1	Short-term effects of thinning, clear-cutting and stump harvesting on				
2	methane exchange in a boreal forest				
3					
4	E. Sundqvist ¹ , P. Vestin ¹ , P. Crill ² , T. Persson ³ , A. Lindroth ¹				
5					
6	¹ Department of Physical geography and Ecosystem Science, Lund University				
7	² Department of Geological Sciences, Stockholm University				
8	³ Department of Ecology, Swedish University of Agricultural Sciences				
9					
10	Abstract				
11	Forest management practices can alter soil conditions, affecting the consumption and				
12	production processes that control soil methane (CH ₄) exchange. We studied the short-				
13	term effects of thinning, clear-cutting and stump harvesting on the CH ₄ exchange				
14	between soil and atmosphere at a boreal forest site in central Sweden, using an				
15	undisturbed plot as the control. Chambers in combination with a high precision laser gas				
16	analyser were used for continuous measurements. Both the undisturbed plot and the				
17	thinned plot were net sinks of CH_{4} , whereas the clear-cut plot and the stump harvested				
18	plot were net CH_4 sources. The CH_4 uptake at the thinned plot was reduced in comparison				
19	to the undisturbed plot. The shift from sink to source at the clear-cut and stump harvested				
20	plots was probably due to a rise of the water table and an increase in soil moisture,				
21	leading to lower gas diffusivity and more reduced conditions which favour CH ₄				
22	production by archea. Reduced evapotranspiration after harvesting leads to wetter soils,				

decreased CH₄ consumption and increased CH₄ production, and should be accounted for
in the CH₄ budget of managed forests.

25

26 **1. Introduction**

27 Methane (CH₄) is the second most important carbon greenhouse gas, with a radiative 28 forcing at least 25 times higher than carbon dioxide from a 100-year perspective 29 (Shindell, et al., 2009). Consumption of CH₄ by methanotrophic bacteria in the aerobic 30 part of the soil profile (Harriss et al., 1982) and production of CH₄ by archaeans in the 31 anaerobic water-saturated part of the profile (Ehhalt, 1974) and at anaerobic micro-sites 32 (von Fischer and Hedin, 2002; Kammann et al., 2009) often occur simultaneously (Le 33 Mer and Roger, 2001; Megonigal and Guenther, 2008). Generally, well-aerated forest 34 soil is a net sink of atmospheric CH₄ (Van Amstel 2012). Consumption in soils is the 35 second largest sink of CH₄ after tropospheric oxidation by hydroxyl radicals with a global sink capacity estimated recently at 28-32 Tg $CH_4 y^{-1}$ (Kirschke et al., 2013). The soil sink 36 capacity is higher in forest soils than in grasslands and arable land (Dutaur and Verchot, 37 38 2007), and therefore the global CH_4 budget is sensitive to disturbances in forests. 39 Conversion of natural forests to arable land, increased N deposition from the atmosphere, 40 and N-fertilization of agricultural lands are estimated to have reduced the global CH₄ soil 41 sink by about 30 % between 1880 and 1980 (Ojima et al., 1993). 42 43 Disturbances, including forest management practices, can also have an impact on the soil

44 CH₄ exchange by altering soil conditions such as soil moisture (Zerva and Menucuccini,

45 2005; Castro et al., 2000), water table depth (Zerva and Menucuccini, 2005) bulk density

46 (Mojeremane et al., 2012), soil temperature (Zerva and Menucuccini, 2005; Thibodeau et al., 2000), nutrient content (Smolander et al., 1998) and pH (Smolander et al., 1998). CH₄ 47 48 oxidation in soil has been observed to be controlled by diffusivity (Koschorreck and 49 Conrad, 1993; Whalen and Reeburgh, 1996; Gulledge and Schimel, 1998). A well-50 drained coarse soil facilitates the exchange of oxygen and CH₄ between the atmosphere 51 and the deeper soil levels where CH₄ is consumed (Verchot et al., 2000). By contrast, 52 increased soil moisture and soil compaction reduce the diffusivity, and promotes anoxic 53 environments in which CH₄ can be produced (Koschorreck and Conrad, 1993; Whalen 54 and Reeburgh, 1996; Gulledge and Schimel, 1998). Changes in water table depth also 55 influence the CH₄ exchange by altering the relative extent of anaerobic and aerobic zones 56 in the soil (Whalen and Reeburgh, 1990). Temperature is also an important driver of CH₄ 57 production, with higher temperatures leading to higher CH₄ production, while 58 consumption by methanotrophs is less strongly enhanced (Dunfield et al., 1993). 59 Increased nitrogen content in the soil has been shown to inhibit CH₄ consumption in 60 several studies (Steudler et al., 1989; Hutsch et al., 1993; Wang and Ineson, 2003). This 61 is due to competition by certain nitrifiers, which might occupy the same niche in the soil. 62 These nitrifiers have an enzyme similar to methanotrophs and are also able to oxidize 63 CH₄, though possibly at a lower rate (Hutsch et al., 1993).

64

Summarizing the effects of forest management practices on CH_4 exchange is difficult since relatively few studies have been made on this topic, and they have covered a range of management practices, soil types and forests. However, several studies reported that clear-cutting led to reduced CH_4 uptake, possibly due to increased soil moisture (Wu et

69	al., 2011), increased nitrogen availability (Steudler et al., 1991; Bradford et al., 2000),
70	changes in pH, (Bradford et al., 2000) and erosion (Kagotani et al., 2001). A shift from
71	soil CH ₄ sink to soil CH ₄ source has been reported due to a rise in water table depth
72	combined with increases in substrate availability (Zerva and Mencuccini, 2005) and due
73	to increases in soil moisture (Castro et al., 2000). The same shift from sink towards
74	emission has been seen following soil compaction by skid trails and machinery, as a part
75	of clear-cutting (Teepe et al., 2004) and thinning (Keller et al., 2005). One study on a
76	clear-cut drained peat soil showed no substantial changes in CH ₄ exchange (Huttunen et
77	al., 2003).
78	
79	Site preparation by mounding at clear-cuts can have a negative impact on CH ₄ exchange

80 from a climate perspective. In one study, compaction of the soil by excavators during 81 mounding increased CH_4 emissions (Mojeremane et al., 2012). CH_4 emissions from 82 stagnant water in hollows created during mounding can sometimes exceed the 83 consumption in the mineral soil on top of the mounds (Mojeremane et al., 2010). 84 However, bedding after clear-cutting has resulted in reduced CH₄ emissions (Castro et al., 85 2000). Drainage can also reduce CH₄ emissions following clear-cutting, but its positive 86 effect on CH₄ emissions was outweighed by increases in CO₂ emissions when drainage 87 was conducted on saturated peaty soils (Mojeremane et al., 2012). 88 89 Stump harvesting for bioenergy production has recently been proposed as a way of

- 90 substituting fossil fuel CO₂ emissions in Sweden. To our knowledge there are no
- 91 publications on the effects of stump harvesting on CH₄ exchange, although it is likely to

92 have a similar effect to other clear-cutting and site preparation actions. There are a few 93 studies on the effect of thinning on CH₄ exchange in a forest. Reduced CH₄ uptake due to 94 increased nitrogen availability has been reported (Thibodeau et al., 2000). A study at 95 three thinned plots in a temperate beech forest reported slightly reduced emissions at one 96 plot, whereas the other two were not significantly different from the control plots 97 (Dannenmann et al., 2007). Another study in a temperate forest actually showed an 98 increased CH₄ uptake after thinning, as opposed to a decrease at two adjacent clear-cut 99 areas (Bradford et al., 2000). Some studies found no significant changes in CH₄ exchange 100 after thinning (Wu et al., 2011; Sullivan et al., 2008).

101

102 The objective of this study was to quantify the short-term CH_4 exchange at four sites: an 103 undisturbed forest plot, a thinned forest plot, a clear-cut plot with stumps remaining, and 104 a clear-cut plot with stumps removed. The comparison between the different treatments is 105 facilitated because all four sites are within a defined area and have a common soil type. 106 We also wanted to investigate how soil moisture, soil temperature and water table depths 107 influenced the soil CH_4 exchange.

108

109 **2. Methods**

110 2.1 Site description

111 The CH₄ exchange measurements took place in a forested area on the southern edge of

112 the boreal zoon, at Norunda research station in central Sweden, $60^{\circ}05'$ N, $17^{\circ}29'$ E.

113 Hourly automated chamber measurements were made using a system that was moved

between 4 differently managed plots (Fig.1). One plot contained undisturbed 120-year-

old mixed pine (*Pinus sylvestris*) and spruce (*Picea abies*) forest, which had not been thinned or fertilized in several decades. The other three plots were recently (2009-2010) impacted by either thinning, clear-cutting or stump harvesting. Thinning was done in order to simulate continuous cover forestry, rather than to increase growth.

119 Measurements were made using four chambers at the thinned plot, and five chambers at each of the other plots. The chamber locations were named U1-U5 at the 120 121 undisturbed plot, T1-T4 at the thinned plot, C1-C5 at the clear-cut plot and S1-S5 at the 122 stump harvested plot. At the clear-cut and stump harvested plots half of the chamber 123 frames were positioned on bare soil, where organic and mineral soil layers were mixed. 124 The disturbance was caused either by stump harvesting, or by site preparation to facilitate 125 the establishment and growth of new plants. The remaining frames were placed on soil surfaces with intact vegetation. The clear-cut and stump harvested plots had been 126 127 fertilized in 1976, 1988 and 1998.

Ground vegetation was sparse and dominated by bilberry (*Vaccinium myrtillus*) and feather mosses (*Hylocomium splendens* and *Pleurozium schreberi*). There were more shrubs and grass at the clear-cut site, following the soil's disturbance. The soil was a glacial till (Lundin et al., 1999) with an organic layer of 3-10 cm depth. For the period 1980-2010, the mean air temperature was 6.5 °C, and the mean annual precipitation was 576 mm (measured 30 km south of Norunda).

134

135 2.2 Timing of measurements

136 Thinning took place in November 2008, the clear-cutting in February 2009 and stump137 harvesting in May 2010. Both the clear-cut plot and the stump harvested plot were

mounded and planted in May 2010. The chamber frames were installed in 2005 at the undisturbed and thinned plots, and in June 2010 at the clear-cut and stump harvested plots, to allow time for soil and vegetation to recover from the disturbance.

Due to equipment limitations, measurements were conducted at one plot at a time. Measurements at the thinned plot were made from 1 August 2009 to 31 May 2010, at the undisturbed plot from 07 July 2010 to 04 October 2010, at the stump-harvested plot from 07 October 2010 to 20 October 2010 and at the clear-cut plot from 21 October to 9 November 2010. Winter data at the thinned plot from 01 December 2009 to 14 April 2010 were not used in the analyses due to uncertainties in the measurements caused by snow and frost.

148

149 2.3 Equipment

150 We used automated, transparent chambers of Polymethyl methacrylate in combination 151 with a high precision off-axis integrated cavity output spectroscopy (ICOS) laser gasanalyser (DLT-100, Los Gatos Research (LGR)) for simultaneous concentration 152 153 measurements of CH₄, CO₂ and H₂O. The chambers had a volume of 110 litres and covered a surface-area of 0.2 m^2 . Gas concentrations in the chambers were measured 154 155 after closure by recirculating the air through the gas analyser for 6 min. The flow rate 156 between chambers and manifolds was 8-10 l/min. This air stream was sub-sampled and 157 passed through the analyser at a flow rate of 1.2 l/min. A fan was installed in each 158 chamber, designed to ensure sufficient mixing of chamber headspace air without disturbing the laminar boundary layer at the ground. Soil moisture was measured in the 159 160 chambers at 0-5 cm depth with a MI-2x thetaProbe from DeltaT Devices. The soil

161 temperature was measured at 5 cm depth inside the chambers using a type T 162 thermocouple. Soil temperature measurements at the thinned plot did not work properly 163 and so temperature data from the undisturbed plot, 125 meters away, was used instead.

164

166 There were differences in height between the chamber frames relative to the ground 167 water table. One pipe with continuous measurements of the ground water table was 168 located 125 m from the thinned plot and 30 meters from the undisturbed plot (Fig. 1). The 169 groundwater table at these plots was treated as horizontal. At the clear-cut and stump harvested plots, the ground water table was measured manually in seven pipes at each 170 plot, on the 8th and 20th of October 2010 and 2nd of November 2010. Some of these pipes 171 172 are shown in Fig.1. An inverse distance-weighting model was used to calculate the height of the ground water table in relation to the ground surface for 40 m^2 areas surrounding the 173 174 chamber frames. The ground water table was also measured continuously at one position 175 on the clear-cut plot.

176

177 2.5 Soil sampling

Soil samples were taken in order to determine organic carbon (C) and nitrogen (N) content and pH in the top 20 cm of the soil including the humus layer, where the chambers had been positioned. The litter layer was not taken into account. Sampling was done in November 2010 at the clear-cut and stump harvested plots and in September 2012 at the undisturbed and thinned plots.

184 At the clear-cut and stump harvested plots, humus layer samples were taken, down to the 185 border between organic and mineral soil layers, using a 10 cm x 10 cm quadratic frame. The mineral soil was sampled with a 15.9 cm^2 steel corer to a depth of 20 cm, but was 186 187 subdivided in the field into 0-10 and 10-20 cm layers. Humus samples were treated 188 individually, while the mineral soil samples were pooled plot-wise for each soil layer. The samples, folded in plastic bags, were transported in cooling boxes to the laboratory, 189 190 where they were kept fresh at 4-5°C during the preparation process before the final 191 analyses.

192

193 Soil samples were passed through either a 5 mm (humus samples) or a 2 mm (mineral 194 soil) mesh. Stones and gravel >2 mm diameter not passing the mesh were always rejected, as were any roots. The sieved soil material from each sample was carefully 195 196 mixed and divided into a number of sub-samples for determination of soil pH (H₂O), and 197 total C and N content. Fresh weight/dry weight ratios were determined after drying the 198 sub-samples at 105 °C for 24 h. Soil layer pH was determined with a glass electrode in 199 the supernatant after shaking for 2 h on a rotary shaker, and sedimentation in an open 200 flask for another 22 h. The proportion of fresh soil to distilled water was 1:1 by volume, 201 compared to about 1:10 for dry matter to water for humus, and 1:2.5 for mineral soil). 202 Total C and N content were determined, using vacuum-dried soil samples at 60 °C for 24 h, in a Carlo-Erba NA 1500 Analyser. Because soil pH was always below 6, we assumed 203 204 that there was no carbonate C, and all C analysed was assumed to be organic C.

At the undisturbed and thinned plots a cylindrical metal corer with an 11 cm² opening was pressed horizontally into the humus layer, and also at 5 cm and 10 cm depth in the mineral layer. At some of the measurement locations (T1, T4, U2, U6) large stones, rocks and roots occupied a large volume of the mineral soil so that sampling at 10 cm depth in the mineral soil was not possible. The soil samples were kept below 5°C until they were analysed.

212

The total amount of C and N in the soil samples at the undisturbed and thinned plots were analysed with an element analyser (Elementar Analysensysteme GmbH, Germany). The pH value was measured after two hours equilibration with a 0.1 M barium chloride solution (Orion Research model Microprocessor ionalyzer/901). The extractions were made on fresh material. Before determining the bulk density, the samples were oven dried for 48 hours at 100 °C and then sieved through a 2 mm mesh.

219

220 2.6 Data analyses

221 The rate of change of CH₄ concentration (dC_CH₄/dt) within the chamber was calculated using a linear fit to the first two minutes of concentration data measured by the gas 222 analyser, beginning immediately after chamber closure. We calculated the r^2 values for 223 224 the fits of five different slopes, which were lagged at 10 seconds intervals after chamber closure. The fit with the highest r^2 value was then selected. The CH₄ flux ($J_{CH4 flux}$) was 225 calculated as $J_{CH4\,flux} = \frac{dC}{dt}\frac{V}{A}$, where C is the molar density (µmol m⁻³), V (m³) is the 226 chamber volume and A (m²) is ground surface area. Fluxes with an r^2 value higher than 227 0.3 were generally kept for further analyses. An r^2 of 0.3 was the limit when the fluxes 228

visually sorted out based on normalized root mean square error. Data kept for further
analyses corresponded to 98 % of the data at the undisturbed plot, 97 % of the data at the
thinned plot, 84 % of the data at the clear-cut plot and 77 % of the data at the stump
harvested plot.

were significantly different from zero. A few outliers that passed the r^2 limit were

234

229

 $MDF = \frac{\sigma}{t}$, where t is the Minimum flux detection limit (MDF) was calculated as 235 measurement time for one specific measurement and σ is the standard deviation for the 236 concentration measurement. For a chamber the size as used in this study, the MDF for a 237 single measurement was 2.8 μ mol m⁻² h⁻¹. For daily average values of hourly 238 measurements this value is reduced to $< 1 \mu mol m^{-2}h^{-1}$ since the MDF value should be 239 divided by the square root of the number of measurements. It is important to note that 240 241 while fluxes below the MDF cannot be securely detected, they must still be considered. 242 For example, consider time series where fluxes decrease smoothly from an emission peak 243 to an uptake. In the transition phase from net emissions to net uptake, fluxes will be close 244 to zero. Removing fluxes <MDF could possible bias the result towards a stronger sink or source than what times series from the individual chambers give support for. Therefore 245 246 also the fluxes within the MDF interval will be kept in the analyses. Removing fluxes within the MDF interval for a single measurement ($\pm 2.8 \ \mu mol \ m^{-2} h^{-1}$) would lead to a 247 248 decrease in the number of flux measurements by 0%, 16%, 13% and 30% for the 249 undisturbed, thinned, clear-cut and stump harvested plots respectively and not change the mean exchange of CH₄ at any of the plots by more than 0.6 μ mol m⁻² h⁻¹. 250 251

252 Correction of the measured CH_4 concentrations for dilution by water vapour was only 253 possible at the undisturbed, clear-cut and stump harvested plots after water vapour 254 measurements started in June 2010. This means that daytime data (global radiation > 20 255 W/m^2) from the thinned plot had to be excluded from the analyses. During night the 256 dilution effect had very little impact.

257

258 The impact of the environmental variables soil temperature, soil moisture, and water table depth on CH₄ exchange was analysed separately by Spearman linear correlations using 259 the corr function, and by multiple linear regression on standardized data using the 260 261 function stepwisefit (both Matlab version R2009b). The stepwise regression analyses 262 were performed by bi-directional elimination. P-values were used in the selection process. The analysis was made on standardized data to adjust for the disparity in 263 264 variable sizes, which makes the outcome of the analyses, the coefficients, comparable. 265 The coefficients would be useful in modelling of CH₄ exchange. A variable with a larger 266 coefficient has a higher impact on the CH_4 exchange. Standardization for a data point x_i was made by $x_i = \frac{x_i - \overline{x}}{\sigma}$ where \overline{x} is the average of all data points and σ is the standard 267 deviation of all data. An R^2 value for the overall model was also calculated showing how 268 much of the variance in CH₄ exchange that is not explained by the environmental 269 270 variables included in the analyses. 271

The significance of mean values at the measurement locations was calculated with thettest function (also Matlab version R2009b).

275 **3. Results**

276 *3.1 Environmental conditions*

277 There were differences in soil moisture and soil temperatures among the plots. On 278 average the undisturbed forest plot, with measurements exclusively from the summer 279 season, July through September, had the driest and warmest records, and also the 280 measurement locations were further above the ground water table than at other plots 281 (Fig.2, Table 1). The clear-cut plot, which was measured in October and November, 282 showed the coldest and wettest conditions including the highest water table. Four of the 283 five measurement locations at this plot were on average less than 15 cm above the ground 284 water table (Fig.2, Table 1). The thinned plot and the stump harvested plot had similar 285 average moisture and temperature conditions, but the measurements at the thinned plot proceeded over a longer time period and thus the conditions varied more. The thinned 286 287 plot also had a generally deeper ground water table than the stump harvested plot (Fig.2, 288 Table 1).

289 Soil N and C content and pH were higher at the clear-cut and stump harvested 290 plots than at the undisturbed and thinned plots (Table 1).

291

3.2 CH₄ exchange

The mean CH_4 exchange of all measurement locations within the plots were as follows: the undisturbed plot and the thinned plot were net CH_4 sinks of -10 µmol m⁻² h⁻¹ and -5 µmol m⁻² h⁻¹ respectively, while the clear-cut plot and at the stump harvested plot were net sources of 13.6 µmol m⁻² h⁻¹ and 17 µmol m⁻² h⁻¹, respectively (Fig.2). However, the CH_4 exchange varied within the plots. At the clear-cut and stump harvested plots, both 298 net sources and net sinks existed (Fig.3). Plot T_3 and T_4 at the thinned plot shifted

between net daily CH₄ sinks and net daily CH₄ sources on a few occasions (Fig.3b).

300 Fluxes ranged from -7.2 to -11.6 μ mol m⁻² h⁻¹ at the undisturbed plot, from -0.3 to -8.6

301 μ mol m⁻² h⁻¹ at the thinned plot, from -3.0 to 32.5 μ mol m⁻² h⁻¹ at the clear-cut plot and 302 from -2.9 to 74.0 μ mol m⁻² h⁻¹ at the stump harvested plot (Fig.3).

303

304 3.3 Drivers of CH₄ exchange at the undisturbed and thinned plots

305 Linear regression analyses between CH₄ exchange and climatic variables showed that for

306 most measurement locations at the undisturbed and thinned plots, consumption

307 significantly (p<0.001) increased with decreasing soil water content, decreasing water

308 table depth and increasing temperatures. Exceptions to this were net CH₄ uptake at

309 locations T₃ and T₄ which decreased with increasing temperatures, and net CH₄ uptake at

310 locations T₂ and T₄, which decreased with decreasing soil moisture (Table 2). Figure 5

311 shows an example of the CH_4 exchange response to temperature and soil water conditions 312 at plot U_4 .

313 Monthly multiple linear regression analyses (Table 3) added some temporal 314 information to the CH₄ exchange at the undisturbed and thinned plots. At the undisturbed 315 plot the water table depth affected CH_4 consumption in August. In September 2010 316 temperature was the most influential variable at all measurement locations. In July 2010 317 the result was less distinct, showing some measurement locations with a higher 318 dependency on water table depth and soil moisture, and some measurement locations 319 with a higher dependency on temperature. The clearest result at the thinned plot was a 320 dependency on soil moisture at measurement locations T1 and T3 in August 2009 and at

321 locations T_1 and T_2 in April 2010 (Table 3). Soils were wetter than average in August and 322 April due to heavy rains in June and July 2009, and snowmelt in spring 2010. However, 323 according to the r² value of the overall model there are lot of unexplained variance in the 324 CH₄ exchange at all measurement locations.

325

326 3.4 Drivers of CH₄ exchange at the clear-cut and stump harvested plots

Generally at the clear-cut and stump harvested plots, the measurement locations with net emissions of CH_4 had either a relatively short distance to water table, or were disturbed by site preparation, or both, although there were exceptions. Plot S_4 and S_5 had the same water table depth and were not disturbed by site preparation, but plot S_4 was a CH_4 sink while plot S_5 was a CH_4 source (Fig.4).

At the majority of the measurement locations on the clear-cut and stump 332 333 harvested plots, higher temperatures correlated significantly (p<0.05) with lower CH₄ 334 emissions, or in one case with a higher net uptake. Both negative and positive significant 335 correlations between CH₄ exchange and soil moisture was found at a few measurement 336 locations but the soil moisture range at those measurement locations was very small. At 337 two measurement locations with net emissions at the clear-cut plot, there was a 338 significant (p<0.05) negative correlation between CH_4 exchange and water table depth, so 339 that a deeper water table depth gave higher CH₄ emissions (Table 2). The multiple linear 340 regression confirmed the significantly negative correlation between CH₄ exchange and 341 temperature at 6 measurement locations.

343 **4. Discussion**

344 All measurement locations at the undisturbed forest plot were sinks of CH₄ throughout 345 the measurement period, which is consistent with the generally drained, drier and warmer 346 soil conditions at the plot (Fig.2). The measurement locations at the thinned plot were 347 also net sinks of CH₄, although reduced in comparison to the undisturbed plot. By 348 contrast, the clear-cut and stump harvested plots were net sources of CH₄. Since the 349 measurements at the different plots were conducted at different times of the year, 350 seasonality and annual variations can probably explain some of the differences in CH₄ 351 exchange and soil conditions. However, it is not likely that differences in water table 352 depth between the plots are due solely to seasonal variations. In the autumn of 2010 the 353 water table was on average more than 1 m higher at the clear-cut and stump harvested 354 plots than at the undisturbed plot. In addition to this, the mean CH_4 exchange for the 355 autumn period October to November at the thinned site did not differ much from the 356 mean CH₄ exchange for the whole measurement period, indicating that average seasonal 357 variations are small (Fig.2). Precipitation was on average higher during the measurement 358 period at the thinned site than during measurements at the other plots, which did not 359 cause a switch from CH₄ sink to CH₄ source. The clear-cut and stump harvested plots are 360 located on a plateau which is uphill from the thinned and undisturbed plots and hence 361 topography should not be responsible for the higher water table at the clear-cut and stump 362 harvested plots (Fig.1).

Water table depth, soil moisture and soil temperature were all shown to be important drivers of CH₄ exchange, as demonstrated by the linear and multiple linear regression analyses. However it appears that the rise of the water table and increased soil

366	moisture caused some of the measurement locations to shift to CH ₄ sources. This is			
367	consistent with results by Zerva and Menucuccini (2005) and Castro et al, (2000).			
368	Temporal shifts to CH ₄ emissions after snowmelt and summer precipitation, as were seen			
369	at measurement locations T_3 and T_4 , were also reported by Wang and Bettany, (1995).			
370	A majority of net emitting measurement locations at the clear-cut and stump			
371	harvested plots (C_2 , C_3 , C_4 , C_5 , S_1) were positioned less than 21 cm above the water table,			
372	and had a volumetric soil moisture content above 40% (Table 1). Also measurement			
373	location T ₃ , when it had temporarily shifted to a CH ₄ source, had volumetric soil moisture			
374	content above 40%. Net emissions were also measured at measurement location $S_{\rm 2}$ and $S_{\rm 5}$			
375	with water table depths at 30-40 cm and volumetric soil moisture contents of 23-40%.			
376	Fiedler and Sommer (2000) found a threshold value of water table depth at 15 cm, below			
377	which only minor annual emissions were measured. The three measurement locations at			
378	the clear-cut and stump harvested plots which showed net consumption of CH_4 were			
379	further than average above the water table for those plots (Fig.4).			
380				
381	Temperature seemed to have a stronger impact on CH ₄ exchange in drier conditions.			
382	Figure 5 illustrates a high correlation, $r^2 = 0.74$, between soil temperature and CH ₄			
383	exchange at measurement location $U_{4,}$ when excluding data points with soil moisture			
384	above 22% and a distance to the water table of less than 1.25 m. The threshold value of			
385	22% was selected after visual inspection of the data. If all the data from wetter conditions			
386	were included (volumetric soil moisture content > 22 % and water table < 1.25 m away),			
387	the corresponding r^2 equals 0.47. This is consistent with the results from the multiple			
388	linear regression analyses showing that water table depth had a significant impact on the			

 CH_4 exchange at all measurement locations in August 2010, when the water table depth varied strongly. In contrast, during September, there were no major precipitation events and soil temperature was the most influential variable. Soil moisture was rarely below 30 % at the thinned plot, thus the temperature dependence was less. In autumn, September to November 2009, all measurement locations at the thinned plot were stable sinks of CH₄ even though the soil temperature was at times below 5°C.

395 At the clear-cut and stump harvested plots, where most measurement locations were net 396 sources of CH₄, we would expect a positive correlation between soil temperature and CH₄ 397 exchange, so that higher temperatures led to higher net emissions of CH_4 . Methanogens 398 generally respond better than methanotrophs to increased temperatures (Dunfield et al., 399 1993). However this was not the case: a majority of the measurement locations showed a 400 significantly negative correlation between temperature and CH₄ exchange. The result is 401 difficult to explain since CH₄ production and oxidation are not measured separately. Soil 402 temperature profiles at the clear-cut and stump harvested plots (data not shown) show 403 that during the measurement period, changes in surface temperature, associated with 404 periods of cloudy conditions and precipitation, at 5 cm depth are larger than at 20 and 40 405 cm depth. Methanotrophs are expected to be located closer to the soil surface than 406 methanogens and the larger temperature increase at the surface might compensate their 407 lower response to temperature, which could explain why net CH₄ exchange is negatively 408 correlated to soil temperature during this period.

409

410 The highest CH_4 emissions were found at four of the five disturbed measurement

411 locations: that is, sites of bare soil where organic and mineral soils were mixed. The soil

412 at disturbed measurement locations seemed less compact than at measurement locations 413 with intact vegetation, so the disturbance probably did not inhibit diffusion. Possibly the 414 availability of fresh organic material was higher at disturbed measurement locations. 415 Fresh, labile organic matter would promote heterotrophic uptake of O_2 and increase the 416 soil's water retention, thereby promoting the activity of methanogenic archeans 417 (Wachinger et al., 2000). The one disturbed measurement location, which showed net 418 CH₄ consumption, S₃, was positioned on top of a mound with relatively large distance to 419 the ground water table (Fig.4). 420 421 Since this is a study of the short-term effects of forest management practices on CH₄ 422 exchange, there are no data on how long-lived these effects are. Sudden shifts from sinks 423 to sources and back again due to changes in soil water conditions are evident, as we have 424 seen at the thinned plot (Fig.3b). It might take years (Tate et al., 2006) to several decades 425 for a soil to regain its full sink capacity. The recovery time for the soil CH₄ sink strength 426 of forests on abandoned agricultural land was more than 100 years (Prieme et al., 1997; 427 Smith et al., 2000). Increasing CH₄ uptake with time after afforestation can be an effect 428 of an increase in the population of CH₄ oxidizing bacteria with time (Barcena et al.,

429 2014) or better soil diffusivity and soil aeration with time (Christiansen & Gundersen,

430 2011; Peichl et al., 2010). A better soil aeration with time could be due to an increase in

431 root biomass, which means that the roots over time loosen the soil and absorb more water

432 (Peichl et al., 2010). Hiltbrunner et al, (2012) found that the soil CH₄ sink capacity of

433 abandoned agricultural land increased with stand age up to 120 years, due to the

- increased transpiration of older forests and their ability to shield the forest floor fromprecipitation, which resulted in more favourable conditions for methanotrophic activity.
- 436

437 Uptake rates by forest landscapes might be overestimated (Grunwald et al., 2012; Fiedler et al., 2005). A study by Grunwald et al, (2012) found that wet forests were as important 438 439 as wetlands for the CH_4 budget of European forests, and Fiedler et al. (2005) found that if 440 2.3% of a forest area consisted of wet soil the forest could turn from a sink to a source of 441 CH₄. As mentioned, after clear-cutting, water table depth decreased and soil moisture 442 increased. Wetter soils and a higher ground water table are common consequences of 443 clear-cutting, and it is therefore important to consider their impact on the CH₄ budget in 444 managed forests, especially if the recovery time for the soil CH₄ sink is several decades. 445 In this study the effects of thinning on the CH₄ exchange were not as pronounced as for 446 clear-cutting, although the plot average consumption was reduced in comparison to the 447 undisturbed plot. Any forest management practice that reduces disturbance and leaves a 448 continuous forest cover might be a better alternative from a global warming perspective. 449

450 **5. Conclusions**

451 Our study on the short term effects of boreal forest management on CH_4 exchange shows 452 that the undisturbed plot and the thinned plot remained net CH_4 sinks, while the clear-cut 453 and stump harvested plots were net CH_4 sources. Linear regression analyses between CH_4 454 exchange and climatic variables showed that for most measurement locations at the 455 undisturbed and thinned plots, net CH_4 uptake increased significantly with decreasing soil 456 moisture, decreasing water table depth and increasing temperatures. A higher water table

457	and increased soil moisture were likely to be responsible for the shift to CH_4 emissions at				
458	the clear-cut and stump harvested plots. At most of the measurement locations, which				
459	showed net emissions, the soil was almost saturated and the water table was within a few				
460	decimetres of the soil surface. Clear-cutting of the forest resulted in a raised ground water				
461	table and in increased soil moisture. These effects should be accounted for in the CH_4				
462	budget of managed forests.				
463					
464	Acknowledgement				
465	Support for this work was provided by Formas and by the Linnaeus Centre LUCCI				
466	(http://www.lucci.lu.se/index.html) funded by the Swedish Research Council. Airborne				
467	LiDAR for the Norunda site, the basis for the digital elevation model, was acquired with				
468	support from the British Natural Environment Research Council (NERC/ARSF/FSF grant				
469	EU10-01 and NERC/GEF grant 933). We thank Anders Båth and Tomas Karlsson for				
470	field assistance.				
471					
472	References				
473	Bárcena, T.G., D'Imperio, L., Gundersen, P., Vesterdal, L., Priemé, A., and Christiansen,				
474	J.R., Conversion of cropland to forest increases soil CH4 oxidation and abundance of				
475	CH4 oxidizing bacteria with stand age: APPL SOIL ECOL, 79, 49-58, 2014				
476					
477	Bradford, M.A., Ineson, P., Wookey, P.A., and Lappin-Scott, H.M., Soil CH4 oxidation:				
478	response to forest clearcutting and thinning. SOIL BIOL BIOCHEM				
479	32(7), 1035-1038, 2000				

481	Castro, M.S., Gholz, H.L., Clark, K.K and Steudler, P.A., Effects of forest harvesting on
482	soil methane fluxes in Florida slash pine plantations. CAN J FOREST RES, 30(10),
483	1534-1542, 2000
484	
485	Christiansen, J.R., and Gundersen, P., Stand age and tree species affect N2O and CH4
486	exchange from afforested soils. BIOGEOSCIENCES, 8, 2535-2546, 2011
487	
488	Dannenmann, M., Gasche, R., Ledebuhr, A., Holst, T,. Mayer, H and Papen, H., The
489	effect of forest management on trace gas exchange at the pedosphere-atmosphere
490	interface in beech (Fagus sylvatica L.) forests stocking on calcareous soils. EUR J
491	FOREST RES, 126(2), 331-346, 2007
492	
493	Dunfield, P., Knowles, R., Dumont, R and Moore, T.R., Methane production and
494	consumption in temperature and sub-arctic pear soils-response to temperature and pH.
495	SOIL BIOL BIOCHEM, 25(3), 321-326, 1993
496	
497	Dutaur, L. and Verchot, L.V., A global inventory of the soil CH(4) sink. GLOBAL
498	BIOGEOCHEM CY, 21(4), 2007
499	
500	Ehhalt, D.H., Atmospheric cycle of methane. TELLUS, 26(1-2), 58-70, 1974
501	

502	Von Fischer, J.C. and Hedin, L.O., Separating methane production and consumption with				
503	a field-based isotope pool dilution technique. GLOBAL BIOGEOCHEM CY, 16(3), 8/1-				
504	8/13, 2002				
505					
506	Fiedler, S. and Sommer, M., Methane emissions, groundwater levels and redox potentials				
507	of common wetland soils in a temperate-humid climate. GLOBAL BIOGEOCHEM CY,				
508	14(4), 1081-1093, 2000				
509					
510	Fiedler, S., Holl, B.S. and Jungkunst, H.F., Methane budget of a Black Forest spruce				
511	ecosystem considering soil pattern. BIOGEOCHEMISTRY, 76(1), 1-20, 2005				
512					
513	Grunwald, D., Fender, A. C., Erasmi, S., and Jungkunst, H. F., Towards improved				
514	bottom-up inventories of methane from the European land surface, ATMOS ENVIRON,				
515	51, 203-211, 2012				
516					
517	Gulledge, J. and Schimel, J.P., Moisture control over atmospheric CH4 consumption and				
518	CO2 production in diverse Alaskan soils. SOIL BIOL BIOCHEM, 30(8-9), 1127-1132,				
519	1998				
520					
521	Harriss, R. C., Sebacher, D. I., and Day, F. P., Methane flux in the great dismal swamp,				
522	NATURE, 297, 673-674, 1982				
523					

- 524 Hiltbrunner, D., Zimmermann, S., Karbin, S., Hagedorn, F., and Niklaus, P. A.,
- 525 Increasing soil methane sink along a 120-year afforestation chronosequence is driven by
- soil moisture, GLOBAL CHANGE BIOL, 18, 3664-3671, 2012
- 527
- 528 Hutsch, B.W., C.P. Webster, and Powlson D.S., Long term effects of nitrogen-
- 529 fertilization on methane oxidation in soil of the broadbalk wheat experiment. SOIL BIOL
- 530 BIOCHEM, 25(10), 1307-1315, 1993
- 531
- 532 Huttunen, J. T., Nykanen, H., Martikainen, P. J., and Nieminen, M., Fluxes of nitrous
- 533 oxide and methane from drained peatlands following forest clear-felling in southern
- 534 Finland, PLANT SOIL, 255, 457-462, 2003
- 535
- 536 Kagotani, Y., Hamabata, E., and Nakajima, T., Seasonal and spatial variations and the
- 537 effects of clear-cutting in the methane absorption rates of a temperate forest soil. NUTR
- 538 CYCL AGROECOSYS, 59(2)169-175, 2001
- 539
- 540 Kammann, C., Hepp, S., Lenhart, K., and Muller, C., Stimulation of methane
- 541 consumption by endogenous CH4 production in aerobic grassland soil, SOIL BIOL
- 542 BIOCHEM, 41, 622-629, 2009
- 543
- 544 Keller, M., Varner, R., Dias, J. D., Silva, H., Crill, P., and de Oliveira, R. C., Soil-
- 545 atmosphere exchange of nitrous oxide, nitric oxide, methane, and carbon dioxide in

546	10g
540	10g

logged and undisturbed forest in the Tapajos National Forest, Brazil, EARTH

- 547 INTERACT. 9, 1-28, 2005
- 548
- 549 Kirschke, S., Bousquet, P., Ciais, P. Saunois, M., Canadell, J.G., Dlugokencky, E.J.,
- 550 Bergamaschi, P., Bergmann, D., Blake, D.R., Bruhwiler, L., Cameron-Smith, P.,
- 551 Castaldi, S., Chevallier, F., Feng, L., Fraser, A., Heimann, M., Hodson, E.L., Houweling,
- 552 S., Josse, B., Fraser, P.J., Krummel1, P.B., Lamarque, J-F., Langenfelds, R.L., Quéré,
- 553 C.L., Naik, V., O'Doherty, S., Palmer, P.I., Pison, I., Plummer, D., Poulter, B., Prinn,
- 554 R.G., Rigby, M., Ringeval1, B., Santini, M., Schmidt, M., Shindell, D.T., Simpson, I.J.,
- 555 Spahni, R., Steele, L.P., Strode, S.A., Sudo, S., Szopa, S., van der Werf, G.R.,
- 556 Voulgarakis, A., van Weele, M., Weiss, R.F., Williams, J.E., and Zeng, G., Three
- decades of global methane sources and sinks. NAT GEOSCI, 6, 813-823, 2013
- 558
- 559 Koschorreck, M. and Conrad, R., Oxidation of atmospheric methane in soil-
- 560 measurements in the field, in soil cores and in soil samples. GLOBAL BIOGEOCHEM
- 561 CY, 7(1), 109-121. 1993
- 562
- 563 Le Mer, J. and Roger, P., Production, oxidation, emission and consumption of methane
- 564 by soils: A review. EUR J SOIL BIOL, 37(1), 25-50, 2001
- 565
- 566 Lundin, L. C., Halldin, S., Lindroth, A., Cienciala, E., Grelle, A., Hjelm, P., Kellner, E.,
- 567 Lundberg, A., Molder, M., Moren, A. S., Nord, T., Seibert, J., and Stahli, M., Continuous

- 568 long-term measurements of soil-plant-atmosphere variables at a forest site, AGR
- 569 FOREST METEOROL, 98-9, 53-73, 1999
- 570
- 571 Megonigal, J.P. and Guenther, A.B., Methane emissions from upland forest soils and
- 572 vegetation. TREE PHYSIOL, 28(4), 491-498, 2008
- 573
- 574 Mojeremane, W., Rees, R.M and Mencuccini, M., Effects of site preparation for
- 575 afforestation on methane fluxes at Harwood Forest, NE England.
- 576 BIOGEOCHEMISTRY, 97(1), 89-107, 2010
- 577
- 578 Mojeremane, W., Rees, R.M and Mencuccini, M., The effects of site preparation
- 579 practices on carbon dioxide, methane and nitrous oxide fluxes from a peaty gley soil.
- 580 FORESTRY, 85(1), 1-15, 2012
- 581
- 582 Ojima, D. S., Valentine, D. W., Mosier, A. R., Parton, W. J., and Schimel, D. S., Effect
- 583 of land-use change on methane oxidation in temperate forest and grassland soils,
- 584 CHEMOSPHERE, 26, 675-685, 1993
- 585
- 586 Peichl, M., Arain, M.A., Ullah, S., and Moore, T.R., Carbon dioxide, methane, and
- 587 nitrous oxide exchanges in an age-sequence of temperate pine forests, GLOB CHANGE
- 588 BIOL, 16 (8), 2198-2212, 2010
- 589

590	Prieme, A., Christensen, S., Dobbie, K. E., and Smith, K. A., Slow increase in rate of				
591	methane oxidation in soils with time following land use change from arable agriculture to				
592	woodland, SOIL BIOL BIOCHEM, 29, 1269-1273, 1997				
593					
594	Shindell, D. T., Faluvegi, G., Koch, D. M., Schmidt, G. A., Unger, N., and Bauer, S. E.,				
595	Improved Attribution of Climate Forcing to Emissions, SCIENCE, 326, 716-718, 2009				
596					
597	Smith, K. A., Dobbie, K. E., Ball, B. C., Bakken, L. R., Sitaula, B. K., Hansen, S.,				
598	Brumme, R., Borken, W., Christensen, S., Prieme, A., Fowler, D., Macdonald, J. A.,				

- 599 Skiba, U., Klemedtsson, L., Kasimir-Klemedtsson, A., Degorska, A., and Orlanski, P.,
- 600 Oxidation of atmospheric methane in Northern European soils, comparison with other
- 601 ecosystems, and uncertainties in the global terrestrial sink, GLOBAL CHANGE BIOL, 6,
- 602 791-803, 2000
- 603
- 604 Smolander, A., Priha, O., Paavolainen, L., Steer, J., and Malkonen, E., Nitrogen and

605 carbon transformations before and after clear-cutting in repeatedly N-fertilized and limed

606 forest soil, SOIL BIOL BIOCHEM, 30, 477-490, 1998

607

- 608 Steudler, P. A., Bowden, R. D., Melillo, J. M., and Aber, J. D., Influence of nitrogen-
- 609 fertilization on methane uptake in temperate forest soils, NATURE, 341, 314-316, 1989

611	Steudler, P. A., Melillo, J. M., Bowden, R. D., Castro, M. S., and Lugo, A. E., The effects
612	of natural and human disturbances on soil-nitrogen dynamics and trace gas fluxes in a
613	Puerto-Rican wet forest. BIOTROPICA, 23, 356-363 1991
614	
615	Sullivan, B., Kolb, T. E., Hart, S. C., Kaye, J. P., Dore, S., and Montes-Helu, M.,
616	Thinning reduces soil carbon dioxide but not methane flux from southwestern USA
617	ponderosa pine forests, FOR ECOL MANAG, 255, 4047-4055, 2008
618	
619	Tate, K. R., Ross, D. J., Scott, N. A., Rodda, N. J., Townsend, J. A., and Arnold, G. C.
620	Post-harvest patterns of carbon dioxide production, methane uptake and nitrous oxide
621	production in a Pinus radiata D. Don plantation, FOR ECOL MANAG, 228, 40-50, 2006
622	
623	Teepe, R., Brumme, R., Beese, F., and Ludwig, B., Nitrous oxide emission and methane
624	consumption following compaction of forest soils, SOIL SCI SOC AM J,68, 605-611,
625	2004
626	
627	Thibodeau, L., Raymond, P., Camire, C., and Munson, A. D., Impact of precommercial
628	thinning in balsam fir stands on soil nitrogen dynamics, microbial biomass,
629	decomposition, and foliar nutrition, CAN J FOREST RES 30, 229-238, 2000
630	
631	Van Amstel, A. Methane. A review, J INTEGR ENVIRON SCI, 9, 5-30, 2012

633	Verchot, L. V., Davidson, E. A., Cattanio, J. H., and Ackerman, I.L., Land-use change				
634	and biogeochemical controls of methane fluxes in soils of eastern Amazonia,				
635	ECOSYSTEMS, 3, 41-56, 2000				
636					
637	Wachinger, G., Fiedler, S., Zepp, K., Gattinger, A., Sommer, M., and Roth, K.,				
638	Variability of soil methane production on the micro-scale: spatial association with hot				
639	spots of organic material and Archaeal populations, SOIL BIOL BIOCHEM 32, 1121-				
640	1130, 2000				
641					
642	Wang, F.L. and Bettany, J.R., Methane emissions from a usually well-drained prairie soil				
643	after snowmelt and precipitation. CAN J SOIL SCI, 75(2), 239-241, 1995				
644					
645	Wang, Z.P. and Ineson, P., Methane oxidation in a temperate coniferous forest soil:				
646	effects of inorganic N. SOIL BIOL BIOCHEM, 35(3) 427-433, 2003				
647					
648	Whalen, S.C. and Reeburgh, W.S., Consumption of atmospheric methane by tundra soils.				
649	NATURE, 346(6280), 160-162, 1990				
650					
651	Whalen, S.C. and Reeburgh, W.S., Moisture and temperature sensitivity of CH4				
652	oxidation in boreal soils. SOIL BIOL BIOCHEM, 28(10-11), 1271-1281, 1996				
653					
654	Wu, X., Bruggemann, N., Gasche, R., Papen, H., Willibald, G., and Butterbach-Bahl, K.,				

655 Long-term effects of clear-cutting and selective cutting on soil methane fluxes in a

656	temperate spruce	e forest in southern	Germany, ENVIRON	POLLUT , 159.	2467-2475.
	1 1		<u> </u>	, , ,	·

- 657 2011
- 659 Zerva, A. and Mencuccini.M., Short-term effects of clearfelling on soil CO2, CH4, and
- 660 N2O fluxes in a Sitka spruce plantation. SOIL BIOL BIOCHEM, **37**(11), 2025-2036,
- 661 2005

Table 1. Information regarding vegetation, C and N pool, pH, soil moisture (5th and 95th percentiles) and

- depth to water table at the individual chamber locations. Chamber locations were named U1-U5 at the
- undisturbed plot, T1-T4 at the thinned plot, C1-C5 at the clear-cut plot and S1-S5 at the stump harvested
- 685 plot.

ID	Time period	Vegetation/Bare	Carbon ^a	Nitrogen ^a	pН ^ь	Soil	Depth to	
		soil	(kg m ⁻²)	(kg m^{-2})		moisture, (%)	water table	
							(cm)	
T ₁	01 August 2009-31 May 2010	Mosses, bilberry	6.7	0.22	3.1	28.8-45.8	54-154	
T_2	01 August 2009-31 May 2010	Mosses, bilberry	5.0	0.17	3.1	25.0-40.0	44-144	
T_3	01 August 2009-31 May 2010	Mosses, bilberry	5.5	0.24	3.5	33.5-55.6	15-116	
T_4	11 December 2009- 31 May 2010	Mosses, bilberry	3.3	0.10	3.0	19.2-36.3	29-129	
\mathbf{U}_1	07 July 2010-04 October 2010	Mosses, bilberry	2.6	0.17	3.3	6.0-27.3	120-173	
U_2	07 July 2010-04 October 2010	Mosses, bilberry	6.1	0.29	3.2	10.0-33.4	107-160	
U_3	07 July 2010-04 October 2010	Mosses, bilberry	no data	no data	no data	9.4-37.0	102-155	
U_4	07 July 2010-04 October 2010	Mosses, bilberry	2.3	0.09	3.3	6.6-32.9	136-190	
U_5	07 July 2010-04	Mosses, bilberry	3.9	0.15	3.4	7.8-23.5	132-185	
\mathbf{S}_1	07 October 2010-20	Mosses, bilberry	14.1	0.45	4.4	42.0-42.9	20-21	
S_2	07 October 2010 07 October 2010-20 October 2010	Bare soil, mixed organic and mineral	6.0	0.19	4.4	23.4-25.3	31-32	
S_3	07 October 2010-20 October 2010	Bare soil, mixed organic and mineral	19.0	0.62	4.4	30.0-33.2	47-48	
\mathbf{S}_4	07 October 2010-20 October 2010	Some vegetation and	no data	no data	no data	35.9-39.4	35-36	
S_5	07 October 2010-20 October 2010-20	No vegetation and thick litter layer	no data	no data	no data	33.7-36.1	37-38	
C_1	21 October 2010-09 November 2010	Mosses, bilberry	4.7	0.16	4.2	41.5-46.2	44-50	
C_2	21 October 2010-09 November 2010	Bare soil, mixed organic and mineral	13.1	0.41	4.2	44.2-50.3	6-12	
C ₃	21 October 2010-09 November 2010	Bare soil, mixed organic and mineral	11.9	0.35	4.2	no data	7-13	
C_4	21 October 2010-09 November 2010	Mosses, bilberry	9.5	0.30	4.2	56.6-57.6	6-13	
C ₅	21 October 2010 November 2010	Bare soil, mixed organic and mineral soil layers	11.5	0.36	4.2	49.5-49.9	0-1	

686

^a C and N pool to a depth of 20 cm in the mineral soil (litter layer excluded).

688 ^b pH (BaCl2) for the undisturbed and thinned plots and pH (H2O) for the clear-cut and stump harvested

689 plots were measured at 0-10 cm depth in the mineral soil.

690

- 692 Table 2. Correlation coefficients *C* and corresponding *P*-values for the linear regressions between CH₄
- 693 exchange and soil temperature, soil moisture and water table depth. The r² shows how well the combined
- 694 variables explain the variance in the CH4 exchange. The correlation analyses are based on data from the
- 695 entire measurement period.
- 696

	С	Р	С	Р	С	Р	r^2
	Soil temperature	Soil temperature	Soil moisture	Soil moisture	Water table	Water table	
	-				depth	depth	
T_1	-0.09	**	0.57	**	0.12	**	0.27
T_2	-0.34	**	-0.23	**	0.46	**	0.26
T_3	0.34	**	0.72	**	0.45	**	0.61
T_4	0.28	**	-0.72	**	0.54	**	0.68
U_1	-0.61	**	0.48	**	0.51	**	0.47
U_2	-0.63	**	0.39	**	0.54	**	0.47
U_3	-0.57	**	0.44	**	0.55	**	0.53
U_4	-0.69	**	0.54	**	0.70	**	0.78
U_5	-0.82	**	0.54	**	0.69	**	0.81
S_1	-0.35	**	-0.18	*	а	а	0.09
S_2	-0.16	*	-0.10	0.09	а	а	0.07
S_3	-0.02	0.82	-0.09	0.21	а	а	0.002
S_4	-0.16	*	-0.009	0.9	а	а	0.05
S_5	-0.34	**	-0.46	*	а	а	0.09
C_1	-0.06	0.29	-0.13	*	0.03	0.54	0.14
C_2	-0.52	**	0.25	**	-0.07	0.13	0.22
C_3	-0.57	**	no data	no data	-0.47	**	0.46
C_4	-0.04	0.57	0.04	0.57	-0.01	0.91	0.01
C ₅	-0.54	**	-0.03	0.54	-0.20	**	0.21

697

698 ** significant, p <0.001

699 * significant, p < 0.05

700 *a*, At the time for measurements on the stump harvested plot, the water table depth was only measured

701 manually on a few occasions and therefore no linear regression could be made for this period.

702

704 Table 3. Coefficients from multiple linear regression analyses. A value is given only if the variable significantly contributes to explain the variation in the CH₄ exchange. The r² shows how well the combined variables explain the variance in the CH4 exchange. S.m represents soil moisture, S.t, soil 705 706 temperature and W.t, water table depth. 707 708

T						T				T				T_4						
	r^2	S.m.	S.t.	W.t.	r^2	S.m.	S.t.	W.t.	\mathbf{r}^2	S.m.	S.t.	W.t.	r^2	S.m.	S.t.	W.t.				
Aug 09	0.58	0.61	-	0.43	0.24	0.14	-0.18	0.44	0.76	0.66	0.33	-	n.d	n.d	n.d	n.d				
Sep 09	0.10	0.31	-	-	0.10	-	-0.28	-	0.28	0.52	-	-	n.d	n.d	n.d	n.d				
Oct 09	0.18	0.27	-0.14	0.22	0.16	-0.28	-0.28	-0.20	0.10	-0.15	0.13	-0.21	n.d	n.d	n.d	n.d				
Nov 09	0.31	0.44	-0.16	0.33	0.23	0.33	-0.44	0.43	0.33	-	0.13	0.50	0.04	-	0.20	-				
Apr 10	0.31	0.54	0.39	0.34	0.27	0.65	-0.34	-0.59	<i>∂</i> 0.10	-	-0.26	-	n.d	n.d	n.d	n.d				
May 10	0.22	0.43	-	-0.18	0.10	-	-0.29	-	0.38	-0.49	-	0.46	0.62	n.d	0.79	n.d	_			
			U ₁			U2				U3			U4						U ₅	
	\mathbf{r}^2	S.m.	S.t.	W.t.	\mathbf{r}^2	S.m.	S. t.	W.t.	\mathbf{r}^2	S.m.	S.t.	W.t.	r ²	S.m.	S.t.	W.t.	\mathbf{r}^2	S.m.	S.t.	W.
Jul 10	0.22	0.42	-	-0.18	0.10	n.d	-0.24	0.22	0.42	0.60	-0.14	0.48	0.12	0.12	-0.28	-	0.49	0.20	-0.27	0.4
Aug 10	0.47	-	-0.12	0.62	0.56	n.d	-0.23	0.61	0.37	0.08	-	0.60	0.80	0.19	-0.18	0.79	0.78	0.15	-0.34	0.6
Sep 10	0.12	0.10	-0.29	-0.10	0.37	n.d	-0.60	-	0.10	0.11	-0.23	-0.10	0.63	0.46	-0.50	-	0.28	-	-0.51	-0.
		S1			S ₂			S ₃		5	54		S5							
	r^2	S.m.	S.t.	r^2	S.m.	S.t.	r^2	S.m.	S.t. r ²	S.	.m. S.t	$\cdot r^2$	S.m.	S.t.						
Oct 10	0.09	_	-0.29	0.07	-0.27	_	0.002	_	- 0.	05 0.	10 0.1	0 0.09	-	-0.2	8					
			C ₁				C ₂				C ₃			С	4				C.	
	r^2	S.m.	S.t.	W.t.	\mathbf{r}^2	S.m.	S.t.	W.t.	r^2	S.m.	S.t.	W.t.	r^2	S.m.	S.t.	W.t.	r ²	S.m.	S.t.	W.
Oct -Nov 10	0.14	-	-0.32	-	0.22	0.29	-0.34	-0.38	0.46	n.d	-0.44	-0.44	0.01	-	-	-	0.21	-0.28	-0.59	0.1









Fig.2. Average CH₄ exchange rates, soil moisture and soil temperature at the four sampling plots. Data from the entire measurement period at each plot is included. The dashed line at the thinned plot represents average values for October and November, since measurements at the clear-cut and stump harvested plots were conducted during this part of the year.

Fig.3a







748 Fig.3c)



752

753 Fig.3. Time series of daily mean CH4 exchange, daily precipitation and daily mean soil temperature at the

754 measurement locations. Fig.3a) Undisturbed plot, Fig.3b) Thinned plot, Fig.3c) Clear-cut plot, Fig.3d)

755 Stump harvested plot.





Fig.4. CH₄ exchange (µmol m⁻² h⁻¹) at all individual measurement locations with associated level of ground water table. The water table depth at plot C_5 is close to zero and that is why the bar is not visible in the

diagram.

¤ Measurement locations where soil surface was disturbed during site preparation.



Fig.5. Correlation between CH_4 exchange (µmol m⁻² h⁻¹) and soil temperature (°C) at measurement location

776 U₄. The different colours represent different soil moisture and water table depths.

- _--