Archean coastal-plain paleosols and life on land

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A B S T R A C T
Coastal-plain paleosols in the 3.0 Ga Farrel Quartzite of Western Australia have organic surface (A horizon) and sulfate-rich subsurface (By) horizons, like soils of the Atacama Desert of Chile, Dry Valleys of Antarctica, and 3.7 Ga paleosols of Mars. Farrel Quartzite paleosols include previously described microfossils, permineralized by silica in a way comparable with the Devonian Rhyne Chert, a well known permineralized Histosol. Five microfossil morphotypes in the Farrel Quartzite include a variety of spheroidal cells (Archeospaeraoides) as well as distinctive large spindles (new genus provisionally assigned to cf. Eopoikilofusa). Previously published cell-specific carbon isotopic analyses of the Farrel Quartzite microfossils, and unusually abundant sulfate considering a likely anoxic atmosphere, allow interpretation of these morphotypes as a terrestrial community of actinobacteria, purple sulfur bacteria, and methanogenic Archaea.

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1. Introduction
The search for evidence of Archean life on Earth has been unusually successful over the past three decades because of a "basic strategy: look in black (carbon-rich) cherts that are in fine-grained (unmetamorphosed) rocks and associated with Cryptozocon-like stromatolites" (Schopf, 1999). Like stromatolites, the Archean fossil record has been interpreted as marine (Allwood et al., 2006; Flannery and Walter, 2012; House et al., 2013), and rarely lacustrine (Awramik and Buchheim, 2009), although lacustrine is apparently controversial (Sakurai et al., 2005). The variety of putative and likely Archean paleosols are reviewed by Rye and Holland (1998), who assumed that all were sterile to simplify estimates of past atmospheric conditions. Possible microfossils in paleosols have been discounted as remnants of ephemeral ponds (Rye and Holland, 2000). Microbial mats in floodplain and intertidal facies (Homann et al., 2015) also appear more like freshwater aquatic (Flannery and Walter, 2012) than terrestrial crusts (Retallack, 2012). Nevertheless, Archean coastal-plain paleosols may have been hiding in plain sight for several decades as coastal sabkha, playa and salina facies. For example, Buick and Dunlop (1990) concluded that 3.5 Ga evaporites were "almost everywhere laid down in a shallow submarine to intermittently exposed environment". Lowe and Worrell (1999) conclude that 3.4 Ga evaporites "represent sedimentation in marginal hypersaline salinas and low-energy coastal lagoons." These are not only sedimentological facies, but also what soil scientists regard as soils, variously classified as Gypsis (Soil Survey Staff, 2014) or Solonchaks (Food and Agriculture Organization, 1974). Playa and salina soils are inundated less commonly than many floodplains, and their development of desiccation cracks, salt crystals, and shear planes is soil-forming rather than a sedimentary process (Retallack, 2001). Evaporitic paleosols of the Farrel Quartzite (3.0 Ga) were also assigned to "shallow to sub-aerial sedimentary environment" by Sugitani et al. (2003). We confirm the discovery by those authors of diagnostic non-marine features and minerals, including sand crystals (Retallack, 2013, 2014; Natham et al., 2016), and nahcolite (Lowe and Worrell, 1999; Jagniecki et al., 2015). This study of evaporitic coastal-plain paleosols with preserved microfossils provides geochemical and microfossil evidence for Archean terrestrial microbial communities.

The 3.0 Ga Farrel Quartzite is famous for microfossils of at least five distinct morphotypes (Sugitani et al., 2007, 2009, 2011), validated as genuine microfossils by a variety of isotopic (House et al., 2013), maturation (Grey and Sugitani, 2009), chemical and ultrastructural tests (Oehler et al., 2010). The Farrel Quartzite has been mapped though complex structures (Sugitani et al., 2006), and its hydrothermal and metamorphic history elucidated using whole rock and trace element chemistry (Sugahara et al., 2010). Its sedimentary facies and structures have also been interpreted over an outcrop strike of 7 km (Sugitani et al., 2003). Virtually every aspect of the Farrel Quartzite has been studied except its paleosols.

2. Materials and methods
Geological and paleosol sections of the Archean (3.0 Ga) Farrel Quartzite near Mt. Grant in the Pilbara region of Western Australia (Fig. 1) were measured and sampled at centimeter scale using a tape measure, and by noting Munsell color and field reaction with dilute hydrochloric acid (Fig. 2). Both color and calcareousness were altered by

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diagenesis and metamorphism, but these observations are useful for locating the sections. Size, stratigraphic limits, and percentage area occupied by nodules and crystals were also measured in the field. Oriented thin sections prepared vertical to bedding were point-counted using a Swift automated stage and Hacker counting box. Point counting was adjusted for pervasive silicification and recrystallization to crystallites minimally 5 μm across, so that this grain size and smaller was counted as clay. Evaporite minerals were all quartz pseudomorphs, and counted from a distinctive chalcedonic texture unlike simple to undulose extinction of quartz and recrystallized chert. Original evaporite mineralogy could not be determined in thin section, but followed previous identifications based on polished slabs and microprobe analyses of Sugitani et al. (2003). Opaque grains include both kerogen and opaque oxides.

Two counts were made from each thin section: one for grain size and one for mineral composition (See Supplementary Information Tables S1 and S2).

Chemical analyses of selected samples (Supplementary Information Table S3) were performed by ALS Chemex of Vancouver, British Columbia, using XRF on glass disk and FeO by Pratt titration. The standard for the analyses was Canadian Certified Reference Materials Project standard SY-4, a diorite gneiss from near Bancroft, Ontario. Errors (2σ) are from 89 replicate analyses of the standard in the same laboratory. This technique failed for analysis of titania, which was below detection (<0.01 wt.%) for two samples (R4336 and R4337), but this sample level was analyzed previously in Japan (Sugahara et al., 2010, sample GFSV6) by XRF at 0.0053 wt.%. Many chemical compositions were identical to those previously determined for these rocks by Sugitani et al. (2003, 2006), and Sugahara et al. (2010). Cores from the abandoned Mt. Goldsworthy Mines were neither used by their studies nor ours, which are based on the freshest possible samples collected from outcrops.

Bulk densities were determined from 20 to 40 g samples using paraffin (Retallack, 1997a) at the University of Oregon, with errors from 10 replicate density determinations of Western Australian Archean chert (specimen R4309) in order to calculate gains and losses (mass transfer of Brimhall et al., 1992) of elements in a soil at a given horizon (τj,wi in moles) from the bulk density of the soil (ρw in g·cm⁻³) and parent material (ρp in g·cm⁻³) and from the chemical concentration of the element in soils (Cj,wi in wt.%) and parent material (Cj,pi in wt.%).

These data are needed to calculate diagenetic and metamorphic alteration as well as soil formation. Also needed are changes in volume within a single bed during weathering, silicification and metamorphism (strain of Brimhall et al., 1992), estimated from an immobile element in soil (such as Ti used here) compared with parent material (εi,wi as a fraction). Parent material is the rock with sedimentary structures immediately below the profile in which such structures are obscured. The relevant Eqs. (1) and (2) (below) are the basis for calculating divergence from parent material composition.

$$\tau_{j,wi} = \frac{\rho_p \cdot C_{j,wi}}{\rho_p - C_{j,pi}} \left[ \frac{\rho_w - C_{j,wi}}{\rho_w} \right] + 1 - 1$$  \hspace{1cm} (1)

$$\epsilon_{i,wi} = \frac{\rho_p \cdot C_{i,pi}}{\rho_p - C_{i,wi}} - 1$$  \hspace{1cm} (2)

Quantitative counts of abundance of five recognized morphotypes of microfossils recognized by Sugitani et al. (2011) were made from ten photomicrographs 1300 by 890 μm in size at subdivisions of oriented thin sections. These ten counts were then averaged to obtain numbers of each morphotype per square millimeter. Separate counts were made for microfossils within matrix and enclosed within sand-sized clasts (Supplementary Information Tables S4 and S5). Also counted were spherical pyrite grains and clusters of spherules like framboids of microbial origin (Wignall and Newton, 1998). The fossiliferous thin
sections and their source rocks are archived as specimens F118310-118312 in the Condon Collection of the Museum of Natural and Cultural History of the University of Oregon.

Microfossils were examined using a transmission electron microscope (Titan 80-300 with EDX and EELS), and a scanning electron microscope (FEI Helios Dual Beam, Focused Ion Beam Microscope) in the Center for Advanced Materials Characterization at Oregon (CAMCOR) of the University of Oregon.

3. Geological setting

The locality studied at S20.38058° E119.44567° is 1.2 km east of Mt. Grant and 2 km west of Warralong Road, on the crest of the ridge 400 m north of the vehicular track (Fig. 1). Attention focused on detailed sampling of a short geological section (Fig. 2) corresponding to unit CE2 of Sugitani et al. (2007) in the uppermost Farrel Quartzite, immediately below conformably overlying shales of the Cleaverville Formation (Fig. 3). The black cherts at this and nearby localities are well known for well preserved microfossils (Sugitani et al., 2006, 2007, 2009).

4. Geochronology

A geological age of 3016 ± 13 Ma by SHRIMP U–Pb method on zircon has been determined from the youngest population of detrital zircon in Farrel Quartzite beneath Cleaverville Formation at Nuyerry Gap (Nelson, 1998) on the western margin of the East Pilbara Terrane (van Kranendonk, 2010). There is a comparable age of 3058 ± 7 Ma by the same method from another volcanic sandstone near the base of the Cleaverville Formation on the Cleaverville Peninsula near Karratha in the West Pilbara Block (Nelson, 1998).

The age of the Farrel Quartzite and Cleaverville Formation of the Gorge Creek Group in the eastern Pilbara near the Mt. Grant microfossil localities is also bracketed by overlying and underlying rocks (van Kranendonk, 2010). The age of the Lalla Rookh Formation of the unconformably overlying Croydon Group is constrained by a metamorphic age of 2907 ± 7 Ma by 40Ar/39Ar on hornblende from the Coolyia Creek Tonalite (Wijbrans and McDougall, 1987), and intrusion ages of 2936 ± 5 Ma by U–Pb SHRIMP on zircon of the Keep It Dark Monzogranite (van Kranendonk, 2000), 2927 ± 3 Ma by U–Pb SHRIMP on zircon of the Woodstock Monzogranite (van Kranendonk, 2010), and 2928 ± 2 Ma by U–Pb SHRIMP on zircon from the Mulgandinnah...
Monzogranite (van Kranendonk, 2010). The Gorge Creek Group, which includes the Farrel Quartzite, is younger than the Kangaroo Caves Formation of the upper Sulfur Springs Group, which is dated by U–Pb TIMS-multigrain on zircon as 3240 ± 2 Ma from a dacite tuff (van Kranendonk, 2000).

5. Sedimentary paleoenvironments

Paleoenvironments of the microfossil locality near Mt. Grant have been interpreted by sedimentary facies analysis as fluvial and coastal plain (Sugitani et al., 2006, 2007). The base of the Farrel Quartzite is trough cross-bedded sandstone and conglomerates filling paleovalleys of the Sunrise Hill angular unconformity with the Kelly and Warrawoona Groups (Dawes et al., 1995). Evaporite sand crystals of nahcolite and barite, now pseudomorphed by silica, have been identified from microprobe analyses and crystallographic studies, and interpreted as evaporites of coastal salinas (Sugitani et al., 2003). The overlying Cleaverville Formation of laminated shale and banded iron formation is interbedded with pillow basalts of a volcanic island arc near Karratha in the western Pilbara craton (Shibuya et al., 2007), and includes submarine fan facies turbidites in the southern Pilbara block (Eriksson, 1982; Eriksson et al., 1994). The East Pilbara lower Cleaverville Formation and underlying Farrel Quartzite, however, share slightly suprachondritic Y/Ho ratios and slight europium anomalies, characteristic of freshwater, and distinct from modern open ocean values and the marine stromatolitic, 3.4 Ga, Strelley Pool Chert (Sugahara et al., 2010). Two of the seventeen Farrel Quartzite cherts analyzed by Sugahara et al. (2010) have high europium anomalies indicative of local hydrothermal springs. Hydrothermal sources may also explain anomalously low Al₂O₃/TiO₂ ratios in a few samples of the Farrel Quartzite (Sugitani et al., 2006). A lacustrine lower Cleaverville Formation near Mt. Grant is also compatible with its proposed tectonic setting within a backarc basin (Eriksson, 1982) or intracontinental sag (Van Kranendonk et al., 2007).

The source terrane of the Farrel Quartzite included felsic and andesitic volcanics, as well as tonalites, trondhjemites and granite, judging from Th-Sc-La values (Sugitani et al., 2006) and from zircons in the Farrel Quartzite as old as 3.71 ± 0.16 to 3.67 ± 0.16 Ga (Sugitani et al., 2006). These deposits of a shallow sea or lake (Cleaverville Formation) and coastal plain (Farrel Quartzite) accumulated on a protocontinent created by the Prinsep Orogeny, which amalgamated the Karratha and East Pilbara crustal blocks into a landmass about 500 km across (van Kranendonk et al., 2007, 2010). The Kaapvaal craton, some 1000 km across, may have been an additional contiguous landmass judging from alignment of greenstone belts, although palaeomagnetic evidence for this Vaalbara supercontinent is only documented by 2.7 Ga (de Kock et al., 2009).

The bedrock landscape represented by the Sunrise Hill unconformity below the Farrel Quartzite (Dawes et al., 1995), had as much as 1 km of relief, judging from local differences in thickness in the fluvial-coastal Farrel Quartzite as old as 3.71 ± 0.16 to 3.67 ± 0.16 Ga (Sugitani et al., 2006). These deposits of a shallow sea or lake (Cleaverville Formation) and coastal plain (Farrel Quartzite) accumulated on a protocontinent created by the Prinsep Orogeny, which amalgamated the Karratha and East Pilbara crustal blocks into a landmass about 500 km across (van Kranendonk et al., 2007, 2010). The Kaapvaal craton, some 1000 km across, may have been an additional contiguous landmass judging from alignment of greenstone belts, although palaeomagnetic evidence for this Vaalbara supercontinent is only documented by 2.7 Ga (de Kock et al., 2009).

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6. Metamorphic-diagenetic alteration

The Farrel Quartzite is folded and faulted, along with older basement rocks, into near-vertical dips and isoclinal folds typical of many Archean greenstone belts (Fig. 1). Like many outcrops in this region, they are metamorphosed to lower greenschist facies, as indicated by metamorphic sericite, chlorite, epidote and apatite (Sugitani et al., 2003). The Cleaverville Formation in the western Pilbara near Karratha shows three distinct phases of metamorphism not observed in Cleaverville Formation in the East Pilbara region near Mt. Grant (Kiyokawa et al., 2002; Hickman, 2004; Shibuya et al., 2007). The Farrel Quartzite and
Cleaverville Formation also escaped contact metamorphism of the Warrawoona and Kelly Groups during emplacement of the Carlini Granite Complex about 6 km to the south (Smithies et al., 2004), mostly 3490–3225 Ma, but with minor intrusions 2955–2830 Ma in age (van Kranendonk, 2010). Microfossil-bearing cherts near Mt. Grant are less veined and microfaulted than cherts of the Warrawoona and Kelly Groups. There is also evidence for less severe hydrothermal alteration of cherts in the Farrel Quartzite from lack of MREE enrichment or pronounced europium anomalies (Sugahara et al., 2010).

Another measure for post-depositional alteration is burial depth, which for microfossil localities near Mt. Grant was not as high as for other areas of the Farrel Quartzite, because the studied outcrops were on a paleotopographic high between the Archean Mallina Basin to the west and the Paleozoic–Mesozoic Canning Basin to the east (Smithies et al., 2004). The Gorge Creek Group is up to 1900 m thick (van Kranendonk, 2010) south of the former shelf edge (Eriksson, 1982). The Cleaverville Formation near Mt. Grant was no more than 1000 m thick (van Kranendonk et al., 2006). The Farrel Quartzite can reach 1000 m thick to the west, but is only 80 m thick at the investigated microfossil localities (Sugitani et al., 2003). The Lalla Rookh Formation of the overlying Croydon Group was confined to fault basins to the south (Krapež, 1984), and its 3000 m did not overlie the microfossil localities, which were covered by Constantine Sandstone of the Croydon Group no thicker than 1300 m, and a comparable thickness of Mallina Formation (van Kranendonk, 2010). The Fortescue Group in the vicinity of Mt. Grant is represented by metagabbro intrusions (Smithies et al., 2004), and these widespread lavas and sediments covered this area to a thickness of at least 620 m as in Balfour Downs to the east (Thorne and Trendall, 2001). Mesozoic and Cenozoic overburden in the Canning Basin revealed by nearby Chirup Well (Lipski, 1994) reached no more than 900 m. Thus the microfossil localities near Mt. Grant may have been buried by at least 4.8 km of overburden, which amounts to about 1.3 kbar or 0.1 GPa pressure in typical continental crust (Thompson, 1992) as envisaged by van Kranendonk et al. (2007), and would reach greenschist facies only at an elevated geothermal gradient of a backarc basin envisaged by Eriksson (1982).

At such depths of burial enrichment in potassium (smectite illitization) is common (Nesbitt and Young, 1989; Fedo et al., 1995; Novoselov and de Souza Filho, 2015), although illitization is promoted at room temperature and pressure by the dissimilatory metal-reducing bacterium Shewanella (Kim et al., 2004). The ternary diagram of Nesbitt and Young (1989) shows weathering trends in mineralogically mature paleosols but no clear illitization (Fig. 4A). Potash enrichment is apparent from mass transfer with respect to aluminum (Fig. 4B), but is modest compared with other Precambrian paleosols enriched in potassium by as much as 8–12 mol (Novoselov and de Souza Filho, 2015). Two samples (R4336 and R4337) were markedly enriched in silica, but without consistent enrichment of alkalies and alkaline earths (Fig. 4B–F). The other samples did not show such off-scale silicification, which must have occurred during or soon after burial to preserve the microfossils so beautifully (Sugitani et al., 2007, 2011). These paleosols may have been protected from extensive diagenetic and metamorphic chemical alteration by early silicification.

7. Criteria for microfossil biogenicity

Putative Archean microfossils from the very same beds of the Farrel Quartzite sampled here (Fig. 5) passed a variety of critical tests for biogenicity by Sugitani et al. (2007, 2009, 2011, 2013). The microfossils are common in both clasts and matrix of silified sediments (Sugitani et al., 2007, 2011). Abundant microfossils show a limited range of similar morphologies in petrographic thin section (Sugitani et al., 2011), are extractable by palynological techniques (Grey and Sugitani, 2009), and have individually distinct $\delta^{13}C$ compositions (House et al., 2013), C/CN ratios and S concentrations (Oehler et al., 2010). Our additional observations support their interpretation as microfossils and shed light on mechanisms of preservation.

Fig. 4. Geochemical indications of limited illitization (A–B) and silicification (C–F) bias in geochemical composition of Archean paleosols from near Mt. Grant. Two samples of black chert in the upper Jurnpa paleosol form outliers (C–F), but there is no silicification trend in the remaining specimens (G).
7.1. Taphonomic series

A key test of biogenicity for Precambrian fossils is an array of preservation (Hofmann, 2004) from well preserved to partly decayed. The best microfossil preservation in the rocks studied was seen at the top of the bed at 1.8 m in the measured section of Fig. 3, but quality of preservation and abundance of microfossils decreased downward within that bed. The other two beds sampled intensively also showed this relationship of preservation better toward at the top of the bed, but also more abundant framboids toward the base of the bed (Fig. 6).

7.2. Late diagenetic chert neomorphism

Examination of the microfossils under the scanning electron microscope and transmission electron microscope (Fig. 7) demonstrates that cell lumens have been infiltrated by silica in the same way as permineralized wood (Retallack, 2011). The cell walls have been partly broken by recrystallization of silica revealed as domains of uniform grey tone in back-scatter scanning electron microscopy (Fig. 7A–B). These neomorphic domains about 2–5 μm across are apparent as a pixelated effect when examined under a crossed polarizers in a petrographic microscope (Fig. 8A), compared with plane light (Fig. 8B). Such recrystallization can give the misimpression of reticulate and spinose walls, widely recognized in permineralized microfossils (Sergeev et al., 2012). Under transmission electron microscopy, however, the cell walls can be traced across neomorphic crystal boundaries (Fig. 7D). Thus, these fossils are not entirely artifacts of organic matter displacement by crystal growth, as suspected for other microfossils (Brasier et al., 2015).

7.3. Organic matter maturity

Under the high-resolution transmission electron microscope, the organic matter is degraded into ropes 1 nm wide, distinct from particulate
**Fig. 6.** Microfossil distribution, petrography and chemical weathering ratios of Archean paleosols from Western Australia: (a) Sections measured in the field with tape and Munsell Soil Color Chart; (b) Microfossil morphotypes after Sugitani et al. (2011) tallied from 10 photomicrographs from each sampled level: (c) Grain size and mineral content from point counting petrographic thin sections: (d) Molecular weathering ratios based on whole rock XRF analyses (See Supplementary Data for tabulated data).

**Fig. 7.** Microfossil preservation in the type Jurnpa paleosol of the Farrel Quartzite, Western Australia: (A–B) backscatter scanning electron micrographs showing degraded organic spheroids larger than and little displaced by neomorphic quartz (small grey fields) and a large pyrite framboid (white in A) and crystals (white in B); (C) transmission electron micrograph of spindle microfossil (upper) showing ropy degradation of organic matter (above), compared with particulate organic matter in matrix (below); (D) microfossil spindle and filaments little deformed by neomorphic quartz crystals. All from Condon Collection of the Museum of Natural and Cultural History of the University of Oregon specimen F18310B = R4336 from A horizon of Jurnpa clay loam.
organic matter in the cherty matrix (Fig. 7D). Ropes of this size are comparable in diameter with 15 fused aromatic rings, and distinct from crystalline graphite (Derenne et al., 2008).

7.4. Mechanisms of permineralization

Preservation of the Mt. Grant microfossils is comparable with permineralization of fossil plants in the Rhynie Chert of the Early Devonian, Old Red Sandstone, Scotland (Rice et al., 1995), as confirmed by comparison with thin sections of Rhynie Chert (Condon Collection Museum of Natural and Cultural History F10670, 107011, 108385). The Rhynie Chert has been interpreted as a sequence of marsh paleosols at the margins of a volcanic hot spring (Rice et al., 1995). The precursor Devonian peats of the Rhynie Chert can be classified in various soil classifications as Fibrists (Soil Survey Staff, 2014), Dystric Histosols (Food and Agriculture Organization, 1974), and Fibric Organosols (Isbell, 1996). From a geological perspective, the Rhynie chert was a sintered peat, in which cellular permineralization was favored by siliceous hot waters and by exclusion of decomposers and oxygen under waterlogged conditions (Rice et al., 1995). Microbial silicification is also apparent in modern hot springs, but occurs in laminated sinters, unlike the Rhynie Chert and Farrel Quartzite (Konhauser et al., 2001; Kyle et al., 2007). Both Farrel and Rhynie cherts are black with structured organic matter, neomorphically recrystallized, and cut by narrow quartz veins (Fig. 8A–B). In both cases, massive fossiliferous chert is distinct from interbedded, laminated, shale and sandstone (Fig. 9).

Independent evidence for local hot spring activity during accumulation of the Farrel Quartzite comes from two samples with unusually low Al₂O₃/TiO₂ ratios (Sugitani et al., 2006) and unusually high europium anomalies (Sugahara et al., 2010). Comparable silica enrichment of the surface horizon of the bed at 1.8 m in Fig. 3 is especially suggested by its extremely low titania and alumina (Supplementary Information Table S3), so that clastic input was overwhelmed by silica enrichment. (Fig. 4C–F).

Differences are also instructive, especially the abundance and variety of early diagenetic evaporite pseudomorphs (Sugitani et al., 2003) in the Mt. Grant cherts (Fig. 10), which are absent from Rhynie Chert. Thus, silicification of Farrel Quartzite also may be related to formation of playa cherts (Hay, 1968; Eugster, 1969) A combination of volcanic glass precursors, very high pH (>9), and silica-accumulating microbes mobilize silica gels in playa lakes (Eugster, 1969; Larsen, 2008). Sulfate-reducing bacteria, evident from frambooids in the Farrel Quartzite paleosols (Fig. 5F–H), excrete silica spicules and are capable of silicification of evaporites (Birnbaum and Wireman, 1985; Birnbaum et al., 1989). In addition, microorganisms can be preserved in flocculant organic oozes, which persist by desiccation and salt-pickling in dry soda lakes for later permineralization (Birnbaum et al., 1989). Microfossiliferous
Cherts (Fig. 8A–B) can be contrasted with silicified sediments poor in microfossils (Fig. 8C–E). The playa chert model for silicification is appealing for many fossiliferous Precambrian intertidal cherts (Maliva et al., 2005), and is especially plausible for beds of the Farrel Quartzite with evaporite pseudomorphs (Sugitani et al., 2003).

Thus two mechanisms are envisaged for the fine preservation of microfossils (Fig. 8C–E). The playa chert model for silicification is appealing for many fossiliferous Precambrian intertidal cherts (Maliva et al., 2005), and is especially plausible for beds of the Farrel Quartzite with evaporite pseudomorphs (Sugitani et al., 2003).

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8. Criteria for recognizing paleosols

Microfossils of the Farrel Quartzite have been considered marine (House et al., 2013), but their matrix lacks lamination, stromatolites, or soft-sediment deformation (Fig. 9) found in Archean lacustrine and marine rocks (Allwood et al., 2006; Flannery and Walter, 2012). Prior facies analysis (Sugitani et al., 2007), concluded that the Farrel Quartzite was deposited in fluvial to coastal-plain paleoenvironments. Furthermore, the terrestrial playa mineral nahcolite (Na2CO3) has been recognized from crystal pseudomorphs of the Farrel Quartzite (Sugitani et al., 2003).

8.1. Field appearance

A first test of the paleosol hypothesis is field observations of sharp tops and gradational changes downward like a soil profile, and a variety of disruptions that obscure bedding and ripple marks of lower layers like soil structure. Especially soil-like structures are near-vertical cracks joining a system of subhorizontal cracks (cutans), and generally massive un laminated structure (especially evident in thin sections cut vertical to bedding: Fig. 8A–B).

Also diagnostic of paleosols are nodularized and etched pseudomorphs of evaporite minerals with clayey seams and many included grains like pedogenic sand crystals (Fig. 10D), comparable with those widespread in Precambrian paleosols (Retallack, 2013; Nabhan et al., 2016). Replacive evaporite sand crystals are characteristic of aridland soils, whereas marine and lacustrine evaporite crystals are clean of inclusions and displacive (Campbell and Claridge, 1987; Amundson et al., 2012; Retallack, 2012, 2014). Familiar examples of non-marine sand crystals are the “barite roses” of Oklahoma (London, 2008), which can be contrasted with transparent marine gypsum in “chrysanthemum stones” of China (Zhao et al., 1998). The large (5–20 mm) sand crystal pseudomorphs of the Farrel Quartzite are scattered within particular beds, unlike hydrothermal barite or nahcolite, which is confined to veins and microinclusions (Samson et al., 1995; Kamenetsky et al., 2002; Rye, 2005).

Finally, black cherty beds suspected to be paleosols were traceable over 7 km laterally (Sugitani et al., 2007), like soils of an extensive coastal plain (Retallack, 1997, 2013).
8.2. Mineral variation within beds

A second test of the paleosol hypothesis was to quantify mineral composition of the beds by petrographic point-counting to see if there was a loss of weatherable minerals upwards within individual profiles. These counts assumed 5 μm neomorphic recrystallization of permineralizing silica documented by transmission electron microscopy (Fig. 7D). There is marked fragmentation and embayment of rock-fragment grains upwards in the profile, as well as a decline in abundance of silicified feldspar, coordinated with increases in clay rims and matrix (Fig. 6). This pattern is compatible with hydrolytic weathering reactions within a paleosol (Amundson et al., 2012; Retallack, 2012, 2014).

8.3. Geochemical mass balance

A third test for the paleosol hypothesis was to analyze chemical data within a framework of chemical mass balance, or “tau analysis”, to see if within-bed changes fall within the collapse-and-loss quadrant of soils or the dilate-and-gain quadrant of sedimentation (Fig. 11), cementation, silicification, illitization, or ore mineralization (Fig. 4). This analysis normalizes to a stable constituent in parent material (titanium was used in this case), and calculates mole fractions from weight percent and bulk density of each sample using Eqs. (1) and (2) (Brimhall et al., 1992). Most sampled beds qualified as paleosols, with the exception of two specimens of the bed at 1.8 m showing off-scale additions (Figs. 4C–F, 11). This anomaly in the surface of the bed at 1.8 m is also seen in molecular weathering ratios of alumina/bases (Fig. 6). These geochemical differences are too large to be explained by sediment addition, and are best explained by silicification of a sediment-poor, organic surface horizon. Tau analysis also reveals a marked depletion of phosphorus in one of the profiles, with modest change in the other two (Fig. 11). Phosphorus requires organic ligands, and thus biological activity for mobilization (Neaman et al., 2005). Furthermore, the degree of phosphorus depletion is proportional to the degree of soil development judged by destruction of sedimentary structures and size of nodules (Fig. 10D). Phosphorus depletion in the least baritic profiles is most instructive, because barite requires pH low enough to leach phosphorus abiotically (Neaman et al., 2005; Jennings et al., 2015).

8.4. Microfossil distribution within beds

A fourth test of the paleosol hypothesis is the distribution of microfossils, which would be evenly distributed if they lived in accreting microbial mats, but show pronounced horizonation (Fig. 6), as in paleosols (Retallack, 2012). Even in the silicified upper part of the bed at 1.8 m in Fig. 2 they are not aligned between microbial mat laminae (Fig. 5A–C), and in other parts of the profiles they are intergranular (Fig. 5D–E) or entirely within dislodged soil peds or grains (Fig. 5F–H).

Five distinct microfossil morphotypes of carbonaceous microfossils have been recognized by Sugitani et al. (2011) and mostly characterized isotopically (House et al., 2013). These morphotypes (Fig. 5A–E) as well as opaque pyrite framboids (Fig. 5F–H) as possible traces of the activity of sulfur reducing microbes, were counted from photomicrographs of known area to determine abundance of each microfossil morphotype (number·mm$^{-2}$). Microfossils in matrix were counted separately from those entirely within clasts, because of erosion and redeposition of fossiliferous clasts (Sugitani et al., 2007). Microfossils within clasts and clasts are erratic in their distribution within beds, but microfossils in matrix show much greater abundance at the surface of beds (just
below 0 levels in Fig. 6), as expected for microbial earth paleosols (Retallack, 2012).

9. Interpreted paleoenvironments of the paleosols

Each kind of paleosol (Table 1) in the Farrel Quartzite, recognized from the evidence outlined above, offers paleoenvironmental information (Table 2). This is justified in the following paragraphs to generate a coherent model for their paleoenvironment (Fig. 12). Non-genetic pedotype names are from the East Pilbara aboriginal language Nyamal (Burgman, 2007), combined with an equally non-genetic textural term to specify a particular paleosol. The following paragraphs follow the five classical soil forming factors (Retallack, 2001), as well as atmospheric CO₂ and O₂.

9.1. Parent material

Paleosols of the Farrel Quartzite are developed on quartzofeldspathic sand with small amounts of clay and rock fragments (Fig. 6), including volcanics, schist and chert. These clasts are evidence of provenance from continental volcanic arc of felsic and andesitic volcanics, and older continental terranes with chert, metamorphic rocks, tonalites,

Table 1
Summary of Farrel Quartzite paleosol definition and classification.

<table>
<thead>
<tr>
<th>Pedotype</th>
<th>Nyamal (52)</th>
<th>Diagnosis</th>
<th>US taxonomy</th>
<th>FAO map unit</th>
<th>Australian classification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Jurnpa</td>
<td>Cold ashes</td>
<td>Black massive cherty surface (A horizon) over barite mottle pseudomorphs in trough cross-beded sandstone (By horizon)</td>
<td>Ustept</td>
<td>Eutric Cambisol</td>
<td>Grey-orthic Tenosol</td>
</tr>
<tr>
<td>Jurl</td>
<td>Salt</td>
<td>Black massive cherty surface (A horizon) over barite crystal pseudomorphs in grey chert (By horizon)</td>
<td>Cambid</td>
<td>Orthic Solonchak</td>
<td>Grey Sodosol</td>
</tr>
<tr>
<td>Kari</td>
<td>Bitter salt</td>
<td>Black massive cherty surface (A horizon) over barite and nahcolite crystal fan pseudomorphs in grey chert (By horizon)</td>
<td>Cambid</td>
<td>Orthic Solonchak</td>
<td>Grey Sodosol</td>
</tr>
<tr>
<td>Ngumpu</td>
<td>Narrow</td>
<td>Black massive cherty surface (A horizon) over laminated grey chert (C horizon).</td>
<td>Fluvient</td>
<td>Eutric Fluvisol</td>
<td>Stratric Rudosol</td>
</tr>
</tbody>
</table>

![Fig. 11. Mass balance geochemistry of Archean paleosols from Western Australia: (a) deviation of paleosol samples (mole fraction) from parent material composition (at cross hairs) due to changes in volume (strain or epsilon) and changes in amounts of constituents (mass transfer or tau of Brimhall et al., 1992); (b) deviation of amounts of constituents (mass transfer or tau) with depth in paleosols. Most samples fall in paleosol field, but two surface samples of the Jurnpa paleosol show off-scale enrichment attributed to early silicification comparable with that found in the Devonian Rhynie Chert (Rice et al., 1995), thus accounting for extraordinary microfossil preservation in that horizon (Sugitani et al., 2007, 2011).](image-url)
trondhjemites, and large expanses of granite. These petrographic observations of Sugitani et al. (2006) are also supported by their analysis of Th–Sc–La values. In this backarc-intracontinental geological setting (Eriksson, 1982; Van Kranendonk et al., 2007), lack of first-cycle volcanic fragments in the form of shards, scoria and pumice is surprising, but devitrification of volcanic glasses may have contributed to the pervasive silicification and neomorphism of the Farrel Quartzite. Detrital zircons dated at 3016 ± 13 and 3058 ± 7 by SHRIMP U–Pb agree with the interpolated geological age of the Farrel Quartzite (Nelson, 1998), and are evidence of active felsic volcanism.

9.2. Topographic setting

The sedimentary setting of the Farrel Quartzite was a coastal plain (Sugitani et al., 2006, 2007), perhaps beside a lake with later marine influence, now represented by the Cleaverville Formation (Sugahara et al., 2010). There may have been as much as 1 km of relief in hinterlands (Dawes et al., 1995; van Kranendonk et al., 2006), but the paleosols developed on flat landforms, as demonstrated by the Kari pedotype, which is traceable for 7 km along strike (Sugitani et al., 2003).

Table 2
Summary of Farrel Quartzite paleosol interpretations.

<table>
<thead>
<tr>
<th>Pedotype</th>
<th>Climate</th>
<th>Topography</th>
<th>Parent material</th>
<th>Soil duration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Jurnpa</td>
<td>Humid (1112 ± 182 mm MAP) temperate (8.3 ± 0.4 °C MAT)</td>
<td>Microbial earth dominated by methanogens and actinobacteria, with purple sulfur bacteria</td>
<td>Streamside levee Quartz-lithic sand</td>
<td>1000–69,000 years</td>
</tr>
<tr>
<td>Jurl</td>
<td>Humid (1443 ± 182 mm MAP) temperate (11.1 ± 0.4 °C MAT)</td>
<td>Microbial earth dominated by purple sulfur bacteria, with actinobacteria and methanogens</td>
<td>Floodplain Quartz-lithic silt</td>
<td>2000–82,000 years</td>
</tr>
<tr>
<td>Kari</td>
<td>Humid (1319 ± 182 mm MAP) temperate (10.1 ± 0.4 °C MAT)</td>
<td>Microbial earth dominated by purple sulfur bacteria and methenogens, with actinobacteria</td>
<td>Floodplain Quartz-lithic silt</td>
<td>6000–116,000 years</td>
</tr>
<tr>
<td>Ngumpu</td>
<td>Not diagnostic for climate</td>
<td>Microbial earth dominated by methanogens and actinobacteria, with purple sulfur bacteria</td>
<td>Streamside bar Quartz-lithic sand</td>
<td>10–1000 years</td>
</tr>
</tbody>
</table>

Note: MAP is mean annual precipitation and MAT is mean annual temperature.

Table 2
Summary of Farrel Quartzite paleosol interpretations.

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</tbody>
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Note: MAP is mean annual precipitation and MAT is mean annual temperature.

Fig. 12. Reconstructed soils and microbes of the 3.0 Ga Farrel Quartzite, Western Australia.
Permanent water table in this floodplain was at least 25 cm below the surface of the paleosols because above that level evaporate sand crystals and nodules show replacive growth, which is characteristic of soils (Retallack, 2012), as revealed by common sedimentary inclusions (Figs. 8C–D, 10D–E), zones of dissolution (Fig. 10D–E), dusty rims (Fig. 10C), and undeformed bedding planes (Fig. 10E), as in playa lakes (Hay, 1968; Renault and Tiecerier, 1994). In contrast, evaporate crystals precipitated from open water or saturated sediments are free of inclusions because they grow displacively: such crystals are well known from marine evaporates (Warren, 2006; Ziegenbalg et al., 2010), and spring tufas (Donovan et al., 1988; Bonny and Jones, 2008a, 2008b). Barites that formed below water table in sulfur springs also differ from those of the Farrel Quartzite in being aligned with original bedding (Donovan et al., 1988; Bonny and Jones, 2008a, 2008b). Hot spring water is discernible in the surface horizon of the Jurnpa paleosol from its off-scale silica enrichment tau value (Fig. 11) and large europium anomalies (Sugahara et al., 2010). However, these geochemical proxies for hot springs are found only in the two upper samples of that paleosol, and not in the other paleosols. Floodplains of the Farrel Quartzite may have been dotted with spring mounds and artesian blisters (Fig. 12), like those on the shore of Lake Bogoria, Kenya (Ashley et al., 2002; Owen et al., 2008). Barite spring waters and their feeder dikes have not yet been found in the Farrel Quartzite, but are documented for the 3.5 Ga Dresser Formation of Western Australia (van Kranendonk and Pirajno, 2004; van Kranendonk et al., 2008).

9.3. Time for soil development

Paleosols of the Farrel Quartzite range from moderately developed for the Jurl and Kari pedotypes to weakly developed for the Jurnpa pedotype and very weakly developed for the Ngumpu pedotype, in a field scale devised for modern soils (Retallack, 2001). A critical field observation is the relative destruction of bedding, which is difficult to trace in moderately developed paleosols because laminations and cross bedding have been disrupted by soil cracking and settling (Fig. 3D–E). These field observations have been confirmed by thin sections cut perpendicular to bedding to show massive fabrics in the paleosols (Fig. 9A–B), compared with laminated fabrics above and below (Fig. 9D–E), and mixed massive-bedded fabrics in between (Fig. 9C).

A more important index of soil development is the formation of discrete nodules or sand crystals, which define moderate development, and take many millennia to form in modern soils (Retallack, 2001). For example, low magnesium calcite nodules in radiocarbon-dated soils of New Mexico grow in diameter (in cm) with soil age (A in kyrs) according to the following equation (Retallack, 2005, $R^2 = 0.57$; S.E. ± 1.8 kyrs).

$$A = 3.92S^{0.34}$$

(3)

Gypsum sand crystals in the Negev and Atacama Deserts increase in abundance (G in area %) with soil age (A in kyrs) according to another equation (Retallack, 2013, $R^2 = 0.95$, S.E. ± 15 kyrs).

$$A = 3.98G + 5.774$$

(4)

If the sand crystals and nodules of Farrel Quartzite paleosols were calcite and had original diameters as now measured in the field (Fig. 2), Eq. (3) would yield $3.6 \pm 1.8$ kyrs for Jurl, $7.9 \pm 1.8$ kyrs for upper Kari, $8.3 \pm 1.8$ kyrs for lower Kari, and $3.3 \pm 1.8$ kyrs for Jurnpa paleosols. If the crystals and nodules were gypsum in volume percent averaged from point counting (Fig. 10), Eq. (4) would yield $67.2 \pm 15$ kyrs for Jurl, $101.5 \pm 15$ kyrs for upper Kari, $80.7 \pm 15$ kyrs for lower Kari, and $44.0 \pm 15$ kyrs for Jurnpa paleosols. These estimates are orders of magnitude different, but provide broad constraints. They are not strictly relevant because pedogenic nodules and crystals in Farrel Quartzite are silica pseudomorphs of barite and naphcoite (Sugitani et al., 2003). Because naphcoite requires much higher than modern CO₂ partial pressures (Eugster, 1966; Jagpieck et al., 2015) and barite nodules require anoxic sulfur bacterial biominalizers (Jennings et al., 2015), there is little hope for obtaining chrononfunctions for formation of either mineral from Holocene soils (Retallack and Kirby, 2007; Jennings et al., 2015). Although quantitative estimates for duration of soil formation in the Archean remain elusive, many millenia were needed for Jurl, Kari and Jurnpa paleosols, and proportionally less for Ngumpu paleosols.

9.4. Paleoclimate

Most pedogenic paleothermometers are based on modern forest to desert shrubland soils (Sheldon et al., 2002; Gallagher and Sheldon, 2013), but a pedogenic paleothermometer based on modern soils under tundra of Iceland (Öskarsson et al., 2012) is more appropriate for Archean paleosols. This linear regression between mean annual temperature (T in °C) and chemical index of weathering (I = $100 \frac{mAl_{2}O_{3}}{mAl_{2}O_{3} + mCaO + mNa_{2}O}$), in molar proportions is given in Eq. (3) ($R^2 = 0.81; \text{S.E.} = \pm 0.4 ^\circ \text{C}$).

$$T = 0.211 - 8.93$$

(5)

These calculations yield temperate mean annual paleotemperatures for the lower A horizons of Jurl (11.1 ± 0.4 °C), Jurnpa (8.3 ± 0.4 °C) and Kari (10.1 ± 0.4 °C) paleosols of the Farrel Quartzite.

These results are comparable with application of this paleothermometer to the thick (6 m) Jericho Dam paleosol below the 3.0 Ga Pongola Supergroup in South Africa (Grandstaff et al., 1986), which may have been within the same paleoclimatic belt as the 3.0 Ga Farrel Quartzite (de Kock et al., 2009). The Jericho Dam sample at 1.7 m depth yielded mean annual temperature of 11.1 ± 0.4 °C for chemical index of weathering (Öskarsson et al., 2012). These results are also consistent with “temperate paleoclimate” (~40 °C inferred for the 3.4 Ga Buck Reef Chert of South Africa), using combined oxygen and hydrogen isotopic evidence (Hren et al., 2009), and revising earlier estimates of 55–85 °C from oxygen isotopic evidence throughout the Archean (Knaath and Lowe, 2003). Temperatures less than 75 °C are also indicated by 16–36 volume % original quartz in the Farrel Quartzite (Fig. 6), because quartz would be prone to dissolution at higher temperature and pH (Sleep and Hessler, 2006). This line of inference is inappropriate for little-weathered sediment of tectonically active terranes (Lowe, 2007), but facies and grain size of the Farrel Quartzite are evidence of subdued tectonic activity (Eriksson, 1982; Sugitani et al., 2003, 2006). A widely used paleohyetometer (Sheldon et al., 2002) based on temperate soils of North America uses chemical index of alteration without potash ($R = 100 \frac{mAl_{2}O_{3}}{(mAl_{2}O_{3} + mCaO + mNa_{2}O)}$), which increases with mean annual precipitation (P in mm) in modern soils ($R_2 = 0.72$; S.E. = ± 182 mm), as follows.

$$P = 221e^{0.0197R}$$

(6)

Eq. (6) is based on the hydrolysis equation of weathering, which enriches alumina at the expense of lime, magnesia, potash and soda. Magnesia is ignored because it is insignificant for most clastic sedimentary rocks (Maynard, 1992), and potash is excluded because it can be enriched during deep burial alteration of sediments (Novoselov and de Souza Filho, 2015). This paleohyetometer is not effective for little-weathered gyspic soils, but works for modern baritic soils and paleosols because they are deeply weathered (Jennings and Driese, 2014; Jennings et al., 2015). This chemical index of alteration paleohyetometer yields humid mean annual precipitation for lower A horizons of Jurl (1443 mm ± 182), Jurnpa (1112 ± 182) and Kari (1319 mm ± 182) paleosols of the Farrel Quartzite.
These data support other indications of deep weathering such as light rare earth enrichment of the Farrel Quartzite and its cherts (Sugahara et al., 2010). The 3.0 Ga Jericho Dam paleosol of South Africa is also deeply weathered (6 m) and indicative of a humid climate (Grandstaff et al., 1986): mean annual precipitation of 1452 mm can be calculated from a sample at 1.7 m using the chemical index of alteration without potash equation (Sheldon et al., 2002).

Evaporites, such as gypsum, which is widespread in desert soils (Retallack and Huang, 2010) are potential evidence against humid paleoclimate. The original composition of Archean evaporites (Fig. 9) has been addressed by crystallographic work on interfacial angles of pseudomorphs in the 3.0 Ga Farrel Quartzite (Sugitani et al., 2003), as well as in the 3.2 Ga Moodies Group (Nabhan et al., 2016), the 3.4 Ga Kromberg Formation (Lowe and Worrell, 1999), the 3.4 Ga Strelley Pool Formation (Lowe, 1983; Allwood et al., 2007), the 3.5 Ga Apex Chert (Pinti et al., 2009) and the 3.5 Ga Dresser Formation (Lambert et al., 1978; Buick and Dunlop, 1990; Runnegar et al., 2001) to suggest that they were barite (BaSO₄), selenite-gypsum (CaSO₄·2H₂O), and nahcolite (NaHCO₃). Evidence that sand crystals and nodules in the Farrel Quartzite were barite comes from glide twinning, and 63–64 wt% BaO in relic cores of some pseudomorphs (Sugitani et al., 2003). Evidence against the former presence of gypsum in the Farrel Quartzite includes the lack of observed selenite fshitall twinning, and absence of pseudomorphs of equant-orthorhombic anhydrite (CaSO₄), which forms readily from elongate-monoclinic gypsum during shallow burial (Klimchouk, 1996). Equant cubes and rhombs in the Farrel Quartzite are iron-stained, and presumed to have been pyrite and siderite, respectively (Sugitani et al., 2003).

Archean silicified barite and nahcolite crystals have been attributed to hydrothermal vein crystallization (Runnegar et al., 2001; Pinti et al., 2009), or to precipitation from saline water (Lambert et al., 1978; Lowe, 1983; Buick and Dunlop, 1990; Lowe and Worrell, 1999; Sugitani et al., 2003; Allwood et al., 2006; Shen et al., 2009). These alternatives may be reasonable for other cases, but are not favored for the Farrel Quartzite because its crystal pseudomorphs have common inclinations (Fig. 8C–E), zones of dissolution (Fig. 8D–E), and dusty rims (Fig. 8C), resembling other Archean pedogenic evaporates (Nabhan et al., 2016). Dissolution planes, birdseye textures, entrolithic folds and nodules are additional field evidence of alternating desiccation and wetting (Sugitani et al., 2003). Nahcolite and barite are widespread as crystals, spherulites and nodules in sodic soils (Kohut and Durdas, 1993; Shahid and Jenkins, 1994; Howari et al., 2002; Jennings and Driese, 2014) and playa lake crusts (Hay, 1968; Renaut and Tiecelin, 1994). Nahcolite also forms from the weathering of natrocarbonatite (Bradley, 1973; Surdam and Stanley, 1979; Larsen, 2008; Garcia-Veigas et al., 2013). In modern soils, nahcolite and barite crystals and spherulites have been attributed to pedogenic precipitation, or by redox changes at the water table, rather than by pedocline (Retallack and Kirby, 2007; Jennings et al., 2015). The growth of barite is favored by acid sulfate weathering at low pH (<3), but that of gypsum at higher pH (4–9: Carson et al., 1982). There is as yet no clear relationship between the depth of barite or nahcolite growth, and mean annual precipitation as there is for gypsum (Retallack and Huang, 2010). Thus it is puzzling that crystals of both nahcolite and barite are confined to a specific horizon (By) within the paleosols, and morphologically comparable with gypsum horizons in paleosols of Earth (Retallack and Huang, 2010) and Mars (Retallack, 2014). The growth of salts may have been suppressed in thin (5–10 cm) uppermost horizons of Archean and Martian paleosols by surficial moisture and carbonic acid from biological productivity, as in desert soils of Earth (Retallack, 2009).

9.5. Atmospheric CO₂

High levels of carbon dioxide are critical for the precipitation and maintenance of nahcolite (Eugster, 1966). It is a rare mineral in surficial environments today because current atmospheric levels of only 408 ppm CO₂ make it prone to recrystallization to trona ([Na₃(CO₃)(HCO₃)·2H₂O] or natron (Na₂CO₃·10H₂O). As for other greenhouse crises (Lowenstein and Demicco, 2006), nahcolite crystal pseudomorphs in the Kari paleosol may be evidence of at least 945 ppm CO₂ in the atmosphere at temperatures of 10–11 °C (Jagniecki et al., 2015).

Much higher Archean CO₂ levels (100–1000 times present atmospheric level of 30,000 to 300,000 ppm) have been inferred by Lowe and Tice (2004) from nahcolite formation in South Africa at assumed Archean temperatures of 70 °C, but such high temperatures are unlikely given more recent oxygen and hydrogen isotopic studies (Hren et al., 2009), preservation of quartz (Sleep and Hesseler, 2006), and application of paleosol paleothermometers used here (Öskarsson et al., 2012). A minimum level of 2500 ppm CO₂ has been inferred from 3.2 Ga ferrous-carbonate weathering rinds (Hesseler et al., 2004), but such fluvial pebbles may have been isolated from the atmosphere by groundwater. Estimates of about 3000 ppm CO₂ comes from iron mineral stability in banded iron formations (Lowe and Tice, 2004; Ohmoto et al., 2004), which also reflect aquatic rather than atmospheric levels. A maximum level of 36,000 ppm CO₂ has been inferred from paleosols ranging in age from 2.75–2.2 Ga because of a presumed lack of siderite in paleosols (Rye et al., 1995). This estimate suffers from choice of meta-morphic rather than pedogenic mineral thermodynamic data (Sheldon, 2006), and there are rhombic pseudomorphs in Farrel Quartzite that may have been siderite (Sugitani et al., 2003). An estimate of 1500 to 9000 ppm atmospheric CO₂ at 3.0 Ga comes from chemical weathering of the Jericho Dam paleosol of South Africa (Grandstaff et al., 1986). This much CO₂ would result in acid rain (pH 4.0–4.5 according to Ohmoto et al., 2004), and soil-microbial CO₂ and H₂SO₄ could drop soil water pH to 3, and thus favor the abundant barite precipitation observed (Carson et al., 1982).

Even the most extreme of these estimates is short of the amount needed for a greenhouse capable of maintaining temperate Archean paleotemperatures given the faint young sun, and other greenhouse gases are required, including water vapor, ammonia, sulfur oxides, and methane (Ohmoto et al., 2004; Rosing et al., 2010; Kasting and Kirschvink, 2012). Much methane would have come from methanogenesis inferred from isotopic values of small spheroids (House et al., 2013) in Farrel Quartzite paleosols and allowed by low (160 ppm) H₂ values likely for the Archean (Kasting and Kirschvink, 2012). An atmosphere with three times the current mass of N₂ and a H₂ mixing ratio of 0.1, would also have created an adequate greenhouse (Wordsworth and Pierrehumbert, 2013), but N₂ in the atmosphere was limited to 1.1 to 0.5 bars, judging from nitrogen and argon isotopic ratios in fluid inclusions of the 3.5 Ga Dresser Formation of Western Australia (Marty et al., 2013). Atmospheric pressure is constrained to no more than twice the modern value by 2.7 Ga raindrop imprints (Som et al., 2012) and no more than half the modern value by 2.7 Ga vesicle sizes in lava flows (Som et al., 2016). The volcanicogenic greenhouse gas CO₂
proposed on theoretical grounds for early Mars (Haley et al., 2007) and early Earth (Claire et al., 2014) is compatible with the abundance of sulfate in paleosols of the Farrel Quartzite.

9.6. Atmospheric O₂

Very low levels of atmospheric oxygen are evident from paleosols of the Farrel Quartzite because of their high ferrous to ferric iron ratios, comparable with swamp soils today exhausted of oxygen by microbial respiration (Retallack, 2001). However, the Farrel Quartzite paleosols were not waterlogged, but exposed to air as indicated by dissolution planes, birdseye textures, entrolledithic folds and nodules of evaporitic mineral pseudomorphs (Sugitani et al., 2003). No highly oxidized minerals such as hematite or goethite were observed in fresh samples of the Farrel Quartzite paleosols.

A reasonable estimate of 0.002 to 0.1% atmospheric O₂ for 3.0 Ga is derived from chemical weathering of the Jerico Dam paleosol of South Africa and associated uraniferous conglomerates (Grandstaff et al., 1986). The Jerico Dam paleosol has been considered altered by hydrothermal fluids or by burial geization of organic acids after formation in an oxidizing atmosphere (Öhmo, 1996) of at least 0.3% O₂. However, there is no clear hydrothermal enrichment of heavy rare earth elements in the Jerico Dam profile (Kimberly and Grandstaff, 1986). Furthermore, burial geization in paleosols of Triassic forest ecosystems (which were much more productive than envisaged for the Archean) extends to a depth of only 30 cm (Retallack, 1997), an order of magnitude less than the 3.3 m required to explain the deep spike in ferrous iron within the Jerico Dam paleosol (Grandstaff et al., 1986; Kimberly and Grandstaff, 1986). These objections do not apply to the Nsuze paleosols of South Africa, geologically younger at 2.9 Ga, and with a lack of oxidative Cr and U recycling suggestive of 0.006 to 0.001% O₂ in the atmosphere (Crowe et al., 2013). Comparable estimates of 0.002 to 0.02% O₂ come from crater anomalies, despite potash were not waterlogged, but exposed to air as indicated by dissolution planes, birdseye textures, entrolledithic folds and nodules of evaporitic mineral pseudomorphs (Sugitani et al., 2003). No highly oxidized minerals such as hematite or goethite were observed in fresh samples of the Farrel Quartzite paleosols.

9.7. Microbiota

Microfossils in paleosols of the Farrel Quartzite have been studied by numerous methods but not taxonomically (Sugitani et al., 2007; Grey and Sugitani, 2009; Oehler et al., 2010; House et al., 2013). This account evaluates the five recognized morphotypes recognized by Sugitani et al. (2011), both from the perspectives of paleobotanical form taxonomy and of biological affinities (Table 3). This is not the full taxonomic study these microfossils deserve, but a provisional study using only comparable fossil taxa already described.

The Farrel Quartzite microflora has some similarities with microfossils described from a locality 4 km west of old Panorama Homestead in the East Pilbara region (Awramik et al., 1983), which includes Archaeaphaeroides pilbarensis, resembling the Farrel Quartzite diffuse-wall spheroids, and Archaeotrichon contortum, resembling Farrel Quartzite filaments. The Panorama Homestead locality is currently mapped within the 3.5 Ga Mt. Ada Basalt (Van Kranendonk, 2006; Hickman, 2013), but the exact location and geological setting of these microfossils is uncertain (Awramik et al., 1983). Some specimens of A. pilbarensis have walls comparable with the folded-wall spheroids of Sugitani et al. (2011), and their other two morphotypes with simple walls and thick walls fit within the same generic, if not specific, concept. The small spheroids in the Farrel Quartzite are comparable with the form species Archaeaphaeroides barbertonensis known from the Fig Tree Group near Barberton, South Africa (Schofield and Barghoorn, 1967), and assigned to the 3.2 Ga Haki Ironstone Member of the Sheba Formation (Lowe and Byerly, 1999).

Spindle-shaped microfossils in the Farrel Quartzite are distinctive, with a bulging cavity containing small internal bodies and a lateral rib that extends to a point at either end (Sugitani et al., 2007, 2009). Comparable spindle-shaped microfossils also have been reported from the 3.4 Ga Strelley Pool Formation of Western Australia (Sugitani et al., 2013), the 3.4 Ga Kromberg Formation of South Africa (Walsh, 2010), the 3.2 Ga Clutha Formation of South Africa (Javaux et al., 2010), and the 1.3 Ga Kendall River Formation of Arctic Canada (Horodyski and Donaldson, 1980; Kah et al., 2006). These spindles deserve their own new form-generic name based on the best studied material of Sugitani et al. (2011), but have been compared with the acritarch genera *Pterospermopsis*, *Pterospermella*, and *Pterospemormiphora* by Sugitani et al. (2009). All three acritarch taxa are double-walled, near spherical with narrow flanges, and have either no projections (Timofeev, 1966), or more than two blunt-ending projections (Eisenack, 1972; Yankauskas, 1980; Samuelsson et al., 1999), unlike the acritarch with simple-walled spindles of the Farrel Quartzite. The Farrel Quartzite spindle-shaped microfossils are commonly clustered with their ends continuous with filaments that appear to join them together, and, in addition, have small internal bodies (Sugitani et al., 2007, 2009). The form genus *Primatifella* of the Ediacaran Kotlin Formation of Russia shares these characters, but has only one end tapering to filaments while the other end is blunt (Gnilovskaya, 1983). The Farrel Quartzite spindles are generally similar to *Euopikofilus claudii* from the Cambrian Gouhou and Xiamaling Formations of China (Xing, 1987; Yin, 1987), and to *Euopikofilus* sp. from the 1.2 Ga Society Cliffs Formation of Baffin Island (Hofmann and Jackson, 1991). Canadian Arctic permineralizations show a similar curvature as the Farrel Quartzite spindles and some have filaments attached at one end. *Euopikofilus* is the most similar valid form genus to the Farrel Quartzite microfossil spindles.

The Farrel Quartzite also contains numerous perfectly spherical opaque grains of pyrite, which are not microfossils, but likely framboïds, and may represent results of microbial sulfur reduction (Sawlowicz, 1993). Archean biological sulfate reduction and atmospheric sources of sulfur come from fractionations of sulfur isotopes in pyrite associated with barite of the 3.5 Ga Dresser Formation of Western Australia (Ueno et al., 2008; Shen et al., 2009). Like microfossils, framboïds were found in both matrix and clasts, but in some cases the framboïds are so regularly arranged just inside the grain edge (Fig. 5F–G) that they may mark cavities occupied by endolithic organisms (McLoughlin et al., 2009).

There also are rare sinuously deformed fragments of microbial mats up to 1 mm long in the Farrel Quartzite (Sugitani et al., 2007, Fig. 10; Sugitani et al., 2011, Fig. 7A–C). For these fragments, the ichnogenus *Eoclatrhus* is suitable, although that genus is based on dimpled impressions rather than permineralizations, and has been regarded as problematic (Hänzschel, 1975). These mat fragments have small internal bodies (Sugitani et al., 2011), and their other two morphotypes with simple walls and thick walls may represent results of microbial sulfur reduction (Sawlowicz, 1993). Archean biological sulfate reduction and atmospheric sources of sulfur come from fractionations of sulfur isotopes in pyrite associated with barite of the 3.5 Ga Dresser Formation of Western Australia (Ueno et al., 2008; Shen et al., 2009). Like microfossils, framboïds were found in both matrix and clasts, but in some cases the framboïds are so regularly arranged just inside the grain edge (Fig. 5F–G) that they may mark cavities occupied by endolithic organisms (McLoughlin et al., 2009).
feature distinguishing them from microbial earths, in which microbes are mixed with unlaminated grains (Retallack, 2012), as envisaged for paleosols of the Farrel Quartzite.

Biological affinities of the Farrel Quartzite microfossils are inferred partly from their morphology and partly from their carbon isotopic composition (Table 3; House et al., 2013). Unusually light carbon isotope composition (−33.8 to −44.2 \( \Delta^{13}C_{org} \)) of small spheroids comparable with \( A. \) barbertonensis are comparable with methanotrophic Proteobacteria and methanogenic Archaea (Schidlowski, 2001). \( A. \) barbertonensis is larger than most methanotrophs, although Archaea such as \( M. \) barkeri have 3 to 4 μm diameter and spherical walled colonies 10 μm in diameter (Sowers et al., 1993). Methanogenesis is also evident from isotopic composition of methane in fluid inclusions from the 3.5 Ga Dresser Formation of Western Australia (Unno et al., 2006).

The large spheroids comparable with \( A. \) pilbarensis also have a distinctive carbon isotope composition (−34.3 to −38.4 \( \Delta^{13}C_{org} \)) comparable with that of purple sulfur bacteria (Chromatiaceae), which is unlike that of other bacteria including Cyanoarchaea, Chloroflexi or Chlororhodaceae (Schidlowski, 2001). Purple sulfur bacteria have an array of morphologies and characteristic arrays (Fjerdingstad, 1979), but modern forms comparable with the Farrel simple wall spheroids are \( C. \) obovatum (Fjerdingstad, 1979), while diffuse wall spheroids superficially resemble \( T. \) mangrovi (Kumar et al., 2008), thick-walled spheroids are like \( R. \) nuda (Fjerdingstad, 1979) and folded-wall spheroids are like \( C. \) densusgranulatum (Fjerdingstad, 1979).

\( E. \) fusiformis microfossil spindles have been regarded as marine plankton because of their wide geographic distribution, "flanged lenticular shape, occasional internal vacuole-like structures, and the possible association with small non-clustered spheroids" (House et al., 2013). A planktonic life style, however, is unlikely considering two non-hydrodynamic features: their connection by filaments into bunches, and their asymmetric curve and lateral ridge like a calzone (Fig. 5C–E). Internal hollows and associated spheroids are more likely spores or other propagules, than internal or remnant vacuoles. Farrel Quartzite filaments have a narrow range of isotopic composition because only three were analyzed (−34.4 to −36.3 \( \Delta^{13}C_{org} \)), but eight spindles from this and other Pilbara Archean cherts (House et al., 2013; Lepot et al., 2013) had a wide spread of values (−35.6 to −40.5 \( \Delta^{13}C_{org} \)), straddling values of both purple sulfur bacteria and methanogens as would be expected of heterotrophic bacteria (Schidlowski, 2001). Attachment of spindles of \( E. \) fusiformis to Archaeotrichon threads (Sugitani et al., 2007) is comparable with \( P. \) flagellata (Gnilovskaya, 1983), and as in that case, can be interpreted as sporangia of actinobacteria. Comparable smooth sporangia with apical spines and attached to the ends of hyphae are found in living \( S. \) roseum (Nolan et al., 2010). If cf. \( E. \) fusiformis from the Farrel Quartzite were actinobacteria, then they are much older than the hitherto earliest known actinobacteria of the 2.2 Ga Gowganda Formation of Canada (Jackson, 1967). Nevertheless, actinobacteria on land would explain the organic-lean nature of paleosols, including the 3.0 Ga Jericho Dam paleosol of South Africa (Grandidge et al., 1986), and a 2.7 ± 0.3 Ma molecular divergence of \( A. \) barbertonensis (Battistuzzi and Hedges, 2008).

Sulfur bacteria (Chromatiaceae and Chlororhodaceae; Youssef et al., 2010) play a major role in precipitation of barite in modern cold sulfur springs within 12 m of the source, as demonstrated by diel fluctuations in aqueous sulfate and incubations, and stable isotopic comparisons from source to sink at Zodletone Spring, Oklahoma (Senko et al., 2004). Soil around Zodletone Spring is black in contact with the water, but grey with alunogen (Al₂(SO₄)₃·17H₂O) and white with gypsum (CaSO₄·2H₂O) above the water (Donovan et al., 1988). Zodletone barite tufa shows a mix of groundwater features, such as cavity filling ponded and erect fans, and soil features, such as such as 6-mm-diameter barite roses with growth chevrons, suggestive of fluctuating water levels. Less similar to Archean paleosol barite is the barite tufa of Flyeby Springs in the Canadian Northwest Territories (Bonny and Jones, 2008a) and of Jasper Cold Sulphur Springs Alberta (Bonny and Jones, 2008b). Both modern springs are more heavily vegetated than Zodletone Spring and their barite tufas are distinctly laminated or stromatolitic with submillimetre barite as isopachous coatings and replacements of microbes, along with small crystals and spherulites. Crystals and nodules are also typical of barite in modern soils (Kohut and Durda, 1993; Shahid and Jenkins, 1994; Howari et al., 2002; Jennings and Drise, 2014), in which both sulfur-oxidizing and sulfur-reducing bacteria are limited in distribution by atmospheric oxygen. Methanogens and methanotrophs also are largely limited to oxygen-poor aquatic environments, such as the guts of ruminants, swamp soils and rice paddy (Conrad, 2007). In the anoxic atmosphere of the Archean, however, aquatic methanotrophs and sulfur bacteria would not have been excluded from the intermittently wet pores of soils that were mostly well drained.

Distinctive microfossil morphology associated with sulfate pseudomorphs in the Farrel Quartzite may be evidence of differences between microbiota of land and sea during the Archean. Archean marine assemblages are characterized by common seafloor filaments (Primin gulivillum; Awramik et al., 1983; Schopf et al., 2007), although some of these may be carbon-displacing vermicular phyllosilicate (Brasier et al., 2015; Wacey et al., 2016). In contrast, likely non-marine assemblages described here are characterized by microfossil spindles (cf. \( E. \) fusiformis; Fig. 5C–E), interpreted as actinobacteria. The many spheroids of \( A. \) pilbarensis in the Farrel Quartzite paleosols (Fig. 5A–B) are

<table>
<thead>
<tr>
<th>Morphotype</th>
<th>Pre cambrian microfossil form taxonomy</th>
<th>Carbon isotopic composition ( (\Delta^{13}C_{org}) )</th>
<th>Plausible taxonomic affiliations</th>
<th>Inferred metabolism</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thread</td>
<td>Archaeotrichon contortum</td>
<td>−35.2 ± 1.0, n = 3</td>
<td>Actinobacteria</td>
<td>Anaerobic fermentation</td>
</tr>
<tr>
<td>Simple wall spheroid</td>
<td>Archaeosphaeroides sp.</td>
<td>−38.7, n = 1</td>
<td>Chromatiaceae (purple sulfur bacteria)</td>
<td>Anaerobic photosynthesis</td>
</tr>
<tr>
<td>(10–60 μm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diffuse wall spheroid</td>
<td>Archaeosphaeroides pilbarensis</td>
<td>−36.5 ± 2.3, n = 4</td>
<td>Chromatiaceae (purple sulfur bacteria)</td>
<td>Anaerobic photosynthesis</td>
</tr>
<tr>
<td>(20–40 μm)</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Thick wall spheroid</td>
<td>Archaeosphaeroides sp.</td>
<td>−34.1 ± 0.1, n = 2</td>
<td>Chromatiaceae (purple sulfur bacteria)</td>
<td>Anaerobic photosynthesis</td>
</tr>
<tr>
<td>(30–40 μm)</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Folded wall spheroid</td>
<td>Archaeosphaeroides sp.</td>
<td>−37.9 ± 3.3, n = 18</td>
<td>Methanosarcinaeae (Archeae)</td>
<td>Methanogenesis</td>
</tr>
<tr>
<td>(20–40 μm)</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Small (&lt;15 μm) spheroid</td>
<td>Archaeosphaeroides barbertonensis</td>
<td>−33.8 to −44.2, n = 13</td>
<td>Mixed microbial mat fragment</td>
<td>Mixed anaerobic microbial mat</td>
</tr>
<tr>
<td>Thick laminated film</td>
<td>“Eoactibactra” sp.</td>
<td>−31.5 ± 0.4, n = 13</td>
<td>Actinobacteria</td>
<td>Anaerobic Fermentation</td>
</tr>
<tr>
<td>Spindle (10–90 μm)</td>
<td>cf. Eupokilofusa sp.</td>
<td>−36.9 ± 1.7, n = 8</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 3: Microfossils of the 3.0 Ga Farrel Quartzite.
interpreted here as photosynthetic sulfur bacteria responsible for the accumulation of barite in the profiles and fueling the activity of sulfate-reducing bacteria represented by frambooids (Fig. 5F–H). This confirms geochemical modeling that envisages evolution of the sulfur cycle on land before it extended to the sea (Stüeken et al., 2012). Archean sulfur isotopic evidence has been used to infer <2.5 μM marine sulfate, which is <0.01% of modern marine sulfate (Crowe et al., 2014; Zhelezinskaia et al., 2014), but microbial sulfate reduction is evident from shallow-marine to lagoonal Archean carbonates (Paris et al., 2014). Sulfate may have been limited to only the shallowest parts of the Archean ocean, but the paleosols described here and by Nabhan et al. (2016) are evidence of widespread Archean sulfate on land.

10. Conclusions

Paleosols of the Farrel Quartzite provide a first quantitative appraisal of Archean microbial earth communities (Retallack, 2012). Morphology and isotopic composition allow educated guesses on the nature of the various microfossil morphotypes. The distinctive microfossil spindles (Fig. 5C–E), provisionally referred to cf. *Eupokilolesus* (Hofmann and Jackson, 1991), are attached to and closely associated with fine filaments comparable with the hyphalike microfossil genus *Archeotrichon* (Awramik et al., 1983). The spindles are hollow structures which have internal bodies (Sugitan et al., 2007), like spores of actinobacterial sporangia (Battistuzzi and Hedges, 2009; Nolan et al., 2010). Both cf. *Eupokilolesus* and *Archeotrichon* have a wide range of isotopic composition (House et al., 2013), perhaps reflecting a varied diet. Archean paleosols of Mt. Grant also include a variety of spherical cells generally similar to the microfossil genus *Archeasphaeroides* (Awramik et al., 1983). Small (5–15 μm diameter) spindles in clusters (Fig. 2A–B), have a distinctive light isotopic composition (−33.8 to −44.2 ε^13C) of methanogens (Schidlowski, 2001; House et al., 2013). Methanogenic bacteria and archaea are mostly smaller than the fossils, but Archaea such as *M. barkeri* have coccoids up to 4 μm in diameter and spherical walled colonies 10 μm in diameter (Sowers et al., 1993). Large (10–60 μm diameter) spindles with heavier isotopic composition (−34.3 to −38.4 ε^13C) than photosynthetic bacteria (Cyanobacteria), green sulfur bacteria (Chlorobiaceae) or green non-sulfur bacteria (Chlororoflexaceae), are isotopically (Schidlowski, 2001) comparable with purple sulfur bacteria (Chromatiaceae).

Such biological affinities support other indications of anaerobic soil forming conditions. High ferrous/ferric metallic weathering ratios (Fig. 6) and iron depletion (Fig. 11) from the paleosols is comparable with that of swamp soils today (Retallack, 2001). These paleosols were not waterlogged because they are dominated by evaporite sand–calcium walled colonies 10 μm in diameter (Sowers et al., 1993). Large (10–60 μm diameter) spindles with heavier isotopic composition (−34.3 to −38.4 ε^13C) than photosynthetic bacteria (Cyanobacteria), green sulfur bacteria (Chlorobiaceae) or green non-sulfur bacteria (Chlororoflexaceae), are isotopically (Schidlowski, 2001) comparable with purple sulfur bacteria (Chromatiaceae).

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Appendix A. Supplementary data

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References


