

Methane consumption and soil respiration by a birch forest soil in West Siberia

By TOMOKO NAKANO^{1*}, GEN INOUE², and MASAMI FUKUDA³, ¹*Department of Geography, Tokyo Metropolitan University, Hachioji, 192-0397 Japan;* ²*National Institute for Environmental Studies, Tsukuba, 305-0053 Japan;* ³*Institute of Low Temperature Science, Hokkaido University, Sapporo, 060-0819 Japan*

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ABSTRACT

Methane and carbon dioxide fluxes were measured in a birch forest soil in West Siberia, in August 1999, June 2000 and September 2000. The study site had a very thick organic horizon that was subject to drought during the observation periods. The soils always took up CH₄, while CO₂ was released from the surface to the atmosphere. CH₄ consumption and CO₂ emission rates ranged from 0.092 to 0.28 mg C m⁻² h⁻¹ and from 110 to 400 mg C m⁻² h⁻¹ respectively. The CH₄ consumption rate and soil temperatures showed significant relationships for individual measurements. The soil respiration rate was weakly correlated with individual soil temperatures. This study examined the effect of current and lagged soil temperatures at a depth of 5 cm on CH₄ consumption and soil respiration. The variation in the correlation coefficient between CH₄ consumption and lagged soil temperature was greatest at a 4-h lag, whereas that for soil respiration showed a gentle peak at lags from several hours to half a day. This difference in the temperature-related lag effect between CH₄ consumption and soil respiration results from differences in the exchange processes. Neither flux showed any correlation with soil moisture. The limited variation in soil moisture during our observation period may account for the lack of correlation. However, the droughty soil conditions resulted in high gas diffusion and, consequently, high CH₄ consumption.

1. Introduction

Terrestrial ecosystems exchange carbon with the atmosphere. Upland boreal forest soils consume atmospheric methane (CH₄) by microbial oxidation and release large amounts of carbon dioxide (CO₂) by means of root respiration and microbial decomposition of soil organic matter. Therefore, boreal forest soils play a potentially important role in atmospheric CH₄ and CO₂ levels, which influence global climate change.

A number of studies have examined consumption of atmospheric CH₄ in boreal and temperate forests. Studies have taken place mainly in North America (Steudler et al., 1989; Crill, 1991; Castro et al., 1994, 1995; Savage et al., 1997; Gullledge and Schimel, 2000; McLain et al., 2002) and Europe (Born et al., 1990; Dörr et al., 1993; Koschorreck and Conrad, 1993; Priemé et al., 1996; Priemé and Christensen, 1997; Brumme and Borken, 1999; Borken et al., 2000). These studies reported consumption rates ranging from 0 to 0.25 mg C m⁻² h⁻¹. Studies based on both laboratory and field measurements have also indicated that CH₄ consumption is controlled by soil tempera-

ture (Born et al., 1990; Crill, 1991; Whalen and Reeburgh, 1996; Priemé and Christensen, 1997; Savage et al., 1997; Borken et al., 2000), soil moisture (Crill, 1991; Koschorreck and Conrad, 1993; Castro et al., 1994; Czepiel et al., 1995; Whalen and Reeburgh, 1996; Borken et al., 2000; McLain et al., 2002), the amount of N in the soil (Steudler et al., 1989; Castro et al., 1995; Brumme and Borken, 1999) and the amount of soil organic matter (Czepiel et al., 1995; Savage et al., 1997; Brumme and Borken, 1999).

The evolution of CO₂ from soils is commonly called soil respiration. Temperature has a strong impact on rates of soil respiration (Dörr and Münnich, 1987; Lessard et al., 1994; Rayment and Jarvis, 1997; Savage et al., 1997; Boone et al., 1998; Davidson et al., 1998). Soil moisture also controls soil respiration. Extremely dry and wet soils reduce CO₂ production and diffusion, respectively (Dörr and Münnich, 1987; Grant and Rochette, 1994; Keith et al., 1997; Davidson et al., 1998, 2000).

This paper presents measurements of CH₄ and CO₂ fluxes between soils and the atmosphere in summer at a birch forest in West Siberia. The West Siberian Plain is an extremely paludinous area that is characterized by enormous carbon stocks due to peat accumulation (Yefremov and Yefremova, 2001). Our study site also has a thick organic soil layer derived from peat, although it is currently well-drained woodland. White birch forest is very common in the southern part of West Siberia, and the volume

*Corresponding author.
e-mail: nakano@comp.metro-u.ac.jp

of growing stock in such forests has been estimated to represent about 40% of the total growing stock in southern West Siberian forests (Alexeyev et al., 1996). However, little up-to-date information exists on CH₄ oxidation and soil respiration in these forests. We aim to (1) evaluate rates of CH₄ consumption and soil respiration from the boreal forest study site in West Siberia and (2) investigate the environmental parameters affecting net CH₄ and CO₂ fluxes from the forest soil.

2. Sites and methods

The research site was located in a boreal forest (lat 56°52'N, long 83°17'E) near Plotnikovo, Tomsk Oblast, in the southern part of the West Siberian Plain. The forest consisted of white birch trees (*Betula pendula*) with occasional Scotch pines (*Pinus sylvestris*). The ground vegetation consisted of shrubs (*Ledum palustre*, *Chamaedaphne calyculata*, *Rubus chamaemorus*) and herbaceous plant communities. The area has an annual mean air temperature of -1.1°C, mean monthly temperatures varying between -18.6°C and 17.6°C throughout the year, and annual precipitation of approximately 530 mm (Lapshina et al., 2001).

A 3 cm thick layer of decomposing litter (L horizon) was underlain with a thick (about 90 cm), black, organic layer (H horizon) derived from bog peat, overlying a silty, mineral horizon (B horizon). Surface soil samples were collected and brought back to Japan in September 2000. The relative gas diffusion coefficient (D/D_0) was determined using Osozawa's (1987) diffusion chamber method with 100 cm³ undisturbed soil cores. After autoclaving at 105°C, soil core porosity was measured with a volume meter (DIK-1000, Daiki Rika Kogyo Co.). The total carbon and nitrogen contents were determined by dry combustion using a CN analyser (Vario-EL, Elementar Analysensysteme GmbH) equipped with a thermal conductivity detector. The soil properties of the surface organic layer at our study site are summarized in Table 1. The values are the averages and standard deviations of three or four samples.

The CH₄ and CO₂ fluxes were measured using a closed-chamber method. The system comprised an acrylic plastic cham-

ber 0.4 m in height, stainless steel collars, a portable CH₄/CO₂ analyser equipped with a semiconductor CH₄ detector and a solid electrolyte CO₂ detector, and a 12 V lead-acid battery. The chamber was equipped with sampling ports, a thermometer and a battery-operated fan. We also placed a Tedlar bag inside the chamber, which was connected to the outside of the chamber with a 0.25 inch tube, to compensate for pressure changes within the chamber. Two collars, which enclosed 0.152 m², were embedded in the soil to a depth of 10 cm before measurements commenced and remained in place throughout the study. The distance between the two collars was about 10 m. Neither collar included live plants, and surface and soil conditions within the collars were similar. The chamber was fitted into a water-filled groove in the collar for the measurements.

Sample air was continuously pumped at a rate of about 1 dm³ min⁻¹ from the chamber through a polyethylene tube to the CH₄/CO₂ analyser and back to the chamber. Signals from the CH₄/CO₂ detectors were outputted as voltage values and recorded at 5 s intervals by a datalogger (NR-1000, Keyence Co.). Inoue et al., (1998) has described the operation and accuracy principles of solid-state CH₄ and CO₂ detectors. To calibrate the detectors, air samples were also taken from the chamber with a syringe before and after flux measurements and placed in 10 ml vacuum vials for analysis. Analysis was conducted with a gas chromatograph equipped with a flame ionization detector for CH₄ and a thermal conductivity detector for CO₂ (GC-14BPTF, Shimadzu Co.). Flux estimates were based on changes in chamber CH₄ and CO₂ concentrations over time, which were measured for 10 min after the chamber enclosed the soil surface. Net fluxes were calculated from the rate constant of the exponential curve fit (Nakano et al., 2004). Observations were conducted over three 1-week periods in August 1999, June 2000 and September 2000. CH₄ and CO₂ fluxes were measured four times per day for each of the two collars: early morning (0300–0600 local time (LT)), morning (0900–1200 LT), afternoon (1500–1800 LT) and night (2100–2400 LT).

The mean differences in the flux values between the two collars and among the three periods were compared statistically using the Mann–Whitney and Kruskal–Wallis tests, respectively (Zar, 1999).

Environmental data were collected concurrently with flux measurements. Air temperature at 1.5 m above ground level and soil temperature at a depth of 5 cm were measured every 30 min using a datalogger with two thermistor probes (SK-L200T, Sato Keiryoki Manufacturing Co.). Soil moisture at the time of the flux measurements was determined using time-domain reflectometry (TDR) probes (0–12 cm depth; HydroSense, Campbell Scientific Inc.) positioned vertically from the surface into the soil. The TDR results were cross-checked using the gravimetric method with 100 cm³ undisturbed soil cores. Soil temperatures (5 and 10 cm depths) at the time of the flux measurements were also measured using thermocouple thermometers (ST-920, Testoterm Co.).

Table 1. Soil properties of the surface organic layer (H horizon) in the West Siberia birch forest

| | Average | SD |
|------------------------------------|---------|--------|
| Thickness (m) | 0.89 | 0.056 |
| Bulk density (Mg m ⁻³) | 0.19 | 0.017 |
| Porosity (%) | 90 | 2.4 |
| D/D_0 * | 0.37 | 0.048 |
| Total C (g g ⁻¹) | 0.43 | 0.016 |
| Total N (g g ⁻¹) | 0.025 | 0.0021 |
| C/N | 17 | 1.8 |

* D/D_0 is the relative gas diffusion coefficient.

Table 2. Summary of net CH₄ and CO₂ fluxes and environmental variables in the West Siberia birch forest

| | | Aug. 1999 | Jun. 2000 | Sep. 2000 |
|---|----------|-----------------|----------------|----------------|
| CH ₄ flux (mg C m ⁻² h ⁻¹) | <i>n</i> | 5 | 14 | 24 |
| | Mean | -0.12 | -0.20 | -0.15 |
| | SE | 0.013 | 0.011 | 0.006 |
| | Range | -0.14 to -0.092 | -0.28 to -0.15 | -0.20 to -0.11 |
| CO ₂ flux (mg C m ⁻² h ⁻¹) | <i>n</i> | | 17 | 24 |
| | Mean | ND | 280 | 200 |
| | SE | ND | 15 | 9.4 |
| | Range | ND | 180 to 400 | 110 to 310 |
| Air temp. (°C) | Mean | 11.1 | 21.9 | 9.1 |
| | Range | 10.3 to 12.0 | 12.7 to 30.7 | 1.2 to 13.8 |
| Soil temp. (°C) at 5 cm depth | Mean | 10.4 | 15.6 | 12.5 |
| | Range | 9.4 to 11.5 | 13.5 to 18.3 | 8.6 to 14.6 |
| Vol. water content (%) | Mean | 11.1 | 10.7 | 11.5 |
| | Range | 8 to 20 | 7 to 15 | 8 to 14 |

SE, standard error; ND, not determined.

3. Results

During all gas samplings, CH₄ and CO₂ fluxes were consistently negative and positive respectively, indicating that atmospheric CH₄ was consumed by the soils and CO₂ was released from the surface to the atmosphere (Table 2). Since the flux values did not differ significantly between the subsites for either CH₄ or CO₂ (Mann–Whitney test, $p > 0.10$), the mean fluxes were computed from all the measurements made in each period. The mean values differed significantly among the measurement periods (Kruskal–Wallis test for CH₄ flux and Mann–Whitney test for CO₂ flux, $p < 0.01$). The mean CH₄ consumption rate ranged from 0.12 to 0.20 mg C m⁻² h⁻¹, and individual values ranged from 0.092 to 0.28 mg C m⁻² h⁻¹. The highest value was observed in June 2000, when air and soil temperatures were also highest. CO₂ fluxes were three orders of magnitude greater than CH₄ influx, and individual measurements ranged from 110 to 400 mg C m⁻² h⁻¹. The mean value was also higher in June 2000 than in September 2000. Soil moisture (volumetric water content) was consistently low (about 10%) during the study periods, although soil temperature showed seasonal variation. Since bulk density was very low and the soil was droughty, the relative gas diffusion coefficient, D/D_0 , was high (Table 1).

Individual CH₄ and CO₂ fluxes showed a negative relationship, indicating high soil respiration at times of high CH₄ consumption (Fig. 1). Savage et al. (1997) found a similar relationship for spatial distributions of CH₄ and CO₂ fluxes. Our results suggest that this negative relationship is appropriate for temporal variations in CH₄ and CO₂ fluxes, as well as for the spatial variation described by Savage et al. (1997).

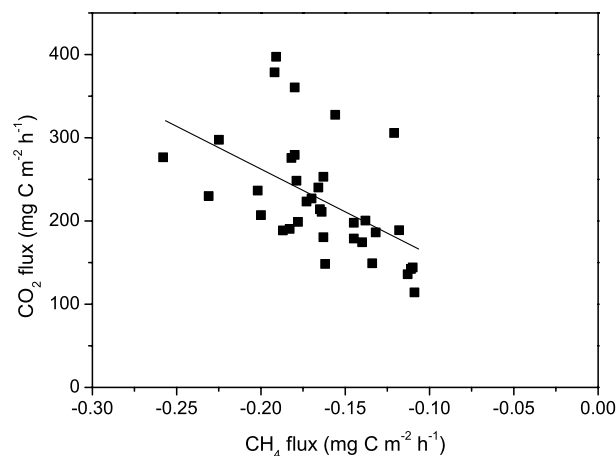


Fig. 1. Relationship between individual CH₄ and CO₂ fluxes.

Significant rank correlations were found between individual CH₄ consumption rates and air and soil (5 cm depth) temperatures, while individual soil respiration rates correlated with the soil temperature at 5 cm depth (Table 3). Mean rates of CH₄ consumption and soil respiration for each measurement period also positively related to the mean soil temperature (Fig. 2).

4. Discussion

4.1. CH₄ consumption

Methane uptake rates in temperate and boreal forests have a large temporal and spatial variation. Researchers have found

Table 3. Spearman's rank correlation coefficient (r_s) for correlations between individual measurements of gas exchange rates and environmental variables

| | CH ₄ consumption | Soil respiration |
|--------------------|-----------------------------|------------------|
| Air temp. | 0.40* | NS |
| Soil temp. (5 cm) | 0.50** | 0.38* |
| Soil temp. (10 cm) | NS | NS |
| Vol. water content | NS | NS |

NS, not significant at $p = 0.05$.

* $p < 0.05$.

** $p < 0.01$.

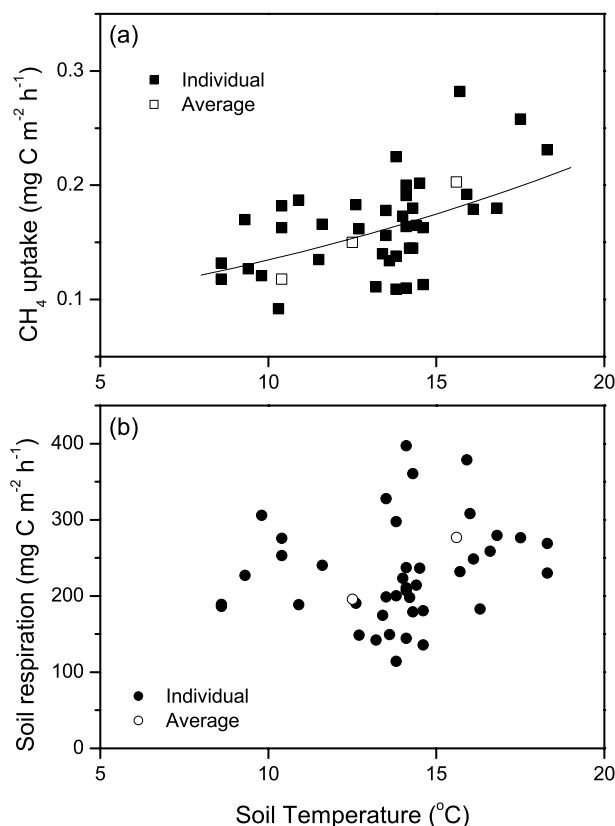


Fig 2. Relationships between soil temperature at 5 cm depth and (a) CH₄ consumption and (b) soil respiration for individual measurements and averages. The exponential regression line (solid line) is significant ($p < 0.01$).

summer-time (June to September) values of 0.084 to 0.11 mg C m⁻² h⁻¹ in a mixed deciduous-coniferous forest in New Hampshire (Crill, 1991), 0.08 to 0.056 mg C m⁻² h⁻¹ in several soils in Germany (Koschorreck and Conrad, 1993), 0.09 to 0.22 mg C m⁻² h⁻¹ in a red pine plantation and a mixed hardwood stand in central Massachusetts (Castro et al., 1995) and 0.01 to 0.04 mg C m⁻² h⁻¹ in an upland birch/aspen forest in interior Alaska (Gulledge and Schimel, 2000). Our results were relatively high compared with the ranges found in these previous studies.

Gas transport is generally considered a major factor affecting the CH₄ consumption rate (Potter et al., 1996), as gas transport capabilities regulate CH₄ diffusion from the atmosphere to the oxidizing sites. Accordingly, a number of studies have found that soil moisture (Koschorreck and Conrad, 1993; Castro et al., 1994; Lessard et al., 1994; Whalen and Reeburgh, 1996; Borken et al., 2000) and soil texture (Born et al., 1990; Dörr et al., 1993) are most important in controlling the CH₄ consumption rate, as these factors influence soil gas diffusivity. We found no significant relationship between measured soil moisture and CH₄ consumption, but the limited variation in soil moisture during the study periods may account for this finding. Throughout the study period, the site had a very low volumetric water content and high relative gas diffusivity (D/D_0). The high diffusivity probably caused high CH₄ consumption rates in our results.

Individual CH₄ fluxes are significantly correlated with soil temperature at 5 cm depth (Fig. 2a and Table 3). The mean fluxes for each measurement period also showed positive relationships with soil temperature. Previous studies have found relatively weak correlations between soil temperature and CH₄ consumption when Q_{10} values ranged from 1.2 to 2.0 (Crill, 1991; Lessard et al., 1994; Priemé and Christensen, 1997; Savage et al., 1997). In this study, the Q_{10} values, calculated by fitting the temperature function to an exponential formula (Davidson et al., 1998), equalled 1.7 for individual CH₄ consumption. This value was comparable with those reported in the studies mentioned above.

Most studies examining the effect of soil temperature on CH₄ consumption have used soil temperatures at the time of flux measurement or daily and monthly average temperatures. We, however, investigated the correlation between CH₄ consumption and time-lagged soil temperature (Fig. 3). Spearman's rank correlation coefficients (r_s) between CH₄ consumption and lagged soil temperature showed an obvious change with the time lag. The correlation coefficient was highest for the temperature 4 h

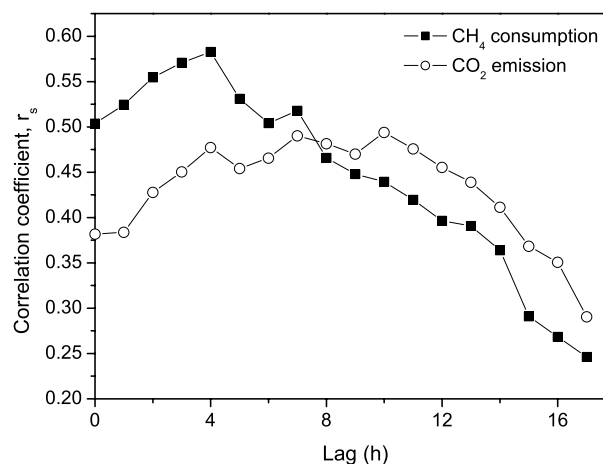


Fig 3. Spearman's rank correlation coefficient between gas exchange rates and lagged soil temperature at a depth of 5 cm.

before the flux measurement ($r_s = 0.58$, $p = 0.0002$) and declined steeply as the lag increased. This result indicates that time-lagged temperatures enabled a more sensitive regulation of CH_4 uptake than did simultaneous temperature measurements. Since CH_4 consumption was caused simply by microbial oxidation, this retardation in the temperature relationship can be interpreted as a lag in the heat transfer from the 5 cm level to the CH_4 oxidation zone. To examine this lagged response to temperature more closely, it is necessary to measure the methane concentration and temperature profiles in soils.

4.2. Soil respiration

Soil respiration is a combination of root respiration and microbial decomposition of soil organic matter. Root respiration represents 33 to 71% of total soil respiration (Nakane et al., 1983; Bowden et al., 1993; Eguchi et al., 1997; Striegl and Wickland, 1998). Although many environmental factors influence the biological and physical processes controlling soil respiration rates, previous studies have shown that respiration rates relate to soil temperature, moisture and/or organic characteristics (Dörr and Münnich, 1987; Skopp et al., 1990; Crill, 1991; Grant and Rochette, 1994; Lessard et al., 1994; Savage et al., 1997; Boone et al., 1998; Davidson et al., 1998, 2000).

Our soil respiration rates were at the high end of the range that has been reported in other studies of boreal and temperate regions. For example, other researchers found summer-time values of 100 to 200 $\text{mg C m}^{-2} \text{ h}^{-1}$ in a New Hampshire forest soil (Crill, 1991), 70 to 160 $\text{mg C m}^{-2} \text{ h}^{-1}$ in upland boreal forests in Manitoba (Savage et al., 1997), 170 to 320 $\text{mg C m}^{-2} \text{ h}^{-1}$ in a mixed-hardwood control-site stand in Massachusetts (Boone et al., 1998), 65 to 230 $\text{mg C m}^{-2} \text{ h}^{-1}$ in a mature jack pine stand in Saskatchewan (Striegl and Wickland, 1998) and 18 to 390 $\text{mg C m}^{-2} \text{ h}^{-1}$ in East Siberian larch forests (Sawamoto et al., 2000). We hypothesize that the greater availability of organic matter in soil and litter resulted in the higher soil respiration rates at our site.

We found only a weak relationship between individual soil respiration rates and soil temperature at 5 cm depth (Fig. 2b and Table 3), although many studies have reported that temperature has a strong impact soil respiration (e.g. Boone et al., 1998; Davidson et al., 1998). This study examined the effect of time-lagged soil temperature at 5 cm depth on soil respiration together with CH_4 consumption (Fig. 3). The rank correlation coefficient increased as the time lag increased from 0 to 4 h and remained high until 10 h. The variation in the relationship between soil respiration and the temperature obviously differed from that with methane uptake, which had a sharp peak at a lag of 4 h. Furthermore, the correlation coefficient for the soil respiration had a small range of variation compared with that of the CH_4 uptake. This suggests that the response to temperature differs between soil respiration and CH_4 oxidation, probably depending on the different gas exchange processes involved.

The process of CH_4 consumption is relatively simple, because it is regulated exclusively by microbial activity and gas diffusion from the atmosphere to the soil, whereas soil respiration is controlled by complicated systems such as plant physiology and phenology and availability of labile C, in addition to soil moisture and temperature controls. Post et al. (2003), suggested that labile soil organic matter significantly affected heterotrophic respiration at short time scales and that this obscures the sensitivity of the respiration to temperature. Eguchi et al. (1997), in diurnal observations of soil respiration in a deciduous broad-leaved forest in Hokkaido, Japan, found that diurnal variation patterns in soil CO_2 efflux differed from diurnal variation patterns in temperature. That is, soil respiration increased at night and decreased in the daytime. Eguchi et al. (1997) deduced that the discrepancy between diurnal variation patterns was caused by tree physiology affecting root respiration, which contributed approximately 70% of the CO_2 production in the H/A horizon at their study site. Generally, when plant roots absorb water, the rate of root respiration increases to replenish energy lost through temporary consumption. Based on measurements of stem circumference and xylem pressure potential presented by Hinckley and Bruckerhoff (1975), Eguchi et al. (1997) suggested that root water uptake by large trees occurred from dusk to night-time. Thus, a night-time increase in soil respiration accompanying high water uptake activity probably contributes to the night-time increase in soil respiration.

By contrast, Rayment and Jarvis (1997) and Davidson et al. (1998) found significant relationships between diurnal variation in soil respiration and soil temperature in an old black spruce forest in Saskatchewan and a mixed hardwood forest in Massachusetts, respectively. Unlike our result, these studies indicated that the soil respiration at their sites responded to soil temperature in phase. Therefore, the temperature sensitivity of soil respiration differs from site to site, and probably depends on the dominant tree species, stand age and soil type. Future research must evaluate the ratio of root respiration to total soil respiration and investigate the physiological conditions of the trees to better understand the factors controlling soil respiration.

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