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Key Points:

- Global hydrological responses to forcings from rising CO₂-induced physiological and radiative effects were decoupled from CMIP6 models' simulations with 1% per year CO₂ increase
- Global hydrological sensitivity to CO₂ physiological forcing is $-0.09 \pm 0.07\%$ (100 ppm)⁻¹ and to CO₂ radiative forcing is $1.54 \pm 0.24\%$ K⁻¹ from transient 4 × CO₂ simulations
- Global hydrological sensitivities to CO₂ physiological and radiative forcings have little changes over transient 2 × CO₂ to 4 × CO₂ scenario

Supporting Information:

Supporting Information may be found in the online version of this article.

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Quantifying Global Hydrological Sensitivity to CO₂ Physiological and Radiative Forcings Under Large CO₂ Increases

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Abstract Prediction of surface freshwater flux (precipitation or evaporation) in a CO₂-enriched climate is highly uncertain, primarily depending on the hydrological responses to physiological and radiative forcings of CO₂ increase. Using the 1pctCO₂ (a 1% per year CO₂ increase scenario) experiments of 12 CMIP6 models, we first decouple and quantify the magnitude of global hydrological sensitivity to CO₂ physiological and radiative forcings. Results show that the direct global hydrological sensitivity (for land plus ocean precipitation) to CO₂ increase only is $-0.09 \pm 0.07\%$ (100 ppm)⁻¹ and to CO₂-induced warming alone is $1.54 \pm 0.24\%$ K⁻¹. The latter is about 10% larger than the global apparent hydrological sensitivity (i.e., including all effects, not only direct responses to warming, $\eta_a = 1.39 \pm 0.22\%$ K⁻¹). These hydrological sensitivities are relatively stable over transient 2× to 4 × CO₂ scenario. The intensification of the global water cycle are dominated by the CO₂ radiative effect ($79 \pm 12\%$) with a smaller positive contribution from the interaction between the two effects ($6 \pm 12\%$), but are reduced by the CO₂ physiological effect ($-10 \pm 8\%$). This finding underlines the importance of CO₂ vegetation physiology in global water cycle projections under a CO₂-enriched and warming climate.

Plain Language Summary Projections of future water resource under rising CO₂ and warming scenarios are highly uncertain, which is mainly due to the poor understanding of hydrological responses within the Earth system. We evaluate two major hydrological responses associated with rising CO₂ related effects: physiological effect (e.g., decrease in plant stomatal conductance per unit leaf area, increase in leaf area index and net available energy, etc.) and radiative effect (e.g., rising CO₂-induced warming, rapid atmospheric adjustment and surface energy budget change, etc.). Using 12 Earth system models' experiments, we find that the CO₂ physiological effect will cause ~0.1% decline for precipitation or evaporation per 100 ppm CO₂ increase at a global scale. The CO₂ radiative effect will cause ~1.5% increase for precipitation or evaporation per Kelvin warming. These changing rates of increase or decline are relatively stable with increasing CO₂ forcing. The modeling experiments under high CO₂ forcing project that the intensification of the global water cycle is mainly controlled by the CO₂ radiative effect, but the CO₂ physiological effect and the interaction between the two effects also play important roles.

1. Introduction

The global water cycle is predicted to intensify in the future, owing to rising atmospheric CO₂ concentration and related surface warming driven by anthropogenic greenhouse gases emissions (Allen & Ingram, 2002; Douville et al., 2023; Huntington, 2006; Wentz et al., 2007). This intensification is generally expressed as increases in surface hydrological fluxes (e.g., precipitation, evapotranspiration, and runoff) with an increase in spatial variability of surface water budget (e.g., precipitation minus evapotranspiration) (Allan, 2023; Zaitchik et al., 2023). Although historical long-term response of precipitation and evapotranspiration to global warming is still ambiguous, recent studies provide some evidence to support an intensifying water cycle, for example, increased trends in extreme river flow (Gudmundsson et al., 2021), observed global surface salinity changes (Durack et al., 2012), and observed increases of global oceanic precipitation and evaporation (Brutsaert, 2017; Wang et al., 2021; Wentz et al., 2007). Climate models also project strong increases in surface hydrological fluxes, for example, global lake evaporation (Wang et al., 2018), global ocean evaporation (Wang et al., 2021), global runoff

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(Alkama et al., 2013; Betts et al., 2007; X. J. Zhang et al., 2014), and global precipitation (Shiogama et al., 2022; Wang et al., 2021) under different atmospheric CO₂ increasing scenarios.

Increased atmospheric CO₂ induces infrared radiative forcing on atmosphere-to-surface energy balance (or “an enhanced greenhouse effect”) which further warms the land and ocean surfaces (Myhre et al., 1998, 2013) and alters hydrological processes around most of the world (Cisneros et al., 2014). This “enhanced greenhouse effect” by rising CO₂ is termed as rising CO₂-induced radiative effect in this study. It induces hydrological responses across all timescales including rapid responses from atmospheric adjustments and slow responses with rising CO₂-induced warming and transient climate change (Allan et al., 2020; Douville et al., 2023). One of the hydrological responses is an increase in lower-tropospheric water vapor with a rate of ~7% per Kelvin warming, driven by increased atmospheric saturation vapor pressure (e_s) and enhanced surface evaporation in response to global warming, as predicted by the Clausius-Clapeyron equation and early global climate models (e.g., Boer, 1993; Held & Soden, 2006). The increased atmospheric water vapor consequently drives an increase of global-mean precipitation with a much smaller growth rate of 1%–3% per Kelvin warming (Allen & Ingram, 2002; Wentz et al., 2007). This smaller rate of increase in precipitation is explained by the suppression from rapid atmospheric adjustments in response to CO₂ radiative effect and cooling effects from scattering aerosols that directly alter the atmospheric energy budget and reduce convective mass fluxes (Allan et al., 2020; Andrews & Forster, 2010; Held & Soden, 2006; Li et al., 2013).

Besides the radiative effect, the increased CO₂ also induces vegetation physiological responses including decrease in stomatal conductance per unit leaf area, and increases in leaf area index (LAI), vegetational cover and growing seasons, and enhanced net available energy at the surface through reduced surface albedo etc. (Guerrieri et al., 2019; Piao, Liu, et al., 2019; Piao, Wang, et al., 2019; X. Z. Zhang et al., 2022). The most significant hydrological response is a decrease in plant transpiration and land evapotranspiration determined by reduced plant stomatal conductance, which further induces regional and large-scale atmospheric vapor transport adjustments and slightly alter global precipitation (e.g., Bounoua et al., 1999; Sellers et al., 1996). This response is also predicted to increase continental runoff (Betts et al., 2007).

The radiative and vegetation physiological effects of rising CO₂ on climate have been proposed and quantified using modeling simulations for an atmospheric CO₂ doubling (abrupt 2 × CO₂) scenario (Sellers et al., 1996). Over the last two decades, climate models or earth system models with fully coupled carbon cycle were used to quantify the rising CO₂-induced radiative and physiological effects on global warming, global carbon cycle and carbon-climate feedback (e.g., Betts et al., 1997; Friedlingstein et al., 2003, 2006; Woodward et al., 1998; X. Zhang, Wang, et al., 2021; X. Z. Zhang, Wang, & Zhang, 2023).

Responses of surface hydrological processes to the forcing of rising CO₂-induced radiative effect (termed as “CO₂ radiative forcing”) or rising CO₂-induced physiological effect (termed as “CO₂ physiological forcing”) also have been studied before (Andrews et al., 2010; Betts et al., 2007; Bonfils et al., 2017; Cao et al., 2010; Richardson et al., 2018; Saint-Lu et al., 2020). For instance, Betts et al. (2007) used an ensemble of experiments with a global climate model (HadSM3) to show that physiological effect of a doubling CO₂ on plant transpiration could increase global mean runoff by 6%. Cao et al. (2010) used another coupled climate model (CCSM3 coupled with CLM3.5/CAM3.5) and found that doubled CO₂ physiological forcing will increase global runoff by ~8.4% and the forcing from combined radiative and physiological effects resulted in ~7.35% increase in land precipitation and ~14.9% increase in runoff. Multi-modeling experiments further emphasized the importance of rising CO₂ physiological effect on global land monsoon rainfall and runoff in a CO₂-enriched climate (Cui et al., 2020).

However, in most previous studies, the hydrological sensitivities of surface hydrological fluxes (e.g., precipitation or evaporation) due to the rising CO₂ radiative and physiological forcings have not been well examined in terms of magnitude and pattern using the state-of-the-art Earth system models. Particularly, the potential interactions between the CO₂ radiative forcing and physiological forcing were ignored in previous studies. The relative roles of the CO₂-radiative and physiological forcings and their interaction remain largely unclear on globe, land or ocean. In this study, we develop a hydrological sensitivity analysis framework to quantify the impacts of CO₂ radiative and physiological forcings and their interaction on global water cycle, using fully or partially coupled simulations by 12 CMIP6 models under a 1% per year increasing atmospheric CO₂ (1pctCO₂) for 140 years (Eyring et al., 2016). With the analysis framework, the strength of the hydrological responses to CO₂ radiative forcing (Friedlingstein et al., 2003; Hansen et al., 1984) is termed as the sensitivity of change in hydrological flux variables (ΔX) to increase in surface air temperature (ΔT). The other is the hydrological responses to the forcing

of surface vegetation, energy and climate changes due to rising CO₂-induced physiological effect. The strength of this hydrological response to CO₂ physiological forcing is termed as the sensitivity of change in hydrological flux variables (ΔX) to increase in atmospheric CO₂ concentration (ΔC).

2. Methods

2.1. Analysis Framework of Rising CO₂-Induced Responses of Global Water Cycle in Earth System

Following the previous frameworks developed for quantifying global carbon cycle—climate feedback (Arora et al., 2013; X. Zhang, Wang, et al., 2021), we develop a similar analysis framework of quantifying global water cycle responding to increasing CO₂ levels by linearly combining the hydrological responses to rising CO₂-induced physiological and radiative forcings and the two forcings' interaction. Potential interactions between the two forcings in the fully coupled Earth system are also considered, which could be very significant at some specific regions. For instance, a potential interaction is that rising CO₂-induced plant growth and LAI increase from CO₂ vegetation physiological effect may further enhance plant transpiration under the surface warming from CO₂-induced radiative forcing (X. Z. Zhang, Zhang, et al., 2021; X. Z. Zhang, Wang, & Zhang, 2023). This effect may increase evapotranspiration, reduce runoff, and alter local precipitation, but how much the interaction affecting hydrological responses is still unknown. Hence, in linear analysis framework of a fully coupled Earth system, change in a hydrological flux variable (ΔX) during a time interval (Δt) since a beginning year (e.g., 1850) can be expressed by

$$\Delta X = B\Delta C + \Gamma\Delta T + \varepsilon, \quad (1)$$

where X is a hydrological flux variable from the surface perspective, which in this study is annual precipitation (P), evapotranspiration on land or evaporation on ocean (E), or surface water availability ($P - E$) on land, ocean, or global scales. C is annual atmospheric CO₂ concentration and T is annual global-averaged surface air temperature. ε is the interaction contribution between CO₂ physiological and radiative forcings. $B \equiv \frac{\partial X}{\partial C}$ is the sensitivity parameter due to CO₂ physiological effect (in a unit of mm yr⁻¹ (100 ppm)⁻¹), and $\Gamma \equiv \frac{\partial X}{\partial T}$ is the sensitivity parameter due to CO₂ radiative effect (in a unit of mm yr⁻¹ K⁻¹). Owing to the different units used for B and Γ , the sensitivity to CO₂ physiological effect and the sensitivity to CO₂ radiative effect are not comparable in this study. Therefore, the hydrological sensitivities to CO₂ concentration only for P , E , and $P - E$ are $B_P = \frac{\partial P}{\partial C}$, $B_E = \frac{\partial E}{\partial C}$, $B_{P-E} = \frac{\partial(P-E)}{\partial C}$, respectively. The hydrological sensitivities to warming alone for P , E , and $P - E$ are $\Gamma_P = \frac{\partial P}{\partial T}$, $\Gamma_E = \frac{\partial E}{\partial T}$, $\Gamma_{P-E} = \frac{\partial(P-E)}{\partial T}$, respectively.

2.2. Calculation of Hydrological Sensitivities From Decoupling Simulations of the CMIP6 1pctCO₂ Experiments

The strength and relative contributions of hydrological responses from the CO₂ physiological forcing and radiative forcing could not be directly estimated, as they interact within the Earth's coupling system of climate, water cycle, and carbon cycle. To date, Earth system models are almost the only effective approach to separate different forcings' contribution to the overall global water cycle responses. To quantify the magnitudes of hydrological sensitivity parameters (i.e., B and Γ) from the global water cycle under rising CO₂-induced forcings, the 1pctCO₂ experiments from the CMIP6 project are ideal decoupling modeling experiments, which provide the fully coupled (COU), the biogeochemically coupled (BGC), and the radiatively coupled (RAD) simulations (Jones et al., 2016). The COU simulations of the “1pctCO₂” configuration include all water cycle and carbon cycle—climate feedbacks driven by transient atmospheric CO₂ increased at a rate of 1% year⁻¹ over a 140-year period. By the end of the 140-year simulation, atmospheric CO₂ concentration is quadrupled ($4 \times \text{CO}_2$) from the pre-industrial level (~ 285 ppm at 1850). The BGC simulations of the “1pctCO₂-bgc” configuration are also driven by the prescribed 1% year⁻¹ CO₂ forcing, but only the biogeochemical cycle is turned on, meaning that only the CO₂ physiological effect on global water cycle is considered. The RAD simulations of the “1pctCO₂-rad” configuration are driven by the prescribed 1% year⁻¹ CO₂ forcing with the CO₂ radiative forcing. Following the approaches for estimating global carbon cycle feedback parameters (Arora et al., 2013; Friedlingstein et al., 2006; X. Zhang, Wang, et al., 2021) and the calculation of apparent hydrological sensitivity (η_a) by Fläschner et al. (2016), the hydrological sensitivity parameter for CO₂ physiological forcing can be estimated by

$$B = \frac{\Delta X^{\text{BGC}} \Delta T^{\text{COU}} - \Delta X^{\text{COU}} \Delta T^{\text{BGC}}}{\Delta C^{\text{BGC}} (\Delta T^{\text{COU}} - \Delta T^{\text{BGC}})} \approx \frac{\Delta X^{\text{BGC}}}{\Delta C^{\text{BGC}}}, \quad (2)$$

and hydrological sensitivity parameter for CO₂ radiative forcing is estimated by

$$\Gamma = \frac{\Delta X^{\text{RAD}}}{\Delta T^{\text{RAD}}}, \quad (3)$$

where X is hydrological flux variable (P , E , and $P - E$). To quantify the contribution of interaction between CO₂ physiological and radiative forcings (the ε term in Equation 1), we calculated differences of the COU, BGC, RAD simulations by

$$\Delta X^{\text{INT}} = \Delta X^{\text{COU}} - \Delta X^{\text{BGC}} - \Delta X^{\text{RAD}}, \quad (4)$$

where X is temperature or hydrological flux variable (P , E , and $P - E$).

Simulations from 12 CMIP6 models of the 1pctCO₂ experiments are used in this study. These models include all required outputs of the three experiments (1pctCO₂, 1pctCO₂-rad, 1pctCO₂-bgc) and are available for download via <https://esgf-node.llnl.gov/search/cmip6/>. They are ACCESS-ESM1-5 (Ziehn et al., 2020), BCC-CSM2-MR (Wu et al., 2019), CESM2 (Danabasoglu et al., 2020), CMCC-ESM2 (Cherchi et al., 2018), CNRM-ESM2-1 (S  f  rian et al., 2019), GISS-E2-1-G (Ito et al., 2020), IPSL-CM6A-LR (Boucher et al., 2020), MIROC-ES2L (Hajima et al., 2020), MPI-ESM1-2-LR (Mauritsen et al., 2019), MRI-ESM2-0 (Yukimoto et al., 2019), NorESM2-LM (Seland et al., 2020), and UKESM1-0-LL (Sellar et al., 2019). We calculate all hydrological sensitivity parameters at transient 4 × CO₂ (i.e., 1% year^{−1} CO₂ increase up to 4 × CO₂) from monthly outputs of the COU, BGC, and RAD simulations of the 12 CMIP6 models, and perform comparative analysis on global, land and ocean scales across models.

3. Results

3.1. Responses of Temperature and Hydrological Fluxes to Rising CO₂-Induced Forcings

Figure 1 shows long-term changes of global mean surface air temperature (T_A) driven by increasing CO₂ concentration at the 1% year^{−1} for 140 years. On a global scale (including land and ocean), T_A is persistently increased to 4.62 ± 0.92 K (on ensemble mean ± standard deviation) at transient 4 × CO₂, as estimated from the fully coupled (COU) simulations of 12 CMIP6 models. The temperature growth curve indicates the non-linear dependence of radiative forcing upon the increasing CO₂ concentration from 1 × CO₂ to 4 × CO₂ (Myhre et al., 1998). Compared to this warming, global land surface warming is higher, with an estimate of 6.51 ± 1.29 K, and the global ocean surface has a smaller warming of 3.93 ± 0.82 K (Figures 1a and 1b). Decoupling experiments show that the CO₂ radiative forcing almost dominates the global warming in fully coupled simulations with ~96% contributions for globe and land and ~98% for ocean (Figures 1e and 1f). By comparison, the CO₂ physiological forcing (from the BGC simulations) induces only small warming of about 0.34 ± 0.28 K on land, and 0.12 ± 0.21 K on ocean at 4 × CO₂ (Figures 1c and 1d). This small warming is mainly contributed by decreased latent heat flux as a result of reduced vegetational stomatal conductance in response to CO₂ increase (Ainsworth & Long, 2005; Bounoua et al., 1999; Cao et al., 2010).

Global mean annual P and E (including land and ocean) increase at the same rate of 8.63 ± 2.89 mm year^{−1} per 100 ppm CO₂ or 15.53 ± 2.48 mm year^{−1} per Kelvin warming (see Figure S1 in Supporting Information S1 for the linear approximations), from the fully coupled simulations of CMIP6 models at the transient 4 × CO₂, due to the overall surface water balance on Earth (Figure 2). This result indicates the global apparent hydrological sensitivity (η_a) is $1.39 \pm 0.22\%$ per Kelvin warming, which includes all rising CO₂-induced effects, not just the effect of warming alone. However, the global land surface shows an increased $\Delta(P - E)$ along with CO₂ increase due to much higher increases in P compared to E on land, with $P - E$ increased at a rate of 8.14 ± 4.01 mm year^{−1} (100 ppm CO₂)^{−1}. Global ocean surface shows a declined $\Delta(P - E)$ along with CO₂ increase at a rate of -3.99 ± 2.00 mm year^{−1} (100 ppm CO₂)^{−1} (Figures 2a, 2d and 2g).

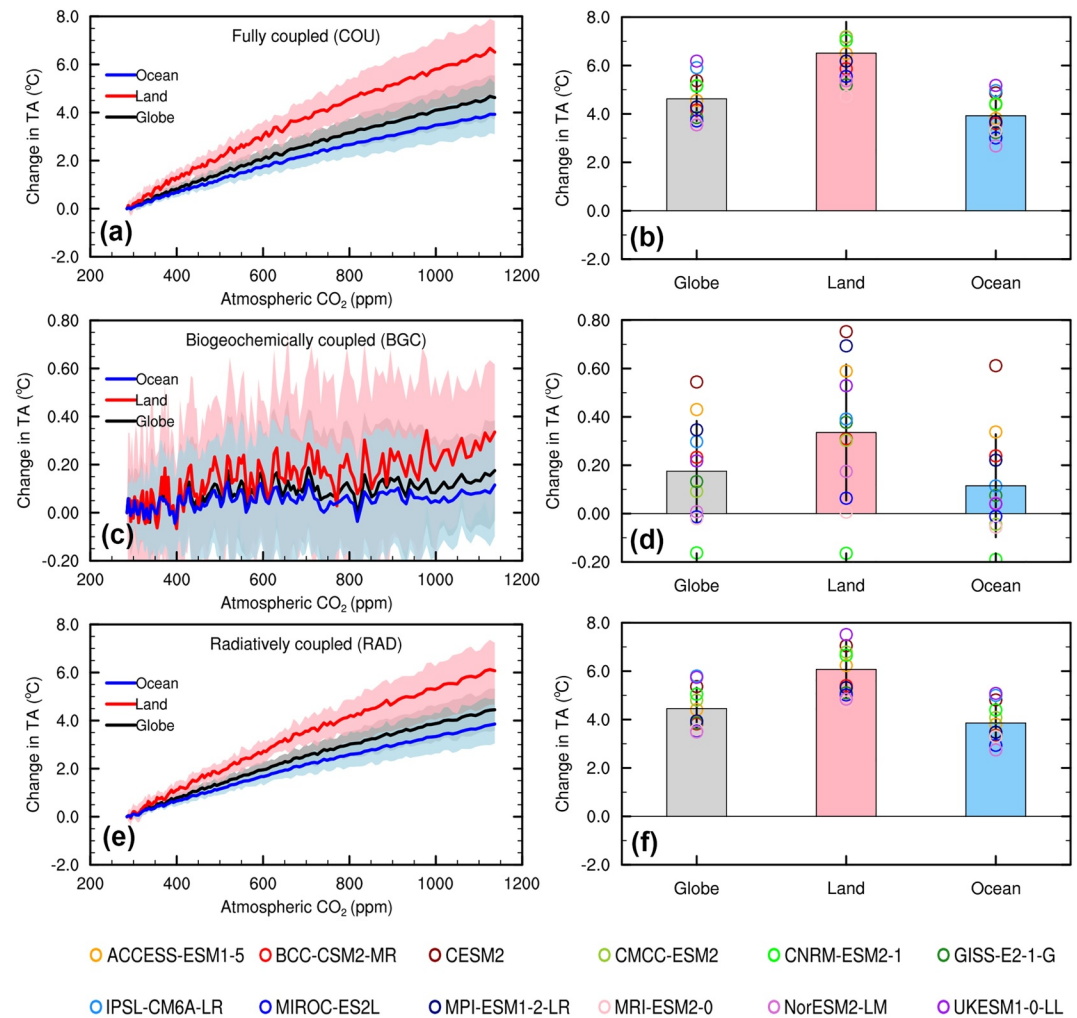


Figure 1. Changes in global annual T_A under transient 1%/year increasing CO_2 forcing scenario. Changes in surface air temperature (ΔT_A) were calculated on global, land and ocean scales, respectively, from (a and b) the fully coupled (COU) simulations, (c and d) the biogeochemically coupled (BGC) simulations, and (e and f) the radiatively coupled (RAD) simulations of the CMIP6 1pct CO_2 experiments. (a, c, e) Time-series of ensemble mean \pm standard deviation ΔT_A of 12 CMIP6 models with atmospheric CO_2 increasing from the preindustrial level of 285 ppm to its quadruple ($4 \times \text{CO}_2$) over a 140-year period. (b, d, f) CMIP6 modeled ΔT_A at $4 \times \text{CO}_2$ for the COU, BGC, RAD simulations, respectively.

Regionally, spatial patterns of the $\Delta(P - E)$ ensemble mean at transient $4 \times \text{CO}_2$ (Figure 3) show that the largest wetting land regions are tropical African (10°S – 10°N , 5°E – 30°E) and Asian rainforests (10°S – 10°N , 90°E – 140°E), South Asia (10°N – 30°N , 70°E – 100°E), eastern North America (30°N – 45°N , 80°W – 90°W) and Western North American area (50°N – 70°N , 140°W – 160°W). The largest drying land regions are Amazonian rainforests (10°S – 10°N , 50°W – 80°W), semi-arid regions over continental Europe (30°N – 55°N , 0°E – 30°E) and southern Africa (10°S – 30°S , 10°E – 35°E) (Figure 3g). Most regions with negative $\Delta(P - E)$ over oceans are latitude zones of -10°S to -40°S and 10°N to 40°N which covers the conventional oceans with low amount of multi-year mean annual precipitation (Figures 3d and 3g). However, the largest increased $\Delta(P - E)$ over oceans are over the intertropical convergence zone (ITCZ), Southern Ocean (-50°S to -70°S) and northern North Atlantic Ocean (Figure 3g). The strong increased $\Delta(P - E)$ over northern North Atlantic Ocean (40°N – 60°N , 20°W – 50°W) driven by a large reduction in E (Figures 3d and 3g) is probably due to the North Atlantic warming hole that is linked to a slowdown of the Atlantic meridional overturning circulation (Keil et al., 2020; Rahmstorf et al., 2015).

Decoupling experiments from the RAD and BGC simulations at $4 \times \text{CO}_2$ show that most drying/wetting regions are primarily due to the rising CO_2 -induced radiative forcing (Figures 3 and 5, and Figure S2 in Supporting

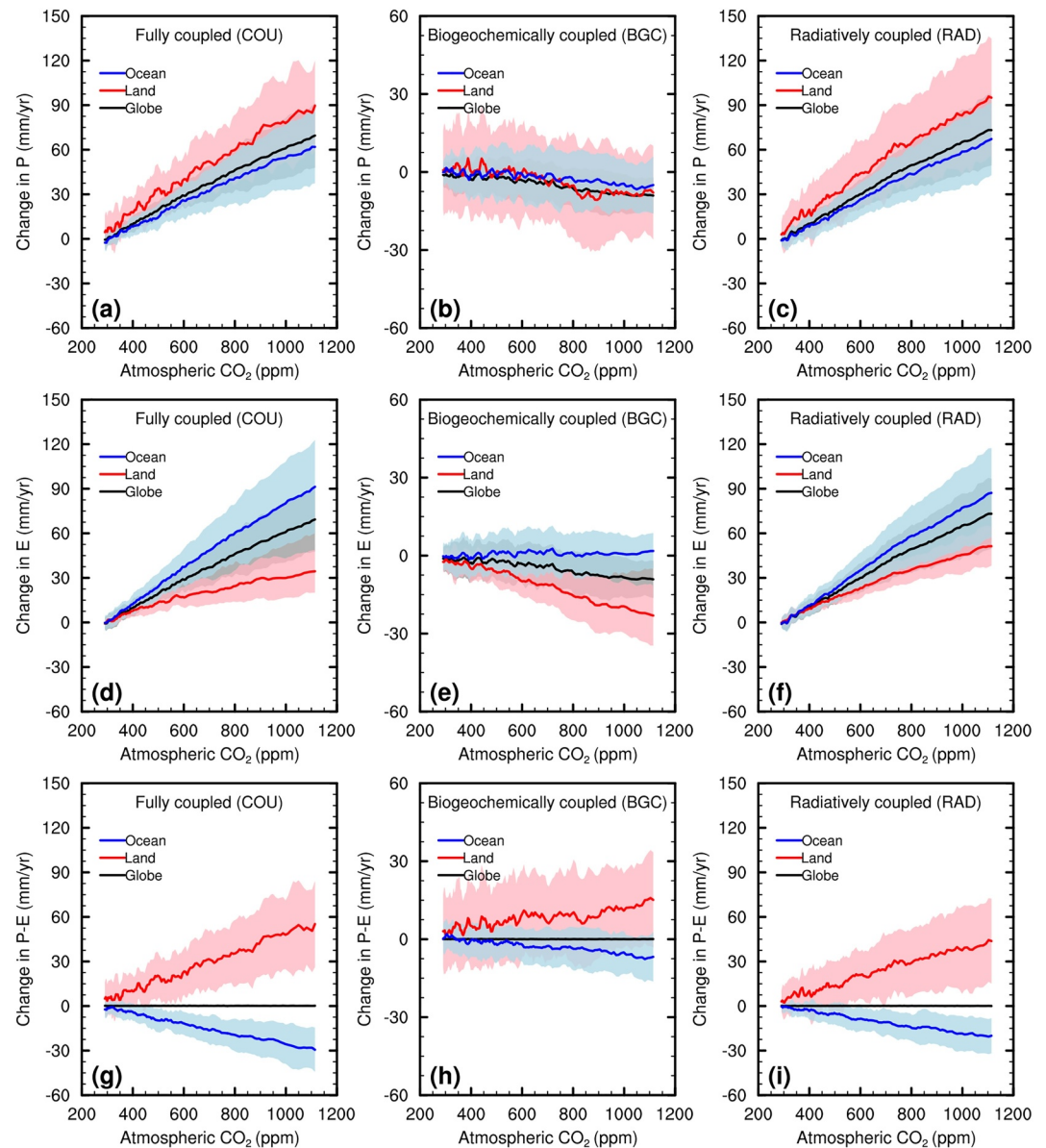


Figure 2. Time series of global annual P , E , and $P - E$ changes under transient 1%/year increasing CO_2 forcing scenario. (a) Changes in precipitation (ΔP) from the COU simulations, and (b) from the BGC simulations due to CO_2 physiological effect, and (c) from the RAD simulations due to CO_2 radiative effect. (d–f) Same as (a–c) but for changes in evaporation (ΔE). (g–i) Same as (a–c) but for differences between precipitation and evaporation ($\Delta(P - E)$). Shaded area represents mean \pm standard deviation from 12 CMIP6 models. All series were smoothed with a 5-year moving average.

Information S1) that induces the expansion of Hadley Cells (Grise & Davis, 2020; Schmidt & Grise, 2017) and alters the strength of tropical atmospheric circulations, cloud feedbacks and vapor transport (Collins et al., 2010; Vecchi et al., 2006) which largely shift the meridional distributions of P by increasing P over -10°S – 10°N around the equator and by reducing P over latitude zones of 10°S – 40°S and 10°N – 40°N (Figure 3). Interestingly, the positive $\Delta(P - E)$ of the northern North Atlantic Ocean (Figure 3g) is due to strong reduction of evaporation derived from the CO_2 radiative forcing, although there is smaller decrease in P over the northern North Atlantic Ocean (Figures 3a–3c). Notably, the CO_2 physiological forcing (BGC) consistently causes a decreased land evaporation by reduced plant transpiration from CO_2 stomatal effects as shown in Figure 2e, but causes a drying up of Amazonian rainforests (10°S – 10°N , 50°W – 80°W) and southern African semi-arid regions (10°S – 30°S ,

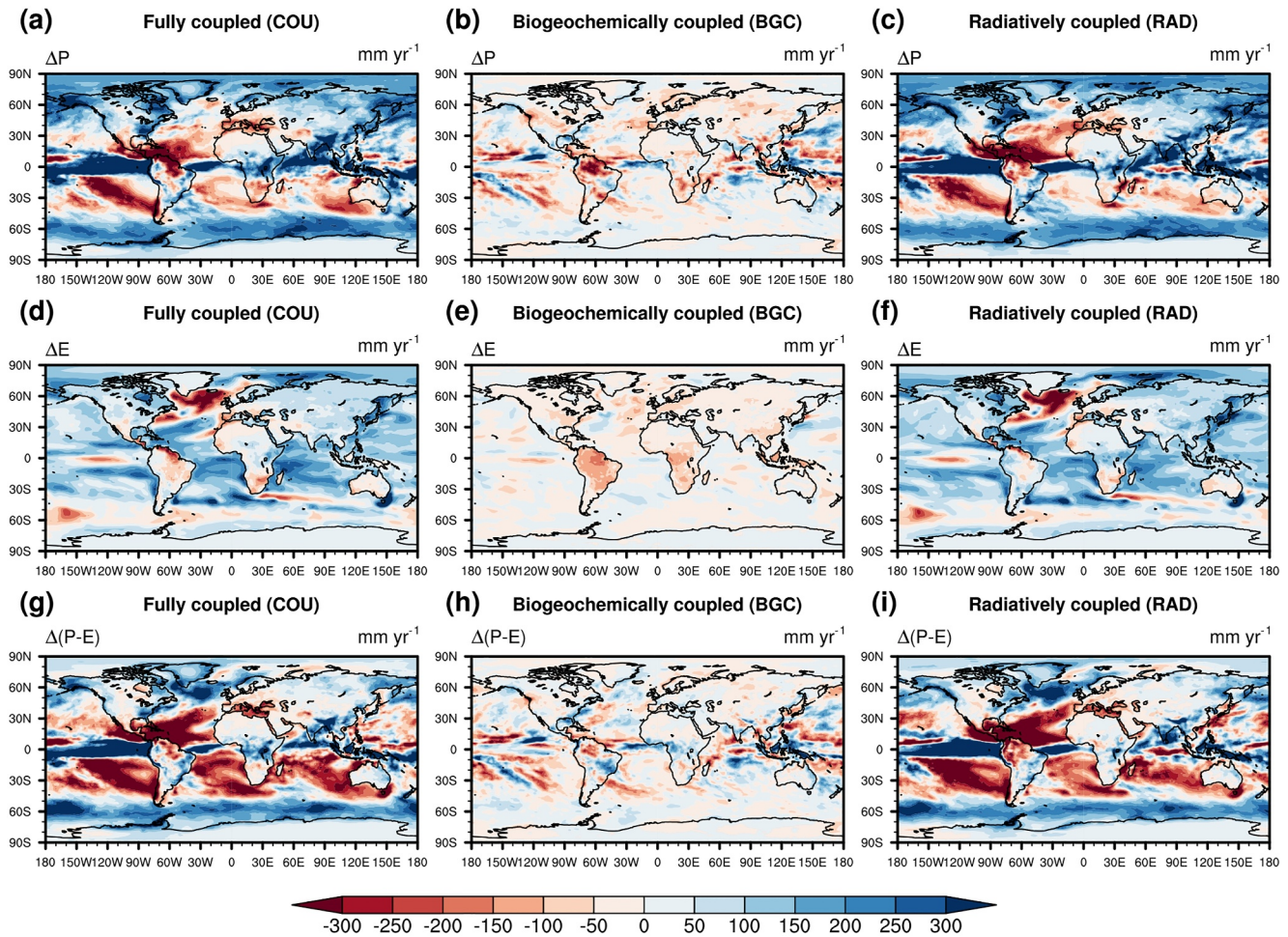


Figure 3. Spatial patterns of long-term changes in P , E , and $P - E$ at transient $4 \times \text{CO}_2$. (a) Changes in precipitation (ΔP) from the COU simulations, (b) from the BGC simulations due to CO_2 physiological effect, and (c) from the RAD simulations due to CO_2 radiative effect. (d–f) Same as (a–c) but for changes in evaporation (ΔE). (g–i) Same as (a–c) but for $\Delta(P - E)$. All results are averaged from 12 CMIP6 models of the 1pct CO_2 experiments.

10°E – 35°E), which is driven by a larger reduction of precipitation (Figures 3b, 3e, and 3h). Previous study suggested that the reduced precipitation due to CO_2 physiological forcing in Amazonian regions is almost entirely due to repartitioning of sensible and latent heat fluxes (Richardson et al., 2018).

The interaction between the CO_2 physiological and radiative forcings (Equation 4), as estimated by the ensemble of 12 CMIP6 models, indicates relatively small changes with atmospheric CO_2 forcing increase for P and E on globe, land and ocean (Figures 4a, 4c, and 4e). However, with CO_2 forcing increased to $4 \times \text{CO}_2$, the impacts of the interaction have comparable contribution as the CO_2 physiological forcing at regional scales (Figures 4 and 5, and Figure S1 in Supporting Information S1). Regionally, the interactions for P and $P - E$ are strong positive anomalies (200 – 300 mm yr^{-1}) in Amazonian rainforests and are strong negative anomalies ($< -300 \text{ mm yr}^{-1}$) over tropical western Pacific Ocean (5°S – 15°N , 120°E – 150°E). The northern (southern) areas of the ITCZ over tropical Indian Ocean (20°S – 10°N , 60°E – 110°E) and tropical Atlantic Ocean (10°S – 10°N , 120°E – 150°E) show strong positive (negative) anomalies of P and $P - E$ (Figures 4b, 4d and 4f). Overall, the interaction-induced drying or wetting effects (anomaly of $P - E$) over land and oceans are mainly driven by the changes of precipitation. This could be explained by that the regional large changes of precipitation triggered by CO_2 radiative forcing were further disturbed by CO_2 physiological forcing-induced responses in many processes, for example, local to large-scale convective activities, atmospheric precipitable water changes, atmospheric circulation anomalies, etc.

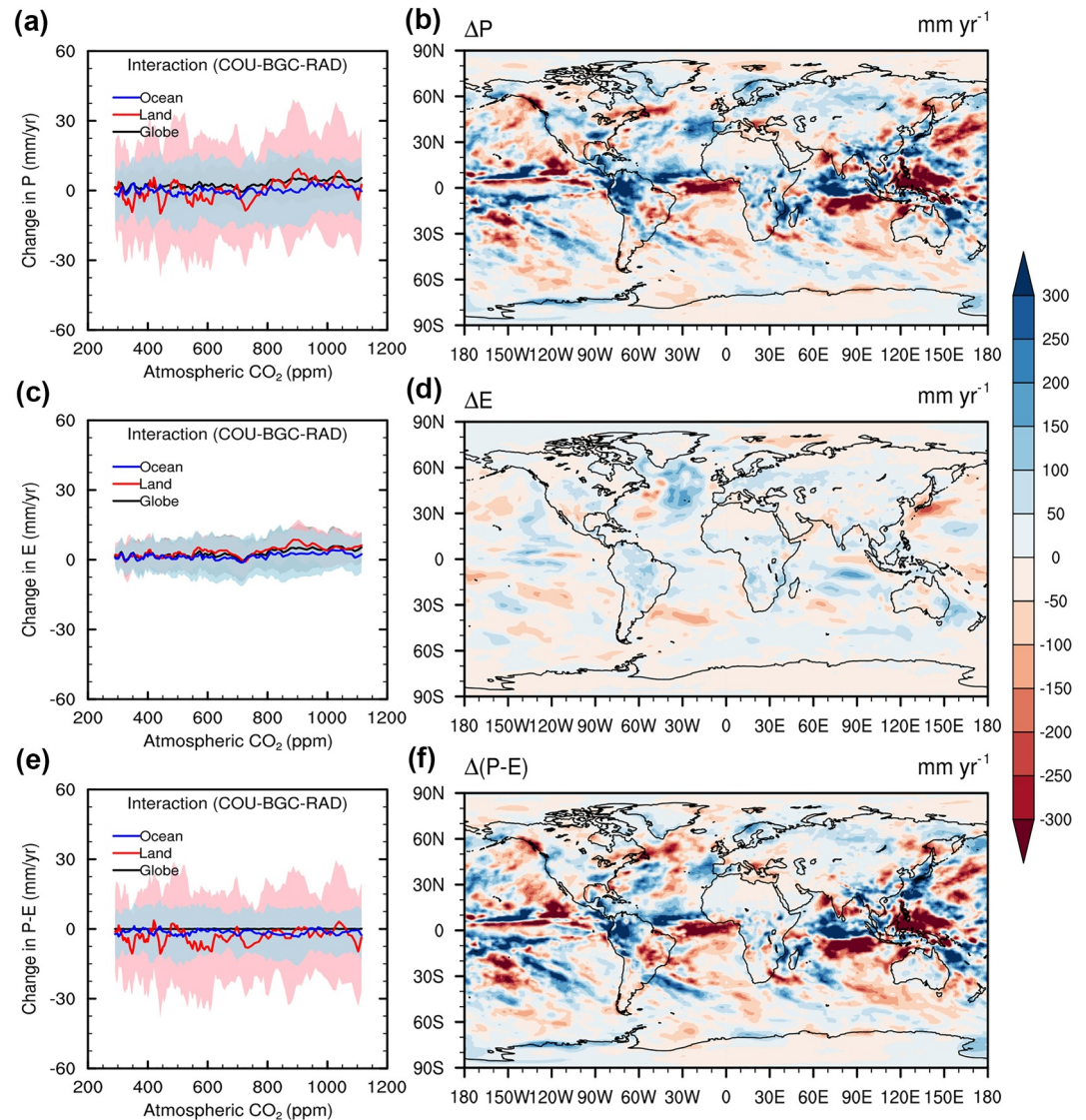


Figure 4. Changes in P , E , and $P - E$ due to interactions between CO_2 physiological and radiative forcings under transient 1%/year increasing CO_2 forcing scenario. (a) Time series of changes in annual precipitation (ΔP) from the COU-BGC-RAD simulations. (b) Spatial pattern of ΔP at transient $4 \times \text{CO}_2$. (c) and (d) Same as (a) and (b) but for changes in evaporation (ΔE). (e) and (f) Same as (a) and (b) but for $\Delta(P - E)$. All results are averaged from 12 CMIP6 models of the 1pctCO₂ experiments.

3.2. Hydrological Sensitivity to Increasing CO_2 Physiological and Radiative Effects

We further estimate the magnitudes of hydrological sensitivity parameters due to the CO_2 physiological forcing and radiative forcing under transient $4 \times \text{CO}_2$. Figure 6 and Table 1 show the multi-model ensemble estimates of the two sensitivity parameters for P , E , and $P - E$ on globe, land and ocean. Due to global surface hydrological flux balance, the hydrological sensitivity to CO_2 physiological forcing for P (i.e., B_P) and E (i.e., B_E) on globe are negative with the same estimate of $-0.97 \pm 0.76 \text{ mm year}^{-1} (100 \text{ ppm})^{-1}$, and the hydrological sensitivity to the CO_2 radiative forcing for P (Γ_P) and E (Γ_E) on globe are positive with $16.84 \pm 2.61 \text{ mm year}^{-1} \text{ K}^{-1}$ (Figure 6). This result indicates that the global hydrological sensitivity to warming is $1.54 \pm 0.24\%$ per Kelvin warming (Table 1). The rising CO_2 radiative forcing intensifies global water cycle, but the CO_2 physiological forcing partly weakens the strength by $-0.09 \pm 0.07\%$ per 100 ppm CO_2 increase. At the global scale, the multi-model ensemble indicates that the CO_2 radiative forcing dominates the overall increases of global P or E at transient $4 \times \text{CO}_2$ with a contribution of $79 \pm 12\%$. The CO_2 physiological forcing has a negative contribution of $-10 \pm 8\%$, but the

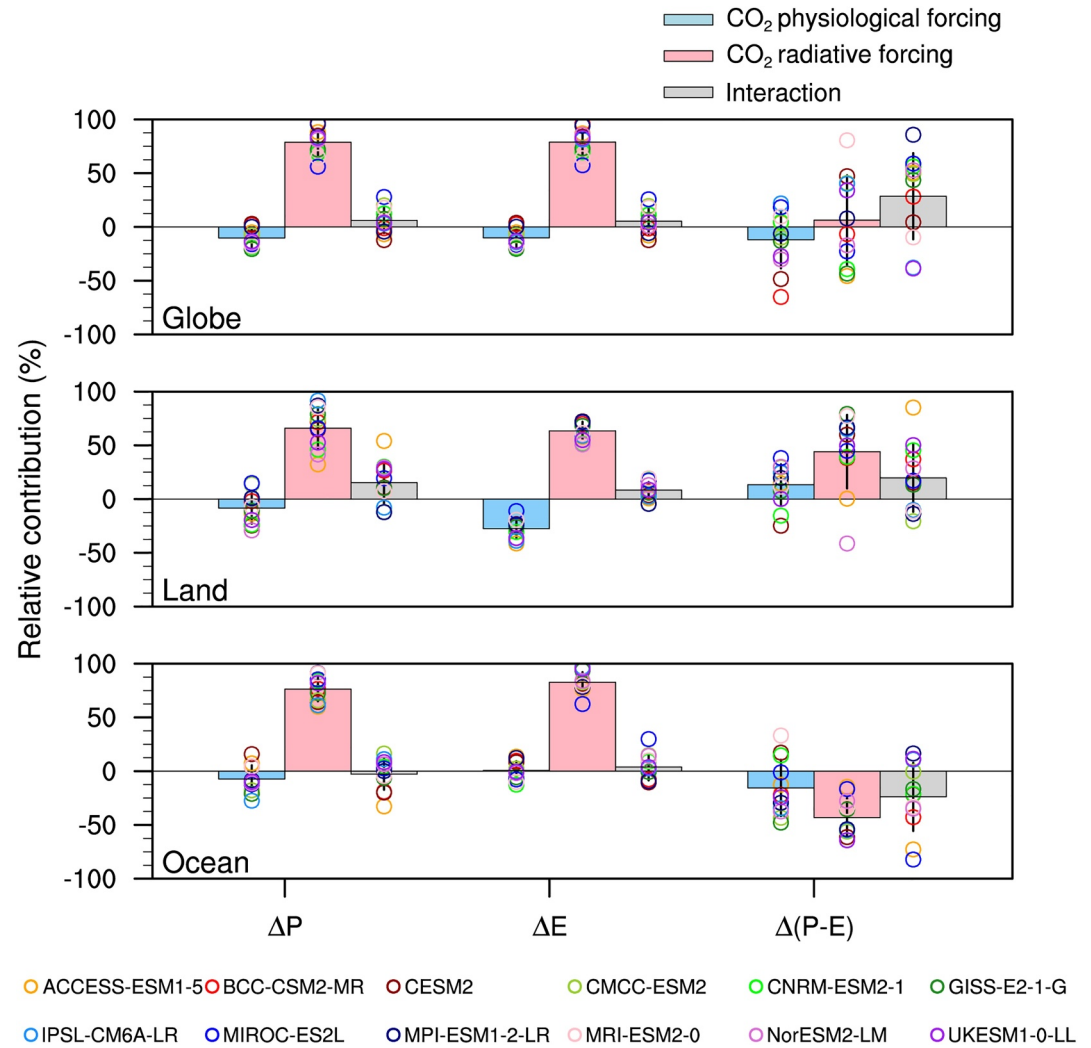


Figure 5. Relative contributions of rising CO₂-induced radiative and physiological forcings to long-term hydrological flux changes at transient 4 × CO₂. Relative contributions to ΔP, ΔE, and Δ(P − E) are calculated on globe, land and ocean, respectively.

interaction between CO₂ radiative and physiological forcing provides a positive contribution of $6 \pm 12\%$ (Figure 5a).

The rising CO₂-induced effects have diverse impacts on water cycles for the land and the ocean surfaces at 4 × CO₂ (Figure 6 and Table 1). On land, the negative response of CO₂ physiological effect for E ($B_{E,L}$) is about 2.5 times larger than the CO₂ physiological effect for P ($B_{P,L}$), which leads to a net positive response of $1.46 \pm 1.34 \text{ mm year}^{-1} (100 \text{ ppm})^{-1}$ for the land water budget ($B_{P-E,L}$) or the CO₂ concentration sensitivity of $0.44 \pm 0.39\%$ per 100 ppm CO₂ increase. The CO₂ radiative effect has stronger positive impact on the land water budget for which the sensitivity parameter ($\Gamma_{P-E,L}$) is $6.62 \pm 3.53 \text{ mm year}^{-1} \text{ K}^{-1}$ or the temperature sensitivity of $1.90 \pm 1.18\%$ per Kelvin warming. The global land surface's wetting (Figures 3g–3i) is dominantly contributed by the CO₂-radiative forcing ($44 \pm 34\%$) and enhanced by the CO₂-physiological forcing ($13 \pm 19\%$) and their interaction with a significant contribution of $20 \pm 32\%$ (Figure 5).

The negative anomalies of water budget ($P-E$) on global ocean surface (Figures 2g–2i) is primarily driven by the negative response due to the CO₂ radiative forcing that contributes $-43 \pm 18\%$ of the overall $P-E$ decrease. This negative hydrological response is further enhanced by the CO₂ physiological forcing ($-16 \pm 26\%$) and the interaction with a significant contribution of $-24 \pm 31\%$. The CO₂-radiative effect on the ocean water budget

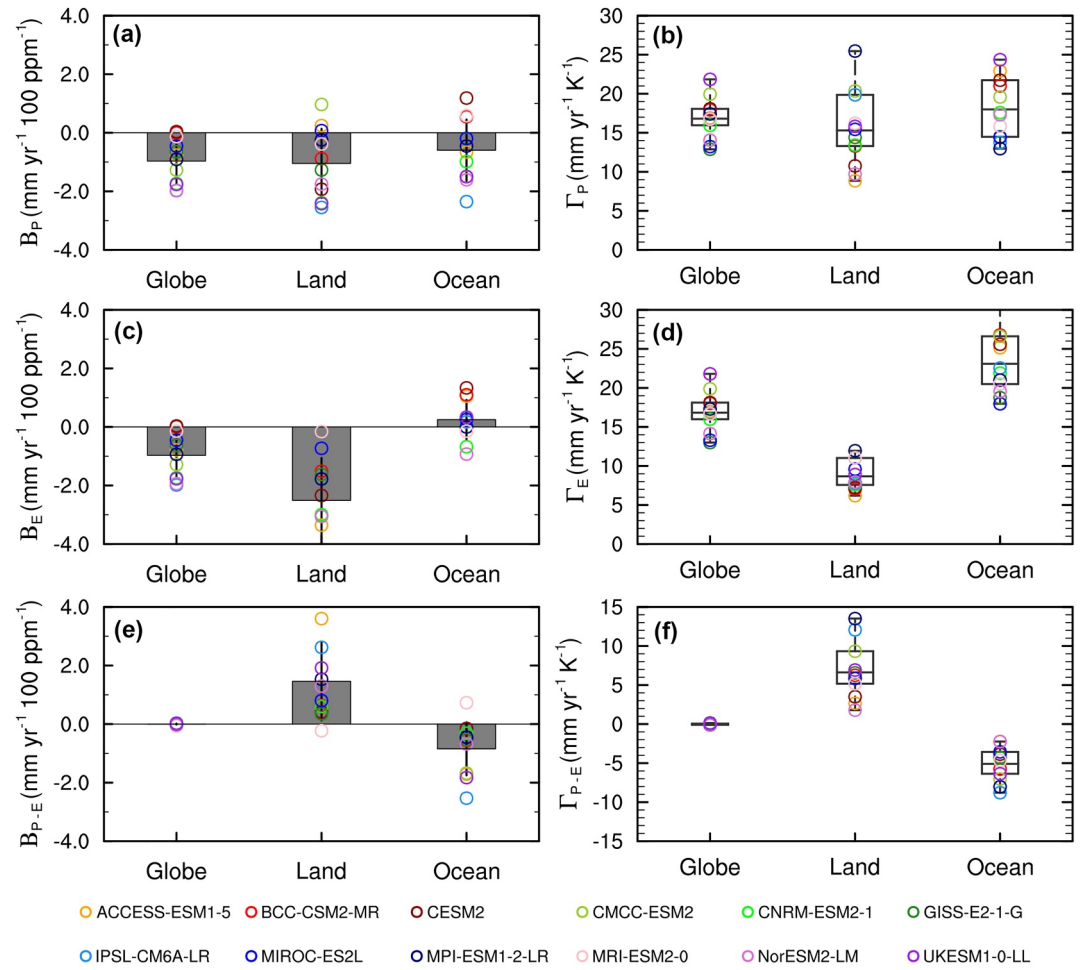


Figure 6. Magnitudes of hydrological sensitivity parameters from rising CO_2 -induced radiative and physiological forcings at transient $4 \times \text{CO}_2$. (a) Sensitivity of precipitation to CO_2 physiological forcing (B_P) on globe, land and ocean, respectively. (b) Sensitivity of precipitation to CO_2 radiative forcing (Γ_P) on globe, land and ocean, respectively. (c, e) Same as (a) but for evaporation (B_E) and precipitation minus evaporation (B_{P-E}), respectively. (d, f) Same as (b) but for evaporation (Γ_E) and precipitation minus evaporation (Γ_{P-E}), respectively.

Table 1

Hydrological Sensitivity Parameters From Global Water Cycle Response to Transient $4 \times \text{CO}_2$ Under 1%/Year CO_2 Forcing Scenario

Parameter ^a	CO ₂ physiological forcing [mm yr ⁻¹ (100 ppm) ⁻¹ or (%/100 ppm)]			CO ₂ radiative forcing [mm yr ⁻¹ K ⁻¹ or (%/K)]		
	B_P	B_E	B_{P-E}	Γ_P	Γ_E	Γ_{P-E}
Globe	-0.97 ± 0.76 (-0.09 ± 0.07)	-0.97 ± 0.76 (-0.09 ± 0.07)	0.00 ± 0.02 (0.00 ± 0.00)	16.84 ± 2.61 (1.54 ± 0.24)	16.83 ± 2.34 (1.54 ± 0.23)	-0.02 ± 0.08 (0.00 ± 0.00)
Land	-1.05 ± 1.18 (-0.13 ± 0.14)	-2.51 ± 1.45 (-0.53 ± 0.30)	1.46 ± 1.34 (0.44 ± 0.39)	15.30 ± 4.79 (1.91 ± 0.78)	8.68 ± 1.83 (1.81 ± 0.46)	6.62 ± 3.53 (1.90 ± 1.18)
Ocean	-0.60 ± 1.05 (-0.05 ± 0.09)	0.25 ± 0.68 (0.02 ± 0.05)	-0.84 ± 0.91 (0.57 ± 0.57)	17.99 ± 3.86 (1.53 ± 0.31)	23.09 ± 3.87 (1.75 ± 0.28)	-5.10 ± 2.13 (3.70 ± 1.61)

Note. Results are ensemble mean \pm s.d. of 12 CMIP6 models. Bold values in () are for percentage changes. ^aNote that B_X is defined as $\frac{\partial X}{\partial \text{CO}_2}$, Γ_X is defined as $\frac{\partial X}{\partial T}$, where X are P , E or $P - E$.

($\Gamma_{P-E,O}$) is $3.70 \pm 1.61\%$ per Kelvin warming, which is due to the larger positive sensitivity for E ($\Gamma_{E,O}$) than the positive sensitivity for P ($\Gamma_{P,O}$) on oceans (Figure 6, Table 1). However, the negative response by the CO_2 -physiological forcing for ocean water budget ($B_{P-E,O}$) is due to the negative sensitivity for P ($B_{P,O}$) and the smaller sensitivity for E ($B_{E,O}$) on ocean (Figure 6).

The above results indicate the global-scale estimates of the two hydrological sensitivities on globe, land and ocean. But how these sensitivities are partitioned to regional scales or latitudinal zones is still unknown. To this end, we further calculated the hydrological sensitivities across latitudinal zones using Equations 2 and 3 from the longitude-averaged ΔP , ΔE , $\Delta(P - E)$, and ΔC , ΔT at transient $4 \times \text{CO}_2$ (Figure S3 in Supporting Information S1). From the zonal average perspective, the global hydrological sensitivity by CO_2 -radiative forcing is positively dominated by P and $P - E$ over high latitudes (50°S – 70°S) and tropical regions (10°S – 10°N) on both land and ocean (Figures S3b and S3f in Supporting Information S1). The hydrological sensitivity by CO_2 -physiological forcing is negatively dominated by E over tropical regions (-30°S – 15°N) on land, but strongly positively dominated by P and $P - E$ over tropical oceans (10°S – 10°N).

3.3. Evolution of Hydrological Sensitivity With Increasing CO_2 Forcing

The above analyses of hydrological sensitivity to CO_2 physiological and radiative effects are based on the long-term changes of hydrological flux variables at transient $4 \times \text{CO}_2$. It is unknown whether the hydrological sensitivities by CO_2 physiological and radiative forcings are consistent over doubling CO_2 forcing levels. To examine whether the global water cycle linearly response to the rising CO_2 -induced physiological and radiative forcing, we calculated annual values of global hydrological sensitivity parameters (B and Γ) for P , E , and $P - E$ over the transient 1%/year increasing CO_2 forcing from $1 \times \text{CO}_2$ to $4 \times \text{CO}_2$.

Figure 7 shows that the global hydrological sensitivities by both CO_2 physiological and radiative forcing have little changes over transient $2 \times \text{CO}_2$ (570 ppm) to $4 \times \text{CO}_2$ (1140 ppm) under the 1% per year CO_2 increase scenario. Compared to the parameter values at transient $4 \times \text{CO}_2$ (Figure 6), the global P and E forced by CO_2 physiological effect (B_P and B_E) increase from about $-8 \text{ mm year}^{-1} (100 \text{ ppm})^{-1}$ at $1 \times \text{CO}_2$ (285 ppm) to a relative stable value (about $-1 \text{ mm year}^{-1} (100 \text{ ppm})^{-1}$) at $2 \times \text{CO}_2$, then have little fluctuations over transient $2 \times \text{CO}_2$ to $4 \times \text{CO}_2$ (Figures 7a and 7c). The global hydrological sensitivities to warming or for CO_2 radiative effect feedback (Γ_P and Γ_E) increase to relative stable values early at $1.75 \times \text{CO}_2$ (500 ppm) and have little changes with CO_2 forcing increased to transient $4 \times \text{CO}_2$. The hydrological sensitivities on land and ocean show the similar results. The large changes and uncertainties in global hydrological sensitivity to CO_2 increase or warming over transient $1 \times \text{CO}_2$ to $2 \times \text{CO}_2$ is likely driven by the strong inter-annual variations of global P and E that are modulated by climate internal variability, which covered up the much slower and smaller changes in P and E due to CO_2 increase or warming only. It is also interesting that the global hydrological sensitivity to warming for P (Γ_P) on land and ocean are basically consistent with the global estimate over transient $2 \times \text{CO}_2$ to $4 \times \text{CO}_2$ (Figure 7b). The increased $\Delta(P - E)$ on land or declined $\Delta(P - E)$ on ocean with CO_2 forcing also nearly linearly response to the CO_2 physiological and radiative effects over transient $2 \times \text{CO}_2$ to $4 \times \text{CO}_2$ (Figures 7e and 7f).

4. Discussion and Conclusions

Most previous studies investigated the hydrological sensitivity to global warming in a fully coupled climate system, which is the so called “apparent hydrological sensitivity” (Fläschner et al., 2016), but few separately quantify the temperature sensitivity of hydrological variables solely due to CO_2 radiative effect, and the CO_2 concentration sensitivity of hydrological variables due to CO_2 vegetation physiological effect. For instance, the early versions of climate models predicted that the apparent hydrological sensitivity of global precipitation is in a very likely range of 1%–3% per Kelvin warming (Covey et al., 2003; Douville et al., 2023; Held & Soden, 2006). This model-based range of hydrological sensitivity is still not largely narrowed by recent estimates from CMIP5/6 models (Andrews & Forster, 2010; Douville et al., 2023; Fläschner et al., 2016; Shioyama et al., 2022). In this study, from the fully coupled simulations of 12 CMIP6 models under transient $4 \times \text{CO}_2$, the estimated apparent hydrological sensitivity (η_a) is 1.17%–1.61% per Kelvin warming. It should be noted that the temperature sensitivity of the hydrological transient responses to atmospheric CO_2 increase in this study could be much smaller than those from the stable climate state or the abrupt $4 \times \text{CO}_2$ scenario due to that a transient climate

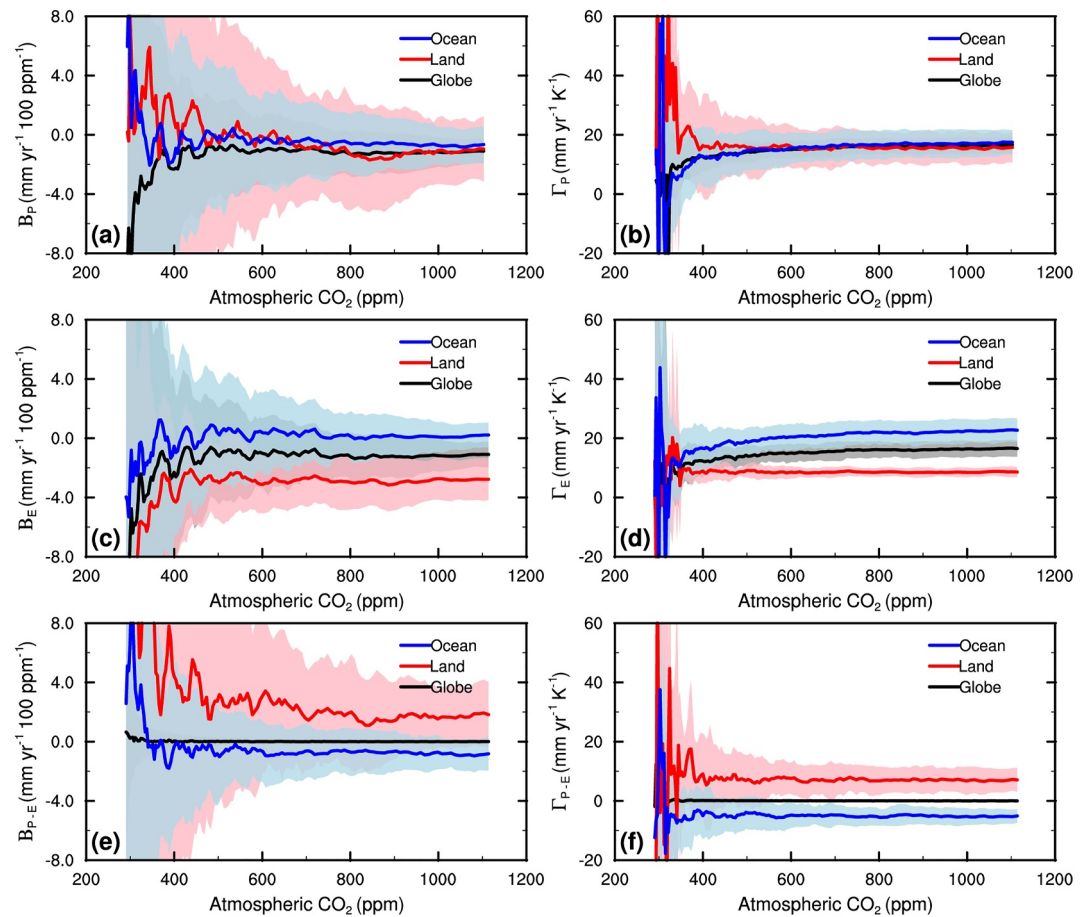


Figure 7. Evolution of hydrological sensitivity parameters from rising CO_2 -induced radiative and physiological forcings over transient 1%/year increasing CO_2 forcing ($1 \times \text{CO}_2$ to $4 \times \text{CO}_2$). (a) Sensitivity of precipitation to CO_2 physiological forcing (B_P) on globe, land and ocean, respectively. (b) Sensitivity of precipitation to CO_2 radiative forcing (Γ_P) on globe, land and ocean, respectively. (c, e) Same as (a) but for evaporation (B_E) and precipitation minus evaporation (B_{P-E}), respectively. (d, f) Same as (b) but for evaporation (Γ_E) and precipitation minus evaporation (Γ_{P-E}), respectively. Shaded area represents mean \pm standard deviation from 12 CMIP6 models of the 1pct CO_2 experiments.

induces a fast decreased precipitation response to CO_2 radiative forcing (Andrews & Forster, 2010; Fläschner et al., 2016).

Using the proposed analysis framework of global water cycle in response to rising CO_2 , this study defined and quantified the global hydrological sensitivity to rising CO_2 -induced physiological and radiative forcings from decoupling simulations of the CMIP6 1pct CO_2 experiments. The CMIP6 models estimated that the global hydrological sensitivity to global warming solely due to the CO_2 radiative forcing is $1.54 \pm 0.24\%$ per Kelvin warming, which about 10% larger than the apparent temperature sensitivity calculated from the fully coupled simulations. The global hydrological sensitivity (precipitation or evaporation) by CO_2 -physiological forcing is $-0.09 \pm 0.07\%$ (100 ppm) $^{-1}$ at transient $4 \times \text{CO}_2$. These global hydrological sensitivities by CO_2 physiological or radiative forcings are relatively stable over transient $2 \times \text{CO}_2$ to $4 \times \text{CO}_2$, which is consistent with the finding that the apparent hydrological sensitivity to warming has little change at large CO_2 forcing between transient $2 \times \text{CO}_2$ and $4 \times \text{CO}_2$ (Raiter et al., 2023).

With the CMIP6 1pct CO_2 experiments, this study isolated the relative contributions of rising CO_2 -induced physiological and radiative forcings and the interaction between the two to the transient responses of global water cycle to increasing CO_2 forcing. At the global scale, the overall increases of global P or E at transient $4 \times \text{CO}_2$ are dominated by the CO_2 radiative forcing ($79 \pm 12\%$) and positively contributed by the interaction ($6 \pm 12\%$) but are reduced by the CO_2 physiological forcing ($-10 \pm 8\%$). On land, the pattern of precipitation minus evaporation

show that the wetting of global land is dominantly contributed by the CO₂ radiative forcing ($44 \pm 34\%$) and enhanced by physiological forcing ($13 \pm 19\%$) and their interaction ($20 \pm 32\%$). In last two decades, a recent study indicated that global water availability ($P-E$) has significantly decreased (Y. Zhang, Li, et al., 2023). The declining trend on land can be reversed for a long period of time, for example, in the transient $4 \times \text{CO}_2$ scenario. However, there are still uncertainties since climate mode changes play a key role in controlling global land water availability. The accuracy of Earth System Models (ESMs) in predicting these climate mode changes is a critical factor for accurately projecting changes in global land water availability. Additionally, 12 model members used in this study may not be the optimal ensemble size to capture robust modeling uncertainty of sensitivity and contribution estimations. Overall, this study quantified hydrological sensitivity to rising CO₂ physiological and radiative forcings and emphasized the importance of CO₂ physiological forcing and interactive effect between physiological and radiative forcings in global water cycle projections under a CO₂-enriched climate.

Data Availability Statement

All simulations of the CMIP6 models for the 1pctCO₂ experiments (Jones et al., 2016) can be freely downloaded from the Earth System Grid Federation (<https://esgf-node.llnl.gov/search/cmip6/>). The supplementary text, tables, and figures are provided in Supporting Information S1. The additional resulting data set is provided at X. Zhang (2023).

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