

# Climate sensitivity from radiative-convective equilibrium: a chalkboard approach

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## **Abstract**

Simple models for Earth's climate sensitivity, or its temperature response to radiative forcing, are developed by combining the time-tested idealization of one-dimensional radiative-convective equilibrium (RCE) with simple yet quantitatively reasonable models for CO<sub>2</sub> forcing and the water vapor feedback. Along the way we introduce key paradigms including the emission level approximation, the forcing-feedback decomposition of climate sensitivity, and 'Simpson's Law' for water vapor thermal emission. We also discuss climate feedbacks unaccounted for in this RCE framework, as well as differing variants of climate sensitivity, all of which may be ripe for their own chalkboard treatments.

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# 1 Introduction

Earth's equilibrium climate sensitivity (ECS), or the long-term global-mean surface temperature change due to a doubling of  $\text{CO}_2$ , is arguably the most central quantity in climate science. First studied by Arrhenius in 1896,<sup>1</sup> ECS sets the overall magnitude and thus the severity of global warming and remains a topic of intense interest to the present day.<sup>2</sup>

The consensus value of ECS has remained close to 3 K for decades, throughout many successive generations of model intercomparisons and literature reviews, most notably the assessment reports from the Intergovernmental Panel on Climate Change (IPCC).<sup>3</sup> These assessments have been based largely on calculations with numerical models of increasing complexity, along with observations and paleoclimate reconstructions. However, the first credible estimate of ECS, given by Manabe and Wetherald<sup>4</sup> in 1967 as 2-3 K, used a highly simplified one-dimensional representation of the climate system known as *radiative-convective equilibrium* (RCE; we refer to this paper hereafter as MW67). This model for Earth's climate makes some drastic simplifications, such as representing the atmosphere in terms of a single, global average column, but then treats other aspects of the climate system in detail, such as the frequency-dependent greenhouse gas radiative transfer of both water vapor ( $\text{H}_2\text{O}$ ) and carbon dioxide ( $\text{CO}_2$ ). These approximations, and the insights they facilitated, have proved remarkably durable, and were recognized with Manabe's receipt of part of the 2021 Nobel Prize in Physics.<sup>5</sup>

The simplicity of Manabe's RCE approach, and the robustness of the 3 K value for ECS, suggests that much of the physics of ECS might actually be understood using a handful of basic physical principles, rather than lying hidden behind the intractable complexity of the climate system. Indeed, recent research has shown that essential topics such as  $\text{CO}_2$  radiative forcing<sup>6</sup> and the  $\text{H}_2\text{O}$  feedback<sup>7</sup> can be described, to a reasonable approximation, with simplified models amenable to analytic description. Taken together, such models allow for an analytic estimate of ECS, at least within the simplified context of RCE. A self-contained formulation of such an estimate is the primary goal of this paper.<sup>8</sup>

The utility of such an estimate, and the motivations for documenting it here, are many-fold. Perhaps the most obvious motivation is for classroom teaching. Current texts in climate science at the advanced undergraduate or beginning graduate level, such as the excellent books by Hartmann<sup>9</sup> and Pierrehumbert,<sup>10</sup> treat the fundamentals extremely well and use them to build intuition, but then tend to jump to empirical observations and numerical simulations to describe real-world phenomena. This is often the best that can be done, but reasonably quantitative chalkboard estimates (where possible) help fill this gap between fundamental theory and empirical observation. Such back-of-the-envelope reasoning is also helpful for everyday practitioners trying to understand and debug the behavior of complex climate models.<sup>11</sup> The analytical approach taken here might also appeal more naturally to physicists, providing a more suitable entry point for physicists seeking to understand or perhaps even contribute to climate science.

But there are also more profound reasons for pursuing a simplified understanding of ECS, and climate science more broadly.<sup>12</sup> Simple models of the kind developed here embody our understanding of the subject at its most basic level. This philosophy is well known to physicists, in the guise of Fermi problems and the Feynman lectures. Given the societal importance of climate change, as well as lingering skepticism about it in some quarters, a chalkboard explanation of ECS, even if only approximate, seems essential for demonstrating the depth of our understanding.

Given this motivation, we develop an estimate of ECS in the simplified framework of RCE pioneered by Manabe. We begin by establishing the paradigm of radiative-convective equilibrium,

as well as the forcing-feedback framework. We then turn to simple models of the greenhouse effect for both  $\text{CO}_2$  and  $\text{H}_2\text{O}$ , which lead to analytical descriptions of both  $\text{CO}_2$  forcing and the water vapor feedback. These ingredients are then combined, via the forcing-feedback framework, to yield an analytical estimate of ECS. We will find that this estimate is around 2 K, within the range found by MW67 but somewhat smaller than the consensus value of 3 K. This discrepancy is due to the approximations inherent in the RCE approach, which neglects clouds and changes in absorbed solar radiation. We then discuss these phenomena, and the prospects for capturing their effects via principled estimates. We close by introducing other metrics of climate sensitivity that account for ocean heat uptake and the carbon cycle, and for which a chalkboard understanding might also be useful.

For classroom instruction, this material might be considered a unit of perhaps eight lectures at the advanced undergraduate or beginning graduate level, for either physics students or students of atmospheric and climate science. Suggestions for exercises are sprinkled throughout the main text and the endnotes.

## 2 Radiative-convective equilibrium (RCE)

We begin by building a very simple model for Earth's surface temperature  $T_s$ , based on the idea of radiative-convective equilibrium (RCE) which we explain in detail below. Before turning to RCE, however, we must consider the even more basic notion of planetary energy balance.

### 2.1 Planetary energy balance

The zeroth order determinant of  $T_s$  is the amount of sunlight Earth receives. The solar flux at Earth's orbit is  $S_0 = 1360 \text{ W/m}^2$ , and this flux is incident on an effective surface area  $\pi R_E^2$  (the projected area of the Earth onto a plane perpendicular to the Sun's rays, where  $R_E$  is Earth's radius). A significant fraction of this incident flux is reflected back to space, primarily via clouds and gaseous atmospheric (Rayleigh) scattering, as well as from bright surfaces such as deserts and ice caps. This fraction is known as Earth's *albedo*  $\alpha$ , measured to be  $\alpha \approx 0.30$ .<sup>13</sup>

Dividing by the Earth's total surface area  $4\pi R_E^2$  then gives the globally averaged net incoming solar radiation

$$S \equiv \frac{S_0(1 - \alpha)}{4} \approx 240 \text{ W/m}^2 . \quad (1)$$

The most basic physical constraint on climate is that of *planetary energy balance*, which says that in steady-state  $S$  must be balanced by outgoing thermal radiation, also known as the 'outgoing longwave radiation', or OLR (here 'longwave' means thermal infrared). We estimate OLR as blackbody emission,

$$\text{OLR} = \sigma T_{\text{em}}^4 , \quad (2)$$

for some 'emission temperature'  $T_{\text{em}}$ . Planetary energy balance thus reads

$$S = \text{OLR} \quad (3a)$$

$$= \sigma T_{\text{em}}^4 . \quad (3b)$$

The observed value of  $S$  yields  $T_{\text{em}} = 255 \text{ K}$ . This is much colder than the observed global average surface temperature  $T_s = 288 \text{ K}$ , but is a reasonable estimate of an effective *atmospheric* temperature, consistent with the fact that OLR largely emanates not from the surface, but from atmospheric

greenhouse gases (GHGs, which are gases that absorb and emit thermal infrared radiation – most prominently water vapor and carbon dioxide). But given the atmospheric  $T_{\text{em}}$ , how can we find  $T_{\text{s}}$ ? How are surface and atmospheric temperatures related?

## 2.2 Single-layer radiative equilibrium

Perhaps the simplest model for estimating  $T_{\text{s}}$  in the presence of GHGs is that of a single-layer atmosphere of unit emissivity, i.e. one that absorbs and emits as a perfect blackbody, pictured on the left hand side of Fig. 1 (ignoring the convective flux for the moment). Such an atmosphere is characterized by an atmospheric temperature  $T_{\text{a}}$ , blackbody emission  $\sigma T_{\text{a}}^4$  from GHGs emanating both upwards to space and downwards to the surface, and transparency to sunlight.<sup>14</sup> The energy budgets at the top of the atmosphere (TOA) and surface then read (Fig. 1)

$$\text{TOA :} \quad S = \sigma T_{\text{a}}^4 \quad (4\text{a})$$

$$\text{surf :} \quad S + \sigma T_{\text{a}}^4 = \sigma T_{\text{s}}^4. \quad (4\text{b})$$

These equations can be immediately solved to yield  $T_{\text{a}} = T_{\text{em}} = 255$  K and  $T_{\text{s}} = 2^{1/4}T_{\text{em}} = 303$  K. This calculation thus overestimates  $T_{\text{s}}$  by 15 K.

Despite this disagreement, this model is standard in introductory texts.<sup>15</sup> Many of these texts remedy this disagreement by introducing a non-unit emissivity for the atmosphere, but the value of this emissivity is typically set by *requiring*  $T_{\text{s}}=288$  K, thus eliminating the model’s predictive power. Furthermore, this approach does not remedy the model’s grossest approximation, which is not that of unit emissivity but rather of neglecting heat transfer by *convection*.

## 2.3 A radiative-convective equilibrium model

Convection brings water vapor and heat from the surface into the interior of the atmosphere, where the water vapor condenses to form cloud and rain drops. This condensation releases the latent heat of vaporization which was drawn from the surface when the molecules first evaporated, and this heat from condensation is then radiated out to space by atmospheric GHGs, closing the loop. These processes are depicted on the right-hand side of Fig. 1. Thus, planetary energy balance is not achieved through purely radiative means but is mediated by convection, with water vapor as the key middleman. The atmosphere is thus better thought of as in a state of *radiative-convective equilibrium*, or RCE for short.<sup>16</sup>

To incorporate this into the simple one-layer model (4), we must add a convective heat flux  $F_{\text{c}}$  to the surface energy balance (4b). An additional variable requires an additional equation, however. To obtain this, we invoke the fact that convective heat fluxes are extremely efficient at transferring heat to the atmosphere, with two simplifying consequences: the surface temperature becomes essentially continuous with the atmospheric temperature near the ground, and atmospheric temperature profiles  $T(z)$  lie more or less along a convective *adiabat*. A convective adiabat is the temperature profile of a parcel lifted adiabatically from the ground; such a parcel will expand and cool as it rises along the  $z$ -axis to lower pressures, much like the cool air escaping from the valve of a pressurized bicycle tire. For a dry air parcel this profile is determined by the *dry adiabatic lapse rate*

$$-\left. \frac{dT}{dz} \right|_{\text{dry}} = \frac{g}{C_p} = 10 \text{ K/km} \quad (\text{dry lapse rate}) \quad (5)$$

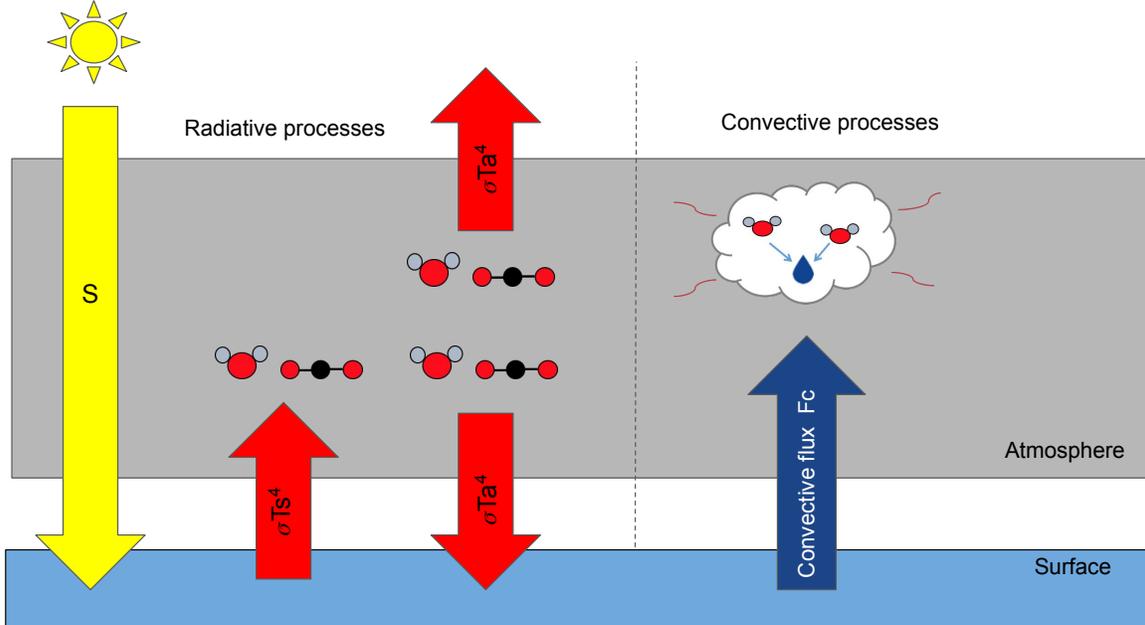


Figure 1: A cartoon of radiative-convective equilibrium. See text for explanation.

where  $g$  is the gravitational acceleration and  $C_p$  is the specific heat of air (see Appendix A, which includes a derivation of (5) from undergraduate thermodynamics). Moisture, along with atmospheric dynamics besides convection, modifies this lapse rate somewhat and makes it variable over the globe, with a global average value of

$$\Gamma \equiv -\frac{dT}{dz} \approx 6.5 \text{ K/km} \quad (\text{avg. lapse rate}) . \quad (6)$$

If we then assume a mean pressure  $p_a = 0.5 \text{ atm}$  for our single-layer atmosphere, corresponding to a height of  $z_a \approx 5 \text{ km}$  (Fig. 2 below),<sup>17</sup> then Eq. (6) yields the *convective constraint*

$$T_s - T_a \equiv \Delta T = (6.5 \text{ K/km})(5 \text{ km}) = 32.5 \text{ K} . \quad (7)$$

Our single-layer RCE model is then

$$S = \sigma T_a^4 \quad (8a)$$

$$S + \sigma T_a^4 = \sigma T_s^4 + F_c . \quad (8b)$$

$$T_s - T_a \equiv \Delta T . \quad (8c)$$

Solving these equations is again straightforward: (8a) again yields  $T_a = T_{em}$ , so then (8c) yields

$$T_s = T_{em} + \Gamma z_a = 287.5 \text{ K} . \quad (9)$$

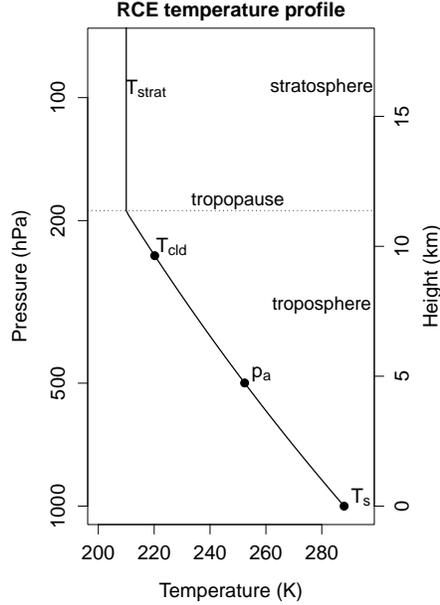


Figure 2: A simplified RCE temperature profile, given by Eq. (10) with  $T_s = 288$  K, along with an isothermal stratosphere with temperature  $T_{\text{strat}} = 210$  K. The RCE profile (10) reaches up to the tropopause at roughly 12 km or 200 hPa, where it intersects the isothermal stratospheric profile. Also shown is the representative height of the single-layer atmosphere  $p_a = 0.5$  atm = 500 hPa, as well as the characteristic cloud-top temperature  $T_{\text{cld}} = 220$  K discussed in Section 6.1.

This is remarkably close to the observed preindustrial value of 288 K. Furthermore, Eq. (9) tells us that this value stems from the radiative energy balance of the planet (as embodied in  $T_{\text{em}}$ ), combined with the effects of convection on the temperature profile of the troposphere (as embodied in  $\Gamma$ ). This single-layer RCE model thus appears to have significant pedagogical advantages over the single-layer radiative equilibrium model (4), but to our knowledge has not yet appeared in textbooks. It has appeared in the journal literature before, for instance in the papers of Hansen et al.<sup>18</sup> and Payne et al.<sup>19</sup>

An important subtlety in the simple calculation above is that it was not necessary to solve the surface energy balance equation (8b); the convective flux  $F_c$  there acts as a Lagrange multiplier, taking on whatever value is required to satisfy Eq. (8b) subject to the constraint (8c); the convective flux is thus analogous to the tension in a pendulum arm, which takes on whatever value is required to satisfy Newton's Laws while keeping the pendulum bob at a fixed distance from its origin.

While this simple one-layer RCE model yields a reasonable estimate of  $T_s$  and also illustrates the essence of how convection couples the surface and atmospheric temperatures, we will also need a vertically-resolved view of the atmosphere for the material that follows. Figure 2 shows the following

temperature profile, obtained by integrating Eq. (6) and also converting to pressure coordinates:

$$T_a(z) = T_s - \Gamma z \quad (10a)$$

$$\text{or } T_a(p) = T_s \left( \frac{p}{p_s} \right)^{R_d \Gamma / g} \quad (10b)$$

where  $R_d$  is the gas constant for dry air and  $p_s$  is the surface pressure (See Appendix A for the definition of  $R_d$  and derivation of Eq. (10b)). Equation (10a) is the vertically-resolved version of the convective constraint Eq. (8c). In general, however, this convective constraint is only valid below an altitude of 10-15 km; this region is known as the *troposphere*. Above the troposphere lies the stratosphere and other air masses, which are heated primarily by ultraviolet solar absorption rather than convection, and are thus closer to radiative equilibrium rather than RCE. For simplicity, the stratosphere is represented here by an isothermal layer with characteristic temperature  $T_{\text{strat}} = 210$  K, which is attached to the troposphere at the *tropopause*, where  $T_a = T_{\text{strat}}$  (Fig. 2). The physics governing the height and temperature of the tropopause, and hence the characteristic temperature  $T_{\text{strat}}$ , is still a subject of active research.<sup>20</sup>

A profound implication of the convective constraint (10a) is that the tropospheric temperature profile  $T_a$  is pegged to  $T_s$ , and the two cannot be varied independently. In other words, the surface and atmosphere should be thought of as a unit, with a single degree of freedom between them.<sup>21</sup> This tight surface-atmosphere coupling is a hallmark of RCE and has important consequences, as we will see below.

### 3 Forcing-feedback preliminaries

Now that we have a picture for Earth’s energy flows and RCE, we can begin to think about climate sensitivity. An extremely useful paradigm for this is the *forcing-feedback framework*, which we describe next.

#### 3.1 The forcing-feedback framework

In this paper, and in much (but certainly not all) climate modeling, the atmospheric CO<sub>2</sub> specific concentration<sup>22</sup>  $q$  (kg CO<sub>2</sub>/kg air) is considered an *external* parameter, which is prescribed and thus does not respond to the internal dynamics of the system.<sup>23</sup> In studying changes in surface temperature  $\Delta T_s$  which result from changes in  $q$ , it turns out to be extremely convenient to decompose the system’s response into two distinct processes, one which occurs at fixed  $T_s$ , and the other which occurs at fixed  $q$ .<sup>24</sup> As we will see, it will be natural sometimes to use  $-\text{OLR}$  as our variable for planetary thermal energy flux, since an increase in  $-\text{OLR}$  indicates increasing planetary thermal energy.

The first process is the decrease in OLR (increase in  $-\text{OLR}$ ) due to an increase in  $q$  from an initial concentration  $q_i$  to a final concentration  $q_f$ , *holding  $T_s$  fixed*; since the lapse rate  $\Gamma$  does not depend on  $q$ , this also fixes  $T_a(z)$  by Eq. (10a).<sup>25</sup> This decrease in OLR due to the change in  $q$  is known as the CO<sub>2</sub> *forcing*

$$\mathcal{F} \equiv -(\text{OLR}(q_f) - \text{OLR}(q_i)) . \quad (11)$$

( $\mathcal{F}$  is the change in  $-\text{OLR}$ , and thus  $\mathcal{F} > 0$  for a CO<sub>2</sub> increase). The forcing  $\mathcal{F}$  breaks the planetary energy balance, causing additional heat to accumulate in the system.

Next, this heat accumulation causes a temperature increase  $\Delta T_s$  and hence a countervailing *increase* in OLR (decrease in  $-\text{OLR}$ ), which persists until the planetary energy balance (3a) is restored and a new equilibrium is established (how long this takes, and what happens along the way, is addressed in section 7.1). This increase in OLR due to increasing  $T_s$  can be approximated as  $(d\text{OLR}/dT_s)\Delta T_s$ , where the derivative is taken at fixed  $q = q_f$ . Consistent with the sign convention in Eq. (11), we then define the ‘feedback parameter’  $\lambda$  as minus this derivative<sup>26</sup>:

$$\lambda \equiv - \left. \frac{d\text{OLR}}{dT_s} \right|_{q=q_f} . \quad (12)$$

Noting that  $(d\text{OLR}/dT_s)\Delta T_s$  must balance the forcing, we have  $\Delta T_s = -\mathcal{F}/\lambda$ . When  $q_f = 2q_i$ , we write the forcing as  $\mathcal{F}_{2\times}$ , and the corresponding  $\Delta T_s$  is defined to be the *equilibrium climate sensitivity* (ECS), that is

$$\text{ECS} = - \frac{\mathcal{F}_{2\times}}{\lambda} . \quad (13)$$

(The choice of a multiplicative rather than additive change in  $q$  for defining ECS will be discussed below.) Equation (13) is known as the *forcing-feedback* decomposition of ECS. It allows us to study ECS by studying  $\mathcal{F}$  and  $\lambda$  separately, which we will do in sections 4 and 5 respectively. Note that we have ignored the possibility that the absorbed solar radiation  $S$  may also depend on  $T_s$ , which would contribute a  $dS/dT_s$  term to the definition of  $\lambda$ . We discuss these so-called ‘shortwave’ feedbacks in Section 6.

### 3.2 Blackbody estimate of ECS

As an exercise we will use Eq. (13) to estimate ECS using the blackbody approximation (2). The blackbody approximation allows us to calculate  $\lambda$ , but we still need a value for the forcing  $\mathcal{F}_{2\times}$ . For the moment, we obtain this by appealing to comprehensive radiative transfer calculations, which for decades<sup>27</sup> have found a fairly consistent value of

$$\mathcal{F}_{2\times} \approx 4 \text{ W/m}^2 . \quad (14)$$

We will provide a theoretical estimate of this number in the next section.

Turning now to the feedback parameter  $\lambda$ , a blackbody estimate for this can be obtained from (2), noting also that  $T_{\text{em}}$  and  $T_s$  vary in a 1-1 fashion according to (9). Recalling that  $T_{\text{em}} = 255$  K, this yields:

$$\lambda_{\text{blackbody}} = - \frac{d\text{OLR}}{dT_s} = -4\sigma T_{\text{em}}^3 \approx -3.75 \text{ (W/m}^2\text{)/K} . \quad (15)$$

According to Eq. (13), this yields  $\text{ECS} = (4 \text{ W/m}^2)/(3.75 \text{ (W/m}^2\text{)/K}) \approx 1$  K, three times smaller than the consensus value of 3 K.

To understand why  $\mathcal{F}_{2\times} \approx 4 \text{ W/m}^2$ , as well as make a better estimate of  $\lambda$  and hence ECS, we need to move beyond the blackbody approximation and account for the *spectral* nature of Earth’s greenhouse effect, i.e. that the emission temperature  $T_{\text{em}}$  of outgoing longwave radiation actually depends rather markedly on frequency, and that  $T_{\text{em}}$  at a given frequency and  $T_s$  may not necessarily exhibit a 1-1 relationship as in Eq. (9). A key ingredient in understanding how these quantities do behave will be the *emission level approximation*, which we turn to next.

### 3.3 Emission level approximation

In Section 2, we defined Earth’s emission temperature  $T_{\text{em}}$  by Eq. (2), and then assumed an average blackbody emission level  $p_a = 0.5$  atm. This led to a 1-1 relationship between  $T_{\text{em}}$  and  $T_s$  which allowed us to estimate  $\lambda$ .

To refine this, we first define a spectrally-resolved emission temperature  $T_{\text{em}}(\nu)$  by the relation

$$\text{OLR}_\nu = \pi B(\nu, T_{\text{em}}(\nu)) \quad (16)$$

where  $\text{OLR}_\nu$  is the spectrally-resolved OLR in units of  $\text{W}/(\text{m}^2 \cdot \text{cm}^{-1})$  and satisfying  $\text{OLR} = \int \text{OLR}_\nu d\nu$ , and  $B(\nu, T)$  is the Planck density in units of  $\text{W}/(\text{m}^2 \cdot \text{cm}^{-1} \cdot \text{sr})$  (the factor of  $\pi$  above accounts for integration over solid angle).<sup>28</sup> The spectral coordinate here is ‘wavenumber’  $\nu$ , defined as inverse wavelength and thus proportional to frequency, with standard unit of  $\text{cm}^{-1}$ . The definition (16) of  $T_{\text{em}}(\nu)$ , as the temperature whose Planck emission yields  $\text{OLR}_\nu$ , straightforwardly generalizes the blackbody definition (2).

The next task is to determine what level(s) in the atmosphere determine  $T_{\text{em}}(\nu)$ , for a given  $\nu$ . This task is aided by the following heuristic, illustrated in Fig. 3. Consider an atmospheric column with GHG molecules whose density  $\rho_{\text{GHG}}$  [ $(\text{kg GHG})/\text{m}^3$ ] decreases exponentially with height; this is true for both  $\text{CO}_2$  and  $\text{H}_2\text{O}$ .<sup>29</sup> Now consider the emission to space (i.e. the contribution to the  $\text{OLR}_\nu$  at a given  $\nu$ ) from these molecules, as pictured in Fig. 3. The top two layers (Fig. 3a) have little difficulty emitting to space because their view is unobstructed, but the density of emitters in these layers is relatively low, so the emission will also be low. In the third layer (Fig. 3b) the molecules’ view of space is still unobstructed (just barely), and their density is higher, so their emission to space is higher. For layers four and five (Fig. 3c) there are plenty of emitters, but their view is almost totally obstructed, so their emission to space is again very low. Thus, emission to space is maximized around a ‘sweet spot’ where the absorbers/emitters above have not yet totally obstructed the view of space, but the density is high enough for emission to be appreciable. This sweet spot will be our emission level.

To formalize this we need the notion of *optical depth*, defined as<sup>30</sup>

$$\tau_\nu(z) \equiv \int_z^\infty \underbrace{\kappa_{\text{ref}}(\nu)}_{(\text{m}^2/\text{kg})} \underbrace{\frac{p}{p_{\text{ref}}}}_{(\text{kg}/\text{m}^2)} \underbrace{\rho_{\text{GHG}} dz'}_{(\text{kg}/\text{m}^2)} = \frac{\text{Effective area of absorbers}}{\text{Actual area of column}}. \quad (17)$$

Here  $\kappa_{\text{ref}}(\nu)$  are the *mass absorption coefficients* of our GHG, which give the effective cross-sectional area of our GHG molecules at wavenumber  $\nu$  per unit mass (units  $\text{m}^2/\text{kg}$ ), evaluated at a reference pressure  $p_{\text{ref}}$ . This effective cross-section depends strongly on frequency (e.g. Figs. 4a and 5a below) but also scales with pressure as  $p/p_{\text{ref}}$ , hence the presence of this factor in Eq. (17).<sup>31</sup> The factor of  $\rho_{\text{GHG}} dz'$  in the integrand in Eq. (17) gives the absorber mass per unit area ( $\text{kg}/\text{m}^2$ ) in an atmospheric layer of differential depth  $dz'$ . Thus, the integral  $\tau_\nu$  in Eq. (17) can be interpreted as the ratio of the integrated effective area of absorbers above height  $z$  to the actual area of the column, as noted in Eq. (17). Applying this notion to Fig. 3, we see that the top two layers with low emissions correspond to  $\tau_\nu < 1$ , where the total effective area of absorbers above is less than the actual area of the column (the ‘optically thin’ regime). Similarly, the bottom two layers with low emissions correspond to  $\tau_\nu > 1$ , where the total effective area of absorbers above is greater than the actual area of the column (the ‘optically thick’ regime). The sweet spot occurs around  $\tau_\nu \approx 1$ .

For simplicity we further assume that *all* emission occurs at exactly  $\tau_\nu = 1$ ; we refer to this as the *emission level approximation*.<sup>32</sup> With this in mind we define our emission pressure  $p_{\text{em}}(\nu)$  by

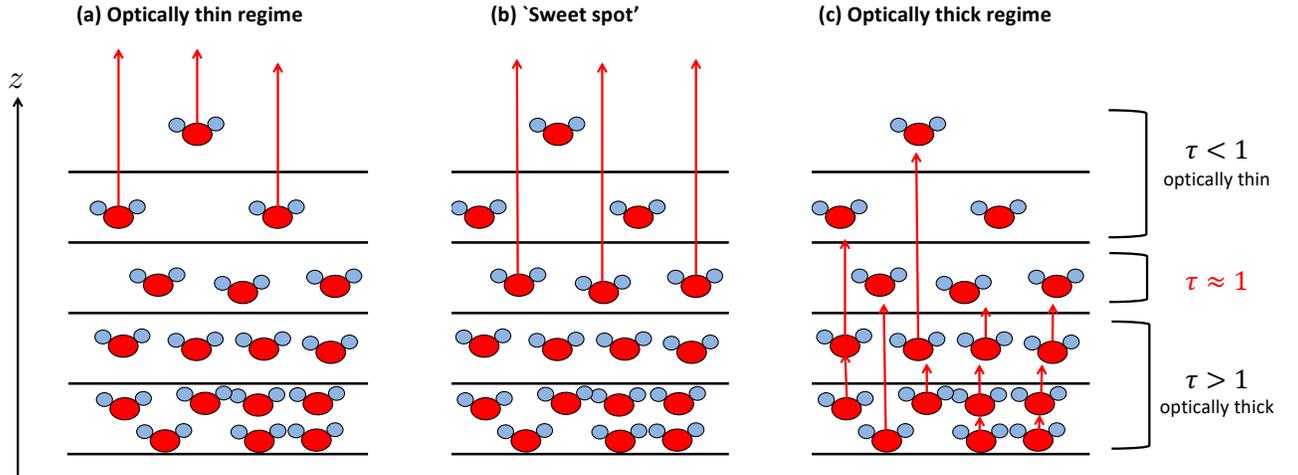


Figure 3: Cartoon of emission to space from different atmospheric layers corresponding to different optical depths. Layer-wise emission to space maximizes at a sweet spot where the optical depth  $\tau \approx 1$ .

the relation

$$\tau_\nu(p_{\text{em}}(\nu)) = 1. \quad (18)$$

This equation will form the basis for reasoning about both  $\text{CO}_2$  forcing as well as the  $\text{H}_2\text{O}$  feedback, which are the subjects of the following sections.

## 4 $\text{CO}_2$ forcing

This section constructs a simple analytic model for the  $\text{CO}_2$  forcing Eq. (11), with the aim of enabling a back-of-the-envelope estimate of the characteristic  $4 \text{ W/m}^2$  value for  $\mathcal{F}_{2\times}$ . The approach here is to consider spectral variations in  $\text{CO}_2$  absorption but to do so in a simplified manner, so as to keep the model analytically tractable. The model we construct has precedent in relatively recent literature, e.g. the works of Wilson and Gea-Banacloche,<sup>33</sup> Jeevanjee et al.,<sup>34</sup> and Romps et al.,<sup>35</sup> but has not yet appeared in textbooks. We present here a very simple version of the model, which can be generalized to include the effects of a non-isothermal stratosphere as well as spectral overlap between  $\text{CO}_2$  and  $\text{H}_2\text{O}$  absorption.<sup>36</sup>

We begin with  $\text{CO}_2$  reference mass absorption coefficients  $\kappa_{\text{CO}_2}$  in the strongly absorbing and climatically relevant  $\nu_2$  bending mode centered on wavenumber  $\nu_0 = 667 \text{ cm}^{-1}$ . Figure 4a displays this absorption spectrum.<sup>37</sup> A key simplification we make is to ignore fine-scale spectral structure and note that on a broad scale,  $\kappa_{\text{CO}_2}(\nu)$  may be parameterized as

$$\kappa_{\text{CO}_2}(\nu) = \kappa_0 \exp\left(-\frac{|\nu - \nu_0|}{l}\right) \quad (19)$$

where  $\kappa_0$  represents a smoothed peak absorption coefficient and  $l$  is a ‘spectroscopic decay parameter’, which sets the rate at which absorption declines away from the peak.

With Eq. (19) in hand, we next evaluate the optical depth integral (17) for CO<sub>2</sub>, for which  $\rho_{\text{GHG}} = q\rho$  where  $q$  is a constant CO<sub>2</sub> specific concentration and  $\rho$  is the density of air. Plugging this into Eq. (17) and invoking hydrostatic balance  $dp/dz = -\rho g$  [see also Eq. (44) below] then yields

$$\tau_{\text{CO}_2}(\nu, p) = \kappa_{\text{CO}_2}(\nu) \int_0^p \frac{p'}{p_{\text{ref}}} \frac{q dp'}{g} = \frac{\kappa_{\text{CO}_2}(\nu) q}{2g} \frac{p^2}{p_{\text{ref}}}. \quad (20)$$

The pressure broadening factor in the integrand above, as well as the fact that the mass of CO<sub>2</sub> in a layer is given by  $qdp'/g$ , conspire to give  $\tau \sim p^2$ .

Invoking the emission level approximation (18), as well as the ‘exponential spectroscopy’ (19), we can set  $\tau_{\text{CO}_2} = 1$  and invert Eq. (20) to find the emission levels  $p_{\text{em}}(\nu)$ :

$$p_{\text{em}}(\nu) = \sqrt{\frac{2gp_{\text{ref}}}{q\kappa_0}} \exp\left(-\frac{|\nu - \nu_0|}{2l}\right). \quad (21)$$

These emissions levels are shown in Fig. 4b for  $(q_i, q_f) = (280, 1120)$  ppm (a quadrupling of CO<sub>2</sub> is easier to visualize than a doubling). The  $p_{\text{em}}$  curves take the form of ‘emission triangles’, which widen and deepen with increased CO<sub>2</sub>. The red and green highlighted regions show that an increase in  $q$  yields a decrease in surface emission, as well as ‘new’ emission from portions of the stratosphere (around 20hPa) that were not emitting significantly before. Also, from Eq. (21) one can deduce that the change in width of the emission triangles with  $q$  is given by

$$\Delta\nu = l \ln\left(\frac{q_f}{q_i}\right). \quad (22)$$

Note the logarithmic dependence on  $q$ , which we comment on further below.

To turn these insights into a formula for CO<sub>2</sub> forcing, Fig. 4c shows  $\text{OLR}_\nu = \pi B(\nu, T_{\text{em}})$  obtained by inserting the emission levels  $p_{\text{em}}$  from Fig. 4b into the temperature profile (10b) and then applying the Planck function. The large values on the outside of the CO<sub>2</sub> band correspond to surface emission  $\pi B(\nu, T_s)$ , and the smaller values in the center correspond to stratospheric emission  $\pi B(\nu, T_{\text{strat}})$  (i.e. where  $p_{\text{em}}(\nu) \lesssim 200$  hPa). Given the definition Eq. (11) of the forcing and the expression (22) for  $\Delta\nu$ , the forcing can then be read off from Fig. 4c as just the difference in the area under solid and dashed curves, i.e.

$$\mathcal{F} = 2l \ln\left(\frac{q_f}{q_i}\right) [\pi B(\nu_0, T_s) - \pi B(\nu_0, T_{\text{strat}})]. \quad (23)$$

This formula can also be interpreted via Fig. 4b as a swap of stratospheric emission for surface emission over spectral regions of total width  $2\Delta\nu$ . For an analytical derivation of Eq. (23), see Appendix B.

We are now in a position to make a back-of-the-envelope estimate of CO<sub>2</sub> forcing. Evaluating Eq. (23) for  $T_s = 288$  K,  $T_{\text{strat}} = 210$  K, and for  $q_f = 2q_i$  yields the estimate

$$\mathcal{F}_{2\times} \approx 2l \ln 2 [\pi B(\nu_0, 288 \text{ K}) - \pi B(\nu_0, 210 \text{ K})] \approx 4.15 \text{ W/m}^2, \quad (24)$$

close to the standard value of 4 W/m<sup>2</sup>.<sup>40</sup>

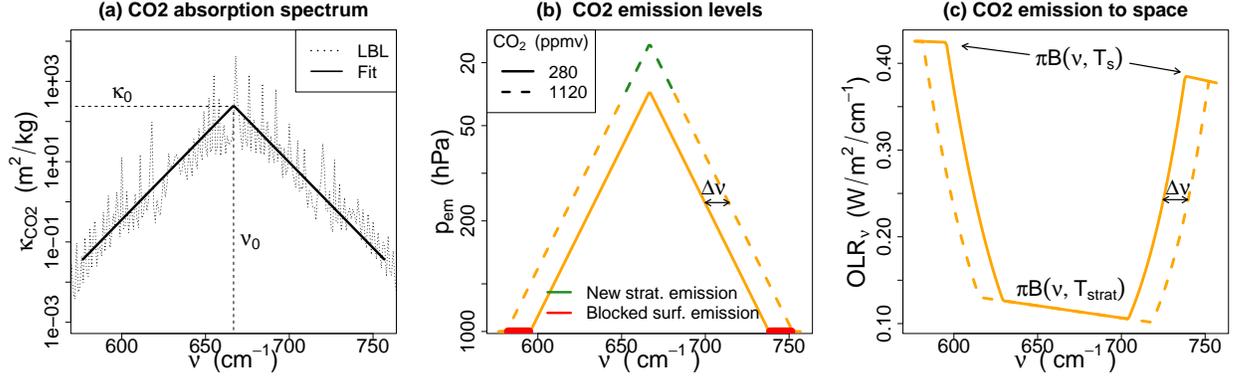


Figure 4: **(a)** CO<sub>2</sub> absorption coefficients  $\kappa_{\text{CO}_2}$  as calculated by a line-by-line (LBL) benchmark radiation code,<sup>39</sup> coarse-grained into 10 cm<sup>-1</sup> bins, along with an exponential fit of the form (19) with  $\kappa_0 = 240$  m<sup>2</sup>/kg and  $l = 10.2$  cm<sup>-1</sup>. Reference pressure and temperature for  $\kappa_{\text{CO}_2}$  are  $(p_{\text{ref}}, T_{\text{ref}}) = (500 \text{ hPa}, 260 \text{ K})$ . **(b)** Graph of emission levels  $p_{\text{em}}$  from Eq. (21), for CO<sub>2</sub> concentrations of 280 and 1120 ppm. **(c)** Graph of  $\text{OLR}_\nu$  calculated as  $\pi B(\nu, T(p_{\text{em}}(\nu)))$ , using the RCE temperature profile from Fig. 2.

The formalism developed here also yields insight into the logarithmic scaling of CO<sub>2</sub> forcing, evident in the  $\ln(q_f/q_i)$  factor in Eqs. (22) and (23). Unwinding the mathematics leading to Eq. (22), one finds that the  $\ln(q_f/q_i)$  factor stems from the exponential spectroscopy  $\kappa_{\text{CO}_2}(\nu)$  in Eq. (19). The fact that  $\kappa_{\text{CO}_2}(\nu)$  appears multiplied by  $q$  in all the relevant physical quantities [e.g. Eqs. (20),(21)] means that a multiplicative change in  $q$  is equivalent to an additive change in  $\nu$  (cf. Appendix B). Thus multiplicative changes in  $q$  always change the width of the  $p_{\text{em}}$  triangles in Fig. 4b by the same  $\Delta\nu$ , and the forcing is proportional to this width.

## 5 The water vapor feedback

Now that we have a handle on CO<sub>2</sub> forcing, we turn towards a better estimate of the feedback parameter  $\lambda$ . For  $\lambda$ , H<sub>2</sub>O will turn out to be the dominant GHG, as its absorption spectrum spans the entirety of the thermal infrared, as shown in Fig. 5 (compare the wavenumber range in Fig. 5 to that of Fig. 4; note also that in analyzing  $\lambda$  we neglect the effects of CO<sub>2</sub> for simplicity). Similar to CO<sub>2</sub>, the H<sub>2</sub>O reference mass absorption coefficients  $\kappa_{\text{H}_2\text{O}}(\nu)$  can be parameterized in terms of exponentials, this time with two exponentials peaked at  $\nu_1 = 150$  cm<sup>-1</sup> and  $\nu_2 = 1500$  cm<sup>-1</sup>, as

$$\kappa_{\text{H}_2\text{O}}(\nu) = \max \left[ \kappa_1 \exp \left( -\frac{|\nu - \nu_1|}{l_1} \right), \kappa_2 \exp \left( -\frac{|\nu - \nu_2|}{l_2} \right) \right]. \quad (25)$$

Note the absorption minimum at roughly 1000 cm<sup>-1</sup>, which will play an important role in what follows.<sup>41</sup>

Next we evaluate the optical depth integral Eq. (17). Here  $\rho_{\text{GHG}}$  is now the water vapor density

$\rho_v$ , which we write as

$$\rho_v = \frac{p_v}{R_v T} = \frac{\text{RH} p_v^*(T)}{R_v T} \quad (26)$$

where the first equality is the ideal gas law (43) for water vapor and the second is just a definition of the relative humidity RH, which we take to have a fixed value of 0.75.<sup>42</sup> Here  $p_v$  is the partial pressure of water vapor,  $R_v = 287 \text{ J}/(\text{kg} \cdot \text{K})$  is the specific gas constant for water vapor, and

$$p_v^*(T) = p_v^{\text{ref}} \exp\left(-\frac{L}{R_v T}\right) \quad (\text{Clausius-Clapeyron}) \quad (27)$$

is the Clausius-Clapeyron relation for the saturation partial pressure of water vapor, with reference pressure  $p_v^{\text{ref}} = 2.5 \times 10^{11} \text{ Pa}$  and  $L$  the latent heat of vaporization (equal to  $2.5 \times 10^6 \text{ J}/\text{kg}$ ). The Clausius-Clapeyron expression is fundamental to our RCE picture of the atmosphere, as it determines how much the water vapor content of a parcel decreases (and hence how much condensation is produced) per degree of cooling.

Inserting these expressions into Eq. (17) and changing the dummy integration variable from  $z'$  to  $T'$  (where the lower limit on  $T'$  is taken to be the tropopause and thus equal to  $T_{\text{strat}} = 210 \text{ K}$ ) yields the  $\text{H}_2\text{O}$  optical depth

$$\begin{aligned} \tau_{\text{H}_2\text{O}}(\nu, T) &= \kappa_{\text{H}_2\text{O}}(\nu) \int_{T_{\text{strat}}}^T \frac{\text{RH} p_v^{\text{ref}}}{R_v T'} \frac{p(T')}{p_{\text{ref}}} \exp\left(-\frac{L}{R_v T'}\right) \frac{dT'}{\Gamma} \\ &\approx \frac{p}{p_{\text{ref}}} \kappa_{\text{H}_2\text{O}}(\nu) M_v^{\text{ref}} \exp\left(-\frac{L}{R_v T}\right) \end{aligned} \quad (28)$$

where  $T_{\text{av}} \equiv (T_s + T_{\text{tp}})/2$ ,  $M_v^{\text{ref}} \equiv \frac{\text{RH} p_v^{\text{ref}} T_{\text{av}}}{\Gamma L}$  is a reference value for column water vapor mass ( $\text{kg}/\text{m}^2$ ), and several approximations were made to make the integral tractable.<sup>43</sup> Note that  $M_v^{\text{ref}} \exp(-L/R_v T)$  is the column mass of water vapor above the isotherm with temperature  $T$ , so Eq. (28) is just the pressure-weighted GHG mass above  $T$  times the cross-sectional area per unit mass, consistent with the interpretation in Eq. (17). Water vapor emission temperatures  $T_{\text{em}}$  can now be diagnosed by setting  $\tau_{\text{H}_2\text{O}} = 1$  and inverting Eq. (28) either numerically or analytically;<sup>44</sup> where  $\tau_{\text{H}_2\text{O}}(\nu, T_s) < 1$ , in the aforementioned optically thin region centered around  $1000 \text{ cm}^{-1}$  (known as the infrared ‘window’), the emission level lies below the surface so we set  $T_{\text{em}} = T_s$ . The results for  $T_s = (280, 290, 300) \text{ K}$  are plotted in Fig. 5b.

A conspicuous feature of Fig. 5b is that the atmospheric emission temperatures  $T_{\text{em}}$  in the optically thick regions seem to be almost entirely insensitive to  $T_s$ , i.e.  $dT_{\text{em}}/dT_s \approx 0$ . This can be deduced from Eq. (28), where vertical variations in  $\tau_{\text{H}_2\text{O}}$  at a given  $\nu$  are dominated by the temperature-dependent exponential, with the pressure-broadening factor playing only a secondary role. Thus, to a good approximation  $\tau_{\text{H}_2\text{O}}$  is a function of temperature alone, and  $\tau_{\text{H}_2\text{O}}(\nu) = 1$  will occur at approximately the same  $T_{\text{em}}$  *regardless of*  $T_s$  (assuming fixed RH). We formalize this fact, first formulated by Simpson<sup>47</sup> in 1928, as ‘Simpson’s Law’:

**Simpson’s Law:** At fixed RH, and for optically thick wavenumbers  $\nu$  dominated by  $\text{H}_2\text{O}$  absorption, emission temperatures are insensitive to surface temperature, i.e.

$$\left. \frac{dT_{\text{em}}(\nu)}{dT_s} \right|_{\text{RH}} \approx 0. \quad (\nu \text{ optically thick})$$

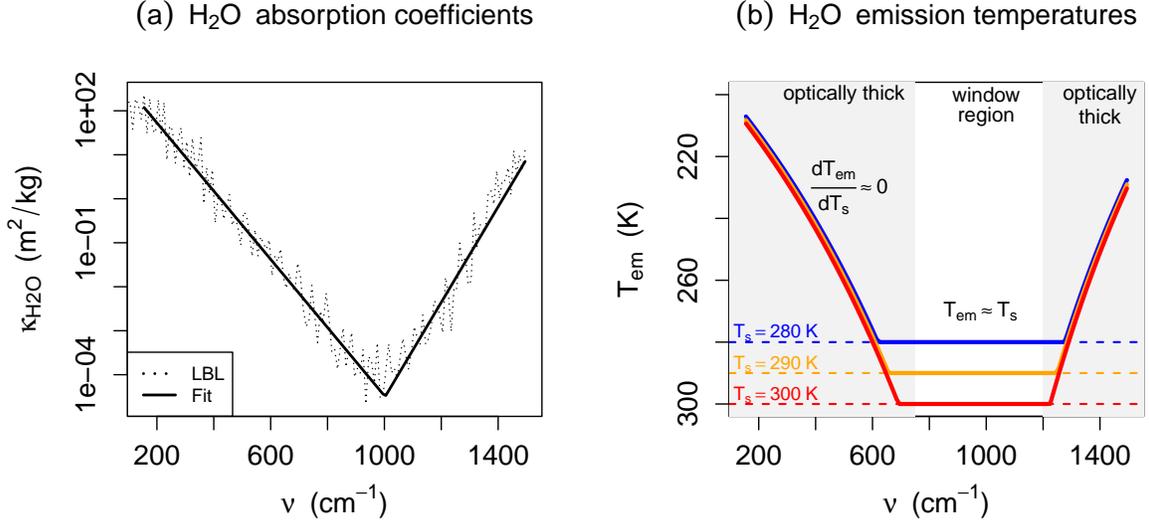


Figure 5: **(a)**  $\text{H}_2\text{O}$  absorption coefficients from a line-by-line (LBL) benchmark calculation<sup>46</sup> at  $(p_{\text{ref}}, T_{\text{ref}}) = (500 \text{ hPa}, 260 \text{ K})$ , along with an exponential fit of the form (25) with  $l_1 = 56 \text{ cm}^{-1}$ ,  $l_2 = 40 \text{ cm}^{-1}$ ,  $\kappa_1 = 130 \text{ m}^2/\text{kg}$ , and  $\kappa_2 = 8 \text{ m}^2/\text{kg}$ . These absorption coefficients have been simplified by neglecting continuum absorption. **(b)** Graph of emission temperatures  $T_{\text{em}}$  diagnosed from Eq. (28) for  $T_s = (280, 290, 300) \text{ K}$ , which are insensitive to  $T_s$  except in the optically thin infrared ‘window’ at  $750 - 1200 \text{ cm}^{-1}$ .

This result is of course only approximate, as the pressure broadening factor (which includes a mild  $T_s$ -dependence via Eq. (10b)) and other effects cause slight deviations,<sup>48</sup> but we nonetheless refer to it as a ‘law’ as it plays a fundamental role in governing the strength of the water vapor feedback, as follows. If  $T_{\text{em}}(\nu)$  and hence  $\text{OLR}_\nu$  (by Eq. (16)) are independent of  $T_s$  for optically thick  $\nu$ , and if we consider the atmosphere perfectly transparent for the optically thin  $\nu$  in the window region so that  $\text{OLR}_\nu$  is given by surface emission  $\pi B(\nu, T_s)$  for those wavenumbers, then the spectrally-resolved feedback parameter  $\lambda_\nu$  (satisfying  $\lambda = \int \lambda_\nu d\nu$ ) is given by

$$\lambda_\nu \equiv -\frac{d\text{OLR}_\nu}{dT_s} \approx \begin{cases} 0 & (\nu \text{ optically thick}) \\ -\pi \frac{\partial B(\nu, T_s)}{\partial T_s} & (\text{window region}) . \end{cases}$$

In other words, the feedback is zero in the optically thick regions due to Simpson’s Law, and in the window region is just given by the surface blackbody response. Given this, it is now straightforward to spectrally integrate  $\lambda_\nu$  and evaluate at  $T_s = 288 \text{ K}$  to obtain  $\lambda$ ; all that is required is an estimate of the limits of the window region. This is typically<sup>49</sup> taken to be  $800 < \nu < 1200 \text{ cm}^{-1}$ , although these limits are not precisely defined. For the sake of obtaining round numbers we take the lower

limit to be  $750 \text{ cm}^{-1}$ , which then yields our RCE estimate of  $\lambda$ :<sup>50</sup>

$$\lambda_{\text{RCE}} = - \int_{750 \text{ cm}^{-1}}^{1200 \text{ cm}^{-1}} \pi \frac{\partial B(\nu, T_s)}{\partial T_s} d\nu \approx -2 \text{ (W/m}^2\text{)/K} , \quad (29)$$

where the integral is computed numerically. This value of  $\lambda$ , which was obtained here in an RCE context but in the literature is known as the ‘longwave clear-sky feedback’ as it ignores cloud feedbacks and shortwave feedbacks, is rather universal and occurs ubiquitously throughout observational and modeling studies.<sup>51</sup> Furthermore,  $\lambda_{\text{RCE}}$  embodies the water vapor feedback discovered by MW67; by holding RH rather than specific humidity fixed, Simpson’s Law becomes applicable and tells us that a *significant portion of the longwave spectrum does not contribute to  $\lambda$  because  $T_{\text{em}}(\nu)$  is fixed* (Fig. 5b). This significantly reduces  $\lambda$  from the naive blackbody estimate (15) by a factor of about 2, consistent with MW67’s early finding that the water vapor feedback doubles climate sensitivity.

Indeed, a feedback parameter of  $\lambda_{\text{RCE}} = -2 \text{ (W/m}^2\text{)/K}$ , combined with  $\mathcal{F}_{2\times} = 4 \text{ W/m}^2$ , yields an estimated equilibrium climate sensitivity of

$$\text{ECS}_{\text{RCE}} = - \frac{\mathcal{F}_{2\times}}{\lambda_{\text{RCE}}} \approx 2 \text{ K} . \quad (30)$$

This estimate is close to those obtained by MW67 (2.3–2.9 K), though somewhat smaller than the consensus value of 3 K. Given the evidence that the neglected cloud and shortwave feedbacks are likely positive, however (see next section), it is unsurprising that  $\text{ECS}_{\text{RCE}}$  is biased low, and it is perhaps best regarded as a lower bound on ECS. Indeed,  $2 \text{ }^\circ\text{C}$  is the lower bound of the very likely range of 2–5  $^\circ\text{C}$  found in the recent sixth assessment report of the IPCC.<sup>52</sup>

## 6 Beyond RCE: Additional feedbacks

In this less detailed section we sketch the phenomena unaccounted for in the RCE framework, discuss their impacts on  $\lambda$  and ECS as assessed with numerical simulations and observations, and discuss prospects for principled estimates similar in spirit to those presented above.

### 6.1 Clouds

A major omission from the framework developed so far is clouds. Clouds exert enormous leverage over the climate system by absorbing and emitting longwave radiation essentially as blackbodies, and also by reflecting shortwave radiation (roughly half of Earth’s albedo is due to clouds<sup>53</sup>). In the longwave, certain aspects of the radiative effects of clouds can be described with a relatively simple formalism, as follows.

The longwave effects of clouds stem primarily from the high clouds arising from thunderstorms and mid-latitude winter storms; consistent with their height these clouds are very cold, with a typical cloud-top temperature  $T_{\text{cld}} = 220 \text{ K}$  (Fig. 2). Their Planck emission is thus much reduced relative to the clear-sky OLR one would see in their absence. If these high clouds cover a fraction  $f$  of the Earth’s surface ( $f \approx 0.18$  in the present-day global mean<sup>54</sup>), then one can write the ‘all-sky’ (i.e. actual) OLR as<sup>55</sup>

$$\text{OLR}_{\text{all}} = (1 - f)\text{OLR}_{\text{clr}} + f\sigma T_{\text{cld}}^4 . \quad (31)$$

In other words, high clouds mask a fraction  $f$  of  $\text{OLR}_{\text{clr}}$ , replacing it with cloud-top emission  $\sigma T_{\text{cld}}^4$ .

To differentiate Eq. (31) and obtain the all-sky longwave feedback  $\lambda_{\text{all}}$ , one needs to know how  $f$  and  $T_{\text{cld}}$  respond to warming. Fortunately, the latter question is answered quite simply by the so-called *fixed anvil temperature* hypothesis,<sup>56</sup> which is related to Simpson’s Law and says that high clouds rise with global warming so as to keep  $T_{\text{cld}}$  fixed, i.e.

$$\frac{dT_{\text{cld}}}{dT_{\text{s}}} \approx 0. \quad (32)$$

Differentiating Eq. (31) thus yields

$$\lambda_{\text{all}} = (1 - f)\lambda_{\text{clr}} + \frac{df}{dT_{\text{s}}}(\text{OLR}_{\text{clr}} - \sigma T_{\text{cld}}^4). \quad (33)$$

As for  $df/dT_{\text{s}}$ ,  $f$  is generally expected to decrease with warming (a feedback known as the ‘iris effect’),<sup>57</sup> but the magnitude of this decrease is uncertain and there is as yet insufficient theory to estimate it from first principles. The terms in Eq. (33) related to  $f$  and  $df/dT_{\text{s}}$  thus tend to compensate, and the value of  $\lambda_{\text{all}}$  ends up not far from  $\lambda_{\text{clr}} \approx -2$  (W/m<sup>2</sup>)/K but with larger error bars.<sup>58</sup>

On the shortwave side, there are highly reflective subtropical marine low clouds whose areal coverage is thought to decrease with global warming, yielding an increase in absorbed sunlight with warming and thus a positive contribution to the total feedback parameter of  $\lambda_{\text{cld}}^{\text{SW}} \sim 0.2$  (W/m<sup>2</sup>)/K.<sup>59</sup> This decrease in coverage is often understood via changes in environmental variables known as ‘cloud-controlling factors’ such as the local sea surface temperature and relative humidity; sophisticated analyses of these dependencies allow us to quantify the associated feedback.<sup>60</sup> Meanwhile, these clouds have also been described by simplified ‘mixed-layer models’.<sup>61</sup> But, these dots so far remain unconnected, and a first principles estimate of the tropical marine low cloud feedback also remains unformulated.

## 6.2 Shortwave feedbacks

In addition to changes in sunlight reflected by clouds, there are other significant shortwave feedbacks (i.e. changes in absorbed solar radiation  $S$  with warming) left unaccounted for in the RCE framework. Perhaps the largest of these is the *surface-albedo* feedback, due primarily to decreasing snow and ice cover with warming, which manifests as changes in albedo  $\alpha$  [cf. Eq. (1)]. This yields a positive feedback  $\lambda_{\text{albedo}} \sim 0.3 - 0.4$  (W/m<sup>2</sup>)/K.<sup>62</sup> While highly idealized models of the ice-albedo feedback have existed for decades,<sup>63</sup> and comprehensive modeling studies reveal a close connection between this feedback and the seasonal cycle,<sup>64</sup> again this gap has not been bridged and a chalkboard estimate of the surface-albedo feedback has yet to be formulated.

Another shortwave feedback which receives less attention but is not insignificant is that due to shortwave absorption by water vapor. Though often neglected in introductory treatments such as that of Section 2, it turns out that water vapor absorbs a rather significant amount of near-infrared sunlight – around 80 W/m<sup>2</sup>.<sup>65</sup> Since the mass of water vapor in the atmosphere increases with warming [at a rate roughly dictated by the Clausius-Clapeyron relation (27)],<sup>66</sup> water vapor shortwave absorption also increases, reducing the amount of (near-infrared) sunlight reflected out to space and thus increasing  $S$ . The end result is a positive shortwave water vapor feedback  $\lambda_{\text{wv}}^{\text{SW}} \sim 0.25$  (W/m<sup>2</sup>)/K.<sup>67</sup> A principled estimate of  $\lambda_{\text{wv}}^{\text{SW}}$  may be fairly easy to obtain, leveraging the fact that water vapor shortwave absorption should be a fixed function of temperature (i.e. obeys its own version of Simpson’s Law), in analogy to water vapor longwave emission.<sup>68</sup>

### 6.3 The total feedback

Putting the RCE (or longwave clear-sky) estimate (29) together with the cloud and shortwave feedbacks outlined above yields an estimate for the total feedback parameter:

$$\lambda_{\text{tot}} \approx \underbrace{\lambda_{\text{RCE}}}_{-2} + \underbrace{\lambda_{\text{clد}}^{\text{SW}}}_{0.2} + \underbrace{\lambda_{\text{albedo}}}_{0.3-0.4} + \underbrace{\lambda_{\text{wv}}^{\text{SW}}}_{0.25} \approx -1.2 \text{ (W/m}^2\text{)/K} . \quad (34)$$

This is close to the value of  $\lambda = -1.3 \text{ (W/m}^2\text{)/K}$  assessed by Sherwood et al.<sup>69</sup> from multiple lines of evidence, which indeed yields the consensus ECS value of  $-\mathcal{F}_{2\times}/\lambda = (4 \text{ W/m}^2)/(1.3 \text{ (W/m}^2\text{)/K}) \approx 3 \text{ K}$ . This more realistic value of  $\lambda$  will be a key ingredient in estimating other measures of climate sensitivity, which we take up in the next section.

## 7 Beyond ECS: other measures of climate sensitivity

In this final section we look beyond ECS to other measures of climate sensitivity. We will find that ECS is a quite idealized notion, and that other measures of climate sensitivity are more relevant for present-day warming and for understanding and defining emissions targets. But we will also see that ECS, as well as its key ingredients  $\mathcal{F}$  and  $\lambda$ , naturally appear in these other measures. Thus, the basic understanding of ECS developed here is necessary for understanding these other measures, and hopefully points the way to a deeper understanding of them.

### 7.1 The deep ocean and timescales of global warming

The equilibrium climate sensitivity is exactly that: an equilibrium quantity. But how *long* does it take the climate system to equilibrate with a given CO<sub>2</sub> concentration, and what does this evolution look like? To address this, we need a time-dependent model of the Earth’s surface temperature  $T_s$ . We proceed by neglecting the dynamics of the land surface (since the Earth is roughly 2/3 ocean covered) and invoking the popular *two-layer* or *two-box* model for the ocean.<sup>70</sup> This model consists of a shallow mixed layer with depth  $h_{\text{ml}} \approx 50 \text{ m}$  and temperature anomaly  $T'_{\text{ml}}$  equal to  $\Delta T_s$ , sitting atop a much larger deep ocean with global average depth  $h_d \approx 2500 \text{ m}$  and temperature anomaly  $T'_d$ . The model is pictured in Fig. 6.

An important characteristic of this model is that if the mixed layer is warmed by a forcing  $\mathcal{F}$ , it both radiates extra energy to space at a rate of  $|\lambda|T'_{\text{ml}}$  and also *exports energy to the deep ocean*, which we parameterize in linearized form as  $\gamma(T'_{\text{ml}} - T'_d)$  (units of W/m<sup>2</sup>). Here  $\gamma$  is the ‘deep ocean heat uptake efficiency’, estimated from models at roughly  $0.7 \text{ (W/m}^2\text{)/K}$ .<sup>71</sup> Setting  $\rho_w$  and  $C_w$  as the densities and specific heat capacities of water, the corresponding equations are

$$\rho_w C_w h_{\text{ml}} \frac{dT'_{\text{ml}}}{dt} = \mathcal{F} - |\lambda|T'_{\text{ml}} - \gamma(T'_{\text{ml}} - T'_d) \quad (35a)$$

$$\rho_w C_w h_d \frac{dT'_d}{dt} = \gamma(T'_{\text{ml}} - T'_d) . \quad (35b)$$

Because the deep ocean has a much larger depth and hence heat capacity than the mixed layer ( $h_d \gg h_{\text{ml}}$ ), we expect  $T'_{\text{ml}}$  to respond to  $\mathcal{F}$  much faster than  $T'_d$  does. Indeed, assuming  $T'_d = 0$

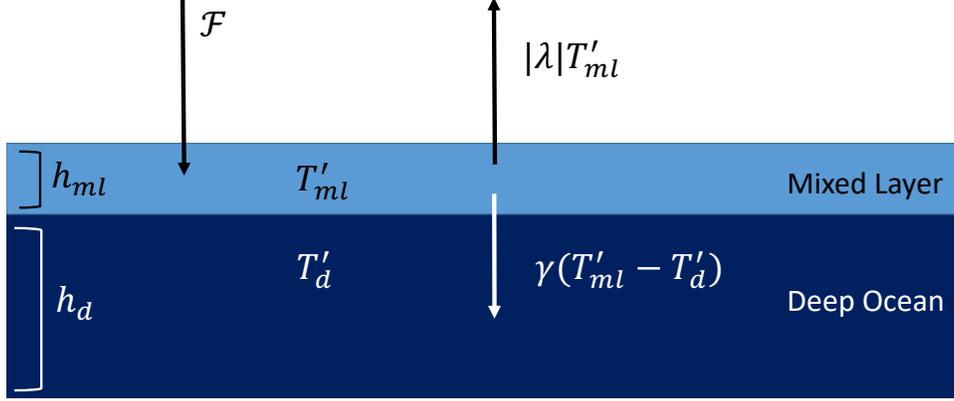


Figure 6: Two-box model for the ocean. See text for discussion.

in Eq. (35a) yields a linear, constant coefficient ordinary differential equation with characteristic timescale

$$\tau_{ml} = \frac{\rho_w h_{ml} C_w}{\gamma + |\lambda|} \approx 3 \text{ years} . \quad (36)$$

If we now consider timescales longer than  $\tau_{ml}$ , we may set  $dT'_{ml}/dt = 0$  in Eq. (35a), solve for  $T'_{ml}$ , plug the result into Eq. (35b), and solve for the characteristic deep ocean timescale  $\tau_d$ . This makes a nice exercise, with result

$$\tau_d = \rho_w h_d C_w \frac{\gamma + |\lambda|}{\gamma |\lambda|} \approx 700 \text{ years} . \quad (37)$$

Thus the vast difference in total heat capacity between the mixed layer and deep ocean indeed leads to two timescales for global warming: a ‘fast’ timescale of about  $\tau_{ml} \approx 3$  years during which the mixed layer equilibrates (really a ‘quasi-equilibrium’, since this equilibrium state will change as the deep ocean slowly responds), and a ‘slow’ timescale of  $\tau_d \approx 700$  years during which the deep ocean equilibrates (a true equilibrium). On intermediate timescales in between  $\tau_{ml}$  and  $\tau_d$  in which the mixed layer is in quasi-equilibrium, we can assume both  $dT'_{ml}/dt = 0$  and  $T'_d \approx 0$ , which from Eq. (35a) yields

$$T'_{ml} \approx \frac{\mathcal{F}}{\gamma + |\lambda|} \quad (\text{quasi-equilibrium}) . \quad (38)$$

This gives the transient warming which occurs before the deep ocean has responded. Indeed, if we specialize to the case of a  $\text{CO}_2$  doubling, the temperature in Eq. (38) becomes a standard sensitivity metric known as the ‘transient climate response’, or TCR:<sup>72</sup>

$$\text{TCR} = \frac{\mathcal{F}_{2\times}}{\gamma + |\lambda|} \approx 2 \text{ K} . \quad (39)$$

This estimate is quite close to the value of 1.8 K found in the recent sixth assessment report of the IPCC.<sup>73</sup> Comparing (39) with (13) shows that  $\text{ECS} > \text{TCR}$ : on the intermediate timescales during

which  $T'_{\text{ml}} = \text{TCR}$ , the mixed layer is both radiating heat to space *and* exporting heat to the deep ocean (Fig. 6), and can thus come to (quasi-)equilibrium at a lower temperature. This is of course not a true equilibrium state, and one can interpret the ratio TCR/ECS as a measure of the ocean’s thermal disequilibrium; this measure will prove useful in Section 7.2.

A key assumption in Eqns. (38) and (39) is that  $T'_d \approx 0$  on intermediate timescales. This approximation turns out to be a reasonable description of the present day, and the quasi-equilibrium formula (38) (which is just a scaling of TCR) can be used to credibly model historical as well as near-term global warming.<sup>74</sup> In these ways TCR is a more relevant metric for present day climate change than ECS, which instead assumes that both the mixed layer *and* deep ocean have reached a mutual equilibrium, which from Eq. (37) would take many hundreds of years.<sup>75</sup>

Before moving on, it should be remarked that the two-box model, while popular, is by no means canonical. Some recent work instead employs three-box models, and the older literature often employed diffusive models.<sup>76</sup> Two-box, three-box, and diffusive models were all employed by the IPCC as emulators of more comprehensive models.<sup>77</sup> All these approaches, however, require empirically determined parameters for heat transfer coefficients and diffusivities, analogous to our heat uptake efficiency  $\gamma$ , and principled estimates for these quantities are still lacking.<sup>78</sup>

## 7.2 The carbon cycle and measures of carbon-climate sensitivity

In addition to only describing very long-term warming, another limitation of ECS is that it assumes that the perturbed  $\text{CO}_2$  concentration  $q$  is *constant* while the Earth system equilibrates. If we stop burning fossil fuels, however,  $q$  will not remain constant; the real Earth has an active carbon cycle in the land and ocean, both of which absorb significant amounts of anthropogenic  $\text{CO}_2$ , which would cause  $q$  to decrease over time. Full consideration of these dynamics leads to two additional measures of climate sensitivity which are fundamental for both climate change science as well as policy: the *transient climate response to cumulative emissions* (TCRE) and the *zero emissions commitment* (ZEC).

TCRE is defined to be the warming at a given time divided by the *cumulative* emissions released prior to that time, in Kelvins per gigaton of carbon (K/GtC). TCRE is found to have a characteristic value of  $\sim 2$  K/1000 GtC, which in simulations turns out to be remarkably invariant over time as well as insensitive to emissions scenario.<sup>79</sup> The robustness of TCRE tells us that any identified temperature target (e.g.  $\Delta T_s = 1.5$  or  $2$  °C) automatically reduces to a cumulative emissions target (e.g. 750 or 1000 GtC), which can only be met if we *cease* emissions prior to reaching the target. This leads directly to the notion of net zero emissions.<sup>80</sup>

It is possible to use the TCR of section 7.1, as well as the logarithmic scaling of  $\text{CO}_2$  forcing, to make a back-of-the-envelope estimate of TCRE as follows. Let  $C_{\text{emit}}$  be our cumulative carbon emissions since the preindustrial era (GtC),  $C_{\text{pre}}$  the preindustrial mass of  $\text{CO}_2$  in the atmosphere (GtC), and  $\alpha$  be the fraction of  $C_{\text{emit}}$  still residing in the atmosphere (the ‘airborne fraction’), so that the total mass of  $\text{CO}_2$  in the atmosphere is  $C_{\text{pre}} + \alpha C_{\text{emit}}$ . Recalling that in the quasi-equilibrium approximation, surface warming  $\Delta T_s$  can be obtained by scaling the TCR by  $\mathcal{F}/\mathcal{F}_{2\times}$

[cf. Eqs. (38) and (39)], we then have

$$\begin{aligned}
\text{TCRE} &\equiv \frac{\Delta T_s}{C_{\text{emit}}} \\
&= \frac{\mathcal{F}}{\mathcal{F}_{2\times}} \frac{\text{TCR}}{C_{\text{emit}}} && \text{by (38) and (39)} \\
&= \frac{\ln\left(\frac{C_{\text{pre}} + \alpha C_{\text{emit}}}{C_{\text{pre}}}\right)}{\ln 2} \frac{\text{TCR}}{C_{\text{emit}}} && \text{by (23)} \\
&\approx \frac{\alpha}{\ln 2} \frac{\text{TCR}}{C_{\text{pre}}} && \text{using } \ln(1+x) \approx x. \quad (40)
\end{aligned}$$

In the last equality  $C_{\text{emit}}$  drops out of this expression, giving some insight into why simulations find TCRE to be roughly invariant over time.<sup>81</sup> Furthermore, evaluating Eq. (40) for  $C_{\text{pre}} = 590$  GtC and the present day value of  $\alpha = 0.4$  indeed yields roughly 2 K/GtC, consistent with simulations.<sup>82</sup> But, here we are simply using the observed value of  $\alpha$ ; we do not yet have ways to make principled estimates of  $\alpha$  from basic carbon cycle dynamics, and thus to understand how and why  $\alpha$  does or does not vary across time and emissions scenarios.

The Zero Emissions Commitment (ZEC), on the other hand, is complementary to TCRE and is defined to be the amount of residual warming which occurs *after emissions cease*. Simulations<sup>83</sup> tend to show that ZEC is small relative to  $\Delta T_s$ , which is a requirement for net zero emissions goals to meaningfully limit  $\Delta T_s$ . Similar to the estimate for TCRE, we may scale TCR and now ECS to estimate ZEC, following an argument due to N. Tarshish.<sup>84</sup> Let  $T_{ze}$  and  $\alpha_{ze}$  be the temperature and airborne fraction of cumulative emissions when emissions cease, and  $T_f$  and  $\alpha_f$  the temperature and airborne fraction at final equilibrium. Normalizing  $ZEC \equiv T_f - T_{ze}$  by  $T_{ze}$ , we have

$$\begin{aligned}
\frac{ZEC}{T_{ze}} &= \frac{T_f}{T_{ze}} - 1 \\
&= \frac{\ln\left(\frac{C_{\text{pre}} + \alpha_f C_{\text{emit}}}{C_{\text{pre}}}\right) \text{ECS}}{\ln\left(\frac{C_{\text{pre}} + \alpha_{ze} C_{\text{emit}}}{C_{\text{pre}}}\right) \text{TCR}} - 1 && \text{by (13), (23), (39)} \\
&\approx \frac{\alpha_f / \alpha_{ze}}{\text{TCR}/\text{ECS}} - 1. \quad (41)
\end{aligned}$$

This tells us that the sign and magnitude of ZEC is determined by a competition between the chemical disequilibrium  $\alpha_f / \alpha_{ze}$  and thermal disequilibrium  $\text{TCR}/\text{ECS}$  when emissions cease. If the chemical disequilibrium is more pronounced than the thermal disequilibrium, i.e.  $\alpha_f / \alpha_{ze} < \text{TCR}/\text{ECS}$ , then the decline in  $\text{CO}_2$  forcing due to equilibrium carbon uptake will dominate over the reduction in the deep ocean heat sink, yielding  $ZEC < 0$ , and vice-versa if  $\alpha_f / \alpha_{ze} > \text{TCR}/\text{ECS}$ . Noting that  $\alpha_f \approx 0.2$  is a characteristic value which may be obtained from principles of carbonate chemistry,<sup>85</sup> the previously cited values  $\alpha_{ze} = 0.4$ ,  $\text{TCR} = 1.8$  K, and  $\text{ECS} = 3$  K then give  $ZEC/T_{ze} \approx -0.17$ , which is indeed small (and, perhaps surprisingly, negative).

While Eq. (41) appears to explain why ZEC is small, it is however only a proximal explanation. The heat uptake efficiency  $\gamma$ , which entered into our estimate Eq. (39) of TCR, as well as the present day airborne fraction  $\alpha \approx 0.4$ , were evaluated via simulations or observations rather than theoretically, so we do not yet have fully principled estimates for TCRE or ZEC. Such estimates would necessarily draw upon physics from across the Earth system, including the dynamics of ocean

heat uptake, ocean carbon uptake, CO<sub>2</sub> fertilization of the biosphere, and more. A chalkboard explanation of these quantities thus poses a grand challenge to climate science, towards which the simple models presented here are simply a first step.

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## A Adiabatic temperature profiles

We here derive the expression (5) for the dry adiabatic lapse rate. For a textbook treatment of this topic, as well as extension to include the effects of moisture, see e.g. the textbook by Wallace and Hobbs<sup>87</sup> or the excellent lecture notes of Romps<sup>88</sup>.

Consider a rising convective parcel that conserves its mass and rises rapidly such that heat exchange with the environment is negligible. This process is governed by the first law of thermodynamics

$$dQ = dU + p dV \quad (42)$$

where  $dQ$  is the heat gained by the system,  $U$  is its internal energy,  $p$  is its pressure and  $V$  its volume. For an ideal gas we have  $U = \rho V C_v T$  as well as the ideal gas law, written in the form typical of atmospheric science

$$p = \rho R_d T \quad (43)$$

where  $\rho$  is the parcel density in kg/m<sup>3</sup>,  $R_d = 287$  J/kg/K is the specific gas constant for dry air (obtained by dividing the universal gas constant by the molar mass of air), and  $C_v = \frac{5}{2} R_d$  is the specific heat capacity at constant volume. Assuming an adiabatic process ( $dQ = 0$ ), we rearrange Eq. (42) into

$$0 = \rho C_p dT - dp$$

where the specific heat at constant pressure  $C_p = C_v + R_d \approx 1000$  J/kg/K. Further assuming hydrostatic balance,<sup>89</sup> which says that the weight of a layer of air is balance by the vertical pressure gradient across it,

$$\frac{dp}{dz} = -\rho g \quad (44)$$

and rearranging then yields Eq. (5):

$$\Gamma_d \equiv -\frac{dT}{dz} = \frac{g}{C_p}.$$

Now consider an arbitrary but constant lapse rate  $\Gamma$ . Then, we have

$$\frac{dT}{dp} = \frac{dT}{dz} \frac{dz}{dp} = \frac{\Gamma}{\rho g} = \frac{\Gamma R_d T}{p g} \quad (45)$$

where Eq. (44) was used in the second equality and Eq. (43) in the third. This relation between  $T$  and  $p$  can be integrated from an arbitrary pressure  $p$  to surface pressure  $p_s$  to obtain Eq. (10b).

## B Analytic derivation of formula for CO<sub>2</sub> forcing

The forcing  $\mathcal{F}$  is usually obtained by numerical integration in  $\nu$  across the CO<sub>2</sub> band shown in Fig. 4. Here we will instead pursue an analytical integration over  $\nu$ , following the thesis of Seeley.<sup>90</sup> We will take advantage of the symmetry evident in Fig. 4 and simply double the integral for  $\nu > \nu_0$ . We will also take advantage of the key fact that  $q$  and  $\kappa_{\text{CO}_2}(\nu)$  appear in Eq. (20) as a product; this, along with the exponential spectroscopy (19) and the fact that  $T_{\text{em}}$  and  $p_{\text{em}}$  are related by Eq. (10b), mean that

$$\frac{dT_{\text{em}}}{d \ln q} = \frac{dT_{\text{em}}}{d \ln \kappa_{\text{CO}_2}} = -l \frac{dT_{\text{em}}}{d\nu}. \quad (46)$$

This equation encapsulates the fact, mentioned towards the end of section 4, that a multiplicative change in  $q$  is equivalent to that of a uniform translation in  $\nu$ . Putting these ingredients together, the forcing Eq. (11) can now be evaluated as an integral over  $\nu$  from band center  $\nu_0$  to the upper limit  $\nu_+ \approx 750 \text{ cm}^{-1}$ , where we ignore spectral variations in the Planck density. This yields

$$\begin{aligned} \mathcal{F} &= -(\Delta \ln q) \frac{d\text{OLR}}{d \ln q} \\ &= -2 \ln(q_f/q_i) \int_{\nu_0}^{\nu_+} \frac{d\text{OLR}_\nu}{d \ln q} d\nu \\ &= -2 \ln(q_f/q_i) \int_{\nu_0}^{\nu_+} \pi \frac{\partial B}{\partial T} \frac{dT_{\text{em}}}{d \ln q} d\nu \quad \text{by Eq. (16)} \\ &= 2l \ln(q_f/q_i) \int_{\nu_0}^{\nu_+} \pi \frac{\partial B}{\partial T} \frac{dT_{\text{em}}}{d\nu} d\nu \quad \text{by Eq. (46)} \\ &= 2l \ln\left(\frac{q_f}{q_i}\right) [\pi B(\nu_0, T_s) - \pi B(\nu_0, T_{\text{strat}})] \end{aligned}$$

where in the last line we invoked the fundamental theorem of calculus, noting also that  $T_{\text{em}}(\nu_+) = T_s$  and  $T_{\text{em}}(\nu_0) = T_{\text{strat}}$ .

## Notes

<sup>1</sup>Arrhenius (1896)

<sup>2</sup>Sherwood et al. (2020); Forster et al. (2021)

<sup>3</sup>The report of Charney et al. (1979) was the first such model intercomparison report. Meehl et al. (2020) reviews ECS over the many generation of IPCC reports. The latest IPCC assessment of ECS is given in Chapter 7 of the sixth assessment report from working group I, Forster et al. (2021) section 7.5; see also the comprehensive ECS review in Sherwood et al. (2020).

<sup>4</sup>Manabe and Wetherald (1967)

<sup>5</sup>Jeevanjee et al. (2022)

<sup>6</sup>see e.g. Wilson and Gea-Banacloche (2012); Jeevanjee et al. (2021b); Romps et al. (2022)

<sup>7</sup>see, e.g. Koll and Cronin (2018); Jeevanjee (2018); Jeevanjee et al. (2021a)

<sup>8</sup>See also Stevens and Kluft (2023) for a similar, but not explicitly pedagogical, approach.

<sup>9</sup>Hartmann (2015)

<sup>10</sup>Pierrehumbert (2011)

<sup>11</sup>For an example of this, see He et al. (2022) in which the formalism for CO<sub>2</sub> forcing developed here is applied to understand the spread in that quantity amongst comprehensive climate models.

<sup>12</sup>See also Emanuel (2020) for a related point of view.

<sup>13</sup>Trenberth and Fasullo (2009); Stephens et al. (2015)

<sup>14</sup>See section 6.2 and note 65 for more on this approximation.

<sup>15</sup>e.g. Petty (2006); Randall (2012); Vallis (2012); Coakley Jr. and Yang (2014); Hartmann (2015); Tziperman (2022)

<sup>16</sup>Thermodynamically speaking this is a steady-state, not an equilibrium, as the net fluxes of e.g. solar radiation, thermal radiation, water vapor at the Earth’s surface etc. are nonzero. Accordingly, the Earth does not come into thermal equilibrium with either the Sun or outer space, but tends towards a temperature in between them. Nonetheless, the use of the term ‘equilibrium’ to describe a steady-state in climate is ubiquitous, so we adopt this terminology here.

<sup>17</sup>It is a nice exercise to derive this height analytically, as follows: assume an isothermal atmosphere with  $T = T_{\text{em}}$ , which from Eqs. (44) and (43) yields the exponential dependence

$$p = (1 \text{ atm}) \exp(-z/H), \quad (47)$$

where  $H \equiv R_d T_{\text{em}}/g \approx 7.5 \text{ km}$  is the atmospheric ‘scale height’. Setting  $p_a = 0.5 \text{ atm}$  then yields  $z_a = H \ln 2 \approx 5 \text{ km}$ . Note also that in this isothermal approximation, and by the ideal gas law (43),  $\rho \sim p/T$  also declines exponentially with height.

<sup>18</sup>Hansen et al. (1981)

<sup>19</sup>Payne et al. (2015)

<sup>20</sup>Stratospheric temperatures are sometimes thought to be governed by the so-called ‘skin temperature’, which arises by considering an optically thin layer of atmosphere sitting atop the troposphere; a standard argument yields a skin temperature of  $T_{\text{em}}/\sqrt[4]{2} = 214 \text{ K}$  (Pierrehumbert, 2011; Hartmann, 2015). This argument ignores the strong dependence of tropopause and stratospheric temperatures on atmospheric composition, however (e.g. Manabe and Strickler, 1964). Recent work instead postulates that  $T_{\text{strat}}$  is governed by the physics of H<sub>2</sub>O radiative transfer (Seeley et al., 2019; Jeevanjee and Fueglistaler, 2020b), but more work on this topic is needed.

<sup>21</sup>At least to first approximation. See discussion in Jeevanjee et al. (2022).

<sup>22</sup>This is the ratio of the mass of a given substance in a parcel to the total mass of the parcel. See Pierrehumbert (2011), pg. 87, for further discussion.

<sup>23</sup>While such a perspective ignores the interactive carbon cycle dynamics of the atmosphere, land, and ocean, it is nonetheless a useful starting point for understanding the impacts of fossil-fuel combustion on Earth’s climate. We discuss how carbon cycle dynamics modify the picture presented here in Section 6.

<sup>24</sup>There are, however, processes known as atmospheric *adjustments* which blur this distinction; see Sherwood et al. (2015).

<sup>25</sup>An important exception to this reasoning is the *stratospheric adjustment*, which is the direct response of stratospheric temperatures to a change in  $q$  resulting from the increased emission to space depicted in Fig. 4b (green dashed lines). This change in stratospheric temperatures occurs independently of changes in  $T_s$ , precisely because the stratosphere is not coupled to the surface in the way the troposphere is. For further discussion, see Hansen et al. (1997); Houghton et al. (1994).

<sup>26</sup>Another useful consequence of this definition is that ‘positive’, amplifying feedbacks make positive contributions to  $\lambda$

<sup>27</sup>Charney et al. (1979); Myhre et al. (1998); Ramaswamy et al. (2019).

<sup>28</sup>We will not need the specific form of  $B(\nu, T)$ , but for completeness it is given by

$$B(\nu, T) = \frac{2hc^2\nu^3}{\exp\left(\frac{hc\nu}{k_b T}\right) - 1} \quad (48)$$

where  $h$  is Planck’s constant,  $k_b$  is Boltzmann’s constant, and  $c$  is the speed of light.

<sup>29</sup>Since CO<sub>2</sub> is well-mixed its density is proportional to the air density  $\rho$ , and  $\rho$  decreases exponentially with height (cf. note 17). For H<sub>2</sub>O, its density is dominated by its Clausius-Clapeyron exponential dependence on temperature (Eqs. (26) and (27)), and temperature is linear in height, so H<sub>2</sub>O density is exponential in height as well.

<sup>30</sup>We here assume a two-stream approximation with unit diffusion coefficient for simplicity; Pierrehumbert (2010)

<sup>31</sup>This pressure scaling is due to collisional pressure broadening away from spectral line centers, and is also accompanied by additional, typically less pronounced temperature scalings. See Pierrehumbert (2010).

<sup>32</sup>For further analysis of the emission level approximation, with analytic calculations suitable for the classroom, see Appendix B of Jeevanjee et al. (2021b) and Appendix B of Jeevanjee and Fueglistaler (2020a).

<sup>33</sup>Wilson and Gea-Banacloche (2012)

<sup>34</sup>Jeevanjee et al. (2021b)

<sup>35</sup>Romps et al. (2022)

<sup>36</sup>Jeevanjee et al. (2021b)

<sup>37</sup>For a pedagogical discussion of the structure of this spectrum and its computation, see Wilson and Gea-Banacloche (2012).

<sup>38</sup>We used the Reference Forward Model (Dudhia, 2017), along with spectroscopic data from HiTRAN2016 (Gordon et al., 2017).

<sup>39</sup>We used the Reference Forward Model (Dudhia, 2017), along with spectroscopic data from HiTRAN2016 (Gordon et al., 2017).

<sup>40</sup>This estimate ignores several effects, each of which generate 20-30% corrections but which offset each other. These effects include the overlapping of CO<sub>2</sub> absorption by H<sub>2</sub>O absorption, the masking of CO<sub>2</sub> forcing by clouds, and stratospheric adjustment. See Jeevanjee et al. (2021b); Huang et al. (2016).

<sup>41</sup>The H<sub>2</sub>O absorption coefficients shown here have been simplified by neglecting so-called ‘continuum’ absorption. Continuum absorption significantly affects absorption in the infrared window at warmer surface temperatures of 300 K and above. See Pierrehumbert (2010); Shine et al. (2012); Koll et al. (2022).

<sup>42</sup>This fixed relative humidity assumption, popularized by Manabe and Wetherald (1967), has been well justified by decades of subsequent simulations and observation (Jeevanjee et al., 2022; Colman and Soden, 2021).

<sup>43</sup>In particular we multiply the integrand by  $T_{\text{av}}/T'$  which does not deviate too far from 1 but which allows the exponential to be integrated. Since the integrand is dominated by values of the exponential evaluated near  $T$ , we also approximate  $p(T') \approx p(T)$  so it can be pulled outside the integral. This expression and these approximations are discussed in more detail in Jeevanjee and Fueglistaler (2020b)

<sup>44</sup>Eq. (28) can be inverted at  $\tau = 1$  using the Lambert W function to obtain

$$T_{\text{em}}(\nu) = \frac{T^*}{W\left(\frac{T^*}{T_{\text{ref}}}(D M_v^{\text{ref}} \kappa_{\text{H}_2\text{O}}(\nu))^{R_d \Gamma/g}\right)}, \quad \text{where } T^* = \frac{L R_d \Gamma}{g R_v}.$$

See also Jeevanjee and Fueglistaler (2020b).

<sup>45</sup>See note 39.

<sup>46</sup>See note 39.

<sup>47</sup>See e.g. Simpson (1928). See also Ingram (2010); Jeevanjee et al. (2021a) for further discussion.

<sup>48</sup>These deviations can be estimated by calculating  $\frac{dT_{\text{em}}}{dT_s}$  implicitly from Eq. (28) and invoking Eq. (10b). This makes a nice exercise, with result

$$\frac{dT_{\text{em}}}{dT_s} = \frac{T_{\text{em}}}{T_s} \frac{1}{\frac{L R_d \Gamma}{g R_v T_{\text{em}}} - 1} \approx \frac{1}{4}. \quad (49)$$

<sup>49</sup>Petty (2006)

<sup>50</sup>This approach has precedent in the work of Koll and Cronin (2018); Ingram (2010).

<sup>51</sup>Koll and Cronin (2018); Zhang et al. (2020); McKim et al. (2021); Colman and Soden (2021)

<sup>52</sup>Forster et al. (2021), Table 7.13

<sup>53</sup>Stephens et al. (2015)

<sup>54</sup>We obtain  $f = 0.18$  as an average of the AIRS and ISCCP satellite data values for global mean high cloud fraction given in Fig. 4.5 of Siebesma et al. (2020).

<sup>55</sup>See McKim et al. (2021) for a validation of this simple approximation for the all-sky OLR.

<sup>56</sup>Hartmann and Larson (2002); Thompson et al. (2017); Jeevanjee and Fueglistaler (2020b)

<sup>57</sup>Mauritsen and Stevens (2015); Bony et al. (2016)

<sup>58</sup>e.g. Andrews et al. (2015); Flynn and Mauritsen (2020)

<sup>59</sup>Zelinka et al. (2016); Klein et al. (2017); Sherwood et al. (2020)

<sup>60</sup>Myers et al. (2021); Ceppi and Nowack (2021)

<sup>61</sup>Lilly (1968); Bretherton and Wyant (1997)

<sup>62</sup>Zelinka et al. (2020); Colman (2013)

<sup>63</sup>North et al. (1981)

<sup>64</sup>Hall (2004)

<sup>65</sup>Trenberth and Fasullo (2009). It is an interesting exercise with the RCE model Eq. (8) to show that subtracting this from the surface energy budget and adding it to the atmospheric energy budget does not affect  $T_s$ , consistent with the interpretation that the atmosphere and surface function as a unit so repartitioning energy between them does not affect their temperatures.

<sup>66</sup>Held and Soden (2006)

<sup>67</sup>Donohoe et al. (2014)

<sup>68</sup>Jeevanjee (2018)

<sup>69</sup>Sherwood et al. (2020)

<sup>70</sup>e.g. Schneider and Thomsson (1981); Gregory (2000); Held et al. (2010); Armour (2017)

<sup>71</sup>Geoffroy et al. (2013).

<sup>72</sup>Strictly speaking, TCR is defined via simulations in which CO<sub>2</sub> is increased from its preindustrial value at 1% year until the concentration has doubled, and TCR is defined as the warming at the time of doubling (around year 70). But, it has been shown that the two-box expression (39) for TCR captures the actual TCR fairly well (Held et al., 2010; Geoffroy et al., 2013)

<sup>73</sup>citeforster2021, section 7.5.5

<sup>74</sup>Held et al. (2010); Geoffroy et al. (2013).

<sup>75</sup>Indeed, it is a simple exercise to show that imposing the full equilibrium conditions  $dT'_{\text{ml}}/dt = dT'_d/dt = 0$  in Eqns. (35) for  $\mathcal{F} = \mathcal{F}_{2\times}$  yields  $T'_{\text{ml}} = T'_d = \mathcal{F}_{2\times}/|\lambda| = \text{ECS}$ .

<sup>76</sup>Three-box models were employed in e.g. Proistosescu and Huybers (2017); Cummins et al. (2020); Tsutsui (2020). Diffusive models were employed in e.g. Hansen (1984); Siegenthaler and Oeschger (1984); Wigley and Schlesinger (1985)

<sup>77</sup>See Chapter 7 of the IPCC sixth assessment report (Forster et al., 2021), and in particular cross-chapter box 7.1. Two popular emulators employed there are MAGICC (Meinshausen et al., 2011), which has a diffusive deep ocean, and FaIR (Leach et al., 2021), which has a three-box ocean.

<sup>78</sup>Though there has been some work in this direction; see e.g. Marshall and Zanna (2014).

<sup>79</sup>MacDougall (2016)

<sup>80</sup>Allen et al. (2022)

<sup>81</sup>Additional insight can be found in the theoretical approaches of e.g. Goodwin et al. (2015); MacDougall and Friedlingstein (2015); Seshadri (2017).

<sup>82</sup>See Friedlingstein et al. (2020) for values of  $C_{\text{pre}}$  and  $\alpha$ . It is also a nice exercise to estimate  $C_{\text{pre}}$  by multiplying the mass of the atmosphere  $4\pi R_E^2 p_s/g$  by the specific CO<sub>2</sub> concentration corresponding to the number concentration of 280 ppm. Note also that for present-day  $C_{\text{emit}}$  of roughly 700 GtC, we have  $\alpha C_{\text{emit}}/C_{\text{pre}} \approx 0.45$ , so the approximation  $\ln(1 + \alpha C_{\text{emit}}/C_{\text{pre}}) \approx \alpha C_{\text{emit}}/C_{\text{pre}}$  used in Eq. (40) is marginally acceptable, but will fail for larger  $C_{\text{emit}}$ .

<sup>83</sup>MacDougall et al. (2020)

<sup>84</sup>See Tarshish et al. (2022)

<sup>85</sup>e.g. Goodwin et al. (2007)

<sup>86</sup>The original notes are available at Jeevanjee (2018) and contain additional material beyond that presented here.

<sup>87</sup>Wallace and Hobbs (2006)

<sup>88</sup>Romps (2020)

<sup>89</sup>There is some sleight-of-hand here, as hydrostatic balance typically applies to the pressure and density of the quiescent *environment*, rather than a rising parcel. For most applications (such as ours) this approximation is permissible, but if we take it to extremes (such as applying the dry adiabatic lapse rate Eq. (5) over large enough distances to generate negative temperatures) it can lead to nonsensical results; see Romps (2020) for further discussion.

<sup>90</sup>See section 2.3 of Seeley (2018).

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