The average cell size of marine phytoplankton is critical for the flow of energy and nutrients from the base of the food web to higher trophic levels. Thus, the evolutionary succession of primary producers through Earth’s history is important for our understanding of the radiation of modern protists ~800 million years ago and the emergence of eumetazoan animals ~200 million years later. Currently, it is difficult to establish connections between primary production and the proliferation of large and complex organisms because the mid-Proterozoic (~1,800–800 million years ago) rock record is nearly devoid of recognizable phytoplankton fossils. We report the discovery of intact porphyrins, the molecular fossils of chlorophylls, from 1,100-million-year-old marine black shales of the Taoudeni Basin (Mauritania), 600 million years older than previous findings. The porphyrin nitrogen isotopes (δ15Npor = 5.6–10.2‰) are heavier than in younger sedimentary sequences, and the isotopic offset between sedimentary bulk nitrogen and porphyrins (εpor = –5.1 to –0.5‰) points to cyanobacteria as dominant primary producers. Based on fossil carotenoids, anoxicogenic green (Chlorobiaceae) and purple sulfur bacteria (Chromatiaceae) also contributed to photosynthate. The low εpor values, in combination with a lack of diagnostic eukaryotic steranes in the time interval of 1,600–1,000 million years ago, demonstrate that algae played an insignificant role in mid-Proterozoic oceans. The paucity of algae and the small cell size of bacterial phytoplankton may have curtailed the flow of energy to higher trophic levels, potentially contributing to a diminished evolutionary pace toward complex eukaryotic ecosystems and large and active organisms.

The succession of primary producers in the oceans shaped marine ecology through Earth’s history (1). Primary producers form the base of the food web. Their cell size, elemental stoichiometry, and cell density influence the flow of energy and nutrients to higher trophic levels (2), presumably setting limits for ecosystem complexity. The composition of phytoplankton communities in Earth’s early oceans, including anoxygenic and oxygenic phototrophic bacteria and eukaryotic algae, may thus have set the pace for the emergence and radiation of different groups of filter feeders, grazers, and predators, including the proliferation of modern protists ~800 Ma and the appearance of eumetazoan animals some 200 My later (1) (Fig. 1B). Based on the fossil record, chlorophyll c (Chl c) algae, including dinoflagellates, coccolithophorids, and diatoms, were the major energy and carbon source in the oceans of the past ~250 My (Fig. 1B). By contrast, Paleozoic (541–251 Ma) and Ediacaran (635–541 Ma) oceans were presumably dominated by primary endosymbiotic algae (Archaeplastida), encompassing the red (Rhodophyta) and green algae (Chlorophyta) (1). Deeper yet in time, reconstructing the succession of primary producers becomes challenging. Phytoplankton without a preserved cuticle or skeleton are rarely preserved in the body fossil record, and there are no uncontentious fossils of planktonic bacteria or algae in the pre-Ediacaran (3). However, biomarkers, the molecular fossils of biological lipids, can provide complementary information about primary producers. For example, hydrocarbon fossils of carotenoid pigments extracted from sedimentary rocks have been used to detect phototrophic green (Chlorobiaceae) and purple sulfur bacteria (PSB) (Chromatiaceae) in 1,640-My-old marine ecosystems (4, 5), while the concentration of eukaryotic steranes, relative to bacterial hopanes, may provide basic information about the ecological relevance of Precambrian algae (6). The relative abundance as well as diversity of steranes dramatically increased in the brief interval between the Sturtian and Marinoan snowball Earth glaciations of 659–645 Ma, heralding the rise of planktonic algae as important primary producers in the oceans (Fig. 1A). Steranes in the time interval 645 to ~500 Ma have a strong predominance of stigmastane (a structure with 29 carbon atoms, C29), revealing Chlorophyta, a division of green algae, as dominant primary producers from the late Cryogenian to early Paleozoic (Fig. 1) (7, 8).

Preceding the Cryogenian rise of planktonic algae, the oldest clearly indigenous eukaryotic steranes appear in the geological record ~900–800 Ma, albeit in low concentrations relative to bacterial biomarkers (7, 8). These Tonian (1,000–720 Ma) steranes display a primordial distribution with a nearly 100% predominance of cholesterol (C27, red circles in Fig. 1A), which may be related to the activity of rhodophytes (9) or heterotrophic eukaryotes (8).

Significance

The oceans of Earth’s middle age, 1.8–0.8 billion years ago, were devoid of animal-like life. According to one hypothesis, the emergence of large, active organisms was restrained by the limited supply of large food particles such as algae. Through the discovery of molecular fossils of the photopigment chlorophyll in 1.1-billion-year-old marine sedimentary rocks, we were able to quantify the abundance of different phototrophs. The nitrogen isotopic values of the fossil pigments showed that the oceans were dominated by cyanobacteria, while larger planktonic algae were scarce. This supports the hypothesis that small cells at the base of the food chain limited the flow of energy to higher trophic levels, potentially retarding the emergence of large and complex life.

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Going yet further back in time into the mid-Proterozoic (1,800–800 Ma), evidence for algae becomes scarce. Based on molecular clocks, the last common ancestor of all algae originated broadly between 1,900 and 1,400 Ma (10, 11), predating the oldest unambiguous fossil of a eukaryotic phototroph, the ∼1,050 Ma (12) benthic red alga *Bangiomorpha* (13) (Fig. 1B). However, clearly indigenous biomarkers for algae have not yet been found in sedimentary rocks >900 Ma despite preservation of bacterial hopanes (Fig. 1). Whether this lack of steranes is caused by preferential degradation of algal organic matter (14) or reflects genuine paucity of eukaryotic phototrophs, remains unresolved.

Here, we present an approach to gauge the activity of bacterial and eukaryotic primary producers in the Precambrian. Based on Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR MS), we identify geoporphyrins in 1,100-Ma-old black shales. Porphyrins are the molecular fossils of (bacterial) chlorophylls. Their nitrogen isotopic composition can provide quantitative information about dominant phototrophs in past ecosystems (15). Some groups of phototrophs possess a characteristic offset ($^{15}$N) between the nitrogen isotopic composition of the whole cell and chlorophylls, independent of the nitrogen source (15), and under suitable conditions this offset is largely unaffected by degradation processes in the water column and bottom sediments (16). Therefore, the nitrogen isotopic composition of porphyrins in sediments and sedimentary rocks may preserve information about primary producers present in an ancient environment, largely independent of physical and chemical conditions such as food source and diagenesis.

Depositional Environment

Eleven black shales studied for their molecular content (Table 1 and SI Appendix, Table S1) come from the En Nesoar and Tourist formations of the 1,100 Ma El Mreïti Group (17) deposited on an epicratonic platform of the Taoudeni Basin on the West African Craton (18). The black shales, with a carbon content of up to 32%, accumulated beneath anoxic ferruginous and occasionally sulfidic waters (19) during maximal flooding of the craton in a quiet subwave base environment possibly protected by offshore stromatolite reefs (18). The black shales contain micrometer-thin laminae of organic matter and pyritized filamentous sheaths that are interpreted as benthic microbial communities, presumably of heterotrophic and/or chemosynthetic microorganisms thriving beneath anoxic waters (19), and irregularly shaped, discrete or bedding-parallel accumulations of organic particles typical of planktonic debris (SI Appendix, SI Geology and Samples).

Results

Biomarker Syngeneity. Potential contaminants introduced in the laboratory were monitored using comprehensive accumulatory system blanks, and compounds that may have permeated into rock samples were quantified and eliminated by comparing the concentration of individual hydrocarbons in the exterior and interior portions of the rock (20–22). All compounds reported here are demonstrably indigenous (SI Appendix, SI Syngeneity Assessment).

Hydrocarbon Biomarkers. All El Mreïti Group shales yielded pentacyclic terpanes, including αβ-hopanes with 27–35 carbon atoms as well as gammacerane (Fig. 2A). A complete series of C$_{31}$ to C$_{50}$ 2α-methylhopanes was also present in low concentrations. The 2α-methylhopane index, defined as the abundance of C$_{31}$ 2α-methylhopane relative to C$_{30}$ αβ-hopane, was low, ranging between 0.35 and 0.77%. Although hopanes were well above detection limits, C$_{25}$ to C$_{30}$ steranes were not detected in any sample (Table 1 and SI Appendix, Fig. S1), consistent with a previous study on Atar Group shales that also failed to detect steranes (23). Based on determination of detection limits, the maximum level of steranes that may be present beneath the noise is extremely low. The ratio of maximum hypothetical C$_{27}$ steranes relative to C$_{27,33}$ hopanes ($^{30}$H$_{max}$) falls into the range of 0.0002–0.0014 (average, 0.00065; n = 11; Table 1). The El Mreïti Group shales were also devoid of the saturated carotenoid derivatives lycopane, β-carotane and γ-carotane. However, two of the most dominant biomarker classes in the aromatic hydrocarbon fraction were C$_{13}$ to C$_{27}$ 2,3,6-trimethyl aryl isoprenoids (2,3,6-Al) and C$_{13}$ to C$_{25}$ 2,3,4-trimethyl aryl isoprenoids (2,3,4-Al) (Fig. 2B), the degradation products of aromatic carotenoids.

Identification of Porphyrins. Geoporphyrins were identified using reversed-phase HPLC-UV/Vis and atmospheric pressure photoionization coupled to FT-ICR MS at 9.4 Tesla. Based on HPLC
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Selected biomarker classes identified in black shales (sample 08, drill core S2). The measured δ15Nbulk values for six porphyrins ranged between −1.1 and −5.2‰ (Table 2). Although δ15Nbulk of organic-rich shales, such as those studied here with total organic carbon (TOC) between 2.1 and 31.7‰, commonly closely reflect the nitrogen isotopic composition of primary bulk organic matter (26), two factors must be considered before interpreting the data: first, the contribution of inorganic nitrogen to δ15Nbulk values, and second, isotopic fractionation effects associated with diagenesis and heterotrophic reworking of photosynthetic organic matter. This may be present in samples beneath the analytical noise (SI Appendix, SI Methods).

Although inorganic nitrogen sources are usually negligible in organic matter-rich samples (16), the El Mreïti Group shales may contain up to 0.1% inorganic nitrogen (see intercept on Nbulk axis in SI Appendix, Fig. S5) (30). However, it is plausible that the conversion of smectite to illite during burial diagenesis of the clay-rich formation with porphyrin standards and comparison with reference material (24, 25) (SI Appendix, Fig. S2), the most abundant porphyrins are nickel Etio porphyrins (1), C30 to C32 deoxophyloerythrophorphy tin (DPEP, 2a–f), C31 and C32 butano (3a–b), and C34 diDPEP (4a) porphyrins, as well as tentatively identified vanadyl (VO) C30–C32 DPEP (2a,h,f) (structure numbers are defined in Fig. 3). FT-ICR MS is abundance-contoured plots confirm C29 to C32 DPEP (2) and C29 to C31 Etio (1) as the most abundant Ni-porphyrins, and C32 DPEP and C30 Etio as the highest concentrated VO-porphyrin (Fig. 4). Also identified were Ni- and VO-based diDPEP (4, 5, and/or 6), rhodo-Etio (7), and rhodo-DPEP (8) (26). Each of these compound classes occurs as homologous series across a wide carbon number distribution with more than 40 carbon atoms (Fig. 4) due to side-chain length variation of core structural motifs. Although these side-chain elongated porphyrins may include bac teriochlorophyll (BChl)-derived structures (27), sample limitation prevented characterization of precise alkyl side-chain configurations (SI Appendix, SI Additional Details on Biomarker Identification and Interpretation).

### Nitrogen and Carbon Isotopic Composition of Individual Ni-Porphyrins.

For the richest sample (#13), it was possible to measure the carbon and nitrogen isotopic compositions of six isolated and purified Ni-porphyrins (SI Appendix, Fig. S3G). Based on molecular masses for isolated compounds determined by HPLC-atmospheric pressure chemical ionization (APCI)–MS, purified porphyrins correspond to C29 to C31 Etio and C30 and C32 DPEP structures (Table 2). Measured C/N ratios of the purified porphyrins (C/N = 12.7–22.1) were higher than predicted stoichiometries (C/N = 7.0–8.0; Table 2), which suggests that isolates still contained other nonporphyrin compounds that were carried over across multiple steps of chromatographic purification (SI Appendix, Fig. S4). However, the APCI mass spectra of the isolated fractions (SI Appendix, Fig. S3A–F) all show relatively clean molecular ions diagnostic for individual Ni-porphyrins and only minor foreign masses. This indicates that the contaminants are poorly ionized by APCI and likely represent hydrocarbons. The presence of nonporphyrin nitrogen-containing compounds is unlikely in these fractions due to their higher polarity relative to porphyrins and absence of significant signals in the APCI mass spectra (see SI Appendix, SI Methods for a detailed discussion about avoidance of artifacts and contaminants). The δ15Npor values measured for six porphyrins ranged between 5.6‰ and 10.2‰, with corresponding values of C/Npor between −0.5 and −5.1‰ (Table 2).

### Discussion

**Bulk Nitrogen Isotopes.** The bulk-rock nitrogen isotopic composition (δ15Nbulk) of the En Nesoar and Tourist Formation shales falls into the narrow range of 4.2–6.1‰ (n = 11, average = 5.1‰, Table 1) and is slightly higher than those of the Mesoproterozoic Belt Basin (−1.1–5.2‰) (28). Although δ15Nbulk of organic-rich shales, such as those studied here with total organic carbon (TOC) between 2.1 and 31.7‰, commonly closely reflect the nitrogen isotopic composition of primary bulk organic matter (29), two factors must be considered before interpreting the data: first, the contribution of inorganic nitrogen to δ15Nbulk values, and second, isotopic fractionation effects associated with diagenesis and heterotrophic reworking of photosynthetic organic matter. This may be present in samples beneath the analytical noise (SI Appendix, SI Methods).

![Fig. 2. Selected biomarker classes identified in black shales (sample 08, drill core S2, 140.25 m) of the El Mreïti Group. (A) Summed GC-MS metastable reaction monitoring (MRM) chromatograms of M+ → 191 transitions of the saturated fraction. A complete homologous series of 17a(H),21b(H)-hopanes with 27–35 carbon atoms (black triangles) were identified. Tγ = 18(α)-22,29,30-trisnorhopane, and γ = gammacerane. (B) m/z 134 selected ion recording chromatogram of the aromatic fraction. Filled circles, 2,3,6-Al; open circles, 2,3,4-Al.](Image)
samples yielded substitution of potassium by ammonium (e.g., ref. 31). As the ammonium would have largely derived from the decomposition of organic matter, and as the substitution process is not coupled to major isotopic fractionation (32), δ15Nbulk probably largely reflects the nitrogen isotopic composition of the organic matter. This is further confirmed by the absence of a correlation between TOC and δ15Nbulk.

Secondary processes in the water column or bottom sediment may alter the original N-isotopic composition of the primary organic matter. In modern environments, aerobic heterotrophic reworking of primary organic matter in marine regions with low organic carbon flux and high oxygen exposure increases δ15Nbulk by +3 to +5‰ (33). However, δ15Nbulk values of high TOC sediments do not have such positive isotopic offsets and faithfully record the N-isotopic composition of primary producers (29). Conversely, in anaerobic waters, δ15Nbulk may decrease by 3‰ under laboratory conditions, and a depletion of 1.2‰ has been observed in anoxic sediments of Lake Luguano (34). However, in two large modern anoxic marine basins, the Black Sea and the Cariaco Trench, no negative N-isotopic offsets were observed (35, 36). Moreover, a study of Black Sea sediments found that the isotopic difference between δ15Nbulk and δ15Npor may be up to 15‰ in the isotopic value of δ15Npor (37). Thus, δ15Nbulk of modern organic-rich shales likely reflect δ15N of contemporaneous phytoplankton.

Our data extend the record of εpor, the offset between δ15Nbulk and δ15Npor, by nearly 1 Gy. Problems associated with the extrapolation of nitrogen isotopic information over such immense periods of time, into an eon with potentially different degradation processes, are discussed in detail in SI Appendix, SI Taphonomic Biases. However, even taking possible nonuniformitarian processes into account, it is likely that εpor of highly organic-rich Proterozoic shales, such as those studied here, reflect the nitrogen isotopic composition of primary biomass. Moreover, even large alterations of δ15Nbulk in the range of −3‰ to +10‰ would not impact our conclusions.

**Porphyryn Nitrogen Isotopes (δ15Npor)**. The measured δ15N values of porphyrins (δ15Npor) range between +5.6‰ and +10.2‰ and are at the isotopically heavy end of heavy data observed in modern and Phanerozoic (541 Ga to present) aquatic systems (Fig. 5). Geoporphyrins are believed to reflect the nitrogen isotopic composition of the precursor chlorophylls (δ15NChl). For example, the isotopic offset between total organic matter and porphyrins remains largely constant during burial (16), and demetallation and metal-complex formation during chlorophyll diagenesis have commonly very little effect on nitrogen isotopic fractionation (37). Thus, the isotopic composition of the Mesoproterozoic geoporphyrins likely represent δ15N of the precursor (bacterio) chlorophylls (SI Appendix, SI Taphonomic Biases).

**Biological Origins of the Geoporphyrins.** The previously oldest geoporphyrins come from the early Cambrian (38), so the structures reported here extend the record by 600 My (SI Appendix, SI Other Reports on Proterozoic Porphyrins). However, the structures of the 1,100 Ma porphyrins, when viewed in isolation, provide limited biological information. The major (bacterio) chlorophylls and their most biogenic sources are summarized in SI Appendix, Table S2. General evidence for photosynthetic activity in the Taoudeni Basin is provided by C30 to C32 DPEP (2), which may originate from Chl a, b, and c, and BChl a, b, and d (39). More specifically, 7-nor DPEP (2d) may originate from Chl b and Chl c3 (40, 41), and 17-nor DPEP (2f) from Chl c (42). Chl b is an accessory pigment in higher plants, green algae, and cyanobacteria (43), although it can also be chemically generated by alteration of Chl a (44), while Chl c is found in chromophyte algae and cyanobacteria (45). Furthermore, the etio (1), butano (3), diDPEP (4, 5, 6), and rhodostructures (7, 8) may have the same biogenic sources as the corresponding DPEP compounds. Hence, the identified porphyrins do not point to any specific chloropigment source and may derive from most groups of oxygenic phototrophs plus a possible contribution from anoxygenic phototrophic bacteria.

More information can be obtained from the nitrogen isotopic composition of the porphyrins. Phototrophic organisms have a specific nitrogen isotopic offset (εpor) between total biomass and chloropigments (εpor = δ15Nbio − δ15NChl) (Fig. 6). In culture experiments, freshwater and marine cyanobacteria, purple nonsulfur bacteria, and green and red algae were grown on different nitrogen sources (15). Despite a wide range of nitrogen isotopic compositions of bulk organic matter (δ15Nbulk = −20 to +20‰) and (bacterio)chlorophylls (δ15NChl = −30 to +25‰), SI Appendix, Fig. S6), the isotopic offset εpor is quite uniform within the same group of organisms but relatively distinct between groups. εpor reflects the difference in nitrogen isotopic fractionation...
between chlorophyll biosynthesis and the synthesis of the bulk of other nitrogen-containing metabolites, mainly proteins (16). Furthermore, εpor is independent of the nitrogen source (NH$_4^+$, N$_2$, or NO$_3^-$) and its isotopic composition (15). Assimilated nitrogen species are initially converted to NH$_3$, which is then incorporated into amino acids from which cellular proteins and chlorophylls are synthesized. Although enzymes converting different nitrogen species to NH$_3$ fractionate nitrogen differently and therefore affect the absolute value of δ$^{15}$N$_{cell}$, the isotopic difference between cellular proteins and chlorophylls remains constant within the clade. Accordingly, εpor is also largely independent of redox conditions during cell growth (15, 46).

The El Mreïti Group porphyrins have εpor values ranging from −5.1 to −10.0‰ (Table 2). Fig. 6 compares these values with literature data from culture experiments and recent and Phanerozoic sediments. Freshwater cyanobacteria have median εpor = −10.9‰ (range, −16.0 to −5.7‰; n = 38); marine planktonic cyanobacteria, −0.6‰ (range, −2.9 to 5.7‰; n = 15); marine benthic cyanobacteria, −10.3‰ (range, −10.4‰ to −10.1‰; n = 2); purple nonsulfur bacteria, +10.1‰ (range, 9.3–15.7‰; n = 3); PSB, +5.8‰ (one environmental sample); and red and green algae, +5.2‰ (range, −0.9 to 10.1‰; n = 42) (15, 47–53). Particular organic matter in modern oceans, recent sediments beneath anoxic waters, and Phanerozoic shales all yield positive εpor values overlapping with the algal signal (Fig. 6), indicating that eukaryotic algae were the dominant export producers in these systems. By contrast, the new Mesoproterozoic εpor values clearly fall into the range of cyanobacteria. However, the assignment of the Mesoproterozoic εpor values to cyanobacteria has some remaining uncertainty. Most organism-specific values in Fig. 6 are based on laboratory data from a limited number of species and may not fully reflect natural conditions or variabilities. Moreover, the compilation is missing some groups of phototrophic bacteria that may have been important in the Mesoproterozoic, particularly Chloroflexi and Chlorobiaceae. Thus, the Mesoproterozoic porphyrins probably have a dominant cyanobacterial source plus an unknown contribution from anoxygenic phototrophic bacteria. Notable eukaryotic algal input, however, is unlikely based on the considerable and consistent offset between the positive εpor of alga-dominated Phanerozoic sediments and the negative εpor of the Mesoproterozoic porphyrins.

Hydrocarbon Biomarker Evidence for Primary Producers. Carotenoid derivatives of the 2,3,4- and 2,3,6-AI series are present in all El Mreïti Group shales (Fig. 2B). 2,3,4-AI may be derived from the biological carotenoid derivatives okenane, renieratane, and renierapurpurane, which are all diagnostic for PSB of the family Chromatiaceae (4). The 2,3,6-AI series is most commonly derived from aromatic carotenoids with the chlorobactane, iso-renieratane, and/or β-isorenieratane skeletons found in green sulfur bacteria (GSB) (Chlorobiaceae). As 2,3,4- and 2,3,6-AI are abundant in all Taoudeni bitumens, PSB and GSB probably played an important ecological role. While both families produce BChl a (54), some PSB also produce BChl b (55) and GSB additionally synthesize BChl c, d, or e (SI Appendix, Table S2). These BChls would all have contributed to the porphyrin pool of the extracts. However, their impact remains uncertain because...
The ratio of eukaryotic steranes over bacterial hopanes (S/H) is a first-order estimate for the relative contribution of eukaryotic to bacterial biomass to sediments. Although hopanes were abundant in the black shales of the El Mreïti Group, steranes remained beneath detection limits. Quantification of background noise shows that steranes, if present, fall below S/H = 0.0002–0.0014 (Table 1), which is one to two orders of magnitude lower than the Tonian average (1,000–2,000 Ma; S/H = 0.06 ± 0.10), and two to four orders of magnitude lower than typical Ediacaran to Phanerozoic values (635 Ma to present; average S/H = 0.75 ± 1.1) (Fig. L4) (8).

Most molecular clocks estimate the origin of the last common ancestor of all extant eukaryotes (LECA) >1,400 Ma (e.g., ref. 10). LECA possessed all enzymes required to biosynthesize most modern common sterols (56). Thus, the absence of steranes in 1,100 Ma El Mreïti Group shales must be related to preferential heterotrophic degradation of sterols or to an ecological insufficiency (or absence) of crown group eukaryotes, particularly eukaryotic algae, during deposition of the black shales. According to the “mat-seal” hypothesis, biomarkers from planktonic algae are generally underrepresented in pre-Ediacaran bitumens due to strong heterotrophic reworking of algal debris on the surface of oxygen-producing, shallow-water microbial mats (14). However, this mechanism cannot apply to the black shales of the Tourist Formation, which formed during maximum flooding of the West African Craton beneath relatively deep, anoxic waters (19). Alternatively, sterols may have suffered severe recycling as planktonic algal debris sank slowly through the water column (14). However, the preservation of cyanobacterial porphyrins also excludes this mechanism. Senescent pico- planktonic cyanobacterial cells sink considerably slower than larger algal cells (57), causing preferential degradation of cyanobacterial over algal porphyrins, a process observed in the modern Black Sea (16). Such preferential degradation of slower sinking bacterial biomass would have elevated, not attenuated, the sterane/hopane ratio. Most importantly, however, porphyrins are degradation sensitive, and their mere existence demonstrates that photosynthetic organic matter suffered little degradation during deposition of the El Mreïti Group shales. Therefore, the lack of detectable steranes is not likely related to severe and selective degradation of algal biomass, but presumably reflects primary paucity of sterol-producing eukaryotes (for further discussion, see SI Appendix, SI Taphonomic Biases).

### Table 2. Molecular weight and stable isotope data for individual porphyrins of 13-Ni

| Peak* | Structure† | [M]+ | m/z² | C/N predicted | C/N measured | δ¹⁵Npor, ‰ | δ¹³Cpor, ‰ | εpor, ‰ |
|-------|------------|------|------|---------------|--------------|------------|-------------|---------|
| A     | C₂₈Etio    | 479  | 389  | 7.00          | 18.5         | 7.8        | -28.6       | -2.7    |
| B     | C₂₉Etio    | 493  | 446  | 7.25          | 16.4         | 7.3        | -28.4       | -2.2    |
| C     | C₃₀Etio    | 507  | N.A. | 7.50          | N.A.         | N.A.       | N.A.        | N.A.    |
| D     | C₃₁DPEP    | 505  | 310  | 7.50          | 13.0         | 8.8        | -29.7       | -3.7    |
| E-1   | C₃₁Etio    | 521  | 150  | 7.75          | 22.1         | 10.2       | -29.2       | -5.1    |
| E-2   | C₃₁Etio    | 521  | 335  | 7.75          | 12.7         | 5.6        | -29.5       | -0.5    |
| F     | C₃₂DPEP    | 533  | 332  | 8.00          | 14.1         | 7.2        | -27.2       | -2.1    |

N.A., not available.
*Peak identification in HPLC chromatogram and APCI mass spectra are given in SI Appendix, Fig. S3.
†Structures tentatively assigned based on molecular weight determined by APCI.
‡Detector response of the elemental analysis–isotope ratio mass spectrometry system (1 mV is equivalent to ∼1 ng N). Procedural HPLC blanks yielded N-background values of ∼10 mV (SI Appendix, SI Methods). The Ni octaethylporphyrin standard was injected in the concentration range of 110–610 mV (nine standard measurements before, middle, and after samples), yielding δ¹⁵N = +0.86 ± 0.4‰, and δ¹³C = -34.17 ± 0.2‰. There are no correlations between compound concentration (detector response) and δ¹⁵Npor of C/N measured.
§Mixed signal of porphyrin and coeluting unknown hydrocarbons (SI Appendix).
¶Peak C in the reversed-phase HPLC chromatogram broke into several smaller peaks during the normal-phase HPLC step, with individual concentrations too low for isotopic measurement.
εpor is unknown for GSB, and only a single value is available from a lacustrine PSB species (εpor = +5.8‰) (11).

The Phototrophic Community and the Dominant Primary Producer.

The biomarker assemblage of the 1,100 Ma El Mreïti Group shales suggests that the community of photothrophs included cyanobacteria, PSB, and GSB. Based on the absence of steranes and low εpor values, eukaryotic algae did not play a detectable role. Although ornamented and process-bearing organic-walled microfossils interpreted as unambiguously eukaryotic are present in El Mreïti Group sediments (58), they do not display evidence of algal affinity. Particularly this process-bearing taxon Trachyhystrichosphaera aimika can be locally abundant in subtidal sediments, and its morphology rather suggests osmotrophy, as was proposed for older Mesoproterozoic microfossils (Tappania) (59).
We thank L. Krajewski for FT-ICR MS measurements, including the Taoudeni Basin (19), were largely ferruginous rather than sulfidic (66), and nutrient limitation scenarios are hard to explain other than sulfide poisoning of eukaryotic algae thus appears to be a common attribute of mid-Proterozoic pelagic environments with phototrophic bacteria. Explanations may include sulfide poisoning of eukaryotic algae as well as low atmospheric oxygen levels (68, 69, 72) and unstable redox conditions (73), the inefficient flow of energy from the base of the microbial loop, leading to diminished transport of nutrients to the deep ocean and upwelling zones, and a reduced flux of organic carbon to bottom sediments (57). As permanent burial of reduced carbon is offset by a net release of molecular oxygen into the atmosphere, the sinking behavior of bacterial phytoplankton may thus have contributed to mid-Proterozoic atmospheric oxygen levels significantly lower than at present. Low atmospheric oxygen concentration and the aerobic degradation of bacterial biomass within the water column may have contributed to the pervasively anoxic nature of mid-Proterozoic marine basins (64, 68–70).

In modern oceans, relatively large eukaryotic phytoplankton cells create food webs with efficient nutrient and energy transfers to higher trophic levels, fueling the proliferation of orders-of-magnitude larger grazers and predators (2, 71). By contrast, in mid-Proterozoic pelagic environments with phototrophic bacteria at the base of the food web, a large proportion of carbon and energy may have been lost within the microbial loop. Based on the fossil record (58), such trophic systems may only have sustained relatively small protists at their apex. In combination with low atmospheric oxygen levels (68, 69, 72) and unstable redox conditions (73), the inefficient flow of energy from the base of the mid-Proterozoic food web to higher trophic levels may have inhibited the evolution of complex eukaryotic ecosystems and large, animal-like organisms.

Materials and Methods
Extended methods are available in SI Appendix, SI Methods. Briefly, biomarker analysis was performed at the Australian National University (ANU) under standard operational procedures and included a thorough syngeneity assessment (20–22). Hydrocarbons were analyzed on a Waters AutoSpec Premier double-sector mass spectrometer interfaced with a 6890 GC (Agilent), fitted with a DB-SMS capillary column. Porphyrin determination using reversed-phase HPLC-UV/Vis followed the standard operating procedure, and the porphyrin injection standards of Geoscience Australia (Canberra) (24, 25), and was conducted at the ANU using an Agilent 1100 system containing a degasser (G1379A, JG 40714), a Quat Pump (G1311A, DE 40926343214), and a DAD (G1315B, DE40521369). Molecular characterization of porphyrins was performed at the National High Magnetic Field Laboratories in Tallahassee, Florida (26), using a 9.4-tesla FT-ICR mass spectrometer. For bulk analyses, ~250 μg of total organic matter of decalcified rock powder was weighed into a tin capsule with 4 mg of vanadyl oxide (VO(O)₃, Alf aesar, 99.9%) and the bulk N and C content as well as their isotopic composition were measured using a Sercon 20–22 IRMS connected to an ANCA-GSL CHN elemental analyzer. The average SD measured on samples was 0.06‰.

Compound-specific isotope analyses on Ni-porphyrins were performed at JAM-TEC (Japan) following the methods of Ogawa et al. (74).

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