Implementation of methane cycling for deep time, global warming simulations with the DCESS Earth System Model (Version 1.2)

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Abstract. Geological records reveal a number of ancient, large and rapid negative excursions of carbon-13 isotope. Such excursions can only be explained by massive injections of depleted carbon to the Earth System over a short duration. These injections may have forced strong global warming events, sometimes accompanied by mass extinctions, for example the Triassic-Jurassic and End-Permian extinctions, 201 and 252 million years ago. In many cases evidence points to methane as the dominant form of injected carbon, whether as thermogenic methane, formed by magma intrusions through overlying carbon-rich sediment, or from warming-induced dissociation of methane hydrate, a solid compound of methane and water found in ocean sediments. As a consequence of the ubiquity and importance of methane in major Earth events, Earth System models for addressing such events should include a comprehensive treatment of methane cycling but such a treatment has often been lacking. Here we implement methane cycling in the Danish Center for Earth System Science (DCESS) model, a simplified but well-tested Earth System Model of Intermediate Complexity. We use a generic methane input function that allows variation of input type, size, time scale and ocean-atmosphere partition. To be able to treat such massive inputs more correctly, we extend the model to deal with ocean suboxic/anoxic conditions and with radiative forcing and methane lifetimes appropriate for high atmospheric methane concentrations. With this new model version, we carried out an extensive set of simulations for methane inputs of various sizes, time scales and ocean-atmosphere partitions to probe model behaviour. We find that larger methane inputs over shorter time scales with more methane dissolving in the ocean lead to ever-increasing ocean anoxia with consequences for ocean life and global carbon cycling. Greater methane input directly to the atmosphere leads to more warming and, for example, greater carbon dioxide release from land soils. Analysis of synthetic sediment cores from the simulations provides guidelines for the interpretation of real sediment cores spanning the warming events. With this improved DCESS model version and paleo-reconstructions, we are now better armed to gauge the amounts, types, time scales and locations of methane injections driving specific, observed deep time, global warming events.

1 Introduction

Analyses of carbonate and organic carbon samples have revealed, with ever-finer time resolution, a number of large and rapid negative carbon isotope excursions since the Cambrian radiation of life about 540 million years ago.. These isotope excursions had typical δ^{13}C amplitudes of > 2 ‰ and typical time scales of < 10000 years. Some prominent examples of this have been found at about 56, 120, 183, 201, 252 and 260 million years ago (Hesslebo et al., 2002; Jenkyns, 2003; McElwain et al., 2005; Retallack & Jahren, 2008; Ruhl et al., 2011; Shen et al., 2011; Wignall et al., 2009; Zachos et al. 2001). Such large, rapid carbon isotope excursions can only be explained by large, rapid injections of light carbon (i.e. carbon-13 depleted) to the ocean-atmosphere-land biosphere system, injections that forced global warming events for each of these excursions. As a starting point the source of this carbon could be some combination of volcanism (δ^{13}C ~ -7 ‰), terrestrial biosphere reduction (δ^{13}C ~ -25 ‰), thermogenic methane input (δ^{13}C ~ -40 ‰) and methane hydrate release (δ^{13}C ~ -60 ‰), but the incorporation of further constraints often allows us to rule out the first two options as important contributors to most if not all of these events.

For example, if we assume an initial carbon pool in the ocean-atmosphere-land biosphere of about 40,000 GtC (1 Gigaton of carbon (GtC) is 10^{15}gC) with a mean δ^{13}C of 1 ‰ as at present, mass balance calculations then show that a large carbon isotope excursion of -5‰, like for example that of the end-Permian event (Shen et. al., 2011), could be explained by injections of ~ 44,400, 8,900, 5,300 and 3,400 GtC of volcanic carbon, biosphere carbon, thermogenic methane or methane hydrate, respectively. The volcanic carbon explanation seems unlikely due to the required size and speed of the input. An explanation in terms of terrestrial biosphere die off seems unlikely since it would require a biomass at least 4 times greater than the present day biomass of around 2000 GtC (Shaffer et al., 2008). Permafrost and peat carbon are other possible sources of terrestrial carbon. Little permafrost could exist for the warm conditions leading up to the warming events considered here (Shaffer et al, 2016). An explanation in terms of global peat conflagration (Kurtz et al., 2003) or degradation seems unlikely since it would require peat carbon reserves almost 20 times greater than present reserves of around 500 GtC (Brigham et al., 2006). Therefore the most likely explanations for the carbon isotope excursions and associated global warming events involve thermogenic methane and/or methane hydrate inputs (Berner, 2002; Hesselbo et al., 2000; McElwain et al., 2005; Retallack & Jahren, 2008; Ruhl et al., 2011; Svensen, H. et al., 2004; Shaffer et al., 2016).

Thermogenic methane and carbon dioxide are produced when magma intrudes into overlying sediments containing old organic carbon. More such intrusions would be expected in association with increased volcanism that creates large igneous provinces and such provinces tend to correlate in time with the ancient carbon isotope excursions (Svensen et al., 2004). Magma contact with shale and coal favor production of CH_4 and CO_2, respectively, whereas such contact with (isotopically-heavy) limestone produces very limited amounts of CH_4 and CO_2 (Arnes et al., 2011). Shale is typically the main host rock through which magma intrudes in large igneous provinces (Svensen et al. 2004; Svensen et al., 2007). Methane hydrate is a

solid compound of methane and water formed for sufficiently high pressure and low temperature and found mainly in ocean sediments. Within a hydrate stability zone, methane hydrate is formed when methane release from bacterial remineralisation of organic matter exceeds that needed to sustain solubility levels in the face of vertical diffusive transport of methane. Warming will lead to methane hydrate destabilization and thereby feedback positively on the warming (Dickens et al., 1995).

5   Methane hydrate dissociation was touted early on in a number of papers as an explanation for some deep-time carbon isotope excursions (e.g. Hesselbo et al., 2000; Berner, 2002; Kemp et al., 2005).

The discussion above underlines the need to consider input and ocean-atmosphere cycling of methane in modelling of such deep time global warming events, yet in most cases models used to study such events have not included a full prognostic and

10   interactively coupled approach incorporating methane into the climate as well as the biogeochemical cycles and isotope components. In addition to the question of the source of the light carbon inputs that force the events, other important aspects for understanding the workings of these events include the amounts, timings and locations of these inputs. For example, over what time scales is this carbon injected and how are the inputs partitioned between the ocean and the atmosphere? In order to address these questions, paleodata reconstructions should be compared with event simulations using Earth System Models

15   that consider methane input and cycling. For such a task, important aspects should be addressed like bacterial oxidation of methane in the ocean, air-sea gas exchange of methane, the dependency of atmospheric methane lifetimes on methane concentration there and the extreme radiative forcing for high atmospheric concentrations of methane. However, to our knowledge, such an Earth System Model does not exist at present. Here we upgrade the Danish Center for Earth System Science (DCESS) Earth System Model to fill this gap by designing appropriate methane input functions and by

20   implementing ocean-atmosphere-biosphere cycling of methane. Since large, rapid inputs of methane to the ocean and oxidation of this methane there can lead to suboxic or anoxic oceanic conditions, we also upgrade our model to deal with ocean denitrification, nitrogen fixation and sulphate reduction (Shaffer 1989). Finally we present some test simulations of deep time, global warming events using this extended model to illustrate ways forward for future model-data analyses.

2 The DCESS model up to now

25   In its present configuration, the DCESS Earth System Model (Shaffer et al., 2008) features modules for the atmosphere, ocean, ocean sediment, land biosphere and lithosphere (Fig. 1). The model has been designed to simulate global change on time scales of years to millions of years. It has been calibrated and tested against Earth System data and against output from more complex models. Model formulation is described in great detail in Shaffer et al., (2008); below follows a condensed description for present purposes.

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The DCESS model consists of one hemisphere, divided by 52° latitude into a low-mid latitude and a high latitude zone (Fig. 2). Sea ice and snow cover are diagnosed from estimated meridional profiles of atmospheric temperature (see below).

Values for global reservoirs, transports and fluxes are obtained by doubling the hemispheric values. For the atmosphere module we use a simple, zonal mean, energy balance model for the near surface atmospheric temperature, forced by yearly-mean insolation, outgoing longwave radiation, meridional sensible and latent heat transports and air-sea heat exchange (Fig. 3). In combination with the simple sea ice and snow parameterizations, the model includes the ice/snow-albedo feedback and the insulating effect of sea-ice. Prognostic equations for mean atmospheric temperature in the each zone are obtained by integrating the energy balance over the zones. Meridional variations of atmospheric temperature are represented by a second order Legendre polynomial in sine of latitude, adjusted to fit zonal mean temperatures. Temperatures and gradients at the 52° dividing latitude from this meridional profile are used in the calculation of meridional transports of sensible heat, latent heat and water vapour.

The atmosphere exchanges heat and gases with the ice-free part of the ocean (Fig. 3). Gases are taken to be well-mixed in the atmosphere and those considered are carbon dioxide and methane for all three carbon isotopes, nitrous oxide and oxygen. Methane is oxidized to carbon dioxide on time scales on the order of ten years, increasing as methane concentrations increase (see below). In this original model, air-sea exchange of methane is not considered. The model also deals with oxygen isotopes in atmospheric water vapor. Prognostic equations for atmospheric gases follow from internal sinks, exchanges with ocean surface layers and the land biosphere as well as input/outputs from weathering and lithosphere outgassing.

The model ocean is 270° wide, extends from the equator to 70° latitude and covers 70.5 % of the Earth surface (Fig. 2). Both the low-mid and high latitude ocean sectors have 55 vertical layers with 100 m resolution. An ocean sediment sector is assigned to each of the ocean layers with widths representing modern day hypsography (Fig. 3). Ocean circulation and mixing is characterized by four parameters: 1. a transport, q, associated with high latitude sinking and the deepest, low-mid latitude upwelling, 2. constant horizontal diffusion between the zones, K_h, associated with the wind-driven circulation and deep recirculation, 3. strong, constant vertical diffusion in the high latitude zone, K_v^{HL}, associated with high latitude convection and, 4. a weak, depth-dependent, vertical diffusion in the low-mid latitude zone, K_v^{LL}. . Tracers considered are temperature, salinity, oxygen isotope in water, phosphate, dissolved oxygen, dissolved inorganic carbon for all three carbon isotopes, and alkalinity. In this original model, ocean methane is not considered. Biogenic production of particulate organic matter depends on surface layer phosphate availability but with lower efficiency in the high latitude zone. The calcite to organic carbon rain ratio depends on surface layer temperature. Remineralisation and dissolution of particles falling down out of the ocean surface are described by exponential functions with depth. Prognostic equations for each of the tracers in each of the ocean layers follow from ocean transport and mixing, internal sinks and sources, and exchanges with the ocean sediment. Ocean surface layers are also acted upon by exchanges with the atmosphere and river inputs associated with weathering.

The semi-analytical, ocean sediment module considers calcium carbonate dissolution and both oxic and anoxic organic matter remineralisation. The sediment is composed of calcite, non-calcite mineral and reactive organic matter. Sediment porosity profiles are related to sediment composition and a bioturbated layer 0.1 m thick is applied. Carbonate and organic carbon burial are calculated from sedimentation velocities at the base of the bioturbated layer. Bioturbation rates and oxic and anoxic remineralisation rates depend on organic carbon rain rates and dissolved oxygen concentrations. The land biosphere module considers leaves, wood, litter and soil. Net primary production depends on atmospheric carbon dioxide concentration and remineralisation rates in the litter and soil depend on mean atmospheric temperatures. Methane production is a small, fixed fraction of the soil remineralisation. The lithosphere module considers outgassing, weathering of carbonate and silicate rocks and weathering of rocks with old organic carbon and phosphorus. Weathering rates are related to mean atmospheric temperatures.

Up to now the DCESS model has been applied mainly to long term, future projections of climate and Earth System response to anthropogenic forcing (Shaffer et al., 2009; Shaffer, 2009; Shaffer, 2010). Model results compare favorably with those of the best other Earth System Models of Intermediate Complexity as seen in recent intercomparison studies of past, present and future carbon cycling and climate (Eby et al., 2013; Joos et al. 2013; Zickfeld et al. 2013). The results from these studies were also included in the recent Fifth Assessment Report of the Intergovernmental Panel for Climate Change (IPCC, 2013). From the 2013 studies onward, the carbon fertilization parameter of the terrestrial biosphere module was decreased somewhat in the model (Eby et al., 2013).

The Paleocene-Eocene Thermal Maximum (PETM) warming event 56 million years ago was addressed in the most recent model application (Shaffer et al., 2016) whereby the very simplified methane cycling sketched above was implemented and the carbon input was taken to occur over a ten thousand year timescale. That is significantly longer than time scales associated with methane oxidation in the atmosphere or ocean overturning and the system response is rather insensitive to input type and location (ocean or atmosphere). Therefore, for simplicity the carbon input was taken to be in the form of carbon dioxide and was injected into the atmosphere. To deal with this deep time warming event, several other modifications were made to the basic DCESS model (Shaffer et al., 2016): 1. The solar constant was decreased by 0.5 % from present day, sea level was raised by 100 m while retaining the modern day hypsography, and mean ocean salinity was taken to be 33.8 (the modern value is 34.7), 2. Background albedo was reduced due to less land at subtropical latitudes and more vegetation cover, 3. Initial land biomass was increased to reflect more land at low and mid-latitudes, 4. To account for high latitude cloud forcing, extra forcing was imposed poleward of the latitude of sea ice extent in the standard model, pre-industrial steady state, 5. Parameters in the simple Budyko-type formulation for outgoing longwave radiation (Budyko 1969) were adjusted in concert with the other forcing changes to achieve late Paleocene temperatures and climate sensitivity estimates, 6. Ocean calcium and magnesium concentrations were adjusted to late Paleocene values, 7. Model weathering formulation was extended to include the size of the weathering substrate, estimated for the late Paleocene to be less than that for the

present, 8. Lithosphere outgassing was increased to conform to late Paleocene estimates, 9. The formulation of the calcite to organic carbon rain ratio was extended to also depend on the calcite saturation state of the ocean surface layer and, 10. The dependence of carbon isotope discrimination during ocean surface layer photosynthesis was extended to depend on concentrations of dissolved carbon dioxide and phosphate there.

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3 Methane cycling implementation

3.1 Atmosphere

For an oxygenated atmosphere the dominant sink for atmospheric methane is oxidation to CO_2 by reaction with the OH radical. Since this reaction depletes the concentration of these radicals, methane atmospheric lifetime, τ, grows as methane

10 concentrations increase. In the original model (Shaffer et al., 2008), this effect and that of associated chemical reactions in the troposphere and stratosphere was addressed by fitting results from a complex atmospheric chemistry model (Schmidt and Shindell, 2003) to the equation

$$\tau = \tau_{PI}(M+b)/\left[(1-a)M+b\right] \tag{1}$$

15 where τ_{PI} is the pre-industrial lifetime, a and b are fitting constants and $M \equiv (pCH_4 - pCH_{4,PI})/pCH_{4,PI}$ where pCH_4 is the atmospheric partial pressure of methane and $pCH_{4,PI}$ is its pre-industrial value taken to be 720 ppb. The Schmidt and Shindell (2003) target results and the model fit to them used in Shaffer et al., (2008) are shown in Fig. 4 (red dots and blue line). For this fit $\tau_{PI} = 6.9$ years, $a = 0.96$ and $b = 6.6$.

20 Fig. 4 also shows results from several additional chemistry modeling studies for the dependency of τ on pCH_4 (blue dots from Lamarque et al. (2006); black dots from Isaksen et al. (2011)). For the range of low to medium methane concentrations considered in Schmidt and Shindell (2003) and Isaksen et al. (2011), these authors found rather low sensitivity of their results to atmospheric water vapor levels citing methane lifetime changes of less than 10% for tropospheric water vapor increases of $30-40\%$. On the other hand, Lamarque et al. (2006) found that atmospheric lifetimes decrease for extreme

25 values of pCH_4 due to much enhanced water vapor and OH production (and thereby greater methane oxidation) of a very warm climate. The red line in Fig. 2 shows a fit to all the chemistry modeling results of the figure using Eq. 1, a fit for which $\tau_{PI} = 9.5$ years, $a = 0.78$ and $b = 11$. This new fit now also captures in part enhanced OH production and associated limitation of τ for very high values of pCH_4 and has been adopted in our enhanced model whereby the model atmospheric methane sink in moles per year is $\upsilon_a pCH_4/\tau$ with υ_a as the atmospheric mole content.

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The standard radiative forcing formulation for methane (including spectral overlap with nitrous oxide) can be used for

methane concentrations up to about 5 ppm (Myhre et al, 1998), much less than the concentrations we deal with here. However, recent work has provided methane radiative forcing formulations, including nitrous oxide overlap, for methane concentrations up to 100 ppm (Byrne and Goldblatt, 2014a) and even up to 10,000 ppm (Byrne and Goldblatt, 2014b). We adopt these results and, for consistency, we also adopt the Byrne and Goldblatt (2014a) expressions for carbon dioxide radiative forcing and for carbon dioxide-nitrous oxide overlap. Figure 5 serves to illustrate these radiative forcings. Note that for very high methane concentrations, radiative forcing levels off and even decreases, an effect caused by enhanced short wave absorption at such concentrations (Byrne and Goldblatt, 2014b). Taken together with the leveling off of methane atmospheric lifetimes at such concentrations (Fig. 4), this limits the role that can be played by methane in enhancing global warming under such extreme conditions.

To illustrate the effects of these new radiative forcing formulations we consider several specific atmosphere compositions and warming associated with them. If we consider an atmosphere with $p\mathrm{CO_2}$, $p\mathrm{CH_4}$ and $p\mathrm{N_2O}$ values of 500, 10 and 0.5 ppm, respectively, this corresponds to radiative forcings relative to pre-industrial conditions of 3.26, 2.32 and 0.73 W/m^2 but with a total forcing overlap of -0.16 W/m^2, leading to a total forcing of 6.15 W/m^2. With the original radiative forcing formulations used in the DCESS model (Myhre et al., 1998; Shaffer et al., 2008) the total radiative forcing is similar, 6.25 W/m^2. For a nominal climate sensitivity of 3°C for a $p\mathrm{CO_2}$ doubling (0.81°C/W/m^2), such total forcings would lead to global mean temperature increases of 4.98 and 5.06 °C, respectively, above the DCESS pre-industrial mean global temperature of 15°C. If we consider an atmosphere with $p\mathrm{CO_2}$, $p\mathrm{CH_4}$ and $p\mathrm{N_2O}$ values of 1000, 100 and 1 ppm, respectively, this corresponds to radiative forcings relative to pre-industrial conditions of 7.45, 6.50 and 1.88 W/m^2 but with a total forcing overlap of -0.93 W/m^2, leading to a total forcing of 14.97 W/m^2. With the original radiative forcing formulations, the total radiative forcing is considerably higher, 18.36 W/m^2. For a 3°C climate sensitivity, such total forcing would lead to global mean temperature increases of 12.13 and 14.87 °C, respectively; for a 5°C climate sensitivity (Shaffer et al, 2016) the increases would be 20.22 and 24.79 °C, respectively.

3.2 Ocean

3.2.1 Simplified nitrogen and sulphur cycling

To be able to deal with suboxic and anoxic ocean conditions that would arise for significant oxidation of methane in the ocean interior, the model has been generalized to consider nitrogen and sulphur cycling with the introduction of nitrate ($\mathrm{NO_3}$), ammonium ($\mathrm{NH_4}$) and hydrogen sulfide ($\mathrm{H_2S}$) as additional ocean tracers (Shaffer, 1989; for simplicity, we often omit ion charges here). Furthermore for simplicity we take the river input of nitrogen to be in the form of nitrate and to be equal to the (climate- and weathering substrate-dependent) river input of phosphate multiplied by r_{NP}, the N:P Redfield

(mole) ratio, taken to be 16. Likewise, organic matter produced, remineralised or buried maintains this ratio of nitrogen to phosphorus. Following Shaffer et al. (2008), we neglect dissolved organic matter such that the rate of export of particulate organic matter down out of the surface layer is equal to new production and we assume an exponential law for the vertical distribution of remineralisation of "nutrient" and "carbon" components of this organic matter, each with a distinct e-folding length, λ_N and λ_C, respectively.

Model new production of organic matter in the surface layer has been generalized to depend on the limiting nutrient, either phosphate or nitrate, such that $NP^{l,h} = \min(NPP^{l,h}, NPN^{l,h})$ where

$$NPP^{l,h}, NPN^{l,h} = A_0^{l,hni} z_{eu}(1, r_{NP})^{-1}(L_f^{l,h}/sy)(PO_4^{l,h}, NO_3^{l,h})\{[PO_4^{l,h}/(PO_4^{l,h}+P_{1/2})][NO_3^{l,h}/(NO_3^{l,h}+N_{1/2})]\} \qquad (2)$$
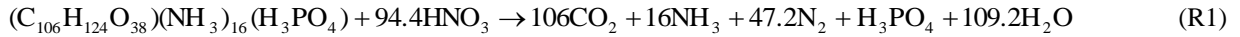
and l,h refer to low-mid latitude and high latitude ocean zones, $A_0^{l,hni}$ are the ice-free ocean surface areas, z_{eu} is the surface layer depth (100 m), r_{NP} is a Redfield ratio (taken to be 16), sy is the number of seconds per year, $PO_4^{l,h}$ and $NO_3^{l,h}$ are the surface layer, phosphate and nitrate concentrations, and $P_{1/2}, N_{1/2}$ are half saturation constants (1 μmol/m^3, 16 μmol/m^3). $L_f^{l,h}$ are efficiency factors, taken to be 1 and 0.36 for the low-mid and high latitude zones, respectively (Shaffer et al., 2008). This is how the model accounts for light and iron limitation in the high latitude zone.

The surface layer sinks due to new production are $-NP^{l,h}$ for phosphate and $-r_{NP}NP^{l,h}$ for nitrate. Other source/sinks and Redfield ratios are adopted from Shaffer et al. (2008). Since we will deal with water column denitrification, we also consider nitrogen fixation as a surface layer source of nitrate when $NO_3^{l,h}$ levels there drop below $r_{NP}PO_4^{l,h}$ or equivalently when $NPP > NPN$. In that case, we take the rate of nitrogen fixation, $NF^{l,h}$, to be

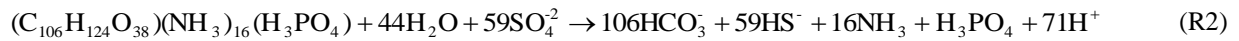$$NF^{l,h} = NF_0[\exp(NPP^{l,h}/NPN^{l,h}-1)-1] \qquad (3)$$

where NF_0 has been chosen to be 1x10^6 moles NO$_3$ per second.

In the water column, remineralisation takes place through oxidation of organic matter with dissolved oxygen as long as oxygen levels are above a certain minimum value O$_{2,\min}$, chosen here to be 3 mmol/m^3. Below that value, remineralisation is assumed to occur by way of denitrification as long as nitrate levels exceed a certain minimum level NO$_{3,\min}$, chosen here to be 0.03 mmol/m^3. The oxidation equation for denitrification is taken to be

$$(C_{106}H_{124}O_{38})(NH_3)_{16}(H_3PO_4) + 94.4HNO_3 \rightarrow 106CO_2 + 16NH_3 + 47.2N_2 + H_3PO_4 + 109.2H_2O \qquad (R1)$$

Here, as for oxidation with dissolved oxygen in the original DCESS model (Shaffer et al., 2008), we consider that organic matter formed in the ocean surface layer consists of proteins and lipids in addition to carbohydrates and adopt the mean composition proposed by Anderson (1995). Such a composition requires more oxygen for complete oxidation than carbohydrate alone would require (Anderson, 1995). This explains the enhanced nitrate sink in denitrification compared to standard Redfield stoichiometry (factors 94.4 and 47.2 compared with 84.8 and 42.4 in Reaction R1).

When oxygen concentrations are below O_{min} and nitrate concentrations are below N_{min}, remineralisation is assumed occur by way of sulphate reduction for which the oxidation equation is taken to be

$$(C_{106}H_{124}O_{38})(NH_3)_{16}(H_3PO_4) + 44H_2O + 59SO_4^{-2} \rightarrow 106HCO_3^- + 59HS^- + 16NH_3 + H_3PO_4 + 71H^+ \qquad (R2)$$

For the same reason as above, the sulphate sink is somewhat enhanced compared to standard Redfield stoichiometry (factor 59 compared with 53 in Reaction R2). Note that the carbon oxidation product in sulphate reduction is bicarbonate that, as opposed to the carbon dioxide produced in denitrification, contributes to alkalinity (ALK) as well as dissolved inorganic carbon (DIC). Both reactions show a minor contribution to alkalinity by way of ammonia production. In addition, for sulfate reduction with our organic matter model, the ALK decrease from the production of hydrogen ion exceeds the ALK increase from the production of bisulfide (HS^- ; from the definition of alkalinity, $ALK \propto HCO_3^- + NH_3 + HS^- - H^+$ (Dickson, 1981)). In summary, for our organic matter stoichiometry, we find ALK/DIC = 0.151 and 1.038 for denitrification and sulfate reduction, respectively.

As shown in Reactions R1 and R2, NH_3 and HS^- are produced by denitrification and/or sulphate reduction. As we show below, additional bisulfide is produced by microbes for anoxic conditions by way of anoxic methane oxidation (AMO). In the model, these species are oxidized to nitrate and sulphate, respectively, when transported by advection and diffusion to oxygenated ocean layers where $O_2 \geq O_{2,min}$. These oxidation reactions are

$$NH_3 + 2O_2 \rightarrow NO_3^- + H^+ + H_2O \qquad (R3)$$

and

$$HS^- + 2O_2 \rightarrow SO_4^{2-} + H^+ \qquad (R4)$$

Note that each of the reactions leads to an ALK decrease from production of hydrogen ions.

Oxygenation rates, $\Gamma_{ox,n}^{l,h}$ (in moles per second), are expressed by

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$$\Gamma_{ox,NH_4,n}^{l,h}, \Gamma_{ox,H_2S,n}^{l,h} = -V_n^{l,h}([NH_3]_n^{l,h}, [HS^-]_n^{l,h}) / \tau_{ox,NS} \qquad (4)$$

where n is the layer number, $V_n^{l,h}$ are the layer volumes, $[NH_3]_n^{l,h}, [HS^-]_n^{l,h}$ are the layer specie concentrations and $\tau_{ox,NS}$ is an ocean lifetime taken to be 200 days.

10   The simple ocean nitrogen and sulphur chemistry sketched above is intended to capture the most important processes that would occur in the ocean following a massive injection of methane without going into further detail of the N and S cycles. Processes that are left by the wayside here include anerobic ammonium oxidation (anammox), oceanic production of nitrous oxide and iron sulphide formation (Kuypers et al. 2003; Freing et al. 2012; Morse et al. 1987). Anammox has been shown to be an important sink of nitrogen nutrient in the ocean but denitrification may still be the most important player here (Ward et

15   al., 2009; Zehr and Kudela, 2011). Treatment of anammox and nitrous oxide would involve in depth consideration of nitrogen cycling including nitrite production and consumption as well as approaches that transcend our simple, horizontally-averaged model geometry. This is beyond the scope of the present paper but is being addressed by our group in other DCESS model developments in progress. Likewise, sulphide formation would require an explicit treatment of iron cycling also beyond the scope of the model at present.

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3.2.2 Methane oxidation and air-sea gas exchange

We now include methane as a tracer in the ocean model acted upon by advection, diffusion, air-sea gas exchange and

25   microbial oxidation in the water column. Here we ignore any weak methane production that may take place in the surface ocean in anoxic microenvironments (Reeburgh, 2007) again with the goal of concentrating on massive methane injections. Oxidation of methane in oxygenated ocean layers ($O_{2,n}^{l,h} \geq O_{2,min}$) produces carbon dioxide and consumes oxygen according to

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$$CH_4 + 2O_2 \rightarrow CO_2 + 2H_2O \qquad (R5)$$
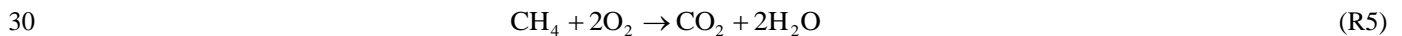
Layer oxidation rates of methane under these conditions, $\Gamma_{ox,CH_4,n}^{l,h}$, are taken to be:

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$$\Gamma_{\mathrm{ox,CH_4},n}^{l,h} = -V_n^{l,h}([\mathrm{CH_4}]_n^{l,h} / \tau_{ox,\mathrm{CH_4}}$$ (5)

where $[\mathrm{CH_4}]_n^{l,h}$ are the layer methane concentrations and $\tau_{ox,\mathrm{CH_4}}$ is an ocean methane lifetime for oxic conditions. From a joint analysis of ocean methane and CFC data, Rehder et al., (1999) found support for modelling methane oxidation as a first order process (Eq. 5) and determined an oceanic methane lifetime of ~ 50 years. A much shorter lifetime of ~ 4 months was deduced for strong, fast and localized methane emissions during the 2010 Deepwater Horizon oil spill in the Gulf of Mexico (Kessler et al., 2011). Below we take $\tau_{ox,\mathrm{CH_4}}$ to be 50 years since this value is likely more applicable to our global approach. But in the following we also check the sensitivity of our results to the much shorter lifetime.

For suboxic/weakly anoxic conditions characterized by $O_2 < O_{2,min}$ and $NO_3 > NO_{3,min}$, we adopt the following reaction for nitrate-dependent, microbial AMO (Raghoebarsing et al. 2006; Cui et al. 2015)

$$5CH_4 + 8HNO_3 \rightarrow 5CO_2 + 4N_2 + 14H_2O$$ (R6)
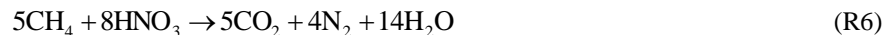
An alternative and slightly more energetic reaction involves nitrite, a species that we however do not consider in our simplified nitrogen cycling approach (Haroon et al. 2013).

For anoxic conditions for which $O_2 < O_{2,min}$ and $NO_3 < NO_{3,min}$, sulphate-dependent, microbial AMO (Treude et al., 2005; Cui et al., 2015) may be expressed as

$$CH_4 + SO_4^{2-} \rightarrow HCO_3^- + HS^- + H_2O$$ (R7)

Note that whereas nitrate-dependent, AMO (Reaction R6) does not produce ALK, the production of bicarbonate and bisulphide in sulphate-dependent, AMO (Reaction R7) is a strong ALK source leading to an ALK/DIC ratio of 2, considerably higher than for sulphate reduction of organic matter (Reaction R2). From carbonate chemistry, an increase in ocean alkalinity tends to depress dissolved carbon dioxide concentrations in the ocean and, by way of air-sea gas exchange, also in the atmosphere.

Layer oxidation rates of methane for sulphate- dependent AMO, $\Gamma_{\mathrm{anox,CH_4},n}^{l,h}$, are taken to be:

$$\Gamma_{\mathrm{anox,CH_4},n}^{l,h} = -V_n^{l,h}([\mathrm{CH_4}]_n^{l,h} / \tau_{anox,\mathrm{CH_4}}$$ (6)

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where τ_{anox,CH_4} is an ocean methane lifetime for anoxic conditions ($O_2 < O_{2,min}$ and $NO_3 < NO_{3,min}$) that may be considerably longer than τ_{ox,CH_4} since Reaction R7 is much less energetic than Reaction R5 (Lam and Kuypers, 2011). To take this into consideration and to be consistent with much lower anoxic remineralization rates compared to oxic rates in the DCESS model sediment module (Shaffer et al., 2008) , we adopt $\tau_{anox,CH_4} = 0.1\tau_{ox,CH_4}$ or 500 years as our standard case. On the other hand, nitrate-dependent AMO (Reaction R6) is energetically similar to oxic methane oxidation (Reaction R5) so we take τ_{ox,CH_4} for the methane lifetime associated with nitrate-dependent AMO.

Air-sea gas exchange for methane, $\Psi_{CH_4}^{l,h}$, is calculated as

$$\Psi_{CH_4}^{l,h} = k_w^{l,h}(\eta_{CH_4}^{l,h} pCH_4 - [CH_4]^{l,h}) \tag{7}$$

where the gas transfer velocities $k_w^{l,h}$ are $0.39u^2(Sc_{CH_4}^{l,h}/660)^{0.5}$ whereby $u^{l,h}$ are mean wind speeds at 10 m above the ocean surface, taken to be 8 m s^{-1} in both zones, and $Sc_{CH_4}^{l,h}$ are Schmidt numbers for methane that depend on ocean surface layer temperature (see below). The methane solubility, $\eta_{CH_4}^{l,h}$, was converted from the Bunsen solubility coefficients that depend on surface layer temperature and salinity (Wiesenburg and Guinasso, 1979) to model units using the ideal gas mole volume. Furthermore, $[CH_4]^{l,h}$ is the dissolved methane concentration in the ocean surface layers.

In recent work it was shown that the traditional calculations of Sc_{CO_2} based on a third order polynomial of surface layer temperature (Wanninkhof, 1992) seriously underestimated Sc_{CO_2} for temperatures (T) above 30°C (Gröger and Mikolajewicz , 2011). These authors proposed the form $Sc = a\exp(-bT) + c$ and fit that equation using experimental data of Jähne et al., (1987) to find best fit values for CO$_2$ of 1956, 0.0663 and 142.6 for a, b and $c,$ respectively. Since we are dealing here with simulations of extreme global warming, we adopt this new Sc_{CO_2} expression and fit here, thereby superseding the traditional Sc_{CO_2} calculation previously used in our model (Shaffer et al., 2008). Gröger and Mikolajewicz (2011) did not consider methane but we directly fit their proposed exponential function to the experimental data of Jähne et al., (1987) for Sc_{CH_4} over 5°C intervals in the temperature range 0 – 40°C. We found best fit values (R^2 = 0.997) for a, b and $c,$ respectively, such that $Sc_{CH_4} = 1771\exp(-0.0650T) + 128.8$. We adopt this relationship here.

4 Sample simulations with methane cycling

4.1 Methane input specification

5   The rate of methane input, F_{CH_4}, we apply to the atmosphere/ocean system is taken to be

$$F_{CH_4}(t) = 404 F_{CH_4}^{tot} / (\tau_{in})^5 \exp(-6.27 t / \tau_{in}) \tag{8}$$

where $F_{CH_4}^{tot}$ is the total input in GtC, τ_{in} is a time scale in years over which the first 75% of the input is added and t is the

10   time in years. This mathematical form has been used before within our group to characterize CO_2 input (Shaffer 2009; Shaffer et al., 2016). As discussed in the Introduction, the great bulk of both possible major F_{CH_4} sources to fuel deep time warming events, methane hydrate and thermogenic methane, are likely to enter via the ocean. Estimates of the time scale for the carbon input driving the deep time global warming events have varied from years to tens of thousands of years (eg. Zeebe et al. 2009; Cui et al., 2011; Wright and Schaller, 2013). In the lower end of this time scale range, Earth System

15   response to $F_{CH_4}^{tot}$ will vary significantly depending on whether this methane is dissolved in the ocean or part or even all of it escapes as gas to the atmosphere, in bubble plumes for example (McGinnis et al, 2006). Such response variation may be explained in part by much shorter time scales for methane oxidation to carbon dioxide (in both the atmosphere and ocean) than ocean diffusive and overturning time scales. Escape to the atmosphere will cause immediate global warming since methane is a powerful greenhouse gas, much more so than carbon dioxide (Myhre et al., 1998). Oxidation of dissolved

20   methane in the ocean will consume dissolved oxygen there (Reaction R5). For oxidation of large amounts of methane, suboxic or anoxic conditions may develop with complex nutrient and carbon cycling consequences upon which we elaborate below. We address these questions here by taking the direct, undissolved methane escape to the atmosphere to be αF_{CH_4} and the methane dissolution in the ocean to be $(1-\alpha) F_{CH_4}$, whereby α is the fraction of total input that directly enters the atmosphere. Furthermore, we must specify how methane dissolution is distributed between the two ocean sectors and among

25   the 55 layers of each sector. Here we simply assume that 84% of the dissolution occurs in the low-mid latitude sector and 16% in the high latitude sector, a division that reflects model sector areas. Furthermore, we assume that the amount assigned to each sector is divided equally among the upper 30 model ocean layers, spanning the depth ranges over which methane hydrate and/or thermogenic methane inputs may be expected (e.g. Buffett and Archer, 2004; Svensen et al., 2004).

4.2 Pre-event steady state and a long sample simulation

In order to investigate the impacts of these model improvements on simulation results, we chose as an initial case study the late Paleocene (pre-PETM) configuration and associated model steady states constructed in Shaffer et al. (2016) but now including some changes like the adaption of the Byrne and Goldblatt (2014a) expressions for carbon dioxide radiative forcing and for carbon dioxide-nitrous oxide overlap. In this configuration we assume a solar constant decrease of 0.5 % from present day, sea level rise of 100 m, mean ocean salinity of 33.8, slightly reduced background albedo (to represent less land in the subtropics and more vegetation cover), extra high latitude, longwave cloud forcing of 30 W/m^2, pre-industrial atmospheric oxygen level, adjustments in ocean calcium and magnesium concentrations and adjustments in initial lithosphere outgassing and weathering inputs. Furthermore, parameters in the model's simplified treatment of outgoing long wave radiation were adjusted to yield a global mean temperature of 25 °C and a climate sensitivity in the range 4 -5 °C for atmospheric pCO$_2$ levels in the range 800-900 ppm (Shaffer et al., 2016). Open system steady states were then sought (by iteration and/or long time integration) such that external inputs (lithosphere outgassing and weathering) balanced external output (burial down out of the ocean sediment). Table 1 lists some properties of the model steady state we chose among various possibilities given the above constraints. This particular model configuration with a global mean temperature of 25.0 °C and an atmospheric pCO$_2$ of 860 ppm has a climate sensitivity of 4.8 °C. While this steady state largely reflects late Paleocene conditions to ease comparison with our prior work, we emphasize that a similar approach can be taken to design appropriate initial conditions for any particular deep time warming event, like the End-Permian event (Shen et. al., 2011), based on paleo-reconstructions for conditions prior to that particular event.

Figure 6 shows time evolutions of some relevant model variables from a 200,000 year simulation that started in equilibrium with the above initial conditions and was forced by methane input of 4000 GtC over a time scale (τ_{in}) of 3 kyr, whereby half of the input reaches the atmosphere and half is dissolved in the ocean (α = 0.5). Furthermore, to be specific it is assumed that the δ^{13}C of the methane input is -40 ‰. Such forcing leads to maximum event temperature and pCO$_2$ rises of 5.7 °C and 1040 ppm, a peak atmospheric pCH$_4$ of 10.3 ppm and a low-mid latitude, Calcite Compensation Depth (CCD) shoaling of 463 m (Fig. 6a,c,h).

The low-mid latitude ocean warms by up to 5.5 °C at the surface, increasing to 6.0 °C at depth, reflecting modest polar amplification of the warming by about 20% (Fig. 6b). Mean low-mid latitude dissolved O$_2$ concentrations at 1000 m depth approach zero after less than 2 kyr and remain so for an additional 3 kyr, driven to a large extent by oxidation of methane dissolved in the ocean (Fig. 6d). Considerable denitrification occurs during this 3 kyr long, anoxic period, reflected in an associated decrease in new production due to nitrogen limitation (Fig. 6e). After this decrease, ocean new production then

steadily increases to about 12% higher than pre-event levels by 24 kyr (due to more ocean phosphate from enhanced weathering) and then slowly decreases again as weathering decreases with temperature. There is only a slight initial decrease of about 100 GtC in land biosphere carbon inventory as more/less new production from CO_2 fertilization was roughly balanced by more/less soil remineralisation from warming/cooling (Fig. 6f). The initial decrease in both ocean new production and land biomass enhance atmospheric pCO_2 and warming while the subsequent ocean new production increase followed by a slow decrease modulate the slow decrease in pCO_2 and slow cooling following peak warming.

Weathering increases over the event are accompanied by an initial drop in carbonate burial as dissolution decreases the amount of sediment $CaCO_3$ as well as the burial velocity (Fig. 6g,h). There is little initial change in organic carbon burial as the opposing effects tend to balance: decreasing burial velocity and increased organic matter preservation from decreasing O_2 concentrations. During this period there is a net source of carbon to the ocean-atmosphere system (in addition to the prescribed methane input) as volcanic/weathering inputs exceed burial outputs. After the methane input event, both $CaCO_3$ and organic carbon burial increase in response to higher new and $CaCO_3$ production. Together with decreasing weathering in response to cooling, this results in weak but long-lived net sink of carbon from the ocean-atmosphere system (Fig. 6g). In combination with decreasing pCO_2 and associated increasing ocean $[CO_3^{2-}]$, the biogenic $CaCO_3$ production increase drives an overshoot of the low-mid latitude CCD to depths greater than the pre-event depth, in agreement with theory and sediment core data (Dickens et al, 1997; Leon-Rodriguez and Dickens, 2010; Penman et al. 2016). This overshoot, that is even more pronounced at high latitudes, peaks by 35 - 45 kyr into the simulation (Fig. 6h).

The low-mid latitude, pelagic $\delta^{18}O$ excursion in biogenic $CaCO_3$ is significantly muted compared to the benthic one due to enhanced $\delta^{18}O$ in the surface layer ($\delta^{18}O_w$) from a more vigorous hydrological cycle during the warming event (Fig. 6i) The carbon isotope excursion is slightly muted in the atmosphere and enhanced in the low-mid latitude, ocean surface layer due to warming and temperature-dependent fractionation in air-sea gas exchange (Fig. 6j). The low-mid latitude, marine organic matter isotope excursion is greater still, in agreement with paleo-reconstructions (McInerney and Wing, 2011), as a consequence of model dependence of carbon isotope discrimination during ocean photosynthesis on concentrations of dissolved carbon dioxide and phosphate in the ocean surface layer (Shaffer et al., 2016). Use of a pCO_2-dependent carbon isotope fractionation leads to a still greater isotope excursion in terrestrial organic matter, also in agreement with paleo-reconstructions (Jahren and Schubert, 2013). Note that the model carbon isotope excursion is more protracted than the initial warming or the $\delta^{18}O$ excursion related to the warming (Fig. 6a,i,j). Relaxation back to about half of maximum carbon isotope excursion values takes place over about 80 kyr compared to about 40 kyr for comparable temperature or $\delta^{18}O$ relaxation. This can be explained by enhanced burial of $\delta^{13}C$-enriched $CaCO_3$ after the methane input event. After 200 kyr, most of model Earth System properties have relaxed back toward pre-event values. This long time scale is dictated by external input/output balances of the global carbon cycle (Fig. 6g), in particular by the slow, negative silicate weathering feedback on climate (Shaffer et al., 2008).

4.3 Sensitivity to input size, time scale and distribution

Figure 7 shows selected model results for the first 20,000 years of simulations like that of Fig. 6 but with methane input amounts of 2000, 4000 and 6000 GtC. Both pCO_2 and especially pCH_4 increase more than linearly in response to the linear increase in methane input (Fig. 7a,c,d). For pCO_2 this is due to positive feedbacks from the land biosphere and ocean. Initial warming from the methane injection leads to greater soil remineralisation, land carbon inventory decrease and input of CO_2 to the atmosphere (Fig. 7f). Increasing methane oxidation in the ocean leads to denitrification that draws down ocean new production leading to increased ocean CO_2 outgassing that was already enhanced in response to decreased ocean CO_2 solubility from warming (Fig. 7e). The non-linear increase in pCH_4 is due mainly to the increase in CH_4 atmospheric lifetime with pCH_4 (Fig. 4). On the other hand, warming increases less than linearly (Fig. 7b) as expected from the radiative forcing dependencies shown in Fig. 5 and from our neglect here for simplicity of climate sensitivity increase with warming (Shaffer et al., 2016). Net carbon input from volcanism/weathering minus burial follows process sequences as described for Fig. 6 above (Fig. 7g). The CCD shoals as more methane is oxidized to CO_2 in the atmosphere and ocean leading to more acidic conditions that drive calcite dissolution in the sediment (Fig. 7h). However, CCD shoaling increase is damped for large methane dissolution in the ocean since this leads to anoxic conditions, methane oxidation with sulphate (Reaction R7) and associated alkalinity inputs that oppose dissolution. This also has had the effect of damping the atmospheric pCO_2 increase. In response to warming, oxygen isotopes of the low-mid latitude ocean surface water are enriched by enhanced evaporation associated with a stronger atmospheric hydrological cycle (Fig. 7i). Low-mid latitude, ocean surface layer salinity (not shown) increases in response to warming for the same reason and the salinity excursion can be approximated very well by multiplying the results in Fig. 7i by 2.8. Atmospheric carbon isotope excursion values are directly proportional to the size of the methane input and exhibit a long time decay scale as noted in Fig. 6 above (Fig. 7j):

Figure 8 shows selected model results for the first 20,000 years of simulations like that of Fig. 6 but with methane injection times τ_{in} of 300, 1000, 3000 and 10,000 years. For the shortest injection times, input rates in terms of carbon equal or exceed those of present day anthropogenic carbon emissions exceeding 10 GtC/yr (Fig. 8a; IPCC, 2013). Global warming spikes to much higher values for short injection times in response to large instantaneous methane radiative forcing and under the influence of slow ocean exchange times (Fig. 8b,d). Warming tops out at 8.0, 6.7, 5.7 and 4.8°C for the shortest to the longest injection times. Atmospheric pCO_2 is also enhanced for shorter injection times in part due to the positive feedbacks from the land biosphere and ocean noted above (Fig. 8c,e,f). Shorter injection times lead to more widespread sub-oxic/anoxic conditions as evidenced by initial decreases in ocean new production in response to denitrification (Fig. 8e). These conditions are a direct response to increased methane oxidation in the ocean due to enhanced methane input rates (Fig. 8a). One consequence of this situation is the muted CCD shoaling for shorter injection times following from enhanced methane oxidation with sulphate (Reaction R7) and associated alkalinity inputs that oppose $CaCO_3$ dissolution (Fig. 8h).

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Both net carbon input and especially oxygen isotopes from low-mid latitude, ocean surface layer water follow global warming due in large part to climate-dependent weathering and temperature-dependent, atmospheric water vapor transport (Fig. 8a,g,i; Shaffer et al., 2008). The atmospheric carbon isotope excursion is much enhanced for short injection times. This can be understood as follows: the carbon isotope signal from the methane injected to the atmosphere, and transferred there to CO$_2$ via oxidation of the methane, builds up in the atmosphere and in the ocean surface layer (not shown) due to long exchange times of about 1 kyr with the deep ocean (Fig. 8j). The atmospheric isotope excursion tops out at -5.24, -3.80, -3.21 and -3.03 for the shortest to the longest injection times.

Figure 9 shows selected model results for the first 20,000 years of simulations like that of Fig. 6 but with 0, 50 and 100% of the methane input reaching the atmosphere ($\alpha = 0$, 0.5 and 1). Global warming, atmospheric pCH$_4$ and, to a lesser extent, atmospheric pCO$_2$ are enhanced for more input directly to the atmosphere (Fig. 9b,c,d). Warming is enhanced by 1.7°C for all methane input to the atmosphere compared to all input dissolved in the ocean. Even with all methane input dissolved in the ocean, atmospheric pCH$_4$ increases initially by more than 2 ppm as methane outgases from the ocean via air-sea gas exchange (Fig. 9d). The positive land biosphere feedback on atmospheric pCO$_2$ is enhanced for all methane to the atmosphere due to enhanced warming (Fig. 9f). On the other hand, ocean feedbacks on atmospheric pCO$_2$ are more nuanced. The positive, solubility feedback is enhanced for all methane to the atmosphere due to enhanced warming whereas the positive feedback from reduced ocean new production is enhanced for all methane to the ocean (Fig. 9e). The latter effect is due to methane oxidation in the ocean leading to denitrification and nitrogen limitation of new production. Increased CaCO$_3$ dissolution in the ocean sediment is a further consequence of methane oxidation in the ocean when all methane is dissolved there. This leads to decreased calcite burial and enhanced net carbon input to the ocean-atmosphere system despite less weathering input from less global warming here (Fig. 9g). This also explains the enhanced CCD shoaling for all methane dissolved in the ocean (although modulated somewhat from increased methane oxidation with sulphate and associated alkalinity inputs as discussed above; Fig. 9h). Again, oxygen isotopes from low-mid latitude, ocean surface layer water follow global warming due to temperature-dependent, atmospheric water vapor transport (Fig, 9i). The atmospheric carbon isotope excursion peaks slightly earlier for all methane to the atmosphere but, due to the long time scale of the methane input (3000 kyr) relative to ocean overturning times, the excursion amplitude is essentially the same for all cases (Fig. 9j). The flow diagram presented in Fig. 10 serves to compare and contrast the pathways and feedbacks associated with 1) methane input to the ocean only and 2) methane input to the atmosphere only. These are the cases plotted as blue and maroon lines, respectively, in Fig. 9.

We also carried out a simulation like the case in Fig. 9 for all methane dissolving in the ocean (blue lines) but for a much shorter oceanic methane lifetime of 4 months (compared to the standard lifetime of 50 years; see Section 3.2.2). Results for this new simulation are plotted as dotted blue lines in Fig. 9. The shorter lifetime leads to much reduced methane concentrations in the ocean and less outgassing to the atmosphere (Fig. 9d). Furthermore, faster methane oxidation in the

ocean decreases dissolved oxygen there slightly, leading to slightly enhanced anoxia, sulphate-dependent, anoxic methane oxidation and alkalinity inputs. As a consequence, CCD shoaling is damped somewhat as is the atmospheric pCO2 increase (Fig. 9c,h). Together with less methane outgassing, this leads to a reduction of maximum global warming of about 7% compared to the standard case (Fig. 9b).

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4.4 Modelled ocean distributions

Figure 11 shows selected, low-mid latitude model ocean results for the first 20,000 years of a simulation forced by a methane input of 6000 GtC with $\delta^{13}C$ = -40 ‰ over a time scale (τ_{in}) of 3 kyr, whereby all the input is dissolved in the ocean ($\alpha =$ 10   0). This forcing configuration was chosen to highlight the workings of the model for suboxic/anoxic conditions and leads to maximum event temperature and pCO2 rises of 5.7 °C and 1095 ppm, a peak atmospheric pCH4 of 6.2 ppm and a low-mid latitude CCD shoaling of 389 m (not shown).

Low-mid latitude ocean warming is greatest at depth, a reflection of polar amplification (Fig. 10a). Maximum warming there 15   (6.2 °C) occurs about 1500 years after maximum surface layer (or atmosphere) warming. Subsurface methane concentrations build up to more than 30 mmol/m^{-3} (Fig. 11b). Oxidation of this methane with dissolved oxygen (Reaction R5) forces suboxic conditions (O_2 < 10 mmol/m^{-3}) for thousands of years at intermediate and mid-depths of the low-mid latitude ocean (Fig. 11c). Furthermore, a combination of reduced solubility due to warming and enhanced remineralisation due to increased ocean new production (Figs. 5-7) maintain suboxic conditions at intermediate depths there well beyond the 20,000 20   simulation years shown here. As dissolved oxygen is forced below $O_{2,min}$ (3 mmol/m^3), methane oxidation proceeds by way of nitrate-dependent, anoxic methane oxidation (Reaction R6), essentially eliminating nitrate for several thousand years at intermediate and mid-depths of the low-mid latitude ocean (Fig. 11d). Over a period of several thousand years after most methane is oxidized, nitrate concentrations recover to and even above pre-event values. This can be explained by a combination of nitrogen fixation (Eq. 3) and enhanced new production due to increased ocean phosphate concentrations 25   from warming-enhanced weathering. As nitrate is forced below $NO_{3,min}$ (0.03 mmol/m^3), methane oxidation proceeds by way of sulphate-dependent, anoxic methane oxidation (Reaction R7). This leads to the production of hydrogen sulphide that reaches concentrations over 10 mmol/m^{-3} at intermediate depths almost 3 kyr into the simulation (Fig. 11e).

Fig. 11f shows the model low-mid latitude, time-space distribution of Omega (Ω) defined as the ratio of carbonate ion 30   concentration to carbonate ion saturation concentration for calcite. Calcite dissolution in the sediment increases as Ω decreases from 1 and model biogenic calcite production in the surface layer is proportional to $(\Omega-1)/\{1+(\Omega-1)\}$ for $\Omega \geq 1$ and zero for Ω < 1 (Shaffer et al., 2016). In response to methane oxidation to CO_2, carbonate ion and thereby Ω initially decrease in intermediate and mid-depths where this oxidation occurs. However, as dissolved oxygen and nitrate are

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consumed and sulphate-dependent, anoxic methane oxidation takes over about 2 kyr into the simulation, Ω increases in these layers in response to alkalinity produced in this reaction. After the period of sulphate-dependent, anoxic methane oxidation, Ω decreases again followed by a slow increase driven by decreasing CO_2 levels from carbonate compensation. Low-mid latitude, oxygen isotope excursions in biogenic carbonate produced in-situ (Fig. 11g) track temperature changes in the deep ocean but are reduced relative to these changes near the surface due to enhanced ambient water isotopes (Fig. 9i). After several thousand years, low-mid latitude, carbon isotope excursions in biogenic carbonate produced in-situ (Fig. 11h) are slightly depressed in the deep ocean. This is due to dilution from additional, less-depleted carbon inputs associated with sediment $CaCO_3$ dissolution.

Figure 12 shows selected, low-mid (LL) and high latitude (HL) model ocean profiles at 0, 3 and 5 kyr into the simulation of Fig. 11 and serves to illustrate differences between the LL and HL ocean zones. Although methane input is scaled to be comparable per unit area for both zones, much higher methane concentrations build up in the LL zone than in the HL zone (Fig. 12a). Much higher HL vertical exchange spreads methane vertically and promotes some outgassing to the atmosphere there. Phosphate concentrations increase with time in response to warming-enhanced weathering (Fig. 12b), explaining the slow increase in ocean new production shown above (Figs. 7-9). Despite sub-oxic and eventually anoxic conditions in the LL zone, the HL zone remains oxygenated due to stronger vertical exchange and the downwelling branch of the overturning circulation there (Fig. 12c). As a consequence, HL nitrate concentrations remain elevated and there is no anoxic remineralisation nor anoxic methane oxidation there (Fig. 12d). This is also explains the essential lack of HL ammonium and hydrogen sulphate, < 0.02 and < 0.1 mmol/m^3, respectively (Figs 12e,f). Ammonium or hydrogen sulphate reaching the HL zone via horizontal exchange with the LL zone is quickly oxidized. By way of carbonate chemistry, addition of CO_2 to the water column via methane oxidation depresses carbonate ion concentrations (Fig. 12g). In terms of Ω (see above) this reduction is relatively stronger in the HL surface layer, tending to depress biogenic carbonate production there. However, this reduction is largely offset by warming enhancement. Addition of CO_2 to the water column via methane oxidation increases ocean acidity (Fig. 12h). For the simulation considered here, surface layer pH is reduced by 0.26 and 0.28 for the LL and HL zones, respectively, from pre-event levels already 0.47 and 0.48 lower than model pre-industrial levels (Shaffer et al., 2008).

4.5 Modelled ocean sediment properties and synthetic sediment cores

DCESS model simulations include prognostic values for burial velocities down out of each of the model's 110 bioturbated sediment layers, each 10 cm thick, as well as sediment properties exported down out of these layers. In the original model, synthetic sediment cores (SSC) produced in this way only contained information on calcite and organic matter content of the buried sediment (Shaffer et al., 2008). We have extended this treatment to include carbon and oxygen isotopes and report the first results of this here for bulk carbonate, in the form of calcite in the model. For each sediment layer we consider

conservation of calcite as well as conservation of carbon and oxygen isotopes in this calcite. This involves tracking the time-dependent inputs (from the rain of biogenic calcite produced in the ocean surface layer) and outputs (dissolution within the sediment layer and export down out the layer by burial). Isotope effects for changes in surface layer CO_3^{2-} are applied in the production of biogenic calcite (Spero et al., 1997). The new model extension also accounts for effects of possible "mining" of buried sediment (i.e. upward directed "burial" velocities) in response to very strong dissolution events (Shaffer et al., 2008). For this, properties buried in each SSC are recorded to provide correct values for properties reentering the active sediment layer from below during any such "mining" event.

Figure 13 shows bulk carbonate results from model low-mid and high latitude SSC's at 3000 m depth together with ocean surface layer properties, all for the simulation of Figure 10. Burial velocities and sediment calcite content are greater in the HL zone compared to the LL zone due to considerably greater model HL rain rates per unit area (Figs. 13a,b). This can be explained in part by temperature-enhanced, HL rain rates in the model: the pre-event HL ocean surface layer is almost 10°C warmer than for the pre-industrial simulation (Shaffer et al., 2008). For this reason and due to slightly higher new production (Table 1), pre-event, HL biogenic calcite rain is more than three times greater than for the pre-industrial case. On the other hand, pre-event, LL biogenic calcite rain is only about 15% greater than for this case. Over the first 20 kyr of the simulation shown here, 38.7 cm of sediment is laid down at 3000 m depth in the HL zone compared to 14.0 cm in the LL zone. Burial velocities and sediment calcite content decrease initially in response to increased dissolution and, to a lesser extent, decreased rain rate, as a consequence of the methane input and its oxidation in the ocean. Later on, both properties increase to exceed their pre-event levels in response to increasing new production from increasing ocean phosphate levels.

In both model zones the carbon isotope excursion in the SSC bulk carbonate is modestly attenuated relative to the ocean surface layer excursions (Figs. 13c,e). This is due to reservoir time scales of up to several thousand years for the active sediment layer and, to a lesser extent, to the CO_3^{2-} isotope effect (a reservoir time scale is the calcite inventory of a sediment layer divided by rate of calcite input to it). The reservoir effect also leads to somewhat distended bulk carbonate excursion maxima centered 3-4 kyr after surface layer maxima (note that the time axis of the figure is "stretched/squeezed" relative to the sediment length axis, a product of time-varying burial rates). Similar conclusions and interpretations also apply to the oxygen isotope excursions in the SSC bulk carbonate and the ocean surface layers in both model zones (Figs. 13d,f). Note that if, in lack of further knowledge, a linear time scale would be assigned to SSC length, the bulk carbonate excursions would appear sharper and shorter, particularly in the HL zone. When used as a paleothermometer, SSC bulk carbonate in the LL zone severely underestimates the model temperature excursion in the LL surface layer, primarily due to the positive ambient water $\delta^{18}O$ excursion demonstrated above, but with a minor contribution from the CO_3^{2-} isotope effect (Fig. 13d). On the other hand, the slightly negative ambient water $\delta^{18}O$ excursion in the HL surface layer and the CO_3^{2-} isotope effect there work to improve the estimate of the surface layer warming from the $\delta^{18}O$ excursion of SSC bulk carbonate, albeit with the 3-4 kyr time lag due to the reservoir effect (Fig. 13f).

Model availability

5   The DCESS model code can be downloaded at www.dcess.dk.

Author contributions

G. Shaffer designed the work, wrote the original draft and led the development of model extensions and code, to which E.
10   Fernandez Villanueva, R. Rondanelli, S. Olsen and J.O.P. Pedersen contributed. E. Fernández Villanueva carried out the
model simulations. All authors discussed model concepts and results and contributed to the final manuscript.

Competing interests

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

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	Pre-event	Pre-industrial
Atmosphere mean temperature (°C)	25.0	15.0
Atmosphere pCO$_2$ (ppm)	860	278
Atmosphere pCH$_4$ (ppm)	1.51	0.72
Atmosphere carbon inventory (GtC)	1826	590
Land carbon inventory (GtC)	2597	2220
Ocean carbon inventory (GtC)	33,473	37,910
Total carbon inventory (GtC)	37,895	40,720
Low-mid latitude CCD depth (m)	4239	4673
Ocean new production (GtC/yr)	6.47	5.40
Ocean biogenic CaCO$_3$ production (GtC/yr)	1.47	0.97
Ocean mean O$_2$ (mmol/m^3)	111.2	183.5

Table 1: Pre-event steady state solution properties compared to pre-industrial properties

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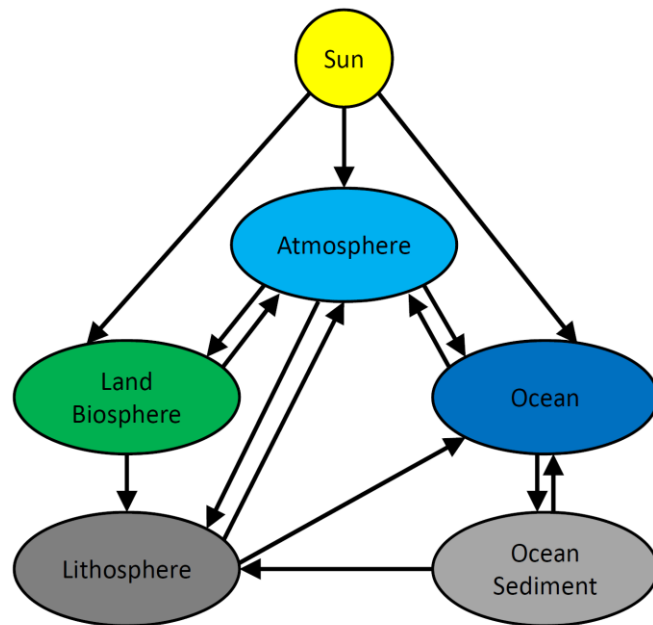
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Figure 1: Modules and interconnections for the Danish Center for Earth System Science (DCESS) Earth System model

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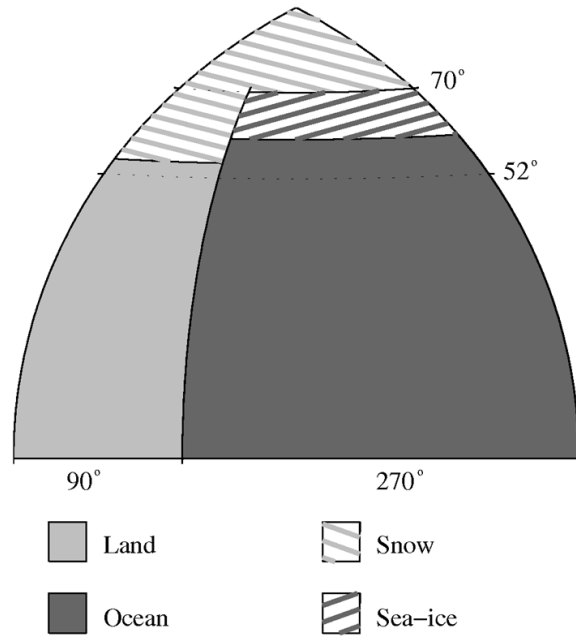
Figure 2. Model configuration. The model consists of one hemisphere, divided by 52° latitude into 360° wide, low-mid latitude and high latitude sectors. Values for global reservoirs, transports and fluxes are twice the hemispheric values. The model ocean is 270° wide and extends from the equator to 70° latitude. The ocean covers 70.5% of the model surface. Low-mid latitude and high latitude ocean sectors have the area ratio 84:16. Sea ice and snow coverage are diagnosed from idealized, meridional profiles of atmospheric temperature.

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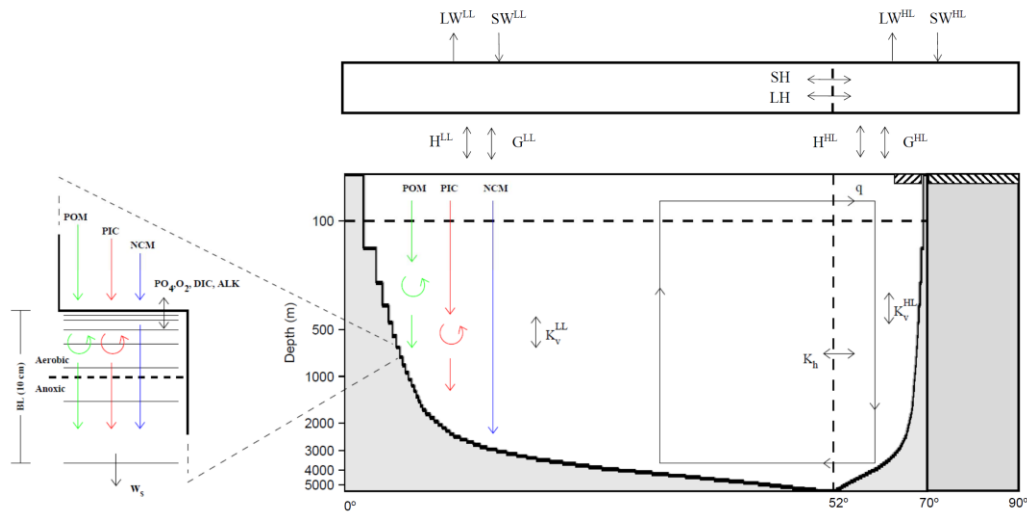
Figure 3. Atmosphere, ocean and ocean sediment configurations. Temperatures of low-mid (LL) and high (HL) latitude atmosphere sectors derive from energy balance forced by yearly-mean insolation (SW), outgoing longwave radiation (LW), meridional transports of sensible (SH) and latent (LH) heat, and air-sea heat exchange (H). In combination with the simple sea ice and snow (hatched bars) parameterizations, the model includes the ice/snow-albedo feedback and the insulating effect of sea-ice. Meridional variations of atmospheric temperature are represented by a second order Legendre polynomial in sine of latitude, adjusted to fit sector mean temperatures. Temperatures and gradients at the 52° dividing latitude from this meridional profile are used to calculate sensible heat, latent heat and water vapour transports. The atmosphere exchanges heat and gases (G) with the ice-free part of the ocean. Gases are well-mixed in the atmosphere and their concentrations follow from internal sinks, exchanges with ocean surface layers and the land biosphere and input/outputs from weathering and lithosphere outgassing. Both low-mid latitude and high latitude ocean sectors are continuously-stratified with 100 m vertical resolution to maximum depths of 5500 m and each sector has a bottom topography based on real ocean depth distribution. Model ocean circulation and mixing are indicated by the black arrows and associated circulation/mixing type labels (see main text). Model ocean biogeochemical cycling is indicated by the colored arrows (acting in both sectors but for simplicity only shown for LL sector) and associated matter (particulate organic (POC), biogenic carbonate (PIC) and inert mineral (NCM)). Each of the 110 model sediment segments has an area set by the real ocean depth distribution and consists of a 10 cm thick, bioturbated layer divided into 7 sublayers. Sediment biogeochemical cycling is indicated by colored arrows and associated matter. Exchange of dissolved species at the sediment-water interface is also indicated. The sedimentation rate down out of the bioturbated layer (w_s) is calculated from mass conservation.
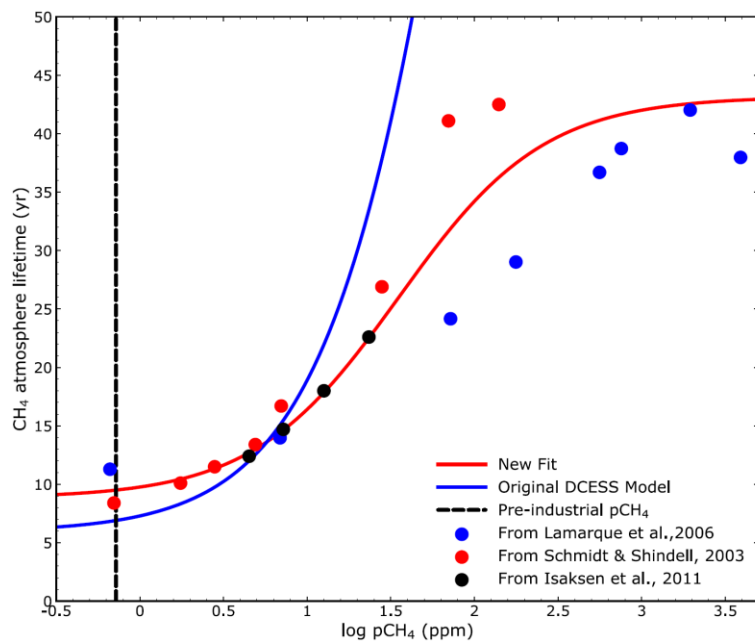
Figure 4: Atmospheric lifetime of methane as a function of atmospheric methane concentration. See text for a discussion of the model results and the functional fit to them.
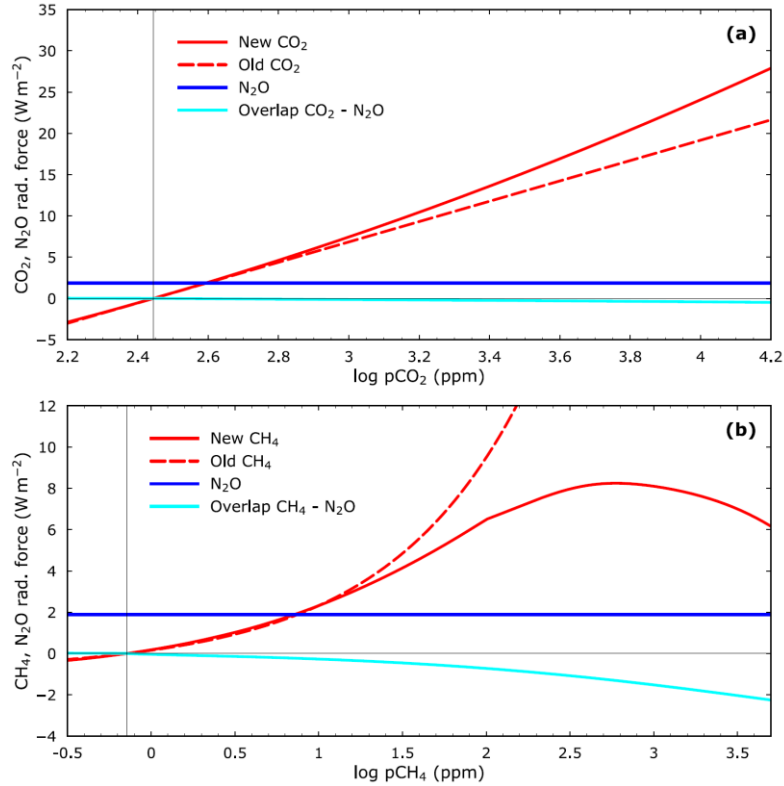
Figure 5: Radiative forcing of carbon dioxide, methane and nitrous oxide as functions of their atmospheric concentrations. a) CO_2 radiative forcing used here (new CO_2; Byrne and Goldblatt, 2014a) compared to CO_2 forcing in the original DCESS model (old CO_2; Shaffer et al., 2008). CO_2-N_2O overlap (Bryne and Goldblatt, 2014a) and N_2O radiative forcing calculated for constant, pre-industrial N_2O partial pressure of 270 ppb., **b**) CH_4 radiative forcing used here (new CH_4; Byrne and Goldblatt, 2014a,b) compared to CH_4 forcing in the original DCESS model (old CH_4). CH_4-N_2O overlap (Byrne and Goldblatt, 2014a) and N_2O radiative forcing calculated for constant N_2O partial pressure as in **a**.
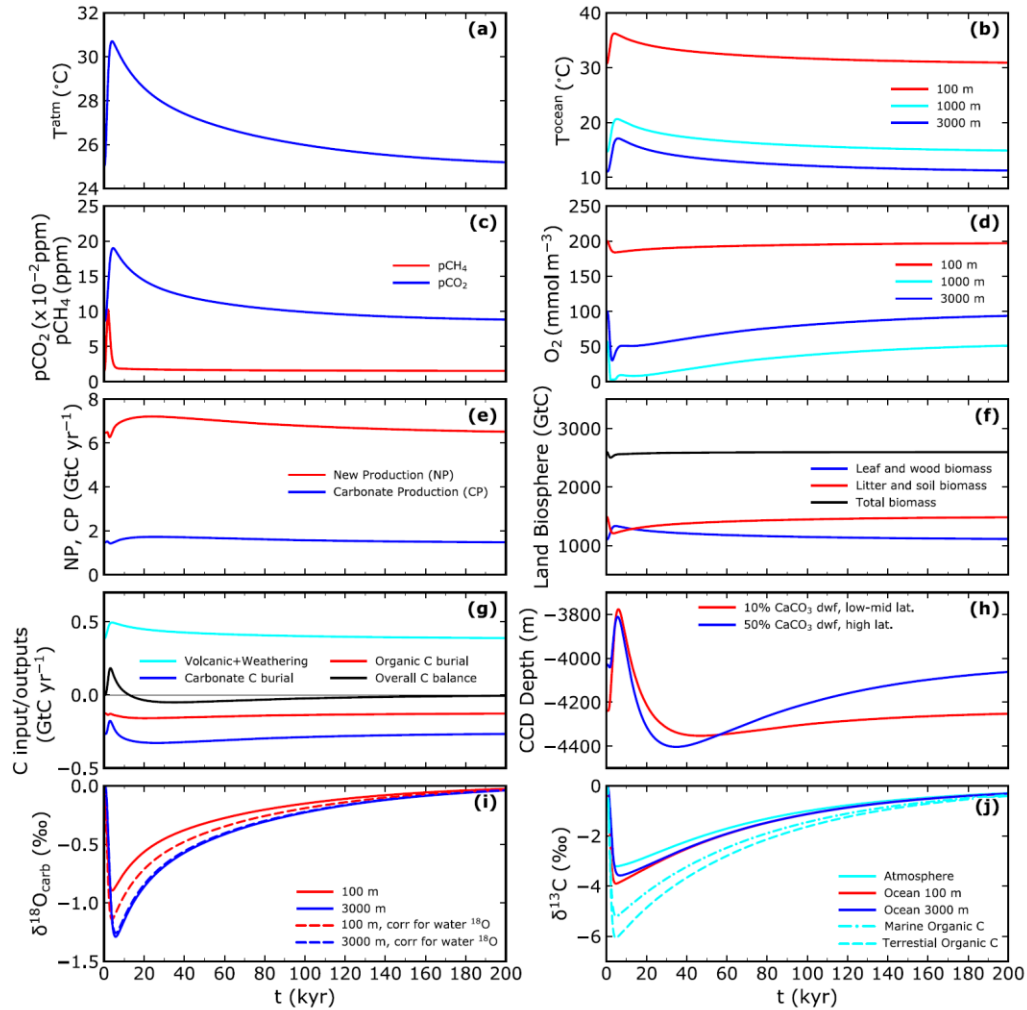
Figure 6: Results from a 200,000 year simulation with a methane input of 4000 GtC over a time scale of 3000 years and with half the input dissolved in the ocean and half escaping as gas to the atmosphere. a) Global mean atmospheric temperature, b) Low-mid latitude ocean temperature c) atmospheric pCO_2 and pCH_4 , d) Low-mid latitude ocean dissolved O_2 concentration, e) Ocean new and biogenic $CaCO_3$ production, f) Land biosphere carbon inventories for Leaves+Wood, Litter+Soil and their total, g) External input/outputs of carbon. The black line is volcanic/weathering inputs minus burial outputs. h) Depths to the low-mid latitude 10 % (CCD) and high latitude 50 % $CaCO_3$ wt%, i) Low-mid latitude excursions of $\delta^{18}O$ in biogenic $CaCO_3$ formed as well as these excursions corrected for excursions in ambient $\delta^{18}O_w$. j) Carbon isotope excursions for the atmosphere, the low-mid latitude ocean, low-mid latitude marine organic carbon and terrestrial organic carbon (see text; the methane input has $\delta^{13}C$ = -40 per mil).

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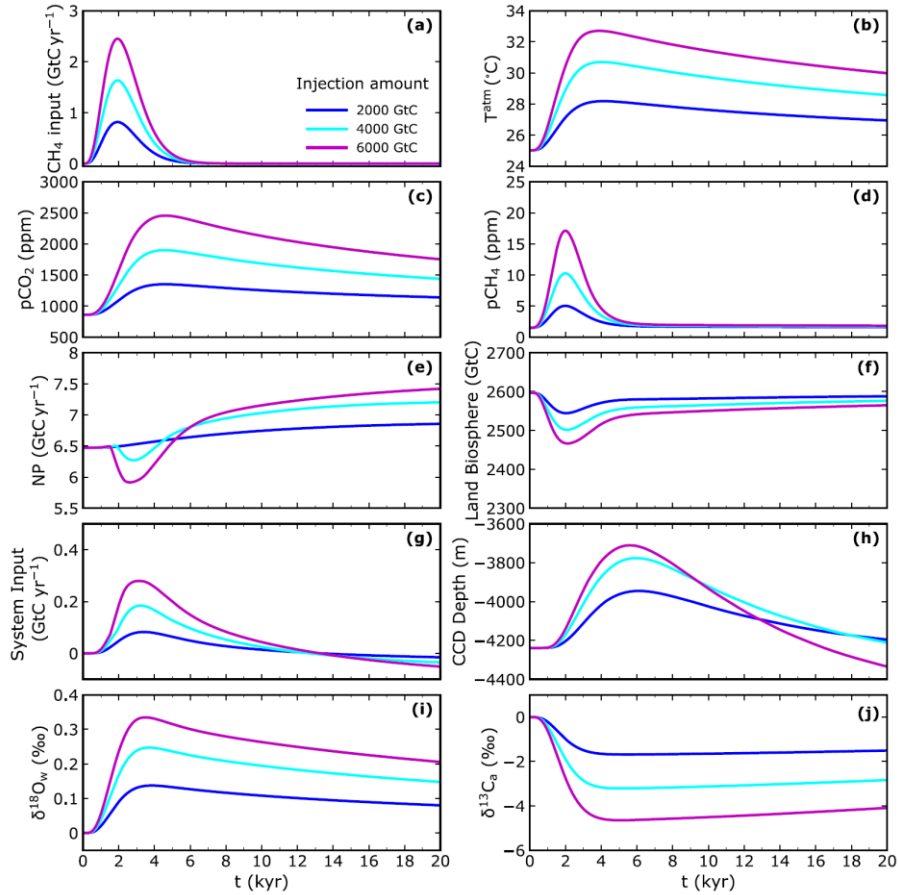
Figure 7: Results for 20,000 year simulations for different methane input amounts over a time scale of 3000 years and with half the input dissolved in the ocean and half escaping as gas to the atmosphere. a) methane input rate, **b**) global mean atmospheric temperature, **c**) atmospheric partial pressure of carbon dioxide, **d**) atmospheric partial pressure of methane, **e**) total ocean new production, **f**) total land biosphere biomass, **g**) volcanic/weathering inputs minus burial outputs (as black line in Fig. 6g), **h**) Low-mid latitude CCD depth (10% $CaCO_3$ dry weight in sediment), **i**) Oxygen isotope excursion of low-mid latitude, ocean surface layer water, **j**) Atmospheric carbon isotope excursion (the methane input has $\delta^{13}C$ = -40 per mil).
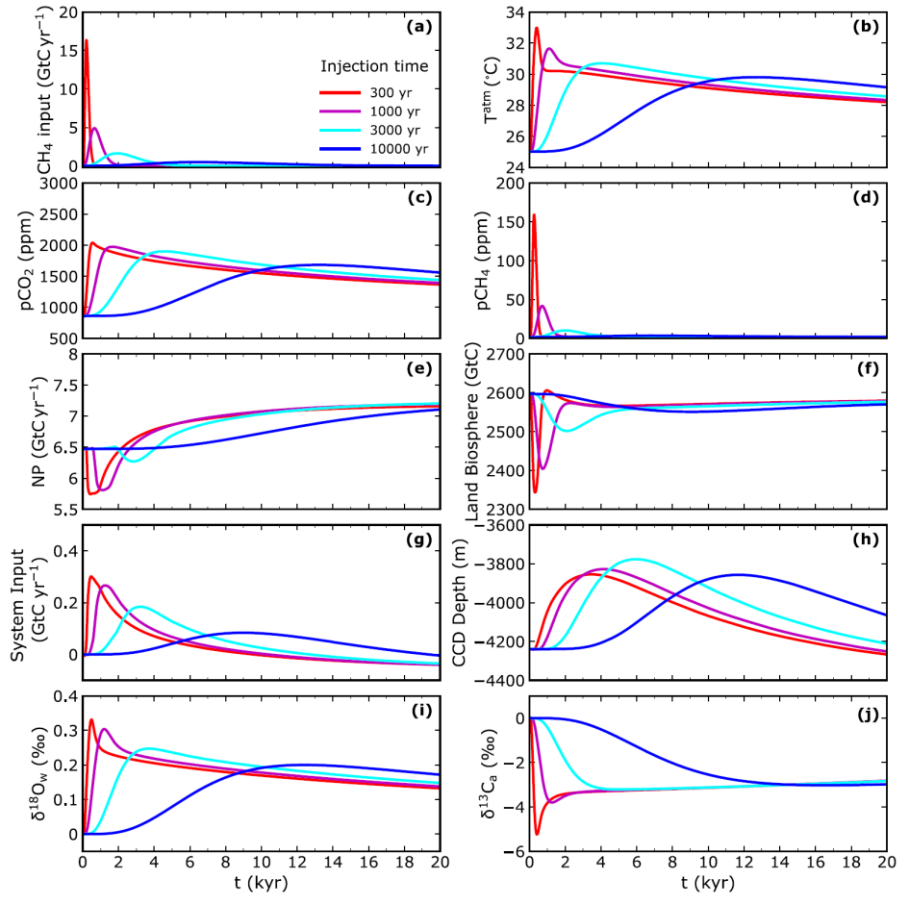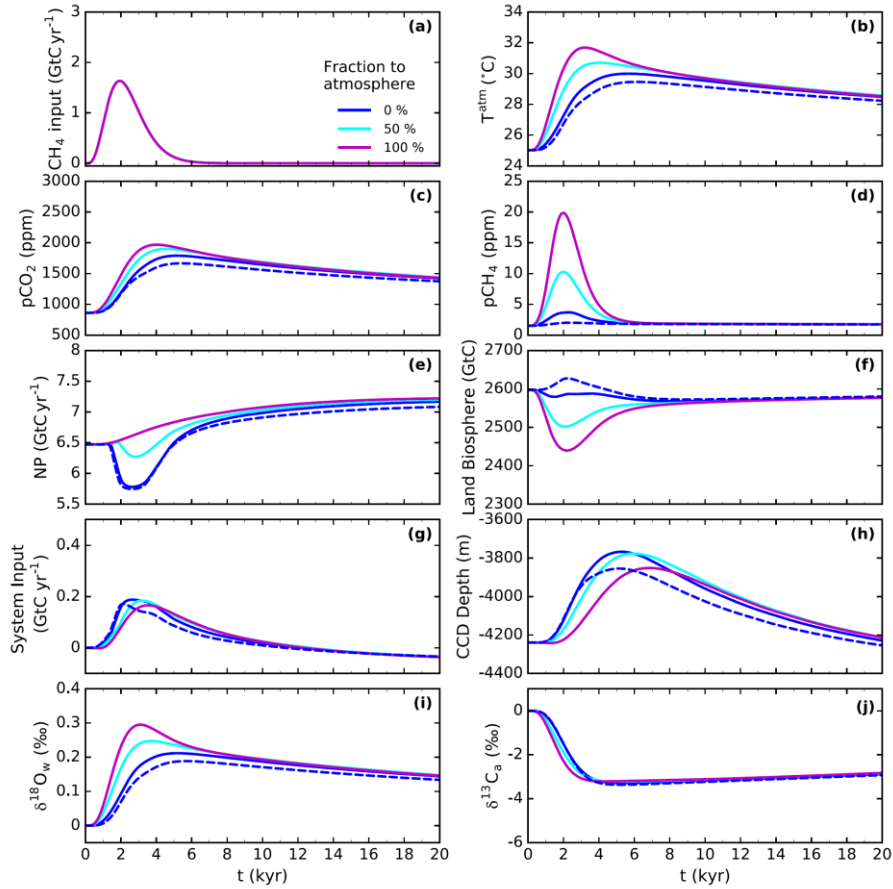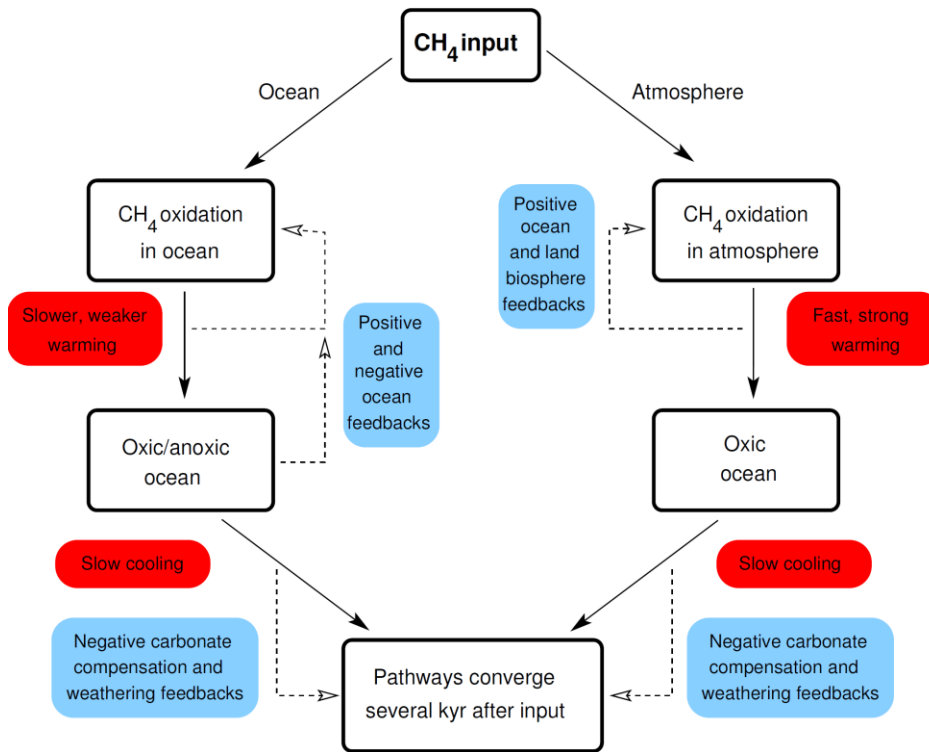
Figure 8: Results for 20,000 year simulations for different methane input time scales for an input of 4000 GtC with half the input dissolved in the ocean and half escaping as gas to the atmosphere. Properties plotted in **a**) − **j**) as in Fig. 7.

Figure 9: Results for 20,000 year simulations for different fractions of methane escape to the atmosphere for a methane input of 4000 GtC over a time scale of 3000 years. Properties plotted in **a**) – **j**) as in Fig. 7. Also shown are results from a simulation with all methane input to the ocean (0 % Fraction to atmosphere) and a much shorter ocean methane lifetime of three months (dashed blue lines; see main text).
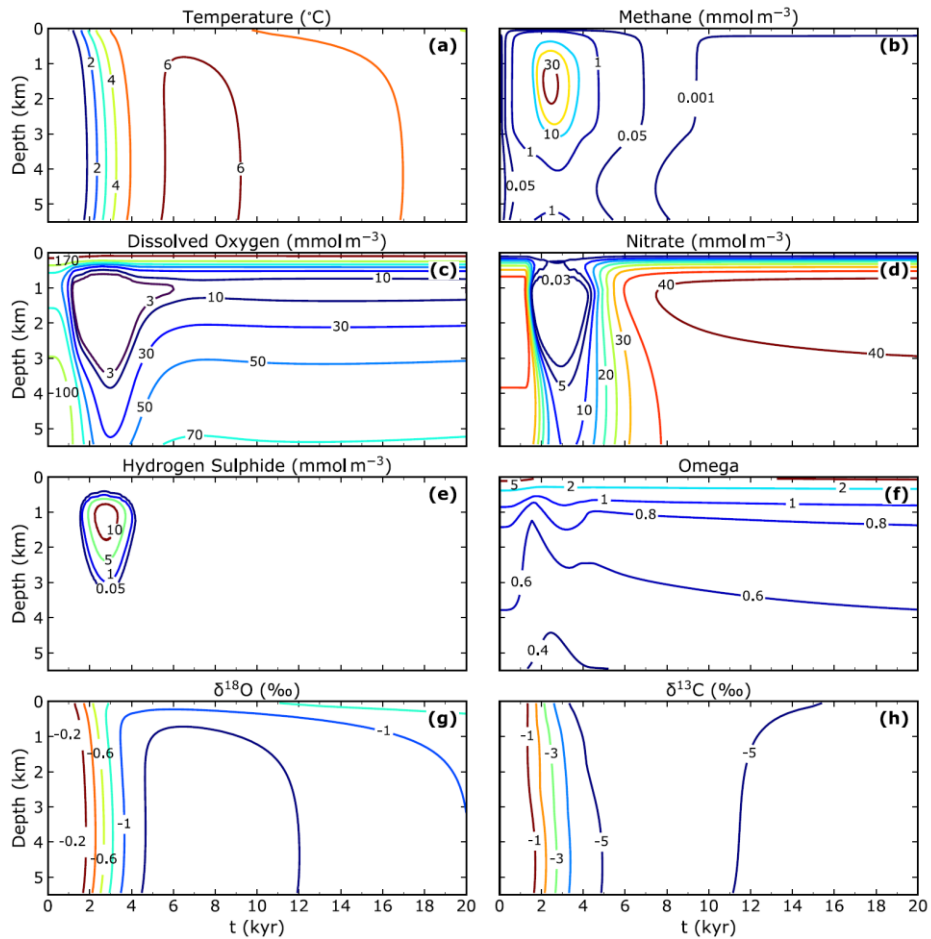
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Figure 10: Comparison of pathways and feedbacks for all methane injection into the ocean or into the atmosphere

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Figure 11: Low-mid latitude ocean property results as functions of water depth and time for a 20,000 year simulation with a methane input of 6000 GtC over a time scale of 3000 years, all of which dissolves in the ocean. a) Temperature anomaly, **b**) methane concentration,, **c**) dissolved oxygen concentration (model denitrification for $O_2 < 3$ mmol m^{-3}), **d**) nitrate concentration (model sulphate reduction for $NO_3 < 0.03$ mmol m^{-3}), **e**) hydrogen sulphide concentration, **f**) Omega, 10 the ratio of carbonate ion concentration to the carbonate ion saturation concentration for calcite (see Fig. 11g), **g**) Oxygen isotope excursion in biogenic carbonate produced in-situ. **h**) carbon isotope excursion. No corrections for the carbonate effect have been applied in **g** and **h** since these corrections were only derived for pelagic foraminifera (Spero et al, 1997),
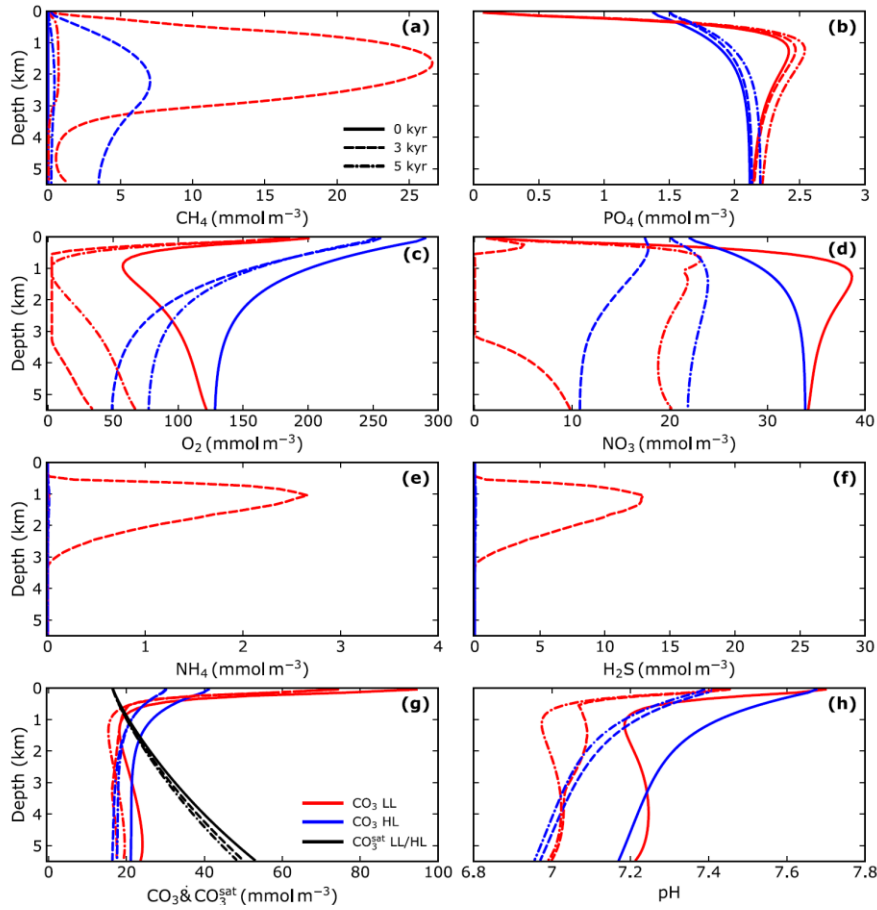
Figure 12: Low-mid and high latitude ocean property profiles at selected times for the simulation of Figure 11. Low-mid latitude (LL) and high latitude (HL) results are plotted with red and blue lines, respectively. Line types for the selected times after the simulation start (0, 3 and 5 kyr) are defined in pane **a**. CO_3^{sat} is the carbonate ion saturation concentration for calcite.
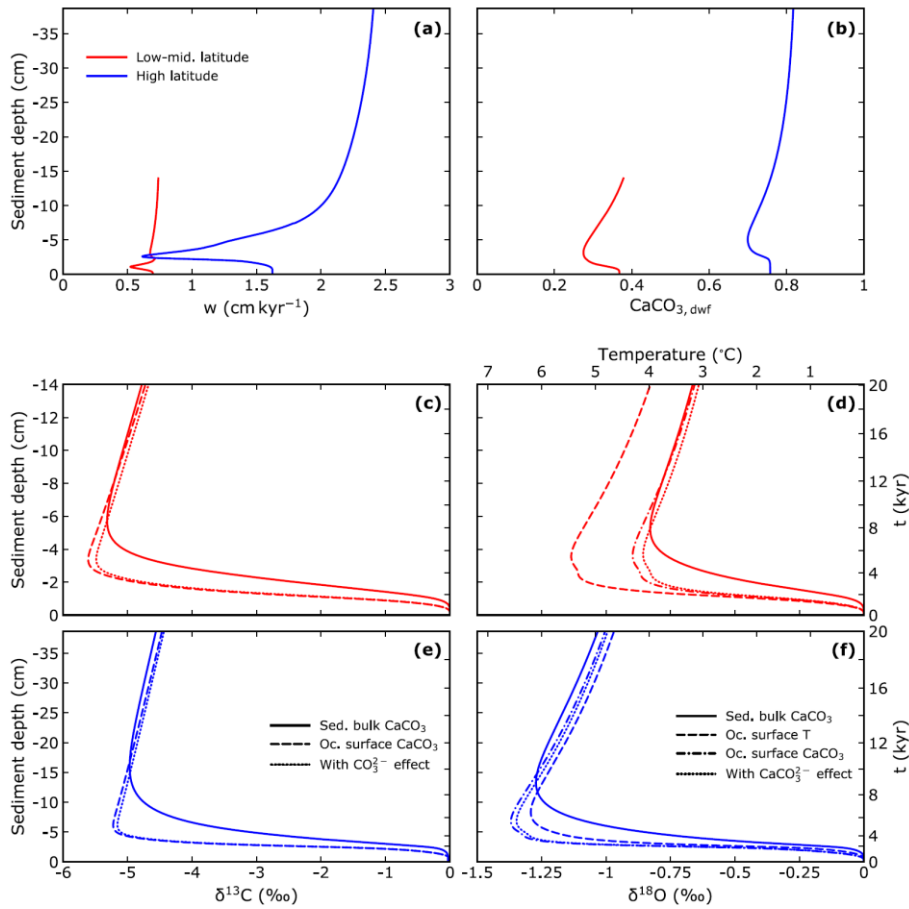
Figure 13: Comparison of bulk carbonate results from model synthetic ocean sediment cores at 3000 m depth with ocean surface layer properties for the simulation of Figure 11. Low-mid latitude (LL) and high latitude (HL) results are plotted with red and blue lines, respectively, and the methane input has $\delta^{13}C$ = -40 per mil. **a**) burial (downward) velocity relative to the base of the bioturbated sediment layer vs. synthetic sediment depth (SSD). SSD is referenced to zero at the start of the simulation and increases downcore by convention, **b**) carbonate dry weight fraction vs. SSD, **c**) LL bulk carbonate $\delta^{13}C$ excursion vs. LL SSD (solid line) . Also shown are the $\delta^{13}C$ excursion in biogenic carbonate in the LL ocean surface layer (dashed line) and this $\delta^{13}C$ including the correction for change in surface layer CO_3^{2-} (dotted line; Spero et al., 1997), both plotted vs. simulation time (right hand side y-axis). The time axis is "stretched/squeezed" relative to the SSD axis, a product of time-varying burial rates, **d**) LL bulk carbonate $\delta^{18}O$ vs. LL SSD (solid line). Also shown are 1. ocean surface layer temperature anomaly (dashed line) on a temperature scale (top x-axis) related to the oxygen isotope scale (bottom x-axis) using temperature-dependence of biogenic carbonate $\delta^{18}O$ (Bemis et al, 1998)), 2. $\delta^{18}O$ in biogenic carbonate produced in the LL ocean surface layer (dashed-dot line) that includes the water $\delta^{18}O$ excursion (see Figs 7-9 i) and 3. this $\delta^{18}O$ including the correction for change in surface layer CO_3^{2-} (dotted line; Spero et al., 1997), all plotted vs. simulation time, **e**) as **c** but for the HL ocean and sediment, **f**) as **d** but for the HL ocean and sediment.