Continuous measurement of soil respiration at a larch forest in Eastern Siberia

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1. INTRODUCTION

Forests dominated by larch (*Larix gmelinii*) are extensively distributed on permafrost in northern East Asia (Gower and Richards, 1990). It is suggested that the boreal forest is a large CO_2 sink (Denning *et al.*, 1995; Otto-Bliesner and Upchurch, 1997). The extensive larch forests probably play an important role in the global carbon cycle. However, field measurement of CO_2 exchange between the atmosphere and larch ecosystems is very limited in Siberia (Dolman *et al.*, 2004; Machimura *et al.* 2005).

We measured soil respiration, which is a major CO_2 source in the terrestrial ecosystem (Schimel, 1995), continuously during the growing seasons in three years, 2003 to 2005, at a larch forest in eastern Siberia.

2. MATERIALS AND METHODS

2.1 Study site

The study site was a mature larch forest located 25 km northeast of Yakutsk, Republic of Sakha, Russian Federation (62.3°N, 129.5°E). The forest was developing on permafrost. The dominant tree species was larch (*Larix gmelinii*) with density of 2100 stems ha⁻¹ (Yajima *et al.*, 1998). The forest floor was covered mainly with shrub and moss. Soil texture was loamy and soil organic content in the top 1-m layer was 13.2 kgC m⁻² (Sawamoto *et al.*, 2003).

2.2 Chamber method

Soil respiration (R_s) was measured continuously with an automated closed/open chamber system during three growing seasons in 2003, 2004 and 2005 on the forest floor.

The automated system consisted of five chambers made from opaque PVC with a covering ground area of 0.0186 m^2 , a data logger (CR10X, CSI), an infrared CO₂ analyzer (LI820, Licor), an air pump, and some electric parts to





switch the airflow between the analyzer and chambers. Each chamber was closed for 6 min one after another. R_s was calculated from an increasing rate of CO₂ concentration in chamber headspace for the last 4 min. Chambers were installed on collars, which were inserted 2 cm deep into the ground. In 2003, the chamber system was shared with another site in shifts of one week each. Also, one chamber broke down throughout the season. Thus, R_s was measured at 4 points at intervals of one week from June 5 to September 18 (47 days). Contrastively, R_s was measured continuously from May 12 to September 29 (141 days) for 2004 and from May 7 to September 27 (144 days) for 2005. In 2004 and 2005, one chamber was fixed on one point and other 4 chambers were moved between 2 points at intervals of 7–10 days (Fig. 1).

2.3 Data analysis

We analyzed temperature responses of daily mean R_s (gC m⁻² d⁻¹) calculated from half-hourly measurements using the following well-known exponential equation:

$$R_s = a \exp(bT), \qquad (1$$

where *a* and *b* are fitted constants, and *T* is soil temperature (°C). In addition, R_s was normalized using equation (2) to eliminate its temperature response:

$$R_{s,b} = R_s \exp(b(T_b - T)), \qquad (2)$$

where $R_{s,b}$ is normalized R_s at T_b , and T_b is base temperature (°C).

For the analysis and discussion, we used meteorological data including soil temperature and soil moisture, which were measured at the same site.

3. RESULTS AND DISCUSSION

3.1. Temperature response

 R_s increased with soil temperature on a daily basis (Fig. 2). In all years, r^2 was highest with T at depth of 10 cm (T_{10}) among depth of 5, 10, 15 and 25 cm. Seasonal variations in R_s can be explained by T_{10} by 75, 89 and 88–94% in 2003, 2004 and 2005, respectively. In 2004, R_s deviated below from its exponential relationship from July 4 to July 26 (DOY186–208) (Fig. 2b). On the other hand, R_s was divided into two groups before and after mid-July in 2005. R_s jumped up from the first group into the second one between July 15 (DOY196) and July 16 with precipitation of 17 mm (Fig. 2c). Although data of 2004 and 2005 almost overlapped, data of 2003 were smaller than other data (Fig. 2d).



Fig. 2. Relationship between daily $R_{\rm s}$ and soil temperature at depth of 10 cm for a) 2003, b) 2004, c) 2005 and d) all years. Exponential curves are fitted.

3.2. Seasonal variation

There were spatial variations in R_s , and the variations increased in July and August (Fig. 3). To investigate temporal variation in R_s during the growing season from May through September, data gaps occurring at one-week intervals were filled from T_{10} using equation (1) at each point on a daily basis.



Fig. 3. Seasonal variations in daily R_s measured in a) 2003, b) 2004 and c) 2005. Different symbols denote different points.

Fig. 4 shows seasonal variations in spatially averaged R_s along with T_{10} , soil moisture and precipitation. In 2003, R_s changed almost in parallel with soil temperature; it peaked at 4.5 gC m⁻² d⁻¹ on July 26 with daily mean T_{10} of 12.7°C. Contrastively, R_s peaked on August 22 at 4.8 gC m⁻² d⁻¹ in 2004, which was about one month later than in 2003. This late peak was caused by low temperature and less precipitation in July. In 2005, R_s suddenly doubled at mid-July from 3.4 to 6.5 gC m⁻² d⁻¹. Precipitation over 10 mm after a dry period of 45 days caused the doubling of R_s . R_s kept a high level until late August.

 $R_{\rm s}$ was normalized at 6°C using equation (2) for 2004 and 2005 to eliminate the temperature effect from seasonal variation (Fig. 5); a temperature coefficient of *b* was obtained from all data in both years (Fig. 2d). We excluded $R_{\rm s}$ in 2003, because its temperature response differed (Fig. 2d). Seasonal variations in normalized $R_{\rm s}$ ($R_{\rm s,6}$) were generally similar to each other. In 2004, $R_{\rm s,6}$ kept low at 2.0–2.5 gC m⁻² d⁻¹ until late May and increased in June with fluctuation between 2.5 and 3.5 gC m⁻² d⁻¹. $R_{\rm s,6}$ decreased again to about 2.0 gC m⁻² d⁻¹ in July and suddenly increased at the end of July. $R_{\rm s,6}$ gradually increased until late August and decreased with fluctuation in September. In 2005, $R_{\rm s,6}$ decreased gradually until mid-July at a low level below 2.5 gC m⁻² d⁻¹ and jumped up above 3.5 gC m⁻² d⁻¹. After mid-July, $R_{\rm s,6}$ kept high until early September and gradually decreased toward the end of September. These seasonal variations were related to the development of the active layer of permafrost, precipitation patterns and larch phenology. Low $R_{\rm s,6}$ measured until late May in 2004 was probably caused by shallow active layer. On the other hand, low $R_{\rm s,6}$ until around mid-June in 2005 was probably related to high soil moisture, which was caused by a larger amount of snow

accumulation during the previous winter than normal winter. Gradual decrease in September was probably related to the phenological change of larch functions, including needle senescence. A sudden drop in September 2004 would be related to soil desiccation, because a positive linear relationship of $R_{s,6}$ was found with soil moisture ($r^2 = 0.55$) in late August and September.



Fig. 4. Seasonal variations in a) daily $R_{\rm s}$ spatially averaged after gap-filling, b) daily mean soil temperature at depth of 10 cm, c) soil moisture at depth of 5 cm, d) daily precipitation in 2003, e) 2004 and f) 2005, from May 1 to September 30.

We focused on the summer period between mid-July and early August, DOY170–220, when $R_{s,6}$ increased considerably (Fig. 6). Increase in $R_{s,6}$ was triggered by precipitation of about 10 mm at late-July in 2004 and mid-July in 2005. Precipitation increases soil moisture and directly affects soil respiration. However, it is difficult to explain the pattern of $R_{s,6}$ by only soil

moisture, because $R_{s,6}$ was independent of decrease in soil moisture between DOY170 and 195 in 2005. In addition, $R_{s,6}$ slowly increased during DOY199–205 in spite of decreasing soil moisture in 2004. For 50 days of midsummer, $R_{s,6}$ showed a decreasing relationship with atmospheric water vapor deficit (VPD) (Fig. 7); the relationship can be fitted to a rectangular hyperbola ($r^2 = 0.45$). It was reported that stomatal conductance of larch trees in eastern Siberia sharply decreased as VPD increased, and it regulated CO₂ uptake of larch forest (Dolman *et al.*, 2004). Also, recent studies showed the coupling of canopy gas exchange with soil respiration (Irvine *et al.*, 2005; Tang *et al.*, 2005). Therefore, high VPD probably decreased larch photosynthesis through stomatal closure, and consequently decreased $R_{s,6}$ indirectly.



Fig. 5. Seasonal variations in a) normalized daily R_s at 6°C and b) daily mean VPD above canopy, from May 1 to September 30.

3.2. Interannual variation

The cumulative change of R_s was shown in Fig. 8 for 153 days from May 1 to September 30. Cumulative R_s was largest in 2005 and smallest in 2003 throughout the period, whereas cumulative R_s was very similar between 2004 and 2005 until the end of June in spite of large difference in soil temperature (Fig. 4).

The sum of R_s for the growing season of 153 days was 272 ± 48 (mean \pm standard deviation), 338 ± 46 and 479 ± 94 gC m⁻² for 2003, 2004 and 2005, respectively; these values are mean and standard deviation of 4 or 9 points (Fig. 9). The growing season R_s was significantly larger in 2005 than other two years by Tukey-Kramer test (p < 0.01), whereas it was not significantly different between 2003 and 2004. The seasonal sum of precipitation was 226, 118 and 191 mm for 2003, 2004 and 2005, respectively. In addition, the seasonal mean of T_{10} was 5.1, 3.4 and 6.3°C for 2003, 2004 and 2005, respectively (Fig. 4). As a result, the climate of the growing season in 2005 was warm and moist compared with that in 2004. This temperature difference mainly caused large difference in R_s between 2004 and 2005, because the relationship between $R_{\rm s}$ and T_{10} was almost identical (Fig. 2d). Also, relatively moist climate would partly increased $R_{\rm s}$ in 2005, which was typically shown in September (Fig. 5). As for the year of 2003, $R_{\rm s}$ was smallest among 3 years in spite of the largest precipitation and second highest soil temperature. Thus, R_s in 2003 may be questionable because of insufficient replication and frequency of measurement. Although winter soil respiration has large variation within vegetation types, it accounts for roughly 20% of annual soil respiration for tundra and boreal forests (Hobbie et al., 2000). Using this information, annual R_s can be estimated at 423 and 599 gC m⁻² y⁻¹ for 2004 and 2005, respectively. These estimated values are slightly larger than annual soil respiration reviewed by Raich and Schlesinger (1992).





Fig. 9. Comparison of R_s summed for the growing season among 3 years. A vertical bar denotes standard deviation.

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