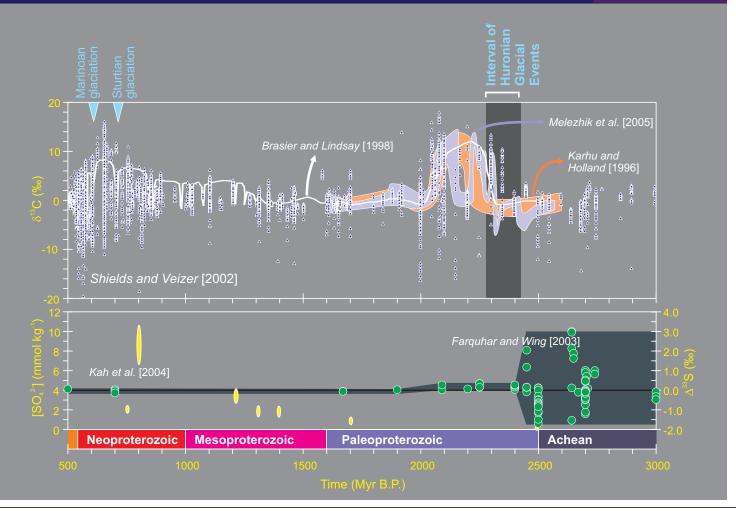
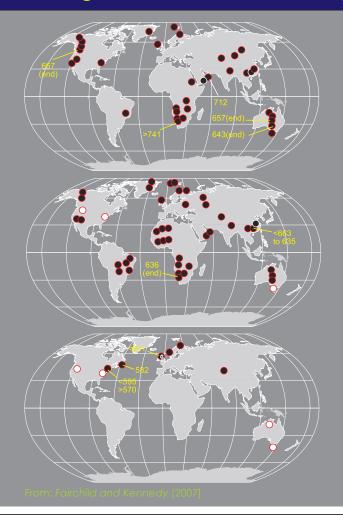
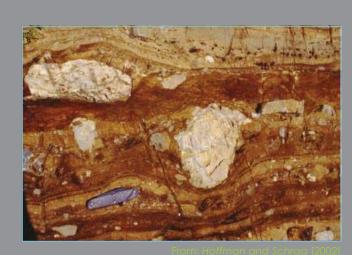


Snowball

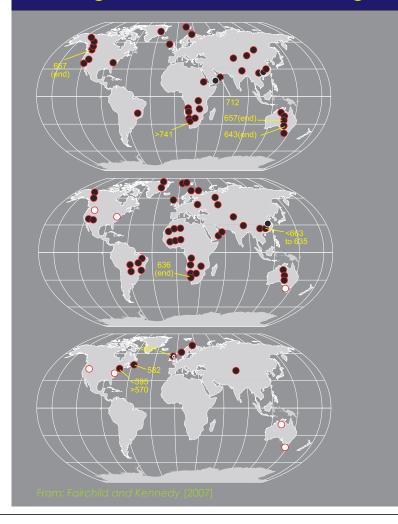


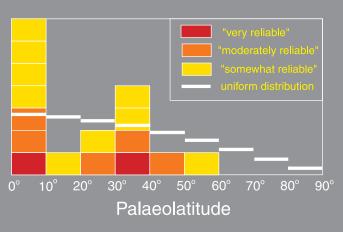
Background – evidence for glaciation



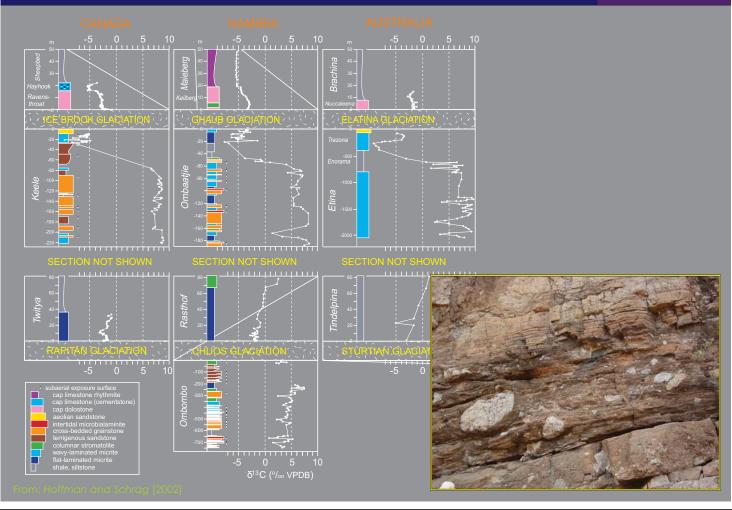


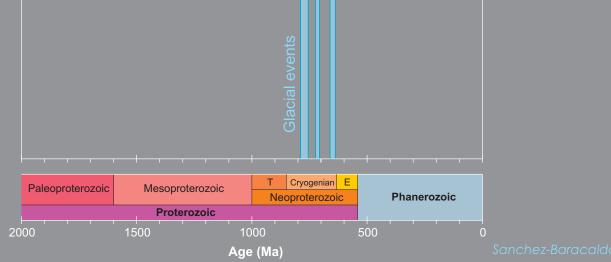
Background – evidence for glaciation



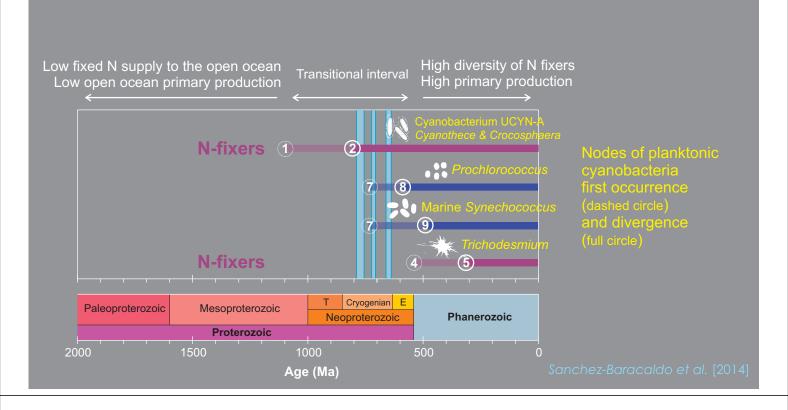


Background – evidence for glaciation

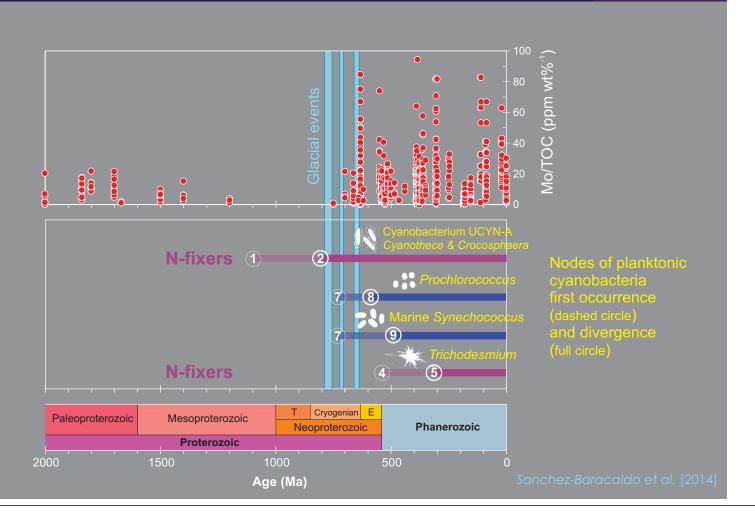




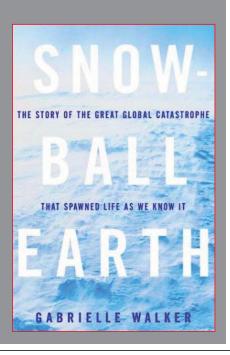
Background – biotic changes



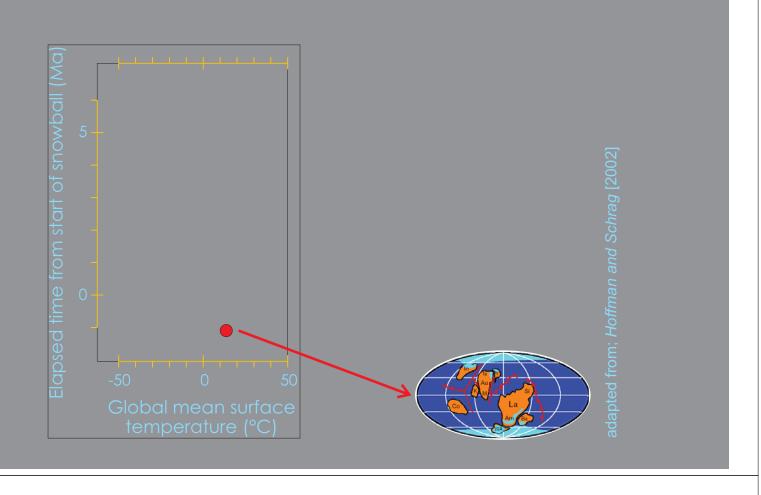
Background – biotic changes

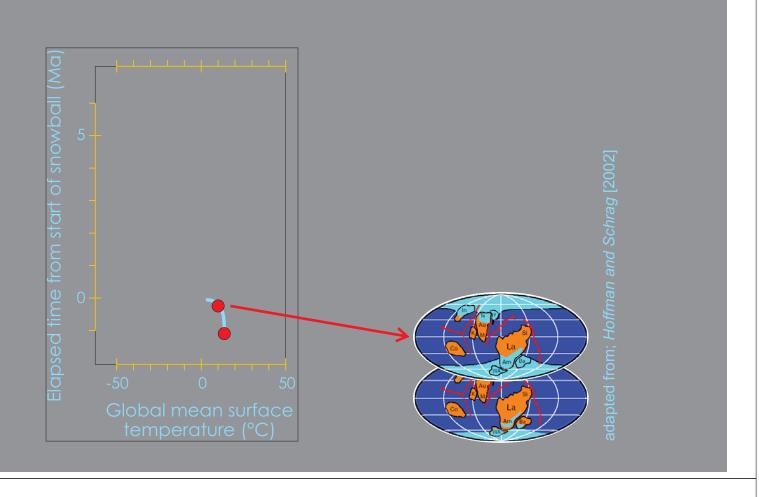


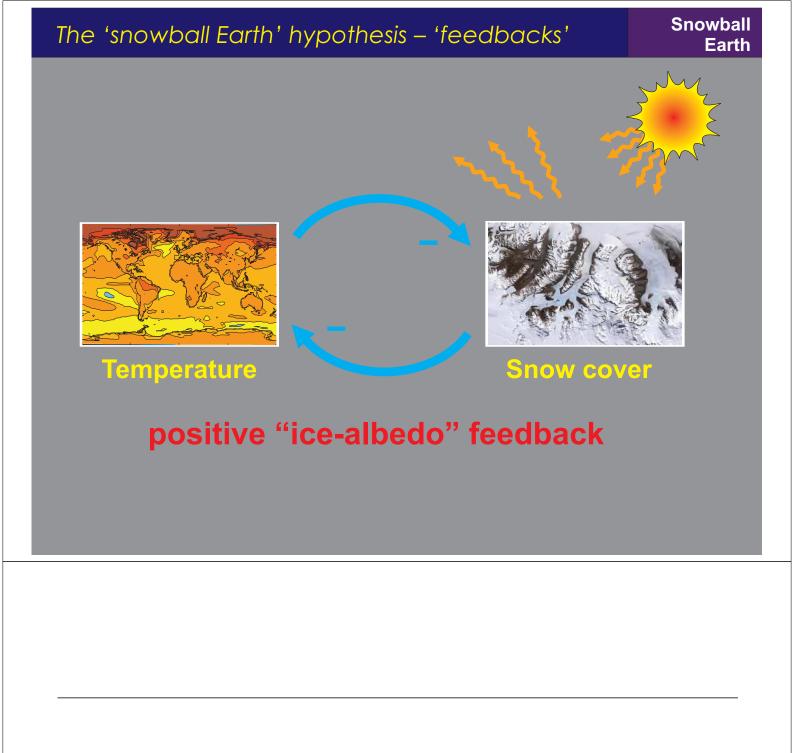
Hoffman et al. [1998] (Science 281

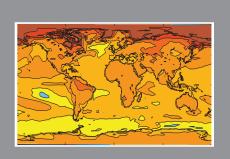


_			
_			







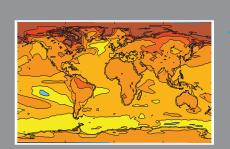


Temperature



Snow cover

TOTAL CHANGE = -1/2°C - 1/4°C

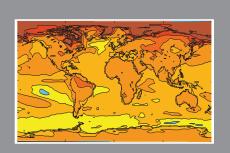


Temperature

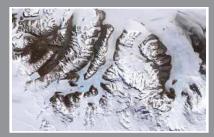


Snow cover

TOTAL CHANGE = -1/2°C - 1/4°C

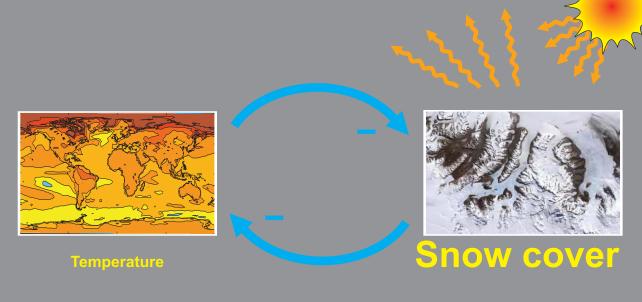




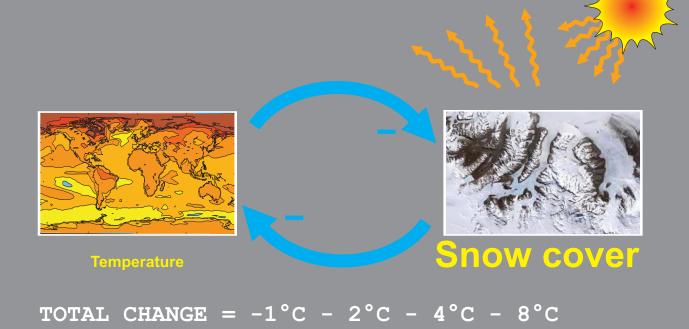


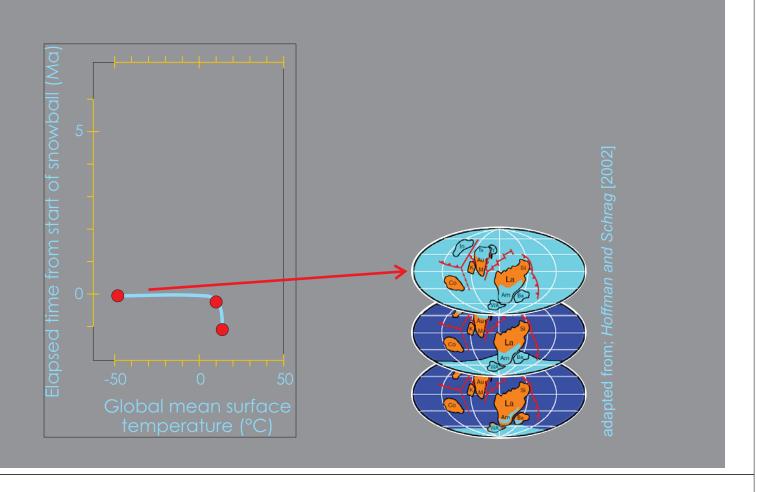
Snow cover

TOTAL CHANGE = -1/2°C - 1/4°C - 1/8°C

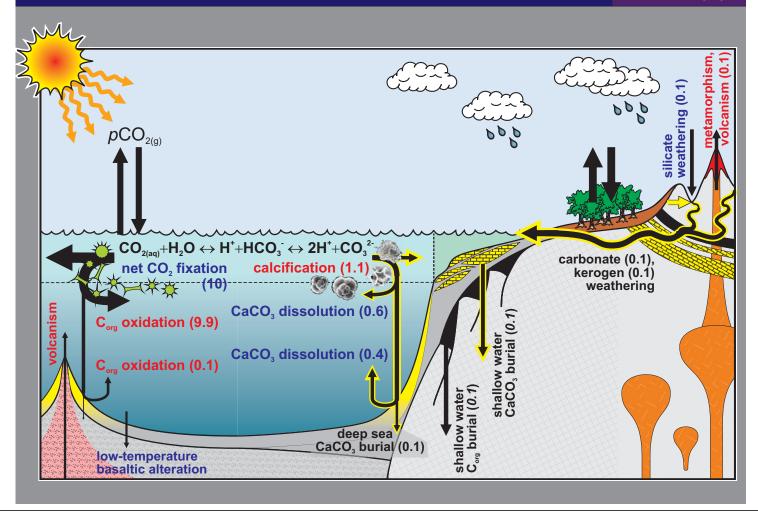


TOTAL CHANGE = -1/2°C - 1/4°C - 1/8°C - 1/16° -





Global carbon cycling (modern)



Global carbon cycling (long-term controls)

Terrestrial weathering can be (approximately equally) divided into carbonate (CaCO₃) and calcium-silicate ('CaSiO₃') weathering:

(1)
$$2CO_{2(aq)} + H_2O + CaSiO_3 \rightarrow Ca^{2+} + 2HCO_3^{-} + SiO_2^{-}$$

(2)
$$CO_{2(aq)} + H_2O + CaCO_3 \rightarrow Ca^{2+} + 2HCO_3$$

Ultimately, the (alkalinity: Ca²⁺) weathering products must be removed through carbonate precipitation and burial in marine sediments:

(3)
$$Ca^{2+} + 2HCO_3^{-} \rightarrow CO_{2(ac)} + H_2O + CaCO_3$$

It can be seen that in (2) + (3), that the CO_2 removed (from the atmosphere) during weathering, is returned upon carbonate precipitation (and burial). In (1) + (3) (silicate weathering) CO_2 is permanently removed to the geological reservoir. This CO_2 must be balanced by mantle (/volcanic) out-gassing on the very long term.

Global carbon cycling (long-term controls)

Terrestrial weathering can be (approximately equally) divided into carbonate (CaCO₃) and calcium-silicate ('CaSiO₃') weathering:

(1)
$$2CO_{2(aq)} + H_2O + CaSiO_3 \rightarrow Ca^{2+} + 2HCO_3^{-} + SiO_2$$

(2)
$$CO_{2(aq)} + H_2O + CaCO_3 \rightarrow Ca^{2+} + 2HCO_3^{-1}$$

Ultimately, the (alkalinity: Ca²⁺) weathering products must be removed through carbonate precipitation and burial in marine sediments:

(3)
$$Ca^{2+} + 2HCO_3^{-} \rightarrow CO_{2(aq)} + H_2O + CaCO_3$$

It can be seen that in (2) + (3), that the CO_2 removed (from the atmosphere) during weathering, is returned upon carbonate precipitation (and burial). In (1) + (3) (silicate weathering) CO_2 is permanently removed to the geological reservoir. This CO_2 must be balanced by mantle (/volcanic) out-gassing on the very long term.

Global carbon cycling (long-term controls)

Terrestrial weathering can be (approximately equally) divided into carbonate (CaCO₃) and calcium-silicate ('CaSiO₃') weathering:

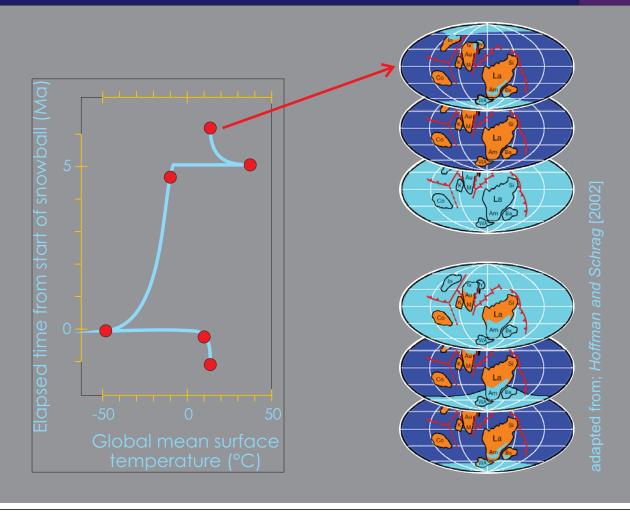
(1)
$$2CO_{2(aq)} + H_2O + CaSiO_3 \rightarrow Ca^{2+} + 2HCO_3^{-} + SiO_2^{-}$$

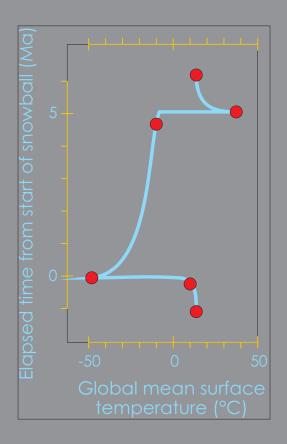
(2)
$$CO_{2(an)} + H_2O + CaCO_3 \rightarrow Ca^{2+} + 2HCO_3$$

Ultimately, the (alkalinity: Ca²⁺) weathering products must be removed through carbonate precipitation and burial in marine sediments:

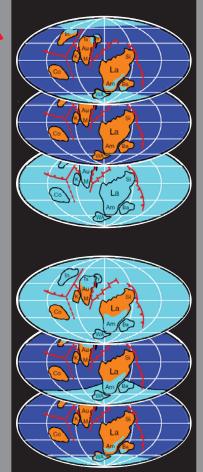
(3)
$$Ca^{2+} + 2HCO_3^- \rightarrow CO_{2(aq)} + H_2O + CaCO_3$$

It can be seen that in (2) + (3), that the CO_2 removed (from the atmosphere) during weathering, is returned upon carbonate precipitation (and burial). In (1) + (3) (silicate weathering) CO_2 is permanently removed to the geological reservoir. This CO_2 must be balanced by mantle (/volcanic) out-gassing on the very long term.



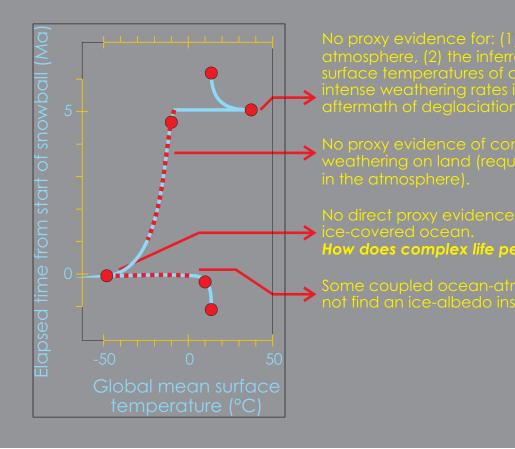


Progression of exotic states of the carbon-climate system

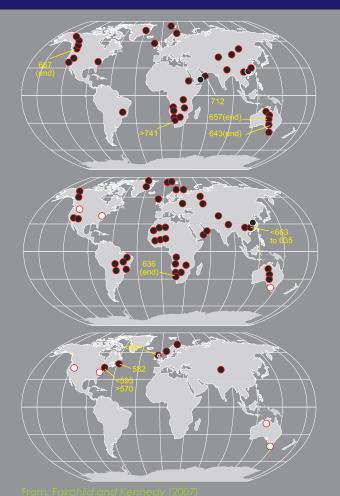


dapted from; Hoffman and Schrag [2002]

All very well, but ...



The 'snowball Earth' hypothesis



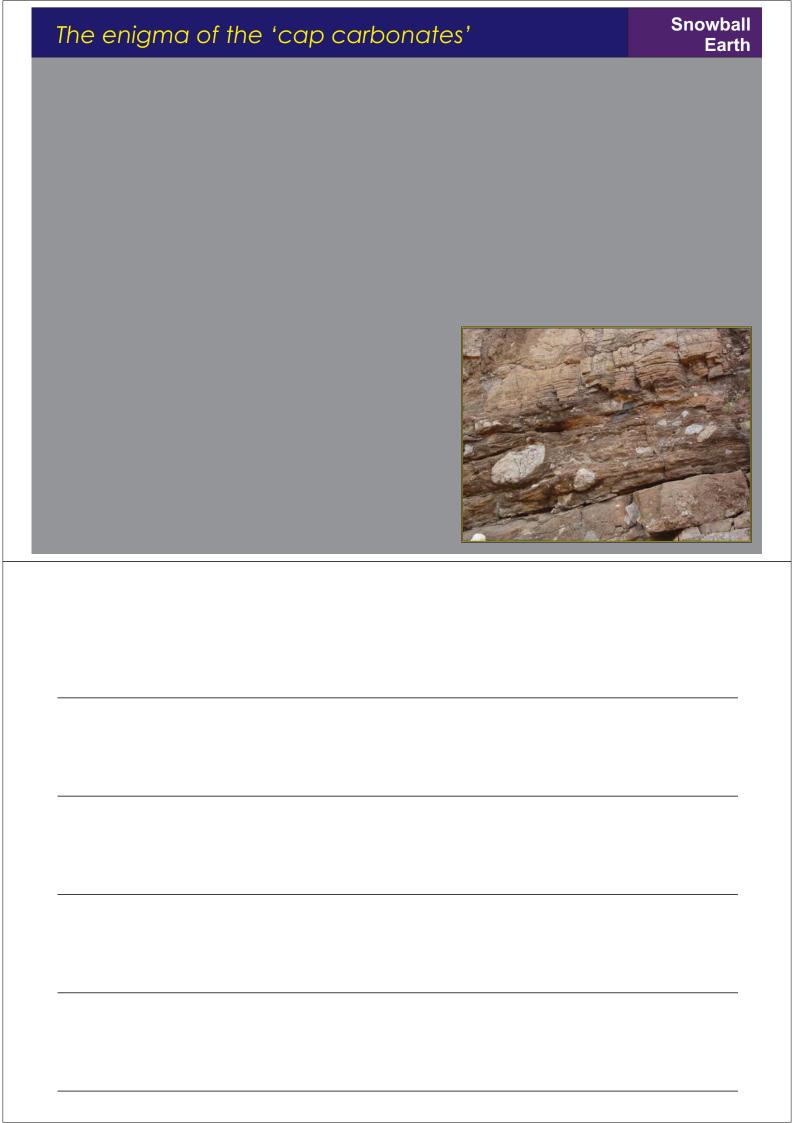
Q 'snowball' or 'slushball'?

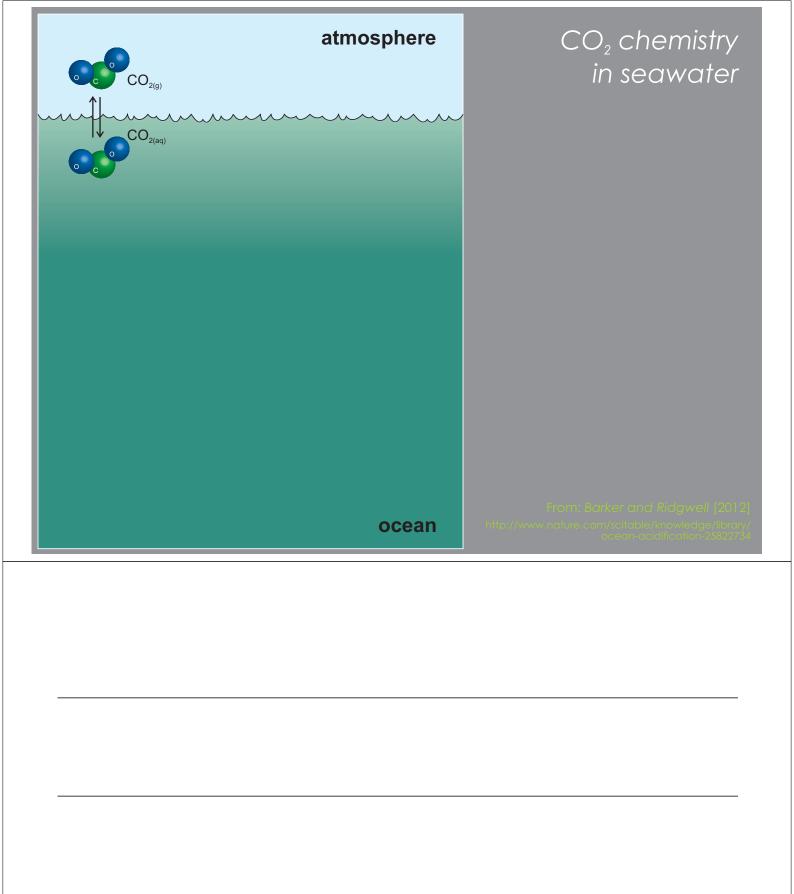
i.e., was the equatorial ocean ice-free, or frozen from pole-to-pole during the glacial? This has profound geochemical and climatic implications, and will constrain the mechanims responsible for going into and coming out of the glaciation.

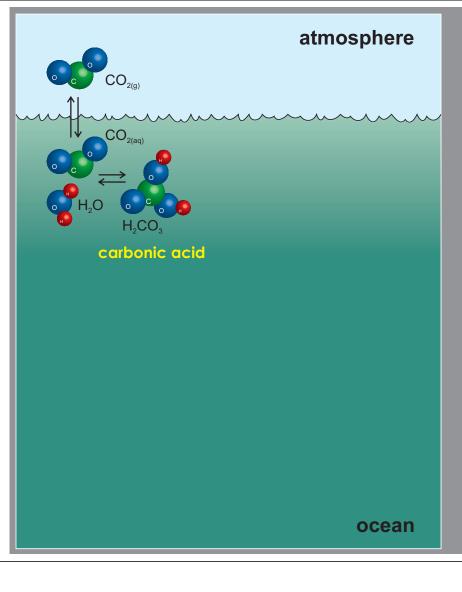
The 'snowball Earth' hypothesis

All (numerical climate) models are wrong. Some may be useful.

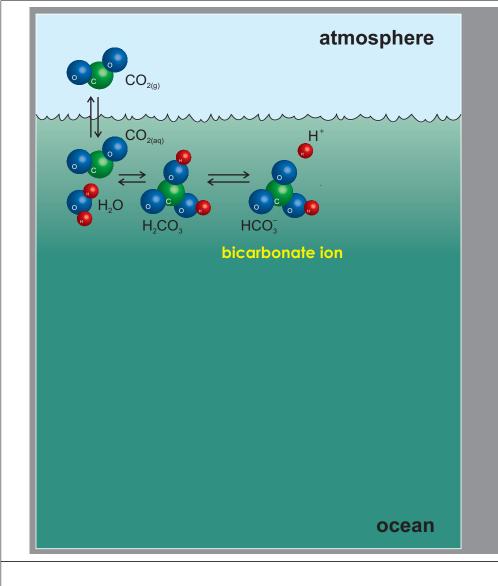
Study	ATM	SEA- ICE	OCN	ICE- SHT	pCO ₂ threshold	(conclusions)
Jenkins and Smith [1999]					1700 ppm	(snowball)
Chandler and Sohl [2000]					<40 ppm	snowball unlikely
Hyde et al. [2000]; Crowley et al. [2001]					130 ppm	slushball probable
Baum and Crowley [2001,2003]					<340 ppm	slushball probable
Poulsen et al. [2001,2]; Poulsen [2003]					n/a	no snowball
Bendtsen [2002]					n/a	snowball less likely
Godderis et al. [2003]					130 ppm	(snowball)
Goodman and Pierrehumbert [2003]					130 ppm	snowball more likely
Donnadieu et al. [2003]					500 - 990	slushball unlikely
Lewis et al. [2003,2004]					1800 ppm	(snowball)
Donnadieu et al. [2004a,b]					<149, 250	(snowball)
Edwards and Ridgwell [unpublished]					200 ppm	(snowball)
KEY: 'ADVANCED' e.g. 3D GCM, thermodynamic sea-ice	e.g. 2 seas	ERMEDI/ 2D EBM, onal mixe ocean			BASIC' .g. 1D EBM, lab ocean	DECOUPLED



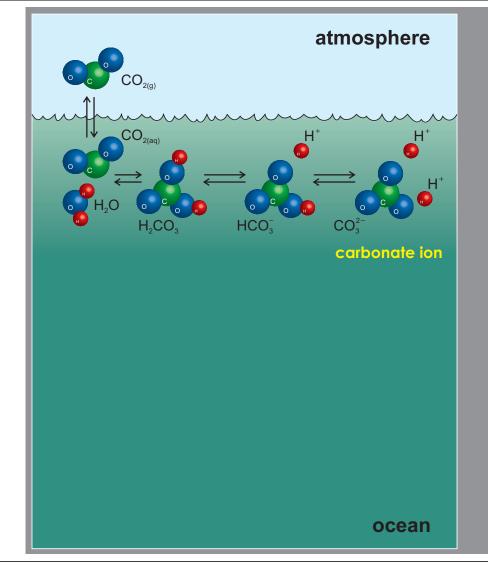




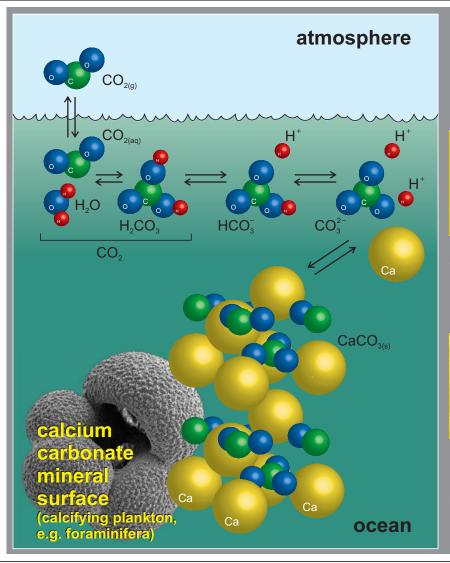
CO₂ chemistry in seawater



CO₂ chemistry in seawater



CO₂ chemistry in seawater



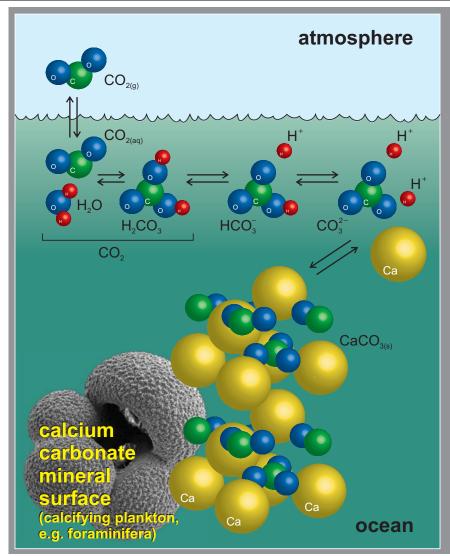
CO₂ chemistry & mineral phases



Aragonite: less stable orthorhombic polymorph (e.g., many corals, pteropods)



Calcite: more stable
(and more abundant)
trigonal polymorph (e.g.,
coccolithophorides, foraminifera)

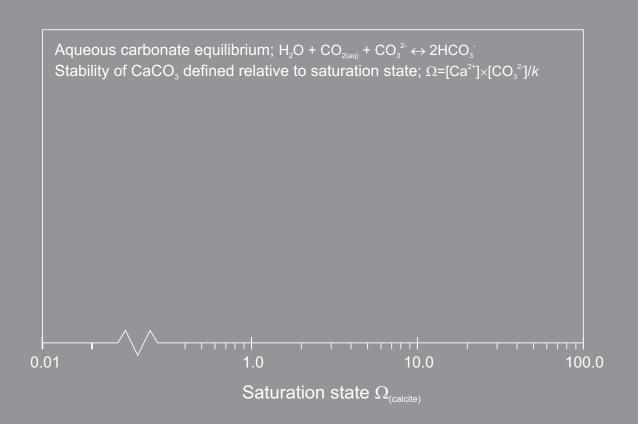


CO₂ chemistry & mineral phases

The addition of CO₂ to seawater results in a decrease in carbonate ion (CO₃²) concentration and ocean acidification. A decrease in CO₃², in turn, suppresses the stability of CaCO₃, defined by its saturation state:

$\Omega = [Ca^{2+}] \times [CO_3^{2-}]/k$

⇒ The thermodynamic efficiency of precipitating CaCO₃ is a function of [CO₃²] (and carbonate 'saturation').



See: GEOG 'World in crisis' lecture on 'Ocean Acidification' (PDF available from www.seao2.org/teaching.html)

