

Turn Your Experiment Run...into a Sprint
New! Fast Chemistries for Fast PCR and Real-Time PCR Instruments
[Learn More >](#)

Science

[AAAS](#) [Magazine](#) [News](#) [Signaling](#) [Careers](#) [Multimedia](#) [Collections](#)

[Current Issue](#) [Previous Issues](#) [Science Express](#) [Science Products](#) [My Science](#)

[About the Journal](#)

[Home](#) > [Science Magazine](#) > [8 September 2000](#) > [Des](#)

Marais , pp. 1703 - 1705

Science 8 September 2000:
Vol. 289, no. 5485, pp. 1703 - 1705
DOI:

PERSPECTIVES

EVOLUTION:

When Did Photosynthesis Emerge on Earth?

David J. Des Marais*

Life began very early in Earth's history, perhaps before 3800 million years ago (Ma) (1), and achieved remarkable levels of metabolic sophistication before the end of the Archean around 2500 Ma (2, 3). The great antiquity of our biosphere might indeed illustrate how easily life can arise on a habitable planet, but it also portends the challenges that confront our efforts to become intimately familiar with our earliest ancestors. The earliest sedimentary rocks have typically undergone extensive alteration by metamorphism, taking a serious toll on microfossils (4). Fortunately, memories of our distant forebears are recorded not only in ancient rocks, but also in biological macromolecules (5) and pathways. The two records are highly complementary: The geologic record offers the absolute timing of evolutionary innovations and their environmental context, while the living biochemical record can reveal the sequence of development of key pathways and biomolecules.

On page 1724 of this issue, Xiong *et al.* (6) have tapped the biological record to study the evolution of photosynthesis. They have obtained new sequence information for genes involved in photosynthesis and performed phylogenetic analyses on the major groups of photosynthetic bacteria. The study better defines the molecular origins of these groups and clarifies the great antiquity of anoxygenic photosynthesis.

When our biosphere developed photosynthesis, it developed an energy resource orders of magnitude larger than that available from oxidation-reduction reactions associated with weathering and hydrothermal activity. The significance of this innovation can be illustrated quantitatively for modern Earth. Hydrothermal sources deliver $(0.13 \text{ to } 1.1) \times 10^{12} \text{ mol year}^{-1}$ globally of reduced S, Fe^{2+} , Mn^{2+} , H_2 , and CH_4 (7); this is estimated to sustain at most about $(0.2 \text{ to } 2.0) \times 10^{12} \text{ mol C year}^{-1}$ of organic carbon production by microorganisms capable of using hydrothermal energy as their energy source (8). In contrast, global photosynthetic productivity is estimated at $9000 \times 10^{12} \text{ mol C year}^{-1}$ (9, 10). Global thermal fluxes were greater in the distant geologic past (11, 12), but the onset of oxygenic photosynthesis most probably increased global organic productivity by at least two to three orders of magnitude. This enormous productivity resulted principally from the ability of oxygenic photosynthetic bacteria to capture hydrogen for organic biosynthesis by cleaving water. This virtually unlimited supply of hydrogen freed life from its sole dependence upon abiotic chemical sources of reducing power, such as hydrothermal sources and weathering. Communities sustained by oxygenic photosynthesis could thrive wherever supplies of sunlight, moisture, and nutrients were sufficient.

ADVERTISEMENT



AAAS
Policy Alert
SEE SAMPLE ISSUE

ADVERTISEMENT



Submit your paper today

CLICK HERE

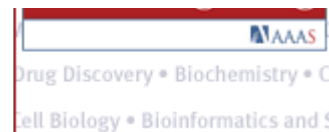
Genetics • Cell Biology • Bioinformatics and
Systems Biology • Molecular Biology
Molecular Biology • Plant Biology
Development • Biology • Immunology
Science Signaling

Photosynthetic microbial communities have left a relatively robust fossil record, in part because their productivity was particularly high on stable submerged continental platforms and margins, and thus contributed to sediments with excellent potential for long-term preservation. The cyanobacterial microfossil record is robust throughout the Proterozoic (around 2500 to 543 Ma) (3). The record of organic biomarkers--molecules that are highly diagnostic for their parent organisms--is consistent with the microfossil record. For example, only cyanobacteria are known to synthesize 2-methyl bacteriohopanepolyols, which are transformed in sediments to 2-methylhopanes. The latter have now been identified in rocks as old as 2500 to 2700 million years (My) (13, 14). The stromatolitic carbonates that were widespread along continental margins throughout the Proterozoic (15) have long been associated with cyanobacterial communities. Few stromatolites contain identifiable cellular fossils, but the large, Paleoproterozoic (2500 My old and younger) stromatolitic reefs that rival modern reefs in size, architecture, and extent (16) compel the present author to cite their development as firm evidence for oxygenic photosynthesis having become well established by 2500 Ma. Buick (17) also concludes that late Archean stromatolites, observed in 2700-My-old lake deposits, required oxygenic photosynthesis to develop abundantly in environmental settings that lacked evidence of hydrothermal activity.

The molecular oxygen released by photosynthesis leads to additional evidence for the antiquity of oxygenic photosynthesis. Before oxygenic photosynthesis arose, ambient oxygen levels were insignificantly low, because chemical sinks such as reduced geothermal outflows and rock weathering greatly exceeded the main abiotic source of oxygen, the photochemical dissociation of water vapor coupled with loss of hydrogen to space (18). The discovery of sterane biomarkers in 2700 Ma sediments (14) demonstrates not only the existence of eukaryotic organisms, but also that free oxygen was available for sterol biosynthesis. The extremely low $^{13}\text{C}/^{12}\text{C}$ values in 2800-My-old kerogens have been attributed to methanotrophic bacteria, which require both oxygen and methane (19). The substantial deposition rates of ferric iron in massive banded iron sediment formations before 2500 Ma are clearly consistent with an abundant biological source of free oxygen (20). Indeed, vast sedimentary deposits of organic carbon, reduced sulfide, ferric iron, and sulfate on continental platforms and along coastal margins are among the most prominent and enduring legacies of billions of years of oxygenic photosynthetic activity (21).

The geologic record thus offers strong evidence for the evolution of oxygenic photosynthesis before 2800 Ma. There are, however, hints of even earlier origins. The microfossil record of cyanobacteria may extend to 3300 to 3500 Ma (22), although the evidence for these early Archean occurrences is controversial (23). Stromatolites occurring in 3460-My-old carbonates and silicified carbonates of the Warrawoona Group, Western Australia, were recently described by Hofman *et al.* (24). These stromatolites developed in a partially restricted, low-energy shallow hypersaline basin (25). Hofman *et al.* (24) conclude that microorganisms were involved in the accretion of these stromatolites. They also surmise that microbial phototaxis (light-stimulated microbial motility) may have played a role in shaping them, but conclude that the evidence for the presence of photosynthetic biota is not yet definitive.

The carbon isotopic record of early Archean carbonates and reduced carbon is consistent with, but not yet compelling for, oxygenic photosynthesis. The isotopic patterns are consistent with isotopic discrimination by some chemoautotrophic bacteria and anoxygenic photoautotrophic bacteria, in addition to oxygenic photoautotrophs (26). But the absence of conclusive evidence should not be interpreted as conclusive evidence of absence. Although the early Archean fossil record contains at best a handful of demonstrated microfossils, none of which are as yet physiologically definitive, this low apparent diversity should not be interpreted too literally (3). It is known that as metamorphic alteration of fossil-containing rocks intensifies, the apparent diversity of remaining microfossil assemblages decreases (4). Conclusive evidence for oxygenic photosynthesis probably also diminishes. Current evidence for early Archean oxygenic photogenesis is perhaps not yet compelling, but it is consistent with it.



[To Advertise](#) [Find Products](#)

ADVERTISEMENT

FEATURED JOBS

AQUATIC BIOGEOCHEMIST
University of Montana
Montana

**Institute for Diabetes,
Obesity and
Metabolism/Department of
Radiology**
University of Pennsylvania
Pennsylvania

**Appointment of Executive
Director**
National University of
Singapore
Singapore

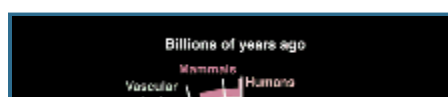
Director
University of Wyoming
Laramie, WY

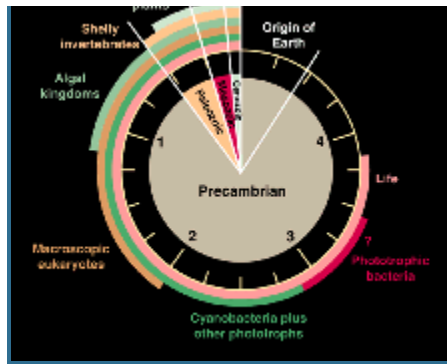
POSTDOCTORAL POSITION
University of Pittsburgh
Cancer Institute
Pittsburgh, PA

Research Specialist
Johns Hopkins University
School of Medicine
Baltimore, MD

Research Associate
Manchester Metropolitan
University
Manchester, United
Kingdom

[More jobs](#)





Earth's biogeologic clock. The great antiquity of our biosphere contrasts sharply with the relative youth of plants and animals. The dual geological and molecular biological records of microorganisms indicate that our early biosphere was remarkably complex. All of the major photosynthetic groups of bacteria arose prior to 2800 Ma, perhaps much earlier.

The report by Xiong *et al.* (6) adds an important constraint to the perspective outlined above. The authors demonstrate conclusively for the first time that the major lineages of pigments involved in anoxygenic photosynthesis arose before the development of oxygenic photosynthesis. This indicates that the six major bacterial lineages had largely developed by the mid-Archean, around 3000 to 2800 Ma, and perhaps much earlier. The study also shows that the early biosphere passed through a stage during which even its photosynthetic populations depended exclusively on abiotic sources of reducing power. Can we recognize such a stage in the geologic record? The fossil record of anoxygenic phototrophic bacteria is poorly known, although ancient populations have been identified in much younger rocks on the basis of organic biomarkers. The presence of Chlorobiaceae in Paleozoic sediments has been inferred based on the identification of a ^{13}C -enriched aromatic polyisoprenoid (27). Geoporphyrins from purple sulfur bacteria have been identified in ancient shales (28). The survival of traces of Archean oil (29) offers the possibility to extend the biomarker record of anoxygenic phototrophs considerably.

As the great antiquity of photosynthesis becomes more and more apparent, it also becomes easier to envision an ancient, global biosphere sustained principally by anoxygenic photosynthesis. The global geothermal heat flow was substantially higher during Earth's first billion years (11), and the vigorous geothermal outgassing probably dispersed reduced chemical species throughout sunlit aquatic environments. Perhaps the substantial decline in thermal activity between 4000 and 3000 Ma created opportunities for oxygenic photosynthesis to develop. Both the geologic and living biological records of our early biosphere promise further key insights into the origin and early evolution of photosynthesis.

References

1. S. J. Mojzsis *et al.*, *Nature* **384**, 55 (1996) [GEOREF].
2. J. W. Schopf and C. Klein, Eds., *The Proterozoic Biosphere: A Multidisciplinary Study* (Cambridge Univ. Press, New York, 1992) [publisher's information].
3. A. H. Knoll, *Palynology: Principles and Applications*, J. Jansonius and D. C. MacGregor, Eds. (American Association of Stratigraphic Palynologists Foundation, Dallas, TX, 1996), pp. 51-80.
4. A. H. Knoll, P. K. Strother, S. Rossi, *Precambrian Res.* **38**, 257 (1988) [GEOREF].
5. C. R. Woese, O. Kandler, M. Wheeler, *Proc. Natl. Acad. Sci. U.S.A.* **87**, 4576 (1990) [Medline].
6. J. Xiong *et al.*, *Science* **289**, 1724 (2000).
7. H. Elderfield and A. Schultz, *Annu. Rev. Earth Planet. Sci.* **24**, 191 (1996) [GEOREF].
8. D. J. Des Marais, *Geomicrobiology*, J. Banfield and K. Nealson, Eds. (Mineralogical Society of America, Washington, DC, 1997), vol. 35, pp. 429-445.
9. J. H. Martin, G. A. Knauer, D. M. Karl, W. W. Broenkow, *Deep Sea Res.* **34**, 267 (1987).

10. J. S. Olson *et al.*, *Atmospheric Carbon Dioxide and the Global Carbon Cycle*, J. R. Trabalka, Ed. (U.S. Department of Energy, Washington, DC, 1985), pp. 175-213.
11. D. L. Turcotte, *Earth Planet. Sci. Lett.* **48**, 53 (1980) [[GEOREF](#)].
12. D. Kadko, J. Baross, J. Alt, *Seafloor Hydrothermal Systems: Physical, Chemical, Biological, and Geological Interactions*, vol. 91 of Geophysical Monograph Series (American Geophysical Union, Washington, DC, 1994), pp. 446-466.
13. R. E. Summons *et al.*, *Nature* **400**, 554 (1999) [[Medline](#)].
14. J. J. Brocks, G. A. Logan, R. Buick, R. E. Summons, *Science* **285**, 1033 (1999).
15. M. R. Walter, Ed., *Stromatolites* (Elsevier, New York, 1976).
16. J. P. Grotzinger, *Controls on Carbonate Platform and Basin Development*, P. D. Crevello, J. L. Wilson, J. F. Sarg, J. F. Read, Eds. (Special Publication 44, Society of Economic Paleontologists and Mineralogists, Tulsa, OK, 1989), pp. 79-106.
17. R. Buick, *Science* **255**, 74 (1992) [[GEOREF](#)].
18. H. D. Holland, *The Chemical Evolution of the Atmosphere and Oceans* (Princeton Univ. Press, Princeton, NJ, 1984).
19. J. M. Hayes, *Early Life on Earth*, S. Bengtson, Ed. (Columbia Univ. Press, New York, NY, 1994), pp. 220-236 [[publisher's information](#)].
20. N. J. Beukes and C. Klein, *The Proterozoic Biosphere*, J. W. Schopf, Ed. (Cambridge Univ. Press, New York, 1992), pp. 147-151 [[publisher's information](#)].
21. D. J. Des Marais, *Org. Geochem.* **27**, 185 (1997).
22. J. W. Schopf, *The Proterozoic Biosphere*, J. W. Schopf, Ed. (Cambridge Univ. Press, New York, 1992), pp. 25-39 [[publisher's information](#)].
23. R. Buick, *Palaios* **5**, 441 (1991).
24. H. J. Hofman *et al.*, *Geol. Soc. Am. Bull.* **111**, 1256 (1999) [[GEOREF](#)].
25. D. R. Lowe, *Precambrian Res.* **19**, 239 (1983) [[GEOREF](#)].
26. M. Schidlowski, J. M. Hayes, I. R. Kaplan, *Earth's Earliest Biosphere*, J. W. Schopf, Ed. (Princeton Univ. Press, Princeton, 1983), pp. 149-186.
27. R. E. Summons and T. G. Powell, *Nature* **319**, 763 (1986) [[GEOREF](#)].
28. M. I. Chicarelli, S. Kaur, J. R. Maxwell, *Metal Complexes in Fossil Fuels; Characterization and Processing*, ACS Symposium Series 344, R. H. Filby and J. F. Branthaver, Eds. (American Chemical Society, Washington, DC, 1987), pp. 40-67.
29. R. Buick, B. Rasmussen, B. Krapez, *Am. Assoc. Petrol Geol. Bull.* **82**, 50 (1998).

The author is at NASA Ames Research Center, Moffett Field, CA 94035, USA. E-mail: ddesmarais@mail.arc.nasa.gov

The editors suggest the following Related Resources on Science sites:

In Science Magazine

LETTERS

On the Origins of Photosynthesis

Kenneth M. Towe; and David J. Des Marais (19 January 2001)
Science **291** (5503), 436. [DOI: 10.1126/science.291.5503.436]

[Full Text »](#)

RESEARCH ARTICLES

Molecular Evidence for the Early Evolution of Photosynthesis

Jin Xiong, William M. Fischer, Kazuhito Inoue, Masaaki Nakahara, and Carl E. Bauer (8 September 2000)

Science **289** (5485), 1724. [DOI: 10.1126/science.289.5485.1724]

[Abstract »](#) [Full Text »](#) [PDF »](#)