Alkenone isotopes show evidence of active carbon concentrating mechanisms in coccolithophores as aqueous carbon dioxide concentrations fall below 7 $\mu molL^{-1}$

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Abstract.

Coccolithophores and other haptophyte algae acquire the carbon required for metabolic processes from the water in which they live. Whether carbon is actively moved across the cell membrane via a carbon concentrating mechanism, or passively through diffusion, is important for haptophyte biochemistry. The possible utilisation of carbon concentrating mechanisms also has the potential to over-print one proxy method by which ancient atmospheric CO_2 concentration is reconstructed using alkenone isotopes. Here I show that carbon concentrating mechanisms are likely used when aqueous carbon dioxide concentrations are below 7 $\mu molL^{-1}$. I ~~use~~ compile published alkenone based CO_2 reconstructions from multiple sites over the Pleistocene and recalculate them using a common methodology, which allows comparison to be made with ice core CO_2 records. Interrogating these records reveal that the relationship between proxy- and ice core- CO_2 breaks down when local aqueous CO_2 concentration falls below 7 $\mu molL^{-1}$. The recognition of this threshold explains why many alkenone based CO_2 records fail to accurately replicate ice core CO_2 records, and suggests the alkenone proxy is likely robust for much of the Cenozoic when this threshold was unlikely to be reached in much of the global ocean.

1 Introduction

Alkenones are long–chain (C_{37-39}) ethyl– and methy– ketones (Figure 1; Brassell et al. (1986); Rechka and Maxwell (1987)) produced by a restricted group of photosynthetic haptophyte algae (Conte et al., 1994). Produced by a narrow group of organisms which live exclusively in the photic zone, alkenones allow probing of algal biogeochemistry, and as alkenones are often preserved in the sedimentary record, alkenones can also provide information about past environmental conditions.

Two main proxy systems based on alkenone geochemistry exist, ~~(1) for~~ one allows reconstruction of sea surface temperature (SST) and relies on the changing degree of unsaturation of the C_{37} alkenone ($U^{K'}_{37}$) (Brassell et al., 1986) ~~and (2)~~ whilst a second for atmospheric CO_2 ~~,~~ concentration is based on reconstructing the isotopic fractionation which takes place during photosynthesis (ε_p) ~~using the carbon istopic~~ (Laws et al., 1995; Bidigare et al., 1997). It is the second system using the stable carbon

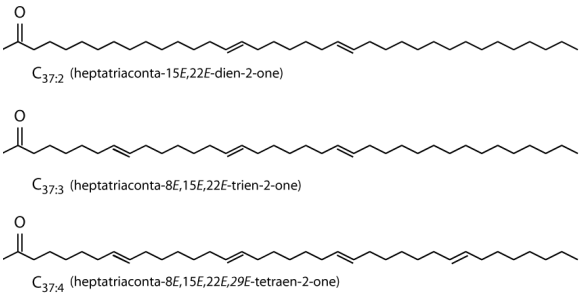

Figure 1. Alkenones are C_{37} unsaturated methyl ketones (Brassell et al., 1986; Rechka and Maxwell, 1987).

isotopic composition of the preserved alkenones for reconstructing atmospheric CO_2 concentration (referred to throughout as $CO_{2(\varepsilon_p - alk)}$) ~~(Laws et al., 1995; Bidigare et al., 1997)~~which is the focus of this study.

In the modern ocean alkenones are produced primarily by two dominant coccolithophore species; *Emiliania huxleyi* and *Gephyrocapsa oceanica*. *E. huxleyi* first appeared 290 kyr ago, and began to dominate over *G. oceanica* around 82 kyrs ago (Gradstein et al., 2012; Raffi et al., 2006). However alkenones are commonly found in sediments throughout the Cenozoic, with the oldest reported detections from mid-Albian aged black shales (Farrimond et al., 1986). Prior to the evolution of *G. oceanica* alkenones were most likely produced by other closely related species from the ~~Noelarhabdaceae~~ Noelaerhabdaceae family (Marlowe et al., 1990; Volkman, 2000). Micropaleontological and molecular data split the coccolith-bearing haptophytes into to disctinct phylogenetic clades; the Isochrysidales which contains the alkenone producers includes modern taxa *E. huxleyi* and *G. oceanica*, and fossil reticulofenestrids, whilst the non-alkeoneproducers are seperated into the order Coccolithales which includes *Coccolthus pelagicus* and *Calcidiscus leptopurus* along with most other coccolithophores.

Proxies for atmospheric CO_2 concentration including $CO_{2(\varepsilon_p - alk)}$, those based on the $\delta^{11}B$ of planktic foraminifera, geochemical modelling and stomatal density, broadly agree that over the Cenozoic atmospheric pCO_2 declined from high levels (>1000 μatm) in the "greenhouse" worlds of the Paleocene and Eocene to close to modern day values (around 400 μatm) in the Pliocene (Pagani et al., 2005, 2011; Pearson et al., 2009; Anagnostou et al., 2016; Foster et al., 2017; Sosdian et al., 2018; Super et al., 2018; Zhang et al., 2013; Beerling and Royer, 2011). However, recently discrepencies have emerged between ~~the alkenone~~ $CO_{2(\varepsilon_p - alk)}$ and other CO_2 proxies at the ~~moderate to low levels of~~ <400 μatm atmospheric CO_2 concentrations of the Pleistocene (i.e. ~~<400 ;~~ Badger et al. (2019, 2013a) and compare Badger et al. (2013b) and Pagani et al. (2009) with Martínez-Botí et al. (2015)). Whilst the longstanding ~~difference~~ differences between alkenone (Pagani et al., 1999), $\delta^{11}B$ (Foster et al., 2012) and stomatal proxies (Kürschner et al., 2008) in the Miocene CO_2 reconstructions ~~appears to be~~ have been partially resolved with new SST records (Super et al., 2018), differences remain in the Pliocene (Pagani et al., 2009; Badger et al., 2013b; Martínez-Botí et al., 2015) and Pleistocene (Badger et al., 2019).

1.1 Carbon Concentrating mechanisms

One plausible reason for ~~these discrepancies is the action~~ the discrepancies between $CO_{2(\varepsilon_p - alk)}$ and other proxies for atmospheric CO_2 is the operation of active carbon concentrating mechanisms (CCMs) in ~~the~~ haptophytes. These are potentially important as $CO_{2(\varepsilon_p - alk)}$ assumes purely passive uptake of carbon into the haptophyte cell purely via diffusion (Laws et al., 1995; Bidigare et al., 1997). The potential for CCMs to ~~be active and to~~ effect $CO_{2(\varepsilon_p - alk)}$ has long been known (Laws et al., 1997, 2002; Cassar et al., 2006) and recent work has refocussed efforts on understanding CCMs in $CO_{2(\varepsilon_p - alk)}$ (Bolton et al., 2012; Bolton and Stoll, 2013; Stoll et al., 2019; Zhang et al., 2019, 2020). Coccolithophores are thought to have low efficiency CCMs, especially compared to diatoms, dinoflagelates and phaeocystis, with evidence that CCMs play a minor role in coccolithohpore biochemistry in the CO_2 replete worlds of the early Cenozoic (Bolton et al., 2012; Reinfelder, 2011). Direct evidence from experimentation with the marine diatom *Phaeodactylum tricornutum* suggest that both passive diffusive uptake and active CCMs operate at the same time, with active uptake used to moderate internal cell CO_2 concentrations to minimise energy use during transport to carboxylation sites (Laws et al., 1997). CO_2, unlike some other nutrients, is replete within the water column, especially when considering the DIC reservoir which includes bicarbonate (HCO_3^-), carbonate (CO_3^{2-}) and dissolved CO_2 ($[CO_2]_{(aq)}$) . However, due to the relatively slow diffusion of dissolved $[CO_2]_{(aq)}$, through water and the slow kinetics of the bicarbonate to $[CO_2]_{(aq)}$ transformation, surface water $[CO_2]_{(aq)}$ can still be depleted by photosynthetic activity. This can become particularly problematic in species which form blooms, and at the cell boundary of species with limited motility. It should be no surprise therefore that many maraine photosynthetic organisms have evolved with mechanisms to concentrate carbon within the cell.

The enzyme carbonic anhydrase (CA) can catalyse the dehydration of HCO_3^- to $[CO_2]_{(aq)}$ to speed up availability of carbon if the $[CO_2]_{(aq)}$ reservoir is depleted and has been observed in several haptophytes including coccolithophores (Rost et al., 2003; Riebesell e and . The exact contribution of CA remains unclear but two possible mechanisms for CCMs have been postulated (Reinfelder, 2011) (1) CA catalyzes dehydration of HCO_3^- at the cell surface, which then allows increased CO_2 to diffuse into the cell passively and (2) HCO_3^- is transported into the cell and then converted by CA. Both of these options will likely impact the $CO_{2(\varepsilon_p - alk)}$ proxy, firstly by changing the effective $[CO_2]_{(aq)}$ within the cell (and so impacting ε_p), and secondly by imparting another carbon isotopic fractionation during CA catalyzation which is not considered by the $CO_{2(\varepsilon_p - alk)}$ proxy system. However CA activity in coccolithophores does not appear to be regulated by CO_2 as it is in diatoms and *Phaeocystis* (Rost et al., 2003) which may indicate a less well developed CCM in coccolithophores.

Calcifying coccolithophores (which include alkenone producers *E. huxleyi* and G. oceanica) may be able to utilize HCO_3^- directly as a carbon source (Trimborn et al., 2007), with precipitation of $CaCO_3$ providing an acid for the dehydration of HCO_3^- but this still requires sufficent HCO_3^- entering the cell and it is unclear whether calcification aids DIC aquisition (Riebesell et al., 2000; Zondervan et al., 2002). The light dependant leak of carbon (as CO_2 and DIC) back from haptophyte cells (including the coccolithophore *E. huxleyi*) to seawater (Tchernov et al., 2003) suggests that CCMs are energy intensive and can concentrate DIC within the cell. Even with active CCMs, it appears that in the ocean coccolithophores are CO_2 limited under some circumstances (Riebesell et al., 2007).

2 Materials and Methods

2.1 Calculating CO_2 from alkenone $\delta^{13}C$ values: The $CO_{2(\varepsilon_p - alk)}$ proxy

In this study I use the now large number of published $CO_{2(\varepsilon_p - alk)}$ records which overlap with ice core records of atmospheric CO_2 concentration (Tables 1 & 2) to explore the relationship between $CO_{2(\varepsilon_p - alk)}$ and CCMs in the Pleistocene, where our understanding of atmospheric CO_2 concentration is best.

3 ~~Materials and Methods~~

2.1 ~~Calculating from alkenone : The proxy~~

Multiple records of $CO_{2(\varepsilon_p - alk)}$ have been published for the Pleistocene (Figure 2, Table 1) allowing direct comparision with ice core based CO_2 records (Table 2). These records are globally distributed in longitude, but are concentrated at low latitude sites, largely as there is a general preference for sites which have (in the modern ocean) surface waters close to equilibrium with the atmosphere (Figure 2, Table 1). In longer term palaeoclimate studies there has also been a preference for low latitude, gyre sites in the belief that these sites are more likely to be oceanographically stable over long time intervals (Pagani et al., 1999). Most of the records included here (Table 1, Figure 2) were generated with the aim to reconstruct atmospheric CO_2 concentration, however one, the MANOP C Site of Jasper et al. (1994), was used to explicitly reconstruct changing disequilibrium due to oceanographic frontal changes over time, and so is excluded from the following analysis.

Whilst these sites do only span a relatively small latitudinal extent, the diversity of settings does allow for investigation of any secondary controls on alkenone $\delta^{13}C$ values ($\delta^{13}C_{alkenone}$). In particular, differences in oceanographic setting and SST to test the hypothesis that low $[CO_2]_{(aq)}$ breaks the relationship between $\delta^{13}C_{alkenone}$ and atmospheric CO_2 concentration, as might be expected if haptophytes are able to actively ~~uptake~~ take up carbon from seawater to meet metabolic demand – i.e. activate CCMs.

To facilitate fair comparision between sites and consistent comparision with the ice core records, all $CO_{2(\varepsilon_p - alk)}$ records ~~I~~ were recalculated using a consistent approach. The approach is based on Bidigare et al. (1997) which updated the initial approach of Jasper and Hayes (1990) to $CO_{2(\varepsilon_p - alk)}$.

This approach removes some additional corrections used in the original publication of the records (such as growth rate adjustment for NIOP 464 (Palmer et al., 2010)) but does allow for direct comparison to be made. For all sites the 'b' term was estimated using modern day surface $[PO_4^{3-}]$ (Bidigare et al., 1997; Pagani et al., 2009)

An overview of how $CO_{2(\varepsilon_p - alk)}$ data are typically generated is given in Badger et al. (2013b).

Briefly, to calculate ε_p requires the stable carbon isotopic composition of the dissolved CO_2 ($\delta^{13}C_{CO_{2(aq)}}$) and haptophyte biomass ($\delta^{13}C_{org}$). The isotopic fractionation between $\delta^{13}C_{alkenone}$ and $\delta^{13}C_{org}$ is first corrected assuming a contstant frac-

Table 1. Sites with Pleistocene $CO_{2(\varepsilon_{p-alk})}$ records. Note that the MANOP Site C record was generated to track changes in surface water–atmosphere equilibrium not atmospheric pCO_2, so although included here for completeness, is not included in the analysis

Site	Age interval (kyr)	Latitude	Longitude	Water depth (m)	Distance from coast (km)	Reference
05PC-21	0.5–188	38° 24'N	131° 33'E	1721	108	Bae et al. (20
DSDP 619	3–92	27° 11.61' N	91° 24.54' W	2259	489	Jasper and Ha
NIOP 464	7.8–29	22° 9' N	63° 21'E	1470	333	Palmer et al. (
ODP 999	111–258	12° 44.639' N	78° 44.360' W	2839	249	Badger et al. (
ODP 925	20–580	4° 12.249' N	43° 29.334' W	~~Zhang et al. (2013) DSDP 619~~ 3042	~~27° 11.61' N~~ 626	~~91° 24.54' W~~
MANOP Site C	0.8–253	0° 57.2" N	138° 57.3' W	~~Jasper et al. (1994) NIOP 464~~ 4287	~~22° 15.4' N~~ 998	~~63° 31.1'E Pa~~
GeoB 1016-3	1.3–196	11° ~~46~~46.2' S	11° ~~40~~40.9' E	~~Andersen et al. (1999) 05PC-21~~ 3410	~~37° 30.11'N~~ 185	~~129° 42.86'E~~

*The full record for ODP Site 925 extends to 38.62 Ma

Table 2. Sources of ice core data, as compiled by Bereiter et al. (2015). WAIS - West Antarctic Ice Sheet, TALDICE - TALos Dome Ice CorE, EDML - EPICA Dronning Maud Land

Age interval (kyr)	Ice core location	Reference
−0.051–1.8	Law Dome	Rubino et al. (2013)
1.8–2	Law Dome	MacFarling Meure et al. (2006)
2–11	Dome C	Monnin et al. (2001, 2004)
11–22	WAIS	Marcott et al. (2014)
22–40	Siple Dome	Ahn and Brook (2014)
40–60	TALDICE	Bereiter et al. (2012)
60–115	EDML	Bereiter et al. (2012)
105–155	Dome C Sublimation	Schneider et al. (2013)
155–393	Vostok	Petit et al. (1999)

tionation ($\varepsilon_{\text{alkenone}}$) of 4.2 ‰ ~~(Popp et al., 1998; Bidigare et al., 1997)~~(Garcia et al., 2013; Popp et al., 1998; Bidigare et al., 1997)
:

$$\varepsilon_{\text{alkenone}} = \frac{\delta^{13}C_{alkenone} + 1000}{\delta^{13}C_{org} + 1000} - 1 \quad (1)$$

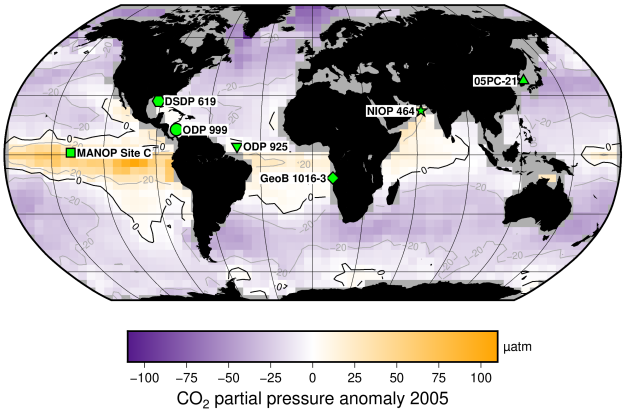

Figure 2. Study sites relative to mean annual surface ocean CO_2 disequilibrium for 2005. Sites are globally distributed in longitude but restricted in latitude, as generally sites are choosen to be close to surface water equilibrium with the atmosphere. Sites used for this study are indicated, over the mean annual surface ocean disequilibrium for 2005 calculated from Takahashi et al. (2014). The MANOP C Site (Jasper et al., 1994) was choosen to study the disequilbrium at that site, so is shown here but not used in the following analyses. Site symbols are used throughout the figures: ODP 999 - circle, 05PC-21 - triangle, ODP 925 - inverted triangle, DSDP 619 - hexagon, MANOP Site C - square, NIOP 464 - star, GeoB 1016-3 - diamond.

The isotopic composition of dissolved inorganic carbon (DIC) is estimated using (ideally) the $\delta^{13}C$ value of planktic foraminifera and the temperature-dependant fractionation between calcite and $[CO_2]_{(g)}$ experimentally determined by Romanek et al. (1992), where T is sea surface temperature in degrees Celsius (SST):

$$\varepsilon_{calcite-CO_{2(g)}} = 11.98 - 0.12T \tag{2}$$

The value the cabon isotopic composition of $CO_{2(g)}$ ($\delta^{13}C_{CO_{2(g)}}$) can then be calculated:

$$\delta^{13}C_{CO_{2(g)}} = \frac{\delta^{13}C_{carbonate} + 1000}{\varepsilon_{calcite-CO_{2(g)}}/1000 + 1} - 1000 \tag{3}$$

From this $\delta^{13}C_{CO_{2(aq)}}$ can be calculated using the relationship experimentally determined by Mook et al. (1974):

$$\varepsilon_{CO_{2(aq)}-CO_{2(g)}} = \frac{-373}{T + 273.15} + 0.19 \tag{4}$$

and

$$\delta^{13}C_{CO_{2(aq)}} = \left(\frac{\varepsilon_{CO_{2(aq)}-CO_{2(g)}}}{1000} + 1\right).\left(\delta^{13}C_{CO_{c(g)}} + 1000\right) - 1000$$

$$\delta^{13}C_{CO_{2(aq)}} = \left(\frac{\varepsilon_{CO_{2(aq)}-CO_{2(g)}}}{1000} + 1\right) \cdot \left(\delta^{13}C_{CO_{c(g)}} + 1000\right) - 1000 \quad (5)$$

Finally ε_{p} can be calculated:

$$\varepsilon_p = \left(\frac{\delta^{13}C_{CO_{2(aq)}} + 1000}{\delta^{13}C_{org} + 1000} - 1\right).1000 \quad (6)$$

and from that $[\mathrm{CO_2}]_{(\mathrm{aq})}$ is calculated using the isotopic fractionation during carbon fixation (ε_{f}) and 'b', which represents the summation of physiological factors:

$$[CO_2]_{(aq)} = \frac{b}{\varepsilon_f - \varepsilon_p} \quad (7)$$

Here ε_{f} is assumed to be a constant 25 ‰ (Bidigare et al., 1997). In the modern ocean the 'b' term, which accounts for physiological factors such as cell size and growth rate, shows a close correlation with $[\mathrm{PO_4^{3-}}]$ (Bidigare et al., 1997; Pagani et al., 2009). However, the relationship between b, growth rate and $[\mathrm{PO_4^{3-}}]$ has recently been questioned (Zhang et al., 2019, 2020) but for the purposes of this analysis is assumed to hold. This is discussed further below. Values for SST, $\delta^{13}\mathrm{C_{alkenone}}$, $\delta^{13}\mathrm{C_{carbonate}}$, salinity and $[\mathrm{PO_4^{3-}}]$ are either taken from the original publications or estimated from modern ocean estimates (Takahashi et al., 2009; Antonov et al., 2010; Garcia et al., 2013; Locarnini et al., 2013).

Providing that the atmosphere is in equilibrium with surface water, the concentration of atmospheric $\mathrm{CO_2}$ can be calculated from $[\mathrm{CO_2}]_{(\mathrm{aq})}$, (and vice versa if atmospheric $\mathrm{CO_2}$ concentration is known) using Henry's law:

$$pCO_2 = \frac{[CO_2]_{(aq)}}{K_{\mathrm{H}}} \quad (8)$$

The solubility coefficient ($\mathrm{K_H}$) is dependant on salinity and SST, and here is calculated following the paramterization of Weiss (1970, 1974).

3 Results

3.1 Multi-site comparisons between $\mathrm{CO_{2(\varepsilon_p-alk)}}$ and the ice core records

Across the six sites included in this analysis, there are 217 $\mathrm{CO_{2(\varepsilon_p-alk)}}$-based estimates of atmospheric $\mathrm{CO_2}$ concentration over the past 260 Ka for comparison with the ice core records (Table 2; Bereiter et al. (2015)). When all $\mathrm{CO_{2(\varepsilon_p-alk)}}$ estimates

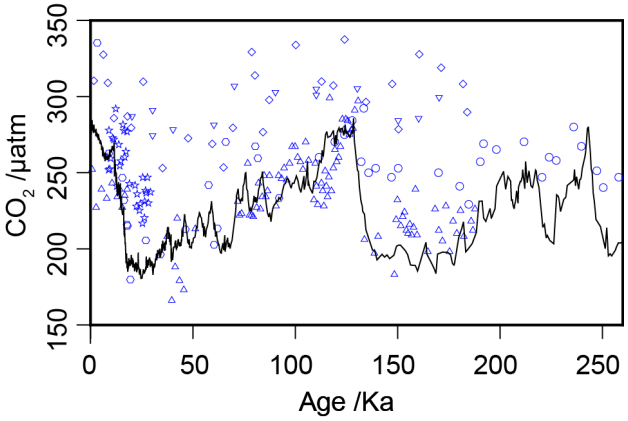

Figure 3. Compiled $CO_{2(\varepsilon_p - alk)}$-based estimates of atmospheric CO_2 concentration over the past 260 Ka (blue circles), with the ice core compilation of Bereiter et al. (2015) shown as the solid black line. Full sources for the ice core and $CO_{2(\varepsilon_p - alk)}$ records are in Tables 2 and 1

are considered together over 260 Ka, this compilation of proxy-based records fails to replicate the ice core record (Figure 3). This has already been noted at specific sites (e.g. Site 999 in the Caribbean Badger et al. (2019)) but this is the first time that all available records coincident with the Pleistocene ice core records have been compiled using a common methodology. Notably the $CO_{2(\varepsilon_p - alk)}$ based estimates are rarely lower than time-equivalent ice core estimate, but frequently higher. Given that haptophytes require carbon to satisfy metabolic demand, this is perhaps unsurprising; if at times of low carbon availibily haptophytes can switch from passive to active uptake to satisfy metabolic demand, it would be times of low atmospheric CO_2 concentration (and so lower $[CO_{2(aq)})$ when the active uptake is most likely to be needed. As $CO_{2(\varepsilon_p - alk)}$-based estimates of atmospheric CO_2 concentration rely on the assumption of a purely ~~difusive~~ diffusive uptake of carbon, it is therefore likely that the proxy would perform ~~least well~~ worse at times of low atmospheric CO_2 concentration.

The ~~haptophyes~~ haptophytes do not directly interact with the atmosphere, obtaining their carbon from dissolved carbon. As it is not only atmospheric CO_2 concentration which controls the concentration of dissolved carbon ($[CO_2]_{(aq)}$), but also temperature, alkalinity and other oceanographic factors which control the equilibrium state between surface waters at the atmosphere, (Figure 2) the multiple sites in different settings now give the opportunity to test whether other factors are important in controlling the accuracy of $CO_{2(\varepsilon_p - alk)}$.

To produce time-equilvalent estimates of atmospheric CO_2 concentration for comparison with the ice core records, a simple linear interpolation of the Bereiter et al. (2015) compilation was initially used (Figure 4). This assumes that both the age model of the ice core and the published age models of the sites are correct and equivalent. This is almost certainly not the case, and so for the calculations below, a ± 3000 year uncertainty is included for ages of both the ice core and $CO_{2(\varepsilon_p - alk)}$ values. Figure 4 shows that $CO_{2(\varepsilon_p - alk)}$-based atmospheric CO_2 concentration agree with ice core CO_2 at some sites and at some times, but not ~~througout. Sites such as~~ throughout. Sites 05-PC21 (Bae et al., 2015) and DSDP Site 619 (Jasper and Hayes, 1990) perform quite well, througout, whilst ~~others~~ ODP Site 999 (Badger et al., 2019) and NIOP 464 (Palmer et al., 2010)

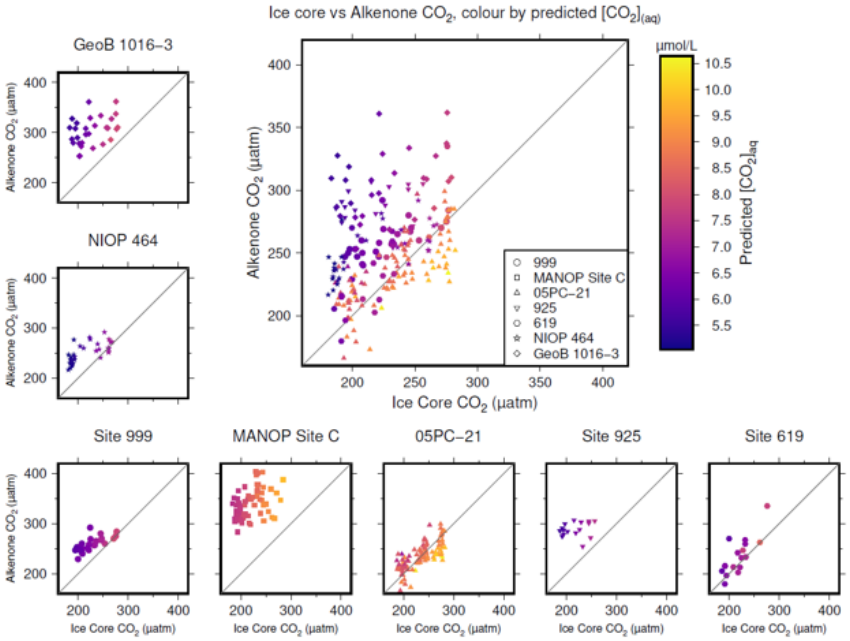

Figure 4. Crossplots of $CO_{2(\varepsilon_p - alk)}$-based atmospheric CO_2 concentration (y-axes) vs the time-equivalent ~~estmiate~~ estimate from ice core records (x-axes; Bereiter et al. (2015); Table 2)). The large panel compiles all sites, with the exception of MANOP Site C, as explained in the text. Symbols are coloured by predicted $[CO_{2(aq)}]$ for each site and time as explained in the text. Full sources for alkenone data are shown in Table 1. A 1:1 line is included in all plots for comparison.

only appear to agree at higher values of CO_2, ~~such as ODP Site 999 (Badger et al., 2019) and NIOP 464 (Palmer et al., 2010) , whilst at some locations~~ at ODP Site 925 (Zhang et al., 2013) and GEoB 1016-3 (Andersen et al., 1999) there is very little overlap between the two methods of reconstructing atmospheric CO_2 ~~(such as ODP Site 925 (Zhang et al., 2013) and GEoB 1016-3 (Andersen et al., 1999))~~concentration.

To explore whether $[CO_2]_{(aq)}$ is an imporant influence on $CO_{2(\varepsilon_p - alk)}$, I calculate predicted $[CO_2]_{(aq)}$ ($[CO_2]_{(aq)-predicted}$) for each of the samples. To calculate $[CO_2]_{(aq)-predicted}$, the ice core value of atmospheric CO_2 concentration at the equivalent time interval is used in combination with Eq. 8 to calculate $[CO_2]_{(aq)}$ at the time of alkenone production. Reconstructed estimates of SST and salinity are used as for $CO_{2(\varepsilon_p - alk)}$ above, along with any estimated surface water-atmosphere disequilibrium. Points in Figure 4 are then coloured by $[CO_2]_{(aq)-predicted}$.

Inspection of Figure 4 suggests a connection between ($[CO_2]_{(aq)-predicted}$) and the skill of $CO_{2(\varepsilon_p - alk)}$ to reconstruct atmospheric CO_2 concentration, with the points clustering around the 1:1 line lighter in colour (so with higher $[CO_2]_{(aq)-predicted}$), whilst points falling away from the 1:1 line have lower $[CO_2]_{(aq)-predicted}$.

To explore this relationship, I progressively restricted the included samples on the basis of $[CO_2]_{(aq)-predicted}$, and at each stage calculated a Pearson correlation co-efficient for each subset. Under this analysis the correlation co-efficient progressively

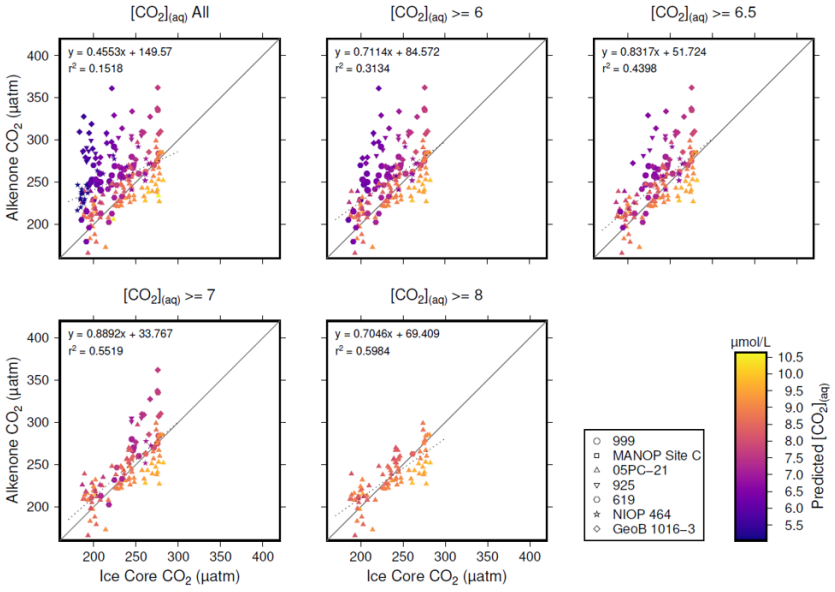

Figure 5. Crossplots of $CO_{2(\varepsilon_p - alk)}$-based atmospheric CO_2 concentration (Table 1; y-axes) vs the time-equivalent estimate from ice core records (x-axes; Bereiter et al. (2015); Table 2)). The sample of published vales of $CO_{2(\varepsilon_p - alk)}$ was progressively restricted by $[CO_2]_{(aq)-predicted}$, indicated by the subplot titles. Individual values are coloured by $[CO_2]_{(aq)-predicted}$, and Sites indicated by shape (see key). Pearson correlation co-efficients and equations of best fit are shown in each panel, along with a 1:1 line.

increased as more of the low $[CO_2]_{(aq)-predicted}$ samples were excluded (Figure 5). All analyses were performed in R (R Core Team, 2020) using RStudio (RStudio Team, 2020).

This suggests that the fidelity of the $CO_{2(\varepsilon_p - alk)}$ depends on the concentration of $[CO_2]_{(aq)}$, improving at higher levels of $[CO_2]_{(aq)}$.

To further investigate this potential relationship, I progressively exclude samples based on $[CO_2]_{(aq)-predicted}$ with a step size of 0.05 $\mu mol L^{-1}$, again calculating Pearson correlation coefficients between ice core and $CO_{2(\varepsilon_p - alk)}$ for each subsample of the population. The result is shown in Figure 6. Here the analysis shows, similar to Figure 5, that as the samples with lowest $[CO_2]_{(aq)-predicted}$ are progressively removed, the correlation between ice core and $CO_{2(\varepsilon_p - alk)}$ increases. Furthermore, this continues only up until $[CO_2]_{(aq)-predicted}$ reaches 7 $\mu mol L^{-1}$. Above this, the correlation coefficient plateaus, until the subsample reaches such a small size that spurious correlations become important (Figure 6b).

3.2 Sensitivity and Uncertainty Tests

~~As it is not impossible that that a similar pattern could emerge if the dataset were particularly shaped so that there was increased density~~ It is possible that the pattern seen in Figure 6b could emerged if the from a dataset shaped with increasing densisity surrounding the 1:1 correlation line ~~,~~without being driven by changes in $[CO_2]_{(aq)-predicted}$. To explore this possiblity I ran

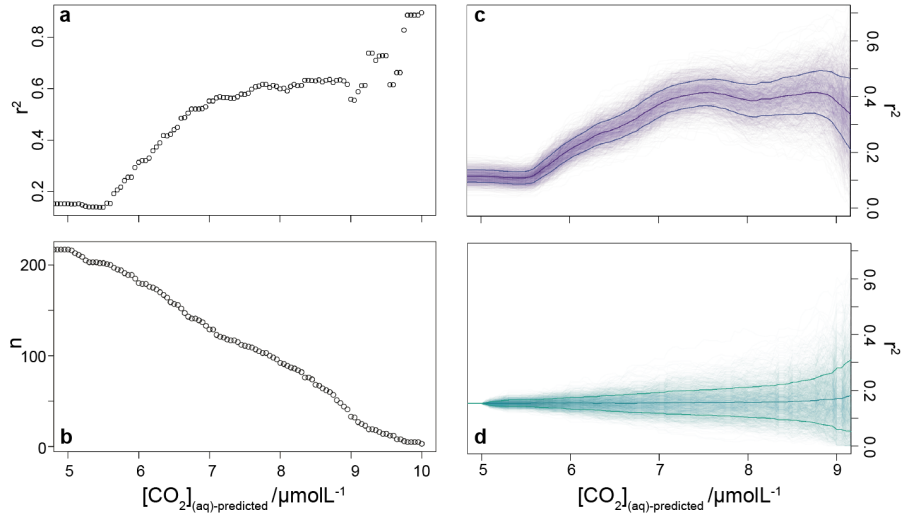

Figure 6. Pearson correlation coefficient (panel a) of a reducing sample of all compiled $CO_{2(\varepsilon_p-alk)}$ (Table 1) vs the time-equivalent estimate from ice core records (Bereiter et al. (2015); Table 2). The sample reduces stepwise by 0.05 $\mu molL^{-1}$, and the number of records in each subsample is shown in panel b. Panel c shows a 1000 member Monte Carlo analysis, whereby uncertainty in $CO_{2(\varepsilon_p-alk)}$ and age is considered, as detailed in the text. Panel d shows a similar 1000 member Monte Carlo analysis, but with random sampling of the whole $CO_{2(\varepsilon_p-alk)}$ population so that the number of samples is equivalent to the dataset shown in panel c, ie the size of the sample follows that shown panel b. Means and one σ uncertainties are shown as the bold lines.

a series of sensitivity experiments. In these, rather than reducing the sample by filtering by $[CO_2]_{(aq)-predicted}$, the whole dataset (Table 1) was randomly ordered, and then stepwise subsampled~~so that the number of samples equalled the number of values for each value of~~ . To make this equivalent to the $[CO_2]_{(aq)-predicted}$ ~~(ie for each point in Figure 6, an equivalently sized but randomly selected~~sample ~~was made such that for any equivalent value of the randomly ordered sample had an equivalent n~~ analysis above, I set the size of each subsample to be equal to each step in the original analysis. This produces a randomly selected, but same sized sub sample such that the size of the subsample reduces in the same way as shown in Figure 6b). Pearson correlation coefficients were ~~calculated~~ calculated for each subsample as above~~.~~ , and I repeated this 1000 times, with the order of each sample randomized each time.

To allow for possible age model uncertainties, a 3000 year (1σ) uncertainty was also applied to each sample. This uncertainty was applied to the age of each sample prior to sampling of the ice core record, and is applied as a normally distributed uncertainty. Uncertainty in $CO_{2(\varepsilon_p-alk)}$ measurements is typically calculated using Monte Carlo modelling of all the parameters (i.e Pagani et al. (1999); Badger et al. (2013a, b)), however this was not done in all the published work (Table 1), and some differences in approach ~~was~~ were found accross the published work. Therefore to create $CO_{2(\varepsilon_p-alk)}$ uncertainty estimates for each value in this study, I emulate the uncertainties based on the $CO_{2(\varepsilon_p-alk)}$ value. I built a simple emulator (Figure 7) by running

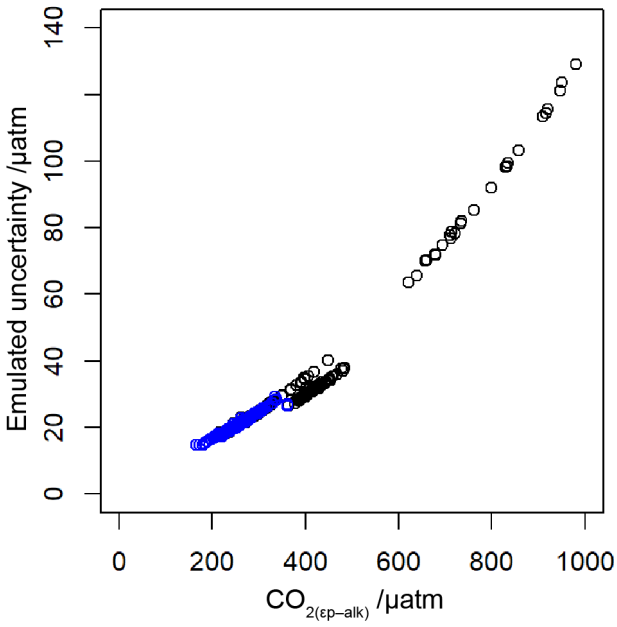

Figure 7. Emulated uncertainty in $CO_{2(\varepsilon_p-alk)}$, generated by running Monte Carlo uncertainty models for all sites in Table 1 applying the same approach to uncertainty as Badger et al. (2013a, b). Estimates used in this study are highlighted in blue.

Monte Carlo uncertainty estimates for all of the included datasets (Table 1) using the same estimates of uncertainty for each variable in the $CO_{2(\varepsilon_p-alk)}$ calculation as applied in Badger et al. (2013a, b). This then allows the uncertainty to be included in the $[CO_2]_{(aq)-predicted}$ calcuation as well as $CO_{2(\varepsilon_p-alk)}$, and allowed for uncertainty estimates to be site-ambivalent.

The result is shown in Figure ~~??~~ 6c,d, and suggests that the 7 $\mu molL^{-1}$ break point remains valid. The absolute value of r^2 is reduced, even at higher $[CO_2]_{(aq)-predicted}$, but this would be expected given the addition of uncertainty in age model, as the published age is most likely to align with the ice core. Given the rapid rate of change at deglaciations, this effect is likely to be particularly pronounced in this dataset as many records have high temporal resolution around deglaciations in order to attempt to resolve them. Any small age model offset introduced by the error modelling in these intervals also clearly has the potential to induce large differences between the $CO_{2(\varepsilon_p-alk)}$ and ice core values. Figure ~~??~~ 6c,d clearly demonstrates that it is the filtering by $[CO_2]_{(aq)-predicted}$ rather than any spurious correlations which determine the shape of the data in ~~Figures 6and ??~~Figure 6a.

~~Pearson correlation coefficient of a reducing sample of all compiled (Table 1) vs the time-equivalent estimate from ice core records (Bereiter et al. (2015); Table 2). As in Figure 6 the sample reduces stepwise by 0.05 . Panel a shows a 1000 member Monte Carlo analysis, whereby uncertainty in and age is considered, as detailed in the text. Panel b shows a similar 1000 member Monte Carlo analysis, but with random sampling of the whole population so that the number of samples is equivalent~~

~~to the dataset shown in panel a, ie the size of the sample follows that shown in Figure 6b. Means and one σ uncertainties are shown as the bold lines.~~

4 Discussion

The plateau in r^2 in Figures 6a and ~~??a~~ 6c suggest that below a $[CO_2]_{(aq)-predicted}$ of ~ 7 $\mu molL^{-1}$ $CO_{2(\varepsilon_p-alk)}$ is no longer as good a predictor of ice core CO_2 as when $[CO_2]_{(aq)-predicted} > 7$ $\mu molL^{-1}$. This is clear from comparing the relationship between samples where $[CO_2]_{(aq)-predicted} < 7$ $\mu molL^{-1}$ with those where $[CO_2]_{(aq)-predicted} > 7$ $\mu molL^{-1}$ in Figure 8. Here the r^2 for the former of 0.15 is substantially less than the latter of 0.55. I suggest that this is because below this threshold, the fundamental assumption of $CO_{2(\varepsilon_p-alk)}$; that carbon is passively taken up by haptophytes, no longer holds true. One obvious explanation for why this would be the case is that at low levels of $[CO_2]_{(aq)}$ haptophytes have to ~~actively uptake carbon~~ rely more on active up take of carbon via CCMs in order to satisfy metabolic demand.

~~Correlations between and ice core where > 7 (black symbols) and < 7 (red symbols).~~

Similar behaviour has been recognised in some culture studies (Laws et al., 1997, 2002; Cassar et al., 2006), with some evidence that the diatom *Phaeodactylum tricornutum* ~~even~~ has a similar CCM threshold ~~(Laws et al., 1997) but this study is the clearest evidence of the behaviour in alkenone based studies of the environment~~of 7 $\mu molL^{-1}$ (Laws et al., 1997). Whilst the evidence for the mechanism of CCM is poorer for coccolithophores than it is for diatoms, any CCM would be expected to compromise the $CO_{2(\varepsilon_p-alk)}$ proxy, either by increased supply of $[CO_2]_{(aq)}$, further carbon isotopic fractionation effects during carbon transport or both (Stoll et al., 2019).

By applying a threshold value for $[CO_2]_{(aq)-predicted}$ of 7 $\mu molL^{-1}$ to the published records (Table 1) values of $CO_{2(\varepsilon_p-alk)}$ which are ~~influence~~ influenced by active CCMs can be eliminated. Recognition of this new threshold value of $[CO_2]_{(aq)-predicted}$ allows for a new record of Pleistocene $CO_{2(\varepsilon_p-alk)}$ to be compiled. This compilation then much better replicates the glacial-interglacial pattern of CO_2 change over the last 260 Ka (Figure 9). Whilst this present compilation does rely on ice core CO_2 records to estimate $[CO_2]_{(aq)-predicted}$, and therefore has little direct utility as a CO_2 record, it does demonstrate that recognition of ~~the CCM threshold~~ a threshold response allows accurate CO_2 reconstruction using $CO_{2(\varepsilon_p-alk)}$. This may represent the point at which isotopic effects of CCMs (plausibly through increased CA activity or HCO_3^- dehydration to meet C demand) overwhelms the assumptions of the $CO_{2(\varepsilon_p-alk)}$ proxy. This, and the behaviour shown in Figures 6a and ~~??a suggests the~~ 6c suggests that from the standpoint of the $CO_{2(\varepsilon_p-alk)}$ proxy CCMs may effectively be considered either active or not, and that when $[CO_2]_{(aq)}$ is plentiful passive uptake dominates, at least sufficiently in most oceanographic settings that $CO_{2(\varepsilon_p-alk)}$ can accurately record atmospheric CO_2 concentration. This implies that if areas of the ocean (or intervals of time) with low $[CO_2]_{(aq)}$ can be avoided, accurate reconstructions of atmospheric CO_2 concentration can be acquired using $CO_{2(\varepsilon_p-alk)}$.

As $[CO_2]_{(aq)}$ is effected by both SST via the temperature dependance of the Henry's law constant and atmospheric CO_2 concentration, for $CO_{2(\varepsilon_p-alk)}$ to be effective in reconstructing atmospheric CO_2 concentration, areas of warm water (i.e. tropical or shallow shelf regions) under relatively low atmospheric CO_2 concentration must be avoided. However, as the atmospheric CO_2 control renders the global surface ocean sufficiently replete in $[CO_2]_{(aq)}$ at Pliocene-like levels of atmospheric

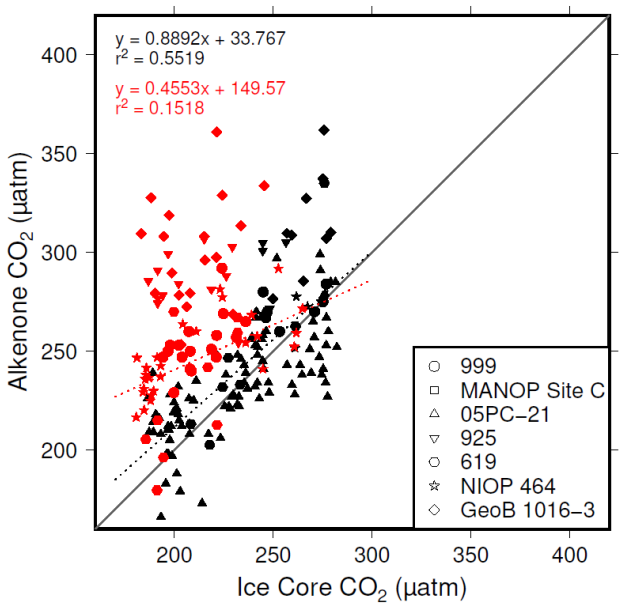

Figure 8. Correlations between $CO_{2(\varepsilon_p-alk)}$ and ice core CO_2 where $[CO_2]_{(aq)-predicted} > 7$ µmolL^{-1} (black symbols) and $[CO_2]_{(aq)-predicted} < 7$ µmolL^{-1} (red symbols).

CO_2 concentration and above (Martínez-Botí et al., 2015) at all but the warmest surface ocean temperatures, $CO_{2(\varepsilon_p-alk)}$ is likely to be a reliable system for most of the Cenozoic. It is only in the Pleistocene that atmospheric CO_2 concentration is low enough for CCMs to be widely active accross the surface ocean, with the low CO_2 glacials providing the most difficulty (Badger et al., 2019). This finding aligns well with evidence that CCMs developed in coccolithophores as a reponse to declining atmospheric CO_2 concentration through the Cenozoic, and were developing in $[CO_2]_{(aq)}$–limited parts of the ocean in the late Miocene at the earliest, and likely not widespread until the Plio-Pleistocene (Bolton et al., 2012; Bolton and Stoll, 2013).

~~Recent has attempted~~ There have been recent attempts to correct for the existance of CCMs in ~~palaeo-records~~ $CO_{2(\varepsilon_p-alk)}$-based reconstructions of atmospheric CO_2 concentration s (Zhang et al., 2019; Stoll et al., 2019; Zhang et al., 2020). However, these assume that CCMs are always active, and ~~that Pleistocene records can~~ crucially do not fundamentally break the relationship between ε_p values and atmospheric CO_2 concentration. However if this is not the case, and the relationship between ε_p values and atmospheric CO_2 concentration fails at Pleistocene levels of atmospheric CO_2 then Pleistocene records cannot be used to ~~correct for them~~ develop corrections of $CO_{2(\varepsilon_p-alk)}$ to be applied throughout the Cenozoic. If, as suggested by the analyses presented here, CCMs *only* act at low $[CO_2]_{(aq)}$, and largely only in conditions prevalent throught the late Pliocene and Pleistocene, it is plausible that corrections based on Pleistocene records could over-compensate for CCMs in the rest of the Cenozoic, when the assumption of passive carbon uptake inherent in $CO_{2(\varepsilon_p-alk)}$ as traditionally applied may still be valid.

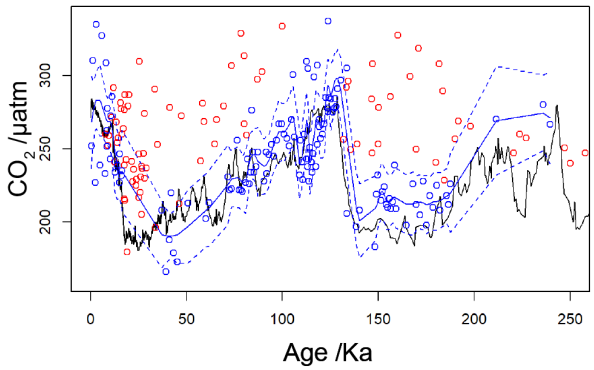

Figure 9. Revised compilation of Pleistocene $CO_{2(\varepsilon_p - alk)}$ vs ice core records. The compiled published records (Table 1) are shown as circles, coloured red where $[CO_2]_{(aq)-predicted}$ is below a threshold of 7 $\mu mol L^{-1}$, and blue where $[CO_2]_{(aq)-predicted} > 7$ $\mu mol L^{-1}$. The solid blue line is a loess filter (span 0.1) through the $[CO_2]_{(aq)-predicted} > 7$ $\mu mol L^{-1}$ values, with 95 % confidence intervals (dashed blue line). The black line is the ice core compilation of Bereiter et al. (2015) (Table 2).

5 Conclusions

Reconstructions of past atmospheric CO_2 concentration with proxy tools like $CO_{2(\varepsilon_p - alk)}$ are critical ~~to~~ for understanding how the Earth's climate system operates, ~~so~~ as long as the tools used can be relied upon to be accurate and precise. This re-analysis of existing Pleistocene $CO_{2(\varepsilon_p - alk)}$ records reveals that below a critical threshold of $[CO_2]_{(aq)}$ of 7 $\mu mol L^{-1}$ the relationship between $\delta^{13}C_{alkenone}$ and atmospheric CO_2 concentration breaks down, plausibly because below this threshold haptophytes are able to actively ~~update~~ take up carbon using CCMs in order to satisfy metabolic demand.

Although reconstructing the low levels of atmospheric CO_2 concentration in the Pleistocene glacials and areas of the global ocean where $[CO_2]_{(aq)}$ is less than 7 $\mu mol L^{-1}$ will be impossible, for much of the Cenozoic, the $CO_{2(\varepsilon_p - alk)}$ proxy retains utility. If care is taken to avoid regions and oceanographic settings where $[CO_2]_{(aq)}$ ~~may be expected~~ is expected to be abnormally low, $CO_{2(\varepsilon_p - alk)}$ remains an important and useful proxy to understand the Earth system.

Code and data availability. This paper relies exclusively on previously published data, available with the original papers and in publicly available repositories. An R notebook supplement is available alongside this manuscript, along with datafiles which allow full replication of all analyses performed.

Author contributions. MPSB conceived the study, designed the methodology, analysed the data, prepared the figures and wrote the manuscript (conceptualization, formal analysis, investigation, methodology, vizualization, writing - original draft, review and editing)

Competing interests. MPSB declares that he has no conflict of interest

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