

1 **Supplementary Information for**

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3 **Global ensemble projections reveal trophic amplification of ocean biomass**  
4 **declines with climate change**

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## 28 **Supplementary Methods**

29 **Ecosystem model selection.** We used six published and peer-reviewed global fisheries and  
30 marine ecosystem models (MEMs) that participated in the first round of the Fisheries and Marine  
31 Ecosystem Model Intercomparison Project (Fish-MIP, Table S1, 1-11). These six MEMs varied  
32 considerably in basic model structure and underlying assumptions, taxonomic scope and key  
33 ecological processes included, and their representation of species, functional groups, size classes  
34 and the proportion of commercial to non-commercial taxa (Tables S1 and S2). Together, this  
35 heterogeneity reflects the diversity of model structures, parameterizations, scopes and purposes,  
36 meaning that our ensemble is more likely to include a greater number of relevant processes in the  
37 ocean than any single model.

38 The MEMs also varied in their representation of fishing, with some models only  
39 representing an unfished ocean (Macroecological) while others were able to incorporate fishing  
40 effort (BOATS, EcoOcean) or fishing mortality (DBEM, DPBM, APECOSM) as forcing  
41 variables, but so far only few models can incorporate feedbacks from the biological to social and  
42 economic systems. For example, BOATS uses an interactive bioeconomic model to determine  
43 spatial and temporal changes in fishing effort (2, 3), and EcoOcean uses a spatially-explicit  
44 fishing dynamics model to simulate spatial patterns of imposed fishing effort (7). Due to these  
45 inherent differences, we could not standardize fishing scenarios across MEMs and therefore  
46 assessed the climate-change effect in an unfished compared to a fished ocean in a subset of  
47 MEMs (see below).

48 In addition to our own empirical model validation (see below), the performance of  
49 individual MEMs has been previously compared with various observed spatial or temporal catch  
50 or biomass data (Fig. S3), and with ecological data from observations or experiments at different

51 stages of model development. Model outputs from BOATS were compared to global historical  
52 time series of catch data from the Sea Around Us Project (SAUP, <http://www.seaaroundus.org/>),  
53 including unreported catches, and trends in the biomass of assessed stocks relative to their  
54 unexploited biomass ( $B/B_0$ ) from the RAM Legacy stock assessment database  
55 (<http://ramlegacy.org/>) (2, 3, Fig. S3A and B). Moreover, parameter sensitivity was extensively  
56 tested with temperature and NPP to optimize the model's ability to reproduce historical patterns  
57 of fish catch from SAUP and RAM (12). Model outputs from EcoOcean were compared to  
58 historic time series of catch data from SAUP (7, Fig. S3C), and to catches for major functional  
59 groups and large marine ecosystems (LMEs) from SAUP in the 1950s (not shown) and 2000s  
60 (shown in Fig. S3D; 7). For DBEM, the mean state projected catch potential among large marine  
61 ecosystems (LMEs) was significantly correlated with observed catch data from SAUP (Fig.  
62 S3E), and this was robust to different underlying calculations for the carrying capacity of each  
63 grid cell (13). Also, the rate of range shifts projected by DBEM correlated significantly with  
64 observed range shifts in the North Sea and Bering Sea (13), and the predicted growth and body  
65 size parameters (that contribute to the calculation of biomass and catch potential) were also well  
66 correlated with observations (13). Moreover, the sensitivity of the model outputs was tested with  
67 different ocean variables, with temperature being the main driver, followed by NPP and then  
68 oxygen, particularly in tropical ecosystems (14). For DPBM, previous work at the scale of  
69 regional seas (5) compared modelled catches (based on a fishing mortality rate of  $0.8 \text{ yr}^{-1}$  that  
70 produced maximum equilibrium catches) to reported landings (excluding discards) from the  
71 Food and Agriculture Organization (FAO) in 11 regional domains based on 78 Exclusive  
72 Economic Zones (EEZ) (Fig. S3F). Additionally, modelled fish and benthic community size  
73 spectra and species averaged size-specific growth rates were consistent with previous empirical

74 observations for EEZs and in the North Sea (5, 15). Model outputs from APECOSM were  
75 compared to biological observations at all stages of model development (e.g. growth,  
76 ingestion/assimilation, reproductive outputs, maintenance, development, size at maturity, aging  
77 mortality; 16). Vertical distributions in APECOSM were assessed using acoustic observations  
78 (using data in 17), and horizontal and size-distributions using fisheries data for large sizes (18)  
79 and MAREDAT for small ones (19). Also, observed and predicted catches were compared for  
80 skipjack tuna fisheries in Indian Ocean across different schools with good correlation (18, Fig.  
81 S3G). The Macroecological model (4) exclusively represents an unexploited ocean and can  
82 therefore not be compared to observed global catch or biomass data; however, comparisons  
83 could be done for some regions, such as the North Sea, where very comprehensive biomass data  
84 exist for non-fished as well as fished species. Here, the median predicted biomass estimate for  
85 fish size classes >100g was 8.3 million tonnes, which compared well with detailed empirical  
86 estimates of 8.6-13.1, 10.0, 3.8-7.5 million tonnes from 3 independent studies (4). Moreover,  
87 almost all parameterization in the Macroecological model was based on empirical analysis and  
88 the underlying functional processes are well supported by many empirical and experimental  
89 studies (4).

90

91 **Climate-change scenarios.** All MEMs were forced with the same standardized set of outputs  
92 from Earth-system models (ESMs) derived from the Coupled Model Inter-comparison Project  
93 Phase 5 (CMIP5 database: <http://cmip-pcmdi.llnl.gov/cmip5/>). We selected NOAA's  
94 Geophysical Fluid Dynamics Laboratory Climate Model (GFDL-ESM2M, 20) and the Institute  
95 Pierre Simon Laplace Climate Model (IPSL-CM5A-LR, 21) because they generated all  
96 necessary physical and biogeochemical outputs needed to run our global marine MEMs,

97 particularly the monthly depth-resolved biogeochemical fields of different size groups of  
98 phytoplankton and zooplankton concentration and productivity (1). While several other ESMs  
99 are available in CMIP5, they did not generate or save all variables required by our MEMs. The  
100 two selected ESMs spanned the range of projections from all CMIP5 models with GFDL-  
101 ESM2M being at the lower end and IPSL-CM5A-LR at the higher end of projected future  
102 changes in SST and NPP (Fig. S1), while other variables, including phytoplankton and  
103 zooplankton biomass (Fig. S2) as well as pH and O<sub>2</sub> concentrations were more similar among all  
104 CMIP5 models (22). Therefore, global annual mean trends of these two ESMs should reflect the  
105 multi-model mean and range of a broad set of CMIP5 models (1, 22).

106 Each ESM was run for each of four Representative Concentration Pathways (RCPs)  
107 representing a standard set of IPCC informed emission scenarios, ensuring that starting  
108 conditions, historical trends, and projections are consistent across models and sectors. RCP2.6  
109 represents a strong mitigation scenario characterized by an emission pathway leading to very low  
110 greenhouse gas (GHG) levels by 2100 (23). RCP4.5 and RCP6.0 represent stabilization emission  
111 scenarios, requiring a stabilization in radiative forcing after 2100, without exceeding the target  
112 value of 4.5 W m<sup>-2</sup> for RCP4.5 and 6.0 W m<sup>-2</sup> for RCP6.0 (24). RCP8.5 represents a business-as-  
113 usual scenario characterized by increasing GHG emissions over time leading to high GHG  
114 emissions in 2100 (25).

115

116 **Standardized model inputs and outputs.** The fundamental goal of Fish-MIP is to compare the  
117 response of a wide range of MEMs to common external forcings, namely standardized climate-  
118 change scenarios. Standardized physical and biogeochemical variables from the 2 ESMs and 4  
119 RCPs were derived on a 1×1-degree global grid for a historical (1970-2005) and a future period

120 (2006-2100) to be used as climate input data in our MEMs (Table S2). Physical variables  
121 included current velocities, water temperature, salinity, dissolved oxygen concentration, pH,  
122 mixed-layer depth and ice coverage. Biogeochemical variables included large and small  
123 phytoplankton and zooplankton concentrations and productivity. All variables were summarized  
124 as monthly, depth-resolved values and then further aggregated or integrated to suit the input  
125 requirements for each MEM, such as providing values for surface, bottom or depth-integrated  
126 layers (Table S2). For more details on input variable selection and requirements of each  
127 ecosystem model see ref. (1).

128 First, we used a no-fishing scenario across all six MEMs to isolate the climate-change effect  
129 on fish or animal biomass. In a subset of three MEMs we also used a simple fishing scenario  
130 based on observed or estimated time-varying data on fishing effort, mortality or exploitation  
131 rates (depending on model requirements) for the historical period (1970-2005) and then kept  
132 constant at 2005 levels for the future period (2005-2100) (1). Thereby, each model relied on its  
133 own mechanism for incorporating fishing data (1). Dynamic projections of future fishing  
134 pressure were not used because appropriate temporally- and spatially-explicit scenarios of future  
135 fishing pressure are not yet available (1), although we do acknowledge that fishing pressure will  
136 change over the projection period. Recently, future fishing scenarios have been developed  
137 conceptually based on the IPCC's Shared Socio-economic Pathways (SSPs) (26). Once  
138 translated into quantitative form, these should be available for future efforts to project changes in  
139 fish and fisheries.

140 All MEMs were required to produce a set of standardized outputs that could be directly  
141 compared and combined into ensemble projections (1). For this study, we selected three outputs:  
142 (i) total consumer biomass density ( $\text{g C m}^{-2}$ ) representing all animals, size classes or trophic

143 groups in each model, (ii) biomass density of all animals, size classes or trophic groups >10 cm  
144 (g C m<sup>-2</sup>), and (iii) biomass density of all animals, size classes or trophic groups >30 cm (g C m<sup>-2</sup>).  
145 Since the taxonomic scope differs among models, we used the terms ‘total animal biomass’,  
146 ‘animals >10 cm’ and ‘animals >30 cm’, respectively, for simplification. All data reported in this  
147 paper are archived and publicly available at <http://doi.org/10.5880/PIK.2018.005>.

148

149 **Simulations.** Several limitations resulted in not all MEMs being able to run the full set of 16  
150 simulations from 2 ESMs, 4 RCPs and 2 fishing scenarios for total animal biomass, biomass of  
151 animals >10 cm and >30 cm. We therefore decided to use comparable subsets of available  
152 MEM-ESM combinations to answer different questions. ESM limitations included the  
153 unavailable monthly, depth- and size-resolved biogeochemical data in GFDL-ESM2M (not run  
154 in APECOSM, DPBM); MEM limitations included substantial simulation run time in some  
155 MEMs (e.g. DBEM did not run all RCPs; DPBM and APECOSM did not perform fishing runs),  
156 the lack of size class differentiation (e.g. DBEM), and the inability to incorporate fishing (e.g.  
157 Macroecological). Therefore, all six MEMs used the IPSL-CM5A-LR no-fishing runs and four  
158 used the GFDL-ESM2M no-fishing runs for RCP2.6 and 8.5 (n = 10), and five and three MEMs,  
159 respectively, for RCP4.5 and 6.0 (n = 8; Table S3). A subset of three MEMs (BOATS,  
160 EcoOcean, DBEM) was used for fishing runs with inputs from both ESMs for RCP2.6 and 8.5 (n  
161 = 6). All six MEMs projected total animal biomass, and five MEMs animal biomass >10 cm and  
162 >30 cm (Table S3). However, since DBEM represents all commercial fish and invertebrate  
163 species covering the full size spectrum, we used DBEM outputs for all three size groups, but  
164 cross-checked results with and without DBEM as a sensitivity analysis.

165

166 **Analyses and model validation.** Because the six MEMs include different species, size classes  
167 and trophic groups, their output for absolute biomass density varied across models. We therefore  
168 calculated time series of relative change in animal biomass defined as percent (%) relative to  
169 1990-99 for each simulation. This reference period was chosen as representing the last decade of  
170 the 20th century, which was later compared to 2090-99 as the last decade of the 21st century.

171 The different time series of relative change were then averaged into a multi-model mean  
172 change, and variability among models was described with the standard deviation (SD) around the  
173 mean (Fig. 1A, Fig. S5) as well as with box and whisker plots (Fig. 1C). To summarize the  
174 results across simulations and size groups, we also calculated the % change in 2090-99 relative  
175 to 1990-99 (Table S3).

176 Given the differences in MEM structure and characterization of fishing (see above) we could  
177 not standardize the fishing scenarios and did not compare the magnitude of the fishing effects  
178 across models. Instead we compared the relative difference in the climate effect (RCP8.5 vs RCP  
179 2.6) in a fished and an unfished ocean. To do so, we calculated  $((RCP8.5 - RCP2.6) / RCP2.6)$   
180 within each MEM in the 2090s and over time which was then compared across MEMs (Fig. 1D,  
181 Fig. S6A).

182 Next, we compared the variability of results due to the different ESMs and MEMs (Fig. 1E).  
183 For ESM variability, we calculated the standard deviation for individual MEM results for  
184 simulations run with the IPSL-CM5A-LR compared to the GFDL-ESM2M (for all MEMs run  
185 with both forcings,  $n = 4$ ). For MEM variability, we calculated the standard deviation of results  
186 across all MEMs for IPSL-CM5A-LR ( $n = 6$ ) and GFDL-ESM2M ( $n = 4$ ) which were then  
187 combined. These calculations were performed separately for the different RCPs and size groups  
188 in an unfished ocean (Fig. 1E).

189 To validate our ensemble model projections with empirical data, we compared our multi-  
190 model mean historical biomass trend of animals >10cm (for better comparison with assessed fish  
191 stocks) with biomass trends of assessed fish stocks in (i) a fished and (ii) an unfished situation  
192 from 1970-2005. For the fished situation, we compared our multi-model mean of all MEM-ESM  
193 combination with fishing ( $n = 6$ ) to the observed biomass of assessed exploited fish stocks  
194 relative to their biomass at MSY ( $B/B_{MSY}$ ) from a database of 331 stocks from ref (27) (Fig. S4).  
195 For the unfished situation, we compared our multi-model mean of all MEM-ESM combination  
196 without fishing ( $n = 10$ ) to the temperature-dependent hindcasts of maximum sustainable yield  
197 (MSY) for 235 stocks from ref (28) that are independent of the effects of fishing (Fig. 2). In  
198 addition to displaying the corresponding temporal trends, we also calculated linear regressions of  
199 projected biomass vs MSY in corresponding years, and found that the models explain a large  
200 fraction of the observed variance without fishing ( $R^2 = 0.44$ , Fig. 2) and an even larger fraction  
201 with fishing ( $R^2 = 0.96$ , Fig. S4). Importantly the linear regression fits with the ensemble mean  
202 were generally better than those with individual MEM-ESM combinations, with  $R^2$  ranging from  
203 0.13-0.43 without fishing (except for BOATS-IPSL,  $R^2 = 0.47$ ) and from 0.80-0.94 with fishing.  
204 This suggests that the ensemble mean is generally better at representing empirical observations  
205 than all but one individual MEMs.

206 To compare the magnitude of biomass declines across trophic levels and evaluate a trophic  
207 amplification effect (29), we calculated the mean and standard deviation of the relative change  
208 (2090s vs 1990s) in net primary production (NPP), total phytoplankton and total zooplankton  
209 biomass (both small and large) across the 2 ESMs ( $n = 2$ ) for each of the 4 RCPs. We then  
210 compared these mean biomass changes to those for higher trophic levels (Fig. 3) represented by

211 total animal biomass (which does not include zooplankton) across all MEM-ESM combinations  
212 ( $n = 10$  for RCP2.6 and 8.5,  $n = 8$  for RCP 4.5 and 6.0).

213 To relate the magnitude of change in total animal biomass (Fig. 4) and biomass of animals  
214  $>10$  cm and  $>30$  cm (Fig. S7) to changes in global air temperature since pre-industrial times, we  
215 obtained global air temperature timeseries from 1861-2100 for both ESMs from the CMIP5  
216 archive. We calculated the average pre-industrial temperature over the period 1861-1870, and  
217 then calculated subsequent changes in air temperature relative to this reference period. We then  
218 plotted the % change in total global animal biomass vs. the change in global air temperature  
219 across the historical and future period including all RCPs in an unfished ocean (Fig. 4). This was  
220 repeated for animals  $>10$  cm and  $>30$  cm (Fig. S7). To assess how this relationship might change  
221 in a fished ocean, we plotted the difference in % biomass change (RCP8.5 vs RCP2.6) in a fished  
222 versus unfished ocean over the difference in global surface air temperature between RCP8.5 vs  
223 RCP2.6 in any given year (Fig. S6B). The resulting positive relationship shows that as warming  
224 increases, the reduction in simulated biomass is slightly less in a fished compared to an unfished  
225 ocean.

226 To visualize spatial patterns of change globally, we mapped the relative (%) change in 2090-  
227 99 compared to 1990-99 for the multi-model mean and standard deviation of total animal  
228 biomass on a  $1 \times 1$ -degree grid across all MEM-ESM-combinations for RCP2.6 and RCP8.5 in an  
229 unfished ocean (Fig. 5). We also mapped the multi-model mean +1SD and -1SD (Fig. S8) to  
230 visually display the full range of minimum to maximum potential changes. As an additional  
231 measure of robustness, we mapped the percentage of MEMs agreeing on the direction of change  
232 (22), where 100% means full model agreement and 50% represents an even split. Next, to  
233 compare the climate effect in a fished and an unfished ocean spatially (Fig. S9), we mapped the

234 multi-model mean relative difference (%) in biomass change (RCP8.5 vs RCP2.6) in 2090-2099  
235 in a fished and in an unfished ocean based on the three MEMs that performed fishing runs  
236 (BOATS, DBEM, EcoOcean). To evaluate the spatial variability due to ESM selection in an  
237 unfished ocean (Fig. S10), we also mapped the multi-model mean biomass change and standard  
238 deviation across all MEMs for the IPSL-CM5A-LR and the GFDL-ESM2M separately. Lastly,  
239 we mapped relative changes in total animal biomass for each MEM for IPSL-CM5A-LR and  
240 GFDL-ESM2M under RCP2.6 and RCP8.5 in an unfished ocean (Fig. S11).

241

**Table S1.** Description of the basic features and key references of each of the six global fisheries and marine ecosystem models. For more detail see ref. (1).

<b>Model name</b>	<b>Basic model structure</b>	<b>Taxonomic scope</b>	<b>Key ecological processes</b>	<b>Spatial &amp; temporal resolution</b>	<b>Vertical resolution</b>	<b>Key references</b>
BiOeconomic mArine Trophic Size-spectrum (BOATS)	Size-structure model that uses macroecological theory and empirical metabolic constraints to calculate the production of commercially-harvested fish across multiple size spectra. Directly coupled to a dynamic fisheries economic model at the grid scale.	3 ‘superspecies’ (small, medium, large) defined by their asymptotic mass, with 50 size classes each, representing all commercial fish.	Applies empirical parameterizations to describe phytoplankton community structure, trophic transfer of primary production from phytoplankton to fish, fish growth rates, natural mortality of fish, stock- and environment-dependent recruitment. Animal movement is not included. Predator-prey relationships are not resolved.	1×1-degree grid Monthly mean timestep	None (2-dimensional domain).	2-3
Macroecological Model	Static size-structure model that uses minimal input parameters together with ecological and metabolic scaling theory to calculate mean size composition and abundance of marine animals (including fish).	180 body mass classes, species are not resolved.	Simple characterization of marine ecosystems in terms of body mass distribution and marine animal abundance based on estimates of predator-prey mass ratios, transfer efficiency and changing metabolic demands with body mass and temperature. Animal movement is not included.	1×1-degree grid Annual mean timestep	Single vertical (surface-integrated) layer.	4
Dynamic Pelagic Benthic Model (DPBM)	Dynamic size- and trait-based model that incorporates a pelagic predator size-spectrum with a benthic detritivore size-spectrum.	1 pelagic predator and 1 benthic detritivore size spectrum, with 100 size classes each.	Individual processes of predation, food-dependent growth, natural mortality, and reproduction give rise to emergent size spectra for each functional group (pelagic predator and benthic detritivore).	1×1-degree grid Monthly mean timestep	2 vertical layers (sea surface and sea floor). No vertical transport or movement.	5

Dynamic Bioclimate Envelope Model (DBEM)	Species distribution model based on bioclimatic envelopes (niche) defined for each species. Simulates changes in species abundance and carrying capacity under environmental change. Carrying capacity is a function of the environment and species' habitat preferences.	892 commercial fish and invertebrate species.	Population dynamics are dependent of habitat suitability and movement of adult species driven by a gradient of habitat suitability and population density. Larval dispersal is driven by currents and temperature. Growth, reproduction, and natural mortality are dependent on oxygen, pH, and temperature.	0.5×0.5-degree grid Annual mean ocean conditions	Vertical layers (sea surface and bottom) defined by species niche preferences.	6
EcoOcean	Trophodynamic model based on species interactions and energy transfer across trophic levels. Ecosim-with-Ecopath (EwE) framework designed to evaluate the impacts of fisheries and climate change on marine resources and ecosystems.	51 trophic biomass groups, including all trophic level and taxonomic groups (marine mammals, birds, fish, invertebrates, primary producers and bacteria).	Combines a food web model comprising a mass-balance component (Ecopath; input: biomass, production/biomass ratio, consumption/biomass ratio, diet composition, catches), a temporal dynamic predator-prey component (Ecosim), and a spatio-temporal dynamic component which is a function of grid cell specific habitat attributes i.e. pH, water depth, temperature, and bottom type (Ecospace).	1×1-degree grid Monthly mean timestep	Vertical layers defined by food web interactions and habitat preferences, vertical movement and transportation through establishment of trophic links and generation and consumption of dead organic matter linking pelagic organisms to demersal and benthic organisms.	7-8
Apex Predators ECOSystem Model (APECOSM)	Composite (hybrid) model. 3D dynamic energy budget Eulerian model of size-structured marine populations and communities, based on individual environmentally driven bioenergetics, trophic interactions and behaviors, that are upscaled to populations and communities.	Explicit size-based communities including 3 communities (epipelagic, migratory, mesopelagic); 95 species length classes and 100 size classes.	Size-based predation, food- and temperature-driven growth, reproduction and senescence. Includes environmental impacts on vertical and horizontal movements and schooling.	1×1-degree grid Monthly mean timestep	3D explicit vertical movement considered.	9-11

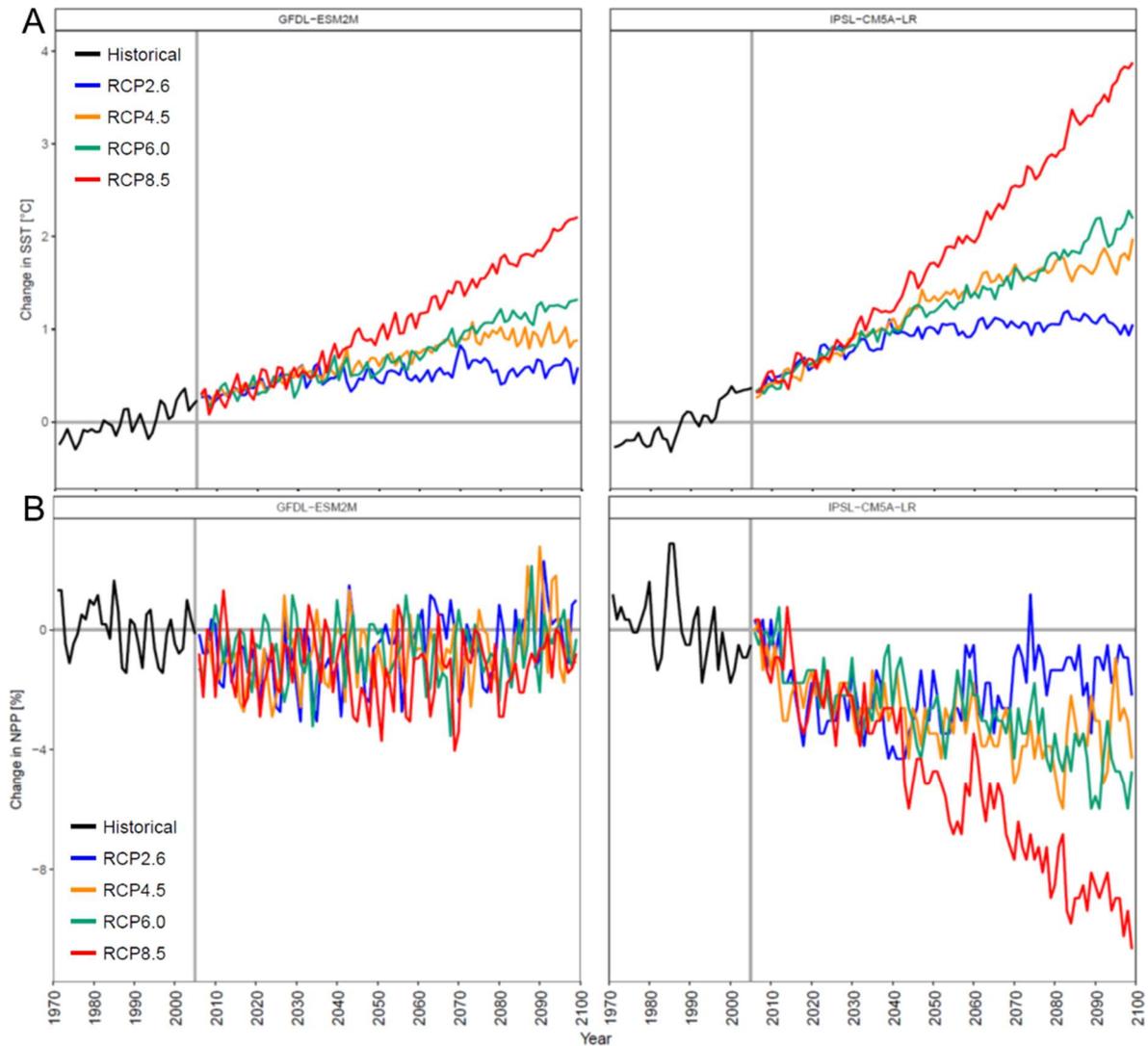
**Table S2.** Overview of layer configuration and selected forcing variables used in the marine ecosystem models included in the ensemble projections. Forcing variables were provided by the two Earth-system models GFDL-ESM2M (1×1-degree grid cell, 50 depth levels) and IPSL-CM5A-LR (1×1-degree grid cell, 31 depth levels). Layers included surface, bottom, depth-integrated surface to bottom, or depth resolved, depending on each ecosystem model's requirements. For more detail see ref. (1).

Model name	Depth integration	Current speed	Sea temperature	Dissolved oxygen concentration	NPP - primary organic carbon production	Phytoplankton carbon concentration	Zooplankton carbon concentration	pH	Salinity	Total alkalinity	Ice coverage	Mixed layer depth
BOATS	Integrated over full water column	-	Upper ocean temperature (average of upper 75m)	-	Depth integrated primary production (full water column)	-	-	-	-	-	-	-
Macro-ecological model	Integrated over full water column	-	Sea surface temperature (0-200m)	-	Depth integrated primary production (assumed to be allocated to the mixed layer depth or euphotic depth if deeper)	Large/small phytoplankton	-	-	-	-	-	Areas shallower than mixed layer depth (or euphotic depth if deeper) treated as productive zone
DPBM	2 layers, surface (0-100m) and bottom	-	Sea surface and sea bottom temperature	-	Depth integrated primary production	Large/small phytoplankton	-	-	-	-	-	Mixed layer depth included

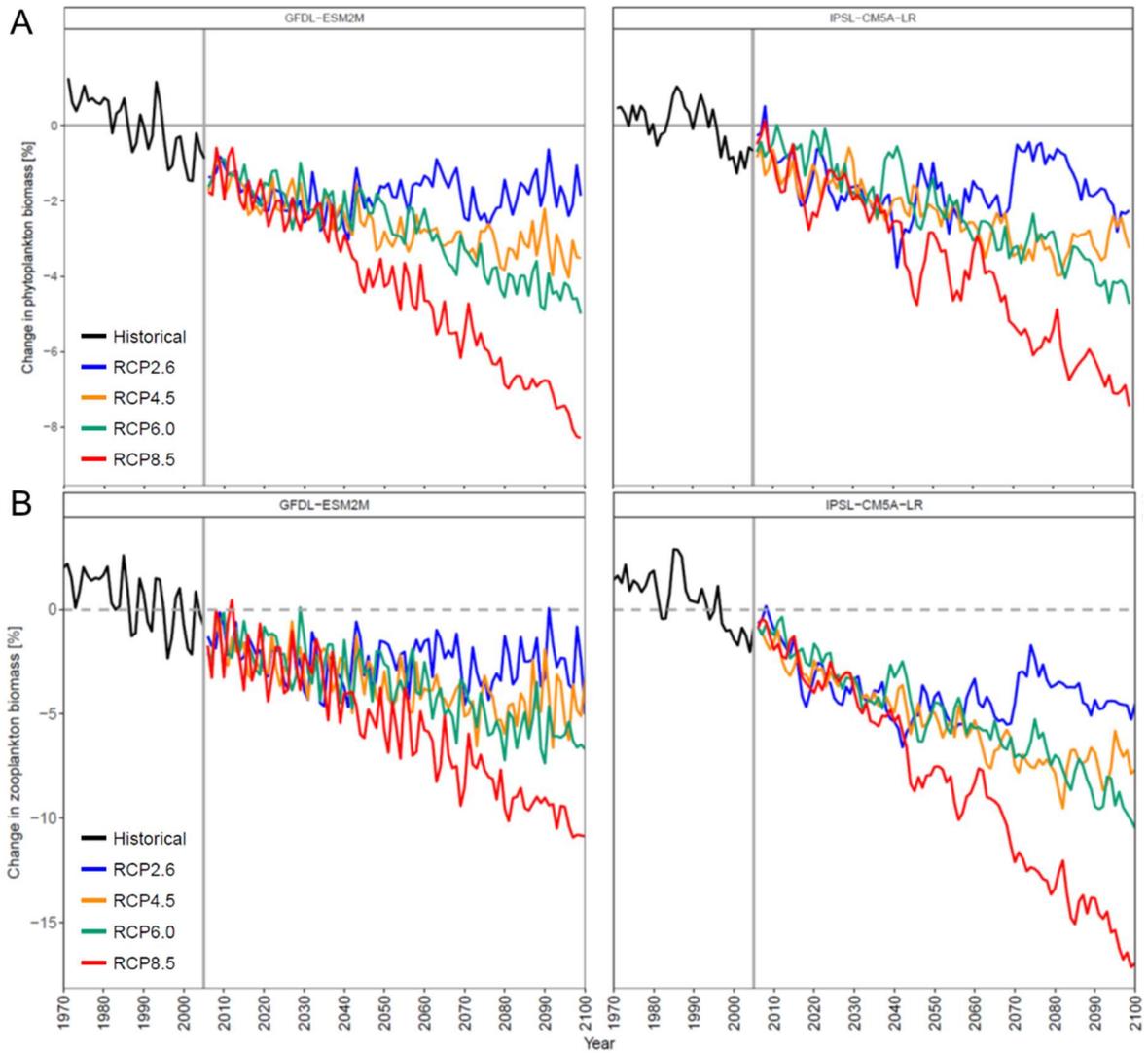
DBEM	Vertical dimension dependent on species specific min/max depth limits	Zonal and meridional velocity	Sea surface and sea bottom temperature	Dissolved oxygen included	Depth integrated primary production	Large/small phytoplankton	-	Sea surface pH	Sea surface and sea bottom salinity	-	Ice coverage incl.	-
EcoOcean	Vertical dimension dependent on depth distribution from species/functional groups	-	Sea surface (150 m)	-	-	Large/small phytoplankton	Large/small zooplankton	-	-	-	Ice coverage included (not floating ice)	-
APECOSM	3D depth-resolved	Zonal, meridional and vertical velocity	Vertically (3D) resolved sea temperature	Dissolved oxygen included	-	Large/small phytoplankton	Large/small zooplankton	pH included	-	Turbulent mixing included	Ice coverage included	-

**Table S3.** Summary of results across all marine ecosystem models (MEMs), Earth-system models (ESMs), emission scenarios (RCPs) and fishing scenarios. Shown is the projected percent (%) change in biomass in 2090-2099 relative to 1990-1999 for total animal biomass and animals >10 cm and >30 cm. Also shown are the mean, standard deviation (SD), and sample size (n) for the full set of available MEM-ESM combinations (n = 10), a reduced set 1 balancing the two ESMs (n = 8), a reduced set 2 balancing the ESMs and RCPs (n = 6), a reduced set 3 balancing the no-fishing and fishing runs (n = 6), and reduced set 4 with the same as set 3 but without DBEM for comparison (n = 4). Note that for DBEM, which does not differentiate between size groups but represents the full size range of commercial fish and invertebrates, we assumed the same relative biomass changes in all size groups.

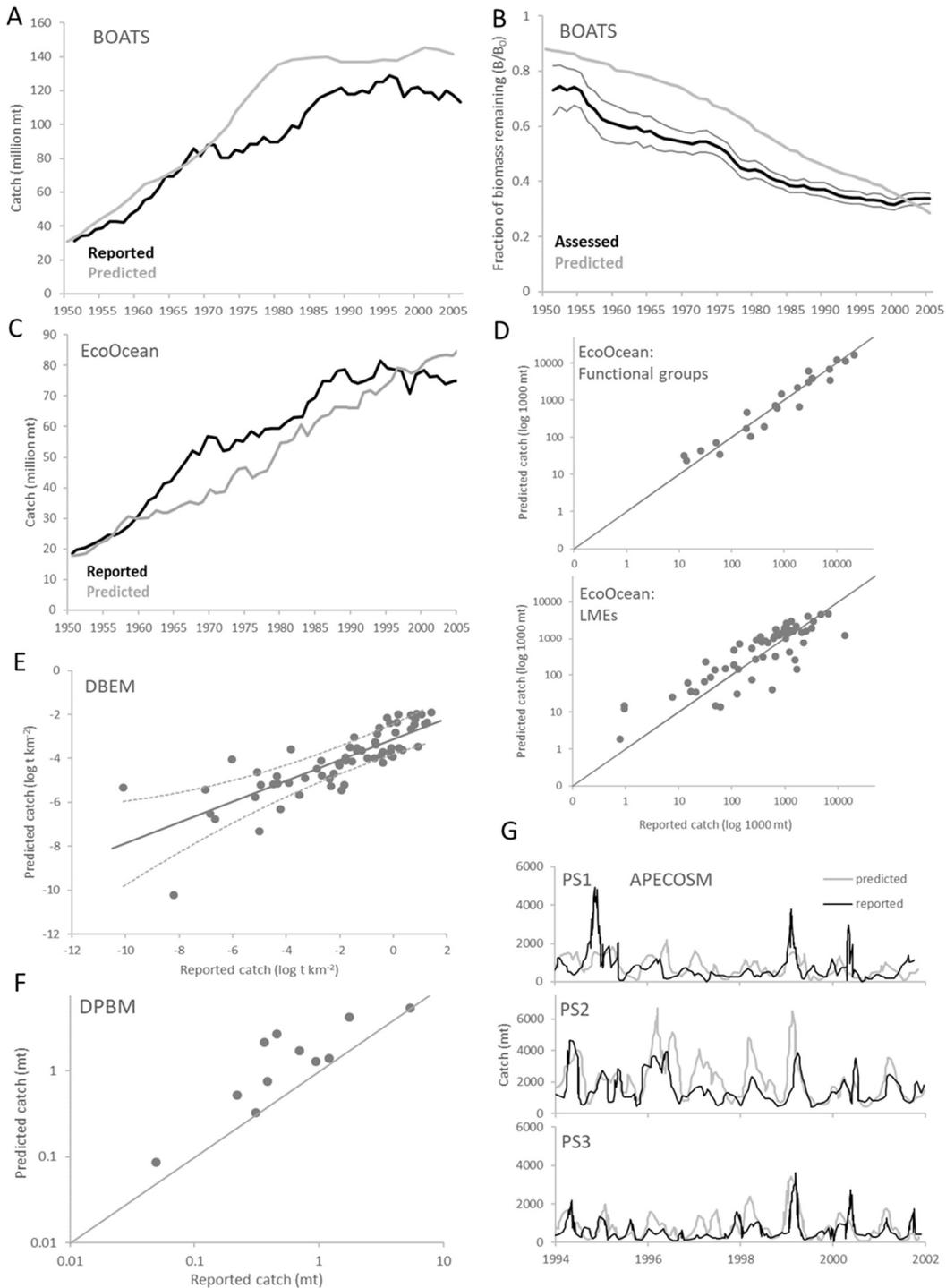
Percent biomass change (2090-2099 relative to 1990-1999)													
Model name	RCP	total		>10cm		>30cm		total		>10cm		>30cm	
		no fishing		no fishing		no fishing		fishing		fishing		fishing	
		GFDL	IPSL										
<b>BOATS</b>	<b>2.6</b>	-5.29	-10.26	-5.29	-10.26	-4.54	-9.67	-66.99	-25.62	-66.99	-25.62	-84.66	-69.22
	<b>4.5</b>	-8.68	-17.83	-8.68	-17.83	-8.37	-17.75	-67.26	-31.22	-67.26	-31.22	-84.49	-71.12
	<b>6.0</b>	-9.75	-20.47	-9.75	-20.47	-9.28	-20.48	-67.64	-33.11	-67.64	-33.11	-84.64	-72.15
	<b>8.5</b>	-15.07	-31.68	-15.07	-31.68	-14.59	-32.47	-68.50	-41.15	-68.50	-41.15	-84.74	-74.71
<b>EcoOcean</b>	<b>2.6</b>	0.23	-2.65	0.57	-4.12	3.55	6.25	-0.76	-3.91	2.99	-5.33	-16.90	-20.24
	<b>4.5</b>	0.07	-6.16	1.66	-6.51	4.45	3.93	-0.90	-7.21	4.41	-6.05	-17.11	-21.81
	<b>6.0</b>	0.07	-7.93	1.53	-8.07	5.51	2.20	-0.83	-9.02	4.35	-7.85	-17.83	-24.21
	<b>8.5</b>	-0.73	-13.60	6.41	-8.81	8.18	-5.28	-1.53	-14.20	10.00	-4.62	-19.54	-30.73
<b>Macroecological</b>	<b>2.6</b>	-4.39	-10.44	-4.39	-10.44	-4.39	-10.44						
	<b>4.5</b>	-7.16	-17.12	-7.16	-17.12	-7.16	-17.12						
	<b>6.0</b>	-9.27	-21.17	-9.27	-21.17	-9.27	-21.17						
	<b>8.5</b>	-15.48	-34.31	-15.48	-34.31	-15.48	-34.31						
<b>APECOSM</b>	<b>2.6</b>		-3.25		-3.54		-3.09						
	<b>4.5</b>		-4.88		-4.23		-3.44						
	<b>6.0</b>		-6.70		-6.45		-5.94						
	<b>8.5</b>		-10.74		-9.63		-9.01						
<b>DBPM</b>	<b>2.6</b>		-4.30		-5.08		-3.63						
	<b>4.5</b>		-6.92		-8.76		-7.61						
	<b>6.0</b>		-7.61		-11.18		-10.20						
	<b>8.5</b>		-12.06		-20.61		-21.68						
<b>DBEM</b>	<b>2.6</b>	-1.23	-6.71	-1.23	-6.71	-1.23	-6.71	-2.87	-10.04	-2.87	-10.04	-2.87	-10.04
	<b>8.5</b>	-10.29	-28.48	-10.29	-28.48	-10.29	-28.48	-11.05	-28.56	-11.05	-28.56	-11.05	-28.56
		<b>Mean</b>	<b>SD</b>										
<b>Full set</b>	<b>n</b>	10		10		10							
all available EM-ESM combinations	<b>2.6</b>	-4.83	3.51	-5.05	3.48	-3.39	5.25						
	<b>4.5</b>	-8.58	6.07	-8.58	6.42	-6.63	8.30						
	<b>6.0</b>	-10.35	7.13	-10.60	7.40	-8.58	9.45						
	<b>8.5</b>	-17.24	10.73	-16.80	12.38	-16.34	13.23						
<b>Reduced set 1</b>	<b>n</b>	8		8		8							
balancing ESMs	<b>2.6</b>	-5.09	3.99	-5.23	4.60	-3.40	24.48						
(without APECOSM and DPBM)	<b>4.5</b>	-9.48	7.33	-9.27	8.18	-7.00	28.21						
	<b>6.0</b>	-11.42	8.53	-11.20	9.78	-8.75	27.92						
	<b>8.5</b>	-18.70	13.37	-17.21	14.15	-16.59	23.00						
<b>Reduced set 2</b>	<b>n</b>	6		6		6							
balancing ESMs & RCPs	<b>2.6</b>	-5.47	4.32	-5.65	5.13	-3.21	29.14						
(without APECOSM, DPBM and DBEM)	<b>4.5</b>	-9.48	7.33	-9.27	8.18	-7.00	28.21						
	<b>6.0</b>	-11.42	8.53	-11.20	9.78	-8.75	27.92						
	<b>8.5</b>	-18.48	14.70	-16.49	15.62	-15.66	27.00						
<b>Reduced set 3</b>	<b>n</b>	6		6		6		6		6		6	
balancing no-fishing and fishing runs	<b>2.6</b>	-4.32	3.87	-4.51	3.88	-2.06	6.11	-18.36	25.47	-17.97	25.89	-33.99	34.15
	<b>8.5</b>	-16.64	11.59	-14.65	13.98	-13.82	15.05	-27.50	24.46	-23.98	28.27	-41.55	30.55
<b>Reduced set 4</b>	<b>n</b>	4		4		4		4		4		4	
set 3 with balanced size groups (without DBEM)	<b>2.6</b>	-4.50	4.46	-4.78	4.45	-1.10	7.32	-24.32	30.52	-23.73	31.24	-47.75	34.31
	<b>8.5</b>	-15.27	12.70	-12.29	15.76	-11.04	17.07	-31.34	29.78	-26.07	35.54	-52.43	32.11



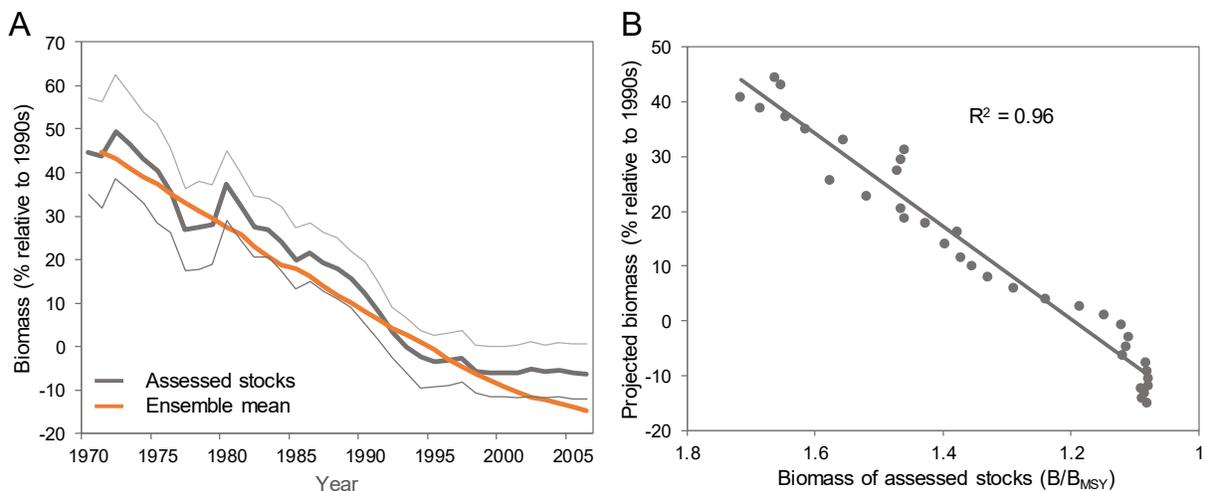
**Fig. S1.** Annual global change in (A) sea surface temperature (SST, °C) and (B) net primary production (NPP, %) relative to the 1990-1999s as projected with GFDL-ESM2M (left) and IPSL-CM5A-LR (right).



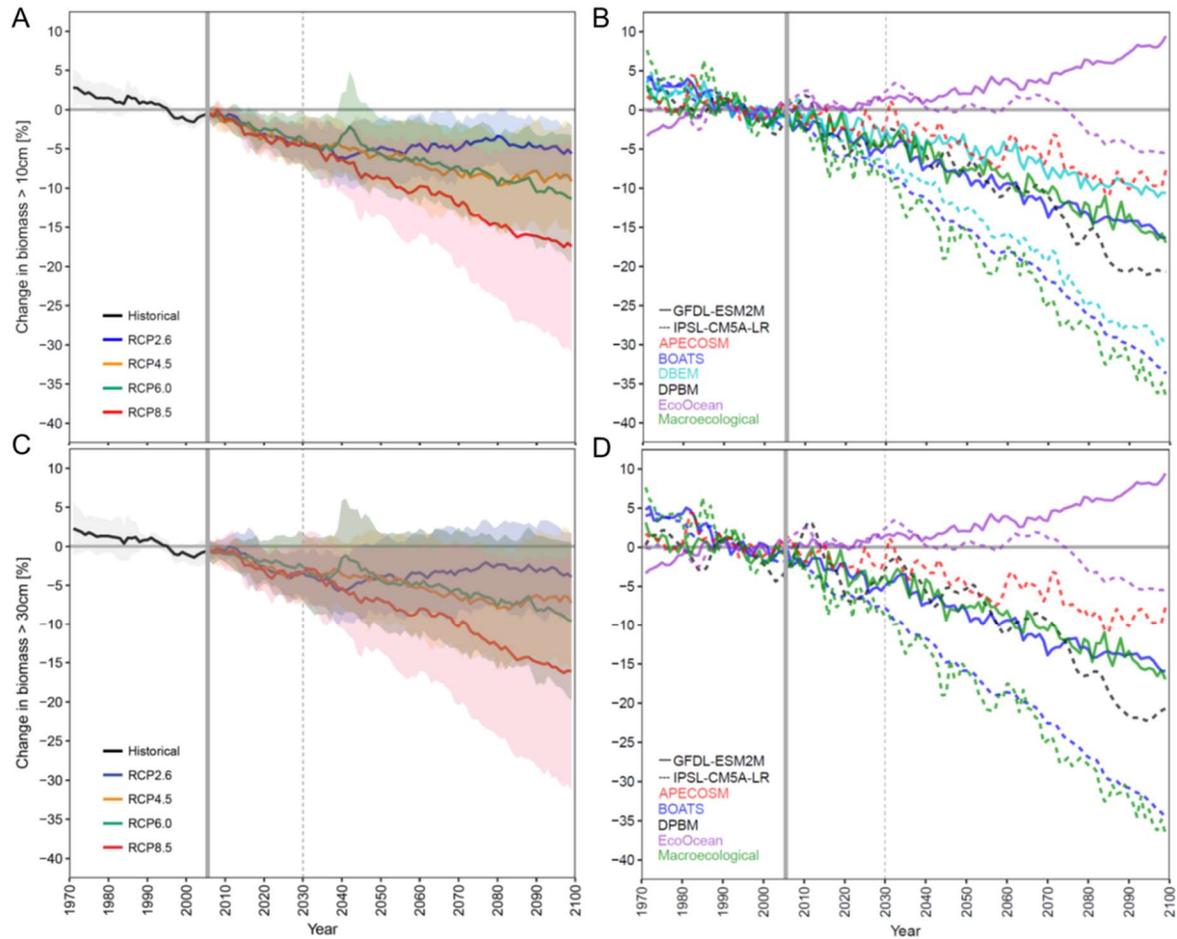
**Fig. S2.** Annual global change in (A) phytoplankton biomass (%) and (B) zooplankton biomass (%) relative to the 1990-1999s as projected with GFDL-ESM2M (left) and IPSL-CM5A-LR (right).



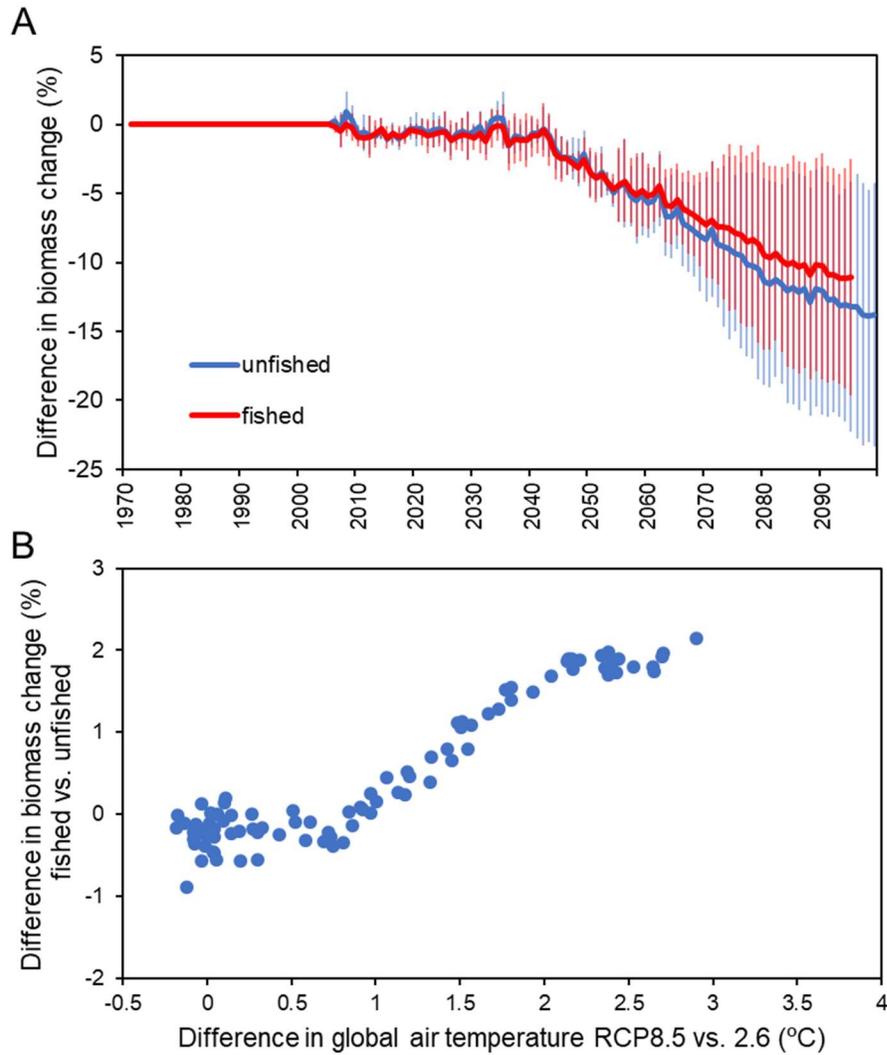
**Fig. S3.** Comparison of previously published model-predicted and observed data for individual MEMs. BOATS predicted and reported global catch (including unreported catch) from SAUP (A) and biomass trends ( $B/B_0$ ) from RAM database (B) (2, 3). EcoOcean predicted and reported global catch from SAUP (C) and catch by functional groups and LMEs (D) (7). DBEM predicted and reported catch by LME from SAUP (E) (13). DPBM time-averaged predicted catch and reported landings of 11 regions based on 78 EEZs from FAO (F) (5). APECOSM predicted and reported catch of 3 skipjack tuna schools (PS1-3) in the Indian Ocean (G) (18). For more detail see *SI Methods*.



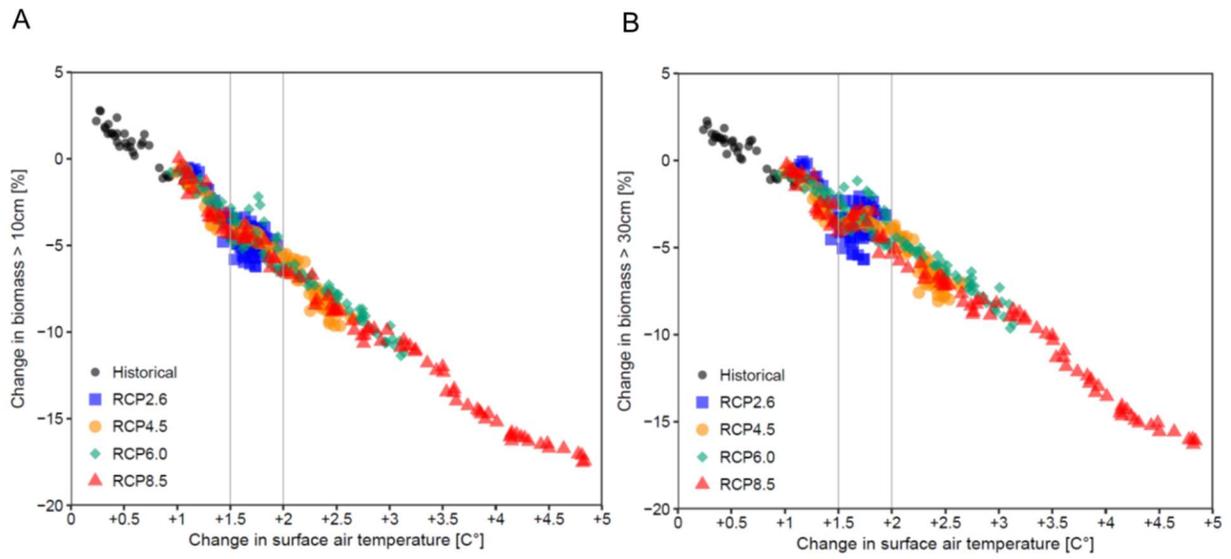
**Fig. S4.** Empirical validation in a fished ocean. Shown are the historical mean global biomass trends for our multi-model ensemble (orange line,  $n = 6$ ) with fishing compared with assessed exploited fish stocks (grey lines,  $\pm 95\%$  CI) (A) and the linear regression fit between the two time series (B). The ensemble mean is based on biomass for animals  $>10$  cm and assessed stocks represent the mean exploited biomass (relative to the biomass at maximum sustainable yield,  $B/B_{MSY}$ ) across 331 stocks (27). The stabilization of biomass since 1995 in assessed stocks is likely representing some successful management efforts for those stocks, which is not included in our global ensemble simulations.



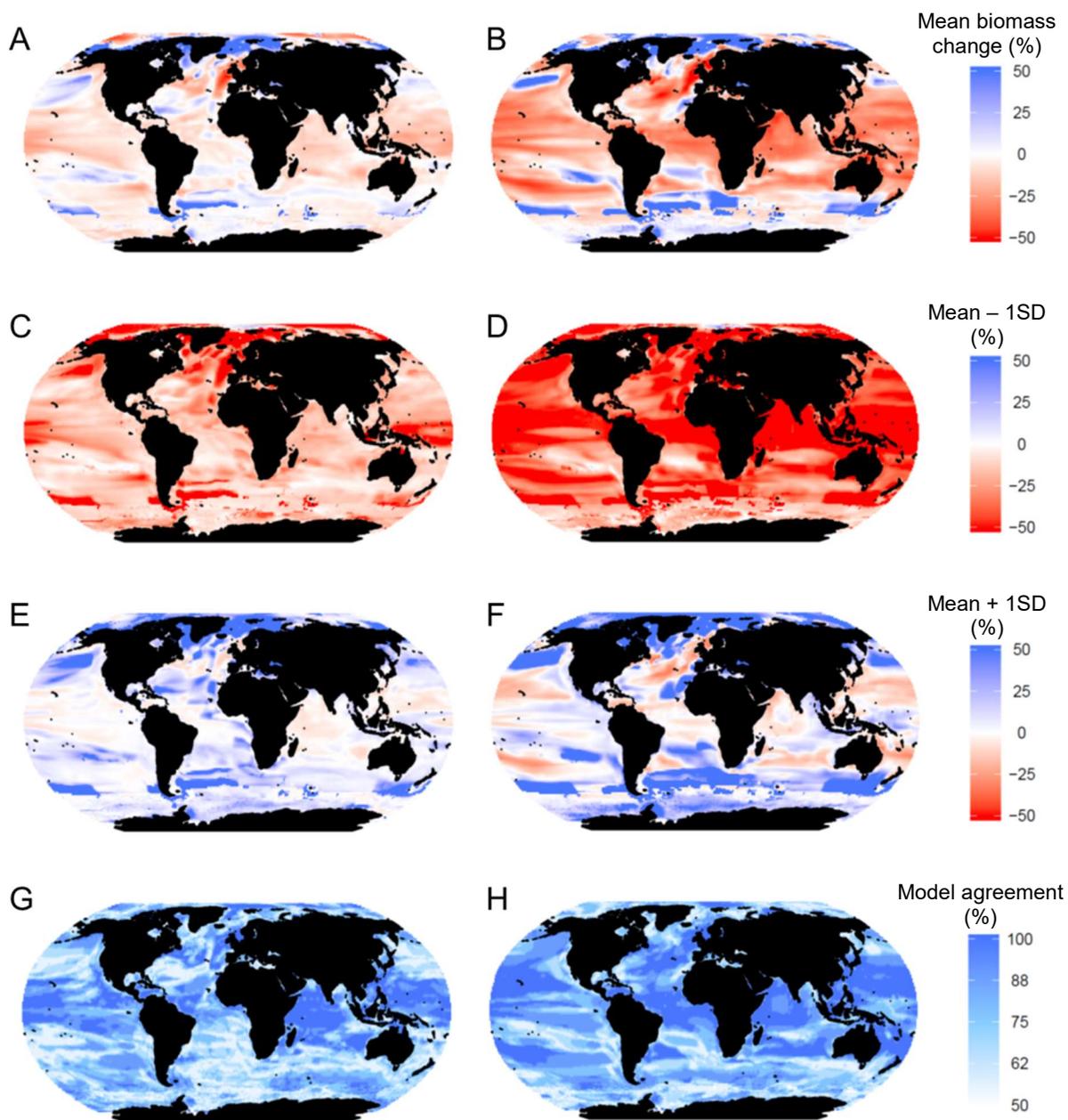
**Fig. S5.** Ensemble projections of global animal biomass with climate change from 1970-2100 for (A) animals >10 cm and (B) animals >30 cm. Left panels: Multi-model mean change in biomass for the historical period and four future emission scenarios (RCPs) with one inter-model standard deviation (shading). Right panels: Individual model projections for one emission scenario (RCP8.5) showing the spread across different ecosystem (color code) and ESMs (dashed/solid) combinations. All trends are relative to 1990-1999. The vertical grey line indicates the separation of historical and future projections, the vertical orange line indicates the SDG target year 2030.



**Fig. S6.** Comparing projected climate-change effects in a fished and an unfished ocean. (A) Mean difference and standard variation in biomass change (%) between RCP8.5 and RCP2.6 in a fished and an unfished ocean over time based on three ecosystem models (BOATS, EcoOcean, DBEM) combined with two ESMs ( $n = 6$ ). (B) Mean difference in biomass change (RCP8.5 vs. 2.6) in fished vs unfished oceans in relation to the differences in global air temperature between RCP8.5 vs. 2.6.

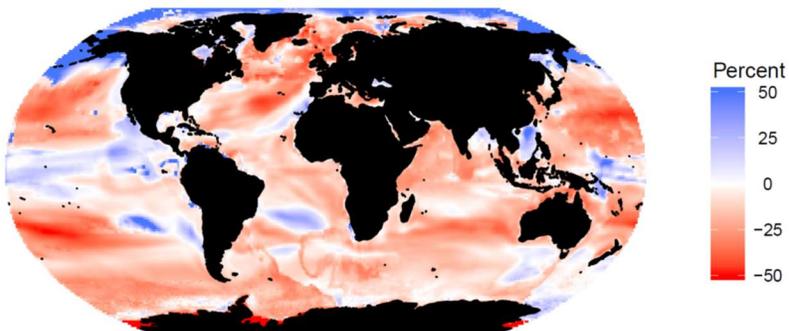


**Fig. S7.** Relationship between the projected change in (A) biomass of animals >10 cm and (B) animals >30 cm in the absence of fishing and increases in global air temperature since pre-industrial times (1860s). Changes in biomass are relative to the 1990s. The vertical lines frame the difference between 1.5 °C and 2 °C warming.

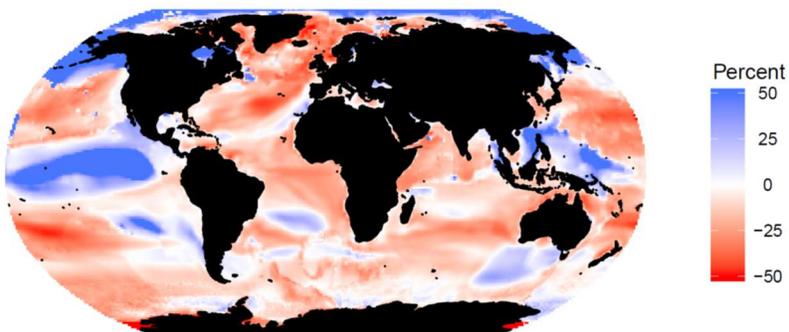


**Fig. S8.** Global spatial patterns of ensemble projection results for RCP 2.6 (left) and RCP 8.5 (right). (A, B) Multi-model mean change (%),  $n = 10$  in total marine animal biomass in 2090-99 relative to 1990-99 without fishing (as in Fig. 1A, B), and the multi-model mean (C, D) minus 1 standard deviation (%) and (E, F) plus 1 standard deviation (%); and (G, H) model agreement (%) on the direction of change.

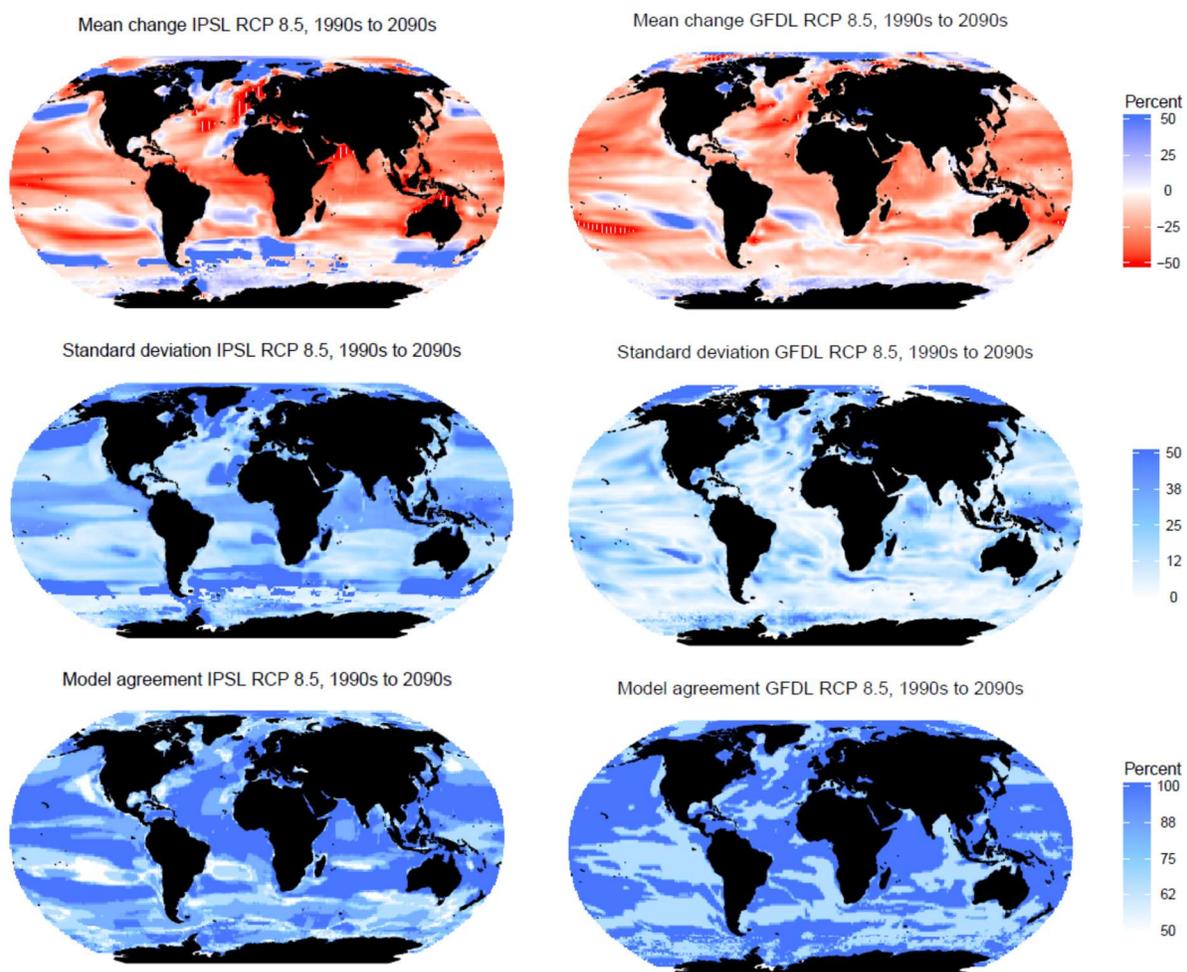
Fished ocean RCP8.5 / RCP2.6 2090s (relative change)



Unfished ocean RCP8.5 / RCP2.6 2090s (relative change)

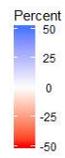
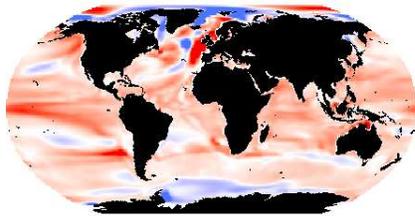


**Fig. S9.** Global spatial patterns of the climate effect defined as the multi-model mean biomass change in RCP8.5 vs. RCP2.6 in a fished (top) compared to an unfished (bottom) ocean based on three ecosystem models (BOATS, EcoOcean, DBEM) and two ESMs.

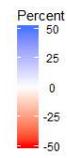
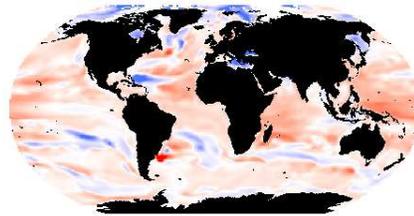


**Fig. S10.** Global spatial patterns of ensemble mean change (%) in total animal biomass in 2090-99 relative to 1990-99 for RCP8.5, the standard deviation, and robustness (% model agreement) among different ecosystem models for the two Earth-system models IPSL-CM5A-LR (left) and GFDL-ESM2M (right).

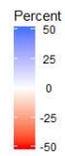
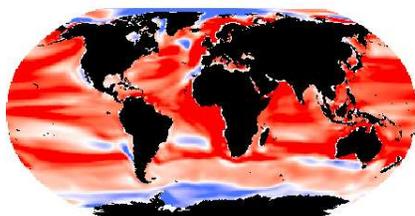
BOATS IPSL RCP2.6 no fishing (relative change)



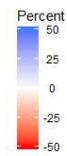
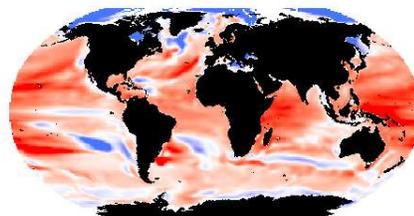
BOATS GFDL RCP2.6 no fishing (relative change)



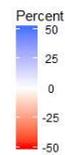
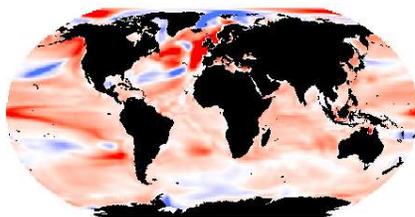
BOATS IPSL RCP8.5 no fishing (relative change)



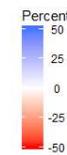
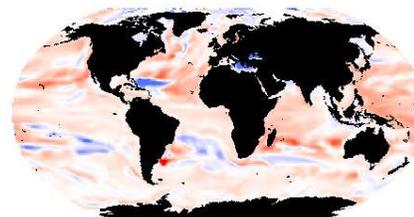
BOATS GFDL RCP8.5 no fishing (relative change)



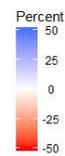
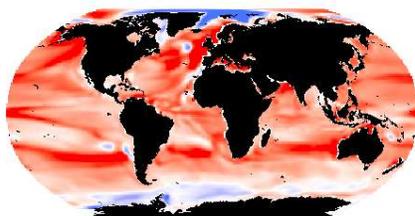
MACROECOLOGICAL IPSL RCP2.6 no fishing (relative change)



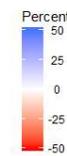
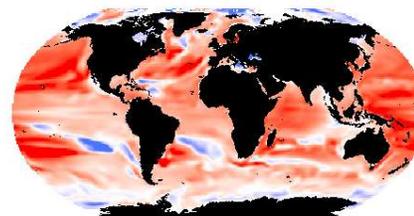
MACROECOLOGICAL GFDL RCP2.6 no fishing (relative change)



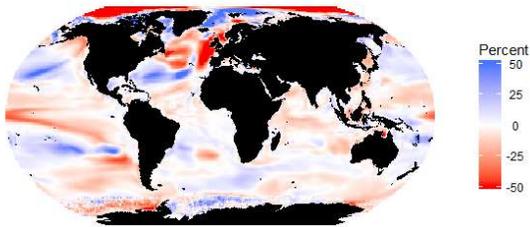
MACROECOLOGICAL IPSL RCP8.5 no fishing (relative change)



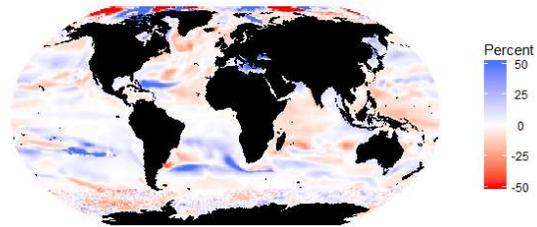
MACROECOLOGICAL GFDL RCP8.5 no fishing (relative change)



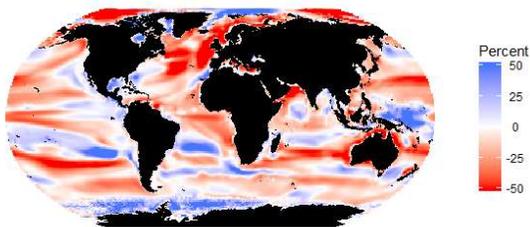
ECOOCEAN IPSL RCP2.6 no fishing (relative change)



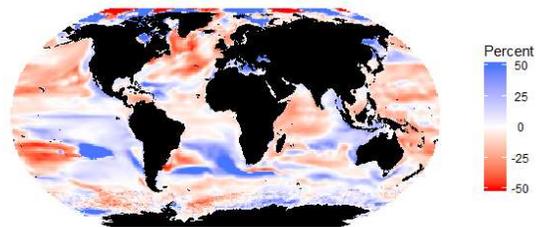
ECOOCEAN GFDL RCP2.6 no fishing (relative change)



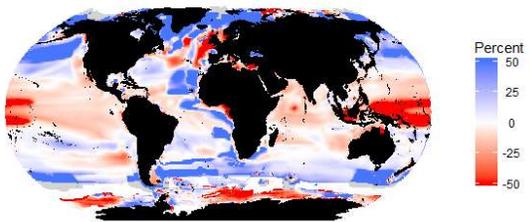
ECOOCEAN IPSL RCP8.5 no fishing (relative change)



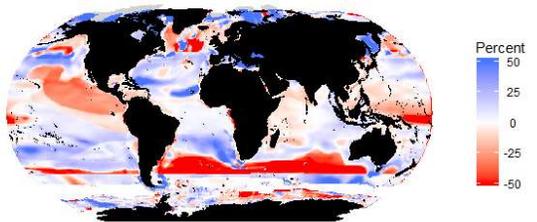
ECOOCEAN GFDL RCP8.5 no fishing (relative change)



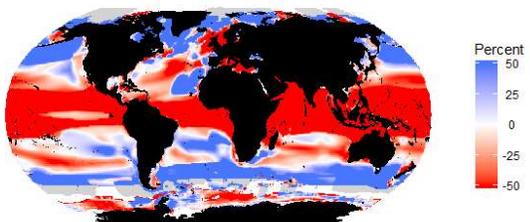
DBEM IPSL RCP2.6 no fishing (relative change)



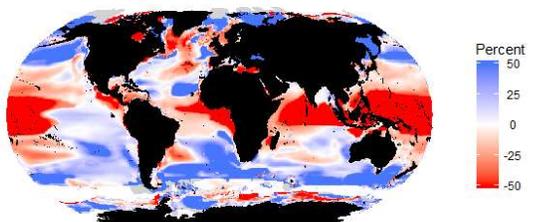
DBEM GFDL RCP2.6 no fishing (relative change)

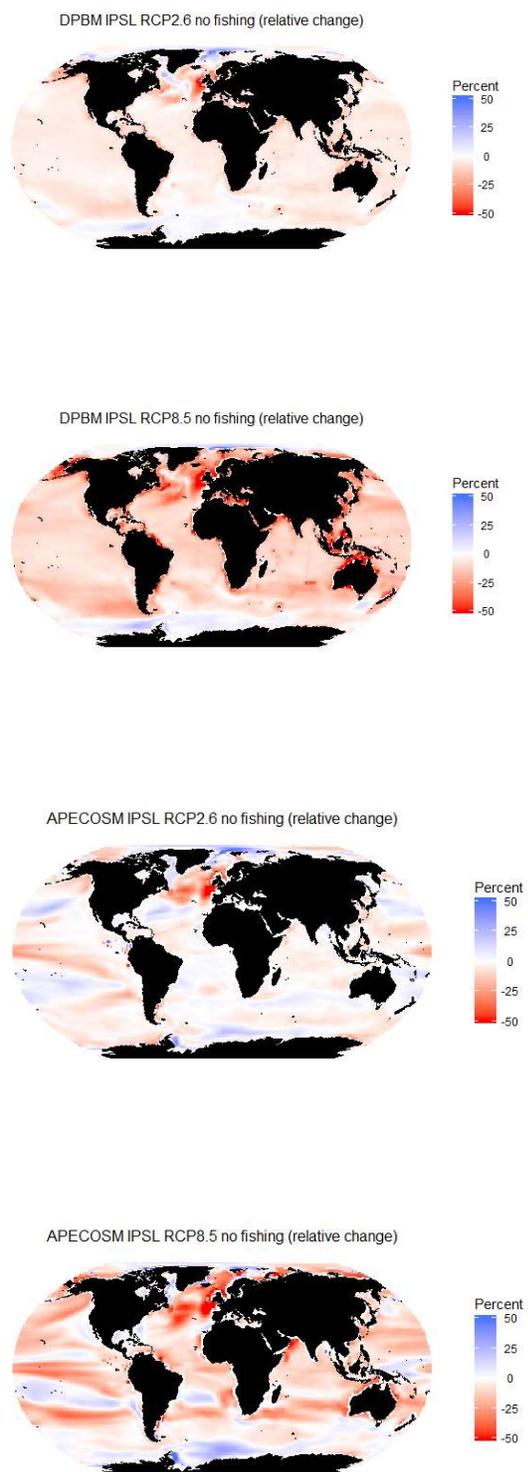


DBEM IPSL RCP8.5 no fishing (relative change)



DBEM GFDL RCP8.5 no fishing (relative change)





**Fig. S11.** Maps for each global ecosystem model highlighting relative changes in total animal biomass without fishing in the 2090s vs 1990s for IPSL-CM5A-LR and GFDL-ESM2M RCP2.6 and RCP8.5.

## References for SI reference citations

1. Tittensor DP, et al. (2018) A protocol for the intercomparison of marine fishery and ecosystem models: Fish-MIP v1.0. *Geosci Model Dev* 11:1421-1442.
2. Galbraith ED, Carozza DA, Bianchi D (2017) A coupled human-Earth model perspective on long-term trends in the global marine fishery. *Nature Comm* 8:14884.
3. Carozza DA, Bianchi D, Galbraith ED (2017) Formulation, general features and global calibration of a bioenergetically-constrained fishery model. *PLoS ONE* 12:e0169763.
4. Jennings S, Collingridge K (2015) Predicting consumer biomass, size-structure, production, catch potential, responses to fishing and associated uncertainties in the world's marine ecosystems. *PLoS One* 10:e0133794.
5. Blanchard JL, et al. (2012) Potential consequences of climate change for primary production and fish production in large marine ecosystems. *Phil Trans R Soc B* 367:2979–2989.
6. Cheung WWL, Dunne J, Sarmiento JL, Pauly D (2011). Integrating ecophysiology and plankton dynamics into projected maximum fisheries catch potential under climate change in the Northeast Atlantic. *ICES J Mar Sci* 68:1008–1018.
7. Christensen V, et al. (2015) The global ocean is an ecosystem: simulating marine life and fisheries. *Glob Ecol Biogeogr* 24:507–517.
8. Christensen V, Walters CJ (2004) Ecopath with Ecosim: methods, capabilities and limitations. *Ecol Model* 172:109–139.
9. Maury O (2010) An overview of APECOSM, a spatialized mass balanced “Apex Predators ECOSystem Model” to study physiologically structured tuna population dynamics in their ecosystem. *Prog Oceanogr* 84:113–117.

10. Maury O, et al. (2007) Modelling environmental effects on the size-structured energy flow through marine ecosystems. Part 2: simulations. *Prog Oceanogr* 74:500-514.
11. Maury O, Poggiale J-C (2013) From individuals to populations to communities: a Dynamic Energy Budget model of marine ecosystem size-spectrum including life history diversity. *J Theoret Biol* 324:52–71.
12. Carozza DA, Bianchi D, Galbraith ED (2016) The ecological module of BOATS-1.0: a bioenergetically-constrained model of marine upper trophic levels suitable for studies of fisheries and ocean biogeochemistry. *Geosci Model Dev* 9:1545-1565.
13. Cheung WWL, et al. (2016) Structural uncertainty in projecting global fisheries catches under climate change. *Ecol Model* 325:57-66.
14. Cheung WWL, Reygondeau G, Frölicher TL (2016) Large benefits to marine fisheries of meeting the 1.5°C global warming target. *Science* 354:1591–1594.
15. Blanchard JL, et al. (2009) How does abundance scale with body size in coupled size-structured food webs? *J Anim Ecol* 78:270–280.
16. Kooijman SALM (2010) *Summary of concepts of dynamic energy budget theory for metabolic organization*. Cambridge University Press, Cambridge.
17. Bianchi D, Galbraith ED, Carozza DA, Mislan KAS, Stock CA (2013) Intensification of open-ocean oxygen depletion by vertically migrating animals. *Nat Geosci* 6:545–548.
18. Dueri S, Faugeras B, Maury O (2012) Modelling the skipjack tuna dynamics in the Indian Ocean with APECOSM-E – Part 2: Parameter estimation and sensitivity analysis. *Ecol Model* 245:55–64.

19. Aumont O, Maury O, Lefort S, Bopp L (2018) Evaluating the potential impacts of the diurnal vertical migration by marine organisms on marine biogeochemistry. *Glob Biogeochem Cycles* 32:1622-1643.
20. Dunne JP, et al. (2012) GFDL's ESM2 global coupled climate-carbon Earth System Models. Part I: Physical formulation and baseline simulation characteristics. *J Climate* 25:6646-6665.
21. Dufresne JL, et al. (2013) Climate change projections using the IPSL-CM5 Earth System Model: from CMIP3 to CMIP5. *Climate Dynamics* 40:2123-2165.
22. Bopp L, et al. (2013) Multiple stressors of ocean ecosystems in the 21st century: projections with CMIP5 models. *Biogeosciences* 10:6225–6245.
23. Van Vuuren DP, et al. (2011) The representative concentration pathways: an overview. *Climatic Change* 109:5-31.
24. Masui T, et al. (2011) An emission pathway for stabilization at 6 Wm<sup>-2</sup> radiative forcing. *Climatic Change* 109:59-76.
25. Riahi K, et al. (2011) RCP 8.5 – A scenario of comparatively high greenhouse gas emissions. *Climatic Change* 109:33-57.
26. Maury O, et al. (2017) From shared socio-economic pathways (SSPs) to oceanic system pathways (OSPs): Building policy-relevant scenarios for global oceanic ecosystems and fisheries. *Glob Environ Change* 45:203–216.
27. Worm B, Branch TA (2012) The future of fish. *Trends Ecol Evol* 27:594–599.
28. Free CM, et al. (2019) Impacts of historical warming on marine fisheries production. *Science* 363:979–983.
29. Kwiatkowski L, Aumont O, Bopp L (2018) Consistent trophic amplification of marine biomass declines under climate change. *Glob Change Biol* 25:218–229.