Use of cosmic ray neutron sensors for soil moisture monitoring in forests

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8 Abstract

Measuring soil moisture with cosmic ray neutrons is a promising technique for intermediate 9 spatial scales. To convert neutron counts to average volumetric soil water content a simple 10 calibration function can be used (the N_0 -calibration of Desilets et al., 2010). The calibration is 11 based on soil water content derived directly from soil samples taken within the footprint of the 12 13 sensor. We installed a cosmic-ray neutron sensor (CRS) in a mixed forest in the lowlands of north-eastern Germany and calibrated it 10 times throughout one calendar year. Each calibration 14 with the N_0 -calibration function resulted a different CRS soil moisture time series, with 15 deviations of up to 24 % of the total range for individual values of soil water content. Also, many 16 17 of the calibration efforts resulted in time series that could not be matched with independent in situ measurements of soil water content. We therefore suggest a modified calibration function with a 18 different shape that can vary from one location to another. A two-point calibration proved to be 19 adequate to correctly define the shape of the modified calibration function if the calibration 20 21 points were taken during both dry and wet conditions spanning at least half of the total range of soil moisture. The best results were obtained when the soil samples used for calibration were 22 23 linearly weighted as a function of depth in the soil profile and non-linearly weighted as a function of distance from the CRS, and when the depth-specific amount of soil organic matter and lattice 24 25 water content was explicitly considered. The annual cycle of tree foliation was found to be a negligible factor for calibration because the variable hydrogen mass in the leaves was small 26

compared to the hydrogen mass changes by soil moisture variations. As a final point, we providea best practice calibration guide for CRS in forested environments.

29

30 1. Introduction

31 Determining average soil moisture content over larger areas is difficult, mainly for two reasons. Firstly, soil moisture can be highly variable even at small spatial scales, especially under 32 intermediate wetness conditions (e.g. Western et al., 2004). Secondly, most common in situ 33 measurement techniques only yield point measurements. To obtain a valid estimate of area-34 35 average soil moisture one needs to collect data from numerous locations within a given area. This can be time-consuming and expensive. More recently, remote sensing of soil moisture at larger 36 scales has become a research focus (e.g. see Ochsner et al., 2013 for a recent review); however, 37 the measurement depth of many of these methods is still limited to the upper 5 cm of the soil. 38 39 Also, both spatial and temporal resolution is rather coarse. A technique that intends to bridge the scale gap between point measurements of soil moisture and remote sensing is the use of cosmic 40 ray neutrons as indicators of soil moisture. A detailed description of the cosmic ray neutron 41 sensors (CRS) can be found in Zreda et al. (2008, 2012), here we will only describe the basic 42 measurement principle. Cosmic ray neutrons on Earth are formed when high-energy protons 43 44 deriving from galactic sources (such as supernovae) enter the Earth's atmosphere. Once in the atmosphere, the protons interact with atomic nuclei (mainly nitrogen and oxygen) producing 45 cascades of secondary neutrons (also called high-energy neutrons) that travel towards the Earth's 46 surface and into the soils. When secondary neutrons interact with air or soil they trigger the 47 release (evaporation) of fast (but low energy) neutrons. The number of fast neutrons above the 48 soil surface depends strongly on the number of hydrogen atoms in the surroundings because 49 50 hydrogen atoms have a very high capacity to moderate fast cosmic ray neutrons (that means to slow them down and turn them into thermal neutrons with even less energy - effectively 51 52 removing the fast neutrons from the system). The number of hydrogen atoms increases with increasing soil water content and hence soils with high water contents re-emit fewer fast neutrons 53 54 than soils with low water content. That leads to fewer fast neutrons being detected above-ground by the CRS which is generally installed 1-2 m above the soil surface. 55

As early as 1966 Hendrick and Edge reported that the intensity of fast (low-energy) neutrons 56 (~1 keV) detected above the ground depended on the hydrogen content of the soil, and Kodama 57 58 (1985) found an inverse correlation of neutron intensity and soil moisture content with a neutron sensor buried in the soil. In 2008, Zreda et al. introduced a method to measure average soil water 59 content over a larger area (~30 ha) with CRS. The footprint of CRS, i.e. the area around the 60 sensor where 86 % of detected neutrons originate from, covers a circle with an approximate 61 62 radius of 300 m (Desilets and Zreda, 2013). However, the radius can decrease with increasing air density and humidity, with increasing vegetation density and with increasing soil moisture to 63 64 about 100 m (Köhli et al., 2015). The effective measurement depth of CRS, i.e. the soil depth where 86 % of detected neutrons originate from, varies between 10 and 70 cm below surface 65 66 (Zreda et al., 2008), depending on soil type, water content and distance from the sensor (Köhli et al., 2015). To account for the contributions of neutrons from different soil depths, various depth-67 68 weighting approaches have been proposed, some of them assuming a linear decrease of weights 69 with depth (Franz et al., 2012a), others assuming a non-linear decrease with depth (Köhli et al., 70 2015).

The original measurement method uses a relationship between neutron flux and volumetric soil 71 72 water content with the shape of the relationship being known from neutron transport simulations. For this relationship, Desilets et al. (2010) presented an equation with three constant shape 73 74 parameters (a_0, a_1, a_2) and one calibration parameter (N_0) which has to be calibrated with soil 75 moisture values determined by the gravimetric method from field soil samples. The influence of soil lattice water and soil organic matter on the signal was investigated by Zreda et al. (2012). 76 77 They found that both lattice water and soil organic matter contain fixed amounts of hydrogen that 78 further attenuate the neutron signal and need to be taken into account. Lattice water and soil organic matter corrections to the original relationship by Desilets et al. (2010) are provided for 79 80 example in Lv et al. (2014).

Other external factors influencing the neutron count that need to be corrected for are (a) atmospheric pressure (Bachelet et al., 1965), (b) incoming neutron flux (see e.g. Zreda et al., 2012, Bogena et al., 2013) and (c) specific humidity (Rosolem et al., 2013). More recently, the effects of biomass on the neutron signal have been discussed. Bogena et al. (2013) noted that aboveground biomass reduced the neutron count rate and thus decreased the sensitivity of the

sensor. To counter this loss of sensitivity they recommended a 24 h integration time for their 86 forested catchment as a compromise between decreased uncertainty and decreased time 87 88 resolution. Hawdon et al. (2013) and Baatz et al. (2015) compared neutron counts for locations with different amounts of biomass. Hawdon et al. (2013) reported that the variation in biomass 89 90 could explain 80 % of the variation in neutron counts when assuming a nonlinear relationship between biomass and neutron counts, Baatz et al. (2015) explained 87 % of the variation 91 92 proposing a linear relationship between the two variables. Baroni and Oswald (2015) suggested that the influence of above-ground biomass between the sensor and the ground which decreases 93 94 the effective measurement depth of the CRS can be incorporated into the weighting approach of 95 Franz et al. (2012a). This is especially important in locations where frequent large biomass 96 changes occur, for example in agricultural fields. Coopersmith et al. (2014) found that soil moisture in a corn crop is often overestimated when the leaf area index (LAI) is relatively high 97 98 while it is underestimated when LAI is relatively low - circumstances which could cause 99 differences in the calibration and resulting soil moisture measurements. The influence of the litter 100 layer in forested environments was investigated by Bogena et al. (2013). Water content in the litter layer changes rapidly and adds additional temporal variability to the CRS time series 101 102 complicating the extraction of the soil moisture signal. Therefore, Bogena et al. (2013) recommended considering the water dynamics in the litter layer explicitly in the calibration 103 104 approach. Franz et al. (2013) introduced a new approach (the universal calibration function) that 105 takes into account all sources of hydrogen thereby requiring estimates of lattice water, soil 106 organic carbon, and vegetation biomass as well as a regression factor that can be derived from 107 calibration or may directly be retrieved from neutron count measurements over a large water body (500 m on all sides and deeper than 1 m). 108

109 Since the launch of the cosmic ray neutron method many changes and corrections have been 110 brought forward that altered the way the method is applied. These changes and corrections can be 111 divided into two groups. On the one hand, there are corrections that are applied to the raw neutron count in order to remove the influence of other variables (such as air pressure and 112 113 humidity variations or fluctuations in incoming neutron counts). On the other hand, changes have 114 been made to the way we average the soil moisture measurements during the calibration 115 campaigns in order to get a representative soil moisture value that corresponds to what the sensor actually "sees" at the time of calibration (changing effective measurement depth, changing 116

footprint diameter, inclusion of lattice water and soil organic matter water equivalent). All this 117 has led to improvements in the method's accuracy for many environments. Most of these studies 118 119 were performed in medium to high-count environments with neutron count rates above 1000 counts per hour, in generally dry environments, at higher elevations and with little vegetation. 120 Only a few studies were performed in low-count environments with count rates below 1000 121 counts per hour (e.g. Rivera Villareyes et al., 2011; Bogena et al., 2013). In the present study, we 122 123 evaluated whether the CRS also provides reliable and consistent soil moisture measurements in a low-count environment, i.e., in a temperate mixed forest close to sea level. We tested several 124 125 weighting approaches to convert gravimetrically determined soil water content of the top 30 cm into an average soil water content that can be used for the calibration of the CRS. Additionally, 126 127 we analyzed whether the annual forest cycle of foliation and defoliation is important to consider for instrument calibration. We furthermore compiled a best-practice for the calibration of CRS in 128 129 forested, low-count environments which is provided in Appendix A.

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131 **2. Field site and instrumentation**

The CRS (CRS-1000 by Hydroinnova) was installed in late 2013 in the Müritz National Park in 132 north-eastern Germany (53°19'49.0"N, 13°11'56.5"E) at an elevation of about 84 m a.m.s.l. (Fig. 133 134 1, inset). Precipitation, temperature and relative humidity data was provided by the climate station Serrahn (1.6 km to the north). Average annual air temperature at the site is 8° C with a 135 maximum in July (17.2°C) and a minimum in January (-0.9°C). Average annual precipitation is 136 137 580 mm with a maximum in June (65 mm) and a minimum in February (28 mm). This makes for 138 a maritime temperate climate (Cfb) in the Köppen climate classification. The sensor is located in a sandy outwash plain, a relic from the last glaciation, which causes the soil texture to be 139 140 homogeneous with sand fractions of about 95% throughout the entire profile. Data from a nearby well shows that the groundwater level at the site is almost 20 m below the terrain surface. The 141 142 vegetation within the sensor footprint consists of both deciduous and coniferous trees. 143 Immediately surrounding the sensor is a mature beech forest (Fagus sylvatica L., older than 100 144 years), also within the footprint (but farther away) with a distance of at least 40 m from the sensor there is young pine (*Pinus sylvestris* L.), oak (*Quercus robur* L.) and spruce (*Picea abies* (L.) 145 146 H.Karst.) forest (all younger than 50 years) as well as a small strip of open grassland (see Fig. 1

and also Fig. 3 for a map of the forest stands and Table 1 for fractions of the different tree stands
within the footprint). Depending on the tree species, the mineral soil is covered by an organic soil
layer and a litter layer of variable depth and water holding capacity.

150 For validation of the CRS soil water content measurements, in May of 2014 we installed 18 soil 151 moisture sensors (TOMST) close to the soil sampling/calibration locations. They are based on the principle of time domain transmission (TDT) and each sensor comes with its own logger and 152 power supply (more information under: http://www.tomst.cz/tms/TMS-3.html). These sensors 153 154 were installed vertically from the terrain surface into the soil so that they continuously measure 155 soil water content averaged over the top 16 cm of the soil. In order to calibrate the sensors we used the gravimetric soil moisture data we collected from the upper 15 cm during the last five 156 157 calibration campaigns which were carried out within the measurement period of the sensors (June-November 2014). The volumetric water content within the upper 15 cm of the CRS 158 footprint was calculated as the mean of all 18 TDT sensors. 159

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161 **3. Methods**

162 **3.1. Calibration**

163 We conducted a total of 10 calibration campaigns throughout one calendar year (2014). The first one (WI) took place in February during winterly conditions with very wet soils. The next four 164 calibrations (S1-4) followed in spring (April-May) and covered the entire period of tree foliation. 165 The sixth calibration (SU) was done under very dry conditions in July and the last four 166 167 calibrations (F1-4) in fall (October-November) covering the trees' defoliation. For all the calibration campaigns we followed the recommended sampling pattern for the calibration of CRS 168 169 which was developed by Zreda et al. (2012) and slightly modified and detailed in Franz et al. (2012b). The sampling pattern prescribes 3 concentric circles around the CRS with radii of 25, 75 170 171 and 200 m, respectively (Fig. 1). The 3 circles are intersected by 6 straight lines that point from the sensor towards north (0°) , north-east (60°) , south-east (120°) , south (180°) , south-west (240°) 172 173 and north-west (300°). Samples are taken in the vicinity of all intersections – the samples do not 174 have to be taken at the exact spot of the intersection. This sampling pattern ensures that each

175 sample has equal weight towards the spatial mean of soil moisture that is detected by the CRS, assuming that the sensitivity of the CRS decreases exponentially with distance. We used a split-176 177 tube sampler to extract 30 cm soil cores at 18 locations within the footprint of the sensor afterwards dividing each soil core into six 5 cm thick soil samples. For each of the 10 calibrations 178 this left us with 108 soil samples which were then transferred in sealed plastic bags to the 179 laboratory where they were immediately weighed, then oven-dried at 105°C for 24 h and then 180 181 weighed again to determine their volumetric water content and bulk density. Afterwards, lattice water, soil organic matter content and root biomass were determined for six depth-representative 182 183 soil samples. To this end the 108 samples (taken from the last calibration campaign in November) were grouped by sampling depth. We extracted 2 g from each of the 18 samples per sampling 184 185 depth and combined them to create one bulk sample per depth. Then, the already oven-dried samples were weighed and put in the oven for another 24 h at a temperature of 400°C. The 186 187 procedure is called 'loss on ignition' since the organic matter is burned off during the process (Ball, 1964; Davies, 1974). This removed most of the soil organic matter and root biomass from 188 189 the samples. After weighing the samples (to compute the fraction of combined soil organic matter and root biomass) they were again placed in the oven for 24 h, this time at a temperature of about 190 191 1000°C. After that, the lattice water was also removed from the samples. A final weighing vielded the fraction of lattice water per soil depth. In order to make soil organic matter and root 192 193 biomass comparable to the influence of pure water we converted them into equivalents of water by multiplying their weight by 0.556 which is the ratio of five times the molecular weight of 194 water to the molecular weight of cellulose (taking into account that cellulose ($C_6H_{10}O_5$) contains 195 196 10 hydrogen atoms per molecule while water (H₂O) only contains two) (Hawdon et al., 2014).

197 The neutron counts from the sensor were smoothed with a 12 h moving window to reduce 198 measurement noise (see Bogena et al., 2013). The next step was to correct the neutron counts for 199 variations in (a) pressure, (b) incoming neutron flux and (c) water vapor in the air. This was done 200 by applying the following corrections:

201 a. Pressure correction:

202
$$N_p = N_{raw} * e^{\left(\frac{x - x_0}{L}\right)}$$
(1).

with $N_{\rm p}$ being the pressure corrected neutron counts (counts h⁻¹), $N_{\rm raw}$ the raw neutron counts 203 (counts h^{-1}), x the atmospheric shielding depth (g cm⁻²) for every time step (derived from 204 atmospheric pressure measured directly inside the CRS case), x_0 the average atmospheric 205 shielding depth (g cm⁻²) for the entire measurement period and L the effective nucleon 206 attenuation length for high-energy neutrons (for our site we assumed a value of 135.9 g cm^{-2} 207 which is equivalent to 133.3 hPa) (Desilets and Zreda, 2003). To convert atmospheric pressure 208 (hPa) into shielding depth (g cm⁻²) the atmospheric pressure has to be multiplied by 1.0194 s² m⁻² 209 1 210

b. Incoming flux correction (Zreda et al., 2012):

212
$$N_{pi} = N_p * \frac{N_{avg}}{N_{nm}}$$
 (2),

with N_{pi} being the sensor neutron count rate corrected for changes in atmospheric pressure and incoming neutrons (counts h⁻¹), N_{avg} the average count rate of incoming neutrons (counts h⁻¹) over the entire measurement period and N_{nm} the neutron count rate of the neutron monitor for each time step (counts h⁻¹).

As the time series of the closest neutron monitor, located in Kiel, Germany, contains several data gaps, we selected the continuous time series of the Jungfraujoch, Switzerland, for this study. We scaled this time series by adjusting its mean (309 counts h^{-1}) to the mean of the Kiel time series (327 counts h^{-1}). The resulting time series resembles the Kiel time series very closely (Fig. S1).

c. Water vapor correction (Rosolem et al., 2013):

222
$$N_{pih} = N_{pi} * \left[1 + 0.0054 * \left(p_{\nu 0} - p_{\nu 0}^{ref}\right)\right]$$
 (3),

with N_{pih} being the sensor neutron count corrected for changes in pressure, incoming neutrons and water vapor (counts h⁻¹), p_{v0}^{ref} the average absolute humidity of the air over the entire measurement period (g m⁻³) and p_{v0} the absolute humidity for each time step (g m⁻³). The constant 0.0054 has units of m³ g⁻¹.

Finally, to convert corrected neutron counts (N_{pih}) into volumetric soil moisture (θ), Desilets et al. (2010) introduced an equation with four parameters – three of which ($a_0 = 0.0808$, $a_1 = 0.372$, a_2 229 = 0.115) were determined via neutron transport simulations and a fourth one (N_0) that serves as a calibration parameter accounting for site and sensor specific variations and representing neutron counts over dry soil at reference conditions during calibration:

232
$$\theta(t) = \left\{ \left[a_0 * \left(\frac{N_{pih}(t)}{N_0} - a_1 \right)^{-1} - a_2 \right] * \rho_{bd} \right\} - W_L - (SOM + B_R)$$
(4).

233 The other parameters ρ_{bd} , W_L , SOM and B_R can be measured directly from the calibration soil samples: the bulk density of the soil (ρ_{bd} in g cm⁻³), the summed volume fraction of lattice water 234 in the soil grains and tightly bound water ($W_{\rm L}$ in m³ m⁻³), the combined volume fraction of soil 235 organic matter and root biomass water equivalent ($SOM+B_R$ in m³ m⁻³). In order to calibrate the 236 237 sensor one first has to determine the depth- (and distance-) weighted averages for ρ_{bd} , W_{L} , $SOM+B_R$ and θ as well as N_{pih} (averaged over 12 h) for the time of calibration. This is necessary 238 239 because several factors can influence the effective measurement depth z^* (which is the depth of the soil layer up to which 86 % of the neutrons that the CRS detects originate from) and the 240 241 footprint size of the sensor (Fig. 2). Afterwards N_0 is adjusted iteratively (e.g. with a simple 242 Solver routine in Microsoft Excel) until the right-hand side of the equation equals the left-hand side. 243

We tested four soil moisture weighting approaches (Table 2), described in detail below, to determine which information is necessary for an accurate calibration.

1. In the first approach (simple depth-weighting, SDW) a linear depth-weighting function was used (Franz et al., 2012b), where wt(z) represents the weight that is applied to the soil moisture measurements from a certain soil depth *z*:

249
$$\begin{cases} wt(z) = a \left[1 - \left(\frac{z}{z^*} \right)^b \right] & 0 \le z \le z^* \\ wt(z) = 0 & z > z^* \end{cases}$$
(5),

250 where

251
$$a = \frac{1}{z^{*} - \frac{z^{*b+1}}{(b+1)z^{*b}}}$$
 (6),

252 and

253
$$z^* = \frac{5.8}{H_p + 0.0829}$$
 (7),

254 and

$$255 H_p = W_L + SOM + B_R + \theta (8).$$

In these equations z is the soil depth below the surface in cm and z^* is the effective measurement 256 depth in cm, a is a parameter that ensures that the weights are conserved, b controls the curvature 257 of the weighting function and equals 1 for linear weighting, $H_{\rm p}$ is the water equivalent of the 258 belowground hydrogen pools (m³ m⁻³), W_L is lattice water (m³ m⁻³), SOM is soil organic matter 259 water equivalent (m³ m⁻³), $B_{\rm R}$ is root biomass water equivalent (m³ m⁻³) and θ is the 260 gravimetrically determined volumetric soil pore water content ($m^3 m^{-3}$). The original approach by 261 262 Franz et al. (2012b) was modified by Bogena et al. (2013) using the total hydrogen content of belowground hydrogen pools H_p instead of just using the volumetric soil water content θ . Since 263 264 $H_{\rm p}$ changes with soil depth we used an iterative approach to determine the appropriate weights. Starting with an average value for the upper 30 cm of the soil we computed an effective 265 measurement depth z^* and weighted H_p of the different soil depths accordingly. With this new 266 value of H_p we then recomputed z^* and the weights. Usually the value of H_p stabilizes after a few 267 iterations. The bulk density (ρ_{bd}) of the soil changes with depth and influences the soil moisture 268 269 measurements too. Therefore it was also being taken into account during the iterative process of determining the effective measurement depth z^* and the weighted soil moisture. In this first 270 weighting approach we did not use our depth-specific measurements of W_L and $SOM+B_R$, instead 271 we assumed an average weight fraction value of combined $W_L+SOM+B_R$ for the entire 30 cm 272 profile. 273

2. The second approach (depth-specific weighting, DSW) was identical to the first one (SDW) 2. Except for using depth-specific measurements of W_L and $SOM+B_R$ (see Table 3 for an example).

3. For the third approach (distance-depth-weighting, DDW), we adopted the weighting approach
described in Köhli et al. (2015). This approach introduces distance-dependent variable depth-

weighting where the effective measurement depth decreases with distance from the sensor. The effective measurement depth z^* is calculated according to:

280
$$z^* = \rho_{bd}^{-1} \left[8.32 + 0.14 * \left(0.97 + e^{\frac{-r}{100}} \right) * \frac{26.42 + H_p}{0.057 + H_p} \right]$$
 (9),

where ρ_{bd} is the bulk density of the soil (g cm⁻³), *r* is the radial distance (in meters) from the CRS and H_p is the water equivalent of the belowground hydrogen pools (m³ m⁻³) (see Eq. 8). This approach also assumes that the footprint size of the sensor varies with soil water content and atmospheric water content. We computed the varying footprint diameter for each calibration campaign and weighted the samples from 25, 75 and 200 m accordingly.

4. The fourth approach (distance-depth-weighting, non-linear, DDWnl) was identical to the third
one (DDW) except for using the non-linear depth-weighting function recommend by Köhli et al.
(2015) instead of the linear one (from Eq. 5):

289
$$wt(z) = e^{\frac{-2z}{z^*}}$$
 (10).

3.2. Estimation of biomass and influence of seasonal changes in biomass

Biomass influences neutron counts due to its hydrogen content. In order to test (and potentially exclude) the influence of seasonal changes in aboveground forest biomass, we estimated living tree biomass and tree biomass changes throughout the year by applying the aboveground dry biomass functions for beech forest (*Fagus sylvatica* L.) from Santa Regina et al. (1997):

295
$$B_S = 0.0894 * DBH^{2.4679}$$
 (11),

296
$$B_B = 0.0317 * DBH^{2.3931}$$
 (12),

297
$$B_L = 0.0145 * DBH^{1.9531}$$
 (13).

298 $B_{\rm S}$ is dry stem biomass (kg tree⁻¹), $B_{\rm B}$ dry branch biomass (kg tree⁻¹), $B_{\rm L}$ dry leaf biomass (kg tree⁻¹) 299 ¹) and *DBH* is the diameter of the tree stem at breast height (cm). Total dry above-ground 300 biomass $B_{\rm ag}$ is the sum of the three components. 301 To apply these functions we conducted a survey of tree diameters and tree density in the beech forest that surrounds the CRS. This allowed us to determine both the total biomass of the beech 302 303 forest, as well as the seasonally variable fraction of biomass (leaf biomass divided by total biomass). We first calculated the water mass (W_{agb}) in stems, branches and leaves (assuming a 304 leaf water content of 0.6 kg per kg of wet biomass (Gravano et al., 1999) and a wood water 305 content of 0.11 kg kg⁻¹ (Bouriaud et al., 2004)). Finally, using the mass fraction of hydrogen in 306 water ($M_w = 0.1119$ kg H per kg H₂O) and in dry biomass ($M_b = 0.0622$ kg H per kg Cellulose: 307 $C_6H_{10}O_5$) the total hydrogen mass (H_{agb}) of above-ground biomass in the beech stand was 308 309 derived:

310
$$H_{agb} = W_{agb} * M_w + B_{ag} * M_b$$
 (14).

We did not conduct surveys on the other tree species. Table 1 shows that the beech stand covers 56% of the footprint area around the CRS (when assuming the exponential distance-weighting from Zreda et al. (2008)). Pine covers 16%, spruce 13%, oak 8%. With the new distance weighting function of Köhli et al. (2015), the cover fractions of the other tree species would decrease even further. Also, the seasonal variation in spruce and pine above-ground biomass is very small and thus we consider it to be constant in this study.

317 **3.3. Validation**

As an objective performance measure to compare the soil moisture time series derived from the CRS with the soil moisture time series from the TDT sensors we used the modified Kling-Gupta efficiency KGE' (Gupta et al., 2009; Kling et al., 2012):

321
$$KGE' = 1 - \sqrt{(r-1)^2 + (\beta - 1)^2 + (\gamma - 1)^2}$$
 (15).

322 With correlation coefficient *r*:

323
$$r = \frac{\sum_{i=1}^{n} (x_i - \bar{x})(y_i - \bar{y})}{\sqrt{\sum_{i=1}^{n} (x_i - \bar{x})^2} \sqrt{\sum_{i=1}^{n} (y_i - \bar{y})^2}}$$
(16),

bias ratio $\beta = \mu_{\text{mod}}/\mu_{\text{obs}}$ and variability ratio $\gamma = (\sigma_{\text{mod}}/\mu_{\text{mod}})/(\sigma_{\text{obs}}/\mu_{\text{obs}})$. The KGE' measures the Euclidian distance in a 3-D space where the correlation coefficient *r* is on one axis, the variability ratio β is on the second axis and the bias ratio γ is on the third axis. KGE' scores range from 1 (representing a perfect fit) to - ∞ . Due to the composite nature of the KGE' it is relatively simple to analyze which feature of the time series (correlation, bias, variability) contributes most to the good/bad performance of a model.

330

331 **4. Results**

4.1. Gravimetric soil water measurements and soil physical characteristics

Soil water content in the sandy soils ranged between 0.03 and 0.37 m³ m⁻³ (absolute minimum 333 and maximum values of individual soil core samples during the 10 sampling campaigns). The 334 spatial distribution of volumetric soil water content for the 10 calibration days is shown in Fig. 3. 335 336 At each location the soil water content is an unweighted average value of the six samples taken from 0 to 30 cm depth. The mean volumetric soil water content for the calibration days over all 337 calibration locations ranged from 0.07 up to 0.16 m³ m⁻³ with standard deviations ranging from 338 0.015 to 0.047 m³ m⁻³. The depth and distance weighted averages used for calibration ranged 339 from 0.08 to 0.24 m³ m⁻³ (see for example Table 4, column: θ_{depthW}). A general soil moisture 340 pattern emerged with the soil moisture under coniferous tree stands being lower and under 341 deciduous tree stands being higher. Especially the uppermost soil layer (0-5 cm) was drier under 342 the coniferous trees – on average about $0.065 \text{ m}^3 \text{ m}^{-3}$ – while the deeper soil layers under 343 coniferous trees were about 0.023 m³ m⁻³ drier. The highest spatial variabilities in soil moisture 344 were encountered during spring and fall seasons and more homogenous soil moisture conditions 345 during winter and summer. The wettest calibration we conducted (WI) yielded an average soil 346 water content of $0.29 \text{ m}^3 \text{ m}^{-3}$ for the top 5 cm. Calibration at higher soil water content is difficult 347 as it only occurs for short periods of time after large precipitation events when significant 348 349 amounts of intercepted water are also present in the canopy and litter layer.

The average bulk density (ρ_{bd}) measurements for the 10 calibration campaigns ranged from 1.16 to 1.22 g cm⁻³ (mean: 1.18 g cm⁻³, standard deviation: 0.02 g cm⁻³). The weight fraction of soil organic matter and root biomass water equivalent (w*SOM*+*B*_R) was determined to be 51.4 g kg⁻¹ in the shallowest soil layer (0-5 cm) with decreasing values at depth. The weight fraction of lattice water (w W_L) was determined to be 3.2 g kg⁻¹ in the shallowest soil layer with slightly increasing values at deeper soil depths.

356 **4.2. Footprint variability**

357 The footprint diameters calculated according to Köhli et al. (2015) and used in approaches 3 and 4 ranged from 185 m for the wettest to 200 m for the driest conditions. This resulted in distance 358 359 weights of ~0.56 (for samples from 25 m distance), ~0.35 (for samples from 75 m distance) and ~ 0.10 (for samples from 200 m distance). These weighting factors varied only marginally 360 between the individual calibration campaigns despite considerable differences in soil and 361 atmospheric water content. Sampling distances with equal weights according to Köhli et al. 362 (2015) would have differed from our sampling pattern (~1 m, ~33 m, ~140 m instead of 25 m, 75 363 m, 200 m), a condition which we balance by adjusting the distance weights. Furthermore the 364 conditions within 30 m around our CRS are quite homogenous since the sensor is located within 365 a pure beech stand and we are expecting little difference in average soil moisture content between 366 locations at 1 and 25 m distance. 367

368 **4.3. Calibration**

The average reference atmospheric pressure (P_0) for the entire measurement period was 369 1005.8 hPa; the average reference incoming neutron flux (N_{avg}) was 328.3 counts h⁻¹; the average 370 reference absolute humidity (p_{v0}^{ref}) was 9.1 g m⁻³. Equations (5) through (10) were used to 371 calculate depth-weighted volumetric soil water content θ_{depthW} and depth-weighted water 372 equivalent of belowground hydrogen pools $(H_p)_{depthW}$ according to the four weighting approaches 373 we applied. Equations (1)-(3) were used to compute N_p , N_{pi} and N_{pih} , and then Eq. (4) to identify 374 N_0 for each calibration. Table 3 provides an example of the depth-weighting following approach 375 2 (DSW with depth-specific values of W_L and $SOM+B_R$). 376

The values in Table 3 result in a depth-weighted average volumetric water content θ_{depthW} of 0.150 m³ m⁻³, a depth-weighted water equivalent of belowground hydrogen pools $(H_p)_{depthW}$ of 0.179 m³ m⁻³ and a depth-weighted bulk density $(\rho_{bd})_{depthW}$ of 0.981 g cm⁻³. If W_L and $SOM+B_R$ were not considered, the values for θ_{depthW} and $(\rho_{bd})_{depthW}$ would change to 0.146 m³ m⁻³ and 1.013 g cm⁻³ respectively, because the effective measurement depth z^* increases when the higher amounts of $SOM+B_R$ in the shallow layers are not considered, thus giving more weight to low soil moisture values in deeper soil horizons.

Table 4 lists the parameters relevant for calibration for all 10 calibration dates (again following approach 2, DSW, with depth-specific values of W_L and $SOM+B_R$).

Following the standard N_0 -calibration approach of Desilets et al. (2010), we should have ended up with the same N_0 value for each of the 10 calibrations. However, the N_0 range we found was considerable – e.g. from 808 to 895 counts h⁻¹ for the DDW approach (mean: 841.9 counts h⁻¹, standard deviation: 13.7 counts h⁻¹). As a consequence, the 10 computed time series also showed differences in volumetric soil water content (Fig. 4 illustrates results for the DDW approach). In the most extreme case, these differences were larger than 0.1 m³ m⁻³ (which is equal to 24 % of the total range of soil water content at the site).

In fact, none of the four weighting approaches was able to solve the problem of determining a unique calibration parameter for our field site. All weighting approaches resulted in largely deviating N_0 -values between the individual calibrations (see means and standard deviations in column 1 and 2 of Table 5). This in turn led to differences in the resulting time series of volumetric soil water content (see means and standard deviations in column 3 and 4 of Table 5).

4.4. Modified calibration function

To include all information of our 10 calibration campaigns into our analysis, we fitted modified 399 calibration functions to four sets of 10 calibration points derived from the four different 400 weighting approaches (see section 3.1). This was done by using the Microsoft Excel Solver 401 402 software to optimize the three shape parameters (a_0, a_1, a_2) and N_0 through the calibration point cloud (solid lines in Fig. 5). Plotting the N_{pih} -values of all 10 calibrations against the 403 404 gravimetrically determined and depth- (and distance-) weighted volumetric soil moisture revealed 405 that the standard shape of the soil moisture-neutron count relation is not valid at our field site. 406 Instead of plotting along functions defined by the standard calibration (Desilets et al., 2010) (examples are dotted lines in Fig. 5) our calibration points are better captured by less steep 407 408 functions (solid lines in Fig. 5 are the best-fit calibration functions for the different approaches). 409 Using the N_0 -calibration function with the standard shape parameters may lead to large soil water 410 content deviations between individual calibration campaigns, especially under wet soil moisture 411 conditions. The slope of the N_0 -calibration function is essentially too steep, which means that in 412 our environment a change in the neutron count is caused by a more subtle change in soil moisture 413 than is assumed by the standard relationship – essentially the sensor has a higher 414 resolution/sensitivity than one would expect.

The optimized parameters for the four approaches are shown in Table 6. The resulting soilmoisture time series are shown in Fig. 6.

417 **4.5. Validation**

418 We tested whether the modified calibration functions improved the performance of the CRS measurements relative to in situ measurements, and if so, which of the weighting approaches 419 420 performed best. In order to do that we compared the soil moisture time series from the CRS 421 (using the standard N_0 -calibration function from Desilets et al. (2010) and applying our newly derived corrected relationships) with the soil moisture time series from the TDT sensors 422 distributed throughout the footprint. As a first step, the CRS measurements had to be converted to 423 a soil water content value representative of the top 15 cm of the soil (the integration depth of the 424 TDT sensors). For this purpose we compared the weighted volumetric water content (θ_{depthW}) 425 426 from the gravimetric measurements of the calibration campaigns (basically what the CRS is 427 supposed to "see") with the unweighted average gravimetric measurements of the top 15 cm (θ_{15cm}) (Fig. S2). We found strong linear correlations for two of the weighting approaches (SDW 428 and DSW) with CRS water content being larger than the θ_{15cm} values and increasing differences 429 for wetter soil conditions (indicating that for higher soil moisture the CRS overestimates soil 430 431 water contents in the top 15 cm while for lower soil moisture the overestimation decreases). For approaches 3 and 4 (DDW and DDWnl) offsets of 0.006 and 0.011 m³ m⁻³ indicated slightly 432 lower weighted soil water content than the unweighted top 15 cm values. The linear correlations 433 for the first two weighting approaches were expected since when it is wetter, the effective 434 435 measurement depth is reduced for the CRS measurements and the wetter shallower soil layers 436 receive more weight. Therefore, the CRS measurements result in higher soil water content than the gravimetric measurements. However, it seems that in approaches 3 and 4 the distance 437 438 weighting counters this effect. A probable explanation is that the formula used for the distance-

depth weighting increases the effective measurement depth. This causes higher weights for 439 deeper (drier) soil layers even under wet conditions and could counteract the trend. We then 440 441 converted the CRS time series by the above relationships into time series that were representative of the top 15 cm and compared them to the TDT measurements. The modified Kling-Gupta 442 efficiency (KGE') was used as a performance measure. The worst performance was achieved by 443 the simple depth weighting approach (KGE'(SDW) = 0.83, Table 7), the performance improved 444 445 when depth-specific weighting was included (KGE'(DSW) = 0.88) and it further improved when including distance weighting (KGE'(DDW) = 0.89). The linear depth weighting worked better 446 447 than the non-linear depth weighting (KGE'(DDWnl) = 0.83). That means that the distance-depthweighting approach (DDW) improved the neutron sensors performance the most. In comparison, 448 449 using the single-point standard N_0 -calibration function and DDW yielded KGE's for the individual calibration campaigns ranging from 0.58 to 0.83 with a mean KGE' of 0.71 (\pm 0.08). It 450 is important to note that all of the modified calibration approaches performed better than their 451 standard calibration counterparts. The improvement of performance of the new N_0 -calibration 452 453 functions compared to the standard calibration functions was caused by the better agreement of both the bias ratios β and the variability ratios γ , i.e. both the means and the variabilities of the 454 CRS time series better matched the TDT observations (see also Fig. 7). This supports the 455 456 hypothesis that at our field site larger than expected changes in neutron count are already caused by subtle changes in soil moisture. 457

458 **4.6. Optimizing calibration efforts**

We further tested whether two or more individual calibration campaigns are required to determine a comprehensive calibration function shape, and under which soil moisture conditions these calibrations should be conducted. We paired each individual calibration point (derived from the best-performing weighting approach, DDW) with all the other calibration points (WI and S1, WI and S2, WI and S3, etc.) and computed best-fit calibration functions for all of these pairings (Fig. 8).

Then we used the resulting calibration functions to convert the measured neutron counts into time series of volumetric soil water content and compared these to the TDT measurements (again using the KGE' as the performance measure). We found that a two-point calibration proved to be

sufficient in case that the difference in soil water content between the two calibrations was larger 468 than 0.1 $\text{m}^3 \text{m}^{-3}$ (i.e. for our sandy soils it covered ~50 % of the observed range of average soil 469 water content). Figure 9 indicates that the calibrated neutron count-soil water content conversion 470 will always perform well if the soil moisture difference between the two calibrations is 471 sufficiently large. Also, it turned out to be more important to capture a calibration point at very 472 dry rather than at very wet soil water contents. This is illustrated in Fig. 9 where predominantly 473 calibrations that involve low soil water contents (red dots) as the minimum value achieve KGE's 474 of 0.9 while these KGE' values are also achieved more frequently with intermediate soil water 475 contents (light blue dots) as the maximum value. 476

477 **4.7. Variability of hydrogen pools**

The tree survey revealed a median diameter of 23.9 cm (Min: 3.2 cm, Q₂₅: 11.5 cm, Q₇₅: 43.7 cm, 478 Max: 93.3 cm) and a tree density of 0.05 stems m^{-2} . With these values at hand and Eqs. (11)-(13) 479 the dry above-ground biomass of the beech stand (B_{ag}) was 63.8 kg m⁻² (with 62.8 kg m⁻² from 480 stem and branches and 1.0 kg m⁻² from leaves) (Fig. 10). These values result in 9.2 kg m⁻² of 481 biomass water (W_{agb}) (with 7.8 kg m⁻² from stem and branches and 1.5 kg m⁻² from leaves). 482 Further calculations yield a hydrogen mass of 4.8 kg m⁻² for stem and branches and a hydrogen 483 mass of 0.22 kg m^{-2} for leaves (Eq.14). Other hydrogen pools within the CRS footprint were also 484 assessed. The thickness of the litter layer was determined to be 5 cm on average. Assuming a 485 porosity of 85 % yields a hydrogen mass of 0.47 kg m⁻² for a dry litter layer. Hence, the hydrogen 486 mass of the static biomass (stem, branches and dry litter) amounted to 5.24 kg m⁻². Beech litter 487 was found to have a maximum interception capacity of 2.8 mm in a forest in Luxembourg 488 (Gerrits et al., 2010) corresponding to an additional 0.31 kg m^{-2} of hydrogen when the litter layer 489 is wet. The canopy interception of beech can be assumed to be up to 1.5 mm (Gerrits et al., 2010) 490 (i.e. another 0.17 kg m^{-2} of hydrogen is added to the system when the canopy is wet). The 491 hydrogen contribution of soil organic matter and root biomass changes with soil water content 492 because the effective measurement depth of the sensor changes. Applying the DDW approach we 493 computed a value of 0.36 kg m⁻² for wet conditions (0.29 m³ m⁻³), a value of 0.44 kg m⁻² for 494 intermediate conditions (0.17 $\text{m}^3 \text{m}^{-3}$) and a value of 0.66 kg m^{-2} for dry conditions (0.05 $\text{m}^3 \text{m}^{-3}$). 495 The hydrogen contribution of lattice water also changes with moisture conditions (wet: 0.05 kg 496 m⁻²; intermediate: 0.07 kg m⁻²; dry: 0.15 kg m⁻²). A pore water content of 0.29 m³ m⁻³ equals a 497

498 hydrogen mass of 4.12 kg m⁻², a pore water content of 0.17 m³ m⁻³ equals a hydrogen mass of 499 3.26 kg m^{-2} and a pore water content of 0.05 m³ m⁻³ reduces the hydrogen mass to 1.77 kg m⁻². 500 Figure 11 and Table 8 give an overview of the different hydrogen pools for varying moisture 501 conditions within the footprint of the CRS.

502

503 5. Discussion

504 **5.1. Potential influences on neutron counts**

The 10 N_0 -calibration parameters derived from our 10 calibrations varied considerably. In a first 505 analysis we found that this was not related to the different soil moisture conditions during 506 calibration. In search of other potentially unaccounted factors that influence the neutron count we 507 compared N_0 -values obtained from the 10 calibrations with apparent atmospheric pressure, 508 specific humidity, temperature and estimates of forest crown cover (derived from photographs 509 510 taken from the ground aiming at the zenith) during the calibration campaigns. No seasonal or other temporal relationships were found. The contributions of different hydrogen pools (Fig 11) 511 512 reveal that a large percentage of hydrogen at our field site stems from the above-ground vegetation (52 to 68 %, depending on moisture conditions). Fortunately, most of this hydrogen is 513 514 static in nature and can be accounted for by the calibration of the CRS. Assuming that the hydrogen content of the stem and branches is constant and only the leaves change seasonally one 515 516 is left with a fraction of variable hydrogen in the above-ground biomass that accounts for 2-3 % of the total hydrogen mass. The variability in hydrogen due to foliation and defoliation in the 517 beech forest surrounding the CRS amounts to 0.22 kg m⁻². This means that it equals a change in 518 soil water content of about 0.031 m³ m⁻³ (under wet conditions) and 0.018 m³ m⁻³ (under dry 519 520 conditions). These differences for wet and dry conditions are due to the fact that the effective measurement depth z^* of the CRS increases for dry conditions: the sensor receives the neutron 521 signal from deeper soil depths and therefore an equal increase in soil water content requires a 522 larger amount of water since a larger soil column has to be filled. At high soil moisture, a 0.01 m³ 523 m^{-3} soil moisture change from 0.28 to 0.29 $m^3 m^{-3}$ equals a change of 0.07 kg m^{-2} of hydrogen in 524 the soil. At low soil moisture the change from 0.05 to $0.06 \text{ m}^3 \text{ m}^{-3}$ is equal to a change in 525 hydrogen of 0.12 kg m⁻². The above calculations with respect to biomass variability disregard the 526

527 fact that fallen leaves still contain hydrogen (which hence is not completely removed from the system immediately and therefore should also reduce the expected variability). At our field site 528 65 % of the distance-weighted area surrounding the CRS is covered by deciduous trees (mainly 529 beech and oak), the other 35 % do not experience a significant annual cycle of leaf growth and 530 fall (pine, spruce and grassland). This should further reduce the influence of seasonally variable 531 biomass on the cosmic ray neutron counts (with a potential maximum influence of leaf-out during 532 wet conditions of 0.020 m³ m⁻³ and only 0.012 m³ m⁻³ in dry conditions). In summary, we do not 533 expect a significant impact of seasonally varying above-ground biomass on the measurements of 534 535 soil water content. Also, we could not find systematic changes in the calibration results connected to the annual cycle of tree foliation/defoliation (i.e. a reduction in counts during summer due to 536 537 higher hydrogen content in the above-ground biomass). Therefore we deem a correction for variable hydrogen from forest canopy biomass at different times of the year unnecessary. 538

539 With regard to other varying hydrogen pools we noticed that the influence of interception storage 540 both in the canopy and in the litter layer can potentially have an impact. When both the canopy and the litter layer are wet, the combined hydrogen amount within these two stores can sum up to 541 almost 5 % of the total hydrogen pool equaling a change in volumetric soil water content of 0.067 542 $m^3 m^{-3}$ (Fig. 11). It is not possible to solve this problem by calibrating during conditions of high 543 interception storage since then the soil water content would be underestimated as soon as the 544 545 canopy is dry. Calibration during conditions of dry canopy and litter layer is recommendable 546 because conditions with an empty interception store are generally prevalent and can be much better defined than conditions with a filled interception store. A potential solution to the influence 547 548 of the variable interception storage filling is the introduction of another neutron count correction 549 using observed, derived or modeled interception storage values (similar to the pressure or the 550 water vapor correction).

The fact that the depth-specific weighting (DSW) approach performed better than the simple depth weighting (SDW) is an indication that the depth variations in lattice water, soil organic matter and root biomass content should be explicitly accounted for during the calibration of the CRS. The best performance was achieved with a weighting approach (DDW) that explicitly takes into account both depth-weighting as well as distance weighting of the soil water content (Table 7). This suggests that the variation in the footprint diameter needs to be considered during

individual calibration campaigns. Linear depth-weighting resulted in a better CRS performance 557 than non-linear depth-weighting since the non-linear depth-weighting basically underestimated 558 559 soil water contents during wet periods (because higher weights of deeper (drier) soil layers were included). This caused both a decrease in the mean soil water content as well as a decrease in the 560 561 variability of the soil water content time series and hence reduced the performance of the CRS. In soils where water content increases with depth the difference between linear and non-linear 562 563 depth-weighting could be smaller (even negligible), at our field site, however, the decrease of water content with depth apparently favors the use of a linear depth-weighting function. 564

565 The differences in calibration results are likely caused by the fact that the shape of the N_0 calibration function is different at our field site. That means that while being temporally stable 566 the shape of the calibration function is spatially variable – there is no standard curve applicable to 567 all sites. At our site the function is less steep than the standard N_0 -calibration function suggested 568 by Desilets et al. (2010), i.e. a similar increase in neutron counts is associated with a smaller 569 decrease in soil moisture. A recalibration of the shape of the curve using all calibration points 570 considerably improved the agreement between in situ measurements and CRS measurements of 571 soil moisture. A two-point calibration already proved to be sufficient to define the correct shape 572 of the calibration function given that the soil moisture states at the two calibration times were 573 sufficiently different. In a recent study Iwema et al. (2015) also investigated temporal field 574 575 sampling strategies for three different calibration methods. They tested combinations of different numbers of random sampling dates and found that using more than six random sampling dates 576 did not improve their calibration results much more. However, for the N_0 -calibration method they 577 found that selecting sampling dates with distinct soil wetness conditions could reduce the 578 579 required number of samplings. In conclusion they also recommended more than one calibration campaign for the N_0 -calibration approach and argued that the shape of the calibration function 580 581 should not be fixed but kept variable during the calibration process. This is in line with our 582 findings on the shape of the calibration function.

We can only speculate about the reasons behind this shape inconsistency of the calibration function for our site since we did not do any theoretical neutron modeling. To our knowledge we are dealing with the lowest number of counts of all published studies (average $N_0 = 878$ counts h⁻¹, Table 4). Although the calibration function was theoretically developed for all environments it

587 has not yet been tested sufficiently in such low-count, forested environments. Moreover, due to the low neutron count the uncertainty in the determination of soil water content during calibration 588 589 has a much higher influence on the calibration results than in high-count environments. Bogena et al. (2013) pointed out another complicating factor that is present in forested environments – the 590 litter layer. They showed that at their sites (N_0 : 913 to 1397 counts h⁻¹) the model-derived water 591 content within the litter layer (under spruce) was subject to much higher variability than the water 592 593 content in the underlying soil. During wet conditions the water within the litter layer contained 36 % of the hydrogen mass within the footprint of the CRS while during dry conditions it contained 594 595 only 10 % of the hydrogen mass. This leads to an increase in the variability of the neutron counts and can thus cause an overestimation of soil water content during wet conditions. Although the 596 597 water within the litter layer at our site accounts for a much smaller fraction of the total hydrogen pool (up to 3 %) it can still have an influence on the neutron counts and the calibration results. 598 599 The occurrence of canopy interception would have the same variability-increasing effect on the CRS signal, although it is expected to be significantly smaller than the influence of the litter 600 layer. Baatz et al. (2014) working also in a low-count environment (N_0 : 936 to 1242 counts h⁻¹) 601 with land use ranging from grassland to agriculture to forest compared the standard N_0 -calibration 602 method to another calibration method developed by Shuttleworth et al. (2013) (the COSMIC 603 operator) and found that the former interpreted dry periods drier and wet periods wetter – which 604 605 is in accordance to our findings that suggest that the standard N_0 -calibration function is too steep. Ly et al. (2014), in a study at a mixed-forest/grassland site also recommended more than one 606 calibration. They operated in a high-count environment in Utah ($N_0 = 2189$ counts h⁻¹) and 607 608 attributed the different shape of their calibration function to binary soil moisture patterns at their site where the grassland soils were much drier than the forest soils under wet conditions but just 609 610 as dry under dry conditions. Our field site is subject to similar spatial variability since it is also comprised of multiple areas with non-uniform soil water content (mean values of soil water 611 contents differ between different forest stands). Following the argumentation of Lv et al. (2014), 612 the fact that distance weighting improved our results can be regarded as an indication that non-613 614 homogeneous soil moisture conditions indeed lead to changes in the shape of the calibration function. At our site, distance weighting reduced the spatial variability within the footprint of the 615 sensor since it assigned higher weights to the closest sampling sites which were all located in the 616 homogenous and relatively wet beech forest, while the influence of the drier soils under the 617 618 coniferous trees was reduced.

619 If it was possible to fully correct for all factors that influence footprint size, depth-weighting and neutron count, a one-time calibration of the CRS would be sufficient. However, the abundance of 620 621 different hydrogen pools and the uncertainties in the sensing depth estimation will always lead to uncertainties in the calibration process. Therefore we argue that for the use of the CRS as a 622 simple tool to measure soil water content at intermediate scales, the efforts of measuring all 623 necessary parameters are not justified. As shown by Iwema et al. (2015) and by the results of this 624 625 study, this issue can be dealt with by using site-specific calibration parameters estimated from in situ samples taken during dry and wet conditions. Hence, we recommend a two-point calibration 626 627 that - although being empirical in nature - inherently incorporates many of the required corrections. 628

629

630 6. Conclusion

Our results suggest that a one-time calibration of the CRS using the available neutron count 631 corrections and weighting approaches is not sufficient at our field site. This is mainly due to the 632 633 fact that the shape of the standard N_0 -calibration function is not able to reproduce the dynamics in soil water content we observed with our network of distributed in situ TDT sensors. Several 634 factors could cause this discrepancy, amongst them the presence of a litter layer and spatially 635 636 heterogeneous soil moisture conditions within the sensor footprint. After calibrating the CRS 10 times in a mixed forest in north eastern Germany we found that a two-point calibration already 637 considerably improved the agreement between soil water content derived from in situ TDT 638 639 measurements and from the CRS, given significantly different moisture conditions during the two 640 calibration periods/campaigns (for a detailed explanation on the procedure see Appendix A). We 641 found that the explicit consideration of depth-specific values of soil organic matter and root 642 biomass improved the calibration results while seasonal changes in above-ground biomass in the forest were found to be negligible. While there is no doubt that further investigations of factors 643 644 that influence the neutron signal are necessary and useful, it is also apparent that it becomes 645 increasingly difficult to distinguish between the effects of the individual correction factors and 646 the uncertainty caused by all the corrections. Therefore our goal was to use empirical data to test available methods and combinations thereof and to provide a guideline on how to easily and 647 648 comprehensively calibrate a CRS in various environments using these methods. Looking beyond that objective, site intercomparison studies along gradients from high to low-count environments and/or from locations with varying litter layers could give rise to the development of simple corrections to the shape of the N_0 -calibration function.

When measuring soil water content with a CRS it is important to note that over time the measurements are hardly ever representative of the exact same soil segment around and below the sensor (Köhli et al, 2015). With the footprint shrinking and expanding and the effective measurement depth in the soil decreasing and increasing we have to be careful when interpreting and using our results. If we keep that in mind, however, this new technology will indeed be able to bridge the gap between point in-situ and areal remote sensing soil moisture measurements and thus provide a valuable tool for the advancement of hydrologic understanding.

659

660 Appendix A: Best practice for calibration in low-count forest environments

We provide an Excel file as a supplement to perform the calculations described in the followingstep-by-step instructions.

- 663 1. Set up (or use) a weather station that monitors air temperature and relative humidity close to664 the CRS.
- Set up the CRS in a location where the conditions within a radius of at least 30 m around
 the sensor are relatively homogeneous (similar soils, tree species, expected soil moisture
 conditions).

Switch on the CRS and come back later for calibration (or set it up before 6 a.m. and start
calibrating on the same day). You should at least have 12 hours of CRS data for one
calibration. Do not switch it off after the calibration, let it record continuously.

4. Choose a day with very dry or very wet soil moisture conditions for the first calibration
campaign and wait for the opposite conditions for your second calibration (this might take a
full year to achieve, but you will not lose any data, you will just not be able to accurately
convert the data immediately).

5. Choose days without rain or snow for your calibrations, litter and canopy should be dry.

- 676 6. Take 108 soil samples from 18 locations (six directions, three distances) and six depths (0677 30 cm). For equal distance weights choose distances according to Köhli et al. (2015) (~1,
 678 ~33 and ~140 m).
- 679 7. Weigh the samples the same day you take them, let them oven-dry for 24 h at 105°C and 680 weigh them again to determine the volumetric water content (θ) and the bulk density (ρ_{bd}).
- 681 8. Create six bulk samples from the six different soil depths (2 g from each of the 18 locations
 682 suffices for each soil depth).
- 683 9. Determine the combined soil organic matter (SOM) and root biomass (B_R) content of the 684 six bulk samples by weighing them (after regular oven-drying at 105°C) and then heating 685 them to a temperature of 400°C for 24 h before weighing them again. Convert SOM and B_R 686 to water equivalents by multiplying the weight by 0.556.
- 687 Caution: In clay-rich soils this method tends to overestimate soil organic matter content
 688 because some of the lattice water is removed already at temperatures around 400°C
 689 (Howard and Howard, 1990).
- 690 10. Determine the lattice water (W_L) content of the six bulk samples by weighing them (after 691 SOM and B_R extraction at 400°C) and then heating them to a temperature of 1000°C for 24 692 h before weighing them again.
- 693 Caution: Carbonate-rich soils experience thermal breakdown of carbonates at temperatures
 694 above 430°C (Ben-Dor and Banin, 1989).
- Determine the water equivalent of the average hydrogen content of belowground hydrogen
 pools (H_p) for each soil depth.

697 Equation (8).

- Apply a linear weighting function to your gravimetrically determined H_p measurements 698 12. accounting for the change in the effective measurement depth z* of the sensor and retrieve a 699 weighted average of H_p within the footprint of the CRS by iteration. Start out by computing 700 the effective measurement depth z* corresponding to your gravimetrically determined 701 values of H_p and ρ_{bd} averaged over the entire 30 cm. Then apply the weights for the 702 703 different soil depths z and update the values. Recalculate the effective measurement depth and continue this procedure until all values stabilize. Do this for each 704 z* sampling/calibration distance (~1, ~33 and ~140 m) separately. 705
- 706 Equations (5), (6) and (9).

- 70713. Apply an additional distance-weight to the depth-weighted volumetric water contents from708the different locations in order to account for variations in the footprint size. Also do this709iteratively adjusting H_p and the distance weights until both become stable.
- Figure 2015 Figure 201
- T12 14. Use the depth-and-distance weights to compute weighted values of soil water content (θ),
 bulk density (ρ_{bd}), lattice water (W_L), soil organic matter and root biomass water equivalent
 T14 (SOM+B_R).
- 715 15. Average raw neutron counts (N_{raw}) from the moderated sensor (measuring fast neutrons)
 716 over 12 h with a moving window.
- 717 16. Retrieve data from the neutron monitor close to your location in order to correct for the
 718 varying intensity of incoming neutrons (you may have to correct this data and fill gaps).
- 719 17. Using the entire time series for the period where cosmic-ray data is available determine 720 average atmospheric pressure (P₀), average incoming neutron intensity (N_{avg}) and average 721 absolute humidity (p_{v0}^{ref}).
- 722 18. Correct raw neutron counts for atmospheric pressure variations (N_p).
- 723 Equation (1).
- 19. Correct raw neutron counts for incoming neutron intensity variations (N_{pi}) .

726 20. Correct raw neutron counts for absolute humidity variations (N_{pih}).

- 72821. Fit a function through the two calibration points altering N0, a0, a1 and a2 (e.g. using729Microsoft Excel solver). When doing this, use average values of the two calibration730campaigns for bulk density (ρ_{bd}), lattice water (W_L), soil organic matter and root biomass731water equivalent (SOM+BR).
- 732 22. Plot the N_{pih} of both calibrations against the gravimetrically measured, distance- and depth-733 weighted volumetric soil water content (θ).
- 734 23. Use best fit parameters to convert time series of N_{pih} to volumetric soil water content.
- 735

736 Acknowledgement

⁷²⁵ Equation (2).

⁷²⁷ Equation (3).

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	Radius 0-50 m	Radius 50-150 m	Radius 150-300 m	Total _
Beech	85.2	32.8	48.7	<mark>55.5</mark> 🔇
Pine	3.0	26.3	17.6	15.6
Spruce	5.8	20.9	11.1	12.6
Oak	0.0	10.3	12.5	7.6
Open (grass)	6.0	9.7	3.9	6.5
Larch	0.0	0.0	5.5	1.8
Birch	0.0	0.0	0.7	0.2

Table 1. Fractions of different tree stands in percent within the footprint of the CRS.

884	Table 2. Overview of the four weighting approaches for other than soil moisture effects on the
885	CRS signal.

Approach	1 SDW	2 DSW	3 DDW	4 DDWnl
consideration of depth-specific W_L and SOM+B _R	no	yes	yes	yes
distance depth-weighting	no	no	yes	yes
non-linear depth-weighting	no	no	no	yes

Table 3. Example of depth weighting (DSW) for an effective measurement depth of $z^* = 22.1$ cm,

888 a = 0.0903 and b = 1. Calibration campaign date 21 November 2014 (F4). Note the difference in

specific weights if only soil water content θ is considered (wt(z, θ)) or if W_L and $SOM + B_R$ is also

890 considered (wt(z, H_p)).

		3 3	$SOM + B_R$	33	
Layer (cm)	θ (m ³ m ⁻³)	$W_{\rm L} ({\rm m}^3{\rm m}^{-3})$	$(m^3 m^{-3})$	$H_{p} (m^{3} m^{-3})$	ρ_{bd} (g cm ⁻³)
0-5	0.187	0.002	0.034	0.223	0.669
5-10	0.136	0.004	0.024	0.163	1.143
10-15	0.117	0.004	0.019	0.140	1.217
15-20	0.109	0.004	0.015	0.129	1.256
20-25	0.106	0.005	0.013	0.124	1.359
25-30	0.100	0.005	0.012	0.118	1.431

891

z (cm)	$wt(z, \theta)$	$\int^{z+5} wt(z,\theta)$	$wt(z, H_p)$	$\int^{z+5} wt(z, H_p)$
	· · · ·	$J_{\mathbf{Z}}$	· F	J _Z
0	0.079	0.356	0.090	0.401
5	0.063	0.278	0.070	0.299
10	0.048	0.200	0.050	0.197
15	0.032	0.122	0.029	0.095
20	0.017	0.044	0.009	0.009
25	0.001	0.000	0.000	0.000
		Σ=1.00		Σ=1.00

893 Table 4. Atmospheric and soil parameters as well as neutron counts for the 10 calibrations. Atmospheric pressure P, absolute humidity p_{v0} , raw neutron count N_{raw} , pressure corrected 894 895 neutron count $N_{\rm p}$, pressure and incoming radiation corrected neutron count $N_{\rm pi}$, pressure, incoming radiation and water vapor corrected neutron count N_{pih} , calibration neutron count N_0 , 896 incoming radiation from the neutron monitor $N_{\rm nm}$, average soil moisture of the top 30 cm $\theta_{\rm 30 cm}$, 897 depth-weighted soil moisture θ_{depthW} , depth-weighted sum of volumetric lattice water content, soil 898 organic matter and root biomass water equivalent $(W_L+SOM+B_R)_{depthW}$, depth-weighted water 899 equivalent of belowground hydrogen pools $(H_p)_{depthW}$, depth-weighted bulk density $(\rho_{bd})_{depthW}$ and 900 average volumetric soil water content θ_{mod} of the resulting time series using the N₀-calibration 901 function with standard parameters. Mean (μ) and standard deviation (σ) values of the 10 902 903 calibration campaigns are given in the two bottom lines.

Calibration	P (hPa)	p _{v0} (g m ⁻³)	N _{raw} (counts h ⁻ ¹)	(counts h^{1})	$(\begin{array}{c} N_{pi} \\ (counts h^{-1}) \end{array}$	$({counts h}^{1})$	$(\text{counts } h^{-1})$
Winter	984.0	5.7	606.2	514.9	518.8	509.4	872.4
Spring1	999.3	8.6	549.2	523.0	527.5	526.2	868.7
Spring2	1021.0	4.9	491.1	550.6	542.8	530.5	871.1
Spring3	1002.9	9.6	544.7	533.1	539.9	541.5	869.2
Spring4	1019.0	8.0	503.4	556.0	549.4	546.1	879.0
Summer	1008.7	14.0	613.3	626.6	623.8	640.5	858.2
Fall1	998.7	11.5	624.7	592.4	593.8	601.5	909.5
Fall2	1014.1	7.8	509.3	542.1	546.7	542.8	876.2
Fall3	990.3	8.5	630.4	561.4	580.4	578.5	892.8
Fall4	1016.7	6.6	544.4	591.0	577.7	569.9	885.7
μ	1005.5	8.5	561.7	559.1	560.1	558.7	878.3
σ	11.9	2.6	50.2	33.1	31.1	37.5	13.8

Calibration	N _{nm} (count s h ⁻¹)	$\begin{array}{c} \theta_{30cm} \\ (m^3 m^{-3}) \end{array}$	θ_{depthW} (m ³ m ⁻ ³)	$(W_L+SOM \\ +B_R)_{depthW} \\ (m^3 m^{-3})$	$(\mathbf{H}_{\mathbf{p}})_{\mathrm{depthW}}$ $(\mathbf{m}^{3} \mathbf{m}^{-3})$	$(ho_{bd})_{depthW}$ (g cm ⁻³)	θ_{mod} (m ³ m ⁻³)
Winter	325.8	0.163	0.228	0.0343	0.262	0.985	0.141
Spring1	325.5	0.153	0.200	0.0340	0.234	1.013	0.143
Spring2	333.0	0.150	0.185	0.0311	0.216	0.955	0.137
Spring3	324.1	0.140	0.175	0.0324	0.207	1.000	0.143
Spring4	332.2	0.139	0.170	0.0302	0.200	0.957	0.145
Summer	329.8	0.073	0.080	0.0278	0.108	1.074	0.151
							26
σ	5.0	0.024	0.038	0.0019	0.039	0.034	0.013
-------	-------	-------	-------	--------	-------	-------	-------
μ	327.7	0.131	0.165	0.0312	0.196	0.997	0.151
Fall4	335.8	0.126	0.150	0.0293	0.179	0.981	0.155
Fall3	317.5	0.119	0.149	0.0316	0.181	1.018	0.166
Fall2	325.5	0.140	0.174	0.0310	0.205	0.970	0.144
Fall1	327.4	0.112	0.137	0.0299	0.167	1.016	0.182

Table 5. Means (μ) and standard deviations (σ) of calibration parameter N_0 and means (μ) and standard deviations (σ) of resulting time series of volumetric soil water content θ_{mod} for the four weighting approaches with 10 calibration campaigns each.

Approach	$(N_0)_{\mu}$ (counts h ⁻¹)	$(N_0)_{\sigma}$ (counts h ⁻¹)	$(\theta_{mod})_{\mu} (m^3 m^{-3})$	$(\theta_{mod})_{\sigma} (m^3 m^{-3})$
1 SDW	855.0	17.3	0.158	0.015
2 DSW	878.3	13.8	0.151	0.013
3 DDW	841.9	13.7	0.139	0.012
4 DDWnl	828.1	13.3	0.134	0.012

	N ₀	\mathbf{a}_0	a_1	\mathbf{a}_2
1 SDW	926.3	0.203	0.109	0.238
2 DSW	1007.8	0.203	0.114	0.267
3 DDW	810.7	0.326	0.001	0.310
4 DDWnl	779.3	0.314	0.001	0.285

910 Table 6. Modified calibration parameters for the four weighting approaches.

912	Table 7. Performance measures for the four weighting approaches – comparison of modified
913	calibration (mdf) with standard calibration (stan). KGE' is the modified Kling-Gupta efficiency,
914	β is the bias ratio and γ is the variability ratio. (KGE')_{\mu} and (KGE')_{\sigma} represent the mean and
915	standard deviation of the KGE' values of the 10 individual single-point standard calibrations.

	KGE'			(KGE'	(KGE'		
	mdf	β mdf	γ mdf	stan) _µ	stan) _σ	$(\beta \ stan)_{\mu}$	$(\gamma \ stan)_{\mu}$
1 SDW	0.830	0.849	0.986	0.675	0.045	1.120	1.258
2 DSW	0.880	0.915	0.964	0.727	0.035	1.032	1.231
3 DDW	0.891	1.076	0.986	0.712	0.081	0.878	1.237
4 DDWnl	0.833	1.148	1.011	0.681	0.096	0.818	1.244

- Table 8. Hydrogen pools (in kg hydrogen per m^2) in the CRS footprint for different moisture
- 918 conditions (wet: $0.29 \text{ m}^3 \text{ m}^{-3}$, full canopy and litter storage; intermediate: $0.17 \text{ m}^3 \text{ m}^{-3}$, dry canopy
- and moist litter storage; dry: $0.05 \text{ m}^3 \text{ m}^{-3}$). Above-ground biomass is split into a static part (AGB
- 920 wet static) comprising stem, branches and dry litter and a variable part (AGB wet variable) that
- 921 represents leaves.

Hydrogen Pool	Wet (kg m ⁻²)	Intermediate (kg m ⁻²)	Dry (kg m ⁻²)
AGB wet static	5.24	5.24	5.24
AGB wet variable	0.22	0.22	0.22
SOM+R _B	0.36	0.44	0.66
Lattice water	0.05	0.07	0.15
Pore water	4.12	3.26	1.77
Litter water	0.31	0.11	0.00
Interception	0.17	0.00	0.00
Total	10.47	9.35	8.04



Figure 1. Soil sampling locations for calibration (blue dots) and forest vegetation around the CRS (red dot in the center). The TDT soil moisture sensors are located in close vicinity to the sampling locations. The larger yellow circle approximates the footprint of the CRS as it was assumed when sampling took place (diameter approximately 300 m). The smaller yellow circle approximates the footprint of the CRS according to newer modeling results by Köhli et al. (2015) (diameter approximately 200 m). Inset: Field site location in Müritz National Park in northeastern Germany.



Figure 2. Simplified representation of factors influencing the raw neutron count (N_{raw}) and the 933 measurement support of the CRS in terms of effective measurement depth and footprint. 934 Temporally variable factors are shown on the left: barometric pressure (P), canopy interception 935 (I), air humidity (H) and litter layer interception (L). Temporally constant factors (for our study 936 site) are shown on the right: vegetation above and below the sensor (V), soil organic matter 937 938 (SOM), root biomass (B_R) and lattice water (W_L). All these factors need to be accounted for in order to isolate the soil water content signal (θ). The time-variable factors require permanent 939 940 monitoring and dynamic correction, the influence of the constant factors is taken into account during calibration. The combination of the time-variable and time-constant factors leads to a site 941 942 specific temporally variable effective measurement depth and footprint diameter.



945 Figure 3. Gravimetrically determined volumetric soil water content patterns in the footprint of the 946 CRS for the 10 calibration dates. The colored dots indicate the unweighted average value from 0 947 to 30 cm at the 18 calibration locations. Background colors represent the unweighted average 948 value of all 108 soil samples. Different forest stands (pine, beech, oak, spruce) are indicated by 949 the patterned background.



Figure 4. Upper panel: volumetric water content derived from CRS data for each of the 10 calibration dates separately (vertical lines indicate calibration dates, colors correspond to time series colors). Filled circles represent the weighted volumetric water content at the time of calibration (according to DDW). Lower panel: differences in water content between calibration S1 and all other calibrations expressed as a percentage of the total possible range of average soil water content – ranging from 0.04 m³ m⁻³ to 0.34 m³ m⁻³at our field site (color coding corresponds to calibration dates in the upper panel).



960

Figure 5. Modified calibration functions (solid lines) for the four different weighting approaches 961 (simple depth-weighting SDW, depth-specific weighting DSW, distance-depth-weighting DDW, 962 distance-depth-weighting, non-linear DDWnl), each one derived from 10 calibration points 963 (circles). Calibration points are better captured by flatter calibration functions (solid lines) with 964 965 modified calibration parameters than by any of the standard calibration functions (dotted lines) based on a single calibration data set only (days S2 and F1 as an example). Black lines illustrate 966 967 that differences in soil moisture between the results of individual calibrations are larger when soil moisture is high. The inset magnifies the area around the calibration points. 968



971 Figure 6. Time series of volumetric water content derived with modified calibration functions 972 using parameters based on the four calibration approaches: simple depth-weighting (SDW), 973 depth-specific weighting (DSW), distance-depth-weighting (DDW) and distance-depth-974 weighting, non-linear (DDWnl). Filled circles represent the weighted average of volumetric water 975 content obtained from soil cores at the time of calibration (weighting according to DDW).



978 Figure 7. Average volumetric water content derived from TDT point measurements (black line) 979 and CRS measurements (orange line) using different calibration functions. Upper panel: the orange line is an average of the volumetric water content derived from the 10 calibration 980 campaigns of the CRS using the standard N_0 -calibration function from Desilets et al. (2010) 981 982 applying the DDW approach. Grey dotted lines are results for 10 individual calibration campaigns (KGE' values range from 0.579 to 0.834). Lower panel: the orange line is the 983 volumetric water content derived from the calibration function with modified calibration 984 parameters applying the DDW weighting approach based on all 10 calibration dates. The colored 985 vertical lines mark the days of the last five calibration campaigns. 986



Figure 8. Best-fit N_0 -calibration functions (red-brown colored lines) for all combinations of twopoint calibrations (blue dots). Best-fit N_0 -calibration function for 10-point calibration (black line). Best-fit two-point N_0 -calibration function derived from calibration points with highest and lowest volumetric water content (yellow line).



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Figure 9. Performance of CRS soil water content data derived from two-point calibrations in relation to difference between soil moisture states ($\Delta \theta$) at the two calibration dates. The color bar indicates volumetric soil water content. Left panel: points are colored according to the soil water content of the drier calibration date. Right panel: points are colored according to the soil water content of the wetter calibration date. Dashed lines indicate that soil moisture differences of less than 0.1 m³ m⁻³ can produce *N*₀-calibration curves with sub-optimal conversions of neutron counts to volumetric soil water content.



Figure 10. Mass of hydrogen in individual beech trees in stem and branches (red diamonds) and
leaves (green triangles) in relation to diameter at breast height (DBH). Fraction of leaf hydrogen
mass to total aboveground tree hydrogen mass (orange line).



Figure 11: Varying hydrogen pools in the beech forest surrounding the CRS for three different site conditions. AGB (above-ground biomass) wet variable represents hydrogen contained in deciduous leaves (both in the biomass and in the leaf water). AGB wet static comprises hydrogen contained in biomass and water of tree stems and branches as well as in biomass of the litter layer.





Figure S1. Incoming neutron flux from the neutron monitors in Kiel, Germany and Jungfraujoch, 1017 Switzerland and synthetic continuous time series of incoming neutron flux combined from these

two and used for the corrections in this study. 1018



Figure S2. Comparison of depth-(and distance-) weighted averages of gravimetrically determined soil water content with unweighted gravimetrically determined soil water content of the upper 15 cm of the soil. The first two weighting approaches overestimate soil water content in the upper 15 cm especially at high soil water contents. The last two approaches have only a slight negative offset and no significant relationship with wetness conditions.