

## Introduction

The ultimate fate of fossil fuel CO<sub>2</sub> emitted to the atmosphere is governed by a range of sedimentological and geological processes operating on timescales of up to hundreds of thousands of years.

However, the response of these carbon cycle processes to increasing total emissions is uncertain, such as whether some sinks might saturate or feedbacks weaken.

The lifetime of an atmospheric CO<sub>2</sub> perturbation will have important implications for components of the Earth system that are temperature-sensitive and have slow response times, such as ice sheets.

## Objectives

- Investigate the atmospheric lifetime of a range of future CO<sub>2</sub> emissions.
- Perform a multi-exponential analysis to identify some of the carbon cycle processes acting to remove excess CO<sub>2</sub> from the atmosphere.
- Derive an empirical response function that can project the response of atmospheric pCO<sub>2</sub> to any arbitrary CO<sub>2</sub> emission.

## Methodology

- Used cGENIE (Fig. 1), a coupled climate-carbon cycle Earth system model<sup>[1]</sup>.

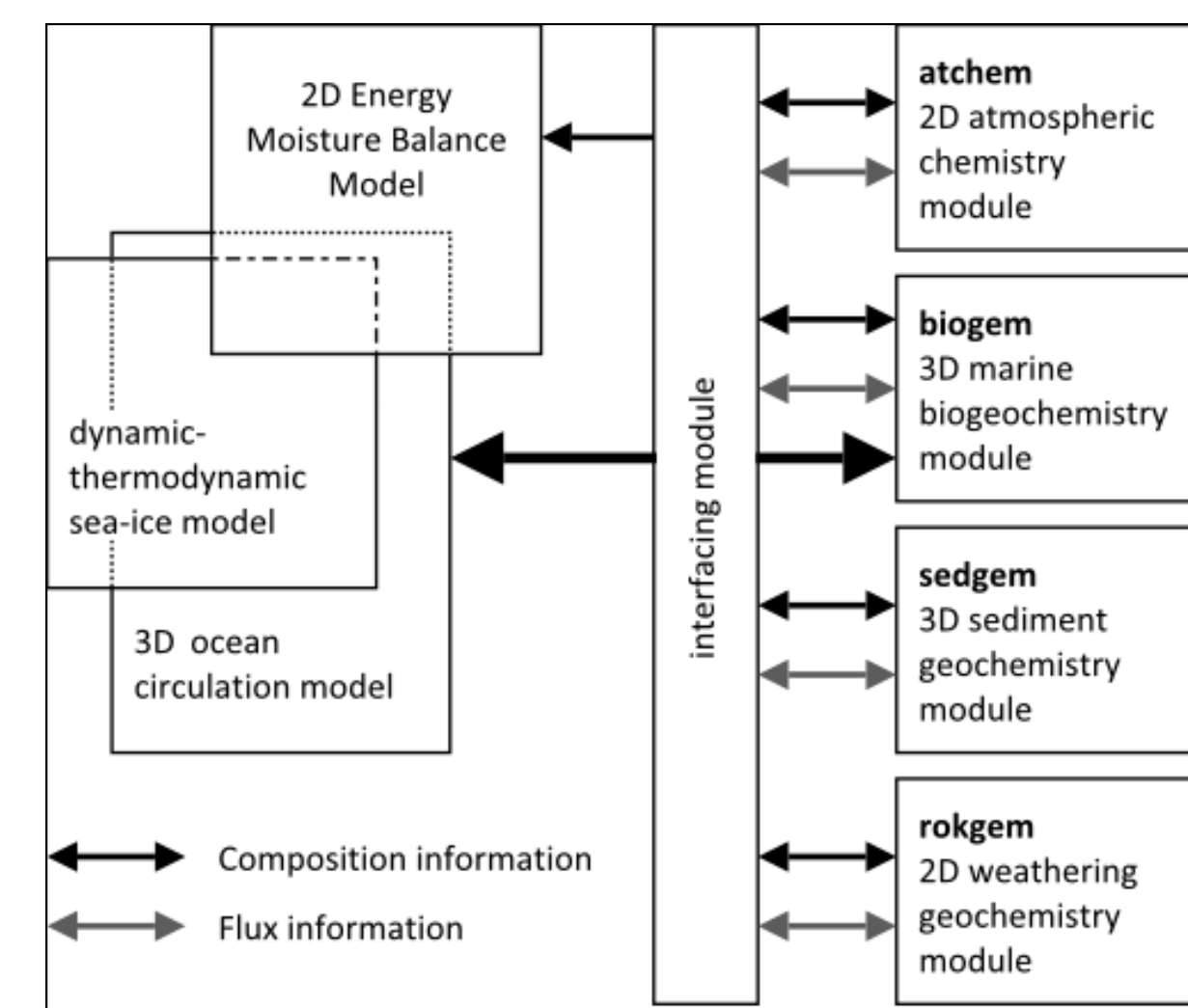


Fig. 1. Schematic of the relationship between the different components of the cGENIE model. Arrows represent coupling.

- Series of 1 Myr experiments (20) with pulse CO<sub>2</sub> emissions ranging from 1,000-20,000 PgC ("Pulse series"; Table 1).
- Series of 1 Myr experiments with time-dependent CO<sub>2</sub> emissions ranging from 1,500-6,300 PgC ("Time-dependent series"; Table 1).
  - Initiated from year 2010 following historical CO<sub>2</sub> emissions<sup>[2]</sup> (1750-2009).

Experiment ensemble	Total emissions (PgC)	Release period (yr)	Release year(s)
Pulse series	1,000 - 20,000	1	1
Time-dependent series	1,500 - 6,300	290	2010-2300

Table 1. Summary of model experiments.

## The Long-Term Carbon Cycle

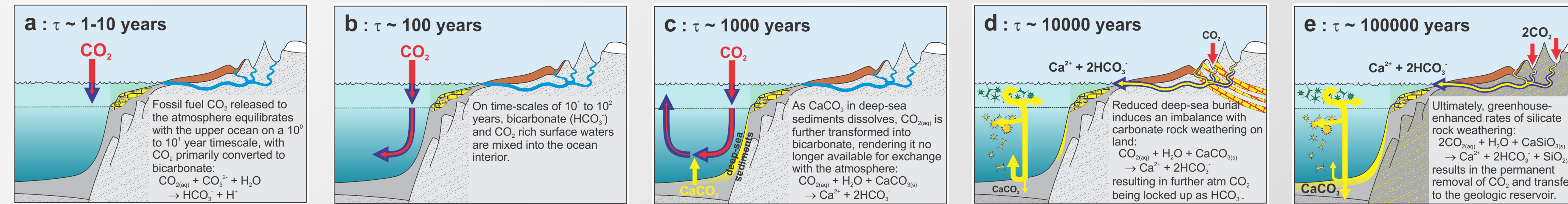


Fig. 2. Illustration of the primary mechanisms of CO<sub>2</sub> sequestration from the atmosphere and their timescales of operation: (a) air-sea gas exchange, (b) ocean invasion, (c) sea-floor CaCO<sub>3</sub> neutralization, (d) terrestrial CaCO<sub>3</sub> neutralization, and (e) the silicate weathering carbon sink.

## Analysis + Results

### Multi-exponential analysis

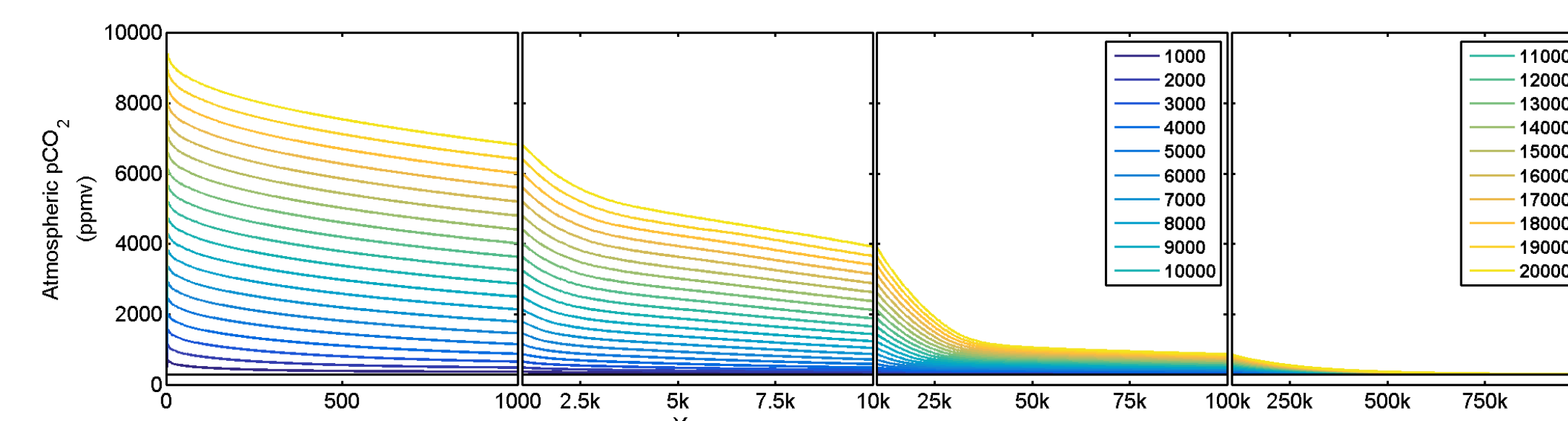


Fig. 3. Atmospheric pCO<sub>2</sub> for the pulse release scenarios predicted by cGENIE. Pre-industrial concentrations are shown in black.

- Pulse series (Fig. 3):
  - Each CO<sub>2</sub> decay curve was fitted with a series of exponentials (4 optimal):  $CO_2(t) = B + E \sum_{i=1}^n A_i \exp^{-(t-t_0)/\tau_i}$
  - Each exponential represents one or more of the carbon cycle processes described in Fig. 2.
  - For each exponential, the fraction of CO<sub>2</sub> (A<sub>i</sub>) and the timescale of decay (τ<sub>i</sub>) was extracted, as shown in Fig. 4.

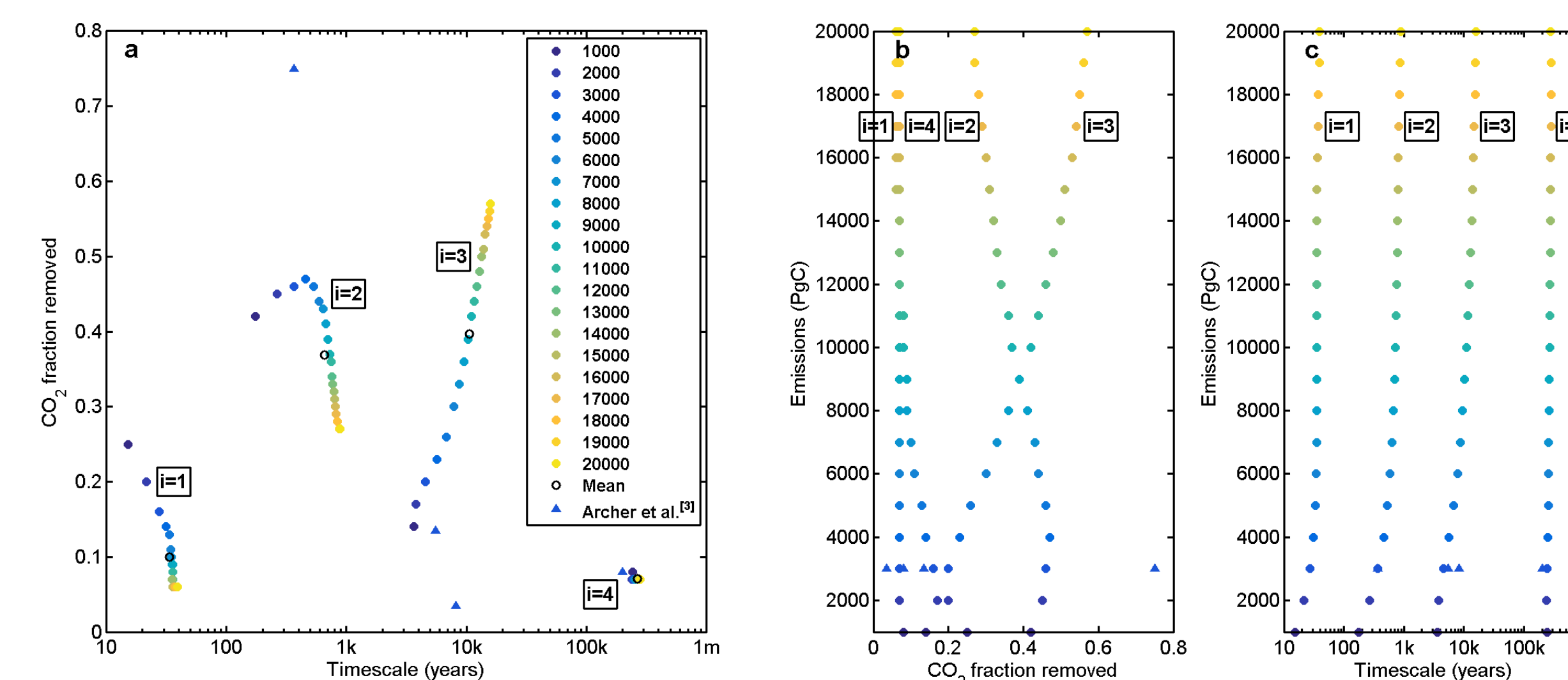


Fig. 4. Change in fitting coefficient values (A<sub>i</sub> and τ<sub>i</sub>) with total CO<sub>2</sub> emissions in PgC (shown as different coloured dots; see legend in panel (a)). (a) CO<sub>2</sub> fraction removed vs timescale, (b) CO<sub>2</sub> fraction removed vs total emissions, and (c) timescale vs total emissions.

### Empirical response function

- The fitting coefficients (A<sub>i</sub> and τ<sub>i</sub>) were made quadratic functions of the total emissions.
- The predictive capability of the response function can be evaluated against the atmospheric pCO<sub>2</sub> predicted by cGENIE for the time-dependent series (Fig. 5a).

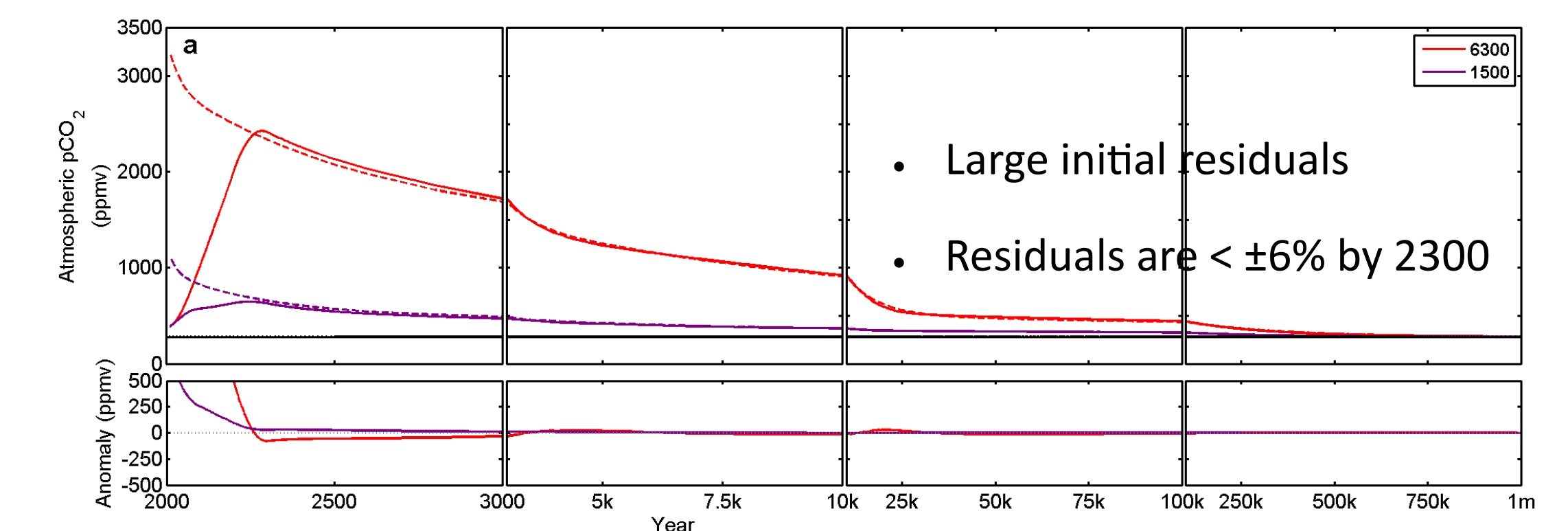


Fig. 5a. Top panel: atmospheric pCO<sub>2</sub> for the time-dependent release scenarios predicted by cGENIE (solid line) and the response function (dashed line). Pre-industrial concentrations are shown in black. Bottom panel: pCO<sub>2</sub> anomaly.

- A convolution analysis approach was then developed to improve the predictive performance of the response function for time-dependent emissions (Fig. 5b).

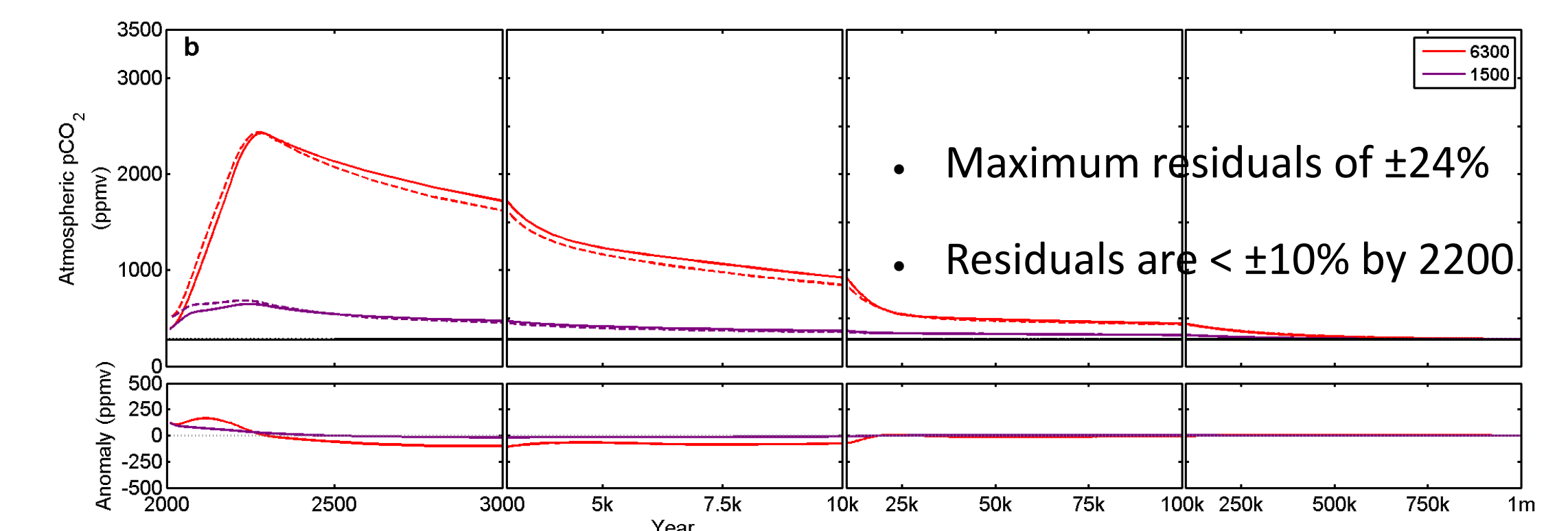


Fig. 5b. Top panel: atmospheric pCO<sub>2</sub> for the time-dependent release scenarios predicted by cGENIE (solid line) and the convoluted response function (dashed line). Pre-industrial concentrations are shown in black. Bottom panel: pCO<sub>2</sub> anomaly.

## Conclusions

- Atmospheric CO<sub>2</sub> perturbations, and hence the resultant climate changes, can last for tens to hundreds of thousands of years.
- As total emissions increase, the response times of all carbon sinks (apart from silicate weathering) lengthen.
- The two processes with the shortest timescales weaken, while the ~10,000 year process of seafloor and terrestrial CaCO<sub>3</sub> neutralization strengthens.
- A number of thresholds may exist, resulting from feedbacks associated with elevated atmospheric pCO<sub>2</sub> levels.
- The empirical model provides a simple and practical tool for rapidly projecting the long-term fate of CO<sub>2</sub> emissions, and can be used in place of a mechanistic model.

For further information please see: Lord et al. (in review) *Global Biogeochemical Cycles*.

## References

- Ridgwell, A. et al. (2007) *Biogeosciences*, 4, pp.87-104.
- Meinshausen, M. et al. (2011) *Climatic change*, 109, pp.213-241.
- Archer, D. et al. (1997) *Geophysical Research Letters*, 24(4), pp.405-408.

## Acknowledgements

This research is funded by RWM via a framework contract with AMEC, who are being supported by Quintessa. It contributes to the MODARIA international research program, coordinated by the International Atomic Energy Agency (IAEA).