This is a repository copy of Precipitation, radiative forcing and global temperature change.

White Rose Research Online URL for this paper:
http://eprints.whiterose.ac.uk/43204/

Article:
Andrews, T, Forster, PM, Boucher, O et al. (2 more authors) (2010) Precipitation, radiative forcing and global temperature change. Geophysical Research Letters, 37. ISSN 0094-8276

https://doi.org/10.1029/2010GL043991

Reuse
See Attached

Takedown
If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.
Precipitation, radiative forcing and global temperature change

Timothy Andrews,¹ Piers M. Forster,¹ Olivier Boucher,² Nicolas Bellouin,² and Andy Jones²

Received 14 May 2010; accepted 24 May 2010; published 21 July 2010.

¹Institute for Climate and Atmospheric Science, School of Earth and Environment, University of Leeds, Leeds, UK.
²Met Office Hadley Centre, Exeter, UK.

Copyright 2010 by the American Geophysical Union.
0094-8276/10/2010GL043991

1. Introduction

[1] Radiative forcing is a useful tool for predicting equilibrium global temperature change. However, it is not so useful for predicting global precipitation changes, as changes in precipitation strongly depend on the climate change mechanism and how it perturbs the atmospheric and surface energy budgets. Here a suite of climate model experiments and radiative transfer calculations are used to quantify and assess this dependency across a range of climate change mechanisms. It is shown that the precipitation response can be split into two parts: a fast atmospheric response that strongly correlates with the atmospheric component of radiative forcing, and a slower response to global surface temperature change that is independent of the climate change mechanism, ~2–3% per unit of global surface temperature change. We highlight the precipitation response to black carbon aerosol forcing as falling within this range despite having an equilibrium response that is of opposite sign to the radiative forcing and global temperature change.


2. Experiments and Method

2.1. Model Setup

[5] We use an atmospheric model based on the Hadley Centre climate model HadGEM1 [Martin et al., 2006] with various improvements coupled to a 50 m thermodynamic mixed-layer ocean and sea-ice model (see Jones et al. [2007] for further description of the improvements and base configuration of the model). A mixed-layer ocean model is a simplification of the climate system, however, Takahashi [2009] found the hydrological sensitivities of CMIP3 models to be representative of their fully dynamic ocean model counterparts. Nine different forcing scenarios (Table 1) have been integrated from a control simulation based on pre-industrial (year 1860) conditions. The nine forcing scenarios were integrated for 10 years in atmosphere-only mode (i.e., using prescribed sea-surface-temperatures (SSTs) and sea-ice extent) and 30 years in the full atmosphere/mixed-layer mode.

2.2. Method: Fast, Slow and Total Responses

[6] Precipitation can respond to both changes in ΔT and to the forcing agent itself. The direct response to the forcing agent can be demonstrated in forcing experiments with SSTs held fixed (as ΔT is largely prohibited). For example when CO₂ is increased but SSTs held fixed, the precipitation and evaporation rate go down [e.g., Yang et al., 2003; Dong et al., 2009; Bala et al., 2009] due to small increases in tropospheric
temperatures above an unchanged surface, increasing atmospheric stability and reducing convection [Dong et al., 2009]. In addition to this response, precipitation also responds to ΔT, through various climate feedbacks that affect precipitation processes, climate models suggest this to be ~2–3% K⁻¹ [Held and Soden, 2006; Lambert and Webb, 2008]. These two responses emerge on different timescales due to the smaller heat capacity of the atmosphere compared to the ocean: for example, in response to increased CO₂ the precipitation rate initially goes down due to the direct atmospheric response, before subsequently increasing on a multi-annual timescale associated with ΔT [e.g., Yang et al., 2003; Andrews et al., 2009; Bala et al., 2009]. We refer to these precipitation responses as the “fast,” ΔP_{fast} and “slow,” ΔP_{slow}, responses, respectively.

To quantify the ΔP_{fast} and ΔP_{slow} terms we follow the method put forward by Bala et al. [2009]. The total change in any climate variable x is the sum of the fast and slow responses, Δx_{total} = Δx_{fast} + Δx_{slow}. The total change is simply the change in the full atmosphere/mixed-layer integrations, from which we discard the first 10-years (the transient part) and calculate equilibrium responses from the average of the remaining 20-years. The fast component is determined from the atmosphere-only integrations where ΔT is largely prohibited; we average over all 10-years. We calculate the slow response from subtracting the fast response from the total response [see Bala et al., 2009]. Note that fast and slow responses can also be estimated from linear regression during transient climate change [Gregory and Webb, 2008], which can also provide useful energetic constraints on how much precipitation increases with ΔT [e.g., Takahashi, 2009], but is inappropriate here as the transient change (ΔT) is not large enough in most of our experiments to constrain the regression lines.

In the following analysis the reported error in any term x is calculated from Monte Carlo simulations. We randomly sample an equal number of annual-means from the original dataset to create 10,000 subsets, and then compute the required term x from each of these sets. The 95% uncertainty in x is then determined from the standard deviation of the 10,000 simulated x values.

### 2.3. Radiative Forcing Calculations

For CO₂, CH₄, O₃ and the solar experiment, radiative forcings are calculated offline using the Edwards-Slingo radiation code [Edwards and Slingo, 1996]. We take the difference between two calls of the radiation code, one based on the control configuration and one with the forcing agent introduced. The influence of stratospheric adjustment is accounted for using the fixed-dynamical-heating approximation [Forster et al., 1997]. For black carbon (BC) aerosol it is important to resolve high-frequency temporal variations because the relative position of cloud and aerosols are important. Therefore, the radiative forcing is calculated online by running a 5-year simulation with the model in atmosphere-only mode. As with the offline calculations, the radiation code is called twice, the call based on the control configuration is passed on to the next model time step. The snow albedo forcing is calculated similarly [see Bellouin and Boucher, 2010]. As SO₄ and biomass burning (BB) aerosol act as cloud condensation nuclei in the model these radiative forcings are calculated from the change in radiation balance in the fixed-SST experiments, which allows the aerosol indirect effects to be incorporated into the forcing estimate [e.g., Lohmann et al., 2010]. Note that as stratospheric adjustment is accounted for (or is negligible for the black carbon and snow albedo forcing) the radiative forcing at the TOA and tropopause is the same, hence any atmospheric forcing only remains in the troposphere.

### 3. Results

The fast, slow and total precipitation responses are shown in Table 2. To compare across the forcing scenarios, we normalise the total precipitation response by total surface-

---

**Table 1. Description of the Forcing Scenarios**

<table>
<thead>
<tr>
<th>Forcing Scenario</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CO₂</td>
<td>CO₂ concentration increased to 2005 levels (286 → 379 ppmv)</td>
</tr>
<tr>
<td>2 x CO₂</td>
<td>CO₂ concentration doubled (286 → 572 ppmv)</td>
</tr>
<tr>
<td>CH₄</td>
<td>CH₄ concentration increased to 2005 levels (805 → 1,774 ppbv)</td>
</tr>
<tr>
<td>Snow albedo</td>
<td>Land snow albedo decreased from 0.80 to 0.75 (^{\text{b}})</td>
</tr>
<tr>
<td>Solar</td>
<td>Solar constant increased by 2% (1365 → 1392.3 W m⁻²)</td>
</tr>
<tr>
<td>O₃</td>
<td>Tropospheric and stratospheric O₃ changed to 1990 levels</td>
</tr>
<tr>
<td>SO₂</td>
<td>SO₂ emissions changed from 1860 to year 2000 (^{\text{c}})</td>
</tr>
<tr>
<td>BB</td>
<td>Biomass burning emissions changed from 1860 to year 2000 (^{\text{c}})</td>
</tr>
<tr>
<td>BC</td>
<td>Black carbon emissions changed from 1860 to year 2000 (^{\text{c}})</td>
</tr>
</tbody>
</table>

\(^{\text{a}}\)The base configuration of the model uses greenhouse gas concentrations and aerosol emissions representative of the year 1860.

\(^{\text{b}}\)See Bellouin and Boucher [2010] for further description.

\(^{\text{c}}\)See Jones et al. [2007] for further description.

---

![Figure 1. Comparison of the hydrological sensitivity (ΔP/ΔT) for a standard method that just considers the total responses only (left), and a method that considers only the slow components of climate change (right). The apparent dependence of the hydrological sensitivity on the forcing agent is largely removed if the fast precipitation response is removed.](image-url)
air-temperature change (fifth column in Table 2). This “hydrological sensitivity” is seen to vary considerably across the forcing mechanisms (range is −0.24 to +3.5% K⁻¹), recovering the result of previous studies that precipitation changes strongly depend on the nature of the forcing mechanism. However, if we compute the hydrological sensitivity for only the slow components of climate change (that is, the component that scales with ΔT), we find a hydrological sensitivity (sixth column in Table 2) that is now in good agreement across the forcing agents; in all experiments the slow precipitation response to ΔT is of the order 2–3% K⁻¹. Previously this has had only been shown for the case of CO₂ and solar forcing [Lambert and Faull, 2007; Andrews et al., 2009; Bala et al., 2009]; our results show it to be a robust result across a range of climate change mechanisms. The comparison is illustrated in Figure 1. It is notable that even black carbon aerosol, which had a negative hydrological sensitivity, falls within the 2–3% K⁻¹ range when only the slow components of climate change are considered.

To further understand the fast and slow precipitation responses we examine the link between precipitation changes and radiative forcing as measured at the TOA, Ftoa, and its partitioning between the surface, Fsurf, and atmosphere, Fatm. Table 3 shows the radiative forcing terms and the ratio R = Fsurf/Ftotal. For climate change mechanisms that force climate through the scattering of solar radiation, such as SO₂ and snow albedo, or changes in the solar constant, R is close to unity, indicating that the atmosphere is largely transparent to these forcings, which are felt at the surface. Black carbon aerosol strongly absorbs solar radiation in the atmosphere, which reduces solar radiation reaching the surface (Fsurf is negative) but also reduces outgoing solar radiation to space (Ftoa is positive). Therefore the surface radiative forcing is of opposite sign to the radiative forcing at the TOA [e.g., Ramanathan et al., 2001], as seen in Table 3 where R = −1.5. Biomass burning aerosol both scatters, which generates a negative radiative forcing at the TOA and surface, and absorbs solar radiation, which generates a small positive atmospheric component, therefore R > 1. O₃ radiative forcing is more complex due to competing effects in the solar and thermal spectra, as well as competing increases and decreases in concentration through the troposphere and stratosphere. The net global-mean effect is an R close to unity. The radiative forcing of both CO₂ and CH₄ increases with height throughout the troposphere [Collins et al., 2006], hence it is mostly absorbed in the atmosphere and R < 1.

Table 2. The Fast, Slow and Total Precipitation Responses

<table>
<thead>
<tr>
<th>Forcing Scenario</th>
<th>ΔPfast (%)</th>
<th>ΔPslow (%)</th>
<th>ΔPtotal (%)</th>
<th>ΔPtotal/ΔTtotal (% K⁻¹)</th>
<th>ΔPslow/ΔTslow (% K⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CO₂</td>
<td>−1.12 ± 0.14</td>
<td>3.67 ± 0.23</td>
<td>2.55 ± 0.18</td>
<td>1.68 ± 0.10</td>
<td>2.49 ± 0.14</td>
</tr>
<tr>
<td>2 x CO₂</td>
<td>−2.53 ± 0.18</td>
<td>8.71 ± 0.26</td>
<td>6.18 ± 0.18</td>
<td>1.60 ± 0.03</td>
<td>2.41 ± 0.06</td>
</tr>
<tr>
<td>CH₄</td>
<td>−0.21 ± 0.23</td>
<td>1.30 ± 0.32</td>
<td>1.09 ± 0.22</td>
<td>1.98 ± 0.52</td>
<td>2.40 ± 0.86</td>
</tr>
<tr>
<td>Snow albedo</td>
<td>−0.04 ± 0.19</td>
<td>1.30 ± 0.24</td>
<td>1.26 ± 0.14</td>
<td>1.75 ± 0.18</td>
<td>2.11 ± 0.37</td>
</tr>
<tr>
<td>Solar</td>
<td>−0.91 ± 0.17</td>
<td>10.23 ± 0.27</td>
<td>9.31 ± 0.21</td>
<td>2.22 ± 0.04</td>
<td>2.53 ± 0.06</td>
</tr>
<tr>
<td>O₃</td>
<td>0.02 ± 0.17</td>
<td>0.53 ± 0.23</td>
<td>0.35 ± 0.15</td>
<td>2.84 ± 0.83</td>
<td>2.61 ± 1.44</td>
</tr>
<tr>
<td>SO₂</td>
<td>0.03 ± 0.13</td>
<td>−2.90 ± 0.19</td>
<td>−2.87 ± 0.13</td>
<td>2.46 ± 0.11</td>
<td>2.56 ± 0.16</td>
</tr>
<tr>
<td>BB</td>
<td>−0.18 ± 0.19</td>
<td>−0.57 ± 0.24</td>
<td>−0.75 ± 0.14</td>
<td>3.50 ± 0.74</td>
<td>3.06 ± 1.05</td>
</tr>
<tr>
<td>BC</td>
<td>−0.68 ± 0.18</td>
<td>0.61 ± 0.26</td>
<td>−0.07 ± 0.19</td>
<td>−0.24 ± 0.72</td>
<td>2.25 ± 0.87</td>
</tr>
</tbody>
</table>

Table 3. Radiative Forcing at the Top-of-Atmosphere and Its Partitioning Between the Surface and Atmosphere

<table>
<thead>
<tr>
<th>Forcing Scenario</th>
<th>Ftoa</th>
<th>Fsurf</th>
<th>Fatm</th>
<th>R = Fsurf/Ftotal</th>
<th>ΔLH</th>
<th>ΔSH</th>
</tr>
</thead>
<tbody>
<tr>
<td>CO₂</td>
<td>1.39</td>
<td>0.28</td>
<td>1.11</td>
<td>0.2</td>
<td>1.02 ± 0.13</td>
<td>0.01 ± 0.06</td>
</tr>
<tr>
<td>2 x CO₂</td>
<td>3.47</td>
<td>0.66</td>
<td>2.81</td>
<td>0.2</td>
<td>2.30 ± 0.18</td>
<td>0.14 ± 0.05</td>
</tr>
<tr>
<td>CH₄</td>
<td>0.51</td>
<td>0.26</td>
<td>0.25</td>
<td>0.5</td>
<td>0.18 ± 0.21</td>
<td>0.07 ± 0.07</td>
</tr>
<tr>
<td>Snow albedo</td>
<td>0.28</td>
<td>0.29</td>
<td>−0.02</td>
<td>1.0</td>
<td>0.02 ± 0.18</td>
<td>−0.06 ± 0.06</td>
</tr>
<tr>
<td>Solar</td>
<td>4.86</td>
<td>3.7</td>
<td>1.16</td>
<td>0.8</td>
<td>0.84 ± 0.16</td>
<td>−0.08 ± 0.07</td>
</tr>
<tr>
<td>O₃</td>
<td>0.16</td>
<td>0.21</td>
<td>−0.05</td>
<td>1.3</td>
<td>−0.02 ± 0.15</td>
<td>−0.06 ± 0.05</td>
</tr>
<tr>
<td>SO₂</td>
<td>−1.15</td>
<td>−2.17</td>
<td>0.01</td>
<td>−0.03 ± 0.13</td>
<td>0.05 ± 0.06</td>
<td></td>
</tr>
<tr>
<td>BB</td>
<td>−0.29</td>
<td>−0.54</td>
<td>0.25</td>
<td>1.9</td>
<td>0.16 ± 0.17</td>
<td>0.09 ± 0.04</td>
</tr>
<tr>
<td>BC</td>
<td>0.36</td>
<td>−0.54</td>
<td>0.90</td>
<td>−1.5</td>
<td>0.62 ± 0.18</td>
<td>0.20 ± 0.05</td>
</tr>
</tbody>
</table>

Notes: Fast, ΔPfast; slow, ΔPslow; total, ΔPtotal = ΔPfast + ΔPslow. Also shown is the hydrological sensitivity (ΔP/ΔT) for i) standard methods that do not separate fast and slow responses (ΔPtotal/ΔTtotal), and, ii) for the slow responses only (ΔPslow/ΔTslow). Errors represent 95% uncertainties.
respectively). $F_{atm}$ does appear to be largely balanced by $\Delta LH$, and $\Delta SH$ is small. However, in general the sum of $\Delta LH$ and $\Delta SH$ is not large enough to meet $F_{atm}$ exactly. This is expected: it is the result of small changes in atmospheric temperatures that act to radiate away some of $F_{atm}$. Changes in temperature are required as a physical mechanism to change $\Delta LH$ and $\Delta SH$, as well as other possible factors such as changes in clouds [e.g., Gregory and Webb, 2008; Andrews and Forster, 2008].

[13] We further note that the residual between $F_{atm}$ and the sum of $\Delta LH$ and $\Delta SH$ is slightly larger for the solar forcing than CO$_2$, despite them having a similar $F_{atm}$ component (Table 3). This supports the findings of Lambert and Faull [2007] that the atmospheric component of solar radiative forcings is balanced more by atmospheric warming than precipitation changes, compared to a CO$_2$ scenario. A thorough explanation would require a detailed analysis of the physical mechanisms, but our central point still holds, that is, the dominant term in eliminating $F_{atm}$ is a change in precipitation (even for solar forcings (Table 3)).

[14] Figure 2 (right) shows the relationship between the slow precipitation response and components of radiative forcing across the experiments. $F_{toa}$ is an excellent predictor of $\Delta P_{slow}$ (Figure 2b), while $F_{srf}$ and $F_{atm}$ are relatively poor predictors (Figures 2d and 2f). The correlation between $F_{toa}$ and $\Delta P_{slow}$ is simply an expression of the radiative forcing concept, i.e., $\Delta P_{slow}$ scales with $\Delta T$ which can be predicted from the radiative forcing.

4. Conclusions and Discussion

[15] We have examined the precipitation response to a range of climate change mechanisms and quantified its
relationship to radiative forcing, including its partitioning between the atmosphere and surface. By distinguishing two different timescale responses, a quick atmospheric response to the forcing and a slower response to global temperature change, we have shown that the apparent forcing dependence of the hydrological sensitivity stems from a fast atmospheric response (see Figure 1). The fast response correlates extremely well with the amount of radiative forcing absorbed in the atmosphere, suggesting that the fast precipitation changes are a mechanism in which the atmosphere regains its energy balance. In all the experiments the subsequent response of precipitation to global surface–air-temperature change is ~ 2–3% K⁻¹ and can be predicted from the radiative forcing as measured at the tropopause or TOA (assuming a sensitivity).

[16] These results improve our understanding of precipitation changes and its relationship to radiative forcing. For example, it shows that climate change mechanisms that largely force climate through the scattering of solar radiation, such as SO₂ and snow albedo, or changes in the solar constant, largely perturb precipitation through changes in global surface temperature which can be predicted through traditional radiative forcing calculations. On the other hand, climate change mechanisms that significantly force climate through absorption of radiation in the troposphere, such as greenhouse gases and black carbon aerosol, have opposing impacts on the precipitation, which require a partitioning of the radiative forcing between the surface and atmosphere to be predicted. In most cases the response to global surface temperature change dominates. However, for black carbon aerosol, the atmospheric component of radiative forcing is sufficiently large that the precipitation response may be of opposite sign to the radiative forcing and global surface temperature change.

[17] We encourage other modelling groups to perform similar experiments to check the robustness of these results. In particular, it has been suggested that fast responses may emerge on different timescales in climate models with full ocean dynamics compared to their mixed-layer counterparts [Williams et al., 2008]. Nevertheless, our model results have practical applications to real world observations and climate predictions. For example, observed estimates of the hydrological sensitivity are not in fact measuring the true precipitation response to global temperature change, they are measuring the fast and slow precipitation responses as they evolve together. Therefore observed estimates of the hydrological sensitivity are likely to change in the future as forcings change and should not be used to project hydrological cycle changes into the future. Our findings offer the opportunity to improve simple climate models used to extrapolate more complex climate models to multiple climate change scenarios. We suggest that observations and projections of precipitation changes will be aided by expanding the radiative forcing concept to include the atmospheric and surface energy budgets, as highlighted here and by the National Research Council [2005].

References
National Research Council (2005), Radiative Forcing of Climate Change, 207 pp., National Acad. Press, Washington, D. C.

T. Andrews and P. M. Forster, Institute for Climate and Atmospheric Science, School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK. (t.andrews@see.leeds.ac.uk)

N. Bellouin, O. Boucher, and A. Jones, Met Office Hadley Centre, FitzRoy Road, Exeter EX1 3PB, UK.