

# WETMETH 1.0: A new wetland methane model for implementation in Earth system models

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**Abstract.** Wetlands are the single largest natural source of methane (CH<sub>4</sub>), a powerful greenhouse gas affecting the global climate. In turn, wetland CH<sub>4</sub> emissions are sensitive to changes in climate conditions such as temperature and precipitation shifts. However, biogeochemical processes regulating wetland CH<sub>4</sub> emissions (namely microbial production and oxidation of CH<sub>4</sub>) are not routinely included in fully coupled Earth system models that simulate feedbacks between the physical climate, the carbon cycle, and other biogeochemical cycles. This paper introduces a process-based wetland CH<sub>4</sub> model (WETMETH) developed for implementation in Earth system models and currently embedded in an Earth system model of intermediate complexity. Here we: (i) describe the wetland CH<sub>4</sub> model; (ii) evaluate the model performance against available datasets and estimates from the literature; (iii) analyze the model sensitivity to perturbations of poorly constrained parameters. Historical simulations show that WETMETH is capable of reproducing mean annual emissions consistent with present-day estimates across spatial scales. For the 2008-2017 decade the model simulates global mean wetland emissions of 158.6 Tg CH<sub>4</sub> yr<sup>-1</sup>, of which 33.1 Tg CH<sub>4</sub> yr<sup>-1</sup> are from wetlands north of 45°N. WETMETH is highly sensitive to parameters for the microbial oxidation of CH<sub>4</sub>, which is the least constrained process in the literature.

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## 25 **1 Introduction**

Wetlands are vegetated locations that are inundated with water on a permanent, seasonal or recurrent basis (Wheeler, 1999). In the context of this study, wetlands are defined following the latest global CH<sub>4</sub> budget report (Saunois et al., 2020): natural ecosystems with inundated or water-saturated soils where anoxic conditions lead to the production of CH<sub>4</sub>. Wetlands across the globe are the single largest natural source of atmospheric CH<sub>4</sub>, accounting for approximately a third of total global emissions (Bridgman et al., 2013; Saunois et al., 2016). Estimates of global wetland CH<sub>4</sub> emissions over the past few decades vary between 140 and 210 Tg CH<sub>4</sub> yr<sup>-1</sup> (Kirschke et al., 2013). Although there exist different types of wetlands such as bogs, fens, swamps, marshes and floodplains (Aselmann and Crutzen, 1989; Saunois et al., 2016), the release of CH<sub>4</sub> from any wetland results from the balance between two biogeochemical processes (Segers, 1998): the production of CH<sub>4</sub> by anaerobic microbes (namely methanogens) and the oxidation of CH<sub>4</sub> primarily by aerobic microbes (namely methanotrophs).

Both CH<sub>4</sub> production and oxidation in wetlands are sensitive to changes in climate conditions. For instance, soil warming accelerates the microbial activity with a higher response for methanogenic than methanotrophic activity (Bridgman et al., 2013; Dunfield et al., 1993; Segers, 1998). At the landscape or larger scale, increased wet conditions tend to enhance methanogenic activity to the detriment of methanotrophic activity (Duval and Radu, 2018; Helbig et al., 2017; Kim, 2015). In turn, wetland CH<sub>4</sub> emissions can affect the global climate through changes in atmospheric CH<sub>4</sub> levels and associated radiative forcing (Dean et al., 2018; O'Connor et al., 2010). Analyses of ice cores suggest that CH<sub>4</sub> emissions from tropical and northern wetlands contributed significantly to climate changes during past glacial-interglacial transitions (Loulergue et al., 2008; Rhodes et al., 2017).

The interactions between climate conditions and wetland CH<sub>4</sub> emissions translate into a positive feedback loop that has the potential to amplify changes in global mean surface air temperature, which is a major concern for future climates (Dean et al., 2018; O'Connor et al., 2010). Research on feedbacks between the physical climate and biogeochemical cycles is generally conducted with 3-dimensional (3-D) fully coupled Earth system models (ESMs) (Arora et al., 2013). Over the past decade, these ESMs have proven very useful to investigate and inform international climate policies such as the accounting of carbon emissions required to avoid the risk of dangerous climate change (Zickfeld et al., 2009) and achieve the goals of the Paris Agreement (Tokarska and Gillett, 2018). Yet, biogeochemical processes regulating CH<sub>4</sub> emissions in wetlands are not commonly included in fully coupled ESM simulations.

In the past, several process-based models have been developed for investigating the response of wetland CH<sub>4</sub> emissions to climate variability and climate change (Hodson et al., 2011; Hopcroft et al., 2011; Pandey et al., 2017; Paudel et al., 2016; Shindell et al., 2004; Zhang et al., 2018; Zhu et al., 2015). These wetland CH<sub>4</sub> models are generally embedded in terrestrial or land surface models and forced with observational datasets or reanalysis products (Melton et al., 2013; Wania et al., 2013; Xu et al., 2016). A second application for wetland CH<sub>4</sub> models has been to quantify the climate response to wetland CH<sub>4</sub> emissions (Gedney et al., 2004, 2019; Zhang et al., 2017b). In this case, results from wetland CH<sub>4</sub> models are used in climate-carbon cycle model emulators to assess their impact on radiative forcing (Gedney et al., 2019; Zhang et al., 2017b).

These modelling studies have contributed to advance research on the possible evolution of wetland CH<sub>4</sub> emissions in the 21<sup>st</sup> century (Koven et al., 2011; Shindell et al., 2004), the magnitude of their impact on the global climate (Gedney et al., 2019; Zhang et al., 2017b), and their implications for international climate policy (Comyn-Platt et al., 2018). However, their quasi-coupling methods do not reflect the complete feedback loop between climate conditions and wetland CH<sub>4</sub> emissions as expected in the natural world. So far, only 1-D and 2-D models of the northern high-latitude regions have been applied for simulating the feedback between climate conditions (temperature changes) and wetland CH<sub>4</sub> emissions in a fully coupled mode (Schneider von Deimling et al., 2012, 2015).

The implementation of process-based wetland CH<sub>4</sub> models in fully coupled ESMs is needed in order to advance research on wetland CH<sub>4</sub>-climate feedbacks in the context of global climate projections (Dean et al., 2018). In particular, this addition to Earth system modelling should be beneficial to ongoing research on the permafrost carbon feedback (Nzotungicimpaye and Zickfeld, 2017; Schuur et al., 2015) and the remaining carbon budget for achieving the goals of the Paris Agreement (Rogelj et al., 2019).

This paper introduces a wetland CH<sub>4</sub> model developed for implementation in ESMs and currently embedded in an Earth system model of intermediate complexity (EMIC). Our study aims at developing a computationally efficient process-based model for simulating large-scale wetland CH<sub>4</sub> emissions constrained with sparse observations. Section 2 gives an overview of processes regulating CH<sub>4</sub> emissions in wetlands. Section 3 provides the model description and an outline of performed model simulations. Section 4 describes the model calibration and choice of parameter values. Section 5 presents the model performance evaluation. Section 6 describes the model sensitivity to poorly constrained parameters. Sections 7 and 8 are for discussions and conclusions, respectively.

## 2 Overview of processes regulating methane emissions in wetlands

### 2.1 Microbial production of methane

Wetlands host several communities of microbes adapted to the predominant anoxic conditions of these environments (Bridgham et al., 2013). Some of these microbes are methanogens, which decompose organic matter for their metabolism and produce CH<sub>4</sub> as a by-product of their respiration (McCalley et al., 2014; Segers, 1998). The organic matter decomposed by methanogens in wetlands originates from litter-fall, root exudates, dead plants and dissolved organic carbon (Bridgham et al., 2013; Conrad, 2009; Girkin et al., 2018; Mitsch and Mander, 2018). In the northern permafrost region, carbon from thawed soils constitutes an additional source of organic matter to methanogens (Kwon et al., 2019; Olefeldt et al., 2013).

There are three pathways through which methanogens produce CH<sub>4</sub> from soil organic matter (Le Mer and Roger, 2001; Segers, 1998; Whalen, 2005). The first pathway (acetotrophic methanogenesis) is operated by methanogens that rely on acetate for their metabolism, resulting in the production of both CH<sub>4</sub> and carbon dioxide (CO<sub>2</sub>) (Bridgham et al., 2013; Whalen, 2005). The second pathway (hydrogenotrophic methanogenesis) is operated by methanogens that produce CH<sub>4</sub> through CO<sub>2</sub> reduction in the presence of hydrogen (Bridgham et al., 2013). The third pathway (methylotrophic methanogenesis) is operated

90 by methanogens that use methylated substrates (e.g. methanol, methylamines, and dimethylsulfide) for their metabolism (Zalman et al., 2018).

Rates of CH<sub>4</sub> production in wetlands are generally highest in upper anoxic layers due to several factors such as the quality of organic matter and the spread of active microbial populations. For instance, in comparison to soil layers at depth where organic matter can be recalcitrant to microbial decomposition, the organic matter in near-surface soil layers is more  
95 labile due to fresh inputs from litter-fall and vegetation mortality (Treat et al., 2015; Walz et al., 2017; Wild et al., 2016). Furthermore, observations at various sites show that methanogenic activity decreases as depth increases (Bridgham et al., 2013; Cadillo-Quiroz et al., 2006).

Increasing soil temperatures stimulate the dynamics and growth of methanogenic communities in wetlands, resulting in an increase of CH<sub>4</sub> production rates (Bridgham et al., 2013; Segers, 1998). However, several studies indicate that there is  
100 an optimal temperature for methanogenic activity between 25°C and 30°C (Dean et al., 2018; Dunfield et al., 1993). Other factors promoting the occurrence of CH<sub>4</sub> production in wetlands include the persistence of anoxic conditions as well as soil pH varying between acidic and neutral (Dunfield et al., 1993; Segers, 1998).

## 2.2 Microbial oxidation of methane

In wetlands, methanotrophs (CH<sub>4</sub>-oxidizing microbes) populate oxic portions of the soil column (Bridgham et al., 2013;  
105 Conrad, 2009; Whalen, 2005). Such oxic portions are primarily soil layers close to the surface which are in contact with the atmosphere, commonly near and above the water table (Bridgham et al., 2013; Le Mer and Roger, 2001; Segers, 1998). In the presence of vascular plants, other oxic portions of the soil column can be found near the roots due to the downward transport of oxygen (O<sub>2</sub>) through plant aerenchyma (Kwon et al., 2019; Whalen, 2005). All these oxic portions of the soil column constitute the so-called oxic zone, which is predominantly made of soil layers near and above the water table (Bridgham et al.,  
110 2013; Conrad, 2009; Segers, 1998). Methanotrophs consume CH<sub>4</sub> that ascends from the zones of production at depth to the overlying oxic zone for their metabolism, and primarily produce CO<sub>2</sub> as part of their respiration (Bridgham et al., 2013; Segers, 1998).

While O<sub>2</sub> has been considered for years to be the only electron acceptor involved in the microbial oxidation of CH<sub>4</sub>, there is a growing evidence of the occurrence of CH<sub>4</sub> oxidation under anoxic conditions operated by anaerobic microbes that  
115 rely on alternate electron acceptors such as nitrate and sulfate (Dean et al., 2018). However, although anaerobic CH<sub>4</sub> oxidation in marine environments has been well established for decades (Hoehler et al., 1994; Reeburgh, 1976), this process remains poorly investigated in wetlands despite its potential importance for the CH<sub>4</sub> cycle (Gauthier et al., 2015; Smemo and Yavitt, 2011).

In analogy to CH<sub>4</sub> production, CH<sub>4</sub> oxidation is influenced by changes in soil temperatures (Bridgham et al., 2013;  
120 Segers, 1998). For instance, CH<sub>4</sub> oxidation rates increase during the summer because of intensified microbial activity but also the availability of substantial CH<sub>4</sub> in response to increased soil temperatures (Segers, 1998). However, the temperature

response for CH<sub>4</sub> oxidation is generally lower than that for CH<sub>4</sub> production (Bridgham et al., 2013; Dean et al., 2018; Dunfield et al., 1993; Segers, 1998).

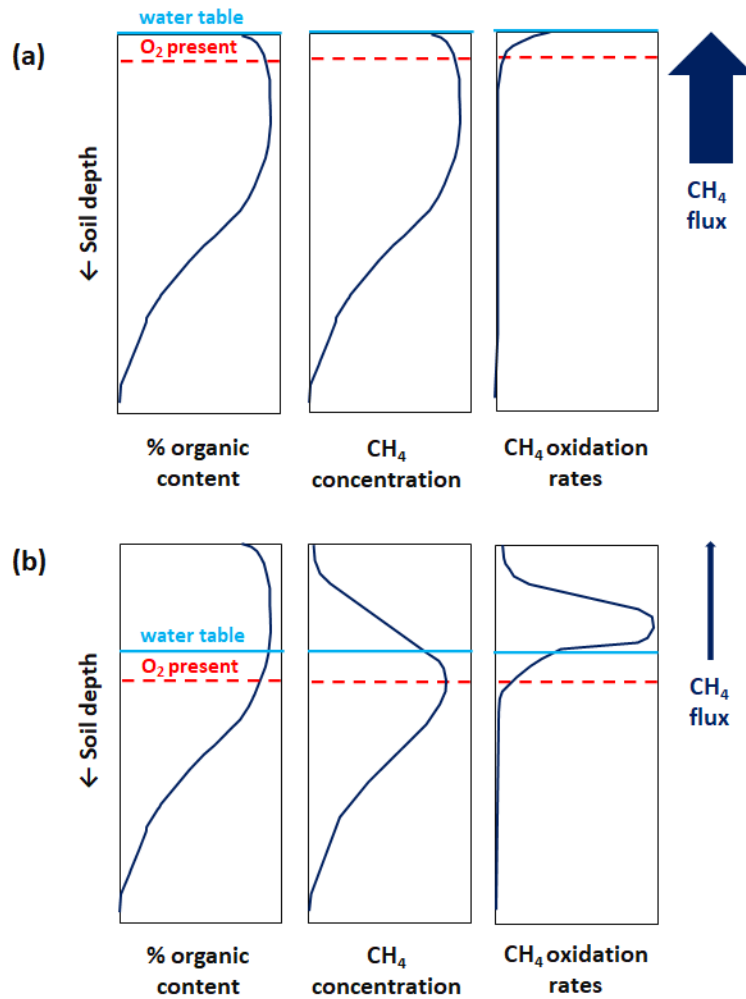
### 2.3 Mechanisms transporting methane to the atmosphere

125 There exist various mechanisms transporting CH<sub>4</sub> produced in wetlands to the atmosphere. Three transport mechanisms are well documented in the literature and generally monitored in situ (Bridgham et al., 2013; Whalen, 2005): the diffusion of CH<sub>4</sub> whereby molecules of CH<sub>4</sub> slowly ascend the overlying water column, the ebullition of CH<sub>4</sub> whereby bubbles of CH<sub>4</sub> rapidly ascend towards the soil surface, as well as the transport of CH<sub>4</sub> through the aerenchyma of vascular plants. However, other transport mechanisms for CH<sub>4</sub> in wetlands have been revealed: the hydrodynamic transport of CH<sub>4</sub> in the form of upwelling  
130 caused by temperature gradients primarily at nighttime (Poindexter et al., 2016), and the transport of CH<sub>4</sub> through tree stems (Bridgham et al., 2013; Conrad, 2009; Pangala et al., 2017) whose driving processes are still not well understood (Barba et al., 2019).

Methane oxidation is highly dependent on the predominant transport mechanism for CH<sub>4</sub>. The water table position plays a crucial role in affecting what fraction of the produced CH<sub>4</sub> reaches the atmosphere (Blodau, 2002; Moore and Roulet,  
135 1993; Segers, 1998). When the water table is well below the surface, methanotrophs may oxidize all of the diffusing CH<sub>4</sub> before the gas reaches the atmosphere (Segers, 1998). In the presence of vascular plants, a lower fraction of the produced CH<sub>4</sub> is oxidized because these plants allow the gas to bypass the oxic zone where methanotrophs are hosted (Blodau, 2002; Bridgham et al., 2013; Segers, 1998). In the case of ebullition, which often occurs episodically, CH<sub>4</sub> may escape to the atmosphere with reduced opportunities for oxidation (Bridgham et al., 2013; Whalen, 2005). How CH<sub>4</sub> oxidation relates to the  
140 transport of CH<sub>4</sub> through tree stems (Barba et al., 2019) or by hydrodynamic processes (Poindexter et al., 2016) is not well established.

### 2.4 A synopsis of wetland methane dynamics

Fig. 1 illustrates vertical profiles of soil organic content, CH<sub>4</sub> concentration, and CH<sub>4</sub> oxidation rates in a soil column with and without inundation at the surface based on principles outlined in the literature (Blodau et al., 2004; Whiticar and Faber, 1985).  
145 In general, the water table position determines the maximum depth at which O<sub>2</sub> is available in the soil column (i.e. the oxic-anoxic interface). When the surface is flooded and the water is stagnant (Fig. 1a), O<sub>2</sub> diffuses slowly into the soil column and may only be present in a portion of the upper soil layer which is in contact with the atmosphere. Under such predominantly anoxic conditions, CH<sub>4</sub> production occurs throughout the soil column and the concentration of CH<sub>4</sub> mirrors soil organic content – eventually with a small reduction near the surface due to CH<sub>4</sub> oxidation. A modest amount of ascending CH<sub>4</sub> may be oxidized  
150 throughout the soil column, but with highest oxidation rates near the surface where some O<sub>2</sub> may be available as an electron acceptor. The combination of high CH<sub>4</sub> production and only modest CH<sub>4</sub> oxidation in the soil column results in large CH<sub>4</sub> emissions into the atmosphere.



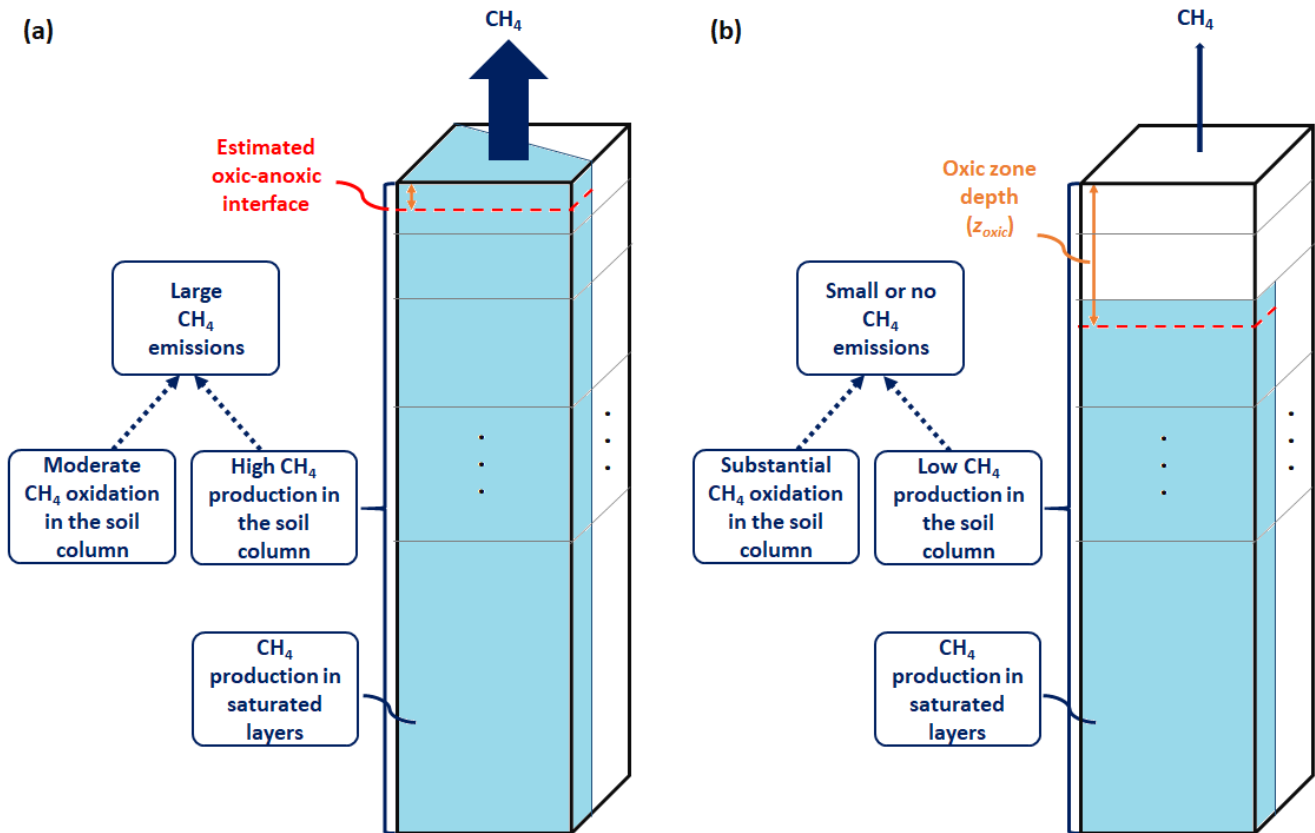
155 Figure 1: Illustrated vertical profiles of soil organic content, CH<sub>4</sub> concentration and oxidation rates in a soil column with inundation  
 at the surface (a) and without inundation at the surface (b). The vertical profiles are based on principles outlined in the literature  
 (Blodau et al., 2004; Whiticar and Faber, 1985). As an illustrative simplification, the soil organic content is assumed to be identical  
 in (a) and (b). Furthermore, the CH<sub>4</sub> concentration profile is assumed to mirror CH<sub>4</sub> production rates at depth within the soil column  
 (see explanatory text in Section 2.4). The blue horizontal line illustrates the water table position and the dashed red horizontal line  
 160 illustrates the oxic-anoxic interface or maximum depth at which O<sub>2</sub> is available in the soil column. The relative magnitude of CH<sub>4</sub>  
 flux in the soil column is shown by the upward arrow to the right, also characterizing the relative magnitude of CH<sub>4</sub> emissions into  
 the atmosphere.

When the flooding recedes, O<sub>2</sub> becomes more prevalent in the upper soil column where CH<sub>4</sub> concentration decreases following a slow down or shut down of CH<sub>4</sub> production as aerobic microbes dominate the competition for organic matter (Fig. 1b). CH<sub>4</sub> production persists below the oxic-anoxic interface where the concentration of CH<sub>4</sub> mirrors soil organic content  
 165 owing to the predominant anoxic conditions. Ascending CH<sub>4</sub> becomes subject to substantial oxidation in the soil column with the highest oxidation rates above the oxic-anoxic interface where O<sub>2</sub> is abundant. The combination of decreased CH<sub>4</sub> production and substantial CH<sub>4</sub> oxidation in the soil column results in small or no CH<sub>4</sub> emissions into the atmosphere.

### 3 Model description and simulations

#### 3.1 The wetland methane model: WETMETH

170 Microbial production and oxidation of  $\text{CH}_4$  are parameterized in WETMETH using a multi-layer ground structure with information on the moisture distribution, the amount of organic matter (carbon content), and the average temperature in each soil layer. These soil variables are commonly simulated by ESMs. Fig. 2 provides a schematic representation of WETMETH for a soil column with and without inundation at the surface. By configuration, it is considered that  $\text{CH}_4$  emissions in WETMETH may occur not only from inundated locations, but also from non-inundated ecosystems with a relatively high level  
175 of soil moisture content (Saunois et al., 2016, 2020).



180 **Figure 2: Illustration of the developed wetland  $\text{CH}_4$  model (WETMETH) and the dynamics of wetland  $\text{CH}_4$  processes as represented in the model. This schematic representation depicts a soil column (model grid box) with inundation at the surface (a) and without inundation at the surface (b). The soil column is shown here with multiple layers of unequal thicknesses. The blue area at the surface of (a) represents the inundated surface area. The blue sections in the different soil layers of (a) and (b) represent water-saturated zones. For both (a) and (b), the dashed red horizontal line illustrates the oxic-anoxic interface and the orange vertical arrow shows the relative thickness of the oxic zone or oxic zone depth ( $z_{\text{oxic}}$ ). Larger  $\text{CH}_4$  emissions are expected to occur when the soil surface is flooded than when it is not due to relatively high  $\text{CH}_4$  production and moderate  $\text{CH}_4$  oxidation in the soil column.**

### 3.1.1 Parameterization of methane production

185 For any land location, the rate of CH<sub>4</sub> production in an underlying soil layer  $i$  ( $P_i$  in kg C m<sup>-3</sup> s<sup>-1</sup>) is parameterized as:

$$P_i = S(\theta_i) C_i r Q_{10}^{\frac{T_i - T_0}{10}} \exp\left(-\frac{z_i}{\tau_{\text{prod}}}\right), \quad (1)$$

where  $S(\theta_i)$  is the fraction of the soil layer that is saturated with water, and  $C_i$  is the amount of soil carbon (in kg C m<sup>-3</sup>) in the layer. Here we consider  $C_i$  to be the aggregate of all sources of soil carbon (i.e. organic matter) such as litter-fall and root exudates. The product of  $S(\theta_i)$  and  $C_i$  represents the organic matter (in kg C m<sup>-3</sup>) available for microbial decomposition under anoxic conditions. When the soil surface is not flooded (Fig. 2b), dry soil layers ( $S(\theta_i) = 0$ ) are assumed to be predominantly oxic and not producing CH<sub>4</sub> ( $P_i = 0$ ) mostly due to aerobic microbes dominating the competition for organic matter which results in the starvation of methanogens (Segers, 1998).

The global factor  $r$  is the specific CH<sub>4</sub> production rate (in kg kg<sup>-1</sup> s<sup>-1</sup>), which can be defined as the mass of CH<sub>4</sub>-C that is produced per kilogram of available soil C per unit of time. A meta-analysis of incubated soil samples from various anaerobic landscapes indicates that  $r$  can vary between 0.3 to 27.2 μg of CH<sub>4</sub>-C per g of soil C per day (equivalent to the range from 3.5 x 10<sup>-12</sup> to 3.1 x 10<sup>-10</sup> kg kg<sup>-1</sup> s<sup>-1</sup>) depending on the landscape type, relative water table position, and soil depth (Treat et al., 2015). In this first version of WETMETH, we combine all possible pathways for CH<sub>4</sub> production in wetlands (see Section 2.1) without distinguishing fast and slow pathways. Section 4.1 discusses the choice of the value for  $r$  as part of the model calibration.

200 The expression  $Q_{10}^{\frac{T_i - T_0}{10}}$ , which depends on the average layer temperature  $T_i$  (in Kelvin, K) and a baseline temperature  $T_0$  (273.15 K), represents the temperature-dependency of CH<sub>4</sub> production expressed with a  $Q_{10}$  coefficient as commonly done to approximate the sensitivity of biological processes to a temperature change of 10 K (Hegarty, 1973). While some biological processes double in rate with a warming of 10 K, several studies report a higher temperature sensitivity for CH<sub>4</sub> production (i.e.  $Q_{10} > 2$ ) although with large uncertainties (Lupascu et al., 2012; Sjögersten et al., 2018; Walz et al., 2017; Whalen, 2005). Nevertheless, a meta-analysis of temperature-response studies suggests an average  $Q_{10}$  of about 4.2 for CH<sub>4</sub> production in pure cultures of methanogens (Hoehler and Alperin, 2014; Yvon-Durocher et al., 2014) in agreement with previous estimates (Blodau, 2002). In order to account for uncertainties with this coefficient and define the occurrence of an optimal temperature for CH<sub>4</sub> production (Dunfield et al., 1993; Metje and Frenzel, 2007; Schipper et al., 2014), a temperature-dependent  $Q_{10}$  is considered in WETMETH. Its mathematical formulation is  $Q_{10}(T_i) = 1.7 + 2.5 \tanh [0.1 (T_{ref} - T_i)]$ , where  $T_{ref} = 308.15$  K is a reference temperature that is used to define an optimal temperature for CH<sub>4</sub> production (Table 1). This formulation is defined in analogy to a mathematical expression used for soil respiration in another study (Wu et al., 2016), and it enables to account for an optimal temperature for CH<sub>4</sub> production of ~300.15 K (i.e. 27°C) which is consistent with previous studies (Dunfield et al., 1993; Metje and Frenzel, 2007). Additional information on this formulation and its implications for the temperature-dependency of CH<sub>4</sub> production are provided in Appendix A1. Furthermore, CH<sub>4</sub> production



215 in WETMETH is assumed to shut down in frozen soil layers although research suggests that slow microbial activity can occur at temperatures below 273.15 K (Panikov and Dedysh, 2000; Rivkina et al., 2004).

The expression  $\exp\left(-\frac{z_i}{\tau_{\text{prod}}}\right)$ , which depends on the depth of the soil layer  $i$  relative to the surface ( $z_i$  in m, positive downwards), describes the declining effect of various environmental controls on CH<sub>4</sub> production with depth that are generally unresolved by ESMs. These environmental factors include the quality of organic matter and the spread of methanogens among other factors (Bridgham et al., 2013; Koven et al., 2015; Treat et al., 2015; Walz et al., 2017; Wild et al., 2016). Here,  $\tau_{\text{prod}}$  (in m) is a scaling parameter for CH<sub>4</sub> production. The choice of the value for  $\tau_{\text{prod}}$  is discussed later as part of the model calibration (see Section 4.1).

**Table 1: Model parameters for methane production and oxidation.**

Parameter	Description	Units	Chosen value
$r$	Specific CH <sub>4</sub> production rate	kg kg <sup>-1</sup> s <sup>-1</sup>	<sup>a</sup> 2.6 x 10 <sup>-10</sup>
$Q_{10}$	Temperature coefficient for CH <sub>4</sub> production	—	<sup>b</sup> 4.2
$T_{\text{ref}}$	Reference temperature for CH <sub>4</sub> production	K	<sup>c</sup> 308.15
$\tau_{\text{prod}}$	Scaling parameter for CH <sub>4</sub> production	m	0.75
$z_{\text{oztz}}$	Thickness of the oxic-anoxic transition zone	m	0.05
$\tau_{\text{oxid}}$	Scaling parameter for CH <sub>4</sub> oxidation	m	0.0146

<sup>a</sup> This value is equivalent to 22.8 μg CH<sub>4</sub>-C produced per g of soil C per day; <sup>b</sup> A temperature-dependent  $Q_{10}$ , approximating 4.2 for a wide range of temperatures, is used instead (see Appendix A1); <sup>c</sup> The reference temperature is used to define an optimal temperature for CH<sub>4</sub> production (see Appendix A1).

The total amount of CH<sub>4</sub> produced in the soil column ( $P$  in kg C m<sup>-2</sup> s<sup>-1</sup>) is calculated as:

$$P = \int_{i=1}^{i=k} P_i dz_i, \quad (2)$$

230 where  $P_i$  (in kg C m<sup>-3</sup> s<sup>-1</sup>) is the rate of CH<sub>4</sub> production in the soil layer  $i$  from Eq. (1),  $dz_i$  (in m) is the thickness of the soil layer  $i$ , and  $k$  represents the bottom-most soil layer. This amount of CH<sub>4</sub> ( $P$ ) is then subject to oxidation in transit to emission into the atmosphere.

### 3.1.2 Parameterization of methane oxidation and net methane emissions

Methane oxidation is parameterized based on the amount of CH<sub>4</sub> produced in the soil column and the relative thickness of the oxic zone. Specifically, the total amount of CH<sub>4</sub> oxidized in the soil column ( $O_x$  in kg C m<sup>-2</sup> s<sup>-1</sup>) and net CH<sub>4</sub> emissions to the atmosphere ( $E$  in kg C m<sup>-2</sup> s<sup>-1</sup>) are calculated as:

$$O_x = P \left(1 - \exp\left(-\frac{z_{\text{oxic}}}{\tau_{\text{oxid}}}\right)\right), \quad (3)$$

$$E = P - O_x, \quad (4)$$

which is equivalent to the following expression:

$$240 \quad E = P \exp\left(-\frac{z_{\text{oxic}}}{\tau_{\text{oxid}}}\right), \quad (5)$$

where  $P$  (in  $\text{kg C m}^{-2} \text{s}^{-1}$ ) is the total amount of  $\text{CH}_4$  produced in the soil column as defined in Eq. (2),  $z_{\text{oxic}}$  (in m) is the relative depth (positive downwards) to the oxic-anoxic interface (Fig. 2), and  $\tau_{\text{oxid}}$  (in m) is a scaling parameter for  $\text{CH}_4$  oxidation. As for  $\tau_{\text{prod}}$ , the choice of the value for  $\tau_{\text{oxid}}$  is discussed as part of the model calibration (see Section 4.2).

Regarding  $z_{\text{oxic}}$ , we assume that  $\text{O}_2$  may be present in soil layers unsaturated with water as well as in a shallow oxic-anoxic transition zone within the upper-most soil layer saturated with water (Fig. 2). In this first development of WETMETH, we consider a constant thickness ( $z_{\text{oatz}}$ ) of 0.05 m for the oxic-anoxic transition zone, with its bottom defined as the oxic-anoxic interface (Frolking et al., 2002; Singleton et al., 2018). When the soil surface is inundated,  $z_{\text{oatz}}$  is identical to  $z_{\text{oxic}}$  (Fig. 2a). Otherwise,  $z_{\text{oatz}}$  is only a fraction of  $z_{\text{oxic}}$  (Fig. 2b). The penetration of  $\text{O}_2$  into the soil and its dynamics with changing moisture conditions can be complex depending on site-specific factors such as the soil composition (Estop-Aragonés et al., 2012) and the presence of vascular plants (Brune et al., 2000). In addition, methanotrophs may be present at depth ( $> 0.05$  m) below the water table probably following some adaptation to low  $\text{O}_2$  conditions (Singleton et al., 2018). Nevertheless, the approach applied here for  $z_{\text{oxic}}$  is reasonable for ESMs not resolving  $\text{O}_2$  dynamics and microbial communities in the soil.

For Eq. (3), the expression  $(1 - \exp(-\frac{z_{\text{oxic}}}{\tau_{\text{oxid}}}))$  represents the fraction of  $P$  that gets oxidized in transit to emission into the atmosphere. Various studies report estimates of  $\text{CH}_4$  oxidation as a fraction of produced  $\text{CH}_4$  in the soil column (Blazewicz et al., 2012; Le Mer and Roger, 2001; Roslev and King, 1996; Segers, 1998; Singleton et al., 2018). From sample-to-sample and site-to-site, however,  $\text{CH}_4$  oxidation exhibits a broad range of values ranging from less than 20% to more than 95% depending on the sampled soil depth ranges, whether or not potential  $\text{CH}_4$  oxidation under anoxic conditions is considered, the monitored transport mechanisms for  $\text{CH}_4$  among many other factors (Blazewicz et al., 2012; Couwenberg et al., 2010; Jauhiainen et al., 2005; Kwon et al., 2019; Le Mer and Roger, 2001; Moosavi and Crill, 1998; Roslev and King, 1996; Segers, 1998; Singleton et al., 2018; Whalen, 2005). Nevertheless, the largest fractions of oxidized  $\text{CH}_4$  are generally associated with the deepest water tables or oxic-anoxic interfaces (Bridgham et al., 2013; Couwenberg et al., 2010; Jauhiainen et al., 2005; Roslev and King, 1996; Segers, 1998; Whalen, 2005).

The parameterization described in Eq. (3) is a simple approach for characterizing  $\text{CH}_4$  oxidation in the soil column. Such a parameterization is practical when there is little knowledge on the soil chemistry (e.g.  $\text{O}_2$  and alternate electron acceptors), the dynamics of methanotrophs and other environmental factors exerting a control on  $\text{CH}_4$  oxidation (Blazewicz et al., 2012; Blodau, 2002; Dean et al., 2018; Kwon et al., 2019; Singleton et al., 2018; Smemo and Yavitt, 2011). Most importantly, this parameterization considers the net effect of all mechanisms transporting  $\text{CH}_4$  from the anoxic soil layers where the gas is produced to the atmosphere. The oxidized  $\text{CH}_4$  is assumed to produce  $\text{CO}_2$  that becomes part of the soil respiration routinely simulated by ESMs.

### 270 3.2 The embedding Earth system model

WETMETH has been embedded in the University of Victoria Earth System Climate Model (UVic ESCM), an Earth system model of intermediate complexity (EMIC) (Weaver et al., 2001). A modified version of the EMIC based on UVic ESCM 2.9

(Eby et al., 2009) is used here. The UVic ESCM consists of a 3-D ocean general circulation model that is coupled to a dynamic-thermodynamic sea ice model, a 2-D (vertically-integrated) energy-moisture balance model for the atmosphere and a land surface model (Weaver et al., 2001). The land surface model is a modified version of the Met Office Surface Exchange Scheme (MOSES) with 14 ground layers of unequal thickness extending down to a depth of 250 m that can simulate permafrost processes such as freeze-thaw dynamics (Avis et al., 2011). The top eight ground layers (~10 m in total depth) are soil layers and contribute to the water cycle, whereas the bottom six ground layers are bedrock layers (Avis et al., 2011). In the hydraulically active layers, porosity and permeability are determined based on the relative abundance of prescribed sand, clay, and silt-sized particles. Water phase changes are determined over a range of soil temperatures to determine the fraction of frozen and unfrozen water in the ground (Avis et al., 2011). All components of the UVic ESCM have a horizontal grid resolution of 3.6° in longitude and 1.8° in latitude (Eby et al., 2009; Weaver et al., 2001).

Wetlands in the UVic ESCM are identified in grid cell areas based on soil moisture content and topography. Model grid cells in which wetlands can occur are those with unfrozen soil moisture contents greater than 65% of the saturated moisture content in the upper soil layer for at least one day in a year (Avis et al., 2011). Instead of using a fixed global threshold value for topography (Avis et al., 2011), the version of the UVic ESCM used here identifies wetland coverage at the sub-grid scale following a TOPMODEL approach for global models (Gedney and Cox, 2003). Appendix A2 describes a minor modification applied to this TOPMODEL approach. Section 5.1 presents an evaluation of wetlands simulated by the UVic ESCM.

The UVic ESCM includes a representation of the global carbon cycle. The terrestrial carbon cycle is simulated using the Top-down Representation of Interactive Foliage and Flora including Dynamics (TRIFFID), a dynamic global vegetation model that is coupled to the land surface model (Avis et al., 2011; Meissner et al., 2003). TRIFFID defines the state of the terrestrial biosphere in terms of soil carbon as well as the structure and coverage of five plant functional types (PFTs): broadleaf trees, needleleaf trees, shrubs, C3 grasses and C4 grasses (Cox, 2001; Matthews et al., 2004; Meissner et al., 2003). Terrestrial carbon gain occurs through photosynthesis that is simulated as a function of atmospheric CO<sub>2</sub> concentration, shortwave radiation, air temperature, humidity, and soil moisture. Soil carbon gain occurs through litter-fall and vegetation mortality. The present-day permafrost carbon pool is simulated by the UVic ESCM following a method that approximates the effect of long-term freeze-thaw cycles on the vertical distribution of carbon in permafrost-affected soils, a process referred to as cryoturbation (MacDougall and Knutti, 2016). Soil carbon can occur in the top six ground layers (~3.35 m in total depth). Terrestrial carbon loss occurs through autotrophic respiration by plants and heterotrophic respiration by soil microbes (Matthews et al., 2004; Meissner et al., 2003). By configuration, permafrost carbon can only be lost through microbial respiration and this heterotrophic respiration is assumed to shut down in frozen soil layers (MacDougall et al., 2012; MacDougall and Knutti, 2016). **Through TRIFFID, all terrestrial carbon fluxes in the UVic ESCM are integrated with a 30-day timestep (Meissner et al., 2003).**

The marine carbon cycle in the UVic ESCM is represented with organic and inorganic carbon cycle models (Eby et al., 2009). The organic carbon cycle is based on marine biology simulated with a nutrient-phytoplankton-zooplankton-detritus (NPZD) ecosystem model (Schmittner et al., 2008). The inorganic carbon cycle model simulates the air-sea exchange of CO<sub>2</sub>

and ocean carbonate chemistry following the protocols of the Ocean Carbon-Cycle Model Intercomparison Project (OCMIP) (Orr, 1999; Weaver et al., 2001). Dissolved inorganic carbon is treated as a passive tracer that is subject to ocean circulation (Weaver et al., 2001). Carbonate dissolution in ocean sediments is simulated with a model of respiration in marine sediments (Archer, 1996; Eby et al., 2009).

### 3.3 Model simulations

For this research, three series of model simulations are performed with the UVic ESCM in its standard fully coupled mode and including WETMETH parameterizations:

1. Firstly, the UVic ESCM is spun up for ~5000 years at year 1850 conditions to allow the model to reach an equilibrium climate state representing the pre-industrial period.
2. Secondly, a transient run from 1850 to 2019 is performed in order to evaluate the model performance. This transient run is based on prescribed CO<sub>2</sub> concentration and other forcing data from the fifth phase of the Coupled Model Intercomparison Project (CMIP5) (Taylor et al., 2012). The UVic ESCM is driven by historical data from 1850 to 2005 and by Representative Concentration Pathway (RCP) 8.5 data from 2006 to 2019. Supplementary Fig. S1 illustrates how the simulated historical climate conditions compare to observations in terms of global mean surface air temperature.
3. Thirdly, a set of transient runs from 2000 to 2009 is performed to analyze the model sensitivity to poorly constrained parameters. This set of model simulations (sensitivity runs) is performed by perturbing values of poorly constrained parameters associated with wetland CH<sub>4</sub> processes.

## 4 Choice of model parameter values

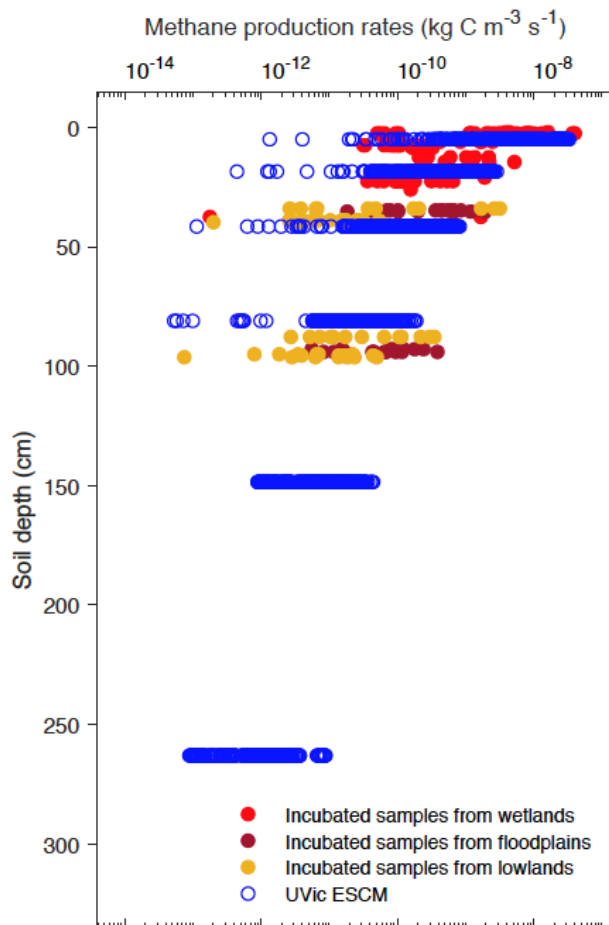
Here, we describe the choice of three WETMETH parameters ( $r$  and  $\tau_{\text{prod}}$  for CH<sub>4</sub> production;  $\tau_{\text{oxid}}$  for CH<sub>4</sub> oxidation) as part of the model calibration. These model parameters are tuned to observations from northern high-latitude regions due to the scarcity of large-scale datasets from other regions. The model calibration against northern observations is based on the assumption that tuned parameter values will be valid across the globe, which is an important limitation as it will be discussed later. Nonetheless, this approach is deemed reasonable given the present state of data availability. Section 5.1 describes northern wetlands simulated by the UVic ESCM as part of the model validation.

### 4.1 Methane production parameters

Parameters for CH<sub>4</sub> production in WETMETH are calibrated against maximum CH<sub>4</sub> production rates measured in laboratory incubations of soil samples from several anaerobic environments across northern high-latitude regions (>50°N). These potential CH<sub>4</sub> production rates are obtained from a synthesis dataset, which includes information on other environmental variables such as the relative depth of the soil samples (Treat et al., 2015).

To allow a fair model-data comparison, measured CH<sub>4</sub> production rates with corresponding soil bulk density from the sites of origin are converted into units of kg C m<sup>-3</sup> s<sup>-1</sup> (see Appendix A3). Furthermore, measurements from landscapes identified as uplands and lakes (in the dataset) are excluded from the dataset used in this model calibration. The remaining  
340 measurements are potential CH<sub>4</sub> production rates in soil samples from landscapes identified (in the dataset) as wetlands, floodplains and lowlands across Alaska.

In order to set values for  $r$  and  $\tau_{\text{prod}}$  from Eq. (1), the depth profile of simulated CH<sub>4</sub> production rates across Alaska for the year 2000 is tuned to that of the measurements. By setting  $r$  to 22.8 μg CH<sub>4</sub>-C produced per g of soil C per C day (equivalent to 2.6 x 10<sup>-10</sup> kg kg<sup>-1</sup> s<sup>-1</sup>) and  $\tau_{\text{prod}}$  to 0.75 m, we obtain a depth profile of simulated CH<sub>4</sub> production rates that  
345 compares fairly well to that of potential CH<sub>4</sub> production rates from the laboratory incubations (Fig. 3). These default values for  $r$  and  $\tau_{\text{prod}}$  are listed in Table 1. Section 6 presents a sensitivity analysis on these model parameters.



**Figure 3: Vertical profiles of simulated and potential CH<sub>4</sub> production rates from wetlands across Alaska. Potential CH<sub>4</sub> production rates are measurements from laboratory incubations of soil samples collected from various anaerobic ecosystems (Treat et al., 2015). Both simulated and measured CH<sub>4</sub> production rates are shown here with a log-transformed axis (base-10 logarithmic scale).**  
350

## 4.2. Methane oxidation parameter

Unlike for CH<sub>4</sub> production, there are no published large-scale measurements of CH<sub>4</sub> oxidation rates that could be used in this research for the calibration of CH<sub>4</sub> oxidation. For that reason, CH<sub>4</sub> oxidation in WETMETH is indirectly calibrated via CH<sub>4</sub> emissions. A synthesis dataset of seasonal and annual CH<sub>4</sub> emissions from various terrestrial sites across temperate, boreal and Arctic regions is used to this end (Treat et al., 2018). The model calibration focuses on annual CH<sub>4</sub> emissions from sites north of 50°N for which many data points are available in the dataset.

While most data points are from direct measurements of CH<sub>4</sub> emissions, some data points are associated with different modelling methods for estimating CH<sub>4</sub> emissions (Treat et al., 2018). To allow a fair model-data comparison, only data points associated with direct measurements of CH<sub>4</sub> emissions are included in the model calibration. Furthermore, measurements from lakes, uplands and alpine landscapes are excluded from this model calibration. In particular, the exclusion of data points from uplands and alpine landscapes sorts out measurements of terrestrial CH<sub>4</sub> uptake (negative CH<sub>4</sub> flux). The retained data points (n = 119) include measurements by chambers (85.7%), flux towers (13.4%) and a combination of flux towers and chambers (0.8%).

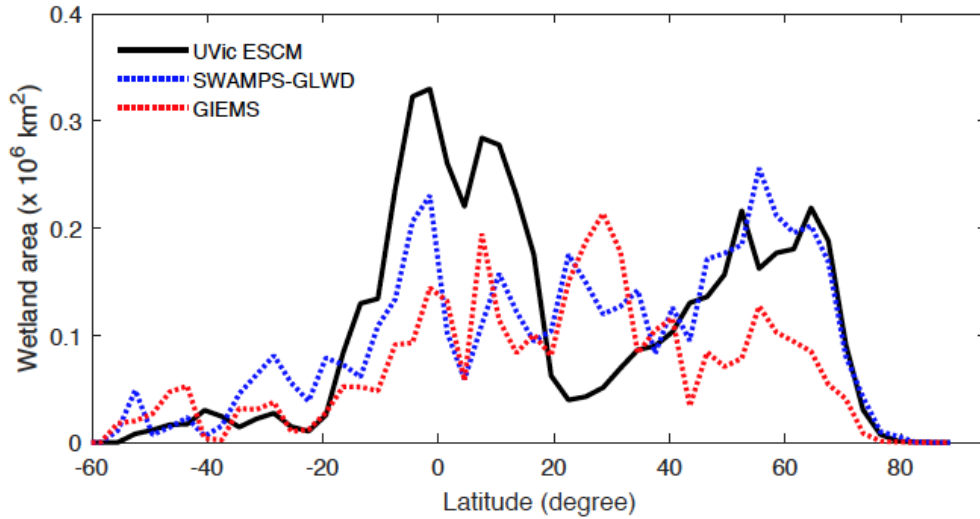
The model calibration in this section aims at choosing a value of  $\tau_{\text{oxid}}$  from Eq. (5) such that the range (minimum - maximum) of annual CH<sub>4</sub> emissions across northern wetlands (>50°N) simulated by the UVic ESCM is comparable to that of annual CH<sub>4</sub> emissions from the data points (0.1-60.6 g CH<sub>4</sub> m<sup>-2</sup> yr<sup>-1</sup>). By setting  $\tau_{\text{oxid}}$  to 0.0146 m, we constrain simulated CH<sub>4</sub> emissions from northern wetlands (specifically, grid-cell CH<sub>4</sub> emissions divided by the inundated fraction of the grid cell) from 2000 to 2009 in the range of 0.04-65.6 g CH<sub>4</sub> m<sup>-2</sup> yr<sup>-1</sup>. This tuned value for  $\tau_{\text{oxid}}$  is listed in Table 1 and implies that ~97% of the CH<sub>4</sub> produced in the soil column gets oxidized in transit to emission when the soil surface is inundated. Section 6 presents a sensitivity analysis on this model parameter.

## 5 Evaluation of the model performance

### 5.1 Wetlands

Fig. 4 shows the latitudinal distribution of wetland areas simulated by the UVic ESCM in comparison to two global datasets. The first dataset is Global Inundation Extent from Multi-Satellites (GIEMS), which is based on remotely sensed inundation areas (Papa et al., 2010; Prigent et al., 2001, 2007, 2012). The second dataset is Surface Water Microwave Product Series-Global Lakes and Wetlands Database (SWAMPS-GLWD), which is based on a combination of information from satellites and maps of inundated areas in order to reduce uncertainties associated with the distribution of global wetlands (Poulter et al., 2017). The comparison between the model and the datasets is done over 2000-2007, which is the overlap period for the datasets. Over this period the UVic ESCM simulates an annual maximal extent of ~12.6 million km<sup>2</sup> for global wetlands, whereas GIEMS and SWAMPS-GLWD estimate ~9.3 and ~10.6 million km<sup>2</sup>, respectively.

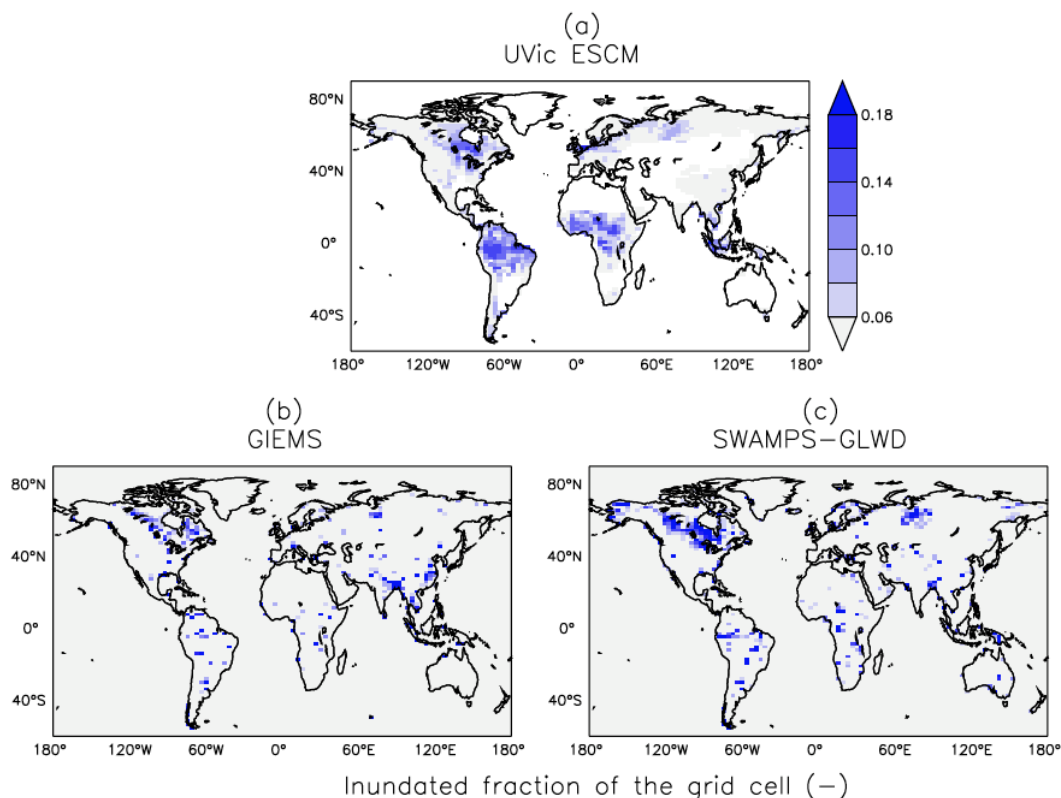
The UVic ESCM agrees better with SWAMPS-GLWD in regions north of 40°N although with some underestimations around 55°N, and relatively well with GIEMS between 20-40°S (Fig. 4). However, the model simulates too small wetland areas between 20-30°N when compared to both GIEMS and SWAMPS-GLWD. While our model could be underestimating wetland areas in this latitude zone, inundated areas estimated by GIEMS include rice paddies which prevail in tropical and sub-tropical regions (Prigent et al., 2007, 2012). Rice paddies are likely not represented in SWAMPS-GLWD as there were efforts to only include natural wetlands during the development of this dataset (Poulter et al., 2017). In comparison to GIEMS and SWAMPS-GLWD, our model simulates small wetland areas in South-East Asia especially near Bangladesh (Fig. 5 and Fig. 6).



390 **Figure 4: Latitudinal distribution of wetland areas simulated by the UVic ESCM over the 2000-2007 period in comparison to two global datasets: GIEMS and SWAMPS-GLWD. The comparison period corresponds to the overlap period for the two datasets. The wetland areas are summed across latitude bins of 3°.**

Between 20°N and 20°S, the UVic ESCM simulates a bimodal distribution of the wetland extent that is consistent with the two datasets although the model simulates too large wetland areas (Fig. 4). Unlike for GIEMS and SWAMPS-GLWD, wetlands simulated by the UVic ESCM are widespread in Amazonia, West and Central Africa (Fig. 5 and Fig. 6). Although 395 the UVic ESCM could be overestimating the extent of wetlands in some of these equatorial regions, it is possible that GIEMS and SWAMPS-GLWD do not detect inundated areas in densely forested regions due to forest canopies. Recent studies suggest that tropical wetlands are commonly underestimated in large-scale datasets (Dargie et al., 2017; Gumbricht et al., 2016).

Conversely, it is possible that the UVic ESCM overestimates tropical wetland areas due to soil hydraulic properties 400 unrepresented in the model. A potential cause for the overestimation of tropical wetlands in our model is the standard approach for simulating global hydrology in land surface models based on the concentration of only sand, clay and silt in the soil. A recent study suggests that the inclusion of ferralsols (weathered soils with micro-aggregated particles that are common in the humid tropics) in a global terrestrial model can help improve the simulation of tropical wetlands (Gedney et al., 2019).



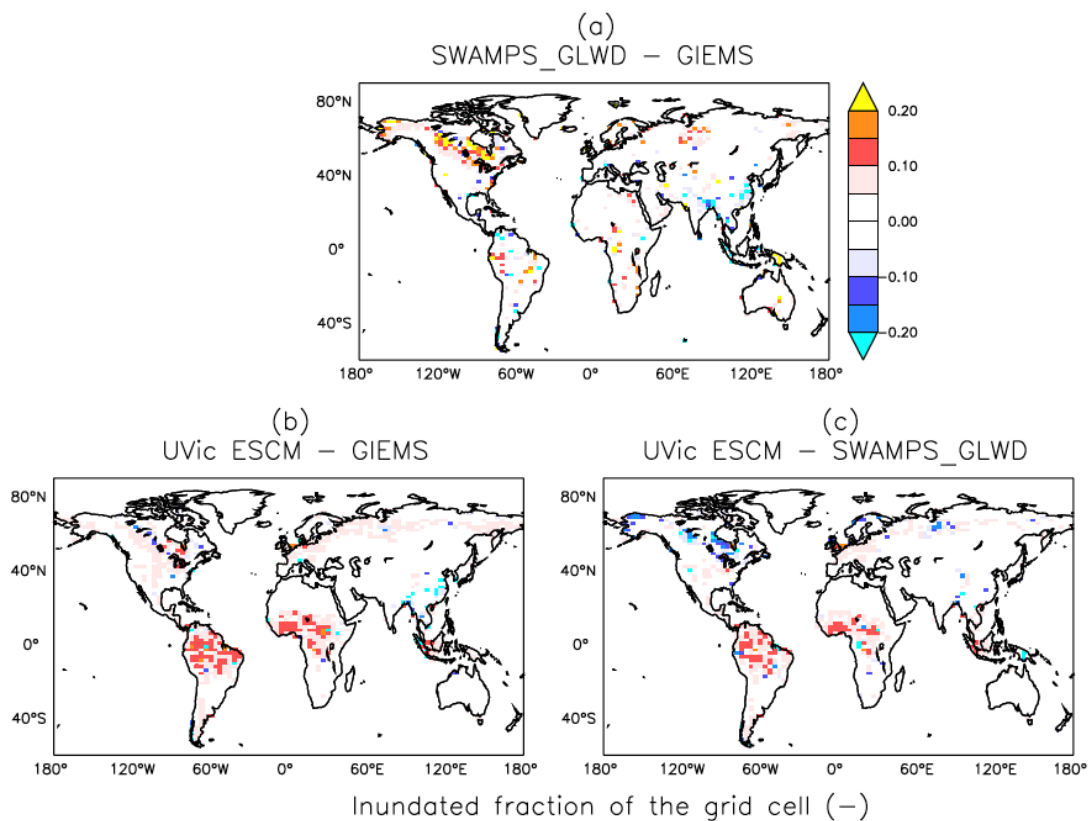
405 **Figure 5: Average wetland extents (inundated fractions of grid cells) across the globe over the 2000-2007 period as simulated by the UVic ESCM (a) in comparison to two datasets: (b) GIEMS and (c) SWAMPS-GLWD. The datasets are regridded to  $3.6^\circ \times 1.8^\circ$  for a fair comparison with the UVic ESCM. The comparison period corresponds to the overlap period for the two datasets.**

Outside of the tropics, the UVic ESCM does a better job at simulating the distribution of wetlands in sub-Arctic and Arctic regions (Fig. 7). The model simulates the occurrence of wetlands (i.e. surface inundation) across the West Siberian Lowlands (WSL) in Russia, the Hudson Bay Lowlands (HBL) in Canada as well as over other parts of northern Canada in agreement with both SWAMPS-GLWD and GIEMS (Fig. 7). However, some disagreements between the UVic ESCM and the two datasets can also be identified: (i) in comparison to GIEMS, the UVic ESCM simulates more wetland area in the Hudson Bay Lowlands (HBL) as well as widespread wetlands in parts of northern Eurasia (Fig. 7b and Fig. S2b); (ii) in comparison to SWAMPS-GLWD, the model simulates less wetland area over the WSL and northern Canada including the HBL and more wetland area in parts of Europe (Fig. 7c and Fig. S2c).

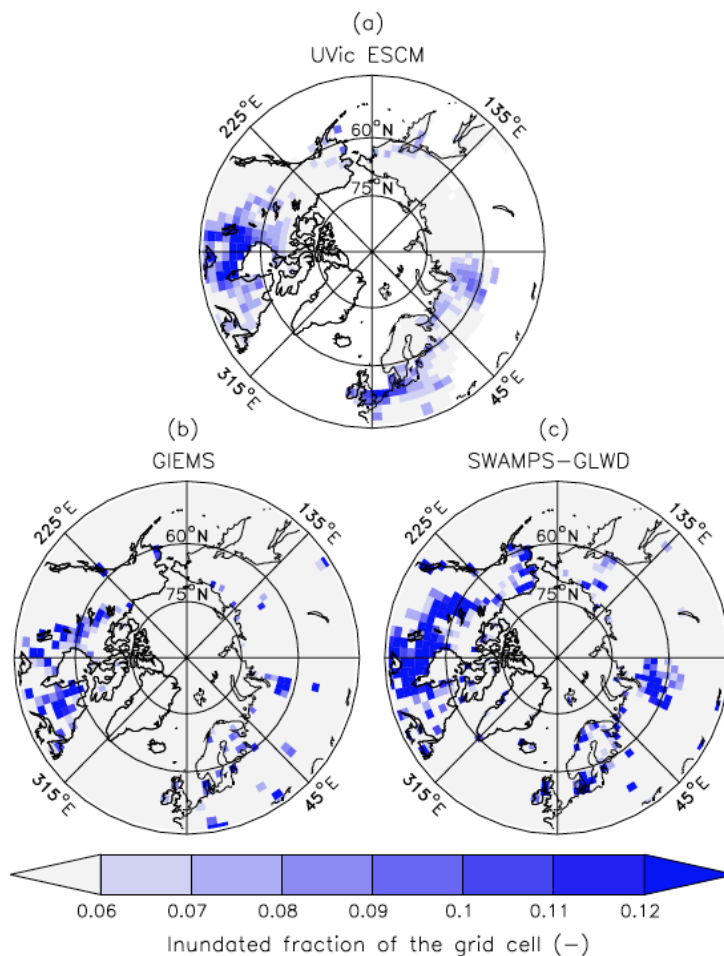
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Statistical analyses show that: (i) the UVic ESCM agrees better with SWAMPS-GLWD than with GIEMS at both the regional and global scale; and (ii) the model compares better with the two datasets across northern regions than at the global scale. For details on the statistical evaluation, please see supplementary Table S1.





420 **Figure 6: Differences in global wetland extents (inundated fractions of grid cells) between two datasets (GIEMS and SWAMPS-GLWD) and the UVic ESCM over the 2000-2007 period: (a) SWAMPS-GLWD – GIEMS, (b) UVic ESCM – GIEMS, and (c) UVic ESCM – SWAMPS-GLWD. The comparison period corresponds to the overlap period for the two datasets.**



425 **Figure 7: Average wetland extents (inundated fractions of grid cells) in the north of 45°N over the 2000-2007 period as simulated by the UVic ESCM (a) in comparison to two datasets: (b) GIEMS and (c) SWAMPS-GLWD. The datasets are regridded to 3.6° x 1.8° for a fair comparison with the UVic ESCM. The comparison period corresponds to the overlap period for the two datasets.**

## 5.2 Wetland methane emissions

Given the relative coarse grid resolution of the UVic ESCM, the model validation with respect to wetland CH<sub>4</sub> emissions focuses on large-scale emissions such as regional, zonal, and global emissions. Moreover, this model validation focuses on northern high-latitude regions because observations and estimates of wetland CH<sub>4</sub> emissions from other regions (e.g. the tropics) are scarce. This focus is further justified by the fact that our model better simulates the distribution of wetlands in northern high-latitude regions than in the tropics (see Section 5.1). Indeed, the extent of wetlands is a major control for wetland CH<sub>4</sub> emissions simulated by process-based models and probably the primary contributor to related uncertainties (Melton et al., 2013; Saunio et al., 2020; Zhang et al., 2017a).

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435 **5.2.1 Northern high-latitude emissions**

The UVic ESCM simulates total CH<sub>4</sub> emissions from northern wetlands that are in the range of recent estimates. Over the 2013-2014 period, the model simulates mean annual emissions of 33.2 Tg CH<sub>4</sub> yr<sup>-1</sup> for wetlands north of 45°N (Table 2). These CH<sub>4</sub> emissions are consistent with estimates from recent upscaled flux measurements (UFMs) over the same period based on a random forest (RF) algorithm and three wetland maps (Peltola et al., 2019): 30.6 ± 9.2 Tg CH<sub>4</sub> yr<sup>-1</sup> (RF-DYPTOP), 31.7 ± 440 9.4 Tg CH<sub>4</sub> yr<sup>-1</sup> (RF-PEATMAP), and 37.6 ± 11.8 Tg CH<sub>4</sub> yr<sup>-1</sup> (RF-GLWD) (Table 2). Supplementary Table S2 shows that the UVic ESCM has no preferential agreement with one of the three UFMs.

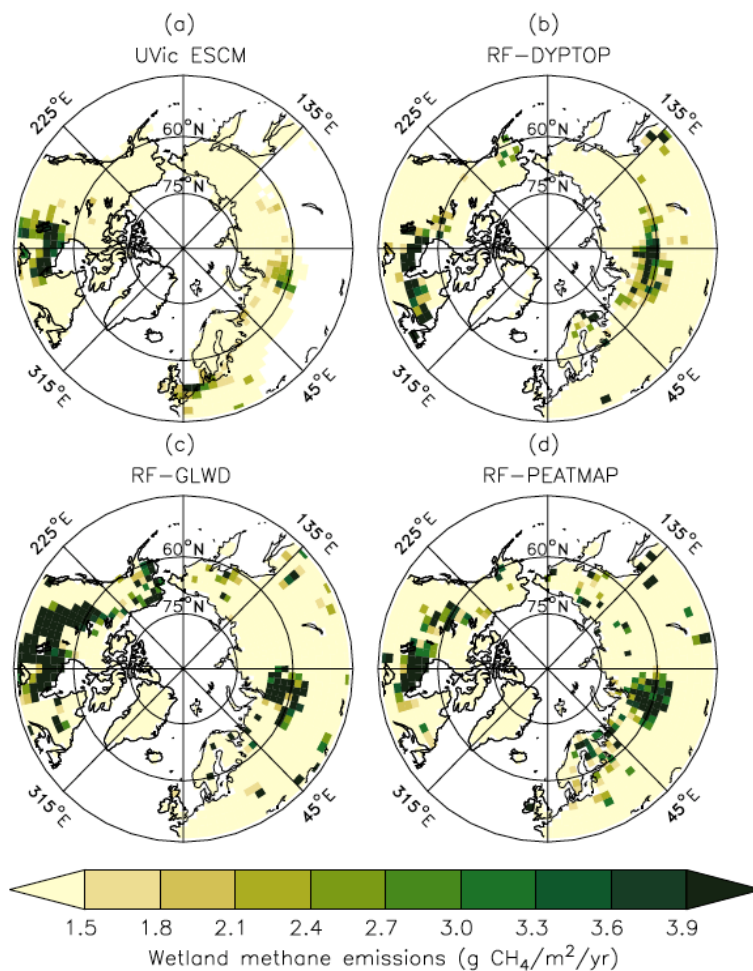
**Table 2: Mean annual wetland CH<sub>4</sub> emissions simulated by the UVic ESCM in comparison to estimated emissions from the literature. All emissions are reported in Tg CH<sub>4</sub> yr<sup>-1</sup> and uncertainties are provided for estimates from the literature. Three periods are used to allow a fair comparison between the UVic ESCM and estimates from the literature where possible: 2008-2017 as in the latest 445 global CH<sub>4</sub> budget report (Saunois et al., 2020), 2013-2014 as for recent upscaled flux measurements across the northern high-latitudes (Peltola et al., 2019), and 1993-2004 as for the WETCHIMP model ensemble (Melton et al., 2013). Principal methods used in the different references for estimates are reported in the last column: Top-down (TD) methods including inverse models (IM), and bottom-up (BU) methods including upscaled measurements (UM) as well as process-based models (PM).**

	Geographical delimitation	UVic ESCM period	UVic ESCM emissions	Estimated emissions	Reference for estimates	Method in reference
Hudson Bay Lowlands	50 – 60°N; 75 – 96°W	2013-2014	2.9	2.3 ± 0.3 2.4 ± 0.3 2.7 - 3.4	Pickett-Heaps et al., 2011 Miller et al., 2014 Thompson et al., 2017	BU IM IM
West Siberian Lowlands	50 – 75°N; 60 – 95 °E	2013-2014	4.1	3.9 ± 1.3 6.1 ± 1.2 6.9 ± 3.6	Glagolev et al., 2011 Bohn et al., 2015 <sup>a</sup> Thompson et al., 2017	UM IM IM
Pan-Arctic Wetlands	60°N – 90°N	2008-2017	17.3	7 – 16 2 – 18	Saunois et al., 2020 Saunois et al., 2020	TD BU
Northern Wetlands	40°N – 90°N 45°N – 90°N	2008-2017 2013-2014	38.5 33.2	37.4 ± 7.2 30.6 ± 9.2 31.7 ± 9.4 37.6 ± 11.8	Treat et al., 2018 Peltola et al., 2019 Peltola et al., 2019 Peltola et al., 2019	BU UM UM UM
Tropical Wetlands	30°S – 30°N	1993-2004	105.5	126 ± 31 90 ± 77	Melton et al., 2013 <sup>a</sup> Sjögersten et al. 2014	PM UM
Global Wetlands	90°S – 90°N	2008-2017	158.6	155 – 200 102 – 182	Saunois et al., 2020 Saunois et al., 2020	TD BU

<sup>a</sup> These reported estimates are model ensemble means. For the West Siberian Lowlands, the range between the inverse models is 3.1–9.8 Tg CH<sub>4</sub> yr<sup>-1</sup> (Bohn et al., 2015). For tropical wetlands, the range between the process-based models is 85–184 Tg CH<sub>4</sub> yr<sup>-1</sup> (Melton et al., 2013). 450

Fig. 8 shows the spatial distribution of simulated CH<sub>4</sub> emissions in comparison to the three UFMs. When compared to each other, the three UFMs exhibit substantial differences primarily attributed to the distinct wetland distributions (Peltola et al., 2019). Considering the general pattern and magnitude of wetland CH<sub>4</sub> emissions, the UVic ESCM agrees with either two or all three UFMs over key source regions such as the Hudson Bay Lowlands (HBL), the West Siberian Lowlands (WSL), 455 western Europe and south-central Canada (Fig. 8).

The UVic ESCM simulates less CH<sub>4</sub> emissions over parts of northeastern Canada and Fennoscandia in comparison to the UFM (Fig. 8). However, the three UFM do not necessarily agree on both the distribution and magnitude of wetland CH<sub>4</sub> emissions in these regions. Furthermore, the UVic ESCM does not simulate wetland CH<sub>4</sub> emissions in southern Eurasia (40-135°E; 45-60°N) while the three UFM suggest that CH<sub>4</sub> can be emitted from sporadic wetlands in this region (Fig. 8). Overall, the mismatch between the UFM and our model in terms of northern CH<sub>4</sub> emissions can be primarily attributed to differences in the wetland extent, but also to the spatial distribution of soil carbon simulated by the UVic ESCM (MacDougall and Knutti, 2016).



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**Figure 8: Average CH<sub>4</sub> emissions from wetlands north of 45°N over the 2013-2014 period as simulated by the UVic ESCM (a) in comparison to three datasets (upscaled flux measurements): (b) RF-DYPTOP, (c) RF-GLWD and (d) RF-PEATMAP. The datasets are regridded to 3.6° x 1.8° for a fair comparison with the UVic ESCM. The comparison period corresponds to the overlap period for the three datasets.**

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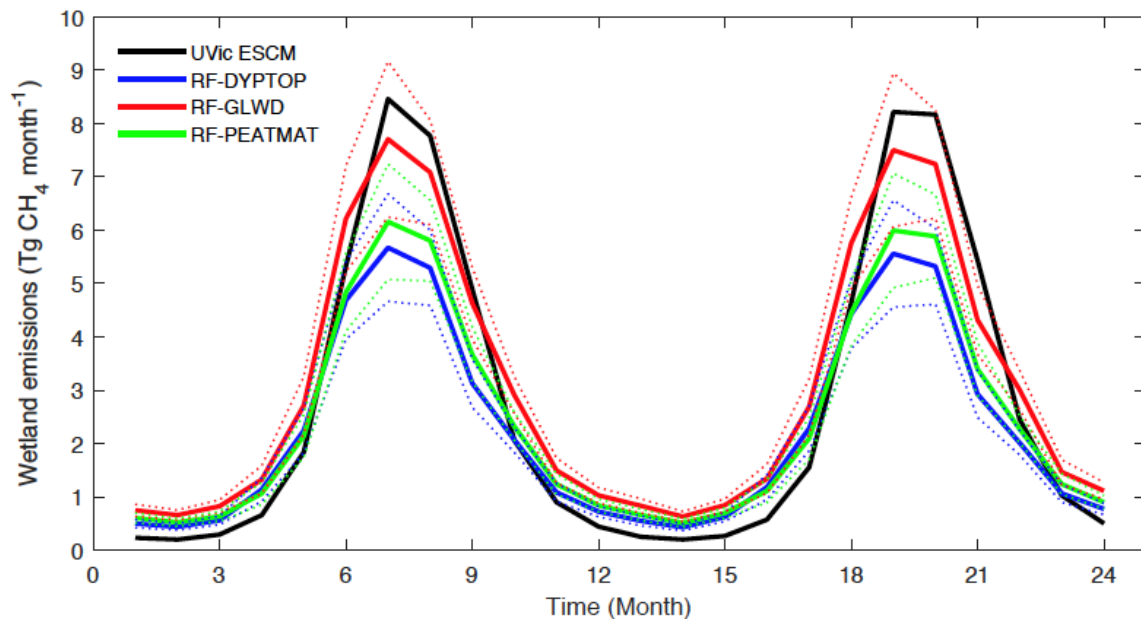
In terms of mean annual emissions from key source regions, the UVic ESCM simulates 2.9 Tg CH<sub>4</sub> yr<sup>-1</sup> for the Hudson Bay Lowlands (HBL) over the 2013-2014 period (Table 2). Although these emissions are lower than estimates by the three

UFMs (3.1-6.5 Tg CH<sub>4</sub> yr<sup>-1</sup>) (Peltola et al., 2019), estimates by inverse models (2.0-3.4 Tg CH<sub>4</sub> yr<sup>-1</sup>) over this region are comparable to our model results (Miller et al., 2014; Pickett-Heaps et al., 2011; Thompson et al., 2017). Furthermore, the UVic ESCM simulates total wetland emissions of 4.1 Tg CH<sub>4</sub> yr<sup>-1</sup> for the West Siberian Lowlands (WSL) over the 2013-2014 period (Table 2). Regional estimates based on the three UFMs are higher (4.9-8.5 Tg CH<sub>4</sub> yr<sup>-1</sup>) than our model results over the same period (Peltola et al., 2019), whereas previous observation-based estimates for the WSL suggest regional wetland emissions (3.9 ± 1.3 Tg CH<sub>4</sub> yr<sup>-1</sup>) that are similar to our model results (Glagolev et al., 2011). Estimates by inverse models over the WSL are relatively high but comparable to our model estimates (Table 2): 6.1 ± 1.2 Tg CH<sub>4</sub> yr<sup>-1</sup> (Bohn et al., 2015) and 6.9 ± 3.6 Tg CH<sub>4</sub> yr<sup>-1</sup> (Thompson et al., 2017).

The UVic ESCM is also evaluated with respect to wetland CH<sub>4</sub> emissions over the 2000-2009 and 2008-2017 decades, which both are reference periods for the latest global CH<sub>4</sub> budget report (Saunois et al., 2020). For wetlands north of 40°N, the UVic ESCM simulates emissions of 37.7 Tg CH<sub>4</sub> yr<sup>-1</sup> over the 2000-2009 decade and 38.5 Tg CH<sub>4</sub> yr<sup>-1</sup> over the 2008-2017 decade. These wetland CH<sub>4</sub> emissions are consistent with recent estimates (37.4 ± 7.2 Tg CH<sub>4</sub> yr<sup>-1</sup>) from data-constrained model ensembles over the same region (Treat et al., 2018). For wetlands north of 45°N, the model simulates total emissions that are in the range of estimates for the 2013-2014 period discussed earlier (32.4 Tg CH<sub>4</sub> yr<sup>-1</sup> over 2000-2009 and 33.1 Tg CH<sub>4</sub> yr<sup>-1</sup> over 2008-2017). For Pan-Arctic wetlands (>60°N), the UVic ESCM simulates emissions of 17.4 Tg CH<sub>4</sub> yr<sup>-1</sup> over the 2000-2009 decade and a similar amount over the 2008-2017 decade (Table 2). These wetland CH<sub>4</sub> emissions correspond to the upper limit of bottom-up estimates (2-18 Tg CH<sub>4</sub> yr<sup>-1</sup>) from the latest global CH<sub>4</sub> budget report (Saunois et al., 2020).

Fig. 9 shows seasonal cycles of CH<sub>4</sub> emissions from wetlands north of 45°N over the 2013-2014 period as simulated by the UVic ESCM and estimated from the three UFMs (Peltola et al., 2019). The pattern and magnitude of simulated seasonal emissions compare well to that of the UFMs. For both the model and UFMs, minimal emissions vary between 0.2-0.6 Tg CH<sub>4</sub> month<sup>-1</sup> and occur in December while peak emissions are well below 10 Tg CH<sub>4</sub> month<sup>-1</sup> and occur in July (Fig. 9). However, simulated peak emissions (~8.5 Tg CH<sub>4</sub> month<sup>-1</sup>) are relatively higher than peak emissions for the UFMs (range of best estimates: 5.6-7.5 Tg CH<sub>4</sub> month<sup>-1</sup>). Moreover, in comparison to the three UFMs, the UVic ESCM simulates lower CH<sub>4</sub> emissions between December and May but higher CH<sub>4</sub> emissions between July and September (Fig. 9).

The UVic ESCM simulates the occurrence of wetland CH<sub>4</sub> emissions during the non-growing season. For wetlands north of 45°N, our model simulates total emissions of 2.1 Tg CH<sub>4</sub> yr<sup>-1</sup> between November and March. The UFMs predict total emissions of 4.6-10.2 Tg CH<sub>4</sub> yr<sup>-1</sup> during these cold months (Peltola et al., 2019). For wetlands north of 60°N, the UVic ESCM simulates emissions of 1.2 Tg CH<sub>4</sub> yr<sup>-1</sup> from October through May in agreement with recent estimates (1.6 ± 0.1 Tg CH<sub>4</sub> yr<sup>-1</sup>) from data-constrained model ensembles for these months (Treat et al., 2018). Based on our calculations, the three UFMs predict about 3.5-4.5 Tg CH<sub>4</sub> yr<sup>-1</sup> emitted from wetlands north of 60°N between October and May. Overall, this analysis shows that WETMETH is capable of simulating non-negligible CH<sub>4</sub> emissions from northern wetlands during cold months as emphasized by recent studies (Treat et al., 2018; Zona et al., 2016).



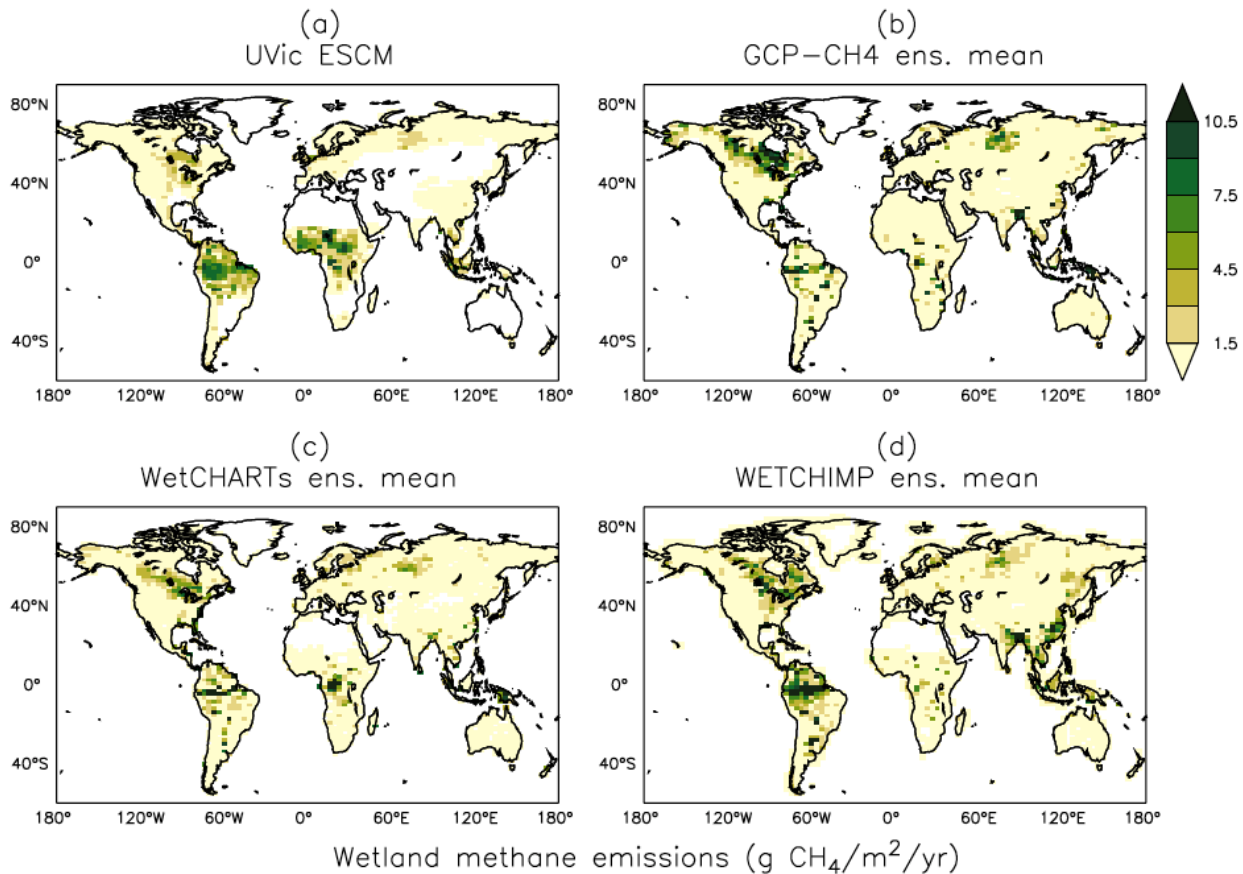
505 **Figure 9: Seasonal variations of CH<sub>4</sub> emissions from wetlands north of 45°N over the 2013-2014 period as simulated by the UVic ESCM in comparison to three upscaled flux measurements (RF-DYPTOP, RF-GLWD and RF-PEATMAP). The dashed lines show the uncertainty range for the upscaled flux measurements.**

### 5.2.2 Global emissions

The UVic ESCM simulates total emissions of 155.1 and 158.6 Tg CH<sub>4</sub> yr<sup>-1</sup> from global wetlands over the 2000-2009 and 2008-  
 510 2017 decades, respectively. According to the latest global CH<sub>4</sub> budget report, these wetland emissions are in the mid-range of bottom-up estimates (102-179 and 102-182 Tg CH<sub>4</sub> yr<sup>-1</sup>) but close to the lower limit of top-down estimates (153-196 and 155-200 Tg CH<sub>4</sub> yr<sup>-1</sup>) over the two decades (Saunois et al., 2020). Previous bottom-up estimates are significantly high (Melton et al., 2013; Saunois et al., 2016) primarily due to possible double counting of emissions from wetlands and other inland water areas (Saunois et al., 2020; Thornton et al., 2016) in addition to uncertainties associated with the extent of wetlands and model  
 515 parameterizations (Melton et al., 2013). Table 2 summarizes the comparison between the model results and estimates from the latest global CH<sub>4</sub> budget report for the 2008-2017 decade.

Fig. 10 shows the spatial distribution of simulated wetland CH<sub>4</sub> emissions over the 2001-2004 period in comparison to three process-based model ensembles: GCP-CH<sub>4</sub> (Poulter et al., 2017), WetCHARTs (Bloom et al., 2017), and WETCHIMP (Melton et al., 2013). The UVic ESCM simulates few CH<sub>4</sub>-emitting areas over South-East Asia in comparison to the three  
 520 model ensembles. The potential underestimation of wetland CH<sub>4</sub> emissions in that region is associated with the relatively few wetland areas simulated by the UVic ESCM (see Section 5.1). In tropical Africa, our model simulates too many CH<sub>4</sub>-emitting locations in comparison to the model ensembles (Fig. 10), which is also associated with the distribution of simulated wetlands (see Section 5.1). Nevertheless, the UVic ESCM simulates the occurrence of wetland CH<sub>4</sub> emissions in key source regions such as the Amazon and Congo River basins, South Sudan (Sudd swamps), and Indonesian islands (Fig. 10). For the Amazon

525 and Congo River basins, however, the UVic ESCM simulates lower wetland CH<sub>4</sub> emissions than predicted by the model ensembles (Fig. 10). This can be due to either the consideration of an optimal temperature for CH<sub>4</sub> production (around 27°C) in our model unlike many other process-based models, or the fact that model parameters in this study are tuned to northern estimates.

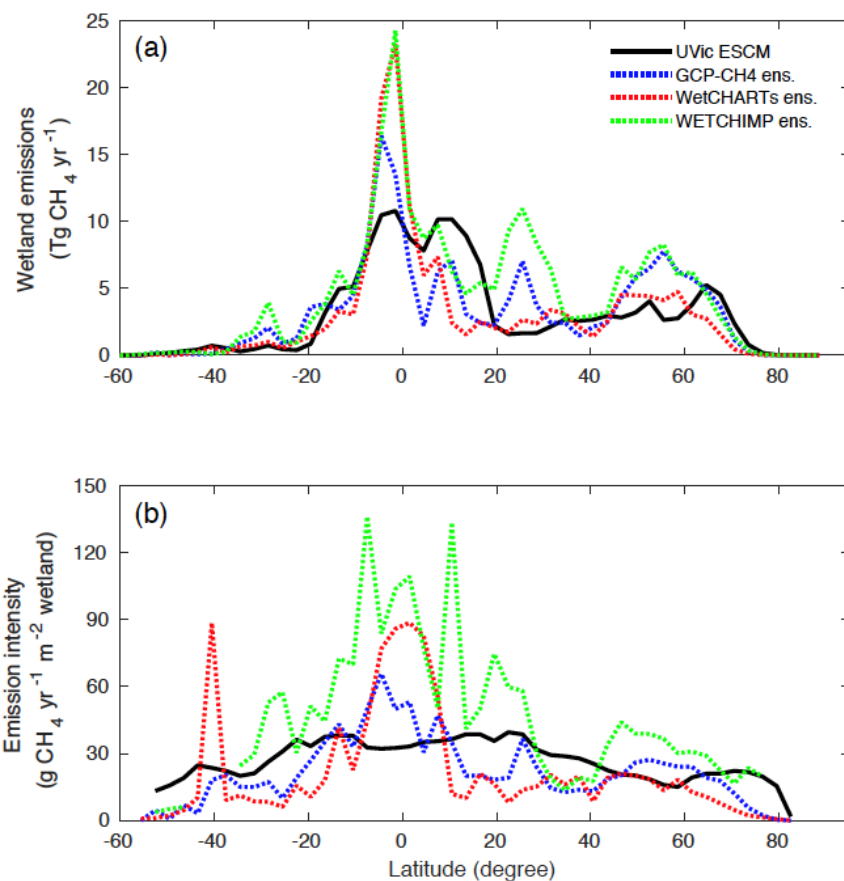


530 **Figure 10: Average methane emissions from global wetlands over the 2001-2004 period as simulated by the UVic ESCM (a) in comparison to three process-based model ensembles: (b) GCP-CH<sub>4</sub>, (c) WetCHARTs, and (d) WETCHIMP. The model ensembles are regridded to 3.6° x 1.8° for a fair comparison with the UVic ESCM. The comparison period corresponds to the overlap period for the three model ensembles.**

Fig. 11a shows the latitudinal distribution of simulated wetland CH<sub>4</sub> emissions in comparison to the model ensembles. Interestingly, although GCP-CH<sub>4</sub> and WetCHARTs are based on the same wetland dataset (SWAMPS-GLWD) (Bloom et al., 2017; Poulter et al., 2017), their zonal wetland CH<sub>4</sub> emissions are very different especially near the Equator and across northern high-latitude regions (Fig. 11a).

Using the three model ensembles as reference, the UVic ESCM simulates significantly lower wetland CH<sub>4</sub> emissions around the Equator (Fig. 11a), despite that the model simulates too large equatorial wetland areas (Fig. 4). In fact, wetland emission intensities (emissions per unit of wetland area) by the UVic ESCM are lower than those by the model ensembles

between 10°S and 10°N (Fig. 11b) due to relatively large wetland areas but small CH<sub>4</sub> emissions in equatorial regions (Fig. 4 versus Fig. 11a). As previously discussed, the relatively small CH<sub>4</sub> emissions simulated by the UVic ESCM in equatorial regions can be associated with either the optimal temperature for CH<sub>4</sub> production considered in WETMETH but not in most other process-based models, or the fact that model parameters in this study are tuned to northern estimates.



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**Figure 11: (a) Latitudinal distribution of wetland methane emissions simulated by the UVic ESCM over the 2001-2004 period in comparison to three process-based model ensembles: GCP-CH<sub>4</sub>, WetCHARTs and WETCHIMP. The comparison period corresponds to the overlap period for the three model ensembles. (b) Latitudinal emission intensity (methane emissions per unit of wetland area) simulated by the UVic ESCM over the 2001-2004 period in comparison to the three process-based model ensembles. GCP-CH<sub>4</sub> and WetCHARTs both use SWAMPS-GLWD as prescribed wetlands. The wetland methane emissions and emission intensities are summed across latitude bins of 3°.**

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Furthermore, the UVic ESCM simulates more wetland CH<sub>4</sub> emissions between 10-20°N than the three model ensembles (Fig. 11a) and this can be attributed to the widespread wetlands in West and Central Africa simulated by our model (Fig. 5 and Fig. 6). In addition, the UVic ESCM simulates significantly less wetland CH<sub>4</sub> emissions between 20-35°N in comparison to the WETCHIMP ensemble (Fig. 11a) and this can be attributed to the relatively small wetland areas simulated by the UVic ESCM in South-East Asia where some models include agricultural wetlands such as rice paddies. Moreover,



wetland emission intensities by the UVic ESCM feature low variability with latitude unlike the three model ensembles (Fig. 11b). Such a relative lack of variability can be attributed to two factors: (i) both wetland areas and CH<sub>4</sub> emissions simulated by the UVic ESCM feature relatively low variability with latitude compared to the datasets and model ensembles (Fig. 4 and Fig. 11a); and (ii) as previously discussed, our model likely simulates too large wetland areas but too small CH<sub>4</sub> emissions around the Equator implying a lack of variability across tropical latitudes.

Despite the various discrepancies between the UVic ESCM and both model ensembles regarding the distribution of wetland CH<sub>4</sub> emissions in the tropics, our model simulates mean annual CH<sub>4</sub> emissions from tropical wetlands that are in the range of estimates from the literature (Table 2). For the 1993-2004 period, the UVic ESCM simulates tropical wetland CH<sub>4</sub> emissions of 105.5 Tg CH<sub>4</sub> yr<sup>-1</sup> whereas the WETCHIMP ensemble predicts 126 ± 31 Tg CH<sub>4</sub> yr<sup>-1</sup> (Melton et al., 2013). Another study suggests a lower mean value (90 ± 77 Tg CH<sub>4</sub> yr<sup>-1</sup>) for wetland CH<sub>4</sub> emissions in the tropics although with large uncertainties (Sjögersten et al., 2014). Indeed, several studies indicate that wetland CH<sub>4</sub> emissions in the tropics are highly uncertain due to limited ground-based measurements and poorly delimited wetland extent (Dargie et al., 2017; Gumbrecht et al., 2016; Hu et al., 2018; Pangala et al., 2017; Saunio et al., 2020).

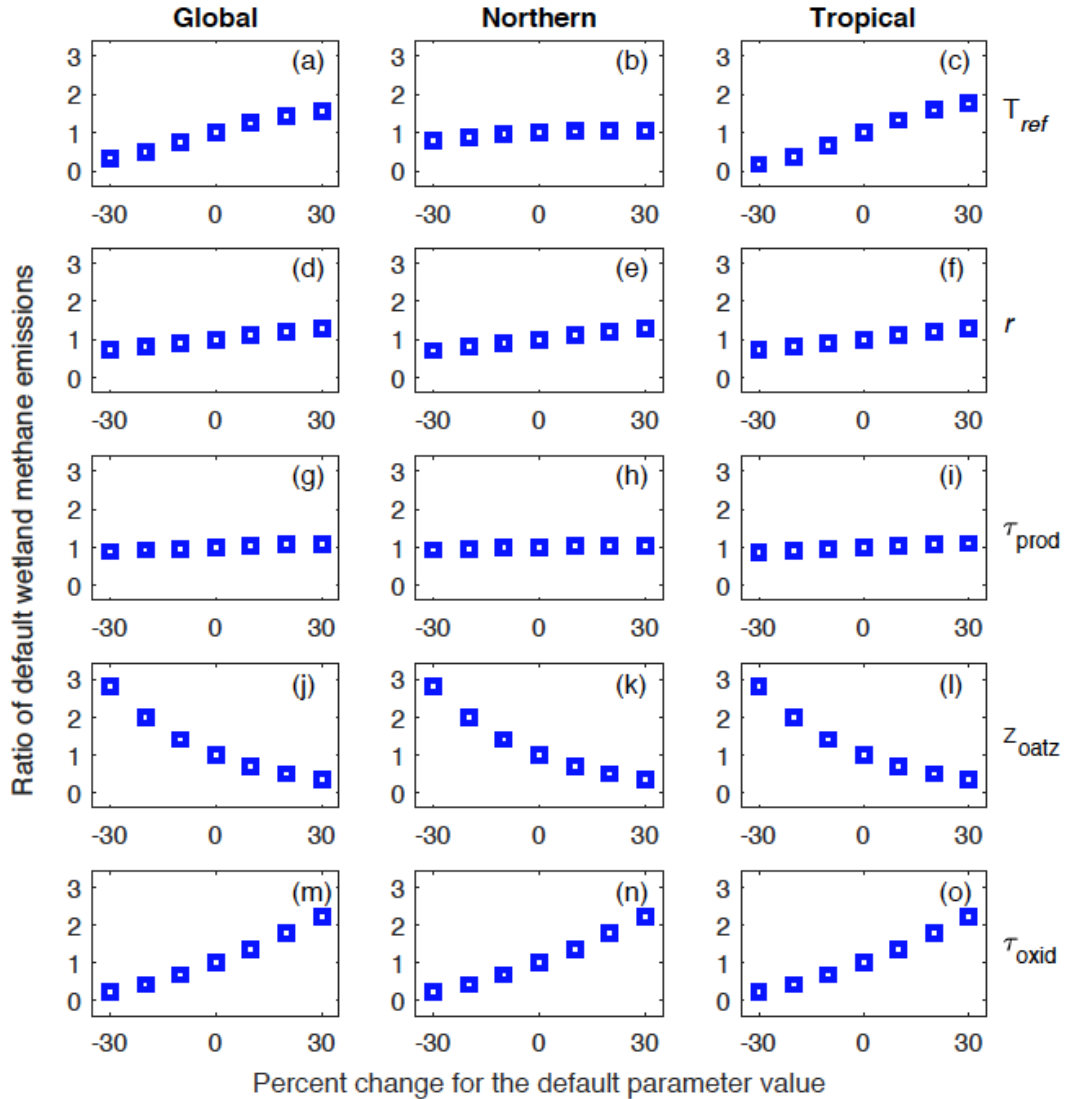
## 570 **6 Model sensitivity to poorly constrained parameters**

We performed a set of 30 model runs with perturbed parameter values (sensitivity runs) over the 2000-2009 decade in order to analyze the model sensitivity to poorly constrained parameters ( $T_{ref}$ ,  $\tau$ ,  $\tau_{prod}$ ,  $z_{oatz}$ , and  $\tau_{oxid}$ ). For each parameter, we increased or decreased the default value by 10, 20, and 30% while holding constant values for other parameters (fixed to default values). We then compared results from the sensitivity runs to the model simulation with all parameter values set to default values (control run). This comparison focuses on the total simulated global (90°S-90°N), northern (45-90°N), and tropical (30°S-30°N) wetland CH<sub>4</sub> emissions over the 2000-2009 decade.

Our results show that the model sensitivity varies with the different parameters and across regions (Fig. 12). Among the five poorly constrained parameters, the UVic ESCM is most sensitive to perturbations of the two parameters for CH<sub>4</sub> oxidation ( $z_{oatz}$  and  $\tau_{oxid}$ ) at both the global and regional scale. For  $z_{oatz}$ , a decrease (increase) of the default parameter value by 10-30% results in an augmentation (reduction) of default wetland CH<sub>4</sub> emissions by 41-179% (29-64%) at both the global and regional scale (Fig. 12j-l). For  $\tau_{oxid}$ , a decrease (increase) of the default parameter value by 10-30% implies a reduction (augmentation) of default wetland CH<sub>4</sub> emissions by 32-77% (37-120%) at both the global and regional scale (Fig. 12m-o).

The UVic ESCM is also very sensitive to perturbations of  $T_{ref}$ , but this sensitivity is more pronounced for tropical regions than northern regions (Fig. 12a-c). We recall that  $T_{ref}$  is used to define an optimal temperature for CH<sub>4</sub> production in WETMETH through the  $Q_{10}$  formulation (see Section 3.1.1 and Appendix A1). For northern regions, a decrease (increase) of  $T_{ref}$  by 10-30% results in a reduction (augmentation) of default wetland CH<sub>4</sub> emissions by 5-21% (3-5%). For tropical regions, however, a decrease (increase) of  $T_{ref}$  by 10-30% results in a reduction (augmentation) of default wetland CH<sub>4</sub> emissions by

34-82% (33-75%). Globally, a decrease (increase) of  $T_{ref}$  by 10-30% results in a reduction (augmentation) of default wetland  $CH_4$  emissions by 26-66% (24-55%). The model sensitivity to perturbations of  $r$  is linear across all regions (Fig. 12d-f). Lastly, the model is least sensitive to perturbation of  $\tau_{prod}$  across the globe (Fig. 12g-i).



595 **Figure 12: Analysis of the model sensitivity to perturbations of poorly constrained parameters:  $T_{ref}$ ,  $r$ ,  $\tau_{prod}$ ,  $z_{oatz}$ , and  $\tau_{oxid}$ .** For each parameter, the default value is increased or decreased by 10, 20, and 30% while values of other parameters are held constant (to default values). The model sensitivity is analyzed with respect to global (90°S-90°N), northern (45-90°N), and tropical (30°S-30°N) wetland methane emissions. Vertical axes show the ratio of the resulting emissions to the default emissions.

## 7 Discussions

### 7.1 WETMETH in the spectrum of wetland methane models

600 A recent study reviewed 40 models of CH<sub>4</sub> emissions in terrestrial ecosystems (predominantly rice paddies and natural wetlands) and classified them into three categories based on their level of complexity: relatively simple models, relatively mechanistic models, and mechanistic models (Xu et al., 2016). Relatively simple models are those that simulate net CH<sub>4</sub> emissions based on soil carbon or other environmental factors without explicit representations for the different CH<sub>4</sub> production and oxidation pathways as well as mechanisms transporting CH<sub>4</sub> to the atmosphere. Relatively mechanistic models are those that account for at least one transport mechanism for CH<sub>4</sub> release in addition to representing CH<sub>4</sub> production and oxidation with simple functions. Mechanistic models are more comprehensive and explicitly simulate different pathways for both CH<sub>4</sub> 605 production and oxidation, more than two mechanisms for CH<sub>4</sub> release, as well as their environmental controls. Based on this classification, WETMETH is a relatively simple model in the sense that it does not distinguish pathways for CH<sub>4</sub> production and oxidation as well as the various mechanisms transporting CH<sub>4</sub> to the atmosphere.

Although some wetland CH<sub>4</sub> models are claimed to be embedded in ESMs (Xu et al., 2016), none of these models are currently run in fully coupled models with feedbacks between climate conditions and the global carbon cycle. Most of 610 these models are rather implemented in dynamic vegetation models or uncoupled land surface components of climate models (Arora et al., 2018; Eliseev et al., 2008; Hodson et al., 2011; Riley et al., 2011; Ringeval et al., 2011; Wania et al., 2009). Nonetheless, relatively simple models present the ideal level of complexity for the current generation of ESMs. More complex models generally imply detailed soil chemistry for O<sub>2</sub> and alternate electron acceptors (Riley et al., 2011; Wania et al., 2010), different carbon substrates and their effects on CH<sub>4</sub> production (Grant, 1998; Lovley and Klug, 1986), an explicit representation 615 of the dynamics of different microbial communities (Grant, 1998; Xu et al., 2015), which all require comprehensive soil chemistry or model parameters that are currently not common in ESMs (Xu et al., 2016). Process parameterizations in mechanistic models generally imply too many degrees of freedom, making it difficult to constrain model parameters against sparse observations. Furthermore, mechanistic models may be too demanding computationally for fully coupled ESM runs without a proportional benefit for large-scale simulations of wetland CH<sub>4</sub> emissions.

620 The particularity of WETMETH among relatively simple models is that the model accounts for an optimum temperature for CH<sub>4</sub> production, a calibration of depth-dependent CH<sub>4</sub> production rates against potential CH<sub>4</sub> production rates from laboratory incubations, dynamic CH<sub>4</sub> oxidation based on the vertical distribution of soil moisture, and the potential for CH<sub>4</sub> emissions in non-inundated ecosystems with relatively high level of soil moisture content. In conclusion, WETMETH is simple enough to be compatible with ESMs and yet complex enough to simulate in an implicit way biogeochemical processes 625 regulating wetland CH<sub>4</sub> emissions.

## 7.2 Limitations for WETMETH

The developed wetland CH<sub>4</sub> model is associated with several limitations, which are linked to either its level of complexity or the scarcity of large-scale datasets for model calibration:

1. The present state of global wetland modelling assumes generic wetlands without distinguishing their different types (Melton et al., 2013; Poulter et al., 2017). Like many other large-scale models of the current generation, WETMETH would not be appropriate for investigating the contribution from particular wetland types to regional or global CH<sub>4</sub> emissions (Aselmann and Crutzen, 1989).
2. Since WETMETH is not based on a comprehensive soil biochemistry module and does not include the different pathways for CH<sub>4</sub> production and oxidation, the model is not suited for investigating the role of specific biological and chemical controls on wetland CH<sub>4</sub> emissions (Bridgham et al., 2013; Kwon et al., 2019).
3. WETMETH does not simulate the contribution from wetland-specific vegetation species to CH<sub>4</sub> emissions, although some of these species can either lead to high emissions (e.g. sedges are vascular plants that can transport CH<sub>4</sub> through their aerenchyma) or low emissions (e.g. mosses are non-vascular plants that have been shown to develop a symbiotic relationship with methanotrophs) (Bridgham et al., 2013; Chen and Murrell, 2010).
4. Ebullition and aerenchyma of vascular plants allow CH<sub>4</sub> produced in wetlands to escape to the atmosphere with little opportunity for oxidation (Segers, 1998; Whalen, 2005). Moreover, stems of woody trees are important conduits for CH<sub>4</sub> emissions in Amazonia, a major source region in the world (Pangala et al., 2017). By considering the net effect of all mechanisms transporting CH<sub>4</sub> to the atmosphere, WETMETH presents a limitation for investigating the relative contribution of transport mechanisms to CH<sub>4</sub> emissions across regions and at the global scale.
5. Methane produced in northern wetlands can be stored underneath frozen soil during the winter and be released abruptly upon spring thaw (Mastepanov et al., 2013; Song et al., 2012). WETMETH does not currently feature such a storage of CH<sub>4</sub> in the soil column, which is probably more relevant for small-scale (site) and short-term (daily) than large-scale (regional) and long-term (seasonal) emissions (Fig. 9).
6. While the existence of an optimal temperature for CH<sub>4</sub> production in wetlands is relatively well established in the literature (Dean et al., 2018), there are currently no estimates of such an optimal temperature for different climate zones across the globe. Previous studies suggest a range of 25-30°C for such an optimal temperature based on measurements of CH<sub>4</sub> production in northern wetlands (Dunfield et al., 1993; Metje and Frenzel, 2007). In WETMETH, we use a global value for this optimal temperature (~27°C) which is assumed to be valid for CH<sub>4</sub> production in both tropical and extra-tropical wetlands (see Section 3.1.1 and Appendix A1). However, our sensitivity analysis suggests that, in the present-day climate, wetland CH<sub>4</sub> emissions in the tropics are much more dependent on the optimal temperature for CH<sub>4</sub> production than wetland CH<sub>4</sub> emissions in the boreal and Arctic regions (see Section 6). The optimal temperature for CH<sub>4</sub> production in WETMETH, along with other factors such as areal wetland extents, contributes to inter-model differences in simulated wetland CH<sub>4</sub> intensities in the tropics (see Figure 11b).

660 7. As presented in this study, poorly constrained WETMETH parameters are tuned to estimates from northern high-latitude regions because large-scale datasets from other regions are scarce (see Section 4). A strong limitation comes with the assumption that the chosen parameter values are representative for CH<sub>4</sub> production and oxidation across the globe. However, the applied model calibration remains a reasonable approach given the scarcity of observations for wetland CH<sub>4</sub> production, oxidation, and emissions at the global scale.

665 Despite these limitations and the model simplicity, WETMETH is skillful when it comes to the simulation of mean seasonal, annual, and decadal wetland CH<sub>4</sub> emissions at the regional, hemispheric, and global scale (see Section 5.2). The implementation of WETMETH in a fully coupled ESM should advance research on the interactions between climate change and wetland CH<sub>4</sub> emissions in the context of global climate projections.

## 8 Conclusions

670 This paper introduces WETMETH – a process-based wetland CH<sub>4</sub> model developed for implementation in ESMs. WETMETH is currently embedded in the UVic ESCM, a fully coupled EMIC. WETMETH is a computationally efficient model, applicable globally and, of appropriate complexity with respect to the current state of wetland CH<sub>4</sub> modelling. Unconstrained model parameters are tuned to potential CH<sub>4</sub> production rates from incubated soil samples and CH<sub>4</sub> emissions from northern wetlands due to the scarcity of large-scale datasets from other regions. Nevertheless, WETMETH reproduces well estimates of mean annual CH<sub>4</sub> emissions over the past few decades at the regional, hemispheric, and global scale.

675 Despite the importance of tropical wetlands in the global CH<sub>4</sub> budget (Kirschke et al., 2013; Sauniois et al., 2016) and climate change (O'Connor et al., 2010; Zhang et al., 2017b), their areal extent and associated CH<sub>4</sub> emissions remain highly uncertain in both the literature and modelling work (including this study) due to a combination of limited ground-based measurements and process understanding (Pangala et al., 2017; Sauniois et al., 2020; Sjögersten et al., 2014), as well as a low accuracy from remotely-sensed products especially over dense rainforests of Indonesia, Amazonia, and the Congo River basin where new peatlands continue to be discovered to date (Dargie et al., 2017). Large-scale wetland mapping is a field of ongoing research (Tootchi et al., 2019) and further model development should focus on the improvement of wetland simulations in the tropics. In parallel, a compilation of tropical wetland CH<sub>4</sub> measurements from various sources into synthesis datasets would be beneficial for constraining wetland CH<sub>4</sub> processes in large-scale models.

685 The inclusion of wetland CH<sub>4</sub> processes in a fully coupled ESM allows to advance the research on the feedback between climate change and wetland CH<sub>4</sub> emissions. The implementation of WETMETH in the UVic ESCM constitutes an ideal tool for investigating interactions between climate conditions and wetland CH<sub>4</sub> emissions from decadal to longer timescales. Of particular importance is the permafrost carbon feedback to climate change, in which CH<sub>4</sub> emissions from northern wetlands are expected to play an important role (Nzotungicimpaye and Zickfeld, 2017).

## 690 **Author contributions**

CMN designed the research under the supervision of KZ, LFWL, JRM, and AHMD. LFWL contributed to the illustrated vertical profiles. CMN developed the wetland methane model with contributions from JRM and KZ. AHMD implemented the TOPMODEL approach in the UVic ESCM to which CMN applied a minor modification. CMN implemented the wetland methane model in the UVic ESCM with contributions from AHMD and ME. CMN performed the model calibration with  
695 contributions from CCT and KZ. CMN carried out the model simulations, evaluated the model performance, interpreted the results, and drafted the manuscript. All authors provided critical feedback on the manuscript and helped shape its final version.

## **Competing interests**

The authors declare that they have no conflict of interest.

## **Code availability**

700 The code for WETMETH 1.0 embedded in the University of Victoria Earth System Climate Model (UVic ESCM) version 2.9 used in this study is available at <https://doi.org/10.5281/zenodo.4066112> (Nzotungicimpaye and Zickfeld, 2020).

## **Data availability**

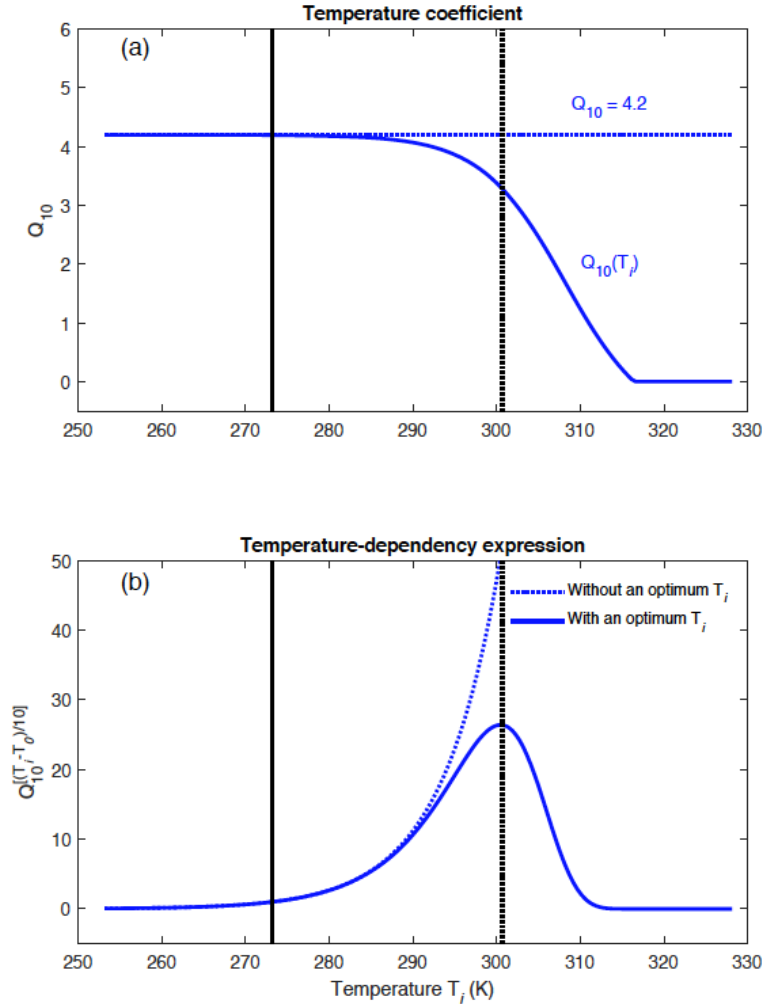
WETMETH output variables analyzed in this study are archived at <https://doi.org/10.20383/101.0215> and will be made accessible upon final publication of the manuscript.

## 705 **Acknowledgements**

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710 Bloom, Jed Kaplan, and Olli Peltola for making their methane emission datasets (WetCHARTs ensemble, WETCHIMP ensemble, and the upscaled flux measurements, respectively) publicly available.

## Appendix A: Temperature-dependent $Q_{10}$ coefficient for methane production

Fig. A1 illustrates the different shapes of the temperature-dependency function for  $\text{CH}_4$  production ( $Q_{10} \frac{T_i - T_0}{10}$ ;  $T_0 = 273.15 \text{ K}$ ) across a range of temperatures when considering: (i) a constant  $Q_{10}$  of 4.2; and (ii) a temperature-dependent  $Q_{10}$  coefficient given by  $Q_{10}(T_i) = 1.7 + 2.5 \tanh [0.1 (T_{ref} - T_i)]$ , where  $T_{ref} = 308.15 \text{ K}$ . The temperature-dependent  $Q_{10}(T_i)$  implies an optimal temperature for  $\text{CH}_4$  production in WETMETH around 300.15 K. When  $Q_{10}(T_i)$  decreases to reach negative values, its value in WETMETH is set to  $10^{-3}$  to represent a very small methanogenic response to temperature changes (Fig. A1).



720 **Figure A1: (a) Differences between a constant  $Q_{10}$  coefficient and a temperature-dependent  $Q_{10}(T_i)$  coefficients and (b) implications for the temperature-dependency expression for  $\text{CH}_4$  production ( $Q_{10}[(T_i - T_0)/10]$ ). The temperature-dependent coefficient  $Q_{10}(T_i) = 1.7 + 2.5 [\tanh(0.1 (308.15 - T_i))]$  allows to account for uncertainties in the  $Q_{10}$  coefficient and to define an optimal temperature for  $\text{CH}_4$  production around 300.15 K (dashed vertical line). The freezing point of water is shown at 273.15 K (continuous vertical line).**

## Appendix B: Applied minor modification to the TOPMODEL approach

The TOPMODEL approach implemented in the UVic ESCM is based on the formulation by Gedney and Cox for global land surface models (Gedney and Cox, 2003). This approach combines the simulated hydrology with a prescribed topographic index to determine the occurrence of wetlands (surface inundation) and soil moisture heterogeneity at the sub-grid scale. The occurrence of wetlands is simulated in an area whose local topographic index ( $\Lambda$ ) satisfies the following condition:

$$\Lambda_{\min} \leq \Lambda \leq \Lambda_{\max} , \quad (\text{B1})$$

where  $\Lambda_{\min}$  is a lower threshold that can be related to under-saturation conditions and  $\Lambda_{\max}$  is an upper threshold that can be related to over-saturation conditions.

In the initial work by Gedney and Cox,  $\Lambda_{\min}$  depends on the transmissivity of the entire soil column ( $T(0)$ ), the transmissivity of the soil column below the mean water table depth ( $z_w$ ) of the grid box ( $T(z_w)$ ) as well as the mean topographic index ( $\Lambda_{\text{mean}}$ ). It is calculated as  $\Lambda_{\min} = \ln \frac{T(0)}{T(z_w)} + \Lambda_{\text{mean}}$ . While  $\Lambda_{\text{mean}}$  is static and prescribed with a topographic index map, both transmissivities ( $T(0)$  and  $T(z_w)$ ) are simulated and non-static for a specific grid cell. Hence,  $\Lambda_{\min}$  is a non-static and grid-dependent threshold. Unlike  $\Lambda_{\min}$ ,  $\Lambda_{\max}$  is a static and global threshold. This threshold is applied to constrain the occurrence of wetlands in areas of stagnant water based on the assumption that locations where the water table rises well above the surface would be characterized by streamflow.

For the current study, a minor modification is applied to the above TOPMODEL approach. The revision consists of using a non-static and grid-dependent  $\Lambda_{\max}$  instead of a static and global threshold. Following the formulation by Comyn-Platt and colleagues (Comyn-Platt et al., 2018), an expression for  $\Lambda_{\max}$  that depends on  $\Lambda_{\min}$  is currently used in the UVic ESCM. This threshold is defined as:

$$\Lambda_{\max} = \Lambda_{\min} + \Lambda_{\text{range}} , \quad (\text{B2})$$

where  $\Lambda_{\text{range}}$  is a global tuning parameter ( $\Lambda_{\text{range}} = 0.93$  in the version of the UVic ESCM used in this study).

In summary, unlike the initial work by Gedney and Cox (Gedney and Cox, 2003), the modified TOPMODEL approach considers two non-static and grid-dependent thresholds ( $\Lambda_{\min}$  and  $\Lambda_{\max}$ ) for the identification of wetlands across the globe.

## Appendix C: Unit conversion for potential methane production rates

Here, we describe steps followed for converting units of maximum  $\text{CH}_4$  production rates measured in laboratory incubations from a soil weight basis ( $\mu\text{g C g DW}^{-1} \text{ hr}^{-1}$ ) to a soil volume basis ( $\text{kg C m}^{-3} \text{ s}^{-1}$ ). This unit conversion relies on the soil bulk density (BD in  $\text{g cm}^{-3}$ ) from the site of origin. The following two steps illustrate the applied unit conversion. In the first step, the potential  $\text{CH}_4$  production rates ( $P_{d,0}$ ) are converted from  $\mu\text{g C g DW}^{-1} \text{ hr}^{-1}$  to  $\mu\text{g C cm}^{-3} \text{ hr}^{-1}$  as follows:

$$P_{d,1} = (\text{BD}) P_{d,0} \quad (\text{C1})$$

Then, the conversion of  $P_{d,1}$  from  $\mu\text{g C cm}^{-3} \text{ hr}^{-1}$  to  $\text{kg C m}^{-3} \text{ s}^{-1}$  is done as follows:



$$P_{d,2} = \frac{\delta}{\gamma} P_{d,1}, \quad (\text{C2})$$

755 where  $\delta$  encompasses the conversion factors from  $\mu\text{g}$  to  $\text{kg}$  and from  $\text{cm}^{-3}$  to  $\text{m}^{-3}$  ( $\delta = 10^{-3} \text{ kg m}^{-3}$ ); and  $\gamma$  is the number of seconds per hour ( $\gamma = 3600 \text{ s}$ ).

## References

- Archer, D.: A data-driven model of the global calcite lysocline, *Global Biogeochem. Cycles*, 10, 511–526, 1996.
- 760 Arora, V. K., Boer, G. J., Friedlingstein, P., Eby, M., Jones, C. D., Christian, J. R., Bonan, G., Bopp, L., Brovkin, V., Cadule, P., Hajima, T., Ilyina, T., Lindsay, K., Tjiputra, J. F. and Wu, T.: Carbon-concentration and carbon-climate feedbacks in CMIP5 earth system models, *J. Clim.*, 26, 5289–5314, 2013.
- Arora, V. K., Melton, J. R. and Plummer, D.: An assessment of natural methane fluxes simulated by the CLASS-CTEM model, *Biogeosciences*, 15, 4683–4709, 2018.
- 765 Aselmann, I. and Crutzen, P. J.: Global distribution of natural freshwater wetlands and rice paddies, their net primary productivity, seasonality and possible methane emissions, *J. Atmos. Chem.*, 8, 307–358, 1989.
- Avis, C. A., Weaver, A. J. and Meissner, K. J.: Reduction in areal extent of high-latitude wetlands in response to permafrost thaw, *Nat. Geosci.*, 4, 444–448, 2011.
- 770 Barba, J., Bradford, M., Brewer, P., Bruhn, D., Covey, K., von Haren, J., Megonigal, J., Mikkelsen, T., Pangala, S., Pihlatie, M., Poulter, B., Rivas-Ubach, A., Schadt, C., Terazawa, K., Warner, D., Zhang, Z. and Vargas, R.: Methane emissions from tree stems: a new frontier in the global carbon cycle, *New Phytol.*, 222, 18–28, 2019.
- Blazewicz, S. J., Petersen, D. G., Waldrop, M. P. and Firestone, M. K.: Anaerobic oxidation of methane in tropical and boreal soils: Ecological significance in terrestrial methane cycling, *J. Geophys. Res. Biogeosciences*, 117, 1–9, 2012.
- Blodau, C.: Carbon cycling in peatlands - A review of processes and controls, *Environ. Rev.*, 10, 111–134, 2002.
- 775 Blodau, C., Basiliko, N. and Moore, T. R.: Carbon turnover in peatland mesocosms exposed to different water table levels, *Biogeochemistry*, 67, 331–351, 2004.
- Bloom, A. A., Bowman, K. W., Lee, M., Turner, A. J., Schroeder, R., Worden, J. R., Weidner, R., McDonald, K. C. and Jacob, D. J.: A global wetland methane emissions and uncertainty dataset for atmospheric chemical transport models (WetCHARTs version 1.0), *Geosci. Model Dev.*, 10, 2141–2156, 2017.
- 780 Bohn, T. J., Melton, J. R., Ito, A., Kleinen, T., Spahni, R., Stocker, B. D., Zhang, B., Zhu, X., Schroeder, R., Glagolev, M. V., Maksyutov, S., Brovkin, V., Chen, G., Denisov, S. N., Eliseev, A. V., Gallego-Sala, A., McDonald, K. C., Rawlins, M. A., Riley, W. J., Subin, Z. M., Tian, H., Zhuang, Q. and Kaplan, J. O.: WETCHIMP-WSL: Intercomparison of wetland methane emissions models over West Siberia, *Biogeosciences*, 12, 3321–3349, 2015.
- Bridgman, S. D., Cadillo-Quiroz, H., Keller, J. K. and Zhuang, Q.: Methane emissions from wetlands: biogeochemical, microbial, and modeling perspectives from local to global scales, *Glob. Chang. Biol.*, 19, 1325–1346, 2013.
- 785 Brune, A., Frenzel, P. and Cypionka, H.: Life at the oxic-anoxic interface: microbial activities and adaptations, *FEMS Microbiol. Rev.*, 24, 691–710, 2000.
- Cadillo-Quiroz, H., Bräuer, S., Yashiro, E., Sun, C., Yavitt, J. and Zinder, S.: Vertical profiles of methanogenesis and methanogens in two contrasting acidic peatlands in central New York State, USA, *Environ. Microbiol.*, 8, 1428–1440, 2006.
- 790 Chen, Y. and Murrell, J. C.: Methanotrophs in moss, *Nature*, 3, 595–596, 2010.
- Comyn-Platt, E., Hayman, G., Huntingford, C., Chadburn, S. E., Burke, E. J., Harper, A. B., Collins, W. J., Webber, C. P., Powell, T., Cox, P. M., Gedney, N. and Sitch, S.: Carbon budgets for 1.5 and 2°C targets lowered by natural wetland and permafrost feedbacks, *Nat. Geosci.*, 11, 568–573, 2018.
- 795 Conrad, R.: The global methane cycle: Recent advances in understanding the microbial processes involved, *Environ. Microbiol. Rep.*, 1, 285–292, 2009.
- Couwenberg, J., Dommain, R. and Joosten, H.: Greenhouse gas fluxes from tropical peatlands in south-east Asia, *Glob. Chang. Biol.*, 16, 1715–1732, 2010.
- Cox, P. M.: Description of the TRIFFID Dynamic Global Vegetation Model, Exeter, UK., 2001.
- 800 Dargie, G. C., Lewis, S. L., Lawson, I. T., Mitchard, E. T. A., Page, S. E., Bocko, Y. E. and Ifo, S. A.: Age, extent and carbon storage of the central Congo Basin peatland complex, *Nature*, 542, 86–90, 2017.
- Dean, J. F., Middelburg, J. J., Röckmann, T., Aerts, R., Blauw, L. G., Egger, M., Jetten, M. S. M., de Jong, A. E. E., Meisel, O. H., Rasigraf, O., Slomp, C. P., in't Zandt, M. and Dolman, A. J.: Methane feedbacks to the global climate system in a warmer world, *Rev. Geophys.*, 56, 207–250, 2018.
- 805 Dunfield, P., Knowles, R., Dumont, R. and Moore, T. R.: Methane production and consumption in temperate and subarctic peat soils: Response to temperature and pH, *Soil Biol. Biogeochem.*, 25, 321–326, 1993.

- Duval, T. and Radu, D.: Effect of temperature and soil organic matter quality on greenhouse gas production from temperate poor and rich fen soils, *Ecol. Eng.*, 114, 167–172, 2018.
- Eby, M., Zickfeld, K., Montenegro, A., Archer, D., Meissner, K. J. and Weaver, A. J.: Lifetime of anthropogenic climate change: Millennial time scales of potential CO<sub>2</sub> and surface temperature perturbations, *J. Clim.*, 22, 2501–2511, 2009.
- 810 Eliseev, A. V., Mokhov, I. I., Arzhanov, M. M., Demchenko, P. F. and Denisov, S. N.: Interaction of the methane cycle and processes in wetland ecosystems in a climate model of intermediate complexity, *Atmos. Ocean. Phys.*, 44, 147–162, 2008.
- Estop-Aragonés, C., Knorr, K.-H. and Blodau, C.: Controls on in situ oxygen and dissolved inorganic carbon dynamics in peats of a temperate fen, *J. Geophys. Res.*, 117, G02002, 2012.
- 815 Frolking, S., Roulet, N. T., Moore, T. R., Lafleur, P. M., Bubier, J. L. and Crill, P. M.: Modeling seasonal to annual carbon balance of Mer Bleue Bog, Ontario, Canada, *Global Biogeochem. Cycles*, 16, 1–21, 2002.
- Gauthier, M., Bradley, R. and Šimek, M.: More evidence that anaerobic oxidation of methane is prevalent in soils: Is it time to upgrade our biogeochemical models?, *Soil Biol. Biochem.*, 80, 167–174, 2015.
- Gedney, N. and Cox, P. M.: The sensitivity of global climate model simulations to the representation of soil moisture heterogeneity, *J. Hydrometeorol.*, 4, 1265–1275, 2003.
- 820 Gedney, N., Cox, P. M. and Huntingford, C.: Climate feedback from wetland methane emissions, *Geophys. Res. Lett.*, 31, L20503, 2004.
- Gedney, N., Huntingford, C., Comyn-Platt, E. and Wiltshire, A.: Significant feedbacks of wetland methane release on climate change and the causes of their uncertainty, *Environ. Res. Lett.*, 14, 084027, 2019.
- Girkin, N. T., Turner, B. L., Ostle, N., Craigan, J. and Sjögersten, S.: Root exudate analogues accelerate CO<sub>2</sub> and CH<sub>4</sub> production in tropical peat, *Soil Biol. Biochem.*, 117, 48–55, 2018.
- 825 Glagolev, M., Kleptsova, I., Filippov, I., Maksyutov, S. and Machida, T.: Regional methane emission from West Siberia mire landscapes, *Environ. Res. Lett.*, 6, 045214, 2011.
- Grant, R. F.: Simulation of methanogenesis in the mathematical model ecosys, *Soil Biol. Biochem.*, 30, 883–896, 1998.
- Gumbrecht, T., Roman-Cuesta, R., Verchot, L., Herold, M., Wittmann, F., Householder, E., Herold, N. and Murdiyarso, D.: An expert system model for mapping tropical wetlands and peatlands reveals South America as the largest contributor, *Glob. Chang. Biol.*, 23, 3581–3599, 2016.
- 830 Hegarty, T.: Temperature coefficient (Q<sub>10</sub>), seed germination and other biological processes, *Nature*, 243, 305–306, 1973.
- Helbig, M., Quinton, W. L. and Sontentag, O.: Warmer spring conditions increase annual methane emissions from a boreal peat landscape with sporadic permafrost, *Environ. Res. Lett.*, 12, 115009, 2017.
- 835 Hodson, E. L., Poulter, B., Zimmermann, N. E., Prigent, C. and Kaplan, J. O.: The El Niño-Southern Oscillation and wetland methane interannual variability, *Geophys. Res. Lett.*, 38, L08810, 2011.
- Hoehler, T. and Alperin, M.: Methane minimalism, *Nature*, 507, 436–437, 2014.
- Hoehler, T. M., Alperin, M. J., Albert, D. B. and Martens, C. S.: Field and laboratory studies of methane oxidation in an anoxic marine sediment: Evidence for a methanogen-sulfate reducer consortium, *Global Biogeochem. Cycles*, 8, 451–463, 1994.
- 840 Hopcroft, P. O., Valdes, P. J. and Beerling, D. J.: Simulating idealized Dansgaard-Oeschger events and their potential impacts on the global methane cycle, *Quat. Sci. Rev.*, 30, 3258–3268, 2011.
- Hu, H., Landgraf, J., Detmers, R., Borsdorff, T., Aan de Brugh, J., Aben, I., Butz, A. and Hasekamp, O.: Toward global mapping of methane with TROPOMI: First results and intersatellite comparison to GOSAT, *Geophys. Res. Lett.*, 45, 3682–3689, 2018.
- 845 Jauhiainen, J., Takahashi, H., Heikkinen, J. E. P., Martikainen, P. J. and Vasander, H.: Carbon fluxes from a tropical peat swamp forest floor, *Glob. Chang. Biol.*, 11, 1788–1797, 2005.
- Kim, Y.: Effect of thaw depth on fluxes of CO<sub>2</sub> and CH<sub>4</sub> in manipulated Arctic coastal tundra of Barrow, Alaska, *Sci. Total Environ.*, 505, 385–389, 2015.
- 850 Kirschke, S., Bousquet, P., Ciais, P., Saunois, M., Canadell, J. G., Dlugokencky, E. J., Bergamaschi, P., Bergmann, D., Blake, D. R., Bruhwiler, L., Cameron-Smith, P., Castaldi, S., Chevallier, F., Feng, L., Fraser, A., Heimann, M., Hodson, E. L., Houweling, S., Josse, B., Fraser, P. J., Krummel, P. B., Lamarque, J.-F., Langenfelds, R. L., Le Quééré, C., Naik, V., Palmer, P. I., Pison, I., Plummer, D., Poulter, B., Prinn, R. G., Rigby, M., Ringeval, B., Santini, M., Schmidt, M., Shindell, D. T., Simpson, I. J., Spahni, R., Paul Steele, L., Strode, S. A., Sudo, K., Szopa, S., Van der Werf, G. R., Voulgarakis, A., Van Weele, M., Weiss, R. F., Williams, J. E. and Zeng, G.: Three decades of global methane sources and sinks, *Nat. Geosci.*, 6, 813–823, 2013.
- 855

- Koven, C. D., Ringeval, B., Friedlingstein, P., Ciais, P., Cadule, P., Khvorostyanov, D., Krinner, G. and Tarnocai, C.: Permafrost carbon-climate feedbacks accelerate global warming, *Proc. Natl. Acad. Sci. U. S. A.*, 108, 14769–14774, 2011.
- Koven, C. D., Lawrence, D. M. and Riley, W. J.: Permafrost carbon–climate feedback is sensitive to deep soil carbon decomposability but not deep soil nitrogen dynamics, *Proc. Natl. Acad. Sci. U. S. A.*, 112, 3752–3757, 2015.
- 860 Kwon, M., Jung, J., Tripathi, B., Göckede, M., Lee, Y. and Kim, M.: Dynamics of microbial communities and CO<sub>2</sub> and CH<sub>4</sub> fluxes in the tundra ecosystems of the changing Arctic, *J. Microbiol.*, 57, 325–336, 2019.
- Loulergue, L., Schilt, A., Spahni, R., Masson-Delmotte, V., Blunier, T., Lemieux, B., Barnola, J., Raynaud, D., Stocker, T. and Chappellaz, J.: Orbital and millennial-scale features of atmospheric CH<sub>4</sub> over the past 800,000 years, *Nature*, 453, 383–386, 2008.
- 865 Lovley, D. R. and Klug, M. J.: Model for the distribution of sulfate reduction and methanogenesis in freshwater sediments, *Geochim. Cosmochim. Acta*, 50, 11–18, 1986.
- Lupascu, M., Wadham, E. R. C. and Pancost, R. D.: Temperature sensitivity of methane production in the permafrost active layer at Stordalen, Sweden: A comparison with non-permafrost northern wetlands, *Arctic, Antarct. Alp. Res.*, 44, 469–482, 2012.
- 870 MacDougall, A. H. and Knutti, R.: Projecting the release of carbon from permafrost soils using a perturbed parameter ensemble modelling approach, *Biogeosciences*, 13, 2123–2136, 2016.
- MacDougall, A. H., Avis, C. A. and Weaver, A. J.: Significant contribution to climate warming from the permafrost carbon feedback, *Nat. Geosci.*, 5, 719–721, 2012.
- Mastepanov, M., Sigsgaard, C., Tagesson, T., Ström, L., Tamstorf, M. P., Lund, M. and Christensen, T. R.: Revisiting factors controlling methane emissions from high-Arctic tundra, *Biogeosciences*, 10, 5139–5158, 2013.
- 875 Matthews, H. D., Weaver, A. J., Meissner, K. J., Gillett, N. P. and Eby, M.: Natural and anthropogenic climate change: Incorporating historical land cover change, vegetation dynamics and the global carbon cycle, *Clim. Dyn.*, 22, 461–479, 2004.
- McCalley, C. K., Woodcroft, B. J., Hodgkins, S. B., Wehr, R. A., Kim, E.-H., Mondav, R., Crill, P. M., Chanton, J. P., Rich, V. I., Tyson, G. W. and Saleska, S. R.: Methane dynamics regulated by microbial community response to permafrost thaw, *Nature*, 514, 478–481, 2014.
- Meissner, K. J., Weaver, A. J., Matthews, H. D. and Cox, P. M.: The role of land surface dynamics in glacial inception: A study with the UVic Earth System Model, *Clim. Dyn.*, 21, 515–537, 2003.
- Melton, J. R., Wania, R., Hodson, E. L., Poulter, B., Ringeval, B., Spahni, R., Bohn, T., Avis, C. A., Beerling, D. J., Chen, G., Eliseev, A. V., Denisov, S. N., Hopcroft, P. O., Lettenmaier, D. P., Riley, W. J., Singarayer, J. S., Subin, Z. M., Tian, H., Zürcher, S., Brovkin, V., Van Bodegom, P. M., Kleinen, T., Yu, Z. C. and Kaplan, J. O.: Present state of global wetland extent and wetland methane modelling: Conclusions from a model inter-comparison project (WETCHIMP), *Biogeosciences*, 10, 753–788, 2013.
- 885 Le Mer, J. and Roger, P.: Production, oxidation, emission and consumption of methane by soils: A review, *Eur. J. Soil Biol.*, 37, 25–50, 2001.
- 890 Metje, M. and Frenzel, P.: Methanogenesis and methanogenic pathways in a peat from subarctic permafrost, *Environ. Microbiol.*, 9, 954–964, 2007.
- Miller, S. M., Worthy, D. E. J., Michalak, A. M., Wofsy, S. C., Kort, E. A., Havice, T. C., Andrews, A. E., Dlugokencky, E. J., Kaplan, J. O., Levi, P. J., Tian, H. and Zhang, B.: Observational constraints on the distribution, seasonality, and environmental predictors of North American boreal methane emissions, *Global Biogeochem. Cycles*, 28, 146–160, 2014.
- 895 Mitsch, W. and Mander, Ü.: Wetlands and carbon revisited, *Ecol. Eng.*, 114, 1–6, 2018.
- Moore, T. and Roulet, N.: Methane flux: Water table relations in northern wetlands, *Geophys. Res. Lett.*, 20, 587–590, 1993.
- Moosavi, S. and Crill, P.: CH<sub>4</sub> oxidation by tundra wetlands as measured by a selective inhibitor technique, *J. Geophys. Res. Atmos.*, 103, 29093–29106, 1998.
- 900 Nzotungicimpaye, C.-M. and Zickfeld, K.: The contribution from methane to the permafrost carbon feedback, *Curr. Clim. Chang. Reports*, 3, 58–68, 2017.
- Nzotungicimpaye, C.-M. and Zickfeld, K.: The first version of WETMETH, a model for wetland methane emissions (WETMETH 1.0), Zenodo, doi:<https://doi.org/10.5281/zenodo.4066112>, 2020.
- 905 O’Connor, F. M., Boucher, O., Gedney, N., Jones, C. D., Folberth, G. A., Coppel, R., Friedlingstein, P., Collins, W. J., Chappellaz, J., Ridley, J. and Johnson, C. E.: Possible role of wetlands, permafrost, and methane hydrates in the methane

- cycle under future climate change: A review, *Rev. Geophys.*, 48, RG4005, 2010.
- Olefeldt, D., Turetsky, M. R., Crill, P. M. and McGuire, A. D.: Environmental and physical controls on northern terrestrial methane emissions across permafrost zones, *Glob. Chang. Biol.*, 19, 589–603, 2013.
- Orr, J. C.: On ocean carbon-cycle model comparison, *Tellus B*, 51, 509–510, 1999.
- 910 Pandey, S., Houweling, S., Krol, M., Aben, I., Monteil, G., Nechita-Banda, N., Dlugokencky, E. J., Detmers, R., Hasekamp, O., Xu, X., Riley, W. J., Poulter, B., Zhang, Z., McDonald, K. C., White, J. W. C., Bousquet, P. and Röckmann, T.: Enhanced methane emissions from tropical wetlands during the 2011 La Niña, *Sci. Rep.*, 7, 45759, 2017.
- Pangala, S. R., Enrich-Prast, A., Basso, L. S., Peixoto, R. B., Bastviken, D., Hornibrook, E. R. C., Gatti, L. V., Ribeiro, H., Calazans, L. S. B., Sakuragui, C. M., Bastos, W. R., Malm, O., Gloor, E., Miller, J. B. and Gauci, V.: Large emissions from floodplains trees close the Amazon methane budget, *Nature*, 522, 230–234, 2017.
- 915 Panikov, N. S. and Dedysh, S. N.: Cold season CH<sub>4</sub> and CO<sub>2</sub> emission from boreal peat bogs (West Siberia): Winter fluxes and thaw activation dynamics, *Global Biogeochem. Cycles*, 14, 1071–1080, 2000.
- Papa, F., Prigent, C., Aires, F., Jimenez, C., Rossow, W. B. and Matthews, E.: Interannual variability of surface water extent at the global scale, 1993-2004, *J. Geophys. Res. Atmos.*, 115, D12111, 2010.
- 920 Paudel, R., Mahowald, N. M., Hess, P. G. M., Meng, L. and Riley, W. J.: Attribution of changes in global wetland methane emissions from pre-industrial to present using CLM4.5-BGC, *Environ. Res. Lett.*, 11, 034020, 2016.
- Peltola, O., Vesala, T., Gao, Y., Rätty, O., Alekseychik, P., Aurela, M., Chojnicki, B., Desai, A. R., Dolman, A. J., Euskirchen, E. S., Friborg, T., Göckede, M., Helbig, M., Humphreys, E., Jackson, R. B., Jocher, G., Joos, F., Klatt, J., Knox, S. H., Kowalska, N., Kutzbach, L., Lienert, S., Lohila, A., Mammarella, I., Nadeau, D. F., Nilsson, M. B., Oechel, W. C., Peichl, M., Pypker, T., Quinton, W., Rinne, J., Sachs, T., Samson, M., Schmid, H. P., Sonnentag, O., Wille, C., Zona, D. and Aalto, T.: Monthly gridded data product of northern wetland methane emissions based on upscaling eddy covariance observations, *Earth Syst. Sci. Data*, 11, 1263–1289, 2019.
- 925 Pickett-Heaps, C. A., Jacob, D. J., Wecht, K. J., Kort, E. A., Wofsy, S. C., Diskin, G. S., Worthy, D. E. ., Kaplan, J. O., Bey, I. and Drevet, J.: Magnitude and seasonality of wetland methane emissions from the Hudson Bay Lowlands (Canada), *Atmos. Chem. Phys.*, 11, 3773–3779, 2011.
- 930 Poindexter, C. M., Baldocchi, D. D., Matthes, J. H., Knox, S. H. and Variano, E. A.: The contribution of an overlooked transport process to a wetland’s methane emissions, *Geophys. Res. Lett.*, 43, 6276–6284, 2016.
- Poulter, B., Bousquet, P., Canadell, J. G., Ciais, P., Peregón, A., Saunio, M., Arora, V. K., Beerling, D. J., Brovkin, V., Jones, C. D., Joos, F., Gedney, N., Ito, A., Kleinen, T., Koven, C. D., McDonald, K., Melton, J. R., Peng, C., Peng, S., Prigent, C., Schroeder, R., Riley, W. J., Saito, M., Spahni, R., Tian, H., Taylor, L., Viogy, N., Wilton, D., Wiltshire, A., Xu, X., Zhang, B., Zhang, Z. and Zhu, Q.: Global wetland contribution to 2000-2012 atmospheric methane growth rate dynamics, *Environ. Res. Lett.*, 12, 094013, 2017.
- Prigent, C., Matthews, E., Aires, F. and Rossow, W. B.: Remote sensing of global wetland dynamics with multiple satellite data sets, *Geophys. Res. Lett.*, 28, 4631–4634, 2001.
- 940 Prigent, C., Papa, F., Aires, F., Rossow, W. B. and Matthews, E.: Global inundation dynamics inferred from multiple satellite observations, 1993-2000, *J. Geophys. Res. Atmos.*, 112, D12107, 2007.
- Prigent, C., Papa, F., Aires, F., Jiménez, C., Rossow, W. B. and Matthews, E.: Changes in land surface water dynamics since the 1990s and relation to population pressure, *Geophys. Res. Lett.*, 39, L08403, 2012.
- Reeburgh, W.: Methane consumption in Cariaco Trench waters and sediments, *Earth Planet. Sci. Lett.*, 28, 337–344, 1976.
- 945 Rhodes, R. H., Brook, E. J., McConnell, J. R., Blunier, T., Sime, L. C., Fäin, X. and Mulvaney, R.: Atmospheric methane variability: Centennial-scale signals in the Last Glacial Period, *Global Biogeochem. Cycles*, 31, 575–590, 2017.
- Riley, W. J., Subin, Z. M., Lawrence, D. M., Swenson, S. C., Torn, M. S., Meng, L., Mahowald, N. M. and Hess, P.: Barriers to predicting changes in global terrestrial methane fluxes: Analyses using CLM4Me, a methane biogeochemistry model integrated in CESM, *Biogeosciences*, 8, 1925–1953, 2011.
- 950 Ringeval, B., Friedlingstein, P., Koven, C., Ciais, P., De Noblet-Ducoudré, N., Decharme, B. and Cadule, P.: Climate-CH<sub>4</sub> feedback from wetlands and its interaction with the climate-CO<sub>2</sub> feedback, *Biogeosciences*, 8, 2137–2157, 2011.
- Rivkina, E., Laurinavichius, K., McGrath, J., Tiedje, J., Shcherbakova, V. and Gilichinsky, D.: Microbial life in permafrost, *Adv. Sp. Res.*, 33, 1215–1221, 2004.
- Rogelj, J., Forster, P. M., Kriegler, E., Smith, C. J. and Séférian, R.: Estimating and tracking the remaining carbon budget for stringent climate targets, *Nature*, 571, 335–342, 2019.
- 955

- Roslev, P. and King, G.: Regulation of methane oxidation in a freshwater wetland by water table changes and anoxia, *FEMS Microbiol. Ecol.*, 19, 105–115, 1996.
- 960 Saunois, M., Bousquet, P., Poulter, B., Peregon, A., Ciais, P., Canadell, J. G., Dlugokencky, E. J., Etiope, G., Bastviken, D., Houweling, S., Mcdonald, K. C., Marshall, J., Melton, J. R., Morino, I., Naik, V., O'Doherty, S., Parmentier, F.-J. W., Patra, P. K., Peng, C., Peng, S., Peters, G. P., Pison, I., Prigent, C., Prinn, R., Ramonet, M., Riley, W. J., Saito, M., Santini, M., Schroeder, R., Simpson, I. J., Spahni, R., Steele, P., Takizawa, A., Thornton, B. F., Tian, H., Tohjima, Y., Viovy, N., Voulgarakis, A., Van Weele, M., Van Der Werf, G. R., Weiss, R., Wiedinmyer, C., Wilton, D. J., Wiltshire, A., Worthy, D., Wunch, D., Xu, X., Yoshida, Y., Zhang, B., Zhang, Z. and Zhu, Q.: The global methane budget 2000–2012, *Earth Syst. Sci. Data*, 8, 697–751, 2016.
- 965 Saunois, M., Stavert, A. R., Poulter, B., Bousquet, P., Canadell, J. G., Jackson, R. B., Raymond, P. A., Dlugokencky, E. J., Houweling, S., Patra, P. K., Ciais, P., Arora, V. K., Bastviken, D., Bergamaschi, P., Blake, D. R., Brailsford, G., Bruhwiler, L., Carlson, K. M., Carrol, M., Castaldi, S., Chandra, N., Crevoisier, C., Crill, P. M., Covey, K., Curry, C. L., Etiope, G., Frankenberg, C., Gedney, N., Hegglin, M. I., Höglund-Isaksson, L., Hugelius, G., Ishizawa, M., Ito, A., Janssens-Maenhout, G., Jensen, K. M., Joos, F., Kleinen, T., Krummel, P. B., Langenfelds, R. L., Laruelle, G. G., Liu, L., Machida, T., Maksyutov, S., McDonald, K. C., McNorton, J., Miller, P. A., Melton, J. R., Morino, I., Müller, J., Murguía-Flores, F., Naik, V., Niwa, Y., Noce, S., O'Doherty, S., Parker, R. J., Peng, C., Peng, S., Peters, G. P., Prigent, C., Prinn, R., Ramonet, M., Regnier, P., Riley, W. J., Rosentreter, J. A., Segers, A., Simpson, I. J., Shi, H., Smith, S. J., Steele, L. P., Thornton, B. F., Tian, H., Tohjima, Y., Tubiello, F. N., Tsuruta, A., Viovy, N., Voulgarakis, A., Weber, T. S., Van Weele, M., Van der Werf, G. R., Weiss, R. F., Worthy, D., Wunch, D., Yin, Y., Yoshida, Y., Zhang, W., Zhang, Z., Zhao, Y., Zheng, B., Zhu, Q., Zhu, Q. and Zhuang, Q.: The global methane budget 2000–2017, *Earth Syst. Sci. Data*, 12, 1561–1623, 2020.
- 970 Schipper, L. A., Hobbs, J. K., Rutledge, S. and Arcus, V. L.: Thermodynamic theory explains the temperature optima of soil microbial processes and high Q10 values at low temperatures, *Glob. Chang. Biol.*, 20, 3578–3586, 2014.
- Schmittner, A., Oeschles, A., Matthews, H. D. and Galbraith, E. D.: Future changes in climate, ocean circulation, ecosystems, and biogeochemical cycling simulated for a business-as-usual CO2 emission scenario until year 4000 AD, *Global Biogeochem. Cycles*, 22, GB1013, 2008.
- 980 Schneider von Deimling, T., Meinshausen, M., Levermann, A., Huber, V., Frieler, K., Lawrence, D. M. and Brovkin, V.: Estimating the near-surface permafrost-carbon feedback on global warming, *Biogeosciences*, 9, 649–665, 2012.
- Schneider von Deimling, T., Grosse, G., Strauss, J., Schirmer, L., Morgenstern, A., Schaphoff, S., Meinshausen, M. and Boike, J.: Observation-based modelling of permafrost carbon fluxes with accounting for deep carbon deposits and thermokarst activity, *Biogeosciences*, 12, 3469–3488, 2015.
- 985 Schuur, E. A. G., McGuire, A. D., Schädel, C., Grosse, G., Harden, J. W., Hayes, D. J., Hugelius, G., Koven, C. D., Kuhry, P., Lawrence, D. M., Natali, S. M., Olefeldt, D., Romanovsky, V. E., Schaefer, K., Turetsky, M. R., Treat, C. C. and Vonk, J. E.: Climate change and the permafrost carbon feedback, *Nature*, 520, 171–179, 2015.
- Segers, R.: Methane production and methane consumption: A review of processes underlying wetland methane fluxes, *Biogeochemistry*, 41, 23–51, 1998.
- 990 Shindell, D. T., Walter, B. P. and Faluvegi, G.: Impacts of climate change on methane emissions from wetlands, *Geophys. Res. Lett.*, 31, L21202, 2004.
- Singleton, C. M., McCalley, C. K., Woodcroft, B. J., Boyd, J. A., Evans, P. N., Hodgkins, S. B., Chanton, J. P., Froelking, S., Crill, P. M., Saleska, S. R., Rich, V. I. and Tyson, G. W.: Methanotrophy across a natural permafrost thaw environment, *ISME J.*, 12, 2544–2558, 2018.
- 995 Sjögersten, S., Black, C., Evers, S., Hoyos-Santillan, J., Wright, E. and Turner, B.: Tropical wetlands: A missing link in the global carbon cycle?, *Global Biogeochem. Cycles*, 28, 1371–1386, 2014.
- Sjögersten, S., Aplin, P., Gauci, V., Peacock, M., Siegenthaler, A. and Turner, B. L.: Temperature response of ex-situ greenhouse gas emissions from tropical peatlands: Interactions between forest type and peat moisture conditions, *Geoderma*, 324, 47–55, 2018.
- 1000 Smemo, K. A. and Yavitt, J. B.: Anaerobic oxidation of methane: An underappreciated aspect of methane cycling in peatland ecosystems?, *Biogeosciences*, 8, 779–793, 2011.
- Song, C., Xu, X., Sun, X., Tian, H., Sun, L., Miao, Y., Wang, X. and Guo, Y.: Large methane emission upon spring thaw from natural wetlands in the northern permafrost region, *Environ. Res. Lett.*, 7, 034009, 2012.
- 1005 Taylor, K. E., Stouffer, R. J. and Meehl, G. A.: An overview of CMIP5 and the experiment design, *Bull. Am. Meteorol. Soc.*,

- 93, 485–498, 2012.
- Thompson, R. L., Sasakawa, M., Machida, T., Aalto, T., Worthy, D., Lavric, J. V., Myhre, C. L. and Stohl, A.: Methane fluxes in the high northern latitudes for 2005–2013 estimated using a Bayesian atmospheric inversion, *Atmos. Chem. Phys.*, 17, 3553–3572, 2017.
- 1010 Thornton, B., Wik, M. and Crill, P. M.: Double-counting challenges the accuracy of high-latitude methane inventories, *Geophys. Res. Lett.*, 43, 12569–12577, 2016.
- Tokarska, K. and Gillett, N.: Cumulative carbon emissions budgets consistent with 1.5°C global warming, *Nat. Clim. Chang.*, 8, 296–299, 2018.
- 1015 Tootchi, A., Jost, A. and Ducharne, A.: Multi-source global wetland maps combining surface water imagery and groundwater constraints, *Earth Syst. Sci. Data*, 11, 189–220, 2019.
- Treat, C. C., Natali, S. M., Ernakovich, J., Iversen, C. M., Lupascu, M., McGuire, A. D., Norby, R. J., Roy Chowdhury, T., Richter, A., Santruckova, H., Schädel, C., Schuur, E. A. G., Sloan, V. L., Turetsky, M. R. and Waldrop, M. P.: A pan-Arctic synthesis of CH<sub>4</sub> and CO<sub>2</sub> production from anoxic soil incubations, *Glob. Chang. Biol.*, 21, 2787–2803, 2015.
- Treat, C. C., Bloom, A. A. and Marushchak, M. E.: Nongrowing season methane emissions—a significant component of annual emissions across northern ecosystems, *Glob. Chang. Biol.*, 24, 3331–3343, 2018.
- 1020 Walz, J., Knoblauch, C., Böhme, L. and Pfeiffer, E. M.: Regulation of soil organic matter decomposition in permafrost-affected Siberian tundra soils - Impact of oxygen availability, freezing and thawing, temperature, and labile organic matter, *Soil Biol. Biochem.*, 110, 34–43, 2017.
- Wania, R., Ross, I. and Prentice, I. C.: Integrating peatlands and permafrost into a dynamic global vegetation model: 1. Evaluation and sensitivity of physical land surface processes, *Global Biogeochem. Cycles*, 23, GB3014, 2009.
- 1025 Wania, R., Ross, I. and Prentice, I. C.: Implementation and evaluation of a new methane model within a dynamic global vegetation model: LPJ-WHyMe v1.3.1, *Geosci. Model Dev.*, 3, 565–584, 2010.
- Wania, R., Melton, J. R., Hodson, E. L., Poulter, B., Ringeval, B., Spahni, R., Bohn, T., Avis, C. A., Chen, G., Eliseev, A. V., Hopcroft, P. O., Riley, W. J., Subin, Z. M., Tian, H., Van Bodegom, P. M., Kleinen, T., Yu, Z. C., Singarayer, J. S., Zürcher, S., Lettenmaier, D. P., Beerling, D. J., Denisov, S. N., Prigent, C., Papa, F. and Kaplan, J. O.: Present state of global wetland extent and wetland methane modelling: methodology of a model inter-comparison project (WETCHIMP), *Geosci. Model Dev.*, 6, 617–641, 2013.
- 1030 Weaver, A. J., Eby, M., Wiebe, E. C., Bitz, C. M., Duffy, P. B., Ewen, T. L., Fanning, A. F., Holland, M. M., MacFayden, A., Matthews, H. D., Meissner, K. J., Saenko, O., Schmittner, A., Wang, H. and Yoshimori, M.: The UVic Earth System Climate Model: Model description, climatology, and applications to past, present and future climates, *Atmosphere-Ocean*, 39, 361–428, 2001.
- Whalen, S. C.: Biogeochemistry of methane exchange between natural wetlands and the atmosphere, *Environ. Eng. Sci.*, 22, 73–94, 2005.
- 1040 Wheeler, B. D.: Water and plants in freshwater wetlands, in *Eco-Hydrology*, edited by A. J. Baird and R. L. Wilby, pp. 127–180, Routledge, London., 1999.
- Whiticar, M. J. and Faber, E.: Methane oxidation in sediment and water column environments-Isotope evidence, *Adv. Org. Geochemistry*, 10, 759–768, 1985.
- 1045 Wild, B., Gentsch, N., Čapek, P., Diáková, K., Alves, R. J. E., Bárta, J., Gittel, A., Hugelius, G., Knoltsch, A., Kuhry, P., Lashchinskiy, N., Mikutta, R., Palmtag, J., Schleper, C., Schneckner, J., Shibistova, O., Takriti, M., Torsvik, V. L., Urich, T., Watzka, M., Šantrůčková, H., Guggenberger, G. and Richter, A.: Plant-derived compounds stimulate the decomposition of organic matter in arctic permafrost soils, *Sci. Rep.*, 6, 25607, 2016.
- Wu, Y., Versegny, D. and Melton, J.: Integrating peatlands into the coupled Canadian Land Surface Scheme (CLASS) v3.6 and the Canadian Terrestrial Ecosystem Model (CTEM) v2.0, *Geosci. Model Dev.*, 9, 2639–2663, 2016.
- 1050 Xu, X., Elias, D. A., Graham, D. E., Phelps, T. J., Carroll, S. L., Wullschlegler, S. D. and Thornton, P. E.: A microbial functional group-based module for simulating methane production and consumption: Application to an incubated permafrost soil, *J. Geophys. Res. G Biogeosciences*, 120, 1315–1333, 2015.
- Xu, X., Yuan, F., Hanson, P. J., Wullschlegler, S. D., Thornton, P. E., Riley, W. J., Song, X., Graham, D. E., Song, C. and Tian, H.: Reviews and syntheses: Four decades of modeling methane cycling in terrestrial ecosystems, *Biogeosciences*, 13, 3735–3755, 2016.
- 1055 Yvon-Durocher, G., Allen, A. P., Bastviken, D., Conrad, R., Gudas, C., St-Pierre, A., Thanh-Duc, N. and Del Giorgio, P. A.:

- Methane fluxes show consistent temperature dependence across microbial to ecosystem scales, *Nature*, 507, 488–491, 2014.
- Zalman, C. A., Meade, N., Chanton, J., Kostka, J. E., Bridgham, S. D. and Keller, J. K.: Methylo-trophic methanogenesis in sphagnum-dominated peatland soils, *Soil Biol. Biochem.*, 118, 156–160, 2018.
- 1060 Zhang, B., Tian, H., Lu, C., Chen, G., Pan, S., Anderson, C. and Poulter, B.: Methane emissions from global wetlands: An assessment of the uncertainty associated with various wetland extent data sets, *Atmos. Environ.*, 165, 310–321, 2017a.
- Zhang, Z., Zimmermann, N. E., Stenke, A., Li, X., Hodson, E. L., Zhu, G., Huang, C. and Poulter, B.: Emerging role of wetland methane emissions in driving 21st century climate change, *Proc. Natl. Acad. Sci. U. S. A.*, 114, 9647–9652, 2017b.
- 1065 Zhang, Z., Zimmermann, N. E., Calle, L., Hurtt, G., Chatterjee, A. and Poulter, B.: Enhanced response of global wetland methane emissions to the 2015-2016 El Niño-Southern Oscillation event, *Environ. Res. Lett.*, 13, 074009, 2018.
- Zhu, Q., Peng, C., Chen, H., Fang, X., Liu, J., Jiang, H., Yang, Y. and Yang, G.: Estimating global natural wetland methane emissions using process modelling: spatio-temporal patterns and contributions to atmospheric methane fluctuations, *Glob. Ecol. Biogeogr.*, 24, 959–972, 2015.
- 1070 Zickfeld, K., Eby, M., Matthews, H. D. and Weaver, A. J.: Setting cumulative emissions targets to reduce the risk of dangerous climate change, *Proc. Natl. Acad. Sci. U. S. A.*, 106, 16129–16134, 2009.
- Zona, D., Gioli, B., Commane, R., Lindaas, J., Wofsy, S. C., Miller, C. E., Dinardo, S. J., Dengel, S., Sweeney, C., Karion, A., Chang, R. Y.-W., Henderson, J. M., Murphy, P. C., Goodrich, J. P., Moreaux, V., Liljedahl, A., Watts, J. D., Kimball, J. S., Lipson, D. A. and Oechel, W. C.: Cold season emissions dominate the Arctic tundra methane budget, *Proc. Natl. Acad. Sci. U. S. A.*, 113, 40–45, 2016.
- 1075