

1 **Inconsistent strategies to spin up models in CMIP5: implications for**
2 **ocean biogeochemical model performance assessment**

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4 **Roland Séférian^{1*}, Marion Gehlen², Laurent Bopp², Laure Resplandy^{3,2}, James C.**
5 **Orr², Olivier Marti², John P. Dunne⁴, James R. Christian⁵, Scott C. Doney⁶,**
6 **Tatiana Ilyina⁷, Keith Lindsay⁸, Paul Halloran⁹, Christoph Heinze^{10,11}, Joachim**
7 **Segschneider¹², Jerry Tjiputra¹¹, Olivier Aumont¹³, Anastasia Romanou^{14,15}**

8 ¹ CNRM-GAME, Centre National de Recherches Météorologiques-Groupe d'Etude
9 de l'Atmosphère Météorologique, Météo-France/CNRS, 42 Avenue Gaspard Coriolis,
10 31057 Toulouse, France

11 ² LSCE/IPSL, Laboratoire des Sciences du Climat et de l'Environnement, Orme des
12 Merisiers, CEA/Saclay 91198 Gif-sur-Yvette Cedex, France

13 ³ Scripps Institution of Oceanography, UCSD, La Jolla, CA, USA

14 ⁴ Geophysical Fluid Dynamics Laboratory, NOAA, Princeton, New Jersey, USA

15 ⁵ Fisheries and Oceans Canada and Canadian Centre for Climate Modelling and
16 Analysis, Victoria, B.C., Canada

17 ⁶ Marine Chemistry and Geochemistry Department, Woods Hole Oceanographic
18 Institution, Woods Hole MA, USA

19 ⁷ Max Planck Institute for Meteorology, Bundesstraße 53, 20146 Hamburg, Germany

20 ⁸ Climate and Global Dynamics Division, National Center for Atmospheric Research,
21 Boulder, Colorado

22 ⁹ College of Life and Environmental Sciences, University of Exeter, Exeter, EX4
23 4RJ, UK

24 ¹⁰ Geophysical Institute, University of Bergen and Bjerknes Centre for Climate
25 Research, Bergen, Norway

26 ¹¹ Uni Research Climate, Allegt. 55, 5007 Bergen and Bjerknes Centre for Climate
27 Research, Bergen, Norway

28 ¹² University of Kiel, Kiel, Germany

29 ¹³ Sorbonne Universités (UPMC, Univ Paris 06)-CNRS-IRD-MNHN, LOCEAN-
30 IPSL Laboratory, 4 Place Jussieu, F-75005 Paris, France

31 ¹⁴ Dept. of Applied Math. and Phys., Columbia University, 2880 Broadway, New
32 York, NY 10025, USA

33 ¹⁵ NASA-GISS, NY, USA

34

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36 *Corresponding author address: Roland Séférian, CNRM-GAME, 42 av. Gaspard
37 Coriolis 31100 Toulouse. E-mail: roland.seferian@meteo.fr

38

39 **Abstract**

40 During the fifth phase of the Coupled Model Intercomparison Project (CMIP5)
41 substantial efforts were made on the systematic assessment of the skill of Earth
42 system models. One goal was to check how realistically representative marine
43 biogeochemical tracer distributions could be reproduced by models. Mean-state
44 assessments routinely compared model hindcasts to available modern biogeochemical
45 observations. However, these assessments considered neither the extent of equilibrium
46 in modeled biogeochemical reservoirs nor the sensitivity of model performance to
47 initial conditions or to the spin-up protocols. Here, we explore how the large diversity
48 in spin-up protocols used for marine biogeochemistry in CMIP5 Earth system models

49 (ESM) contribute to model-to-model differences in the simulated fields. We take
50 advantage of a 500-year spin-up simulation of IPSL-CM5A-LR to quantify the
51 influence of the spin-up protocol on model ability to reproduce relevant data fields.
52 Amplification of biases in selected biogeochemical fields (O_2 , NO_3 , Alk-DIC) is
53 assessed as a function of spin-up duration. We demonstrate that a relationship
54 between spin-up duration and assessment metrics emerges from our model results and
55 is consistent when confronted against a larger ensemble of CMIP5 models. This
56 shows that drift has implications on performance assessment in addition to possibly
57 aliasing estimates of climate change impact. Our study suggests that differences in
58 spin-up protocols could explain a substantial part of model disparities, constituting a
59 source of model-to-model uncertainty. This requires more attention in future model
60 intercomparison exercises in order to provide realistic ESM results on marine
61 biogeochemistry and carbon cycle feedbacks.

62

63 **1- Introduction**

64 **1-1 Context**

65 Earth system models (ESM) are recognized as the current state-of-the-art global
66 coupled models used for climate research (e.g., Hajima et al., 2014; IPCC, 2013).
67 They expand the numerical representation of the climate system used during the 4th
68 IPCC assessment report (AR4) that was limited to coupled physical general
69 circulation models, to the inclusion of biogeochemical and biophysical interactions
70 between the physical climate system and the biosphere. ESMs that contributed to
71 CMIP5 substantially differ in terms of their simulations of physical and
72 biogeochemical components. These differences in design translate into a significant
73 variability of the models' ability to reproduce the observed biogeochemistry and

74 carbon cycle, which in turn may impact projected climate change responses (IPCC,
75 2013).

76

77 In the typical objective evaluation and intercomparison of these models, a suite of
78 standardized statistical metrics (e.g., correlation, root-mean-squared errors) is applied
79 to quantify differences between modeled and observed variables (e.g., Doney et al.,
80 2009; Rose et al., 2009; Stow et al., 2009; Romanou et al., 2014; 2015). With the goal
81 of constraining future projections, statistical metrics are often used for model ranking
82 (e.g., Anav et al., 2013), weighting of model projections (e.g., Steinacher et al., 2010)
83 or selection of the most skillful models across a wider ensemble (e.g., Cox et al.,
84 2013; Massonnet et al., 2012; Wenzel et al., 2014). Most of these approaches can be
85 considered as “blind” given that they are routinely applied without considering
86 models’ specific characteristics and treat models *a priori* as equivalently independent
87 of observations. However, since these models are typically initialized from
88 observations, the spin-up procedure of climate variables are the most model-
89 dependent protocols that could introduce errors or drifts in modeled fields with
90 consequences on skill score metrics.

91

92 **1-2 Initialization of biogeochemical fields and spin-up protocols in CMIP5**

93 Ocean initialization protocols aim at obtaining stable and equilibrated distributions of
94 model state variables, such as temperature or concentrations of dissolved tracers. Most
95 commonly used initialization protocols consist of initializing both physical and
96 biogeochemical variables with either climatologies of the observed fields or constant
97 values before running the model to equilibrium. In theory, equilibrium corresponds to
98 steady-state and, hence, temporal derivatives of tracer fields close to zero. The time

99 needed to equilibrate tracer distributions or, in other words, the integration time
100 needed by the model to converge towards its own attractor (which is different from
101 the true state of the climate system) varies greatly between components of the climate
102 system. It spans from several weeks for the atmosphere (e.g., Phillips et al., 2004) to
103 several centuries for ocean and sea ice components (e.g., Stouffer et al., 2004). The
104 equilibration of ocean biogeochemical tracers across the entire water column amounts
105 to several thousands of years (e.g., Heinze et al., 1999; Wunsch and Heimbach, 2008)
106 and depends on the state of background ocean circulation as well as the turbulent
107 mixing and eddy stirring parameterizations (e.g., Aumont et al., 1998; Bryan, 1984;
108 Gnanadesikan, 2004; Marinov et al., 2008). In practice, these simulations, called
109 “spin-up”, span in general only several hundreds of years at the end of which a quasi-
110 equilibrium state is assumed for the interior ocean tracers.

111

112 The present degree of complexity and increasing spatial as well as temporal resolution
113 of marine biogeochemical ESM components, however, often precludes a spin-up to
114 reach adequate equilibration of biogeochemical tracers. This is a consequence of the
115 increasing number of state variables present in most of the current generation of
116 biogeochemical models (e.g., for each tracer a separate advection equation has to be
117 solved via a numerical CPU time demanding algorithm), more complex process
118 descriptions (e.g., including more plankton functional types than before), and
119 increasing spatial as well as temporal resolution. This number has continuously
120 increased from simple biogeochemical models (e.g., HAMOCC3, Maier-Reimer and
121 Hasselmann (1987)) to marine biodiversity models (e.g., Follows et al., 2007).
122 Current generation biogeochemical models embedded in CMIP5 ESMs contain
123 roughly two to four times more state variables than the physical models (e.g.,

124 atmosphere, ocean, sea-ice), which makes their equilibration computationally costly
125 and difficult. The initialization of biogeochemical state variables is further
126 complicated by the scarcity of biogeochemical observations as compared to
127 observations of physical variables (e.g., temperature, salinity). While three-
128 dimensional observation-based climatologies exist for macro-nutrients, oxygen,
129 dissolved carbon and alkalinity, for other tracers such as dissolved iron, dissolved
130 organic carbon and biomass of the various plankton functional types data are still
131 sparse and represent measurements done over different time periods and climate
132 conditions (in spite of considerable efforts such as the GEOTRACES program for
133 trace elements, or MAREDAT for biomasses of plankton functional types). The latter
134 are initialized either with constant values (e.g. global average estimates) or with
135 output from a previous model run. An additional difficulty stems from the use of
136 modern climatologies to initialize the ocean state, implicitly assuming a long-term
137 steady state, which does not necessarily represent the preindustrial state of the ocean.
138 These climatologies incorporate the ongoing anthropogenic perturbation of marine
139 biogeochemical fields, be it the uptake of anthropogenic CO₂ or the excess of
140 nutrients inputs and pollutants (e.g., Doney, 2010). Although methods exist to remove
141 the anthropogenic perturbation from observed ocean carbon tracer fields, their use is
142 still debated since they lead to non-unique results (e.g., Tanhua et al., 2007; Yool et
143 al., 2010).

144
145 The equilibration of marine biogeochemical tracer distributions is driven not only by
146 the ocean circulation but also by numerous internal biogeochemical processes acting
147 at various time scales. For example, while the transport and degradation of sinking
148 organic matter spans days to perhaps several months, the associated impact on deep

149 water chemistry accumulates over several decades to centuries as zones of differential
150 remineralization are mixed across water masses and follows the ocean circulation
151 (Wunsch and Heimbach, 2008). For models including interactive sediment modules,
152 the sediment equilibration takes even longer ($O(10^4)$ years; e.g., Archer et al. (2009)
153 and Heinze et al. (1999)). As a consequence of the interplay between ocean
154 circulation and biogeochemical processes, biogeochemical models require long spin-
155 up times to equilibrate (e.g., Khatiwala et al., 2005; Wunsch and Heimbach, 2008).
156 Modeling studies of paleo-oceanographic passive tracers such as $\delta^{18}\text{O}$ or $\Delta^{14}\text{C}$
157 (Duplessy et al., 1991), or global ocean passive tracers (Wunsch and Heimbach,
158 2008), as well as more recently available modern global scale data compilations (e.g.,
159 Key et al., 2004; Sarmiento and Gruber, 2006) and GEOTRACES Intermediate Data
160 product 2014 (Version 2) <http://www.bodc.ac.uk/geotraces/data/idp2014/>) provide an
161 estimate of the time required for the ocean biogeochemical reservoir to equilibrate
162 with the climate systems (excluding continental weathering and reaction with marine
163 sediments). Depending on ocean circulation, it ranges from 1500 years for subsurface
164 water masses to 10000 years for the deep water masses (Wunsch and Heimbach,
165 2008).

166

167 In a context of model-to-model intercomparison, this time range contributes to the
168 model uncertainty. Lessons from the previous OCMIP-2 exercise have demonstrated
169 that some models required $\sim 10,000$ years to equilibrate to a global sea-air carbon flux
170 of 0.01 Pg C y^{-1} .

171

172 While it is recognized that long time-scale processes influence the length of spin-up to
173 equilibrium, the spin-up duration is usually defined *ad hoc* based on external

174 constraints or internal biogeochemical criteria. The computational cost is commonly
175 invoked as external constraint to shorten and limit the spin-up duration. It is directly
176 related to model complexity (e.g., Tjiputra et al., 2013; Vichi et al., 2011; Yool et al.,
177 2013) and spatial resolution (Ito et al., 2010). The internal biogeochemical criteria
178 applied to derive the duration of the spin-up simulations are generally defined by (i)
179 reaching a steady-state, quasi equilibrium of the long-term global-mean CO₂ fluxes
180 between the ocean and the atmosphere (e.g., Dunne et al., 2013; Ilyina et al., 2013;
181 Lindsay et al., 2014; Romanou et al., 2013; Séférian et al., 2013), (ii) determining the
182 amount of carbon stored into the ocean at preindustrial state (e.g., Dunne et al., 2013;
183 Vichi et al., 2011) or (iii) representing relevant biogeochemical tracer patterns (e.g.,
184 oxygen minimum zone in Ito and Deutsch (2013)).

185

186 Despite its importance, only limited information on spin-up procedures is available
187 through the CMIP5 metadata portal (<http://metaforclimate.eu/trac>). Information on
188 spin-up protocols and model initialization is usually not taken into account in model
189 intercomparison studies (e.g., Andrews et al., 2013; Bopp et al., 2013; Cocco et al.,
190 2013; Frölicher et al., 2014; Gehlen et al., 2014; Keller et al., 2014; Resplandy et al.,
191 2013; 2015; Rodgers et al., 2014; Séférian et al., 2014). This information, if available,
192 can only be found separately in the reference papers of individual models (e.g.,
193 Adachi et al., 2013; Arora et al., 2011; Collins et al., 2011; Dunne et al., 2013; Ilyina
194 et al., 2013; Lindsay et al., 2014; Romanou et al., 2013; Séférian et al., 2013; Séférian
195 et al., 2015; Tjiputra et al., 2013; Vichi et al., 2011; Volodin et al., 2010; Watanabe et
196 al., 2011; Wu et al., 2013). The duration of spin-up simulations of CMIP5 ocean
197 biogeochemical components spans from one hundred years (e.g., CMCC-CESM) to
198 several thousand years (e.g., MPI-ESM-LR, MPI-ESM-MR) (Figure 1 and Table 1).

199 Model initialization and spin-up procedures are equally variable across the model
200 ensemble (Figure 1 and Table 1). Four different sources of initialization and four
201 different procedures of model equilibration emerge from the 24 ESMs reviewed for
202 this study.

203

204 Biogeochemical state variables were mostly initialized from observations, although
205 from various releases of the same World Ocean Atlas global climatology (WOA1994,
206 WOA2001, WOA2006, WOA2010). A small subset of ESMs relied either on a mix
207 between previous model output and observations or solely on model output from a
208 previous simulation for initialization. Similarly, spin-up procedures fall into two
209 categories. The first one may be called “sequential”: it consists in decomposing the
210 spin-up integration into one long offline simulation (~200-10000 years) and one
211 shorter online simulation (~100-1000 years). During the offline simulation, the
212 biogeochemical model is forced by dynamical fields from the climate model or from
213 reanalysis (CanESM2, MRI-ESM, Figure 1 and Table 1). Some modeling groups have
214 adopted a “direct” strategy, which consists in running solely one online or coupled
215 spin-up simulation (e.g., CNRM-ESM1, GFDL-ESM2M, GFDL-ESM2G, GISS-E2-
216 H-CC, GISS-E2-R-CC, NorESM1-ME). Finally, a spin-up “acceleration” procedure is
217 used by CMCC-CESM. This technique consists of enhancing the ocean carbon
218 outgassing to remove anthropogenic carbon from the ocean, a legacy from
219 initialization with modern data (Global Data Analysis Project or GLODAP following
220 Key et al., 2004). None of these spin-up procedures, durations and sources of
221 initialization can be considered as “standard”; each of them is unique and subjectively
222 employed by one modeling group.

223

224 Objective arguments and hypotheses justifying the choice of one method of spin-up
225 rather than the others have been the focus of previous studies (e.g., Dunne et al., 2013;
226 Heinze and Ilyina, 2015; Tjiputra et al., 2013). Similarly, modeling groups discussed
227 impacts of their particular spin-up procedure on model performance (e.g., Dunne et
228 al., 2013; Lindsay et al., 2014; Séférian et al., 2013; Vichi et al., 2011). However, no
229 study has addressed the potential for the large diversity of spin-up procedures found
230 across the CMIP5 ensemble to translate into model-to-model differences in terms of
231 comparative model performance assessments or model evaluations in terms of future
232 projections.

233

234 **1-3 Objectives of this study**

235 This study assesses the role of the spin-up protocol in the representation of
236 biogeochemical fields and subsequent model skill assessment, providing a
237 complementary analysis from the studies of Sen Gupta et al. (2012; 2013). It relies on
238 a 500-year long spin-up simulation from a state-of-the-art Earth system model, IPSL-
239 CM5A-LR to investigate the impacts of spin-up strategy on selected biogeochemical
240 tracers and residual model drift across the various ESMs of the CMIP5 ensemble. We
241 demonstrate that the duration of the spin-up has implications for the determination of
242 robust and meaningful skill-score metrics that should improve future intercomparison
243 studies such as CMIP6 (Meehl et al., 2014).

244

245 Section 2 describes the model, the observations, the model experiments, as well as the
246 methods used for assessing the impacts of spin-up protocols on the representation of
247 biogeochemical fields in IPSL-CM5A-LR, as well as across the ensemble of CMIP5
248 ESMs. Section 3 presents the analysis developed for the assessment of the impact of

249 spin-up duration on the representation of biogeochemical structures. Implications and
250 recommendations are discussed in Sections 4 and 5, respectively.

251

252 **2- Methods**

253 **2-1- Model simulations**

254 This study exploits in particular results from one simulation performed with IPSL-
255 CM5A-LR (Dufresne et al., 2013) as representative for other CMIP5 Earth system
256 models. As a typical representative of the current generation of ESMs, IPSL-CM5A-
257 LR combines the major components of the climate system (Chap 9, Table 9.1, (IPCC,
258 2013). The atmosphere is represented by the atmospheric general circulation model
259 LMDZ (Hourdin et al., 2006) with a horizontal resolution of $3.75^\circ \times 1.87^\circ$ and 39
260 levels. The land surface is simulated with ORCHIDEE (Krinner et al., 2005). The
261 oceanic component is NEMOv3.2 in its ORCA2 global configuration (Madec, 2008).
262 It has a horizontal resolution of about 2° with enhanced resolution at the equator
263 (0.5°) and 31 vertical levels. NEMOv3.2 includes the sea-ice model LIM2 (Fichefet
264 and Maqueda, 1997), and the marine biogeochemistry model PISCES (Aumont and
265 Bopp, 2006). PISCES simulates the biogeochemical cycles of oxygen, carbon and the
266 main nutrients with 24 state variables. The model simulates dissolved inorganic
267 carbon and total alkalinity (carbonate alkalinity + borate + water) and the distributions
268 of macronutrients (nitrate and ammonium, phosphate, and silicate) and micronutrient
269 iron. PISCES represents two sizes of phytoplankton (i.e., nanophytoplankton and
270 diatoms) and two zooplankton size-classes: microzooplankton and mesozooplankton.
271 PISCES simulates semi-labile dissolved organic matter, and small and large sinking
272 particles with different sinking speeds (3 m d^{-1} and $50 \text{ to } 200 \text{ m d}^{-1}$, respectively).
273 While fixed elemental stoichiometric C:N:P-O₂ ratios after Takahashi et al. (1985) are

274 imposed for these three compartments the internal concentrations of iron, silica and
275 calcite are simulated prognostically . The carbon system is represented by dissolved
276 inorganic carbon, alkalinity and calcite. Calcite is prognostically simulated following
277 Maier-Reimer (1993) and Moore et al. (2002). Alkalinity in the model system
278 includes the contribution of carbonate, bicarbonate, borate, protons, and hydroxide
279 ions. Oxygen is prognostically simulated. The model distinguishes between oxic and
280 suboxic remineralization pathways, the former relying on oxygen as electron acceptor,
281 the latter on nitrate. For carbon and oxygen pools, air-sea exchange follows the
282 Wanninkhof (1992) formulation.

283 The boundary conditions account for nutrient supplies from three different sources:
284 atmospheric dust deposition for iron, phosphorus and silica (Jickells and Spokes,
285 2001; Moore et al., 2004; Tegen and Fung, 1995), rivers for nutrients, alkalinity and
286 carbon (Ludwig et al., 1996) and sediment mobilization for sedimentary iron (de Baar
287 and de Jong, 2001; Johnson et al., 1999). To ensure conservation of nitrogen in the
288 ocean, annual total nitrogen fixation is adjusted to balance losses from denitrification.
289 For the other macronutrients, alkalinity and organic carbon, the conservation is
290 ensured by tuning the sedimental loss to the total external input from rivers and dust.
291 In PISCES, an adequate treatment of external boundary conditions has been
292 demonstrated to be essential for the accurate simulation of nutrient distributions
293 (Aumont and Bopp, 2006; Aumont et al., 2003). Riverine carbon inputs induce a
294 natural outgassing of carbon of 0.6 Pg C y^{-1} which has been shown essential to model
295 the inter-hemispheric gradient of atmospheric CO_2 under preindustrial state (Aumont
296 et al., 2001).

297

298 The core simulation of this study is a 500-year long coupled preindustrial run. It uses
299 the same atmospheric, land surface and ocean configurations as IPSL-CM5A-LR
300 (Dufresne et al., 2013) for which the marine biogeochemistry has been extensively
301 evaluated (see e.g., Séférian et al. (2013) for modern-state evaluation). The only
302 difference between the “standard” preindustrial simulation contributed to CMIP5 and
303 the present one is the initial conditions. While the CMIP5 preindustrial simulation
304 starts from an ocean circulation after several thousand years of online physical
305 adjustment, the present simulation starts from an ocean at rest using the January
306 temperature and salinity fields from the World Ocean Atlas (Levitus and Boyer,
307 1994). Biogeochemical state variables were initialized from data compilations or
308 climatologies as explained in the following section. Atmospheric CO₂ and other
309 greenhouse gases, as well as natural aerosols, were set to their 1850 preindustrial
310 values. The simulation is extensively described in terms of ocean physics by Mignot
311 et al. (2013). Mignot and coworkers show that the strength of the Atlantic meridional
312 overturning circulation and the Antarctic circumpolar current as well as the upper 300
313 m ocean heat content stabilize after 250 years of simulation.

314

315 Although the spin-up protocol used to conduct this 500-year long simulation is not
316 readily comparable to the one used to produce the initial conditions for the CMIP5
317 preindustrial simulation, its duration is greater than the median length of on-line
318 adjustment computed from the multiple spin-up protocols applied during CMIP5
319 (~395 years, Figure 1 and Table 1). Besides, the methodology of initializing
320 biogeochemical state variables from data fields is not broadly employed by the
321 various modeling groups that have contributed to CMIP5. Despite the above-
322 mentioned methodological shortcuts, we take this 500-year long preindustrial

323 simulation as a representative example of a spin-up protocol for the diversity of
324 approaches used by CMIP5 models.

325

326 **2-2- Observations for initialization and evaluation**

327 Two streams of data sets were used in this study. The first stream combines data from
328 the World Ocean Atlas 1994 (WOA94, Levitus and Boyer (1994) and Levitus et al.,
329 (1993)) for the initialization of 3-dimensional fields of temperature and salinity,
330 dissolved nitrate, silicate, phosphate and oxygen, and data from GLODAP (Key et al.,
331 2004) for preindustrial dissolved inorganic carbon and total alkalinity. This stream of
332 data was chosen purposely in our experimental setup to be slightly different than the
333 second stream of data, World Ocean Atlas 2013 (WOA2013, Levitus et al. (2013)),
334 the evaluation data set.

335

336 A second stream of data was used to compare modeled biogeochemical fields. It
337 includes up-to-date observed climatologies of nitrate and oxygen from the WOA2013.
338 This database is based on samples collected since 1965, and incorporates also data
339 from WOA94 onwards. For the concentrations of preindustrial dissolved inorganic
340 carbon and total alkalinity, we still use GLODAP. The second stream of data was
341 selected to be as close as possible to the “standard” evaluation procedure of skill-
342 assessment protocols found in CMIP5 model reference papers (Adachi et al., 2013;
343 Arora et al., 2011; Collins et al., 2011; Dunne et al., 2013; Ilyina et al., 2013; Lindsay
344 et al., 2014; Romanou et al., 2013; Séférian et al., 2013; Séférian et al., 2015; Tjiputra
345 et al., 2013; Vichi et al., 2011; Volodin et al., 2010; Watanabe et al., 2011; Wu et al.,
346 2013). Differences between these two streams of data are minor and are further
347 detailed below.

348

349 **2-3- Approach and statistical analysis**

350 To quantify the impacts of a large diversity of spin-up procedures on the
351 representation of biogeochemical fields in CMIP5, we employ a three-fold approach.

352 (1) The 500-year long spin-up simulation described in Section 2.1 is used to
353 determine the influence of the spin-up procedure on the representation of
354 biogeochemical fields in IPSL-CM5A-LR.

355 (2) In the next step, relationships between biases in modeled fields, model-data
356 mismatches and the duration of the spin-up simulation are identified across the
357 CMIP5 ensemble. For this step, drifts in biogeochemical fields are determined from
358 the first century of the preindustrial simulation (referred to as *piControl*) of each
359 CMIP5 ESM.

360 (3) Finally, the various ensemble of modern hindcast (referred to as *historical*) from
361 each available CMIP5 ESM are used to estimate the impact of these drifts in
362 biogeochemical fields on the ability of models to replicate modern observations. For a
363 given model, we use the ensemble average of the available ‘historical’ members if
364 several realizations are available.

365 For this purpose, several statistical skill score metrics are computed following Rose et
366 al. (2009) and Stow et al. (2009) from model fields interpolated on a regular 1° grid
367 and to fixed depth levels. The skill score metrics are (1) the global averaged
368 concentrations for overall drift; (2) the error or bias between modeled and observed
369 fields at each grid-cell; (3) spatial correlation between model and observations to
370 assess mismatches between modeled and observed large-scale structures; (4) the root-
371 mean squared error (RMSE) to assess the total cumulative errors between modeled
372 and observed fields. These statistical metrics are computed across the water column,

373 but for clarity we focus on surface, 150 m (thermocline) and 2000 m (deep) levels.
374 These statistical metrics were chosen among those described in the literature, because
375 they proved to yield the most indicative scores for tracking model errors or
376 improvement along the various intercomparison exercises (IPCC, 2013).

377

378 The drift is determined for either concentrations in simulated biogeochemical fields or
379 for skill score metrics (e.g., RMSE) using a linear regression fit over a time window
380 of 100 years. This time window of 100 years was chosen as a trade off between a
381 longer time window (>200 years) that smoothes the drift signal and a shorter time
382 window (<100 years) that introduces fluctuations due to internal variability and hence
383 impacting the quality of the fit (see the assessment performed with the millennial-long
384 CMIP5 *piControl* simulation of IPSL-CM5A-LR in Figure S1).

385 The drift is assumed to decrease exponentially during the spin-up simulation and is
386 described by a simple drift model:

$$387 \quad drift(t) = drift(t = 0) \times \exp\left(-\frac{1}{\tau} t\right) \quad (1)$$

388 where τ is the relaxation time of the respective field at a given depth level. It
389 corresponds to the time required to nullify the drift.

390

391 Our analyses focus on the global distribution of nitrate (NO_3), dissolved oxygen (O_2)
392 and the difference between total alkalinity and dissolved inorganic carbon (Alk-DIC).

393 The latter serves as an approximation of carbonate ion concentration following Zeebe
394 and Wolf-Gladrow (2001). We use this approximation of the carbonate ion
395 concentration rather than its concentration, $[\text{CO}_3^{2-}]$, since the latter was poorly
396 assessed in CMIP5 reference papers and was not provided by a majority of ESMs.

397 These three biogeochemical tracers were chosen because (1) most current

398 biogeochemical models simulate Alk, DIC, NO₃ and O₂ prognostically and (2) they
399 are frequently used in state-of-the-art model performance assessment (e.g., Anav et
400 al., 2013; Bopp et al., 2013; Doney et al., 2009; Friedrichs et al., 2009; 2007; Stow et
401 al., 2009), and (3) DIC and Alk are both used as “master tracers” for the carbonate
402 system in the ocean biogeochemistry models (while [CO₃²⁻], e.g., is not explicitly
403 advected as a tracer but diagnosed from temperature, salinity, DIC, Alk, [H⁺], and
404 pCO₂ when needed) . Modeled distributions of NO₃, O₂ and Alk-DIC reflect the
405 representation of biogeochemical processes related to the biological pump (CO₂, NO₃,
406 O₂), the air-sea gas exchange and ocean ventilation (CO₂ and O₂), as well as carbonate
407 chemistry (Alk-DIC). These biogeochemical processes are of particular relevance for
408 investigating the impact of climate change on marine productivity (e.g., Henson et al.,
409 2010), ocean deoxygenation (e.g., Gruber, 2011; Keeling et al., 2009) and the ocean
410 carbon sink, processes for which future projections with the current generation of
411 ESMs yield large inter-model spreads (e.g., Friedlingstein et al., 2013; Resplandy et
412 al., 2015; Séférian et al., 2014; Tjiputra et al., 2014).

413

414 **3 Results**

415 **3-1 Comparison of observational datasets**

416 Our review of spin-up protocols for CMIP5 ESM shows that several modeling groups
417 have employed different streams of datasets to initialize their biogeochemical models
418 (e.g., WOA1994, WOA2001), while model evaluation relies on the most up-to-date
419 stream of data. Differences between the two data streams used for initializing and
420 assessing, respectively, NO₃ and O₂ concentrations are analyzed. Table 2 summarizes
421 RMSE and correlation between WOA1994 and WOA2013 for these two
422 biogeochemical fields.

423

424 Table 2 indicates that differences between the two streams of data are fairly small.

425 The total difference (RMSE) represents a departure between 5 to 10% from the global

426 average concentrations of WOA2013 across depth levels. It is generally lower in

427 regions where the sampling density has not increased markedly between the two

428 releases. These values can be used as a baseline for model-to-model comparison

429 assuming that errors attributed to the various sources of initialization cannot be larger

430 than 10%. Considering that some models have used outputs from previous model

431 simulations or globally averaged concentrations as initial conditions, we acknowledge

432 that this baseline is not a perfect criterion for benchmarking model performance.

433 There is, however, no ideal solution to address this issue since there is no standardized

434 set of initial conditions in CMIP5 except some recommendations for the decadal

435 prediction exercise in which specific attention was paid to initialization (e.g.,

436 Keenlyside et al., 2008; Kim et al., 2012; Matei et al., 2012; Meehl et al., 2013; 2009;

437 Servonnat et al., 2014; Smith et al., 2007; Swingedouw et al., 2013).

438

439 **3-2 Equilibration state metrics in IPSL-CM5A-LR**

440 The global mean sea surface temperature (SST) is a common metric to quantify the

441 energetic equilibrium of the model. This metric has been widely used in various

442 papers referenced in this study to determine the equilibration of ESM physical

443 components. Figure 2a shows the evolution of this metric during the 500-year long

444 spin-up simulation. The global average SST sharply decreases during the first 250

445 years of the simulation. In the last 250 years of the simulation, the global averaged

446 SST displays a small residual drift of $\sim 10^{-4} \text{ }^\circ\text{C y}^{-1}$ which falls into the range of the

447 drifts reported for CMIP5 ESMs. The evolution over the last 250 years is comparable

448 to those of other physical equilibration metrics, such as the ocean heat content or the
449 meridional overturning circulation (Mignot et al., 2013).

450

451 The temporal evolution of sea-to-air CO₂ fluxes was used in phase 2 of the Ocean
452 Carbon Model Intercomparison Project (OCMIP-2, Orr (2002)) as an equilibration
453 metric for the marine biogeochemistry and was still widely used during CMIP5.

454 Figure 2b presents its evolution in the 500-year long spin-up simulation. The global
455 ocean sea-to-air CO₂ flux is $\sim -0.7 \text{ Pg C y}^{-1}$ over the last decades of the spin-up
456 simulation (negative values indicate ocean CO₂ uptake).

457 To assess the global sea-to-air carbon flux, we use the range of values estimated from
458 preindustrial natural ocean carbon flux inversions (e.g. Gerber and Joos (2010) or
459 Mikaloff Fletcher et al. (2007)). Since, these estimates do not account for the
460 preindustrial carbon outgassing induced by the river input, while our model does, we
461 have added a constant outgassing of 0.45 Pg C y^{-1} to the range of $0.03 \pm 0.08 \text{ Pg C y}^{-1}$
462 (Mikaloff Fletcher et al. 2007). This value of 0.45 Pg C y^{-1} corresponds to the global
463 open-ocean river-induced carbon outgassing accordingly to IPCC (2013) or Le Quéré
464 et al. (2015). Consequently, in our modeling framework, the target value of the global
465 sea-to-air carbon flux ranges between 0.4 and 0.56 Pg C y^{-1} .

466

467 Figure 2b shows that the global sea-to-air carbon flux does not fit our range of values
468 estimated from preindustrial natural ocean carbon flux inversions. Besides, Figure 2b
469 shows that the drift in the global sea-to-air carbon flux reduces more slowly after a
470 strong decline during the first 50 years of the spin-up simulation. While this drift is
471 about $0.001 \text{ Pg C y}^{-2}$ from year 250 to 500, it is weaker over the last century of the
472 simulation ($7 \times 10^{-4} \text{ Pg C y}^{-2}$). Using a linear fit over the last century of the simulation

473 with a drift of 7×10^{-4} Pg C y^{-2} , we estimate that the simulated sea-to-air carbon flux
474 would reach the range of 0.4-0.56 Pg C y^{-1} after 1100 to 1300 supplemental years of
475 spin-up simulation. Our simple drift model (Equation 1) gives a relaxation time of
476 around 160 years, which indicates that drift in ocean carbon flux should range
477 between 2×10^{-7} and 7×10^{-7} Pg C y^{-2} after this 1100 to 1300 supplemental years of spin-
478 up simulation.

479

480 These estimates do not account for the non-linearity of the ocean carbon cycle and the
481 associated process uncertainties (Schwinger et al., 2014), and hence potentially
482 underestimate the time required to equilibrate the ocean carbon cycle and sea-to-air
483 carbon fluxes in the range of inversion estimates. The drift of 0.001 Pg C y^{-2} is,
484 however, much smaller than the oceanic sink for anthropogenic carbon. Even if not
485 fully equilibrated in terms of carbon balance, it is likely that this run would have
486 given consistent estimates of anthropogenic carbon uptake in transient historical
487 hindcasts.

488

489 **3-3 Temporal evolution of model errors in IPSL-CM5A-LR**

490 Figure 3 shows the temporal evolution of globally averaged concentrations for O₂,
491 NO₃ and Alk-DIC at the surface (panels a, b and c), 150 m (panels d, e and f) and
492 2000 m (panels g, h, and i). Globally averaged concentrations of O₂, NO₃ and Alk-
493 DIC (solid lines) reach steady state after 100 to 250 years of spin-up at the surface.
494 While modeled nominal values for O₂ concentration converge toward the observed
495 concentration (i.e., 172.3 $\mu\text{mol L}^{-1}$), that of NO₃ and to a lesser extent Alk-DIC
496 present persistent deviations from WOA2013 and GLODAP. At the surface, the
497 convergence of the simulated oxygen to observed value is expected since the

498 dominant governing process of thermodynamic saturation (through the air-sea gas
499 exchange) is well understood and modeled. The deviation in surface NO_3 highlights
500 uncertainty related to near surface biological processes and upper ocean physics.
501 Below the surface, concentrations of biogeochemical tracers drift away from the
502 globally averaged concentrations computed from WOA2013 or GLODAP (Figure 3,
503 panels d-i). At 150 and 2000 meters, the drift in global averaged concentrations for
504 these fields, computed over the last 250 years, is still significant with $p < 10^{-4}$ (Table 3).
505 Dashed lines in Figure 3 indicate the temporal evolution of RMSE, which quantifies
506 the total mismatch between simulated and observed fields. Except for the surface
507 fields, Figure 3 shows that RMSE globally increases with time for all biogeochemical
508 fields. The linear drift in RMSE over the last 250 years of the spin-up simulation falls
509 within the 2-3 % ky^{-1} range at the surface. It is much larger at 2000 m (144-280 % ky^{-1}
510 ; Table 3). This is also the case regionally, because the latitudinal maximum in RMSE
511 (RMSE_{max}) is similar to the global RMSE. Table 3 also shows that the magnitude of
512 drift in RMSE for O_2 , NO_3 and Alk-DIC differs at a given depth as different processes
513 affect the interior distribution of these biogeochemical fields.

514

515 **3-4 Evolution of geographical mismatches in IPSL-CM5A-LR**

516 To further explore the evolution of mismatch in biogeochemical distributions, we
517 analyze differences (ϵ) between simulated and observed fields of O_2 , NO_3 from
518 WOA2013 and Alk-DIC from GLODAP after the initialization and at the end of the
519 spin-up, i.e., the first year and the last year of the core spin-up simulation performed
520 with the IPSL-CM5A-LR model (Figures 4, 5 and 6).

521

522 Figure 4 (panels a, c, and e) shows that surface concentrations of biogeochemical

523 fields are associated with small biases at initialization. This error represents less than
524 5% of the observed surface concentrations for O₂, NO₃ and Alk-DIC and reflects the
525 weak difference between the data stream employed for initialization and validation.
526 After 500 years of spin-up, deviations between the modeled and observed fields at the
527 surface have increased locally by up to ~40% (Figure 4, panels b, d, and f). The
528 largest deviations are found in high-latitude oceans for O₂ and NO₃ and also to some
529 extent in the tropics for NO₃ and Alk-DIC.

530

531 Below the surface, distributions of modeled biogeochemical fields compare well to
532 the observations at 150 m at initialization with averaged errors close to zero (Figure 5,
533 panels a, c, and e). This result was expected since WOA2013 and WOA1994 differ
534 weakly at these depth levels. Subsurface distributions at initialization strongly contrast
535 with the concentrations that resulted from 500 years of spin-up (Figure 5, panels b, d,
536 and f). After 500 years of spin-up, strong mismatches characterize the distribution of
537 O₂, NO₃ and Alk-DIC fields in the high-latitude oceans and in the tropics. Figure 5
538 illustrates that pattern of errors are well correlated. It directly translates the
539 assumptions employed in the biogeochemical model (here the elemental C:N:-O₂
540 stoichiometry of PISCES). Figure 6 shows that model-data deviations at 2000 m have
541 substantially increased regionally after 500 years of simulation, showing large errors
542 in the southern hemisphere oceans. This appears clearly in Figure 6, panels d and f for
543 NO₃ and Alk-DIC fields, respectively.

544

545 The temporal evolution of the total mismatch between modeled and observed fields of
546 O₂, NO₃ and Alk-DIC over the whole water column is presented in Figure 7 in terms
547 of RMSE (Figure 7, panels a-c). As expected, Figure 7 illustrates that there is a good

548 match during the first years of simulation for all biogeochemical fields at all depth
549 levels with low RMSE. After a few centuries, patterns of error evolve differently
550 across depth for O₂, NO₃ and Alk-DIC.
551 The temporal evolution of RMSE shows that patterns of error have reached a steady
552 state after few decades within the upper hundred meters of the ocean but continue to
553 evolve at greater depths, even after 500 years. Patterns of errors within the
554 thermocline and deep water masses evolve at time scales of few decades and few
555 centuries, respectively in relation with the structure of the large-scale ocean
556 circulation. Mid-depth (~1500-2500m) RMSE evolves much slower because this
557 depth corresponds to the depth of the very old radiocarbon age (e.g., Wunsch and
558 Heimbach, 2007; 2008) whose characteristics time scale spans over thousand of years.
559 At the end of the spin-up simulation, two maxima of comparable amplitude are found
560 for RMSE at 150 and 3750 m for O₂ and at 50 m and 3800 m for Alk-DIC.

561

562 **3-5 Drifts in IPSL-CM5A-LR spin-up simulation**

563 With the evolution of the RMSE established, we can use the simple drift model
564 (Equation 1) to determine the relaxation time, τ , required to reach equilibration after a
565 longer of spin-up simulation. To use this simple drift model, we compute the drift in
566 RMSE determined from time segments of 100 years distributed evenly every 5 years
567 from year 250 to 500 for O₂, NO₃ and Alk-DIC tracers. The drift model (magenta
568 lines in Figure 8) is fitted level to the 80 drift values for each field and each depth
569 (colored crosses in Figure 8).

570

571 The simple drift model fits well the evolution of the drift in RMSE for the
572 biogeochemical variables along the spin-up simulation of IPSL-CM5A-LR (Figure 8).

573 Correlation coefficients are mostly significant at 90% confidence level ($r^*=0.14$
574 determined with a student distribution with significance level of 90% and 80 degrees
575 of freedom), except for NO_3 at surface and Alk-DIC at 150 m. Another exception is
576 found for NO_3 at 150 m where the drift does not correspond to an exponential decay
577 of the drift as function of time. The large confidence interval of the fit indicates that
578 the fit would have been considered as non-significant given a longer spin-up
579 simulation or a higher confidence threshold.

580

581 When significant, estimates of τ for O_2 RMSE are $\approx 90, 564$ and 1149 y at the surface
582 150 m and 2000 m, respectively. These values match reasonably well τ estimated for
583 NO_3 RMSE at 2000 m (1130 y) and those for Alk-DIC RMSE at surface and 2000 m
584 (137 and 1163 y). However, these estimates are sensitive to the time windows used to
585 compute the drift. For a subset of time windows between 100 and 250 years by step of
586 50 years, τ estimates for O_2 RMSE are $\approx 114\pm 67, 375\pm 140$ and 1116 ± 527 y at the
587 surface 150 m and 2000 m depth. These large uncertainties associated with τ
588 estimates are essentially due to the length of the spin-up simulation. A longer spin-up
589 simulation would improve the quality of the fit (see Figure S1).

590

591 **3-6 Drifts in CMIP5 ESMs preindustrial simulations**

592 In this subsection, the analysis is extended to the CMIP5 archive. We focus on oxygen
593 fields in the long preindustrial simulation, *piControl*, for the 15 available CMIP5
594 ESMs. From these simulations that span from 250 to 1000 years, we compute the drift
595 in O_2 RMSE across depth from several time segments of 100 years distributed evenly
596 every 5 years from the beginning until the end of the *piControl* simulation. These
597 drifts are used as a surrogate for drift computed from the spin-up of each model since

598 such simulations are not available through the data portal.

599

600 Figure 9 represents the drift in O₂ RMSE versus the spin-up duration for each CMIP5
601 ESM. The analysis shows that the drift in O₂ RMSE differs substantially between
602 models. For a given model, drifts in other biogeochemical tracers (NO₃ and Alk-DIC)
603 display similar features (not shown). The between-model differences in drift are not
604 surprising since there are no reasons for different models to exhibit similar drift for a
605 given field. Yet, Figure 9 shows that a global relationship emerges from this ensemble
606 when using the simple drift model to fit the drift in O₂ RMSE as function of the spin-
607 up duration (solid green lines in Figure 9). With a 90% confidence level, this
608 relationship suggests a general decrease of the drift as a function of spin-up duration
609 for all depth levels. At the surface and at 2000 m depth, the quality of fits is low with
610 correlation coefficients of about ~0.4. These are however significant at 90%
611 confidence level ($r^*=0.34$ determined with a student distribution with significance
612 level of 90% and 15 models as degree of freedom). The weakest correlation
613 coefficient is found for the fit at 150 m depth and hence indicating that there is no link
614 between the drift in O₂ RMSE and the duration of the spin-up simulation. This low
615 significance level must be put into perspective given the large diversity of spin-up
616 protocols and initial conditions (Figure 1 and Table 1) that can deteriorate the drift-
617 spin up duration relationship in this ensemble of models.

618

619 The drift versus spin up duration relationship established from the 15 CMIP5 ESMs is
620 nonetheless consistent with the results obtained with IPSL-CM5A-LR (The results in
621 Figure 8 have been reported in Figure 9 with magenta crosses). Consistency is
622 indicated by the sign of the drift versus spin up duration relationship of the IPSL-

623 CM5A-LR model at the various depth levels, although their magnitudes differ. This
624 difference in magnitude is not surprising if one considers that drift is highly model
625 and protocol dependent and that the length of the IPSL-CM5A-LR spin-up simulation
626 is potentially too short to determine accurate estimates of the long-term drift in O₂
627 RMSE. Despite these differences, our analyses show that a relationship between the
628 drift in O₂ RMSE versus the spin-up duration emerges from an ensemble of models
629 and is broadly consistent with our theoretical framework of a drift model established
630 from the results of the IPSL-CM5A-LR model (Figure 8).

631 **3-7 Impact of the drift on model skill score assessment metrics across CMIP5**

633 **ESMs**

634 In the following, we investigate the influence of model drift on skill score assessment
635 metrics that are routinely used to benchmark model performance. For this purpose, we
636 use the ensemble-mean O₂ RMSE as a metrics to assess the distance between the
637 biogeochemical observations and model results. For this purpose, we compute O₂
638 RMSE from each ensemble member of the CMIP5 models averaged from 1986 to
639 2005 with respect to WOA2013 observations. The model-data distance is then
640 determined for each CMIP5 model using the mean across the available ensemble
641 members.

642
643 The left hand side panels of Figure 10 present the performance of available CMIP5
644 models in terms of distance to oxygen observations at the surface, 150 m and 2000 m,
645 respectively. In these panels, the various CMIP5 models are ordered as function of
646 their distance to the oxygen observations. Following Knutti et al. (2013), either the
647 ensemble mean or the ensemble median is used to identify groups of models with
648 similar skill within the CMIP5 ensemble. The left hand side panels of Figure 10 show

649 that the ability of models to reproduce oxygen observations varies across depth levels.
 650 The RMSE in the simulated O₂ fields in CESM1-BGC, HadGEM2-ES, HadGEM2-
 651 CC, GFDL-ESM2M, MPI-ESM-LR and MPI-ESM-MR is generally smaller than the
 652 ensemble mean or ensemble median RMSE across the various depth levels (Figure 10
 653 panels a, b and c). On the other side of the ranking, CMCC-CESM, CNRM-CM5,
 654 CNRM-CM5-2, IPSL-CM5B-LR and NorESM1-ME exhibit RMSE generally higher
 655 than the ensemble mean and median RMSE across the various depth levels. The other
 656 models, i.e., CNRM-ESM1, GFDL-ESM2G, IPSL-CM5A-LR and IPSL-CM5A-MR
 657 display O₂ RMSE that is generally close to the ensemble mean or the ensemble
 658 median.

659

660 To assess the impact of model's drift inherited from the diversity of spin-up strategies
 661 (Figure 1 and Table 1) on the performance metrics, we use a simple additive
 662 assumption to incorporate an incremental error due to the drift, $\Delta RMSE$, to the above-
 663 mentioned RMSE. This incremental error due to the drift is computed using the
 664 relaxation time τ determined from the *piControl* simulations of each CMIP5 model at
 665 each depth level (Equation 1 and Figure 9) and a common duration of T=3000 years
 666 for all models (m):

$$667 \quad \Delta RMSE_m(z) = \int_0^T drift_m(z, t = 0) \times \exp\left(-\frac{1}{\tau(z)} t\right) dt \quad (2)$$

668 where $\Delta RMSE$ has the same unit as RMSE.

669 The common duration T is used to bring model drift close to zero and hence to make
 670 models comparable to each other.

671 We employ $\Delta RMSE$ to penalize the distance from the observations assuming that this
 672 drift-induced deviation in tracer fields can be added to RMSE. This means that the

673 effect of the penalty is to increase the distance giving a consistent measure of the
674 equilibration error.
675
676 Right hand side panels of Figure 10 show the influence of this penalization approach
677 on the model ranking at the various depth levels. They show that several models have
678 been upgraded in the ranking while others have not. For example, both MPI-ESM-LR,
679 MPI-ESM-MR have been upgraded at the surface and 2000 m. On the other hand, the
680 rank of HadGEM2-ES and HadGEM2-CC has been downgraded to the 5th and 3th
681 position due to the large drift in surface oxygen concentrations in comparison to that
682 of the other models. The surface drift might be attributed to drivers in oxygen fluxes
683 (e.g., SST, SSS). The ranking of GFDL-ESM2G and GFDL-ESM2M slightly changes
684 with penalization but both models stay close to the ensemble mean or the ensemble
685 median. At the bottom of the ranking, models with large deviation from the oxygen
686 observations (i.e., CMCC-CESM, IPSL-CM5B-LR, NorESM1-ME, CNRM-CM5) are
687 found. For these models, the computed Δ RMSE and RMSE result in similar ranking,
688 because even a small drift and hence relatively low Δ RMSE cannot compensate for
689 their large RMSE.

690

691 **4- Discussion**

692 **4-1 Implications for biogeochemical processes**

693 Our results show that errors in ocean biogeochemical fields amplify during the spin-
694 up simulation but not at the same rate at all depths. These differences in error
695 evolution are consistent with an increasing contribution of biogeochemical processes
696 in setting the distribution of tracers at depth. Indeed, Mignot et al. (2013) with the
697 same model simulation showed that the large-scale ocean circulation reaches quasi-

698 equilibrium after 250 years of spin-up, but our analyses indicate that biogeochemical
699 tracers do not (Figure 3).

700

701 Besides, our analysis demonstrates that error propagation and biogeochemical drift are
702 highly model dependent. For example, despite having the same initialization strategy
703 and comparable spin up duration, the GFDL-ESM2G, GFDL-ESM2M, and
704 NorESM1-ME models display considerable difference in drift (Figures 9 and 10) that
705 mirror large differences in model performance and properties (e.g., resolution,
706 simulated processes).

707

708 The identification of the dynamical or biogeochemical processes responsible for these
709 errors is not within the scope of this study and would required additional long
710 simulations with additional tracers targeted for attribution of the various
711 biogeochemical processes and the underlying ocean physics (e.g., Doney et al., 2004)
712 involved (e.g. using abiotic, passive tracers as suggested in Walin et al. (2014)). Some
713 mechanisms can be nonetheless invoked to explain differences or similarities in
714 behavior between biogeochemical fields. For example, the evolution of surface
715 concentrations for O₂ and Alk-DIC is controlled by the solubility of O₂ and CO₂ in
716 seawater and the concentration of these gases in the atmosphere (set to the observed
717 values and kept constant in all experiments performed with IPSL-CM5A-LR
718 discussed here) and the biological soft-tissue and calcium carbonate counter pumps
719 (in relation with the vertical transport of nutrients and alkalinity). Therefore, the
720 equilibration of the O₂ and Alk-DIC surface fields once the physical equilibrium is
721 reached (~250 years of spin-up) is expected (Figure 3, panels a and c and Figure 7).
722 Nevertheless, spatial errors could increase depending on the physical state of the

723 model (Figure 4, panels b and f). By contrast, the evolution of NO_3 concentration is
724 predominantly determined by ocean circulation, biological processes, and to a lesser
725 extent by external supplies from rivers and atmosphere. Below the surface,
726 concentrations of O_2 , NO_3 , and Alk-DIC evolve in response to the combined effect of
727 ocean circulation and biogeochemical processes. The combination of dynamical and
728 biogeochemical processes on the one hand, and the spin-up strategy on the other hand
729 both shape the modeled distributions of large-scale biogeochemical tracers.

730

731 Consequences of the difficulty in achieving the correct equilibration procedure are
732 even larger for biogeochemical features that are defined by regional characteristics in
733 tracer concentrations, such as high nutrient/low chlorophyll regions, oxygen minimum
734 zones and nutrient-to-light colimitation patterns. This point is illustrated by recent
735 studies focusing on future changes in phytoplankton productivity (e.g. Vancoppenolle
736 et al. (2013) and Laufkötter et al. (2015). Vancoppenolle and co-workers report a
737 wide spread of surface mean NO_3 concentrations (1980-1999) in the Arctic with a
738 range from 1.7 to 8.9 $\mu\text{mol L}^{-1}$ across a subset of 11 CMIP5 models. The spread in
739 present day NO_3 concentrations translates into a large model-to-model uncertainty in
740 future net primary production. Laufkötter and colleagues determined limitation terms
741 of phytoplankton production for a subset of CMIP5 and MAREMIP (Marine
742 Ecosystem Model Intercomparison Project) models. The authors demonstrate that
743 nutrient-to-light colimitation patterns differ in strength, location and type between
744 models and arise from large differences in the simulated nutrient concentrations.
745 Although large differences between models were reported by Vancoppenolle et al.
746 (2013) and Laufkötter et al. (2015) such as the spatial resolution and the complexity
747 of biogeochemical models, differences in nutrient concentrations were identified as

748 the largest source of model-to-model spread in addition to simply model error. The
749 authors of both studies qualitatively invoked differences in spin-up duration to explain
750 this spread. Besides, a recent assessment of interannual to decadal variability of ocean
751 CO₂ and O₂ fluxes in CMIP5 models, suggests that decadal variability can range
752 regionally from 10 to 50% of the total natural variability among a subset of 6 ESMs
753 (Resplandy et al., 2015). In that study, the authors demonstrate that, despite the
754 robustness of driving mechanisms (mostly related to vertical transport of water
755 masses) across the model ensemble, model-to-model spread can be related to
756 differences in modeled carbon and oxygen concentrations. In light of present results,
757 it appears likely that differences in spin-up strategy and sources of initialization could
758 also contribute to the amplitude of the natural variability of the ocean CO₂ and O₂
759 fluxes.

760

761 **4-2 Implications for future projections**

762 The inconsistent strategy to spin-up models in CMIP5 is a significant source of model
763 uncertainty. It needs to be better constrained in order to draw robust conclusions on
764 the impact of climate change on the carbon cycle as well as its climate feedback (e.g.,
765 Arora et al., 2013; Friedlingstein et al., 2013; Roy et al., 2011; Schwinger et al., 2014;
766 Séférian et al., 2012) and on marine ecosystems (e.g., Bopp et al., 2013; Boyd et al.,
767 2015; Cheung et al., 2012; Doney et al., 2012; Gattuso et al., 2015; Lehodey et al.,
768 2006). So far, the most frequent approach relies on the use of long preindustrial
769 control simulations to ‘remove’ the drift embedded in the simulated fields over the
770 historical period or future projections (e.g., Bopp et al., 2013; Cocco et al., 2013;
771 Friedlingstein et al., 2013; 2006; Frölicher et al., 2014; Gehlen et al., 2014; Keller et
772 al., 2014; Steinacher et al., 2010; Tjiputra et al., 2014). Although this approach allows

773 to determine relative changes, it does not allow to investigate the underlying reasons
774 of the spread between models in terms of processes, variability and response to
775 climate change. The “drift-correction” approach, much as the one used for this study,
776 assumes that drift-induced errors in the simulated fields can be isolated from the
777 signal of interest. Verification of this fundamental hypothesis would require a specific
778 experimental set-up consisting of the perturbation of model fields (e.g., nutrients or
779 carbon-related fields) to assess by how much the model projections would be
780 modified. So far, several modeling groups have generated ensemble simulation in
781 CMIP5 using a similar approach. However, the perturbations were applied either to
782 physical fields only or to both the physical and marine biogeochemical fields. To
783 assess impacts of different spin-up strategies and/or initial conditions on future
784 projections of marine biogeochemical tracer distributions, ensemble simulations in
785 which only biogeochemical fields are perturbed would be needed.

786

787 **4-3 Implications for multi-model skill-score assessments.**

788 While the importance of spin-up protocols is well accepted in the modeling
789 community, the link between spin-up strategy and the ability of a model to reproduce
790 modern observations remains to be addressed.

791

792 Most of the recent CMIP5 skill assessment approaches were based on *historical*
793 hindcasts that were started from preindustrial runs of varying duration and from
794 various spin-up strategies. Therefore, in typical intercomparison exercises, Earth
795 system models with a short spin-up, and hence modeled distributions still close to
796 initial fields, are confronted with Earth system models with a longer spin-up duration
797 and modeled distributions that have drifted further away from their initial states. Our

798 study highlights that such inconsistencies in spin-up protocols and initial conditions
799 across CMIP5 Earth system models (Figure 1 and Table 1) could significantly
800 contribute to model-to-model spread in performance metrics. The analysis of the first
801 century of CMIP5 *piControl* simulations demonstrated a significant spread of drift
802 between CMIP5 models (Figure 9). An approximate exponential relationship between
803 the amplitude of drift and the spin up duration emerges from the ensemble of CMIP5
804 models, which is consistent with results from IPSL-CM5A-LR. For example, while
805 the global average root-mean square error increased up to 70% during a 500-year
806 spin-up simulation with IPSL-CM5A-LR, its rate of increase (or drift) decreased with
807 time to a very small rate ($0.001 \text{ Pg C y}^{-1}$). Combining a simple drift model and this
808 relationship, we propose a penalization approach in an effort to assess more
809 objectively the influence of documented model differences on model-data biases.
810 Figure 10 compares the state-of-the-art approach to assess model performance (left
811 hand side panels) to the drift-penalized approach (right hand side panels). This novel
812 approach penalizes models with larger drift without affecting the models with smaller
813 drift. Taking into account drift in modeled fields results in subtle adjustments in
814 ranking, which reflect differences in spin-up and initialization strategies.

815

816 **4-4 Limitations of the framework**

817 In this work, the analyses focus on the globally averaged O_2 RMSE across a diverse
818 ensemble of CMIP5 models, which differ in terms of represented processes, spatial
819 resolution and performance in addition to differences in spin-up protocols. Major
820 limitations of the framework are presented below.

821

822 Due to their specificities in terms of processes and resolution (e.g., Cabré et al.,

823 (2015), Laufkötter et al. (2015)), regional drift in CMIP5 models may differ from the
824 drift computed from globally averaged skill-score metrics (see Figure S2 and S3).
825 These differences may lead to different estimates of the relaxation time τ at regional
826 scale. Moreover, the combination of regional ocean physics and biogeochemical
827 processes in each individual model may drive an evolution of regional drift in RMSE
828 that does not fit the hypothesis of an exponential decay of the drift during the course
829 of the spin-up simulation.

830

831 The above-mentioned remark can explain the relatively low confidence level of the fit
832 to drift across the multi-model CMIP5 ensemble (Figure 9). The relatively low
833 significance level of the fit directly reflects not only the large diversity of spin-up
834 protocols and initial conditions (Figure 1 and Table 1) but also the large diversity of
835 processes and resolution of the CMIP5 models. An improved derivation of the
836 penalization would require access to output from spin-up simulations for each
837 individual model or, at least, a better quantification of model-model differences in
838 terms of initial conditions.

839

840 Finally, it is unlikely that model fields drift at the same rate along the spin-up
841 simulation, even under the same spin-up protocols. Indeed, as shown in Kriest and
842 Oeschies (2015), various parameterizations of the particles sinking speeds in a
843 common physical framework may lead to a similar evolution of the globally averaged
844 RMSE in the first century of the spin-up simulation but display very different
845 behaviour within a time-scale of $O(10^3)$ years. As such, drift and τ estimates need to
846 be used with caution when computed from short spin-up simulation because they can
847 be subject to large uncertainties.

848

849 **5- Conclusions and recommendation for future intercomparison exercises**

850 Skill-score metrics are expected to be widely used in the framework of the future
851 CMIP6 (Meehl et al., 2014) with the development of international community
852 benchmarking tools like the ESMValTool (<http://www.pa.op.dlr.de/ESMValTool>), see
853 also Eyring et al. (2015)). The assessment of model skill to reproduce observations
854 will focus on the modern period. Complementary to this approach, our results call for
855 the consideration of spin-up and initialization strategies in the determination of skill
856 assessment metrics (e.g., Friedrichs et al., 2009; Stow et al., 2009) and, by extension,
857 to model weighting (e.g., Steinacher et al., 2010) and model ranking (e.g., Anav et al.,
858 2013). Indeed, the use of equilibrium-state metrics of the model like the 3-
859 dimensional growth rate or drift of relevant skill score metrics (e.g. RMSE) could be
860 employed to increase the reliability of these traditional metrics and, as such, should be
861 included in the set of standard assessment tools for CMIP6.

862

863 In an effort to better represent interactions between marine biogeochemistry and
864 climate (Smith et al., 2014), future generations of Earth system models are likely to
865 include more complex ocean biogeochemical models, be it in terms of processes (e.g.,
866 Tagliabue and Völker, 2011; Tagliabue et al., 2011) or interactions with other
867 biogeochemical cycles (e.g., Gruber and Galloway, 2008) or increased spatial
868 resolution (e.g., Dufour et al., 2013; Lévy et al., 2012) in order to better represent
869 mesoscale biogeochemical dynamics. These developments will go along with an
870 increase in the diversity and complexity of spin-up protocols applied to Earth system
871 models, especially those including an interactive atmospheric CO₂ or interactive
872 nitrogen cycle (e.g., Dunne et al., 2013; Lindsay et al., 2014). The additional

873 challenge of spinning-up emission-driven simulations with interactive carbon cycle
874 will also require us to extend the assessment of the impact of spin-up protocols to the
875 terrestrial carbon cycle. Processes such as soil carbon accumulation, peat formation as
876 well as shift in biomes such as tropical and boreal ecosystems for dynamic vegetation
877 models require several long time-scales to equilibrate (Brovkin et al., 2010; Koven et
878 al., 2015). In addition, the terrestrial carbon cycle has large uncertainties in terms of
879 carbon sink/source behavior (Anav et al., 2013; Dalmonech et al., 2014; Friedlingstein
880 et al., 2013) which might affect ocean CO₂ uptake (Brovkin et al., 2010). A novel
881 numerical algorithm to accelerate the spin-up integration time for computationally
882 expensive ocean biogeochemical models has emerged (Khatiwala, 2008), which could
883 further complicate the determination of inter-model spreads.

884

885 To evaluate the contribution of variable spin-up and initialization strategies to model
886 performance, these should be documented extensively and the corresponding model
887 output should be archived. Ideally, for future coupled model intercomparison
888 exercises (i.e., CMIP6, CMIP7, Meehl et al., (2014)), the community should agree on
889 a set of simple recommendations for spin-up protocols, following past projects such
890 as OCMIP-2. In parallel, any trade-off between model equilibration and
891 computationally efficient spin-up procedures has to be linked with efforts to reduce
892 model errors due to the physical and biogeochemical parameterizations.

893

894

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916 References:

- 917 Adachi, Y., Yukimoto, S., Deushi, M., Obata, A., Nakano, H., Tanaka, T. Y., Hosaka,
918 M., Sakami, T., Yoshimura, H., Hirabara, M., Shindo, E., Tsujino, H., Mizuta, R.,
919 Yabu, S., Koshiro, T., Ose, T. and Kitoh, A.: Basic performance of a new earth
920 system model of the Meteorological Research Institute (MRI-ESM1), Papers in
921 Meteorology and Geophysics, 64, 1–18, doi:10.2467/mripapers.64.1, 2013.
- 922 Anav, A., Friedlingstein, P., Kidston, M., Bopp, L., Ciais, P., Cox, P., Jones, C., Jung,
923 M., Myneni, R. and Zhu, Z.: Evaluating the Land and Ocean Components of the
924 Global Carbon Cycle in the CMIP5 Earth System Models, J. Climate, 26(18), 6801–
925 6843, doi:10.1175/JCLI-D-12-00417.1, 2013.
- 926 Andrews, O. D., Bindoff, N. L., Halloran, P. R., Ilyina, T. and Le Qu 'er 'e, C.:
927 Detecting an external influence on recent changes in oceanic oxygen using an optimal
928 fingerprinting method, Biogeosciences, 10(3), 1799–1813, doi:10.5194/bg-10-1799-
929 2013, 2013.
- 930 Archer, D., Buffett, B. and Brovkin, V.: Ocean methane hydrates as a slow tipping
931 point in the global carbon cycle, Proceedings of the National Academy of Sciences,
932 106(49), 20596–20601, 2009.
- 933 Arora, V. K., Boer, G. J., Friedlingstein, P., Eby, M., Jones, C. D., Christian, J. R.,
934 Bonan, G., Bopp, L., Brovkin, V., Cadule, P., Hajima, T., Ilyina, T., Lindsay, K.,
935 Tjiputra, J. F. and Wu, T.: Carbon–Concentration and Carbon–Climate Feedbacks in
936 CMIP5 Earth System Models, J. Climate, 26(15), 5289–5314, doi:10.1175/JCLI-D-
937 12-00494.1, 2013.
- 938 Arora, V. K., Scinocca, J. F., Boer, G. J., Christian, J. R., Denman, K. L., Flato, G.
939 M., Kharin, V. V., Lee, W. G. and Merryfield, W. J.: Carbon emission limits required
940 to satisfy future representative concentration pathways of greenhouse gases, Geophys.

- 941 Res. Lett., 38(5), L05805, doi:10.1029/2010GL046270, 2011.
- 942 Aumont, O. and Bopp, L.: Globalizing results from ocean in situ iron fertilization
943 studies, *Global Biogeochem. Cycles*, 20(2), GB2017, doi:10.1029/2005GB002591,
944 2006.
- 945 Aumont, O., Maier-Reimer, E., Blain, S. and Monfray, P.: An ecosystem model of the
946 global ocean including Fe, Si, P colimitations, *Global Biogeochem. Cycles*, 17(2),
947 1060, doi:10.1029/2001GB001745, 2003.
- 948 Aumont, O., Orr, J. C., Monfray, P., Ludwig, W., Amiotte-Suchet, P. and Probst, J.-
949 L.: Riverine-driven interhemispheric transport of carbon, *Global Biogeochem. Cycles*,
950 15(2), 393–405, doi:10.1029/1999GB001238, 2001.
- 951 Aumont, O., Orr, J., Jamous, D., Monfray, P., Marti, O. and Madec, G.: A degradation
952 approach to accelerate simulations to steady-state in a 3-D tracer transport model of
953 the global ocean, *Climate Dynamics*, 14(2), 101–116, 1998.
- 954 Bopp, L., Resplandy, L., Orr, J. C., Doney, S. C., Dunne, J. P., Gehlen, M., Halloran,
955 P., Heinze, C., Ilyina, T., Séférian, R., Tjiputra, J. and Vichi, M.: Multiple stressors of
956 ocean ecosystems in the 21st century: projections with CMIP5 models,
957 *Biogeosciences*, 10(10), 6225–6245, doi:10.5194/bg-10-6225-2013, 2013.
- 958 Boyd, P. W., Lennartz, S. T., Glover, D. M. and Doney, S. C.: Biological
959 ramifications of climate-change-mediated oceanic multi-stressors, *Nature Clim.*
960 *Change*, 5(1), 71–79, 2015.
- 961 Brovkin, V., Lorenz, S. J., Jungclauss, J., Raddatz, T., Timmreck, C., Reick, C. H.,
962 Segschneider, J. and Six, K.: Sensitivity of a coupled climate-carbon cycle model to
963 large volcanic eruptions during the last millennium, *Tellus B*, 62(5), 674–681,
964 doi:10.1111/j.1600-0889.2010.00471.x, 2010.
- 965 Bryan, K.: Accelerating the Convergence to Equilibrium of Ocean-Climate Models, *J.*
966 *Phys. Oceanogr.*, 14(4), 666–673, doi:10.1175/1520-
967 0485(1984)014<0666:ATCTEO>2.0.CO;2, 1984.
- 968 Cheung, W. W. L., Sarmiento, J. L., Dunne, J. P., Frölicher, T. L., Lam, V. W. Y.,
969 Palomares, M. L. D., Watson, R. and Pauly, D.: Shrinking of fishes exacerbates
970 impacts of global ocean changes on marine ecosystems, *Nature Climate change*,
971 2(10), 1–5, doi:10.1038/nclimate1691, 2012.
- 972 Cabré, A., Marinov, I., Bernardello, R. and Bianchi, D.: Oxygen minimum zones in
973 the tropical Pacific across CMIP5 models: mean state differences and climate change
974 trends, *Biogeosciences*, 12(18), 5429–5454, doi:10.5194/bg-12-5429-2015, 2015.
- 975 Cocco, V., Joos, F., Steinacher, M., Frölicher, T. L., Bopp, L., Dunne, J., Gehlen, M.,
976 Heinze, C., Orr, J., Oschlies, A., Schneider, B., Segschneider, J. and Tjiputra, J.:
977 Oxygen and indicators of stress for marine life in multi-model global warming
978 projections, *Biogeosciences*, 10(3), 1849–1868, doi:10.5194/bg-10-1849-2013, 2013.
- 979 Collins, W. J., Bellouin, N., Doutriaux-Boucher, M., Gedney, N., Halloran, P.,
980 Hinton, T., Hughes, J., Jones, C. D., Joshi, M., Liddicoat, S., Martin, G., O'Connor,

- 981 F., Rae, J., Senior, C., Sitch, S., Totterdell, I., Wiltshire, A. and Woodward, S.:
 982 Development and evaluation of an Earth-System model – HadGEM2, *Geosci. Model*
 983 *Dev*, 4(4), 1051–1075, doi:10.5194/gmd-4-1051-2011, 2011.
- 984 Cox, P. M., Pearson, D., Booth, B. B., Friedlingstein, P., Huntingford, C., Jones, C. D.
 985 and Luke, C. M.: Sensitivity of tropical carbon to climate change constrained by
 986 carbon dioxide variability, *Nature*, 494(7437), 341–344, doi:10.1038/nature11882,
 987 2013.
- 988 Dalmonech, D., Foley, A. M., Anav, A., Friedlingstein, P., Friend, A. D., Kidston, M.,
 989 Willeit, M. and Zaehle, S.: Challenges and opportunities to reduce uncertainty in
 990 projections of future atmospheric CO₂: a combined marine and terrestrial biosphere
 991 perspective, *Biogeosciences Discuss.*, 11(2), 2083–2153, doi:10.5194/bgd-11-2083-
 992 2014, 2014.
- 993 de Baar, H. J. W. and de Jong, J. T. M.: *The biogeochemistry of iron in seawater*,
 994 edited by D. R. Turner and K. A. Hunter, John Wiley, Hoboken, N. J., 2001.
- 995 Doney, S. C., Lindsay, K., Caldeira, K., Campin, J.-M., Drange, H., Dutay, J.-C.,
 996 Follows, M., Gao, Y., Gnanadesikan, A., Gruber, N., Ishida, A., Joos, F., Madec, G.,
 997 Maier-Reimer, E., Marshall, J. C., Matear, R. J., Monfray, P., Mouchet, A., Najjar, R.,
 998 Orr, J. C., Plattner, G.-K., Sarmiento, J., Schlitzer, R., Slater, R., Totterdell, I. J.,
 999 Weirig, M.-F., Yamanaka, Y. and Yool, A.: Evaluating global ocean carbon models:
 1000 The importance of realistic physics, *Global Biogeochem. Cycles*, 18(3),
 1001 doi:10.1029/2003GB002150, 2004.
- 1002 Doney, S. C.: The Growing Human Footprint on Coastal and Open-Ocean
 1003 Biogeochemistry, *Science*, 328(5985), 1512–1516, doi:10.1126/science.1185198,
 1004 2010.
- 1005 Doney, S. C., Lima, I., Moore, J. K., Lindsay, K., Behrenfeld, M. J., Westberry, T. K.,
 1006 Mahowald, N., Glover, D. M. and Takahashi, T.: Skill metrics for confronting global
 1007 upper ocean ecosystem-biogeochemistry models against field and remote sensing
 1008 data, *Journal of Marine Systems*, 76(1-2), 95–112,
 1009 doi:10.1016/j.jmarsys.2008.05.015, 2009.
- 1010 Doney, S. C., Ruckelshaus, M., Emmett Duffy, J., Barry, J. P., Chan, F., English, C.
 1011 A., Galindo, H. M., Grebmeier, J. M., Hollowed, A. B., Knowlton, N., Polovina, J.,
 1012 Rabalais, N. N., Sydeman, W. J. and Talley, L. D.: Climate Change Impacts on
 1013 Marine Ecosystems, *Annu. Rev. Marine. Sci.*, 4(1), 11–37, doi:10.1146/annurev-
 1014 marine-041911-111611, 2012.
- 1015 Dufour, C. O., Sommer, J. L., Gehlen, M., Orr, J. C., Molines, J.-M., Simeon, J. and
 1016 Barnier, B.: Eddy compensation and controls of the enhanced sea-to-air CO₂ flux
 1017 during positive phases of the Southern Annular Mode, *Global Biogeochem. Cycles*,
 1018 27(3), 950–961, doi:10.1002/gbc.20090, 2013.
- 1019 Dufresne, J.-L., Foujols, M. A., Denvil, S., Caubel, A., Marti, O., Aumont, O.,
 1020 Balkanski, Y., Bekki, S., Bellenger, H., Benshila, R., Bony, S., Bopp, L., Braconnot,
 1021 P., Brockmann, P., Cadule, P., Cheruy, F., Codron, F., Cozic, A., Cugnet, D., Noblet,
 1022 N., Duvel, J. P., Ethe, C., Fairhead, L., Fichet, T., Flavoni, S., Friedlingstein, P.,

1023 Grandpeix, J. Y., Guez, L., Guilyardi, E., Hauglustaine, D., Hourdin, F., Idelkadi, A.,
1024 Ghattas, J., Joussaume, S., Kageyama, M., Krinner, G., Labetoulle, S., Lahellec, A.,
1025 Lefebvre, M.-P., Lefèvre, F., Lévy, C., Li, Z. X., Lloyd, J., Lott, F., Madec, G.,
1026 Mancip, M., Marchand, M., Masson, S., Meurdesoif, Y., Mignot, J., Musat, I.,
1027 Parouty, S., Polcher, J., Rio, C., Schulz, M., Swingedouw, D., Szopa, S., Talandier,
1028 C., Terray, P., Viovy, N. and Vuichard, N.: Climate change projections using the
1029 IPSL-CM5 Earth System Model: from CMIP3 to CMIP5, *Clim Dyn*, 40(9-10), 2123–
1030 2165, doi:10.1007/s00382-012-1636-1, 2013.

1031 Dunne, J. P., John, J. G., Adcroft, A. J., Griffies, S. M., Hallberg, R. W., Shevliakova,
1032 E., Stouffer, R. J., Cooke, W., Dunne, K. A., Harrison, M. J., Krasting, J. P.,
1033 Malyshev, S. L., Milly, P. C. D., Phillipps, P. J., Sentman, L. A., Samuels, B. L.,
1034 Spelman, M. J., Winton, M., Wittenberg, A. T. and Zadeh, N.: GFDL’s ESM2 Global
1035 Coupled Climate–Carbon Earth System Models. Part I: Physical Formulation and
1036 Baseline Simulation Characteristics, *J. Climate*, 25(19), 6646–6665, doi:doi:
1037 10.1175/JCLI-D-11-00560.1, 2013.

1038 Duplessy, J. C., Bard, E., Arnold, M., Shackleton, N. J., Duprat, J. and Labeyrie, L.:
1039 How fast did the ocean–atmosphere system run during the last deglaciation? *Earth
1040 and Planetary Science Letters*, 103(1-4), 27–40, doi:10.1016/0012-821X(91)90147-A,
1041 1991.

1042 Eyring, V., Righi, M., Evaldsson, M., Lauer, A., Wenzel, S., Jones, C., Anav, A.,
1043 Andrews, O., Cionni, I., Davin, E. L., Deser, C., Ehbrecht, C., Friedlingstein, P.,
1044 Gleckler, P., Gottschaldt, K. D., Hagemann, S., Juckes, M., Kindermann, S., Krasting,
1045 J., Kunert, D., Levine, R., Loew, A., Mäkelä, J., Martin, G., Mason, E., Phillips, A.,
1046 Read, S., Rio, C., Roehrig, R., Senftleben, D., Sterl, A., van Ulft, L. H., Walton, J.,
1047 Wang, S. and Williams, K. D.: ESMValTool (v1.0) – a community diagnostic and
1048 performance metrics tool for routine evaluation of Earth System Models in CMIP,
1049 *Geosci. Model Dev. Discuss.*, 8(9), 7541–7661, 2015.

1050 Fichet, T. and Maqueda, M. A. M.: Sensitivity of a global sea ice model to the
1051 treatment of ice thermodynamics and dynamics, *J. Geophys. Res.*, 102(C6), 12609–
1052 12646, 1997.

1053 Follows, M. J., Dutkiewicz, S., Grant, S. and Chisholm, S. W.: Emergent
1054 Biogeography of Microbial Communities in a Model Ocean, *Science*, 315(5820),
1055 1843–1846, doi:10.1126/science.1138544, 2007.

1056 Friedlingstein, P., Cox, P., Betts, R., Bopp, L., Bloh, Von, W., Brovkin, V., Cadule,
1057 P., Doney, S., Eby, M., Fung, I., Bala, G., John, J., Jones, C., Joos, F., Kato, T.,
1058 Kawamiya, M., Knorr, W., Lindsay, K., Matthews, H. D., Raddatz, T., Rayner, P.,
1059 Reick, C., Roeckner, E., Schnitzler, K. G., Schnur, R., Strassmann, K., Weaver, A. J.,
1060 Yoshikawa, C. and Zeng, N.: Climate–Carbon Cycle Feedback Analysis: Results from
1061 the C 4MIP Model Intercomparison, *J. Climate*, 10(14), 3337–3353,
1062 doi:10.1175/JCLI3800.1, 2006.

1063 Friedlingstein, P., Meinshausen, M., Arora, V. K., Jones, C. D., Anav, A., Liddicoat,
1064 S. K. and Knutti, R.: Uncertainties in CMIP5 climate projections due to carbon cycle
1065 feedbacks, *J. Climate*, 130917124100006, doi:doi: 10.1175/JCLI-D-12-00579.1,
1066 2013.

- 1067 Friedrichs, M. A. M., Carr, M.-E., Barber, R. T., Scardi, M., Antoine, D., Armstrong,
 1068 R. A., Asanuma, I., Behrenfeld, M. J., Buitenhuis, E. T., Chai, F., Christian, J. R.,
 1069 Ciotti, A. M., Doney, S. C., Dowell, M., Dunne, J. P., Gentili, B., Gregg, W.,
 1070 Hoepffner, N., Ishizaka, J., Kameda, T., Lima, I., Marra, J., Mélin, F., Moore, J. K.,
 1071 Morel, A., O'Malley, R. T., O'Reilly, J., Saba, V. S., Schmeltz, M., Smyth, T. J.,
 1072 Tjiputra, J., Waters, K., Westberry, T. K. and Winguth, A.: Assessing the
 1073 uncertainties of model estimates of primary productivity in the tropical Pacific Ocean,
 1074 *Journal of Marine Systems*, 76(1-2), 113–133, doi:10.1016/j.jmarsys.2008.05.010,
 1075 2009.
- 1076 Friedrichs, M. A. M., Dusenberry, J. A., Anderson, L. A., Armstrong, R. A., Chai, F.,
 1077 Christian, J. R., Doney, S. C., Dunne, J. P., Fujii, M., Hood, R., McGillicuddy, D. J.,
 1078 Jr., Moore, J. K., Schartau, M., Spitz, Y. H. and Wiggert, J. D.: Assessment of skill
 1079 and portability in regional marine biogeochemical models: Role of multiple
 1080 planktonic groups, *J. Geophys. Res.*, 112(C8), doi:10.1029/2006JC003852, 2007.
- 1081 Frölicher, T. L., Sarmiento, J. L., Paynter, D. J., Dunne, J. P., Krasting, J. P. and
 1082 Winton, M.: Dominance of the Southern Ocean in anthropogenic carbon and heat
 1083 uptake in CMIP5 models, *J. Climate*, 141031131835005, doi:10.1175/JCLI-D-14-
 1084 00117.1, 2014.
- 1085 Gattuso, J. P., Magnan, A., Bille, R., Cheung, W. W. L., Howes, E. L., Joos, F.,
 1086 Allemand, D., Bopp, L., Cooley, S. R., Eakin, C. M., Hoegh-Guldberg, O., Kelly, R.
 1087 P., Portner, H. O., Rogers, A. D., Baxter, J. M., Laffoley, D., Osborn, D., Rankovic,
 1088 A., Rochette, J., Sumaila, U. R., Treyer, S. and Turley, C.: Contrasting futures for
 1089 ocean and society from different anthropogenic CO₂ emissions scenarios, *Science*,
 1090 349(6243), aac4722–aac4722, doi:10.1126/science.aac4722, 2015.
- 1091 Gehlen, M., Séférian, R., Jones, D. O. B., Roy, T., Roth, R., Barry, J., Bopp, L.,
 1092 Doney, S. C., Dunne, J. P., Heinze, C., Joos, F., Orr, J. C., Resplandy, L.,
 1093 Segschneider, J. and Tjiputra, J.: Projected pH reductions by 2100 might put deep
 1094 North Atlantic biodiversity at risk, *Biogeosciences*, 11(23), 6955–6967, 2014.
- 1095 Gerber, M. and Joos, F.: Carbon sources and sinks from an Ensemble Kalman Filter
 1096 ocean data assimilation - Gerber - 2010 - *Global Biogeochemical Cycles* - Wiley
 1097 Online Library, *Global Biogeochem. Cycles*, 24, GB3004,
 1098 doi:[10.1029/2009GB003531](https://doi.org/10.1029/2009GB003531), 2010.
- 1099 Gnanadesikan, A.: Oceanic ventilation and biogeochemical cycling: Understanding
 1100 the physical mechanisms that produce realistic distributions of tracers and
 1101 productivity, *Global Biogeochem. Cycles*, 18(4), doi:10.1029/2003GB002097, 2004.
- 1102 Gruber, N.: Warming up, turning sour, losing breath: ocean biogeochemistry under
 1103 global change, *Philosophical Transactions of the Royal Society A: Mathematical,*
 1104 *Physical and Engineering Sciences*, 369(1943), 1980–1996,
 1105 doi:10.1098/rsta.2011.0003, 2011.
- 1106 Gruber, N. and Galloway, J. N.: An Earth-system perspective of the global nitrogen
 1107 cycle, *Nature*, 451(7176), 293–296, doi:doi:10.1038/nature06592, 2008.
- 1108 Sen Gupta, A. S., Muir, L. C., Brown, J. N., Phipps, S. J., Durack, P. J., Monselesan,

- 1109 D. and Wijffels, S. E.: Climate Drift in the CMIP3 Models, *J. Climate*, 25(13), 4621–
1110 4640, doi:10.1175/JCLI-D-11-00312.1, 2012.
- 1111 Sen Gupta, A. S., Jourdain, N. C., Brown, J. N. and Monselesan, D.: Climate Drift in
1112 the CMIP5 models, *J. Climate*, 26(21), 8597–8615. <http://doi.org/10.1175/JCLI-D-12-00521.s1>.
1113
- 1114 Hajima, T., Kawamiya, M., Watanabe, M., Kato, E., Tachiiri, K., Sugiyama, M.,
1115 Watanabe, S., Okajima, H. and Ito, A.: Modeling in Earth system science up to and
1116 beyond IPCC AR5, *Progress in Earth and Planetary Science*, 1(1), 29–25,
1117 doi:10.1186/s40645-014-0029-y, 2014.
- 1118 Heinze, C., Maier-Reimer, E., Winguth, A. and Archer, D.: A global oceanic sediment
1119 model for long-term climate studies, *Global Biogeochem. Cycles*, 13(1), 221–250,
1120 1999.
- 1121 Heinze, M. and Ilyina, T.: Ocean biogeochemistry in the warm climate of the late
1122 Paleocene, *Climate of the Past*, 11(1), 1933–1975, doi:10.5194/cp-11-63-2015, 2015.
- 1123 Henson, S. A., Sarmiento, J. L., Dunne, J. P., Bopp, L., Lima, I., Doney, S. C., John,
1124 J. and Beaulieu, C.: Detection of anthropogenic climate change in satellite records of
1125 ocean chlorophyll and productivity, *Biogeosciences*, 7(2), 621–640, doi:10.5194/bg-7-621-2010, 2010.
1126
- 1127 Hourdin, F., Musat, I., Bony, S., Braconnot, P., Codron, F., Dufresne, J.-L., Fairhead,
1128 L., Filiberti, M.-A., Friedlingstein, P., Grandpeix, J.-Y., Krinner, G., LeVan, P., Li,
1129 Z.-X. and Lott, F.: The LMDZ4 general circulation model: climate performance and
1130 sensitivity to parametrized physics with emphasis on tropical convection, *Climate*
1131 *Dynamics*, 27, 787–813, doi:10.1007/s00382-006-0158-0, 2006.
- 1132 Ilyina, T., Six, K. D., Segschneider, J., Maier-Reimer, E., Li, H. and Núñez-Riboni, I.:
1133 Global ocean biogeochemistry model HAMOCC: Model architecture and
1134 performance as component of the MPI-Earth system model in different CMIP5
1135 experimental realizations, *J. Adv. Model. Earth Syst.*, 5(2), 287–315,
1136 doi:10.1029/2012MS000178, 2013.
- 1137 IPCC: Climate Change 2013: The Physical Science Basis, edited by: Stoker, T. F.,
1138 Qin, D., Plattner, G., Tignor, M., Allen, S. K., Boschung, J., Nauels, A., Xia, Y.,
1139 Bex, V., and Midgley, P. M., Cambridge Univ. Press, Cambridge, UK, and New
1140 York, NY, USA, 2013..
- 1141 Ito, T. and Deutsch, C.: Variability of the Oxygen Minimum Zone in the Tropical
1142 North Pacific during the Late 20th Century, *Global Biogeochem. Cycles*, n/a–n/a,
1143 doi:10.1002/2013GB004567, 2013.
- 1144 Ito, T., Woloszyn, M. and Mazloff, M.: Anthropogenic carbon dioxide transport in the
1145 Southern Ocean driven by Ekman flow, *Nature*, 463(7277), 80–83,
1146 doi:10.1038/nature08687, 2010.
- 1147 Jickells, T. and Spokes, L.: The biogeochemistry of iron in seawater, edited by D. R.
1148 Turner and K. A. Hunter, John Wiley, Hoboken, N. J., 2001.

- 1149 Johnson, K., Chavez, F. and Friederich, G.: Continental-shelf sediment as a primary
1150 source of iron for coastal phytoplankton, *Nature*, 398(6729), 697–700, 1999.
- 1151 Keeling, R. F., Körtzinger, A. and Gruber, N.: Ocean Deoxygenation in a Warming
1152 World, *Annu. Rev. Marine. Sci.*, 2(1), 199–229,
1153 doi:10.1146/annurev.marine.010908.163855, 2009.
- 1154 Keenlyside, N. S., Latif, M., Jungclaus, J., Kornbluh, L. and Roeckner, E.:
1155 Advancing decadal-scale climate prediction in the North Atlantic sector, *Nature*,
1156 453(7191), 84–88, doi:10.1038/nature06921, 2008.
- 1157 Keller, K. M., Joos, F. and Raible, C. C.: Time of emergence of trends in ocean
1158 biogeochemistry, *Biogeosciences*, 11(13), 3647–3659, doi:10.5194/bgd-10-18065-
1159 2013, 2014.
- 1160 Key, R., Kozyr, A., Sabine, C., Lee, K., Wanninkhof, R., Bullister, J., Feely, R.,
1161 Millero, F., Mordy, C. and Peng, T.: A global ocean carbon climatology: Results from
1162 Global Data Analysis Project (GLODAP), *Global Biogeochem. Cycles*, 18(4),
1163 doi:10.1029/2004GB002247, 2004.
- 1164 Khatiwala, S., Visbeck, M. and Cane, M. A.: Accelerated simulation of passive
1165 tracers in ocean circulation models, *Ocean Modelling*, 9(1), 51–69,
1166 doi:10.1016/j.ocemod.2004.04.002, 2005.
- 1167 Khatiwala, S.: Fast spin up of ocean biogeochemical models using matrix-free
1168 Newton-Krylov, *Ocean Modelling*, 23, 121-129, 2008.
- 1169 Kim, H.-M., Webster, P. J. and Curry, J. A.: Evaluation of short-term climate change
1170 prediction in multi-model CMIP5 decadal hindcasts, *Geophys. Res. Lett.*, 39(10),
1171 L10701, doi:10.1029/2012GL051644, 2012.
- 1172 Knutti, R., Masson, D. and Gettelman, A.: Climate model genealogy: Generation
1173 CMIP5 and how we got there, *Geophys. Res. Lett.*, 40(6), 1194–1199,
1174 doi:10.1002/grl.50256, 2013.
- 1175 Koven, C. D., Chambers, J. Q., Georgiou, K., Knox, R., Negron-Juarez, R., Riley, W.
1176 J., Arora, V. K., Brovkin, V., Friedlingstein, P. and Jones, C. D.: Controls on
1177 terrestrial carbon feedbacks by productivity vs. turnover in the CMIP5 Earth System
1178 Models, *Biogeosciences Discuss.*, 12(8), 5757–5801, 2015.
- 1179 Kriest, I. and Oschlies, A.: MOPS-1.0: towards a model for the regulation of the
1180 global oceanic nitrogen budget by marine biogeochemical processes, *Geosci. Model
1181 Dev*, 8(9), 2929–2957, doi:10.5194/gmd-8-2929-2015, 2015.
- 1182 Krinner, G., Viovy, N., de Noblet-Ducoudré, N., Ogée, J., Polcher, J., Friedlingstein,
1183 P., Ciais, P., Sitch, S. and Prentice, I. C.: A dynamic global vegetation model for
1184 studies of the coupled atmosphere-biosphere system, *Global Biogeochem. Cycles*,
1185 19(1), 1–33, 2005.
- 1186 Laufkötter, C., Vogt, M., Gruber, N., Aita-Noguchi, M., Aumont, O., Bopp, L.,
1187 Buitenhuis, E., Doney, S. C., Dunne, J., Hashioka, T., Hauck, J., Hirata, T., John, J.,
1188 Le Quéré, C., Lima, I. D., Nakano, H., Séférian, R., Totterdell, I., Vichi, M. and

- 1189 Völker, C.: Drivers and uncertainties of future global marine primary production in
1190 marine ecosystem models, *Biogeosciences*, 12(23), 6955–6984, 2015.
- 1191 Le Quéré, C., Moriarty, R., Andrew, R. M., Peters, G. P., Ciais, P., Friedlingstein, P.,
1192 Jones, S. D., Sitch, S., Tans, P., Arneeth, A., Boden, T. A., Bopp, L., Bozec, Y.,
1193 Canadell, J. G., Chini, L. P., Chevallier, F., Cosca, C. E., Harris, I., Hoppema, M.,
1194 Houghton, R. A., House, J. I., Jain, A. K., Johannessen, T., Kato, E., Keeling, R. F.,
1195 Kitidis, V., Klein Goldewijk, K., Koven, C., Landa, C. S., Landschützer, P., Lenton,
1196 A., Lima, I. D., Marland, G., Mathis, J. T., Metz, N., Nojiri, Y., Olsen, A., Ono, T.,
1197 Peng, S., Peters, W., Pfügel, B., Poulter, B., Raupach, M. R., Regnier, P., Rödenbeck,
1198 C., Saito, S., Salisbury, J. E., Schuster, U., Schwinger, J., Séférian, R., Segsneider,
1199 J., Steinhoff, T., Stocker, B. D., Sutton, A. J., Takahashi, T., Tilbrook, B., van der
1200 Werf, G. R., Viovy, N., Wang, Y. P., Wanninkhof, R., Wiltshire, A. and Zeng, N.:
1201 Global carbon budget 2014, *Earth Syst. Sci. Data*, 7(1), 47–85, doi:10.5194/essd-7-
1202 47-2015, 2015.
- 1203 Lehodey, P., Alheit, J. and Barange, M.: Climate variability, fish, and fisheries,
1204 *Journal of Climate*, 2006.
- 1205 Levitus, S. and Boyer, T.: World ocean atlas 1994, volume 4: Temperature, PB--95-
1206 270112/XAB, National Environmental Satellite, Data, and Information Service,
1207 Washington, DC (United States). 1994.
- 1208 Levitus, S., Antonov, J. I., Baranova, O. K., Boyer, T. P., Coleman, C. L., Garcia,
1209 H. E., Grodsky, A. I., Johnson, D. R., Locarnini, R. A., Mishonov, A. V., Reagan, J.
1210 R., Sazama, C. L., Seidov, D., Smolyar, I., Yarosh, E. S., and Zweng, M. M.: The
1211 World Ocean Database TI, *Data Science Journal*, 12, WDS229–WDS234, 2013.
- 1212 Levitus, S., Conkright, M. E., Reid, J. L., Najjar, R. G. and Mantyla, A.: Distribution
1213 of nitrate, phosphate and silicate in the world oceans, *Progress in Oceanography*,
1214 31(3), 245–273, 1993.
- 1215 Lévy, M., Lengaigne, M., Bopp, L., Vincent, E. M., Madec, G., Ethe, C., Kumar, D.
1216 and Sarma, V. V. S. S.: Contribution of tropical cyclones to the air-sea CO₂ flux: A
1217 global view, *Global Biogeochem. Cycles*, 26(2), doi:10.1029/2011GB004145, 2012.
- 1218 Lindsay, K., Bonan, G. B., Doney, S. C., Hoffman, F. M., Lawrence, D. M., Long, M.
1219 C., Mahowald, N. M., Moore, J. K., Randerson, J. T. and Thornton, P. E.:
1220 Preindustrial Control and 20th Century Carbon Cycle Experiments with the Earth
1221 System Model CESM1(BGC), *J. Climate*, 141006111735008, doi:10.1175/JCLI-D-
1222 12-00565.1, 2014a.
- 1223 Ludwig, W., Probst, J. and Kempe, S.: Predicting the oceanic input of organic carbon
1224 by continental erosion, *Global Biogeochem. Cycles*, 10(1), 23–41, 1996.
- 1225 Madec, G.: NEMO ocean engine, Institut Pierre-Simon Laplace (IPSL), France.
1226 Institut Pierre-Simon Laplace (IPSL). [online] Available from: [http://www.nemo-
1227 ocean.eu/About-NEMO/Reference-manuals](http://www.nemo-ocean.eu/About-NEMO/Reference-manuals), (last access: November 2013) 2008.
- 1228 Maier-Reimer, E.: Geochemical cycles in an ocean general circulation model.
1229 Preindustrial tracer distributions, *Global Biogeochem. Cycles*, 7(3), 645,

- 1230 doi:10.1029/93GB01355, 1993.
- 1231 Maier-Reimer, E. and Hasselmann, K.: Transport and storage of CO₂ in the ocean —
1232 —an inorganic ocean-circulation carbon cycle model, *Clim Dyn*, 2(2), 63–90–90,
1233 doi:10.1007/BF01054491, 1987.
- 1234 Marinov, I., Gnanadesikan, A., Sarmiento, J. L., Toggweiler, J. R., Follows, M. and
1235 Mignone, B. K.: Impact of oceanic circulation on biological carbon storage in the
1236 ocean and atmospheric pCO₂, *Global Biogeochem. Cycles*, 22(3), GB3007,
1237 doi:10.1029/2007GB002958, 2008.
- 1238 Massonnet, F., Fichet, T., Goosse, H., Bitz, C. M., Philippon-Berthier, G., Holland,
1239 M. M. and Barriat, P. Y.: Constraining projections of summer Arctic sea ice, *The*
1240 *Cryosphere*, 6(6), 1383–1394, doi:10.5194/tc-6-1383-2012, 2012.
- 1241 Matei, D., Baehr, J., Jungclauss, J. H., Haak, H., Muller, W. A. and Marotzke, J.:
1242 Multiyear Prediction of Monthly Mean Atlantic Meridional Overturning Circulation at
1243 26.5 N, *Science*, 335(6064), 76–79, doi:10.1126/science.1210299, 2012.
- 1244 Meehl, G. A., Goddard, L., Boer, G., Burgman, R., Branstator, G., Cassou, C., Corti,
1245 S., Danabasoglu, G., Doblas-Reyes, F., Hawkins, E., Karspeck, A., Kimoto, M.,
1246 Kumar, A., Matei, D., Mignot, J., Msadek, R., Pohlmann, H., Rienecker, M., Rosati,
1247 T., Schneider, E., Smith, D., Sutton, R., Teng, H., van Oldenborgh, G. J., Vecchi, G.
1248 and Yeager, S.: Decadal Climate Prediction: An Update from the Trenches, *Bull.*
1249 *Amer. Meteor. Soc.*, doi:doi: 10.1175/BAMS-D-12-00241.1, 2013.
- 1250 Meehl, G. A., Goddard, L., Murphy, J., Stouffer, R. J., Boer, G., Danabasoglu, G.,
1251 Dixon, K., Giorgetta, M. A., Greene, A. M., Hawkins, E., Hegerl, G., Karoly, D.,
1252 Keenlyside, N., Kimoto, M., Kirtman, B., Navarra, A., Pulwarty, R., Smith, D.,
1253 Stammer, D. and Stockdale, T.: Decadal Prediction, *Bull. Amer. Meteor. Soc.*, 90(10),
1254 1467–1485, doi:10.1175/2009BAMS2778.1, 2009.
- 1255 Meehl, G. A., Moss, R., Taylor, K. E., Eyring, V., Stouffer, R. J., Bony, S. and
1256 Stevens, B.: Climate Model Intercomparisons: Preparing for the Next Phase, *Eos*
1257 *Trans. AGU*, 95(9), 77–78, doi:10.1002/2014EO090001, 2014.
- 1258 Mignot, J., Swingedouw, D., Deshayes, J., Marti, O., Talandier, C., Séférian, R.,
1259 Lengaigne, M. and Madec, G.: On the evolution of the oceanic component of the
1260 IPSL climate models from CMIP3 to CMIP5: A mean state comparison, *Ocean*
1261 *Modelling*, 72 IS -(0 SP - EP - PY - T2 -), 167–184, 2013.
- 1262 Mikaloff Fletcher, S. E., Gruber, N., Jacobson, A. R., Gloor, M., Doney, S. C.,
1263 Dutkiewicz, S., Gerber, M., Follows, M., Joos, F., Lindsay, K., Menemenlis, D.,
1264 Mouchet, A., Müller, S. A. and Sarmiento, J. L.: Inverse estimates of the oceanic
1265 sources and sinks of natural CO₂ and the implied oceanic carbon transport, *Global*
1266 *Biogeochem. Cycles*, 21(1), GB1010, doi:10.1029/2006GB002751, 2007.
- 1267 Moore, J., Doney, S. and Lindsay, K.: Upper ocean ecosystem dynamics and iron
1268 cycling in a global three-dimensional model, *Global Biogeochem. Cycles*, 18(4), –,
1269 doi:10.1029/2004GB002220, 2004.
- 1270 Moore, J., Doney, S., Kleypas, J., Glover, D. and Fung, I.: An intermediate

- 1271 complexity marine ecosystem model for the global domain, *Deep Sea Research Part*
1272 *II: Topical Studies in Oceanography*, 49, 403–462, 2002.
- 1273 Orr, J. C.: *Global Ocean Storage of Anthropogenic Carbon*, Gif-sur-Yvette, France.
1274 2002.
- 1275 Phillips, T. J., Potter, G. L., Williamson, D. L., Cederwall, R. T., Boyle, J. S., Fiorino,
1276 M., Hnilo, J. J., Olson, J. G., Xie, S. and Yio, J. J.: Evaluating Parameterizations in
1277 General Circulation Models: Climate Simulation Meets Weather Prediction, *Bull.*
1278 *Amer. Meteor. Soc.*, 85(12), 1903–1915, doi:10.1175/BAMS-85-12-1903, 2004.
- 1279 Resplandy, L., Bopp, L., Orr, J. C. and Dunne, J. P.: Role of mode and intermediate
1280 waters in future ocean acidification: Analysis of CMIP5 models, *Geophys. Res. Lett.*,
1281 40(12), 3091–3095, 2013.
- 1282 Resplandy, L., Séférian, R. and Bopp, L.: Natural variability of CO₂ and O₂ fluxes:
1283 What can we learn from centuries-long climate models simulations? *Journal of*
1284 *Geophysical Research-Oceans*, 120(1), 384–404, doi:10.1002/2014JC010463, 2015.
- 1285 Rodgers, K. B., Lin, J. and Frölicher, T. L.: Emergence of multiple ocean ecosystem
1286 drivers in a large ensemble suite with an earth system model, *Biogeosciences*
1287 *Discuss.*, 11(12), 18189–18227, doi:10.5194/bgd-11-18189-2014, 2014.
- 1288 Romanou, A., Gregg, W. W., Romanski, J. and Kelley, M.: Natural air–sea flux of
1289 CO₂ in simulations of the NASA-GISS climate model: Sensitivity to the physical
1290 ocean model formulation, *Ocean Modelling*, 66 IS -, 26–44,
1291 doi:10.1016/j.ocemod.2013.01.008, 2013.
- 1292 Romanou, A., J. Romanski, and W.W. Gregg, 2014: Natural ocean carbon cycle
1293 sensitivity to parameterizations of the recycling in a climate model. *Biogeosciences*,
1294 11, 1137-1154, doi:10.5194/bg-11-1137-2014.
1295
- 1296 Romanou, A., W.W. Gregg, J. Romanski, M. Kelley, R. Bleck, R. Healy, L.
1297 Nazarenko, G. Russell, G.A. Schmidt, S. Sun, and N. Tausnev, 2013: Natural air-sea
1298 flux of CO₂ in simulations of the NASA-GISS climate model: Sensitivity to the
1299 physical ocean model formulation. *Ocean Model.*, 66, 26-44,
1300 doi:10.1016/j.ocemod.2013.01.008.
- 1301 Rose, K. A., Roth, B. M. and Smith, E. P.: Skill assessment of spatial maps for
1302 oceanographic modeling, *Journal of Marine Systems*, 76(1-2), 34–48,
1303 doi:10.1016/j.jmarsys.2008.05.013, 2009.
- 1304 Roy, T., Bopp, L., Gehlen, M., Schneider, B., Cadule, P., Frölicher, T. L.,
1305 Segschneider, J., Tjiputra, J., Heinze, C. and Joos, F.: Regional Impacts of Climate
1306 Change and Atmospheric CO₂ on Future Ocean Carbon Uptake: A Multimodel Linear
1307 Feedback Analysis, *J. Climate*, 24(9), 2300–2318, doi:10.1175/2010JCLI3787.1,
1308 2011.
- 1309 Sarmiento, J. L. and Gruber, N.: *Ocean Biogeochemical Dynamics*, Princeton
1310 University Press, Princeton, New Jersey, USA, 526 pp., 2006.
- 1311 Schwinger, J., Tjiputra, J. F., Heinze, C., Bopp, L., Christian, J. R., Gehlen, M.,

- 1312 Ilyina, T., Jones, C. D., Salas-Mélia, D., Segsneider, J., Séférian, R. and Totterdell,
 1313 I.: Nonlinearity of Ocean Carbon Cycle Feedbacks in CMIP5 Earth System Models, *J.*
 1314 *Climate*, 27(11), 3869–3888, doi:10.1175/JCLI-D-13-00452.1, 2014.
- 1315 Servonnat, J., Mignot, J., Guilyardi, E., Swingedouw, D., Séférian, R. and Labetoulle,
 1316 S.: Reconstructing the subsurface ocean decadal variability using surface nudging in a
 1317 perfect model framework, *Clim Dyn*, 44(1-2), 1–24–24, doi:10.1007/s00382-014-
 1318 2184-7, 2014.
- 1319 Séférian, R., Bopp, L., Gehlen, M., Orr, J., Ethé, C., Cadule, P., Aumont, O., Salas y
 1320 Mélia, D., Voldoire, A. and Madec, G.: Skill assessment of three earth system models
 1321 with common marine biogeochemistry, *Climate Dynamics*, 40(9-10), 2549–2573,
 1322 doi:10.1007/s00382-012-1362-8, 2013.
- 1323 Séférian, R., Iudicone, D., Bopp, L., Roy, T. and Madec, G.: Water Mass Analysis of
 1324 Effect of Climate Change on Air–Sea CO₂ Fluxes: The Southern Ocean, *J. Climate*,
 1325 25(11), 3894–3908, doi:10.1175/JCLI-D-11-00291.1, 2012.
- 1326 Séférian, R., Ribes, A. and Bopp, L.: Detecting the anthropogenic influences on recent
 1327 changes in ocean carbon uptake, *Geophys. Res. Lett.*, 2014GL061223,
 1328 doi:10.1002/2014GL061223, 2014.
- 1329 Séférian, R., Delire, C., Decharme, B., Voldoire, A., Salas y Mélia, D., Chevallier,
 1330 M., Saint-Martin, D., Aumont, O., Calvet, J.-C., Carrer, D., Douville, H.,
 1331 Franchistéguy, L., Joetzjer, E. and Sénési, S.: Development and evaluation of CNRM
 1332 Earth-System model – CNRM-ESM1, *Geosci. Model Dev. Discuss.*, 8(7), 5671–5739,
 1333 2015.
- 1334 Smith, D. M., Cusack, S., Colman, A. W., Folland, C. K., Harris, G. R. and Murphy,
 1335 J. M.: Improved Surface Temperature Prediction for the Coming Decade from a
 1336 Global Climate Model, *Science*, 317(5839), 796–799, doi:10.1126/science.1139540,
 1337 2007.
- 1338 Smith, M. J., Palmer, P. I., Purves, D. W., Vanderwel, M. C., Lyutsarev, V.,
 1339 Calderhead, B., Joppa, L. N., Bishop, C. M. and Emmott, S.: Changing how Earth
 1340 System Modelling is done to provide more useful information for decision making,
 1341 science and society, *Bull. Amer. Meteor. Soc.*, 140224132934008,
 1342 doi:10.1175/BAMS-D-13-00080.1, 2014.
- 1343 Steinacher, M., Joos, F., Frölicher, T. L., Bopp, L., Cadule, P., Cocco, V., Doney, S.
 1344 C., Gehlen, M., Lindsay, K., Moore, J. K., Schneider, B. and Segsneider, J.:
 1345 Projected 21st century decrease in marine productivity: a multi-model analysis,
 1346 *Biogeosciences*, 7(3), 979–1005, doi:10.5194/bg-7-979-2010, 2010.
- 1347 Stouffer, R. J., Weaver, A. J. and Eby, M.: A method for obtaining pre-twentieth
 1348 century initial conditions for use in climate change studies, *Clim Dyn*, 23(3-4), 327–
 1349 339, doi:10.1007/s00382-004-0446-5, 2004.
- 1350 Stow, C. A., Jolliff, J., McGillicuddy, D. J. J., Doney, S. C., Allen, J. I., Friedrichs, M.
 1351 A. M., Rose, K. A. and Wallhead, P.: Skill assessment for coupled biological/physical
 1352 models of marine systems, *Journal of Marine Systems*, 76, 4–15,

- 1353 doi:10.1016/j.jmarsys.2008.03.011, 2009.
- 1354 Swingedouw, D., Mignot, J., Labetoulle, S., Guilyardi, E. and Madec, G.:
1355 Initialisation and predictability of the AMOC over the last 50 years in a climate
1356 model, *Clim Dyn*, 40(9-10), 2381–2399, doi:10.1007/s00382-012-1516-8, 2013.
- 1357 Tagliabue, A. and Völker, C.: Towards accounting for dissolved iron speciation in
1358 global ocean models, *Biogeosciences*, 8(10), 3025–3039, 2011.
- 1359 Tagliabue, A., Bopp, L. and Gehlen, M.: The response of marine carbon and nutrient
1360 cycles to ocean acidification: Large uncertainties related to phytoplankton
1361 physiological assumptions, *Global Biogeochem. Cycles*, 25(3), GB3017–n/a,
1362 doi:10.1029/2010GB003929, 2011.
- 1363 Takahashi, T., Broecker, W. and Langer, S.: Redfield Ratio Based on Chemical-Data
1364 From Isopycnal Surfaces, *Journal of Geophysical Research-Oceans*, 90, 6907–6924,
1365 1985.
- 1366 Tanhua, T., Koertzing, A., Friis, K., Waugh, D. W. and Wallace, D. W. R.: An
1367 estimate of anthropogenic CO₂ inventory from decadal changes in oceanic carbon
1368 content, *P Natl Acad Sci Usa*, 104(9), 3037–3042, doi:10.1073/pnas.0606574104,
1369 2007.
- 1370 Tegen, I. and Fung, I.: Contribution to the Atmospheric Mineral Aerosol Load From
1371 Land-Surface Modification, *J Geophys Res-Atmos*, 100, 18707–18726, 1995.
- 1372 Tjiputra, J. F., Olsen, A., Bopp, L., Lenton, A., Pfeil, B., Roy, T., Segschneider, J.,
1373 Totterdell, I. and Heinze, C.: Long-term surface pCO₂ trends from observations and
1374 models, *Tellus B; Vol 66 (2014)*, 66(2-3), 151–168, doi:10.1007/s00382-007-0342-x,
1375 2014.
- 1376 Tjiputra, J. F., Roelandt, C., Bentsen, M., Lawrence, D. M., Lorentzen, T., Schwinger,
1377 J., Seland, Ø. and Heinze, C.: Evaluation of the carbon cycle components in the
1378 Norwegian Earth System Model (NorESM), *Geosci. Model Dev*, 6(2), 301–325,
1379 doi:10.5194/gmd-6-301-2013, 2013.
- 1380 Vancoppenolle, M., Bopp, L., Madec, G., Dunne, J. P., Ilyina, T., Halloran, P. R. and
1381 Steiner, N.: Future Arctic Ocean primary productivity from CMIP5 simulations:
1382 Uncertain outcome, but consistent mechanisms, *Global Biogeochem. Cycles*, 27(3),
1383 605–619, 2013.
- 1384 Vichi, M., Manzini, E., Fogli, P. G., Alessandri, A., Patara, L., Scoccimarro, E.,
1385 Masina, S. and Navarra, A.: Global and regional ocean carbon uptake and climate
1386 change: sensitivity to a substantial mitigation scenario, *Climate Dynamics*, 37(9-10),
1387 1929–1947, doi:10.1007/s00382-011-1079-0, 2011.
- 1388 Volodin, E. M., Dianskii, N. A. and Gusev, A. V.: Simulating present-day climate
1389 with the INMCM4.0 coupled model of the atmospheric and oceanic general
1390 circulations, *Izv. Atmos. Ocean. Phys.*, 46(4), 414–431,
1391 doi:10.1134/S000143381004002X, 2010.
- 1392 Walin, G., Hieronymus, J. and Nycander, J.: Source-related variables for the

- 1393 description of the oceanic carbon system, *Geochem. Geophys. Geosyst.*, 15(9), 3675–
1394 3687, doi:10.1002/2014GC005383, 2014.
- 1395 Wanninkhof, R.: A relationship between wind speed and gas exchange over the ocean,
1396 *J. Geophys. Res.*, 97(C5), 7373–7382, 1992.
- 1397 Wassmann, P., Duarte, C. M., AGUSTÍ, S. and SEJR, M. K.: Footprints of climate
1398 change in the Arctic marine ecosystem, *Global Change Biol*, 17(2), 1235–1249,
1399 doi:10.1111/j.1365-2486.2010.02311.x, 2010.
- 1400 Watanabe, S., Hajima, T., Sudo, K., Nagashima, T., Takemura, T., Okajima, H.,
1401 Nozawa, T., Kawase, H., Abe, M., Yokohata, T., Ise, T., Sato, H., Kato, E., Takata,
1402 K., Emori, S. and Kawamiya, M.: MIROC-ESM 2010: model description and basic
1403 results of CMIP5-20c3m experiments, *Geosci. Model Dev*, 4(4), 845–872,
1404 doi:10.5194/gmdd-4-1063-2011, 2011.
- 1405 Wenzel, S., Cox, P. M., Eyring, V. and Friedlingstein, P.: Emergent constraints on
1406 climate-carbon cycle feedbacks in the CMIP5 Earth system models, *J. Geophys. Res.*
1407 *Biogeosci.*, 2013JG002591, doi:10.1002/2013JG002591, 2014.
- 1408 Wu, T., Li, W., Ji, J., Xin, X., Li, L., Wang, Z., Zhang, Y., Li, J., Zhang, F., Wei, M.,
1409 Shi, X., Wu, F., Zhang, L., Chu, M., Jie, W., Liu, Y., Wang, F., Liu, X., Li, Q., Dong,
1410 M., Liang, X., Gao, Y. and Zhang, J.: Global carbon budgets simulated by the Beijing
1411 Climate Center Climate System Model for the last century, *J Geophys Res-Atmos*,
1412 118(10), 4326–4347, doi:10.1002/jgrd.50320, 2013.
- 1413 Wunsch, C. and Heimbach, P.: Practical global oceanic state estimation, *Physica D:*
1414 *Nonlinear Phenomena*, 230(1-2), 197–208, doi:10.1016/j.physd.2006.09.040, 2007.
- 1415 Wunsch, C. and Heimbach, P.: How long to oceanic tracer and proxy equilibrium?
1416 *Quaternary Science Reviews*, 27(7-8), 637–651, doi:10.1016/j.quascirev.2008.01.006,
1417 2008.
- 1418 Yool, A., Oschlies, A., Nurser, A. J. G. and Gruber, N.: A model-based assessment of
1419 the TrOCA approach for estimating anthropogenic carbon in the ocean,
1420 *Biogeosciences*, 7(2), 723–751, 2010.
- 1421 Yool, A., Popova, E. E. and Anderson, T. R.: MEDUSA-2.0: an intermediate
1422 complexity biogeochemical model of the marine carbon cycle for climate change and
1423 ocean acidification studies, *Geosci. Model Dev*, 6(5), 1767–1811, doi:10.5194/gmd-6-
1424 1767-2013-supplement, 2013.
- 1425 Zeebe, R. E. and Wolf-Gladrow, D. A.: *CO₂ in seawater: equilibrium, kinetics,*
1426 *isotopes*, Elsevier Science Ltd. 2001.
- 1427
- 1428
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Models	spin-up procedure	initial conditions	offline time	online time	total spin-up duration	References
BCC-CSM1-1	sequential	WOA2001, GLODAP	200	100	300	(Wu et al., 2013)
BCC-CSM1-1-m	sequential	WOA2001, GLODAP	200	100	300	(Wu et al., 2013)
CanESM2	sequential (forced w/ obs.)	OCMIP profiles, CanESM1	6000	600	6600	(Arora et al., 2011)
CESM1-BGC	direct	CCSM4	0	1000	1000	(Lindsay et al., 2014)
CMCC-CESM	sequential (w/ acc.)	WOA2001, GLODAP	100	100	200	(Vichi et al., 2011)
CNRM-CM5	sequential	WOA1994, GLODAP, IPSL	3000	100	3100	(Séférian et al., 2013)
CNRM-CM5-2	sequential	WOA1994, GLODAP, CNRM	3000	100	3100	(Schwinger et al., 2014)
CNRM-ESM1	sequential	CNRM-CM5	0	1300	1300	(Séférian et al., 2015)
GFDL-ESM2G	direct	WOA2005,	0	1000	1000	(Dunne et al.,

		GLODAP				2013)
GFDL-ESM2M	direct	WOA2005, GLODAP	0	1000	1000	(Dunne et al., 2013)
GISS-E2-H-CC	direct	WOA2005, GLODAP DIC*	0	3300	3300	(Romanou et al., 2013)
GISS-E2-R-CC	direct	WOA2005, GLODAP DIC*	0	3300	3300	(Romanou et al., 2013)
HadGEM2-CC	sequential	HadCM3LC , WOA2011	400	100	500	(Collins et al., 2011; Wassmann et al., 2010)
HadGEM2-ES	sequential	HadCM3LC , WOA2010	400	100	500	(Collins et al., 2011)
INMCM4	sequential	Uniform DIC	3000	200	3200	(Volodin et al., 2010)
IPSL-CM5A-LR	sequential	WOA1994, GLODAP, IPSL	3000	600	3600	(Séférian et al., 2013)
IPSL-CM5A-MR	sequential	WOA1994, GLODAP, IPSL	3000	300	3300	(Dufresne et al., 2013)
IPSL-CM5B-LR	sequential	IPSL- CM5A-LR	0	300	300	(Dufresne et al., 2013)
MIROC-ESM	sequential	GLODAP/c onstant values	1245	480	1725	(Watanabe et al., 2011)
MIROC-ESM- CHEM	sequential	GLODAP/c onstant values	1245	484	1729	(Watanabe et al., 2011)
MPI-ESM-LR	sequential	HAMOCC/ constant values	10000	1900	11900	(Ilyina et al., 2013)
MPI-ESM-MR	sequential	HAMOCC/ constant values	10000	1500	11500	(Ilyina et al., 2013)
MRI-ESM1	sequential (forced w/ obs.)	GLODAP	550	395	945	(Adachi et al., 2013)
NorESM	direct	WOA2010, GLODAP	0	900	900	(Tjiputra et al., 2013)

1445

1446 **Table 1:** Summary of spin-up strategy, sources of initial conditions, offline/online

1447 durations and references used to equilibrate ocean biogeochemistry in CMIP5 ESMs.

1448 The so-called direct and sequential strategies inform whether the spin-up of the ocean

1449 biogeochemical model is run directly in online/coupled mode or first in offline (ocean
 1450 biogeochemistry only) and then in online/coupled mode. DIC* refers to the
 1451 observation-derived estimates of preindustrial dissolved inorganic carbon
 1452 concentration using the ΔC^* method. w/ acc. and forced w/ obs. indicates the strategy
 1453 using ‘acceleration’ and observed atmospheric forcings during the spin-up,
 1454 respectively.

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 1456

	O ₂			NO ₃		
Depth	surface	150 m	2000 m	surface	150 m	2000 m
RMSE	7.19	8.75	5.50	2.07	2.90	2.08
R ²	0.98	0.98	0.99	0.96	0.92	0.94

1457

1458 **Table 2:** Differences between the oxygen (O₂, $\mu\text{mol L}^{-1}$) and nitrate (NO₃, $\mu\text{mol L}^{-1}$)
 1459 datasets used for initializing IPSL-CM5A-LR (WOA1994) and the datasets used for
 1460 assessing its performances (WOA2013).

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	O ₂			NO ₃			Alk-DIC		
metrics	mean	RMSE	RMSE _{max}	mean	RMSE	RMSE _{max}	mean	RMSE	RMSE _{max}
Surf	-0.2	2.6	55.8	-0.1	-0.1	34.2	1.6	-0.1	-0.1
150 m	3.4	39.0	31.5	-15.9	33.4	55.2	6.1	27.9	24.7

2000 m									
	-30.4	144.3	-40.1	2	51.8	-34.8	-69.6	281.8	47.5

1463 **Table 3:** Drift in % ky⁻¹ for oxygen (O₂), nitrate (NO₃) and total alkalinity minus DIC
1464 (Alk-DIC) at surface, 150 and 2000 meters as simulated by the IPSL-CM5A-LR
1465 model. The drift has been computed over the last 250 years of the spin-up simulation
1466 using a linear regression fit of the globally averaged concentrations, root-mean
1467 squared error (RMSE) and latitudinal maximum root-mean squared error (RMSE_{max})
1468 with respect to the values at year 250.

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1471

1472 **Figure 1:** Spin-up protocols of CMIP5 Earth system models. Color shading represents
1473 strategies of the various modeling groups. *Online* and *Offline* steps refer to runs
1474 performed with coupled climate model and with stand-alone ocean biogeochemistry
1475 model, respectively. Sources of initial conditions for biogeochemical component of
1476 CMIP5 Earth system models are indicated as hatching below the barplot.

1477

1478 **Figure 2:** Time series of two climate indices over the 500-year spin-up simulation of
1479 IPSL-CM5A-LR. They represent the global averaged sea surface temperature (a) and
1480 the global mean sea-air carbon flux (b). For sea-air carbon flux, negative value
1481 indicates uptake of carbon. Steady state equilibrium of physical components as
1482 described in Mignot et al., (2013) is reached at ~250 years and is indicated with a
1483 vertical dashed line. Drifts in sea surface temperature and global carbon flux are
1484 indicated with dashed blue lines. They are computed using a linear regression fit over
1485 years 250 to 500. Hatching on panel (b) represents the range of inverse modeling
1486 estimates for preindustrial global carbon flux as described in Mikaloff Fletcher et al.,
1487 (2007), i.e., 0.03±0.08 Pg C y⁻¹ plus 0.45 Pg C y⁻¹ corresponding to the riverine-
1488 induced natural CO₂ outgassing outside of near-shore regions consistently with Le
1489 Quéré et al. (2015).

1490

1491 **Figure 3:** Time series of globally averaged concentration ($[X]$ in solid lines) and
1492 globally averaged root-mean squared error (RMSE in dashed lines) for dissolved
1493 oxygen (O_2), nitrate (NO_3) and difference between alkalinity and dissolved inorganic
1494 carbon (Alk-DIC) as simulated by IPSL-CM5A-LR. $[X]$ and RMSE are given at
1495 surface (a,b and c), 150 m (d, e and f), and 2000 m (g, h and i) for these three
1496 biogeochemical fields. Their values are indicated on the left-side and right-side y-axis,
1497 respectively. Hatching represents the $\pm\sigma$ observational uncertainty due to optimal
1498 interpolation of in situ concentrations around the observed $[X]$.

1499

1500 **Figure 4:** Snap-shots of spatial biases, ϵ , in surface concentrations ($\mu\text{mol L}^{-1}$) in
1501 biogeochemical fields during the 500-year spin-up simulation of IPSL-CM5A-LR. ϵ
1502 in dissolved oxygen (O_2), nitrate (NO_3) and difference between alkalinity and
1503 dissolved inorganic carbon (Alk-DIC) is given for the first year (a, c and e,
1504 respectively) and for the last year of spin-up simulation (b, d and f, respectively).

1505

1506 **Figure 5:** As Figure 4 but for concentrations at 150 m. Note that color shading does
1507 not represent the same amplitude in spatial biases as in Figures 4 and 6.

1508

1509 **Figure 6:** As Figure 4 but for concentrations at 2000 m. Note that color shading does
1510 not represent the same amplitude in spatial biases as in Figures 4 and 5.

1511

1512 **Figure 7:** Temporal-vertical evolution in root-mean squared error (RMSE) for
1513 biogeochemical tracers during the 500-year-long spin-up simulation of IPSL-CM5A-
1514 LR. RMSE is given for (a) dissolved oxygen O_2 , (b) nitrate NO_3 and (c) difference
1515 between alkalinity and dissolved inorganic carbon Alk-DIC.

1516

1517 **Figure 8:** Temporal evolution of drift in root-mean squared error (RMSE) for
1518 dissolved oxygen (O_2 , blue crosses), nitrate (NO_3 , green crosses) and difference
1519 between alkalinity and dissolved inorganic carbon (Alk-DIC, orange crosses) during
1520 the 500-year-long spin-up simulation of IPSL-CM5A-LR. Drift in RMSE is given at
1521 surface (a,b and c), 150 m (d, e and f), and 2000 m (g, h and i) for these three
1522 biogeochemical fields. Drift in RMSE is computed from time segments of 100 years
1523 beginning every 5 years from the beginning until year 400 of the spin-up simulation
1524 for O_2 , NO_3 and Alk-DIC tracers. The best-fit linear regressions between drifts in

1525 RMSE and spin-up duration over year 250 to 500 are indicated in solid magenta lines;
1526 their 90% confidence intervals are given by thin dashed envelopes.

1527

1528 **Figure 9:** Scatterplot of drifts in root-mean squared error (RMSE) in O₂ concentration
1529 versus the duration of the spin-up simulation for the available CMIP5 Earth system
1530 models. Drifts in O₂ RMSE are respectively given for surface (a), 150 m (b) and 2000
1531 m (c) for oxygen concentrations. Drift in O₂ RMSE is computed from several time
1532 segments of 100 years beginning every 5 years from the beginning until the end of the
1533 piControl simulation for the available CMIP5 models. Coloured symbols indicate the
1534 mean drift in O₂ RMSE while vertical lines represent the associated 90% confidence
1535 interval. The best-fit linear regressions between models' mean drifts in RMSE and
1536 spin-up duration are indicated as solid green lines; their 90% confidence intervals are
1537 given by thin dashed envelopes. Fits are assumed robust if correlation coefficients are
1538 significant at 90% (i.e., $r^* > 0.34$). For comparison, drift in O₂ RMSE from our spin-up
1539 simulation with IPSL-CM5A-LR (Figure 8) are represented by magenta crosses.

1540

1541 **Figure 10:** Rankings of CMIP5 Earth system models based on standard and penalized
1542 version of the distance from oxygen observations. The standard distance metric is
1543 calculated as the ensemble-mean root-mean squared error (RMSE) for O₂
1544 concentrations at surface (a), 150 m (b) and 2000 m (c). The penalized distance metric
1545 incorporates drift-induced changes in O₂ RMSE (Δ RMSE) to O₂ RMSE at surface (d),
1546 150 m (e) and 2000 m (f). Ensemble-mean RMSE are calculated using available
1547 ensemble members of Earth system models oxygen concentrations averaged over the
1548 1986-2005 historical period relative to WOA2013 observations. Δ RMSE is
1549 determined using Equation 2 and fits derived from first century of the CMIP5
1550 piControl simulations. Solid red and magenta lines indicate the multi-model mean
1551 standard and penalized distance from O₂ observations, respectively. With the same
1552 colour pattern, dashed lines are indicative of the multi-model median for the standard
1553 and penalized distance from O₂ observations.

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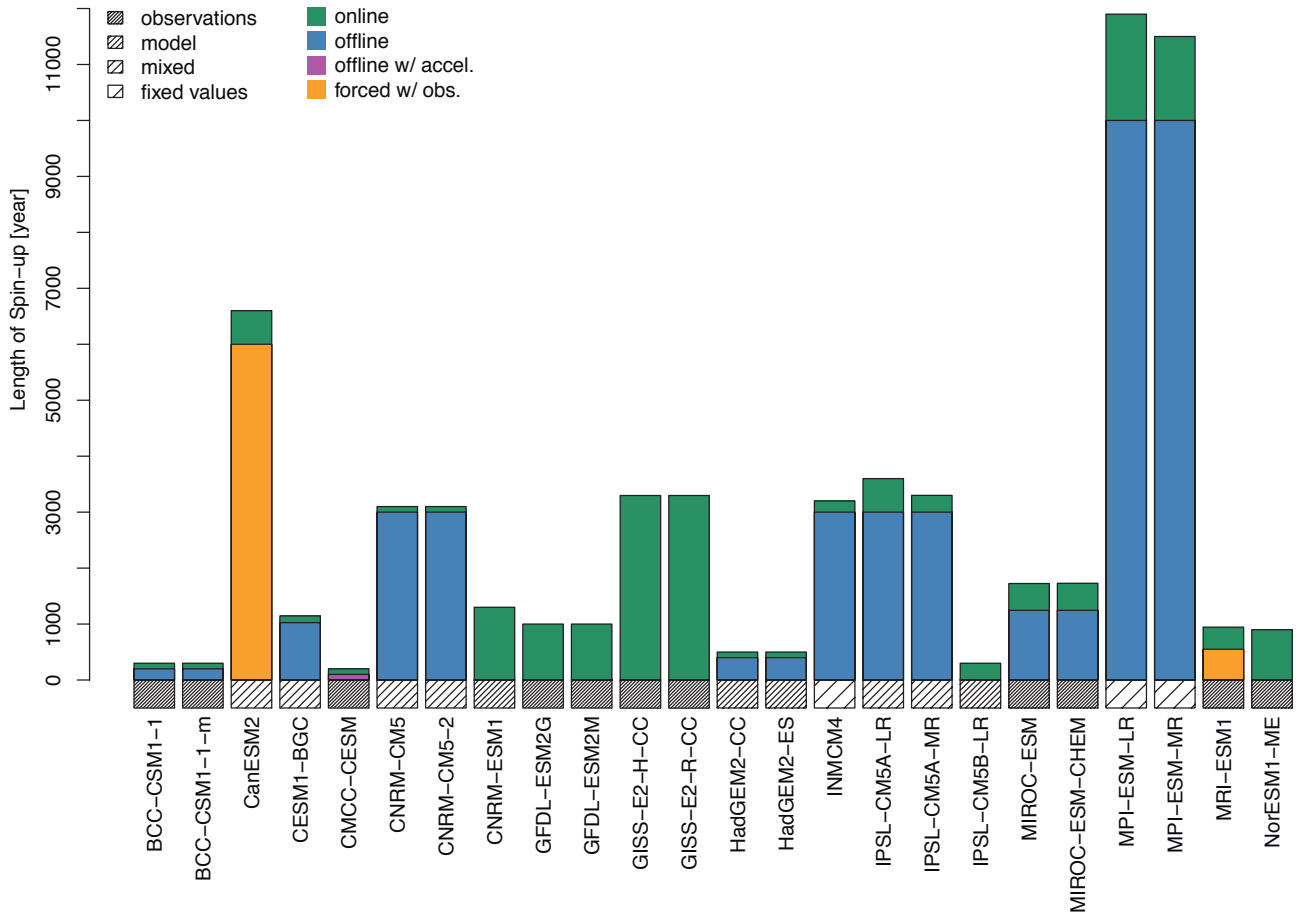


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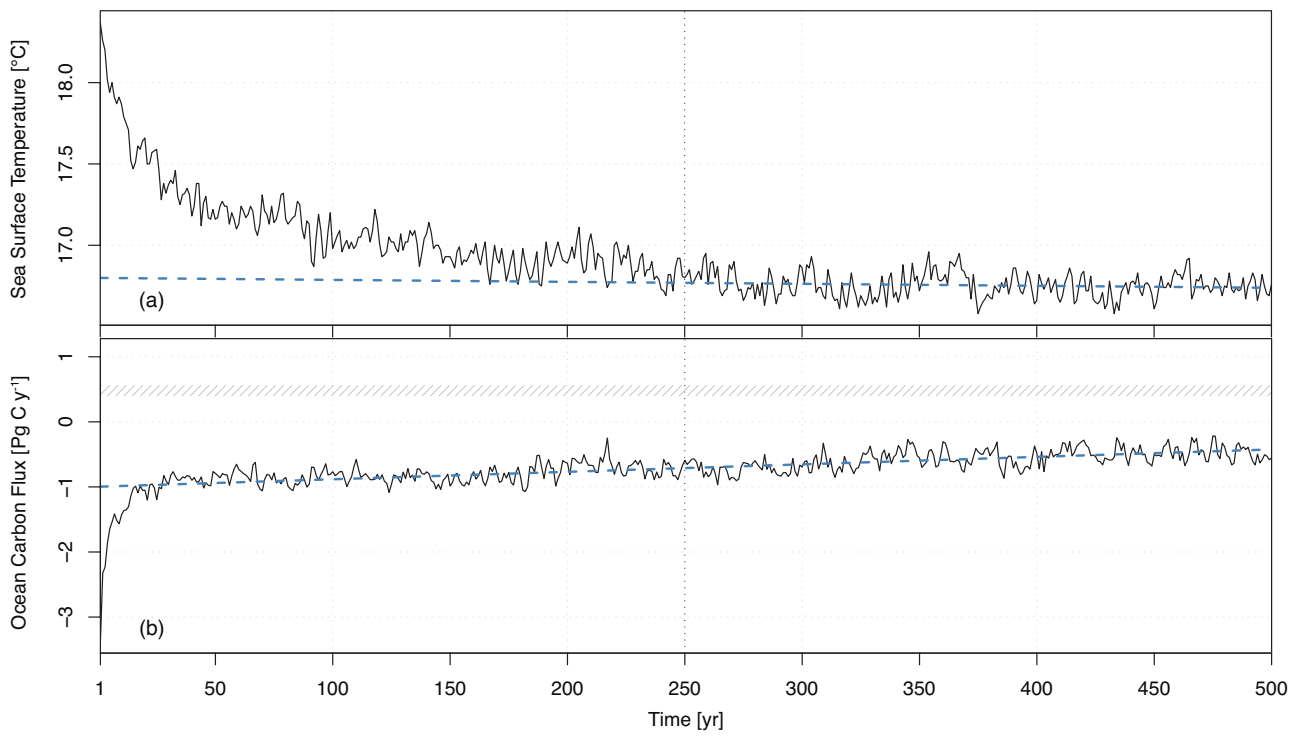


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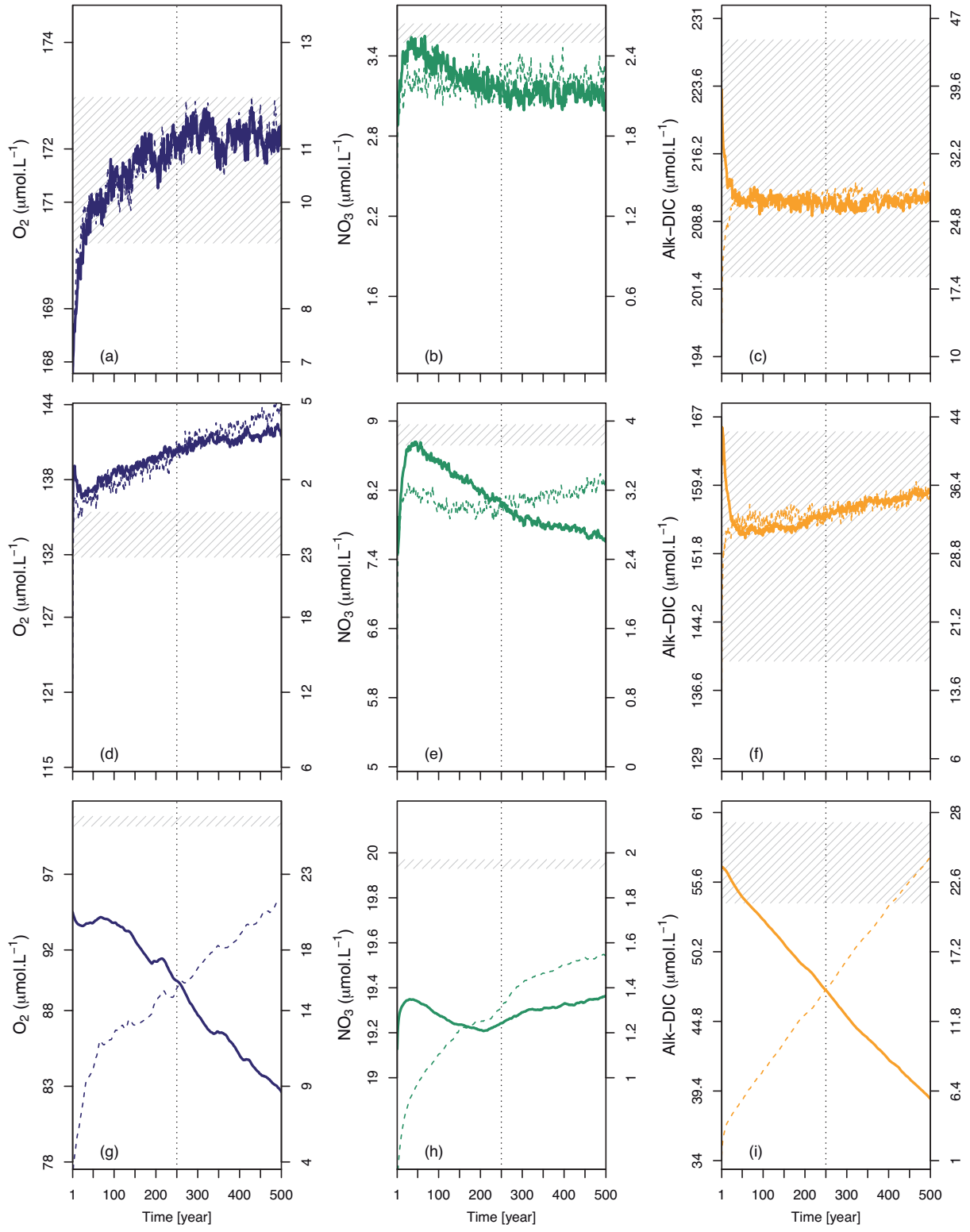


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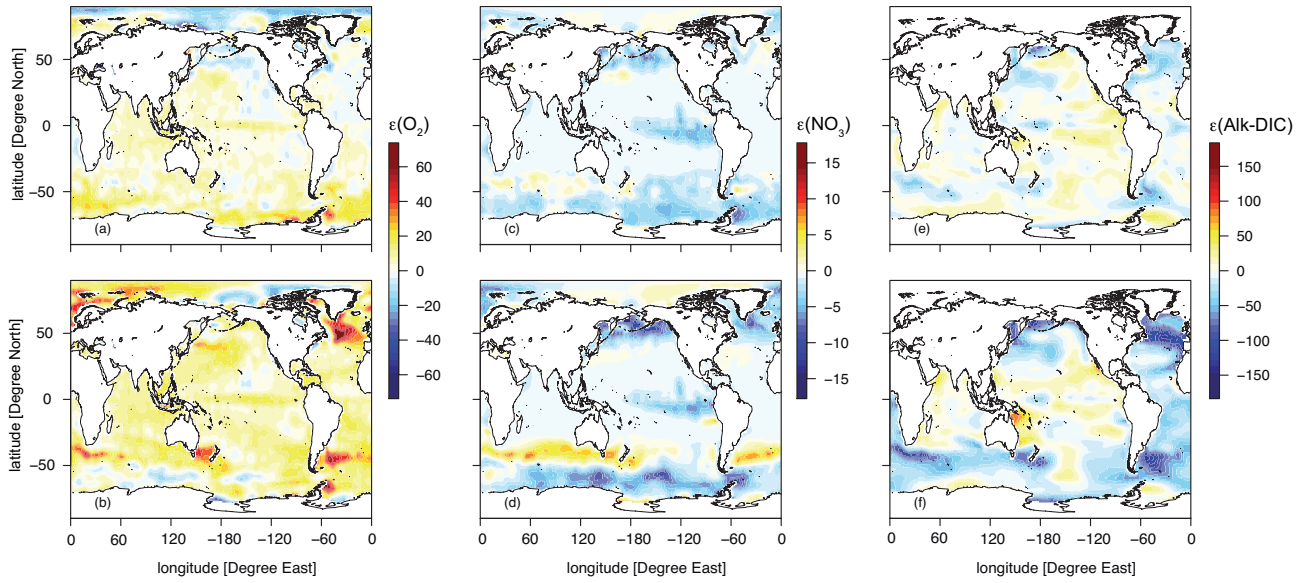


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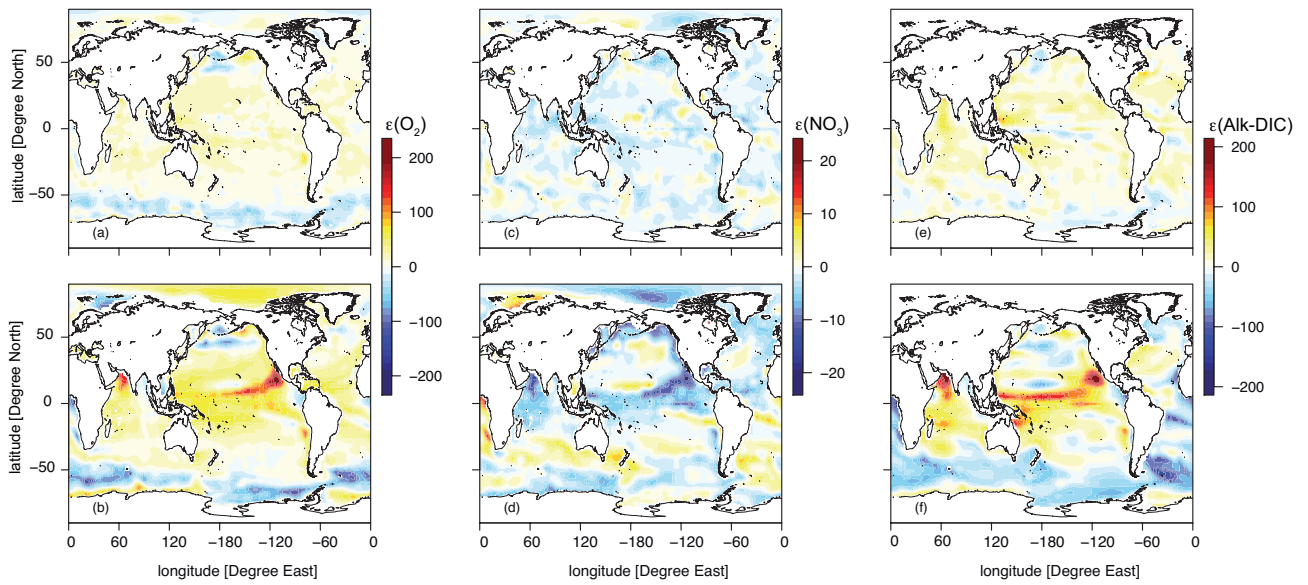


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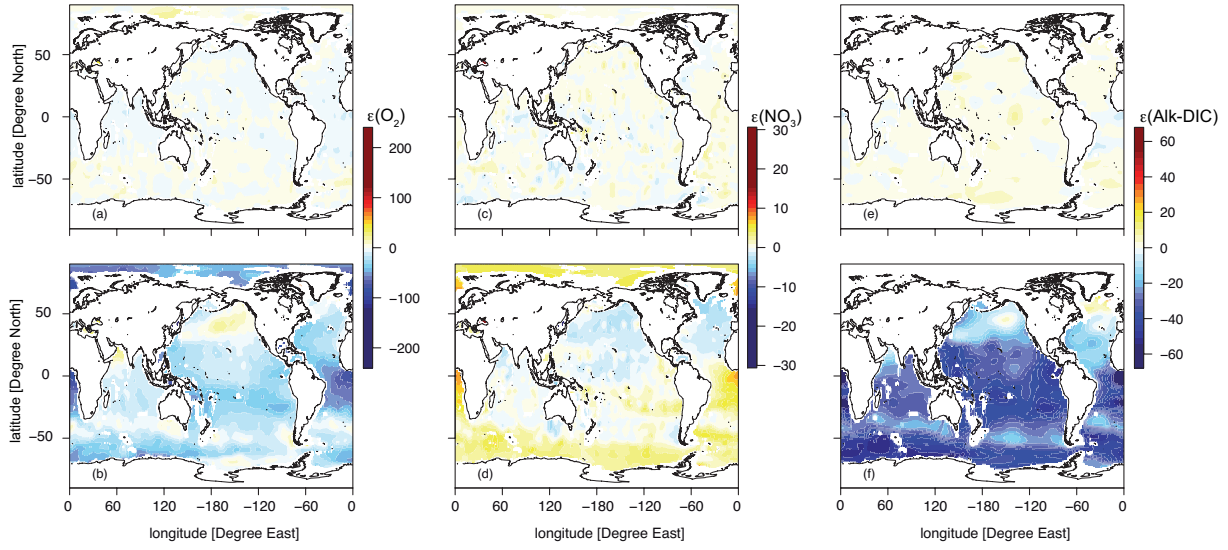


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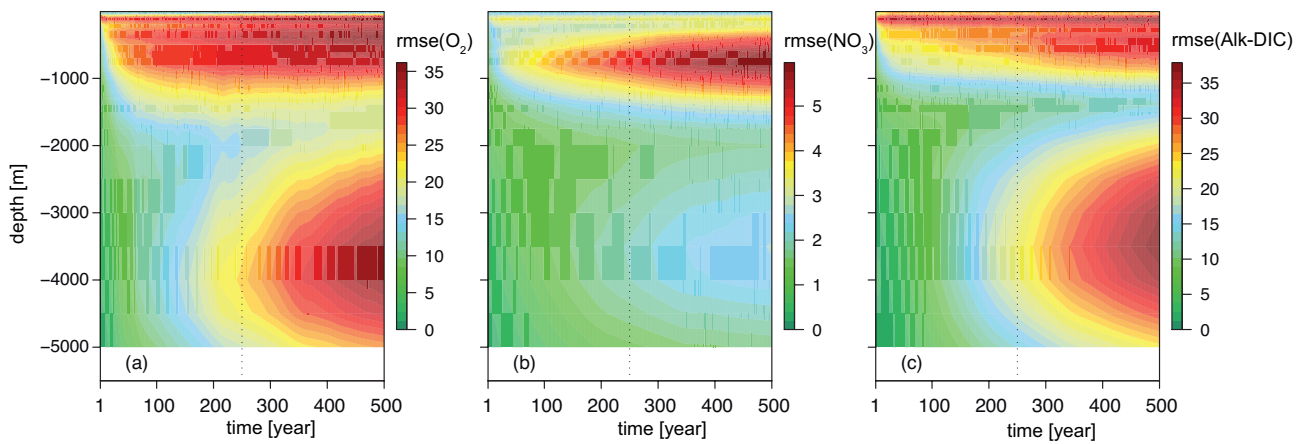


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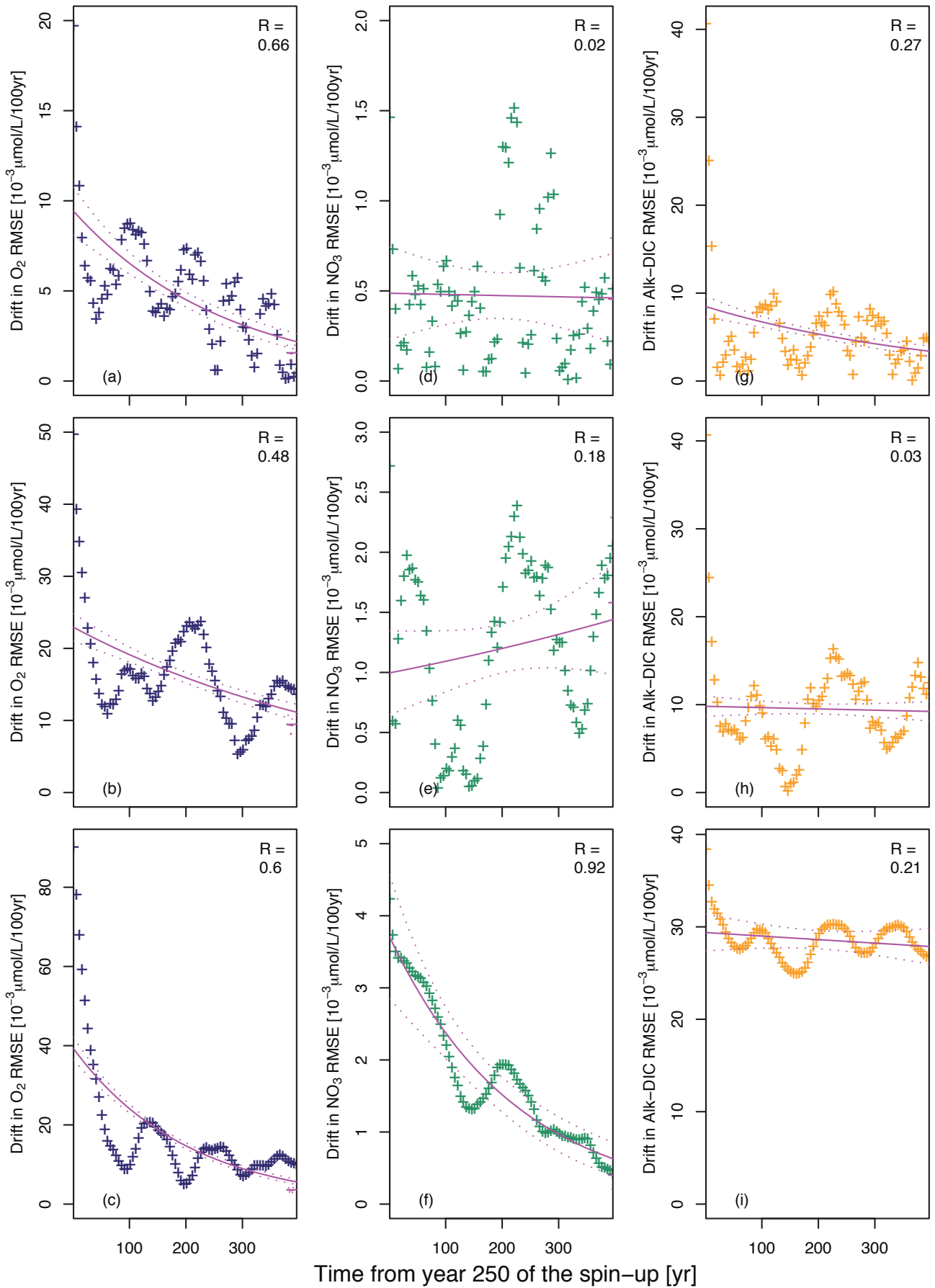


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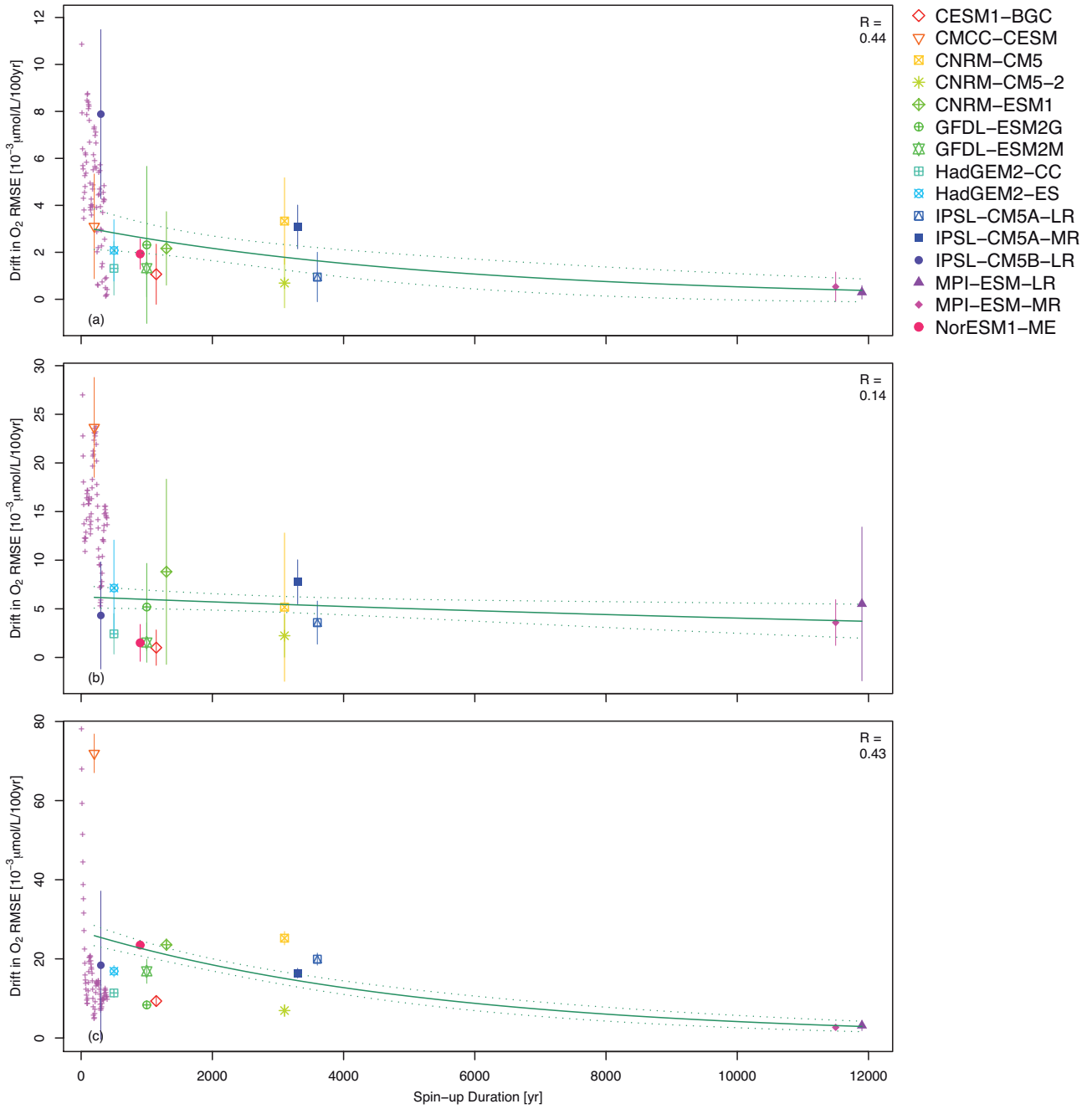


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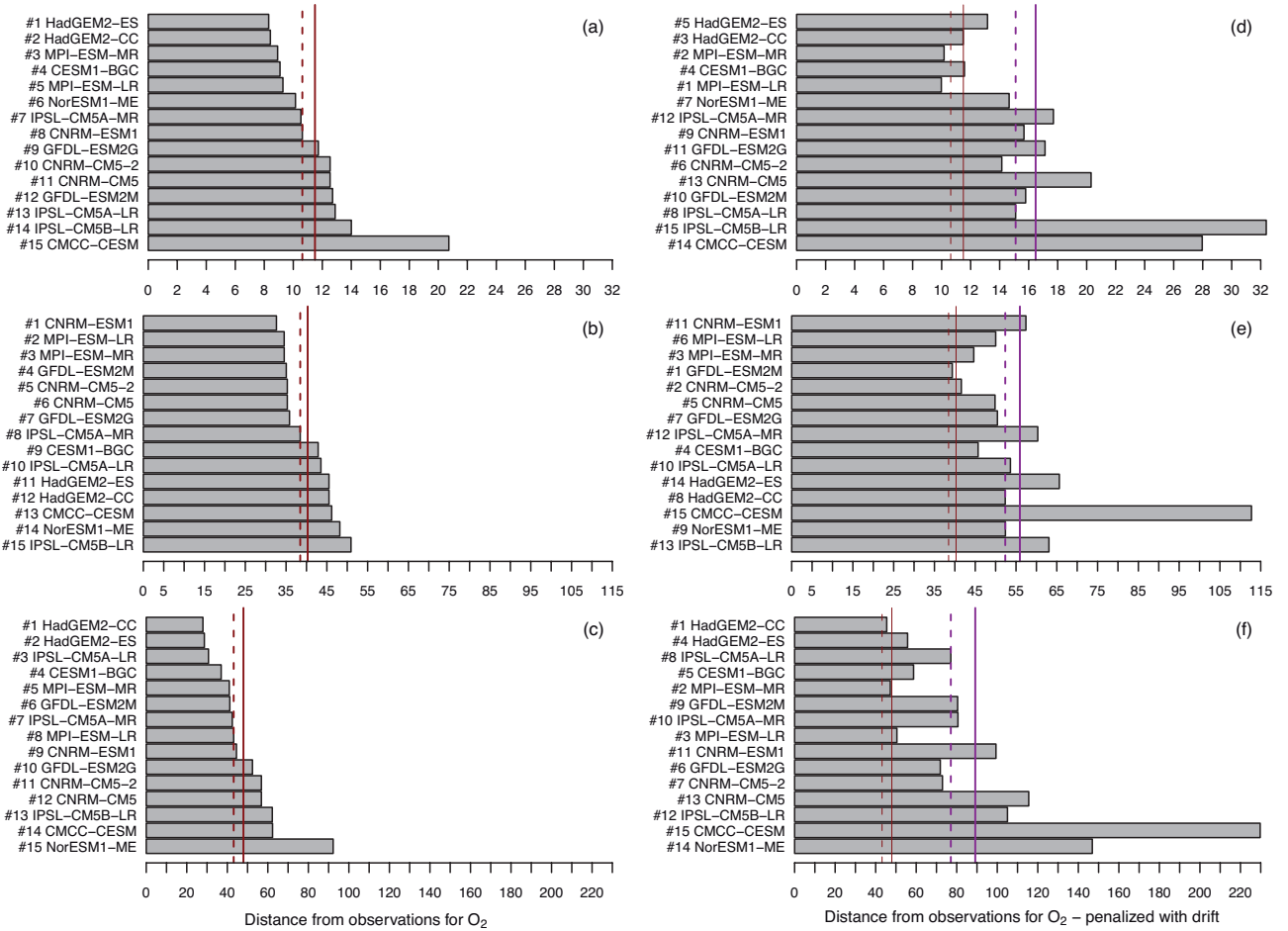


Figure 10: