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

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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 was highest during early summer in June (the average of the two branches $0.55 \mu\text{g g}_{(\text{dw})}^{-1} \text{h}^{-1}$) and lowest during early autumn in September (the average of the two branches $0.12 \mu\text{g g}_{(\text{dw})}^{-1} \text{h}^{-1}$). 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 indicating the 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-

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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 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 cuvettes in March and remained there until October. The other end of the cuvette remained open except during measurements, to ensure sufficient ventilation for the shoot in other times. 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 Teflon enclosures was approximately 20 l and they were equipped with inlet and outlet ports and a thermometer. 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. The air samples were collected onto adsorbent tubes simultaneously from both the inlet and outlet ports. The emission rate (E) 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 cuvette. 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

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Carbopack-B. The sampling time was 60 min. 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 293 K and $1000 \mu\text{mol m}^{-2} \text{ s}^{-1}$) 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 be-

low 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 growing 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 VOC emission rates

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.

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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 than the monoterpenes. In addition to β -caryophyllene, two other sesquiterpenes were detected, but they were not identified and they are referred to here as sesquiterpene1 and sesquiterpene2. 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).

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. 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 defence-related compounds and re-

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sistance 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 Hyytiälä 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 cuvette.

3.3. The effect of debudding

The increased monoterpene emissions caused by the debudding of branch A are studied in Figs. 3 and 4, where the normalized (20°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. 3). 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. 4). It is well known that in pines 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. The July average emission potentials (20°C) were 0.18 and 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 monoterpene emission potentials between the two branches did not show large deviations until autumn. In October the branch which was not debudded (B) exhibited the highest emission potential of the measurement period. It was more than six times as high as the emission potential of branch A. 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. A nonlinear regression fit similar to the one done by Tarvainen et al. (2005) of the July sesquiterpene emissions to the temperature dependent emission algorithm of Guen-

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ther et al. (1993) (Fig. 5), henceforth called 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 observed higher emissions of the debudded branch A were reflected in the standardized emission potentials (at 303.15 K and $1000 \mu\text{mol photons m}^{-2} \text{s}^{-1}$), which were $407 \pm 47 \text{ ng g}^{-1} \text{ h}^{-1}$ for branch A and $150 \pm 19 \text{ ng g}^{-1} \text{ h}^{-1}$ for branch B. When normalized to 20°C these values are close to the ones given in Table 1 for the average sesquiterpene emission potentials of branches A and B in July.

Tarvainen et al. (2005) found that the 1,8-cineole emissions measured in 2003 in Hyytiälä were equally well described by a temperature and light dependent algorithm (Guenther, 1997) and by a temperature-only dependent pool algorithm (Guenther et al., 1993). 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 the simple exponential TEMP algorithm, we tested both the light and temperature dependent emission algorithm 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 parameterisation. The results of the nonlinear regression analysis of the emission rates of branches A and B are presented in Table 2 for these three algorithms. In addition to being able to describe the emissions equally well, all three algorithms also yield almost identical standardized emission potentials (at 303.15 K and $1000 \mu\text{mol m}^{-2} \text{s}^{-1}$) for each branch. When normalised to 20°C , these values 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 remarkable similarity of the emission potentials of 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 algorithm 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 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 (20°C) was highest in June (the average of the two branches 0.55 $\mu\text{g g}_{(\text{dw})}^{-1} \text{h}^{-1}$) and lowest in September (the average of the two branches 0.12 $\mu\text{g g}_{(\text{dw})}^{-1} \text{h}^{-1}$).

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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 indicating the defensive role of the sesquiterpene emissions.

It has been shown earlier that the monoterpene emissions in Hyytiälä 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 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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Table 1. The monthly mean VOC emission potentials (20°C) of two branches (A and B) of the same *Pinus sylvestris* ($\text{ng g}_{(\text{needle dry weight})}^{-1} \text{h}^{-1}$). The emission potentials of the temperature dependent compounds are standardized to 20°C according to Guenther et al. (1993) using β value 0.09 K^{-1} . The MBO emission was standardized using the light and temperature dependent isoprene algorithm. The effect of the debudding of branch A on 13 May has been removed from the monthly mean by ignoring the measurements a week after debudding.

	April	May	June	July	August	September	October
MBO A	2	12	3	6	3	5	2
MBO B	4	14	35	15	7	8	4
1,8-Cineol A	1	18	18	27	8	0	0
1,8-Cineol B	2	12	14	29	7	0	0
linalool A	2	0	0	26	1	0	0
linalool B	3	0	0	6	1	0	0
Δ^3 -Carene A	128	361	368	232	124	72	76
Δ^3 -Carene B	306	370	460	351	131	80	434
other monoterpenes A	55	98	146	108	55	28	34
other monoterpenes B	140	101	118	99	58	52	286
β -caryophyllene A	1	1	21	60	9	1	0
β -caryophyllene B	1	4	5	19	4	2	5
other sesquiterpenes A	0	0	4	14	4	1	0
other sesquiterpenes B	0	1	1	5	2	1	2

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Table 2. The results of the nonlinear regression analysis of the 1,8-cineole and β -caryophyllene 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., 1997) and SCHUH (Schuh et al., 1997).

	Branch A			Branch B		
	E0	β	R2	E0	β	R2
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
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						

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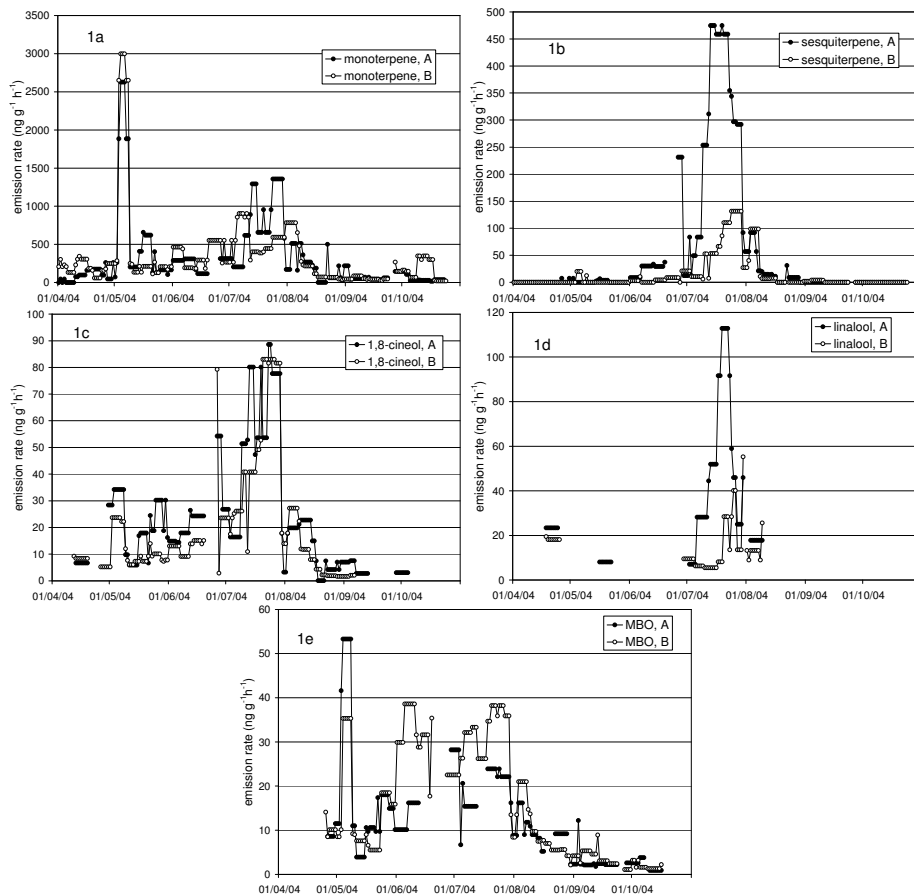


Fig. 1. 7 day running medians of the VOC emission rates from Scots pine in Hyttiälä, Finland in 2004. The samples were taken at midday from two different branches (A and B) in the same tree.

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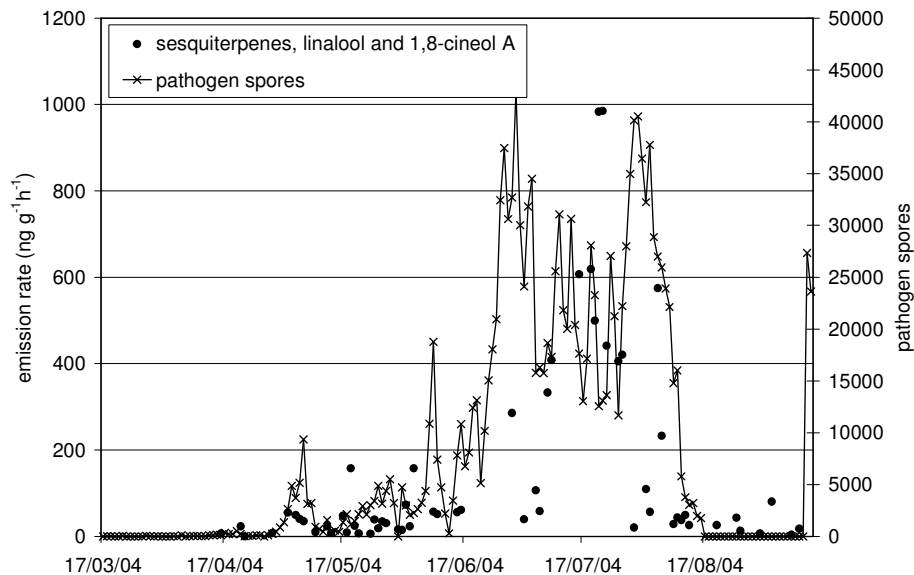


Fig. 2. The sum of the emission rates of sesquiterpenes, linalool and 1,8-cineole from branch A and the concentrations of pathogen spores measured at Hyytiälä.

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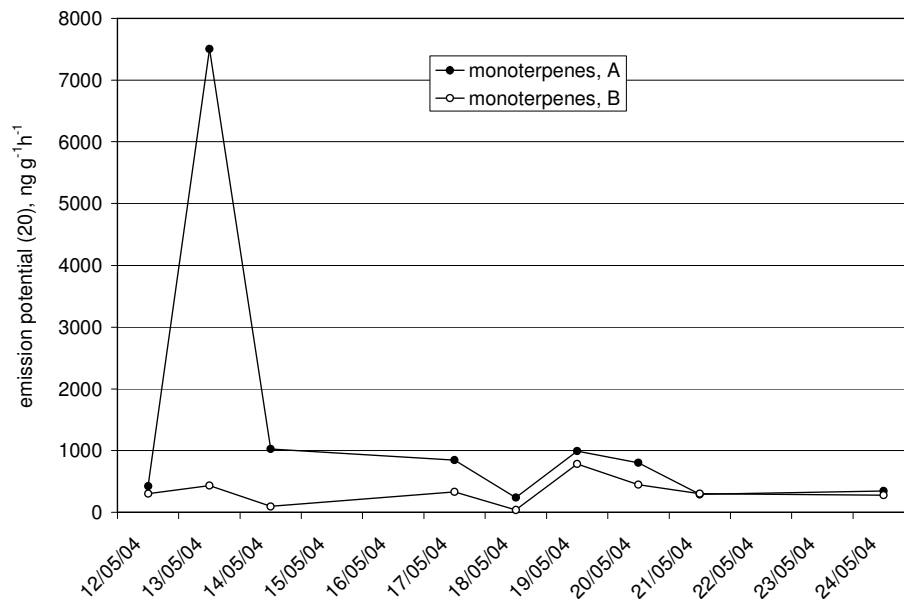


Fig. 3. The effect of the debudding of branch A on 13 May on the monoterpene emissions. Branch B was left untouched.

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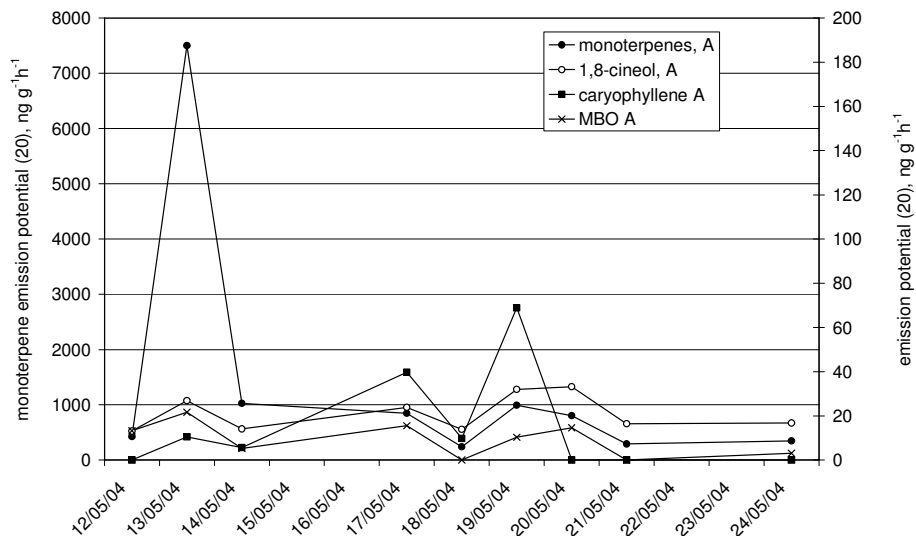


Fig. 4. The effect of the debudding of branch A on the emission potential of total monoterpenes, 1,8-cineole, β -caryophyllene and MBO.

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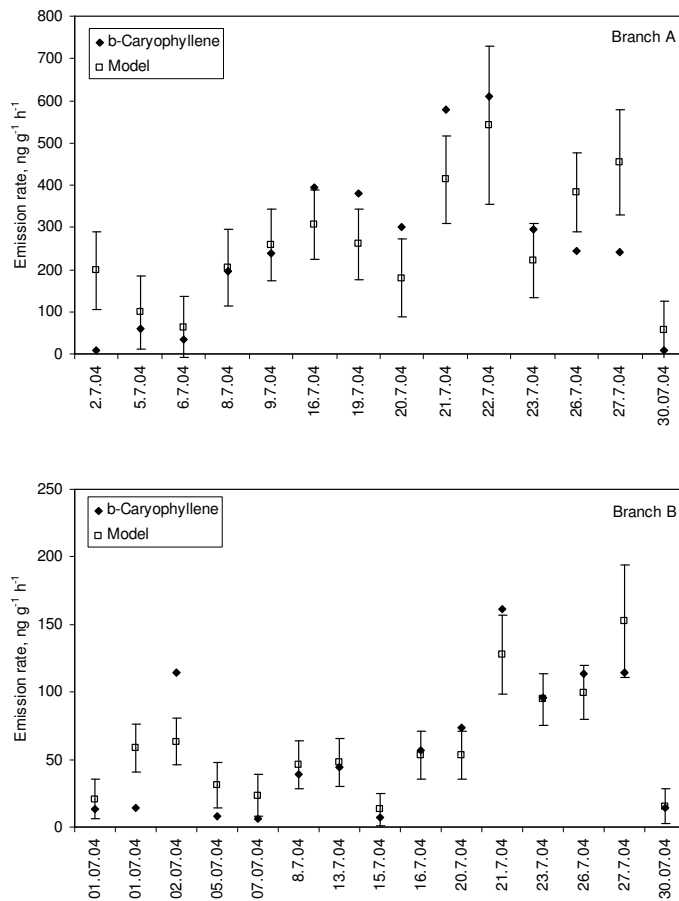


Fig. 5. Measured and modelled β -caryophyllene emission rates in Hyytiälä in July 2004. The modeled emissions were obtained by a linear regression fit of the temperature dependent emission algorithm of Guenther et al. (1993) to the measured data.

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