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The Vostok ice core showed that, over the last 420 kyr, Antarctic climate and concentrations of greenhouse gases (CO<sub>2</sub> and CH<sub>4</sub>) were tightly coupled. In particular, CO<sub>2</sub> seemed to be confined between bounds of about 180 ppmv in glacial periods to 280 ppmv in interglacials. The Vostok data are telling us how the coupled Earth system behaves – now we have to understand why it behaves this way. During 2004, The European Project for Ice Coring in Antarctica (EPICA) published new Antarctic temperature and dust records extending back to 740 kyr. The early part of the record shows a changed behaviour, with much weaker but longer interglacials. Greenhouse gas data for this extended period will soon be available, and obvious questions arise: Will CO<sub>2</sub> retain its interglacial bound in weak interglacials or will a new upper bound be seen? Is there any trend in average CO<sub>2</sub> over 800 kyr?

This provides an unusual opportunity for modellers and others to predict what the data will look like in advance and to allow us to explore the underlying assumptions behind ideas and models. Several groups took up this “EPICA challenge”, using models, concepts and correlations. In these posters we summarise their predictions. There is no prize and we will not be declaring a winner. But we hope that the results will lead to a greater understanding and better models.

### The EPICA Challenge: Inversion of CO<sub>2</sub> radiative forcing from the deuterium temperature proxy record

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We judge it as elusory to attempt for a mechanistic carbon cycle-biogeochemical modelling explanation in face of the difficulties to explain the CO<sub>2</sub> rise during the well-documented last transition. Instead, we utilize the climate sensitivity concept to link radiative forcing and temperature. The idea is that the EPICA δD record reflects the temperature in Antarctica caused by radiative forcing from CO<sub>2</sub> and other components. The δD record is inverted for atmospheric CO<sub>2</sub> taking into account the estimated influence of radiative forcing by ice sheets and atmospheric aerosols.

The following relationship is applied to obtain a normalized CO<sub>2</sub> record from a normalized δD record:

$$CO_{2,N}(t) = \frac{1}{1 - \alpha_{ice} - \alpha_{dust}} \cdot (\delta D_N(t) - \alpha_{ice} \cdot RF_{ice,N} - \alpha_{dust} \cdot RF_{dust,N}) \quad (1)$$

$\alpha_{ice}$  and  $\alpha_{dust}$  are the fraction of ice and dust forcing of the total radiative forcing by ice and dust and CO<sub>2</sub>.

Normalized radiative forcings by ice sheets,  $RF_{ice,N}$ , and by aerosols,  $RF_{dust,N}$ , relative to interglacial conditions are estimated from the benthic δ<sup>18</sup>O record, a proxy for ice volume, and the EPICA dust record, a proxy for atmospheric aerosol loading assuming logistic relationships. The proxy records have been normalized (δD<sub>N</sub>, δ<sup>18</sup>O<sub>N</sub>, dust<sub>N</sub>) between 0, for typical interglacial levels, and -1, for glacial values. Then, we obtain for atmospheric CO<sub>2</sub>:

$$CO_2(t) = CO_{2,IG} + (CO_{2,G} - CO_{2,IG}) \cdot \frac{1}{1 - \alpha_{ice} - \alpha_{dust}} \cdot \left( \frac{\delta D_N(t) - \alpha_{ice} \cdot (-1 + K_{ice})}{\delta D_N(t) + K_{ice}} - \alpha_{dust} \cdot \frac{dust_N(t)}{dust_N(t) + K_{dust}} \right) \quad (2)$$

An example solution is given in Figure 1.

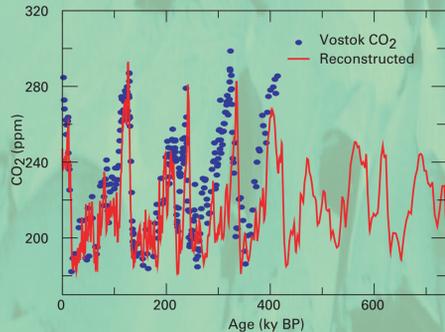
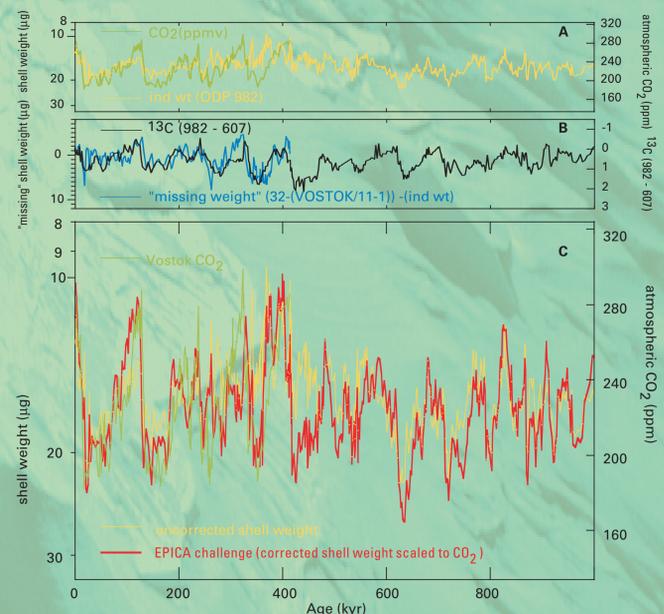


Figure 1: A comparison between the reconstructed CO<sub>2</sub> values (red) and the Vostok ice core record (blue). Atmospheric CO<sub>2</sub> has been reconstructed from equation (2) using the EPICA δD and dust record (EPICA community members, Nature, 2004) and the benthic δ<sup>18</sup>O record from Karner et al., Paleoclimatology, 2002. The parameter values used are  $\alpha_{ice}=0.3$ ,  $K_{ice}=-0.15$ ,  $\alpha_{dust}=0.05$ ;  $K_{dust}=0.2$ . The interglacial and glacial values used for the normalization of the proxy records are 269 (CO<sub>2,G</sub>) and 185 ppm (CO<sub>2,IG</sub>) for CO<sub>2</sub>, -368 and -444 permil for δD, -0.9 and +0.65 permil for δ<sup>18</sup>O, and 0 and 1400 microgram per litre meltwater for dust.

### EPICA CO<sub>2</sub> prediction using “dissolution corrected” foraminiferal shell weights

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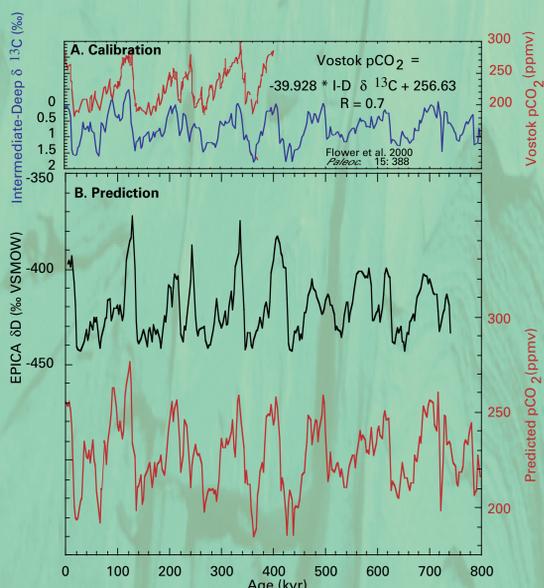
Working on the premise that the shell weights of planktonic foraminifera reflect the carbonate chemistry of surface waters and therefore respond to changes in atmospheric CO<sub>2</sub> (Barker and Elderfield, 2002) a 1Myr shell weight record for *G. bulloides* was produced from core ODP 982 in the North Atlantic (57.5°N, 15.9°W, 1134m). Despite its shallow depth, we discovered that the core appeared to have been subject to dissolution during the Mid-Brunhes (Fig. A) and as such needs to be corrected for shell weight loss at the seafloor. Correction has been made using a correlation between the hypothetical “missing weight” (scaled Vostok minus measured weight) and the difference between benthic δ<sup>13</sup>C at the sites of ODP 982 and ODP 607 in the deep North Atlantic (Fig. B) as a simple measure of foraminiferal corrosion. The resulting “corrected” shell weight record is shown in Fig. C and scaled to CO<sub>2</sub> from Vostok. The reconstruction takes no account of changing temperature, salinity or alkalinity, which will all affect the shape and amplitude of variability through their effect on carbonate chemistry. It also ignores other possible “vital” controls on shell weight such as salinity, nutrients etc. and assumes that our simple dissolution correction is valid. Allowing for such uncertainties, the features which best characterise our predicted pre-Vostok CO<sub>2</sub> are: (a) generally low interglacial CO<sub>2</sub> prior to MIS 11 (reminiscent of the pre-Vostok EPICA temperature record) and (b) particularly low CO<sub>2</sub> during glacial stage 16.



### Prediction of EPICA pCO<sub>2</sub> by Atlantic chemical stratification

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Chemical stratification and associated carbonate compensation is an important means of sequestering CO<sub>2</sub> in the deep ocean and has a strong relation to Vostok pCO<sub>2</sub> (Flower et al., 2000, Paleoclimatology, 15: 388). Here I develop an empirical calibration relating the intermediate-to-deep Atlantic δ<sup>13</sup>C gradient (ODP Sites 982 - 665) to Vostok pCO<sub>2</sub> (panel A). I derive a predicted pCO<sub>2</sub> curve for the past 800 ka compared to EPICA δD of precipitation (EPICA community members, 2004, Nature, 429: 623), which is a proxy for Antarctic air temperature (panel B). The reconstruction suggests (1) a somewhat smaller range of glacial-interglacial pCO<sub>2</sub> (~50 ppmv) for 800-400 ka compared to 400-0 ka, (2) similar levels during interglacials throughout the past 800 ka, and (3) higher pCO<sub>2</sub> levels during glaciations 800-400 ka (~210 ppmv) compared to the lowest levels from 400-0 ka. This pattern is in contrast to the EPICA δD record, which indicates cooler interglacial Antarctic air temperatures from 800-400 ka compared to subsequent interglacials. If confirmed by actual measurements, these relationships have the interesting implication that interglacial Antarctic air temperature has become more sensitive to pCO<sub>2</sub> levels during the past 400 ka.



### EPICA - CO<sub>2</sub> prediction

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I used a box model (Köhler et al., 2004, submitted to GBC) to reconstruct pCO<sub>2</sub>. Taken the same processes identified as possible causes for observed changes in pCO<sub>2</sub>, δ<sup>13</sup>C and Δ<sup>14</sup>C during Termination I especially previous terminations were matched by the simulation results with good accuracy. Due to timing uncertainties of our forcing data sets (10-20 kyr) and their coarse temporal resolution (1-3 kyr) fast fluctuations are probably not matched very well. The identification of Heinrich events with reduced NADW formation on the basis of the IRD in one North Atlantic sediment core might lead to mismatches, I therefore show simulation results with and without the Heinrich events matching Vostok ice core data on their GT4 age scale with similar accuracy (r = 0.70 and r = 0.72). The main processes determining pCO<sub>2</sub> are exchange processes of alkalinity and DIC between ocean and sediment, changes in the vertical mixing of the water column and iron fertilization of the marine export production in the Southern Ocean followed by changes in oceanic temperature and the strength of the NADW formation. The dynamics of sea level, terrestrial biosphere and gas exchange through sea ice work in the opposite direction increasing a measured rise in pCO<sub>2</sub> of 80 ppmv by another 30 ppmv. Coral reef growth impacting only during sea level high stands is not considered.

