8, 32.0, 27.9%), respectively. Average AFP soil depth 0-10 cm in treatments NP-40 and NP-80 were only significantly different from NP-0 ($p=0.05$). All other treatments NP (10-80) in all depths (0-30 cm) were not significantly different from NP-0. This indicates soil water content was significantly evaporated from all treatments NP- (10-80) and all soil depths (0-30 cm). CH₄ uptake increased

significantly in NP-(10-80) as compare to P- (10-80) due to increase in AFP. CH₄ uptake was positively correlated with AFP in NP-(0-40) and negatively correlated in NP-80 in all soil depths 0-30 cm. Negative correlation in NP-80 was due to lowest AFP as compared to NP-(0-40). Soil CH₄ uptake significantly increased as AFP increased (Díaz et al., 2018).

Relationship between WFPS and CH₄ uptake in all treatments was exactly opposite to relationship between

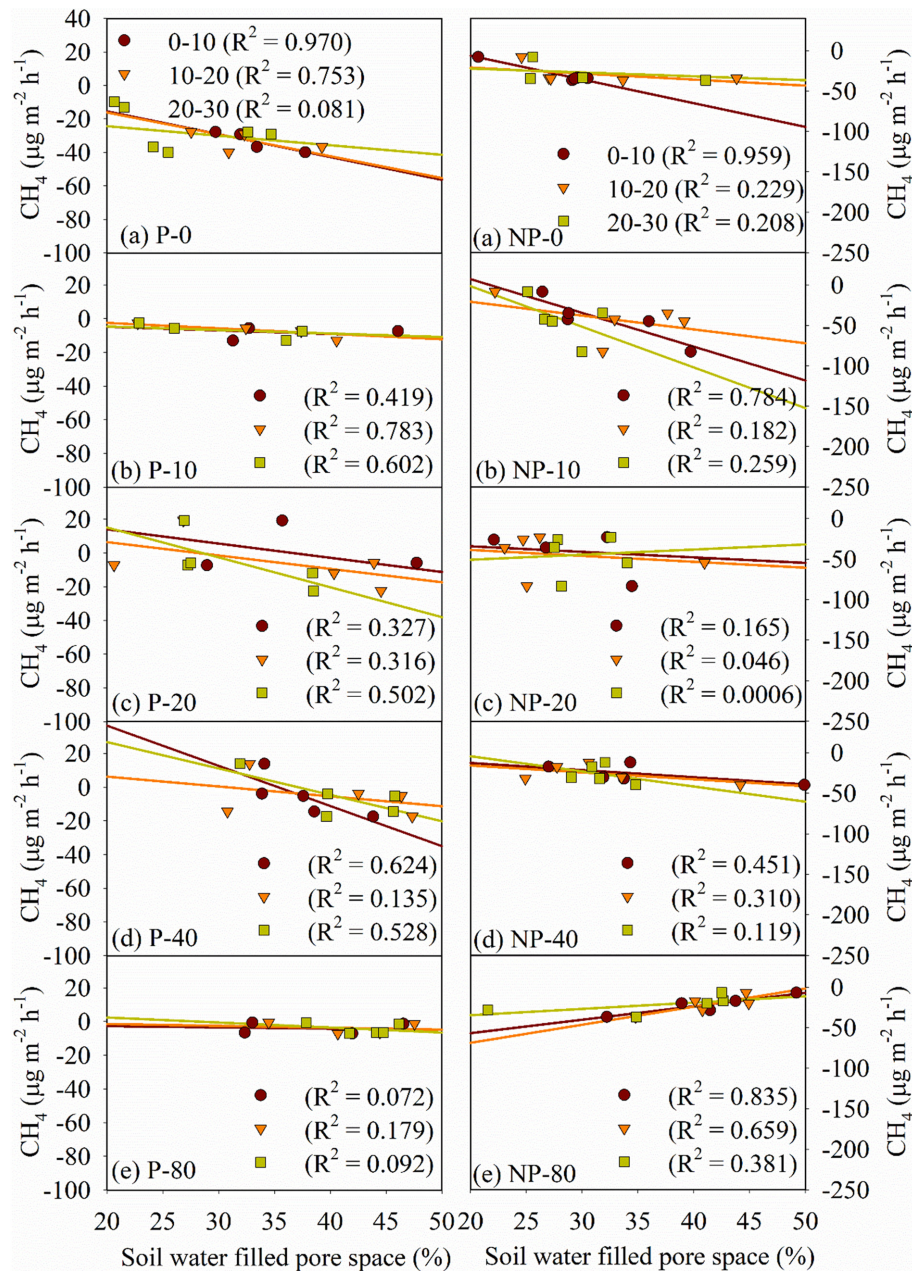


Fig. 4. Relationships (a-e, P-(0-80) and NP-(0-80)) between CH_4 emission and soil water filled pore space in different soil depths (0-10, 10-20, and 20-30 cm).

AFP and CH_4 uptake (Fig. 4). CH_4 uptake was negatively correlated with WFPS in all treatments P-(0-80) and NP-(0-40). CH_4 uptake was positively correlated with WFPS in all treatments NP-80). Positive correlation in NP-80 was due to highest WFPS as compared to NP-(0-40). WFPS in all soil depths (0-30 cm) of P-(10-80) and NP-(10-80) was not significantly different from P-0 and NP-80, respectively ($p=0.05$).

Immediate effect of water application on CH_4 reduction was prominent. CH_4 uptake in P-(10-80) was extrapolated to actual precipitation in 2017 (Fig. 5). Hourly precipitation varied between 0.0042 to 6.021 $\text{mm m}^{-2} \text{h}^{-1}$ (Korea Metrological Administration 2017). Estimated daily CH_4 uptake due to precipitation was calculated by extrapolation of field results. Since, average CH_4 uptake in P-(20-80) was not significantly different from each other, CH_4 uptake at P-

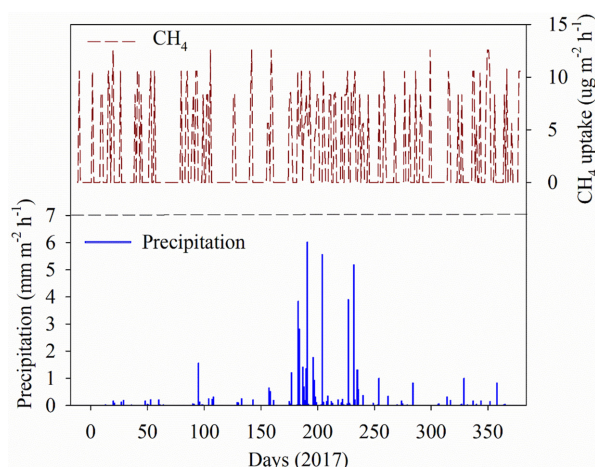


Fig. 5. Average daily precipitation, estimated average daily CH₄ uptake, South Korea.

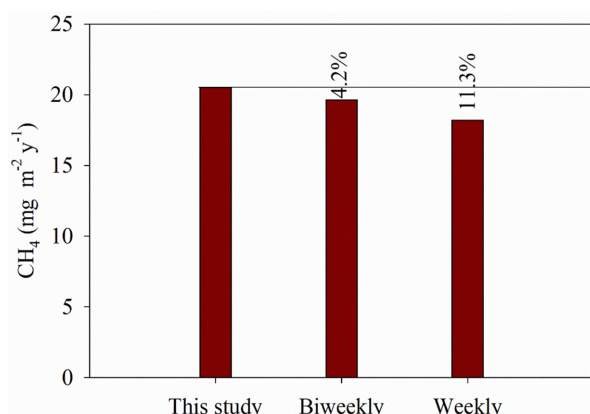


Fig. 6. Estimated total CH₄ uptake (2017) in this study and its difference (%) with weekly and biweekly sampling frequencies.

80 was assumed same for precipitation above $80 \text{ mm m}^{-2} \text{ h}^{-1}$. Minimal and maximal CH₄ uptake 4.4 and $12.6 \mu\text{g m}^{-2} \text{ h}^{-1}$ was observed at precipitation 80 and $0.1 \text{ mm m}^{-2} \text{ h}^{-1}$, respectively.

Effect of CH₄ sampling frequency (weekly and biweekly) on estimated total CH₄ uptake 2017 was compared (Fig. 6). The most common CH₄ sampling frequencies in temperate forests have been reported weekly and biweekly (Brumme and Borken., 1999; Kim et al., 2016; Kirschke et al., 2013). We assumed that CH₄ uptake was measured on weekly or biweekly from the field. Weekly and biweekly data of daily CH₄ uptake was subtracted from estimated CH₄ uptake in 2017. After subtracting weekly and biweekly CH₄ uptake, total estimated CH₄ uptake was 18.2 and $19.7 \text{ mg m}^{-2} \text{ y}^{-1}$, respectively. Total estimated CH₄ uptake in 2017 in this

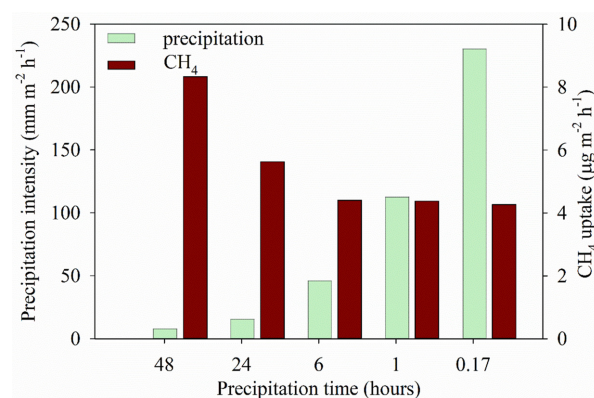


Fig. 7. CH₄ uptake at different intensities of precipitation.

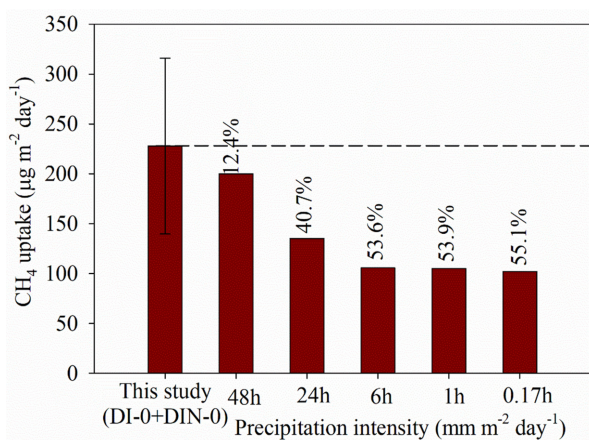
study was decreased by 11.3 and 4.2% . Total estimated CH₄ uptake in this study was not significantly different from weekly and biweekly CH₄ uptake subtracted data ($p=0.05$).

CH₄ uptake at variable intensity of precipitation was calculated by interpolation of CH₄ uptake results in this study (Fig. 7). Maximal total precipitation in 2017 was 372.7 , 370.1 , 275 , 112.5 , and 39.2 mm in 48 h , 24 h , 6 h , 1 h , and 0.17 h , respectively (Korea Metrological Administration 2017). Precipitation intensity increased with decreasing total precipitation time. Maximal and minimal precipitation was observed at 0.17 and 48 h , respectively. CH₄ uptake decreased with increasing precipitation intensity. CH₄ uptake in 24 , 6 , 1 , and 0.17 h was 32.5 , 47.2 , 47.5 , and 48.9% lower than CH₄ uptake in 48 h , respectively. CH₄ uptake in 6 , 1 and 0.17 h was not significantly different from each other. CH₄ uptake response to precipitation intensity was in agreement with CH₄ uptake in this study.

Average annual CH₄ uptake in temperate forests of different countries varied between 240 to $5890 \mu\text{g m}^{-2} \text{ day}^{-1}$ as shown in Table 1. Annual CH₄ uptake in the temperate forest of South Korea has been reported 1960 to $2920 \mu\text{g m}^{-2} \text{ day}^{-1}$ with an average uptake $2440 \mu\text{g m}^{-2} \text{ day}^{-1}$. Minimal and maximal daily CH₄ uptake in this study was 186.2 and $957.0 \mu\text{g m}^{-2} \text{ day}^{-1}$, respectively. This indicates that the experimental results from this research is not very different from the previous reports. It also confirms that the experimental procedure in this research is sound and comparable to the others. Average CH₄ uptake in (P-0+NP-0) was compared with reduced CH₄ uptake due to variable precipitation intensities as shown in Fig. 8. Hourly CH₄ uptake was converted to daily uptake by multiplied with

Table 1. Summary of published CH₄ uptake in temperate forests

Countries	CH ₄ uptake rate ($\mu\text{g m}^{-2} \text{ day}^{-1}$)		Forest types	References
	Range	Mean		
Canada	40-1100	570	Coniferous Deciduous	Lessard et al. (1994)
Denmark	140-330	240	Coniferous	Ambus and Christensen (1995)
	700-1070	890	Deciduous	Dobbie et al. (1996b)
	670-1370	1020	Coniferous	Priemé and Christensen (1997)
	170-2180	1180	Deciduous	Smith et al. (2000)
Germany	120-960	540	Coniferous	Butterbach-Bahl et al. (1998)
	1970	1970	Coniferous	Steinkamp et al. (2001)
	20-1030	530	Coniferous	Smith et al. (2000)
	30-680	360	Coniferous	Brumme and Borken (1999)
	250-3560	1910	Coniferous	Born et al. (1990)
Ireland	1340	1340	Coniferous	Butterbach-Bahl et al. (1998)
South Korea	1960	1960	Deciduous	Jang et al. (2006)
	2920	2920	Deciduous	Jang et al. (2011)
Norway	800-1400	1100	Coniferous	Sitaula et al. (1995)
	50-1900	970	Coniferous	Smith et al. (2000)
Poland	1000	1000	Mixed	Dobbie et al. (1996b)
	100-4580	2340	Deciduous	Smith et al. (2000)
Scotland	860-1060	960	Coniferous	Macdonald et al. (1996)
	10-3300	1400	Deciduous	Dobbie et al. (1996b)
	2190-2970	2580	Deciduous	Dobbie and Smith (1996a)
	70-1170	620	Coniferous	Macdonald et al. (1997)
Sweden	380	380	Coniferous	Klemetsson and Klemetsson (1997)
	2280-3670	700	Coniferous	Smith et al. (2000)
	0-600	300	Deciduous	Yavitt et al. (1993)
	2100-6900	4500	Deciduous	Goldman et al. (1995)
Japan	7600	7600	Mixed	Ishizuka et al. (2000)
	5890	5890	Coniferous	Tamai et al. (2003)
Russia	2950-8960	5010	Coniferous	Nakano et al. (2004)
UK	1080-3240	1080	Mixed	Smith et al. (2000)
	1060-2650	1610	Deciduous	Bradford et al. (2001)

**Fig. 8.** Comparison of CH₄ uptake in this study with reduced CH₄ uptake due to variable precipitation intensity. Error bar represent ± 1 standard error of mean.

twentyfour hours. Daily CH₄ uptake decreased with increasing precipitation intensity from 48 h to 6 h. CH₄ uptake reduction in 6 h, 1 h, and 0.17 h was not significantly different from eachother. Maximal and minimal decreased in CH₄ uptake was observed at 0.17h and 48h precipitation intensity, respectively.

4. Conclusion

We measured CH₄ uptake in temperate plantation from different treatments of variable precipitation, i.e., 0, 10, 20, 40, and 80 $\text{mm m}^{-2} \text{ day}^{-1}$. CH₄ flux was observed immediately after water application in P-(10-80) and observed after two

days interval when water was not applied NP-(0-80). CH₄ uptake in P-(10-80) was significantly lower than NP-(10-80). In our first hypothesis we assumed CH₄ emission may occur because rain water will replace CH₄ present in subsoil. Throughout the experimental period temperate forest soil was CH₄ sink rather than source. CH₄ uptake decreased significantly due to increasing water application in P-(10-80). We also hypothesized that CH₄ uptake will decrease with increasing WFPS, in P-80 WFPS was 53% higher than P-0 CH₄ uptake decreased 85.6% in P-80. This decrease in CH₄ uptake was positively correlated with air filled porosity and negatively correlated with water filled pore space. Soil texture at experimental site was sandy loam, which is relatively coarse texture further studies needed to establish if the relationship between variable precipitation to CH₄ uptake holds true across different soil texture classes. Our results can be used as a reference for regions with similar conditions. This study demonstrated the effects of variable precipitation on net daily CH₄ uptake and it may help in calculating more accurate net annual CH₄ sink in temperate forests in the world.

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References

- Ambus, P. and Christensen, S., 1995, Spatial and seasonal nitrous oxide and methane fluxes in Danish forest-, grassland-, and agroecosystems, *J Environ Qual*, **24**(5), 993-1001.
- Borken, W., Davidson, E.A., Savage, K., Gaudinski, J., and Trumbore, S.E., 2003, Drying and wetting effects on CO₂ release from organic horizons, *Soil Sci Soc Am J*, **67**, 1888-1897.
- Born, M., Dorr, H., and Levin, I., 1990, Methane consumption in aerated soils of the temperate zone, *Tellus B*, **42**(1), 2-8.
- Bradford, M., Wookey, P., Ineson, P., and Lappin-Scott, H., 2001, Controlling factors and effects of chronic nitrogen and sulphur deposition on methane oxidation in a temperate forest soil, *Soil Biol Biochem*, **33**(1), 93-102.
- Brumme, R. and Borken, W., 1999, Site variation in methane oxidation as affected by atmospheric deposition and type of temperate forest ecosystem, *Global Biogeochem Cy*, **13**(2), 493-501.
- Butterbach-Bahl, K., Gasche, R., Huber, C., Kreutzer, K., and Papen, H., 1998, Impact of N-input by wet deposition on N-trace gas fluxes and CH₄-oxidation in spruce forest ecosystems of the temperate zone in Europe, *Atmos Environ*, **32**(3), 559-564.
- Castro, M.S., Melillo, J.M., Steudler, P.A., and Chapman, J.W., 1994, Soil moisture as a predictor of methane uptake by temperate forest soils, *Can J For Res*, **24**(9), 1805-1810.
- Castro, M.S., Steudler, P.A., Melillo, J.M., Aber, J.D., and Bowden, R.D., 1995, Factors controlling atmospheric methane consumption by temperate forest soils, *Global Biogeochem Cycles*, **9**(1), 1-10.
- Czepiel, P., Crill, P., and Harriss, R., 1995, Environmental factors influencing the variability of methane oxidation in temperate zone soils, *J Geophys Res Atmos*, **100**(D5), 9359-9364.
- Dalal, R., Allen, D., Livesley, S., and Richards, G., 2008, Magnitude and biophysical regulators of methane emission and consumption in the Australian agricultural, forest, and submerged landscapes: a review, *Plant and Soil*, **309**(1-2), 43-76.
- Dastane, N.G., 1978, Effective Rainfall in irrigated agriculture, Irrigation and Drainage Paper No. 25, Food and Agriculture Organization, Rome, Italy, www.fao.org/docrep/X5560E/x5560e00.htm#Contents.
- Davidson, E.A., Ishida, F.Y., and Nepstad, D.C., 2004, Effects of an experimental drought on soil emissions of carbon dioxide, methane, nitrous oxide, and nitric oxide in a moist tropical forest, *Global Change Biol*, **10**(5), 718-730.
- Díaz, M.A., Bown, H.E., Fuentes, J.P., and Martínez, A.M., 2018, Soils act as sinks or sources of CH₄ depending on air-filled porosity in sclerophyllous ecosystems in semiarid central Chile, *Appl Soil Ecol*, **130**, 13-20.
- Dobbie, K. and Smith, K., 1996a, Comparison of CH₄ oxidation rates in woodland, arable and set aside soils, *Soil Biol Biochem*, **28**(10-11), 1357-1365.
- Dobbie, K., Smith K., Prieme, A., Christensen, S., Degorska, A., and Orlanski, O., 1996b, Effect of land use on the rate of methane uptake by surface soils in northern Europe, *Atmos Environ*, **30**(7), 1005-1011.
- Dutaur, L. and Verchot, L.V., 2007, A global inventory of the soil CH₄ sink, *Global Biogeochem Cycles*, **21**(4), 1-9.
- Epron, D., Plain, C., Ndiaye, F.-K., Bonnaud, P., Pasquier, C., and Ranger, J., 2016, Effects of compaction by heavy machine traffic on soil fluxes of methane and carbon dioxide in a temperate broadleaved forest, *Forest Ecol Manag*, **382**, 1-9.
- Gao, J., Zhou, W., Liu, Y., Zhu, J., Sha, L., Song, Q., Ji, H., Lin,

- Y., Fei, X., Bai, X., Zhang, X., Deng, Y., Deng, X., Yu, G., Zhang, J., Zheng, X., Grace, J., and Zhang, Y., 2018, Effects of Litter Inputs on N₂O Emissions from a Tropical Rainforest in Southwest China, *Ecosystems*, **21**(5), 1013-1026.
- Goldman, M.B., Groffman, P.M., Pouyat, R.V., McDonnell, M.J., and Pickett, S.T., 1995, CH₄ uptake and N availability in forest soils along an urban to rural gradient, *Soil Biol Biochem*, **27**(3), 281-286.
- Gütlein, A., Zistl-Schlingmann, M., Becker, J.N., Cornejo, N.S., Detsch, F., Dannenmann, M., Appelhans, T., Hertel, H., Kuzyakov, Y., and Kiese, R., 2017, Nitrogen turnover and greenhouse gas emissions in a tropical alpine ecosystem, Mt. Kilimanjaro, *Tanzania, Plant and soil*, **411**(1-2), 243-259.
- IPCC, 2014, The Scientific Basis. Contribution of Working Group I to the Fifth Assessment Report of the *Intergovernmental Panel on Climate Change* Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 1535.
- Ishizuka, S., Sakata, T., and Ishizuka, K., 2000, Methane oxidation in Japanese forest soils, *Soil Biol Biochem*, **32**(6), 769-777.
- Itoh, M., Ohte, N., and Koba, K., 2009, Methane flux characteristics in forest soils under an East Asian monsoon climate, *Soil Biol Biochem*, **41**(2), 388-395.
- Jang, I., Lee, S., Hong, J.-H., and Kang, H., 2006, Methane oxidation rates in forest soils and their controlling variables: a review and a case study in Korea, *Ecol Res*, **21**(6), 849-854.
- Jang, I., Lee, S., Zoh, K.-D., and Kang, H., 2011, Methane concentrations and methanotrophic community structure influence the response of soil methane oxidation to nitrogen content in a temperate forest, *Soil Biol Biochem*, **43**(3), 620-627.
- Keller, M. and Reiners, W.A., 1994, Soil-atmosphere exchange of nitrous oxide, nitric oxide, and methane under secondary succession of pasture to forest in the Atlantic lowlands of Costa Rica, *Global Biogeochem Cycles*, **8**(4), 399-409.
- Kessavalou, A., Doran, J.W., Mosier, A.R., and Drijber, R.A., 1998, Greenhouse gas fluxes following tillage and wetting in a wheat-fallow cropping system, *J Environ Qual*, **27**(5), 1105-1116.
- Kim, D.G., Thomas, A.D., Pelster, D., Rosenstock, T.S., and Sanz-Cobena, A., 2016, Greenhouse gas emissions from natural ecosystems and agricultural lands in sub-Saharan Africa: synthesis of available data and suggestions for further research, *Biogeosciences*, **13**(16), 4789-4809.
- Kim, D.-G., Vargas, R., Bond-Lamberty, B., and Turetsky, M., 2012, Effects of soil rewetting and thawing on soil gas fluxes: a review of current literature and suggestions for future research, *Biogeosciences*, **9**(7), 2459-2483.
- Kirschke, S., Bousquet, P., Ciais, P., Saunois, M., Canadell, J.G., Dlugokencky, E.J., Bergamaschi, P., Bergmann, D., Blake, D.R., Bruhwiler, L., Cameron-Smith, P., Castaldi, S., Chevallier, F., Feng, L., Fraser, A., Heimann, M., Hodson, E.L., Houweling, S., Josse, B., Fraser, P.J., Krummel, P.B., Lamarque, J., Langenfelds, R.L., Quéré, C.L., Naik, V., O'Doherty, S., Palmer, P.I., Pison, I., Plummer, D., Poulter, B., Prinn, R.G., Rigby, M., Ringeval, B., Santini, M., Schmidt, M., Shindell, D.T., Simpson, I.J., Spahni, R., Steele, L.P., Strode, S.A., Sudo, K., Szopa, S., van der Werf, G.R., Voulgarakis, A., Weele, M.V., Weiss, R.F., Williams, J.E., and Zeng, G., 2013, Three decades of global methane sources and sinks, *Nat Geosci*, **6**(10), 813.
- Klemetsson, Å.K. and Klemetsson, L., 1997, Methane uptake in Swedish forest soil in relation to liming and extra N-deposition, *Biol. Fertil. Soils*, **25**(3), 296-301.
- Lee, X., Wu, H.J., Sigler, J., Oishi, C., and Siccama, T., 2004, Rapid and transient response of soil respiration to rain, *Global Change Biol*, **10**, 1017-1026.
- Le Mer, J. and Roger, P., 2001, Production, oxidation, emission and consumption of methane by soils: A review, *Eur J Soil Biol*, **37**(1), 25-50.
- Lessard, R., Rochette, P., Toppt, E., Pattey, E., Desjardins, R.L., and Beaumont, G., 1994, Methane and carbon dioxide fluxes from poorly drained adjacent cultivated and forest sites, *Can J Soil Sci*, **74**(2), 139-146.
- MacDonald, J.A., Skiba, U., Sheppard, L.J., Hargreaves, K.J., Smith, K.A., and Fowler, D., 1996, Soil environmental variables affecting the flux of methane from a range of forest, moorland and agricultural soils, *Biogeochemistry*, **34**(3), 113-132.
- MacDonald, J.A., Skiba, U., Sheppard, L.J., Ball, B., Roberts, J.D., Smith, K.A., and Fowler, D., 1997, The effect of nitrogen deposition and seasonal variability on methane oxidation and nitrous oxide emission rates in an upland spruce plantation and moorland, *Atmos. Environ*, **31**(22), 3693-3706.
- Moore, B., Kaur, G., Motavalli, P., Zurweller, B., and Svoma, B., 2018, Soil greenhouse gas emissions from agroforestry and other land uses under different moisture regimes in lower Missouri River Floodplain soils: a laboratory approach, *Agroforest Syst*, **92**(2), 335-348.
- Nakano, T., Inoue, G., and Fukuda, M., 2004, Methane consumption and soil respiration by a birch forest soil in West Siberia, *Tellus B Chem Phys Meteorol*, **56**(3), 223-229.
- Nakano, T., Sawamoto, T., Morishita, T., Inoue, G., and Hatano, R., 2004, A comparison of regression methods for estimating soil-atmosphere diffusion gas fluxes by a closed-chamber technique, *Soil Biol Biochem*, **36**(1), 107-113.
- Ojima, D., Valentine, D., Mosier, A., Parton, W., and Schimel, D., 1993, Effect of land use change on methane oxidation in temperate forest and grassland soils, *Chemosphere*, **26**(1-4), 675-685.

- Prieme, A. and Christensen S., 2001, Natural perturbations, drying-wetting and freezing-thawing cycles, and the emission of nitrous oxide, carbon dioxide and methane from farmed organic soils, *Soil Biol Biochem*, **33**, 2083-2091.
- Priemé, A. and Christensen, S., 1997, Seasonal and spatial variation of methane oxidation in a Danish spruce forest, *Soil Biol Biochem*, **29**(8), 1165-1172.
- Pulleman, M. and Tietema A., 1999, Microbial C and N transformation during drying and rewetting of coniferous forest floor material, *Soil Biol Biochem*, **31**, 275-285.
- Reeburgh, W.S., 2003, Global methane biogeochemistry, *Treatise on geochemistry*, **4**, 347.
- Rolston, D.E., 1986, Gas Flux, In: Klute, A. (ed.), *Methods of Soil Analysis: Part 1-Physical and Mineralogical Methods*, Soil Science Society of America, p. 1103-1119.
- Sitaula, B.K., Bakken, L.R., and Abrahamsen, G., 1995, CH₄ uptake by temperate forest soil: effect of N input and soil acidification, *Soil Biol Biochem*, **27**(7), 871-880.
- Smith, K.A., Dobbie, K.E., Ball, B.C., Bakken, L.R., Sitaula, B.K., Hansen, S., Brumme, R., Borken, W., Christensen, S., Priemé, A., Fowler, D., Macdonald, J.A., Skiba, U., Klemmedtsson, L., Kasimir-Klemmedtsson, A., Degórska, A., and Orlanski, P., 2000, Oxidation of atmospheric methane in Northern European soils, comparison with other ecosystems, and uncertainties in the global terrestrial sink, *Global Change Biol*, **6**(7), 791-803.
- Smith, K.A., Ball, T., Conen, F., Dobbie, K.E., Massheder, J., and Rey, A., 2003, Exchange of greenhouse gases between soil and atmosphere: interactions of soil physical factors and biological processes, *Eur J Soil Sci*, **54**(4), 779-791.
- Sponseller, R.A., 2007, Precipitation pulses and soil CO₂ flux in a Sonoran Desert ecosystem, *Global Change Biol*, **13**, 426-436.
- Steinkamp, R., Butterbach-Bahl, K., and Papen, H., 2001, Methane oxidation by soils of an N limited and N fertilized spruce forest in the Black Forest, Germany, *Soil Biol Biochem*, **33**(2), 145-153.
- Tamai, N., Takenaka, C., Ishizuka, S., and Tezuka, T., 2003, Methane flux and regulatory variables in soils of three equal-aged Japanese cypress (*Chamaecyparis obtusa*) forests in central Japan, *Soil Biol Biochem*, **35**(5), 633-641.
- Teh, Y.A., Silver, W.L., and Conrad, M.E., 2005, Oxygen effects on methane production and oxidation in humid tropical forest soils, *Global Change Biol*, **11**(8), 1283-1297.
- Vasconcelos, S.S., Zarin, D.J., Capanu, M., Littell, R., Davidson, E.A., Ishida, F.Y., Santos, E.B., Araújo, M.M., Aragão, D.V., Rangel-Vasconcelos, L.G.T., Oliveira, F.D.A., McDowell, W.H., and de Carvalho, C.J.R., 2004, Moisture and substrate availability constrain soil trace gas fluxes in an eastern Amazonian regrowth forest, *Global Biogeochem Cycles*, **18**(2), 1-10.
- Von Fischer, J.C. and Hedin, L.O., 2007, Controls on soil methane fluxes: Tests of biophysical mechanisms using stable isotope tracers, *Global Biogeochem Cycles*, **21**(2), 1-9.
- Wang, Y., Hu, C., Ming, H., Oenema, O., Schaefer, D.A., Dong, W., Zhang, Y., and Li, X., 2014, Methane, carbon dioxide and nitrous oxide fluxes in soil profile under a winter wheat-summer maize rotation in the North China Plain, *PLoS one*, **9**(6): e98445.
- Wei, H., Peng, C., Liu, S., Liu, X., Li, P., Song, H., Yuan, M., and Wang, M., 2018, Variation in Soil Methane Fluxes and Comparison between Two Forests in China, *Forests*, **9**(4), 204.
- Weitz, A.M., Keller, M., Linder, E., and Crill, P., 1999, Spatial and temporal variability of nitrogen oxide and methane fluxes from a fertilized tree plantation in Costa Rica, *J Geophys Res Atmos*, **104**(D23), 30097-30107.
- Whalen, S.C. and Reeburgh, W., 1996, Moisture and temperature sensitivity of CH₄ oxidation in boreal soils, *Soil Biol Biochem*, **28**, 1271-1281.
- Yavitt, J., Simmons, J., and Fahey, T., 1993, Methane fluxes in a northern hardwood forest ecosystem in relation to acid precipitation, *Chemosphere*, **26**(1-4), 721-730.
- Zhu, J., Mulder, J., Wu, L.P., Meng, X.X., Wang, Y.H., and Dörsch, P., 2013, Spatial and temporal variability of N₂O emissions in a subtropical forest catchment in China, *Biogeosciences*, **10**(3), 1309-1321.