Exploiting the Abrupt $4 \times CO_2$ Scenario to Elucidate Tropical Expansion Mechanisms

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ABSTRACT

Future emissions of greenhouse gases into the atmosphere are projected to result in significant circulation changes. One of the most important changes is the widening of the tropical belt, which has great societal impacts. Several mechanisms (changes in surface temperature, eddy phase speed, tropopause height, and static stability) have been proposed to explain this widening. However, the coupling between these mechanisms has precluded elucidating their relative importance. Here, the abrupt quadrupled-CO$_2$ simulations of phase 5 of the Coupled Model Intercomparison Project (CMIP5) are used to examine the proposed mechanisms. The different time responses of the different mechanisms allow us to disentangle and evaluate them. As suggested by earlier studies, the Hadley cell edge is found to be linked to changes in subtropical baroclinicity. In particular, its poleward shift is accompanied by an increase in subtropical static stability (i.e., a decrease in temperature lapse rate) with increased CO$_2$ concentrations. These subtropical changes also affect the eddy momentum flux, which shifts poleward together with the Hadley cell edge. Transient changes in tropopause height, eddy phase speed, and surface temperature, however, were found not to accompany the poleward shift of the Hadley cell edge. The widening of the Hadley cell, together with the increase in moisture content, accounts for most of the expansion of the dry zone. Eddy moisture fluxes, on the other hand, are found to play a minor role in the expansion of the dry zone.

1. Introduction

The thermal structure of the atmosphere, and its circulation, are projected to change over the next few decades (IPCC 2013; Vallis et al. 2015). One of the most robust signals to emerge from climate models is the poleward shift of the latitudinal edges of the tropical belt (e.g., Hu et al. 2013; Kang et al. 2013). Such a widening of the tropical belt would directly affect the hydrological cycle in semiarid regions (IPCC 2014), as it pushes the dry zone (where most of the world’s deserts are concentrated) poleward, further drying the subtropics (e.g., Lu et al. 2007; Scheff and Frierson 2012).

In spite of the robust tropical expansion projected by climate models, large uncertainty remains in the observed trends over the last several decades (Davis and Rosenlof 2012; Allen et al. 2014; Lucas et al. 2014). Studies have reported that the tropical belt has been expanding poleward at a rate of 0.3°–2° decade$^{-1}$ (Fu et al. 2006; Hudson et al. 2006; Hu and Fu 2007; Archer and Caldeira 2008; Seidel et al. 2008; Nguyen et al. 2013; D’Agostino and Lionello 2017). The wide range of observed trends is mostly due to the wide range of metrics used to define the tropical belt (e.g., tropopause height, subtropical jet, streamfunction, precipitation and evaporation, and outgoing longwave radiation). Furthermore, except for changes in mean meridional mass streamfunction, all other metrics were found to produce trends that are statistically insignificant (Davis and Rosenlof 2012). More recently, it has been shown that many of the metrics that were originally used do not correlate with the streamfunction expansion (Solomon et al. 2016; Davis and Birner 2017) and hence are not representative of a widening Hadley cell.

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One of the main difficulties in observing a significant trend is the presence of both the internal variability of the climate system (e.g., Pacific decadal oscillation), which obscures the ongoing anthropogenic forced signal (Allen et al. 2014; Quan et al. 2014; Lucas and Nguyen 2015; Allen and Kovilakam 2017; Mantsis et al. 2017), and the multiplicity of forcing agents (Allen et al. 2012, 2014; Waugh et al. 2015) that are able to affect the width of the tropical circulation. For example, while enhanced greenhouse gas emissions were found to expand the tropical circulation in recent decades (e.g., Bony et al. 2013; Hu et al. 2013), the buildup of anthropogenic aerosols in the Northern Hemisphere were found to contract the circulation, and thus offset the expansion signal (Allen and Ajoku 2016). In the Southern Hemisphere, ozone depletion was found to be the main forcing of tropical expansion in the second half of the twentieth century (Son et al. 2009; Polvani et al. 2011b), and the ongoing recovery of ozone hole will therefore cancel the effect of greenhouse gases throughout the first half of the twenty-first century (Polvani et al. 2011a; Barnes et al. 2014).

The unabated emissions of greenhouse gases into the atmosphere is one of the main contributors to the Hadley cell expansion over the twenty-first century. Therefore, it is important to understand what controls tropical expansion under increased CO$_2$ concentrations. To avoid complications from the presence of other forcings, we will here confine our attention to the 4-fold quadrupling of CO$_2$ is instantaneous, it conveniently with unambiguous attribution. Furthermore, because the quadrupling of CO$_2$ is instantaneous, it conveniently separates different components of the climate system, which respond with different time scales: this scenario, therefore, allows us to learn a great deal regarding the mechanism of tropical expansion.

Over the last decades, several mechanisms have been proposed to explain the tropical expansion, which we now briefly review. Enhanced CO$_2$ concentrations warm both the troposphere and sea surface, while cooling the stratosphere. While several studies have claimed that changes in surface temperature $T_s$ are mostly responsible for widening the tropical belt (e.g., Frierson et al. 2007; Adam et al. 2014; Son et al. 2018), others have argued that it is the atmospheric warming that matters the most (e.g., Lu et al. 2009; Bony et al. 2013). Part of this confusion stems from the fact that different time periods were chosen in different studies, as well as the different ozone forcings (Waugh et al. 2015).

Other studies (e.g., Lu et al. 2008; Ceppi and Hartmann 2013) have suggested that changes in eddy phase speed may drive the expansion of the Hadley cell under global warming. The rise of the tropopause height in warmer climates (Thuburn and Craig 1997; Santer et al. 2003) may affect the meridional temperature gradient aloft (Lorenz and DeWeaver 2007), and thus the eddy phase speed (Wittman et al. 2007). Because the mean zonal winds decrease from the midlatitudes equatorward, higher eddy phase speeds may push the subtropical critical latitude poleward (Chen and Held 2007). The dissipation of eddies at critical latitudes implies a poleward shift of the latitude of zero eddy momentum flux divergence. Since the latter balances the mean flow in steady state (e.g., Vallis 2006), this would push the latitudinal extent of the Hadley cell poleward.

While the above studies have proposed possible mechanisms, none of them constitutes an actual theory for why the tropics are expanding with increased CO$_2$ concentrations. To date, there have been two main theories that provide scalings for the Hadley cell width. The first, proposed by Held and Hou (1980), posits that the Hadley cell is a thermally closed circulation, and that poleward moving air at its upper branch conserves angular momentum. In this theory, the Hadley cell width ($\phi_{HH}$) scales as follows:

$$\phi_{HH} \propto \left( \frac{g H_0 \Delta \theta}{\Omega a^2 \theta_0} \right)^{1/2},$$

where $g$ is gravity, $\Omega$ is Earth’s rotation rate, $a$ is Earth’s radius, $\theta_0$ is a reference potential temperature, $\Delta \theta$ is the equator-to-pole potential temperature difference in radiative equilibrium, and $H_0$ is the tropical tropopause height.

The second theory, proposed by Held (2000), posits that the Hadley cell edge corresponds to the latitude where the vertical shear of the angular momentum–conserving flow becomes baroclinically unstable, using the two-layer model criterion of Phillips (1954). In this theory the Hadley cell width ($\phi_{H00}$) scales as follows:

$$\phi_{H00} \propto \left( \frac{N^2}{\Omega a} \right)^{1/2},$$

where $N^2 = (g/\theta)(\partial \theta/\partial z)$ is the subtropical static stability and $H_0$ is the subtropical tropopause height.

Using idealized GCMs (e.g., Walker and Schneider 2006; Frierson et al. 2007; Levine and Schneider 2015), full atmospheric-only GCMs (e.g., Frierson et al. 2007), and coupled GCMs (e.g., Lu et al. 2007, 2008), it has been shown that, under global warming, the latitudinal extent of the Hadley cell follows the Held (2000) scaling. The tropical expansion was found to correlate with both the increase in subtropical static stability (e.g., Lu et al. 2008; Son et al. 2018) and the subtropical tropopause height, which pushes the baroclinic zone poleward (e.g., Lu et al. 2007).
In all above studies, the long-term response of the tropical expansion was studied (e.g., under statistically steady-state or slowly varying forcing) by correlating it with the above suggested components (e.g., $T_s$, tropopause height, static stability, and eddy phase speed). Since correlation alone need not imply a casual relation, especially in a highly coupled system, the relative importance of the different components that may drive the widening of the tropical belt could not be estimated.

Studying the evolution of the atmospheric circulation to abrupt change in CO$_2$, on the other hand, will allow us to elucidate the relative importance of the different components in affecting the steady-state response. For instance, Wu et al. (2012) showed that under abrupt double CO$_2$ forcing the stratosphere is first to respond, leading to changes in tropospheric eddies that alter the zonal mean circulation. While eddies were found to play an important role in these simulations, changes in eddy phase speed were found not to correlate with circulation changes (Wu et al. 2013; Staten et al. 2014). Using abrupt 4 $\times$ CO$_2$ forcing in CMIP5 models, Grise and Polvani (2017) showed that while the Hadley cell expands and reaches 90% of its final location after the first few years, global-mean $T_s$ monotonically increases reaching 90% of its final value only after several decades, implying that global-mean $T_s$ is not the main driver of the Hadley cell expansion.

While the Hadley cell edge and global-mean $T_s$ show different response times to quadrupling CO$_2$, the poleward expansion of the dry zone (the latitude where precipitation equal evaporation) was found to follow the slower changes in global-mean $T_s$, rather than the rapid changes in the Hadley cell edge (Grise and Polvani 2017). This implies that in spite of the high correlation between the edge of the Hadley cell (streamfunction expansion) and the edge of the dry zone (Quan et al. 2014; Solomon et al. 2016), different mechanisms control them. At steady state, changes in precipitation and evaporation are in balance with moisture flux divergence changes. Both long-term (Seager et al. 2010) and transient (Grise and Polvani 2017) responses point to the importance of the time-mean circulation and transient eddies in shifting the edge of the dry zone poleward. However, the effects of stationary eddy moisture flux, as well as the zonal mean flow, remain unclear.

Here, following Grise and Polvani (2017), the evolution of the Hadley cell expansion, the dry zone’s edge, and the above suggested mechanisms, are examined under abrupt 4 $\times$ CO$_2$ forcing in CMIP5 models. The different time scales of the response of the relevant components will allow us to determine the relative importance of the different proposed mechanisms in widening the tropics. Since the steady-state result in these simulations stems from the transient response to the abrupt 4 $\times$ CO$_2$ forcing, studying the transient response is a key for understanding the equilibrium widening of the Hadley. Because all mechanisms discussed above assume zonal symmetry, and because most of the continents reside in the Northern Hemisphere, the tropical response of increased CO$_2$ is here studied only in the Southern Hemisphere, which provides the most appropriate conditions for validating the proposed mechanisms.

The paper is structured as follows. Section 2 describes the different simulations used to study the widening of the tropics, and the different metrics in question. The roles of the different atmospheric components and the two scaling theories on the Hadley cell expansion are examined in sections 3 and 4, respectively. The analysis of the mechanism behind the poleward shift of the dry zone edge is conducted in section 5. Finally, section 6 summarizes the results.

2. Methods

To study the evolution of the different components that affect the Hadley cell width, and the edge of the dry zone, three sets of the “r1i1p1” experiments from the CMIP5 multimodel ensemble (Taylor et al. 2012) are analyzed: first, 200 years of preindustrial control run with a constant 1850 forcing; second, 150 years of abrupt (set as initial conditions) 4 $\times$ CO$_2$ forcing (relative to preindustrial values); and third, 140 years of 1% yr$^{-1}$ increased CO$_2$ run (starting from preindustrial values). For each experiment we analyze 23 coupled models, which are listed in Table 1.

Two key metrics are used for estimating the width of the tropical belt, and are illustrated in Figs. 1a and 1b, respectively, from the preindustrial control run of the GFDL-ESM2G (Dunne et al. 2012):

(i) The edge of the Hadley cell $\phi_{500}$ is calculated as the latitude where the meridional mass streamfunction (colors in Fig. 1) $\Psi$, defined by

$$\Psi = \frac{2\pi a \cos \phi}{g} \int_0^P [\nu] dp,$$

first changes sign poleward of its maximum value at 500 hPa (green cross in Fig. 1a); $\phi$ is latitude, $P$ is pressure, $\nu$ is the meridional velocity, the overbar represents the annual mean, and the square brackets represent the zonal mean.

(ii) The edge of the dry zone $\phi_{P-E}$ (black cross in Fig. 1b) is evaluated as the latitude where zonal- and annual-mean precipitation $P$; (red line in Fig. 1b) equals evaporation $E$ (blue line in Fig. 1b) poleward of the latitude of maximum $\Psi$. 
The above metrics are calculated after a 0.1° latitudinal cubic interpolation (which preserves the metrics’ latitudinal structure) of $C$ and $P^2E$.

In addition, to properly analyze the different mechanisms of tropical expansion discussed in section 1, two other quantities, related to eddy components, are calculated as follows:

(i) The latitude of maximum eddy momentum flux $u_{\text{max}}$ (orange cross in Fig. 1a), where $u$ is zonal wind, and crosses represent deviation from both zonal and monthly means.

(ii) The phase speed of maximum eddy momentum flux $c_{\text{phase}}$ (black cross in Fig. 1c). The eddy momentum flux phase speed is calculated following Randel and Held (1991) at 250 hPa, in segments of consecutive 60 days averaged over each year.

The above eddy quantities are calculated after a 0.1° latitudinal quadratic interpolation (to capture the maximum value of eddy momentum fluxes). Only four models in Table 1 (GFDL-ESM2G, GFDL-ESM2M, MIROC5, and IPSL-CM5B-LR) have available daily data, which are required for calculating eddy fluxes. Thus, the eddy analysis in this study is performed for only these four models.

Finally, $\phi_{100}$ and $\phi_{100}$ are computed as follows:

(i) The different components in Eq. (1) [the Held and Hou (1980) scaling] are defined as follows; $\theta_0$ as the Southern Hemisphere tropospheric-mean (at 500 hPa) potential temperature, $\Delta \phi$ as the equator-to-pole tropospheric-mean potential temperature difference (the difference between the tropical average, $0°S–20°S$, and the high-latitude average, $75°–90°S$) and the tropical tropopause height $H_t$ as the lowest level where the vertical temperature gradient crosses the 2 K km$^{-1}$ value, in accordance with the World Meteorological Organization (WMO).

(ii) The different components in Eq. (2) [the Held (2000) scaling] are defined as follows: The subtropical static stability $N^2$ is averaged between 400–850 hPa, and the subtropical tropopause height $H_e$ follows the above WMO definition. Vertically averaging the static stability up to a constant level, and not up to
of the Hadley cell in order to properly examine the Held (2000) scaling, which includes extratropical quantitates. To estimate the response time of each of the above metrics and components to the abrupt $4 \times CO_2$ forcing, we fit the following function, $y = c_o (1 - e^{-\tau t})$, to their time evolution, where the parameters $c_o$ and $\tau$ are numerically obtained using a least squares algorithm. The response time is then defined by calculating the three $e$-folding time ($3\tau$). The later provides an estimate of the time when the response of each component reaches ~95% of its steady-state value. Statistically steady-state values are defined as the average over the last 50 years of the 150-yr-long $4 \times CO_2$ simulations.

3. Assessing the importance of surface temperature and eddy phase speed

As discussed in section 1, two atmospheric components have been proposed to exert a strong influence on the widening of the Hadley cell: 1) the surface temperature $T_s$ and 2) the eddy phase speed $c_{\text{phase}}$. To examine each of these components, their evolution, under the abrupt $4 \times CO_2$ forcing, is examined and compared to the poleward shift of $\phi_{500}$.

a. Surface temperature changes

Several previous studies argued for the importance of $T_s$ in affecting the Hadley cell width. Both the global-mean $T_s$ (e.g., Staten et al. 2012; Quan et al. 2014; Son et al. 2018) and the meridional $T_s$ gradient (e.g., Frierson et al. 2007; Adam et al. 2014) have been suggested to affect the Hadley cell width. However, as these suggestions are based on correlations in the long-term response across many models, no causality arguments can be made. To examine whether changes in $T_s$ vary together with the widening of the Hadley circulation, the evolutions of $\phi_{500}$ (black), the Southern Hemisphere mean $T_s$ (blue), and the meridional $T_s$ gradient ($\Delta \phi T_s$; purple; tropical minus subtropical temperature difference) are plotted in Fig. 2.

The evolution of $\phi_{500}$ to the abrupt forcing, does not coincide with the response of $T_s$ (Fig. 2a). As discussed in Grise and Polvani (2017), while the response time of the Hadley cell width is ~7 years, $T_s$ monotonically increases through the entire simulation, implying that $T_s$ is not the main driver of the Hadley cell expansion: through most of the simulation $T_s$ increases while the Hadley cell width does not. Nevertheless, because the atmospheric circulation is driven by vertical temperature gradients (e.g., Charney 1947; Eady 1949), the importance of $T_s$ reported in previous studies might be related to changes in the vertical temperature profile, and not changes in $T_s$ alone, as further discussed in section 4.

Changes in $\Delta \phi T_s$ also do not coincide with the response of the Hadley cell width to the abrupt forcing

The multimodel mean of each of the above quantities is calculated by first taking its annual and zonal average for each model separately, and then averaging across all models. All subtropical quantities are calculated between 35° and 45°S, and tropical quantities between 0° and 20°S. The subtropical latitudes are defined outside of the Hadley cell in order to properly examine the Held (2000) scaling, which includes extratropical quantitates.

**Fig. 1.** Metrics for evaluating tropical expansion. (a) Zonal- and time-mean meridional mass streamfunction (kg s$^{-1}$; colors) and eddy momentum flux ($[u'v'] \cos \phi$; m$^2$ s$^{-1}$; black contours) as a function of pressure and latitude. The green and orange crosses respectively represent the latitude where the meridional mass streamfunction first changes sign poleward of its maximum value at 500 hPa $\phi_{500}$ and the latitude of maximum eddy momentum flux $\phi_{egm}$. The eddy momentum flux contours correspond to $[-39, -34, -30, -25, 0.5, 1, 2, 4] \text{m}^2 \text{s}^{-1}$. (b) The black cross represents the latitude where precipitation equals evaporation at 500 hPa $P_{500}$ (black), the Southern Hemisphere mean $T_s$ (blue), and the meridional $T_s$ gradient ($\Delta \phi T_s$; purple; tropical minus subtropical temperature difference) are plotted in Fig. 2. The evolution of $\phi_{500}$ to the abrupt forcing, does not coincide with the response of $T_s$ (Fig. 2a). As discussed in Grise and Polvani (2017), while the response time of the Hadley cell width is ~7 years, $T_s$ monotonically increases through the entire simulation, implying that $T_s$ is not the main driver of the Hadley cell expansion: through most of the simulation $T_s$ increases while the Hadley cell width does not. Nevertheless, because the atmospheric circulation is driven by vertical temperature gradients (e.g., Charney 1947; Eady 1949), the importance of $T_s$ reported in previous studies might be related to changes in the vertical temperature profile, and not changes in $T_s$ alone, as further discussed in section 4.

Changes in $\Delta \phi T_s$ also do not coincide with the response of the Hadley cell width to the abrupt forcing
In contrast to the monotonic widening of the Hadley cell, $DhTs$ shows a nonmonotonic behavior, reaching a maximum value after approximately 10 years and then actually decreasing through the rest of the simulation. Thus, while there is a direct relation between $DhTs$ and $f_{C500}$ during the fast response (up to 10 years), in the slow response they show an inverse relation. This might explain the direct relation between $DhTs$ and $f_{C500}$ found in an atmosphere-only model (Frierson et al. 2007), where slow-response processes, associated with the ocean circulation, are absent, in contrast to the inverse relation found in reanalysis data (Adam et al. 2014).

### b. Eddy phase speed

Several studies (e.g., Lu et al. 2008; Ceppi and Hartmann 2013) have suggested that the increased eddy phase speed found under global warming (Chen and Held 2007) results in the poleward shift of the Hadley cell edge. Larger eddy phase speeds push the critical latitude poleward, so that eddies dissipate at higher latitudes. The associated shift of the latitude of zero eddy momentum flux divergence (i.e., the latitude where the mean meridional flow is zero) results in the widening of the Hadley cell. Again, since the above mechanism was examined by correlating the long-term response fields, the cause and effect relation remains unclear.

To determine if indeed the eddy phase speed causes the changes in latitude of maximum eddy momentum flux (which corresponds to the latitude of zero eddy momentum flux divergence) and the poleward shift of the Hadley cell edge, the evolution of $f_{C500}$ (black), $f_{[u'y']}$ (red), and $c_{phase}$ (blue) are plotted in Fig. 3. One can see that $f_{[u'y']}$ shifts poleward together with $f_{C500}$, and they are highly correlated ($r = 0.68$) through the entire run (Fig. 4a) [as shown by Ceppi and Hartmann (2013) in reanalyses]. As in Chen and Held (2007), we find that with increased levels of CO$_2$ $c_{phase}$ is larger compared to preindustrial values; however, note that $c_{phase}$ shows an instantaneous increase, while $f_{C500}$ and $f_{[u'y']}$ show a gradual increase in the first years followed by a more moderate change through the rest of the simulation. As a result, changes in $c_{phase}$ have a low correlation with the poleward shift of $f_{C500}$ and $f_{[u'y']}$ (Fig. 4b; $r = 0.09$). This low correlation may stem from the increase in mean zonal wind, which acts to push the critical latitude equatorward (e.g., Lu et al. 2014). This result confirms the one reported in previous studies (Wu et al. 2013; Staten et al. 2014), which also found low correlation in the time evolution of the eddy phase speed and the mean meridional circulation. Thus, the high correlation between the poleward shift of $f_{C500}$ and $f_{[u'y']}$ (Fig. 4a) is not due to changes in the eddies’ dissipation region inside the tropics (critical latitude, as shown in Figs. 3 and 4), but rather is due to changes in
the eddies’ source region (extratropical baroclinicity), as further discussed in the next section.

4. The relative importance of the Hadley cell width scalings and their components

Having shown that $T_s$ and $c_{\text{phase}}$ occur on different time scales than the Hadley cell expansion, we next turn to the theories, and analyze the Held and Hou (1980) and Held (2000) scalings for the Hadley cell width. Studying the evolution of these scaling, along with their different components, will provide further insights into the physical mechanism behind the Hadley cell expansion at steady state (note that the balance at the upper branch of the Hadley cell between the annual mean meridional advection of zonal momentum and Coriolis acceleration holds through the entire simulation).

a. Comparing Hadley cell width scalings

Previous studies found that while changes in $\phi_{\text{HH}}$ [Eq. (2)] along with its components (subtropical static stability and tropopause height) are highly correlated with changes in $\phi_{\text{F500}}$ (Frierson et al. 2007; Lu et al. 2007, 2008; Son et al. 2018), changes in the tropical tropopause height in $\phi_{\text{HH}}$ [Eq. (1)] have a low correlation with changes in $\phi_{\text{F500}}$ (Lu et al. 2007). To properly examine whether indeed $\phi_{\text{F500}}$ better predicts $\phi_{\text{F500}}$ than $\phi_{\text{HH}}$, it is crucial to first compare the entire scaling [Eqs. (1) and (2)], when all components are taken into account. As shown in Fig. 5, across all models, the zonal mean statistically steady-state poleward shifts of $\phi_{\text{F500}}$ indeed shows a better correlation with $\phi_{\text{F500}}$ ($r = 0.88$) than $\phi_{\text{HH}}$ ($r = 0.24$). Because $\phi_{\text{F500}}$ accounts for the role of eddies in affecting the mean circulation in the subtropics, its high correlation with $\phi_{\text{F500}}$ is not surprising, given the high correlation between $\phi_{\text{F500}}$ and $\phi_{\text{uu}}$ shown above in Figs. 3 and 4a.

As the poleward shift of $\phi_{\text{F500}}$ better correlates with the poleward shift of $\phi_{\text{F500}}$, the transient evolution of the latter is further examined in order to obtain a deeper understanding of the mechanisms behind the transient and steady-state widening of the Hadley cell. As one can see in Fig. 6a, $\phi_{\text{F500}}$ shows a similar evolution under the abrupt forcing as $\phi_{\text{F500}}$ (cf. red and black lines). It shows a rapid response in the first years that accounts for most of the expansion, followed by a more moderate poleward shift. In spite of the similar overall behavior, the evolution of $\phi_{\text{F500}}$ also deviates from the behavior of $\phi_{\text{F500}}$ (note that the overlapping of the confidence interval of $\phi_{\text{F500}}$ and $\phi_{\text{HH}}$ starts only after 7 years, when $\phi_{\text{F500}}$ has already reached 95% of its steady-state value). In Fig. 6b, we zoom in on the first 40 years of evolution: whereas the response time of $\phi_{\text{F500}}$ is 7 years (black line), the response time of $\phi_{\text{HH}}$ is ~20 years (red line). This slower increase of $\phi_{\text{F500}}$ relative to $\phi_{\text{F500}}$ indicates that different physical mechanisms control them, which we proceed to elucidate next.

b. The roles of static stability and tropopause height

Previous studies found that both components of the Held (2000) scaling [the tropopause height and static stability; Eq. (2)] correlate with $\phi_{\text{F500}}$ (Lu et al. 2007, 2008); this makes it difficult to quantify their relative
importance in shifting $\phi_{500}$ poleward. However, studying their evolution allows us to disentangle their relative roles, and sheds new light on the mechanism behind the transient and steady-state tropical expansion. In particular, we wish to understand which of these components is responsible for the rapid poleward shift that results in the similar evolution as $\phi_{C500}$, and which for the slower increase that causes $\phi_{H00}$ to deviates from $\phi_{C500}$.

Dividing Eq. (2) by its preindustrial value and taking the logarithm enables us to disentangle the components in the Held (2000) scaling, the subtropical tropopause height $H_e$ and static stability $N$:

$$\log \left( \frac{\phi_{H00}}{\phi_{H00,PI}} \right) = 0.5 \log \left( \frac{N}{N_{PI}} \right) + 0.5 \log \left( \frac{H_e}{H_{e,PI}} \right).$$

The evolution of the different terms in Eq. (3) is plotted in Fig. 7a. By construction, the sum of the tropopause height term (blue line) and static stability term (green line) equals the response of $\phi_{H00}$ (red line). In the first ~5 years, changes in static stability and tropopause height contribute equally to the rapid poleward shift of $\phi_{H00}$. While the response time of the static stability is ~10 years (similar to poleward shift of the Hadley cell), the response time of tropopause height is ~25 years (similar to poleward shift of $\phi_{H00}$). Thus, after the fast response of the first years, the slower monotonic increase of the tropopause height alone contributes to the increase of $\phi_{H00}$. This increase results in the deviation between the evolution of $\phi_{500}$ and $\phi_{H00}$ (cf. black and red lines in Fig. 6a).

What changes in the temperature field are responsible for the rapid and slow responses of $\phi_{H00}$? Both an increase in tropospheric temperature, relative to the lower stratosphere, and a decrease in temperature lapse rate increase the tropopause height [e.g., Thuburn and Craig 2000; see also Eq. (9) in Vallis et al. (2015)]. In addition, a decrease in temperature lapse rate stabilizes the atmosphere, thus increasing the static stability. Plotting the evolution of subtropical temperature at various levels shows that the upper troposphere (400–600 hPa; blue line in Fig. 7b) warms faster than the lower troposphere (600–850 hPa; red line in Fig. 7b) in the first ~15 years, after which they both warm at approximately the same rate. As a consequence, the response time of the tropospheric vertical temperature difference (green line in Fig. 7b), calculated between the upper and lower troposphere, is ~14 years. This explains the rapid response of both the static stability (green line in Fig. 7a) and tropopause height (blue line in Fig. 7a). The continued monotonic increase in tropospheric temperature (relative to the slower increase in lower-stratospheric temperature, black line) further increases the tropopause height (blue line in Fig. 7a), leading $\phi_{H00}$ to a response time of ~20 years, and thus to deviate from $\phi_{500}$.

To more clearly illustrate the effect of the tropopause height on $\phi_{H00}$, the latter is recalculated using constant preindustrial tropopause height values $\phi_{H00|\eta_{pre}}$ (green line in Fig. 6), so that only changes in static stability affect the widening of the Hadley cell. One can see that $\phi_{H00|\eta_{pre}}$ better follows the poleward shift of $\phi_{500}$ (cf. black and green lines in Fig. 6). Unlike $\phi_{H00}$, which has a
response time of ~20 years, $\phi_{H00}$, has a response time of ~10 years, similar to the response time of $\phi_{C500}$. This shows the importance of the static stability, and not the tropopause height, in shifting the Hadley cell poleward.

c. Changes in subtropical baroclinicity

How come the tropopause height causes the response of $\phi_{H00}$ to deviate from the response of $\phi_{C500}$? In the Held (2000) theory, an increase in tropopause height should increase the critical shear necessary for baroclinic instability, according to the two-layer model (Phillips 1954), and thus push poleward the latitude where it is equal to the angular momentum conserving shear. One possibility is that the subtropical baroclinicity in the atmosphere is insensitive to the tropopause height.

To examine this possibility, a linear normal-mode instability analysis is conducted following Smith (2007), which was found to capture the baroclinic properties of eddies in both GCMs and reanalysis data (Jansen and Ferrari 2012; Chemke and Kaspi 2015, 2016a,b; Chemke et al. 2016). In this analysis the eigenvalue problem of the linearized quasigeostrophic potential vorticity (QGPV) equation is numerically solved at each latitude, using each model’s mean zonal wind, static stability, and tropopause height. The tropopause height sets the upper boundary of the problem. The resulting growth rates of the eddies are used as a measure for baroclinicity. First, the subtropical growth rate (the mean of the growth rate over all wavenumbers averaged over the subtropics) is calculated for each model using the zonal- and annual-mean fields of the preindustrial and of the statistically steady-state abrupt $4 \times CO_2$ runs (last 50 years). Then, the growth rate of the statistically steady-state abrupt $4 \times CO_2$ run is recalculated, but with the tropopause height in preindustrial values. The difference in the growth rate between the $4 \times CO_2$ and preindustrial runs is plotted against the difference between the $4 \times CO_2$ run using the preindustrial tropopause height and preindustrial run (Fig. 8a). First, most of the models (17) show a decrease in the growth rate relative to preindustrial values (y axis in Fig. 8), with a multimodel mean value of $-1.1 \times 10^{-7} \text{s}^{-1}$ (red
This indicates that the baroclinicity decreases under the abrupt $4 \times CO_2$ forcing. Second, if the tropopause height affects the growth rate, the dots in Fig. 8a should not fall on the 1:1 line (black line). However, in most models changes in growth rate are similar (blue dots on the 1:1 line; $R^2 = 0.93$) whether or not the tropopause height is allowed to increase from preindustrial values. Thus, changes in tropopause height in the $4 \times CO_2$ simulations have a minor effect on baroclinicity.

In contrast, static stability changes have a large effect: the difference in the growth rate between the $4 \times CO_2$ and preindustrial runs is plotted against the difference between the $4 \times CO_2$ and preindustrial runs using the $4 \times CO_2$ static stability values (Fig. 8b). Using fixed $4 \times CO_2$ static stability values rather than preindustrial values, as was done for the tropopause height, precludes including stratospheric static stability values for the $4 \times CO_2$ calculation, when the tropopause height increases. Keeping the static stability at abrupt $4 \times CO_2$ values yields changes in baroclinicity that are different than when allowing the static stability to change (dots do not fall on the 1:1 line in Fig. 8b; $R^2 = -0.05$). In addition, keeping the static stability constant results in minor changes in baroclinicity in most models (dots fall close to the x-axis zero line, with multimodel mean value of $2.8 \times 10^{-8}$ m$^{-1}$s$^{-1}$; red dot). Thus, the increase in subtropical static stability explains most of the decrease in subtropical baroclinicity as the climate becomes warmer.

In summary, the rapid increase of static stability (i.e., rapid decrease of the subtropical lapse rate) in the first ~10 years decreases the baroclinicity, and explains most of the poleward shift of the Hadley cell and of the eddy momentum flux. The tropopause height, on the other hand, seems to have a minor effect on baroclinicity, and thus shows different response time than the Hadley cell width. This result is in agreement with the results of Son et al. (2018), who found high correlation between changes in subtropical static stability and Hadley cell shift in past and projected climates.

d. Applicability to more realistic scenarios

The above analyses are conducted using the abrupt $4 \times CO_2$ forcing. The abrupt change in CO$_2$ helps disentangle the different response times of the different components that affect the Hadley cell expansion, and thus provides better understanding of the controlling mechanisms. In a more realistic scenario, however, CO$_2$ increases continuously. To examine the above results in such a realistic scenario, the 1% CO$_2$ increase per year scenario is now investigated. Figure 9a is similar to Fig. 6a, only showing the evolution under the 1% CO$_2$ increase per year scenario, relative to preindustrial values, of the poleward shift of $\phi_{4500}$ (black), $\phi_{1800}$ [red; Eq. (2)], and $\phi_{1800}\mid_{\Delta CO_2}$ (green). Unlike for the abrupt CO$_2$ forcing, $\phi_{4500}$ does not reach a statistically steady state value, but rather monotonically shifts poleward, due to the gradual increase in CO$_2$. Toward the end of the simulation, $\phi_{1800}$ shifts more poleward than $\phi_{4500}$ and...
Note that in spite of the overlapping confidence intervals of \( \phi_{500} \) and \( \phi_{100} \) there is a clear separation of the multimodel ensemble means that increases with time. Similar to the abrupt \( 4 \times CO_2 \) experiment, this shows that the disagreement of \( \phi_{100} \) and \( \phi_{500} \) stems from tropopause height changes.

To see the different responses of static stability and tropopause height in the 1% CO2 increase per year scenario, the different terms of Eq. (3) are plotted in Fig. 9b. By construction, the sum of the change in static stability (green) and tropopause height (blue) equals the change in \( \phi_{100} \) (red). Similar to the abrupt \( 4 \times CO_2 \) experiment, at the beginning of the run (up to year \( \sim 70 \)) both static stability and tropopause height contribute equally to the poleward expansion of \( \phi_{100} \). Then, the tropopause height increases faster than the static stability (as it is also affected by the warming of the troposphere relative to the lower stratosphere), which further pushes poleward \( \phi_{100} \), causing it to deviate from the poleward shift of \( \phi_{500} \). This again shows the importance of the static stability, and not the tropopause height, in shifting the Hadley cell edge poleward.

5. Edge of the dry zone

Another important metric for the edge of the tropics is the latitude where precipitation equals evaporation \( \phi_{P-E} \) (edge of the dry zone). Unlike the Hadley cell edge, this metric holds information regarding the hydrological cycle, as it separates regions of net evaporation (the dry zone) from regions of net precipitation (midlatitudes). Thus, this metric is important in assessing the effects of climate change on the hydrological cycle. Recently, Grise and Polvani (2017) and Seviour et al. (2018) showed that the evolution of \( \phi_{500} \) deviates from the poleward shift of \( \phi_{P-E} \). While several studies found the response of these two metrics to be highly correlated across a large set of models (e.g., Lu et al. 2007; Polvani et al. 2011b; Quan et al. 2014; Solomon et al. 2016), their different time evolutions suggest that different physical mechanisms control them.

The evolution of \( \phi_{500} \) (black line) and \( \phi_{P-E} \) (blue line) to the abrupt \( 4 \times CO_2 \) forcing are plotted in Fig. 10a. In the first years \( \phi_{P-E} \) shows a rapid poleward shift as \( \phi_{500} \), after which it shows a slower poleward shift. As discussed in Grise and Polvani (2017), whereas the response time of \( \phi_{500} \) is \( \sim 7 \) years, the response time of \( \phi_{P-E} \) is \( \sim 20 \) years. By studying the zonal- and annual-mean vertically integrated moisture budget, Grise and Polvani (2017) showed that the poleward shift of the Hadley cell edge (mean meridional velocity) and changes in transient eddy moisture flux divergence, contribute, respectively, to the fast and slow responses of the dry zone edge. Here, we further elaborate on the results of Grise and Polvani (2017) and analyze the subtropical moisture budget by including the effects of stationary eddies and zonal mean flow.

Following Trenberth and Guillemot (1995) and Seager et al. (2010), changes of the zonal- and annual-mean vertically integrated subtropical moisture budget, can be written as follows:

\[
\delta [E - P] = \frac{1}{ga} \frac{\partial}{\partial \phi} \frac{\partial}{\partial \phi} \left( \delta [\mathbf{q}] \cos \phi + \delta [\mathbf{u} \cdot \mathbf{q}] \cos \phi \, dp \right),
\]

where \( \delta \) represents the transient response, \( E \) and \( P \) are the evaporation from the surface and precipitation, respectively, and \( q \) is specific humidity. While the first term on the right-hand side accounts for changes in mean moisture flux, the second term accounts for changes in eddy moisture flux (deviation from both zonal and monthly means).
Plotting the evolution of the contribution of the right-hand side terms in Eq. (4) to the shift of \( \Phi_{P-E} \) shows that most of the response of \( \Phi_{P-E} \) (blue line in Fig. 10b) is associated with changes \([\bar{y}\delta q]]\) alone (red line in Fig. 10b). The meridional divergence of \([v^+ q^+]\) (yellow line in Fig. 10b), on the other hand, shows minor changes to quadrupling CO2, and thus does not contribute to the increase in \( \Phi_{P-E} \). The response of \([\bar{y}\delta q]]\) can be further decomposed as follows:

\[
\delta[\bar{y}\delta q] \approx [\bar{y}]\delta[q] + [\bar{y}\delta[q]],
\]

where the first term on the right-hand side accounts for changes in circulation and the second term accounts for changes in moisture itself (Held and Soden 2006; Seager et al. 2010).

Plotting the evolution of the contribution of the meridional divergence of the right-hand side of Eq. (5) to the shift of \( \Phi_{P-E} \) shows that changes in circulation (red
line in Fig. 10c) and in moisture (yellow line in Fig. 10c) have different time evolutions, and thus play different roles in shifting $\phi_{P-E}$. While changes in circulation have a rapid response, with a response time of $\sim 4$ years (as discussed in section 3), changes in moisture are slower and constantly increasing, with a response time of $\sim 100$ years. This suggests that the initial fast poleward shift of $\phi_{P-E}$ is driven by the rapid expansion of the Hadley cell (which is driven by the fast response of the subtropical lapse rate, as discussed in section 4b), and that its slower monotonic shift is driven by changes in moisture content. At the end of the simulation, $\sim 75\%$ of the poleward shift of the dry zone edge is due to the widening of the Hadley cell, and $\sim 25\%$ due to the increase in subtropical moisture. This is similar to the results of Scheff and Frierson (2012), who found that most of the drying in subtropical regions stems from the poleward shift of the mean meridional circulation, and not from changes in moisture content.

To better understand the minor contribution of eddy moisture flux to the poleward shift of the dry zone edge (yellow line in Fig. 10b), the response of the eddy moisture flux is further decomposed as follows:

$$\delta[v^* q^*] = \delta[v^* q] + \delta[v^* q^*],$$  

(6)

where the first and second terms represent changes in transient (purple line in Fig. 10d) and stationary (green line in Fig. 10d) eddies, respectively. As discussed in Grise and Polvani (2017) and Seager et al. (2010), the divergence of transient eddy moisture flux (purple) acts to push $\phi_{P-E}$ poleward. Here we show that the divergence of stationary eddy moisture flux (green), on the other hand, acts to push $\phi_{P-E}$ equatorward. The cancelation of the transient and stationary eddy moisture fluxes (as was found under regional warming in the Northern Hemisphere: Shaw and Voigt 2016) results in a minor contribution of the eddy fields to the poleward shift of $\phi_{P-E}$.

What causes the different behavior of transient and stationary eddy moisture fluxes? Figure 11 shows the vertically integrated, multimodel, zonal-mean latitudinal structure of the transient (Figs. 11a,c) and stationary (Figs. 11b,d) eddy moisture flux in the preindustrial run (red lines) and the statistically steady-state $4 \times CO_2$ run (blue lines). Both transient and stationary eddy moisture flux are negative, as they act to transfer moisture from the tropics toward the South Pole (Figs. 11a,b). While the transient eddy moisture flux reaches maximum value at midlatitudes ($\sim 40^\circ$S), the stationary eddy moisture flux reaches maximum value in the lower subtropics ($\sim 25^\circ$S), where most of the Southern Hemisphere continents reside. Because of their different latitudinal structures, the transient eddy moisture flux tend to diverge moisture in the subtropics (positive values in Fig. 11c), while the stationary eddy moisture flux tends to converge moisture in the subtropics (negative values in Fig. 11d). Under increased CO$_2$ concentrations, both the transient and stationary eddy moisture fluxes shift poleward (cf. red and blue lines in Figs. 11a and 11b). Thus, a poleward shift of the transient eddy moisture flux increases its divergence in the subtropics, while a poleward shift of the stationary eddy moisture flux increases its convergence in the subtropics (cf. red and blue lines in Figs. 11c and 11d), resulting in opposite tendencies, as shown in Fig. 10d, that lead to the total cancelation in the subtropics.

6. Summary

The projected widening of the tropical circulation with increased greenhouse gases (e.g., Lu et al. 2007; Vallis et al. 2015) and the associated shift of the dry zone edge have important societal impacts. Several mechanisms have been suggested to explain this poleward shift, but the coupling among the different atmospheric components did not allow for a quantification of their relative importance in expanding the tropics. Here, using the abrupt quadruple-CO$_2$ scenario runs from the CMIP5, we have been able to separate the relevant mechanisms of tropical expansion. The different response times of the various mechanisms has enabled us to disentangle the different components in question and elucidate their relative importance.

Upon abrupt $4 \times CO_2$, the Hadley cell rapidly expands poleward, with a response time of $\sim 7$ years (Grise and Polvani 2017). Confirming several earlier studies (e.g., Frierson et al. 2007; Lu et al. 2007, 2008), we find that this expansion can be explained with the Held (2000) theory, where changes in subtropical baroclinicity control the width of the Hadley cell. In addition, we have shown that a rapid increase in static stability (i.e., decrease in the subtropical lapse rate) in the first few years of the simulation leads to the rapid poleward shift of the Hadley cell edge. This was found to occur in both the abrupt $4 \times CO_2$ and the 1% yr$^{-1}$ increase forcing. Changes in tropopause height, however, were found to divert the poleward shift of the Held (2000) theory from that of the Hadley cell edge. Unlike the static stability, the increase in tropopause height was found to have a minor effect on subtropical baroclinicity, and thus on the widening of the tropical belt.

Changes in subtropical baroclinicity also affect subtropical eddy fields. As in Ceppi and Hartmann (2013), we find that the poleward shift of the Hadley cell is indeed highly correlated with the poleward shift of subtropical eddy momentum flux. Several studies have suggested that this correlation might not be solely due to
changes in eddy generation (i.e., baroclinicity) outside the tropics, but also due to an increase in eddy phase speed, which pushes the critical latitude poleward (where eddies dissipate) together with the subtropical eddy momentum flux. Here, however, we have shown that the increase of the eddy phase speed under abrupt $4 \times CO_2$ (e.g., Chen and Held 2007) does not correlate with the poleward shift of the Hadley cell edge. And, more importantly, changes in global-mean surface temperature and meridional surface temperature gradient were also found not to accompany changes in Hadley cell width.

Unlike the poleward shift of the Hadley cell edge, which is relatively fast, the poleward shift of the dry zone edge has a slower time response, and monotonically increases throughout the simulation (Grise and Polvani 2017; Seviour et al. 2018). In the first few years after CO$_2$ quadrupling the rapid poleward movement of the Hadley cell edge shifts the dry zone edge poleward. This accounts for most of the poleward shift of the dry zone edge. The slower increase in moisture further pushes the dry zone edge poleward through the rest of the run. Eddy moisture flux was found to have a minor contribution to shifting the dry zone edge poleward, due to the cancelling effects of transient and stationary eddies, which, respectively, diverge and converge moisture in the subtropics.

The widening of the Hadley cell is only one example of circulation changes under increased CO$_2$ concentrations. For each of these circulation changes, one can find several mechanisms in the literature. Having several mechanisms for one physical phenomenon stems from the difficulty of isolating any causality in such a coupled and complex system, which leads to develop mechanisms based on correlating the long-term fields. However, studying the evolution of the different components in the atmosphere to an abrupt and strong forcing allows taking a step toward causality, and thus having better physical understanding of the climate system.

It is important to note that the key role of static stability that we have found in our study, that of modulating...
the Hadley cell edge under increased greenhouse gases, does not necessarily apply under different forcings. For example, analyzing different idealized warming patterns, Tandon et al. (2013) argued that the wind shear, and not static stability, is most important for changing subtropical baroclinicity and the edge of the Hadley cell. Thus, the widening of the circulation in recent years, which is likely driven by both internal variability (Allen et al. 2014; Quan et al. 2014; Lucas and Nguyen 2015; Allen and Kovilakam 2017; Mantisis et al. 2017) and different forcing agents (Allen et al. 2012, 2014; Waugh et al. 2015), may stem from other components and not only from static stability. Nonetheless, the main result of our study is that subtropical static stability is key to tropical expansion under increased CO2 concentrations. In accordance with our results, Shaw and Tan (2018) have recently demonstrated the importance of subtropical changes in modulating the width of the Hadley cell under increased greenhouse gases.

Finally, the next step for fully understanding the widening of the Hadley cell under increased greenhouse gases should focus on elucidating the different warming rates in the lower and upper subtropical troposphere (i.e., the increase in static stability). To accomplish that, as was done in Chemke and Polvani (2018), one has to analyze the thermodynamic equation and quantify for the different terms that affect the temperature field. Currently, only one of the CMIP5 models provides all the terms needed to analyze the thermodynamic equation, but this is not sufficient for disentangling the natural variability and the model biases from the forced response.

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