Biogeochemical Controls on the Redox Evolution of Earth’s Oceans and Atmosphere

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The redox state of Earth’s atmosphere has undergone a dramatic shift over geologic time from reducing to strongly oxidizing, and this shift has been coupled with changes in ocean redox structure and the size and activity of Earth’s biosphere. Delineating this evolutionary trajectory remains a major problem in Earth system science. Significant insights have emerged through the application of redox-sensitive geochemical systems. Existing and emerging biogeochemical modeling tools are pushing the limits of the quantitative constraints on ocean–atmosphere redox that can be extracted from geochemical tracers. This work is honing our understanding of the central role of Earth’s biosphere in shaping the long-term redox evolution of the ocean–atmosphere system.

Keywords: biogeochemistry, oxygenation, biosphere, redox, evolution

REDOX ON A PLANETARY SCALE

Earth’s Modern Oxygen Cycle

On the modern Earth, molecular oxygen (O₂) is produced through photosynthesis in sunlit surface environments in which cyanobacteria, algae, and plants use energy from the Sun to transfer electrons from water to biomass. Much of the biomass produced by the photosynthetic biosphere is consumed rapidly through aerobic respiration, which gains energy by transferring electrons from biomass back to O₂. These coupled processes can be denoted as:

\[ \text{CO}_2 + \text{H}_2\text{O} \leftrightarrow \text{CH}_2\text{O} + \text{O}_2 \] (1)

Any photosynthetic biomass removed from the surface system via burial in marine or terrestrial sediments leads to a net accumulation of O₂ in the ocean–atmosphere system (moving from left to right in Equation 1). On geologic timescales, this organic matter can be consumed through exhumation and oxidation in terrestrial weathering environments and through thermal breakdown of organic carbon during metamorphism. Both of these processes lead, ultimately, to a net consumption of O₂ from the ocean–atmosphere system (moving from right to left in Equation 1). Thus, Equation (1) can be thought of conceptually as showing a “fast” O₂ cycle (photosynthesis/respiration), responding on timescales of less than ~10³ years, and a “slow” O₂ cycle (organic carbon burial and weathering/thermal breakdown), responding on timescales of ~10⁶ years.

There is an additional long-term cycle that links O₂ to the cycling of sulfur (S) and iron (Fe) at Earth’s surface:

\[ 2\text{Fe}_2\text{O}_3 + 8\text{SO}_4^{2-} + 16\text{H}^+ \leftrightarrow 15\text{O}_2 + 4\text{FeS}_2 + 8\text{H}_2\text{O} \] (2)

Moving from left to right in Equation (2) denotes the sum of three processes: (1) photosynthesis; (2) consumption of biomass through microbial sulfate reduction, producing reduced sulfur (S²⁻); (3) formation of pyrite (FeS₂) via reaction of reduced sulfur and reactive iron in the environment (shown here as hematite, Fe₂O₃). In essence, the overall effect of this process is to transfer reducing power from water to pyrite via organic biomass, causing a net release of O₂. The subsequent uplift/weathering and/or volatilization of reduced sulfur species in the crust balances this release of O₂ on geologic timescales (moving from right to left in Equation 2).

Oxygen can also be consumed via reaction with reduced volcanic volatiles that originate from Earth’s crust and mantle and during the oxidation of reduced minerals in seafloor basalt as oxygen-rich seawater percolates through the oceanic lithosphere. In addition, the escape of hydrogen to space allows oxygen atoms produced by photochemistry to recombine as O₂, irreversibly oxidizing Earth surface environments. Similarly, oxidized or reduced species can be subducted into Earth’s mantle, leading to secular net oxidation/reduction of Earth’s surface environments that is not necessarily irreversible but that can persist on timescales of mantle overturn (that is, on the order of ~10⁹ years). For example, organic carbon deposited in deep-sea sediments can potentially be preserved as graphite during subduction, leading to a net export of reducing power from Earth’s surface (Fig. 1).

Nutrients as a Driver of Earth’s Redox Balance

The global oxygen cycle is fundamentally a story of the redox cycling of carbon, sulfur, and iron through the Earth system. However, O₂ cycling is also fundamentally controlled by the availability of bio-essential elements ("nutrients") in oceanic and terrestrial ecosystems. All organisms require a range of major and trace nutrients to produce biomass, which can be seen by elaborating on Equation (1) to include the major nutrients involved in oxygenic photosynthesis:
The stoichiometric coefficients in Equation (3) denote the bulk stoichiometry of photosynthetic biomass (the classical “Redfield ratio”, initially observed by US oceanographer Alfred Redfield in 1934), which can vary significantly as a function of taxonomic identity and organism growth status. A variety of trace elements—for example, Fe and Zn—are also incorporated into biomass, but for simplicity are neglected here. An important result is that every 138 moles of O$_2$ produced by the photosynthetic biosphere requires a mole of bioavailable phosphorus (P, shown here as phosphate) and 16 moles of bioavailable nitrogen (N, shown here as nitrate). These nutrient elements can then either be recycled by respiration back to inorganic, bioavailable forms (“ remineralized”) or buried in sediments. In the ocean, nutrients are generally taken up in sunlit surface waters and exported to depth with organic matter, leading to a vertical “biological pump” of organic carbon and nutrients into the ocean interior, with a corresponding demand for O$_2$.

There are strong stabilizing feedbacks that link the oceanic inventories of bioavailable nitrogen and phosphorus (Tyrrell 1999) and that draw together the long-term cycling of nutrients into the ocean interior, with a corresponding demand for O$_2$.

Processes that regulate the mean ocean nutrient budgets. (A) The Redfield line (in red) is the ratio of N to P in marine phytoplankton and in the deep oceans. A point on the line (red circle) is taken as an example of what happens under two scenarios: enhanced weathering/recycling of P and elevated burial/scavenging of P. Dashed and solid arrows denote adjustments following the perturbations to the ocean phosphate (PO$_4^{3-}$) budget from an initial state. (B) Long-term regulation of the mean ocean N and P budgets is driven by competition between N fixing ($\mu_f$) and nonfixing phytoplankton ($\mu_o$) (Tyrrell 1999). When P is in excess relative to N, the N fixers outcompete other phytoplankton (low $\mu_f/\mu_o$) and push the ocean N/P ratio back toward that required for the growth of nonfixing plankton (the Redfield ratio).

continued loss of nitrogen from the ocean interior through microbial denitrification and the sedimentary burial of organic nitrogen will draw the nitrogen reservoir down until the N/P set point is reached (Fig. 2). This simple thermostatic behavior can be modulated by the taxonomically diverse nutrient requirements of plankton and ocean circulation, but requires only that nitrogen fixation incur an energetic cost due to the difficulty of enzymatically breaking down N$_2$ (Tyrrell 1999). In this framework, the long-term fertility of the biosphere (and, thus, the capacity of Earth’s biosphere to produce O$_2$) is controlled most directly by the availability of P in Earth’s oceans. Earth’s surface O$_2$ cycle must, therefore, be considered in the context of the processes that regulate oceanic P cycling.

**The Importance of Ocean Redox Structure for Oxygen and Nutrient Cycling**

The impacts of atmospheric chemistry and planetary nutrient cycling on Earth’s surface O$_2$ are tightly linked through ocean redox structure. First, it is the abundance of O$_2$ in the atmosphere and the cycling of nutrients through the biological pump that structure the ocean...
redox landscape. At the same time, this redox landscape feeds back on marine nutrient cycling through a range of mechanisms that can, ultimately, act as either positive or negative feedbacks on long-term biospheric O2 production. Deep ocean O2 is governed by the balance between O2 supply through gas exchange at the ocean surface and through downward mixing of O2-rich waters, and by O2 demand through aerobic respiration once a water mass is no longer in communication with the atmosphere. Thus, under a constant ocean circulation regime, O2 supply is controlled principally by atmospheric O2 abundance, whereas O2 demand is controlled principally by the intensity of the biological carbon pump (and, hence, nutrient abundance). This relationship can be understood heuristically by considering a simple three-box ocean model of ocean and nutrient biogeochemistry (Fig. 3). In this model, we can derive a simple expression for the global average O2 concentration in the ocean interior at steady state ([O2]int):

\[
[O_2]_{int} = [O_2]_{vent} - \lambda([PO_4^{3-}]_{int} - [PO_4^{3-}]_{vent})
\]  

(4)

where [O2]vent and [PO4^{3-}]vent denote the oxygen and phosphate concentrations of cold, high-latitude waters ventilating the ocean interior, [PO4^{3-}]int denotes the phosphate abundance of the ocean interior, and \( \lambda \) denotes the stoichiometric relationship between the amount of phosphate released during consumption of O2 during aerobic respiration (see Equation 3). The term in parentheses describes the fraction of nutrients mixed into the high-latitude surface ocean that is consumed during photosynthesis (currently with a value of ~1 µmol kg\(^{-1}\)) and can be thought of as depicting the efficiency of the biological pump. Using this approach, one can estimate the steady state concentration of O2 in the deep ocean as a function of both atmospheric O2 abundance and biological pump efficiency (Fig. 3). This analysis shows that the ocean interior can become anoxic by either dropping atmospheric O2 abundance or by increasing the strength/efficiency of the biological pump. For example, with a modern biological pump this model predicts that the ocean interior would become pervasively anoxic at an atmospheric O2 abundance ~40% that of the modern Earth. Alternatively, with an atmospheric O2 abundance of 10% of the modern Earth even a relatively weak biological pump operating at ~20% of modern intensity would provide sufficient O2 demand in the ocean interior to drive the ocean toward extensive anoxia. Despite being incapable of capturing the impact of differences in ocean circulation, remineralization length scale of organic carbon in the water column, or spatial variability in deep O2, this model illustrates an important principle—it is the combination of atmospheric O2 abundance and the activity level of the ocean biosphere (as governed by ocean nutrient inventory) that fundamentally dictates the marine redox landscape.

There is, however, an important caveat to this analysis. It implicitly assumes that the abundance of O2 in the ocean–atmosphere system and the marine inventory of bioavailable phosphorus can be varied independently of one another. In reality, the throughput of bioavailable phosphorus fundamentally controls the ocean–atmosphere O2 abundance at steady state (Lenton and Watson 2000), while ocean redox state controls the recycling and bioavailability of phosphorus in marine systems (Van Cappellen and Ingall 1994; Reinhard et al. 2017). To gain intuition for this, one can consider the dynamics regulating the most significant O2 production and removal fluxes on the modern Earth—the burial and oxidation of organic carbon (Fig. 1). The efficiency of organic carbon burial in marine sediments, and, thus, the amount of O2 released by the biosphere at a given nutrient abundance, increases as O2 decreases. At the same time, all of the processes that consume O2, including oxidation of organic carbon in the crust, slow down as O2 drops. This combination provides strong negative feedback against decreasing atmospheric O2—as atmospheric O2 drops, the sources of O2 tend to increase while the sinks tend to decrease, pushing atmospheric O2 back up. Lowering atmospheric O2 beyond a certain level requires either a decrease in the inputs of bioavailable P to the marine system or an increase in the inorganic sinks of P from the oceans (e.g., less efficient use of bioavailable P by the marine biosphere).

At the same time, ocean redox plays an important role in controlling the recycling and bioavailability of phosphorus, which, in turn, impacts long-term biospheric O2 production. When the ocean interior is pervasively oxygenated, as on the modern Earth, bioavailable phosphorus is removed primarily through burial in marine sediments either as an authigenic mineral phase, or as a sorbed species on sedimentary Fe-oxides, or as a constituent of organic matter, with the remainder removed largely in association with Fe-oxide minerals in deep-sea hydrothermal systems (Ruttenberg 2014). In contrast, anoxic conditions can lead to a range of recycling processes that can dramatically impact phosphorus bioavailability. Anoxic and sulfidic...

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**Figure 3**  (A) Schematic depiction of a three-box model for oxygen and phosphorus distributions within the ocean. Oxygen is introduced through gas exchange at the ocean surface and then mixed throughout the ocean via overturning circulation and deep convection. Organic matter produced by photosynthesis sinks into the ocean interior (the ocean “biological pump”), and creates respiratory O2 demand. (B) Steady state dissolved O2 concentrations in the deep ocean as a function of both atmospheric O2 level and the strength of the biological pump relative to that of the modern ocean. Contours are labeled according to dissolved O2 concentration; the grey shaded region denotes conditions under which the deep ocean is expected to become anoxic. PAL = present atmospheric level.
conditions, either in the water column (" euxinia") or in surface marine sediments, result in remobilization of Fe-oxide-bound phosphorus and can change the style of organic phosphorus recycling, both of which can potentially lead to more effective recycling of bioavailable phosphorus under sulfidic anoxia (Van Cappellen and Ingall 1994). In contrast, anoxic but iron-rich (" ferruginous") conditions can lead to effective scavenging and removal of bioavailable phosphorus through two principal mechanisms: the formation of Fe-phosphate mineral phases and coreplication of P with other Fe-bearing minerals (see Reinhard et al. 2017 and references therein).

**THE EVOLUTION OF OCEAN–ATMOSPHERE REDOX ON EARTH**

Earth’s modern atmosphere is strongly oxidizing, being composed of ~21% O2 by volume, with reducing gases generally present at trace abundance. Indeed, this abundance of O2 (and its photochemical by-product, ozone, O3) gives rise to a series of compelling "biosignatures" in the atmosphere that are, in principle, detectable from remote distances through atmospheric spectroscopy (Meadows et al. 2018). An important consequence of this high O2 level in the atmosphere is that the oceans are generally very well-oxygenated, despite a large and robust biosphere and attendant respiratory demand in the ocean’s interior. However, atmospheric O2 has changed by many orders of magnitude throughout Earth’s history, in concert with dramatic shifts in biogeochemical cycling, tectonic events, perturbations throughout Earth’s history, in concert with dramatic shifts in biogeochemical cycling, tectonic events, perturbations to global climate, and biological innovations, all of which have impacted ocean redox structure.

Prior to the evolution of oxygenic photosynthesis, the abundance of O2 in Earth’s atmosphere would have been extremely low, with production largely through CO2 photolysis, yielding ground-level O2 abundances of ~10^{−12} times the present atmospheric level (PAL) (Kasting et al. 1979). This means that the redox state of the atmosphere would once have been dominated by some combination of H2 and CH4; O2 would have been virtually absent from oceanic environments. Following the emergence of oxygenic photosynthesis, the potential for locally nontrivial dissolved O2 abundances in the shallowest marine systems would have emerged (so-called "oxygen oases"). This conceptual framework finds support from both theoretical models (Olson et al. 2013) and geochemical observations (e.g., Planavsky et al. 2014). Nevertheless, the preservation of significant sulfur mass-independent fractionation (S-MIF) isotope signals in marine sediments deposited prior to ~2.3 Ga indicates a pervasively anoxic atmosphere (Farquhar et al. 2000), likely on the order of ~10^{−7} PAL (Zahnle et al. 2006). Such low atmospheric O2 abundances are consistent with a wide range of independent geologic evidence (Holland 1984). Atmospheric O2 would be regulated in this Earth system state via consumption by reduced volcanic gases and reaction with reduced Fe and S in the Earth’s crust, with localized regions of elevated O2 in shallow seas (Fig. 4). An important wrinkle to this framework comes in the form of transient "whiffs" of atmospheric O2 during the Neorarchean, as deduced from trace element, isotopic, and mineralogical records (see Lyons et al. 2014).

Although it seems clear that O2 was present at trace abundance for most, if not all, of Hadean and Archean time (at levels of ~10^{-12} – 10^{-10} PAL), the abundances of the most important reducing gases, H2 and CH4, are not well-constrained. Photochemical models predict that a nontrivial abundance of reducing atmospheric gas (either H2 or CH4) must be present in order to preserve distinct exit channels for photolytic sulfur from the atmosphere and, thus, support the preservation of S-MIF signals. But the precise quantitative requirements are unclear. More recently, it has been suggested that the apparent mass fractionation of xenon isotopes in the Archean atmosphere requires a total hydrogen mixing ratio in the atmosphere above ~1% (corresponding to an atmospheric CH4 abundance of ~0.5%) in order to support the requisite atmospheric escape rates (Fahnle et al. 2019). This is broadly consistent with photochemical and coupled biosphere–atmosphere models for the Archean Earth system (Kharecha et al. 2005; Ozaki et al. 2018).

A variety of geochemical and geologic observations, including the disappearance of S-MIF signals from the rock record, indicate that the initial accumulation of O2 in Earth’s atmosphere occurred at ~2.4–2.3 Ga. This is often referred to as the Great Oxidation Event (GOE). How “great” this event was, however, remains an open question. The disappearance of S-MIF signals from the sedimentary record only requires a relatively small change in the overall size of the atmospheric O2 reservoir. For example, a shift in atmospheric pO2 from 10^{-7} to 10^{-4} PAL, more than sufficient to shut down S-MIF signals, would require the net accumulation of only ~0.01% of the modern atmospheric O2 inventory. Nevertheless, it is manifestly evident in a range of sedimentological and geochemical records that this geologically brief interval marks an important shift in the background redox state of Earth’s atmosphere (Holland 1984).

More recently, it has been suggested that there was a protracted—but ultimately transient—period of elevated atmospheric O2 abundance between ~2.2 Ga and 2.0 Ga (Fig. 4). This period coincides with a striking perturbation to Earth’s carbon cycle, evident in an extended period of 13C-enriched carbonates deposited worldwide—the so-called “Lomagundi Event” (Schidlowski et al. 1976). Early estimates suggested that this carbon cycle perturbation represented a long-term pulse of organic matter. But the precise quantitative requirements are unclear. More recently, it has been suggested that the apparent mass fractionation of xenon isotopes in the Archean atmosphere requires a total hydrogen mixing ratio in the atmosphere above ~1% (corresponding to an atmospheric CH4 abundance of ~0.5%) in order to support the requisite atmospheric escape rates (Fahnle et al. 2019). This is broadly consistent with photochemical and coupled biosphere–atmosphere models for the Archean Earth system (Kharecha et al. 2005; Ozaki et al. 2018).

**A FIGURE 4** Evolving redox state of Earth’s atmosphere in terms of the two major redox-active species in Earth’s atmosphere: 1) partial pressures of oxygen (pO2) relative to the atmospheric level (PAL); 2) partial pressures of methane (pCH4) in parts per million by volume (ppmv). Shaded boxes (Archean in orange; Proterozoic in green; Phanerozoic in grey) show ranges derived from the geologic record and theoretical models, while the black line on each of the two graphs shows one plausible trajectory through these models. Upward arrows denote possible Mesozoic to Neorarchean “whiffs” of atmospheric O2, though their timing, duration, and magnitude are not fully known. A putative transient high-O2 interval is also shown with the dashed shaded box (in the Paleoproterozoic), though the atmospheric composition during this period is not well-constrained. Values for the modern atmosphere are shown as grey lines. GOE = Great Oxidation Event.
O₂ was significantly more abundant in the atmosphere than today (Planavsky et al. 2016). In any case, the dependent variability in atmospheric redox during the ~2.5 Ga interval can be inferred from pervasively anoxic oceans. In any case, it seems reasonable that between ~2.2 Ga and 2.0 Ga the ocean redox landscape would have featured a pervasively well-oxygenated mixed layer with dissolved O₂ concentrations controlled largely by gas exchange with the atmosphere, and possibly widespread oxygenation of the ocean interior. Fully elucidating these features of the Earth system during this period awaits more precise quantitative constraints on atmospheric O₂ abundance (Fig. 4).

Though it is apparent that atmospheric O₂ increased after ~2.5 Ga, atmospheric abundances of O₂ over the ensuing ~2 Gy are uncertain. A lack of significant stable chromium (Cr) isotope fractionation in some ironstones, marine shales, and ancient soil horizons (paleosols) has been interpreted to reflect very low atmospheric pO₂, perhaps as low as 10⁻³ PAL, for much of the period between ~1.8 Ga and 0.8 Ga. This is consistent with the observed mobilization behavior of iron (Fe) and manganese (Mn) in paleosols, records of rare earth element (REE) behavior in marine carbonates, and the stable oxygen isotope composition of Proterozoic evaporite minerals (summarized in Planavsky et al. 2018). Some workers have suggested much higher levels of atmospheric pO₂ during this period (Zhang et al. 2016), raising the possibility of significant time-dependent variability in atmospheric redox during the Mesoproterozoic (cf. Planavsky et al. 2016). In any case, the paradigmatic view is that atmospheric pO₂ was significantly lower than that of the modern Earth prior to ~600–800 Ma, somewhere between ~0.1% PAL and 10% PAL (Fig. 4). Regulation of atmospheric O₂ in this state is still uncertain, but it is likely to have been controlled by incomplete oxidation of reduced C, Fe, and S in Earth’s crust together with muted biospheric O₂ production as a result of P scavenging from pervasively anoxic oceans.

Atmospheric CH₄ during Earth’s middle history is less well-constrained, because there are currently no available geologic proxies for the quantitative abundance of CH₄ in Earth’s atmosphere at intermediate redox states. Indeed, even under a mildly reducing atmosphere, proxy constraints are indirect and plagued by significant uncertainty. Photochemical models predict that once atmospheric O₂ is high enough to support significant ozone, CH₄ can be shielded from photochemical destruction and tends to “rebound” to relatively high values following an initial drop during the Great Oxidation Event (Claire et al. 2006), perhaps to as high as ~100 ppm. However, models of ocean biogeochemistry that take into account the potential for effective microbial sinks for CH₄ in marine environments tend to predict steady-state atmospheric CH₄ abundances that are much lower than ~100 ppm under Proterozoic conditions, more on the order of ~1–10 ppm (Olson et al. 2016), unless CH₄ abundances had been supplemented by significant fluxes from nascent terrestrial ecosystems (Zhao et al. 2018).

In any case, evidence from the rock record indicates a dramatic reorganization of Earth surface biogeochemistry and ocean–atmosphere redox state at some point in the late Proterozoic, perhaps as early as ~800 Ma (Lyons et al. 2014). This period may have marked a pervasive—though perhaps episodic—oxygenation of Earth’s deepest oceans, broadly in step with major shifts in biospheric complexity and perturbations to global climate. Earth’s ocean–atmosphere system has been largely oxidizing subsequent to this tumultuous interval. There have been very occasional excursions to more pervasively reducing environments in the ocean interior, but against a backdrop of strongly oxidizing atmospheric chemistry (Fig. 4). Regulation of atmospheric O₂ in this state has been principally through the incomplete oxidation of organic carbon in Earth’s crust and the increased efficiency of organic carbon burial when oxygenation of the ocean interior decreases (Daines et al. 2017).

**MAJOR REMAINING FRONTIERS**

The history of ocean–atmosphere redox on Earth has been understood in broad strokes for many decades (Holland 1962), and this long-term evolutionary texture is very likely robust. However, placing quantitative constraints on ancient atmospheric composition and ocean chemistry is challenging, and many fascinating questions remain. In particular, the coming years will witness many exciting advances in efforts to place more precise constraints on background atmospheric O₂ during the Paleo- and Mesoproterozoic, to constrain the overall size and scope of Earth’s photosynthetic biosphere through time, better understand the mechanisms regulating the amount of atmospheric O₂ during different periods of Earth’s history and the transitions between them, and to develop a better mechanistic understanding of the processes that link major nutrient cycles to evolving ocean–atmosphere O₂.

Many of the geochemical proxies for the abundance of O₂ in surface environments that have been the most useful in recent years are still being developed, and more direct coupling of proxy records with quantitative models will be central to the further refinement of quantitative estimates for atmospheric pO₂. Issues of particular interest include constraining the duration, causes, and magnitude of Neoproterozoic “whiffs” of O₂; estimating atmospheric pO₂ during the putative “oxygen overshoot” in the Paleoproterozoic; and more definitively constraining O₂ levels in surface environments during the Mesoproterozoic in the lead up to the rise of complex life. In some cases, proxy reconstructions for atmospheric O₂ will rely on independent constraints on other atmospheric properties, such that the development of better estimates for other atmospheric gases (such as CO₂) will also be crucial for constraining O₂.

Much the same can be said for attempts to constrain the productivity of Earth’s biosphere through time, a critical component of the O₂ cycle. A particularly exciting development in this regard is the exploration of the triple oxygen isotope composition of marine and lacustrine sediments during Earth’s history, which can potentially be used to constrain global photosynthetic productivity (Crockford et al. 2018) or be inverted to estimate atmospheric pO₂ (Planavsky et al. 2018). However, a number of challenges remain, including the potentially sporadic stratigraphic distribution of the sedimentary archives for the rare oxygen isotope signal, the need for more mechanistic models that couple surface isotope signals to atmospheric production and downstream recycling, and limits of precision that may ultimately be controlled by the need for independent constraints on atmospheric CO₂.

**ElEmEnts**

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Further development of empirical constraints from Earth’s rock record will provide crucial tie points for theoretical models aimed at understanding the basic processes that regulate Earth’s oxygen cycle, and the processes that link the surface redox landscape to nutrient recycling and the productivity of the biosphere. A striking example is the Earth system during mid-Proterozoic time, ~1.8–0.8 Ga. Modelled redox states are well above those inferred for the Hadean, with atmospheric O₂ well below that of the modern Earth, but how such ‘weakly oxygenated’ conditions at Earth’s surface would be regulated on long timescales is a major outstanding question. This question is also part in part to uncertainties in empirical reconstructions of atmospheric O₂, but it is also related to the fact that Earth’s O₂ cycle at intermediate redox states is likely to be strongly controlled by a range of processes not currently well represented in biogeochemical models. Foremost, parameterizations of nutrient scavenging under reducing conditions; the competitive dynamics between oxygenic and anoxygenic photosynthetic organisms; and the redox cycling of reduced C, S, and Fe in Earth’s crust under low O₂ conditions all need to be improved. Some of these improvements will only be possible with new kinetic data.

Interesting questions remain even deeper into Earth’s past. For example, it is still not definitively known whether the reducing capacity of Earth’s mantle (as described in Stagno and Fei 2020) has changed over time, with potentially dramatic implications for processes controlling atmospheric abundances of O₂, CH₄, H₂, and a range of other redox-active species. As a result, our understanding of the background redox state of Earth’s atmosphere during Hadean/Archean time would benefit significantly from better empirical constraints on mantle redox state during Earth’s early history and better empirical and/or theoretical constraints on degassing fluxes throughout Earth’s evolution. In data supporting the hypothesis that oxygenic photosynthesis first emerged and began to influence the redox structure of Earth’s surface oceans. Constraining the timing of this foundational biological novelty with confidence will ultimately require the leveraging of new geochemical redox proxies, the combination of new observations with existing and well-established approaches, and better quantitative models for evaluating “false positives” for the presence of O₂ in Earth’s surface environments when Earth’s atmosphere was reducing.

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REFERENCES
Planavsky NJ and 6 coauthors (2018) A case for low atmospheric oxygen levels during Earth’s middle history. Emerging Topics in Life Sciences 2: 149-159
Planavsky NJ and 8 coauthors (2016) No evidence for high atmospheric oxygen levels 1,400 million years ago. Proceedings of the National Academy of Sciences of the United States of America 113: E2550-E2551

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