Effect of organic carbon addition on paddy soil organic carbon decomposition under different irrigation regimes

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Abstract. Anaerobic decomposition of organic carbon (OC) in submerged rice paddies is coupled to the reduction of alternative soil electron acceptors, primarily Fe^{3+}. During reductive dissolution of Fe^{3+} from pedogenic oxides, previously adsorbed native soil organic carbon (SOC) could be co-released into solution. Incorporation of crop residues could hence indirectly, i.e. through the stimulation of microbially mediated Fe^{3+} reduction, promote the loss of native SOC via enhanced dissolution and subsequent mineralisation to CO_2 and CH_4. Our aim was to estimate the relevance of such a positive feedback during the degradation of added OC, and to investigate the impact of irrigation management on this mechanism and on priming effects on native SOC decomposition in general. In a six-week pot experiment with rice plants, two Bangladeshi soils with contrasting SOC to oxalate-extractable Fe ($SOC : Fe_{ox}$) ratios were kept under a regime of alternate wetting and drying (AWD) or continuous flooding (CF), and were either amended with maize shoots or not. The $\delta^{13}C$ signatures of dissolved organic C and emitted CH_4 and CO_2 were used to infer the decomposition of added maize shoots ($\delta^{13}C = -13.0\,‰$) versus native SOC ($\delta^{13}C = -25.4\,‰$ and $-22.7\,‰$). Addition of maize residues stimulated the reduction of Fe as well as the dissolution of native SOC, and the latter to a larger extent under CF, especially for the soil with the highest $SOC : Fe_{ox}$ ratio. Estimated Fe-bound SOC contents denote that stimulated SOC co-release during Fe reduction could explain this positive priming effect on SOC dissolution after the addition of maize. However, priming effects on SOC mineralisation to CO_2 and CH_4 were lower than for SOC dissolution, and were even negative under AWD for one soil. Enhanced reductive dissolution of Fe-bound SOC upon exogenous OC addition therefore does not necessarily lead to stimulated SOC mineralisation. In addition, AWD irrigation was found to decrease the above-mentioned priming effects.

1 Introduction

Anaerobiosis in flooded paddy fields thoroughly affects soil chemical processes, as in the absence of oxygen, the decomposition of organic carbon (OC) requires alternative terminal electron acceptors like manganese (Mn^{4+}), iron (Fe^{3+}), sulphate, acetate or carbon dioxide (CO_2). Anaerobic reduction of the latter two electron acceptors, which also depends on the production of dissolved organic carbon (DOC) as an electron donor, leads to the production and emission of methane (CH_4) (Kögel-Knabner et al., 2010; Ponnamperuma, 1972). Obviously, irrigation management strongly affects the prevalence of aerobic versus anaerobic OC decomposition. The impact of irrigation management on soil processes is an extensively studied topic because of the demand to grow rice more efficiently with less water (Carrijo et al., 2017). In particular, the adoption of alternate wetting and drying (AWD), a periodic drying and reflooding irrigation practice, is increasingly promoted (Lampayan et al., 2015). However, our understanding of how the alternation of redox conditions under AWD irrigation affects the decomposition of native soil organic carbon (SOC) and exogenous OC in comparison with more consistent anaerobic conditions is still rather limited. This limited insight mainly stems from difficulties in discerning native SOC and exogenous OC mineralisation based

on soil gaseous emissions. In upland soils, assessments usually rely on $\delta^{13}C$ isotopic signatures of emitted CO_2 derived from C sources with distinct $\delta^{13}C$ (Hayes, 1983; Werth and Kuzyakov, 2010). However, in submerged soils, emission of CH_4 also needs to be accounted for. Since most CH_4 and CO_2 is emitted through rice plant aerenchyma transport, a realistic experiment necessitates the presence of growing rice plants, which further complicates the tracking of OC mineralisation in paddy soils. It is then important to account for C isotope fractionation during the OC decomposition process, mainly caused by the microbial community discriminating for or against ^{13}C. In particular, during anaerobic CH_4 production, the shift in $\delta^{13}C$ is typically large and moreover, production pathway-dependent (Conrad, 2005; Sugimoto and Wada, 1993; Schweizer et al., 1999; Werth and Kuzyakov, 2010; Conrad et al., 2012). The $\delta^{13}C$ of emitted CO_2 is on the other hand influenced by CO_2 consumption during hydrogenotrophic CH_4 production (i.e. with H_2/CO_2 as substrate) and its partial storage as dissolved bicarbonate and CO_2. As a result, in the absence of oxygen, it is much less straightforward to link the $\delta^{13}C$-CO_2 and $\delta^{13}C$-CH_4 of emissions to the $\delta^{13}C$ of the original C substrate (Conrad et al., 2012).

Furthermore, the incorporation of crop residues is a common practice in paddy soil (Ponnamperuma, 1984; Mandal et al., 2004). It is known that exogenous OC addition can modify the rate of SOC decomposition in upland soils via so-called priming effects. Positive priming could for example result from soil microbial growth upon the addition of fresh OC as an energy source, resulting in the co-metabolism of biologically more recalcitrant SOC (Kuzyakov et al., 2000; Blagodatskaya and Kuzyakov, 2008). Positive as well as negative priming effects of OC addition on the dissolution of native SOC and on its further mineralisation into CO_2 and CH_4 have also been reported for flooded paddy soils (Bertora et al., 2018; Conrad et al., 2012; Ye et al., 2015; Ye and Horwath, 2017; Yuan et al., 2014), but again these processes are much less well understood. Indeed, considering the unique interlinkage between anaerobic microbial activity and redox reactions with mineral soil oxidants, priming mechanisms could be quite different than in upland soils. For example, Bertora et al. (2018) and Ye and Horwath (2017) observed that rice straw addition enhances the reductive dissolution of pedogenic Fe, and found indications that this in turn led to a stimulated co-release of OC originating from the Fe-OC complex. Pedogenic Fe^{3+} oxides, hydroxides and oxyhydroxides (hereafter collectively referred to as "Fe oxides"), which are often abundant in tropical soils, are indeed relevant sorbents of SOC owing to their high reactive specific surface area or through complexation (Wagai et al., 2013; Chen et al., 2020; Kaiser and Guggenberger, 2000). Dissolution of OC from these minerals following the establishment of anaerobic conditions could thus be a considerable C-releasing process and may take place (i) by desorption due to a pH increase, caused by the consumption of protons during reduction reac-

tions; or (ii) by co-release of sorbed OC along with reductive dissolution of short-range-ordered Fe^{3+} minerals such as ferrihydrite and goethite (Grybos et al., 2009; Said-Pullicino et al., 2016). However, it is still unknown whether such stimulated native SOC dissolution would in turn significantly promote further conversion into CO_2 and CH_4. This would seem plausible as DOC is assumed to be one of the most bioavailable SOC fractions, especially under submerged conditions (Marschner and Kalbitz, 2003; Said-Pullicino et al., 2016). The indirect stimulation of SOC mineralisation by OC amendment through enhanced reduction of Fe in paddy soil may furthermore be expected to depend on the water regime, as this overridingly determines the extent of pedogenic Fe reduction and increase in pH. Lastly, opposite to the co-dissolution of Fe and OC, DOC can likewise be adsorbed onto or can co-precipitate with the newly formed Fe^{3+} oxides as soon as the E_h again increases and Fe^{2+} is reoxidised (Sodano et al., 2017). Such C removal pathways might again strongly depend on irrigation management.

Our aim was to investigate how the addition of exogenous OC influences the dissolution and mineralisation of native SOC in paddy soils in function of water management, with particular attention to the role of the co-release of Fe-bound SOC. To this end, patterns of DOC and the emission of CO_2 and CH_4 were compared in a six-week pot experiment with living rice plants in a tropical greenhouse. Two common irrigation practices were compared, i.e. the water-saving AWD practice next to continuous flooding (CF). We hypothesised that maize shoot addition would result in positive priming of native SOC mineralisation. This priming should be stronger under CF than under AWD, owing to enhanced pedogenic Fe^{3+} reduction with stimulated net co-release of Fe-bound native SOC. Maize shoots were used as an external OC source, because of the contrast of their $\delta^{13}C$ (C_4 crop) with the $\delta^{13}C$ of the native SOC (largely C_3-derived due to long-term rice cultivation) in both soils. A secondary aim was to see if the proportion of SOC compared to pedogenic Fe, with the latter considered to be the main pool of readily reducible Fe, interacts with the effect of OC amendment on native SOC decomposition. Therefore, we compared two soils from Bangladeshi young floodplain paddy fields that were specifically chosen for their contrasting SOC to oxalate-extractable Fe (Fe_{ox}, as a proxy for short-range-ordered Fe oxides that can be considered easily reducible; Postma, 1993; van Bodegom et al., 2003) ratio. We expected a stronger stimulation of native SOC release from the soil with the higher SOC : Fe_{ox} ratio.

2   Material and methods

2.1   Soils

Two young floodplain paddy soils (from soil series Balina and Sonatala) from northern Bangladesh were selected from

a larger set of soils previously used by Akter et al. (2018) and Kader et al. (2013). Per soil series, the puddle layer soil (0–15 cm) of 15 locations within one field was sampled and then bulked by means of a clean spade in May 2014, after which the soil was stored in air-dried, ground and sieved form. Both soils were selected based on their contrasting $SOC : Fe_{ox}$ ratios with other traits remaining largely similar (Table 1).

X-ray diffraction analyses confirmed that both soils contained mica, vermiculite, chlorite and kaolinite, and Sonatala also some crystalline goethite (Kader et al., 2013). The presence of Fe^{3+} in chlorite and vermiculite was confirmed by Mössbauer spectroscopy (Akter et al., 2018), but ferrihydrite and poorly crystalline goethite were the main Fe^{3+} pools in both soils.

2.2 Pot experiment

A six-week pot experiment with living rice plants was run in the tropical greenhouse of the Faculty of Bioscience Engineering (Ghent University) in Melle, Belgium, from 23 April to 4 June 2018. The mean ambient temperature was 28 °C and the relative humidity was 82.2 %. PVC tubes (diameter: 18.5 cm; height: 25.0 cm) were filled with soil from either Balina (4.8 kg dry soil) or Sonatala (4.0 kg dry soil) until a height of 17.5 cm, matching their respective bulk densities, and were subjected to two contrasting irrigation treatments, as described below. Ground maize shoots were mixed into the soil at a dose of 4 g dry mass kg^{-1} soil (i.e. 1.90 g C kg^{-1}) to three replicate pots per soil type and per irrigation regime. Maize shoots had an OC content of 474.4 g kg^{-1}, a $\delta^{13}C$ of $-13.04 ‰$ and a total N content of 13.0 g kg^{-1}. Two replicate pots were installed as controls without maize added, and two more replicates per irrigation and soil combination received maize (same dose of 4 g kg^{-1}) but had no living rice plants. Macronutrients were added once as basal fertilisation to all pots as urea, KCl and $Ca_3(PO_4)_2$, at doses of 60 kg N ha^{-1}, 40 kg K ha^{-1} and 10 kg P ha^{-1}, respectively. Five 15 d-old rice seedlings were transplanted to each pot in a single hill. These seedlings were obtained by germinating rice seeds of BINA dhan14 (*Oryza sativa* L.), a Bangladeshi short-duration (maturity 120–130 d) dry season rice variety, and sowing those in seeding beds that were kept saturated in the tropical greenhouse for 15 d.

A standing water table of 2.5 cm was initially maintained in all pots by adding demineralised water every one or two days to enable seedling establishment. Starting from 11 d after transplanting (DAT), the two different irrigation regimes were imposed. In pots under CF, the 2.5 cm water table was maintained until the end of the experiment. Pots under AWD, in contrast, were left to dry out until the water table dropped to between -8 and -14 cm, after which they were reflooded until a 2 cm standing water table was reached. Water table depths were monitored using a perforated tube (diameter: 3 cm) that was permanently installed in the pots. Soil drying was only due to evapotranspiration. Drying cycles took on average 6 (± 3) d and shortened towards the end of the experiment.

2.3 Monitoring of biochemical soil parameters

The soil redox potential (E_h) was measured manually daily or every two days at 4 and 12 cm depth in eight pots with plants (i.e. 2 irrigation treatments × 2 soil series × with versus without maize). This was achieved by connecting a redox mV meter (Paleo Terra, the Netherlands) to permanently installed platinum (Pt) probes and one Ag|AgCl reference electrode (Paleo Terra, the Netherlands) per pot, and expressed relative to the standard hydrogen electrode. Soil pH was measured six times during the experiment in every pot at about 2 cm depth by inserting a glass pH electrode into the saturated soil.

To place CH_4 and CO_2 flux measurements in context, KCl-extractable ammonium (NH_4^+) and nitrate (NO_3^-) were determined at the start of the experiment (0 DAT), and at 11, 23, 30 and 45 DAT. For this, soil samples were taken by carefully inserting a tube (diameter: 1 cm) until 10 cm depth about four times at different locations near the side of each PVC pot. Soil samples were then stored at -20 °C until analysis. The contents of NH_4^+ and NO_3^- were determined with a continuous flow auto-analyser (Skalar, the Netherlands) after extracting soil samples with 1 M KCl at a 1 : 5 ratio.

At the end of the experiment, the above- and below-ground biomass of the rice plants was determined after drying at 60 °C for three days. The $\delta^{13}C$ of SOC was determined before the experiment for solid native SOC and maize, and after the experiment in SOC, with an EA-IRMS (automated nitrogen carbon analyser – solids and liquids – coupled to a SerCon 20-20 IRMS, Syscon Electronics, the Netherlands).

2.4 Soil solution sampling and analyses of Fe^{2+}, Mn^{2+}, DOC and $\delta^{13}C$-DOC

To track dissolved Fe, Mn, Ca, Mg and DOC, soil solution samples (2×9 mL per pot) were taken ten times throughout the experiment. Solution samples were collected by connecting a pre-evacuated 12 mL glass vial through a needle, 2-way stopcock and extension tube to a porous MacroRhizon soil moisture sampler (Rhizosphere Research Products, the Netherlands) that was permanently installed in each pot in a vertical position until about 9 cm depth. Solutions were analysed for their Fe, Mn, Ca and Mg concentrations by ICP-OES (Thermo Scientific, Unites States) without filtration, after acidifying the solutions with some drops of concentrated HNO_3 to redissolve precipitates. The sum of the molar concentrations of dissolved Fe and the increment in dissolved Ca and Mg (compared to their initial level) was taken as a proxy for total reduced Fe since Fe^{2+} dissolves easily, but readily exchanges with Ca^{2+} and Mg^{2+} on colloid surfaces (Saeki, 2004). In the same samples, the DOC concentration and $\delta^{13}C$-DOC were measured by FIA-IRMS (flow injec-

Table 1. Initial properties of the selected two paddy soil series and maize shoots used in the soil incubation experiment.

Parameter	Balina	Sonatala
Soil texture[a]	Silty clay loam	Silt loam
Soil type[a]	Mollic Haplaquept	Aeric Haplaquept
Yearly cropping pattern[a]	Rice – fallow – fallow	Rice – fallow – rice
OC content ($g\,kg^{-1}$)	14.1	22.4
$\delta^{13}C$ (‰)	−25.37	−22.70
Fe_{ox} ($g\,kg^{-1}$)[b,c]	8.4	4.8
$SOC:Fe_{ox}$ ratio (–)[d]	1.7	4.7
pH-KCl[c]	4.0	5.5
pH-H_2O[e]	6.1	6.9
Total N ($g\,kg^{-1}$)[c]	1.9	2.1
NH_4^+-N ($mg\,kg^{-1}$)[f]	8	3
NO_3^--N ($mg\,kg^{-1}$)[f]	0	1
C:N ratio	7.4	10.7

[a] Data taken from Kader et al. (2013). [b] Oxalate-extractable Fe. [c] Data taken from Akter et al. (2018). [d] Ratio of SOC to oxalate-extractable Fe. [e] pH-H_2O by inserting a glass pH electrode in 1 : 5 soil–water extracts after 18 h equilibration. [f] Exchangeable mineral N determined in 1 M KCl extracts.

tion analysis with isotope ratio mass spectrometry, DELTA V Plus Advantage, Thermo Scientific, United States) after dilution in 0.85 % H_3PO_4, evacuation and ultrasonication to remove carbonates and dissolved gases. To account for small differences in soil to soil solution ratios, soil solution concentrations of Fe equivalents and DOC were for each time point converted to mmol dissolved Fe equivalents kg^{-1} soil and mg C kg^{-1} soil, respectively.

2.5 Estimation of SOC bound to pedogenic Fe and its $\delta^{13}C$

We furthermore estimated the content and $\delta^{13}C$ signature of SOC bound to weakly crystalline pedogenic Fe in the initial Balina and Sonatala soils based on the release of SOC after reduction with hydroxylamine ($NH_2OH \cdot HCl$), i.e. by assessing the change in OC content and $\delta^{13}C$ of the solid soil samples with versus without $NH_2OH \cdot HCl$ treatment. In all samples, particulate organic matter was firstly removed through ultrasonication at $60\,J\,mL^{-1}$ and subsequent wet sieving at $53\,\mu m$ (Sleutel et al., 2007). After drying the sieved suspensions, 0.8 g soil and 40 mL of a solution of 0.25 M $NH_2OH \cdot HCl$ and 0.25 M HCl were mixed in a 85 mL Nalgene centrifuge tube, similarly to Chao and Zhou (1983). The tubes were then kept in a water bath at 50 °C for 30 min with regular stirring. After centrifuging at 1000 g for 15 min, the supernatant was decanted, and traces of $NH_2OH \cdot HCl$ and HCl were removed by washing the residue soil three times with 40 mL demineralised water. Samples without the $NH_2OH \cdot HCl$ treatment were treated identically, but the $NH_2OH \cdot HCl$ solution was replaced by demineralised water. Finally, all samples were dried for four days at 40 °C, ground and analysed for their C content and $\delta^{13}C$ by means of elemental analyser – isotope ratio mass

spectrometry (EA-IRMS), i.e. an ANCA-SL (automated nitrogen carbon analyser – solids and liquids) interfaced with a 20-22 IRMS (Sercon Ltd., United Kingdom). The content of Fe-bound SOC was then estimated as the difference in SOC between treated samples and samples without $NH_2OH \cdot HCl$ treatment. Analogous to Keeling (1958), the $\delta^{13}C$ of Fe-bound SOC ($\delta^{13}C$-SOC_{Fe}) was calculated based on the following mass balance: TS1

$$\delta^{13}C\text{-}SOC_{Fe} = \frac{\begin{array}{c}\delta^{13}C\text{-}SOC_{untreated} \cdot [C]\text{-}SOC_{untreated} \\ -\delta^{13}C\text{-}SOC_{NH_2OH} \cdot [C]\text{-}SOC_{NH_2OH}\end{array}}{[C]\text{-}SOC_{untreated} - [C]\text{-}SOC_{NH_2OH}}, \quad (1)$$

where $\delta^{13}C$-SOC_{NH_2OH} and [C]-SOC_{NH_2OH} refer respectively to the $\delta^{13}C$ and C concentration of samples treated with $NH_2OH \cdot HCl$, and $\delta^{13}C$-$SOC_{untreated}$ and [C]-$SOC_{untreated}$ to those of samples without $NH_2OH \cdot HCl$ treatment.

2.6 Gaseous C fluxes

In order to estimate the decomposition of OC to gaseous CH_4 and CO_2, we measured emission fluxes and $\delta^{13}C$ isotopic signatures of CO_2 and CH_4 at ten different moments for pots under CF (most frequently after the onset of the experiment) and at twenty-six moments for pots under AWD (spread evenly over the course of the experiment). These extra measurements were required to sufficiently capture changing emissions during soil wetting and drying cycles. Gaseous efflux measurements were conducted by connecting a cavity ring-down spectrometer (G2201-i CRDS isotopic CO_2/CH_4 analyser, Picarro, United States) in a loop to an opaque PVC gas flux chamber (diameter: 18.5 cm; height: 25.0 cm) with an internal battery-operated fan. For each measurement, this flux chamber was secured onto the pots by

means of a PVC connection ring (diameter: 18.5 cm; height: 16.0 cm), and the headspace accumulation of CO_2 and CH_4 as well as the evolution of their $^{13}C/^{12}C$ ratio (as opposed to the initial headspace air) was recorded every 4 s during 12 min. The CRDS was calibrated using standard gases with known $\delta^{13}C$ signatures of $-35.95\%o$ and $-26.43\%o$. Starting from 28 DAT, an extra PVC extension ring (diameter: 18.5 cm; height: 25.0 cm) was added to the chambers because rice plants had outgrown the initial chamber.

CH_4 and CO_2 fluxes were determined as the slope of the accumulating headspace CH_4 and CO_2 concentrations as a function of time, and were converted into a mass-based unit (mg C kg^{-1} soil h^{-1}) by means of the ideal gas law. Based on the conservation of mass, $^{13}C/^{12}C$ ratios of the emitted CH_4 and CO_2 were determined as the intercept of the linear regression from the isotopic signature of the headspace gas in function of the reciprocal of the headspace concentration, also known as the Keeling plot method (Keeling, 1958). All $^{13}C/^{12}C$ ratios were expressed relative to the international VPDB (Vienna Pee Dee Belemnite) standard as $\delta^{13}C$ (‰). When rice seedlings were young and pots were flooded, a considerable part of the produced CO_2 and CH_4 appeared to be transported to the atmosphere through ebullition. In those cases, their accumulation in the headspace did not follow a linear course but was stepwise, with alternating time intervals of high and low accumulation rates, which were assumed to be dominated by ebullition versus by plant-mediated transport and molecular diffusion, respectively. When this was the case, additional C emission fluxes and $\delta^{13}C$ signatures were derived for shorter time intervals during which emissions were dominated by ebullition or by plant-mediated transport and diffusion.

Average daily fluxes were estimated from the measured fluxes by accounting for diurnal fluctuation of the actual temperature using a temperature dependency model:

$$\text{Relative correction daily rate} = 1 - Q_{10}^{((T_{avg} - T_{meas})/10)}, \quad (2)$$

where T_{avg} and T_{meas} (in °C) respectively represent the average daily soil temperature and the temperature during measurement. The Q_{10} temperature coefficient was set to 3.1 for CH_4 (Wang et al., 2015; Wei et al., 2021; Hattori et al., 2001) and to 2.3 for CO_2 (Huang et al., 2015; P. Zhou et al., 2014; Zhang et al., 2007; Wei et al., 2021). Overall, T_{avg} was 29.5 ± 0.6 °C while T_{meas} was 32.6 ± 1.0 °C, as gas efflux measurements were always carried out around noon.

2.7 Source partitioning of gaseous and dissolved C

The fractions of emitted CO_2 and CH_4 derived from added maize shoots (C_4) ($f_{CO_2 \,|\, maize}$ and $f_{CH_4 \,|\, maize}$) versus from native SOC (C_3) ($f_{CO_2 \,|\, SOC} = 1 - f_{CO_2 \,|\, maize}$ and $f_{CH_4 \,|\, SOC} = 1 - f_{CH_4 \,|\, maize}$) were inferred from $\delta^{13}C$ signatures of emissions from (i) the "mixed pool" pots (to which maize was added); (ii) control pots without maize added ($\delta^{13}C$-$CO_{2 \,|\, SOC}$ and $\delta^{13}C$-$CH_{4 \,|\, SOC}$ as reference for

the C_3-derived emission endmember); and (iii) a shorter ancillary incubation with maize shoots as the only OC source ($\delta^{13}C$-$CO_{2 \,|\, maize}$ and $\delta^{13}C$-$CH_{4 \,|\, maize}$ as reference for the C_4-derived emission endmember) (see Supplement). The latter two were used instead of the $\delta^{13}C$ of respectively native SOC and maize-C in order to account for isotopic fractionation during microbial decomposition and emission of CO_2 and CH_4. Any effect of the presence of rice plants (plant respiration or decomposition of plant photosynthates to CO_2 and CH_4) is accounted for by the fact that the control pots without maize added (i.e. $\delta^{13}C$-$CO_{2 \,|\, SOC}$ and $\delta^{13}C$-$CH_{4 \,|\, SOC}$) contained rice plants too. Based on the conservation of mass, the fraction of maize-derived CO_2 is for example (Werth and Kuzyakov, 2010; Rochette and Flanagan, 1997; Hayes, 1983):

$$f_{CO_2 \,|\, maize} = \frac{\delta^{13}C\text{-}CO_2 - \delta^{13}C\text{-}CO_{2 \,|\, SOC}}{\delta^{13}C\text{-}CO_{2 \,|\, maize} - \delta^{13}C\text{-}CO_{2 \,|\, SOC}}. \quad (3)$$

The CO_2 or CH_4 fluxes derived from native SOC ($CO_{2 \,|\, SOC\text{-derived}}$ or $CH_{4 \,|\, SOC\text{-derived}}$) or maize ($CO_{2 \,|\, maize\text{-derived}}$ or $CH_{4 \,|\, maize\text{-derived}}$) in the mixed pots were then obtained by multiplying emissions with $f_{CO_2 \,|\, SOC}$ or $f_{CH_4 \,|\, SOC}$ versus $f_{CO_2 \,|\, maize}$ or $f_{CH_4 \,|\, maize}$. Then, cumulative emissions of CH_4, CO_2, maize-derived C and native SOC-derived C were calculated.

We calculated a dimensionless priming effect (PE) coefficient of maize addition on SOC-derived CO_2 or CH_4 emission by comparing SOC-derived CO_2 and CH_4 fluxes with those from control pots without maize ($CO_{2 \,|\, SOC}$ or $CH_{4 \,|\, SOC}$) (Conrad et al., 2012). For example, relative priming of CO_2 emission was defined as

$$PE_{CO_2} = \frac{CO_{2 \,|\, SOC\text{-derived}}}{CO_{2 \,|\, SOC}} - 1. \quad (4)$$

In addition, the fraction of DOC derived from maize ($f_{DOC \,|\, maize}$) was calculated as well based on $\delta^{13}C$ signatures of DOC in amended pots ($\delta^{13}C$-DOC), of control pots without amendment ($\delta^{13}C$-DOC_{SOC}), of solid maize-C ($\delta^{13}C_{maize}$) and of solid SOC ($\delta^{13}C_{SOC}$), as follows:

$$f_{DOC \,|\, maize} = \frac{\delta^{13}C\text{-}DOC - \delta^{13}C\text{-}DOC_{SOC}}{\delta^{13}C_{maize} - \delta^{13}C_{SOC}}. \quad (5)$$

DOC contents derived from SOC ($DOC_{SOC\text{-derived}}$) in amended soils were obtained by multiplying their total DOC contents with $(1 - f_{DOC \,|\, maize})$, and the relative priming effect of maize addition on SOC-derived C dissolution was then calculated in an analogous way to Eq. (4).

Because of the presence of rice plants, DOC contents as well as CH_4 and CO_2 emissions originate not only from native SOC versus maize shoots, but also from rice plant photosynthates. Having a similar $\delta^{13}C$ as SOC, rice plant-derived DOC, CH_4 and CO_2 will contribute to what is partitioned as SOC-derived C, leading to overestimation of SOC decomposition. However, this contribution is believed to be very

low in our experiment considering the young age of the rice plants. An estimate of the plant-derived DOC contents in our pots based on literature (Lu et al., 2004; Nguyen, 2003) was as low as $14 \, \text{mg} \, \text{C} \, \text{kg}^{-1}$ soil, with therefore indeed low potential contributions to soil C emissions, in agreement with observations of Yuan et al. (2012) at 41 d, and comparison of our emissions from maize-amended pots with and without rice plants. SOC-derived emission estimates based on isotopic mixing (Eq. 3) are henceforth assumed to be derived from native SOC decomposition.

2.8   Statistical analyses

Statistical tests were conducted with R 3.6.1 for Windows (R Core Team, 2019), with $P = 0.05$ taken as the default significance level. Mostly, results for both soils (Balina and Sonatala) were analysed separately. The effect of irrigation treatment (starting from 11 DAT in the case of time series) on measured variables was assessed by two-way or three-way ANOVA or by Kruskal-Wallis tests with irrigation treatment and maize amendment as factors, and DAT as a third factor in the case of time series. In the case of interaction between the irrigation management and maize amendment factors, the tests were repeated for data from maize-amended versus control pots separately. In the case of interaction with DAT, differences between irrigation treatments were also assessed separately for individual points in time. Student's t-tests or Wilcoxon-Mann-Whitney tests were used to test the significance of priming effects for dissolution (i.e. comparing averages after integration over time from SOC-derived DOC in maize-amended and unamended soil) as well as emission (i.e. comparing SOC-derived emissions from maize-amended and unamended soil), to compare if dissolution and emission priming effects differed from one another, and to test if the $\delta^{13}C$ of CH_4 and CO_2 differed when emissions were dominated versus not dominated by ebullition.

3   Results

3.1   Redox potential and iron reduction

Within one to two weeks the soil redox potential (E_h) generally decreased to ca. -250 to $-150 \, \text{mV}$ (Fig. 1). The effect of the irrigation treatment on E_h seemed to depend on both soil and addition of maize, but temporary increases in E_h were observed upon soil drying. The E_h was more responsive to wetting and drying at 4 cm than at 12 cm depth. Maize addition did not particularly induce a lower E_h, irrespective of soil type, though it did stimulate reductive dissolution of Fe ($P < 0.001$ for both soils between 2 and 11 DAT) during the observed initial increase in dissolved Fe equivalents (Fig. 2). Similar trends were observed for Mn^{2+} (data not shown). When maize was added, Fe concentrations after 11 DAT dropped to lower levels in the case of AWD compared to CF ($P < 0.001$ for Balina and $P = 0.04$ for Sonatala). In

control pots without maize shoots added, there was no significant effect of irrigation treatment on the decrease in dissolved Fe.

3.2   Exchangeable NH_4^+, pH and rice plant biomass

Concentrations of exchangeable NH_4^+ increased until 11 DAT, partly owing to the initial urea additions of 39 and $47 \, \text{mg} \, \text{N} \, \text{kg}^{-1}$ to the Balina and Sonatala soils, respectively. Thereafter they remained more or less constant (Fig. S2 in the Supplement). Overall, exchangeable NH_4^+ levels reached in the Balina soil were about double those for Sonatala. The effects of the presence of rice plants or irrigation treatment were at times significant but tended to be small and are not further discussed in detail. NH_4^+ levels were mostly significantly higher with maize added than without, probably because of net N mineralisation. Concentrations of NO_3^- were always minor, i.e. below $1 \, \text{mg} \, \text{N} \, \text{kg}^{-1}$ (data not shown).

The soil solution pH of the Balina soil increased from an initial value of 6.1 to 6.8 (± 0.1) after 17 DAT and then remained constant onwards (Fig. S3). In the Sonatala soil, pH remained nearly constant throughout (initially 6.9, then 6.8 ± 0.1 starting from 17 DAT).

After 42 d, rice plants had reached a height of $76 \pm 5 \, \text{cm}$, with an above-ground dry biomass of $2.7 \pm 0.7 \, \text{g}$ per pot and a below-ground dry biomass of $0.6 \pm 0.3 \, \text{g}$ per pot. There was no effect from soil type or irrigation practice on the above-ground biomass. Below-ground biomass was not affected by soil type either but was significantly higher for plants grown under CF ($0.8 \pm 0.3 \, \text{g}$ per pot) than under AWD ($0.4 \pm 0.2 \, \text{g}$ per pot) ($P < 0.001$).

3.3   Organic carbon dissolution

Extremely high total DOC peaks were reached 1 to 2 weeks after submergence in all treatments (maximally 1912 and $1329 \, \text{mg} \, \text{C} \, \text{kg}^{-1}$ soil for maize-amended Balina and Sonatala soils, respectively), which were also much higher with maize added ($P < 0.001$ for time series until 11 DAT). Not only maize-derived DOC (data not shown), but also enhanced dissolution of native SOC caused these DOC peaks (Fig. 3). Over the course of the experiment, SOC-derived DOC was indeed higher with maize added than without in both soils, which was significant for Sonatala under CF (averaged concentrations over the course of the season: $P = 0.003$). In other words, there was a positive priming effect on SOC dissolution (i.e. stimulation) during most of the incubation. This priming effect was irrespective of the irrigation regime in the Balina soils ($P = 0.97$), while it was stronger under CF than under AWD in the Sonatala soils ($P < 0.001$) starting from 16 DAT. Priming of SOC dissolution was also stronger for Sonatala than for Balina ($P < 0.001$). Dissolution of maize-C was, in contrast, nearly identical in both soils, with an average peak concentration of $811 \, \text{mg} \, \text{C} \, \text{kg}^{-1}$ at 7 DAT, fol-
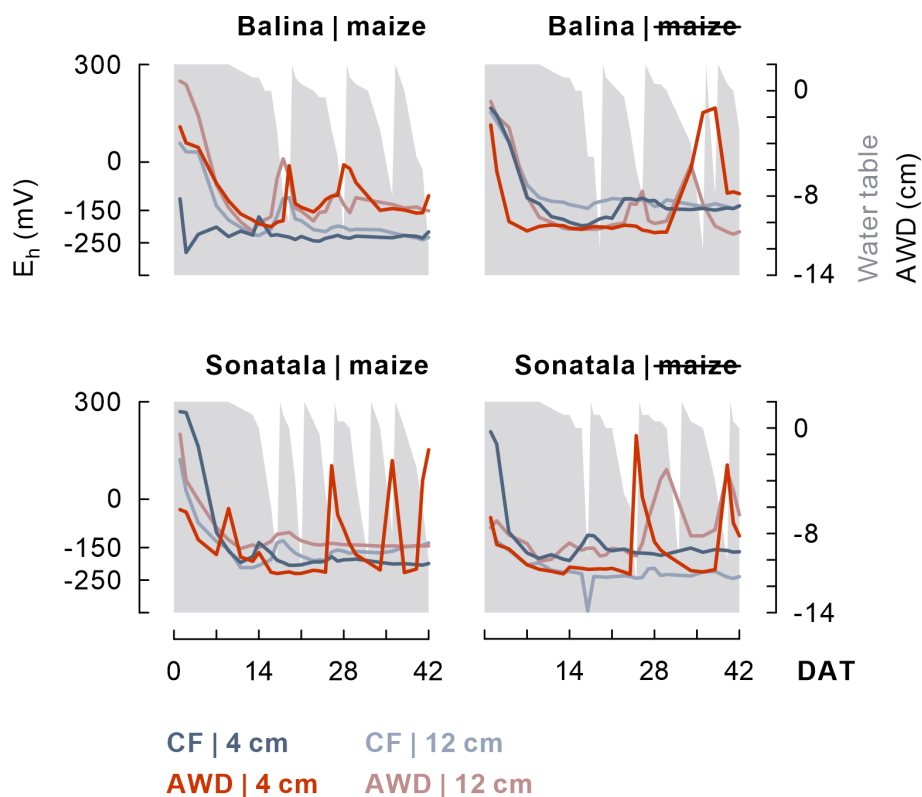
Figure 1. After a general initial drop, the soil redox potential (E_h) (values per replicate at 4 and 12 cm depth) depended to some extent on the irrigation treatment in both paddy soils (Balina and Sonatala). Water table fluctuations in the case of AWD are indicated in grey.
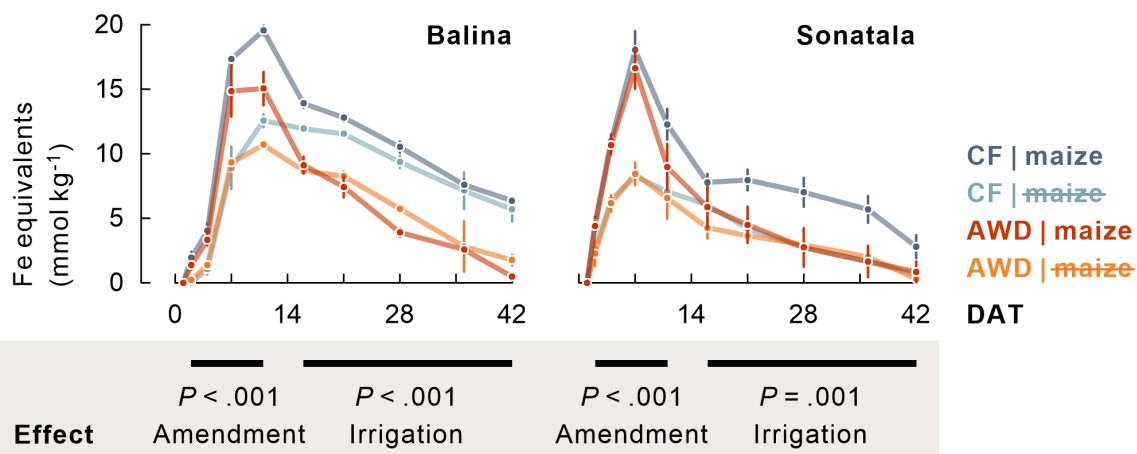
Figure 2. Maize addition stimulated reductive dissolution of Fe (average $\pm$ standard error of three or two replicates, calculated as sum of dissolved Fe and increments in dissolved Ca and Mg), and Fe^{2+} disappeared to a larger extent from the soil solution under AWD than under CF in maize-amended pots. The P-values indicate the effect of maize addition on the time series between 2 and 11 DAT, and the effect of irrigation treatment in maize-amended pots on the time series between 16 and 42 DAT.

lowed by a near-complete disappearance of maize-derived DOC around 16 DAT (< 79 mg C kg^{-1}) (data not shown).

3.4   Gaseous carbon emission

Gaseous C emissions consisted on average of 31 % CH$_4$ and 69 % CO$_2$. Irrigation management did not affect the cumulative C emissions (CO$_2$ + CH$_4$), and neither CO$_2$ nor CH$_4$ emissions individually in the case of the Balina soil (Fig. 4). For the Sonatala soil, however, the effect of irrigation depended on whether or not the soils were amended with maize. With maize added, total gaseous C ($P = 0.04$) and CH$_4$ ($P = 0.03$) emissions were higher under CF than AWD, but CO$_2$ emissions were not significantly affected. On the other hand, in unamended Sonatala pots, total C ($P = 0.02$) and CO$_2$ ($P = 0.003$) emissions were higher under AWD than under CF, while CH$_4$ emission was not significantly different. CH$_4$ and CO$_2$ emissions were in general higher with maize added than without ($P < 0.001$). Total gaseous C emission was furthermore lower from the Sonatala than from the Balina soil ($P = 0.01$). In general, gaseous C emissions were similar with or without living rice plants in pots with maize added ($P = 0.23$). Both the addition of maize ($P < 0.001$) and the implementation of CF ($P < 0.001$) led to more gaseous C emission via ebullition: 60 ± 15 % of C was emitted through ebullition under CF compared to only 37 ± 11 % under AWD. With maize added, 56 ± 16 % of C was emitted via ebullition, compared to only 37 ± 12 % for unamended control pots.

In addition, the course of CH$_4$ and CO$_2$ fluxes seemed linked to the wetting and drying cycles under AWD (Fig. 5) as (i) more gaseous C was emitted when the soils dried; (ii) most ebullition transport occurred when the water table was at the soil surface, usually a few days before rewetting; and (iii) δ^{13}C-CO$_2$ signatures increased as water tables declined.

CH$_4$ and CO$_2$ emissions were clearly more enriched in ^{13}C when maize was added (Figs. 6 and S1), evidently indicating the contribution of maize C decomposition. The δ^{13}C of CH$_4$ increased to a distinct maximum at 16 DAT (Balina) or 13 DAT (Sonatala) (Fig. S1). In control pots without maize, the δ^{13}C of emitted CH$_4$ was higher for Sonatala than for Balina, in line with the difference in δ^{13}C of native SOC (Fig. 6). Implementation of AWD to control pots resulted in higher δ^{13}C signatures of emitted C (CO$_2$ + CH$_4$) compared to CF ($P = 0.02$). Isotopic fractionation took place during mineralisation of maize and native SOC into CH$_4$ and CO$_2$ (Figs. 6 and S1). Lastly, the δ^{13}C of CH$_4$ ($P = 0.02$) and CO$_2$ ($P < 0.001$) was on average 2.62 ‰ and 3.45 ‰ higher when emissions were dominated by ebullition than when they were dominated by plant transport and diffusion (data not shown).

3.5   Source partitioning of emitted CO$_2$ and CH$_4$

The fractions of maize-derived and SOC-derived CO$_2$ and CH$_4$ were calculated based on isotopic mixing, considering the δ^{13}C of SOC and maize, and accounting for isotopic fractionation during decomposition to CO$_2$ and CH$_4$ (Fig. S1). Surprisingly, the δ^{13}C-CH$_4$ of emissions from an ancillary C$_4$-reference incubation with only maize and no native SOC as OC (δ^{13}C-CH$_{4 \,|\, \text{maize}}$) was usually lower (instead of higher) than from control pots with no maize added (δ^{13}C-CH$_{4 \,|\, \text{SOC}}$) (Fig. S1). The difference in isotopic signature between maize and SOC was therefore not further reflected in a likewise difference in δ^{13}C of the CH$_4$ derived from both sources. In addition, the δ^{13}C-CH$_4$ was often higher than the δ^{13}C of both endmembers in the case of the maize-amended Balina and Sonatala soils. For these reasons, the fraction of CH$_4$ emission derived from maize ($f_{\text{CH}_4 \,|\, \text{maize}}$) could unfortunately not be reliably deduced, in spite of the high measurement frequency across the experiment. As a best estimate, we therefore assumed that the contributions of emitted CH$_4$ from maize decomposition would equal $f_{\text{CO}_2 \,|\, \text{maize}}$. After about 16 DAT, this derived fraction of maize-derived CO$_2$ increased to about 100 % for most of the pots (data not shown). The calculated cumulative maize-derived CO$_2$ and CH$_4$ emissions ranged between 1737 and 2757 mg C kg^{-1}, in most cases exceeding the theoretical maximum of 1898 mg C kg^{-1}, i.e. the amount of initially added maize. Therefore, in those cases we instead consider complete degradation of all the initially added maize for further calculation of cumulative SOC-derived C emissions, which seems reasonable considering the complete disappearance of dissolved maize-C around 16 DAT (Fig. 3), and the fact that the δ^{13}C of SOC was not higher after than before the experiment (Fig. 6).

The effect of maize addition on SOC-derived C emission was negligible in the case of Balina, while for Sonatala, it depended on the irrigation treatment (Fig. 7). CF irrigation led to a positive priming ($P = 0.05$) (i.e. stimulated SOC-derived gaseous C emissions upon maize addition), whereas negative priming on SOC-derived C emissions in maize-amended soil occurred under AWD ($P = 0.01$). In addition, priming on SOC-derived C emission was higher under CF than AWD for Sonatala ($P = 0.004$).

4   Discussion

4.1   Are Fe reduction and SOC dissolution linked?

With the onset of anaerobic conditions, exemplified by a decreasing soil E_h (Fig. 1), there was a strong increase in dissolved Fe (Fig. 2), which shows that part of the soil Fe was quickly reduced and dissolved. At the same time, DOC concentrations increased rapidly until maxima at 11 DAT (Balina) and at 4–7 DAT (Sonatala). The synchrony of OC
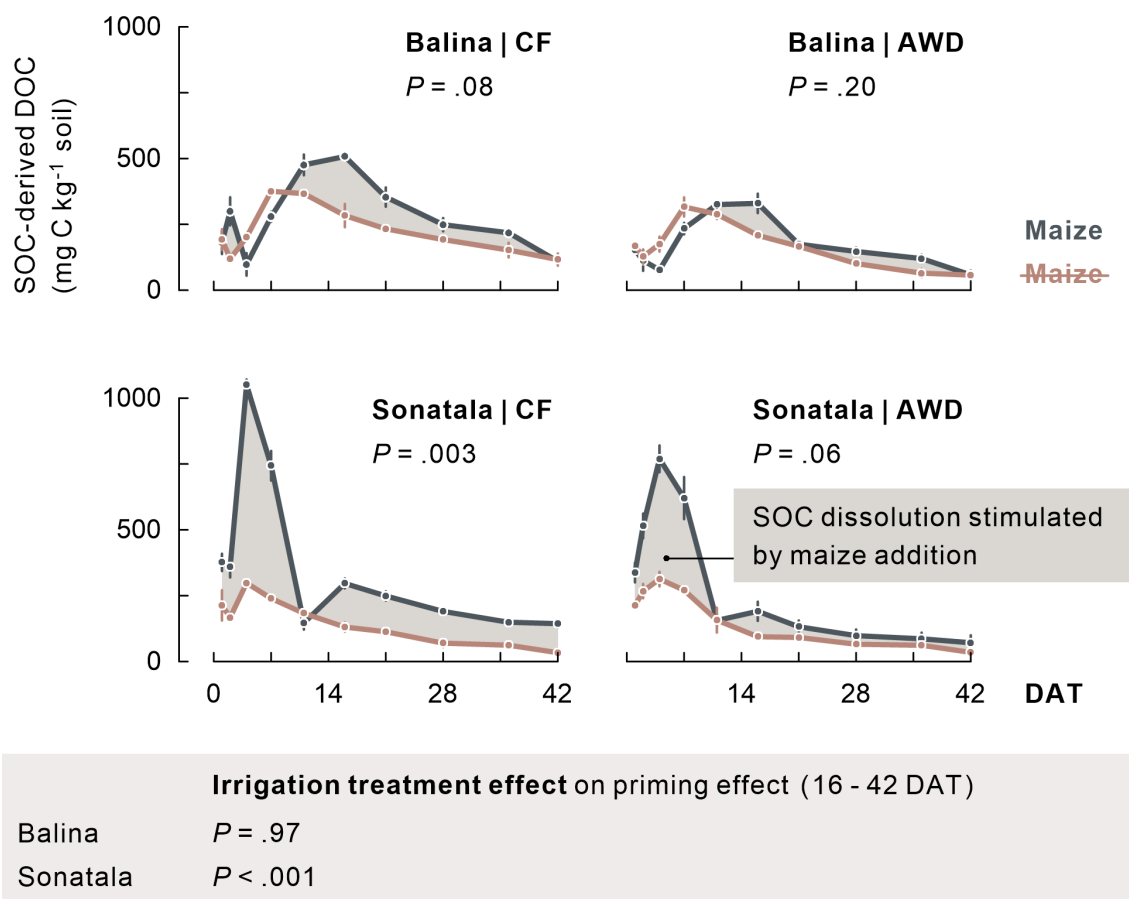
Figure 3. Dissolution of native SOC (average ± standard error of three or two replicates) was stimulated upon addition of maize, as it appears when comparing with DOC in unamended pots (indicated in brown). P-values of the priming effect significance using average values per season (in the figure) and of the irrigation treatment effect on the priming effect time series starting from 16 DAT (below the figure) are indicated.

dissolution and Fe reduction patterns prompts the question of whether both processes are physically linked. Said-Pullicino et al. (2016) suggested that observed DOC increases could have been due to the co-release of OC during reductive dis-
5 solution of Fe oxides. With an adapted $NH_2OH \cdot HCl$ extraction procedure, we estimated that the amount of C bound onto reducible Fe^{3+} minerals was $1367 \, mg \, kg^{-1}$ for Balina and $2000 \, mg \, kg^{-1}$ for Sonatala (Fig. 7), constituting approximately 9 % of SOC. Release of this substantial Fe-bound
10 SOC pool might have been a crucial source of DOC, but other important C dissolution pathways need to be considered as well. Firstly, a rise in pH upon soil reduction under anaerobic conditions can lead to SOC desorption (Grybos et al., 2009) but this mechanism was most likely unimportant
15 here because of the initially high pH that did not considerably change during the incubation, especially in the Sonatala soil (Fig. S3). Secondly, rhizodeposition from rice plants can also cause an increase in DOC (Said-Pullicino et al., 2016; Lu et al., 2004). However, based on values from Lu et al. (2004)
20 and Nguyen (2003), maximum DOC contents derived from

rice plant photosynthates in our soils are estimated to be around $14 \, mg \, C \, kg^{-1}$ soil, which is again limited since rice plants were juvenile during the experiment. Next to the direct contribution of rhizodeposition, some root photosynthates (e.g. oxalate and citric acid) can also indirectly increase DOC 25 levels by promoting the release of Fe-bound SOC in the rhizosphere through their strong metal-complexing capacity (Keiluweit et al., 2015; Yu et al., 2017). This mechanism was, however, likewise restricted here considering its local impact and the juvenile age of the rice plants. Lastly, fermentation of 30 SOC and added maize into water-soluble organic metabolites must obviously have largely contributed to the accumulation of DOC. As such, based on DOC patterns alone it is not possible to unequivocally state that reductive dissolution of Fe with co-release of Fe-bound OC drove DOC build-up. 35

The stable isotope approach then allowed to discriminate between SOC-derived and maize-derived DOC. Maize addition roughly doubled dissolution of native SOC (Fig. 3). Since also Fe reduction was stimulated upon the addition of maize (Fig. 2), it seems likely that, next to stimu- 40
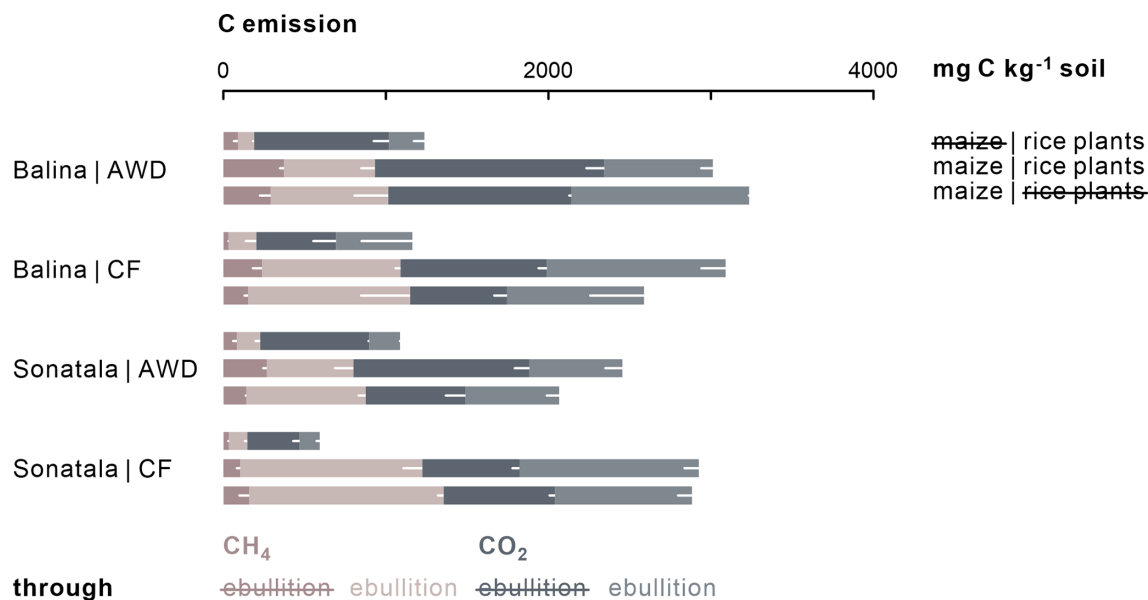
Figure 4. Total cumulative C emissions (average − standard error of three or two replicates, represented by white bars) from maize-amended soils were two to fourfold of those from unamended pots ($P < 0.001$). In addition, relatively more C was emitted through ebullition under CF ($P < 0.001$) and upon addition of maize shoots ($P < 0.001$).

lated SOC fermentation, promoted SOC co-release upon enhanced Fe reduction was indeed a potentially important priming mechanism, in agreement with the observations of Ye and Horwath (2017) and Bertora et al. (2018). Dissolution of all Fe-bound SOC present might produce a DOC peak of approximately $584 \, \text{mg C kg}^{-1}$ (Balina) or $854 \, \text{mg C kg}^{-1}$ (Sonatala), when roughly comparing it to the DOC production and the DOC mineralisation kinetics of maize (where we know that $1898 \, \text{mg added maize-C kg}^{-1}$ yielded a peak of $811 \, \text{mg C kg}^{-1}$ maize-derived DOC). While just a tentative approximation, it becomes clear that the magnitude of SOC-derived DOC peaks suffices to explain the observed increment in SOC-derived DOC with maize added (Fig. 3).

Comparison of the $\delta^{13}C$ of Fe-bound SOC and DOC could further contribute to the quantification of DOC derived from pedogenic Fe. Attempts to measure $\delta^{13}C$ of $NH_2OH \cdot HCl$ extracts directly by means of FIA-IRMS unfortunately failed owing to interference of the matrix, even after trying to remove $NH_2OH \cdot HCl$ and Cl^- by oxidation and precipitation. However, from the difference in soil $\delta^{13}C$ before and after $NH_2OH \cdot HCl$ extraction, the $\delta^{13}C$ of Fe-bound SOC could be estimated indirectly with reasonable uncertainty, i.e. $-27.8 \pm 0.7 \, \permil$ for Balina and $-23.3 \pm 3.3 \, \permil$ for Sonatala. Apparently, Fe-C was a bit more depleted in ^{13}C than bulk SOC (Table 1). In control pots, with only Fe-bound SOC and bulk SOC as C sources, $\delta^{13}C$ signatures of DOC were also lower, supporting the assumption that DOC partly consists of released Fe-bound C. To robustly derive the contribution of potentially enhanced Fe-bound C release to stimulated SOC dissolution in maize-amended pots, three-source partitioning

with an additional contrast in ^{14}C next to the existing variation in $^{13}C/^{12}C$ would be required (when rice plants can be disregarded).

4.2 Extent of priming of SOC mineralisation as compared to SOC dissolution

Our objective was then to more specifically assess whether the stimulated SOC dissolution would be reflected in enhanced native SOC-derived mineralisation to CO_2 and CH_4. As it turns out, native SOC-derived gaseous C (i.e. $CO_2 + CH_4$) emissions were not significantly modified by the addition of maize in the Balina soil (Fig. 7). For Sonatala, there was however positive priming (i.e. more SOC-derived C emissions in soil with maize addition than in soil without) under CF irrigation ($P = 0.05$) and a negative priming under AWD ($P = 0.01$). For the CF treatment, the discussed stimulation of native SOC dissolution, probably partially by enhanced co-release of Fe-bound SOC, may have promoted SOC mineralisation upon maize addition. Yet, priming effect coefficients were overall larger for dissolution than for gaseous C emission in Sonatala ($P < 0.001$) and not significantly for Balina ($P = 0.18$) (Fig. 8), which suggests that the enhanced SOC dissolution would not have resulted in a proportionate SOC mineralisation. Hanke et al. (2013) likewise found that extra solubilised C after paddy soil submergence did not cause a proportional rise in gaseous C emissions. They assumed that the availability of alternative terminal electron acceptors was limiting mineralisation. However, as SOC mineralisation (indicated by SOC-derived CO_2 and CH_4 emission) was actually slower under AWD than
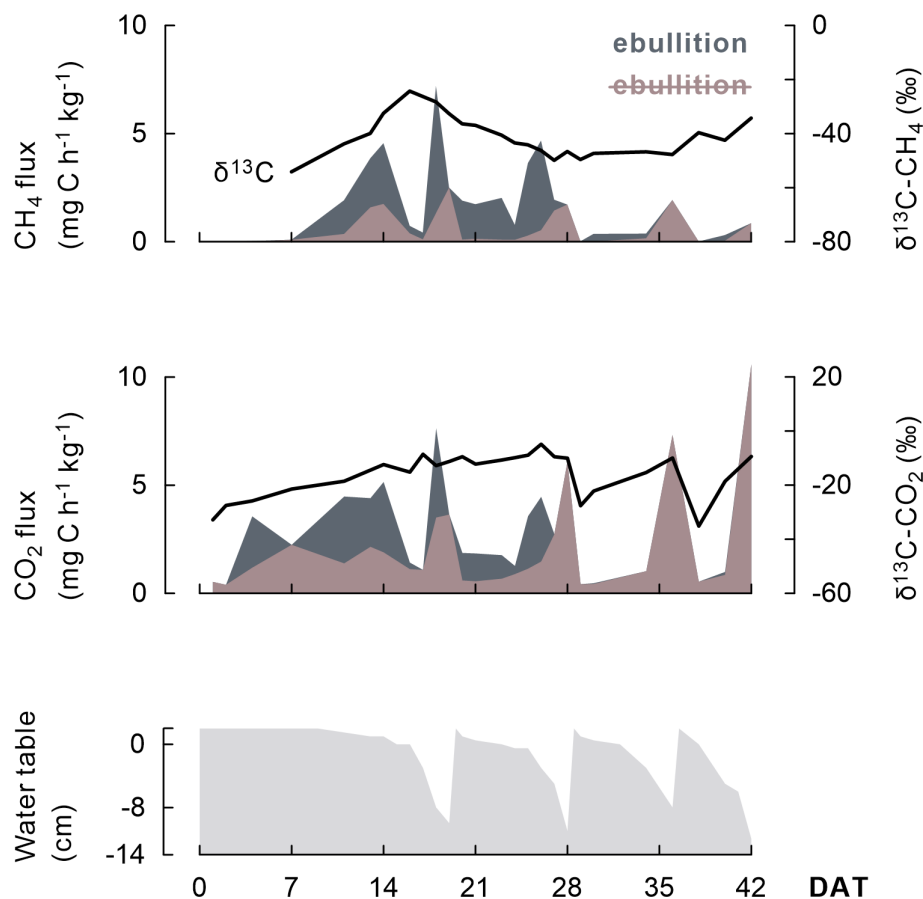
Figure 5. As this representative example (one replicate of the Balina soil with maize shoots added and under AWD) shows, the progression of CH_4 and CO_2 emission fluxes (shaded) and even their $\delta^{13}C$ signature (black line) appear linked to the change in water table depth under AWD. A distinction is made between emission fluxes dominated by ebullition (dark grey shaded) versus by diffusion or plant transport (brown shaded). Transport through ebullition was particularly considerable when rice plants were juvenile.
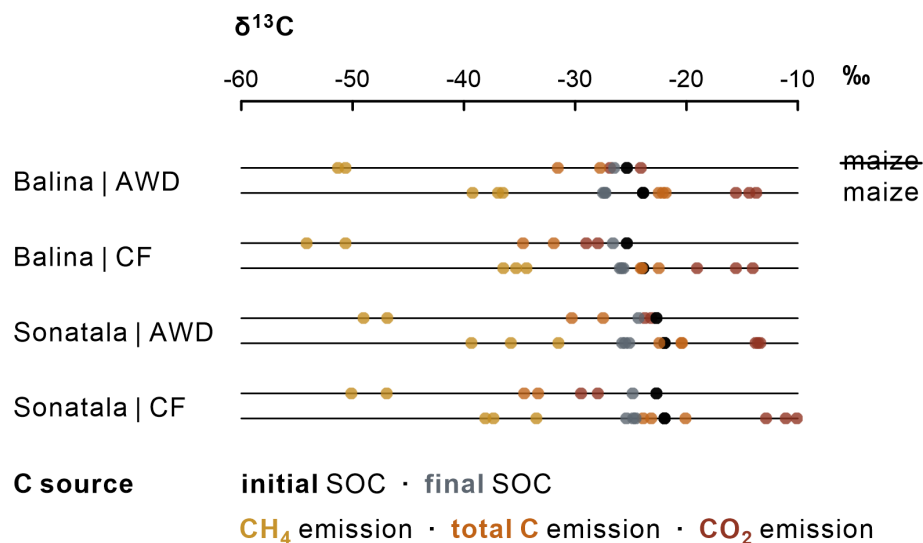
Figure 6. In the absence of maize, the overall $\delta^{13}C$ signature of C emissions (values per replicate) was lower (i.e. less ^{13}C-enriched) under CF than under AWD ($P = 0.02$).

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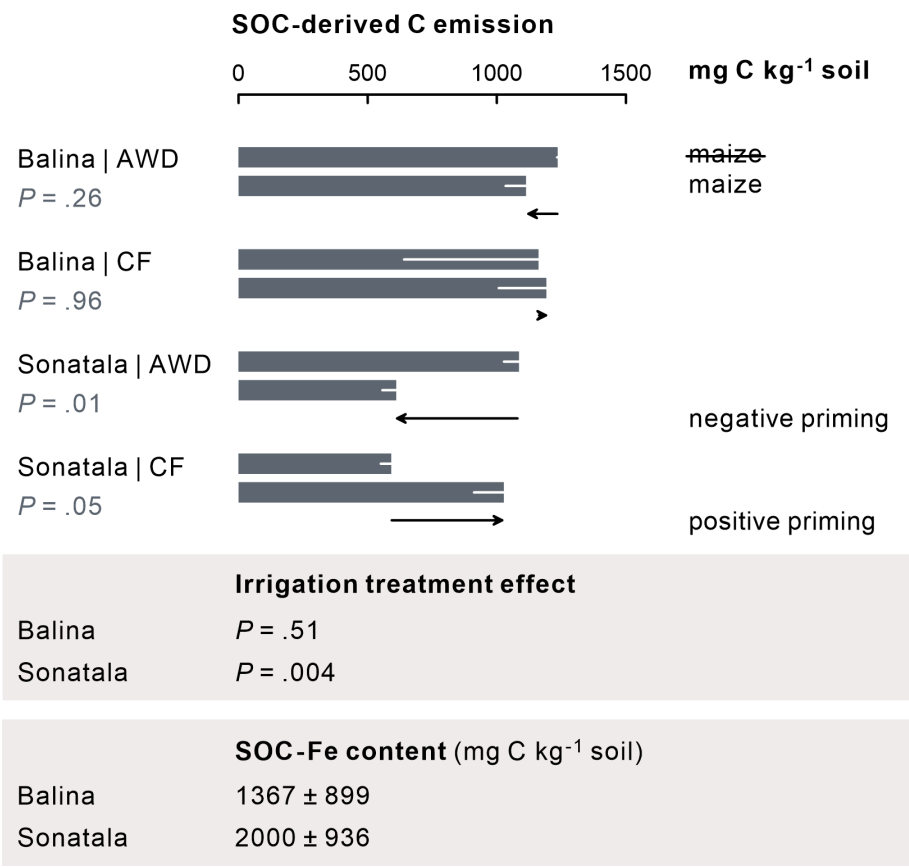
Figure 7. For the Sonatala soil, maize addition led to positive priming of gaseous C ($CO_2 + CH_4$) emission under CF and negative priming under AWD. P-values of the priming effect significance (in the figure) and of the irrigation treatment effect on the priming effect (below the figure) are indicated. Standard errors (three or two replicates) are indicated. Fe-bound SOC levels (average $\pm$ standard error, estimated by means of $NH_2OH \cdot HCl$) exceed the positive priming effect of Sonatala under CF, so stimulated Fe-C dissolution and emission has the potential to explain this priming.

under CF in the Sonatala soil, electron acceptor availability must not really have limited priming of SOC-derived C emission. It instead seems more likely that preferences of microorganisms that decompose DOC explain the observed smaller priming of soil C emission compared to dissolution. DOC-decomposing microorganisms must have preferred the abundantly available maize-derived DOC at the expense of SOC-derived DOC mineralisation in maize-amended soils. This hypothesis is supported by the faster removal of maize-derived compared to SOC-derived DOC (data not shown). Maize shoots and native SOC obviously differ in their biochemical quality, where SOC in paddy soils is generally quite aromatic with an accumulation of hardly degradable lignin components (Olk et al., 2002; Z. Zhou et al., 2014). In fact, phenol accumulation in paddy soil in part explains their larger topsoil OC content compared to upland counterparts (Chen et al., 2021). The energy yield of oxidation of such reduced SOC compounds (phenols and lipids) becomes low when coupled to Fe reduction (Keiluweit et al., 2017), and therefore it is logical that maize-derived DOC would

have been preferred over SOC-derived DOC as a microbial substrate. Lastly, enhanced Fe reduction might also abiotically mediate positive priming of native SOC mineralisation through Fenton reactions that lead to the production of reactive hydroxyl radicals (Yu and Kuzyakov, 2021). However, as priming effect coefficients for gaseous C emissions were low compared to those for dissolution (Fig. 8), and since the relative contribution of CO_2 to gaseous C emissions in maize-amended pots was lower than in control pots (Fig. 4), it seems unlikely that the potentially enhanced production of hydroxyl radicals would have played a considerable role in stimulating SOC decomposition in maize-amended pots.

4.3 Influence of irrigation management and soil type

Furthermore, we wanted to verify whether irrigation management plays a role in the priming effect on SOC-derived dissolution and mineralisation. It is of interest that the enhancement of SOC dissolution by maize addition was stronger under CF than AWD, especially for the Sonatala soil (Fig. 3). Under permanently flooded conditions, the addition of maize
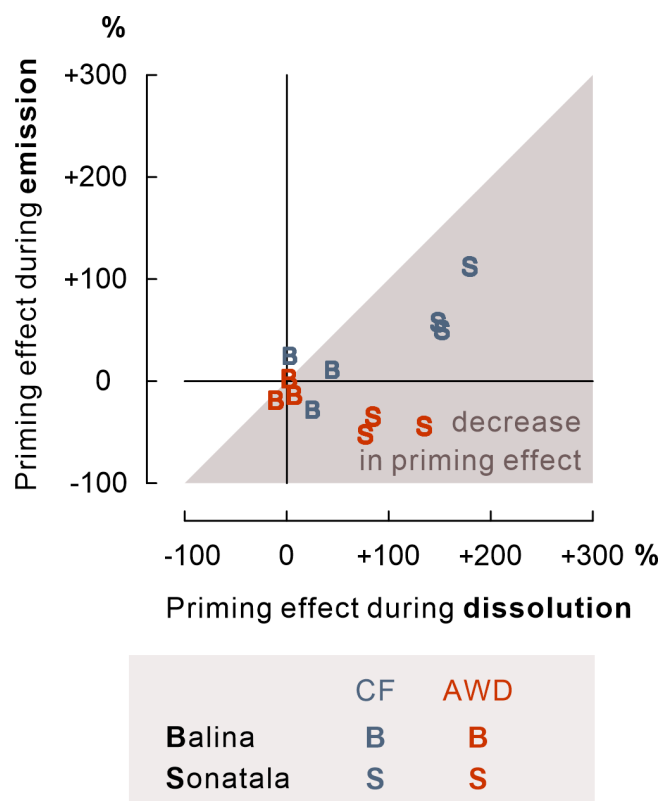
Figure 8. For Sonatala ($P < 0.001$) and not significantly for Balina ($P = 0.18$), the priming effect on gaseous C ($CO_2 + CH_4$) emission was lower than on SOC dissolution.

was probably more effective in promoting anaerobic microbial fermentation resulting in dissolution of SOC, as the biomass of anaerobes could grow under less disturbed conditions than in the case of a regime with repeated soil drying. This irrigation treatment effect on DOC priming was also manifested in higher priming on SOC emission under CF than AWD in the case of Sonatala ($P = 0.004$) (Fig. 7). Furthermore, the decrease in priming effect was slightly more pronounced for AWD than CF for Sonatala ($P = 0.08$) (Fig. 8). The negative priming of gaseous C emission under AWD in Sonatala was therefore possibly not only due to a lesser stimulation of SOC-derived dissolution under AWD than under CF. On top of that, the microbial community under AWD might have been less adapted to anaerobic mineralisation than under CF, resulting in an even larger preference for maize-derived than SOC-derived DOC.

Furthermore, the effect of irrigation management on overall gaseous emissions was rather limited, especially for Balina soil, and surprisingly partially determined by ebullition. With no maize added, CO_2 emission ($P = 0.003$) was lower under CF than under AWD for Sonatala (Fig. 4). Indeed, SOC mineralisation is often retarded under anaerobic conditions (Kögel-Knabner et al., 2010; Sahrawat, 2004). However, CH_4 effluxes were also surprisingly lower under

CF, while the more reducing conditions compared to under AWD were expected to stimulate CH_4 production (Jiang et al., 2019; Peyron et al., 2016). One cause for this contrasting outcome may be the often considerable temporal storage of produced CH_4 and CO_2 in submerged CF pots. Indeed, transport of CH_4 and CO_2 in paddy soils usually proceeds predominantly via rice plant aerenchyma, whereas diffusion-based outgassing is retarded by permanently standing water. With rice plants still juvenile and relatively limited CO_2 and CH_4 production in the unamended pots, it is thus possible that a larger part of these gases was stored dissolved or as gas bubbles in the CF pots. With AWD on the other hand, the capacity of the standing water to store gases decreases as the water table goes down regularly (Green, 2013; Tokida et al., 2013). Transport of the produced CH_4 and CO_2 to the atmosphere was indeed clearly facilitated by episodic flushes, e.g. through ebullition, when the water table decreased (Fig. 5), in line with the observations of Yagi et al. (1996) and Han et al. (2005). With maize added, on the other hand, CH_4 emission ($P = 0.03$) was 53 % higher under CF than AWD in the case of the Sonatala soil. The much faster production of CH_4 compared to in unamended pots likely led to oversaturation of the dissolved CH_4 pool, resulting in less restricted CH_4 emission. For Balina, on the contrary, the irrigation regime did not significantly affect CO_2 and CH_4 emission. This all in all low effect of water management on gaseous C emissions in the Balina soil was in line with its limited impact on the progression of E_h at 12 cm depth (Fig. 1), pH (Fig. S3) and the fact that exchangeable NH_4^+ levels (Fig. S2) only differed during the final two weeks.

Lastly, the contrast in Fe-bound SOC estimates between Sonatala and Balina corresponded with the difference in their respective $SOC : Fe_{ox}$ ratios (4.7 versus 1.7), and moreover with the higher SOC-derived DOC levels and stronger priming of SOC dissolution in Sonatala compared to Balina. Ye et al. (2016) found that a soil with less electron acceptors gave rise to stronger priming on CH_4 emission in a laboratory incubation under submerged conditions, which corresponds with our findings on C ($CH_4 + CO_2$) emission priming for the CF treatment. However, the higher $SOC : Fe_{ox}$ ratio cannot immediately explain the more pronounced decrease in priming effect on C emission compared to dissolution for Sonatala than for Balina ($P < 0.001$), except if the DOC derived from pedogenic Fe would be more recalcitrant, leading to a bigger preference of DOC decomposers for maize-derived DOC. Soil type also influenced temporal evolutions of biochemical parameters, but only to some extent. For Sonatala, redox-related processes took place three to five days earlier than for Balina, such as: (i) the initial decline of E_h (Fig. 1); (ii) the peak in dissolved Fe (Fig. 2); (iii) the peak in DOC, in particular in SOC-derived DOC (Fig. 3); and (iv) the "peak" in emission $\delta^{13}C$-CH_4 signatures (Fig. S1). This probably all relates to the higher electron surplus (i.e. higher amount of electron donors as compared to the main electron acceptor Fe^{3+}) in combination with a

lower Fe_{ox} content for Sonatala, with its higher $SOC:Fe_{ox}$ ratio, resulting in a faster decrease in soil E_h. In Sonatala, it is likely that microbes quickly needed to couple OC oxidation with the reduction of alternative electron acceptors other than Fe^{3+}, so that E_h dropped below the level enabling CH_4 production ($-150\,\text{mV}$) after 4 d on average for Sonatala while only after 9 d for Balina. In general, the impact of irrigation management and maize residue addition on DOC release and emissions seemed to depend on the soil type (in particular the $SOC:Fe_{ox}$ ratio), and soil type therefore seems to be an equally important factor controlling the progression of reductive processes.

4.4 Using an isotope mass balance to infer native SOC mineralisation in paddy soils

Because of different types of isotopic fractionation, most importantly caused by microbial discrimination for or against ^{13}C as a substrate, source partitioning of gaseous emissions was not straightforward. Unlike the results of Conrad et al. (2012) but in agreement with the observations of Ye et al. (2016), isotopic fractionation was different for maize than for SOC. For example, we observed a very consistent initial enrichment in ^{13}C-CH_4 over time until strikingly high $\delta^{13}C$-CH_4 signatures of $-22.4\pm2.9\,‰$ (Balina; 16 DAT) and $-26.4\pm4.4\,‰$ (Sonatala; 13 DAT), with thereafter ^{13}C depletion (Figs. S1 and 5). As such trends were inexistent for $\delta^{13}C$-CO_2, these $\delta^{13}C$ evolutions are likely the result of gradually changing dominant CH_4 production pathways. Indeed, CH_4 can be produced with acetate, CO_2 or other methylated compounds as precursors. Among any of these precursors, the lowest fractionation (i.e. resulting in CH_4 with the least negative $\delta^{13}C$) occurs with the acetate-dependent pathway, resulting in a $\delta^{13}C$-CH_4 as high as $-27\,‰$ (relative to VPDB) (Conrad, 2005). The even further enrichment of ^{13}C-CH_4 in the maize-amended plots can be explained by the higher $\delta^{13}C$ of maize (Fig. 6). Another explanation for the constantly changing $\delta^{13}C$-CH_4 could be that the contributions of various gas transport mechanisms and associated isotopic fractionation would have changed over time. The $\delta^{13}C$-CO_2 increased as water tables declined during AWD cycles, particularly towards the end of the experiment (Fig. 5). In the unamended pots, the $\delta^{13}C$ of CO_2 ($P = 0.02$) was also higher under AWD than under CF (Fig. 6). Both observations clearly suggest that change in transport fractionation affected the evolving $\delta^{13}C$-CO_2 and $\delta^{13}C$-CH_4 patterns. Interestingly, continuous CRDS-based recording of CO_2, CH_4 and their $\delta^{13}C$ allowed to discern sudden ebullition events over the course of a measurement. Emissions dominated by ebullition indeed displayed higher $\delta^{13}C$-CH_4 but also higher $\delta^{13}C$-CO_2 signatures than emissions during which ebullition appeared absent. The evolving contribution of diffusion and plant-mediated versus ebullition transport throughout the experiment thus must have altered the $\delta^{13}C$ of emitted CO_2 and CH_4. In the case of CH_4, it is indeed well known that

$\delta^{13}C$ isotopic fractionation not only takes place during plant-mediated transport (Zhang et al., 2014, 2015) but also during gas bubble formation (Zhang et al., 2014). Our results suggest that the $\delta^{13}C$ of emitted CO_2 is also impacted by the contribution of ebullition transport. Using pots without maize added, we quantified the overall isotopic fractionation between SOC and SOC-derived CO_2 and CH_4. We then also attempted to quantify isotopic fractionation between maize-C and maize-derived CO_2 and CH_4 by means of an ancillary incubation experiment (Fig. S1), but this was only successful for CO_2. In addition, maize addition lifted the contribution of ebullition towards 56 % of all emitted gaseous C compared to 37 % without maize. Because of that, the estimated $\delta^{13}C$-$CO_{2\,|\,SOC}$ and $\delta^{13}C$-$CH_{4\,|\,SOC}$ used in Eq. (3) were perhaps not representative for the maize-amended pots, since the proportion of ebullition transport was particularly large in our experiment. However, decreasing ebullition by retarding mineralisation, e.g. by lowering soil temperatures or by working with a less labile substrate than the finely ground maize, would after all have but a limited impact. To improve the source partitioning of emissions, the use of exogenous OC with a more strongly contrasting $\delta^{13}C$ would help the most, as then the relative effect of isotopic fractionation declines.

5 Conclusion

Using a stable isotope approach, it was confirmed that the addition of high-quality OC like maize shoots stimulates Fe reduction and dissolution of native SOC. The synchrony of both processes suggests that priming of SOC dissolution after addition of maize could result from increased net co-release of OC during reductive dissolution of Fe oxides, next to enhanced fermentation of native SOC to DOC. In support of this hypothesis, the pool of Fe-bound C (1.3–$2.0\,\text{g C kg}^{-1}$), estimated by means of reduction with $NH_2OH \cdot HCl$, was large enough to explain this. Moreover, we found that maize addition stimulated SOC dissolution more strongly in the soil with the highest $SOC:Fe_{ox}$ ratio and Fe-bound SOC content. However, positive priming of SOC mineralisation into CO_2 and CH_4 only occurred under CF in one out of the two investigated soils. Priming of SOC dissolution therefore does not necessarily result in proportional priming of native SOC-derived emissions, most likely owing to an overall preference of microbes for maize-derived DOC in anaerobic conditions. However, it turned out that it is particularly difficult to assess priming effects on native SOC mineralisation in paddy soils, because source partitioning of C emissions based on a stable isotope mass balance is inherently complicated due to various isotopic fractionation effects. Nevertheless, our study indicated that particularly in soil with a high $SOC:Fe_{ox}$ ratio, the addition of labile OC can result in substantial extra dissolution and mineralisation of native SOC under CF irrigation, with a possible adverse

impact on the SOC balance in the longer term. The adoption of water-saving irrigation can instead successfully decrease the stimulation of SOC dissolution caused by OC addition and inhibit the positive priming of SOC mineralisation.

Code availability. The codes used in the preparation of this paper are available from the corresponding author upon request.

Data availability. The raw data obtained in this study are available from the corresponding author upon request.

Supplement. The supplement related to this article is available on-line at: https://doi.org/10.5194/bg-18-1-2021-supplement.

Author contributions. SS, HD and PB designed the experiment, with input of MA. MA and SS selected and provided paddy soils from Bangladesh. HD, OM and HL conducted the incubation experiment with the support of MA. HD did the lab analyses with the support of SS, MA, OM and HL. SB and HD performed the dissolved organic C content and δ^{13}C analyses, and SS, SB and HD developed and performed the procedure to quantify Fe-bound soil organic C. HD and SS conducted the data analyses and wrote the manuscript, with input from all other authors. CE2

Competing interests. The authors declare that they have no conflict of interest.

Acknowledgements. We would like to genuinely thank the involved staff of the tropical greenhouse, the lab of the Soil Fertility and Nutrient Management (SoFer) group and the Isotope Bioscience Laboratory (ISOFYS) from Ghent University (Belgium).

Financial support. This research has been supported by the European Research Area Network (ERA-Net) and the Research Foundation Flanders (FWO) in the framework of the project "Greenhouse gas emissions from paddy rice soils under alternative irrigation management" (GreenRice) (grant no. GA.010.14N).

Review statement. This paper was edited by Luo Yu and reviewed by Guanghui Yu and one anonymous referee.

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Remarks from the language copy-editor

Remarks from the typesetter