



**A simple open source  
model for analyses of  
the global carbon  
cycle – Hector v0.1**

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# A simple object-oriented and open source model for scientific and policy analyses of the global carbon cycle – Hector v0.1

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## Abstract

Simple climate models play an integral role in policy and scientific communities. They are used for climate mitigation scenarios within integrated assessment models, complex climate model emulation, and uncertainty analyses. Here we describe Hector v0.1, an open source, object-oriented, simple global climate carbon-cycle model. This model runs essentially instantaneously while still representing the most critical global scale earth system processes. Hector has three main carbon pools: an atmosphere, land, and ocean. The model's terrestrial carbon cycle includes respiration and primary production, accommodating arbitrary geographic divisions into, e.g., ecological biomes or political units. Hector's actively solves the inorganic carbon system in the surface ocean, directly calculating air–sea fluxes of carbon and ocean pH. Hector reproduces the global historical trends of atmospheric [CO<sub>2</sub>] and surface temperatures. The model simulates all four Representative Concentration Pathways with high correlations ( $R > 0.7$ ) with current observations, MAGICC (a well-known simple climate model), and the Coupled Model Intercomparison Project version 5. Hector is freely available under an open source license, and its modular design will facilitate a broad range of research in various areas.

## 1 Introduction

Projecting future impacts of anthropogenic perturbations on the climate system relies on understanding the interactions of key earth system processes. To accomplish this, a hierarchy of climate models with differing levels of complexity and resolution are used, ranging from simple energy balance models to fully-coupled atmosphere–ocean–general circulation models (AOGCMs) (Stocker, 2011).

Simple climate models (SCMs) represent only the most critical global scale earth system processes with low spatial and temporal resolution, e.g., carbon fluxes between the ocean and atmosphere, and respiration and primary production on land.

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## 2.2 Model coupler

Hector's control flow starts with the coupler, which is responsible for: (1) parsing and routing input data to the model components, (2) tracking how the components depend on each other, (3) passing messages and data between components, (4) providing facilities for logging, time series interpolation, etc., and (5) controlling the main model loop as it progresses through time. Any errors thrown by the model are caught by the wrapper, which prints a detailed summary of the error.

Input data are specified in flat text files, and during startup are routed to the correct model component for its initialization. Some of the key initial model conditions are summarized in Tables 1 and 2. For more details of initial model conditions we urge the reader to download Hector v0.1 (<https://github.com/JGCRI/hector>). Components can send messages to each other during the model run, most often requesting data. The coupler handles message routing (via the *capability* mechanism, below) and enforces mandatory type checking: e.g., if a component requests mean global temperature in °C but the data are provided in K, an error will be thrown (i.e., execution halts) unless the receiving component can handle this situation.

Visitor patterns are units of code that traverse all model components and handle model output (Martin et al., 1997). Two visitors currently exist: one saves an easily-readable summary table to an output file, while the other writes a stream of model data (both standard outputs and internal diagnostics). After the model is finished running, this "stream" file can be parsed and summarized by R scripts included with the code (R Development Core Team, 2014). Log files may also be written by any model entity, using facilities provided by the coupler. The full sequence of events during a model run is summarized in Fig. 1.

## 2.3 Components

Model components are submodels that communicate with the coupler. From the coupler's point of view, components are fully defined by their *capabilities* and

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### 3 Main carbon cycle

In the model's default terrestrial carbon cycle, terrestrial vegetation, detritus, and soil are linked with each other and the atmosphere by first-order differential equations (Fig. 2). Vegetation net primary production is a function of atmospheric  $[\text{CO}_2]$  and temperature. Carbon flows from the vegetation to detritus and then soil, losing fractions to heterotrophic respiration on the way. Land-use emissions are specified as inputs. An "earth" pool debits carbon emitted as anthropogenic emissions, allowing a continual mass-balance check across the entire carbon cycle.

More formally, any change in atmospheric carbon, and thus  $[\text{CO}_2]$ , occurs as a function of anthropogenic emissions, land-use change emissions, and the atmosphere-ocean carbon flux. The atmosphere is treated as a single well-mixed box whose rate of change is:

$$\frac{dC_{\text{atm}}}{dt} = F_A(t) + F_{\text{LC}}(t) - F_O(t) - F_L(t) \quad (1)$$

where,  $F_A$  is the anthropogenic emissions,  $F_{\text{LC}}$  is the land use change emissions and  $F_O$  and  $F_L$  are the atmosphere ocean and atmosphere land fluxes. The overall terrestrial carbon balance at time  $t$  is the difference between net primary production (NPP) and heterotrophic respiration (RH). This is summed over user-specified  $n$  groups (each typically regarded as a latitude band, biome, or -political units), with  $n \geq 1$ :

$$F_L(t) = \sum_{i=1}^n \text{NPP}_i(t) - \text{RH}_i(t) \quad (2)$$

Note that NPP here is assumed to include disturbance effects, for which there is currently no separate term. For each biome  $i$ , NPP and RH are computed as functions of their preindustrial values  $\text{NPP}_0$  and  $\text{RH}_0$ , current atmospheric carbon  $C_{\text{atm}}$ , and the

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biome's temperature anomaly  $T_i$ :

$$\text{NPP}_i(t) = \text{NPP}_0 \cdot f(C_{\text{atm}}, \beta_i) \quad (3)$$

$$f(C_{\text{atm}}, \beta_i) = 1 + \beta_i(C_{\text{atm}}/C_0) \quad (4)$$

$$\text{RH}_i(t) = \text{RH}_0 \times Q_{10i}^{T_i(t)/10} \quad (5)$$

$$T_i(t) = T_G(t) \times \delta_i \quad (6)$$

These are commonly used formulations: NPP is modified by the user-specified carbon fertilization parameter,  $\beta$  (Piao et al., 2013). RH changes are controlled by a biome-specific  $Q_{10}$  value. Biomes can experience temperature changes at rates that differ from the global mean  $T_G$ , controlled by a user specified temperature factor  $\delta_i$ .

Land carbon pools (vegetation, detritus, and soil) change as a result of NPP, RH, and land-use change fluxes, whose effects are partitioned among these carbon pools. In addition, carbon flows from vegetation to detritus and soil (Fig. 2). Partitioning fractions ( $f$ ) control the flux quantities between pools (Table 2). For simplicity Eqs. (8)–(10) omit the time  $t$  and biome-specific  $i$  notations, but each pool is tracked separately for each biome at each time step:

$$\frac{dC_V}{dt} = \text{NPP} f_{\text{nv}} - C_V(f_{\text{vd}} + f_{\text{vs}}) - F_{\text{LC}} f_{\text{lv}} \quad (7)$$

$$\frac{dC_D}{dt} = \text{NPP} f_{\text{nd}} + C_V f_{\text{vd}} - C_D f_{\text{ds}} - \text{RH} f_{\text{rd}} - F_{\text{LC}} f_{\text{ld}} \quad (8)$$

$$\frac{dC_S}{dt} = \text{NPP} f_{\text{ns}} + C_V f_{\text{vs}} + C_D f_{\text{ds}} - \text{RH} f_{\text{rs}} - F_{\text{LC}} f_{\text{ls}} \quad (9)$$

The ocean–atmosphere carbon flux is the sum of the ocean's surface fluxes (currently  $n = 2$ , high and low latitude surface box):

$$F_O(t) = \sum_{i=1}^n F_i(t) \quad (10)$$

The surface fluxes of each individual box are calculated from an ocean chemistry model described in detail by Hartin et al. (2014) based on equations from Zeebe and Wolf-Gladrow, (2001). The flux of CO<sub>2</sub> for each box *i* is calculated by:

$$F_i(t) = k\alpha\Delta p\text{CO}_2 \quad (11)$$

Where *k* is the CO<sub>2</sub> gas-transfer velocity, *α* is the solubility of CO<sub>2</sub> in water based on salinity, temperature, and pressure, and Δ*p*CO<sub>2</sub> is the atmosphere–ocean gradient of *p*CO<sub>2</sub> (Takahashi et al., 2009). At steady state, the cold high latitude surface box (> 55°, subpolar gyres) acts as a sink of carbon from the atmosphere, while the warm low latitude surface box (< 55°) off gases carbon back to the atmosphere. Temperatures of the surface boxes are linearly related to atmospheric global temperatures (see Sect. 4.1), *T*<sub>HL</sub> = Δ*T* – 13 and *T*<sub>LL</sub> = Δ*T* + 7 (Lenton, 2000). The ocean model, modeled after Lenton et al. (2000) and Knox and McElroy (1984), circulate carbon through four boxes (two surface, one intermediate depth, one deep), via water mass advection and exchange, simulating a simple thermohaline circulation (Fig. 2). At steady state, approximately 100 Pg of carbon are transferred from the high latitude surface box to the deep box based on the volume of the box and transport in Sv (10<sup>6</sup> m<sup>3</sup> s<sup>-1</sup>) between the boxes. The change in carbon of any box *i* is given by the fluxes in and out:

$$\frac{dC_i}{dt} = \sum_{j=1}^{\text{in}} F_{j \rightarrow i} - \sum_{j=1}^{\text{out}} F_{i \rightarrow j} + F_i \quad (12)$$

As the model advances, the carbon values or DIC change in each box. The new DIC values are used within the chemistry submodel to calculate *p*CO<sub>2</sub> values at the next time step.

### 3.1 Adaptive-time step solver

The fundamental time step in Hector is currently one year, and most model components are solved at this resolution. The carbon cycle, however, can operate on a variable time step, helping to stabilize it under particularly high-emissions scenarios. This

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## 4.2.1 Halocarbons

The halocarbon component of the model can accept an arbitrary number of gas species, each characterized by a name, a lifetime  $\tau$  (yr), a radiative forcing efficiency  $\rho$  ( $\text{W m}^{-2} \text{pptv}^{-1}$ ), a user-specified preindustrial concentration (pptv), and a molar mass (g). For each gas, its concentration ( $C_i$ ) at time  $t$  is then computed based on a specified emissions time series  $E$ , assuming an exponential decay from the atmosphere:

$$C_i(t) = C_i(t-1) \left(1 - \frac{1}{\tau}\right) + E_i(t) \quad (15)$$

The default model input files include these parameters and a time series of emissions for C2F6, CCl4, CF4, CFC11, CFC12, CFC113, CFC114, CFC115, CH3Br, CH3CCl3, CH3Cl, HCF22, HCF141b, HCF142b, HFC23, HFC32, HFC125, HFC134a, HFC143a, HFC227ea, HFC245ca, HFC245fa, HFC4310, SF6, halon1211, halon1301, and halon2402.

Radiative forcing by halocarbons, other gases controlled under the Montreal Protocol, SF<sub>6</sub>, and ozone are calculated via:

$$\text{RF} = \alpha[C(t) - C(t_0)] \quad (16)$$

where  $\alpha$  is the radiative efficiency in  $\text{W m}^{-2} \text{ppbv}^{-1}$ , and  $C$  is the atmospheric concentration.

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## 4.2.2 Ozone

Tropospheric ozone concentrations are calculated by the CH<sub>4</sub> concentration and the emissions of three primary pollutants: NO<sub>x</sub>, CO, and NMVOCs:

$$\begin{aligned} O_3(t) = O_3(2000) + 5.0 \ln \left[ \frac{CH_4(t)}{CH_4(2000)} \right] + 0.125[eNO_x(t) - eNO_x(2000)] \\ + 0.0011[eCO(t) - eCO(2000)] \\ + 0.0033[eVOC(t) - eVOC(2000)] \end{aligned} \quad (17)$$

where the constants are the ozone sensitivity factors for each of the precursors (Ehhalt et al., 2001). The radiative forcing of tropospheric ozone is calculated from a linear relationship using a radiative efficiency factor (Joos et al., 2001) and a pre-industrial value of ozone of 25 DU (IPCC, 2001):

$$RF_{O_3} = 0.042 \text{ W m}^{-2} \text{ DU}^{-1} \times [[O_3] - [O_3]_{pre}] \quad (18)$$

## 4.2.3 BC and OC

The radiative forcing from black carbon is a function of the black carbon and organic carbon emissions (*eBC* and *eOC*).

$$RF_{BC} = 0.0743 \times 10^{-9} \text{ W m}^{-2} \text{ kg}^{-1} \times eBC \quad (19)$$

$$RF_{OC} = -0.0128 \times 10^{-9} \text{ W m}^{-2} \text{ kg}^{-1} \times eOC \quad (20)$$

The coefficients  $0.0743 \times 10^{-9}$  and  $-0.0128 \times 10^{-9}$  include both indirect and direct forcings of black and organic carbon (fossil fuel and biomass) (Bond et al., 2013).

## 4.2.4 Sulphate aerosols

The radiative forcing from sulphate aerosols is a combination of the direct and indirect forcings (Joos et al., 2001).

$$RF_{SO_x \text{ Direct}} = -0.4 W m^{-2} \times \frac{eSO_x(t)}{eSO_x(2000)} \quad (21)$$

$$5 \quad RF_{SO_x \text{ Indirect}} = -0.8 W m^{-2} \times \frac{(\ln(eSN) + eSO_x(t))}{eSN} \times \left( \ln \frac{eSN + eSO_x(2000)}{eSN} \right)^{-1} \quad (22)$$

The direct forcing by sulphate aerosols is proportional to the anthropogenic sulphur emissions ( $GgS yr^{-1}$ ) divided by the sulphate emissions from 2000. The indirect forcing by sulphate aerosols is a function of the anthropogenic and natural sulphur emissions. Natural sulphur emissions denoted by  $eSN$  is equal to 42 000  $GgS$ . A time series of annual mean volcanic stratospheric aerosol forcing ( $W m^{-2}$ ) is supplied from Meinshausen et al. (2011b) and is added to the indirect and direct forcing for a total sulphate forcing.

## 4.2.5 $N_2O$ and $CH_4$

15 The radiative forcing equations for  $CH_4$  and  $N_2O$  (Joos et al., 2001) are a function of the concentrations (ppbv) and their radiative efficiency:

$$RF_{CH_4} = 0.036 W m^{-2} \left[ \sqrt{CH_4(t)} - \sqrt{CH_4(t_0)} \right] - f[CH_4(t), N_2O(t_0)] - f[CH_4(t_0), N_2O(t_0)] \quad (23)$$

$$15 \quad RF_{N_2O} = 0.12 W m^{-2} \left[ \sqrt{N_2O(t)} - \sqrt{N_2O(t_0)} \right] - f[CH_4(t_0), N_2O(t)] - f[CH_4(t_0), N_2O(t_0)] \quad (24)$$

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details or parameterizations typically found in large scale complex models, but instead to represent only the most critical global processes. This allows for fast execution times, ease of understanding and straightforward analysis of the model output. To help with the analysis of Hector we included within the online database of Hector, R scripts to process Hector's output as well as the comparison data.

Hector's two key features are its open source license and modular design. This allows the user to manipulate the input files, enable/disable/replace components, or include components not found within the core version of Hector. For example, the user can design a new submodel (e.g., sea-ice) to answer specific climate questions relating to that process. Because of these critical features, Hector has the potential to be a key analytical tool in both the policy and scientific communities. We welcome user input and encourage use, modifications, and collaborations with Hector.

While Hector has many strengths, there are a few limitations that later versions of Hector hope to address. For example, Hector does not have differential radiative forcing and atmospheric temperature calculations over land and ocean. The land responds to changes in emissions of greenhouse gases, and aerosols much quicker than the ocean, leading to different temperature responses over the land and ocean. Also, Hector does not explicitly deal with oceanic heat uptake. Surface temperatures are calculated based on a linear relationship with atmospheric temperature and heat uptake by the ocean is parameterized by a constant heat uptake efficiency. While Hector can reproduce global trends in atmospheric CO<sub>2</sub>, and temperature, we cannot investigate ocean heat uptake in the deep ocean using Hector. Currently, there is placeholder in Hector for a more sophisticated sea-level rise submodel. The current edition of Hector uses inputs of concentrations of CH<sub>4</sub> and N<sub>2</sub>O to calculate radiative forcing from CH<sub>4</sub> and N<sub>2</sub>O. Ideally we would like Hector to calculate concentrations from emissions of CH<sub>4</sub> and N<sub>2</sub>O. This would allow for quick integration within IAMs.

Future plans with Hector include addressing some of the above limitations and conducting numerous scientific experiments, using Hector as a stand-alone simple climate carbon-cycle model. Also, Hector will be incorporated into Pacific Northwest National

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Laboratory's Global Change Assessment Model to begin running policy relevant experiments. Hector has the ability to be a key analytical tool used across many scientific and policy communities due to its modern software architecture, open source, and object-oriented structure.

## 5 Code availability

Hector is freely available at <https://github.com/JGCRI/hector>. The specific Hector v0.1 referenced in this paper is available at <https://github.com/JGCRI/hector/releases/tag/v0.1>

*Author contributions.* C. A. Hartin and B. P. Bond-Lamberty developed the ocean and terrestrial carbon models, respectively, and led the overall development of Hector. R. P. Link and P. Patel wrote critical code for Hector's coupler and carbon cycle solver. A. Schwarber helped with the development of the atmospheric forcing components. C. A. Hartin wrote the manuscript with contributions from all co-authors.

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**Table 1.** Initial model conditions prior to spinup, assuming a pre-industrial steady state.

Variable	Description	Initial value	Units
$C_{\text{atm}}^*$	Atmospheric Carbon	588.1	PgC
$C_{\text{D}}^*$	Detritus Carbon	55	PgC
$C_{\text{S}}^*$	Soil Carbon	1782	PgC
$C_{\text{V}}^*$	Vegetation Carbon	550	PgC
$C_{\text{DO}}$	Deep Ocean	26 000	PgC
$C_{\text{HL}}$	Surface Ocean High Latitude	140	PgC
$C_{\text{IO}}$	Intermediate Ocean	8400	PgC
$C_{\text{LL}}$	Surface Ocean Low Latitude	770	PgC
$F_{\text{L}}$	Atmosphere–Land Carbon Flux	0	PgC yr <sup>-1</sup>
$F_{\text{O}}$	Atmosphere–Ocean Carbon Flux	0	PgC yr <sup>-1</sup>
$\text{NPP}_0$	Net Primary Production	50	PgC yr <sup>-1</sup>
$T_{\text{G}}$	Global Temperature Anomaly	0	°C
$T_{\text{HL}}$	Temperature of high latitude surface ocean box	2	°C
$T_{\text{LL}}$	Temperature of low latitude surface ocean box	22	°C

\* Parameters appearing in the input file.

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**Table 2.** Model parameters for the land and ocean carbon components.

Variable	Description	Value
$f_{ds}$	annual fraction of detritus carbon that is transferred to soil	0.60
$f_{ld}^*$	annual fraction of land use change flux from detritus	0.01
$f_{ls}$	annual fraction of land use change flux from soil	0.89
$f_{lv}^*$	annual fraction of land use change flux from vegetation	0.10
$f_{nd}^*$	annual fraction of NPP carbon that is transferred to detritus	0.60
$f_{ns}$	annual fraction of NPP carbon that is transferred to soil	0.05
$f_{nv}^*$	annual fraction of NPP carbon that is transferred to vegetation	0.35
$f_{rd}$	annual fraction of respiration carbon that is transferred to detritus	0.25
$f_{rs}$	annual fraction of respiration carbon that is transferred to soil	0.02
$f_{vd}$	annual fraction of vegetation carbon that is transferred to detritus	0.034
$f_{vs}$	annual fraction of vegetation carbon that is transferred to soil	0.001
$\beta^*$	Beta	0.36
$Q10^*$	$Q10$ respiration	2.45
$T_H^*$	High-latitude circulation	$4.9e7 \text{ m}^3 \text{ s}^{-1}$
$T_T^*$	Thermohaline circulation	$7.2e7 \text{ m}^3 \text{ s}^{-1}$
$E_{ID}^*$	Water mass exchange – intermediate – deep	$1.25e7 \text{ m}^3 \text{ s}^{-1}$
$E_{LI}^*$	Water mass exchange – low latitude – intermediate	$2.0e8 \text{ m}^3 \text{ s}^{-1}$

\* Parameters appearing in the input file.



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**Table 4a.** Skill of Hector vs. observations, CMIP5, and MAGICC, correlation coefficients ( $R$ ) and root mean square error (RMSE) for atmospheric  $[\text{CO}_2]$ , surface temperature anomaly, radiative forcing, fluxes of carbon (ocean and land), and low latitude surface ocean pH.

Variable	Skill	Historical 1850–2004			Units
		observations	MAGICC	CMIP5	
$[\text{CO}_2]^*$	$R$	0.99	0.99	1.0	ppmv
	RMSE	2.73	2.84	2.11	
temperature	$R$	0.85	0.85	0.81	deg C
	RMSE	0.11	0.11	0.13	
Forcing	$R$	–	0.79	–	$\text{W m}^{-2}$
	RMSE	–	0.36	–	
Ocean Flux	$R$	–	–	0.95	$\text{Pg Cyr}^{-1}$
	RMSE	–	–	0.27	
Land Flux	$R$	–	–	0.55	$\text{Pg Cyr}^{-1}$
	RMSE	–	–	1.30	
pH	$R$	–	–	0.99	unitless
	RMSE	–	–	0.003	

\*  $[\text{CO}_2]$  observations are an average of Law Dome and Mauna Loa.

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**Table 4b.** Continued.

RCP 8.5 1850–2300				
Variable	Skill	MAGICC	CMIP5	Units
[CO <sub>2</sub> ]	<i>R</i>	1.0	0.99	ppmv
	RMSE	7.69	10.62	
temperature	<i>R</i>	1.0	0.98	°C
	RMSE	0.33	0.91	
Forcing	<i>R</i>	1.0	–	W m <sup>-2</sup>
	RMSE	0.25	–	
Ocean Flux	<i>R</i>		0.80	Pg Cyr <sup>-1</sup>
	RMSE		1.45	
Land Flux	<i>R</i>	–	0.60	Pg Cyr <sup>-1</sup>
	RMSE	–	4.01	
pH	<i>R</i>	–	1	unitless
	RMSE	–	0.004	

\* CMIP5 [CO<sub>2</sub>] only to 2100.

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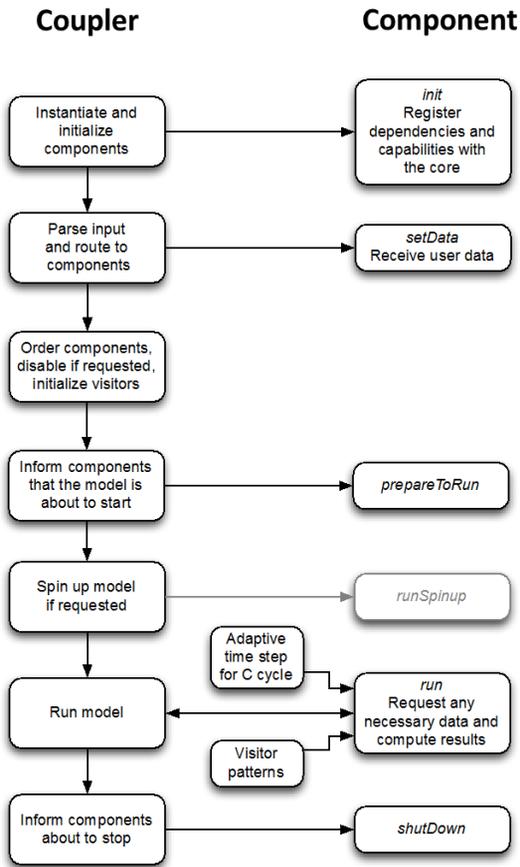
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**Table 4c.** Continued.

RCP 8.5 2005–2300				
Variable	Skill	MAGICC	CMIP5	Units
[CO <sub>2</sub> ]*	<i>R</i>	1.0	1.0	ppmv
	RMSE	8.64	10.23	
temperature	<i>R</i>	1.0	0.97	°C
	RMSE	0.17	0.86	
Forcing	<i>R</i>	1.0	–	W m <sup>-2</sup>
	RMSE	0.04	–	
Ocean Flux	<i>R</i>	–	0.10	Pg Cyr <sup>-1</sup>
	RMSE	–	1.42	
Land Flux	<i>R</i>	–	0.65	Pg Cyr <sup>-1</sup>
	RMSE	–	4.73	
pH	<i>R</i>	–	1.0	unitless
	RMSE	–	0.003	

\* CMIP5 [CO<sub>2</sub>] only to 2100.



**Figure 1.** Model phases for the coupler (left) and a typical component (right). Arrows show flow of control and data. The greyed spinup step is optional.

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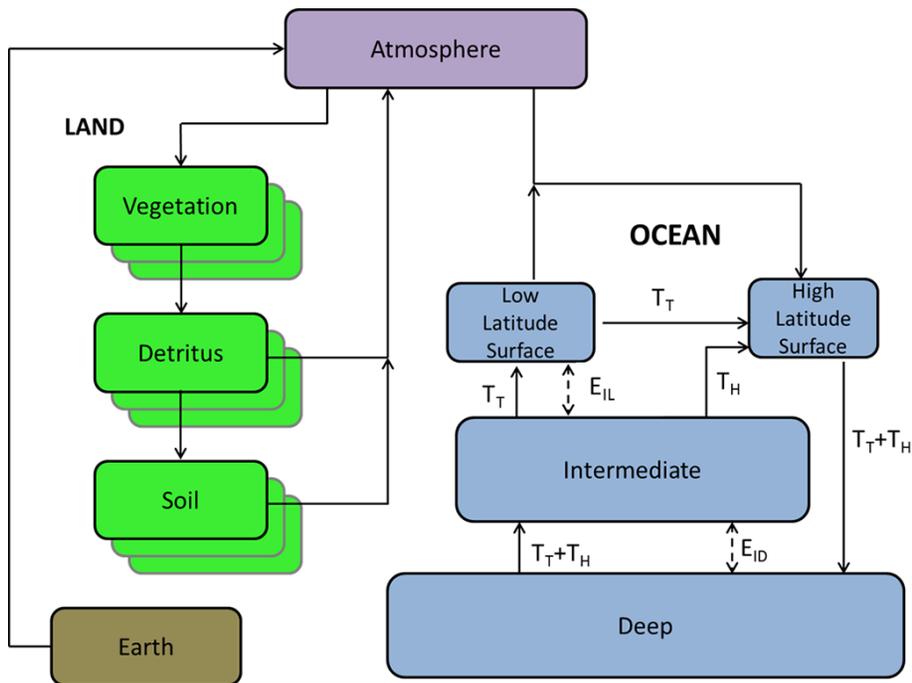
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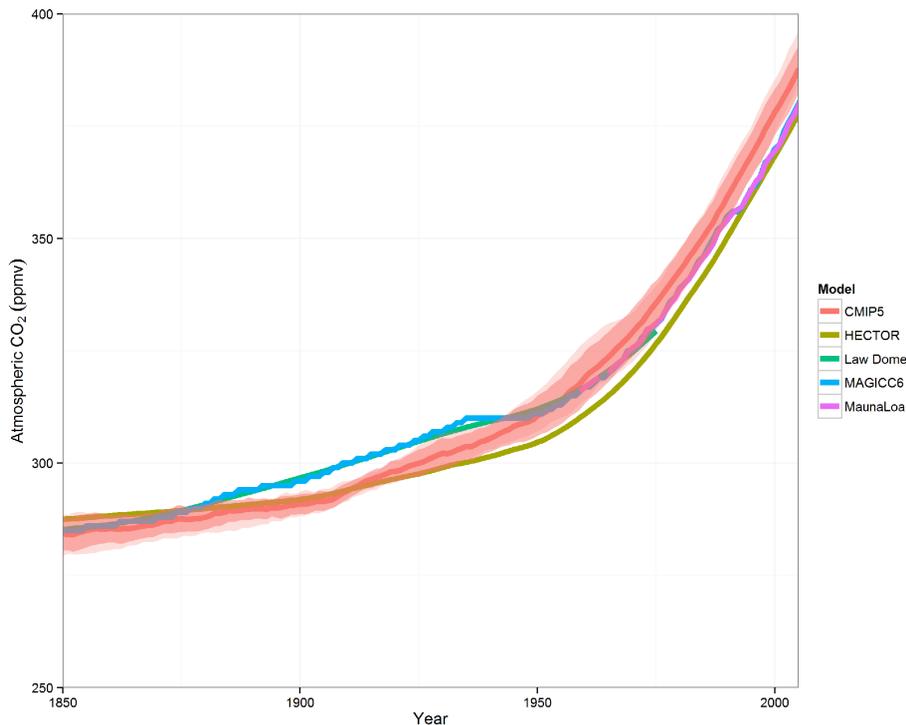




**Figure 2.** Representation of Hector's carbon cycle, land, atmosphere, and ocean. The atmosphere consists of one well mixed box. The ocean consists of four boxes, with advection and water mass exchange simulating thermohaline circulation. At steady state, the high latitude surface ocean takes up carbon from the atmosphere, while the low latitude surface ocean off gasses carbon to the atmosphere. The land consists of a user defined number of biomes or regions for vegetation, detritus and soil. At steady state the vegetation takes up carbon from the atmosphere while the detritus and soil release carbon back into the atmosphere. The earth pool is continually debited with each time step to act as a mass balance check on the carbon system.

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**Figure 3.** Historical atmospheric [CO<sub>2</sub>] from 1850 to 2005 for Hector (brown), CMIP5 (pink), MAGICC6 (blue), Law Dome (green), and Mauna Loa (purple). Note CMIP5 data are from the emissions driven historical scenario (esmHistorical). Notice that MAGICC6 matches the observational record. We have the capabilities of running Hector under numerous constraints. Within this study we are running Hector unconstrained to highlight the full performance of the model.

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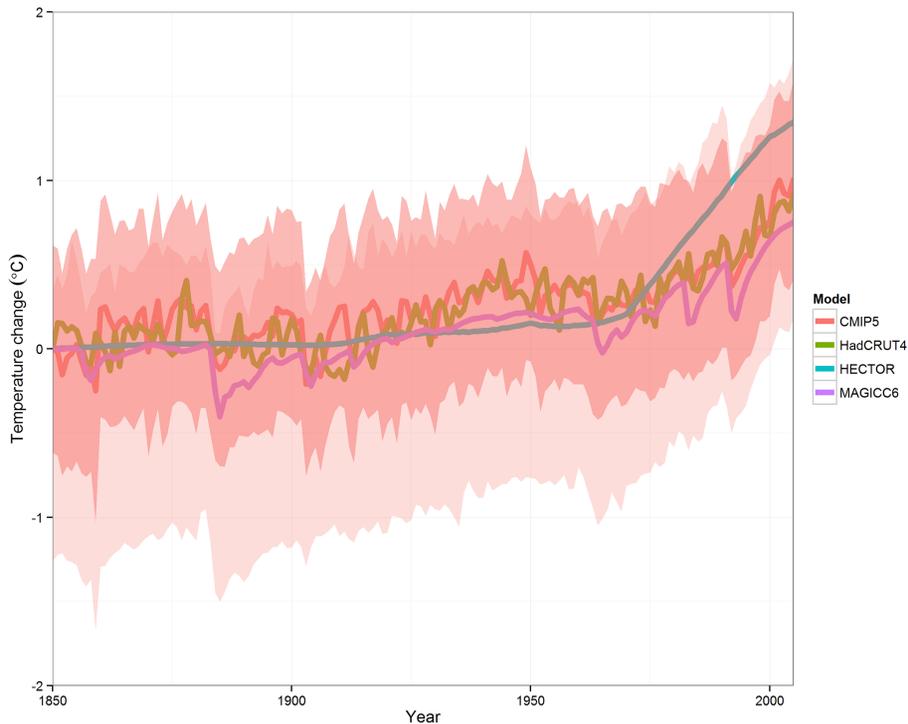
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**Figure 4.** Historical global temperature anomaly relative to 1850 for Hector (blue), MAGICC6 (purple), CMIP5 median, SD and model spread (pink), and historical observations from HadCRUT4 (green).

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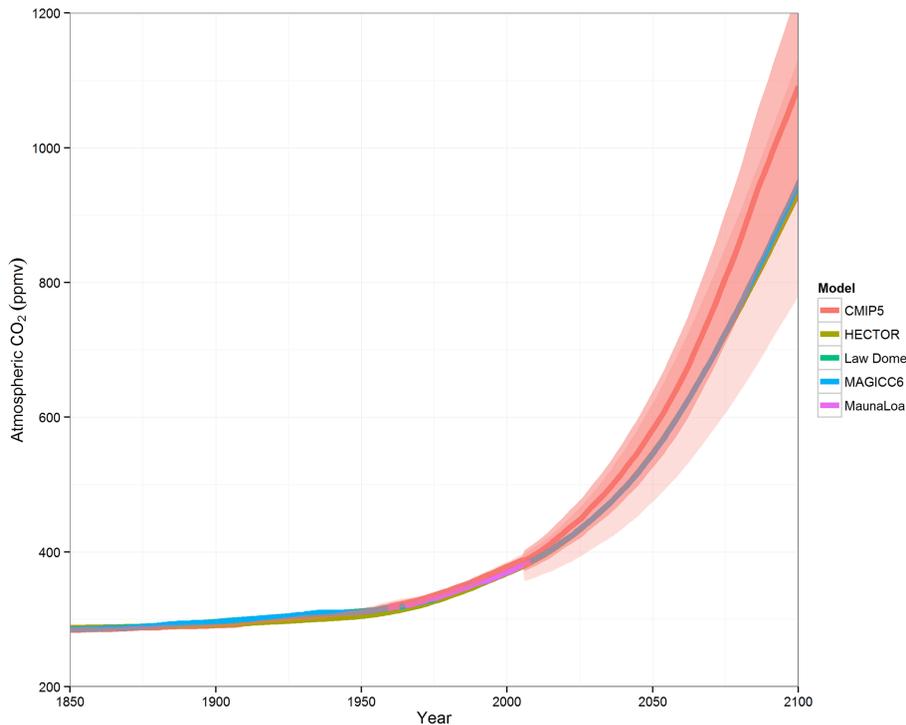
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**Figure 5.** Atmospheric  $[CO_2]$  from 1850 to 2100 under RCP 8.5 for Hector (yellow), MAGICC6 (blue), Mauna Loa (purple), Law Dome (green) and esmRCP 8.5 CMIP5 median, one SD and model spread (pink).

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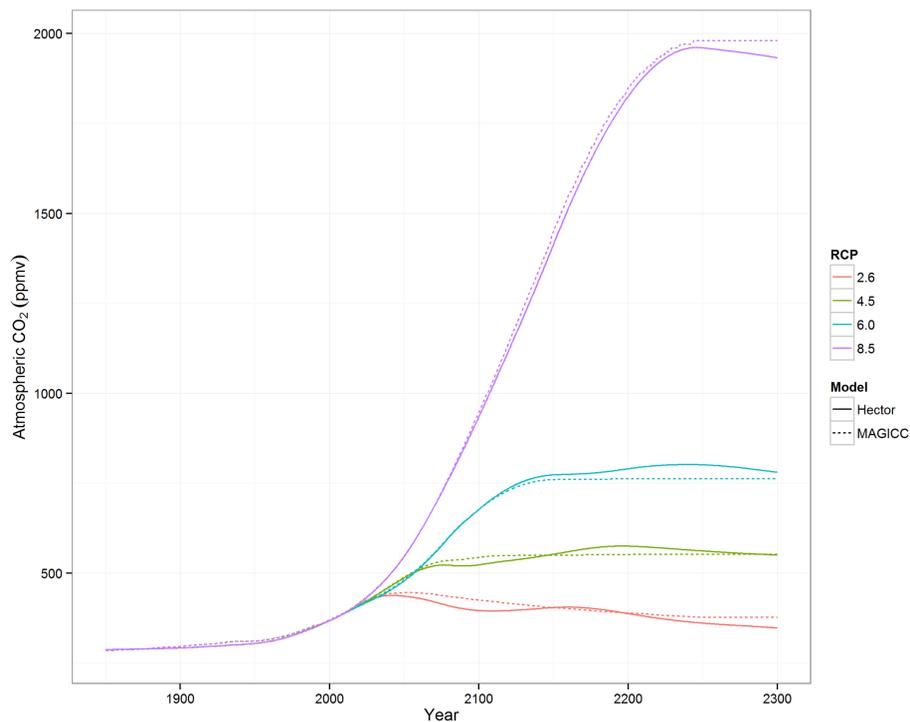
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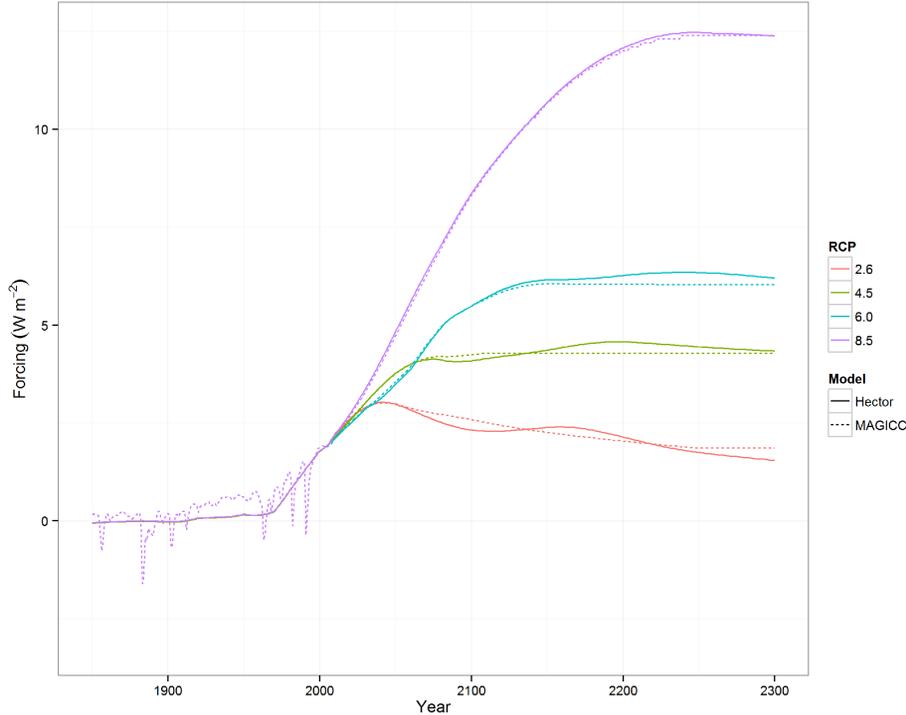
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**Figure 6.** Atmospheric [CO<sub>2</sub>] from 1850 to 2300 for RCP 2.6 (yellow), RCP 4.5 (green), RCP 6.0 (blue), RCP 8.5 (purple), Hector (solid) and MAGICC6 (dashed).

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**Figure 7.** Relative radiative forcing from 1850 to 2300 for Hector (solid) and MAGICC6 (dashed) for all four RCP scenarios, 2.6 (red), 4.5 (green), 6.0 (blue), 8.5 (purple). Hector has the option to enable or disable radiative forcing from historical volcanic emissions. We have opted to disable this for ease of comparison across all RCPs.

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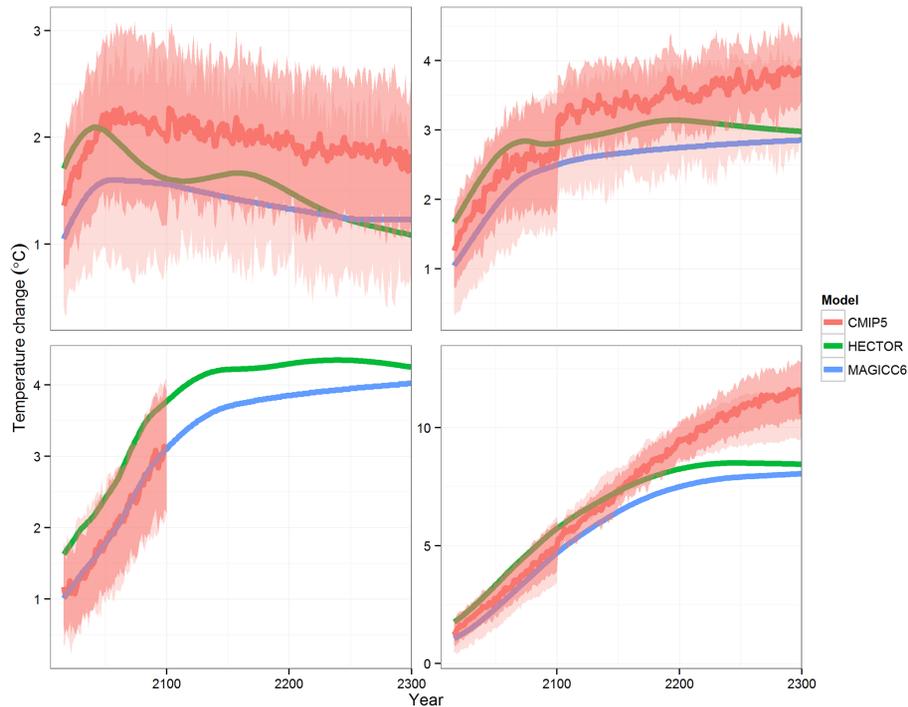
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**Figure 8.** Global temperature anomaly relative to 1850 for **(a)** RCP 2.6, **(b)** RCP 4.5, **(c)** RCP 6.0, and **(d)** RCP 8.5, comparing Hector (green), MAGICC6 (blue), and CMIP5 median, SD and model spread (pink). The CMIP5 models under RCP 6.0 used in this study do not extend to 2300. Note the change in scales between the four panels.

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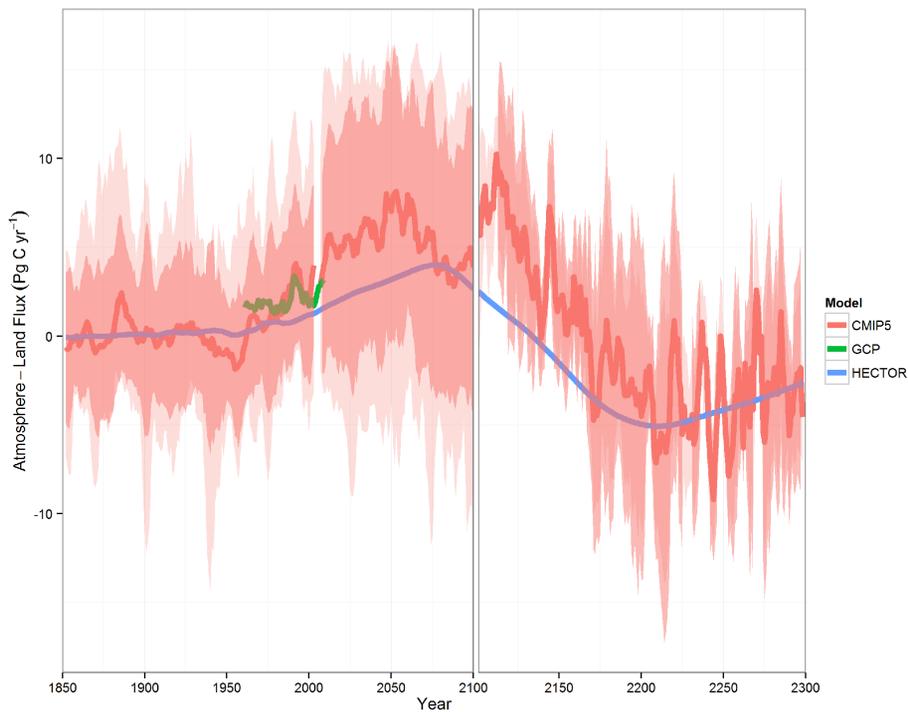
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**Figure 11.** Global air-land fluxes of carbon under RCP 8.5, Hector (blue), CMIP5 (red), and observations from GCP (green) (Le Quéré et al., 2013). The break in the graph at 2100 signifies a change in the number of models that ran the RCP 8.5 extension.

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