

## Predicting the in situ rate constant of soil carbon mineralisation from laboratory-based measurements in tropical soils under contrasting tillage management systems

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- Postprint version 1
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- Predicting the in situ rate constant of soil carbon mineralisation from laboratory-3
- 4 based measurements in tropical soils under contrasting tillagemanagement
- 5 systems
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- 12

#### 13 Abstract

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Predicting soil organic carbon (SOC) mineralisation in the field is crucial to 15 16 assessing the impact of changes in land use and tillage practices. Although soil 17 data from long-term field experiments are essential to calibrate model parameters 18 for SOC mineralisation, these are frequently lacking regarding some parts of the 19 tropics. The aim of this study was to develop and test an indicator for the in situ 20 SOC mineralisation rate constant using data obtained during soil incubation in the 21 laboratory. Sixty-nine plots corresponding to four soil types (andosol, nitisol, 22 ferralsol and vertisol) under six crops (banana, sugarcane, vegetables, yam, melon 23 and pineapple) involved in 12 tropical cropping systems were analysed. Soil 24 sampling and laboratory incubation procedure were designed to reflect the long-25 term impact of cropping systems on SOC dynamics. Data from long-term 26 experiments were obtained from a previous study performed on the same cropping 27 systems in Guadeloupe (Caribbean). While SOC contents and stocks were only 28 controlled by soil type, mineralised SOC in the laboratory (C<sub>min</sub>) was con- trolled to 29 the same degree by both soil type and crop. Moreover, the C<sub>min</sub>/SOC ratio, which is an indicator of the more active SOC fraction, was only controlled by the crop. 30 31 Although C<sub>min</sub> and C<sub>min</sub>/SOC were not affected by recent C inputs from crop 32 residues, they were negatively correlated with the rate constant of SOC mineralisation observed in the field (k<sub>field</sub>). These results indicate that differences 33 34 between the cropping systems (i.e. perennial vs. annual crops) concerning the 35 management of soil tillage markedly affected depletion of the mineralisable SOC 36 fraction. Overall, the results showed that C<sub>min</sub> measured under regulated laboratory

conditions could be helpful to estimate k<sub>field</sub>. This is important in tropical regions in
 order to anticipate the effects of changes in land use and tillage management

- 39 before changes in SOC stocks become noticeable in the field.
- 40
- 41 Keywords: Caribbean, Cropping system, Indicator, Long-term effect, Soil
- 42 incubation, Soil type
- 43
- 44

#### 45 **1. Introduction**

46

47 The accurate prediction of soil organic carbon (SOC) mineralisation under field conditions is of crucial importance from both the agronomic and environmental points of 48 49 view (Oelbermann et al., 2017). In the long-term (e.g. decade or century), predicting 50 SOC mineralisation is necessary to assess the impact of changes in land use, farming 51 practices and tillage management on soil quality and C emissions (Lal, 2004). Although 52 some authors used soil incubations to estimate the rate constants to be included in 53 models devoted to conceptualising and de- scribing SOC mineralisation in the field 54 (Brisson et al., 2003), an alternative procedure involves the statistical fitting of data 55 obtained from diachronic measurements of SOC content in the field (e.g. Andriulo et 56 al., 1999; Sierra et al., 2015). The advantage of this method over laboratory incubations 57 is that it integrates all aspects of managing the plant-soil system over many years, including the impacts on SOC turnover of crop residue management and soil tillage 58 59 (Cerri et al., 2007). However, this method requires a relatively important soil database 60 including information on the farming practices that affect SOC turnover (e.g. rotations, amount and quality of crop residues, management of organic amendments, intensity of 61 62 soil tillage), which may be unavailable, particularly in tropical regions (Sierra et al., 63 2017).

Although changes in agricultural management may influence the mineralisation 64 65 rate constant, the ability of the models of SOC dynamics to assess the impact of such 66 changes depends on including of farming practices in the model framework. For example, only a few models of SOC dynamics have explicitly considered the increase 67 in SOC decomposition following soil tillage (Bortolon et al., 2009), which is a major 68 process that affects the SOC balance (Martínez et al., 2017). In the case of the 69 70 Century model adapted to the conditions of Southern Brazil, soil tillage is represented 71 by a cultivation parameter which operates as a multiplier to enhance the mineralisation 72 rate constant of SOC fractions during the two months that follow tillage operations

73 (Bortolon et al., 2011). However, it may be difficult to extrapolate this empirical approach based on local information about farming practices to other tropical regions such as the 74 75 Caribbean, where these practices may vary considerably as a function of farmer's 76 resources and the orientation of agricultural production; e.g. perennial crops such 77 as sugarcane and banana vs. annual food and vegetable crops (IFAD, 2014). For 78 example, Sierra et al. (2017) found that vegetable crop systems in Guadeloupe, an 79 archipelago in the Caribbean, could be represented by five farm types that applied 80 contrasting tillage systems on very different soil types, ranging from manual cultivation 81 to more labour-intensive strategies which include up to six tillage operations per year. 82 By contrast, soil tillage under perennial crops is only applied once every five or six 83 years, at the time of planting (Raphael et al., 2012).

Compared with complex models of SOC dynamics, simple models of the 84 85 annual SOC balance require minimal data inputs and few parameters (Saffih-Hdadi 86 and Mary, 2008) and are therefore well suited to situations with scarce agricultural data, such as in the Caribbean. This also concerns the lack of suitable model 87 parameters regarding most tropical food and vegetable crops (Nin-Pratt et al., 88 89 2011). Undoubtedly, such models may be less robust than process-based models, so 90 that their accuracy in assessing the impact on SOC dynamics of new cropping 91 systems or changes of farming practices needs to be tested (Coucheney et al., 92 2015). In this context, the use of soil indicators of the SOC mineralisation rate 93 constant calibrated from long-term field experiments may be helpful to predict SOC 94 changes under new agricultural scenarios. This is important in tropical regions in 95 order to anticipate the effects of changes in land use (e.g. conversion of food to 96 energy crops) and practices (e.g. manual vs. mechanical soil tillage) (FAO Subregional Office for the Caribbean, 2013), before changes in SOC stocks may be 97 98 noticed in the field.

99 In a previous study, Sierra et al. (2015) designed the MorGwanik model of 100 SOC dynamics adapted to the pedoclimatic conditions of the Caribbean in order to 101 assess SOC changes under export and diversified agriculture at the level of the 102 Agro-Ecological Region (AER). The aim of the present study was to evaluate the 103 ability of an indicator based on SOC mineralisation under laboratory conditions to 104 predict the mineralisation rate constants obtained by these authors from long-term field 105 studies, which involved twelve cropping systems under contrasting tillage 106 management. To achieve this, four conditions were imposed to the laboratory 107 experiment: i- only soils under monoculture were selected for the trial in order to 108 best reflect the impact of the management of each crop, ii- soil sampling from each

109	selected plot was carried out on sub plots that did not receive crop residues following
110	the harvest of the previous crop, in order to reduce their effect on the mineralisable
111	C fraction, iii- soil samples were neither dried nor disrupted to reduce the presence of
112	artefacts linked to sample handling during soil incubation (e.g. the Birch effect;
113	Jarvis et al., 2007) and, iv- relatively long-term soil incubation periods (i.e. 15 wk)
114	were applied to obtain a substantial amount of mineralised C so as to ensure the
115	reliability of the analysis. Under these conditions, we hypothesise that SOC
116	mineralisation during the laboratory experiment reflects the impact of the cropping
117	systems on SOC dynamics in the field. The goal was therefore to develop and to test an
118	indicator that could be used during future research concerning the impact of new
119	agricultural practices on SOC stocks in the soils of the Caribbean region. The study was
120	carried out in the Guadeloupe archipelago (Lesser Antilles) which within a small area
121	displays nearly every physical landscape and cropping system found in the
122	Caribbean.
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125	2. Materials and methods
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127	2.1. Study location
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129	The study was carried out in the Guadeloupe archipelago in the eastern
130	Caribbean. Guadeloupe is composed of two main islands (Basse- Terre: 848 km <sup>2</sup> and
131	Grande-Terre: 586 km <sup>2</sup> ) and several smaller islands. Only the soils of the two main
132	islands were analysed during this study. Although Grande-Terre is characterised by a
133	gently undulating surface, Basse-Terre Island is dominated by a volcanic mountain
134	chain oriented northwest to southeast with elongated hills with convex slopes in the
135	lowlands. We analysed four soil types (FAO, 2006):
136	i- Andosols located in the uplands of southern Basse-Terre, which had developed on
137	young ash deposits. These soils are acidic (Table 1) and characterised by high
138	allophanic clay contents (75%). The mean temperature in this region is 23.9 °C and the
139	mean annual rainfall is 3800 mm.
140	ii- Nitisols located in the lowlands of southern Basse-Terre, which had developed on old
141	ash deposits. These soils are acidic and rich in halloysite clay (70%). The mean
142	temperature is 25.0 °C and the mean annual rainfall is 2200 mm.
143	iii- Ferralsols located in the northern and eastern parts of Basse-Terre, which had
144	developed on old ash deposits. These soils are acidic and rich in kaolinite clay (70%)

•

and active aluminium and iron hydrous oxides. The mean temperature is 25.4 °C and
the mean annual rainfall is 2300 mm.

147 iv- Vertisols located in Grande-Terre, which had developed on coral reef limestone.

148 These soils are relatively alkaline and characterized by a high clay content (80%)

dominated by smectite. The mean temperature is 26.5 °C and the mean annual rainfallis 1100 mm.

In all of these regions there is a dry season from December to May; it is less
pronounced in the uplands of southern Basse-Terre (i.e. 45% of annual rainfall during
the dry season) and more marked in Grande- Terre (i.e. only 30% of annual rainfall
during the dry season).

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156 2.2. Management o the cropping systems

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In this section we present the principal characteristics of the management of the
 cropping systems analysed in this study, mainly those that directly affect SOC
 dynamics.

Sugarcane is the dominant crop in Guadeloupe and it is cultivated on ferralsols and vertisols as a monoculture or in rotation with tuber and vegetable crops. When sugarcane is cultivated in rotation this is under a 5–6-yr cycle. The growth season is 12 months and harvesting is mechanical, where residues (tops and leaves) are shredded before being returned to the soil surface. After harvesting, the remaining stubble grows new shoots (ratoon crop). For this reason, soil tillage is only applied before planting in the first year of the cycle. Most sugar production is exported to European markets.

Banana is mainly cultivated on andosols, ferralsols and nitisols, as a monoculture or in rotation with vegetable crops with a 5–6-yr cycle. Harvesting is manual and each plant is harvested separately. Residues (leaves and stems) are cut and placed on the soil surface. After harvesting, a sucker from a lateral shoot of the mother plant grows a new stem. Soil tillage is therefore only applied before planting in the first year of the cycle. Most banana production is exported to European markets.

Most diversification crops (i.e. vegetables, yam, pineapple, melon) are cultivated in rotation with export crops (i.e. sugarcane and banana), but vegetable crops and pineapple are also cultivated as monocultures. Vegetables are cultivated on andosols, nitisols and vertisols, yam on ferralsols and vertisols, melon is only cultivated on vertisols, and pineapple is only grown on ferralsols. All these crops are managed as annual crops and their cycle within the rotation varies from 3 yr for yam to 6 yr for pineapple. Soil tillage is quite intensive in these cropping systems, with 4–6 tillage operations per year. Yam is cultivated on mechanically-prepared ridges. Except for
yam, harvesting is manual and the residues (leaves and stems) are placed on the soil
surface or buried at a depth of 0.1–0.2 m. Pineapple residues are shredded before
being returned to the soil. Yam is harvested mechanically and the residues (leaves and
stems) are partially buried during ridge removal. Most production from diversification
crops supply local markets, except for melon which is mainly exported to Europe.

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#### 188 2.3. Laboratory experiment

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190 Sixty-nine plots representing 12 soil x crop situations were selected using the 191 database developed for the TropEmis Project carried out in Guadeloupe (Sierra et al., 192 2015). The soil x crop situations and mean values of soil properties are presented in 193 Table 1. This database contains information on soils, crop management and farm 194 characteristics that cover all AERs of the archipelago. Only plots under monoculture 195 were selected for the sugarcane, banana and vegetable crops. For other crops, the 196 length of the cycle at the time of soil sampling was 3 yr for yam, 6 yr for pineapple and 197 5 yr for melon. To reduce the effect of recent crop residues, soil samples for each selected plot were collected from five sub-plots of 10 m<sup>2</sup> that had not received crop 198 199 residues following the last harvest. Soil samples were collected after harvest in the last 200 year of the cycle of each crop, from January 2015 to March 2015. For all the selected 201 plots, soil sampling was performed after a rainfall, thus raising soil moisture to a level slightly higher than field capacity. This was designed to prevent the effect of drying and 202 203 rewetting the soil sample on the burst of C respiration following the start of the 204 laboratory experiment (i.e. the Birch effect; Wade et al., 2016). Soil samples were 205 taken from the upper 0.25 m soil layer on a systematic grid using a 0.06-m diameter 206 auger and collecting three sub- samples per sub-plot. Although it is well known that the 207 contribution of deeper layers to SOC mineralization can be important (e.g. Zhu et al., 208 2017), in this study we focused on the topsoil because SOC in that layer is very 209 sensitive to changes in land use and tillage management in the tropical soils 210 analysed here (Sierra et al., 2015).

- Soil organic C (determined with a C analyser) and organic N (Kjeldahl
  method) analyses were performed on a composite sample at the SADEF Soil
  Testing Laboratory in France. The stock of SOC (SOC<sub>stock</sub>, Mg C ha<sup>-1</sup>) was
  calculated as:
- 215

 $216 \qquad SOC_{stock} = SOC_{cont} \times (BD \times L \times 10,000)$ 

- where  $SOC_{cont}$  (Mg C Mg<sup>-1</sup> soil) is the SOC content, BD (Mg soil m<sup>-3</sup> soil) is the bulk density of the soil, L (m) is the depth of the soil layer analysed for  $SOC_{cont}$  (i.e. 0.25 m in this study) and 10,000 (m<sup>2</sup> ha<sup>-1</sup>) is used to express  $SOC_{stock}$  by hectare. Bulk density was determined using the soil core method (NRCS, 2014) with 5 replicates per plot.
- A part of the composite sample was used for laboratory incubations but the 222 223 sample was not sieved (aggregate size 5 mm) to reduce the effect of soil disturbance 224 (Balesdent et al., 2000). Soil moisture was set at -30 kPa using a regulated vacuum 225 pump. At this water potential the gravimetric water contents were on average 0.66 kg kg<sup>-1</sup> for andosols, 0.40 kg kg<sup>-1</sup> for nitisols, 0.43 kg kg<sup>-1</sup> for ferralsols and 0.46 kg kg<sup>-1</sup> 226 for vertisols. Laboratory incubations were carried out as described by Raphael et al. 227 228 (2012). Briefly, a glass vial containing 20 g (dry matter) of soil was placed in a glass 229 jar and incubated at 30 °C in the dark for 15 wk using four replicates per plot. The CO<sub>2</sub> 230 released from SOC mineralisation (i.e. microbial respiration) was trapped in a 0.3 M NaOH solution and determined by titrimetry. The traps containing the NaOH solution 231 232 were changed after each measurement which was performed seven times during the experiment. 233

Differences in SOC stocks and SOC mineralisation between soil-crop situations were assessed using ANOVA under an incomplete two-way design with the version 19.5 of the XLSTAT program.

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#### 238 2.4. Field rate constant of SOC mineralisation

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The SOC mineralisation rate constants in the field ( $k_{field}$ , in yr<sup>-1</sup>) for each soilcrop situation analysed during the present study had been reported by Sierra et al. (2015), who determined changes of SOC stocks using the following equation:

244 
$$dSOC_{stock}/dt = [C_{res} \times h_{res}] - [SOC_{stock} \times (k_{AER} \times k_{crop})]$$
(2)

245

where dSOC<sub>stock</sub>/dt is the annual change in SOC stock (Mg C ha<sup>-1</sup> yr<sup>-1</sup>), C<sub>res</sub> (Mg C ha<sup>-1</sup> yr<sup>-1</sup>) is the annual C input from crop residues,  $h_{res}$  (unitless) is the humification coefficient of crop residues,  $k_{AER}$  (yr<sup>-1</sup>) is the mineralisation rate constant for each AER (i.e. for each soil type in the present study), and  $k_{crop}$  (unitless) is the coefficient accounting for the effect of the cropping system on  $k_{AER}$ . Although  $k_{AER}$  reflects the effect of pedoclimatic conditions on SOC mineralisation,  $k_{crop}$  links with the impact of soil tillage on SOC mineralisation (Sierra et al., 2015). Therefore,  $k_{crop}$  is lower for perennial crops where soil tillage is applied every 5–6 yr (e.g. banana and
sugarcane), and higher for annual crops that involve several tillage operations per year
(e.g. vegetables and yam).

Using this approach, the rate constant of SOC mineralisation in the field ( $k_{field}$ , yr<sup>-1</sup>) can be calculated as:

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259  $k_{\text{field}} = k_{\text{AER}} \times k_{\text{crop}}$ 

260

Sierra et al. (2015) estimated k<sub>AER</sub> and k<sub>crop</sub> using 253 plots with diachronic measurements of SOC stocks between 1998 and 2014, and information concerning the rotations and management of C inputs included in the database. The k<sub>AER</sub>, k<sub>crop</sub> and k<sub>field</sub> values for each soil-crop situation included in the present study are presented in Table S1 of the Supplementary Data file. The characteristics of crop residues are presented in Table S2 (e.g. residue biomass, N and lignin content, C/N ratio, coefficient of humification and C input from crop residues).

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#### **3. Results and discussion**

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#### 3.1. SOC stocks and C mineralisation under laboratory conditions

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274 The organic C and organic N contents were highest in andosols, and there 275 were no significant differences between the other soil types (P < 0.05) (Table 1). 276 SOC stocks were mainly affected by soil type (Table 2), which was due to differences 277 in SOC content (Table 1). SOC stocks in the 0–0.25 m soil layer decreased 278 significantly in the following order: andosols (74 Mg C ha<sup>-1</sup>) > vertisols (60 Mg C ha<sup>-1</sup>) = ferralsols (58 Mg C ha<sup>-1</sup>) > nitisols (46 Mg C ha<sup>-1</sup>) (P < 0.05) (Table 1). It is well known 279 280 that andosols store a great amount of SOC because allophanic minerals incorporate 281 organic compounds that develop amorphous gels, which in turn provide physical 282 protection for SOC against microbial decomposition (Kramer and Chadwick, 2016). By 283 contrast, in lowlands dominated by nitisols under a less humid climate, some gel minerals 284 gradually reorganise into crystalline 1:1 clay minerals (e.g. halloysite), which adsorb organic matter on their surfaces thus decreasing the degree of physical protection 285 286 (Colmet-Daage and Lagache, 1965). Aluminium and iron hydrous oxides in ferralsols and 287 smectite clay in vertisols offer a degree of physical SOC protection that is intermediate between the allophanic minerals of andosols and the halloysite clay of nitisols, which 288

(3)

explain their intermediate SOC stocks compared to these soil types (Powers and
Schlesinger, 2002). ANOVA indicated that SOC contents and SOC stocks were not
affected by the crop (Table 2), which confirms that although crop management may
affect the annual SOC balance (Sierra et al., 2015), it does not affect the differences in
SOC between soils, which are dependent on the inherent pedoclimatic characteristics
of each AER (Cabidoche et al., 2004).

295 The procedure applied to evaluate SOC mineralisation under laboratory 296 conditions was suited to preventing artefacts due to sample handling. Indeed, SOC 297 mineralisation displayed a linear time course without evidence of an initial stimulation of 298 CO<sub>2</sub> release due to rewetting and soil disruption during the preparation of soil samples 299 (Jarvis et al., 2007) (Fig. 1). Such a linear time course differed from the results of SOC 300 mineralisation under laboratory experiments reported by other authors who observed an 301 exponential pattern of C release indicating first-order kinetics and a dependence of C 302 mineralisation upon the size of the mineralisable SOC pool (e.g. Benbi and Khosa, 303 2014). However, other authors found that in some cases, SOC mineralisation could be 304 described by applying different kinetics models (i.e. zero-order, first-order, mixed zero-305 and first-order) (Saviozzi et al., 2014). In our laboratory experiment the rate of SOC 306 mineralisation was constant in time, which suggests that the decrease in the size of the 307 mineralisable SOC fraction during soil incubation did not affect that rate. This apparent 308 zero-order kinetics was probably linked to the relatively small size of the mineralised 309 SOC fraction when compared with the total amount of SOC (e.g. the C<sub>min</sub>/SOC ratio 310 averaged 3.7% for the 69 soils; Table 1).

311 Unlike the findings with respect to SOC contents and SOC stocks, soil type 312 and crop affected in a similar extent the amount of mineralised SOC after 15 wk of laboratory incubation ( $C_{min}$ ) (P < 0.01; Table 2). On average,  $C_{min}$  ranged from 1.4 313  $q C kq^{-1}$  in andosols (4.0% of the SOC content) to 0.8  $q C kq^{-1}$  in vertisols (3.5%), 314 and from 1.3 g C kg<sup>-1</sup> in soils under banana (4.8%) to 0.6 g C kg<sup>-1</sup> under sugarcane 315 316 (3.0%) (Table 1). It appears that the effect of soil type on C<sub>min</sub> was driven by the 317 SOC content, as suggested by the highest values being observed in andosols 318 (Table 1) and the significant correlations found between the two variables in 319 andosols and vertisols (Fig. 2). However, this correlation was induced by small 320 number of extreme high SOC values in both soil types (i.e. two extreme high values in andosols and one extreme high value in vertisols; Fig. 2a and d). In fact, when 321 322 these extreme values were excluded from the analysis, the correlation between C<sub>min</sub> 323 and SOC content was not significant in either andosols or vertisols (P < 0.05). The weakness of the relationship between C<sub>min</sub> and SOC was induced by the impact of 324

325 the crop on C<sub>min</sub>, which was particularly noticeable in ferralsols and nitisols (Fig. 2b and c). Indeed, soils under banana in nitisols and ferralsols, and under pineapple in 326 ferralsols, presented the highest C<sub>min</sub> values regardless of their SOC content. Some 327 328 authors have reported that crop residues recently returned to the soil may increase 329 the more biodegradable SOC fraction and then boost soil respiration (e.g. Austin et 330 al., 2017; Lian et al., 2017). Our results indicated than the relationship between  $C_{min}$ 331 and C input from crop residues was not significant (P < 0.05; Fig. 3), and the same 332 was found between C<sub>min</sub> and other chemical and biological characteristics of the 333 residues for the six crops analysed during this study (i.e. residue biomass, N and 334 lignin contents, C/N ratio and coefficient of humification; Table S2) (these 335 relationships are not shown in Fig. 3). Similarly, although soils under sugarcane 336 presented simultaneously the highest C input from residues and the lowest  $C_{min}$ , 337 soils under banana exhibited the highest C<sub>min</sub> and a C input value that was 40% 338 lower than that from sugarcane (Fig. 3). As discussed in Section 3.2, it appears that 339 C<sub>min</sub> was dependent on the global impact of the cropping system rather than on recent C inputs from crop residues. This supports the suitability of the procedure 340 341 used for soil sampling, whose aim was to reduce the impact of residues from the 342 previous crop.

343

# 344 3.2. Relationship between SOC mineralisation under field and laboratory conditions345

Negative and significant correlations were observed between Cmin and kfield and 346 347 between  $C_{min}$  and  $k_{crop}$  (P < 0.05) (Fig. 4). The relationship between  $C_{min}$  and  $k_{AER}$ 348  $(R^2 = 0.22)$  was not significant (P < 0.05; data not shown in Fig. 4). These negative 349 relationships support our hypothesis concerning a direct link between k<sub>field</sub> and SOC 350 mineralisation under the conditions imposed to the laboratory experiment. These 351 results suggest that the organic C pool involved in SOC mineralisation under 352 laboratory conditions was depleted in soil-crop systems with high SOC mineralisation 353 in the field (i.e. high  $k_{\text{field}}$  and  $k_{\text{crop}}$ ), which induced a lower  $C_{\text{min}}$ . It is interesting to note 354 that this C pool could not be expressed in terms of SOC content because its 355 relationship with C<sub>min</sub> was guite inconsistent for most of the soil-crop systems 356 analysed in this study (Fig. 2). Further, although the negative relationship between C<sub>min</sub> and k<sub>crop</sub> indicated that the cropping system affected C<sub>min</sub> (Fig. 4b), ANOVA 357 358 revealed that the crop did not affect the SOC content (Table 2). It thus follows that the 359 organic C pool involved in SOC mineralisation during the laboratory experiment was probably associated with the more labile fractions of organic matter, which were more 360

361 or less depleted as a function of the level of C mineralisation in the field. These 362 results disagree with those of some studies which reported a significant correlation 363 between total C and N contents and the mineralised N observed under laboratory 364 conditions (e.g. Ros et al., 2011; Wade et al., 2016). However, our results are in line 365 with those obtained from long-term field experiments concerning the impact of tillage systems on SOC dynamics (Martínez et al., 2017; Muñoz-Romero et al., 2017). 366 367 These authors found that the labile SOC fraction was larger under a no-tillage 368 system than with conventional tillage, and that this fraction controlled SOC 369 mineralisation under laboratory conditions. These results were attributed to the effect of 370 tillage on the degree of exhaustion of the less stable forms of SOC in the field. This 371 proposal could be utilised to interpret some of the differences between cropping 372 systems observed during the present study. For example, in banana systems, soil 373 tillage is applied every 5 yr and its impact on the labile C fraction would be smaller than 374 that in yam and vegetable crop systems including several operations of tillage per year. 375 Therefore, the marked differences in C<sub>min</sub> observed between these cropping systems 376 reflected the effects of farming practices on SOC turnover. Under our approach, the 377 effect of soil tillage was represented by the  $k_{crop}$  coefficient (Eq. (2)).

378 A more interesting finding was that the C<sub>min</sub>/SOC ratio was markedly 379 affected by the crop but not by the soil type (Table 2). This ratio is frequently referred as being an indicator of the more active SOC fraction involved in SOC 380 381 mineralisation under laboratory conditions (e.g. Campbell and Souster, 1982). 382 During our study, the  $C_{min}$ /SOC content ratio decreased significantly in the following 383 order (Table 1): banana (5.0%) > pineapple (4.2%) > melon (3.4%) >384 vegetables (3.2%) = yam (3.2%) > sugarcane (3.0%) (P < 0.05). The lack of an 385 effect of the soil on this ratio was associated with the strong impact of banana and 386 sugarcane, which respectively presented the highest and lowest  $C_{min}$ /SOC values 387 regardless of soil type. Martínez et al. (2017) reported significant correlations 388 between the active SOC fraction determined under laboratory conditions and C input 389 from previous crops. As cited above, this relationship was not significant in the present 390 study (Fig. 3). We therefore concluded that SOC mineralisation as determined in the 391 laboratory reflected the overall impact of the soil-crop system on SOC over several 392 years of crop monoculture. This conclusion was supported by the significant 393 relationship between C<sub>min</sub> and k<sub>field</sub>; the latter was determined from a long-term field 394 study (i.e. 17 yr; Sierra et al., 2015).

Whilst the soil and crop could explain almost 80% of total  $C_{min}$  variation (Table 2), some results still need to be explained. Firstly, because the soil x crop

397 interaction was not significant some apparent interactions concerning vertisols (i.e. similar C<sub>min</sub> values for four contrasting crops characterised by a broad range of soil 398 tillage intensities; Fig. 4) could not be accurately assessed. This was probably due 399 400 to the small number of degrees of freedom attributed to the interactions in our incomplete two-way design. In other words, because the crops are not cultivated in 401 all the AERs concerned, only a small number of interactions were included in the 402 403 ANOVA. Secondly, soils under sugarcane presented the lowest values for C<sub>min</sub> and 404 C<sub>min</sub>/SOC despite its high rate of annual C input from residues and low intensity soil 405 tillage. We feel these two issues might be linked to the biochemical characteristics of 406 crop residues that could affect the level of the labile pool of SOC and then C<sub>min</sub>. 407 Several authors found that when lignocellulosic material decomposes in soil, the 408 lignin fraction is not incorporated into microbial biomass but is directly transferred 409 into SOC (e.g. Corbeels et al., 2005). Taking account of the fact that the turnover of 410 microbial metabolites contributes to the labile SOC fraction (Paul, 2016), it could be 411 hypothesised that part of the residues from sugarcane, with the highest lignin 412 content observed during the present study (Table S2), was not recycled into the 413 more labile C fraction, thus causing a relatively low C<sub>min</sub>. By contrast, banana 414 residues have low lignin content (Table S2), which together with a relatively large 415 quantity of residues induced the highest values for C<sub>min</sub> and the C<sub>min</sub>/SOC content ratio observed in this study. It is possible that the similar C<sub>min</sub> values observed in 416 417 vertisols under the four contrasting cropping systems (i.e. sugarcane, vegetables, 418 yam and melon) were linked to the combined effects of the quantity and quality of 419 residues on the size of the labile C pool throughout the cycle of monocultures. 420 Despite some discrepancies between the field and laboratory experiments, our 421 results suggest that C<sub>min</sub> could be used as an indicator of the impact of new cropping 422 systems and practices on SOC, and to assess  $k_{field}$  before SOC changes become 423 noticeable in the field. However, the impact of residue quality on the size of the 424 mineralisable C fraction is not currently included explicitly in the k<sub>field</sub> and k<sub>crop</sub> 425 coefficients. Further work is therefore necessary to determine the relationship between 426 the biochemical properties of crop residues and the distribution of SOC fractions in 427 tropical soils under contrasting tillage management. 428

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#### 430 **4. Conclusions**

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- The major finding of this study was that SOC mineralisation under the

433 conditions imposed to the laboratory experiment (C<sub>min</sub>) was a suitable indicator of the impact of cropping systems on the mineralisation rate constant ( $k_{field}$  and  $k_{crop}$ ) 434 estimated from long-term field experiments. The negative relationship between C<sub>min</sub> 435 436 and k<sub>field</sub> was mainly controlled by the degree of depletion of the labile SOC fraction induced by the management of the soil-crop system. Differences in the intensity of 437 438 soil tillage between cropping systems (i.e. represented by  $k_{crop}$ ) were probably the 439 key factor affecting the degree of this depletion. It is important to point out that a low 440 C<sub>min</sub> does not imply that k<sub>field</sub> and k<sub>crop</sub> values will fall in the near future due to 441 exhaustion of the labile SOC fraction. In fact, C<sub>min</sub> indicates an earlier impact of the 442 cropping system but it does not predict a change to k<sub>field</sub>, which is dependent on the 443 management of the cropping system and the pedoclimatic conditions that prevail in 444 each AER. Although laboratory soil incubation is a time-consuming method, our 445 results indicate that C<sub>min</sub> could be useful to quantify the impact of new agricultural 446 practices (e.g. manual vs. mechanical tillage) before SOC changes become noticeable in the field. To achieve this, further work is necessary to assess the effect 447 of residue quality on  $k_{crop}$  under contrasting tillage management in tropics. 448 449

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#### 451 **Acknowledgements**

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#### 462 Appendix A. Supplementary data

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464 Supplementary material related to this article can be found, in the online 465 version, at

- 466
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- 468References

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#### 587 **Table 1**

588 Some soil properties of the 0–0.25m soil layer corresponding to the 12 cropping

589 systems analysed during this study. For each property, mean values followed by

<sup>590</sup> different letters are significantly different at P < 0.05. SOC: soil organic C; SON: soil

<sup>591</sup> organic N; Cmin: mineralised C accumulated during the 15-wk laboratory experiment.

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	Soil	Crop	Number of plots	Bulk density Mg m <sup>-3</sup>	pH	SOC g kg <sup>-1</sup>	SON	SOC/SON ratio	SOC stock Mg ha <sup>-1</sup>	C <sub>min</sub> g Ckg <sup>-1</sup>	C <sub>min</sub> /SOC ratio %
_	Andosol	Banana	6	0.81d	5.9 b	40.8 a	4.0 a	10.0 a	82 a	1.63a	4.2bc
	Andosol	Vegetables	5	0.82d	5.9 b	32.8 a	3.2 a	10.0 a	66 a	1.12b	3.4c
	Nitisol	Banana	5	0.92c	5.1 c	19.9 b	2.0 b	9.9 a	45 c	1.20b	6.2a
	Nitisol	Vegetables	5	0.92c	5.0 c	21.2 b	2.1 b	9.8 a	48 c	0.51e	2.4d
	Ferralsol	Banana	5	1.05b	5.2 c	24.1 b	2.4 b	9.9 a	63 ab	1.05bc	4.6b
	Ferralsol	Sugarcane	6	1.05b	5.2 c	21.4 b	2.1 b	10.0 a	56 b	0.55e	2.5d
	Ferralsol	Yam	5	1.07b	5.2 c	21.3 b	2.1 b	9.9 a	56 b	0.59e	2.8d
	Ferralsol	Pineapple	6	1.03b	5.1 c	21.1 b	2.1 b	10.0 a	55 b	0.89cd	4.2bc
	Vertisol	Sugarcane	8	1.14a	7.1 a	21.8 b	2.1 b	9.9 a	60 b	0.74de	3.4c
	Vertisol	Vegetables	6	1.12a	7.2 a	21.4 b	2.2 b	10.0 a	59 b	0.78de	3.6bc
	Vertisol	Yam	5	1.14a	7.0 a	21.6 b	2.1 b	10.0 a	59 b	0.80de	3.7bc
	Vertisol	Melon	7	1.15a	7.2 a	21.6 b	2.2 b	10.0 a	59 b	0.73de	3.4c

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#### **Table 2**

Results of the analysis of variance. SOC: soil organic carbon; Cmin: mineralised C accumulated during the 15-wk

604 laboratory experiment. The % variance refers to the contribution of each source of variation to the total variance observed605 for each soil parameter.

607 -														
608	Source of variation	Degrees of freedom	SOC content		SOC stock			Cmin			Cmin/SOC content			
609			% variance	F-value	P-value	% variance	F-value	P-value	% variance	F-value	P-value	% variance	F-value	P-value
610	Soil	3	67	46.3	< 0.01	45	18.9	< 0.01	44	39.3	< 0.01	5	2.3	0.09
611	Crop	5	3	1.1	0.35	5	1.2	0.32	34	18.1	< 0.01	45	10.3	< 0.01
	Soil $\times$ Crop	3	3	2.3	0.09	5	2.1	0.11	1	0.7	0.56	6	2.4	0.09
612	Error	57	27			45			21			44		



**Fig. 1.** Time course of SOC mineralisation during the laboratory experiment ( $C_{min}$ ). The two soil-crop situations presented correspond to the extreme mineralised C values observed during this study. Vertical bars indicate the standard deviation.



Fig. 2. Relationship between accumulated mineralised C after 15 wk of soil incubation in the laboratory (C<sub>min</sub>) and the SOC
 content for each soil type and crop.





Fig. 4. Relationship between accumulated mineralised C after 15 wks of soil incubation in the laboratory (C<sub>min</sub>) and (a) the
 SOC mineralisation rate constant (k<sub>field</sub>), and (b) the coefficient reflecting the impact of tillage management on SOC
 mineralisation (k<sub>crop</sub>). k<sub>field</sub> and k<sub>crop</sub> were obtained by Sierra et al. (2015) from long-term field experiments. Vertical bars

indicate the standard deviation.