# Response to Reviewers 1 and 2 on Manuscript, bg-2020-273

# Estimating maximum fine fraction organic carbon in UK grasslands

Kirsty C. Paterson, Joanna M. Cloy, Robert. M. Rees, Elizabeth M. Baggs, Hugh Martineau, Dario Fornara, Andrew J. Macdonald, and Sarah Buckingham

We thank the reviewers for their comments and evaluation of our manuscript. Please find below our response to comments made by reviewers 1 and 2. Reviewer's comments are in black text, and our responses are in blue text, changes to the manuscript are highlighted in yellow. Line numbers refer to the revised manuscript (marked version) below.

#### Reviewer 1

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The authors further improved the paper considering the suggestion/comments raised, therefore I think that can be accepted for publication. I have only some minor suggestions and an issue not completely taken from the last review.

One of the take-home message of the paper is about the unsuitability of Hassink's equation in UK conditions but, then, the recommended Quantile Regression (90th percentile) gives similar absolute and mean ratio of unsaturated sites (Table 4). Is there any contradiction in the message?

The absolute number of unsaturated samples are similar, 105 and 99 for Hassink's equation in the UK and the Quantile Regression (90<sup>th</sup> percentile) equation, generated using the data from this study. However, as shown in figure 4, the samples deemed oversaturated, above the solid red line in panel A, and above the green dashed line in panel B, differ. For instance, the samples from Overton with a fine fraction % between 20 and 30 are deemed oversaturated using the Hassink equation, whilst they are undersaturated when the maximum fine fraction organic carbon is estimated by quantile regression. We therefore do not believe this to be contradictory as the samples that make up the absolute numbers presented in table 4 differ.

Moreover, I still think there is nothing wrong to fit saturation with the intercept (as done by Hassink, Six and others), since the saturation equation is likely a type of function where the x domain is always >0. At least, the recommended QR90 may be fitted also with the intercept; maybe it does not change too much when selecting the 90th percentile (as indeed regressing all the data). Unless the Authors considered the intercept approach totally wrong, this aspect can be mentioned.

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We agree that with large data sets it is unlikely that a sample will have no fine fraction, and therefore x is always > 0. However, in this instance we have opted to follow the logic that it is not possible to detect organic carbon in the fine fraction, when there is none, and chose to use a forced zero intercept.

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Line 33: likely 'protection' in aggregates is more appropriate than stabilization Changed

Line 77: The sentence about priming is very generic without any reference.

The sentence has been changed to the following, lines 72 and 73.

"Organo-mineral associations form the basis of microaggregates (Baldock and Skjemstad, 2000), and thus the destruction of aggregates makes the organic carbon protected within the aggregates more accessible for mineralisation by the soil microbial community. This may result in the increased mineralisation of existing SOC, known as the priming effect (Kuzyakov et al., 2000)"

Line 191 and Fig.1. As C:N of fine fraction was inserted, I would suggest to add also C:N of bulk soil since there is room in the figure layout.

The C:N of the bulk soil has been added to the figure 1, panel F and the caption has been updated. Line 161 has been changed to the following.

"The fine fraction and bulk soil C:N ratios were significantly different between the sites, Figure 1. However, the mean value of the fine fraction showed little deviation,  $9.84 \pm 0.94$  (mean  $\pm$  standard deviation). Full details of all the measured properties of bulk and fine fraction, per field are presented in Table A1."

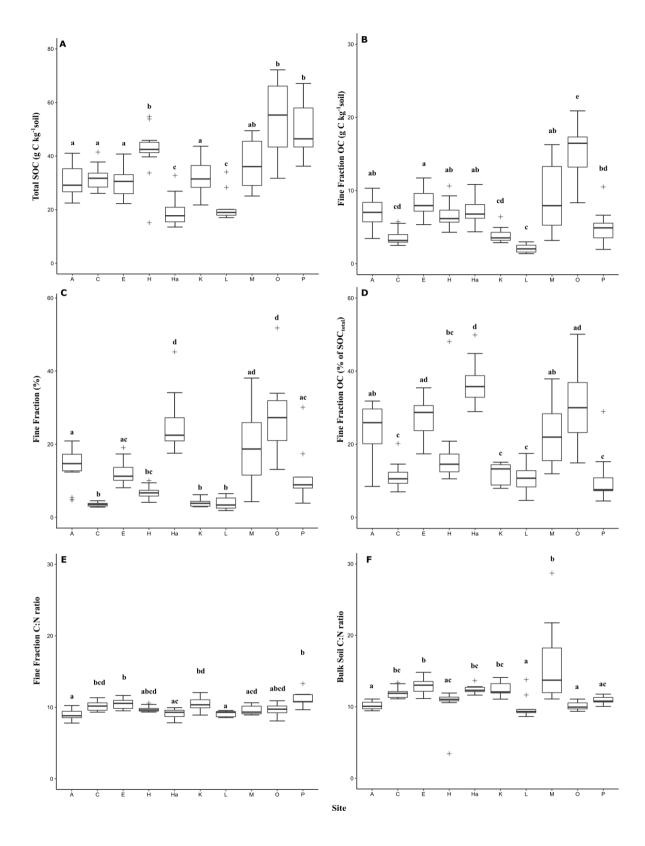


Figure 1. Measured total SOC (g C kg<sup>-1</sup>soil) (A) total fine fraction organic carbon (g C kg<sup>-1</sup>soil) (B), mass proportion of fine fraction (< 20  $\mu$ m, %) (C), relative proportion of measured fine fraction organic carbon of the total SOC content of the bulk soil (D), fine fraction C:N ratio (E) and bulk soil C:N ratio (F), for each of the grassland sites; Aberyswyth (A), Crichton (C), Easter Bush (E), Hillsborough (H), Harpenden (Ha), Kirkton (K), Llangorse (L), Myerscough (M), Overton (O) and Plumpton (P). Boxes represent the 25<sup>th</sup> and 75<sup>th</sup> percentile, with lines showing the median value. Whiskers show the lowest and highest values with outliers indicates as crosses (> 1.5 times the interquartile range). Lettering indicates significant differences between soils (P < 0.05).

Line 270: at least in the appendix (figure caption), a small explanation of how C inputs are calculated is needed if authors want to comment this figure. By the way, is not clear what the unit "estimated fine fraction OC input OC (g kg-1 C kg-1)" is.

The calculation method has been added to the caption of figure 1A.

"Figure A1. Estimated fine fraction OC input (g C kg<sup>-1</sup> soil) compared to measured fine fraction OC (g C kg<sup>-1</sup> soil) in each of the sites studied. The estimated fine fraction OC input (g C kg<sup>-1</sup> soil) was calculated by subtracting the maximum fine fraction OC (g C kg<sup>-1</sup> soil) from the current fine fraction OC (g C kg<sup>-1</sup> soil). The maximum fine fraction OC (g C kg<sup>-1</sup> soil) was estimated using the quantile regression equation ( $\tau = 0.90$ ), where, maximum fine fraction OC = 0.92 multiplied by the mass proportion fine fraction (%)."

Lines 351-2: isn'it the quantile regression applied also linear?

85 Yes, this is correct, the words "least squares" have been added to correct the sentence.

#### 90 Reviewer 2

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The authors have improved many aspects of the manuscript, providing further details, and strengthening the structure and conclusions on this compelling topic. I have some further comments, which I offer as suggestions, towards a foreseeable acceptance of this work for publication.

95 It would be nice to see a stronger case made for the quantile and boundary line expression based on its much higher empirical estimates of a maximum fine fraction OC. If only the higher data points are used, in what property do they differ from the lower data points? Is there a correlation of the soil properties with the differences from the maximum fine fraction OC? As the authors emphasize the regression method has "repercussions for decisions made regarding land management". If only the higher data points are considered, how can this be translated into practice?

100 Why should the maximum fine fraction OC be ranked even higher considering that its variability also increases? Can you identify specific soil types or properties that can help to adapt and improve local decisions on land management?

In the conclusions we summarise our support for the use of quantile regression to estimate maximum fine fraction OC. The reasons for this are outlined in the discussion, primarily due to its statistical robustness.

We have explored the data to try and identify soil properties which could help explain why a sample is further or closer to saturation, as suggested by examining those at either end. However, it was not possible to elucidate any properties which may be causing this. This was discussed in lines 229 to 240, from the perspective of additional OC input to reach saturation.

In terms of land management, we felt that the sites were too diverse to give some practical recommendations. In order to do so it would be good to examine paired grasslands, in terms of soil and environmental characteristics, but with contrasting management, for example organic versus conventional to tease out what management processes can be altered to maximise fine fraction OC. The following was added to lines 254 to 257.

"Further work investigating grasslands with similar soil types and textures, and environmental conditions, but contrasting management in terms of fertiliser regimes, grazing densities, sward composition and management, may help to elucidate management factors which can be used to increase fine fraction OC and explain the differences observed in this work."

Given the high variability of the maximum fine fraction OC found in larger datasets and how it responds to a multitude of soil properties and management factors, the applicability of a linear regression estimate may be debated but still provide a valid approach depending on the individual research questions. From my perspective, the authors should reword the statements in 1. 66 and 231 towards more specific and constructive details when referring to previous studies that used the Hassink linear regression equation. Can you quantify the mentioned over or underestimation as well as the variability of the data in the 90th percentile?

Overall, nice work.

The sentences have been changed to the following, in order to convey the aim of this work to assess the suitability of the Hassink, (1997) equation for estimating maximum fine fraction OC in UK grasslands.

## Lines 57 to 61.

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"To our knowledge no comparisons between the equation developed by Hassink, (1997) and one generated in the same way with a different data set has been done within the UK. This style of assessment would help to determine the suitability of the Hassink, (1997) linear regression equation to predict maximum fine fraction OC in UK soils, without this sequestration potentials may be both over and underestimated."

## 140 Lines 196 to 201.

"Determining the potential C sequestration capacity of soils is essential to predict the influence of land management for climate change mitigation. The determination of saturation deficit using the mass proportion of the fine fraction and current fine fraction OC content is an established method with a strong grounding in correlation between the variables. As mentioned earlier, previous studies have examined methods to improve estimates of maximum fine

fraction OC, however thus far no comparison has been made between the Hassink, (1997) linear regression equation, and one developed using grassland soils in the UK."

Standard deviation of 90<sup>th</sup> percentile added to line 239.

"The measured fine fraction OC in the UK sites lacks homogeneity of variance (Fig. 4), where the variation in the measured fine fraction OC increases with the proportion of fine fraction. Standard deviation of the proportion of fine fraction in the 10<sup>th</sup> percentile is 0.4 compared to 90<sup>th</sup> percentile of 6.9."

# Estimating maximum fine fraction organic carbon in UK grasslands

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Abstract. Soil organic carbon (SOC) sequestration across agroecosystems worldwide can contribute to mitigate the effects of climate change by reducing levels of atmospheric CO<sub>2</sub>. Stabilisation of organic carbon (OC) in the fine soil fraction (< 20 µm) is considered an important long-term store of SOC and the saturation deficit (difference between measured OC and estimated maximum OC in the fine fraction) is frequently used to assess SOC sequestration potential following the linear regression equation developed by Hassink, (1997). However, this approach is often taken without any assessment of the fit of the equation to the soils being studied. The statistical limitations of linear regression have previously been noted, giving rise to the proposed use of boundary line analysis and quantile regression to provide more robust estimates of maximum SOC stabilisation. The objectives of this work were to assess the suitability of the Hassink, (1997) equation to estimate maximum fine fraction OC in UK grassland soils of varying sward ages and to evaluate the linear regression, boundary line and quantile regression methods to estimate maximum fine fraction OC. A chronosequence of 10 grasslands was sampled, in order to assess the relationship between sward age (time since last reseeding event) and the measured and predicted maximum fine fraction OC. Significantly different regression equations show that the Hassink, (1997) equation does not accurately reflect maximum fine fraction OC in UK grasslands when determined using the proportion of fine soil fraction (< 20 µm, %) and measured fine fraction OC (g C kg<sup>-1</sup> soil). The OR estimate of maximum SOC stabilisation was almost double that of linear regression and BL analysis (0.89  $\pm$  0.074, 0.43  $\pm$  0.017 and 0.57  $\pm$  0.052 g C kg<sup>-1</sup> soil, respectively). Sward age had an inconsistent effect on the measured variables and potential maximum fine fraction OC. Fine fraction OC across the grasslands made up 4.5 to 55.9% of total SOC, implying that there may be either high potential for additional C sequestration in the fine fraction of these soils, or stabilisation protection in aggregates is predominant in these grassland soils. This work highlights the need to ensure that methods used to predict maximum fine fraction OC reflect the soil *in situ*, resulting in more accurate assessments of carbon sequestration potential.

## 1. Introduction

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Carbon (C) sequestration in soils offers a significant opportunity to remove CO<sub>2</sub> from the atmosphere and store it into long lived C pools (Lal, 2004; Powlson et al., 2011), with co-benefits for soil structure and functioning (Lorenz and Lal, 2018; Smith, 2012; Soussana et al., 2004). However, to utilise soils as a CO<sub>2</sub> drawdown mechanism, accurate estimates of their storage capability are required. With respect to soil organic carbon (SOC) sequestration, organic carbon (OC) stabilised via adsorption to mineral surfaces in the fine soil fraction (< 20 µm) is often regarded as the most important due to its longer residence time (Baldock and Skjemstad, 2000; Six et al., 2002). There is empirical evidence that there is an upper protective capacity limit, or saturation point of the mineral stabilised OC pool (Six et al., 2002; Stewart et al., 2007). Potential SOC sequestration (or saturation deficit) can be estimated by subtracting the current fine fraction OC from the estimated maximum fine fraction OC (Angers et al., 2011).

Hassink, (1997) compared pairs of Dutch arable and grassland soils and found that while soil bulk SOC contents significantly differed among soils, fine fraction OC did not. These findings led to the idea that the saturation point of the fine soil fraction could be estimated by linear regression using the mass proportion of fine fraction in a soil sample (%) and the current fine fraction OC (g C kg<sup>-1</sup> soil). Several iterations of the concept have been proposed to overcome the limitations of linear regression. For example, boundary line analysis uses a defined upper or lower subset of a data set to estimate the boundary line, when a limiting response to an independent variable(s) along a boundary is supported (Lark and Milne, 2016; Schmidt et al., 2000). Using the upper 90<sup>th</sup> percentile of a data set, boundary line analysis overcomes the limitation of linear regression depicting the mean response to the independent variable (Feng et al., 2013; Shatar and Mcbratney, 2004), which is thought to cause an underestimation of sequestration potential. Quantile regression estimates the response of a specific quartile using the entire data set. It also makes no assumptions regarding homogeneity of variance, thus increasing the robustness of the estimated maximum fine fraction OC, as sample size is not reduced as in BL analysis (Beare et al., 2014; Cade and Noon, 2003). Using a forced zero intercept overcomes the contradiction of a positive intercept indicating the presence of fine fraction OC without any fine soil fraction (Beare et al., 2014; Feng et al., 2013; Liang et al., 2009). To our knowledge no comparisons between the equation developed by Hassink, (1997) and one generated in the same way with a different data set has been done within the UK. This style of assessment would help to determine the suitability of the Hassink, (1997) linear regression equation to predict maximum fine fraction OC in UK soils, without this sequestration potentials may be both over and underestimated. These suggestions have been proposed to improve estimates of maximum fine fraction OC. However, several studies use the original equation presented by Hassink, (1997) to estimate sequestration potential at different scales (e.g. Angers et al., 2011; Chen et al., 2019; Lilly and Baggaley, 2013; Wiesmeier et al., 2014). There is a need for validity checks to determine the suitability of the Hassink, (1997) linear regression equation to predict maximum fine fraction OC of the soils in the respective studies. Without this sequestration potentials may be both over and underestimated.

Within the UK, human-managed grasslands are the dominant land use, covering 36% of the land area (Ward et al., 2016). Managed grasslands are planted and maintained to increase agricultural productivity through fertiliser and liming applications, and the re-seeding of swards. The high levels of disturbance associated with re-seeding events by mould board ploughing and harrowing in particular, result in changes in soil structure, notably the breaking up of aggregates, nutrient cycling and SOC mineralisation (Carolan and Fornara, 2016; Drewer et al., 2017; Soussana et al., 2004). Organo-mineral associations form the basis of microaggregates (Baldock and Skjemstad, 2000), and thus the destruction of aggregates makes the organic carbon protected within the aggregates more accessible for mineralisation by the soil microbial community. This may result in the increased mineralisation of existing SOC, known as the priming effect (Kuzyakov et al., 2000) organo-mineral stabilised OC in the fine fraction, more accessible for mineralisation by the soil microbial community. Additionally, the release of other organic carbon pools may induce a priming effect, potentially enhancing the losses from the typically stable mineral associated OC in the fine fraction. The long-term effect of such re-seeding event on SOC dynamics is understudied, it is therefore important to understand how disturbance might affect OC in the fine fraction, and thus the SOC sequestration ability of managed grasslands.

The objectives of this study were (i) to assess the suitability of the Hassink, (1997) equation to estimate maximum fine fraction OC in UK grassland soils, (ii) to evaluate the linear regression, boundary line and quantile regression methods to estimate maximum fine fraction OC, and (iii) to explore the relationship between sward age (time since last reseeding event), and current and predicted maximum fine fraction OC. We hypothesised that i) the linear regression equation developed using UK grassland soils would be significantly different to that of Hassink, (1997), and that ii) grasslands with an older sward age, would have a greater proportion of total SOC stabilised in fine fraction (< 20 µm) and a lower sequestration potential.

## 2. Materials and Methods

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## 2.1 Site Description and Sampling

Ten grassland chronosequences covering a wide range of soil types, land use and climatic conditions were identified across the UK in 2016. The sites included the range of agricultural activity associated with UK grasslands (upland grazing, dairy, and mixed grazing), variations in soil type (organo-mineral, mineral and chalk) and the majority of UK climatic zones (Table 1). At each location, five to eight individual fields of different sward age (represented by years since a ploughing and reseeding

event), ranging from 1 to 179 years, were identified for sampling. In each field, areas were avoided which had different applications of manure, soil types or topography, headlands, areas near gates, where lime or manure had previously been dumped, or where livestock congregate. Two replicate soil cores were collected to a depth of 30 cm using a soil auger with a 2.5 cm diameter steel core and bulked to give a single composite sample. This was repeated 10 times in each field at regular intervals in a 'W' shape across the field totalling 10 replicate samples per field per site. Intact soil cores for determining bulk densities were collected at three locations in each field at two depths (10 to 15 cm and 20 to 25 cm) using intact rings (7.5 cm diameter, 5 cm height). Replicate samples were sieved to 2 mm and fresh subsamples were used to determine soil pH in water. Remaining sieved soils were dried at 40°C and ball milled prior to determination of total C and N contents (% by mass) using a Flash 2000 elemental analyser. Intact soils were dried at 107°C and weighed to calculate dry bulk densities, any stones were removed.

#### 2.2 Soil fractionation

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The fine fraction (< 20 µm) of the soil was separated using a combined ultrasonic dispersion and sedimentation method adapted from Hassink, (1997). Briefly, 20 g of dried sieved soil was soaked in 100 mL of deionised water for 24 hours. The suspension was then sonicated with a Microson XL2000 Ultrasonicator for 20 minutes at 20 W in 50 mL centrifuge tubes, surrounded by ice to prevent overheating. The separated samples were recombined in 150 mL tubes, and shaken end over end to disperse the soil water suspension. Sedimentation times were determined using a table applying Stokes Law, for 20 µm particles, a particle density of 2.65 g cm<sup>-3</sup> and sedimentation depth of 5 cm at temperatures between 20°C and 35°C (Jackson, 2005). After the appropriate sedimentation time, the fine fraction was siphoned off the soil suspension. The fine fraction was dried for 24 hours at 107°C and ball milled prior to total C and N analysis (% by mass) using a Flash 2000 Elemental Analyser, to determine the current OC content of the fine fraction. At each site, a minimum of 3 fields varying in age (young, intermediate, and old at that location) were selected, and 3 of the 10 replicate field samples were selected at random for fractionation.

Hydrochloric acid (HCl) fumigation was used to remove carbonates from the Plumpton samples. Ball-milled samples, in silver capsules, were moistened with deionised water (1:4 sample:water ratio) to aid the efficiency of carbonate removal by HCl fumes (Dhillon et al., 2015). The samples were placed in a vacuum desiccator with a beaker of 100 mL of 12 M HCl, for 24 hours and subsequently dried in a ventilated oven at 60°C for 16 hours, to remove excess moisture and HCl (Dhillon et al., 2015). Total C and N contents were determined as outlined above.

# 120 2.3 Statistical analyses

All statistical analyses were carried out using R software version 3.5.3 (Team, 2019). Significant differences were determined by ANOVA's and by post-hoc Tukey tests ( $\alpha = 0.05$ ). Where assumptions of normality and variance were not satisfied by testing (Shapiro Wilkens and Levenes Test) significant differences were identified using Kruskal test and post hoc Dunn test. A Kendal tau ( $\tau$ ) correlation matrix was produced using the 'corrplot' package (Wei and Simko, 2017).

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#### 2.3.1 Regression analyses

Linear regression was used to predict maximum fine fraction OC, with the mass proportion of fine fraction ( $< 20 \mu m$ , %) in a sample and the measured OC of the fine fraction (g C kg<sup>-1</sup> soil) as the independent and dependent variables, respectively. Regression equations were developed for the combined UK data set, and the individual sites. Linear regression with a forced zero intercept was used with data from this study and the data published in Hassink (1997).

Boundary line analyses were performed as an alternative to linear regression, both with and without a forced zero intercept to predict maximum fine fraction OC for all UK sites. The data was organised by mass proportion of the fine fraction (%) and divided into subgroups at 5, 10 and 15% intervals. The 10% interval reflects the method of Feng et al. (2013), whilst the 5 and 15% intervals were used to assess the effect of interval on estimation of maximum fine fraction OC. The groups were then ordered by measured fine fraction OC (g C kg<sup>-1</sup> soil), and the values in the 90th percentile were used to plot the boundary line. Boundary line analysis was not used for individual sites as it resulted in too few data points. Quantile regression analysis was performed in Rstudio using the 'quantreg' package (Koenker, 2019), for the 90th and median percentiles ( $\tau = 0.90$  and  $\tau = 0.50$ ). Forcing the intercept to zero overcomes the paradox of having C stabilised as MAOC without any fine fraction in the soil. Significant differences between slopes were identified using the 'lsmeans' package (Lenth, 2016), followed by post-hoc Tukey tests ( $\alpha = 0.05$ ).

#### 2.3.2 Carbon saturation ratio

The carbon saturation ratio was determined in order to identify the degree of saturation across the sites, when estimating maximum fine fraction OC, using the Hassink (1997), UK, and site-specific linear regression equations both with and without a forced zero intercept, and the equations generated by boundary line and quantile regression analyses. The carbon saturation ratio was calculated by dividing the current fine fraction OC by the estimated maximum fine fraction OC content. Values < 1 were deemed under saturated, = 1 as at saturation and > 1 as oversaturated.

## 3. Results

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## 3.1 Current C concentrations

The measured total SOC and fine fraction OC concentrations exhibited variation within the grassland sites (Fig. 1). Total SOC varied from 8.2 to 85.8g C kg<sup>-1</sup> soil, with a median of 32.7 g C kg<sup>-1</sup> soil. Hillsborough, Overton and Plumpton had significantly higher total SOC, whilst Harpenden and Llangorse had the lowest total SOC (P < 0.05) (Fig. 1). The measured fine fraction OC ranged from 1.4 to 20.9 g C kg<sup>-1</sup> soil, with a median of 6.2 g C kg<sup>-1</sup> soil. Overton had the highest total fine fraction OC (P < 0.05) and was the only organically managed site (Fig. 1). The proportion of OC stabilised in the fine fraction ( $< 20 \mu m$ ) had high variability across the UK sites accounting for 4.5 to 50.1% of total SOC with a median of 17.5%. The proportion of total SOC stabilised in the fine fraction ( $< 20 \mu m$ ), and proportion of fine fraction in a sample did not significantly differ in Harpenden and Overton, however they have significantly different measured fine fraction OC contents (g C kg<sup>-1</sup> soil) (P < 0.05), indicating different saturation potentials (Fig. 1). Soil C:N ratio was positively correlated with fine fraction C:N (0.30, P < 0.0001), Table 2, however there was no relationship between bulk soil C:N ratio and proportion of fine fraction (data not shown). The fine fraction and bulk soil C:N ratios wereas significantly different between the sites, Figure 1.5 hHowever the mean value of all the fine fraction data-showed little deviation, 9.84  $\pm$  1.00 (mean  $\pm$  standard deviation). Full details of all the measured properties of bulk and fine fraction, per field are presented in Table A1.

The significance of correlations between the measured soil properties, time since reseeding and known environmental factors were analysed. The matrix of Kendall tau ( $\tau$ ) correlation coefficients in Table 2, revealed that measured fine fraction OC was positively correlated with median annual temperature ( $\tau$  = 0.13, P < 0.05), %N ( $\tau$  = 0.26, P < 0.0001) and %C ( $\tau$  = 0.27, P < 0.0001) in the bulk soil, and negatively correlated with mean annual rainfall ( $\tau$  = -0.36, P < 0.0001), and %N ( $\tau$  = 0.15, P < 0.05) in the fine fraction. Mass proportion of fine fraction and measured fine fraction OC (g C kg<sup>-1</sup> soil) were positively correlated in cambisols ( $R^2$  = 0.61, P < 0.05), gleysols ( $R^2$  = 0.76, P < 0.05), podzols ( $R^2$  = 0.93, P < 0.05), and stagnosols ( $R^2$  = 0.88, P < 0.05) (Fig. 2). However, the proportion of total SOC stabilised in the fine fraction (< 20 µm), was greatest in luvisols (P < 0.05) (Fig. 3).

## 3.2 Estimated maximum fine fraction organic carbon

The slope generated from the UK data used to estimate MAOC (Table 3) was significantly different (P < 0.05) to the slope reported in Hassink (1997). There was no significant difference between the slopes generated from the UK data, the data from Hassink (1997) when estimated by linear regression with a forced zero intercept. Significantly different (P < 0.05) slopes were found between the individual UK sites, owing to the range in the proportion of the fine fraction within each sample, from 1.85 to 51.8%, (Tables A3 and A4).

Coefficients from boundary line analysis are presented in (Table 3). There was no significant difference in slopes between the 5, 10, and 15% fine fraction intervals used. The median percentile quantile regression analysis had a similar slope to the boundary line and linear regression with forced zero intercept. Quantile regression using the 90<sup>th</sup> percentile resulted in the steepest slope of all estimation methods (Table 3). The C saturation ratios revealed the difference in number of samples with potential to sequester more C (Table 4). The Hassink (1997) linear regression equation, without a forced zero intercept, predicted the greatest number of unsaturated sites, followed by the 90<sup>th</sup> percentile quantile regression, with a forced zero intercept. There was no clear relationship between oversaturated sites and proportion of silt and clay contents as oversaturation occurred across all proportions, indicated by points above the lines in Fig. 4.

#### 3.3 Effect of sward age on current C concentrations and estimated maximum MAOC

Sward age (years since last reseeding event) had a weak positive correlation with the mass proportion of the fine fraction (%) (Table 2). When grouped in five year intervals, significant differences were found between age group and the mass proportion of the fine fraction (%), measured fine fraction OC (< 20 µm), (g C kg<sup>-1</sup> soil), and the C:N ratio of the fine fraction (Table 5), however there was no consistent increase or decrease with sward age. At the individual sites, significant differences were observed between fields, with some properties, but again there was no consistent effect of sward age (Tables A3 and A4).

## 4. Discussion

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## 4.1 Estimation of maximum fine fraction organic carbon

Determining the potential C sequestration capacity of soils is essential to predict the influence of land management for climate change mitigation. The determination of saturation deficit using the mass proportion of the fine fraction and current fine fraction OC content is an established method with a strong grounding in correlation between the variables. As mentioned earlier, previous studies have examined methods to improve estimates of maximum fine fraction OC, however thus far no comparison has been made between the Hassink, (1997) linear regression equation, and one developed using grassland soils in the UK.

Determining the potential C sequestration capacity of soils is essential to predict the influence of land management for climate change mitigation. The determination of saturation deficit using the mass proportion of the fine fraction and current fine fraction OC content is an established method with a strong grounding in correlation between the variables. However, the Hassink (1997) linear regression equation has been used to estimate sequestration potentials, without prior testing to determine its applicability to the soils in question (e.g. Angers et al., 2011; Chen et al., 2019; Lilly and Baggaley, 2013; Wiesmeier et al., 2014). This may have potentially over or underestimated sequestration potential, which may have repercussions for decisions

made regarding land management. The significantly different slopes for the linear regression equations (Table 3) shows that the Hassink (1997) regression equation is not suitable for estimating maximum fine fraction OC in UK grasslands. Previous concerns have focused on the potential for the equation developed by Hassink (1997) to underestimate maximum fine fraction OC, as least squares linear regression represents the mean response of the independent variable, rather than the maximum. For the UK grasslands in this study estimating maximum fine fraction OC using the Hassink (1997) regression approach resulted in a significant overestimation of fine fraction OC sequestration potential. Future work using maximum fine fraction OC predictions equations reported in the literature (e.g. Beare et al., 2014; Feng et al., 2014; Hassink, 1997; Six et al., 2002) should first conduct a validity test, and determine if the regression equations match the soils in question or a subset of the data, to ensure results are not significantly over or underestimated.

To overcome the contradiction of an intercept greater than zero, indicating that C is stabilised in the fine fraction without any fine fraction, a forced zero intercept was used. The linear regression slopes with a forced zero intercept were not significantly different, and were similar to that of Feng et al. (2013),  $0.42 \pm 0.002$ . Liang et al. (2009) reported a lower slope of 0.36 in Chinese black soils, whilst Beare et al. (2014) reported a slope of  $0.70 \pm 0.03$  in long-term New Zealand pastures. The range of reported values, and differences across the UK sites (Tables A3 and A4), suggest that the effect of the proportion of fine fraction of a sample on fine fraction OC is not consistent and likely reflects differences in pedogenic and environmental conditions, land management and possibly the fine fraction OC isolation method. It may be that the use of the mass proportion of fine fraction to predict maximum fine fraction OC is only suited on larger scales, rather than smaller, site specific scales, as indicated by the variability in this study.

Boundary line analysis and quantile regression have been suggested as alternatives to overcome the limitations of linear regression. The estimation of maximum fine fraction OC was greatest when using quantile regression ( $\tau$  = 0.90), whereas boundary line estimates at 5 and 10% intervals were similar to quantile regression ( $\tau$  = 0.50), and those estimated from linear regression (Table 3). The use of the median percentile quantile regression highlights the closeness of linear regression predictions being more indicative of mean values, thus underestimating SOC sequestration potential. The boundary line estimate of Feng et al. (2013), 0.89  $\pm$  0.05, was nearly double their linear regression; this was not the case in our study. Boundary line analysis uses a subset of data to estimate, in this case, an upper limit, the data set used by Feng et al. (2013) had a wider spread of measured fine fraction OC of 0.9 to 71.7 g C kg<sup>-1</sup> soil, compared to 1.72 to 18.29 g C kg<sup>-1</sup> soil in our UK soils. Therefore, the upper subset of data was composed of higher values giving a steeper slope and demonstrates that the C sequestration estimate generated by boundary line analysis is biased by the range of data.

The strength of using quantile regression analysis is that it makes no assumptions of homogeneity of variance and uses the entire data set to estimate the upper limit of a response. The measured fine fraction OC in the UK sites lacks homogeneity of variance (Fig. 4), where the variation in the measured fine fraction OC increases with the proportion of fine fraction. Standard deviation of the proportion of fine fraction in the 10<sup>th</sup> percentile is 0.4 compared to 90<sup>th</sup> percentile of 6.9.

Therefore, of the methods explored in this study for our grassland soils, we consider the quantile regression at the 90<sup>th</sup> percentile estimate of maximum fine fraction OC to be the most robust. This method results in the greatest number of unsaturated samples (Table 4) suggesting great potential for additional sequestration.

When examining the estimated OC input versus existing fine fraction OC using estimates generated by quantile regression at the 90<sup>th</sup> percentile, a positive correlation between current fine fraction OC and estimated C input (Kendall tau ( $\tau$ );0.323, P < 0.001), was observed for the entire data set. However, this was not the case at the site level (Fig. A1). Where in some instances increasing fine fraction OC (g C kg<sup>-1</sup> soil) was associated with increased estimated C input until saturation, such as Aberyswyth, Myerscough and Plumpton. Therefore, despite a higher fine fraction OC content these samples are furthest from saturation. In contrast the opposite was true for Crichton and Hillsborough (and Harpenden, Kirkton and Overton, although not statistically significant) implying that for these sites, samples with a higher fine fraction OC are closer to, or over saturation. It is unclear why this is the case particularly as in all sites, bar Harpenden, there is a positive regression between mass proportion of the fine fraction and fine fraction OC (Table A3). Meaning that higher fine fraction OC is also associated with higher mass proportion of the fine soil fraction. It is likely that the OM input to the soils with the higher mass proportion of fine fraction is insufficient to bridge the gap between current and estimated maximum fine fraction OC, as it is not possible to identify any other effect due to pedogenic or environmental conditions, but contrasting management in terms of fertiliser regimes, grazing densities, sward composition and management, may help to elucidate management factors which can be used to increase fine fraction OC and explain the differences observed in this work.

Estimating maximum fine fraction OC on the basis of mass proportion of fine fraction is likely to be an oversimplification of the dynamics of fine fraction OC accrual. Other parameters such as mineralogy, soil microbial community, environmental conditions (e.g precipitation, Table 2) and land management, can significantly influence fine fraction OC stabilisation (Cotrufo et al., 2015; Kallenbach et al., 2016). This work has identified some soil and environmental properties that may play a role in fine fraction OC stabilisation such as median annual temperature, %N and %C in the bulk soil, mean annual rainfall and %N in the fine fraction (Table 2). Warmer median annual temperatures may enhance plant productivity and microbial processing, the by-products of which are important precursors to fine fraction OC (Cotrufo et al., 2013). It would be interesting to know at which point higher temperatures have a deleterious effect on fine fraction OC accumulation. Mean annual rainfall and %N in the fine fraction were negatively correlated to fine fraction OC. It was anticipated that fine fraction OC would be positively correlated with fine fraction N, as nitrogen rich microbial by-products have been found to form new organo-mineral associations onto which OC preferentially binds (Kopittke et al., 2018). These bonds may have been disturbed during the fractionation process, resulting in an N rich fine fraction with less OC content.

The influence of soil type of fine fraction OC was also evident in our results as all soil types had statistically significant positive correlations between the mass proportion of fine fraction and measured fine fraction OC, except for leptosols and

luvisols (Fig. 2). However, these soil types exhibited the greatest proportion of total SOC stabilised in the fine fraction (Fig.3). Luvisols have a high base saturation facilitating more fine fraction OC stabilisation via complexation of organic ligands by free Ca<sup>2+</sup> (Chen et al., 2020). Identifying soils where a greater proportion of total SOC is stored in the fine fraction is important to identify where fine fraction OC needs to be protected, but also where it can be enhanced.

Whilst we consider the quantile regression at the 90th percentile method to provide the most robust estimate of maximum fine fraction OC in the sites studied, further experimental work to test the saturation level of these soils, would help to validate this. Incubation studies which force an unsaturated soil to its 'saturation' level and the effect of influencing variables, mentioned above will help to elucidate the factors controlling fine fraction OC saturation. In particular further empirical evidence of how to manipulate fine fraction OC stabilisation processes in a way that is practical for grassland management to promote the formation of new organo-mineral associations, and understanding their stability will be important for establishing the true potential of additional carbon sequestration across managed grasslands.

# 4.3 Effect of sward age on fine fraction OC

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It was anticipated that for fields of an older sward age, a greater proportion of total SOC would be stabilised as fine fraction OC, as tillage breaks up macroaggregates making OC in the fine fraction available for mineralisation. Alternatively, fine fraction OC is less sensitive to disturbance than particulate organic matter (POM), resulting in the accumulation of POM as the fine fraction OC pool remains stable, if sufficiently saturated. The results seem to support neither hypothesis. The proportion of total SOC stabilised in the fine fraction was not consistently higher in the oldest field, and in some instances was significantly less, such as Aberystwyth (Table A2). When grouped in five year intervals, significant differences in C:N ratio of the fine fraction, the proportion of fine fraction in a sample (%) by mass, measured fine fraction OC (g C kg<sup>-1</sup> soil) and the relative proportion of measured fine fraction OC of the total SOC content of the bulk soil were found between age groups (Table 5), however, there was no consistent trend in the results. This data does not support the hypothesis that older swards will have a greater proportion of SOC stabilised in the fine soil fraction, and a reduced potential for additional C sequestration. Equally there was no negative correlation between sward age and the proportion of total SOC stabilised which would be supportive of the alternate hypothesis. From the data it appears that fine fraction OC makes up a greater proportion of SOC with increasing sward age when comparing the less than 5 years, 6 to 10 and 11 to 15 years age groups. However there is a significant decrease in the amount of SOC that is stabilised in the fine fraction in the 16 to 20 years group, this is likely due to fields in this age range originating from Crichton, Hillsborough and Plumpton, which have some of the lowest mass proportion of the fine fraction (Figure 1, C). The sward age analysis may also be confounded by the variation of the proportion of fine fraction, particularly on soil properties influenced by mass proportion of fine fraction such as %C and %N and current fine fraction OC (g C kg<sup>-1</sup> soil). However, it was not possible to conduct robust ANCOVA's with a grouping variable with more than two levels. It may be possible to elucidate the relationship better from a wider study with more samples per age group as our 16 to 20 years group only has 9 values compared to 48 in the less than 5 years group.

Fine fraction OC only accounted for 4.5 to 50.12% indicating high OC storage in other soil pools such as POM, or different aggregate fractions. The fine roots of grassland flora species promote aggregate formation (O'Brien and Jastrow, 2013; Rasse et al., 2005), which may be a dominant stabilisation process in grasslands. However previous work has found no effect of sward age or the frequency of grassland reseeding on the % C in differing aggregate fractions (> 2000 μm, 250–2000 μm, 53–250 μm and < 53 μm) (Carolan and Fornara, 2016; Fornara et al., 2020). The impact of reseeding disturbance may be offset due to the high density of roots in grasslands by facilitating aggregate reformation. Additionally, dissolved organic carbon from below ground inputs is more efficiently stabilised in organo-mineral associations than above ground dissolved organic carbon (litter leachate) (Sokol and Bradford, 2019). The narrow rhizosphere to bulk soil ratio in grasslands, may make the fine fraction OC in grasslands more resilient to disturbance events.

#### 5. Conclusions

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315 Estimating the long-term sequestration of soil C in the fine fraction is difficult due to the lack of reliable methodologies that can be widely applied to all soils. Our study has demonstrated that the Hassink (1997) linear regression equation is not suitable to estimate maximum fine fraction OC in a range of UK grassland soils. Therefore, caution should be applied to estimates of maximum fine fraction OC obtained using the Hassink (1997) equation, in instances where it may not accurately reflect fine fraction OC of the soil in situ. After exploring various univariate estimation methods, we recommend the use of quantile 320 regression at the 90<sup>th</sup> percentile to overcome the shortfalls of least squares linear regression. However, such a simple estimate is unlikely to accurately reflect the dynamics of fine fraction OC stabilisation. This work has helped to identify some key parameters which play a role in fine fraction OC stabilisation, such median annual temperature, mean annual precipitation, bulk soil %C and %N and fine fraction %N. Further work to understand how these parameters influence fine fraction OC dynamics, will help to accurately assess the feasibility of achieving soil carbon sequestration targets. Our results showed little 325 evidence of the impact of time since last reseeding event on the OC in the fine soil fraction. However, improving our understanding of SOC stabilisation processes, and their resilience to grassland management is essential to ensure that current SOC is not only enhanced but also protected.

#### **Author contribution**

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KCP, SB, JMC, RMR and EMB formulated the research question and study design. KCP conducted the experimental work, data analysis, and prepared the manuscript draft. All authors contributed to editing and reviewing of the manuscript.

# **Data Availability**

All data resulting from this study are available from the authors upon request to Sarah Buckingham (sarah.buckingham@sruc.ac.uk)

#### **Competing Interest**

335 The authors declare that they have no conflict of interest.

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Table 1 Summary of UK grassland site characteristics.

Site	Age range (years)	Land Use <sup>a</sup>	Mean Annual Temperature (°C) <sup>b</sup>	Mean Annual Rainfall (mm) <sup>a</sup>	Elevation (m.a.s.l)	WRB Soil Type <sup>c</sup>	Soil Texture <sup>c</sup>
Aberystwyth (52°25'N 04°02'W)	2 to 33	UpG	9.5 to 11	1000	20 to 65	ST, CM	Clay to sandy loam
Crichton (55°02'N 03°35'W)	1 to 20	DP	9.5 to 9.9	1100	5 to 50	CM	Clay loam to sandy loam
Easter Bush (55°51'N 03°52'W)	3 to 6	MG	6 to 9	< 700	215 to 265	GL	Clay loam to sandy loam
Harpenden (51°48'N 00°22'W)	22 to 179	UnG	9.5 to 10.5	700	120 to 130	LV	Silty clay loam
Hillsborough (54°27' N 6°04' W)	1 to 37	DP	8.5 to 10	900	120	СМ	Clay loam
Kirkton (56°25'N 04°39'W)	1 to 35	UpG	8 to 9.4	2528	163 to 170	PZ	Clay loam to sandy loam
Llangorse (51°55'N 03°16'W)	2.5 to 25	MG	8 to 10	1000		СМ	Loam/ Clay to Silty loam
Myerscough (53°51'N 02°46'W)	2 to 48.4	MG	9 to 10.5	1000	8 to 15	GL	Clay to sandy loam
Overton (51°48'N 02°08'W)	3 to 50	MGO	9 to 11	800	240 to 276	LP	Clay loam to silty loam
Plumpton (50°54'N 00°04'W)	1 to 20	MG	9.5 to 11	800	49 to 85 &160 to 215	ST	Clay to clay loam, Chalky clay to chalky loam

<sup>&</sup>lt;sup>a</sup> Land Use; DP; Dairy pasture, MG; Mixed grazing; MGO; Mixed grazing organic, UpG; Upland grazing, UnG; Ungrazed.

<sup>&</sup>lt;sup>b</sup> Mean annual temperature and rainfall estimated from Met Office climatic region summaries, averaged over 1981 to 2010.

<sup>&</sup>lt;sup>c</sup> World Reference Base (WRB) Soil Type: ST; Stagnosols, CM; Cambisols, GL; Gleysol, LV; Luvisols; PZ; Podzol; LP;

<sup>455</sup> Leptosol .Soil type and texture determined from GPS locations and UK Soil Observatory Map viewer.

Table 2. Correlation matrix of Kendal tau ( $\tau$ ) coefficients for bulk and fine fraction ( $<20~\mu m$ ) soil properties, sward age and known environmental parameters.

							Bulk S	Soil			Fine	Fracti	on
		Temp.	Prec.	Age	%N	%C	C:N	pН	%SC	Fine fraction OC	%N	%C	C:N
	Temp.	1											
	Prec.	-0.05	1										
	Age	0.15	-0.11	1									
	%N	0.23***	-0.07	0	1								
	%C	0.16*	0.06	0.04	0.73***	1							
Soil	C:N	-0.25***	-0.05	0.02	_a	_a	1						
Bulk soil	pН	0.07	-0.30***	-0.07	-0.04	0.03	0.02	1					
	%SC	0.26***	-0.43***	0.14*	0.12	0.12	-0.01	0.21***	1				
	Fine fraction OC	0.13*	-0.36***	0.1	0.26***	0.27***	0.01	0.12	_a	1			
u	%N	-0.32***	0.28***	-0.07	0.17**	0.14*	-0.02	-0.27***	-0.47***	-0.15**	1		
Fine Fraction	%C	-0.33***	0.25***	-0.09	0.18**	0.17**	0.06	-0.25***	-0.47***	_a	0.87***	1	
	C:N	-0.21***	-0.08	-0.16	0.11**	0.21***	0.30***	-0.05	-0.15*	-0.02	_a	_a	1

<sup>460</sup> a No correlation calculated as one variable used to calculate the other.

Age; years since last reseeding event, Temp; median value from the mean annual temperature range (°C), Prec.; mean annual rainfall (mm), %SC; mass proportion of fine fraction in a sample (%), Fine fraction OC; measured fine fraction OC (g C kg<sup>-1</sup> bulk soil).

Level of significance: \* P < 0.05, \*\* P < 0.01, \*\*\*P < 0.0001

Table. 3. Analyses coefficients for the estimation of maximum fine fraction organic carbon by linear regression (LR), linear regression with forced zero intercept (LR\_0), boundary line (BL) and quantile regression (QR). Lettering indicates slopes which were significantly different within a method (P < 0.05).

Method		Slope (± 1 SEM)	P slope	Intercept (± 1 SEM)	P intercept	RMSE	n	$\mathbb{R}^2$
LR	Hassink, (1997)	$0.37^{a}$		4.09			40	
	All UK	$0.32 \pm 0.023^{b}$	***	$2.86 \pm 0.368$	***	2.58	129	0.61
LR_0	Hassink, (1997) <sup>a</sup>	$0.45 \pm 0.02$	***			4.97	40	0.94
	All UK	$0.47 \pm 0.017$	***			3.13	129	0.85
	5% intervals	$0.48 \pm 0.058$	***			5.89	19	0.79
BL	10% intervals	$0.48 \pm 0.070$	***			6.36	15	0.77
	15% intervals	$0.56 \pm 0.056$	***			4.77	14	0.89
	QR ( $\tau = 0.90$ )	$0.92 \pm 0.071$	***			7.90	129	0.90
QR	$QR \ (\tau = 0.50)$	$0.49 \pm 0.032$	***			3.15	129	0.66

<sup>475</sup> RMSE, root mean square error.

Level of significance: \*\*\* P < 0.001

<sup>&</sup>lt;sup>a</sup> Data extracted from Hassink, (1997) used to generate slope value with forced zero intercept.

Table 4. Carbon saturation ratios calculated from the estimated maximum fine fraction organic carbon by linear regression (LR), linear regression with forced zero intercept (LR $_0$ ), boundary line (BL) and quantile regression (QR). Values < 1 indicate unsaturated, = 1 at saturation and > 1 are oversaturated samples.

Method		No. of unsaturated	Mean ratio	Median	
		samples $(n = 129)$			
LR	Hassink, (1997)	105	0.77	0.73	
	UK	75	0.98	0.94	
	UK site specific	71	1	0.99	
Forced 0 intercept					
LR_0	Hassink (1997)	30	1.52	1.44	
	UK	34	1.47	1.39	
	UK site specific	57	1.09	1.04	
BL	5%	38	1.42	1.34	
	10%	36	1.43	1.35	
	15%	50	1.22	1.15	
QR	50 <sup>th</sup>	38	1.4	1.32	
	90 <sup>th</sup>	99	0.74	0.7	

Table 5. Effect of sward age grouped at five year intervals on selected soil properties. Values are means  $\pm$  standard error of the mean, and different letters indicate age groups which are significantly different (P < 0.05), by columns.

Age	n	C:N	% SC	Fine fraction organic carbon (g C kg <sup>-1</sup> soil)	Fine fraction organic carbon (% of SOC <sub>total</sub> )
0 to 5	48	$10.18 \pm 0.15^{a}$	$10.00 \pm 1.41^{a}$	$5.68 \pm 0.49^{a}$	$18.32 \pm 1.5^{ab}$
6 to 10	18	$9.79\pm0.26^{ab}$	$14.47 \pm 1.69^{b}$	$8.58 \pm 0.59^{b}$	$24.94 \pm 1.35^{c}$
11 to 15	15	$9.33\pm0.11^{b}$	$15.27 \pm 2.98^{ab}$	$9.17\pm1.66^{ab}$	$20.66\pm2.55^{abc}$
16 to 20	9	$10.41\pm0.31^a$	$6.10\pm0.77^a$	$4.68\pm0.44^a$	$11.54 \pm 1.12^{a}$
21+	39	$9.50\pm0.11^{b}$	$15.19 \pm 1.69^{b}$	$7.44 \pm 0.69^{ab}$	$23.21 \pm 1.94^{bc}$

Age; years since last reseeding event, C:N ratio of the fine fraction, %SC; proportion of fine fraction in a sample (%) by mass, Fine fraction organic carbon (% of SOC<sub>total</sub>); relative proportion of measured fine fraction OC of the total SOC content of the bulk soil.

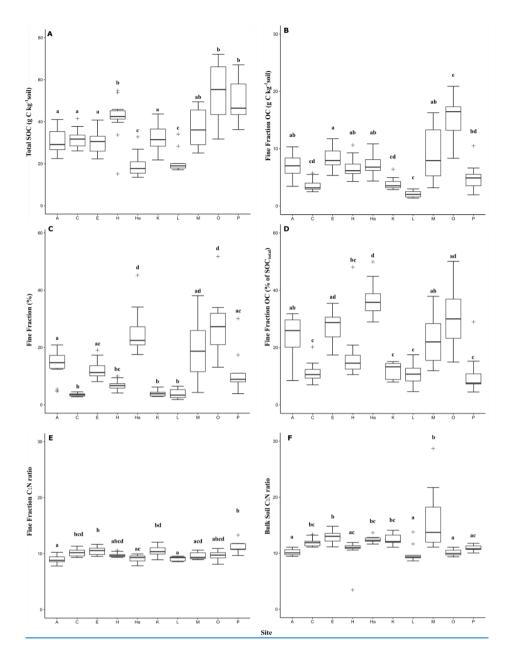


Figure 1. Measured total SOC (g C kg<sup>-1</sup>soil) (A) total fine fraction organic carbon (g C kg<sup>-1</sup>soil) (B), mass proportion of fine fraction ( $< 20 \,\mu m$ , %) (C), relative proportion of measured fine fraction organic carbon of the total SOC content of the bulk soil (D), and fine fraction C:N ratio (E) and bulk soil C:N ratio (F), for each of the grassland sites; Aberyswyth (A), Crichton (C), Easter Bush (E), Hillsborough (H), Harpenden (Ha), Kirkton (K), Llangorse (L), Myerscough (M), Overton (O) and Plumpton (P). Boxes represent the  $25^{th}$  and  $75^{th}$  percentile, with lines showing the median value. Whiskers show the lowest and highest values with outliers indicates as crosses (> 1.5 times the interquartile range). Lettering indicates significant differences between soils (P < 0.05).



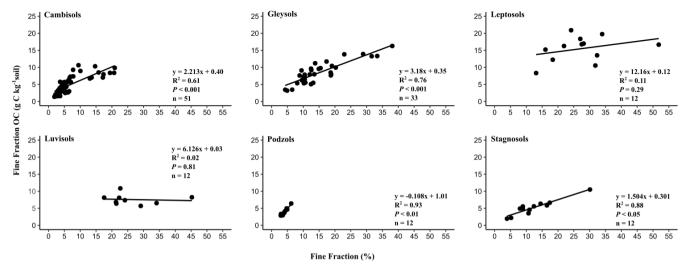
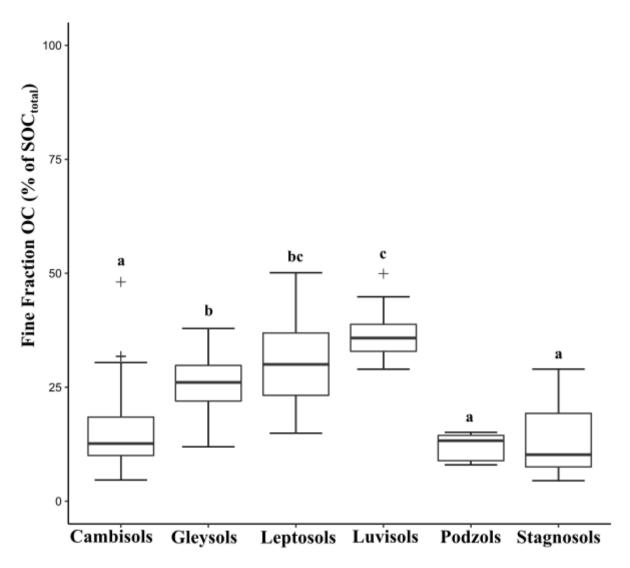


Figure 2. Relationships between mass proportion of the fine fraction (%) and fine fraction organic carbon (g  $C \ kg^{-1} \ soil$ ) in the soil types used in this study.



# World Reference Base Soil Type

Figure 3. Relative proportion of measured measured fine fraction organic carbon of the total SOC content of the bulk soil for the different soil types used in this study. Lettering indicates significant differences at P < 0.05.

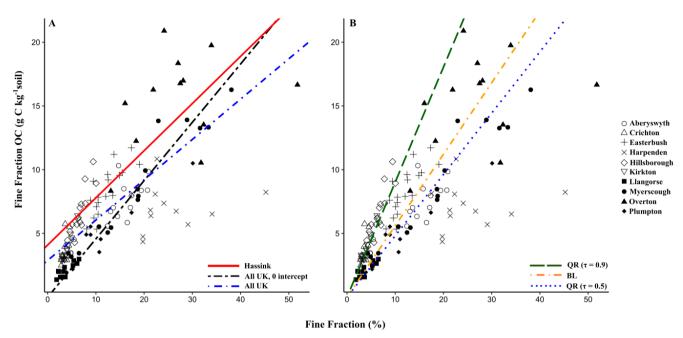


Figure 4. Measured fine fraction organic carbon (g C kg<sup>-1</sup> soil) in relation to mass proportion of fine fraction of a soil sample (%).

Line of best fit represent (A) linear regression method of Hassink, (1997) and data from this study, and (B) boundary line (BL) using 15% intervals, and quantile regression analysis (QR) at 90<sup>th</sup> and 50<sup>th</sup> percentiles.

# Appendix A

Table A1. Bulk soil properties for each UK site. Values are means of ten replicates in each field,  $\pm$  one standard error of the mean. Except Harpenden where values are means of five replicates per field. Lettering indicates values which are significantly different, within a site (P < 0.05).

	Age				С	C stock	N stock
Site	(year	$BD^a$	pН	C:N	(g C kg <sup>-1</sup> soil)	(t C ha <sup>-1</sup> )	(t N ha <sup>-1</sup> )
	s)				(g C kg Soil)	(сспа)	(t IV IIa )
Aberyswyth	2	1 ± 0.01 <sup>a</sup>	$5.20 \pm 0.05^{a}$	$9.70 \pm 0.05^{b}$	$26.95 \pm 0.63^{b}$	$73.61 \pm 1.73^{b}$	$7.59 \pm 0.16^{b}$
	6	$0.98\pm0.04^a$	$4.70\pm0.04^{bc}$	$9.68\pm0.08^b$	$26.7\pm0.82^b$	$73.86 \pm 2.28^{b}$	$7.62\pm0.18^b$
	11	$0.82\pm0.05^{b}$	$5.12\pm0.06^a$	$10.46\pm0.09^a$	$29.72 \pm 0.83^{b}$	$76.97 \pm 2.15^{b}$	$7.36\pm0.20^b$
	31	$0.74 \pm 0.05^{b}$	$4.99\pm0.09^{ab}$	$10.54 \pm 0.22^{a}$	$29.4 \pm 1.63^{b}$	$74.23 \pm 4.12^{b}$	$7.01\pm0.31^{b}$
	33	$0.69\pm0.03^{b}$	$4.18\pm0.02^{c}$	$10.59 \pm 0.10^{a}$	$38.19 \pm 1.97^{b}$	$95.67 \pm 4.92^{a}$	$9.01\pm0.40^a$
Crichton	1	$0.92 \pm 0.03$	$5.14 \pm 0.03^{ab}$	$12.19\pm0.08^{ab}$	$34.66\pm0.66^a$	$82.40 \pm 1.56$	$6.76\pm0.11^b$
	3	$0.99 \pm 0.07$	$5.65\pm0.06^b$	$11.73 \pm 0.11^{bc}$	$29.94\pm1.37^{ab}$	$74.63 \pm 3.41$	$6.36\pm0.26^b$
	15	$0.93 \pm 0.05$	$4.77\pm0.04^{ac}$	$9.90\pm0.88^{c}$	$30.85 \pm 3.06^b$	$79.73 \pm 7.91$	$7.98 \pm 0.23^a$
	20	$0.93 \pm 0.04$	$4.54\pm0.03^{c}$	$13.21\pm0.14^a$	$27.26\pm0.87^b$	$66.62 \pm 2.11$	$5.04\pm0.15^{c}$
Easter Bush	3	$1.02\pm0.04^{abc}$	$5.45\pm0.06~^{ab}$	$13.03 \pm 0.11^{bc}$	$32.46 \pm 1.29^a$	$93.52 \pm 3.72^a$	$7.17\pm0.26^a$
	5	$1.19\pm0.03^a$	$5.44 \pm 0.06^{ab}$	$12.84 \pm 0.21^{bc}$	$26.41 \pm 0.54^{b}$	$74.45 \pm 1.52^b$	$5.80\pm0.10^{bc}$
	5	$0.84\pm0.06^{c}$	$5.67\pm0.04^a$	$11.74\pm0.17^{d}$	$27.50\pm1.0^b$	$58.15 \pm 2.12^{c}$	$4.94\pm0.14^{d}$
	6	$0.96\pm0.05^{bc}$	$5.32\pm0.06^b$	$12.45 \pm 0.13^{c}$	$30.46 \pm 1.93^{ab}$	$71.16 \pm 4.50^{bc}$	$5.72\pm0.36^{cd}$
	6	$1.12\pm0.05^{ab}$	$5.81\pm0.20^{\rm a}$	$14.15\pm0.13^{\rm a}$	$28.95 \pm 0.99^{ab}$	$75.69 \pm 2.59^{b}$	$5.35\pm0.17^{cd}$
	8	$1.12\pm0.03^{ab}$	$4.99 \pm 0.04^{c}$	$13.43 \pm 0.11^{b}$	$33.03\pm0.50^a$	$89.43 \pm 1.34^a$	$6.66\pm0.10^{ab}$
Harpenden	22	$1.37 \pm 0.07$	$7.37\pm0.04^{\rm a}$	$12.09 \pm 0.2$	$16.06 \pm 0.59^{c}$	$25.37 \pm 0.93^{c}$	$3.3\pm0.12^{c}$
	68	$1.12 \pm 0.08$	$5.85\pm0.12^{ab}$	$12.34\pm0.08$	$19.8\pm0.63^{b}$	$50.49 \pm 1.59^{b}$	$4.06\pm0.13^b$
	179	$1.09 \pm 0.14$	$5.63\pm0.06^b$	$12.8 \pm 0.26$	$28.7\pm1.47^{\rm a}$	$72.98 \pm 3.74^{\rm a}$	$5.89\pm0.30^a$
Hillsborough	1	$1.79 \pm 0.10$	$6.31\pm0.07^a$	$11.25 \pm 0.12^{ab}$	$46.68 \pm 2.04$	$120.16 \pm 5.26^{ab}$	$10.69\pm0.51^{ab}$
	7	$1.88 \pm 0.08$	$5.10\pm0.04^{\rm b}$	$11.46\pm0.11^{b}$	$42.85 \pm 1.52$	$108.86 \pm 3.87^{b}$	$9.51\pm0.34^{bc}$
	16	$1.79 \pm 0.05$	$5.33\pm0.08^b$	$10.87 \pm 0.06^{c}$	$42.36 \pm 1.98$	$111.63 \pm 5.21^{ab}$	$10.27 \pm 0.47^{ab}$
	23	$1.75\pm0.05$	$4.76\pm0.03^c$	$11.33\pm0.09^{ab}$	$46.44 \pm 1.78$	$125.43 \pm 4.82^{a}$	$11.08\pm0.45^a$
	37	$1.69 \pm 0.06$	$5.13\pm0.06^b$	$10.34\pm0.77^{ac}$	$40.90 \pm 3.10$	$86.04 \pm 6.52^{c}$	$8.38 \pm 0.24^{c}$
Kirkton	1	$0.9 \pm 0.04$	$4.78\pm0.04^c$	$12.13 \pm 0.11^{c}$	$27.90 \pm 0.81^{c}$	$82.03 \pm 2.39^{b}$	$6.77 \pm 0.22$
	3	$0.95 \pm 0.04$	$5.49\pm0.06^a$	$12.61 \pm 0.15^{b}$	$36.67 \pm 1.56^a$	$98.19 \pm 4.17^{ab}$	$7.79 \pm 0.31$

	5	$0.83 \pm 0.06$	$5.15 \pm 0.03^{b}$	$13.56 \pm 0.08^{a}$	$34.83 \pm 1.84^{ab}$	$103.03 \pm 5.45^{a}$	$7.59 \pm 0.38$
	35	$0.97 \pm 0.06$	$4.72\pm0.07^{c}$	$11.67 \pm 0.13^{d}$	$30.51 \pm 1.48^{bc}$	$90.50 \pm 4.38^{ab}$	$7.72 \pm 0.32$
Llangorse	2.5	$1.01\pm0.04$	$5.14\pm0.08^c$	$9.21 \pm 0.09$	$17.83 \pm 0.42$	$49.75 \pm 1.18$	$5.40\pm0.11^{ab}$
	5	$0.93 \pm 0.04$	$5.44\pm0.03^b$	$9.40 \pm 0.07$	$18.60\pm0.45$	$50.80\pm1.22$	$5.40\pm0.10^b$
	15	$0.94 \pm 0.06$	$5.68 \pm 0.03^a$	$9.36 \pm 0.17$	$19.42\pm0.38$	$53.70 \pm 1.06$	$5.74\pm0.12^{ab}$
	25	$1.06 \pm 0.03$	$5.54 \pm 0.07^{ab}$	$9.16 \pm 0.87$	$19.73 \pm 2.52$	$55.10 \pm 7.05$	$6.18\pm0.34^a$
Myerscough	2	$1.22\pm0.02^{ab}$	$4.97\pm0.05^b$	$13.58 \pm 0.24^{bc}$	$27.47\pm0.65^c$	$82.25 \pm 1.96^{c}$	$6.07 \pm 0.15^{bc}$
	6	$1.10\pm0.04^{b}$	$5.59\pm0.05^a$	$11.79 \pm 0.76^{c}$	$41.44 \pm 2.73^a$	$124.05 \pm 8.17^{a}$	$10.56\pm0.28^a$
	13	$0.93\pm0.05^{b}$	$5.00\pm0.20^b$	$13.12 \pm 0.43^{\circ}$	$44.82 \pm 2.34^a$	$134.45 \pm 7.01^a$	$10.30\pm0.71^a$
	34	$1.29\pm0.02^{\rm a}$	$5.99 \pm 0.13a$	$17.20\pm1.12^{ab}$	$37.58 \pm 1.45^{ab}$	$112.46 \pm 4.36^{ab}$	$6.71\pm0.30^b$
	48.4	$1.44\pm0.06^{\rm a}$	$5.77\pm0.02^a$	$22.10\pm1.46^{\rm a}$	$29.86 \pm 1.96^{bc}$	$88.97 \pm 5.85^{bc}$	$4.03\pm0.08^c$
Overton	3	$0.98\pm0.09^a$	$6.58\pm0.12^b$	$9.76\pm0.05^{b}$	$32.77 \pm 0.84^{c}$	$83.02 \pm 2.13^{b}$	$8.51\pm0.23^b$
	12	$0.38\pm0.03^{b}$	$6.83\pm0.03^b$	$10.18\pm0.12^{ab}$	$70.18 \pm 1.92^a$	$81.20 \pm 2.23^{b}$	$7.99\pm0.23^b$
	22	$0.71\pm0.07^{ab}$	$7.36\pm0.04^a$	$10.68 \pm 0.39^{a}$	$59.88 \pm 3.86^{b}$	$132.75 \pm 8.56^a$	$12.33\pm0.39^a$
	50	$1.74\pm0.9^a$	$4.63\pm0.08^{c}$	$10.14\pm0.14^{ab}$	$51.18\pm2.84^b$	$153.08 \pm 8.50^{a}$	$15.08\pm0.80^a$
Plumpton	1	$0.99\pm0.02^{\rm a}$	$6.34\pm0.08^b$	$10.85\pm0.08ab$	$40.92 \pm 1.21^{b}$	$122.21 \pm 3.61^{b}$	$11.26\pm0.28^b$
	5	$1.08\pm0.03a$	$7.15 \pm 0.06a$	$11.27\pm0.41a$	$45.55 \pm 0.61^b$	$132.09 \pm 1.77^{b}$	$11.87\pm0.48^b$
	20	$0.72 \pm 0.04b$	$5.38 \pm 0.21c$	$10.54 \pm 0.17$ b	$58.08 \pm 2.36^{a}$	$163.23 \pm 6.62^{a}$	$15.47 \pm 0.56^{a}$

<sup>&</sup>lt;sup>a</sup>Bulk density (BD), means and SEM of six samples, except Harpenden with two samples per field, corrected for stones.

Location	Age (years)	%N	%C	C:N	%Fine Fraction	Organic Carbon (g C kg <sup>-1</sup> bulk soil) <sup>a</sup>	Organic Carbon (% of SOCtotal)
Aberyswyth	2	$0.48 \pm 0.01^{b}$	$4.16 \pm 0.08$	$8.62 \pm 0.08^{ab}$	$19.08 \pm 1.04^{a}$	$7.93 \pm 0.45^{a}$	$0.30 = 0.01^{a}$
	6	$0.52\pm0.04^{b}$	$4.14 \pm 0.30$	$8.05 \pm 0.17^{b}$	$14.47 \pm 1.18^{ab}$	$5.92 \pm 0.22^{ab}$	$0.21\pm0.02^{ab}$
	11	$0.55\pm0.03^{\rm b}$	$4.89 \pm 0.25$	$8.86 \pm 0.06^{ab}$	$18.02 \pm 1.51^{ab}$	$8.77 \pm 0.56^{a}$	$0.29 \pm 0.02^{ab}$
	31	$0.61\pm0.05^{ab}$	$5.78 \pm 0.62$	$9.51 \pm 0.30^{ab}$	$13.78 \pm 0.49^{b}$	$8.02 \pm 1.15^{a}$	$0.27\pm0.02^{ab}$
	33	$0.76\pm0.03^a$	$7.57 \pm 0.37$	$9.96 \pm 0.23^{a}$	$5.02 \pm 0.22^{c}$	$3.81 \pm 0.36^{b}$	$0.10\pm0.01^{b}$
Crichton	1	$1.01\pm0.06$	$10.53 \pm 0.83$	$10.40 \pm 0.23^{ab}$	$4.00 \pm 0.45$	$4.24 \pm 0.69$	$0.12 \pm 0.02$
	3	$1.15\pm0.27$	$11.17 \pm 2.28$	$9.84 \pm 0.35^{b}$	$3.28 \pm 0.23$	$3.75 \pm 1.00$	$0.13 \pm 0.04$
	15	$1.02\pm0.12$	$9.76 \pm 1.20$	$9.52 \pm 0.12^{b}$	$3.52 \pm 0.26$	$3.38 \pm 0.31$	$0.10 \pm 0.02$
	20	$0.82 \pm 0.05$	$9.07 \pm 0.72$	$11.03 \pm 0.24^{a}$	$3.37 \pm 0.3$	$3.01 \pm 0.09$	$0.11 \pm 0.01$
Easter Bush	3	$0.65 \pm 0.04$	$7.15 \pm 0.50$	$11.00 \pm 0.13^{ab}$	$14.38 \pm 1.56^{ab}$	$10.27 \pm 1.19^{a}$	$0.30 \pm 0.01$
	5	$0.65 \pm 0.04$	$6.91 \pm 0.50$	$10.57 \pm 0.06^{bc}$	$12.17\pm0.9^{ab}$	$8.34 \pm 0.43^{ab}$	$0.32 \pm 0.02$
	5	$0.67 \pm 0.02$	$6.62 \pm 0.23$	$9.83 \pm 0.07^{c}$	$9.55 \pm 0.73^{b}$	$6.32 \pm 0.51^{b}$	$0.23 \pm 0.03$
	6	$0.68 \pm 0.03$	$7.81 \pm 0.43$	$9.85 \pm 0.24^{c}$	$9.75 \pm 0.23^{b}$	$6.88 \pm 1.13^{ab}$	$0.23 \pm 0.01$
	6	$0.72 \pm 0.12$	$7.11 \pm 1.31$	$11.43 \pm 0.16^{a}$	$10.58 \pm 1.04^{b}$	$8.22 \pm 0.70^{ab}$	$0.27 \pm 0.02$
	8	$0.59 \pm 0.04$	$6.07 \pm 0.30$	$10.35 \pm 0.28^{bc}$	$16.47 \pm 1.3^{a}$	$9.91 \pm 0.26^{ab}$	$0.30 \pm 0.01$
Harpenden	22	$0.23\pm0.01^{b}$	$1.90 \pm 0.04^{c}$	$8.20 \pm 0.26^{b}$	$36.15 \pm 4.77^{a}$	$6.82 \pm 0.75^{ab}$	$0.42 \pm 0.04$
	68	$0.32\pm0.01^{b}$	$3.08 \pm 0.06^{b}$	$9.54 \pm 0.21^{a}$	$22.27 \pm 0.92^{b}$	$6.86 \pm 0.28^{ab}$	$0.36 \pm 0.02$
	179	$0.46\pm0.03^a$	$4.35 \pm 0.36^{a}$	$9.54 \pm 0.12^{a}$	$20.83 \pm 1.64^{b}$	$9.02 \pm 0.91^{a}$	$0.32 \pm 0.01$
Hillsborough	1	$0.90 \pm 0.08$	$8.97 \pm 1.14$	$9.91 \pm 0.34$	$7.37 \pm 0.12$	$6.86 \pm 1.93$	$0.14 \pm 0.03$
	7	$1.04 \pm 0.06$	$10.23 \pm 0.91$	$9.80 \pm 0.31$	$8.05 \pm 0.08$	$8.15 \pm 0.96$	$0.17 \pm 0.02$
	16	$0.99 \pm 0.04$	$9.36 \pm 0.32$	$9.46 \pm 0.03$	$6.33 \pm 0.19$	$5.92 \pm 0.13$	$0.15 \pm 0.01$
	23	$1.15 \pm 0.01$	$11.11 \pm 0.13$	$9.70 \pm 0.18$	$4.58 \pm 0.27$	$5.10 \pm 0.36$	$0.12 \pm 0.01$
	37	$1.04 \pm 0.04$	$10.12 \pm 0.35$	$9.76 \pm 0.04$	$7.15 \pm 0.33$	$7.22 \pm 0.10$	$0.27 \pm 0.11$
Kirkton	1	$0.91 \pm 0.03$	$9.27 \pm 0.12^{b}$	$10.15 \pm 0.24^{b}$	$3.90 \pm 0.1$	$3.62 \pm 0.13$	$0.14 \pm 0.01$
	3	$1.01 \pm 0.04$	$10.63 \pm 0.33^{a}$	$10.56\pm0.27^{ab}$	$3.02 \pm 0.03$	$3.20 \pm 0.07$	$0.08 \pm 0.00$
	5	$0.88 \pm 0.03$	$10.23 \pm 0.16^{ab}$	$11.66 \pm 0.31^{a}$	$4.62 \pm 0.95$	$4.75 \pm 1.03$	$0.13 \pm 0.02$

	35	$0.96 \pm 0.03$	$9.22 \pm 0.40^{b}$	$9.64 \pm 0.39^{b}$	$4.23 \pm 0.42$	$3.93 \pm 0.51$	$0.14 \pm 0.00$
Llangorse	2.5	$0.51\pm0.03^{b}$	$4.76\pm0.29^b$	$9.36 \pm 0.07$	$6.00\pm0.32^{\rm a}$	$2.83\pm0.10^a$	$0.16\pm0.01^a$
	5	$0.88 \pm 0.08^a$	$8.29\pm0.80^a$	$9.43 \pm 0.07$	$2.65 \pm 0.43^{b}$	$2.13\pm0.11^{ab}$	$0.11\pm0.01^{ab}$
	15	$0.67\pm0.10^{ab}$	$6.06\pm0.78^{ab}$	$9.11 \pm 0.24$	$3.23 \pm 1.03^{b}$	$1.81 \pm 0.34^{b}$	$0.09\pm0.02^b$
	25	$0.62\pm0.06^{ab}$	$5.32 \pm 0.54^{b}$	$8.60 \pm 0.05$	$3.27\pm0.22^{b}$	$1.72\pm0.14^{b}$	$0.07\pm0.01^{b}$
Myerscough	2	$0.63 \pm 0.08$	$6.60 \pm 0.76$	$10.43 \pm 0.12$	$5.23 \pm 0.66^{c}$	$3.35 \pm 0.09^{c}$	$0.12\pm0.00^b$
	6	$0.49 \pm 0.03$	$4.57\pm0.29$	$9.31 \pm 0.23$	$27.50\pm3.85^a$	$12.39 \pm 1.24^{a}$	$0.31\pm0.04^a$
	13	$0.50 \pm 0.05$	$4.83 \pm 0.60$	$9.52 \pm 0.32$	$30.88 \pm 4.39^a$	$14.45 \pm 0.92^{a}$	$0.30\pm0.02^a$
	34	$0.47 \pm 0.01$	$4.28 \pm 0.13$	$9.12 \pm 0.07$	$18.72 \pm 0.04^{ab}$	$8.02\pm0.24^b$	$0.21\pm0.02^{ab}$
	48.4	$0.47 \pm 0.02$	$4.47 \pm 0.36$	$9.58 \pm 0.36$	$12.08 \pm 0.74^{bc}$	$5.35\pm0.14^{bc}$	$0.17\pm0.02^b$
Overton	3	$0.42\pm0.03^{c}$	$3.57 \pm 0.31^{c}$	$8.45\pm0.18^{b}$	$38.65 \pm 6.58$	$13.57 \pm 1.77$	$0.41 \pm 0.05$
	12	$0.88\pm0.05^{\rm a}$	$8.52 \pm 0.59^{a}$	$9.64 \pm 0.11^{a}$	$20.70 \pm 2.41$	$17.45 \pm 1.75$	$0.25 \pm 0.02$
	22	$0.61\pm0.02^{b}$	$6.36\pm0.18^b$	$10.36\pm0.15^{\mathrm{a}}$	$19.85 \pm 4.39$	$12.52 \pm 2.50$	$0.23 \pm 0.07$
	50	$0.63\pm0.05^{\text{b}}$	$6.23 \pm 0.29^{b}$	$10.04 \pm 0.45^{a}$	$29.50 \pm 2.23$	$18.29 \pm 0.86$	$0.34 \pm 0.04$
Plumpton	1	$0.35 \pm 0.02^{b}$	$3.81\pm0.18^{b}$	$10.87 \pm 0.09$	$19.50 \pm 5.61$	$7.23 \pm 1.74$	$0.18 \pm 0.05$
	5	$0.36\pm0.04^b$	$4.19\pm0.49^b$	$11.76 \pm 0.91$	$6.60 \pm 2.08$	$2.56 \pm 0.49$	$0.06 \pm 0.01$
	20	$0.56\pm0.02^a$	$5.96 \pm 0.23^{a}$	$10.75 \pm 0.62$	$8.60 \pm 0.3$	$5.11 \pm 0.21$	$0.08 \pm 0.01$

<sup>%</sup>Fine Fraction; mass proportion of fine fraction in a sample (%).

<sup>&</sup>lt;sup>a</sup> MAOC (g C kg<sup>-1</sup> bulk soil) accounts for the proportion of fine fraction per kilogram of bulk soil.

Table A3. Linear regression coefficients for the estimation of maximum fine fraction organic carbon (g C kg $^{-1}$  soil). Lettering indicates slopes which are significantly different (P < 0.05).

Site	Slope	P slope	Intercept	P intercept	RMSE	n	$\mathbb{R}^2$	
Site	(± 1 SEM)	7 slope	(± 1 SEM)	1 miercepi	KWISE	11		
Aberyswyth	$0.33 \pm 0.059^{bc}$	***	$2.28 \pm 0.892$	*	1.11	15	0.70	
Crichton	$1.14 \pm 0.470^{abcd}$	*	$-0.44 \pm 1.684$	Ns	0.79	12	0.37	
Easter Bush	$0.49 \pm 0.094^{d}$	***	$2.33 \pm 1.172$	Ns	1.10	18	0.63	
Harpenden	$-0.02 \pm 0.07^{a}$	Ns	$8.01 \pm 1.837$	**	1.42	9	0.01	
Hillsborough	$0.97 \pm 0.148^{d}$	***	$0.16 \pm 1.02$	Ns	0.84	15	0.77	
Kirkton	$1.01 \pm 0.088^{abcd}$	***	$-0.11 \pm 0.357$	Ns	0.26	12	0.93	
Llangorse	$0.29 \pm 0.055^{\rm abc}$	***	$1.03 \pm 0.225$	***	0.27	12	0.73	
Mysercough	$0.40 \pm 0.031^{\rm bcd}$	***	$1.07 \pm 0.669$	Ns	1.14	15	0.93	
Overton	$0.12 \pm 0.109^{cd}$	Ns	$12.16 \pm 3.142$	**	3.35	12	0.11	
Plumpton	$0.30 \pm 0.042^{ab}$	***	$1.45 \pm 0.573$	*	0.82	9	0.88	

585 RMSE: Root mean square error.

Level of significance: \* P < 0.05, \*\* P < 0.01, \*\*\*P < 0.001

Table A4. Linear regression coefficients for the estimation of maximum fine fraction organic carbon (g C kg $^{-1}$  soil) with a forced zero intercept. Lettering indicates slopes, that are significantly different (P < 0.05).

Site	Slope (± 1 SEM)	P	RMSE	n	R <sup>2</sup>
Aberyswyth	$0.47 \pm 0.024^{bc}$	***	1.357	15	0.96
Crichton	$1.02 \pm 0.067^{cdef}$	***	0.796	12	0.95
Easter Bush	$0.67 \pm 0.024^{e}$	***	1.231	18	0.98
Harpenden	$0.26\pm0.035^a$	***	2.739	9	0.87
Hillsborough	$0.99 \pm 0.033^{\rm f}$	***	0.842	15	0.99
Kirkton	$0.98 \pm 0.0197^{\mathrm{def}}$	***	0.265	12	0.99
Llangorse	$0.52 \pm 0.035^{abcdef}$	***	0.474	12	0.95
Mysercough	$0.45 \pm 0.016^b$	***	1.255	15	0.98
Overton	$0.52\pm0.055^{bcd}$	***	5.297	12	0.89
Plumpton	$0.39\pm0.030^{ab}$	***	1.141	9	0.96

RMSE: Root mean square error.

595

600

Level of significance: \* P < 0.05, \*\* P < 0.01, \*\*\*P < 0.001

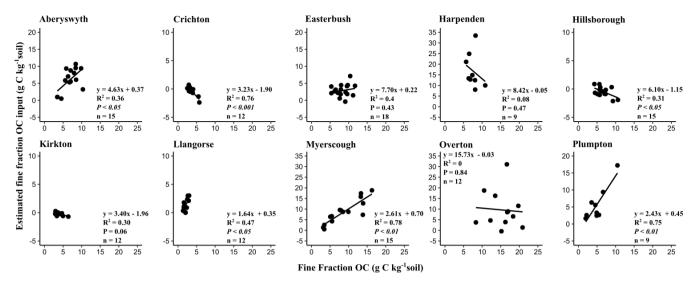


Figure A1. Estimated fine fraction OC input (g C kg<sup>-1</sup> soil) compared to measured fine fraction OC (g C kg<sup>-1</sup> soil) in each of the sites studied. The estimated fine fraction OC input (g C kg<sup>-1</sup> soil) was calculated by subtracting the maximum fine fraction OC (g C kg<sup>-1</sup> soil) from the current fine fraction OC (g C kg<sup>-1</sup> soil). The maximum fine fraction OC (g C kg<sup>-1</sup> soil) was estimated using the quantile regression equation ( $\tau = 0.90$ ), where, maximum fine fraction OC = 0.92 multiplied by the mass proportion fine fraction (%).

Figure A1. Estimated fine fraction OC input (g C kg<sup>-1</sup>soil) compared to measured fine fraction OC (g C kg<sup>-1</sup>soil) in each of the sites studied. The estimated OC input was predicted using quantile regression at the 90<sup>th</sup> percentile.