Trends in soil solution dissolved organic carbon (DOC) concentrations across European forests

3

Marta Camino-Serrano¹, Elisabeth Graf Pannatier², Sara Vicca¹, Sebastiaan 4 Luyssaert^{3,22}, Mathieu Jonard⁴, Philippe Ciais³, Bertrand Guenet³, Bert Gielen¹, 5 Josep Peñuelas^{5,6}, Jordi Sardans^{5,6}, Peter Waldner², Sophia Etzold², Guia 6 Cecchini⁷, Nicholas Clarke⁸, Zoran Galić⁹, Laure Gandois^{10,11}, Karin Hansen¹², 7 Jim Johnson¹³, Uwe Klinck¹⁴, Zora Lachmanová¹⁵, Antti-Jussi Lindroos¹⁶, 8 Henning Meesenburg¹⁴, Tiina M. Nieminen¹⁶, Tanja G.M. Sanders¹⁷, Kasia 9 Sawicka¹⁸, Walter Seidling¹⁷, Anne Thimonier², Elena Vanguelova¹⁹, Arne 10 Verstraeten²⁰, Lars Vesterdal²¹, Ivan A. Janssens¹ 11

- 12 [1]{Department of Biology, PLECO, University of Antwerp, Belgium}
- 13 [2] {WSL, Swiss Federal Institute for Forest, Snow and Landscape Research, Zürcherstrasse
- 14 111, 8903, Birmensdorf, Switzerland}
- 15 [3] {Laboratoire des Sciences du Climat et de l'Environnement, LSCE/IPSL, CEA-CNRS-
- 16 UVSQ, Université Paris-Saclay, F-91191 Gif-sur-Yvette, France}
- 17 [4] {UCL-ELI, Université catholique de Louvain, Earth and Life Institute, Croix du Sud 2,
- 18 BE-1348 Louvain-la-Neuve, Belgium}
- 19 [5] {CREAF, Cerdanyola del Vallès, Catalonia, Spain}
- 20 [6] {CSIC, Global Ecology Unit CREAF-CSIC-UAB, Cerdanyola del Vallès, Catalonia,
- 21 Spain}
- 22 [7] {Earth Sciences Department, University of Florence, Italy}
- 23 [8] {Norwegian Institute of Bioeconomy Research, N-1431, Ås, Norway}
- 24 [9] {University of Novi Sad-Institute of Lowland Forestry and Environment, Serbia}
- 25 [10] {Université de Toulouse: UPS, INP, EcoLab (Laboratoire Ecologie fonctionnelle et
- 26 Environnement), ENSAT, Castanet-Tolosan, France}
- 27 [11] {CNRS, EcoLab, Castanet-Tolosan, France}

- [12] {IVL Swedish Environmental Research Institute, Natural Resources & Environmental
 Effects, SE-100 31, Stockholm, Sweden}
- 30 [13] {UCD School of Agriculture and Food Science, University College Dublin, Belfield,31 Ireland}
- 32 [14] {Northwest German Forest Research Institute, Grätzelstr. 2, D-37079, Göttingen,
 33 Germany}
- 34 [15] {FGMRI, Forestry and Game Management Research Institute, Strnady 136, 252 02
 35 Jíloviště, Czech Republic}
- 36 [16] {Natural Resources Institute Finland (Luke), P.O. Box 18, 01301 Vantaa, Finland }
- 37 [17] {Thünen Institute of Forest Ecosystems, Alfred-Möller-Strasse 1, D-16225, Eberswalde,
 38 Germany}
- 39 [18] {Soil Geography and Landscape Group, Wageningen University, PO Box 47, 6700 AA
 40 Wageningen, NL}
- [19] {Centre for Ecosystem, Society and Biosecurity, Forest Research, Alice Holt Lodge,
 Wrecclesham, Farnham, Surrey GU10 4LH, UK}
- 43 [20] {Research Institute for Nature and Forest (INBO), Kliniekstraat 25, BE-1070 Brussels,
 44 Belgium}
- 45 [21] {University of Copenhagen, Department of Geosciences and Natural Resource
 46 Management, Rolighedsvej 23, DK-1958 Frederiksberg C, Denmark}
- 47 [22] now at {Free University of Amsterdam, Department of Ecological Science, Boelelaan
 48 1085, NL-1081HV, the Netherlands}
- 49 Correspondence to: M. Camino-Serrano (marta.caminoserrano@uantwerpen.be)
- 50

51 Abstract

52 Dissolved organic carbon (DOC) in surface waters is connected to DOC in soil solution 53 through hydrological pathways. Therefore, it is expected that long-term dynamics of DOC in 54 surface waters reflect DOC trends in soil solution. However, a multitude of site-studies has 55 failed so far to establish consistent trends in soil solution DOC, whereas increasing 56 concentrations in European surface waters over the past decades appear to be the norm, 57 possibly as a result of recovery from acidification. The objectives of this study were therefore 58 to understand the long-term trends of soil solution DOC from a large number of European 59 forests (ICP Forests Level II plots) and determine their main physico-chemical and biological 60 controls. We applied trend analysis at two levels: 1) to the entire European dataset and 2) to 61 the individual time series and related trends with plot characteristics, i.e., soil and vegetation 62 properties, soil solution chemistry and atmospheric deposition loads. Analyses of the entire 63 dataset showed an overall increasing trend in DOC concentrations in the organic layers, but, 64 at individual plots and depths, there was no clear overall trend in soil solution DOC. The rate change of soil solution DOC ranged between -16.8% yr^{-1} and +23% yr^{-1} (median= +0.4% yr^{-1} 65 ¹) across Europe. The non-significant trends (40%) outnumbered the increasing (35%) and 66 67 decreasing trends (25%) across the 97 ICP Forests Level II sites. By means of multivariate 68 statistics, we found increasing trends in DOC concentrations with increasing mean nitrate 69 (NO₃) deposition and increasing trends in DOC concentrations with decreasing mean sulphate (SO_4^{2}) deposition, with the magnitude of these relationships depending on plot 70 deposition history. While the attribution of increasing trends in DOC to the reduction of SO_4^{2-} 71 72 deposition could be confirmed in low to medium N deposition areas, in agreement with 73 observations in surface waters, this was not the case in high N deposition areas. In 74 conclusion, long-term trends of soil solution DOC reflected the interactions between controls acting at local (soil and vegetation properties) and regional (atmospheric deposition of SO_4^{2-} 75 76 and inorganic N) scales.

77 **1** Introduction

78 Dissolved organic carbon (DOC) in soil solution is the source of much of the terrestrially 79 derived DOC in surface waters (Battin et al., 2009; Bianchi, 2011; Regnier et al., 2013). Soil 80 solution DOC in forests is connected to streams through different hydrological pathways: 81 DOC mobilized in the forest floor may be transported laterally at the interface of forest floor 82 and mineral soil to surface waters or percolates into the mineral soil, where additional DOC 83 can be mobilized and/or DOC is partly adsorbed on particle surfaces and mineralized 84 thereafter (Fig. 1). From the mineral soil DOC may be either leached laterally or vertically 85 via groundwater into surface waters (McDowell and Likens, 1988). Therefore, it could be expected that long-term dynamics of DOC in surface waters mirror those observed in 86 87 ecosystem soil solutions.

B8 Drivers related to climate change (temperature increase, precipitation change, atmospheric B9 CO_2 increase), the decrease in acidifying deposition or land use change and management may

90 individually or jointly explain trends in surface water DOC concentrations (Evans et al., 91 2012; Freeman et al., 2004; Oulehle et al., 2011; Sarkkola et al., 2009; Worrall and Burt, 92 2004). Increasing air temperatures warm the soil, thus stimulating soil organic matter (SOM) 93 decomposition through greater microbial activity (Davidson and Janssens, 2006; Hartley and 94 Ineson, 2008; Kalbitz et al., 2000). Other drivers, such as increased atmospheric CO_2 and the 95 accumulation of atmospherically deposited inorganic nitrogen are thought to increase the 96 sources of DOC by enhancing primary plant productivity (i.e., through stimulating root 97 exudates or, increased litterfall) (de Vries et al., 2014; Ferretti et al., 2014; Sucker and 98 Krause, 2010). Changes in precipitation, land use and management (e.g. drainage of 99 peatlands, changes in forest management or grazing systems) may alter the flux of DOC 100 leaving the ecosystem but no consistent trends in the hydrologic regime or land use changes 101 were detected in areas where increasing DOC trends have been observed (Monteith et al., 102 2007).

103 Recent focus was mainly on decreasing acidifying deposition as an explanatory factor for 104 DOC increases in surface waters in Europe and North America by means of decreasing ionic 105 strength (De Wit et al., 2007; Hruška et al., 2009) and increasing the pH of soil solution, 106 consequently increasing DOC solubility (Evans et al., 2005; Haaland et al., 2010; Monteith et 107 al., 2007). Although the hypothesis of an increase in surface water DOC concentration due to 108 a recovery from past acidification was confirmed in studies of soil solution DOC in the UK 109 and northern Belgium (Sawicka et al., 2016; Vanguelova et al., 2010; Verstraeten et al., 110 2014), it is not consistent with trends in soil solution DOC concentrations reported from 111 Finnish, Norwegian, and Swedish forests (Löfgren and Zetterberg, 2011; Ukonmaanaho et 112 al., 2014; Wu et al., 2010). This inconsistency between soil solution DOC and stream DOC 113 trends could suggest that DOC in surface water and soil solution responds differently to 114 (changes in) environmental conditions in different regions (Akselsson et al., 2013; Clark et 115 al., 2010; Löfgren et al., 2010). Alternatively, other factors such as tree species and soil type, 116 may be co-drivers of organic matter dynamics and input, generation and retention of DOC in 117 soils.

Trends of soil solution DOC not only vary among forests but often also within the same site (Borken et al., 2011; Löfgren et al., 2010). Forest characteristics such as tree species composition, soil fertility, texture or sorption capacity may affect the response of soil solution DOC to environmental controls, for instance, by controlling the rate of soil acidification through soil buffering and nutrient plant uptake processes (Vanguelova et al., 2010). Within a 123 site, DOC variability with soil depth is typically caused by different intensity of DOC 124 production, transformation, and sorption along the soil profile (Fig. 1). Positive temporal 125 trends in soil solution DOC (increasing concentrations over time) were frequently reported 126 for the organic layers and shallow soils where production and decomposition processes 127 control the DOC concentration (Löfgren and Zetterberg, 2011). However, no dominant trends 128 are found for the mineral soil horizons, where physico-chemical processes, such as sorption, 129 become more influential (Borken et al., 2011; Buckingham et al., 2008). Furthermore, 130 previous studies have used different temporal and spatial scales which may have further 131 added to the inconsistency in the DOC trends reported in the literature (Clark et al., 2010).

132 In this context, the International Co-operative Programme on Assessment and Monitoring of 133 Air Pollution Effects on Forests (ICP Forests, 2010) compiled a unique dataset containing data from more than 100 intensively monitored forest plots (Level II) which allow to unravel 134 135 regional trends in soil solution DOC of forests at a European scale, and perform statistical 136 analysis of the main controls behind these regional trends. Long-term measurements of soil solution DOC are available for these plots, along with information on aboveground biomass, 137 soil properties, and atmospheric deposition of inorganic N and SO₄²⁻, collected using a 138 harmonized sampling protocol across Europe (Ferretti and Fischer, 2013). This dataset has 139 140 previously been used to investigate the spatial variability of DOC in forests at European scale 141 (Camino-Serrano et al., 2014), but an assessment of the temporal trends in soil solution DOC 142 using this large dataset has not been attempted so far.

The main objective of this study is to understand the long-term temporal trends of DOC concentrations in soil solution measured at the ICP Forests Level II plots across Europe. Based on the increasing DOC trends in surface waters, we hypothesize that temporal trends in soil solution DOC will also be positive, but with trends varying locally depending on plot characteristics. We further investigated whether plot characteristics, specifically climate, inorganic N and SO_4^{2-} deposition loads, forest type, soil properties, and changes in soil solution chemistry can explain differences across sites in DOC trends.

150 2 Materials and Methods

151 **2.1 Data description**

Soil solution chemistry has been monitored within the ICP Forests Programme since the1990s on most Level II plots. The ICP Forests data were extracted from the pan-European

154 Forest Monitoring Database (Granke, 2013). A list of the Level II plots used for this study 155 can be found in Supplementary material, Table S1. The methods for collection and analysis 156 of soil solution used in the various countries (Switzerland: Graf Pannatier et al. (2011); 157 Flanders: Verstraeten et al. (2012); Finland: Lindroos et al. (2000); UK: Vanguelova et al. 158 (2010), Denmark: Hansen et al. (2007)) follow the ICP Forests manual (Nieminen, 2011). 159 Generally, lysimeters were installed at several fixed depths starting at 0 cm, defined as the 160 interface between the surface organic layer and underlying mineral soil. These depths are 161 typically aligned with soil "organic layer", "mineral topsoil", "mineral subsoil", and "deeper 162 mineral soil" but sampling depths vary among countries and even among plots within a 163 country. Normally, zero-tension lysimeters were installed under the surface organic layer and tension lysimeters within the mineral soil. However, in some countries zero-tension 164 lysimeters were also used within the mineral layers and in some tension lysimeters below the 165 166 organic layer. Multiple collectors (replicates) were installed per plot and per depth to assess 167 plots spatial variability. However, in some countries, samples from these replicates were pooled before analyses or averaged prior to data transmission. The quality assurance and 168 169 control procedures included the use of control charts for internal reference material to check 170 long-term comparability within national laboratories as well as participation in periodic 171 laboratory ring tests (e.g., Marchetto et al., 2011) to check the international comparability. 172 Data were reported annually to the pan-European data center, checked for consistency and 173 stored in the pan-European Forest Monitoring Database (Granke, 2013).

174 Soil water was usually collected fortnightly or monthly, although for some plots sampling 175 periods with sufficient soil water for collection were scarce, especially in prolonged dry 176 periods or in winter due to snow and ice. After collection, the samples were filtered through a 177 0.45 µm membrane filter, stored below 4 °C and then analyzed for DOC, together with other soil solution chemical properties (NO₃⁻, Ca, Mg, NH₄⁺, SO₄⁻², total dissolved Al, total 178 179 dissolved Fe, pH, electrical conductivity). Information on the soil solution chemistry at the 180 studied plots can be found in Supplementary material (Table S4-S11). The precision of DOC analysis differed among the laboratories. The coefficient of variation of repeatedly measured 181 182 reference material was 3.7% on average. The time span of soil solution time series used for 183 this study ranged from 1991 to 2011, although coverage of this period varied from plot to plot 184 (Supplementary material, Table S1).

Soil properties, open field bulk deposition and throughfall deposition of NO_3^- , NH_4^+ , and SO_4^{-2-} , are measured at the same plots as well as stem volume increment. The atmospheric

deposition of NO_3^- , NH_4^+ and SO_4^{2-} data covers the period 1999-2010 (Waldner et al., 2014). 187 Stem volume growth was calculated by the ICP Forests network from diameter at breast 188 189 height (DBH), live tree status, and tree height which were assessed for every tree (DBH > 5190 cm) within a monitoring plot approximately every five years since the early 1990s. Tree stem 191 volumes were derived from allometric relationships based on diameter and height 192 measurements according to De Vries et al. (2003), accounting for species and regional differences. Stem volume growth (in m³) between two consecutive inventories was calculated 193 194 as the difference between stem volumes at the beginning and the end of one inventory period 195 for living trees. Stem volume data were corrected for all trees that were lost during one 196 inventory period, including thinning. Stem volume at the time of disappearance (assumed at 197 half of the time of the inventory period) was estimated from functions relating stem volume 198 of standing living trees at the end of the period vs volume at the beginning of the period. The 199 methods used for collection of these data can be found in the Manuals of the ICP Forests 200 Monitoring Programme (ICP Forests, 2010). The soil properties at the plots used for this 201 study were derived from the ICP Forests aggregated soil database (AFSCDB.LII.2.1) (Cools 202 and De Vos, 2014).

203 Since continuous precipitation measurements are not commonly available for the Level II plots, precipitation measurements for the location of the plots were extracted from the 204 205 observational station data of the European Climate Assessment & Dataset (ECA&D) and the 206 ENSEMBLES Observations (E-OBS) gridded dataset (Haylock et al., 2008). We used 207 precipitation measurements extracted from the E-OBS gridded dataset to improve the 208 temporal and spatial coverage and to reduce methodological differences of precipitation 209 measurements across the plots. The E-OBS dataset contains daily values of precipitation and 210 temperature from stations data gridded at 0.25 degrees resolution. When E-OBS data were 211 not available, they were gap-filled with ICP Forests precipitation values gained by deposition 212 measurements where available.

213 2.2 Data preparation

We extracted data from plots with time series covering more than 10 years and including more than 60 observations of soil solution DOC concentrations of individual or groups of collectors. Outliers, defined as \pm 3 interquartile range of the 25 and 75 quantiles of the time series, were removed from each time series to avoid influence of few extreme values in the long-term trend (Schwertman et al., 2004). Values under 1 mg L⁻¹, which is the detection

- limit for DOC in the ICP Level II plots, were replaced by 1 mg L⁻¹. After this filtering, 529 time series from 118 plots, spanning from Italy to Norway, were available for analysis. Soil solution, precipitation, and temperature were aggregated to monthly data by the median of the observations in each month and by the sum of daily values in the case of precipitation. Data of inorganic N (NH₄⁺ and NO₃⁻) and SO₄²⁻ throughfall and open field bulk deposition measured at the plots were interpolated to monthly data (Waldner et al., 2014).
- 225 The plots were classified according to their forest type (broadleaved/coniferous dominated), soil type (World Reference Base, (WRB 2006)), their stem growth (slow, $< 6 \text{ m}^3 \text{ ha}^{-1} \text{ yr}^{-1}$, 226 intermediate, 6–12 m³ ha⁻¹ yr⁻¹; and fast, > 12 m³ ha⁻¹ yr⁻¹), and soil solution pH (low, <4.2, 227 228 intermediate, 4.2–5, high, >5). Plots were also classified based on mean throughfall inorganic N (NO₃⁻ +NH₄⁺) deposition level, defined as: high deposition (HD, >15 kg N ha⁻¹ vr⁻¹). 229 medium deposition (MD, 5–15 kg N ha⁻¹ yr⁻¹), and low deposition (LD, <5 kg N ha⁻¹ yr⁻¹) and 230 mean throughfall SO_4^{2-} deposition level, defined as: high deposition (HD, >6 kg S ha⁻¹ yr⁻¹), 231 and low deposition (LD, $< 6 \text{ kg S ha}^{-1} \text{ yr}^{-1}$). 232

233 2.3 Statistical methods

234 Time series can typically decomposed into random noise, seasonal, and trend components (Verbesselt et al., 2010). In this paper, we used methods to detect the actual trend (change in 235 236 time) after removing the seasonal and random noise components. The sequence of methods 237 applied is summarized in Fig. 2. The analysis of temporal trends in soil solution DOC 238 concentrations was carried out at two levels: 1) the European level and 2) the plot level. 239 While the first analysis allows an evaluation of the overall trend in soil solution DOC at a 240 continental scale, the second analysis indicates whether the observed large scale trends are occurring at local scales as well, and tests whether local trends in DOC can be attributed to 241 242 certain driver variables.

243 Linear mixed-effects models (LMM) were used to detect the temporal trends in soil solution DOC concentration at the European scale (Fig. 2). For these models, the selected 529 time 244 245 series were used. For the trend analysis of individual time series, however, we focused on the 246 long-term trends in soil solution DOC at European forests that show monotonicity. Therefore, 247 DOC time series were first analyzed using the Breaks For Additive Seasonal and Trend 248 (BFAST) algorithm to detect the presence of breakpoints (Verbesselt et al., 2010; Vicca et al., 249 2016) with the time series showing breakpoints, i.e., not monotonic, being discarded (see Description of the statistical methods in Supplementary material). In total, 258 monotonic 250

251 time series from 97 plots were used for our analysis after filtering (Fig. 2). Then, monotonic 252 trend analyses were carried out from the filtered dataset using the Seasonal Mann Kendall 253 (SMK) test for monthly DOC concentrations (Hirsch et al., 1982; Marchetto et al., 2013). 254 Partial Mann Kendall (PMK) tests were also used to test the influence of precipitation as a 255 co-variable to detect if the trend might be due to a DOC dilution/concentration effect 256 (Libiseller and Grimvall, 2002). Sen (1968) slope values were calculated for SMK and PMK. 257 Moreover, LMMs were performed again with the filtered dataset to compare results with and 258 without time series showing breakpoints (Fig. 2).

- 259 For this study, five soil depth intervals were considered: the organic layer (0 cm), topsoil (0-260 20 cm), intermediate (20-40 cm), subsoil (40-80 cm) and deep subsoil (> 80 cm). The slopes of each time series were standardized by dividing them by the median DOC concentration 261 262 over the sampling period (relative trend slope), aggregated to a unique plot-soil depth slope 263 and classified by the direction of the trend as significantly positive, i.e., increasing DOC over time (P, p < 0.05), significantly negative, i.e., decreasing DOC over time (N, p < 0.05), and 264 non-significant, i.e., no significant change in DOC over time (NS, $p \ge 0.05$). When there was 265 266 more than one collector per depth interval, the median of the slopes was used when the 267 direction of the trend (P, N, or NS) was similar. After aggregation per plot-depth combination, 191 trend slopes from 97 plots were available for analysis (Supplementary 268 Material, Table S2). Trends for other soil solution parameters (NO₃⁻, Ca²⁺, Mg²⁺, NH₄⁺, SO₄²⁻ 269 270 , total dissolved Al, total dissolved Fe, pH, electrical conductivity), precipitation and 271 temperature were calculated using the same methodology as for DOC. Since the resulting standardized Sen slope in % yr⁻¹ (relative trend slope) was used for all the statistical analysis, 272 273 from here on we will use the general term "trend slope" in order to simplify.
- 274 Finally, Structural Equation Models (SEM) were performed to determine the capacity of the several factors (SO_4^{2-} and/or NO_3^{-} deposition, stem growth and soil solution chemistry) in 275 276 explaining variability in the slope of DOC trends among the selected plots (Fig. 2). We evaluated the influence of both the annual mean (kg ha⁻¹ yr⁻¹) and the trends (% yr⁻¹) in 277 deposition and soil solution parameters.. All the statistical analyses were performed in R 278 279 software version 3.1.2 (R Core Team, 2014) using the "rkt" (Marchetto et al., 2013), 280 "bfast01" (de Jong et al., 2013) and "sem" (Fox et al., 2013) packages, except for the LMMs that were performed using SAS 9.3 (SAS institute, Inc., Cary, NC, USA). More detailed 281 282 information on the statistical methods used can be found in Supplementary material.

283 **3 Results**

3.1 Soil solution DOC trends at European scale

285 First, temporal trends in DOC were analyzed for all the European DOC data pooled together 286 by means of LMM models to test for the presence of overall trends. A significantly increasing 287 DOC trend (p < 0.05) in soil solution collected with zero-tension lysimeters in the organic 288 layer was observed mainly under coniferous forest plots (Table 1). Similarly, a significantly 289 increasing DOC trend (p < 0.05) in soil solution collected with tension lysimeters was found 290 in deep mineral soil (> 80 cm) for all sites, mainly for coniferous forest sites (Table 1), but 291 this trend is based on a limited number of plots which are not especially well distributed in 292 Europe (75 % of German plots). By contrast, non-significant trends were found in the other 293 mineral soil depth intervals (0-20 cm, 20-40 cm, 40-80 cm) by means of the LMM models. 294 When the same analysis was applied to the filtered European dataset, i.e., without the time 295 series showing breakpoints, fewer significant trends were observed: only an overall positive 296 trend (p < 0.05) was found for DOC in the organic layer using zero-tension lysimeters, again 297 mainly under coniferous forest sites but no statistically significant trends were found in the 298 mineral soil (Table 1).

3.2 Soil solution DOC concentration trend analysis of individual time series

We applied the BFAST analysis to select the monotonic time series in order to assure that the detected trends were not influenced by breakpoints in the time series. Time series with breakpoints represented more than 50% of the total time series aggregated by soil depth interval (245 out of 436).

304 The individual trend analysis using the SMK test showed trend slopes of soil solution DOC concentration ranging from -16.8% yr⁻¹ to +23% yr⁻¹ (median= +0.4% yr⁻¹, interquartile range 305 = +4.3% yr⁻¹). Among all the time series analyzed, the majority were not statistically 306 significant trends (40%, 104 time series), followed by significantly positive trends (35%, 91 307 308 time series) and significantly negative trends (24%, 63 time series) (Table 1). There was, 309 thus, no uniform trend in soil solution DOC in forests across a large part of Europe. 310 Furthermore, the regional trend differences were inconsistent when looking at different soil 311 depth intervals separately (Fig. 3 and 4), which made it difficult to draw firm conclusions 312 about the spatial pattern of the trends in soil solution DOC concentrations in European 313 forests.

The variability in trends was high, not only at continental scale, but also at plot level (Fig. 5). We found consistent within-plot trends only for 50 out of the 97 sites. Moreover, some plots even showed different trends (P, N or NS) in DOC within the same depth interval, which was the case for 17 plot-depth combinations (16 in Germany and one in Norway), evidencing a high small-scale plot heterogeneity.

319 Trend directions (P, N or NS) often differed among depths. For instance, in the organic layer, 320 we found mainly non-significant trends and, if a trend was detected, it was more often 321 positive than negative, while positive trends were the most frequent in the subsoil (below 40 322 cm) (Table 1). Nevertheless, it is important to note that a statistical test of whether there was 323 a real difference in DOC trends between depths was not possible as the set of plots differed 324 between the different soil depth intervals. However, a visual comparison of trends for the few 325 plots in which trends were evaluated for more than three soil depths showed that there was no 326 apparent difference in DOC trends between soil depths (Supplementary material, Fig. S1 and 327 S2).

Finally, for virtually all plots, including precipitation as a co-variable in the PMK test gave the same result as the SMK test, which indicates that precipitation (through dilution or concentration effects) did not affect the DOC concentration trends. Dilution/concentration effect was only detected in four plots (Supplementary material, Table S1).

332 3.3 Factors explaining the soil solution DOC trends

333 **3.3.1 Effects of vegetation, soil and climate**

There was no direct effect of forest type (broadleaved vs. coniferous) on the direction of the 334 statistically significant trends in soil solution DOC (Fig. 6A). Both positive and negative 335 trends were equally found under broadleaved and coniferous forests ($\chi^2(1, n = 97) = 0.073$, p 336 = 0.8). Increasing DOC trends, however, occurred more often under forests with a mean stem 337 growth increment below 6 m³ ha⁻¹ yr⁻¹ over the study period, whereas decreasing DOC trends 338 were more common in forests with a mean stem growth increment between 6 and 12 m^3 ha⁻¹ 339 $vr^{-1}(\gamma^2(2, n = 53) = 5.8, p = 0.05)$ (Fig. 6B). Only six forests with a mean stem growth above 340 12 m³ ha⁻¹ yr⁻¹ were available for this study (five showing increasing DOC trends and one 341 342 showing a decreasing DOC trend) and, thus, there is not enough information to draw 343 conclusions about the relationship between stem growth and soil solution DOC trends for forests with very high stem growth (> $12 \text{ m}^3 \text{ ha}^{-1} \text{ yr}^{-1}$). 344

345 The DOC trends also varied among soil types, more than half of the plots showing a consistent increasing DOC trend at all evaluated soil depth intervals were located in 346 347 Cambisols, (6 out of 11 plots), which are rather fertile soils, whereas plots showing consistent 348 negative trends covered six different soil types. Other soil properties, like clay content, cation 349 exchange capacity or pH, did not clearly differ between sites with positive and negative DOC 350 trends (Table 2). It is remarkable that trends in soil solution pH, Mg and Ca concentrations 351 were similar across plots with both positive and negative DOC trends. Soil solution pH 352 increased distinctly in almost all the sites, while Ca and Mg decreased markedly (Table 2).

Finally, no significant correlations were found between trends in temperature or precipitation and trends in soil solution DOC, with the exception of a positive correlation between trends in soil solution DOC in the soil depth interval 20-40 cm and the trend in temperature (r = 0.47, p = 0.03).

357 3.3.2 Effects of mean and trends in atmospheric deposition and soil solution 358 parameters

Analyzing different models that could explain the DOC trends using the overall dataset 359 indicated both direct and indirect effects of the annual mean SO_4^{2-} and NO_3^{-} throughfall 360 atmospheric deposition on the trend slopes of DOC. The Structural Equation Model 361 362 accounted for 32.7% of the variance in DOC trend slopes (Fig. 7A). According to this model, lower mean throughfall SO_4^{2-} deposition resulted in increasing trend slopes of DOC in soil 363 solution and higher mean throughfall NO_3^- deposition resulted in increasing trend slopes of 364 DOC (Fig. 7A). When considering trends in SO_4^{2-} and NO_3^{-} deposition, there was no 365 apparent spatial correlation with soil solution DOC trends, with deposition mainly decreasing 366 or not changing over time (Fig. 8) and the DOC trends varying greatly across Europe (Fig. 3) 367 and 4). However, when SEM was run using the trend slopes in SO_4^{2-} and NO_3^{-} deposition 368 instead of the mean values, we found that trend slopes of DOC significantly increased with 369 increasing trend in NO_3^- and decreased with increasing trend in SO_4^{2-} deposition, but the 370 latter was a non-significant relationship (Supplementary Material, Fig. S3). However, the 371 372 percentage of variance in DOC trend slopes explained by the model was much twice lower 373 (16%).

374 Sites with low and medium N deposition

The variables in the model that best explained the temporal changes in DOC were the same for the forests with low and medium N deposition; for both groups, NO_3^- deposition and

 SO_4^{2-} deposition (directly, or indirectly through its influence on plant growth) influenced the 377 trend in DOC (Fig. 7B). Lower mean SO_4^{2-} deposition again resulted in a significant increase 378 in trend slopes, while increasing NO_3^- deposition resulted in increasing DOC trend slopes. 379 The percentage of variance in DOC trend slopes explained by the model was 33%. The SEM 380 run with the trends in SO_4^{2-} and NO_3^{-} throughfall deposition for forests with low and medium 381 N deposition explained 24.4% of the variance in DOC trends, and showed a significant 382 increase of trend slopes of DOC with decreasing trend in SO_4^{2-} deposition (Supplementary 383 384 Material, Fig. S3).

385 Sites with high N deposition

For the plots with high N deposition, however, we found no model for explaining the trends 386 in DOC using the mean annual SO_4^{2-} and NO_3^{-} throughfall deposition. In contrast, the best 387 model included the relative trend slopes in SO_4^{2-} and NO_3^{-} deposition as well as in median 388 soil solution conductivity (% yr⁻¹) as explaining variables (Fig. 7C). Increasing the relative 389 trend slopes of NO_3^- deposition resulted in increasing the DOC trend slopes. Also both the 390 391 trend slopes of SO_4^{2-} and NO_3^{-} deposition affected the trend slopes of DOC indirectly through an effect on the trends in soil solution conductivity, although acting in opposite directions: 392 while increasing NO_3^- deposition led to decreasing soil solution conductivity, increasing 393 SO₄²⁻ deposition resulted in increasing trends in soil solution conductivity, but the latter 394 relationship was only marginally significant (p=0.06). Increasing trends in conductivity, in 395 396 turn, resulted in increasing trend slopes of DOC. The percentage of the variance in DOC trend slopes explained by the model was 25% (Fig. 7C). Nevertheless, trends in soil solution 397 DOC were not directly affected by trends in SO_4^{2-} deposition in forests with high N 398 399 deposition.

400 **4 Discussion**

401 **4.1** Trend analysis of soil solution DOC in Europe

402 **4.1.1** Evaluation of the trend analysis techniques

A substantial proportion (40%) of times series did not indicate any significant trend in sitelevel DOC concentrations across the ICP Forests network. Measurement precision, strength
of the trend, and the choice of the method may all affect trend detection (Sulkava et al., 2005;
Waldner et al., 2014). Evidently, strong trends are easier to detect than weak trends. To detect
a weak trend, either very long time series or very accurate and precise datasets are needed.

The quality of the data is assured within the ICP Forests by means of repeated ring tests that are required for all participating laboratories and the accuracy of the data has been improved considerably over an eight years period (Ferretti and König, 2013; König et al., 2013). However, the precision and accuracy of the dataset still varies across countries and plots. We enhanced the probability of trend detection by the SMK, PMK, and BFAST tests by removing time series with breakpoints caused by artifacts (such as installation effects).

414 Nevertheless, we found a majority of non-significant trends. For these cases, we cannot state 415 with certainty that DOC did not change over time: it might be that the trend was not strong 416 enough to be detected, or that the data quality was insufficient for the period length available 417 for the trend analysis (more than 9 years in all the cases). For example, the mixed-effects 418 models detected a positive trend in the organic layer, and while many of the individual time 419 series measured in the organic layer also showed a positive trend, most were classified as 420 non-significant trends (Table 1; Fig. 3). This probably led to an underestimation of trends that 421 separately might not be strong enough to be detected by the individual trend analysis but 422 combined with the other European data these sites may contribute to an overall trend of 423 increasing DOC concentrations in soils of European forests. Nevertheless, the selected trend 424 analysis techniques (SMK and PMK) are the most suitable to detect weak trends (Marchetto 425 et al., 2013; Waldner et al., 2014), thus reducing the chances of hidden trends within the non-426 significant trends category.

427 On the other hand, evaluating hundreds of time series may introduce random effects that may 428 cause the detection of false significant trends. This multiple testing effect was controlled by 429 evaluating the trends at a 0.01 significance level: Increasing the significance level hardly 430 changed the number of detected significant trends (positive trends: 91 (p<0.05) vs. 70 431 (p<0.01); negative trends: 63 (p<0.05) versus 50 (p<0.01)). Since the detected trends at 0.01 432 significance level outnumbered those expected just by chance at the 0.05 level (13 out of 258 433 cases), it is guaranteed that the detected positive and negative trends were real and not a 434 result of a multiple testing effect.

435 **4.1.2** Analysis of breakpoints in the time series

436 Soil solution DOC time series measured with lysimeters are subject to possible interruptions 437 of monotonicity, which is manifested by breakpoints. For instance, installation effect, 438 collector replacement, local forest management, disturbance by small animals, or by single or 439 repeated canopy insect infestations may disrupt DOC concentrations through abrupt soil 440 disturbances and/or enhanced input from the canopy to the soil (Akselsson et al., 2013; Kvaalen et al., 2002; Lange et al., 2006; Moffat et al., 2002; Pitman et al., 2010). In general, 441 442 detailed information on the management history and other local disturbances was lacking for 443 the majority of Level II plots, which hinders assigning observed breakpoints to specific site 444 conditions. The BFAST analysis allowed us to filter out time series affected by local 445 disturbances (natural or artefacts) from the dataset and to solely retain time series with 446 monotonic trends. By applying the breakpoint analysis, we reduced the within-plot trend 447 variability, while most of the plots showed similar aggregated trends per plot-depth 448 combinations (Supplementary material, Fig. S4). Thereby, we removed some of the within-449 plot variability that might be caused by local factors not directly explaining the long-term 450 monotonic trends in DOC and thus complicating or confounding the trend analysis (Clark et 451 al., 2010).

452 In view of these results, we recommend that testing for monotonicity of the individual time 453 series is a necessary first step in this type of analyses and that the breakpoint analysis is an 454 appropriate tool to filter large datasets prior to analyzing the long-term temporal trends in 455 DOC concentrations. It is worth mentioning that, by selecting monotonic trends, we selected 456 a subset of the trends for which it is more likely to relate the observed trends to 457 environmental changes. A focus on monotonic trends does not imply that the trends with 458 breakpoints are not interesting, further work is needed to interpret the causes of these abrupt 459 changes and verify if these are artefacts or mechanisms, since they may also contain useful 460 information on local factors affecting DOC trends, such as forest management or extreme 461 events (Tetzlaff et al., 2007). This level of detail is, however, not yet available for the ICP 462 Forests Level II plots.

463 **4.1.3** Variability in soil solution DOC trends within plots

Even after removing sites with breakpoints in the time series, within-plot trend variability remained high (median within-plot range: 3.3% yr⁻¹), with different trends observed for different collectors from the same plot (Fig. 5). This high small-scale variability in soil solution DOC makes it difficult to draw conclusions about long-term DOC trends from individual site measurements, particularly in plots with heterogeneous soil and site conditions (Löfgren et al., 2010).

The trends in soil solution DOC also varied across soil depth intervals. The mixed-effectmodels suggested an increasing trend in soil solution DOC concentration in the organic layer,

472 and an increasing trend in soil solution DOC concentration under 80 cm depth only when the 473 entire dataset (with breakpoints) was analyzed. The individual trend analyses confirmed the 474 increasing trend under the organic layer (Table 1), while more heterogeneous trends in the 475 mineral soil were found, which is in line with previous findings (Borken et al., 2011; Evans et al., 2012; Hruška et al., 2009; Löfgren and Zetterberg, 2011; Sawicka et al., 2016; 476 477 Vanguelova et al., 2010). This difference has been attributed to different processes affecting 478 DOC in the organic layer and top mineral soil and in the subsoil. External factors such as acid 479 deposition may have a more direct effect in the organic layer where interaction between DOC 480 and mineral phases is less important compared to deeper layers of the mineral soil (Fröberg et al., 2006). However, DOC measurements are not available for all depths at each site, 481 482 complicating the comparison of trends across soil depth intervals. Hence, the depth-effect on 483 trends in soil solution DOC cannot be consistently addressed within this study (see 484 Supplementary material, Fig. S1, Fig. S2).

Finally, the direction of the trends in soil solution DOC concentrations did not follow a clear 485 regional pattern across Europe (Fig. 3 and 4) and even contrasted with other soil solution 486 parameters that showed widespread trends over Europe, such as decreasing SO_4^{2-} and 487 488 increasing pH. This finding indicates that effects of environmental controls on soil solution 489 DOC concentrations may differ depending on local factors like soil type (e.g., soil acidity, 490 texture) as well as site and stand characteristics (e.g., tree growth or acidification history). 491 Thus, the trends in DOC in soil solution appear to be an outcome of interactions between 492 controls acting at local and regional scales.

In order to compare soil solution DOC trends among sites, trends of DOC concentrations are always expressed in relative trends (% yr⁻¹). By using the relative trends, we removed the effect of the median DOC concentration at the "plot-depth" combination and, consequently, the results do not reflect the actual magnitude of the trend, but their importance in relation with the median DOC concentration at the "plot-depth" combination. It implies that the interpretation of our results was done only in relative terms (see Supplementary Material, Table S3, Fig S5).

500 **4.2** Controls on soil solution DOC temporal trends

501 **4.2.1 Vegetation**

502 Biological controls on DOC production and consumption, like net primary production (NPP), 503 operating at site or catchment level, are particularly important when studying soil solution as 504 plant-derived carbon is the main source of DOC (Harrison et al., 2008). Stem growth was 505 available as a proxy for NPP only for 53 sites and was calculated as the increment between 506 inventories carried out every five years. Similarly to what has been found for peatlands (Billett et al., 2010; Dinsmore et al., 2013), the results suggest that vegetation growth is an 507 508 important driver of DOC temporal dynamics in forests.. Differences in DOC temporal trends 509 across all soil depths were strongly related to stem growth, with more productive plots, as indicated by higher stem volume increment (6-12 m^3 ha⁻¹ yr⁻¹), more often exhibiting 510 511 decreasing trends in DOC (Fig. 6 and 7). .

512 The drivers of variation in forest productivity and its relationship with trends in DOC 513 concentrations are yet unclear. Forest productivity might indirectly affect DOC trends 514 through changes in soil solution chemistry (via cation uptake) (Vanguelova et al., 2007), but 515 the available data do not allow to test this. Alternatively, variation in plant carbon allocation 516 and therefore in the relationship between aboveground productivity and belowground C 517 inputs can strongly influence the relationship between forest productivity and DOC trends. 518 For example, nutrient availability strongly influences plant C allocation (Poorter et al., 2012; 519 Vicca et al., 2012), with plants in nutrient rich soils investing more in aboveground tissue at 520 the expense of belowground C allocation. Assuming that more productive forests are located 521 in more fertile plots, the decreasing trends in DOC concentrations may result from reduced C 522 allocation to the belowground nutrient acquisition system (Vicca et al., 2012), hence, 523 reducing an important source of belowground DOC.

524 Further research assessing nutrient availability and determining the drivers of variation in 525 forest productivity, allocation and DOC is needed to verify the role of nutrients and other 526 factors (e.g., climate, stand age, management) in DOC trends and disentangle the 527 mechanisms behind the effect of forest productivity on soil solution DOC trends.

528 **4.2.2** Acidifying deposition

529 Decreased atmospheric SO_4^{2-} deposition and accumulation of atmospherically deposited N 530 were hypothesized to increase DOC in European surface waters over the last 20 years (Evans 531 et al., 2005; Hruška et al., 2009; Monteith et al., 2007). Sulphate and inorganic N deposition decreased in Europe over the past decades (Waldner et al., 2014) but trends in soil solution 532 533 DOC concentrations varied greatly, with increases, decreases, as well as steady states being 534 observed across respectively 56, 41 and 77 time series in European forests (Fig. 3,4 and 8). Although we could not demonstrate a direct effect of trends in SO_4^{2-} and inorganic N 535 deposition on the trends of soil solution DOC concentration, the multivariate analysis 536 537 suggested that the hypothesis of increased DOC soil solution concentration as a result of decreasing SO_4^{2-} deposition may apply only at sites with low or medium mean N deposition 538 539 over the last decades.

540 Our results show that DOC concentrations in the soil solution are positively linked to 541 inorganic N deposition loads at sites with low or medium inorganic N deposition, and to N 542 deposition trends at sites with high inorganic N deposition (Fig. 7). The role of atmospheric 543 inorganic N deposition in increasing DOC leaching from soils has been well documented 544 (Bragazza et al., 2006; Liu and Greaver, 2010; Pregitzer et al., 2004; Rosemond et al., 2015). 545 The mechanisms behind this positive relationship are either physico-chemical or biological. 546 Chemical changes in soil solution through the increase of NO_3^{-1} ions can trigger desorption of 547 DOC (Pregitzer et al., 2004), and biotic forest responses to inorganic N deposition, namely, enhanced photosynthesis, altered carbon allocation, and reduced soil microbial activity 548 549 (Bragazza et al., 2006; de Vries et al., 2009; Janssens et al., 2010; Liu and Greaver, 2010), 550 can increase the final amount of DOC in the soil. As the most consistent trends are found in 551 organic layers, where production/decomposition control DOC concentration (Löfgren and 552 Zetterberg, 2011), effects of inorganic N deposition through increase of primary productivity 553 (de Vries et al., 2014; de Vries et al., 2009; Ferretti et al., 2014) are likely drivers of 554 increasing DOC trends. One proposed mechanism is incomplete lignin degradation and 555 greater production of DOC in response to increased soil NH_4^+ (Pregitzer et al., 2004; Zech et al., 1994). Alternatively, N-induced reductions of forest heterotrophic respiration (Janssens et 556 557 al., 2010) and reduced microbial decomposition (Liu and Greaver, 2010) may lead to greater 558 accumulation of DOC.

559 Moreover, our results suggested that only at sites with lower and medium inorganic N 560 deposition, decreasing trends in $SO_4^{2^-}$ deposition coincided with increasing trends in soil 561 solution DOC (Supplementary Material, Fig. S3), as previously hypothesized for surface 562 waters, indicating an interaction between the inorganic N deposition loads and the 563 mechanisms underlying the temporal change of soil solution DOC.

Similar to our observation for soil solution DOC, decreasing SO_4^{2-} deposition has been linked 564 to increasing surface water DOC (Evans et al., 2006; Monteith et al., 2007; Oulehle and 565 Hruska, 2009). Sulphate deposition triggers soil acidification and a subsequent release of Al³⁺ 566 in acid soils. The amount of Al^{3+} is negatively related to soil solution DOC due to two 567 plausible mechanisms: 1) The released Al³⁺ can build complexes with organic molecules, 568 enhancing DOC precipitation and, in turn, suppressing DOC solubility, thereby decreasing 569 570 DOC concentrations in soil solution (de Wit et al., 2001; Tipping and Woof, 1991; Vanguelova et al., 2010), and 2) at higher levels of soil solution Al^{3+} in combination with low 571 pH, DOC production through SOM decomposition decreases due to toxicity of Al³⁺ to soil 572 organisms (Mulder et al., 2001). Consequently, when SO_4^{2-} deposition is lower, increases of 573 574 soil solution DOC concentration could be expected (Fig. 7A, B). Finally, an indirect effect of 575 plant response to nutrient-limited acidified soil could also contribute to the trend in soil 576 solution DOC by changes to plant belowground C allocation (Vicca et al., 2012) (see Sect. 4.2.1.). 577

Nevertheless, increasing DOC soil solution concentration as a result of decreasing SO_4^{2-} 578 579 deposition occurred only at sites with low or medium mean N deposition. Therefore, our 580 results indicate that the response of DOC to changes in atmospheric deposition seems to be 581 controlled by the past and present inorganic N deposition loads (Clark et al., 2010; Evans et al., 2012; Tian and Niu, 2015). It suggests that the mechanisms of recovery from SO_4^{2-} 582 583 deposition and acidification take place only in low and medium N deposition areas, as has been observed for inorganic N deposition effects (de Vries et al., 2009). In high inorganic N 584 deposition areas, it is likely that impacts of N-induced acidification on forest health and soil 585 condition lead to more DOC leaching, even though SO_4^{2-} deposition has been decreasing. 586 Therefore, the hypothesis of recovery from acidity cannot fully explain overall soil solution 587 588 DOC trends in Europe, as was also previously suggested in local or national studies of longterm trends in soil solution DOC (Löfgren et al., 2010; Stutter et al., 2011; Ukonmaanaho et 589 al., 2014; Verstraeten et al., 2014). Collinearity between SO_4^{2-} deposition and inorganic N 590 deposition was low (variance inflation factor <3) for both the mean values and temporal 591 trends. We therefore assumed that the proposed response of DOC to the decline in SO_4^{2-} 592 deposition in low to medium N areas is not confounded by simultaneous changes in SO_4^{2-} and 593 594 NO_3^- deposition, even more so because the statistical models account for the covariation in SO_4^{2-} and NO_3^{-} deposition (Figure 7). Nonetheless, as SO_4^{2-} and NO_3^{-} deposition are 595

- 596 generally decreasing across Europe (Figure 8), concomitant changes in NO_3^- deposition may 597 still have somewhat confounded the attribution of DOC changes solely to SO_4^{2-} deposition.
- 598 Ultimately, internal soil processes control the final concentration of DOC in the soil solution. 599 The solubility and biological production and consumption of DOC are regulated by pH, ionic strength of the soil solution and the presence of Al^{3+} and Fe (Bolan et al., 2011; De Wit et al., 600 2007; Schwesig et al., 2003). These conditions are modulated by changes in atmospheric 601 602 deposition but not uniformly across sites: soils differ in acid-buffering capacity (Tian and Niu, 2015), and the response of DOC concentrations to changes in SO_4^{2-} deposition will thus 603 be a function of the initial soil acidification and buffer range (Fig. 7). Finally, modifications 604 605 of soil properties induced by changes in atmospheric deposition are probably an order of 606 magnitude lower than the spatial variation of these soil properties across sites, making it 607 difficult to isolate controlling factors on the final observed response of soil solution DOC at 608 continental scale (Clark et al., 2010; Stutter et al., 2011).
- 609 In conclusion, our results confirm the long-term trends of DOC in soil solution as a 610 consequence of the interactions between local (soil properties, forest growth), and regional 611 (atmospheric deposition) controls acting at different temporal scales. However, further work is needed to quantify the role of each mechanism underlying the final response of soil 612 613 solution DOC to environmental controls. We recommend that particular attention should be 614 paid to the biological controls (e.g., net primary production, root exudates or litterfall and 615 canopy infestations) on long-term trends in soil solution DOC, which remains poorly 616 understood.

617 **4.3** Link between DOC trends in soil and streams

618 An underlying question is how DOC trends in soil solution relate to DOC trends in stream 619 waters. Several studies have pointed out recovery from acidification as a cause for increasing 620 trends in DOC concentrations in surface waters (Dawson et al., 2009; Evans et al., 2012; 621 Monteith et al., 2007; Skjelkvåle et al., 2003). Overall, our results point to a noticeable 622 increasing trend in DOC in the organic layer of forest soils, which is qualitatively consistent 623 with the increasing trends found in stream waters and in line with positive DOC trends 624 reported for the soil organic layer or at maximum 10 cm depth of the mineral soil in Europe 625 (Borken et al., 2011; Hruška et al., 2009; Vanguelova et al., 2010). DOC from the organic 626 layer may be transferred to surface waters via hydrologic shortcuts during storm events, when

shallow lateral flow paths are activated. On the other hand, trends in different soil layersalong the mineral soil were more variable and responded to other soil internal processes.

629 It is currently difficult to link long-term dynamics in soil and surface water DOC. Large scale 630 processes become more important than local factors when looking at DOC trends in surface 631 waters (Lepistö et al., 2014), while the opposite seems to apply for soil solution DOC trends. 632 Furthermore, stream water DOC mainly reflects the processes occurring in areas with a high 633 hydraulic connectivity in the catchment, such as peat soils or floodplains, which normally yield most of the DOC (Ledesma et al., 2016; Löfgren and Zetterberg, 2011). Further 634 635 monitoring studies in forest soils with high hydraulic connectivity to streams are needed to be 636 able to link dynamics of DOC in forest soil with dynamics of DOC in stream waters.

637 Finally, stream water DOC trends are dominantly controlled by catchment hydrology 638 (Sebestyen et al., 2009; Stutter et al., 2011; Tranvik and Jansson, 2002), since an increase in 639 DOC concentration does not necessarily result in increased DOC transport, which is the 640 product of DOC concentration and discharge. Differences in hydrology among sites may 641 (partly) explain the inconsistent patterns found in soil solution DOC concentration trends at 642 different sites and depths, as previously proposed (Stutter et al., 2011), but data to verify this 643 statement are currently not available. Hence, while this study of controls on trends in DOC 644 concentrations in soil provides key information for predictions of future C losses to stream 645 waters, future studies at larger scale that include catchment hydrology (precipitation, runoff 646 and drainage) are crucial to relate soil and stream DOC trends.

647 **5** Conclusions

Different monotonic long-term trends of soil solution DOC have been found across European 648 649 forests at plot scale, with the majority of the trends for specific plots and depths not being 650 statistically significant (40%), followed by significantly positive (35%) and significantly 651 negative trends (25%). The distribution of the trends did not follow a specific regional 652 pattern. A multivariate analysis revealed a negative relation between long-term trends in soil solution DOC and mean SO_4^{2-} deposition and a positive relation to mean NO_3^{-} deposition. 653 While the hypothesis of increasing trends of DOC due to reductions of SO_4^{2-} deposition could 654 be confirmed in low to medium N deposition areas, there was no significant relationship with 655 SO_4^{2-} deposition in high N deposition areas. There was evidence that an overall increasing 656 657 trend of DOC concentrations occurred in the organic layers and, to a lesser extent, in the deep 658 mineral soil. However, trends in the different mineral soil horizons were highly heterogeneous, indicating that internal soil processes control the final response of DOC in soil solution. Although correlative, our results suggest that there is no single mechanism responsible for soil solution DOC trends operating at large scale across Europe but that interactions between controls operating at local (soil properties, site and stand characteristics) and regional (atmospheric deposition changes) scales are taking place.

664 Acknowledgements

We want to thank the numerous scientists and technicians who were involved in the data 665 666 collection, analysis, transmission, and validation of the ICP Forests Monitoring Programme 667 across the UNECE region from which data have been used in this work. The evaluation was 668 mainly based on data that are part of the UNECE ICP Forests PCC Collaborative Database 669 (see <u>www.icp-forests.net</u>) or national Databases (e.g., Eberswalde Forestry State Center of 670 Excellence (Landeskompetenzzentrum Forst Eberswalde, LFE) for parts of the data for 671 Germany). For soil, we used and acknowledge the aggregated forest soil condition database 672 (AFSCDB.LII.2.1) compiled by the ICP Forests Forest Soil Coordinating Centre. The long-673 term collection of forest monitoring data was to a large extent funded by national research 674 institutions and ministries, with support from further bodies, services and land owners. It was partially funded by the European Union under the Regulation (EC) No. 2152/2003 675 676 concerning monitoring of forests and environmental interactions in the Community (Forest 677 Focus) and the project LIFE 07 ENV/D/000218. SV is a postdoctoral research associate of 678 the Fund for Scientific Research – Flanders. IAJ, JP and PC acknowledge support from the European Research Council Synergy grant ERC-2013-SyG-610028 IMBALANCE-P. 679 Finally, we acknowledge the E-OBS dataset from the EU-FP6 project ENSEMBLES 680 (http://ensembles-eu.metoffice.com) and the data providers in the ECA&D project 681 682 (http://www.ecad.eu). We want to thank three anonymous reviewers for helpful comments on previous manuscript versions. 683

684 **References**

Akselsson, C., Hultberg, H., Karlsson, P. E., Karlsson, G. P., and Hellsten, S.: Acidification
trends in south Swedish forest soils 1986-2008: -Slow recovery and high sensitivity to seasalt episodes, Sci Total Environ, 444, 271-287, 2013.

Battin, T. J., Luyssaert, S., Kaplan, L. A., Aufdenkampe, A. K., Richter, A., and Tranvik, L.
J.: The boundless carbon cycle, Nature Geoscience, 9, 2009.

Bianchi, T. S.: The role of terrestrially derived organic carbon in the coastal ocean: A
changing paradigm and the priming effect, P Natl Acad Sci USA, 108, 19473-19481, 2011.

- 692 Billett, M. F., Charman, D. J., Clark, J. M., Evans, C. D., Evans, M. G., Ostle, N. J., Worrall,
- F., Burden, A., Dinsmore, K. J., Jones, T., McNamara, N. P., Parry, L., Rowson, J. G., and
 Rose, R.: Carbon balance of UK peatlands: current state of knowledge and future research
 challenges, Clim Res, 45, 13-29, 2010.
- 696 Bolan, N. S., Adriano, D. C., Kunhikrishnan, A., James, T., McDowell, R., and Senesi, N.:
- Dissolved Organic Matter: Biogeochemistry, Dynamics, and Environmental Significance in
 Soils. Adv Agron, 2011.
- 699 Borken, W., Ahrens, B., Schulz, C., and Zimmermann, L.: Site-to-site variability and 700 temporal trends of DOC concentrations and fluxes in temperate forest soils, Global Change 701 Biol, 17, 2428-2443, 2011.
- Bragazza, L., Freeman, C., Jones, T., Rydin, H., Limpens, J., Fenner, N., Ellis, T., Gerdol, R.,
 Hajek, M., Hajek, T., Lacumin, P., Kutnar, L., Tahvanainen, T., and Toberman, H.:
 Atmospheric nitrogen deposition promotes carbon loss from peat bogs. P Natl Acad Sci USA,
 51, 2006.
- Buckingham, S., Tipping, E., and Hamilton-Taylor, J.: Concentrations and fluxes ofdissolved organic carbon in UK topsoils. Sci Total Environ, 1, 2008.
- 708 Camino-Serrano, M., Gielen, B., Luyssaert, S., Ciais, P., Vicca, S., Guenet, B., De Vos, B.,
- 709 Cools, N., Ahrens, B., Arain, M. A., Borken, W., Clarke, N., Clarkson, B., Cummins, T.,
- 710 Don, A., Pannatier, E. G., Laudon, H., Moore, T., Nieminen, T. M., Nilsson, M. B., Peichl,
- 711 M., Schwendenmann, L., Siemens, J., and Janssens, I.: Linking variability in soil solution
- dissolved organic carbon to climate, soil type, and vegetation type, Global Biogeochem Cy,28, 497-509, 2014.
- Clark, J. M., Bottrell, S. H., Evans, C. D., Monteith, D. T., Bartlett, R., Rose, R., Newton, R.
 J., and Chapman, P. J.: The importance of the relationship between scale and process in understanding long-term DOC dynamics, Sci Total Environ, 408, 2768-2775, 2010.
- 717 Cools, N. and De Vos, B.: A harmonised Level II soil database to understand processes and 718 changes in forest condition at the European Level. In: Michel, A., Seidling, W., editors. 2014.
- Forest Condition in Europe: 2014 Technical Report of ICP Forests., Vienna, 72-90 pp., 2014.
- Davidson, E. A. and Janssens, I. A.: Temperature sensitivity of soil carbon decompositionand feedbacks to climate change, Nature, 440, 165-173, 2006.
- Dawson, J. J. C., Malcolm, I. A., Middlemas, S. J., Tetzlaff, D., and Soulsby, C.: Is the Composition of Dissolved Organic Carbon Changing in Upland Acidic Streams?, Environ Sci
- 724 Technol, 43, 7748-7753, 2009.
- de Jong, R., Verbesselt, J., Zeileis, A., and Schaepman, M. E.: Shifts in Global Vegetation
 Activity Trends, Remote Sens, 5, 1117-1133, 2013.
- de Vries, W., Reinds, G. J., Posch, M., Sanz, M. J., Krause, G. H. M., Calatayud, V., Renaud,
- J. P., Dupouey, J. L., Sterba, H., Vel., E. M., Dobbertin, M., Gundersen, P., and Voogd, J. C.
- H.: Intensive Monitoring of Forest Ecosystems in Europe, 2003 Technical Report. EC,
- 730 UNECE, Brussels, Geneva, 163 pp., 2003.
- de Vries, W., Solberg, S., Dobbertin, M., Sterba, H., Laubhann, D., van Oijen, M., Evans, C.,
- 732 Gundersen, P., Kros, J., Wamelink, G. W. W., Reinds, G. J., and Sutton, M. A.: The impact
- of nitrogen deposition on carbon sequestration by European forests and heathlands, Forest
- 734 Ecol Manag, 258, 1814-1823, 2009.

- de Vries, W., Du, E., and Butterbach-Bahl, K.: Short and long-term impacts of nitrogen
 deposition on carbon sequestration by forest ecosystems, Current Opinion in Environmental
 Sustainability, 9–10, 90-104, 2014.
- de Wit, H. A., Groseth, T., and Mulder, J.: Predicting Aluminum and Soil Organic Matter
 Solubility Using the Mechanistic Equilibrium Model WHAM, Soil Sci. Soc. Am. J., 65,
 1089-1100, 2001.
- 741 de Wit, H. A., Mulder, J., Hindar, A., and Hole, L.: Long-term increase in dissolved organic
- 742 carbon in streamwaters in Norway is response to reduced acid deposition, Environ Sci
- 743 Technol, 41, 7706-7713, 2007.
- Dinsmore, K. J., Billett, M. F., and Dyson, K. E.: Temperature and precipitation drive
 temporal variability in aquatic carbon and GHG concentrations and fluxes in a peatland
 catchment, Global Change Biol, 19, 2133-2148, 2013.
- Evans, C. D., Monteith, D. T., and Cooper, D. M.: Long-term increases in surface water
 dissolved organic carbon: Observations, possible causes and environmental impacts, Environ
 Pollut, 137, 55-71, 2005.
- 750 Evans, C. D., Chapman, P. J., Clark, J. M., Monteith, D. T., and Cresser, M. S.: Alternative
- explanations for rising dissolved organic carbon export from organic soils, Global ChangeBiol, 12, 2044-2053, 2006.
- 753 Evans, C. D., Jones, T. G., Burden, A., Ostle, N., Zielinski, P., Cooper, M. D. A., Peacock,
- M., Clark, J. M., Oulehle, F., Cooper, D., and Freeman, C.: Acidity controls on dissolved organic carbon mobility in organic soils, Global Change Biol, 18, 3317-3331, 2012.
- Ferretti, M. and Fischer, R.: Forest monitoring, methods for terrestrial investigation in Europewith an overview of North Ameria and Asia, Elsevier, UK, 2013.
- Ferretti, M. and König, N.: Chapter 20 Quality Assurance in International Forest
 Monitoring in Europe. In: Developments in Environmental Science, Ferretti, M. and Fischer,
 R. (Eds.), Elsevier, 2013.
- 761 Ferretti, M., Marchetto, A., Arisci, S., Bussotti, F., Calderisi, M., Carnicelli, S., Cecchini, G.,
- Fabbio, G., Bertini, G., Matteucci, G., de Cinti, B., Salvati, L., and Pompei, E.: On the tracks
 of Nitrogen deposition effects on temperate forests at their southern European range an
- 764 observational study from Italy, Global Change Biol, 20, 3423-3438, 2014.
- Fox, J., Nie, Z., and Byrnes, J.: sem: Structural Equation Models., 2013.
- Freeman, C., Fenner, N., Ostle, N. J., Kang, H., Dowrick, D. J., Reynolds, B., Lock, M. A.,
 Sleep, D., Hughes, S., and Hudson, J.: Export of dissolved organic carbon from peatlands
 under elevated carbon dioxide levels, Nature, 430, 195-198, 2004.
- Fröberg, M., Berggren, D., Bergkvist, B., Bryant, C., and Mulder, J.: Concentration and
 fluxes of dissolved organic carbon (DOC) in three Norway spruce stands along a climatic
 gradient in Sweden, Biogeochemistry, 77, 1-23, 2006.
- 772 Graf Pannatier, E. G., Thimonier, A., Schmitt, M., Walthert, L., and Waldner, P.: A decade of
- 773 monitoring at Swiss Long-Term Forest Ecosystem Research (LWF) sites: can we observe
- trends in atmospheric acid deposition and in soil solution acidity?, Environmental monitoring
- and assessment, 174, 3-30, 2011.
- Granke, O.: Chapter 23 Methods for Database Quality Assessment. In: Developments in
 Environmental Science, Ferretti, M. and Fischer, R. (Eds.), Elsevier, 2013.

- Haaland, S., Hongve, D., Laudon, H., Riise, G., and Vogt, R. D.: Quantifying the Drivers of
- the Increasing Colored Organic Matter in Boreal Surface Waters, Environ Sci Technol, 44,2975-2980, 2010.
- Hansen, K., Vesterdal, L., Bastrup-Birk, A., and Bille-Hansen, J.: Are indicators for critical
 load exceedance related to forest condition?, Water Air Soil Poll, 183, 293-308, 2007.
- Harrison, A. F., Taylor, K., Scott, A., Poskitt, J., Benham, D., Grace, J., Chaplow, J., and
 Rowland, P.: Potential effects of climate change on DOC release from three different soil
 types on the Northern Pennines UK: examination using field manipulation experiments,
 Global Change Biol, 14, 687-702, 2008.
- Hartley, I. P. and Ineson, P.: Substrate quality and the temperature sensitivity of soil organic
 matter decomposition, Soil Biol Biochem, 40, 1567-1574, 2008.
- Haylock, M. R., Hofstra, N., Tank, A. M. G. K., Klok, E. J., Jones, P. D., and New, M.: A
 European daily high-resolution gridded data set of surface temperature and precipitation for
 1950-2006, J Geophys Res-Atmos, 113, 2008.
- Hirsch, R. M., Slack, J. R., and Smith, R. A.: Techniques of Trend Analysis for Monthly
 Water-Quality Data, Water Resour Res, 18, 107-121, 1982.
- Hruška, J., Kram, P., Mcdowell, W. H., and Oulehle, F.: Increased Dissolved Organic Carbon
 (DOC) in Central European Streams is Driven by Reductions in Ionic Strength Rather than
- 796 Climate Change or Decreasing Acidity, Environ Sci Technol, 43, 4320-4326, 2009.
- ICP Forests: Manual on methods and criteria for harmonized sampling, assessment,
 monitoring and analysis of the effects of air pollution on forests., UNECE ICP Forests
 Programme Co-ordinating Centre, Hamburg, Germany, [http://www.icpforests.org/Manual.htm], 2010.
- Janssens, I. A., Dieleman, W., Luyssaert, S., Subke, J. A., Reichstein, M., Ceulemans, R.,
 Ciais, P., Dolman, A. J., Grace, J., Matteucci, G., Papale, D., Piao, S. L., Schulze, E. D.,
 Tang, J., and Law, B. E.: Reduction of forest soil respiration in response to nitrogen
 deposition, Nature Geoscience, 3, 315-322, 2010.
- Kalbitz, K., Solinger, S., Park, J. H., Michalzik, B., and Matzner, E.: Controls on the dynamics of dissolved organic matter in soils: A review, Soil Sci, 165, 277-304, 2000.
- König, N., Cools, N., Derome, K., Kowalska, A., De Vos, B., Fürst, A., Marchetto, A.,
 O'Dea, P., and Tartari, G. A.: Chapter 22 Data Quality in Laboratories: Methods and Results
 for Soil, Foliar, and Water Chemical Analyses. In: Developments in Environmental Science,
- 810 Ferretti, M. and Fischer, R. (Eds.), Elsevier, 2013.
- 811 Kvaalen, H., Solberg, S., Clarke, N., Torp, T., and Aamlid, D.: Time series study of 812 concentrations of SO_4^{2-} and H⁺ in precipitation and soil waters in Norway, Environ Pollut, 813 117, 215-224, 2002.
- Lange, H., Solberg, S., and Clarke, N.: Aluminum dynamics in forest soil waters in Norway,
 Sci Total Environ, 367, 942-957, 2006.
- 816 Ledesma, J. L. J., Futter, M. N., Laudon, H., Evans, C. D., and Kohler, S. J.: Boreal forest
- riparian zones regulate stream sulfate and dissolved organic carbon, Sci Total Environ, 560,
 110-122, 2016.
- Lepistö, A., Futter, M. N., and Kortelainen, P.: Almost 50 years of monitoring shows that climate, not forestry, controls long-term organic carbon fluxes in a large boreal watershed,
- 821 Global Change Biol, 20, 1225-1237, 2014.

- Libiseller, C. and Grimvall, A.: Performance of partial Mann-Kendall tests for trend detection in the presence of covariates, Environmetrics, 13, 71-84, 2002.
- 824 Lindroos, A.-J., Derome, J., Starr, M., and Ukonmaanaho, L.: Effects of Acidic Deposition
- 825 on Soil Solution Quality and Nutrient Leaching in Forest Soils. In: Forest Condition in a 826 Changing Environment, Mälkönen, E. (Ed.), Forestry Sciences, Springer Netherlands, 2000.
- Liu, L. and Greaver, T. L.: A global perspective on belowground carbon dynamics under nitrogen enrichment, Ecol Lett, 13, 819-828, 2010.
- Löfgren, S., Gustafsson, J. P., and Bringmark, L.: Decreasing DOC trends in soil solution along the hillslopes at two IM sites in southern Sweden - Geochemical modeling of organic matter solubility during acidification recovery, Sci Total Environ, 409, 201-210, 2010.
- Löfgren, S. and Zetterberg, T.: Decreased DOC concentrations in soil water in forested areas
 in southern Sweden during 1987-2008, Sci Total Environ, 409, 1916-1926, 2011.
- 834 Marchetto, A., Mosello, R., Tartari, G., Derome, J., Derome, K., König, N., Clarke, N., and
- 835 Kowalska, A.: Atmospheric Deposition and Soil Solution Working Ring Test 2009, Project
- 836 FutMon, Verbania Pallanza, 41 pp., 2011.
- Marchetto, A., Rogora, M., and Arisci, S.: Trend analysis of atmospheric deposition data: A
 comparison of statistical approaches, Atmos Environ, 64, 95-102, 2013.
- McDowell, W. H. and Likens, G. E.: Origin, Composition, and Flux of Dissolved OrganicCarbon in the Hubbard Brook Valley, Ecol Monogr, 58, 177-195, 1988.
- Moffat, A. J., Kvaalen, H., Solberg, S., and Clarke, N.: Temporal trends in throughfall and
 soil water chemistry at three Norwegian forests, 1986-1997, Forest Ecol Manag, 168, 15-28,
 2002.
- Monteith, D. T., Stoddard, J. L., Evans, C. D., de Wit, H. A., Forsius, M., Hogasen, T.,
 Wilander, A., Skjelkvale, B. L., Jeffries, D. S., Vuorenmaa, J., Keller, B., Kopacek, J., and
 Vesely, J.: Dissolved organic carbon trends resulting from changes in atmospheric deposition
 chemistry, Nature, 450, 537-U539, 2007.
- Mulder, J., De Wit, H. A., Boonen, H. W. J., and Bakken, L. R.: Increased levels of aluminium in forest soils: Effects on the stores of soil organic carbon, Water Air Soil Poll, 130, 989-994, 2001.
- Nieminen, T. M.: Soil Solution Collection and Analysis. Manual Part XI, In: ICP Forests,
 2010: Manual on methods and criteria for harmonized sampling, assessment, monitoring and
 analysis of the effects of air pollution on forests. UNECE ICP Forests Programme Coordinating Centre, Hamburg, 30 pp., [http://www.icp-forests.org/Manual.htm], 2011.
- Oulehle, F. and Hruska, J.: Rising trends of dissolved organic matter in drinking-water
 reservoirs as a result of recovery from acidification in the Ore Mts., Czech Republic, Environ
 Pollut, 157, 3433-3439, 2009.
- Oulehle, F., Evans, C. D., Hofmeister, J., Krejci, R., Tahovska, K., Persson, T., Cudlin, P., and Hruska, J.: Major changes in forest carbon and nitrogen cycling caused by declining sulphur deposition, Global Change Biol, 17, 3115-3129, 2011.
- Pitman, R. M., Vanguelova, E. I., and Benham, S. E.: The effects of phytophagous insects on
 water and soil nutrient concentrations and fluxes through forest stands of the Level II
- 863 monitoring network in the UK, Sci Total Environ, 409, 169-181, 2010.

- Poorter, H., Niklas, K. J., Reich, P. B., Oleksyn, J., Poot, P., and Mommer, L.: Biomass allocation to leaves, stems and roots: meta-analyses of interspecific variation and environmental control, New Phytologist, 193, 30-50, 2012.
- Pregitzer, K. S., Zak, D. R., Burton, A. J., Ashby, J. A., and MacDonald, N. W.: Chronic
 nitrate additions dramatically increase the export of carbon and nitrogen from northern
 hardwood ecosystems, Biogeochemistry, 68, 179-197, 2004.
- R Core Team: R: A language and environment for statistical computing., R Foundation for
 Statistical Computing, Vienna, Austria, 2014.
- 872 Regnier, P., Friedlingstein, P., Ciais, P., Mackenzie, F. T., Gruber, N., Janssens, I. A.,
- 873 Laruelle, G. G., Lauerwald, R., Luyssaert, S., Andersson, A. J., Arndt, S., Arnosti, C.,
- Borges, A. V., Dale, A. W., Gallego-Sala, A., Goddéris, Y., Goossens, N., Hartmann, J.,
 Heinze, C., Ilyina, T., Joos, F., LaRowe, D. E., Leifeld, J., Meysman, F. J. R., Munhoven, G.,
- 876 Raymond, P. A., Spahni, R., Suntharalingam, P., and Thullner, M.: Anthropogenic
- 877 perturbation of the carbon fluxes from land to ocean, Nature Geoscience, 6, 2013.
- 878 Rosemond, A. D., Benstead, J. P., Bumpers, P. M., Gulis, V., Kominoski, J. S., Manning, D.
- W. P., Suberkropp, K., and Wallace, J. B.: Experimental nutrient additions accelerate terrestrial carbon loss from stream ecosystems, Science, 347, 1142-1145, 2015.
- 881 Sarkkola, S., Koivusalo, H., Lauren, A., Kortelainen, P., Mattsson, T., Palviainen, M.,
- 882 Piirainen, S., Starr, M., and Finer, L.: Trends in hydrometeorological conditions and stream
- 883 water organic carbon in boreal forested catchments, Sci Total Environ, 408, 92-101, 2009.
- Sawicka, K., Monteith, D. T., Vanguelova, E. I., Wade, A. J., and Clark, J. M.: Fine-scale
 temporal characterization of trends in soil water dissolved organic carbon and potential
 drivers, Ecological Indicators, 68, 36-51, 2016.
- Schwertman, N. C., Owens, M. A., and Adnan, R.: A simple more general boxplot method
 for identifying outliers, Comput Stat Data An, 47, 165-174, 2004.
- Schwesig, D., Kalbitz, K., and Matzner, E.: Effects of aluminium on the mineralization of
 dissolved organic carbon derived from forest floors, Eur J Soil Sci, 54, 311-322, 2003.
- 891 Sebestyen, S. D., Boyer, E. W., and Shanley, J. B.: Responses of stream nitrate and DOC
- 892 loadings to hydrological forcing and climate change in an upland forest of the northeastern
- 893 United States, J Geophys Res-Biogeo, 114, 2009.
- Sen, P. K.: Estimates of the Regression Coefficient Based on Kendall's Tau, Journal of the
 American Statistical Association, 63, 1379-1389, 1968.
- Skjelkvåle, B. L., Evans, C., Larssen, T., Hindar, A., and Raddum, G. G.: Recovery from
 acidification in European surface waters: A view to the future, Ambio, 32, 170-175, 2003.
- Stutter, M. I., Lumsdon, D. G., and Rowland, A. P.: Three representative UK moorland soils
 show differences in decadal release of dissolved organic carbon in response to environmental
 change, Biogeosciences, 8, 3661-3675, 2011.
- 901 Sucker, C. and Krause, K.: Increasing dissolved organic carbon concentrations in 902 freshwaters: what is the actual driver?, Iforest, 3, 106-108, 2010.
- Sulkava, M., Rautio, P., and Hollmen, J.: Combining measurement quality into monitoring
 trends in foliar nutrient concentrations, Lect Notes Comput Sc, 3697, 761-767, 2005.

- Tetzlaff, D., Malcolm, I. A., and Soulsby, C.: Influence of forestry, environmental change
 and climatic variability on the hydrology, hydrochemistry and residence times of upland
 catchments, J Hydrol, 346, 93-111, 2007.
- Tian, D. and Niu, S.: A global analysis of soil acidification caused by nitrogen addition,
 Environmental Research Letters, 10, 024019, 2015.
- Tipping, E. and Woof, C.: The distribution of humic substances between the solid and
 aqueous phases of acid organic soils; a description based on humic heterogeneity and chargedependent sorption equilibria, Journal of Soil Science, 42, 437-448, 1991.
- 913 Tranvik, L. J. and Jansson, M.: Climate change Terrestrial export of organic carbon, Nature,
 914 415, 861-862, 2002.
- 915 Ukonmaanaho, L., Starr, M., Lindroos, A. J., and Nieminen, T. M.: Long-term changes in
- acidity and DOC in throughfall and soil water in Finnish forests, Environ Monit Assess, 186,
 7733-7752, 2014.
- 918 Vanguelova, E. I., Hirano, Y., Eldhuset, T. D., Sas-Paszt, L., Bakker, M. R., Puttsepp, U.,
- 919 Brunner, I., Lohmus, K., and Godbold, D.: Tree fine root Ca/Al molar ratio Indicator of Al
- 920 and acidity stress, Plant Biosyst, 141, 460-480, 2007.
- 921 Vanguelova, E. I., Benham, S., Pitman, R., Moffat, A. J., Broadmeadow, M., Nisbet, T.,
- 922 Durrant, D., Barsoum, N., Wilkinson, M., Bochereau, F., Hutchings, T., Broadmeadow, S.,
- 923 Crow, P., Taylor, P., and Houston, T. D.: Chemical fluxes in time through forest ecosystems
- in the UK Soil response to pollution recovery, Environ Pollut, 158, 1857-1869, 2010.
- Verbesselt, J., Hyndman, R., Newnham, G., and Culvenor, D.: Detecting trend and seasonal
 changes in satellite image time series, Remote Sens Environ, 114, 106-115, 2010.
- Verstraeten, A., Neirynck, J., Genouw, G., Cools, N., Roskams, P., and Hens, M.: Impact of
 declining atmospheric deposition on forest soil solution chemistry in Flanders, Belgium,
 Atmos Environ, 62, 50-63, 2012.
- Verstraeten, A., De Vos, B., Neirynck, J., Roskams, P., and Hens, M.: Impact of air-borne or
 canopy-derived dissolved organic carbon (DOC) on forest soil solution DOC in Flanders,
 Belgium, Atmos Environ, 83, 155-165, 2014.
- Vicca, S., Balzarolo, M., Filella, I., Granier, A., Herbst, M., Knohl, A., Longdoz, B., Mund,
 M., Nagy, Z., Pintér, K., Rambal, S., Verbesselt, J., Verger, A., Zeileis, A., Zhang, C., and
 Peñuelas, J.: Remotely-sensed detection of effects of extreme droughts on gross primary
 production, Scientific Reports, 6, 28269, 2016.
- 937 Vicca, S., Luyssaert, S., Penuelas, J., Campioli, M., Chapin, F. S., Ciais, P., Heinemeyer, A.,
- 938 Hogberg, P., Kutsch, W. L., Law, B. E., Malhi, Y., Papale, D., Piao, S. L., Reichstein, M.,
- Schulze, E. D., and Janssens, I. A.: Fertile forests produce biomass more efficiently, Ecol
 Lett, 15, 520-526, 2012.
- 941 Waldner, P., Marchetto, A., Thimonier, A., Schmitt, M., Rogora, M., Granke, O., Mues, V.,
- 942 Hansen, K., Karlsson, G. P., Zlindra, D., Clarke, N., Verstraeten, A., Lazdins, A.,
- 943 Schimming, C., Iacoban, C., Lindroos, A. J., Vanguelova, E., Benham, S., Meesenburg, H.,
- 944 Nicolas, M., Kowalska, A., Apuhtin, V., Napa, U., Lachmanova, Z., Kristoefel, F., Bleeker,
- A., Ingerslev, M., Vesterdal, L., Molina, J., Fischer, U., Seidling, W., Jonard, M., O'Dea, P.,
- Johnson, J., Fischer, R., and Lorenz, M.: Detection of temporal trends in atmospheric
- deposition of inorganic nitrogen and sulphate to forests in Europe, Atmos Environ, 95, 363-374, 2014.

- 949 Worrall, F. and Burt, T.: Time series analysis of long-term river dissolved organic carbon 950 records, Hydrological Processes, 18, 893-911, 2004.
- Wu, Y. J., Clarke, N., and Mulder, J.: Dissolved Organic Carbon Concentrations in 951 Throughfall and Soil Waters at Level II Monitoring Plots in Norway: Short- and Long-Term 952 Variations, Water Air Soil Poll, 205, 273-288, 2010.
- 953
- 954 Zech, W., Guggenberger, G., and Schulten, H. R.: Budgets and Chemistry of Dissolved
- Organic-Carbon in Forest Soils Effects of Anthropogenic Soil Acidification, Sci Total 955
- Environ, 152, 49-62, 1994. 956
- 957

Table 1. Temporal trends of DOC concentrations obtained with the linear mixed models (LMM) built for different forest types, soil depth intervals and collector types with the entire dataset (with breakpoints) and with the dataset without time series showing breakpoints (without breakpoints) and the Seasonal Mann Kendal tests (SMK). The table shows the median DOC concentrations in mg L⁻¹ ([DOC]), relative trend slope (rslope in % yr-1), the number of observations (n) and the p value. For the SMK tests, the number of time series showing significant negative (N), non-significant (NS) and significant positive (P) trends are shown and the interquartile range of the rslope is between brackets. LMMs for which no statistically significant trend was detected (p>0.1) are represented in grey, the LMMs for which a significant trend is detected are in bold (p<0.05) and in italics (0.05<p<0.1). (O: organic layer, M02: mineral soil 0-20 cm, M24: mineral soil 20-40 cm, M48: mineral soil 40-80 cm, M8: mineral soil > 80 cm/ TL: tension lysimeter, ZTL: zero-tension lysimeter/ n.s.: no significant.)

In broadleaved and coniferous forests:

Collector	Layer	[DOC]		LMM			LMM			SMŀ	K	
type			(with	n breakp	oints)	(without breakpoints)			(without breakpoints)			
			n	rslope	p value	n	rslope	p value	rslope	N	NS	Р
TL	0	47.3	3133	6.75	0.078	1168	-0.30	n.s.	-1.03	1	3	1
									(±1.65)			
	M02	12.9	19311	0.10	n.s.	8917	-1.06	n.s.	0.16	17	29	21
									(±4.78)			
	M24	4.93	7700	2.69	n.s.	3404	3.66	n.s.	0.6	11	12	11
									(±9.03)			
	M48	3.66	24614	0.95	n.s.	11065	0.80	n.s.	0.67	22	30	32
									(±4.76)			
	M8	3.27	9378	6.78	0.0036	3394	3.41	n.s.	1.007	8	9	16
									(±8.79)			
ZTL	0	37.9	8136	3.75	<0.001	4659	1.63	0.0939	1.7	3	16	8
									(± 4.28)			
	M02	30.7	3389	-0.54	n.s.	445	0.17	n.s.	-0.7	0	3	1
									(±1.85)			
	M24	17.3	739	0.36	n.s.					0	0	0

M48	4.73	654	-3.37	n.s.	336	1.05	n.s.	1.07	1	2	1
								(±3.08)			
M8	3.7	118	1.39	n.s.					0	0	0

Collector type	Layer	[DOC]	(wi	LMM th breakpo	oints)	(with	LMM (without breakpoints)			SMK (without breakpoints)				
			n	rslope	p value	n	rslope	p value	rslope	N	NS	Р		
TL	0	41.4	637	-5.96	n.s.	475	-0.17	n.s.	-0.3 (±0.9)	0) 2	0		
	M02	8.80	8397	3.07	0.0764	3104	0.51	n.s.	0.89 (±5.94)	4	7	10		
	M24	3.78	2584	-0.05	n.s.	928	6.01	n.s.	1.03 (±11.31)	3	5	4		
	M48	2.60	10635	-0.93	n.s.	4634	2.46	n.s.	1.51 (±5.31)	11	8	16		
	M8	2.60	4354	-6.85	0.0672	1797	-0.10	n.s.	0.3 (±6.28)	4	5	6		
ZTL	0	33.3	4057	0.37	n.s.	1956	-0.90	n.s.	0.96 (±5.47)	2	7	3		
	M02	4.26	608	0.26	n.s.	192	1.88	n.s.	2.72	0	0	1		
	M24	20.4	94	11.80	0.026					0	0	0		
	M48	3.42	427	-2.84	n.s.				0	0	1	0		
	M8	2.42	34	-36.18	<0.001					0	0	0		

In broadleaved forests:

In coniferous forests:

Collector	Layer	[DOC]		LMM			LMM			SMK		
type			(with	h breakp	breakpoints) (without breakpoints)				(without breakpoints)			
			n	rslope	p value	n	rslope	p value	rslope	N	NS	Р
TL	0	49.0	2496	8.15	0.0633	693	1.33	n.s.	-1.06	1	1	1
									(±2.25)			
	M02	15.7	10914	-0.97	n.s.	5813	-1.60	n.s.	-0.04	13	22	11

									(±3.98)			
	M24	5.72	5116	2.71	n.s.	2476	3.66	n.s.	-0.3	7	7	8
									(±7.82)			
	M48	4.44	13979	1.24	n.s.	6431	0.05	n.s.	0.3	16	22	11
									(±4.32)			
	M8	3.70	5024	9.93	<0.001	1597	7.58	n.s.	2.89	4	4	10
									(±10.28)			
ZTL	0	42.9	4079	3.59	0.0018	2703	3.09	0.0045	1.85	1	9	5
									(± 2.88)			
	M02	36.9	2781	-0.60	n.s.	253	-1.44	n.s.	-0.83	0	3	0
									(±0.4)			
	M24	16.3	645	0.23	n.s.					0	0	0
	M48	44.0	227	-0.39	n.s.	251	-0.55	n.s.	2.14	1	1	1
									(±3.66)			
	M8	4.14	84	13.87	0.0995					0	0	0

Table 2. Site properties for the 13 plots showing consistent negative trends (N) of DOC concentrations and for the 12 plots showing consistent positive trends (P) of DOC concentrations. Soil properties (clay percentage, C/N ratio, pH(CaCl₂), cation exchange capacity (CEC)) are for the soil depth interval 0-20 cm. Mean atmospheric deposition (inorganic N and SO_4^{2-}) is throughfall deposition from 1999 to 2010. When throughfall deposition was not available, bulk deposition is presented with an asterisk. Relative trend slopes (rslope) in soil solution pH, Ca²⁺ and Mg²⁺ concentrations were calculated using the Seasonal Mann-Kendall test.

Code Plot	Trend	Soil Type (WRB)	Clay (%)	C/N	Hd	$CEC (cmol_+ kg^{-1})$	MAP (mm)	MAT (°C)	N depos. (kg N ha ⁻¹ yr ¹)	SO_4^{2-} deposition (kg S ha ⁻¹ yr ⁻¹)	rslope pH (%yr ⁻¹)	rslope $Ca^{2+}(\% \text{ yr}^{-1})$	rslope Mg^{2+} (% yr^{-1})
Franc	ce (co	ode = 1)											
30	N	Cambic Podzol	3.79	16.8	3.96	1.55	567	11.9	7.28	4.25	0.10	-0.90	-1.00
41	N	Mollic Andosol	23.9	16.6	4.23	7.47	842	10.6	4.43	4.15	0.00	-1.10	-1.30
84	N	Cambic Podzol	4.09	22.8	3.39	4.07	774	10.5	7.66	3.77*	0.50	2.00	1.00
Belgi	um (code =2)											
11	Р	Dystric Cambisol	3.54	17.7	2.81	6.22	805	11.0	18.7	13.2	0.40	-11.0	-8.00
21	Р	Dystric Podzo- luvisol	11.2	15.4	3.59	2.41	804	10.3	16.8	13.2	0.00	-9.00	-5.00
Germ	nany ((code:= 4)											
303	N	Haplic Podzol	17.3	16.5	3.05	8.77	1180	9.10	17.5		0.40	-5.00	-2.00
304	N	Dystric	21.3	17.7	3.63	6.14	1110	6.20	16.4		0.00	-3.00	-0.40

Code Plot	Trend	Soil Type (WRB)	Clay (%)	C/N	Hd	CEC (cmol ₊ kg ⁻¹)	MAP (mm)	MAT (°C)	N depos. (kg N ha^{-1} yr ¹)	SO_4^{2-} deposition (kg S ha ⁻¹ yr ⁻¹)	rslope pH (%yr ⁻¹)	rslope Ca^{2+} (% yr^{-1})	rslope Mg^{2+} (% yr^{-1})
308	N	Cambisol Albic Arenosol	3.80	16.5	3.41	1.63	816	9.20	14.2*		0.00	-5.00	-2.00
802	N	Cambic Podzol	6.00	25.7	3.35	4.33	836	11.9	25.2	13.2	0.50	-2.40	-1.50
1502	N	Haplic Arenosol	4.40	23.8	3.78	2.35	593	9.40	9.79	5.66		-16.0	-14.0
306	Р	Haplic Calcisol					782	10.2	13.9		0.50	2.00	2.00
707	Р	Dystric Cambisol					704	10.7	18.3	8.49	0.00	-10.0	-2.00
806	Р	Dystric Cambisol					1349	8.30	23.0	6.81	0.30	-7.00	-6.00
903	Р	Dystric Cambisol					905	9.60			0.20	-5.00	-3.00
920	Р	Dystric Cambisol					908	8.90			-1.00	-6.00	-0.50
1402	Р	Haplic Podzol	8.65	26.2	3.24	9.04	805	6.90	13.5	24.3	1.20	-6.00	9.00
1406	Р	Eutric Gleysol	15.9	23.1	3.59	6.67	670	8.80	15.3	6.23	1.11	-4.00	-3.00
Italy (code	= 5)											
1	N	Humic Acrisol	3.14	12.2	5.32	31.6	670	23.3			-0.30	-10.0	-10.0
United	d Kir	igdom (code	= 6)										
922	Р	Umbric Gleysol	34.8	15.6	3.31	10.8	1355	9.50			0.40	-9.00	2.00

Code Plot	Trend	Soil Type (WRB)	Clay (%)	C/N	рН	$CEC (cmol_+ kg^{-1})$	MAP (mm)	MAT (°C)	N depos. (kg N ha ⁻¹ yr ¹)	SO_4^{2-} deposition (kg S ha ⁻¹ yr ⁻¹)	rslope pH (% yr ⁻¹)	rslope Ca^{2+} (% yr ⁻¹)	rslope Mg^{2+} (% yr^{-1})
Austr	ia (co	ode = 14)											
9	N	Eutric Cambisol	20.1	12.8	5.26	25.9	679	10.8		3.80*	0.40	-1.50	-0.60
Switz	erlan	d (code = 50))										
15	N	Dystric Planosol	17.6	14.7	3.73	7.76	1201	8.90	15.1	4.67	-0.10	-13.0	-4.00
2	Р	Haplic Podzol	14.7	18.3	3.17	3.59	1473	4.40			-0.80	-5.00	-3.00
Norw	ay (c	ode =55)											
14	N	Cambic Arenosol	9.83	25.4	3.46				14.7	21.9	0.10	-1.70	-3.30
19	N		10.5	18.7	3.79		836	4.60	1.54	2.61	0.50	-7.00	-4.00
18	Р		3.05	29.5	3.69		1175	0.35		2.40	-0.90	0.00	0.00

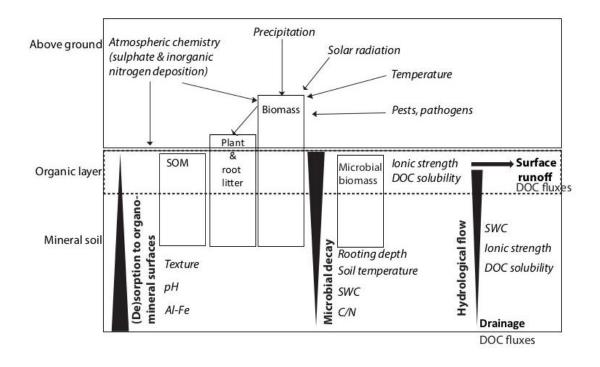
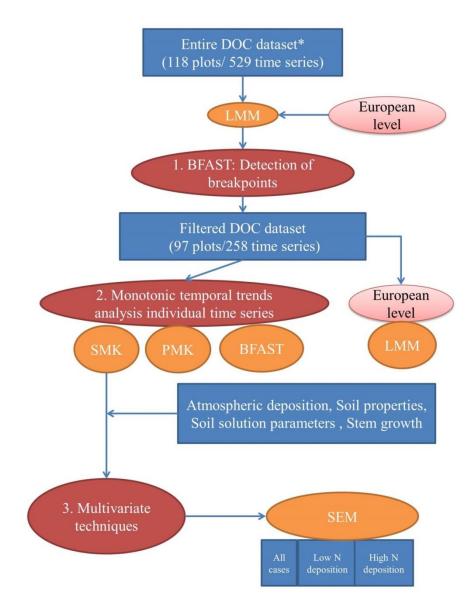


Figure 1. Schematic diagram illustrating the main sources (in boxes) of dissolved organic carbon (DOC) and the main processes (in bold) and factors (in italics) controlling DOC concentrations in soils.



* Time series > 10 years and > 60 obs.

Acronym	Model	Type of analysis
LMM	Linear mixed-effects models	Temporal trends
BFAST	Breaks For Additive Seasonal and Trend	Analysis of breakpoints in time series
SMK	Seasonal Mann Kendall test	Monotonic temporal trends
PMK	Partial Mann Kendall test	Monotonic temporal trends
SEM	Structural equation Model	Multivariate analysis (direct/indirect effects)

Figure 2. Flow-diagram of the sequence of methods applied for analysis of temporal trends of soil solution DOC and their drivers.

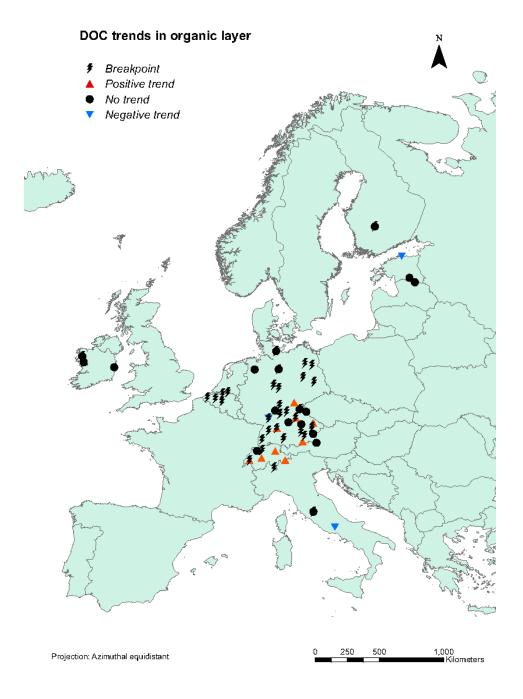


Figure 3. Directions of the temporal trends in soil solution DOC concentration in the organic layer at plot level. Trends were evaluated using the Seasonal Mann-Kendall test. Data span from 1991 to 2011.

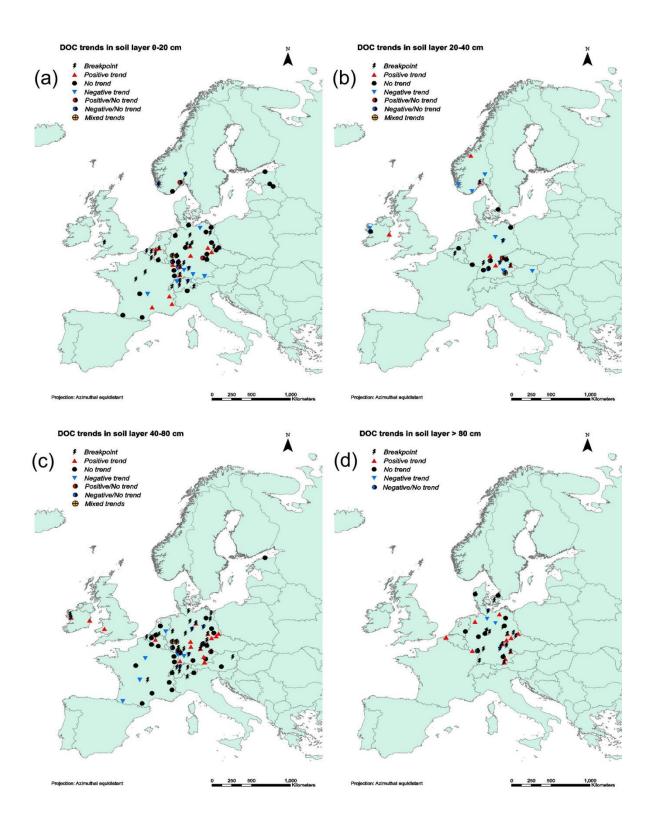


Figure 4. Directions of temporal trends in soil solution DOC concentration at plot level in the mineral soil for soil layers: (a) topsoil (0-20 cm), (b) intermediate (20-40 cm), (c) subsoil (40-80 cm) and (d) deep subsoil (> 80 cm). Trends were evaluated using the Seasonal Mann-Kendall test. Data span from 1991 to 2011.

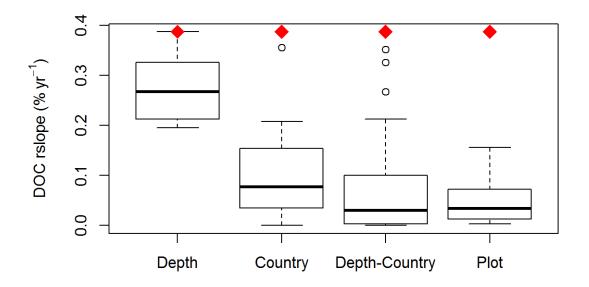


Figure 5. Range of relative trend slopes (max-min) for trends of DOC concentration in soil solution within each 1) depth interval, 2) country, 3) depth interval per country, and 4) plot. The boxplots show the median, 25% and 75% quantiles (box), minimum and 1.5 times the interquartile range (whiskers) and higher values (circles). The red diamond marks the maximum range of slopes in soil solution DOC trends in the entire dataset.

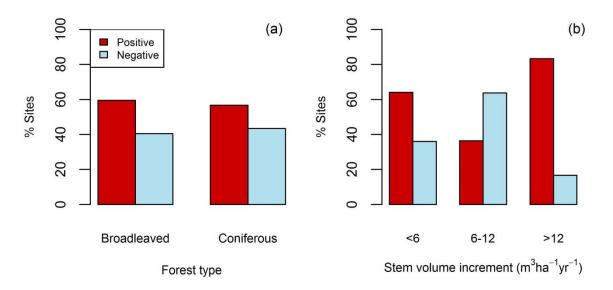


Figure 6. Percentage of occurrence of positive and negative trends of DOC concentration in soil solution separated by A) forest type and B) stem volume increment ($m^3 ha^{-1} yr^{-1}$).

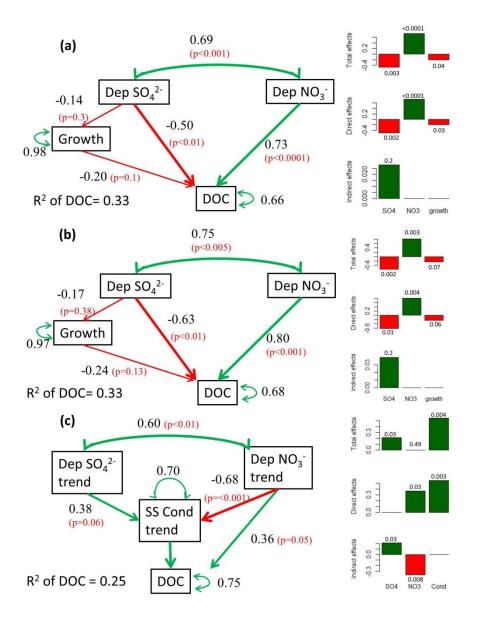


Figure 7. Diagrams of the structural equation models (SEM) that best explain the maximum variance of the resulting trends of DOC concentrations in soil solution for: A) all the cases , B) cases with low or medium throughfall inorganic N deposition (< 15 kg N ha⁻¹ yr⁻¹), and C) cases with high throughfall inorganic N deposition (>15 kg N ha⁻¹ yr⁻¹) with mean or trends in annual SO_4^{2-} and NO_3^{-} deposition (% yr⁻¹) with direct and indirect effects through effects on soil solution parameters (trends of conductivity in μ S cm⁻¹) and mean annual stem volume increment (growth) in m³ ha⁻¹ yr⁻¹). p-values of the significance of the corresponding effect are between brackets. Green arrows indicate positive effects and red arrows indicate negative effects. Side bar graphs indicate the magnitude of the total, direct and indirect effects and their p-values.

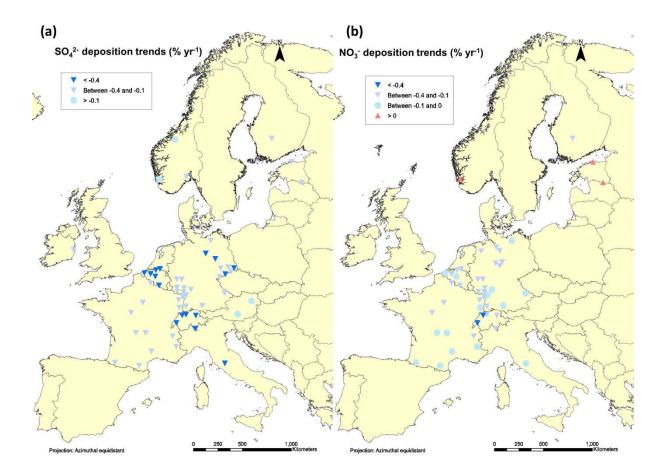


Figure 8. Temporal trends in a) throughfall SO_4^{2-} deposition and b) throughfall NO_3^{-} deposition at plot level. Trends were evaluated using the Seasonal Mann-Kendall test. Data span from 1999 to 2010.