Exploring the impact of Cenomanian paleogeography and marine gateways on oceanic oxygen

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Abstract

The Cenomanian-Turonian period recorded one of the largest disruptions to the oxygen and carbon cycles, the Oceanic Anoxic Event 2 (OAE2, 94 Ma). This event is global, yet paleo-reconstructions document heterogeneous ocean oxygenation states and sedimentary carbon contents, both temporally and spatially, suggesting that several mechanisms are at play. To better understand the long-term controls on oceanic oxygen and the initial oxygenation conditions prevailing at the beginning of OAE2, we perform numerical simulations of the Cenomanian using the IPSCL-CM5A2 Earth System Model, which includes a marine biogeochemistry component. We examine the control of the biogeochemical states of the global and Central Atlantic oceans by the depth of the Central American Seaway (CAS). The simulations show that a vigorous ocean circulation existed during the Cenomanian and that dysoxia/anoxia was caused by paleogeography rather than by ocean stagnation. The existence of restricted basins, disconnected from the deep global circulation and supplied with oxygen-depleted waters from Oxygen Minimum Zones of the surrounding basins, played a key role in the development of dysoxic/anoxic regions. A comparison with redox-proxy data suggests that a deep connection existed between the Pacific and Central Atlantic during OAE2. The paleogeographic configuration and that of gateways and submarine topographic barriers appear as major long-term controllers of the ocean as prerequisite conditions for OAEs to occur.

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14 KEY POINTS

- Cenomanian paleogeography appears as a major long-term controller to install
 prerequisite conditions for Oceanic Anoxic Event 2 to occur.
- A deep connection between the Pacific and Central Atlantic oceans is needed before
 OAE2 to explain the Central Atlantic redox data.
- The reconstructed Cenomanian oceanic circulation is vigorous but does not prevent
 large parts of the ocean to be anoxic.

24 ABSTRACT

The Cenomanian-Turonian period recorded one of the largest disruptions to the oxygen 26 27 and carbon cycles, the Oceanic Anoxic Event 2 (OAE2, 94 Ma). This event is global, yet 28 paleo-reconstructions document heterogeneous ocean oxygenation states and sedimentary 29 carbon contents, both temporally and spatially, suggesting that several mechanisms are at 30 play. To better understand the long-term controls on oceanic oxygen and the initial 31 oxygenation conditions prevailing at the beginning of OAE2, we perform numerical 32 simulations of the Cenomanian using the IPSCL-CM5A2 Earth System Model, which 33 includes a marine biogeochemistry component. We examine the control of the 34 biogeochemical states of the global and Central Atlantic oceans by the depth of the Central 35 American Seaway (CAS). The simulations show that a vigorous ocean circulation existed 36 during the Cenomanian and that dysoxia/anoxia was caused by paleogeography rather than by ocean stagnation. The existence of restricted basins, disconnected from the deep global 37 38 circulation and supplied with oxygen-depleted waters from Oxygen Minimum Zones of the 39 surrounding basins, played a key role in the development of dysoxic/anoxic regions. A 40 comparison with redox-proxy data suggests that a deep connection existed between the 41 Pacific and Central Atlantic prior to OAE2. A shallowing of the CAS may have contributed to 42 the establishment of enhanced anoxia in the Central Atlantic during OAE2. The 43 paleogeographic configuration and that of gateways and submarine topographic barriers 44 appear as major long-term controllers of the oceanic circulation and oxygen distribution, 45 leading to low-oxygen concentrations in extended parts of the ocean as prerequisite conditions 46 for OAEs to occur.

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1. Introduction

50 The Cenomanian-Turonian is a key period to study the oxygen and carbon cycles as it 51 recorded one of their greatest disturbance, known as the Oceanic Anoxic Event 2 (OAE2) and 52 characterized by the widespread deposition of organic-rich sediments (Arthur et al., 1987; 53 Jenkyns, 2010; Owens et al., 2018). A positive carbon isotopic excursion (CIE) associated 54 with OAE2 delimits the event in the lithological succession and shows that this event was 55 global and rapid (<1 Myr; Sageman et al., 2006; Jenkyns, 2010; Gangl et al., 2019; Jones et 56 al., 2019). However, the associated carbon enrichment is strongly variable, both spatially, 57 with content of total organic carbon (TOC) varying between 0 to 50% during OAE2 (Owens 58 et al., 2018), but also temporally, with sites documenting the onset of deposition of organic-59 rich sediments prior to OAE2, particularly in the Southern Central Atlantic (Kuypers et al., 2002; Montoya-Pino et al., 2010; Owens et al., 2012; Trabucho Alexandre et al., 2010; 60 61 Westermann et al., 2014).

Proxy data of redox sensitive trace elements also document heterogeneous 62 63 oxygenation state before and during OAE2 (Niels A. G. M. van Helmond et al., 2014), with 64 some areas exhibiting for example anoxia before OAE2. A diachronism between CIE and 65 carbon enrichment was also shown with highest TOC values prevailing before and after CIE, 66 but not during (i.e indicating oxic conditions during OAE2), suggesting a partial decoupling 67 of CIE and anoxia (Eldrett et al., 2014; Lowery et al., 2018). It has been suggested that the CIE was triggered by an abrupt and massive volcanism episode (Turgeon & Creaser, 2008; 68 69 Du Vivier et al., 2014), driving an increase in atmospheric pCO_2 (Barclay et al., 2010; Bice et 70 al., 2006) and an amplified hydrological cycle leading to increased nutrient inputs to the 71 ocean (Blättler et al., 2011; N. A.G.M. Van Helmond et al., 2015; Niels A.G.M. van Helmond 72 et al., 2014; Jenkyns et al., 2017; Nederbragt et al., 2004; Pogge Von Strandmann et al., 2013; 73 Ruvalcaba Baroni et al., 2014). Other studies have shown that the CIE was also modulated by 74 orbital controls determining the exact timing of its beginning and termination (Batenburg et al., 2016; Li et al., 2017; Mitchell et al., 2008; Wagner et al., 2004). The decoupling of CIE
and anoxia, as well as the spatial and temporal variations in carbon content and ocean redox
conditions suggest that other mechanisms, in the longer term, are at play and act on temporal
and spatial scales different from those suggested for CIE alone.

79 Unravelling the short-term controls on oxygenation and carbon burial during the 80 OAEs appears thus challenging without a good understanding of the long-term control on 81 oceanic oxygen and of the oxygenation conditions that existed at the beginning of the OAEs. 82 The Cretaceous ocean has long been considered stratified and stagnant, due to the coeval 83 greenhouse climate (Degens & Stoffers, 1976; Erbacher et al., 2001; Sinninghe Damste & 84 Koster, 1998). Such a sluggish circulation was invoked to explain sea-floor anoxia and black 85 shale deposition, but more recent observational and modelling studies have shown that a 86 vigorous oceanic circulation was in place, with an active ventilation of the deep-ocean 87 (Yannick Donnadieu et al., 2016; Ladant et al., 2020; MacLeod et al., 2008; Martin et al., 88 2012; Monteiro et al., 2012; Mourlot et al., 2018; Chris J Poulsen et al., 1998; Soares et al., 89 2014; Thiéblemont et al., 2020; Trabucho Alexandre et al., 2010; Du Vivier et al., 2014; 90 Zheng et al., 2013). Other controls, such as long-term changes in paleogeography and in 91 particular in the geometry of marine gateways, have thus been invoked to explain changes in 92 oceanic circulation and ocean deoxygenation (Yannick Donnadieu et al., 2016; W. Dummann 93 et al., 2020; Christopher J. Poulsen et al., 2001; Trabucho Alexandre et al., 2010). Dummann 94 et al. (2020b) have in particular demonstrated that the shallow Falkland Plateau and Maurice 95 Ewing Bank exerted a first-order control on ocean anoxia during the OAE1 in the Southern 96 Atlantic, by providing favorable conditions without which organic carbon burial would not be 97 possible during OAE1 (Wolf Dummann et al., 2020).

98 During the Cenomanian-Turonian, the Central Atlantic basin, from where numerous 99 data document OAE2, is surrounded by five gateways, namely the Equatorial Atlantic 100 Gateway (EAG) in the South, the CAS in the West, the Western Interior Seaway (WIS) in the 101 North-West, the East Greenland Seaway (EGS) in the North-East and the Tethys Seaway in 102 the East (Fig. 1). The paleodepths of the WIS, EGS and EAG are relatively well-constrained, 103 with very shallow bathymetry for the WIS and EGS (<200 meters; Martinson et al., 1998; 104 Gernigon et al., 2020) and shallow bathymetry for the EAG (<800 meters; Ye et al., 2017). 105 The Tethys Seaway paleodepth is more uncertain (Nouri et al., 2016), as well as the 106 paleodepth of the CAS that is suggested to vary concomitantly to OAE2 due to the Caribbean 107 Large Igneous Province (CLIP) activity and the formation of the Caribbean Plateau (Buchs et 108 al., 2018). There is some evidence supporting a causal relationship between the intense

109 volcanic activity associated with the CLIP development in the Late Cenomanian and the 110 OAE2 (Joo et al., 2020; Turgeon & Creaser, 2008). Recent work also demonstrated that the 111 depth of the CAS can exert a significant role on the oceanic circulation during the Cretaceous 112 (Y. Donnadieu et al., 2006; Yannick Donnadieu et al., 2016; Ladant et al., 2020; Monteiro et 113 al., 2012; Topper et al., 2011; Trabucho Alexandre et al., 2010) and, consequently, can 114 strongly impact the distribution of dissolved oxygen in the ocean (Kerr & Kerr, 1998) and 115 affect rates of organic matter preservation (Monteiro et al., 2012). Although it is unlikely that 116 changes in gateway bathymetry are directly responsible for triggering OAE2 because they act 117 on longer timescales, changes in the CAS depth and geometry on the runup to OAE2 may 118 have preconditioned and/or contributed to the establishment of anoxic conditions in the deep 119 Central Atlantic Ocean through a reorganization of deep-ocean currents. To date, however, 120 the dynamics of oceanic oxygen during the Cenomanian and OAE2 has only been 121 investigated using box models (Flögel et al., 2011; Ruvalcaba Baroni et al., 2014) and Earth System Models of Intermediate Complexity (EMIC; Monteiro et al., 2012). 122

123 Here, we examine the sensitivity of oceanic dissolved oxygen to a mid-Cretaceous 124 paleogeography and to the CAS geometry using numerical simulations of the Cenomanian 125 ocean with the IPSL-CM5A2 Earth System Model (ESM) (Sepulchre et al., 2020), which 126 includes the marine biogeochemistry model PISCES (Aumont et al., 2015). We first analyze 127 the long-term impact of a baseline Cenomanian-Turonian paleogeography (with the CAS 128 open to deep circulation) on oceanic circulation and oxygen at the global scale. In a second 129 step, we test the impact of changes in ocean dynamics on dissolved oxygen by performing 130 two sensitivity experiments with intermediate and shallow CAS configurations. Results are 131 focused on the Central Atlantic to investigate the consequences on dissolved oxygen of 132 oceanic circulation changes arising from the different gateway geometries. Finally, the 133 comparison with redox proxy data provides a better understanding of the long-term 134 paleogeographic control on oceanic oxygen during the Cenomanian, as well as an estimate of 135 the pre-OAE2 oxygenation state, which will serve as a basis for further unraveling the 136 triggering mechanisms of OAE2 and the associated strong carbon burial.

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2. Model description and experimental design

- 140 2.1 IPSL-CM5A2 Earth System Model
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142 The IPSL-CM5A2 Earth System Model (Sepulchre et al., 2020) is an updated version 143 of IPSL-CM5A-LR developed at IPSL (Institut Pierre-Simon Laplace) within the CMIP5 144 framework (Dufresne et al., 2013). IPSL-CM5A2 is composed of the LMDZ atmospheric 145 model (Hourdin et al., 2013), the ORCHIDEE land surface and vegetation model (including 146 the continental hydrological cycle, vegetation and the land carbon cycle; Krinner et al., 147 2005), and the NEMO ocean model (Madec and the NEMO Team, 2008), including the LIM2 148 sea-ice model (Fichefet & Maqueda, 1997) and the PISCES-v2 marine biogeochemistry model (Aumont et al., 2015). The ocean and atmospheric components are synchronized via 149 150 the OASIS coupler (Valcke et al., 2006) and the XIOS input/output parallel library is used to 151 read and write data. The atmospheric grid has a horizontal resolution of 96x95, (equivalent to 152 3.75° in longitude and 1.875° in latitude) and 39 uneven vertical levels. ORCHIDEE shares 153 the same horizontal resolution, whereas the oceanic grid has 31 uneven vertical levels (from 154 10m width at the surface to 500m at the bottom) and a nominal horizontal resolution of 2° , 155 enhanced to up to 0.5° in latitude in the 20°S-20°N latitudinal band. NEMO uses a tripolar 156 grid to overcome the North Pole singularity (Madec & Imbard, 1996). For a more detailed 157 description of the model and its different components, see Sepulchre et al. (2020).

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2.2 PISCES marine biogeochemistry model

161 The PISCES model (Pelagic Interactions Scheme for Carbon and Ecosystem Studies, 162 Aumont et al., 2015) simulates the lower trophic levels of marine ecosystems 163 (nanophytoplankton, diatoms, microzooplankton and mesozooplankton), carbonate chemistry 164 and the biogeochemical cycles of carbon, oxygen, and of the main nutrients: phosphorus, 165 nitrogen, iron and silica. In PISCES, dissolved oxygen is produced in the ocean according to 166 phytoplankton net primary production and consumed by zooplankton heterotrophic 167 respiration, oxic remineralization of organic matter and nitrification (see Aumont et al. [2015] 168 for detailed equations of all these terms). Dissolved oxygen is also exchanged at the air-sea interface using the parameterization of Wanninkhof (1992). Note that the atmospheric 169 170 concentration of di-oxygen is set to a fixed ratio of 0.21. As in Bopp et al. (2017), we 171 decompose dissolved oxygen into the O₂ saturation (O₂sat) and Apparent Oxygen Utilization 172 (AOU) components. O₂sat only depends on seawater temperature and salinity. At the surface 173 of the ocean, the concentration of dissolved oxygen is very close to its saturation state, such 174 that $[O_2] = O_2$ sat. AOU represents the quantity of oxygen that has been consumed by 175 biogeochemical processes on the trajectory of a given water-mass from the ocean's surface,

and thus integrates both a biological and a ventilation signature. The three terms are linked as follows : $[O_2] = O_2 \text{sat} - \text{AOU}$ (Bopp et al., 2017). We also used an additional inert artificial tracer allowing to compute the age of water masses, corresponding to the time spent since last surface contact (Bopp et al., 2017).

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2.3 Experimental design

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2.3.1 Boundary and Initial conditions

184 The baseline and sensitivity simulations share the Cenomanian-Turonian (CT) land-185 sea configuration (Fig.1) of Sewall et al. (2007), in which the bathymetry from Müller et al. 186 (2008) is implemented, in order to represent deep-oceanic topographic features, such as 187 ridges, that are absent from the Sewall paleogeographic configuration. The pCO_2 188 concentration is fixed at 1120 ppm, i.e. 4 times the pre-industrial atmospheric level (PAL), 189 which is a value suggested by CT pCO_2 reconstructions (e.g., Wang et al., 2014), and which 190 has been shown to reasonably reproduce the mean CT climate in IPSL-CM5A2 simulations 191 (Laugié et al., 2020). Vegetation is assigned along latitudinal bands as described in Laugié et 192 al. (2020) and a mean soil color and texture is prescribed uniformly to all continents. Polar ice 193 sheets are removed and replaced by a brown bare soil after adjusting the topography to 194 account for isostatic rebound. The solar constant is reduced to a CT value of 1353.36 W.m⁻² 195 (Gough, 1981). Orbital parameters are kept at their modern value. The global budgets of all 196 macro-nutrients (nitrate, phosphate and silicate) in the oceans are also kept identical to the 197 modern ones. We deliberately choose to keep these budgets constant in order to isolate the 198 effect of ocean dynamics on the intensity of ocean anoxia while mimicking the tectonic drift 199 of the CLIP across the Central American seaways by shallowing the depth of the gateway.

200 The three simulations present different paleobathymetric configurations for the Central 201 American Seaway (CAS; Fig. 1). The baseline DeepCAS simulation exhibits a deep 202 connection (~ 4000 m depth) between the Pacific and Central Atlantic (Fig. 1). The CAS 203 depth is reduced to intermediate (~ 2500 m) and shallow depths (~ 300 m) in the 204 IntermediateCAS and ShallowCAS simulations, respectively (Fig. 1). All other boundary 205 conditions are identical for the three simulations. The simulations are initialized with warm 206 idealized conditions adapted from those described in Lunt et al. (2017). The constant initial 207 salinity field is set to 34.7 PSU, and ocean temperatures are initialized with the depth-208 dependent distribution of

209 Laugié et al. (2020).

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212 2.3.2 Equilibrium state

214 The baseline DeepCAS simulation is first run for 3000 years with IPSL-CM5A2. 215 From the end state of this first simulation, we branch the two CAS sensitivity simulations and 216 integrate them for 3500 years. The baseline simulation is also extended for another 3500 217 years. Finally, because the biogeochemical equilibrium of the ocean typically requires a 218 longer spin-up (Séférian et al., 2016), we extend each simulation over an additional 3000-year 219 period by performing "offline" PISCES-only simulations. In those offline simulations, 220 PISCES is forced by using the last 100-year climatology (e.g., monthly means of ocean 221 temperature, salinity, currents, incoming radiation,) from each of the 3 coupled climate 222 simulations. This strategy allows deep-ocean biogeochemistry, and in particular seafloor 223 oxygen concentrations, to reach a final state closer to equilibrium. Climate and 224 biogeochemical results are discussed using climatological averages calculated over the last 225 100 years of IPSL-CM5A2 and PISCES integrations, respectively.

The equilibrium for the three IPSL-CM5A2 simulations is assessed using surface and deep oceanic temperatures (Supplementary Fig.S1a,b) and the intensity of the global meridional stream function (Supplementary Fig.S1c). Both surface and deep temperatures have reached a near equilibrium state with temporal drifts estimated at less than 0.05°C/century (Supplementary Table 1) during the last 1000 years of the climate model integration. The meridional streamfunction is also well equilibrated with stable maximum intensity of around 17 Sv during the last 2000 years of the simulation.

Furthermore, the three PISCES simulations exhibit negligible drifts in deep ocean oxygen concentrations (Supplementary Fig.1d) after 3000 years of offline simulations (Supplementary Table 1; <0.1 mmol/m³/century).

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237 2.4 Redox data

The simulated results are compared to reconstructions of oceanic oxygen state inferred from pre-OAE2 and OAE2 data. Our data compilation is built from the databases of Monteiro et al. (2012) and Owens et al. (2018). The oxygenation states documented in these two databases are essentially inferred from TOC values, which are a function of bottom water oxygen, but are also influenced by other parameters such as the organic matter flux to the seafloor, the dilution with inorganic matter or diagenetic processes. In our compilation, we thus selected only data points for which other indicators than TOC were available, such as analysis of redox sensitive trace elements or presence/absence of benthic biota, which are
direct indicators of bottom water oxygenation. We also chose to only select data for which an
estimation of the paleowater depth exists in order to be comparable with our model results.
The data points are distributed to their paleolocation using the GPlates software (Qin et al.,
2012) and the global plate rotation model of Scotese (2016). In case of an inconsistency
between the calculated paleoposition and the paleowater depth, data points are then moved to
the nearest location with a correct paleodepth in the model (See Supplementary information).

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3. Results

In a first step, we describe the major patterns of ocean circulation and oxygen distribution at the global scale, using the *DeepCAS* simulation. Results from the *IntermediateCAS* and *ShallowCAS* simulations are very similar to DeepCAS at the global scale, with the change in CAS depth mostly affecting the Central Atlantic. In a second step, we focus on the Central Atlantic and analyze the regional impact of the CAS on ocean circulation and oxygen distribution in the three simulations.

3.1 Ocean circulation and oxygen distribution in the global ocean

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3.1.1 Upper and intermediate ocean

266 Upper ocean waters (0-100 meters) are generally close to dissolved oxygen saturation, 267 which means that oxygen concentrations are mainly determined by surface temperature and 268 its influence on oxygen solubility (Fig. 2a). The Cenomanian upper ocean is characterized by 269 elevated temperatures and a reduced equator-to-pole gradient (Laugié et al., 2020; Norris et 270 al., 2002; O'Brien et al., 2017; Robinson et al., 2019; Tabor et al., 2016). In the tropical area, O_2 saturation is reduced to an average of 198 mmol/m³ because oxygen solubility decreases 271 272 with temperature. In comparison, the mean temperature in the tropical ocean in the pre-273 industrial simulation of Laugié et al. (2020) is 22.2° C, with a corresponding O₂ saturation at 228 mmol/ m^3 . In the high latitudes, O2 saturation reaches higher values (~300 mmol/ m^3) than 274 in the low latitudes but, again, with a large offset compared to preindustrial high-latitude 275 276 saturation levels (~370 mmol/m³). Subsurface oxygen concentrations (< 600 meters of water 277 depth) also exhibit a latitudinal gradient (Fig. 2b) but they are more impacted by ocean 278 dynamics, in particular in regions of strong vertical currents (e.g., equatorial upwelling, 279 Fig. 2b). Specifically, in the Central Atlantic and Eastern Equatorial Pacific, the model 280 simulates complete anoxia in the subsurface and up to the base of the photic zone. At

intermediate depths (600-1600 meters of water depth), oxygen concentration is globally more
homogeneous in the ocean except in the South Pacific and Central Atlantic (Fig. 2c). South
Pacific waters remain well-oxygenated from the surface to the bottom because this region is
the source of deep-water in our Cenomanian simulations. In contrast, the Central Atlantic is
bathed by oxygen-depleted waters (Fig. 2c), whose origin is described in more details in
Section 3.2.

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288 3.1.2 Deep Ocean

289 The deep-sea oxygen distribution is controlled at first order by the location of deep-water 290 formation sites and by the bathymetry (Fig. 3a). In our simulations, deep-water formation 291 sites are only located in the South Pacific Ocean, although intermediate-water formation takes 292 place in the North Pacific Ocean (Fig. 2c). The South Pacific deep convection feeds a global 293 meridional overturning circulation (MOC) of maximum intensity at ~18 Sv, similar to the 294 modern Atlantic MOC (18 Sv; Talley et al., 2003). Note that the intensity of the MOC appears 295 only weakly sensitive to the depth of the CAS (Supplementary Fig.S2). The simulated 296 Cenomanian MOC reaches the abyssal ocean and suggests that the Cenomanian global 297 circulation was vigorous, contrary to the well-anchored image of a sluggish ocean but in 298 agreement with results from other recent model simulations (Yannick Donnadieu et al., 2016; 299 Ladant et al., 2020; Trabucho Alexandre et al., 2010).

300 From their origin in deep-water formation sites in the South Pacific, deep waters flow 301 northward into the Neotethys and Equatorial Pacific Oceans (Fig.3a). The deep Neotethys 302 waters then circulate back to the Pacific to flow north of the mid-Pacific ridge and toward the 303 North-Eastern Pacific. Pacific deep waters ultimately end up in the Central Atlantic whereas 304 the deep South Atlantic is restricted from the global circulation by four submarine 305 topographic barriers that are the Kerguelen Plateau, the Davie Ridge, the Equatorial Atlantic 306 Gateway and the Drake passage (Fig. 3a). This pattern of deep circulation leads to interbasin 307 gradients in deep ocean oxygen concentrations and water mass age (Fig. 3b); the degree of 308 oxygenation of deep waters appearing well correlated with their age (Fig.4; Supplementary 309 Fig.S3). The younger and most oxygenated waters are indeed found in the South Pacific and 310 in the Neotethys Ocean. The South Atlantic and North-Eastern Pacific exhibit reduced 311 oxygenation and older water masses, and the Central Atlantic is the oldest and most oxygen-312 depleted deep basin. The disconnection of the South Atlantic from the global deep circulation 313 also explains the presence of old and oxygen-depleted water masses in this basin (Fig. 3b).

Interestingly, a similarly restricted deep circulation, confined to the Pacific and Neotethys Oceans due to bathymetric barriers, is also simulated in a Cenomanian simulation with the CCSM4 ESM (Ladant et al., 2020), suggesting that submarine topographic barriers are major controllers of the deep circulation and thus of dissolved oxygen in the deep ocean.

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3.1.3 Relationship between oxygen and marine productivity

In our simulations, there is an obvious correlation between ocean dynamics and deep ocean oxygenation (Fig. 4), indicating that older waters are the most oxygen-depleted ones. There is not such a clear correlation between marine productivity and deep oxygen concentrations, but interesting insights can be drawn from oxygen-productivity cross plots calculated in different regions of the ocean (shown on Fig. 4b) with distinct oxygen dynamics in the water column (Fig. 5).

The meridional surface oxygen gradient is evident from the vertical oxygen profiles (Fig. 5a). High latitudes locations (Profiles 1 and 3) exhibit high oxygen levels in the surface due to low temperatures whereas lower-latitude profiles show lower surface values (Profiles 2,4,5,6).

332 Profile 1 is located in a deep-water formation area, with a low marine productivity. Deep-333 convection brings oxygen down to seafloor, and high oxygen values are found throughout the 334 whole water column (Fig. 5a). In contrast, albeit at the same latitude but in the Northern 335 Hemisphere, the Profile 3 is located close to a region of intermediate water formation with 336 seasonal primary production (Fig. 5b,c). Limited convection and slightly higher productivity 337 values impose a strong vertical gradient in oxygen between 300 and 1000 m and reduces 338 intermediate to deep oxygen levels. Nonetheless, these two regions can be viewed as "low 339 productivity - high oxygen" areas.

340 Profile 2 and 6 are located in the subtropics in low productivity areas (Fig.5c). Marine 341 productivity is low because of poor nutrient availability, which limits carbon export and 342 oxygen consumption at depth. However, elevated upper ocean temperatures reducing O_2 343 saturation leads to moderate surface and intermediate oxygen levels ("low productivity – low 344 oxygen" region). In the deep ocean, those 2 oxygen profiles display a divergent trend. Profile 345 2 is located directly north of the deep-water formation area, from where oxygenated waters 346 circulate. Oxygen concentrations in Profile 2, therefore, increase below ~ 2300 m (Fig. 5a). 347 Profile 6 is instead located in the South Atlantic away from well-oxygenated deep-water sources, which explains the continuous decrease of oxygen concentrations with depth (Fig.5a).

350 In contrast to the profiles discussed above, which are located in open ocean sectors of 351 the Pacific and South Atlantic Oceans, Profiles 4 and 5 are more representative of coastal 352 settings of the Central Atlantic basin. At these two locations, marine productivity is very high 353 and leads to an intense export of organic matter and high oxygen utilization at depth. Hence, 354 intermediate and deep ocean oxygen concentrations fall below the hypoxic threshold and, for 355 Profile 5, even below the anoxia threshold ("high productivity – low oxygen" regions). The 356 differences between both profiles are explained by vertical ocean dynamics (Supplementary 357 Fig. S4). Profile 5 is indeed located in a major upwelling area whereas the site of Profile 4 358 experiences seasonal deepening of the mixed-layer and is thus a sinking area for upper ocean 359 waters. Waters remain oxic only above 100 m in Profile 5 because upwelling supplies 360 nutrient-rich and oxygen-depleted waters to the surface (Supplementary Fig.S5), which fuels 361 productivity and enhances oxygen depletion. At the location of Profile 4, deep coastal 362 upwellings also allow nutrient-rich waters to reach the photic zone but the seasonal MLD 363 deepening carries dissolved oxygen down the water column, which increases the 364 remineralization potential. Increased remineralization increases the nutrient supply and boosts 365 marine primary production to levels higher than at Profile 5. Deeper in the water column, both 366 sites are bathed by intermediate and deep waters originating from the Pacific and exhibit 367 similar oxygen concentrations. However, these last findings about Profiles 4 and 5, and the 368 oxygen dynamics in the Central Atlantic more generally, are meaningful in a context of CAS 369 open to deep circulation but are significantly altered with a shallower CAS configuration, as 370 we now examine.

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372 3.2 Central Atlantic circulation

We now focus on the Central Atlantic circulation and distinguish three depth intervals: upper ocean circulation (<600 meters of water depth), intermediate circulation (600 – 1600 meters of water depth) and deep circulation (> 1600 of water depth). Inflow and ouflow water fluxes are computed along the five gateways surrounding the Central Atlantic Ocean: the CAS, the EAG, the WIS, the EGS and the Tethys Seaway (Fig.1).

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3.2.1 Upper ocean circulation (0-600 meters)

Fig.6 depicts the upper ocean circulation of the Central Atlantic, showing the directions of major currents in and out of the Central Atlantic (Fig.6a) and the corresponding 382 water fluxes (Fig.6b). Most of the horizontal circulation is dominated by a westward current, 383 the circum-equatorial current, flowing through the Central Atlantic from the Tethys Seaway 384 (main inflow) to the CAS (main outflow). Similar patterns of upper ocean circulation were 385 simulated in previous modeling studies (Yannick Donnadieu et al., 2016; Topper et al., 2011; 386 Trabucho Alexandre et al., 2010). Minor counter-currents also flow into the Central Atlantic 387 across the CAS and the EAG. The circum-equatorial upper ocean current separates the basin 388 into two distinct areas: an upwelling area in the southern Central Atlantic, strongly influenced 389 by underlying older waters, and a downwelling area in the northern Central Atlantic, in which 390 the thermocline deepens with mixed layer depth reaching 350 m. High marine productivity is 391 simulated in these areas of strong water mixing regardless of the CAS configuration (See 392 Supplementary Fig.S6). In contrast, though the upper ocean circulation is very similar, the 393 intensity of water fluxes through the CAS is smaller for the ShallowCAS simulation, due to 394 the reduced water depth (Fig.6b). Incoming and outgoing fluxes are approximatively halved 395 in the *ShallowCAS* simulation compared to the *IntermediateCAS* and *DeepCAS* simulations.

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3.2.2 Intermediate circulation (600-1600 meters)

398 The intermediate currents system in the Central Atlantic is similar in the DeepCAS and 399 IntermediateCAS simulations (Fig. 7a and 7c). The largest incoming water flux comes from 400 the Pacific through the CAS in the deeper intermediate layers. This estuarine circulation 401 between the Pacific and the Central Atlantic is comparable to that suggested by previous 402 model studies (Yannick Donnadieu et al., 2016; Monteiro et al., 2012; Topper et al., 2011; 403 Trabucho Alexandre et al., 2010) and explains the latitudinal gradient in water mass 404 properties in the Central Atlantic (Supplementary Fig.S7). The southern part of the basin is 405 under the influence of both intermediate waters coming from the eastern Equatorial Pacific 406 and upwelled waters from the deep Central Atlantic (Fig.7 and Supplementary Fig.S7). As a 407 result, the intermediate southern Central Atlantic is bathed by colder, fresher and older waters 408 than in the northern Central Atlantic, whose intermediate water mass composition includes a 409 greater proportion of warmer, saltier and younger waters originating from the Neotethys and 410 from sinking waters (Supplementary Fig.S7).

411 On the contrary, the *ShallowCAS* simulation exhibits a different intermediate circulation 412 (Figs. 7b and 7d) because the depth of the CAS prevents intermediate currents to flow across 413 it. The inflow water transport from the Neotethys more than doubles compared to the 414 DeepCAS and IntermediateCAS, and intermediate waters originating from the EAG start 415 flowing into the Central Atlantic. Without the influence of Pacific intermediate and deep 416 waters, Central Atlantic intermediate waters are then globally younger and warmer in the
417 *ShallowCAS* simulation than they are in the two other simulations (Supplementary Fig.S7).

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419 3.2.3 Deep circulation (> 1600 meters)

420 The deep circulation is different for the three CAS configurations (Fig. 8). The 421 DeepCAS and IntermediateCAS simulations exhibit water exchange only through the CAS 422 because the depth of the other gateways is less than 1600 meters. The deep Central Atlantic is 423 thus bathed by deep and cold waters from the Pacific. The IntermediateCAS configuration 424 (2500 m depth maximum) limits inflows and outflows to half that of the DeepCAS 425 simulation, and prevents the deepest waters (below 2500 meters) of the Pacific from entering 426 the Central Atlantic. In contrast, in the ShallowCAS configuration, each gateway enclosing 427 the Central Atlantic is closed to deep circulation. The deep Central Atlantic is instead supplied 428 with waters from intermediate levels. In particular, warm intermediate waters coming from 429 the EAG represent a significant contribution to deep Central Atlantic waters, as indicated by 430 the water age (Fig. 9). As a consequence, the deep Central Atlantic presents much warmer 431 waters compared to the two previous simulations (13.6°C vs 8.7°C and 9°C for DeepCAS and 432 IntermediateCAS simulations, respectively).

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3.3 Central Atlantic oxygenation

435 The changes in ocean circulation with CAS configuration in the Central Atlantic leads 436 to different oxygenation states at the seafloor (Fig. 10). Regions of intermediate and deep 437 depths present a decreasing oxygen content with the shallowing of the CAS. They evolve 438 from a hypoxic or oxic state in the *DeepCAS* simulation, to an anoxic or hypoxic state in the 439 IntermediateCAS simulation and to complete anoxia in the ShallowCAS simulation (oxic = $[O_2] > 62.5 \text{ mmol/m}^3$; hypoxic = $6.5 < [O_2] < 62.5 \text{ mmol/m}^3$; anoxic = $[O_2] < 6.5 \text{ mmol/m}^3$). 440 This oxygen-depletion trend is directly linked to the origin of waters bathing the intermediate 441 442 and deep parts of the basin (Fig. 10 and Supplementary Fig.S8). With a deep CAS, the Central 443 Atlantic is supplied with well-oxygenated deep waters from the Pacific. With an intermediate 444 CAS, the Central Atlantic is supplied with more hypoxic intermediate waters from the Pacific. 445 Finally, with a shallow CAS, the Central Atlantic is mostly supplied with anoxic shallow waters from the Equatorial Atlantic Gateway. The CAS, therefore, acts as a submarine 446 447 topographic barrier for both currents and oxygen and, if shallow enough, can be responsible 448 for complete marine anoxia at a basin-scale despite the existence of an active circulation.

450 4. Discussion

452 Our compilation of redox data (See supplementary data) indicates the ocean oxygenation 453 state before (Fig. 11a) and during OAE2 (Fig. 11b), allowing identifying heterogeneous 454 conditions both spatially and temporally. Before OAE2, most of the ocean was oxic, except 455 some areas located in the Central Atlantic (Fig.11a, Southern Central Atlantic, Southern WIS, 456 deepest part of Gulf of Mexico). During OAE2 (Fig.11b), redox data indicates an expansion 457 of the anoxia to the whole Central Atlantic, with some locations showing fluctuations from 458 oxic to hypoxic/anoxic conditions (e.g. in the WIS, Northern Central Atlantic, Moroccan 459 margin). Outside the Central Atlantic, the hypoxia/anoxia extends to the Southern Atlantic 460 and to the Neotethys (deep outer shelves or even some parts of the deep basin), with 461 fluctuations between oxic and anoxic conditions also reconstructed in some locations. In the 462 easternmost part of the South Neotethys (e.g North Indian, Kerguelen plateau) as well as in 463 the Northern Pacific coasts, hypoxia is sometimes reached during OAE2, but in alternation 464 with oxic conditions. Finally, the Southern Pacific always stays oxic.

In the following, we compare our modelling results to these oxygenation trends for the pre-OAE and OAE2 periods, in a first step at the global scale, and in a second step with a focus on the Central Atlantic, in order to investigate the question of the CAS paleobathymetry.

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1.1 The pre-OAE2 oxygenation state: a data/model comparison

472 Outside the Central Atlantic, the results predicted by our three Cenomanian 473 simulations agree well with reconstructions of oceanic oxygen state inferred from data for 474 pre-OAE2, as illustrated in the DeepCAS simulation (Fig. 11a). Most of the sites suggest 475 sedimentary burial in oxic conditions and the model consistently reproduces a well-476 oxygenated seafloor at the site locations. Simulated dissolved oxygen concentrations show 477 that these oxic environments exhibit a large range of values, from 62.5 mmol/m³ (dysoxia limit) to more than 300 mmol/m³ but, unfortunately, proxies only discriminate anoxic/dysoxic 478 479 from oxic conditions and rarely allow to reconstruct the absolute oxygen concentration in oxic 480 environments. However, it is interesting that all sites that shift from an oxic to a hypoxic or 481 anoxic environment during OAE2 (Fig 11.b) are located in areas where the model simulates 482 moderate oxygenation for the pre-OAE2 (e.g., Southern Atlantic, Tethyan continental slopes, North-Eastern Africa, $[O_2] \sim 120 \text{ mmol/m}^3$) driven by basin restriction and/or reduced O_2 483

484 saturation due to elevated temperatures. Other sites that remain oxic or dysoxic during OAE2 485 are found in more-oxygenated areas during pre-OAE2 (North-Eastern/North-Western-Central Pacific, shallow Tethyan and Tibetan platforms, Kerguelen plateau, $[O_2] > 150 \text{ mmol/m}^3$). 486 487 Finally, the New Zealand site is interesting in that neither the model nor proxy records 488 document any major change in oxygen concentrations during OAE2, which agrees with the 489 site location in an area of deep-water formation ensuring high simulated oxygen values (New-490 Zealand, $[O_2] \sim 250 \text{ mmol/m3}$). One site in the Neotethys doesn't follow this trend with data 491 documenting anoxic conditions at the location of which the model simulates elevated oxygen 492 concentrations (Fig.11). This inconsistency could be due to the complex paleogeography of 493 the western Neotethys that is compartmented in numerous restricted deep basins separated by 494 shallow platforms (Golonka et al., 2000; Nouri et al., 2016). Anoxia could locally exist in 495 such basins but this degree of complexity is not represented at the spatial resolution of our 496 model.

497 Seafloor oxygen concentrations in the Central Atlantic basin are, in contrast, highly 498 sensitive to the configuration of the CAS and a specific comparison to data is made in section 499 4.3 for the Central Atlantic. For areas where the oxygen level is relatively insensitive to CAS 500 depths, i.e. the southern Central Atlantic and the northern Gulf of Mexico, the model 501 reproduces the dysoxic and anoxic areas inferred from proxy data for the pre-OAE2 interval 502 (Figs. 10 and 11a). Finally, the WIS represents another location of model-data mismatch in 503 the three simulations because the model suggests oxygenated conditions at odds with 504 observations of anoxia at this site (2 dots, see Fig.11a). However, the oxygenation state has 505 been shown to be related to sea-level variations in this region (Lowery et al., 2018), which are 506 not taken into account in the simulations and may provide an explanation to the model/data 507 discrepancy.

508 At the global scale, the spatial patterns of simulated seafloor oxygen are consistent 509 with the modelling study of Monteiro et al. (2012) that uses the GENIE EMIC. The most 510 severe anoxic regions are the Central/Equatorial Atlantic and dysoxia is simulated in the 511 Neotethys and Eastern Pacific oceans whereas the Southern Pacific remains more oxygenated. 512 The seafloor is however globally less oxygenated in Monteiro et al. (2012), likely due to a 513 less intense overturning circulation (around 6 Sv in the intermediate ocean vs. 18 Sv in our 514 study). Another major difference is observed in the South Atlantic where the model of 515 Monteiro et al. (2012) simulates higher oxygen concentrations, probably due to deep-water 516 formation in this area, which is absent in our model. This inconsistency is hard to resolve at 517 the moment as there is no data available in this region for the Cenomanian. From a physical

518 perspective, Donnadieu et al. (2016) also suggested weak deep-water formation in the South 519 Atlantic using the coarse-resolution FOAM global climate model. On the contrary, the more 520 complex CCSM4 ESM Cenomanian simulation of Ladant et al. (2020) does not produce 521 deep-water formation in the South Atlantic, indicating that these inconsistencies could be a 522 problem of model complexity and resolution, and/or employed paleogeography.

523 Our results illustrate a long-term control of paleogeography on oceanic oxygen beyond 524 OAE2 shorter-term triggers. For example, the anoxia located in the Southern Central and 525 Equatorial Atlantic is a robust feature of our and previous simulations (Monteiro et al. 2012) 526 under pre-OAE2 conditions and is also inferred from pre-OAE2 data, suggesting that this 527 localized anoxia is driven by paleogeography and is decoupled from the specific triggers of 528 the OAE2. In addition, regions where anoxia spreads over during OAE2 were those already 529 characterized by reduced oxygenation in our model. Our results therefore suggest a significant 530 preconditioning of the global ocean oxygenation on the runup to the OAE2 driven by 531 paleogeographic setting of the Cenomanian. More generally, we can hypothesize that ocean 532 preconditioning is required for OAEs to occur (See also Song et al., 2019) and that shorter-533 term controllers (such as an abrupt rise of pCO_2 or an increase in nutrient supply) would then 534 turn preconditioned areas into hypoxic or anoxic environments (See also Dummann et al., 535 2020a).

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1.2 Constraining Oceanic Gateways paleobathymetry.

539 The CLIP is estimated to have mainly formed between 95 and 83 Ma in the Pacific Ocean 540 westward of the CAS and to have moved through the CAS during the Late Cretaceous (Andjić et al., 2019; Dürkefälden et al., 2019; Loewen et al., 2013; Romito & Mann, 2020), 541 542 thereby shallowing the connection between the Central Atlantic and Pacific oceans (Buchs et 543 al., 2018). Numerous evidence suggest a significant role for the CLIP volcanism in the 544 initiation of OAE2 (Joo et al., 2020; Kerr & Kerr, 1998; Turgeon & Creaser, 2008). However, 545 the timing of its formation and progressive eastward movement is not well constrained. This 546 complexity limits inferences about the potential contribution of a CLIP-induced shallowing of 547 the CAS to OAE2. In an attempt to better constrain the CAS paleobathymetry during the Late 548 Cenomanian, we compare the study sites of the Central Atlantic to our three scenarios of 549 seafloor oxygen simulated for the different CAS configurations (Deep, Intermediate, Shallow; 550 Fig.10).

551 Seven sites document anoxic conditions before OAE2. They are located in the Southern 552 Central Atlantic and in the Gulf of Mexico (Fig 10; Perez-Infante et al., 1996; Kuypers et al., 553 2002; Owens et al., 2012; van Helmond et al., 2014b; Westermann et al., 2014; Lowery et al., 554 2017). In particular, previous studies suggested that the anoxic area surrounding the 555 Equatorial Atlantic Gateway was driven by the estuarine circulation established between the 556 Pacific and Central Atlantic (Topper et al., 2011; Trabucho Alexandre et al., 2010) but our 557 results demonstrate that this anoxic area is stable even in the ShallowCAS simulation despite 558 the disappearance of the estuarine circulation. Our findings imply that this anoxic zone would 559 rather be driven by the global paleogeography than regionally controlled by the CAS. In fact, 560 all these pre-OAE2 anoxic sites are located in regions simulated as hypoxic to anoxic 561 regardless of the CAS configuration and, thus, a comparison between our simulations does not 562 help constraining the CAS depth.

563 In the deep Central Atlantic, five sites record oxic conditions before OAE2 (Fig. 10; 564 Owens et al., 2012; van Helmond et al., 2014b). The comparison with modelling results is 565 partially consistent only for the *DeepCAS* simulation (Fig. 10a). In this scenario, the deep connection (> 2500 meters depth) between the Pacific and Central Atlantic basins allows 566 567 oxygenated deep waters from the Pacific to flow into and fill the western part of the deep 568 Central Atlantic (Supplementary Fig.S8), resulting in simulated oxic deep waters that 569 correlate with the proxy record. In the IntermediateCAS simulation, the shallower CAS (< 570 2500 meters of water depth) prevents these oxygenated deep waters to penetrate into the 571 Central Atlantic, which is supplied by oxygen-poor intermediate waters resulting in hypoxic 572 conditions, which is not consistent with data in the deep Central Atlantic for the pre-OAE2 573 (Fig. 10b). The same conclusion is made for the ShallowCAS scenario, for which the deep 574 Central Atlantic is fully anoxic due to very high sea-bottom temperature and subsidence of 575 anoxic waters from the Equatorial Atlantic Gateway (Fig. 10c).

576 Our numerical results thus suggest that a CAS opened only to upper ocean circulation is 577 unlikely in the Cenomanian because a long-term complete anoxia does not characterize the 578 Central Atlantic basin besides during the OAE2. Our simulations underline that deep 579 connections are required between the two basins prior to the OAE2 in order to fill the Central 580 Atlantic with well-oxygenated deep waters. This result suggests either a motion of the CLIP 581 through the CAS coeval, at the earliest, with the beginning of the OAE2, or the existence of 582 multiple volcanic arc systems separated by deep channels around the CLIP (Romito & Mann, 583 2020). A tantalizing implication of our results is that the early eastward movement of the 584 CLIP across the CAS at the end of the Cenomanian may have participated in the formation of

585 a complete anoxia in the deep Central Atlantic during the OAE2. The transition from abyssal 586 (4000 m, the DeepCAS) to bathyal depths (2500 m, IntermediateCAS) in our simulations 587 indeed contribute to decreasing seafloor oxygen concentrations in the Central Atlantic. 588 However supplementary simulations are needed to corroborate this hypothesis as other 589 factors, such as an enhanced supply of nutrients to the ocean consecutive to volcanism or 590 pCO2 increase (Adams et al., 2010; Barclay et al., 2010; Joo et al., 2020; Pogge Von 591 Strandmann et al., 2013; Turgeon & Creaser, 2008) may force the deep Central Atlantic to 592 become anoxic despite deep ocean connections across the CAS.

593 The eastern part of the Central Atlantic remains hypoxic to anoxic in the deep, regardless 594 of the CAS depth, probably because of the combination of its bottleneck configuration for 595 deep circulation and of elevated marine productivity above (Supplementary Fig.S6). These 596 reduced oxygen conditions are inconsistent with pre-OAE2 data for the eastern Central 597 Atlantic that document oxic environments (Niels A. G. M. van Helmond et al., 2014). This 598 discrepancy suggests that an intermittent source of oxygenated deep waters may have existed 599 in the northern part of the Cenomanian Central Atlantic, thereby creating oxic conditions in 600 the deep eastern side of the basin (Fig. 10; Sites 1276 and 641) prior to the OAE2 and also 601 providing an explanation to the temporary re-oxygenation of the deep Central Atlantic during 602 the Plenus Cold Event within the OAE2 (van Helmond et al. 2014b). The recent Cenomanian 603 simulation of Ladant et al. (2020) indicate possible deep-water formation in the Cenomanian 604 North Atlantic but this source does not exist in our simulations, and we note that evidence is 605 lacking to unambiguously confirm its existence. A deeper connection with the Neotethys is 606 also suggested during the Late Cretaceous (Buchs et al., 2018; Nouri et al., 2016) and could 607 supply oxygen to the North-Eastern Central Atlantic. However, reconstructing 608 paleogeography and paleobathymetry in this area is particularly challenging because the 609 Neotethyan region is composed of numerous shallow carbonate platforms separated by deep 610 and narrow corridors (Stampfli and Borel 2002). As such, we cannot draw from our results 611 anything but speculative conclusions about a possible role of the Tethys Seaway on the 612 evolution of deep Central Atlantic oxygenation.

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1.3 How to make anoxic an ocean despite an active circulation?

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615 Very weak renewal of deep-water masses is often needed to generate anoxia in an ocean 616 basin. Silled basins are a prime example as they are affected by a strong circulation restriction 617 and stratification with long water renewal rate (Algeo & Lyons, 2006). It was for instance 618 calculated that the renewal rate in the anoxic Cleveland basin during the Toarcian Oceanic

619 Anoxic Event was between 4000 and 40 000 years (McArthur et al., 2008). During the OAE2, 620 the renewal rate in the deep Central Atlantic is estimated between 500 and 4000 years (Niels 621 A. G. M. van Helmond et al., 2014), which is similar to that of the modern Black Sea (Algeo 622 & Lyons, 2006). This estimation is consistent with the three simulated scenarios that exhibit 623 water ages in the deep Central Atlantic comprised between ~500 years (in the ShallowCAS 624 simulation) and ~2000 years (in the IntermediateCAS simulation). The ShallowCAS scenario 625 provides interesting insights in terms of ocean dynamics and oxygenation mechanisms by 626 showing that the degree of oxygen depletion can be linked to the provenance of water masses 627 rather than a sluggish circulation. Indeed, this silled basin in the ShallowCAS simulation 628 exhibits an active deep circulation (around 5-6 Sv), with relatively short renewal age, and is 629 fully anoxic at the same time because incoming subsiding waters that flow into the deep 630 Central Atlantic come from the anoxic OMZ of the Equatorial Atlantic Gateway. These 631 results propose an alternative to the common view that ocean stagnation is required to create 632 anoxia and enhance organic carbon burial in deep basins.

633 Previous model studies have also suggested that ocean stagnation was not sufficient to 634 create intermediate to shallow anoxia/euxinia (Ozaki et al. 2011, Monteiro et al. 2012). (Ozaki 635 et al., 2011) showed that ocean stagnation only promoted deep-water anoxia and that massive 636 phosphorus inputs were additionally required to create a global anoxia by enhancement of 637 marine productivity. Monteiro et al. (2012) observed as well that enhanced marine 638 productivity due to higher nutrient content was the most efficient process to expand oceanic 639 anoxia. In contrast, our findings demonstrate that full basin-scale anoxia can co-exist with an 640 active ocean circulation and without the need for enhanced productivity.

- 641
- 642 5. Conclusion
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The activity of the CLIP in the Late Cenomanian have impacted the paleobathymetry 644 645 of the CAS via the formation of the Caribbean plateau. In this study, we use the IPSLCM5A2 646 earth system model, which includes the marine biogeochemistry model PISCES, to explore 647 the impact of the paleobathymetry of the CAS on oceanic circulation and dissolved oxygen 648 distribution during the Cenomanian. Regardless of the depth of the CAS, the model simulates 649 an active oceanic circulation, dominated by the Pacific Ocean. The spatial variations of the 650 global oxygen distribution compare well with pre-OAE2 proxy records and exhibit oceanic 651 environments ranging from fully oxic to fully anoxic. These spatial variations are strongly 652 driven by the global paleobathymetry and by the configuration of oceanic gateways that 653 control the pathways of deep currents and the associated oxygen supply. Restricted basins 654 such as the Central and South Atlantic are oxygen-poor and even locally anoxic in all 655 simulations, while the deep Southern Pacific is well-oxygenated because it is a region of 656 deep-water formation. The depth of the CAS, however, exerts substantial control on the 657 seafloor oxygen concentration of the Central Atlantic basin. In the DeepCAS configuration, 658 most of the deep western Central Atlantic is oxic whereas the deep eastern part of the basin is 659 dysoxic. Dysoxia spreads westward in the IntermediateCAS configuration while a complete anoxia develops in the whole basin in the ShallowCAS configuration. A comparison with 660 661 redox proxy data suggests that a deep connection (> 2500 meters) existed before the OAE2 between the Pacific and Central Atlantic. A shallowing of the CAS to an intermediate depth 662 663 (~2500 meters of water depth) due to the eastward movement of the Caribbean LIP may have 664 participated in the establishment of complete hypoxic to anoxic environments in the deep 665 Central Atlantic, by reducing the oxygen supply from Pacific bottom waters. These results illustrate how long-term paleobathymetric forcings may precondition the ocean for favorable 666 667 low-oxygen conditions allowing OAEs to develop and high quantity of organic carbon to be preserved in the oceans and eventually buried. 668

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670 DATA AVAILABILITY STATEMENT

- 671 Code availability:
- 672 LMDZ, XIOS, NEMO and ORCHIDEE are released under the terms of the CeCILL license. OASIS-
- 673 MCT is released under the terms of the Lesser GNU General Public License (LGPL). IPSL-CM5A2
- 674 code is publicly available through svn, with the following command lines: svn co
- 675 http://forge.ipsl.jussieu.fr/igcmg/svn/modipsl/branches/publications/IPSLCM5A2.1_11192019
- 676 modipsl
- 677 cd modipsl/util;./model IPSLCM5A2.1
- 678 The mod.def file provides information regarding the different revisions used, namely:
- 679 NEMOGCM branch nemo_v3_6_STABLE revision 6665
- 680 XIOS2 branchs/xios-2.5 revision 1763
- 681 IOIPSL/src svn tags/v2_2_2
- 682 LMDZ5 branches/IPSLCM5A2.1 rev 3591
- 683 branches/publications/ORCHIDEE_IPSLCM5A2.1.r5307 rev 6336
- 684 OASIS3-MCT 2.0_branch (rev 4775 IPSL server)
- 685 The login/password combination requested at first use to download the ORCHIDEE component is
- anonymous/anonymous. We recommend to refer to the project website:
- 687 http://forge.ipsl.jussieu.fr/igcmg_doc/wiki/Doc/Config/IPSLCM5A2 for a proper installation and
- 688 compilation of the environment.

- 690 Data availability: Datasets from previously published sources have been used for this research. The
- 691 complete list of references is detailed in the Supporting Data D1.
- 692

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948 SUPPLEMENTARY INFORMATION

- Supporting Information S1
- 950 Supporting Data

D1

Figures







Figure 2: Oxygen concentration for the Deep CAS simulation (in mmol.m⁻³); (a) At the surface level of the model (0-10 m)
with contours of sea-surface temperatures (°C); (b) At 180 meters depth with areas of strong upwelling (grey shaded areas, vertical speed > 8 cm/day), and (c) At 700 meters depth with the mixed layer depth as contours (meters).





Figure 3: Oceanic circulation and oxygenation for the Deep CAS simulation. (a) Deep- currents velocity on the seafloor for areas where water depth > 3000 meters (m/s). Blue crosses correspond to areas of deep-water formation. Submarine topographic features are labelled in red: TP: Tethyan platforms, CAS: Central American Seaway, EAG: Equatorial Atlantic Gateway, DR: Davie Ridge, DP: Drake Passage, KP: Kerguelen Plateau. (b) Oxygen concentrations on the seafloor for areas where water depth > 3000 meters (mmol/m3). Black contours correspond to the water age at the same depth (years).



983 984 985 Figure 4: (a) Plot of water age (years) vs. oxygen concentration (mmol/m3) for water depth > 3000. Dots are colored by basin. Deep-water formation occurs in the Southern Pacific (Black cross) and black arrows indicate the two main circulation pathways through the basin. (b) Map showing the extent of different basins.



Figure5: (a) Vertical profiles in oxygen concentrations for the DeepCAS simulation. Locations are indicated on (c). (b) Marine productivity integrated over the whole water column $(gC.m^{-3}.yr^{-1})$ vs. oxygen concentration (integrated over the first 200 meters of water depth) plotted for locations around each profile of (a), covering -2/+2 decimal degrees in longitude and latitude around the profile. (c) Marine productivity integrated over the whole water column (gC.m⁻³.yr⁻¹) and locations of profiles.



Figure 6: (a) Schematic representation of surface circulation in the Central Atlantic. See localization on Fig.1. Nth.Am.: North America, Sth.Am.: South America, Af.: Africa. (b) Incoming and outgoing fluxes (Sverdrups – Sv) into the Central Atlantic, computed through sections described in Fig.1. The histogram is calculated for the DeepCAS simulation, values for IntermediateCAS and ShallowCAS simulations are very similar, except for the incoming and outgoing fluxes through the CAS: the dark blue histogram corresponds to fluxes consistent in the three simulations, the shaded part of the histogram corresponds to supplementary fluxes observed only in IntermediateCAS and DeepCAS simulations.



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Figure 7: Schematic representation of intermediate circulation in the Central Atlantic for (a) DeepCAS and IntermediateCAS simulations and (b) ShallowCAS simulation. (c) Incoming/outgoing water fluxes (Sverdrups – Sv) for DeepCAS simulations. 1025 Fluxes for IntermediateCAS simulations are similar. (d) Incoming/outgoing water fluxes for ShallowCAS simulation.

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Figure 8: Schematic representation of deep circulation in the Central Atlantic for (a) DeepCAS, (b) IntermediateCAS and (c) ShallowCAS simulations.



Figure 9: Vertical sections of the water age (years) for the ShallowCAS simulation. The Section A is through the Tethys seaway and the section B through the Equatorial Atlantic Gateway (EAG). Blue arrow North of the EAG (section B) indicates intermediate water subsidence from the EAG to the Central Atlantic. No subsidence is observed near the Tethys Seaway.





1032 1033 1034 1035 1036 Figure 10: Seafloor dissolved oxygen concentration (mmol/m³) for (a) DeepCAS, (b) IntermediateCAS and (c) ShallowCAS simulations. Red, and black lines correspond to hypoxia and anoxia limits, respectively. Waters are hypoxic for oxygen concentrations below 62.5 mmol.m⁻³ and anoxic below 6.5 mmol.m⁻³ (see color scale)

Crosses and dots correspond to Pre-OAE2 redox data (see discussion). Sites IODP 1276 and ODP 641 exhibit a data-model mismatch addressed in the discussion.



Figure 11: Seafloor dissolved oxygen concentration for the DeepCAS simulation (mmol.m⁻³) compared with redox data for
(a) the pre-OAE2 period and (b) the OAE2. The black and red lines indicate the limits of anoxia and dysoxia, respectively.
Except in the central Atlantic, oxygen modelling results for IntermediateCAS and Shallow CAS are similar (See
Supplementary Figure S9). Dots emphasized with asterisks correspond to model-data mismatches discussed.

Figure 1.



Figure 2.



Figure 3.



Figure 4.



Figure 5.



Figure 6.



Figure 7.



Figure 8.



Figure 9.



Figure 10.



Figure 11.



Supplementary Online Material for:

Exploring the impact of Cenomanian paleogeography and marine gateways on oceanic oxygen

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Supplementary Figure 1: Time series for (a) sea surface temperature, (b) deep oceanic temperatures (2300 meters of water depth), (c) maximum intensity of the meridional stream function (MSF) and (d) deep dissolved oceanic oxygen (2300 meters of water depth; PISCES offline simulations). The simulation was started with a low atmospheric time step for technical reasons, the SST diminution observed around 3000 years corresponds to a return to the standard atmospheric time step.



Supplementary Table 1: Temperature and $[O_2]$ drifts during the last 1000 years of simulation for the surface and the deep ocean (3000 meters of water depth), in °C/century and mmol/m³/century, respectively.

	DeepCAS	IntermediateCAS	ShallowCAS
SST (°C/century)	0.011	0.022	0.025
Deep ocean temperature (°C/century)	0.024	0.025	0.03
Surface dissolved oxygen (mmol/m³/century)	0.0001	0.0002	0.00015
O2 bottom (mmol/m ³ /century)	0.027	0.065	0.049



Supplementary Figure S2: Global meridional stream function (Sv). Positive values (red) indicate a clockwise circulation and negative values (blue) an anticlockwise circulation. Annual average over the last 100 years of simulation. (a) DeepCAS simulation, (b) IntermediateCAS simulation and (c) ShallowCAS simulation.

Supplementary Figure S3: Meridional vertical section in the Eastern Pacific showing the supply of oxygen to the deep ocean via deep-water formation. Color shading corresponds to simulated oceanic dissolved oxygen concentration (mmol/m³, annual average for longitudes between 130 and 180°E. Black contours indicate the simulated water age (years, annual average for longitudes between 130 and 180°E). Water age is the number of years after last surface contact. Black arrows show water sinking and oxygen supply.



Supplementary Figure S4: Plot of marine productivity (mmol/m3, integration for the whole water column) vs. oxygen concentration (mmol/m3) at 90-100 meters of water depth in the Central Atlantic (DeepCAS simulation, annual average over the last 100 years of simulation), colored by vertical velocity.



Supplementary Figure S5: Surface oceanic trends for the Deep CAS simulations (Annual average for 1 to 100 meters of water depth). (a) Water age (years). (b) Vertical velocity (m/day).



Supplementary Figure S6: Marine productivity ($gC/m^2/year$) for Deep, Intermediate and Shallow CAs simulations (Integrated over the whole water column, Annual average over the last 1000 years of simulation). Nth.Am.: North America, Sth.Am.: South America, Af.: Africa.



Supplementary Figure S7: Oceanic trends for intermediate waters (Annual average between 600 and 1600 meters of water depth over the last 100 years of simulation for DeepCAS simulation (left) and ShallowCAS simulation (right). From top to bottom: Water age (years, Water temperature (°C), Water salinity (psu).



90°% 80°% 70°% 60°% 50°% 40°% 30°% 20°% 10°% 0° 10°E 20°E 30°E 90°% 80°% 70°% 60°% 50°% 40°% 30°% 20°% 10°% 0° 10°E 20°E 30°E



Supplementary Figure S8: Vertical sections of $[O_2]$ (mmol/m³, Annual average over the last 100 years of simulation) through the CAS (latitude of 7°N) for (a) DeepCAS, (b) IntermediateCAS and (c) ShallowCAS simulations.

Supplementary Figure S9: Seafloor dissolved oxygen concentration (mmol/m³, Annual average over the last 100 years of simulation and comparison with redox data for (a) IntermediateCAS simulation / pre-OAE2 data, (b) IntermediateCAS simulation / OAE2 data, (c) ShallowCAS simulation / pre-OAE2 data and (d) ShallowCAS simulation / OAE2 data. The black and red lines indicate the limits of anoxia and dysoxia, respectively.

