# CO<sub>2</sub>-Equivalence and the Transition to Stabilisation

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#### **Abstract**

The relations between the various greenhouse gases are specified in terms of two distinct forms of CO<sub>2</sub>-equivalence: emission-equivalence and Emission-equivalence is usually expressed in concentration-equivalence. terms of radiative effect over a 100-year lifetime (i.e. a 100-year global warming Concentration-equivalence reflects the instantaneous radiative potential). effect of the concentrations and so corresponds to using the infinite-time limit of the global warming potential. These considerations are reflected in the concept that for stabilisation at any target concentration of CO<sub>2</sub>, there is a 'carbon budget', i.e. a limit to the total amount of carbon that can ever be emitted [1, 9, 10]. Such a restriction does not apply for shorter-lived gases such as N<sub>2</sub>O and CH<sub>4</sub>. A more detailed analysis of the stabilisation profiles for CO<sub>2</sub> [4, 2] goes beyond the 'fixed CO<sub>2</sub> budget' and the allowable residual emissions provide a basis for defining 'CO<sub>2</sub>-equivalent emissions' under stabilisation. The discussion draws on experience in contributing the Garnaut Review of Climate Change [6] in Australia.









## 1 Greenhouse gas equivalence?

Anthropogenic global warming is the consequence of a range of trace gases. Since mitigation measures entail costs, some quantitative measure is required for determining the relative importance of the gases in order to avoid costs that are disproportionate to the benefits. Two forms of CO<sub>2</sub>-equivalence have been defined [7: glossary] in order to address such needs:

concentration equivalence which is defined as equivalence in radiative forcing and is the property of a mixture of gases, and

emission equivalence which looks at the integrated radiative effect of emissions, and compares gases using the 'Global Warming Potential' (GWP) defined by:

$$GWP_X(\tau) = \frac{a_X}{a_{CO2}} \frac{\int_0^{\tau} R_X(t) dt}{\int_0^{\tau} R_{CO2}(t) dt}$$
 (1)

where  $a_X$  is the radiative forcing per kg of gas X and  $R_X(t)$  is a response function giving the proportion of gas X remaining in the atmosphere after time t — see 'Notation' section at end of poster.

Thus 'emissions equivalence' is really a family of definitions whose relations depend on the time horizon,  $\tau$ . The Kyoto Protocol defines CO<sub>2</sub>-equivalence using a 100-year GWP.

'Concentration-equivalence' also has a degree of variation in the definition, depending on whether or not various short-lived components such as aerosols and tropospheric ozone are included. One approach [2] is to include all components when analysing past and present conditions, and consider only the long-lived components when considering future commitments.

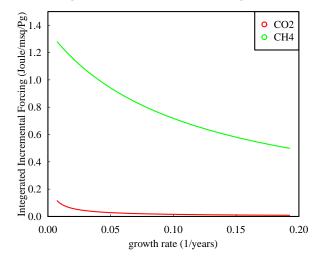
The limitations of setting targets on the basis of emission-equivalence are illustrated in specific examples by Reilly et al. [11], in particular in their figure 2a. Case 2' is stringent reductions of  $CO_2$  only, while 3' is the "economically optimum"  $CO_2$ -emission-equivalent reductions. Using  $CO_2$ -emission-equivalence leads to large differences in temperature outcomes (presumably because it leads to large differences in  $CO_2$ -equivalent concentrations).

# 2 Analysis

The Laplace Transform formalism provides a powerful way of analysing systems that are linear (or which can be linearised). In particular, in a linear system with fixed emission growth rate,  $\lambda$ ,  $r_X(p)$  (which is the Laplace transform of the response function  $R_X(t)$ ) can define an exact equivalence between greenhouse gases with a scaling  $\alpha$  given by

$$\alpha_X(\lambda) = \frac{a_X}{a_{CO2}} \frac{r_X(\lambda)}{r_{CO2}(\lambda)} = \frac{a_X}{a_{CO2}} \frac{\int_0^\infty e^{-\lambda t} R_X(t) dt}{\int_0^\infty e^{-\lambda t} R_{CO2}(t) dt}$$
(2)

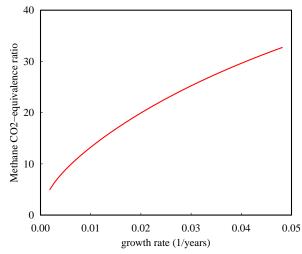
Thus when emissions (and thus the increase in concentrations) exhibit exponential growth of  $100\lambda$  percent per annum, emissions of each kg of  $CO_2$  are equivalent to emitting  $\alpha_X(\lambda)$  kg of gas X. While the time-horizon,  $\tau$ , in the GWP definition (1) is generally motivated in terms of time-scales of interest, treating the GWP definition (1) as an approximation to (2) suggests that the equivalence is most meaningful when the 'time-scale of interest' corresponds the e-folding time of emissions growth.



Laplace transform of radiative forcing from pulses of CO<sub>2</sub>, CH<sub>4</sub>.

Ratios of Laplace transform of radiative forcing from pulses of  $CH_4$  and  $CO_2$ , as function of 'growth rate', p

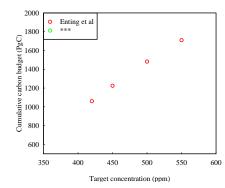
— the factor  $\alpha_{\rm CH4}$  defined by equation 2.



## 3 Carbon budgets and beyond

A number of analyses [e.g. 1, 6, 9, 10] have considered emission targets in terms of a 'carbon budget' — a limit to the total amount of carbon that can ever be emitted consistent with a given target concentration. There are two related considerations:

- the 'carbon budget' concept is a 'diagnostic statistic' of behaviour on times scales of order 100 years — it is not a mechanistic description of carbon cycle dynamics;
- it gives only an approximate description of stabilisation.



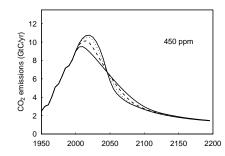
**Estimates** the 'carbon budget' for various target concentrations [2].

For others gases the budget concept is a much poorer approximation. Concentrations of non CO<sub>2</sub> gases can be stabilised with fixed emissions related to target mass,  $M_X^{\rm target}$ , and atmospheric lifetime,  $T_X$ , by:  $E_X^{\rm stabilisation} = M_X^{\rm target}/T_X$ 

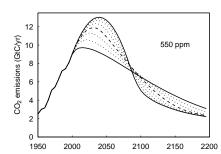
$$E_X^{\text{stabilisation}} = M_X^{\text{target}}/T_X$$

Thus equivalence between emissions of non-CO<sub>2</sub> gases corresponds to equivalence of  $a_X/T_X$ . This corresponds to the GWP with infinite time horizon, in which case, the GWPs of all non-CO<sub>2</sub> gases become zero — in the very long-term, only  $CO_2$  matters in this approximation.

CO<sub>2</sub> can be re-integrated into this framework by appreciating that the 'carbon budget' concept is indeed an approximation. The examples below show that a small residual emission rate is consistent with stabilising concentrations.



Estimates of carbon emissions for various target concentrations (from [2]).



In these terms, an 'emission equivalence' factor, appropriate for stabilisation can be represented as:

$$\alpha_X' = a_X/T_X / a_{\text{CO2}} \frac{\partial M_{\text{CO2}}^{\text{target}}}{\partial E_{\text{CO2}}^{\text{residual}}}$$
 (3)

#### 4 Feedbacks

Enting [3] used the Laplace Transform formalism to capture climate-to carbon feedbacks using response functions, U(t), giving the warming response to radiative forcing, and H(t) describing an additional  $\mathrm{CO}_2$  source from warming which is denoted W(t). Denoting their respective Laplace Transforms, u(p), h(p) and w(p)) gives:

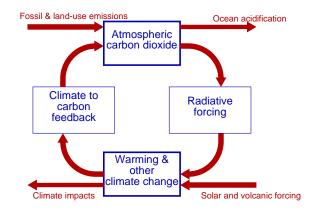
$$w(p) = u(p)[f(p) + a_{CO2} q(p)]$$
 (4)

$$q(p) = r(p)[s(p) + h(p) w(p)]$$
 (5)

whence

$$w(p) = \frac{u(p)f(p) + a_{CO2} u(p) r(p) s(p)}{1 - a_{CO2} u(p) r(p) h(p)}$$
(6)

$$q(p) = \frac{r(p)[s(p) + f(p) h(p) u(p)]}{1 - a_{CO2} u(p) r(p) h(p)}$$
(7)



Forcing from emissions, S(t), or radiative change, F(t), (with respective Laplace transforms s(p), f(p)) is amplified by a feedback factor:

$$\kappa_{\rm FB} = 1/[1 - a_{CO2} u(p) r(p) h(p)].$$

For multi-decadal time-scales, the C4MIP study [5] gives  $\kappa = 1.18 \pm 0.11$  from 11 models, ranging from  $\kappa =$  1.04 to 1.44 (with only the CO<sub>2</sub> term included).

Generalising this to consider  $H_X(t)$ , the emissions of gas X in response to warming shows that the effect of radiative forcing, f(p), or greenhouse gas emissions,  $s_X(p)$ , is amplified by feedback factor:

$$\kappa_{\rm FB} = \frac{1}{1 - u(p) \sum_X a_X r_X(p) h_X(p)} \tag{8}$$

Notionally, the carbon fluxes from feedbacks would need to be subtracted from emission targets in order to match targets calculated without reference to feedbacks.

However, to the extent that the  $CO_2$  response, R(t), r(p) has been calibrated against 20th century changes, some climate-carbon feedbacks will have been implicitly included [2], in which case, the  $CO_2$  feedbacks, at least on multi-decadal timescales, should be excluded from the sum in equation (8).

# **5 Beyond linearity?**

With anthropogenic climate change already occurring, and a commitment to further change due to past emissions, mitigation needs to be considered in terms of reducing, rather than avoiding, dangerous impacts.

There is considerable uncertainty in aspects of the carbon-climate system, both in the feedbacks and the underlying response, parameterised above through U(t), u(p). [u(0) is proportional to the climate sensitivity, while u(p>0) parameterises transient climate sensitivities.] Consequently, mitigation analysis needs to be in terms of ranges that span the uncertainty [e.g. 13].

There is great asymmetry in the distribution of risk associated with these ranges — damage functions grow faster than linearly and there is the possibility of instabilities, irreversibility and 'tipping points'.

For example although the 450 ppm ( $CO_2$  concentration equivalent) target, is often thought of as notionally 'safe' with 2°C warming, 450 ppm of actual  $CO_2$  increases pH in ocean to a point where aragonite becomes unstable even at the surface.

An 'extreme' possibility is reflected in Lovelock's view of a third hotter climate state [8], (in addition to the glacial and interglacial states). In a self-regulated system (which is how Lovelock's Gaia theory envisages the geosphere-biosphere system), once self-regulation fails, the establishment of a new conditions can be very abrupt.

Do these considerations change the way in which equivalence should be considered?

For example, can short-term reductions in  $CH_4$  (at the expense of delaying  $CO_2$  reductions) provide pathway to avoid tipping point? i.e. should emission-equivalence be defined with a shorter  $\tau$  ( $\ll 100$  years) rather than longer ( $\gg 100$  years) as is implied by previous analysis?

There would seem to be two related pre-conditions for adopting such a changed definition of CO<sub>2</sub>-equivalence:

- we know where the tipping point is; and
- in particular, the tipping point has not already been passed.

Neither of these can be asserted with any certainty.

# 6 Summing up

The two different forms of emission-equivalence each capture legitimate aspects of greenhouse gas forcing, but the existence of two approaches complicates discussion. In either case, CO<sub>2</sub>-equivalence must be regarded as a diagnostic statistic. There are severe constraints on using CO<sub>2</sub> emission-equivalence (based in GWPs) as a computational tool. The limitations are discussed in generic terms by Smith and Wigley [12] and illustrated in specific examples by Reilly et al. [11]. (The Garnaut economic analysis [6] used GWP-based 'emission equivalence' with the 100 year time horizon).

When GWPs are defined with increasingly longer time horizons and used to define CO<sub>2</sub>-equivalent emissions, the equivalence relations formally approach those defined by 'concentration-equivalence', but focus all constraints on CO<sub>2</sub>. This result that 'only CO<sub>2</sub> matters' is equivalent to noting that CO<sub>2</sub> has a fixed 'budget' while other gases do not. Appreciating that the CO<sub>2</sub> budget is only an approximate description of carbon cycle responses has the potential to give a practical quantification of CO<sub>2</sub> emission equivalence under conditions of stabilisation via equation (3). These principles remain applicable when linear feedbacks are considered. The location of potential 'tipping points' in the earth system remains too uncertain to be the basis of any more 'targetted' trade-off between emissions of the different gases.

#### **Notation**

- $a_X$  Radiative forcing per kg of gas X.
- $H_X(t)$  Response function specifying emissions of gas X from unit (global) temperature increase, with Laplace Transform  $h_X(p)$ .
- p Transform variable in Laplace Transform, in years<sup>-1</sup>.
- $R_X(t)$  Impulse response function of emissions of gas X, with Laplace Transform  $r_X(p)$ .
- $S_X(t)$  Emission history of gas X, with Laplace Transform  $S_X(p)$ .
- t Time, in years.
- $T_X$  Atmospheric lifetime of gas X.
- U(t) Temperature response for unit pulse of radiative forcing, with Laplace Transform u(p).
- $\kappa_{\rm FB}\,$  Amplification factor from feedback loop(s).
- $\lambda$  Growth rate for emissions, concentrations etc in linear approximation.
- $\xi \dots \xi_k$  Gain around feedback loop ... from process k.
- au Time horizon for defining GWPs.

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