

This is an Open Access document downloaded from ORCA, Cardiff University's institutional repository: <https://orca.cardiff.ac.uk/id/eprint/179339/>

This is the author's version of a work that was submitted to / accepted for publication.

Citation for final published version:

Song, Yafang, Mills, Benjamin J. W., Bowyer, Fred T., Andersen, Morten B. , Ossa Ossa, Frantz, Dickson, Alexander J., Harvey, Jason, Zhang, Shuichang, Wang, Xiaomei, Wang, Huajian, Canfield, Donald E., Shields, Graham A. and Poulton, Simon W. 2025. Tracking the spatial extent of redox variability in the mid-Proterozoic ocean. *Geology* 10.1130/G53447.1

Publishers page: <https://doi.org/10.1130/G53447.1>

Please note:

Changes made as a result of publishing processes such as copy-editing, formatting and page numbers may not be reflected in this version. For the definitive version of this publication, please refer to the published source. You are advised to consult the publisher's version if you wish to cite this paper.

This version is being made available in accordance with publisher policies. See <http://orca.cf.ac.uk/policies.html> for usage policies. Copyright and moral rights for publications made available in ORCA are retained by the copyright holders.



# Tracking the spatial extent of redox variability in the mid-Proterozoic ocean

**Yafang Song<sup>1,2\*</sup>, Benjamin J. W. Mills<sup>1</sup>, Fred T. Bowyer<sup>1</sup>, Morten B. Andersen<sup>3</sup>, Frantz Ossa Ossa<sup>3,4</sup>, Alexander J. Dickson<sup>5</sup>, Jason Harvey<sup>1</sup>, Shuichang Zhang<sup>6</sup>, Xiaomei Wang<sup>6</sup>, Huajian Wang<sup>6</sup>, Donald E. Canfield<sup>6,7</sup>, Graham A. Shields<sup>8</sup> and Simon W. Poulton<sup>1,9</sup>**

*<sup>1</sup>School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK*

*<sup>2</sup>Deep Space Exploration Laboratory/State Key Laboratory of Lithospheric and Environmental Coevolution, University of Science and Technology of China, Hefei 230026, China*

*<sup>3</sup>School of Earth and Environmental Sciences, Cardiff University, Cardiff CF10 3AT, UK*

*<sup>4</sup>Department of Earth Sciences, Khalifa University of Science and Technology, Abu Dhabi 127788, United Arab Emirates*

*<sup>5</sup>Centre of Climate, Ocean and Atmosphere, Department of Earth Sciences, Royal Holloway University of London, Egham TW20 0EX, UK*

*<sup>6</sup>Key Laboratory of Petroleum Geochemistry, Research Institute of Petroleum Exploration and Development, China National Petroleum Corporation, Beijing 100083, China*

*<sup>7</sup>Nordcee, Department of Biology, University of Southern Denmark, Odense 5230, Denmark*

*<sup>8</sup>Department of Earth Sciences, University College London, London WC1E 6BT, UK*

*<sup>9</sup>International Research Frontiers Initiative, Earth-Life Science Institute, Tokyo Institute of Technology, Tokyo 152-8550, Japan*

\*Corresponding author: Yafang Song ([yf.song@ustc.edu.cn](mailto:yf.song@ustc.edu.cn))

## 23 ABSTRACT

24 Emerging geochemical evidence suggests considerable redox heterogeneity in the mid-  
25 Proterozoic ocean. However, quantitative estimates of the extent of different modes of anoxia remain  
26 poorly constrained. Due to their complementary redox-related behaviour, uranium and molybdenum  
27 isotopes can be combined to reconstruct ancient marine redox landscapes, but this approach has not  
28 been applied to the mid-Proterozoic. Here, we present new  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  data for marine rocks  
29 from the  $\sim 1.4$  Ga Xiamaling Formation, North China Craton, together with independent redox  
30 indicators (Fe speciation and redox-sensitive trace metals). We find that most samples deposited  
31 under oxic or dysoxic conditions retain low U and Mo contents, with  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  values  
32 indistinguishable from continental crust, demonstrating a dominant detrital signal. By contrast,  
33 euxinic samples with authigenic enrichments in U and Mo record the highest authigenic  $\delta^{238}\text{U}$  and  
34  $\delta^{98}\text{Mo}$  values, consistent with efficient reduction of U and Mo. Samples deposited under ferruginous  
35 conditions exhibit a wider range of intermediate  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  values that generally fall between  
36 the (dys)oxic and euxinic end-members. Using a coupled U-Mo isotope mass balance model, we infer  
37 limited euxinia ( $<0.5\%$  of the global seafloor area) but extensive low-productivity (dys)oxic and  
38 ferruginous settings in  $\sim 1.4$  Ga oceans. This redox landscape would have provided potentially  
39 habitable environments for eukaryotic evolution in the mid-Proterozoic.

40

## 41 INTRODUCTION

42 Reconstructing the oxygenation history of Earth's surface environment is crucial to understand  
43 the trajectory of Earth's habitability. The mid-Proterozoic (1.8–0.8 Ga) was a critical interval for  
44 early eukaryote evolution (Knoll and Nowak, 2017), and while emerging evidence suggests that

45 biological innovation at this time occurred under heterogeneous ocean redox conditions (Sperling et  
46 al., 2014; Zhang et al., 2016; Luo et al., 2021; Song et al., 2023), the global extent of different modes  
47 of anoxia remains poorly constrained. As such, a quantitative assessment of the global redox  
48 landscape may provide critical insight into the spatial extent of habitable conditions, thus ultimately  
49 enabling improved consideration of potential controls on early eukaryote evolution.

50 Uranium and molybdenum isotopes ( $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$ ) are useful tools for reconstructing global  
51 redox conditions because of their redox sensitivity and long oceanic residence times (Andersen et al.,  
52 2017; Kendall et al., 2017). The largest U isotope fractionations occur during U reduction in anoxic  
53 environments, with heavy U isotopes preferentially sequestered in sediments (Weyer et al., 2008).  
54 The largest Mo isotope fractionations occur during adsorption to Mn-Fe (oxyhydr)oxides under oxic  
55 conditions (Kendall et al., 2017). By contrast, minimal isotopic difference is expected between Mo  
56 in sediments and coeval seawater during near-quantitative drawdown under strongly euxinic  
57 conditions (Nägler et al., 2011). Since rapid Mo burial specifically requires relatively high dissolved  
58 sulfide concentrations, but U burial only requires anoxia, a particularly robust reconstruction of the  
59 extent of different ocean redox conditions can be achieved when considering  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$   
60 together (Andersen et al., 2020; Kendall et al., 2020). Although there are several studies using either  
61  $\delta^{238}\text{U}$  or  $\delta^{98}\text{Mo}$  to investigate ocean redox variability in the mid-Proterozoic (Arnold et al., 2004;  
62 Yang et al., 2017; Gilleaudeau et al., 2019; Luo et al., 2021), there have been no studies of U-Mo  
63 isotope co-variation during this period.

64 Here, we present new U and Mo isotope data for drill core samples from the ~1.4 Ga Xiamaling  
65 Formation, North China Craton. In combination with new Re concentration data, which enables  
66 specific identification of dysoxic conditions (Crusius et al., 1996; Song et al., 2023; Li et al., 2025),

67 and existing Fe speciation,  $\delta^{98}\text{Mo}$  and U-Mo concentration data, we first explore  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$   
68 systematics in the context of the local redox state. We then utilize an isotope mass balance model to  
69 reconstruct the spatial extent of different redox conditions in the  $\sim 1.4$  Ga ocean.

70

## 71 **GEOLOGIC SETTING**

72 The  $\sim 1.4$  Ga Xiamaling Formation (Zhang et al., 2015) study site is located in the Xiahuayuan  
73 region of Hebei Province, north China (Fig. 1A), where the formation can be divided into six units  
74 (Wang et al., 2017; see Supplemental Material for detailed geologic background). Here, we focus on  
75 fresh drill core material from the upper four units (Fig. 1B), which are dominantly composed of low  
76 thermal maturity mudstones and black shales, representing deep-water deposition (Wang et al., 2017).

77

## 78 **RESULTS**

79 In total, 50 samples were analyzed for  $\delta^{238}\text{U}$  compositions, while an additional 15 samples were  
80 analysed to augment existing  $\delta^{98}\text{Mo}$  data (Zhang et al., 2019; see Supplemental Material for methods  
81 and data). Bulk  $\delta^{238}\text{U}$  values show relatively constant values through unit 4 ( $-0.22 \pm 0.04\text{‰}$ , 2SD;  
82 Fig. 2). An excursion to lower values (as low as  $-0.41\text{‰}$ ) and then to higher values (up to  $0.08\text{‰}$ )  
83 occurs in unit 3, followed by a progressive increase to values approaching  $0.2\text{‰}$  at the top of unit 2  
84 (Fig. 2). Samples in unit 1 show more scatter, but with an initial drop to lower values, followed by a  
85 general increase up-section (Fig. 2). Authigenic  $\delta^{238}\text{U}$  ( $\delta^{238}\text{U}_{\text{auth}}$ ) compositions (see Supplemental  
86 Material for detrital corrections; note that samples with low U and Mo contents were excluded from  
87 authigenic correction) range from  $-0.44\text{‰}$  to  $+0.26\text{‰}$ , and exhibit a similar trend to bulk  $\delta^{238}\text{U}$  (Fig.  
88 3). Our  $\delta^{98}\text{Mo}$  data range from  $0.15\text{‰}$  to  $1.26\text{‰}$ , with negligible difference relative to authigenic

89  $\delta^{98}\text{Mo}$  ( $\delta^{98}\text{Mo}_{\text{auth}}$ ) (Fig. 2; Table S1). In almost all cases, Re enrichment factors ( $\text{Re}_{\text{EF}}$ ; see  
90 Supplemental Material for the enrichment factor (EF) calculation) are above 1 (Fig. 2).

91

## 92 **DISCUSSION**

### 93 **Local Redox conditions**

94 Previous detailed reconstructions have invoked a generally anoxic, but dynamic redox setting  
95 for the Xiamaling Formation (Zhang et al., 2016; Wang et al., 2017; Song et al., 2023). Based on  
96 mostly low but variable highly reactive Fe to total Fe ( $\text{Fe}_{\text{HR}}/\text{Fe}_{\text{T}}$ ) ratios and  $\text{U}_{\text{EF}}\text{-Mo}_{\text{EF}}$  values, coupled  
97 with generally low pyritization ( $\text{Fe}_{\text{py}}$ ) of the  $\text{Fe}_{\text{HR}}$  pool (Fig. 2), units 3 and 4 have been interpreted to  
98 record orbital-scale variability in the spatial extent of a ferruginous oxygen minimum zone (OMZ),  
99 which became more productive through unit 3 (Wang et al., 2017; Song et al., 2023). More expansive  
100 deeper water anoxia developed in unit 2, with a progression from ferruginous to euxinic conditions,  
101 while unit 1 documents continued euxinia, punctuated by more oxygenated conditions in the upper  
102 half of the unit (Fig. 2; Wang et al., 2017).

103 Persistent enrichments in Re (Fig. 2), including for samples with low  $\text{Fe}_{\text{HR}}/\text{Fe}_{\text{T}}$  ratios, further  
104 suggest that deeper waters below the OMZ in units 3 and 4 were likely dysoxic (Crusius et al., 1996;  
105 Song et al., 2023; Li et al., 2025), although intervals of fully oxygenated conditions (Zhang et al.,  
106 2016; Wang et al., 2017) cannot be discounted (hence we refer to such samples as being (dys)oxic).

### 107 **Controls on U and Mo Isotope Compositions**

108 To explore  $\delta^{98}\text{Mo}$ - $\delta^{238}\text{U}$  variability, we plot  $\delta^{98}\text{Mo}$  and  $\delta^{238}\text{U}$  profiles in the context of local  
109 redox conditions (Fig. 2). The (dys)oxic samples have  $\delta^{238}\text{U}$  values that are essentially  
110 indistinguishable from average continental crust (-0.3‰), consistent with generally low U contents

111 and a dominant detrital contribution (Fig. 2; Fig. S1) (Andersen et al., 2017). Ferruginous samples  
112 have  $\delta^{238}\text{U}_{\text{auth}}$  values that span a relatively wide range (Fig. 2), consistent with a previous  $\delta^{238}\text{U}$  study  
113 of ferruginous lakes (Cole et al., 2020). In more detail, ferruginous samples with low TOC and  $\text{U}_{\text{EF}}$   
114 values (termed F2 samples) tend to have  $\delta^{238}\text{U}$  similar to the detrital composition, whereas higher  
115  $\delta^{238}\text{U}_{\text{auth}}$  values are observed for samples with higher TOC and  $\text{U}_{\text{EF}}$  (F1 samples) (Fig. 2). Euxinic  
116 samples are generally enriched in authigenic U and record the highest  $\delta^{238}\text{U}_{\text{auth}}$  values (Fig. 2; see  
117 Supplemental Material for discussion of two euxinic samples with anomalously low  $\delta^{238}\text{U}_{\text{auth}}$ ).

118 Integrating our new  $\delta^{98}\text{Mo}$  analyses with previously published data (Zhang et al., 2019) shows  
119 that (dys)oxic and F2 samples are dominantly characterized by low  $\delta^{98}\text{Mo}$  values ( $0.34 \pm 0.14\text{‰}$ ),  
120 close to continental crust ( $\sim 0.3\text{‰}$ ) (Voegelin et al., 2014), while the highest  $\delta^{98}\text{Mo}$  values are  
121 observed for euxinic and F1 sediments (Fig. 2). It is noteworthy that euxinic samples have  $\delta^{98}\text{Mo}$   
122 values spanning a wide range, possibly suggesting variable, and relatively low, water column sulfide  
123 levels (Neubert et al, 2008). In addition, sulfidic pore waters may also have exerted an important  
124 control on the observed  $\delta^{98}\text{Mo}$  variability (Kendall et al., 2017).

125 While  $\delta^{238}\text{U}$  systematics are commonly used to evaluate global ocean redox changes, the  
126 potential impact of local redox conditions and organic carbon loading on fractionations recorded in  
127 the sediments needs to be considered (Lau et al., 2022; Rutledge et al., 2024). Although bulk  $\delta^{238}\text{U}$   
128 values show a general positive correlation with TOC (when OMZ samples are excluded) and U  
129 contents,  $\delta^{238}\text{U}_{\text{auth}}$  values display no such correlation (Fig. 3), suggesting a negligible local  
130 environmental control on  $\delta^{238}\text{U}_{\text{auth}}$  fractionations. We also note that our  $\delta^{238}\text{U}_{\text{auth}}$  values are generally  
131 comparable to shale  $\delta^{238}\text{U}_{\text{auth}}$  data ( $-0.27\text{‰}$  to  $0.16\text{‰}$ ) from the  $\sim 1.36$  Ga Velkerri Formation,

132 northern Australia (Yang et al., 2017), suggesting a consistent isotopic offset from global seawater  
133 and relatively stable seawater  $\delta^{238}\text{U}$ .

134 Neither  $\delta^{98}\text{Mo}$  or  $\delta^{98}\text{Mo}_{\text{auth}}$  compositions show a systematic covariation with TOC or Mo  
135 contents (Fig. 3), therefore local changes in productivity or sedimentation rate appear to have exerted  
136 minimal impact on the observed  $\delta^{98}\text{Mo}$  variability. Diamond et al. (2018) and Zhang et al. (2019)  
137 reported two distinct  $\delta^{98}\text{Mo}$  maxima for euxinic sediments of the Xiamaling Formation at two  
138 different localities, which likely suggests heterogenous sulfide availability in the relatively low-  
139 sulfate mid-Proterozoic ocean (Fakhraee et al., 2019). Nevertheless, a combined evaluation of  $\delta^{238}\text{U}$ -  
140  $\delta^{98}\text{Mo}$  can be used to estimate global ocean redox variability (Andersen et al. 2020; Kendall et al.,  
141 2020).

142

### 143 **Reconstructing Global Redox Conditions**

144 Generally, fluctuating  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  values in the Xiamaling Formation are consistent with  
145 local redox dynamics, and their ranges across different redox conditions are relatively constant  
146 throughout the formation (Fig. 2), suggesting relatively stable oceanic U-Mo inventories and isotopic  
147 compositions. To quantitatively evaluate the global oceanic redox distribution at this time, we employ  
148 a stochastic isotope mass balance model, assuming that both U and Mo are dominantly sourced from  
149 rivers and are buried under (dys)oxic, ferruginous and euxinic conditions. In this model, U-Mo burial  
150 is a function of the areal proportion of each redox sink, and seawater  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  are calculated  
151 from U-Mo burial and corresponding isotope fractionations for each redox sink, utilising specific  
152 fractionation factors from previous studies (Table S2; see Supplemental Material for detailed  
153 modelling approach). By running the model 10,000 times from a modern-day initialization through a



154 random selection of ferruginous and euxinic areal fractions, model outputs (i.e., U and Mo isotope  
155 compositions for each redox sink) are produced that fit both the  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  data for the  
156 Xiamaling Formation (Fig. 4). An advantage of our model is that specific isotope compositions for  
157  $\sim 1.4$  Ga seawater are not required, as the model calculates coeval seawater isotope compositions in  
158 association with randomly selected isotope compositions and fractionation factors for each sink.

159 The resultant area that intersects all coloured areas (area b in Fig. 4) suggests that, at most,  $\sim 0.5\%$   
160 of the global seafloor was overlain by euxinic waters at  $\sim 1.4$  Ga, while the areal fraction of  
161 ferruginous conditions was at least 20%, leaving the rest of the seafloor in a (dys)oxic state. Our  
162 estimate of more expansive ferruginous conditions is consistent with ferruginous-dominated redox  
163 models proposed for the mid-Proterozoic (Poulton et al., 2010; Planavsky et al., 2011; Poulton and  
164 Canfield, 2011). By contrast, our estimate for the extent of euxinia at  $\sim 1.4$  Ga is relatively small  
165 compared to estimates for other intervals of the mid-Proterozoic ( $< 10\%$ ), which were constrained by  
166 either  $\delta^{238}\text{U}$  or  $\delta^{98}\text{Mo}$  (Gilleaudeau et al., 2019; Luo et al., 2021). This may reflect either temporal  
167 variability in water column euxinia, or model limitations when considering one isotope system in  
168 isolation.

169 Recent studies have suggested increased ocean oxygenation at  $\sim 1.4$  Ga, relating to enhanced  
170 nutrient-driven primary productivity (Cox et al., 2016). However, our results suggest that large  
171 expanses of the ocean were more likely dysoxic than oxic, and as such, elevated productivity was  
172 likely restricted to local settings experiencing increased nutrient availability, driven either by locally-  
173 enhanced nutrient influxes or recycling under euxinic conditions (Song et al., 2023). The limited  
174 extent of euxinia in  $\sim 1.4$  Ga oceans would have diminished the specific ‘toxicity’ control of sulfide  
175 on eukaryote evolution (Anbar and Knoll, 2002), which together with the expansive extent of at least

176 mildly oxygenated oceans (Heard et al., 2023), may have exerted a significant control on the evolution  
177 of the biosphere in the mid-Proterozoic.

178

## 179 CONCLUSIONS

180 Uranium and Mo isotope systematics in the ~1.4 Ga Xiamaling Formation provide constraints  
181 on global ocean redox evolution in the mid-Proterozoic. Our  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  data exhibit distinct  
182 ranges related to different redox conditions, with (dys)oxic samples having  $\delta^{238}\text{U}$ - $\delta^{98}\text{Mo}$  values  
183 indistinguishable from the detrital input. By contrast, the highest  $\delta^{238}\text{U}_{\text{auth}}$ - $\delta^{98}\text{Mo}_{\text{auth}}$  values are  
184 preferentially recorded in euxinic samples. Ferruginous sediments with low TOC tend to have low  
185  $\delta^{238}\text{U}$ - $\delta^{98}\text{Mo}$  values, while higher-TOC ferruginous samples tend to have intermediate to high  $\delta^{238}\text{U}$ -  
186  $\delta^{98}\text{Mo}$ . Coupled U-Mo modelling results suggest that eutrophic euxinic settings covered ~0.5% of the  
187 global seafloor at most, while less-productive ferruginous and (dys)oxic settings were much more  
188 extensive at ~1.4 Ga. This redox partitioning may account for the moderate diversification of  
189 eukaryotes observed in the Xiamaling Formation. Similar studies linking mid-Proterozoic isotopic  
190 and fossil records will help to determine whether the Xiamaling Formation documents a relatively  
191 static mid-Proterozoic redox environment, or one stage in a temporal progression towards increased,  
192 stable ecological niche space that promoted enhanced eukaryotic diversification.

193

## 194 ACKNOWLEDGMENTS

195 This work was funded by a China Scholarship Council-University of Leeds Scholarship to YS, NERC  
196 grant NE/R010129/1 to SWP, BJWM and GAS, NERC grant NE/T008458/01 to SWP and FTB, and  
197 NERC grant NE/V004824/1 to MBA and FOO. SWP was supported in part by the World Research

198 Hub (WRH) Program of the International Research Frontiers Initiative, Tokyo Institute of  
199 Technology.

200

## 201 REFERENCES CITED

- 202 Anbar, A. D., and Knoll, A. H., 2002, Proterozoic ocean chemistry and evolution: a bioinorganic  
203 bridge?: *Science*, v. 297, p. 1137-1142, <https://doi.org/10.1126/science.1069651>
- 204 Andersen, M.B., Matthews, A., Bar-Matthews, M. and Vance, D., 2020. Rapid onset of ocean  
205 anoxia shown by high U and low Mo isotope compositions of sapropel S1: *Geochemical*  
206 *Perspectives Letters*, 15, p. 10-14, <https://doi.org/10.7185/geochemlet.2027>.
- 207 Andersen, M.B., Stirling, C.H., and Weyer, S., 2017, Uranium isotope fractionation: Reviews in  
208 Mineralogy and Geochemistry, v. 82, p. 799-850, <https://doi.org/10.2138/rmg.2017.82.19>.
- 209 Arnold, G.L., Anbar, A.D., Barling, J., and Lyons, T.W., 2004, Molybdenum isotope evidence for  
210 widespread anoxia in mid-Proterozoic oceans: *Science*, v. 304, p. 87-90,  
211 <https://doi.org/10.1126/science.109178>.
- 212 Cole, D.B., Planavsky, N.J., Longley, M., Böning, P., Wilkes, D., Wang, X., Swanner, E.D.,  
213 Wittkop, C., Loydell, D.K., Busigny, V., and Knudsen, A.C., 2020, Uranium isotope  
214 fractionation in non-sulfidic anoxic settings and the global uranium isotope mass  
215 balance: *Global Biogeochemical Cycles*, v. 34, e2020GB006649,  
216 <https://doi.org/10.1029/2020GB006649>.
- 217 Cox, G. M., Jarrett, A., Edwards, D., Crockford, P. W., Halverson, G. P., Collins, A. S., Poirier, A.,  
218 and Li, Z. X., 2016, Basin redox and primary productivity within the Mesoproterozoic Roper  
219 Seaway: *Chemical Geology*, v. 440, p. 101-114,  
220 <https://doi.org/10.1016/j.chemgeo.2016.06.025>.
- 221 Crusius, J., Calvert, S., Pedersen, T., and Sage, D., 1996, Rhenium and molybdenum enrichments in  
222 sediments as indicators of oxic, suboxic and sulfidic conditions of deposition: *Earth and*  
223 *Planetary Science Letters*, v. 145, p. 65-78, [https://doi.org/10.1016/S0012-821X\(96\)00204-X](https://doi.org/10.1016/S0012-821X(96)00204-X).
- 224 Diamond, C.W., Planavsky, N.J., Wang, C. and Lyons, T.W., 2018, What the ~ 1.4 Ga Xiamaling  
225 Formation can and cannot tell us about the mid-Proterozoic ocean. *Geobiology*, v. 16, p. 219-  
226 236, <https://doi.org/10.1111/gbi.12282>.
- 227 Fakhraee, M., Hancisse, O., Canfield, D.E., Crowe, S. A., and Katsev, S., 2019, Proterozoic  
228 seawater sulfate scarcity and the evolution of ocean–atmosphere chemistry: *Nature Geoscience*,  
229 v. 12, p. 375-380, <https://doi.org/10.1038/s41561-019-0351-5>.
- 230 Gilleaudeau, G.J., Romaniello, S.J., Luo, G., Kaufman, A.J., Zhang, F., Klabe, R.M., Kah, L.C.,  
231 Azmy, K., Bartley, J.K., Zheng, W., and Knoll, A.H., 2019, Uranium isotope evidence for  
232 limited euxinia in mid-Proterozoic oceans: *Earth and Planetary Science Letters*, v. 521, p. 150-  
233 157, <https://doi.org/10.1016/j.epsl.2019.06.012>.
- 234 Heard, A.W., Wang, Y., Ostrander, C.M., Auro, M., Canfield, D.E., Zhang, S., Wang, H., Wang,  
235 X., and Nielsen, S.G., 2023, Coupled vanadium and thallium isotope constraints on  
236 Mesoproterozoic ocean oxygenation around 1.38-1.39 Ga. *Earth and Planetary Science*  
237 *Letters*, v. 610, p. 118127, <https://doi.org/10.1016/j.epsl.2023.118127>.

238 Kah, L.C., Lyons, T.W., and Frank, T.D., 2004, Low marine sulphate and protracted oxygenation of  
 239 the Proterozoic biosphere: *Nature*, v. 431, p. 834-838, <https://doi.org/10.1038/nature02974>.

240 Kendall, B., Dahl, T.W., and Anbar, A.D., 2017, The stable isotope geochemistry of molybdenum:  
 241 *Reviews in Mineralogy and Geochemistry*, v. 82, p. 683-732,  
 242 <https://doi.org/10.2138/rmg.2017.82.16>.

243 Kendall, B., Wang, J., Zheng, W., Romaniello, S.J., Over, D.J., Bennett, Y., Xing, L., Kunert, A.,  
 244 Boyes, C., and Liu, J., 2020, Inverse correlation between the molybdenum and uranium  
 245 isotope compositions of Upper Devonian black shales caused by changes in local depositional  
 246 conditions rather than global ocean redox variations. *Geochimica et Cosmochimica Acta*, v.  
 247 287, p. 141-164, <https://doi.org/10.1016/j.gca.2020.01.026>.

248 Knoll, A.H., and Nowak, M.A., 2017, The timetable of evolution: *Science advances*, v. 3,  
 249 e1603076, <https://doi.org/10.1126/sciadv.1603076>.

250 Lau, K.V., Hancock, L.G., Severmann, S., Kuzminov, A., Cole, D.B., Behl, R.J., Planavsky, N.J.,  
 251 and Lyons, T.W., 2022, Variable local basin hydrography and productivity control the uranium  
 252 isotope paleoredox proxy in anoxic black shales: *Geochimica et Cosmochimica Acta*, v. 317,  
 253 p. 433-456, <https://doi.org/10.1016/j.gca.2021.10.011>.

254 Li, S., Wignall, P.B., and Poulton, S.W., 2025, Co-application of rhenium, vanadium, uranium and  
 255 molybdenum as paleo-redox proxies: Insight from modern and ancient environments:  
 256 *Chemical Geology*, v. 674, 10.1016, <https://doi.org/10.1016/j.chemgeo.2024.122565>.

257 Luo, J., Long, X., Bowyer, F.T., Mills, B.J., Li, J., Xiong, Y., Zhu, X., Zhang, K., and Poulton,  
 258 S.W., 2021, Pulsed oxygenation events drove progressive oxygenation of the early  
 259 Mesoproterozoic ocean: *Earth and Planetary Science Letters*, v. 559, 116754,  
 260 <https://doi.org/10.1016/j.epsl.2021.116754>.

261 McLennan, S.M., 2001, Relationships between the trace element composition of sedimentary rocks  
 262 and upper continental crust: *Geochemistry, Geophysics, Geosystems*, v. 2, 2000GC000109,  
 263 <https://doi.org/10.1029/2000GC000109>.

264 Nägler, T. F., Neubert, N., Böttcher, M. E., Dellwig, O., and Schnetger, B., 2011, Molybdenum  
 265 isotope fractionation in pelagic euxinia: Evidence from the modern Black and Baltic  
 266 Seas: *Chemical Geology*, v. 289, p. 1-11, <https://doi.org/10.1016/j.chemgeo.2011.07.001>.

267 Neubert, N., Nägler, T.F., and Böttcher, M.E., 2008, Sulfidity controls molybdenum isotope  
 268 fractionation into euxinic sediments: Evidence from the modern Black Sea: *Geology*, v. 36, p.  
 269 775-778, <https://doi.org/10.1130/G24959A.1>.

270 Planavsky, N.J., McGoldrick, P., Scott, C.T., Li, C., Reinhard, C.T., Kelly, A.E., Chu, X., Bekker,  
 271 A., Love, G.D., and Lyons, T.W., 2011, Widespread iron-rich conditions in the mid-  
 272 Proterozoic ocean: *Nature*, v. 477, p. 448-451, <https://doi.org/10.1038/nature10327>.

273 Poulton, S.W., Fralick, P.W., and Canfield, D.E., 2010, Spatial variability in oceanic redox structure  
 274 1.8 billion years ago: *Nature Geoscience*, v. 3, p. 486-490, <https://doi.org/10.1038/ngeo889>.

275 Poulton, S.W., and Canfield, D.E., 2011, Ferruginous conditions: a dominant feature of the ocean  
 276 through Earth's history: *Elements*, v. 7, p. 107-112, <https://doi.org/10.2113/gselements.7.2.107>.

277 Rutledge, R.L., Gilleaudeau, G.J., Remírez, M.N., Kaufman, A.J., Lyons, T.W., Bates, S., and  
 278 Algeo, T.J., 2024, Productivity and organic carbon loading control uranium isotope behavior in  
 279 ancient reducing settings: Implications for the paleoredox proxy: *Geochimica et*  
 280 *Cosmochimica Acta*, v. 368, p. 197-213, <https://doi.org/10.1016/j.gca.2024.01.007>.

281 Song, Y., Bowyer, F.T., Mills, B.J., Merdith, A.S., Wignall, P.B., Peakall, J., Zhang, S., Wang, X.,  
 282 Wang, H., Canfield, D.E., and Shields, G.A., 2023, Dynamic redox and nutrient cycling  
 283 response to climate forcing in the Mesoproterozoic ocean: *Nature Communications*, v. 14, p.  
 284 6640, <https://doi.org/10.1038/s41467-023-41901-7>.  
 285 Sperling, E.A., Rooney, A.D., Hays, L., Sergeev, V.N., Vorob'Eva, N.G., Sergeeva, N.D., Selby,  
 286 D., Johnston, D.T., and Knoll, A.H., 2014, Redox heterogeneity of subsurface waters in the  
 287 Mesoproterozoic ocean: *Geobiology*, v. 12, p. 373-386, <https://doi.org/10.1111/gbi.12091>.  
 288 Voegelin, A.R., Pettke, T., Greber, N.D., von Niederhäusern, B., and Nägler, T.F., 2014, Magma  
 289 differentiation fractionates Mo isotope ratios: evidence from the Kos Plateau Tuff (Aegean  
 290 Arc): *Lithos*, v. 190, p. 440-448, <https://doi.org/10.1016/j.lithos.2013.12.016>.  
 291 Wang, X., Zhang, S., Wang, H., Bjerrum, C.J., Hammarlund, E.U., Haxen, E.R., Su, J., Wang, Y.,  
 292 and Canfield, D.E., 2017, Oxygen, climate and the chemical evolution of a 1400 million year  
 293 old tropical marine setting: *American Journal of Science*, v. 317, p. 861-900,  
 294 <https://doi.org/10.2475/08.2017.01>.  
 295 Weyer, S., Anbar, A.D., Gerdes, A., Gordon, G.W., Algeo, T.J., and Boyle, E.A., 2008, Natural  
 296 fractionation of <sup>238</sup>U/<sup>235</sup>U: *Geochimica et Cosmochimica Acta*, v. 72, p. 345-359,  
 297 <https://doi.org/10.1016/j.gca.2007.11.012>.  
 298 Yang, S., Kendall, B., Lu, X., Zhang, F., and Zheng, W., 2017, Uranium isotope compositions of  
 299 mid-Proterozoic black shales: Evidence for an episode of increased ocean oxygenation at 1.36  
 300 Ga and evaluation of the effect of post-depositional hydrothermal fluid flow: *Precambrian*  
 301 *Research*, v. 298, p. 187-201, <https://doi.org/10.1016/j.precamres.2017.06.016>.  
 302 Zhang, S., Wang, X., Hammarlund, E. U., Wang, H., Costa, M. M., Bjerrum, C. J., Connelly, J. N.,  
 303 Zhang, B., Bian, L., and Canfield, D. E., 2015, Orbital forcing of climate 1.4 billion years  
 304 ago: *Proceedings of the National Academy of Sciences*, v. 112, E1406-E1413.  
 305 Zhang, S., Wang, X., Wang, H., Bjerrum, C.J., Hammarlund, E.U., Costa, M.M., Connelly, J.N.,  
 306 Zhang, B., Su, J., and Canfield, D.E., 2016, Sufficient oxygen for animal respiration 1,400  
 307 million years ago: *Proceedings of the National Academy of Sciences*, v. 113, p. 1731-1736,  
 308 <https://doi.org/10.1073/pnas.1523449113>.  
 309 Zhang, S., Wang, X., Wang, H., Bjerrum, C.J., Hammarlund, E.U., Haxen, E.R., Wen, H., Ye, Y.,  
 310 and Canfield, D.E., 2019, Paleoenvironmental proxies and what the Xiamaling Formation tells  
 311 us about the mid-Proterozoic ocean: *Geobiology*, v. 17, p. 225-246,  
 312 <https://doi.org/10.1111/gbi.12337>.  
 313 Zhao, G., Sun, M., Wilde, S. A., and Li, S., 2005, Late Archean to Paleoproterozoic evolution of the  
 314 North China Craton: key issues revisited: *Precambrian Research*, v. 136, p. 177-202,  
 315 <https://doi.org/10.1016/j.precamres.2004.10.002>.

321 FIGURE CAPTIONS

322 Figure 1. A: Geological map of the North China Craton, after Zhao et al. (2005) and Wang et al.  
323 (2017). Red square shows the location of the study area. B: Simplified stratigraphy of the Xiamaling  
324 Formation, with age constraints from Zhang et al. (2015).

325

326 Figure 2. Stratigraphic geochemical profiles for the Xiamaling Formation. Total organic carbon  
327 (TOC), Fe speciation, U and Mo concentration data are from Wang et al. (2017). Re data are from  
328 this study (closed circles) and Song et al. (2023) (open circles). The bulk  $\delta^{98}\text{Mo}$  profile comprises  
329 data from Diamond et al. (2018) (red open circles), Zhang et al. (2019) (dark open circles for core  
330 samples and grey open circles for outcrop rocks), and this study (closed circles). All  $\delta^{238}\text{U}$  data are  
331 from this study. Blue shading on the  $\delta^{98}\text{Mo}_{\text{auth}}$  and  $\delta^{238}\text{U}_{\text{auth}}$  profiles represent isotope ranges for  
332 (dys)oxic and F2 samples, while grey shading indicates the euxinic isotope range. EF–enrichment  
333 factor. (U)CC–(upper) continental crust. F1–high-TOC ferruginous. F2–low-TOC ferruginous.

334

335 Figure 3. Cross-plots of  $\delta^{238}\text{U}$  and  $\delta^{98}\text{Mo}$  *versus* their respective elemental and TOC contents.  $\delta^{238}\text{U}_{\text{det}}$   
336 and  $\delta^{98}\text{Mo}_{\text{det}}$  represent average continental crust values of -0.3‰ (Andersen et al., 2017) and 0.3‰  
337 (Voegelin et al., 2014), respectively. Note that the regression line on the  $\delta^{238}\text{U}$  *versus* TOC plot  
338 excludes the OMZ data (an  $R^2$  of 0.12 is obtained when the OMZ data are included).

339

340 Figure 4. Simplified U-Mo isotope mass balance model outputs. Most likely isotope compositions  
341 are mapped as a function of the relative areal fraction in different redox sinks. Area (a) represents the  
342 solution space that satisfies the  $\delta^{238}\text{U}$  data for the three sinks, while area (b) represents the solution

343 space that additionally satisfies the  $\delta^{98}\text{Mo}$  data. Thus, area (b) represents the inferred spatial extent of  
344 ferruginous and euxinic conditions, as satisfied by both isotope systems.

345

346

347 <sup>1</sup>Supplemental Material. Detailed descriptions of the geological background, analytical methods,  
348 detrital corrections, further discussion, modelling approach and geochemical data. Please visit  
349 <https://doi.org/10.1130/XXXX> to access the supplemental material, and contact  
350 [editing@geosociety.org](mailto:editing@geosociety.org) with any questions.

351