

# Major Portion of the Increased Atmospheric Concentration of Carbon Dioxide Emitted by the Oceans

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## To cite this article:

Jyrki Kauppinen, Pekka Malmi. (2025). Major Portion of the Increased Atmospheric Concentration of Carbon Dioxide Emitted by the Oceans. *International Journal of Atmospheric and Oceanic Sciences*, 9(1), 66-69. <https://doi.org/10.11648/j.ijaos.20250901.17>

**Received:** 15 April 2025; **Accepted:** 10 May 2025; **Published:** 21 June 2025

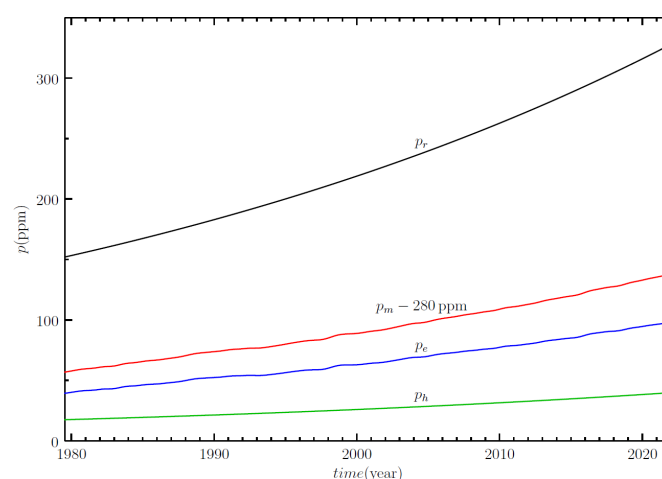
**Abstract:** This paper continues the application of Henry's Law to climate change. The basic ideas have been covered in our previous paper. We separately calculated the concentrations of carbon dioxide in the atmosphere as a function of time emitted by the oceans and human contribution. We know that the sum of those two concentrations was equal to the observed one. Note that the carbon dioxide emitted by the oceans depends only on the temperature. We can verify our theory by calculating the global temperature using only the observed atmospheric carbon dioxide emitted by the oceans, which is convincing evidence of Henry's Law. In this paper, we will present additional evidence using the measured abundance of  $^{13}\text{CO}_2$  in the atmosphere starting from the 18<sup>th</sup> century. We will also show that the concentrations of  $\text{N}_2\text{O}$  and  $\text{SF}_6$  follow Henry's Law and that human contributions are insignificant. Note that these concentrations can be used as a thermometer to measure global temperature. However, as a thermometer,  $\text{CO}_2$  is better due to its higher concentration in the atmosphere.

**Keywords:** Climate Change, Henry's Law,  $\delta^{13}\text{C}$

## 1. Introduction

At equilibrium, the concentration of any gas depends on the temperature. Thus, concentrations can be used as a global thermometer as shown in our previous paper [1]. The main problem is a big release of the gas directly into the atmosphere, because we must estimate the portion that stays in the atmosphere. Usually, the majority of this released gas,  $p_r$ , dissolves in the oceans. Note that we will use the word 'release' instead of 'emission' in cases where gases go directly into the atmosphere not emitted by water. Thus, we will use the term human release not human emission. In addition, all the chemical reactions in the atmosphere and in the oceans can make the application of Henry's Law more difficult. Figure 1 from our previous paper presents partly experimental  $\text{CO}_2$  curves  $p_e$  and  $p_h$ , emitted by the oceans and the portion of the human release remaining in the atmosphere, respectively. The curve  $p_e$  also gives the observed temperature change. This is the first experimental evidence of our theory concerning Henry's Law. The source of  $\text{CO}_2$  in the atmosphere can be studied measuring the abundance of  $^{13}\text{C}$  in air samples. The

measurements are available in the unit  $\delta^{13}\text{C}$ .



**Figure 1.** Human  $\text{CO}_2$  release  $p_r$  (black), measured  $p_m - 280$  ppm (red),  $\text{CO}_2$  emitted by water  $p_e$  (blue) and human  $\text{CO}_2$  contribution to the atmosphere  $p_h$  (green). [1]

The curves  $p_e$  and  $p_h$  explain the observed  $\delta^{13}\text{C}$

measurements very well. This will be the second experimental proof of our theory. In principle, all gases must obey Henry's Law. Thus, as new examples, we look at  $N_2O$  and  $SF_6$  in the atmosphere. We will find that increases in those gases are merely due to the temperature increasing and human influences are small.

## 2. Explanation of Per-mil ( $\delta^{13}C$ ) Measurements in the Atmosphere

Per-mil (per-mille, promille...) values define as follows:

$$\delta^{13}C = \left( \frac{(^{13}C/^{12}C)_{\text{sample}}}{(^{13}C/^{12}C)_{\text{standard}}} - 1 \right) \times 1000, \quad (1)$$

where the standard  $^{13}C/^{12}C$  is 0.0112372. The per-mil value gives the abundance ratio of carbon dioxide isotopes  $^{13}CO_2$  and  $^{12}CO_2$  compared with the standard ratio. Using the curves  $p_m$ ,  $p_e$  and  $p_h$  shown in Figure 1, we can calculate the per-mil curve as follows:

$$\delta^{13}C = \frac{-6.5\text{‰} \cdot 280 \text{ ppm} - 26\text{‰} p_h - \delta^{13}C_{\text{ocean}} p_e}{p_m} \quad (2)$$

The first term in the above equation is a constant per-mil  $-6.5\text{‰}$  and concentration 280 ppm before 1750, as shown in Figure 2. The next term is the contribution of burning fossil fuel, which has an average per-mil value of approximately  $-26\text{‰}$ . The last term is an effect of carbon dioxide emitted by the oceans. The per-mil value  $\delta^{13}C_{\text{ocean}}$  of the emitted  $CO_2$  is little smaller than  $-6.5\text{‰}$ , because in the oceans, the dissolution of the major part of human release  $p_r$  decreases the abundance of  $^{13}C$ , see for example [2].

Note that  $p_r$  has the per-mil value  $-26\text{‰}$ . The per-mil value of the oceans  $\delta^{13}C_{\text{ocean}}$  has been derived by fitting Equation (2) with the observed per-mil values  $\delta^{13}C$ , assuming a linear decrease in  $\delta^{13}C_{\text{ocean}}$ , see Figure 3. According to the fit, per-mil values of  $\delta^{13}C_{\text{ocean}}$  changed from  $7.1\text{‰}$  to  $7.8\text{‰}$  in the time interval between 1980 and 2020. We also fit Equation (2) with exponential decay in  $\delta^{13}C_{\text{ocean}}$ . However, the difference between the linear and exponential fits was negligible.

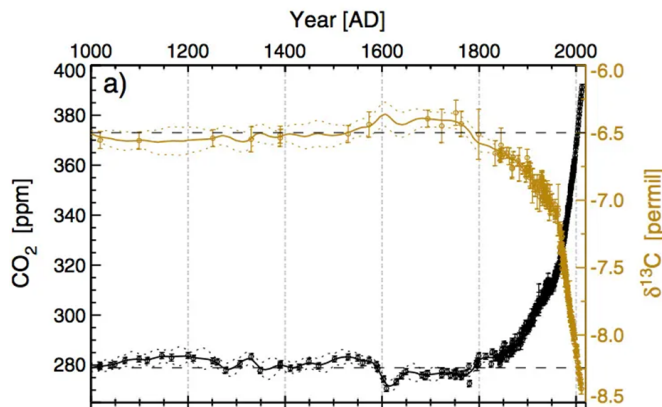


Figure 2.  $CO_2$  concentration (black circles) and the  $\delta^{13}C$  (brown circles) [15]

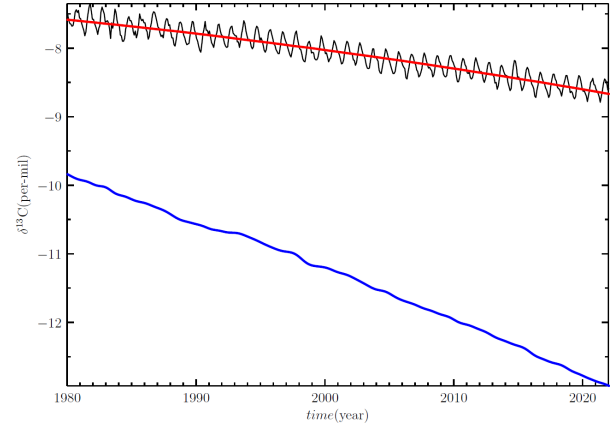


Figure 3. Observed per-mil values [16] (black), Equation (2) fitted to the observed values (red) and per-mil values calculated assuming total increase of  $CO_2$  in the atmosphere ( $p_e + p_h$ ) is anthropogenic (blue).

The total  $CO_2$  concentration,  $p_m$ , measured in Mauna Loa is given by:

$$p_m = 280 \text{ ppm} + p_h + p_e \quad (3)$$

Equations (2) and (3) include both  $p_e$  and  $p_h$ , so we can solve these values using the observed  $\delta^{13}C$  and  $p_m$  values also in equations (2) and (3). If we define, according to the fit,  $\delta^{13}C_{\text{ocean}} = -7.06\text{‰} - 0.0177\text{‰/yr} (t - t_0)$ , where  $t_0 = 1980 \text{ yr}$ ,  $\delta^{13}C_{\text{fossil}} = -26\text{‰}$ ,  $\delta^{13}C_0 = -6.5\text{‰}$  and  $p_0 = 280 \text{ ppm}$ , we get:

$$p_e = \frac{p_m(\delta^{13}C_{\text{fossil}} - \delta^{13}C) - p_0(\delta^{13}C_{\text{fossil}} - \delta^{13}C_0)}{\delta^{13}C_{\text{fossil}} - \delta^{13}C_{\text{ocean}}} \quad (4)$$

and

$$p_h = p_m - p_0 - p_e. \quad (5)$$

Next we compare these curves with those in Figure 1. First the measured  $\delta^{13}C$  is smoothed out using Fourier filtering, see Figure 4. It is now possible to compare the curves  $p_e$  and  $p_h$  derived using Henry's Law [1] with those derived using the measured  $\delta^{13}C$ , see Figure 5. In 2020, the values of  $p_e$  and  $p_h$  were 95 ppm and 37 ppm, respectively. Note that all the information of Henry's Law is in the per-mil values.

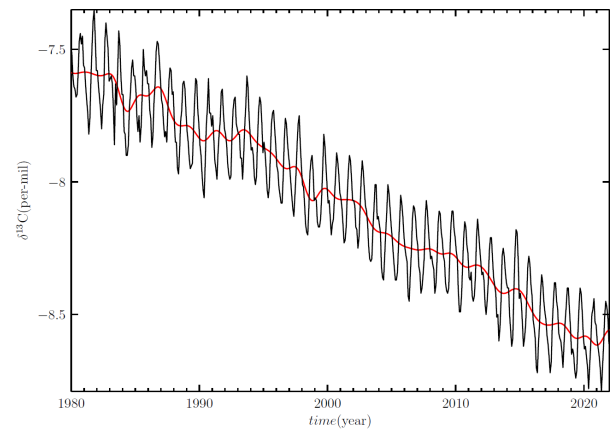
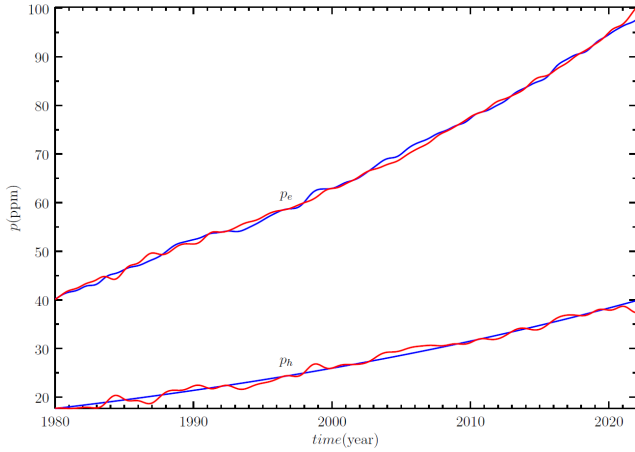


Figure 4. The observed per-mil curve  $\delta^{13}C$  between 1980 and 2022 in Mauna Loa (black) [16], and a Fourier smoothed [3,4] red curve, which does not show seasonal variations.



**Figure 5.** CO<sub>2</sub> emitted by ocean  $p_e$  (upper curves) and human CO<sub>2</sub> contribution to the atmosphere  $p_h$  (lower curves). The blue curves are derived using Henry's law and the red curves, Eqs. (4) and (6), using per-mil data with Eqs. (2) and (3).

In Intergovernmental Panel on Climate Change (IPCC) Fifth Assessment Report (AR5) [5] (p. 467), referring to paper [6] by Joos et al., it is stated that in 2011 “about half of the emissions remained in the atmosphere  $240 \text{ PgC} \pm 10 \text{ PgC}$  since 1750”. In this statement, emission means human release, or our  $p_r$ . This statement means  $112.5 \text{ ppm CO}_2$  and is close to the sum of  $p_e + p_h$  in 2011, see also Ollila [7]. This implies that the total increase of CO<sub>2</sub> should be anthropogenic. This is simple to test by replacing  $\delta^{13}\text{C}_{\text{ocean}}$  with  $-26 \text{ ‰}$  in Equation (2). Figure 3 shows the result compared with the measured per-mil curve. As can be seen, the per-mil value was  $-12.1 \text{ ‰}$  in 2011, compared with the measured value of  $-8.32 \text{ ‰}$ . It is clear that the IPCC case is not realistic and no explanation has been provided for these conflicting values. Figure 5 thus proves, experimentally, that human contribution is surprisingly small.

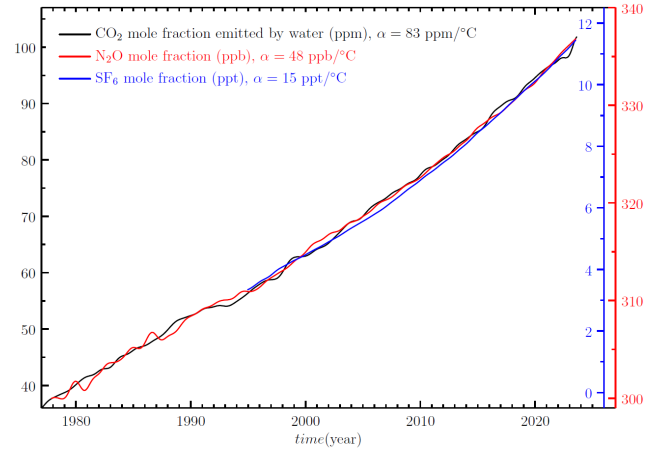
### 3. Discussion

In our previous paper [1] we introduced a global thermometer using CO<sub>2</sub> emitted from the oceans,  $p_e$ , because  $p_e$  depends on the temperature. However, in this case, we have to calculate  $p_h$  in order to get  $p_e$  from Equation (3). The human contribution in the atmosphere is approximately given by:

$$p_h = \tau \frac{dp_r}{dt}, \quad (6)$$

where  $p_r$  is the total human release into the atmosphere and  $\tau$  is the response time of Henry's Law. The derivative of  $p_r$  is now large enough that  $p_h$  is significant, see Figure 1. In principle all gases in the atmosphere can work a global thermometer because gases must obey Henry's Law and its temperature dependence. In Figure 6, the measured concentrations of CO<sub>2</sub>, N<sub>2</sub>O and SF<sub>6</sub> in the atmosphere are shown. The scales of concentrations have been adjusted so that the curves overlap and as a result, they are remarkably similar. This behaviour is real because the same temperature change  $\Delta T$  controls the abundance. If we apply Equation (7)

from paper [1]:



**Figure 6.** Measured concentrations of CO<sub>2</sub> (black), N<sub>2</sub>O (red) and SF<sub>6</sub> (blue) [17]. The scale of each curve adjusted so the curves overlap each other's so good as possible.

$$\Delta T = \frac{1}{\alpha} \left( p_e + \tau \frac{dp_e}{dt} \right), \quad (7)$$

where  $\alpha$  is the emissions strength of a given gas in Henry's Law. We can then get the total change of the temperature between 1980 and 2020 calculated from CO<sub>2</sub> and N<sub>2</sub>O. The results are  $0.78^\circ\text{C}$  and  $0.79^\circ\text{C}$ , respectively. Note that in the case of N<sub>2</sub>O other sources like human release are small. In other words, the derivatives of other releases, using Equation (6), are so small that they cannot be seen in the figures.

It is useful to note that solving  $p_e$  and  $p_h$  from Equations (2) and (3) is a simple task. We do not need to know the human release  $p_r$ , the emission strength  $\alpha$ , response time  $\tau$ , or temperature. The information about these parameters is in per-mil values.

### 4. Conclusion

Our main results [1, 8-14] concerning climate change can be summarized by the following: the total temperature increase from 1750 to 2020 was about  $1.3^\circ\text{C}$ , which consists of roughly  $1.2^\circ\text{C}$  due to the decrease of clouds or relative humidity [8, 9, 12, 13] and  $0.1^\circ\text{C}$  of greenhouse gases due to  $p_e$  and  $p_h$ . The human contribution was about  $0.03^\circ\text{C}$  due to  $p_h$ . Our climate sensitivity [8-12] for CO<sub>2</sub> is  $0.24^\circ\text{C}$ , which is about one order of magnitude smaller than the value given by IPCC. Our climate sensitivity has been theoretically derived, and it has proved robust in at least five different experimental methods [8-12]. On the other hand, IPCC value is based only on theoretical circulation models without any experimental verification [13].

We have shown that the increase of CO<sub>2</sub> levels is not the cause of global warming but a consequence. Future research should look for other causes of global warming than increasing CO<sub>2</sub> levels. The most promising line of research is to focus on the causes of the variation of cloudiness or relative humidity.

## Abbreviations

IPCC      Intergovernmental Panel on Climate Change  
AR5      Fifth Assessment Report

## Conflicts of Interest

No conflict of interest for this article.

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