



**Abstract**

The successful application of simple statistical methods in economic modelling suggest that similar methods could provide insight into global climate. We apply an autoregressive method to observed time series to determine the dependence of atmospheric CO<sub>2</sub> concentration on carbon emissions and, in turn, the dependence of globally averaged temperature on atmospheric CO<sub>2</sub> concentration. We ascribe physical meaning to the regression parameters in terms of first order differential equations describing the diffusion of CO<sub>2</sub> and heat between reservoirs, viz.: the diffusion of CO<sub>2</sub> between the atmosphere and the deep ocean and the transport of heat from the mixed layer. A strong feature of regression models is their built-in mechanism for deciding when a model provides an adequate description of the given data. Two implications of this statistically robust approach are that CO<sub>2</sub> diffuses from the atmosphere within a time scale of decades and that global average temperatures are unlikely to exceed 2°C above pre-industrial values.

**Plain Language Summary**

We use a conventional, statistical method to estimate the dependence of carbon concentration on emissions, and, in turn, to estimate the dependence of temperature on carbon concentration. Then, based on general assumptions about future emissions, we make realistic forecasts of global temperature and show that human caused CO<sub>2</sub> and its associated forcing are creating a temperature pulse and not a permanent plateau or self-amplifying feedback loop. There is no observational evidence that global climate is heading towards a catastrophe nor that carbon dioxide remains in the atmosphere indefinitely.

## 1 Introduction

In the early days of economic modelling it was found that complicated models performed poorly when used for out-of-sample forecasting. Regression models with few independent variables often produced better forecasts than did the big models (Nelson, 1972) (Ashley, 1988). Global Circulation Models are no less complicated than economic models but, until now, this aspect of them has been ignored. Furthermore, there are issues with the fundamental physics of GCMs because they are deterministic and do not allow for the Gibbs entropy associated with turbulence (Reid, 2019).

The fundamental question such models are designed to answer is the extent to which increases in atmospheric greenhouse gas concentrations affect the Earth's climate. We develop a simple physical model of the ocean-atmosphere system based on the diffusion of heat and carbon dioxide between reservoirs. This model has the advantage that regression methods can be used to estimate parameters from well-established time series data of emission, concentration and temperature. Once these parameters are known, they can be used to predict future CO<sub>2</sub> concentrations and global average temperature based on reasonable assumptions about future carbon emissions.

## 2 Diffusion and the ARX Model

By convention, a fluid is assumed to be a Newtonian continuum describable by differential equations such as the Navier-Stokes equations and diffusion equations. In practice, observations of fluids usually comprise time series, i.e. sequences of discrete numbers sampled or averaged over equal intervals of time. Time series can be related to continuous differential equations by means of finite difference approximations. Time series readily lend themselves to a statistical interpretation and allow high frequency noise and seasonal effects to be eliminated by choosing a suitable sampling interval.

The diffusion of a chemical constituent or heat between two reservoirs is described by Fick's Law. In one dimension

$$J = -D \frac{dc}{dz} \quad (1)$$

where  $J$  is the diffusion flux,  $D$  is the diffusion coefficient,  $c$  is the concentration and  $z$  is the spatial coordinate. Together with the continuity equation

$$\frac{\partial c}{\partial t} = F(t) - \oint J dA \quad (2)$$

this leads to the one dimensional diffusion equation

$$\frac{\partial c}{\partial t} + \oint \frac{D(c - c_0)}{\delta z} dA = F(t) \quad (3)$$

where  $F(t)$  is the rate of pumping into the source reservoir,  $c_0$  is the concentration in the sink reservoirs,  $\delta z$  is the thickness of the boundary layer between the two reservoirs and the integral is taken over the entire boundary layer area. (3) can be written

$$\frac{\partial c}{\partial t} + \frac{c}{\tau} = F(t) + \frac{c_0}{\tau} \quad (4)$$

where  $c$  is spatially homogeneous and  $\tau$  is a constant which has the units of time. The second term on the right is generally negligibly small and can be subsumed into the forcing function,  $F$ , or ignored. In finite difference terms, with  $c_0 = 0$ , (4) becomes

$$\frac{c_i - c_{i-1}}{\Delta t} + \frac{c_i}{\tau} = F_i \quad (5)$$

where  $\Delta t$  is the time step or sampling interval,  $c_i$  and  $F_i$  designate values at time  $i\Delta t$ .

The above equations are deterministic and apply equally well to a numerical fluid dynamical model as they do to the regression model we are developing here. However, in order estimate model parameters from the data we need to develop a proper regression model and introduce random variables. We do this by making the forcing function,  $F(t)$ , a random variable as the sum of a deterministic component,  $f(t)$ , and a random component,  $\epsilon_i$ , viz.:

$$F_i = f_i + \xi_i \quad (6)$$

where the expectation mean of  $\epsilon_i$  is zero, i.e.  $\mathbf{E}\xi_i = 0$  and  $\mathbf{E}F_i = f_i$ .

We now change notation so as to make a clear distinction between random variables (upper case) and constants (lower case). Equation (5) becomes

$$Y_i = \alpha_0 x_i + \alpha_1 y_{i-1} + \Xi_i, \quad i = 1, \dots, N \quad (7)$$

where  $Y_i = c_i$ ,  $x_i = f_i$ ,  $y_{i-1} = c_{i-1}$ ,

$$\alpha_0 = \frac{\Delta t}{1 + \Delta t/\tau} \quad (8)$$

and

$$\alpha_1 = \frac{1}{1 + \Delta t/\tau} \quad (9)$$

This can be further generalized to

$$Y_i = \alpha_0 x_i + \sum_{j=1}^p \alpha_j y_{i-j} + \sum_{k=1}^q \beta_j \Xi_{i-k}, \quad i = 1, \dots, N \quad (10)$$

where the atmospheric carbon concentration,  $c_i$  in (5), is represented by both  $Y$  and  $y$ ,  $x_i = \Gamma_i$  is the exogenous variable and the  $\Xi_i$  are unselfcorrelated random variables with zero mean. The regression coefficients  $\alpha_0$ ,  $\alpha_j$  and  $\beta_j$  are to be estimated from the data and  $p$  and  $q$  are small positive integers.

This is the ARMAX(p,q) model (for ‘autoregressive moving average with exogenous variable’). There are software packages for parameter estimation available under the aegis of the major programming languages. Unfortunately some of these, such as the Python *Statsmodels* package, are flawed, because they estimate the exogenous parameter,  $\alpha_0$ , prior to estimating the other parameters, leading to omitted-variable bias (Greene, 2003).

Note the distinction between the random variable  $Y_i$  and the sample values  $y_{i-j}$  which are constants. Equation (10) is a state space representation (Hamilton, 1994) describing states of the system at a succession of discrete instants; the random variable,  $Y_i$ , at one instant becomes the constant,  $y_i$ , in the following instant. The direction of time is important in regression, which, unlike correlation, allows causality to be inferred.

Estimation of the MA coefficients,  $\{\beta_i\}$ , requires an iterative Kalman filter method which does not always converge. The second, moving average summation in (10) is a blurring function, so that  $q > 1$  when the sampling interval,  $\Delta t$ , is too small. The given time series can be downsampled or ‘decimated’ by  $q$  to give a new time series with sampling interval  $q\Delta t$  with little loss of information and (10) becomes

$$Y_m = \alpha'_0 x_m + \sum_{n=1}^p \alpha'_n y_{m-n} + \Xi'_m, \quad m = 1, \dots, M \quad (11)$$

102 where  $m = qi$ ,  $qM \leq N$  and

$$\Xi'_m = \sum_{k=1}^q \beta_j \Xi_{m-k} \quad (12)$$

103 so that  $E(\Xi'_m \Xi'_n) = 0$  for  $m \neq n$  because the summations do not overlap. The  $\{\Xi'_m\}$   
 104 in (11) are therefore unselfcorrelated with zero mean. This is a simple case of a Renor-  
 105 malization Group Transformation (Wang et al., 2013). The aim here is to find the opti-  
 106 mum scale for the given problem. In our case the optimum scale is considered to be  
 107 the finest scale which satisfies the statistical test for the validity of the model.

108 The model summarized by (11) can be termed an ARX(p) model for ‘autoregres-  
 109 sive with exogenous variable’. Its parameters and their confidence limits can be estimated  
 110 using ordinary least squares. The sequence of residuals,  $\{\xi'_m\}$ , is given by

$$\xi'_m = y_m - \left( \hat{\alpha}'_0 x_m + \sum_{n=1}^p \hat{\alpha}'_n y_{m-n} \right), \quad m = 1, \dots, M \quad (13)$$

111 where  $y_m$  is the sample value or ‘realization’ of  $Y_m$ ,  $\hat{\alpha}'_0$  and  $\hat{\alpha}'_n$  are the coefficient esti-  
 112 mates and the  $\{\xi'_m\}$  are to be tested for self-correlation.

### 113 3 The Bomb Test Curve

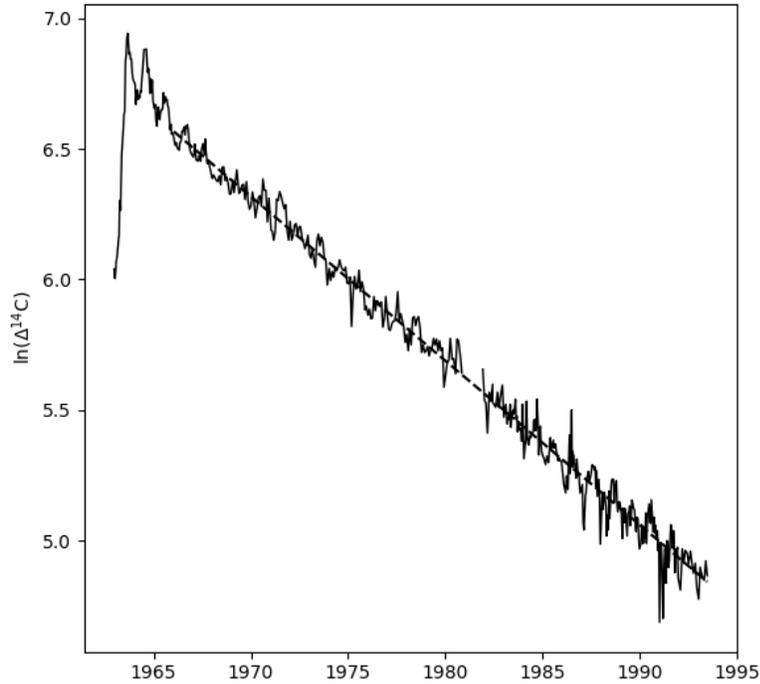
114 Given the complexity of ocean-atmosphere interactions, the diffusion relationship  
 115 (4) appears remarkably simple. However there is strong observational evidence that such  
 116 a simple equation does indeed describe diffusion of atmospheric carbon into the ocean.

117 The testing of nuclear weapons during the 1950s and 1960s injected significant amounts  
 118 of the radioactive  $^{14}\text{C}$  isotope of carbon into the atmosphere. More importantly, the abrupt  
 119 cessation of atmospheric testing following the Nuclear Test Ban Treaty of 5 August 1963,  
 120 meant that the rate of production of the  $^{14}\text{C}$  isotope reverted to the constant natural  
 121 background level. This allows the movement of carbon dioxide between natural reser-  
 122 vairs to be assessed in much the same way that radioactive isotopes are used to assess  
 123 metabolic processes in nuclear medicine.

124 The decrease in  $\Delta^{14}\text{C}$  is known as “The Bomb Test Curve”. Numerous observa-  
 125 tions were made in the decades following the cessation of testing following the Nuclear  
 126 Test Ban Treaty. Here we look at a single high quality data set from Fruholmen, Nor-  
 127 way (Nydal & Lövseth, 1983) shown in Figure 1. The natural logarithm,  $\ln(\Delta^{14}\text{C})$ , is  
 128 plotted on the vertical axis rather than  $\Delta^{14}\text{C}$  itself so that exponential behaviour be-  
 129 comes linear. A regression line was fitted between January 1966 and the end of the data  
 130 set in June 1993.

Statistic	Value
$r$	-0.9939
slope	-.06289 year <sup>-1</sup>
$\tau$	15.9 years
$t_{1/2}$	11.02 years

**Table 1.** Regression Statistics related to Figure 1



**Figure 1.** The natural logarithm of  $\Delta^{14}\text{C}$  values recorded at Fruholmen, Norway as a function of time Dashed line: regression line fitted between January 1966 and June 1993.

Regression statistics are shown in Table 1. The fit is remarkably good and accounts for 98.8 percent of the variance. Hence, with a high degree of accuracy:

$$\Delta^{14}\text{C} = Ae^{-t/\tau} \quad (14)$$

131 where  $A$  is the value of  $\Delta^{14}\text{C}$  at  $t = 0$ . Thus half of the bomb test  $^{14}\text{CO}_2$  disappears  
 132 from the atmosphere every 11 years<sup>1</sup>.

133 Equation (14) is the solution of (4) with  $c = \Delta^{14}\text{CO}_2$  and  $F(t)$  specifies the rate  
 134 at which concentration increases due to new material being introduced into the reser-  
 135 voir. There was a spike in  $F(t)$  at the time of nuclear testing, after which it reverted to  
 136 a constant background value due to the bombardment of upper atmosphere Nitrogen by  
 137 cosmic rays.

138 Carbon dioxide reacts with water to form carbonate and bicarbonate ions. Hence  
 139 the diffusion rate of carbon *per se* involves reaction rates and diffusion rates for each of  
 140 these three species. These are almost completely independent of atomic mass (Zeebe, 2011)  
 141 and so all the isotopes of carbon,  $^{12}\text{C}$ ,  $^{13}\text{C}$ ,  $^{14}\text{C}$ , in the form of  $\text{CO}_2$  and its radicles, dif-

<sup>1</sup> Note that this half-life of atmospheric  $^{14}\text{CO}_2$  is due to diffusion of this gas from the atmosphere and is unrelated to the radioactive half-life of  $^{14}\text{C}$  of 5730 years

142 fuse through water at the same rate and the time constant,  $\tau$ , applies equally to all iso-  
143 topic species of  $\text{CO}_2$ .

144 It is therefore reasonable to assume that  $\text{CO}_2$  diffuses from the atmosphere into  
145 some other reservoir or sink. The excellent fit of a single regression line indicates that  
146 any diffusion process must be dominated by a single sink with a single time constant and  
147 that this sink is not approaching saturation. Furthermore the fact that the atmospheric  
148  $\Delta^{14}\text{CO}_2$  has, by now, returned to its pre-bomb background level implies that the sink  
149 is much larger than the source, the atmosphere. The only candidate sink which fulfils  
150 these conditions is the deep ocean.

## 151 4 Heat Transport

152 Environmental temperatures are largely dependent on the transport of heat and  
153 other forms of energy by turbulent processes (Hasselmann, 1976). Heat diffuses in the same  
154 way as chemical constituents described by (1) and (2) leading to Fourier's heat equation:

$$\rho_0 c_p \frac{\partial T}{\partial t} = - \oint_A K_0 \frac{\partial T}{\partial z} dA + Q(x, t) \quad (15)$$

155 where  $\rho_0$  and  $c_p$  are the density and specific heat of a reservoir,  $T$  is its temperature,  $t$   
156 is the time,  $z$  is a spatial variable with the units of length,  $K_0$  is the conductivity or sim-  
157 ilar constant,  $Q$  is the flux of heat or other form of energy and  $A$  is the surface area of  
158 the reservoir.

159 In the present case the reservoir is the mixed layer of the ocean. Because the spe-  
160 cific heat of water is much larger than that of air, the top 2.5 m of the mixed layer holds  
161 as much heat as the entire atmosphere above it. The globally averaged mixed layer depth (de  
162 Boyer Montgut et al., 2004) is about 100m, so that the heat capacity of the mixed layer  
163 is 40 times that of the atmosphere. Seventy percent of the Earth's surface is ocean. Ef-  
164 fectively, mixed layer temperatures comprise the bulk of globally averaged surface air tem-  
165 perature measurements and the two measurements are highly correlated.

166 Integrating (15) over the entire mixed layer and including a radiative forcing term,  
167  $G(t)$ , for the heat trapping effect of greenhouse gases gives Newton's Law of Cooling for  
168 the mixed layer:

$$\rho c_p \frac{\partial T}{\partial t} = -KT + G(t) + q(t) \quad (16)$$

169 where  $\rho$ ,  $c_p$  are the density and specific heat of sea water and  $K$  is a global average of  
170 all forms of cooling which are approximately proportional to temperature differences, i.e.  
171 conduction via the thermocline into the deep ocean and convection to the top of the at-  
172 mosphere to be radiated into space.  $q(t)$  is the random component of heat gained and  
173 lost by the mixed layer due to clouds and similar unpredictable phenomena.

174 In finite difference form (16) becomes

$$T_i = a.T_{i-1} + b.\Gamma_i + \xi_i, \quad i = 1, \dots, N \quad (17)$$

175 where  $t = i\Delta t$ ,  $\xi_i = \Delta t q_i / (\rho c_p + K\Delta t)$  and  $a$  and  $b$  are constants to be estimated from  
176 the  $N$  data values.  $\Gamma_i$  is given by

$$\Gamma = \ln([\text{CO}_2]) \quad (18)$$

177 where  $[\text{CO}_2]$  is the atmospheric carbon dioxide concentration in parts per million (Huang  
178 & Shahabadi, 2014). This too can be further generalized to the form (10).

Model	Exogenous Variables	$Q$	$P$
ARX(0)	$E_i$ only	1424.6	0.0000
ARX(1)	$E_i, C_{i-1}$	95.4	0.0000
ARX(2)	$E_i, C_{i-1}, C_{i-2}$	120.0	0.0000
ARX(3)	$E_i, C_{i-1}, \dots, C_{i-3}$	91.3	0.0000
ARX(4)	$E_i, C_{i-1}, \dots, C_{i-4}$	71.0	0.0000
ARX(5)	$E_i, C_{i-1}, \dots, C_{i-5}$	71.4	0.0000

**Table 2.** Ljung-Box parameter,  $Q$ , and its probability,  $P$ , for ARX models of undecimated CO<sub>2</sub> concentration,  $C_i$ , and CO<sub>2</sub> emissions data,  $E_i$ .

## 5 Finding Models

We investigated the relationships between the three time series: anthropogenic emissions, atmospheric CO<sub>2</sub> concentration and global average temperature. We did so by estimating the coefficients  $\alpha'_0$  and  $\alpha'_n$  in the ARX model, (11), using Ordinary Least Squares. All coefficients were estimated simultaneously to avoid omitted-variable bias and the complexities of de-convoluting a moving average component were avoided by decimating the data when necessary.

We used the model twice; to estimate firstly, (i) the regression of atmospheric carbon concentration,  $C$ , on anthropogenic carbon emissions,  $E$ , and then, (ii) the regression of global average temperature anomaly,  $T$ , on the logarithm of atmospheric carbon concentration,  $\Gamma$ , estimated from step (i). In step (i),  $E$  is the exogenous variable. In step (ii)  $\Gamma$  is the exogenous variable.

In each case we used the model with  $q = 0, 1, \dots, 5$  ( $q = 0$  implying that the second term on the RHS of (11) is omitted) in order to determine the smallest value of  $q$  for which the residuals given by (13) could be deemed unselfcorrelated. The successful models were then used to forecast global average temperature from Hubbert curves of emission estimates (Hubbert, 1962).

The *sine qua non* of all regression models is that the innovation,  $\Xi$ , be unselfcorrelated. It is an assumption which can be validated by testing whether the sample residuals,  $\{\xi'_m\}$ , given by (13), are self-correlated. If they are, then the random variables,  $\{\Xi'_m\}$ , in (11) cannot be assumed to be unselfcorrelated and (11) is not a valid regression model.

The Ljung-Box test parameter,  $Q$ , is computed from the sample autocorrelation of  $\{\xi'_m\}$  at lag  $k$  out to some maximum lag,  $k_{max}$  (Ljung & Box, 1978). Under the null hypothesis that the residuals are unselfcorrelated,  $Q$  has a  $\chi^2$  distribution with  $k - n$  degrees of freedom where  $n$  is the number of regression coefficients fitted. From this a probability,  $P$ , can be found and suitable values of  $q$ , the decimation factor, and  $p$ , the number of autoregressive coefficients. By Occam's Razor we chose the smallest values which satisfy this test.

Step (i): applying Ljung-Box to the residuals given by (13) for ARX( $p$ ),  $p = 0, \dots, 5$  gives the results shown in Table 2. Probabilities are zero for all values of  $p$  indicating that the null hypothesis that the residuals are unselfcorrelated can be rejected. Both time series were then decimated by 2. The results are shown in Table 3. The probability,  $P$ , for the ARX(1) model has a value of 0.4259 indicating that the null hypothesis that the residuals are unselfcorrelated cannot be rejected. The ARX(1) case, is a good fit to the decimated data and constitutes a valid regression model for  $C$  on  $E$ .

Step (ii): an ARX model of the regression of global average temperature,  $T$ , on the logarithm of atmospheric CO<sub>2</sub> concentration is also required. These time series were also

Model	Exogenous Variables	$Q$	$P$
ARX(0)	$E_i$ only	513.5	0.0000
ARX(1)	$E_i, C_{i-1}$	28.5	0.4359
ARX(2)	$E_i, C_{i-1}, C_{i-2}$	28.6	0.3830
ARX(3)	$E_i, C_{i-1}, \dots, C_{i-3}$	24.5	0.5483
ARX(4)	$E_i, C_{i-1}, \dots, C_{i-4}$	24.3	0.5049
ARX(5)	$E_i, C_{i-1}, \dots, C_{i-5}$	22.0	0.5796

**Table 3.** Ljung-Box parameter,  $Q$ , and its probability,  $P$ , for ARX models of CO<sub>2</sub> concentration,  $C_i$  and CO<sub>2</sub> emissions,  $E_i$ , when both time series have been decimated by 2

Model	Exogenous Variables	$Q$	$P$
ARX(0)	$T_i$ vs $\ln(C_i)$ only	158.4	0.000
ARX(1)	$T_i$ vs $\ln(C_i), T_{i-1}$	60.9	0.0003
ARX(2)	$T_i$ vs $\ln(C_i), T_{i-1}, T_{i-2}$	32.9	0.1995
ARX(3)	$T_i$ vs $\ln(C_i), T_{i-1}, \dots, T_{i-3}$	31.2	0.2214
ARX(4)	$T_i$ vs $\ln(C_i), T_{i-1}, \dots, T_{i-4}$	32.5	0.1436
ARX(5)	$T_i$ vs $\ln(C_i), T_{i-1}, \dots, T_{i-5}$	34.2	0.0807

**Table 4.** Ljung-Box parameter,  $Q$ , and its probability,  $P$ , for ARX models of global average temperature and CO<sub>2</sub> concentration data decimated by 2

Model	$\hat{\alpha}'_0$	$\hat{\alpha}'_1$	$\hat{\alpha}'_2$
$C_i$ vs $E_i, C_{i-1}$	0.210	0.969	
$T_i$ vs $\ln(C_i), T_{i-1}, T_{i-2}$	1.549	0.213	0.299

**Table 5.** Regression coefficients estimated from the selected models

216 decimated by 2 to keep the same sampling interval throughout. The results of the Ljung-  
 217 Box test for these ARX models are shown in Table 4. The ARX(2) model was chosen  
 218 as the simplest model with unselfcorrelated residuals. Estimates of the regression coef-  
 219 ficients of (11),  $\hat{\alpha}'_0$ ,  $\hat{\alpha}'_1$  and  $\hat{\alpha}'_2$  for the two selected models are shown in Table 5

220 The ARX(1) model of Step (i) could equally well have been derived by simply es-  
 221 timating the regression of the first difference of observed CO<sub>2</sub> concentration on CO<sub>2</sub> emis-  
 222 sions. However the present formalism led to recognition of the need to decimate these  
 223 time series by two in order to satisfy the Ljung-Box test.

## 224 6 The Impulse Response Function of CO<sub>2</sub> Concentration

225 The impulse response of a system is its output in response to a brief input signal.  
 226 It completely characterises a linear time-invariant system; once the impulse response is  
 227 known, the output corresponding to any input can be found by convolution. For con-  
 228 tinuous time systems the impulse response is related to the characteristic equation of the  
 229 differential equations describing the system. In discrete time systems the impulse response  
 230 can be found experimentally by applying appropriate regression methods to the mea-  
 231 sured input and output time series.

232 In assessing the effect of various greenhouse gases on global climate, the impulse  
 233 responses of climate parameters such as global average temperature to various greenhouse  
 234 gas inputs are of major importance. This has been done for a number of gases in order  
 235 to determine their ‘‘Global Warming Potential’’, for example by Joos et al. (2013). Var-  
 236 ious computed impulse response functions of the atmospheric concentration of CO<sub>2</sub> in  
 237 response to global emissions are shown in their Figure 1a. In every case more than 20  
 238 percent of the emitted pulse supposedly remains in the atmosphere after 1000 years. Sim-

239 ilar results have been used by modellers as far back as the First IPCC Assessment Re-  
 240 port as shown in their Figure 8.

241 Although presented as such, the long-duration impulse responses for atmospheric  
 242 CO<sub>2</sub> described by Joos et al are not derived from observations of the real world. They  
 243 are not based on observation; they are all derived from global circulation models. The  
 244 use of a fluid dynamical model as a surrogate for the real world only works if the model  
 245 is homologous with the real world, i.e. there must be an *exact* correspondence in *every*  
 246 aspect of the model; the correspondence between a handful of model averages and their  
 247 real world values is not sufficient. In the present case the long-duration of the model-  
 248 derived, CO<sub>2</sub> impulse response is most likely due to there being too little interaction be-  
 249 tween the atmosphere and the deep ocean in the GCMs. The sequestration of CO<sub>2</sub> in  
 250 this very large, deep-ocean reservoir would be greatly inhibited in such a model.

251 The duration of the CO<sub>2</sub> impulse response is crucial in forecasting future climate  
 252 and in planning emission reductions. It is important that it be estimated from real world  
 253 observations.

254 It is clear from Figure 1 and Table 1 that the impulse response of atmospheric  $\Delta^{14}\text{C}$   
 255 is exponential with a half time in the atmosphere of 11 years. The presence of bomb test  
 256  $\Delta^{14}\text{C}$  can no longer be detected because it has fallen below the level of the  $\Delta^{14}\text{C}$  cre-  
 257 ated naturally by cosmic rays. There is no remaining twenty percent left over to last for  
 258 a thousand years.

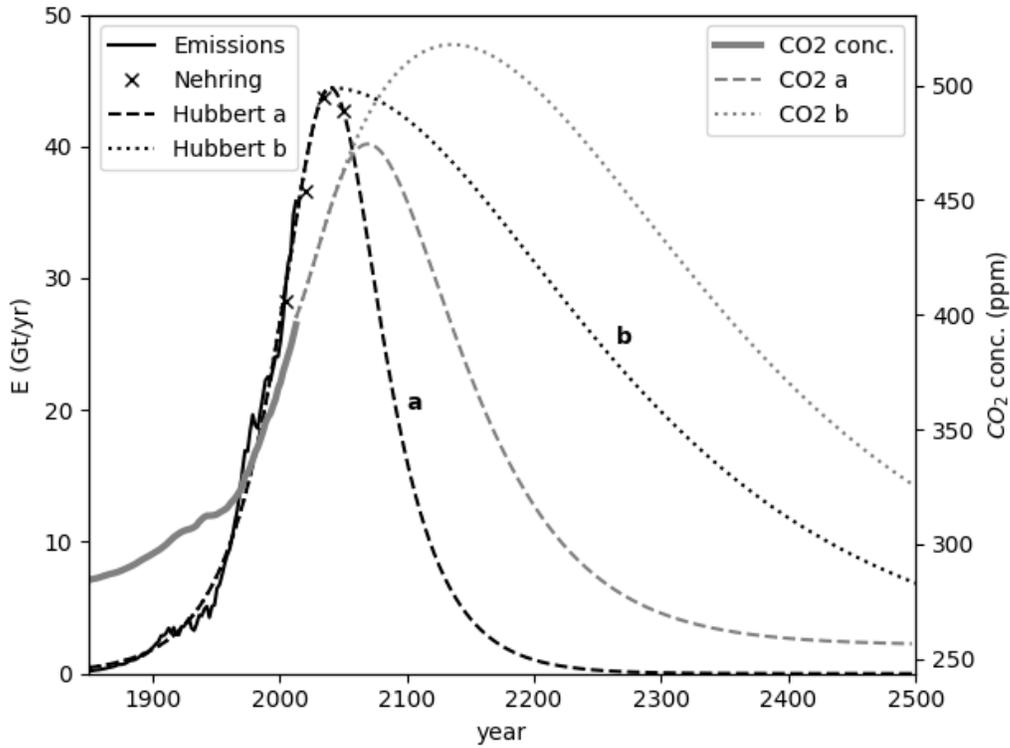
259 It might be argued that Bomb Test Curve is not a fair test of the Joos et al im-  
 260 pulse response because it only shows the carbon being absorbed. There are only very  
 261 small pre-existing  $\Delta^{14}\text{C}$  values in the deep ocean, whereas there are regions of the ocean  
 262 where CO<sub>2</sub> exceeds atmospheric values and diffusion will occur in the opposite direction.  
 263 In the first case the integrand,  $D(c-c_0)/\delta z$ , in (3) is always positive whereas in the sec-  
 264 ond case the integrand can be either positive or negative leading to a different net dif-  
 265 fusion rate and a different time constant.

266 The impulse response function of atmospheric CO<sub>2</sub> concentration is the character-  
 267 istic equation of (4) which is exponential. The time constant,  $\tau$ , is found by substitut-  
 268 ing  $\alpha_1$  from Table 5 (0.969) into (9) which gives  $\tau = 62.5$  years, once again, much less  
 269 than the millennial time scales postulated by Joos et al. The difference between the half  
 270 life time from the bomb tests of 11 years and the time constant from the regression is  
 271 explained by the obvious fact, that the bomb test curve only measures absorption of CO<sub>2</sub>  
 272 in the ocean, whereas our regression also measures the delaying effect of outgassing CO<sub>2</sub>  
 273 from the ocean to the atmosphere.

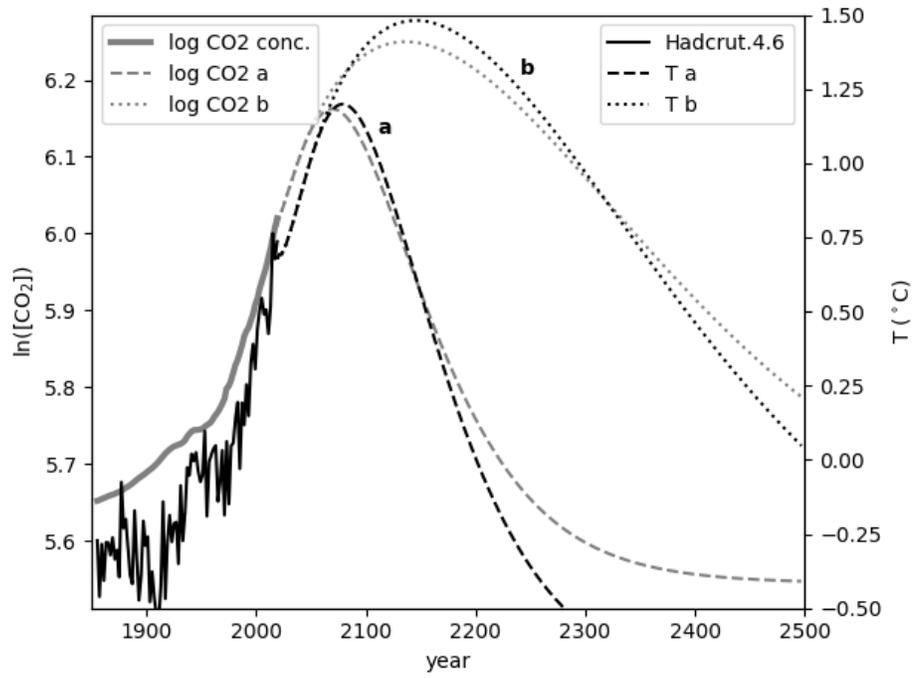
## 274 7 Forecasting Global Average Temperature

275 Numerical fluid dynamical models such as GCMs are not well suited to prediction  
 276 because they are fundamentally unstable and must often be damped by setting artifi-  
 277 cially high values of model parameters such as viscosity. Their value lies in providing in-  
 278 sight into underlying physical processes. On the other hand, regression models are well  
 279 suited to prediction when future values of exogenous variables are known or can be es-  
 280 timated. Their statistical nature allows hypothesis testing and the provision of confidence  
 281 limits whereas GCMs require computationally expensive and statistically dubious “en-  
 282 sembles” of model runs for this purpose.

Using regression models for prediction purposes is based on a single assumption  
 which may or may not be warranted depending on the circumstances: it is the assump-  
 tion of covariance stationarity. Under this assumption regression coefficients estimated  
 from the data, such as those of Table 5, can be substituted back into (11) to yield ex-  
 pectation values of Y outside the domain of the original data over the domain in which



**Figure 2.** The solid black curve shows the recorded total carbon dioxide emission rate (Gt/yr). The crosses show recorded and predicted emission rates quoted by Nehring (Nehring, 2009). Curve **a** is the symmetrical Hubbert curve which best fits Nehring's data. Curve **b** is an asymmetrical Hubbert curve with decay time constant set to five times the onset constant. The grey dashed and dotted curves show the predicted CO<sub>2</sub> concentrations forecast from the two Hubbert emission curves.



**Figure 3.** The grey curves show the logarithm of carbon concentrations resulting from the emission curves shown in Figure 2. The solid black curve is the observed (HadCRUT.4.6.0.0) global average temperature anomaly. The dashed black curves show the predicted global average temperature anomalies corresponding to the Hubbert curves **a** and **b** in Figure 2.

the exogenous variable,  $x$ , is known or can be estimated. This can be done by iteration. Thus, from (11)

$$E(Y_{M+1}) = \alpha'_0 x_M + \sum_{n=1}^p \alpha'_n y_{M+1-n} \quad (19)$$

283 since  $E(\Xi'_m) = 0$  by definition. If  $y_{M+1}$  is now defined as  $E(Y_{M+1})$  for substitution into  
 284 the right hand side of (19), the process can be repeated and the future behaviour of the  
 285 endogenous variable,  $y$ , forecast over the domain for which the exogenous variable,  $x$ ,  
 286 is known.

287 In order to forecast global average temperature we need to know the endogenous  
 288 variable,  $\Gamma$ , a function of atmospheric carbon dioxide concentration,  $C$ , given by (18).  
 289 In order to forecast  $C$  we need to know the carbon dioxide emission rate,  $E$ .

290 Although controversial, the Hubbert curve (Hubbert, 1962) provides a canonical start-  
 291 ing point. The Hubbert curve for projected carbon emissions, fitted to known emissions  
 292 and to the data of Nehring (Nehring, 2009) is shown as the dashed black curve, **a**, of Fig-  
 293 ure 2. The atmospheric CO<sub>2</sub> concentration forecast using the iterative method is shown  
 294 as the grey dashed line.

295 A major criticism of the Hubbert Curve is its symmetry. In reality, the decline in  
 296 resources is usually significantly slower than the onset of the curve. The dotted black  
 297 curve, **b**, in Figure 2, shows an asymmetrical Hubbert curve with the same onset and  
 298 maximum as curve **a**, but with a decay time which is five times slower. The correspond-  
 299 ing, atmospheric CO<sub>2</sub> forecast is shown as the grey dotted line.

300 The emissions-generated, grey CO<sub>2</sub> concentration curves in Figure 2 were used via  
 301 (18) to generate the (grey) log concentration curves **a** and **b** shown in Figure 3.

## 302 8 Data Sources

303 Time series in the form of annual averages of the relevant variables,  $E$ ,  $C$  and  $T$   
 304 were downloaded from the Web in November 2020.

305 The global average temperature anomaly data,  $T$ , were taken from the HadCRUT  
 306 .4.5.0.0 data set (Morice et al., 2012).

307 Carbon dioxide concentrations,  $C$ , were taken from the University of Melbourne  
 308 Greenhouse Gas Factsheet (Mainshausen et al., 2017) supplemented with recent values  
 309 from Mauna Loa.

310 Global fossil fuel emissions,  $E$ , were downloaded from CDIAC, the Carbon Diox-  
 311 ide Information Analysis Center (Boden et al., 2017).

## 312 9 Results

313 The temperature anomaly forecasts generated from the two Hubbert scenarios are  
 314 shown as the black dashed and dotted curves labelled **a** and **b** in Figure 3. The dashed  
 315 curve, **a**, predicted from the symmetrical Hubbert curve has a maximum in the year 2079  
 316 with an anomaly value of 1.20°C (i.e. 1.57°C above the value in 1850 and 0.46°C above  
 317 the value in 2020). The dotted curve, **b**, predicted from the asymmetrical Hubbert curve  
 318 has a maximum in the year 2145 with an anomaly value of 1.48°C (i.e. 1.85°C above the  
 319 value in 1850 and 0.74°C above the value in 2020).

320 The total emissions for the two Hubbert curves for the 650 years displayed in Fig-  
 321 ure 2 are also of interest, viz.: **a** 1824 Gt CO<sub>2</sub> and **b** 6223 Gt CO<sub>2</sub>. Thus the total emis-  
 322 sions for the Hubbert (b) curve are more than three times those for the Hubbert (a) curve

323 but the resulting temperature maxima differ by only 0.28°C. It follows that peak global  
324 average temperature is only a weakly dependent on total carbon emissions.

## 325 10 Conclusions

326 This work began as an attempt to estimate Climate Sensitivity using rigorous re-  
327 gression methods. Climate Sensitivity is defined as the temperature response to a sus-  
328 tained doubling of atmospheric CO<sub>2</sub> concentration and is used for the inter-comparison  
329 of GCMs. We soon realized that the concept itself is unrealistic because, according to  
330 (4), CO<sub>2</sub> diffuses into the ocean at a rate proportional to its concentration so that, in  
331 order to sustain the higher concentration, a high rate of emissions would need to be sus-  
332 tained indefinitely. Given the finite nature of viable hydrocarbon resources, this is an  
333 unrealistic scenario.

334 Viewed on a time scale of centuries, human exploitation of fossil fuels in the indus-  
335 trial era is generating a pulse in atmospheric carbon concentration termed “Peak Car-  
336 bon”. This, in turn, generates a pulse in global average temperature. The world is presently  
337 in the onset phase of this pulse; reasonable estimates of recoverable fossil fuel reserves  
338 suggest that the Peak Carbon pulse will reach a maximum within the next century or  
339 so. Global average temperature will follow suit with a maximum value which is less than  
340 2°C above pre-industrial values.

341 The impulse response of atmospheric CO<sub>2</sub> concentration is the response caused by  
342 a hypothetical, short variation in CO<sub>2</sub> emissions. The supposed, long-lived impulse re-  
343 sponse, widely accepted by the climate modelling community, is the most egregious flaw  
344 in the application of numerical global circulation models to climate. It implies that CO<sub>2</sub>  
345 emitted now will linger in the atmosphere for millennia. It sets the scene for the vari-  
346 ous catastrophes and tipping points presented in IPCC reports and justifies stringent emis-  
347 sion regulations. It is based on the unwarranted assumption that GCMs provide a pre-  
348 cise description of the ocean/atmosphere system as if it were some sort of clockwork mech-  
349 anism. In contrast, our statistical model implies that the atmospheric concentration of  
350 CO<sub>2</sub> is self-regulated by diffusion into the deep ocean and that the small perturbation  
351 of the global environment caused by the combustion of fossil fuels will be brief.

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Figure 1.

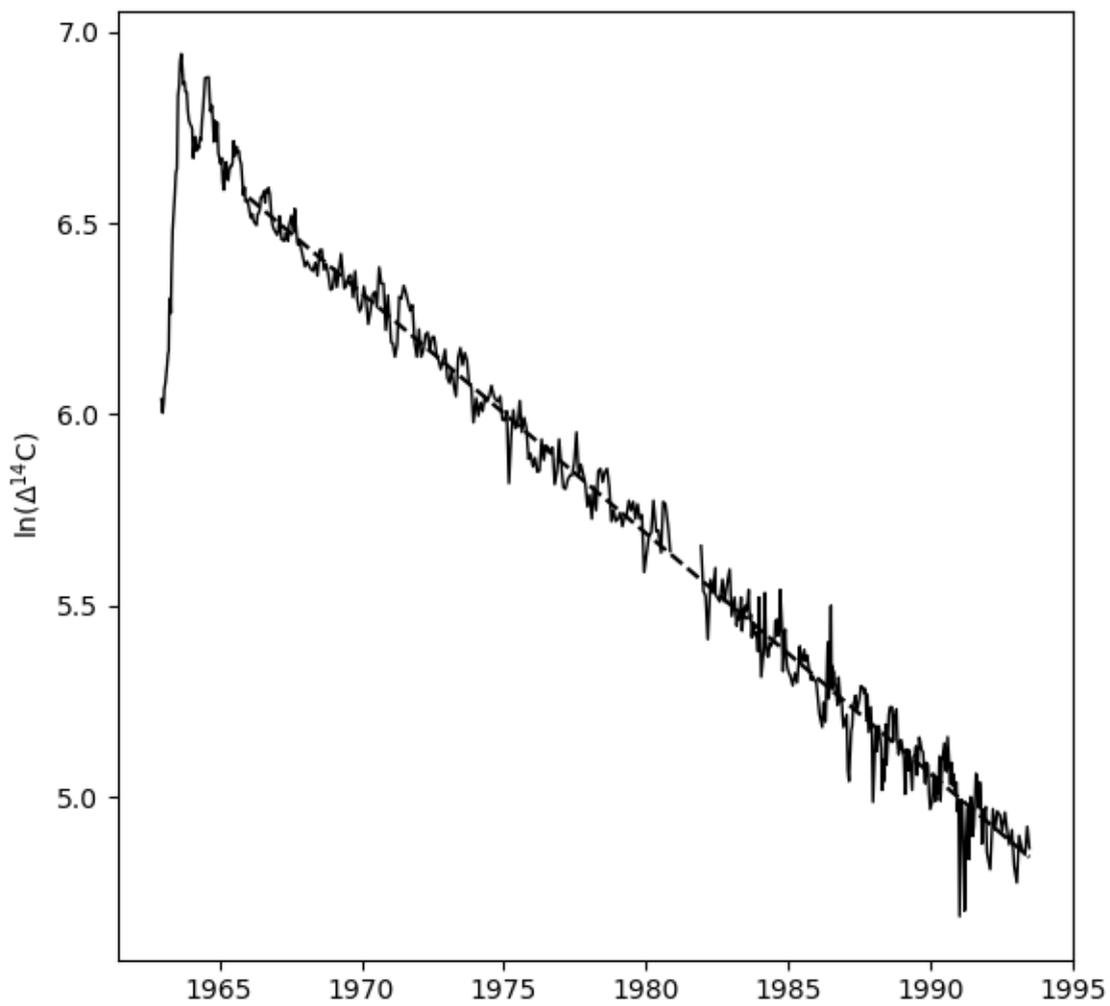


Figure 2.

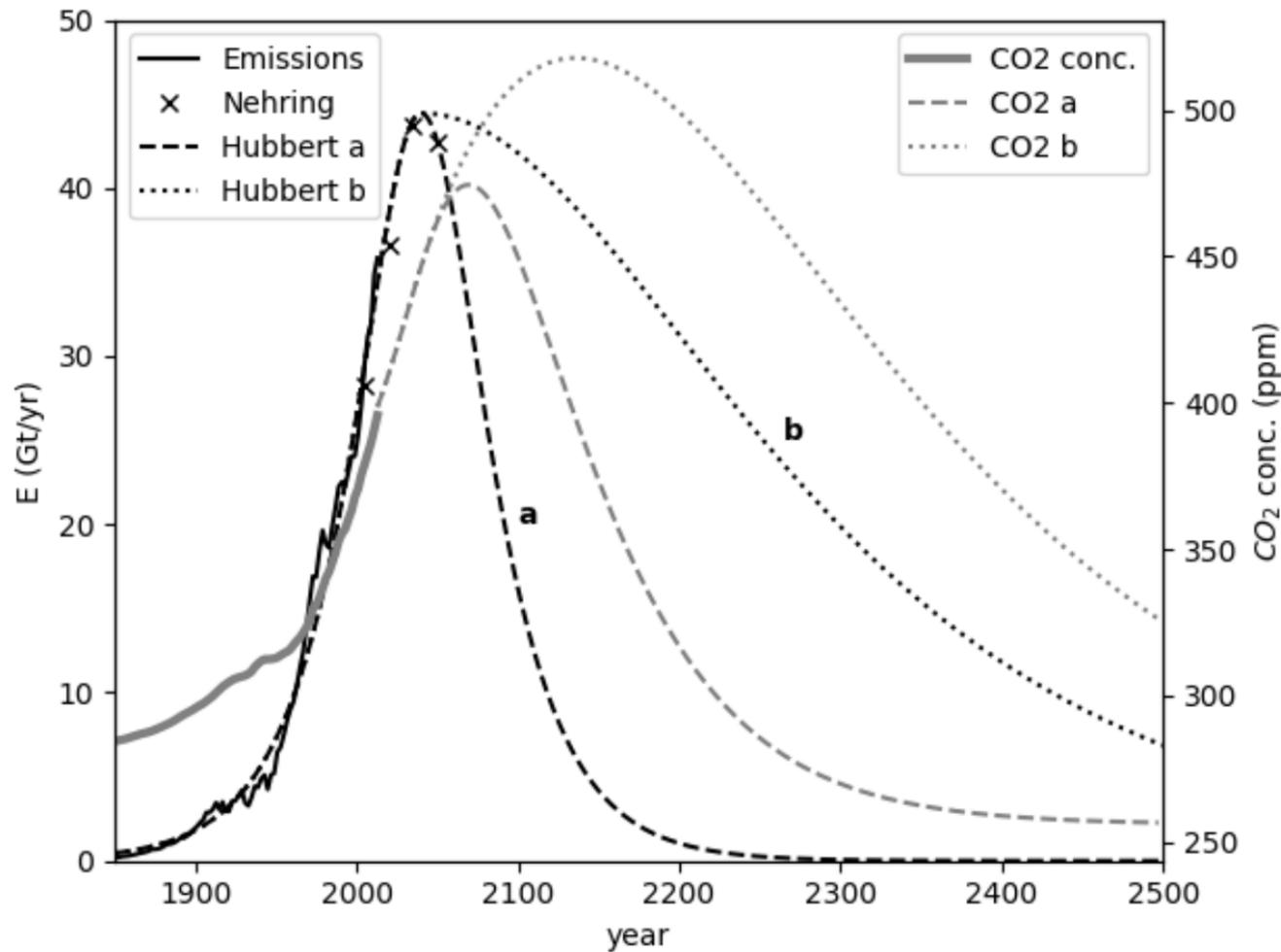


Figure 3.

