## **Supplementary Material for "Biotic replacement and mass extinction of the Ediacara biota" Darroch et al. (submitted)**

- **S1** Scatterplot illustrating positive correlation (spearman rho) between 'worker effort'
- (quantified as the number of original taxonomic papers), and overall generic richness for
- Ediacaran fossil localities. Note that we include skeletal fossils (e.g., *Namacalathus* and
- *Cloudina*), likely form taxa (e.g., *Aspidella*), and enigmatic tubular taxa (e.g.,

*Shaanxilithes*) in counts.



 **S2** – Measured sections (see also Fig. 1), showing the stratigraphic distribution of Ediacaran fossils encountered during measurement. Strata exposed as three prominent ridges or breaks in slope, which are interspersed with scree material, can be traced around the top of the koppe. The first ridge is equivalent to fossil bed 'A' of Narbonne et al. 21 (1997) while the second ridge (located stratigraphically  $\sim$  2 m above Bed 1) is equivalent 22 to fossil bed 'B' (of Narbonne et al. 1997) The third ridge is located ~8.5 m above ridge 2, and to our knowledge has not previously been identified as a fossiliferous horizon. *In situ* Ediacaran macrofossils were recovered from five individual horizons within the siliciclastic deposits (including the three ridges). From our section 1 in particular, we recovered *in*-*situ* fossils from ~1 m above the base of the siliciclastic horizons (thin- bedded green siltstone), the top surface of ridge 1 (weakly rippled coarse sandstone with 28 abundant microbial mat texture),  $\sim$  50 cm above ridge 1 (thin-bedded green siltstone), the top surface of ridge 2 (rippled medium sandstone), and within ridge 3 (thin yellow-green medium sandstone horizons with minor carbonate). Fossils recovered from float material occurred in a number of different lithologies, suggesting the existence of numerous other fossiliferous horizons not identified in this survey. Microbial mat textures are developed throughout the section, but particularly well on the top surface of ridge 1 where a large proportion of *in*-*situ* fossils are recorded. Similar to previous studies, we find a dramatic change in bedding, from horizontal to sub-horizontal/sub-vertical at the contact between fossil-bearing siliciclastic horizons and underlying carbonate, interpreted by Narbonne et al. (1997) as a 'mega slump'. Due to the discovery of Ediacaran macrofossils preserved in-situ on bedding planes (i.e., not jumbled and/or preserved in 3 dimensions, similar to transported assemblages elsewhere in Namibia – see e.g., Vickers-Rich et al., 2013) immediately above the basal contact, we agree with these previous workers that deformation was likely not syn-depositional, but rather the result of faulting or relatively recent slumping. Consequently, fossils are most likely autochthonous (or parauthocthonous) rather than transported as part of mass-flow facies. 



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 **S3** – Google Earth image of the koppe at Farm Swartpunt (top), and showing the approximate outline of surveyed area shaded in white (bottom).





- 61 **S4** Fossil database and description of sampling methods. Our database was built
- 62 through intensive survey of the fossiliferous horizons at Farm Swartpunt, over three days
- 63 in June 2014 (4 dedicated spotters, working 5 hours per day; equivalent to  $~60$  hours
- 64 total). The identities and contexts of all identifiable fossils are listed in Supplementary<br>65 Table 1. The total surveyed area at Farm Swartpunt, within the relevant horizons is
- Table 1. The total surveyed area at Farm Swartpunt, within the relevant horizons is
- 66 estimated at 20358.68 meters<sup>2</sup> (= 0.02 km<sup>2</sup>) (based on calculations using the polygon tool
- 67 in Google Earth) see S3.









 

 

 

 

 

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- **S5** Representative Ediacara biota recovered from Farm Swartpunt: a-c) *Pteridinium*
- *simplex*; d-f) *Swartpuntia germsi*; g) *Nasepia* sp.; h-i) *Aspidella*-type holdfasts, possibly
- 103 belonging to *Swartpuntia* (see S5b); j) unidentified rangeomorph taxon, provisionally<br>104 assigned to *Bradgatia*; k) unidentified Erniettomorph taxon, provisionally assigned to
- assigned to *Bradgatia*; k) unidentified Erniettomorph taxon, provisionally assigned to *Ernietta*.
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- **S5 (cont.)** Collected slab preserving specimens of *Aspidella* (SWP-99); holdfast 110 structure ('Hf'), *Swarptuntia*-type segmented stem ('St'), and *Swartpuntia*-type per
- 110 structure ('Hf'), *Swarptuntia*-type segmented stem ('St'), and *Swartpuntia*-type petaloid<br>111 ('Pet') are clearly visible. The positions of multiple other suspected holdfast structures
- 111 ('Pet') are clearly visible. The positions of multiple other suspected holdfast structures are marked with 'x'. A poorly preserved additional petaloid, possibly belonging to
- 112 are marked with 'x'. A poorly preserved additional petaloid, possibly belonging to another *Swartpuntia*, is circled.
- another *Swartpuntia*, is circled.





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**S6** – Facies, taphonomic, and geochronologic summaries for analysed assemblages

Farm Swartpunt (southern Namibia)

 *Palaeoenvironmental and stratigraphic setting* – The fossil-bearing horizons at Farm Swartpunt are part of the latest Ediacaran Nama Group, Urusis Formation (Spitskopf Member), deposited into the southernmost (Witputs) of two subbasins. Fossil beds are contained within siliciclastic horizons overlying brecciated horizons that contain slumped intervals, interpreted as a post-depositional 'megaslump' by Narbonne et al. (1997; see also S2). Fossils occurring above the slumped horizons are preserved in-situ (our 'Bed 1' – see S2); within the slumped horizons fossils may have been moved from their original positions, but are most likely parautocthonous (rather than allocthonous). The palaeoenvironment is interpreted as a quiet and open-marine setting at or near fair weather wave base, and shows evidence for occasional disruption by storms (Narbonne et al., 1997).

 *Taphonomic mode* – Fossils from all horizons are preserved as 2D casts and molds on the top- and bottom-surfaces of beds. Fossiliferous horizons frequently also preserve

evidence for microbial mats, and thus were likely preserved in the "death mask" style

common to many other Ediacaran localities, including Mistaken Point (Narbonne, 2005),

and South Australia (Gehling, 1999). The 3D taphonomic mode, characterized by moldic

infills (which can be isolated from the surrounding matrix) is not evident here, despite

being frequently seen elsewhere in Namibia (see e.g., Vickers-Rich et al., 2013),

reinforcing inference that these organisms represent an autocthonous accumulation.

*Geochronology* – An ash bed in the lower carbonate package of the Urusis Formation has

148 been dated by U-Pb geochronology at  $545.1 \pm 1$  Ma (recalculated to  $542.58 \pm 1.25$  Ma by

149 Schmitz, 2012), and an ash bed ~85 meters below the investigated fossil beds at  $543.3 \pm 1$ 

150 Ma (Grotzinger et al., 1995 - see Fig. 1; recalculated to  $540.61 \pm 0.67$  Ma by Narbonne et

al., 2012). Strata from the overlying Nomtsas Formation in the Swartkloofberg Farm

152 directly north of Swartpunt contain an ash bed dated to  $539.4 \pm 1$  Ma (i.e., Cambrian;

153 Grotzinger et al., 1995; recalculated to  $538.18 \pm 1.11$  Ma by Narbonne et al., 2012).

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- Nilpena (southern Australia)

 *Palaeoenvironmental and stratigraphic setting* – Fossils from Nilpena occur within the Ediacara member of the Rawnsley Quartzite (Flinders Ranges), broadly consisting of a

variety of shallow marine and deltaic facies, preserving evidence for wave action and

occasional storms (Gehling and Droser, 2013). More detailed sedimentological studies

(Droser et al., 2006; Gehling and Droser, 2013) have identified a complex series of

taxon-restricted paleoecosystems representing distinct sedimentary facies (i.e.

paleoenvironments) including shoreface sands, wave-base sands, delta-front sands, sheet-

flow sands, and mass-flow sands. The shoreface-, wave-base-, and delta-front sands are

all interpreted to reflect in-situ and untransported assemblages. By contrast, the sheet-

flow- and mass-flow sands preserve (largely) transported assemblages.

*Taphonomic mode* – Ediacaran preservation across the Flinders Ranges is typically

- represented by 2D casts and molds (i.e., 'death mask' preservation; Gehling, 1999) on the
- bottom surfaces of coarse-grained sandy storm event beds. However, rare sedimentary
- facies from Nilpena have resulted in 3D-preservation of large Ediacaran fronds (Gehling and Droser, 2013).

 *Geochronology* – The fossil assemblages from Nilpena are most similar to assemblages from Russia, Siberia, Ukraine, and northwestern Canada, and so assignment to the 'White

- Sea' assemblage (i.e., 555-550 Ma) is established mostly on a biostratigraphic basis. U-
- 174 Pb dates from Russia indicate ages between  $552.85 \pm 0.3$  Ma (Zimnie Gory Formation) to
- 175  $550.2 \pm 4.6$  Ma (base of the Yorga Formation; Iglesia-Llanos et al., 2005), however
- Nilpena lacks any notable volcanic ash beds, and so accurate dating has been difficult.
- This correlation receives some moderate support from a single U-Pb detrital zircon date
- 178 of  $556 \pm 24$  Ma from the Bonney Sandstone (Preiss, 2000).
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- Mistaken Point (Newfoundland)

 *Palaeoenvironmental and stratigraphic setting* – Fossiliferous horizons at Mistaken Point are dominated by relatively deepwater (> 500 m) turbiditic sandstones and mudstones

- (Wood et al., 2003; Ichasso et al., 2007; Mason et al., 2013; Liu et al., 2014). The fossil horizons analyzed here belong to the Drook (PC surface), Briscal (BC surface), Mistaken
- Point (E and D surfaces) and Trepassey (SH surfaces) Formations; for stratigraphic
- sectiosn see Wood et al. (2003) and Clapham et al. (2003). Previous studies infer a
- deepwater (toe of slope to lower slope) paleobathymetry well below storm wave base and
- the photic zone (Wood et al. 2003; Ichaso et al. 2007; Mason et al., 2013). Turbidite beds are typically overlain by a thin (1–2 cm) mudstone interpreted as pelagic fallout (Wood et
- al. 2003). This thin pelagic mudstone is characterized by thin, black, 'crinkly' and
- discontinuous silt laminae that may represent diagenetically altered microbial mats (e.g.
- Wood et al. 2003; Narbonne et al. 2005). Thin (1–2 cm) beds interpreted as deepwater
- contourite deposits (Wood et al. 2003) are found above terminal-stage turbidite beds and,
- when overlain by volcanic ash, typically contain Ediacaran fossils.
- *Taphonomic mode* Organisms at Mistaken Point were preserved in-situ after being smothered by volcanic ash ('Conception-style' preservation of Narbonne 2005). The
- local presence of seafloor microbial mats and rapid onset of anaerobic decay led to early
- lithification of the soles of overlying ash beds, effectively casting fine-scale morphology
- (Narbonne 2005; Laflamme et al. 2011; Liu et al. 2011).
- *Geochronology* Ash beds bracketing the fossiliferous horizons analysed in this study have been dated using U-Pb geochronology at 580 Ma, 578 Ma, and 565 Ma (see Benus, 1988; Bowring et al., 2003). See Darroch et al., 2013 (figure 1) for stratigraphic section with dated horizons in context.
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- White Sea (Russia)
- *Palaeoenvironmental and stratigraphic setting* Ediacaran deposits in the White Sea area of Russia are represented a thick (500 m) succession of sandstones, siltstones, and
- mudstones deposited in shallow basin at high palaeolatitudes; the studied assemblage
- comes from the Verkhovhka Formation, which underlies the Zimnie Gory Formation in
- the vicinity of the Solza River (Zakrevskaya, 2013). The analysed community comes
- from a single bed, and flourished in relatively shallow (at or within fair weather wave
- base) palaeonvironment, likely representing an alluvial fan or delta-front type setting
- disrupted by periodic mass sedimentation events which buried Ediacaran organisms
- (Zakrevskaya, 2013).
- *Taphonomic mode* White Sea fossils from the analysed horizon are preserved in
- 'Flinders-style' (of Narbonne, 2005), most likely as "death masks" after being smothered
- by transported sediment, similar to modes of preservation described for other Ediacaran
- localities worldwide (Zakrevskaya, 2013). This indicates that fossils are untransported,
- and most likely represent an in-situ accumulation largely in life-position (Narbonne,
- 2005; Zakrevskaya, 2013).
- *Geochronology* U-Pb dates from the White Sea area of Russia indicate ages between
- 222  $552.85 \pm 0.3$  Ma (Zimnie Gory Formation) to  $550.2 \pm 4.6$  Ma (base of the Yorga
- Formation; Llanos et al., 2005). The base of the Verhkovhka Formation has a U-Pb date
- 224 of  $558 \pm 1$  Ma (Grazhdankin, 2004).
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 **S7** – Palaeoecological indices for all studied (raw) datasets; 'SR' = Species Richness, 240 'Dom.' = Dominance  $(1 - \text{Simpson's Index})$ , 'M-Div' = Margalef's Diversity, 'S-W' = Shannon-Weiner Index, 'B&G Even.' = Buzas and Gibson's Evenness. Note that no surveyed-area estimates have been published for Nilpena datasets (Gehling and Droser, 2013), raising the possibility (however unlikely) that some of the elevated diversity seen in these sites may be due to richness-area effects. Also note that given the incomplete outcrop and geometry of our 'Bed 1' at Swartpunt, no reliable estimates of surveyed area could be obtained. For Mistaken Point datasets, '*Charnia*' on the Mistaken Point BC, D, E, and SH surfaces is now assigned to *Beothukis* (Brasier and Antcliffe, 2009). 'Networks' on the D surface are now assigned to *Hapsidophyllas* (Bamforth and Narbonne, 2009). 'Charnia I' on the LMP surface is now assigned to *Trepassia* (Narbonne and Gehling, 2003), while 'Charnia II' and 'ostrich feathers' on the same surface are both assigned to *Culmofrons* (Laflamme et al., 2012). *Hiemalora* on the LMP surface is now assigned to *Primocandelabrum*, on the basis of observations by Hoffmann et al. (2008). Similar to Darroch et al. (2013) we exclude Ivesheadiomorphs from analyses, as these may not represent body fossils (see for example Liu et al., 2012). For Nilpena datasets, numbers of *Aspidella* and *Funisia* are listed as '>999' in some facies. In these cases we have standardized the number of these taxa at 1000 individuals. All indices were calculated

257 using the open-access statistical software R.



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 **S8** – Results of rarefaction analyses at species (rather than genus) level; note that patterns are virtually identical between analyses at both taxonomic resolutions.



## **Species**

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 **S9** – Results of rarefaction analyses excluding *Aspidella* for both genus- and species-level analyses; results are identical to those of raw data, illustrating that patterns are not controlled by frondose taxa. Top panels illustrate all datasets. Middle panels illustrate contrasts between Swartpunt and Mistaken Point datasets, and lower panels illustrate contrasts between Swartpunt and Nilpena datasets; error bars have been added to these panels as 95% confidence intervals around mean diversity values. Areas of low sampling intensity (shaded in grey) have been expanded in adjacent panels to better illustrate







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- **S10** Supp. Geochemical data Table 1 (as .xls file)
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- **S11** Supp. Geochemical data Table 2 (as .xls file)
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 **S12** – Although these geochemical tests provide no evidence for a stressed environment, caveats do exist. First, current geochemical proxies can fingerprint anoxic conditions with certainty, but have difficulty unambiguously distinguishing oxic from ferruginous conditions under certain conditions (Sperling et al., 2014). The sum of multi-proxy data from the Spitskopf Member, however, makes a ferruginous Palaeoenvironment unlikely. More pertinently, available geochemical proxies distinguish oxic from anoxic conditions in an essentially binary fashion, and cannot inform us about degrees of dysoxia that are biologically relevant (Poulton and Canfield, 2011). Along these lines, it is also possible 300 that the Nama Group Ediacarans were living in close proximity to a chemocline, and 301 were periodically flooded by low- $O_2$  waters. The biological relevance of this is mitigated. were periodically flooded by low- $O_2$  waters. The biological relevance of this is mitigated by the observation that in modern environments where metazoans are subjected to periodic upwelling of anoxic and even euxinic waters, such as off the coast of Namibia, a well-established and moderate diversity (albeit lower diversity than in very nearshore waters) community continues to exist (Zettler et al., 2009; 2013). This illustrates that in the modern ocean, relatively diverse communities of aerobic multicellular heterotrophs can exist in the face of periodic dysoxic to anoxic waters (although many Ediacaran organisms may not actually have been animals – see Erwin et al., 2011; Laflamme et al., 2013). Finally, it is noted that organic carbon contents of the fossiliferous strata are not just low, but essentially nonexistent. In conjunction with a complete absence of pyrite, it suggests that these rocks have been subject to oxidative weathering processes. While this will not unduly affect the ability of the iron speciation proxy to distinguish an oxic from anoxic water column, as pyrite will weather into iron oxides and remain in the highly reactive pool, it does indicate that the original organic carbon values were likely higher than measured.

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