

Seasonal variation of mono- and sesquiterpene emission rates of Scots pine

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Abstract. The seasonal variation of mono- and sesquiterpene emission rates of Scots pine was measured from April to October in 2004. The emission rates were measured daily in the afternoons with the exception of weekends. Emissions were measured from two branches; one of them was debudded in May (branch A), while the other was allowed to grow new needles (branch B). The monoterpene emission pattern remained almost constant throughout the measurement period, Δ^3 -carene being the dominant monoterpene (50–70% of the VOC emission). The standard monoterpene emission potential (30°C) was highest during early summer in June (the average of the two branches $1.35 \mu\text{g g}^{-1} \text{h}^{-1}$) and lowest during early autumn in September (the average of the two branches $0.20 \mu\text{g g}^{-1} \text{h}^{-1}$). The monoterpene emission potential of branch A remained low also during October, whereas the emission potential of branch B was very high in October. The sesquiterpenes were mainly emitted during mid summer, the dominant sesquiterpene being β -caryophyllene. Branch A had a higher sesquiterpene emission potential than branch B and the emission maximum occurred concomitant with the high concentration of airborne pathogen spores suggesting a potential defensive role of the sesquiterpene emissions. The sesquiterpene emissions were well correlated with linalool and 1,8-cineol emissions, but not with monoterpenes. Sesquiterpene and 1,8-cineol emissions were equally well described by the temperature dependent and the temperature and light dependent algorithms. This is due to the saturation of the light algorithm as the measurements were always conducted during high light conditions.

1 Introduction

During the past decade the characteristics of the biogenic VOC emissions from European ecosystems have been intensively studied, both in the boreal forests and in the South European regions. So far the studies have mainly concentrated on the short-term emissions of monoterpenes (Janson 1993; Janson et al., 2001; Rinne et al., 1999, 2000; Komenda and Koppmann, 2002) and isoprene (Hakola et al., 1998), even though the VOC emissions have large seasonal variations. Downy birch has been shown to emit large amounts of linalool and sesquiterpenes early in the growing season, while their later emissions consist mainly of sabinene and ocimene (Hakola et al., 2001). Norway spruce emits monoterpenes in May, whereas their emission mainly consists of isoprene in June and sesquiterpenes in July (Hakola et al., 2003). Due to their high reactivity the sesquiterpenes have drawn a lot of attention lately. Their atmospheric lifetimes are only a few minutes and therefore they cannot be measured from the ambient air samples (Hakola et al., 2000, 2003), yet they have a high potential to form secondary organic aerosol (Hoffmann et al., 1997; Jaoui et al., 2003). Bonn and Moortgat (2003) suggest that sesquiterpene ozonolysis could be responsible for the atmospheric new particle formation observed frequently in several rural locations (Mäkelä et al., 1997; Kulmala et al., 2004). Sesquiterpenes may also affect tropospheric ozone concentrations – they may participate in ozone formation when enough nitrogen oxides are present and in a very clean environment some of the ozone deposition may be attributed to sesquiterpene reactions (Kurpius and Goldstein, 2003). The oxidation processes of these very reactive compounds can also produce hydroxyl radicals (Holzinger et al., 2004).

The present study is focused on the seasonal development of the VOC emission rates of Scots pine. The temperature and light dependence of the VOC emissions of Scots pine

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has been studied by Tarvainen et al. (2005) during the growing seasons of 2002 and 2003. However the data set in the study was concentrated on spring and it was found that especially for sesquiterpenes more measurements were needed during high summer to assess the seasonality. The knowledge on seasonal variation of biogenic VOC emissions is needed for several purposes, e.g. for the evaluation of the effect of biogenic VOCs on the ozone concentrations in the boreal forested areas and the estimation of the contribution of sesquiterpenes to the ozone deposition as well as the new particle formation and growth processes in clean areas.

2 Experimental

The VOC emission rates of Scots pine (*Pinus sylvestris* L.) were measured in southern Finland in Hyytiälä (61°51' N, 24°17' E) from April to October in 2004. The emission rates were measured daily in the afternoons with the exception of weekends, when no measurements were conducted. The measured tree was growing in a natural forest environment, with an average tree height of 14 m. The samples were collected at a height of about 13 m from two fully sunlit branches. Two branches of the same tree were enclosed in two Teflon enclosures in March and remained there until October in order to prevent losses to the surfaces that can take place due to insufficient equilibrium time (Helmig et al., 2004). The other end of the enclosures remained open except during measurements, to ensure sufficient ventilation for the shoot in other times. The purge flow was maintained only during the measurements. One of the measured shoots (henceforth called branch A) was debudded in May before budbreak, while the other (branch B) one was allowed to grow new needles.

The emission rates were measured using a dynamic flow through technique. The volume of the cylinder shape Teflon enclosures was approximately 20 l and they were equipped with inlet and outlet ports and a thermometer inside the enclosure. The cover of the enclosures was made of transparent Teflon film. The photosynthetically active photon flux density (PPFD) was measured just above the enclosure. The flow through the enclosure was about 8 l per minute. Ozone was removed from the inlet air using MnO₂-coated copper nets (Pollmann et al., 2005). The efficiencies of the ozone scrubbers were checked prior to and after the measurement campaign and they removed ozone completely. The air samples were collected onto adsorbent tubes simultaneously from both the inlet and outlet ports. The emission rate is determined as the mass of compound per needle dry weight and time according to

$$E = \frac{(C_2 - C_1)F}{m} \quad (1)$$

Here C_2 is the concentration in the outgoing air, C_1 is the concentration in the inlet air, and F is the flow rate into the

enclosure. The dry weight of the biomass (m) was determined by drying the needles at 75°C until consistent weight was achieved, and the mass of the new needles was estimated using a regression from a comparable adjacent shoot where daily growth measurements were performed.

The samples were collected using adsorbent tubes filled with Tenax-TA and Carbopack-B. The sampling time was 60 min and sampling flow rate about 100 ml min⁻¹ resulting in approximately 3 liter sample volumes. The adsorbent tubes were analyzed using a thermodesorption instrument (Perkin-Elmer ATD-400), connected to a gas chromatograph (HP 5890) with HP-1 column (60 m, i.d. 0.25 mm) and a mass-selective detector (HP 5972). The detection limits were 32 ng m⁻³ for isoprene and 2-methyl-3-buten-2-ol (MBO). For monoterpenes the detection limits were: camphene 11 ng m⁻³, carene 42 ng m⁻³, 1,8-cineole 84 ng m⁻³, limonene 60 ng m⁻³, α -pinene 30 ng m⁻³, β -pinene 36 ng m⁻³, sabinene 59 ng m⁻³, and terpinolene 29 ng m⁻³. The detection limit for β -caryophyllene was 79 ng m⁻³.

The standardized emission potentials (at temperature 293 K and light intensity 1000 μ mol photons m⁻² s⁻¹) of 2-methyl-3-buten-2-ol (MBO) were calculated according to the temperature and light dependent emission algorithm and those of mono- and sesquiterpenes according to the temperature dependent emission algorithm presented by Guenther et al. (1993) and Guenther (1997). The light and temperature dependence of the emissions of some of the compounds was further tested as described in Tarvainen et al. (2005) using the emission algorithms presented by Guenther (1997) and Schuh et al. (1997).

Sampling of fungal spores was undertaken with volumetric Burkard-spore trap (Hirst, 1952) on the level of tree canopy in Hyytiälä. The spores were counted during 1 March–14 August 2004, and identified on stratified randomised fields with microscopic observation (Mäkinen, 1981). The daily average counts of airborne spores belonging to orders Uredinales and Ustilaginales, and subdivisions Basidiomycotina and Ascomycotina (one-, two- and more than two-celled spore types) were used in analyses. Uredinales and Ustilaginales are biotrophic pathogens. Marked fraction of Basidiomycotina spores dispersing in spring-time and summer are pathogenic or saprophytic wood-rotting fungi. Group Ascomycotina contains numerous pathogenic and endophytic species invading living hosts (Manion, 1991).

According to the statistics of the Finnish Meteorological Institute, year 2004 was close to or slightly warmer than the long term average. In the southwestern and western parts of the country the growing season started already in April, which is earlier than average. April was, however, also very dry, with the precipitation amounts far below the long term average. The beginning of May was exceptionally warm. However, there were severe cold spells later in May and even in June in the southern parts of the country, which slowed down or even completely stopped the progress of the grow-

Table 1. The monthly mean VOC emission potentials (30°C) of two branches (A and B) of the same Scots pine tree ($\text{ng g}_{(\text{needledryweight})}^{-1} \text{h}^{-1}$). The emission potentials are standardized to 30°C according to Guenther et al. (1993) using β value 0.09 K^{-1} for monoterpenes and 0.19 K^{-1} for sesquiterpenes (see Table 2). The MBO emission was standardized using the light and temperature dependent isoprene algorithm. (Guenther, 1997). 95% confidence intervals (in parenthesis) are calculated as $2 \cdot \text{std}(\text{emission potential}) / \sqrt{(\text{number of measurements})}$. The effect of the debudding of branch A on 13 May has been removed from the monthly mean by ignoring the measurements for one week after debudding.

	April	May	June	July	August	September	October
MBO A	58 (8)	67 (23)	33 (11)	28 (7)	16 (3)	21 (13)	16 (8)
MBO B	14 (16)	52 (20)	126 (39)	56 (19)	26 (5)	27 (7)	14 (15)
1,8-Cineol A	2 (4)	44 (12)	45 (18)	66 (18)	19 (7)	1 (1)	1 (2)
1,8-Cineol B	5 (5)	30 (11)	35 (14)	71 (20)	18 (4)	0 (0)	0 (0)
linalool A	4 (7)	0 (0)	0 (0)	64 (31)	2 (3)	0 (0)	0 (0)
linalool B	7 (9)	0 (0)	0 (0)	14 (10)	2 (3)	0 (0)	0 (0)
3-Carene A	314 (141)	896 (395)	905 (904)	572 (152)	306 (98)	176 (86)	188 (89)
3-Carene B	752 (289)	910 (380)	1132 (640)	863 (404)	322 (152)	196 (106)	619 (375)
other monoterpenes A	141 (54)	263 (96)	360 (236)	266 (89)	136 (42)	69 (29)	84 (77)
other monoterpenes B	369 (171)	282 (82)	295 (132)	243 (62)	142 (55)	129 (112)	404 (227)
caryophyllene A	9 (17)	3 (4)	139 (79)	385 (112)	56 (23)	9 (12)	0 (0)
caryophyllene B	5 (11)	28 (47)	33 (24)	127 (35)	28 (16)	12 (10)	38 (77)
other sesquiterpenes A	0 (0)	0 (0)	26 (40)	92 (19)	23 (11)	6 (11)	0 (0)
other sesquiterpenes B	0 (0)	5 (9)	4 (6)	32 (16)	14 (7)	7 (7)	15 (31)

ing season. The summer was exceptionally rainy, with record high precipitation amounts in southern and central Finland in July. May, June and July were also cloudier than the long term average. September was again quite rainy, completing the exceptional wetness of thermal growing season, which then ended after the first week of October in the southern and central parts of the country.

3 Results and discussion

3.1 Seasonal variability of the emissions

The 7-day running medians of the measured emission rates are presented in Figs. 1a–e. Small amounts of monoterpenes were emitted already in March (not shown in the figure) when the measurements were started. During the first week of May the temperature was exceptionally high for the time of the year and the maximum monoterpene emission rate for the growing season of 2004 was observed during that period (Fig. 1a). In general, the monoterpene emissions from the two measured branches did not differ much, although in April, branch B emitted almost twice as much monoterpenes as branch A. In order to investigate the differences in the emission behaviour of new and old needles branch A was

debudded on 13 May, while branch B was allowed to grow the new needles. The effect of debudding is discussed in more detail below. Monoterpene emission rates increased at the end of June, but they decreased already in August and remained low until the emission rates from branch B increased at the end of October. Unexpectedly high emission rates from Scots pine during autumn have been measured also earlier (Tarvainen et al., 2005). The monoterpene emission pattern remained almost the same throughout the growing season (Table 1). The main compound was Δ^3 -carene, composing more than 50% of the emission during almost the whole period. The normalized emission potentials of Δ^3 -carene and other monoterpenes from both branches were significantly lower in the late summer and early fall (August–September) than earlier in the summer.

Sesquiterpenes, mainly β -caryophyllene (Fig. 1b) and the oxygen containing compounds 1,8-cineole (Fig. 1c) and linalool (Fig. 1d) were detected in the emission later in the year than the monoterpenes. In addition to β -caryophyllene, two other sesquiterpenes were detected, according to NIST mass spectra library these compounds could be α -farnesene and α -caryophyllene, but these identifications are only tentative. 1,8-cineole was identified for the first time at the end of April and linalool was not observed until July. The

sesquiterpene emissions were initiated at the end of June and they ceased already in the beginning of August, although sesquiterpenes were found in a few occasional samples also during spring. The emissions of β -caryophyllene were well correlated with those of linalool and 1,8-cineole, but not with the monoterpene emissions. In the middle of the summer (15 June–15 August), at the time of the sesquiterpene emissions, the correlation coefficients (r) between the β -caryophyllene and linalool, 1,8-cineole, α -pinene and Δ^3 -carene emissions from branch A were 0.9, 0.8, 0.3 and 0.3, respectively. The sesquiterpenes and linalool have been found to be well correlated also in the VOC emission from downy birch (Hakola et al., 2001). The contribution of β -caryophyllene was significant in July, when it was the most abundant compound after Δ^3 -carene in the emission from branch A (Table 1). The normalized emission potentials of β -caryophyllene and other sesquiterpenes from both branches were significantly higher in July than the rest of the measurement period.

MBO emissions were detected for the first time at the end of April (Fig. 1e). Simultaneously, a small isoprene emission was also observed. The isoprene emission rate was well correlated with the MBO emission rate and it is possible that this is an artefact as MBO can be dehydrated – resulting in isoprene formation – in the analysis. It is possible that our MBO results are underestimated by up to 50%. Still the MBO emissions were quite low, although MBO has been found to be one of the major VOC compounds in pine forests in the United States (Goldan et al., 1993; Harley et al., 1998).

3.2 Sesquiterpene emissions and pathogen spores

Martin et al. (2003) have shown that methyl jasmonate causes a 5-fold increase in the total terpenoid emission of Norway spruce, with an emission pattern shift towards oxygenated monoterpenes (linalool) and sesquiterpenes (e.g. (E)- β -farnesene). Methyl jasmonate is known to induce the production of defense-related compounds and resistance against several herbivores, and has also been used to induce defensive responses without causing physical damage to the plant. Since daily measurements of pathogen spores from ambient air in Hyttälä were available, we plotted them together with the sum of the emission rates of the stress-induced compounds (linalool, 1,8-cineole and sesquiterpenes) (Fig. 2). The sesquiterpene emissions were first detected at about the same time as spores with potential pathogens were measured for the first time. Also the maximum emissions of sesquiterpenes, linalool and 1,8-cineole occur concomitant with the maximum of the pathogen spores (Fig. 2). This finding would support the theory that the sesquiterpenes and oxygenated monoterpenes are released by the plant for defensive purposes (Wei, 2003). Factor analysis conducted with the data also showed at least two groups of compounds with emissions behaving in distinctive manner. The first group comprised of α -pinene, sabinene, β -pinene, Δ^3 -carene and limonene, and the second one of 1,8-cineole, linalool,

caryophyllene and α -farnesene. The pathogen spores had their highest factor loading on the same factor with the latter group.

There are also other stress factors that could initiate sesquiterpene emissions, ozone for example, and one could speculate if the sesquiterpene emissions are affected when removing ozone from the air entering the enclosure. Removal of ozone is a standard procedure, however, to avoid chemical reactions between ozone and emitted compounds, and thus our results are comparable with others that have been gained using shoot enclosures in field conditions.

3.3 The effect of debudding

The increased monoterpene emissions caused by the debudding of branch A are studied in Figs. 3a, b, where the normalized (30°C) emission potentials of various compounds for both branches are shown. The increase of the emission occurred immediately after the cutting of the terminal buds of the branch A on 13 May and lasted for approximately a week (Fig. 3a). The emission rates of all monoterpenes from branch A increased after the debudding, but those of MBO, 1,8-cineole and β -caryophyllene did not (Fig. 3b). It is well known that in pine MBO is not stored in needles, but is released to the air immediately after it is synthesized (Harley et al., 1998), and this might be the case for 1,8-cineol, too. According to Niinemets et al. (2002), the pools of oxygenated VOCs with low Henry's law constant, preferably partitioning to the leaf aqueous phase, may rapidly adjust to changes in stomatal aperture, and their storage pools therefore are relatively small when the stomata are open.

Later in the growing season these two branches behaved somewhat differently. The most important difference in the emission rates between the branches was found in the emissions of sesquiterpenes. The debudded branch (A) had much higher sesquiterpene emission rates reflecting to emission potentials normalized to 30°C, which differ significantly from each other (Table 1). The July average emission potentials (30°) were 0.39 ± 0.11 and $0.13 \pm 0.04 \mu\text{g g}^{-1} \text{h}^{-1}$ for branches A and B, respectively. The only visible difference between the branches was that after debudding branch A started to grow new buds. However, it is not likely that the sesquiterpenes were emitted from the developing buds, since high emissions related to bud sprouting could not be seen in the beginning of the summer, neither in fall during the current year's bud development in branch B.

Contrary to sesquiterpene emissions, higher MBO emission rates were measured from branch B which was not debudded and had new needles. This could be due to maturation of new needle tissue, as the enzyme activities for terpenoid biosynthesis correlate with the developmental stage of leaf tissues (Lehning et al., 2001).

The seasonal variation of the mono- and sesquiterpene emission potentials of the two branches is depicted in Fig. 4. The monoterpene emission potentials between the two

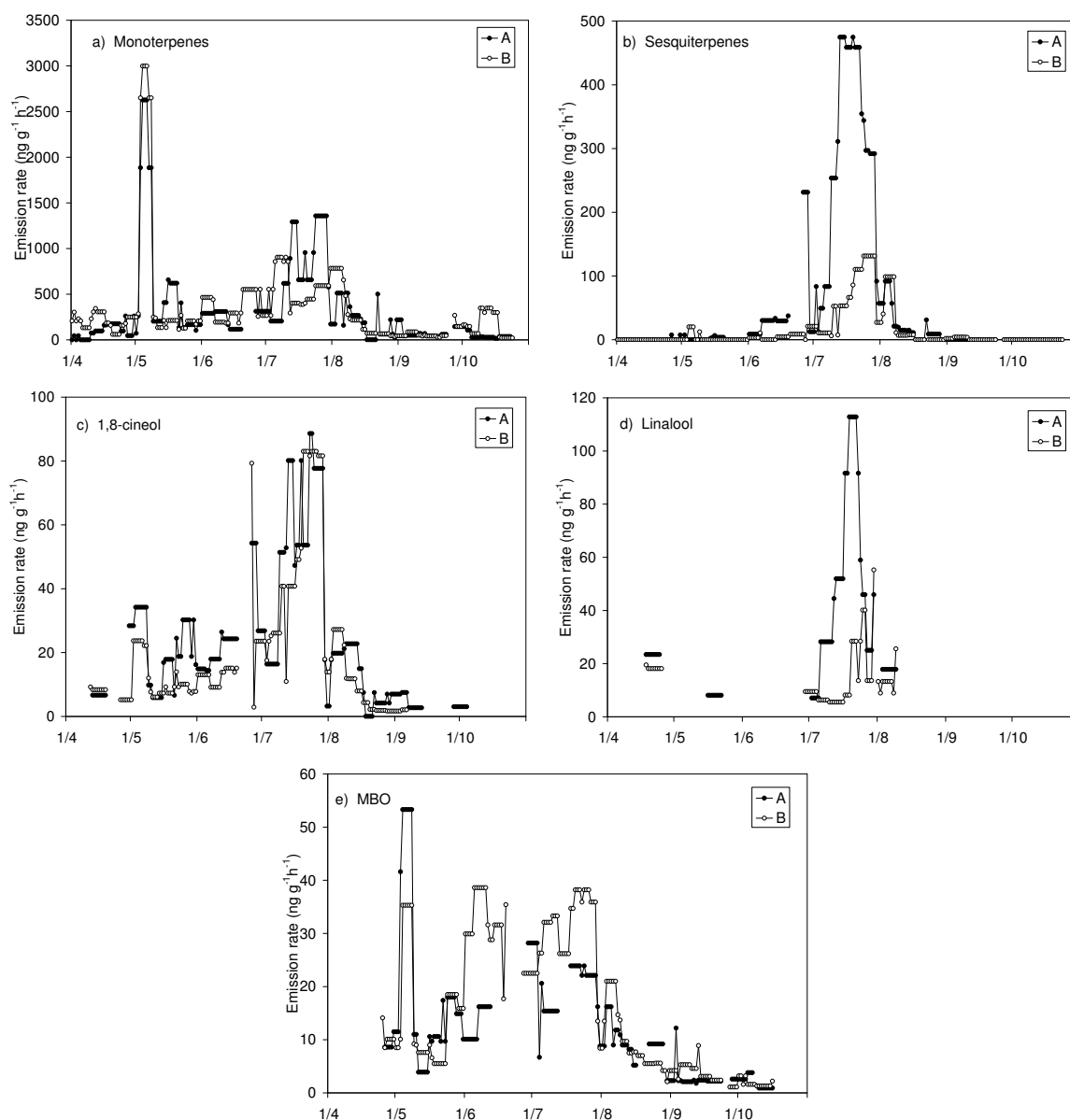


Fig. 1. 7-day running medians of the measured VOC (sum of monoterpenes, sum of sesquiterpenes, MBO, 1,8-cineol, linalool) emission rates from Scots pine in Hyttiälä, Finland in 2004. The samples were taken at midday from two different branches (A and B) of the same tree.

branches did not show significant differences until autumn. In October the branch which was not debudded (B) showed significantly higher emission potential than the debudded branch (Fig. 4). The contribution of the new needle year class to the whole branch monoterpene emissions can thus be significant, especially during the winter hardening period in late fall.

3.4 Light and temperature dependence of the emissions

As has been shown in a number of studies (e.g. Janson, 1993; Komenda and Koppmann, 2002; Tarvainen et al., 2005) the

VOC emission rates of Scots pine are dependent on temperature and the present study again confirmed the earlier results. As our data set included the high summer measurements which were lacking in the work of Tarvainen et al. (2005), we have further investigated the temperature and light dependence of sesquiterpenes and 1,8-cineole during this intense emission period. In addition to the algorithms commonly used in biogenic emission modeling (Guenther et al., 1993; Guenther, 1997) which were also applied by Tarvainen et al. (2005), an algorithm presented by Schuh et al. (1997) was tested for the light dependence.

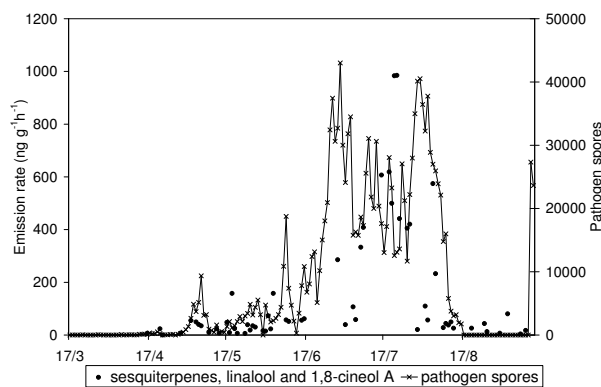


Fig. 2. The sum of the emission rates of sesquiterpenes, linalool and 1,8-cineole from branch A and the concentrations of pathogen spores measured in Hyytiälä.

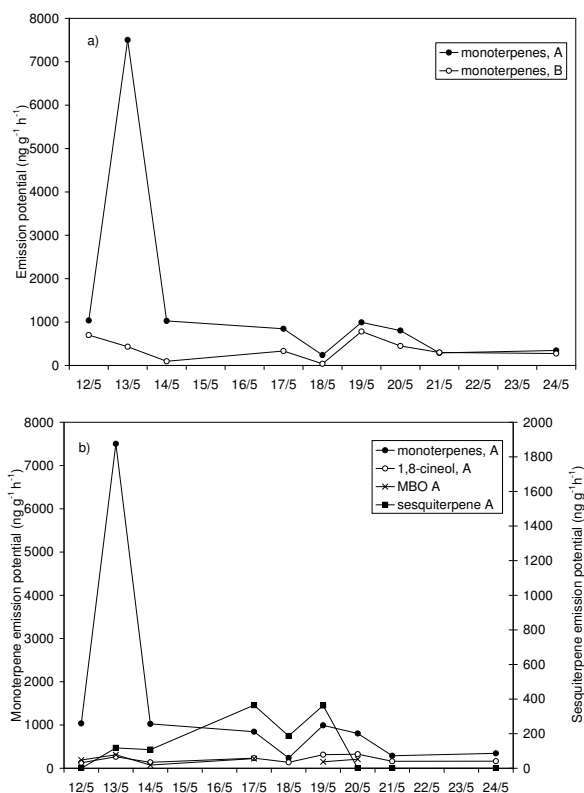


Fig. 3. (a) The effect of the debudding of branch A on 13 May on the monoterpene emissions. Branch B was left untouched. (b) The effect of the debudding of branch A on the emission potential of total monoterpenes, 1,8-cineole, β -caryophyllene and MBO.

Nonlinear regression was used to fit the observed emissions to the temperature dependent emission algorithm, Eq. (5) of Guenther et al. (1993), henceforth called the TEMP algorithm. In addition to the simple exponential TEMP algorithm, we tested the light and temperature dependent emission algorithm, Eq. (1) of Guenther et al. (1993), with cor-

Table 2. The results of the nonlinear regression analysis of the β -caryophyllene and 1,8-cineole emission rates of branches A and B in July 2004 using three emission algorithms; the temperature dependent TEMP algorithm (Guenther et al., 1993) and two temperature and light dependent algorithms G97 (Guenther et al., 1993, Guenther, 1997) and SCHUH (Schuh et al., 1997). E_0 is the emission potential (at 30°C and 1000 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$), β is the coefficient describing the strength of the temperature dependence in the TEMP algorithm and R^2 is the regression statistic.

	Branch A			Branch B		
	E_0	β	R^2	E_0	β	R^2
β -caryophyllene						
TEMP	407±47	0.175	0.63	150±19	0.201	0.76
G97	373±41	–	0.61	122±12	–	0.70
SCHUH	391±42	–	0.62	133±13	–	0.70
1,8-cineole						
TEMP	90.1±6.4	0.180	0.82	113±11	0.180	0.82
G97	82.3±5.9	–	0.79	97.9±7.0	–	0.80
SCHUH	86.4±5.8	–	0.81	107±8	–	0.80

rected formulation from Eq. (2) of Guenther (1997), and the slightly different formulation of the light dependence presented by Schuh et al. (1997), henceforth called the G97 and SCHUH algorithms, respectively. The latter two algorithms have identical formulation, except that Schuh et al. (1997) suggest a slightly more moderate light dependence, with the light dependent environmental correction factor used in the G97 algorithm raised to the power of two in the SCHUH parameterization. The results of the regression analysis for both studied compounds and all three tested algorithms are presented in Table 2 which gives the standard emission potential E_0 (at 30°C and 1000 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$), the β coefficient indicating the strength of the temperature dependence when applicable, and the regression statistic R^2 .

The nonlinear regression fit of the July sesquiterpene data to the TEMP algorithm resulted in a reasonably good agreement, with an R^2 of 0.63 and 0.76 for branches A and B, respectively. The modelled β -caryophyllene emission rates obtained from this fit are presented in Fig. 5 together with the observations. The observed higher emissions of the debudded branch A are reflected in the standardized emission potentials, which were 407±47 $\text{ng g}^{-1} \text{h}^{-1}$ and 150±19 $\text{ng g}^{-1} \text{h}^{-1}$ for branches A and B, respectively (Table 2). These values are close to the ones given in Table 1 for the β -caryophyllene emission potentials of branches A and B in July obtained as averages from the measurements. Tarvainen et al. (2005) were not able to present late summer emission potentials for β -caryophyllene, but their values for early summer and autumn, 160±160 $\text{ng g}^{-1} \text{h}^{-1}$ and 158±295 $\text{ng g}^{-1} \text{h}^{-1}$, respectively, are very close to the values obtained in this study for the undisturbed branch B. The β coefficient values obtained in our study were of 0.18 for branch A and 0.20 for branch B, which is also in agreement

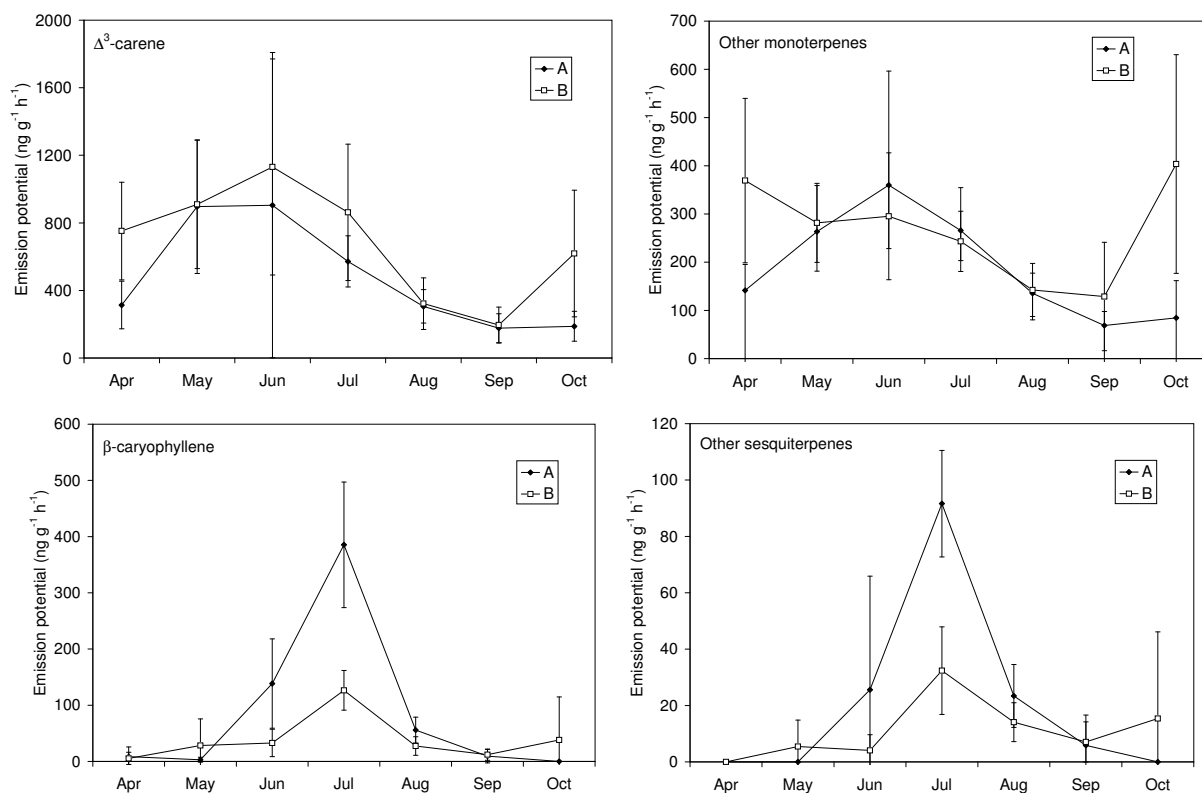


Fig. 4. Monthly mean mono- and sesquiterpene emission potentials (30°C) of the two branches with the estimated 95% confidence intervals. The main monoterpene, Δ^3 -carene is plotted separately from the rest of the monoterpenes and the main sesquiterpene, β -caryophyllene, separately from the rest of the sesquiterpenes.

with the results of Tarvainen et al. (2005) who found values 0.18 and 0.16 for early summer and autumn, respectively.

Regression fits of the July β -caryophyllene data to the G97 and SCHUH algorithms also resulted in a reasonably good agreement, with only slightly lower standard emission potentials for both branches (Table 2).

Tarvainen et al. (2005) found that the 1,8-cineole emissions measured in 2003 in Hyytiälä were equally well described by the temperature and light dependent algorithm G97 and by the temperature-only dependent TEMP algorithm. They concluded that more measurements of the 1,8-cineole emissions should be made, especially during the high summer in order to resolve the light dependence of the emissions. Our data set of 2004 had a better coverage of this high emission period but it also yielded the same ambiguous results. In addition to being able to describe the emissions equally well, all three algorithms also yielded almost identical standard emission potentials, with values ranging from $82 \pm 6 \text{ ng g}^{-1} \text{ h}^{-1}$ to $90 \pm 6 \text{ ng g}^{-1} \text{ h}^{-1}$ for branch A and from $98 \pm 7 \text{ ng g}^{-1} \text{ h}^{-1}$ to $113 \pm 11 \text{ ng g}^{-1} \text{ h}^{-1}$ for branch B (Table 2). The standard emission potentials are again very close to those given in Table 1 for the average emission potentials of 1,8-cineole for branches A and B in July. The standard emission potential reported for 1,8-cineole by Tarvainen

et al. (2005) was $68 \text{ ng g}^{-1} \text{ h}^{-1}$ over the whole vegetation period which is slightly lower than the values obtained in this study for the intense emission period in July. According to our results the β coefficient for 1,8-cineole was 0.18 for both branches, while Tarvainen et al. (2005) again found a slightly lower value of 0.14.

The remarkable similarity of the emission potentials of especially 1,8-cineole and the high R^2 values obtained by the nonlinear regression analysis using the different algorithms is due to the saturation of the light algorithm as the measurements were always conducted during high light conditions. Diurnal measurements would be needed for the validation or development of emission algorithms. In the data set collected during these experiments in Hyytiälä, the dependence of the temperature (T) of the photosynthetically active photon flux density (PPFD) was logarithmic with the form $T = 11.885 \cdot \ln(\text{PPFD}) - 59.614$ ($R^2 = 0.72$). In July 2004 the dependence was even stronger: $T = 7.8851 \cdot \ln(\text{PPFD}) - 28.085$ ($R^2 = 0.83$). The maximum temperatures in July were approximately 30°C , i.e. well below the turning point of the exponential growth of the temperature dependent correction factor of the G97 and SCHUH algorithms. Furthermore, the PPFD values in the July data ranged approximately between 500 and $1500 \mu\text{mol m}^{-2} \text{ s}^{-1}$ leading to saturation of the light al-

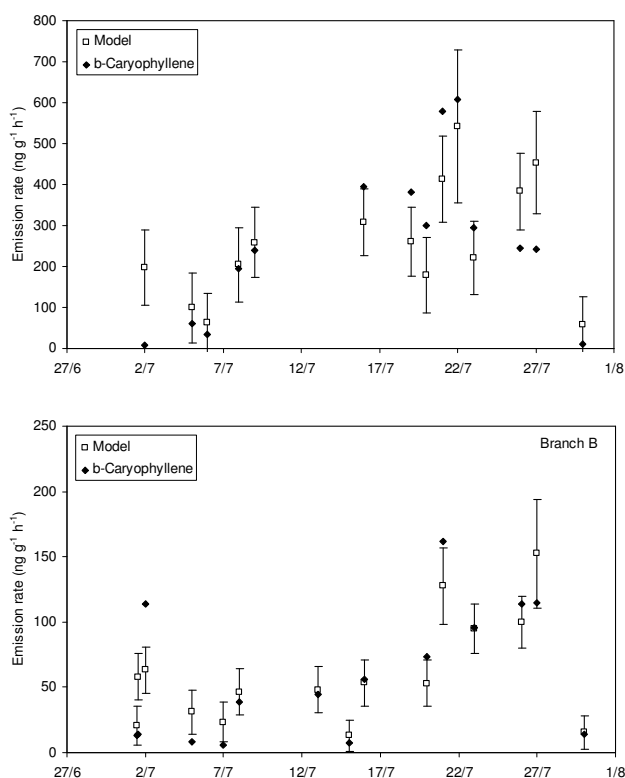


Fig. 5. Measured and modelled β -caryophyllene emission rates in Hyttiälä in July 2004. The modeled emissions were obtained by a nonlinear regression fit of the temperature dependent emission algorithm of Guenther et al. (1993) to the measured data.

gorithm in most cases. This explains why the results using the G97 and SCHUH algorithms were almost identical as in this region the light dependent environmental correction factor is quite close to unity and the effect of the power of two remains small.

4 Conclusions

The seasonal variation of monoterpene emission rates of Scots pine confirmed the earlier results; a high emission potential during early summer, declining emission potentials later in summer and early autumn and higher again later in autumn (Komenda and Koppmann, 2002; Tarvainen et al., 2005). However, this study showed that the increase of the emission potential during autumn is due to the new needles, as the emission potential of the debudded branch did not increase. The monoterpene emission pattern remained almost constant throughout the measurement period, Δ^3 -carene being the dominant monoterpene (50–70% of the VOC emission). The standard monoterpene emission potential (30°C) was highest in June (the average of the two branches $1.35 \mu\text{g g}^{-1} \text{h}^{-1}$) and lowest in September (the average of the two branches $0.20 \mu\text{g g}^{-1} \text{h}^{-1}$).

The new needles also had a much higher MBO emission potential than the needles from the previous years. The standardized MBO emission potential was more than ten times higher in the branch with new needles in June and continued to be a little higher also later in the growing season.

The sesquiterpenes were mainly emitted in the middle of the summer, the dominant sesquiterpene being β -caryophyllene. The sesquiterpene emissions were well correlated with the linalool and 1,8-cineole emissions, but not with the monoterpenes. The emission maximum occurred concomitant with the maximum concentration of the pathogen spores suggesting a potential defensive role of the sesquiterpene emissions.

It has been shown earlier that the monoterpene emissions in Hyttiälä are usually well explained by a temperature dependent emission algorithm (Tarvainen et al., 2005). In the present study, the temperature dependence of the sesquiterpene and 1,8-cineole emissions was further studied and they were found to be equally well described by the temperature dependent and the temperature and light dependent algorithms. This ambiguity of the algorithm fitting results is due to the saturation of the light algorithm as the measurements were always conducted during high light conditions.

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