



Reviews and syntheses: Four Decades of Modeling Methane Cycling in Terrestrial Ecosystems

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Abstract

Over the past four decades, a number of numerical models have been developed to quantify the magnitude, investigate the spatial and temporal variations, and understand the underlying mechanisms and environmental controls of methane (CH₄) fluxes within terrestrial ecosystems. These CH₄ models
20 are also used for integrating multi-scale CH₄ data, such as laboratory-based incubation and molecular analysis, field observational experiments, remote sensing, and aircraft-based measurements across a variety of terrestrial ecosystems. Here we summarize 39 terrestrial CH₄ models to characterize their strengths and weaknesses and to suggest a roadmap for future model improvement and application. Our key findings are that: (1) the focus of CH₄ models has shifted from theoretical to site- and regional-level
25 applications over the past four decades, (2) large discrepancies exist among models in terms of representing CH₄ processes and their environmental controls, and (3) significant data-model and model-model mismatches are partially attributed to different representations of landscape characterization and inundation dynamics. Three areas for future improvements and applications of terrestrial CH₄ models are: (1) CH₄ models should more explicitly represent the mechanisms underlying land-atmosphere CH₄



30 exchange, with emphasis on improving and validating individual CH₄ processes over depth and
horizontal space, (2) models should be developed that are capable of simulating CH₄ emissions across
highly heterogeneous spatial and temporal scales, particularly hot moments and hot spots, and (3)
efforts should be invested to develop model benchmarking frameworks that can easily be used for
model improvement, evaluation, and integration with data from molecular to global scales.

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1. Introduction

Methane (CH₄) has ~26 times the warming potential of carbon dioxide (CO₂) over a 100-year
horizon and is the second most important anthropogenic greenhouse gas, accounting for ~15% of
anthropogenic forcing to climate change (Forster et al., 2007; IPCC, 2013; Rodhe, 1990). Therefore, an
40 accurate estimate of CH₄ exchanges between land and the atmosphere is fundamental for understanding
climate change (Bridgham et al., 2013; Nazaries et al., 2013; Spahni et al., 2011). The ecosystem
modeling approach has been one of the most broadly used integrative tools for examining mechanistic
processes, quantifying the budget of CH₄ flux across spatial and temporal scales (Arah and Kirk, 2000;
Arah and Stephen, 1998; Cao et al., 1995; Curry, 2007; Fung et al., 1991; Huang et al., 1998b; Nouchi
45 et al., 1994; Potter, 1997; Riley et al., 2011; Walter et al., 1996; Xu et al., 2007; Zhuang et al., 2004),
and predicting future flux (Anisimov, 2007). Specifically, many CH₄ models have been developed to
integrate data, improve process understanding, quantify budgets, and project exchanges with the
atmosphere under a changing climate (Cao et al., 1995; Grant, 1998; Huang et al., 1998a; Potter, 1997;
Riley et al., 2011; Tian et al., 2010; Zhuang et al., 2004). In addition, model sensitivity analyses help to
50 design field and laboratory experiments by identifying the most uncertain processes and parameters in
the models (Massman et al., 1997; Xu, 2010).

Based on the complexity of the CH₄ processes represented, CH₄ models fall into two broad
categories: (1) empirical models to estimate and extrapolate measured methanogenesis, methanotrophy,
or CH₄ emission at plot, country, or continental scales (Christensen et al., 1996; Eliseev et al., 2008;
55 Mokhov et al., 2007; Wania et al., 2010, 2009); and (2) process-based models used for prognostic



understanding of individual CH₄ processes in response to multiple environmental drivers and budget quantification (reviewed below).

Over the past decades, many empirical and process-based models have been developed, for example CASA (Potter, 1997), CH₄MOD (Huang et al., 1998b), CLM4Me (Riley et al., 2011), 60 DAYCENT (Del Grosso et al., 2000), DLEM (Tian et al., 2010; Xu and Tian, 2012), DNDC (Li, 2000a), *ecosys* (Grant, 1998), HH (Cresto-Aleina et al., 2015), MEM (Cao et al., 1995), TEM (Zhuang et al., 2004), etc. However, recent analyses and model inter-comparisons have shown that most of these models poorly reproduce regional- to global-scale observations (Bohn and Lettenmaier, 2010; Bohn et al., 2015; Melton et al., 2013; Wania et al., 2013). A comprehensive synthesis and evaluation of the 65 mechanisms incorporated into these models is lacking. In this paper we summarize CH₄ models published over the past four decades, their evolution in terms of process representation, and their coupling with Earth System Models. We pay special attention to the key processes in CH₄ cycling, specifically CH₄ substrate cycling, methanogenesis, methanotrophy, and transport in the soil profile, and their environmental controls. Emphasis is given to how these mechanisms were simulated in 70 various models and how they were categorized in terms of complexity and ecosystem function. Models for understanding reactions in bioreactors (Bhadra et al., 1984; Pareek et al., 1999), mining plots (De Visscher and Van Cleemput, 2003), and marine systems (Elliott et al., 2011) were excluded.

2. Primary CH₄ Processes

Biological methane production in sediments was first noted in the late 18th century (Wolfe, 75 2004), and the microbial oxidation of methane was proposed at the beginning of the 20th century (Hanson and Hanson, 1996). Since then, methane cycling processes have been intensively studied and documented (Christensen et al., 1996; Hakemian and Rosenzweig, 2007; Lai, 2009; Melloh and Crill, 1996; Mer and Roger, 2001), and most have been described mathematically and incorporated into ecosystem models (Table 1). Herein, we do not attempt to review all CH₄ processes, as a number of 80 reviews have been published on this topic (Barlett and Harriss, 1993; Blodau, 2002; Bridgham et al., 2013; Cai, 2012; Chen et al., 2012; Conrad, 1995; Conrad, 1996; Hakemian and Rosenzweig, 2007; Higgins et al., 1981; Lai, 2009; Monechi et al., 2007; Segers, 1998; Wahlen, 1993). Rather, we focus on



85 primary CH₄ processes in terrestrial ecosystems, and their environmental controls from a modeling
perspective. In this context there exist three major methanogenesis mechanisms, two CH₄
methanotrophy mechanisms, and (depending on how one counts) seven CH₄ transport pathways in
plants and soils. We note that most models do not explicitly represent all of these transport pathways,
and that the relative importance of these pathways varies substantially in time and space. We also pay
attention to several other modeling features including capability for plot- or regional-level simulations,
vertical representation of biogeochemical processes, and whether the model is embedded in an Earth
90 System Model.

The published literature concludes that two processes dominate biological CH₄ production
(Conrad, 1999; Krüger et al., 2001): acetoclastic and hydrogenotrophic methanogenesis, which account
for ~50% - 90% and ~10% - 43% of global annual CH₄ produced, respectively (Conrad and Klose,
1999; Kotsyurbenko et al., 2004; Mer and Roger, 2001; Summons et al., 1998). Methylo-trophic
95 methanogenesis (producing CH₄ from methanol, methylamines, or dimethylsulfide) is usually
considered a minor contributor of CH₄, but may be significant in marine systems (Summons et al.,
1998). The proportion of CH₄ produced via any of these pathways varies widely in time, space, and
across ecosystem types.

Methanotrophy occurs under aerobic (Gerard and Chanton 1993) and anaerobic (Smemo and
00 Yavitt 2011) conditions. These oxidative processes can occur in several locations in soil and plants
(Frenzel and Rudolph 1998, Heilman and Carlton 2001, Ström et al. 2005) and using CH₄ either
produced in the soil column or transported from the atmosphere (Mau et al. 2013). Large variation in
the relative magnitudes of these pathways as a percentage of total methanotrophy has been observed:
aerobic oxidation of CH₄ in soil contributes 1% - 90% (King, 1996; Ström et al., 2005), anaerobic
05 oxidation of CH₄ within the soil profile contributes 0.3% - 5% (Blazewicz et al., 2012; Murase and
Kimura, 1996), oxidation of CH₄ during transport in plant aerenchyma contributes <1% (Frenzel and
Karofeld, 2000; Frenzel and Rudolph, 1998), and oxidation of atmospheric CH₄ contributes ~10 – 100%
(~10% for wetland and ~100% for upland) (Gulledge and Schimel, 1998a; Gulledge and Schimel,
1998b; Topp and Pattey, 1997) to total methanotrophy. CH₄ is transported from the soil profile to the



10 atmosphere in typical open-water wetlands by several pathways: diffusive and advective plant-mediated
transport accounts for 12~98% (Butterbach-Bahl et al., 1997; Mer and Roger, 2001; Morrissey and
Livingston, 1992), soil gaseous diffusion accounts for ~5% for wetlands and > 90% for upland systems
(Barber et al., 1988; Mer and Roger, 2001) (soil gaseous advection and aqueous diffusion and advection
are typically relatively small (Tang et al., 2013), and ebullition accounts for 10~60% (Chanton et al.,
15 1989; Tokida et al., 2007) of the CH₄ produced in the soil that is emitted to the atmosphere. These
processes vary significantly, depending on temporal and spatial scales.

Environmental factors affecting CH₄ processes have many direct and indirect controls. The
dominant direct factors controlling methanogenesis and methanotrophy in most systems include oxygen
availability, dissolved organic carbon concentration, soil pH, soil temperature, soil moisture, nitrate and
20 other reducers, ferric iron, microbial community structure, active microbial biomass, wind speed
(Askaer et al. 2011), plant root structure (Nouchi et al. 1990), etc. Indirect factors include soil texture
and mineralogy, vegetation, air temperature, soil fauna, nitrogen input, irrigation, agricultural practices,
sulfate reduction, and carbon quality, etc. (Banger et al., 2012; Bridgham et al., 2013; Hanson and
Hanson, 1996; Higgins et al., 1981; Mer and Roger, 2001). The complicated effects induced by a few
25 key factors on CH₄ processes have been mathematically described and incorporated in many CH₄
models; for example, direct factors such as temperature, moisture, oxygen availability, soil pH, and soil
redox potential (Grant, 1998; Riley et al., 2011; Tian et al., 2010; Zhuang et al., 2004). The indirect
factors such as nitrogen input (Banger et al., 2012), irrigation (Wassmann et al., 2000), and agricultural
practices were not reviewed in this study as their impacts are indirect and were modeled through
30 impacts on vegetation and hydrology (Li, 2000a; Ren et al., 2011; Xu et al., 2010).

3. Model Representation of CH₄ Processes

[Insert Figure 1 here]

We reviewed 39 CH₄ models (Fig. 1 & Table 1), which were developed for a variety of
purposes. The first CH₄ model was published in 1986 by Lovley & Klug (1986) to simulate
35 *methanogenesis* in water sediments, and since then a number of CH₄ models have been developed and
applied at numerous scales (Table 1). For example, *Cao et al.* developed the Methane Emission Model



(MEM) and applied it to quantify the global CH₄ source in rice paddies and the sensitivity of the global CH₄ budget's response to climate change (Cao et al., 1995; Cao et al., 1998). Walter *et al.* developed and applied an ecosystem CH₄ model to quantify global CH₄ emission in response to water level
40 fluctuations (Walter and Heimann, 2000; Walter et al., 1996). Grant *et al.* (1998) developed the *ecosys* model, which is currently the ecosystem-scale model that most mechanistically represents the many kinetic processes and microbial mechanisms for methanogenesis, methanotrophy, and CH₄ emission (Grant and Roulet, 2002). Huang *et al.* developed the CH₄MOD model to investigate CH₄ processes and flux in rice paddies (Huang et al., 1998b; Huang et al., 2004; Li et al., 2012). Zhuang *et al.*
45 developed a CH₄ module within the terrestrial ecosystem model (TEM) framework and applied it at site- and regional-levels (Zhuang et al., 2004; Zhuang et al., 2006), particularly across high latitudes (Fan et al., 2013; Zhu et al., 2011, 2013a, b, 2014). Tian et al. (2010) developed the dynamic land ecosystem model (DLEM), which is capable of simulating methanogenesis, methanotrophy, and CH₄ transport in terrestrial ecosystems; the scale of its application has ranged widely (i.e. plot, country,
50 continent, and global) for budget estimation and attribution analysis (Ren et al., 2011; Tian et al., 2015; Tian et al., 2011; Xu and Tian, 2012; Xu et al., 2010). Riley et al (2011) developed CLM4Me, a CH₄ module for the Community Land Model, which is incorporated in the Community Earth System Model. The family of LPJ models (LPJ-Bern, LPJ-WHyMe, LPJ-WSL) was developed under the LPJ framework to simulate CH₄ processes, but with different modules for CH₄ cycling; for example, LPJ-
55 Bern and LPJ-WHyMe incorporate Walter's CH₄ module (Walter and Heimann, 2000; Walter et al., 1996; Wania et al., 2009) while LPJ-WSL incorporates the CH₄ module from Christensen et al (Christensen et al., 1996).

[Insert Figure 2 here]

The number of CH₄ models has steadily increased since the 1980s (Figs. 1 & 2): 1 in the 1980s,
60 12 in the 1990s, 14 in the 2000s, and 12 for 2010-2015. This rapid increase in CH₄ model development indicates the rapidly growing effort to analyze CH₄ cycling and quantify CH₄ budgets across spatial scales. Meanwhile, the key mechanisms represented in the models have changed at a slower pace (Fig. 2). The most important changes are representation of vertically-resolved processes within the soil and



regional model simulation. For example, the percentage of the newly developed models with vertically-
65 resolved CH₄ biogeochemistry has increased from 54% before 2000 to ~83% in the recent decade
(2010-2015). The proportion of models with regional simulation capability has doubled from ~50%
before the 2010s to almost 100% afterwards (Fig. 2).

[Insert Tables 1, 2, and 3 here]

The majority of these models were designed to simulate land-surface exchanges in saturated
70 ecosystems (primarily natural wetlands and rice paddies) (Huang et al., 1998b; Li, 2000a; Walter et al.,
1996) (Table 1). Not all of the models explicitly represented the belowground mechanistic processes for
CH₄ production and consumption and the primary carbon biogeochemical processes (Christensen et al.,
1996; Ding and Wang, 1996). The land-atmosphere CH₄ exchange is a net balance of many processes
75 including production, oxidation, and transport, which are represented in models with different
complexities (Table 2). Some models are very complicated, while some are relatively simple. The
obvious tradeoff in modeling CH₄ cycling is to represent mechanisms as accurately as possible while
managing complexity (Evans et al., 2013), and ensuring that additional complexity enhances
predictability (Tang and Zhuang, 2008).

4. CH₄ Model Classification

80 *[Insert Figure 3 here]*

Current CH₄ models can be classified into three groups based on their representation of primary
mechanistic processes for methanogenesis, methanotrophy (Fig. 3), and CH₄ transport (Fig. 4). The first
group of CH₄ models uses a very simple framework for land-surface CH₄ flux, and most were
developed before the 2000s (e.g., Christensen's model, CASA, etc.) (Fig 3A). These models treated
85 land-surface CH₄ flux as an empirical function and link it to environmental controls, or soil organic
carbon; this group of models ignored the mechanistic processes of methanogenesis, methanotrophy, and
CH₄ transport. The second group of CH₄ models considers processes in a relatively simple manner (e.g.,
one or two primary CH₄ transport pathways, methanogenesis as a function of DOC, oxidation of
atmospheric CH₄, etc.); however, the methanogenesis and methanotrophy mechanisms are still not
90 mechanistically represented (Fig. 3B). The third group of CH₄ models explicitly simulates the processes



for methanogenesis, methanotrophy, and CH₄ transport as well as their environmental controls, which allows comprehensive investigation of physical, chemical, or biological processes' contribution to land-surface CH₄ flux (Fig. 3C). Of the models in the third group, none of them fully represent all these processes (although some have most of the features described); for example, the *ecosys* model is one of the few models to represent most of the CH₄ cycling processes shown in Fig. 3C, although it has not been embedded in an Earth System Model. We recommend that the third group of CH₄ models be the focus of mechanistic studies, and the basis for improving reduced form models applicable to Earth System Model applications.

4.1. Methanogenesis

Models make use of four types of modeling frameworks (Table 3) to relate methanogenesis to substrate requirements. Similar to Eqs (1) – (4) in Table 3, the model representation of methanogenesis can be classified into four types: (1) empirical association between methanogenesis and environmental condition, including temperature and water table; (2) empirical correlation of methanogenesis with biological variables (particularly heterotrophic respiration and soil organic matter); (3) methanogenesis as a function of concentration of substrate (DOC); and (4) a suite of mechanistic processes simulated for methanogenesis.

Representation of the substrate for methanogenesis may be a key aspect of simulating CH₄ cycling in terrestrial ecosystems; however, more than half of the models we examined do not explicitly simulate substrates for methanogenesis. We note, however, that explicit representation of substrates and their effects on methanogenesis requires additional model parameters, and therefore degrees of freedom in the model, which can lead to increased equifinality (Tang and Zhuang, 2008). The optimum complexity level for methanogenesis and consumption models remains to be determined.

The first group of models correlates methanogenesis with environmental factors and ignores substrate production and its influence on methanogenesis [Eq. (1)] (Table 3). This group of models includes Christensen's model (Christensen et al., 1996), which simulates the net flux of CH₄ based on fraction of saturated soil column and soil temperature, and the IAP-RAS model (Mokhov et al., 2007), which calculates methanogenesis as an empirical equation of soil temperature. This group of models has



a role in site-specific interpolation of observations for scaling over time at a given site, but does not explicitly represent carbon or acetate substrate.

20 The second group of models directly links methanogenesis with heterotrophic respiration or soil organic matter content, but does not explicitly represent carbon or acetate substrate availability [Eq. (2)]; examples are the LPJ model family (Hodson et al., 2011; Spahni et al., 2011; Wania et al., 2010, 2009) and CLM4Me (Riley et al., 2011).

25 The third group of models simulates dissolved organic carbon (DOC) or different pools of soil organic carbon, which are treated as a substrate pool influencing CH₄ production [Eq. (3)]; examples are the MEM model (Cao et al., 1995; Cao et al., 1998) and DLEM (Tian et al., 2010).

30 The fourth group of CH₄ models considers the primary substrates for methanogenesis, that is, acetate and single-carbon compounds [Eq. (4)]; examples are Kettunen's model (Kettunen, 2003), Segers' model (Segers and Kengen, 1998; Segers and Leffelaar, 2001a, b; Segers et al., 2001), van Bodegom's model (van Bodegom et al., 2000; Van Bodegom et al., 2001), and the *ecosys* model (Grant, 1998).

35 Methanogenesis is a fundamental process for CH₄ cycling, and a majority of models simulate methanogenesis in either implicit or explicit ways (Tables 2 & 3). For example, 31 models (i.e. Arah's model, "Cartoon" model, CASA, CH4MOD, Christensen's model, CLM4Me, Ding's model, DLEM, DNDC, *ecosys*, Gong's model, HH model, IAP-RAS, Kettunen's model, Lovley's model, LPJ-Brn, LPJ-WHyMe, LPJ-WSL, Martens' model, MEM, MERES, ORCHIDEE, SDGVM, Segers' model, TEM, TRIPLEX-GHG, UW-VIC, van Bodegom's model, VISIT, Walter's model, and Xu's model) simulate methanogenesis as one individual process. As a comparison, only three out of 39 CH₄ models reviewed explicitly simulate two methanogenesis pathways (acetoclastic methanogenesis and
40 hydrogenotrophic methanogenesis) (Table 3). As mentioned earlier, it is well-recognized that there are two dominant methanogenesis pathways and their relative combination changes significantly across environmental gradients, for example, along the soil profile (Falz et al., 1999) and across landscape types (McCalley et al., 2014). This lack of representation of two methanogenesis mechanisms might have caused dramatic bias and needs to be address in future model improvements.



45 4.2. Methanotrophy

Methanotrophy is another important process for simulating the land-atmosphere exchange of CH₄ (Table 2). Aerobic and anaerobic methanotrophy occurs in different locations in the soil profile, and affect both methanogenesis in the profile and CH₄ diffusing in from the atmosphere. For example, the oxidation of atmospheric CH₄, rhizosphere and bulk soil oxidation, and oxidation during CH₄ transport from soil to the atmosphere have been measured and modeled (Tables 1 & 2). Anaerobic CH₄ oxidation has been measured (Blazewicz et al., 2012) and has proposed to be incorporated into ecosystem models (Gauthier et al., 2015).

It has been confirmed that the aerobic oxidation of CH₄ produced in the soil profile and aerobic oxidation of atmospheric CH₄ play a major role in CH₄ consumption in the system, and that anaerobic oxidation of CH₄ is a minor contributor. Currently, no models explicitly simulate the anaerobic oxidation of CH₄ in soil, although a few recent studies highlighted the importance of this process (Blazewicz et al., 2012; Caldwell et al., 2008; Conrad, 2009; Smemo and Yavitt, 2011; Valentine and Reeburgh, 2000). The key reasons for this omission are that the process has not been mathematically described, the key parameters are uncertain (Gauthier et al., 2015), and the biochemical mechanism is not fully understood.

The Michaelis-Menten-like equations, widely used for simulating CH₄ production and oxidation, consider substrates limiting factors (Segers and Kengen, 1998). A few CH₄ models in the third category (linking methanogenesis with a substrate) use the Michaelis-Menten-like equation to compute methanogenesis and methanotrophy rates (Eqs. 3, 5, & 6). For example, DLEM simulates methanogenesis as a function of DOC concentration and other environmental controls, and Michaelis-Menten-like functions were used to compute methanogenesis on the basis of DOC as substrate. Methanotrophy has been simulated with dual Monod Michaelis-Menten-like equations with CH₄ and oxygen as limiting factors (Table 3). We note that the Michaelis-Menten-like relations may be inaccurate when representing multi-substrate, multi-consumer networks (Tang and Riley 2013, 2015). Although their approach (Equilibrium Chemistry Approximation, ECA) has not been applied for simulations of CH₄ emissions, CH₄ dynamics are inherently multi-consumer, including transformations



associated with methanogens, heterotrophs, ebullition, advection, diffusion, and aerenchyma transport, even if only one substrate is considered.

4.3. CH₄ within the Soil/Water Profile

75 CH₄ produced in the soil profile or below the water table is not transported immediately into the atmosphere. The time required for CH₄ to migrate from deep soil profile to the atmosphere ranges from minutes to days (depending on temperature, water, soil texture, and emissivity of plant roots), or even a season if the surface is frozen. The majority of current CH₄ models assume that CH₄ transport to the atmosphere occurs immediately after CH₄ is produced, and a portion is oxidized (Tian et al., 2010; 80 Zhuang et al., 2004); for models simulating CH₄ flux over minutes to days, the lack of modeled transport may produce unrealistic simulations.

Some models do simulate CH₄ dynamics within the soil and water profile (e.g., ecosys, CLM4Me), which produces a lag between methanogenesis and emission, allowing for oxidation to be explicitly represented during transport, and is valuable for simulating the seasonality of CH₄ flux (Table 85 2). For example, the recently observed CH₄ burst in the spring season in some field experiments confirms that the storage of CH₄ produced in winter will likely produce a strong emission outburst (Song et al., 2012). Without the mechanism of CH₄ storage beneath the soil surface, this phenomenon is impossible to simulate. In most of the models considering CH₄ storage, the CH₄ is treated as a simple gas pool, under the water table, which will be transported to the atmosphere through several transport 90 pathways.

4.4. CH₄ Transport from Soil to the Atmosphere

The transport of CH₄ produced and stored in soil column is the final bottleneck for CH₄ leaving the system; therefore, this process is an important control on the instantaneous land-surface CH₄ flux. Several important pathways of CH₄ transport to the atmosphere are identified: plant-mediated diffusive and advective transport, aqueous and gaseous diffusion, and ebullition (Beckett et al., 2001; Chanton, 95 2005; Mer and Roger, 2001; Whiting and Chanton, 1996). Model simulation of these transport pathways uses direct control of simulated land surface CH₄ flux, with CH₄ transport simulation considered in a manner similar to Eq. (7) (Table 3).



00 The majority (77%) of the current models simulate at least one transport pathway. Specifically,
66% of the models simulate CH₄ transport via aerenchyma, 77% simulate gaseous diffusive transport,
and 54% simulate ebullition transport (Table 1). More than 50% of models simulated these three
transport pathways. Some models simulate explicitly the aqueous and gaseous diffusion of CH₄ (Riley
et al., 2011), while most models do not simulate advective transport. Many models simulate diffusion
and plant-mediated transport in very simple ways. For model improvement in this area, three issues
05 remain as challenges:

(1) Most models treat transport implicitly; for example, the diffusion processes is treated
simply as an excessive release of CH₄ when its concentration exceeds a threshold (Tian et al.,
2010). This treatment prevents the model from simulating the lag between methanogenesis and
its final release to the atmosphere, which has been confirmed to be the key mechanism for hot-
10 moment and hot-spot of CH₄ flux (Song et al., 2012) and for oxidation during transport.

(2) The parameters for plant species capable of transporting gas (i.e., *aerenchyma*) are
poorly constrained (Riley et al. 2011), although plant-mediated transport has been identified as
the dominant pathway for CH₄ emission in most natural wetlands (Aulakh et al., 2000; Colmer,
2003).

(3) Simultaneously representing aqueous and gaseous phases of CH₄ is one potentially
important issue for simulating CH₄ transport from soil to the atmosphere (Tang and Riley,
2014). However, these processes are only explicitly represented in a few extant CH₄ models
15 (Riley et al., 2011; Grant et al., 1998).

4.5. Environmental Controls on CH₄ Processes

20 Although a suite of environmental factors affects various CH₄ processes, many of these factors
are not explicitly simulated in many models. These factors include soil temperature, soil moisture,
substrate, soil pH, soil redox potential, and oxygen availability. Many other factors not directly
incorporated in the models, could indirectly affect CH₄ cycling. For example, nitrogen fertilizer affects
methanogenesis through its stimulating impacts on ecosystem productivity, which in turn affects DOC,
25 soil moisture and soil temperature (Xu et al., 2010). The CLM4Me model simulates permafrost and its



effects on CH₄ dynamics, and has a simple relationship for soil pH impacts on methanogenesis (Riley et al., 2011). In this review, we specifically focus on temperature, moisture, and pH because these factors directly affect CH₄ processes in all environments, and they have been explicitly simulated in the many of the models.

30 Three types of mathematical functions have been used to simulate the temperature dependence of CH₄ processes: (1) linear functions of air or soil temperature (Eq. 10 in Table 3), (2) Q₁₀ function (Eq. 10 in Table 4), and (3) Arrhenius type function (Eq. 12 in Table 3). Of these three model representations of temperature dependence, the Q₁₀ equation is the most common mathematical description. However, the parameters for these empirical functions vary widely across the models
35 (Table 4). Actual temperature responses may diverge significantly from the models at low temperatures, close to the freezing point of water, and high temperatures, close to the denaturation point of enzymes.

[Insert Table 4 here]

Soil moisture is another important factor controlling CH₄ processes, because water limits O₂ diffusion from the air through the soil column and because microbes can become stressed at low matric
40 potential. CH₄ is produced typically under conditions with a low reduction potential, which is normally associated with long-term inundation. Although methanogenesis occurs solely under reducing conditions (methanogenesis within plant biomass under aerobic condition has never been simulated although it has been reported in experiments (Keppler et al., 2006)), methanotrophy occurs under drier, aerobic conditions. A low water content can also limit microbial activity in frozen soils or soils with
45 high osmolarity (Watanabe and Ito, 2008). Therefore, soil moisture has different impacts on different CH₄ processes. Four types of model representation are used to simulate moisture's effects on CH₄ processes (Eqs. 12-15 in Table 3).

- (1) Methanogenesis occurs only in the saturated zone and an exponential function for soil moisture is used to control methanotrophy (e.g., CLM4Me);
- 50 (2) Linear function for moisture impacts (e.g., CLASS use linear function for moisture impact on methanotrophy) (Curry, 2007);



- (3) Reciprocal responsive curves for moisture impacts on methanogenesis and methanotrophy (e.g., DLEM) (Tian et al., 2010);
- (4) A bell-shaped curve for methanogenesis (e.g., TEM uses a function similar to Eq. (16) for moisture impacts) (Zhuang et al., 2004).

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The pH is another important factor that has been included in a number of CH₄ models (Cao et al., 1995; Zhuang et al., 2004). Methanogens and methanotrophs depend on proton and sodium ion translocation for energy conservation, thus they are directly affected by pH. The pH impacts on CH₄ processes are simulated as a bell-shaped curve although the mathematical functions used to describe pH impacts are different (Eq. 17a, 17b, and 17c). Moreover, even when the same functions were used in different models, they were associated with different parameter values; for example, the MEM model sets pH_{min} (minimum pH value for CH₄ processes being active), pH_{opt} (optimal pH value for CH₄ processes being most active), and pH_{max} (maximum pH value for CH₄ processes being active) values of 5.5, 7.5, and 9 (Cao et al., 1995). This set of parameter values was adopted in the TEM model (Zhuang et al., 2004), whereas the DLEM model uses values of 4, 7, and 10 (Tian et al., 2010). The CLM4Me model uses a different function while keeping the impact curve at the same shape, but its peak has an optimal pH of 6.2 (Meng et al., 2012).

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For the other environmental factors, model representation is still in its infancy; however, several models consider oxygen availability as an electron acceptor for methanotrophy (e.g., Arah's model, Beckett's model, "Cartoon" model, CLM4Me, *ecosys*, Kettunen's model, MERES, Segers' model, van Bodegom's model, De Visscher's model, and Xu's model). In addition, only a few models simulate the impacts of the electron acceptor (i.e. nitrate, sulfate, etc.) on CH₄ processes (Table 2). For example, van Bodegom's model simulates iron biogeochemistry, and Lovley's, Marten's, and van Bodegom's models simulate sulfate as the electron acceptor and its impacts on methanogenesis and methanotrophy (Lovley and Klug, 1986; Martens et al., 1998; Van Bodegom et al., 2001). Explicitly representing these processes enables future coupling of CH₄ cycling to processes that are regionally significant, such as iron reduction on the Alaskan North Slope (Miller et al., 2015). These models' representation has the



advantage of more accurately simulating biogeochemical processes of carbon and ions, although large uncertainties still exist because of the lack of data for constraining model parameters.

80 5. Summary

Through the four decades of modeling CH₄ cycling in terrestrial ecosystems, consensus has been reached on several fronts. First, CH₄ cycling includes a suite of complicated processes, and both the simple and complex models are able to estimate land-surface CH₄ flux to a certain level of confidence, although models of different complexity do provide different results (Tang et al., 2010). Second, 85 although a number of CH₄ models have been developed, several gaps remain that need new model representations (e.g., dynamic linkage between inundation dynamics and the CH₄ module (Melton et al., 2013), anaerobic oxidation of CH₄ (Gauthier et al., 2015)).

Two recent CH₄ model-model inter-comparison projects raised several important points (Bohn et al., 2015; Melton et al., 2013): (1) the distribution of the inundation area is important for accurately 90 simulating global CH₄ emissions, but was poorly represented in CH₄ models; (2) the modeled response of land-surface CH₄ emission to elevated CO₂ is likely biased as a number of global change factors were missing, which indicates the need for modeling with multiple global environmental factors; and (3) the need for comparison with high-frequency observational data is identified as an important task for future model-model inter-comparison. These lessons will be helpful for, and likely addressed during, 95 model improvements and applications of more mechanistic CH₄ models.

Although the primary individual CH₄ processes have been studied and quantified at a certain level of confidence, only a few modeling studies have reported these individual processes. For example three pathways of CH₄ transports were represented in Kettunen, 2003 and Walter et al., 1996, but none of those modeled results have been evaluated against observational results for those individual 00 processes. One reason is that measurements rarely distinguish among individual processes; another reason is that the majority of CH₄ models do not explicitly represent all processes (Table 2). However, a number of studies report significant shifts in the processes contributing to the surface CH₄ flux along environmental gradients or across biomes (Conrad, 2009; Krumholz et al., 1995; McCalley et al., 2014). Projecting CH₄ fluxes into future changing climate conditions requires not only accurate simulations of



05 CH₄ processes, but also shifts among the various processes. In addition, CO₂ flux has been evaluated
within the Earth System Modeling framework, but only a few studies have evaluated the CH₄ flux and
its contribution to climate dynamics. Given the much higher warming potential and relatively faster rate
of increase of atmospheric CH₄, fully coupled simulations are needed to represent the feedbacks
between terrestrial CH₄ exchanges and climate. We note that a few recent studies reported a relatively
10 small climate warming-methane feedback from global wetlands and permafrost (Gao et al., 2013;
Gedney et al., 2004; Riley et al., 2011). A fully mechanistic CH₄ model that accounts for all the
important features is critically needed. In addition, a modeling framework to integrate multiple sources
of data, such as microbial community structure and functional activities, ecosystem-level
measurements, and global scale satellite measurements of gas concentration and flux is needed with
15 these mechanistic CH₄ models.

6. Needs for Mechanistic Methane Models

During the recent few years, the scientific community has continued to improve and optimize
models to better simulate methanogenesis, methanotrophy, CH₄ transport, and their environmental and
biological controls (Xu et al., 2015; Zhu. Q. et al., 2014). A number of emerging tasks have been
20 identified, and progress in these directions is expected. First, linking genomic data with large-scale CH₄
flux measurements will be an important, while challenging, task for the entire community; for example,
some work has been carried out in this direction (De Haas et al., 2011; Larsen et al., 2012). An effort
has been initialized to develop a new microbial functional group-based CH₄ model, which has the
advantages of linking genomic information for each individual process with the four microbial
25 functional groups (Xu et al., 2015). Second, data-data and model-model comparisons are another
important effort for model comparison and improvement. One ongoing encouraging feature that all
recently developed CH₄ models possess is the capability for regional simulations as well as the
possibility to be run at the site level (Riley et al., 2011; Zhu. Q. et al., 2014).

Third, microbial processes need to be considered for incorporation into ecosystem models for
30 simulating carbon cycling and CH₄ processes (DeLong et al., 2011; Xu et al., 2014). Although a few
models explicitly simulate the microbial mechanisms of CH₄ cycling (Arah and Stephen, 1998; Grant,



1998; Li, 2000a; Segers and Kengen, 1998), none of them have been used for regional- or global-scale estimation of microbial contributions to the CH₄ budget. A reasonable experimental design and a well-validated microbial functional group-based CH₄ model should be combined to enhance our capability to apply models to estimate a regional CH₄ budget and to investigate the combination of microbial and environmental contributions to the land surface CH₄ flux (DeLong et al., 2011). Fourth, incorporating well-validated CH₄ modules into Earth System Modeling frameworks will allow a fully coupled simulation that provides a holistic understanding of the CH₄ processes, with its connections to many other processes and mechanisms in the atmosphere. Several recently developed models fall in the framework of Earth System Models (Riley et al., 2011; Ringeval et al., 2010), which provide a foundation for this application in a relatively easy way. This effort will likely contribute not only to the CH₄ modeling community, but also to the entire global change science community (Koven et al., 2011). The iron and sulfate biogeochemistry that has been simulated in a few models was not included in any of the three groups because that effort will likely be achieved over the long term, owing to poor understanding of the mechanisms and the lack of observational data.

[Insert Figure 4 here]

Based on the above-mentioned needs and model features as well as the mechanisms for the CH₄ models, the next generation of CH₄ models will likely include several important features (Fig. 4). The models should (1) be embedded in an Earth System Model, (2) consider the vertical distribution of thermal, hydrological, and biogeochemical transport and processes, (3) represent mechanistic processes for microbial CH₄ production, consumption, and transport, and (4) support data assimilation and a model benchmarking system as auxiliary components.

7. Challenges for Developing Mechanistic CH₄ Models

Knowledge Gaps - Modeling CH₄ cycling is a dynamic process. As new mechanisms are identified the modeling community should ensure that the mechanisms are well studied and mathematically described, as has occurred over the past decades (Conrad, 1989; McCalley et al., 2014; Schütz et al., 1989; Xu et al., 2015). However, a number of knowledge gaps need to be filled before a full modeling framework of CH₄ processes within terrestrial ecosystems can be achieved. The first gap



is either confirmation or rejection of a few recently observed CH₄ mechanisms; these mechanisms need
60 to be fully vetted before being considered for incorporation into a model. The first most well-known
mechanism still under debate is aerobic CH₄ production within plant tissue (Beerling et al., 2008;
Keppler et al., 2006). Since its first report in 2006 (Keppler et al., 2006), a few studies have confirmed
the mechanism in multiple plant species (Wang et al., 2007). While its existence in nature is still under
debate (Dueck et al., 2007), this mechanism will likely not be incorporated into an ecosystem model
65 before solid evidence is presented and consensus is reached. The second new mechanism is fungi as a
microbial group carrying out CH₄ production (Lenhart et al., 2012). More field- or lab-based
experiments are needed to investigate this mechanism and its contribution to the global CH₄ budget,
probably through a data model integration approach. Third, the aerobic production of methane from the
cleavage of methylphosphonate has been demonstrated in marine systems (Karl et al., 2008), but the
70 significance of this process in terrestrial systems is unknown.

Another knowledge gap is the missing comprehensive understanding of spatial and temporal
variations in CH₄ flux; particularly, the “hot spots” and “hot moments” of observed CH₄ flux are still
not completely understood (Becker et al., 2008; Mastepanov et al., 2008; Song et al., 2012). The
traditional static chamber method of measuring CH₄ emissions could underestimate the CH₄ flux
75 because sparse sampling is unlikely to detect these foci or pulses of unusually high emissions. Better
methods are also needed to measure CH₄ cycling during the shoulder seasons in the Arctic and subarctic
when fluxes may be most variable (Zona et al. 2016). These knowledge gaps are key hurdles for CH₄
model development efforts. No model has yet been tested for simulating hot spots or hot moments over
large spatial or long temporal scales. However, the high range (usually of order 1-10) of these processes
80 might cause regional budgets to vary substantially (Song et al., 2012); therefore, mechanistic model
representations of these mechanisms are highly needed.

Modeling Challenges - Better simulation of CH₄ cycling in terrestrial ecosystems requires
improvement in the model structure to represent mechanistic CH₄ processes. First is the challenge for
better simulating the vertical profile of soil biogeochemical processes and validating these models with
85 observational results. Although some models have a capability for vertical distribution of carbon and



nitrogen (Koven et al., 2013; Tang et al. 2013; Mau et al., 2013), a better framework for CH₄ and extension to cover the majority of CH₄ models are needed. This vertical distribution of biogeochemistry is necessary for simulating the vertical distribution of CH₄ processes and CH₄ transport through the soil profile before reaching the atmosphere. A second challenge is incorporating tracer capability. Isotopic tracers (¹³C, ¹⁴C) have been widely used for quantifying the carbon flow and partitioning among individual CH₄ processes (Conrad, 2005; Conrad and Claus, 2005), but for ecosystem models this capability has not been represented even though it is very important to understanding CH₄ processes and integrating field observational data. A third challenge is to simulate microbial functional groups. Microbial processes are carried out by different functional groups of microbes (Lenhart et al., 2012; 95 McCalley et al., 2014). Therefore, model comparison with individual processes requires representing the microbial population sizes (or active biomass) for specific functional groups (Tveit et al., 2015). This goal has proved more difficult than representing plant functional types or traits in models, because not all microbial taxonomic groups have ecologically coherent functions (Philippot et al., 2010). A fourth challenge is to simulate the lateral transport of dissolved and particulate biogeochemical 00 variables that are necessary to better simulate the storage and transport of CH₄ within heterogeneous landscapes (Weller et al., 1995). A fifth challenge is modeling CH₄ flux across spatial scales. Although a few studies have been used to demonstrate the approach for simulating CH₄ budget at plot scale and eddy covariance domain scale (Zhang et al., 2012), a mechanistic framework to link CH₄ processes at distinct scales is still lacking while highly valuable.

05 Data Needs - First, a comprehensive dataset of field measurements of CH₄ fluxes across various landscape types is needed. Although a number of datasets have been compiled (Aronson and Helliker, 2010; Chen et al., 2012; Liu and Greaver, 2009; Mosier et al., 1997; Yvon-Durocher et al., 2014), some landscape types are still not fully covered. Meanwhile, high-frequency field observational data are also needed, particularly long-term observational data in some less-studied ecosystems. It is well-known that 10 inter-annual variation of climate may turn an ecosystem from a CH₄ sink to a CH₄ source (Nauta et al., 2015; Shoemaker et al., 2014); therefore, a long-term observational dataset that covers these shifts in CH₄ flux and its associated ecosystem information would improve our understanding of the processes



and our representation of them in CH₄ models. Second, microbial community shifts and their role in CH₄ processes are important, although information is incomplete for model representation of this mechanism (McCalley et al., 2014; Schimel and Gullledge, 1998). Although a number of studies have reported the microbial community structure and its potential association with changes in CH₄ processes (Monday et al., 2014; Schimel, 1995; Wagner et al., 2005), none of this progress has been documented in a mathematical manner suitable for a modeling representation.

Last but not least, a comprehensive dataset of all primary CH₄ processes within an individual ecosystem would be valuable for model optimization and validation. Although some datasets exist, no study has investigated all primary individual CH₄ processes within the same plot over the long term. Given the substantial spatial heterogeneity of CH₄ processes, this lack of process representation may cause bias in CH₄ simulations at regional scale. It should be noted that land surface net CH₄ flux is a measurable ecosystem-level process, whereas many individual CH₄ processes are difficult to accurately measure. Therefore, designing field- or lab-based-experiments suitable for measuring these processes is a fundamental need. For example, the anaerobic oxidation of CH₄ has been identified as a critical process for some ecosystem types, but no comprehensive dataset on it is available for model development or improvement.

Data-Model Integration - Model development and data collection are two important, but historically independent scientific approaches; the integration between model development and data collection is much stronger for advancing science (De Kauwe et al., 2014; Luo et al., 2012; Peng et al., 2011). Although data-model integration is recognized as very important for understanding and predicting CH₄ processes and some progress has been made, integrating experiments and models presents multiple challenges, particularly, 1) the methods for integrating data with the models are not well developed for CH₄ cycling; 2) the metrics for evaluating data-model integration are not consistent in the scientific community; and 3) the regular communication between data scientists and modelers on various aspects of CH₄ processes and their model representation is lacking.

Methods for data-model integration have been recently created, for example, Kalman Filter (Gao et al., 2011), Bayesian (Ogle and Barber, 2008; Ricciuto et al., 2008; Schleip et al., 2009; Van Oijen et



40 al., 2005), and Monte Carlo (Casella and Robert, 2005). However, no studies have evaluated these
methods for integrating CH₄ data with models. In addition, the metric for evaluating the data-model
integration is still not well developed. A very helpful strategy for data-model integration is to solicit
timely input from modelers when designing a field experiment. A good example of this is the U.S.
Department of Energy-sponsored project Next Generation Ecosystem Experiments - Arctic (ngee-
45 arctic.ornl.gov), which was planned with inputs from field scientists, data scientists, and modelers.
Another successful example is the U.S. DOE-sponsored project, SPRUCE (mnspruce.ornl.gov), in
which the experiment design for data-model integration created an opportunity for modeling needs to be
adopted by the field scientists.

8. Concluding Remarks

50 CH₄ dynamics in terrestrial ecosystems have been intensively studied, but model representation
of CH₄ cycling has lagged. Currently, the primary mechanisms for CH₄ processes in terrestrial
ecosystems are implicitly represented in many, but not all, ecosystem models. Development of CH₄
models began in the late 1980s, and the pace of growth has been fast since the 1990s. Model
development shifted from theoretical analysis in the 1980s and 1990s to being more applied in the
55 2000s and 2010s, expressed as being more focused on regional CH₄ budget quantification and
integration with multiple sources of observational data. Although some current CH₄ models consider
most of the relevant mechanisms, none of them consider all the processes for methanogenesis,
methanotrophy, CH₄ transport, and their primary environmental controls. Further, evidence
demonstrating that incorporating all of these processes would lead to more accurate prediction is
60 needed. Incorporating sophisticated parameter assimilation, uncertainty quantification, equifinality
quantification, and metrics of the benefits associated with increased model complexity are therefore
required.

The CH₄ models for accurate projection of land-climate feedback in the next few decades
should: (1) use mechanistic formulations for primary CH₄ processes, (2) be embedded in Earth System
65 Models for the global evaluation of terrestrial-climate feedback associated with CH₄ fluxes, (3) have the
capacity to integrate multiple sources of data, which makes the model not only a prediction tool but also



an integrative tool, and (4) be developed in association with model benchmarking frameworks. These four characteristics pave the way for examining CH₄ processes and flux in the context of global change.



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Table 1. Terrestrial ecosystem models for CH₄ cycling and the model representation of three pathways of CH₄ transport (models are in alphabetical order; author's last name is used if the model name is not available)

Model	Aerenchynma	Diffusion	Ebullition	References
Arah's model	Yes	Yes	No	(Arah and Stephen, 1998)
Beckett's model	Yes	Yes	No	(Beckett et al., 2001)
"Cartoon" model	Yes	Yes	Yes	(Arah and Kirk, 2000)
CASA	Yes	Yes	Yes	(Potter, 1997; Potter et al., 1996)
CH4MOD	Yes	Yes	Yes	(Huang et al., 1998b; Huang et al., 2004; Li et al., 2012)
Christensen's model	No	No	No	(Christensen et al., 1996)
CLASS	No	Yes	No	(Curry, 2009; Curry, 2007)
CLM4Me	Yes	Yes	Yes	(Riley et al., 2011)
CLM-Microbe	Yes	Yes	Yes	(Xu et al., 2015; Xu et al., 2014)
DAYCENT	No	Yes	No	(Del Grosso et al., 2002; Del Grosso et al., 2009; Del Grosso et al., 2000)
Ding's model	Yes	No	No	(Ding and Wang, 1996)
DLEM	Yes	Yes	Yes	(Tian et al., 2010; Xu and Tian, 2012)
DNDC	Yes	Yes	Yes	(Li, 2000b)
<i>ecosys</i>	No	Yes	Yes	(Grant, 2001, 1998)
Gong's model	Yes	Yes	Yes	(Gong et al., 2013)
HH model	Yes	Yes	Yes	(Cresto-Aleina et al., 2015)
IAP-RAS	No	No	No	(Eliseev et al., 2008; Mokhov et al., 2007)



Kettunen's model	Yes	Yes	Yes	(Kettunen, 2003)
Lovley's model	No	No	No	(Lovley and Klug, 1986)
LPJ-Bern	Yes	Yes	Yes	(Spahni et al., 2011)
LPJ-WHyMe	Yes	Yes	Yes	(Wania et al., 2010, 2009)
LPJ-WSL	No	No	No	(Hodson et al., 2011)
Martens' model	Yes	Yes	Yes	(Martens et al., 1998)
MEM	No	No	No	(Cao et al., 1995; Cao et al., 1998)
MERES	Yes	Yes	No	(Matthews et al., 2000)
Nouchi's model	Yes	Yes	No	(Hosono and Nouchi, 1997; Nouchi et al., 1994)
ORCHIDEE	Yes	Yes	Yes	(Ringeval et al., 2010; Ringeval et al., 2011)
Ridgwell's model	No	Yes	No	(Ridgwell et al., 1999)
SDGVM	No	No	No	(Hopcroft et al., 2011)
Segers' model	Yes	Yes	Yes	(Segers and Kengen, 1998; Segers and Leffelaar, 2001a, b; Segers et al., 2001)
Tagesson's model	No	No	No	(Tagesson et al., 2013)
TEM	Yes	Yes	Yes	(Zhuang et al., 2004)
TRIPLEX-GHG	Yes	Yes	Yes	(Zhu Q. et al., 2014)
UW-VIC	Yes	Yes	Yes	(Bohn and Lettenmaier, 2010; Bohn et al., 2007)
van Bodegom's model	Yes	Yes	Yes	(van Bodegom et al., 2000; Van Bodegom et al., 2001)
VISIT	Yes	Yes	Yes	(Inatomi et al., 2010; Ito and Inatomi, 2012)
De Visscher's	No	Yes	No	(De Visscher and Van



model				Cleemput, 2003)
Walter's model	Yes	Yes	Yes	(Walter and Heimann, 2000; Walter et al., 1996)
Xu's model	Yes	Yes	Yes	(Xu et al., 2007)

Table 2. Key mechanisms/features of CH₄ processes and their representations in CH₄ models

Key mechanisms	Models
Methanogenesis	Arah's model, "Cartoon" model, CASA, CH4MOD, Christensen's model, CLM4Me, CLM-Microbe, Ding's model, DLEM, DNDC, <i>ecosys</i> , Gong's model, IAP-RAS, Kettunen's model, Lovley's model, LPJ-Brn, LPJ-WHyMe, LPJ-WSL, Martens' model, MEM, MERES, ORCHIDEE, SDGVM, Segers' model, TEM, TRIPLEX-GHG, UW-VIC, van Bodegom's model, VISIT, Walter's model, Xu's model
Methanotrophy	Arah's model, "Cartoon" model, CASA, CLASS, CLM4Me, CLM-Microbe, DAYCENT, DLEM, DNDC, <i>ecosys</i> , Gong's model, Kettunen's model, LPJ-Bern, LPJ-WHyMe, Martens' model, MEM, MERES, ORCHIDEE, Ridgwells model, SDGVM, Segers' model, TEM, TRIPLEX-GHG, UW-VIC, van Bodegom's model, VISIT, De Visscher's model, Wlatter's model, Xu's model
Anaerobic oxidation of CH ₄	CLM-Microbe
Substrate (Acetate/DOC)	CH4MOD, CLM-Microbe, DLEM, DNDC, <i>ecosys</i> , Gong's model, Kettunen's model, Lovley's model, Martens' model, MEM, MERES, SDGVM, Segers' model, van Bodegom's model, Xu's model
Microbial functional groups	CLM-Microbe, DNDC, <i>ecosys</i>
CH ₄ storage in soil profile	Arah's model, Beckett's model, "Cartoon" model, CLM4Me, CLM-Microbe, <i>ecosys</i> , Kettunen's model, Martens' model, MERES, Nouchi's model, ORCHIDEE, Segers' model, UW-VIC, van Bodegom's model, VISIT, De Visscher's model, Walter's model
O ₂ availability for CH ₄ oxidation	Arah's model, Beckett's model, "Cartoon" model, CLM4Me, CLM-Microbe, <i>ecosys</i> , Kettunen's model, MERES, Segers' model, van Bodegom's model, De Visscher's model, Xu's model
Iron biogeochemistry	van Bodegom's model
Sulfate biogeochemistry	Lovley's model, Martens' model, van Bodegom's model
Frozen trapped CH ₄	None
Embedded in Earth System Model	CLM4Me, CLM-Microbe, IAP-RAS, ORCHIDEE, SDGVM



Vertical resolved biogeochemistry	Arah's model, Beckett's model, "Cartoon" model, CLASS, CLM4Me, CLM-Microbe, DNDC, <i>ecosys</i> , Gong's model, HH model, IAP-RAS, Kettunen's model, Lovley's model, LPJ-Bern, LPJ-WHyMe, LPJ-WSL, Martens' model, MERES, ORCHIDEE, Ridgwell's model, SDGVM, Segers' model, TRIPLEX-GHG, UW-VIC, VISIT, De Visscher's model, Walter's model, Xu's model
Regional-scale, capacity for up-scaling	CASA, CH4MOD, Christensen's model, CLASS, CLM4Me, CLM-Microbe, DAYCENT, DLEM, <i>ecosys</i> , Gong's model, HH model, IAP-RAS, LPJ-Bern, LPJ-WHyMe, LPJ-WSL, Martens' model, MEM, MERES, ORCHIDEE, Ridgwell's model, SDGVM, Tagesson's model, TEM, TRIPLEX-GHG, UW-VIC, VISIT, Walter's model



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Table 3. The mathematical equations used to described the CH₄ processes used in representative models (P_{CH_4} is the CH₄ production rate; $Oxid_{CH_4}$ is the CH₄ oxidation rate; T_{CH_4} is the CH₄ transport rate; D_{CH_4} is the CH₄ diffusion rate; some parameter may have been changed from original publication to keep relatively consistent in this table)

CH ₄ processes		Equations	Ecological description	Model examples
CH ₄ substrate and CH ₄ production	1	$P_{CH_4} = f(T, W)$	A function of temperature (T) and moisture (W)	Christensen's model, IAP-RAS, DAYCENT
	2a	$P_{CH_4} = r \times HR \times f(T, W)$	A portion of heterotrophic respiration, affected by temperature (T) and moisture (W)	LPJ family, CLM4Me, Ding's model, MERES, TRIPLEX-GHG
	2b	$P_{CH_4} = r \times SOM \times f(T, W)$	A portion of soil organic matter (SOM), affected by temperature (T) and moisture (W); Walter's model use indirect association with NPP	CH4MOD, Gong's model, HH model, Walter's model
	3	$P_{CH_4} = V \times \frac{[DOC]}{K_{DOC} + [DOC]} \times f(T, W)$	A portion of dissolved organic carbon (DOC), affected by temperature (T) and moisture (W)	MEM, DLEM
	4	$P_{CH_4} = f(DOC, Acetate, CO_2) \times f(T, W)$	Mechanistic processes for CH ₄ production are considered, affected by temperature (T) and moisture (W)	Kettunen's model, Segers' model, van Bodegoms model, and <i>ecosys</i>
CH ₄ oxidation	5	$Oxid_{CH_4} = V \times \left(\frac{[CH_4]}{K_{CH_4} + [CH_4]} \right) \times f(T, W)$	Oxidation as a function of CH ₄ concentration and temperature and moisture	DLEM, TRIPLEX-GHG, VISIT



	6	$Oxid_{CH_4} = V \times \left(\frac{[CH_4]}{K_{CH_4} + [CH_4]} \right) \left(\frac{[O_2]}{K_{O_2} + [O_2]} \right) \times f(T, W)$	Oxidation as a function of CH ₄ and O ₂ concentration, temperature and moisture	Arah's model, Cartoon model, CLM4Me, CLM-Microbe, Kettunen's model
CH ₄ transport	7	$T_{CH_4} = V * ([CH_4] - \overline{[CH_4]})$	V is the parameter for distance, diffusion coefficient, etc.; [CH ₄] is the concentration of CH ₄ in the soil/water profile (dissolvability for DLEM, 0 for DNDC); and $\overline{[CH_4]}$ is the threshold of CH ₄ concentration above which CH ₄ will be transported to the atmosphere via either of the three transport pathways	DLEM, DNDC, Walter's model
	8a	$A = \frac{C(z) - C_a}{r_L z / D + r_a} p T \rho_r$	<i>Aerenchyma transport</i>	CLM4Me
	8b	Moves to first unsaturated layer and then released to gaseous phase	<i>Ebullition</i>	CLM4Me
	8c	$D_{CH_4} = D \times \frac{\Delta[CH_4]}{\Delta z}$	Diffusion of CH ₄ was simulated following Fick's law; CLM4Me separate aqueous and gaseous diffusion	CLM4Me, CLM-Microbe, <i>ecosys</i> , Ridgwell's model, TRIPLEX-GHG; Sergers' model
Temperature effects	9	$f(T) = a \times T + b$ $f(T) = a \times T^2 + b \times T + c$ $f(T) = b \times e^{0.2424 \times T}$	Linear regression on temperature or degree days; DNDC simulate temperature impact on production not on oxidation	DAYCENT, DNDC, IAP-RAS, LPJ family



	10	$f(T) = Q_{10}^{\frac{(T-T_{ref})}{10}}$	Q_{10} equations; T_{ref} is the reference temperature	CH4MOD, CLM-Microbe, CLM4Me, DLEM, VISIT, Kettunen's model
	11a	$V_T = V^0 \times \exp\left(\frac{\Delta E}{R} \left[\frac{1}{T^0} - \frac{1}{T}\right]\right)$	Arrhenius equation	Arah's model, Ding's model
	11b	$f_T = \frac{T_s \times \exp\left(A - \frac{H_a}{R \times T_s}\right)}{\left[1 + \exp\left(\frac{H_{dl} - S \times T_s}{R \times T_s}\right) + \exp\left(\frac{H_{dh} - S \times T_s}{R \times T_s}\right)\right]}$	Modified Arrhenius equation; T_s is soil temperature at K; A is the parameter for $f_T = 1.0$ at $T_s = 303.16$ K; H_a is the energy of activation (J mol^{-1}); R is universal gas constant ($\text{J mol}^{-1} \text{K}^{-1}$); H_{dl} and H_{dh} are energy of low and high temperature deactivation (J mol^{-1})	<i>ecosys</i>
Moisture effects on methanogenesis and methanotrophy	12	No moisture effect is simulated, rather inundation area is simulated	No equation, while a temporal and spatial variation of inundation and saturation impacts	CASA
	13	$F_\theta = e^{-P/P_c}$	Water stress for oxidation, where P is soil moisture and $P_c = -2.4 \times 10^5$ mm	CLM4Me
	14	$f(SM) = \begin{cases} 1, \\ \left[1 - \frac{\log_{10}\phi - \log_{10}(0.2)}{\log_{10}(100) - \log_{10}(0.2)}\right]^\beta \\ 0, \end{cases}$	β is an arbitrary constant, ϕ is the soil water potential	CLASS



	15	$f_{prod}(SM) = \left(\frac{SM - SM_{fc}}{SM_{sat} - SM_{fc}} \right)^2 \times 0.368 \times e^{\frac{SM - SM_{fc}}{SM_{sat} - SM_{fc}}}$ $f_{oxid}(SM) = 1 - f_{prod}(SM)$	Different impacts on CH ₄ production and consumption; SM: soil moisture; SM _{fc} : field capacity; SM _{sat} : saturation soil moisture	DLEM
	16	$f(SM) = \frac{(M_V - M_{min}) \times (M - M_{min})}{(M_V - M_{min}) \times (M_V - M_{max}) - (M - M_{min}) \times (M - M_{max})}$	Bell-shape curve	TEM
pH effects	17a	$f(pH) = \frac{(pH - pH_{min}) \times (pH - pH_{max})}{(pH - pH_{min}) \times (pH - pH_{max}) - (pH - pH_{min}) \times (pH - pH_{max})}$	Bell-shape curve	CLM-Microbe, MEM, TEM,
	17b	$f(pH) = 10^{-0.2335 \times pH^2 + 2.7727 \times pH - 8.6}$	Bell-shape curve	CLM4Me
	17c	$f(pH) = \begin{cases} 0 & pH \leq 4 \\ \frac{1.02}{1 + 1000000 \times e^{(-2.5 \times pH)}} & 4 < pH < 7 \\ \frac{1.02}{1 + 1000000 \times e^{(-2.5 \times (14 - pH))}} & pH \geq 7 \end{cases}$	Bell-shape curve	DLEM



Table 4. Temperature dependence of CH₄ processes in various models (blank indicates the Q₁₀ function is not used; all temperatures are expressed as °C, 273.15 was used for unit conversion)

Model	Q ₁₀	Reference temperature (°C)	Note	Sources
CASA			Based on a linear equation with temperature	(Potter, 1997)
DAYCENT			Linear equation $y = 0.209 * T + 0.845$	(Del Grosso et al., 2000)
LPJ family LPJ-Bern LPJ-WHyMe LPJ-WSL			Linear function was used for temperature impacts on diffusion	(Hodson et al., 2011; Spahni et al., 2011; Wania, 2007)
Christensen's model	2	2	For temperature > 0, the temperature impact is set to zero when < 0	(Christensen and Cox, 1995)
CH4MOD	3	30	T=30 for 30 < T ≤ 40	(Huang et al., 1998b)
CLM4Me	2	2	Parameters for baseline simulation	(Riley et al., 2011)
CLM-Microbe	1.5	13.5		(Xu et al., 2015)
DLEM	2.5	30	For a temperature range of [-5, 30]; temperature impact is set to zero when < -5 or > 30	(Tian et al., 2010)
Kettunen's model	4.0 for production, 2.0 for oxidation	10	Standard Q ₁₀ function	(Kettunen, 2003)
ORCHIDEE	Abisko site, 2.6; Michigan site, 3.2; Panama site, 1.2	Mean annual temperature	Q ₁₀ function with different parameters across biomes	(Ringer et al., 2010)



TEM	Alpine tundra: wetland, 3.5; upland, 0.8. Wet tundra: wetland, 2.2; upland, 1.1. Boreal forest: wetland, 1.9; upland, 1.5	Alpine tundra: wetland, -3.0; upland, 8.0. Wet tundra: wetland, -5.5; upland, 8.0. Boreal forest: wetland, 1.0; upland, 7.0	Q ₁₀ function with different parameters across biomes	(Zhuang et al., 2004)
TRIPLEX-GHG	1.7-16 for production, 1.4-2.4 for oxidation	25 for optimal, 45 for highest temperature	Modified Q ₁₀ equation	(Zhu et al., 2014a)
VISIT		Mean annual temperature		(Ito and Inatomi, 2012)
Walter's model	2	Ombrotrophic bog, 12; poor fen, 6.5; oligotrophic pine fen, 3.5; Arctic tundra, 0; swamp, 27	Q ₁₀ function with different parameters across biomes	(Walter and Heimann, 2000)
Arah's model		10	Arrhenius equation	(Arah and Stephen, 1998)
<i>ecosys</i>		30	Modified Arrhenius equation	(Grant et al., 1993)



Figure legend

Figure 1. The published CH₄ models and modeling trends in terms of applicability and mechanistic representation of CH₄ cycling processes at decadal-scale and the envisioned CH₄ model capability

10 Figure 2. Percentage of CH₄ models with consideration of some key CH₄ mechanisms. The percentage was calculated as the number of models considering each mechanisms divided by the total number of published models in each time period.

15 Figure 3. Three types of models with key mechanisms for CH₄ production and oxidation (*SOM*: Soil organic matter; *NPP*: net primary production; *DOC*: dissolved organic carbon; *O_{atm}*: oxidation of atmospheric CH₄; *P*: plant-mediated transport; *D*: diffusion transport; *E*: ebullition transport; *O_{xid}*: oxidation; *O_{trans}*: oxidation of CH₄ during transport)

Figure 4. Key features of future mechanistic CH₄ models with a full representation of primary CH₄ processes in the terrestrial ecosystems. The data assimilation system and model benchmarking system are also shown as auxiliary components to the future CH₄ models.

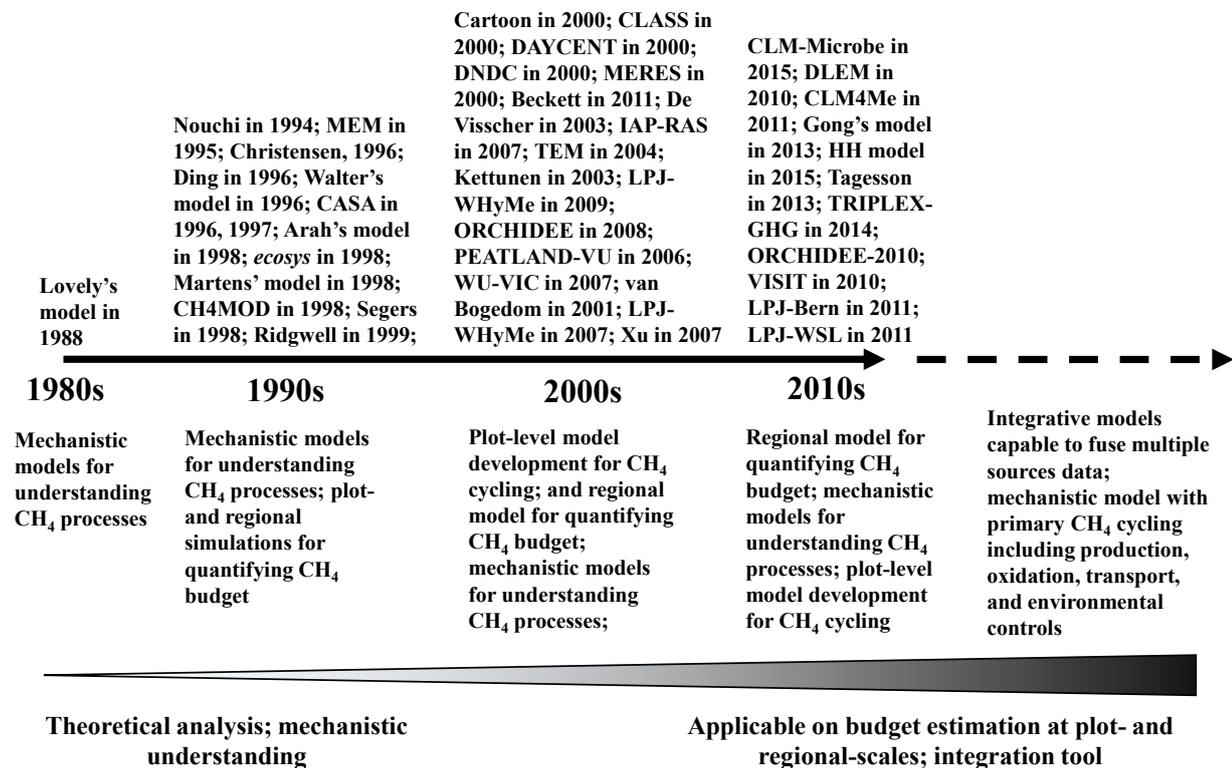


Fig. 1.

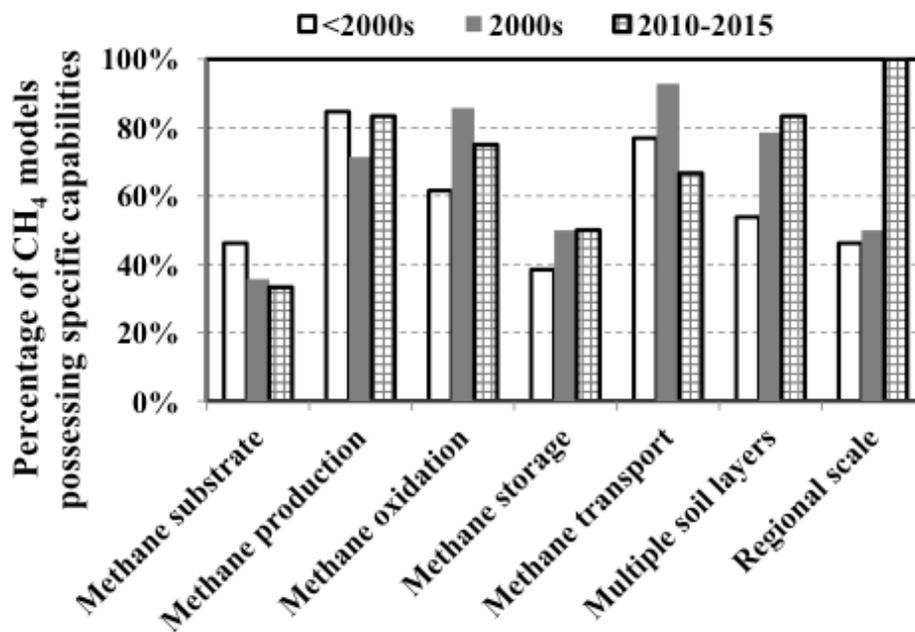


Fig. 2.

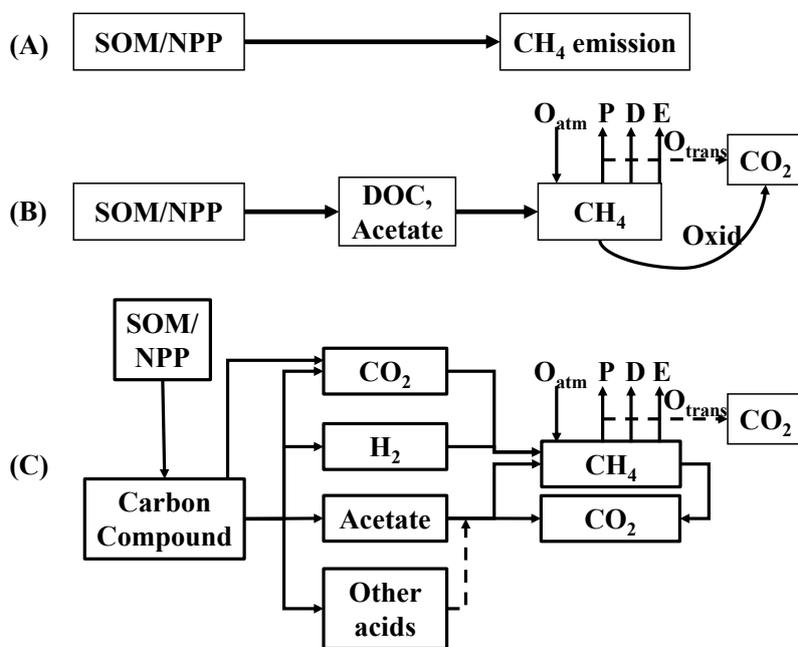


Fig. 3.

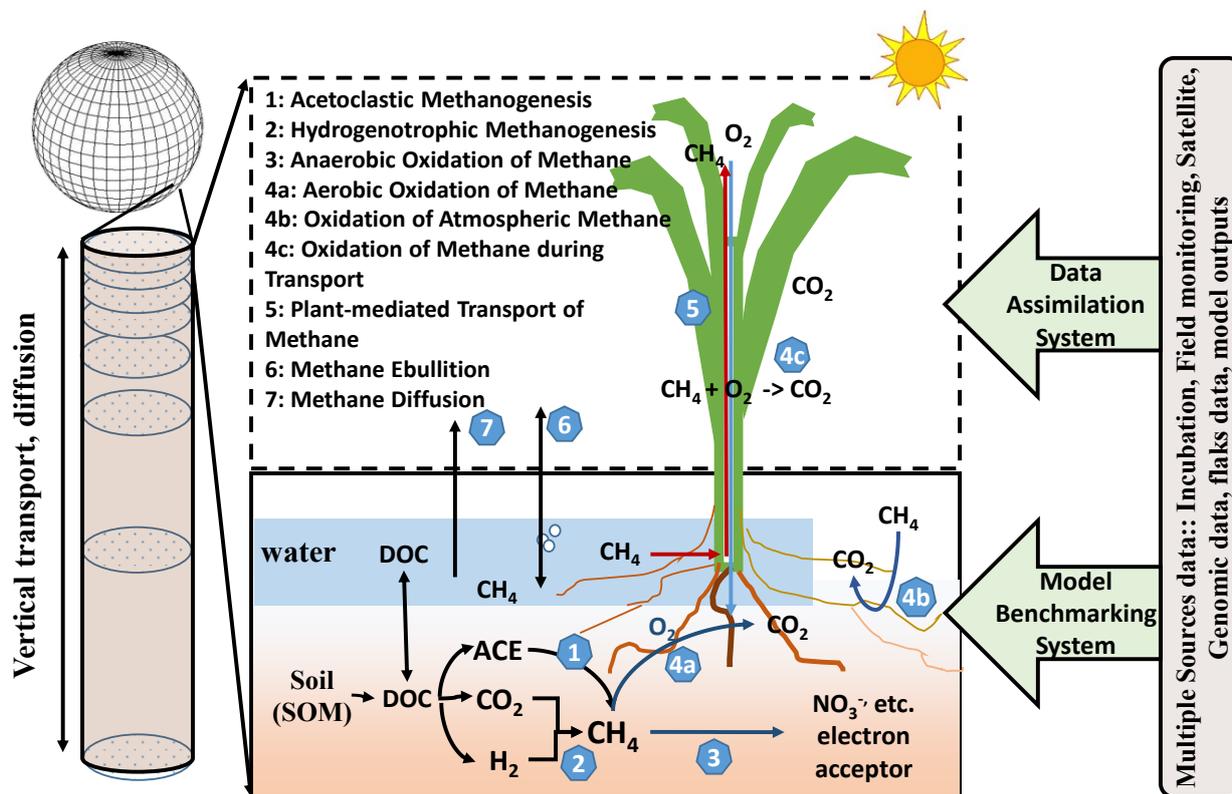


Fig. 4