

Interpreting palaeofire evidence from fluvial sediments; A case study from Santa Rosa Island, California with implications for the Younger Dryas Impact Hypothesis

Journal:	<i>Journal of Quaternary Science</i>
Manuscript ID	JQS-15-0070.R2
Wiley - Manuscript type:	Research Article
Date Submitted by the Author:	n/a
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Keywords:	Fluvial sedimentology, stratigraphy, charcoal, Younger Dryas Impact Hypothesis, extra-terrestrial impact

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2 1 **Interpreting palaeofire evidence from fluvial sediments; A case study from**
3 2 **Santa Rosa Island, California with implications for the Younger Dryas**
4 3 **Impact Hypothesis**
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21 17

22 18 Received: xx June 2015

23 19 Revised 28 Sept 2016
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ABSTRACT: Fluvial sequences from the late Pleistocene to the Holocene are exposed in Arlington Canyon, Santa Rosa Island, Northern Channel Islands, California, USA, including one outcrop that features centrally in the controversial hypothesis of an extra-terrestrial impact at the onset of the Younger Dryas. The fluvial sequence in Arlington Canyon contains a significant quantity and range of organic material, much of which has been charred. The purpose of this study was to systematically describe the key outcrop of the Arlington sequence, provide new radiocarbon age control and analyse organic material in the Arlington sediments within a rigorous palaeobotanical and palaeo-charcoal context. These analyses provide a test of previous claims for catastrophic impact-induced fire in Arlington Canyon. Carbonaceous spherular materials were identified as predominantly fungal sclerotia; ‘carbon elongates’ are predominantly arthropod coprolites, including termite frass. Glassy carbon formed from the precipitation of tars during charcoalification. None of these materials indicate high-temperatures formation or combustion. Charcoal and other materials in Arlington Canyon document widespread and frequent fires both before and after the onset of the Younger Dryas, recording predominantly low-temperature surface

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2 40 fires. In summary, we find no evidence in Arlington Canyon for an extra-terrestrial impact or catastrophic
3 41 impact-induced fire.
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7 43 KEYWORDS: Fluvial sedimentology; stratigraphy; charcoal; Younger Dryas Impact Hypothesis; extra-
8 44 terrestrial impact;
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10 46 11 47 **Introduction** 12 48

13 48
14 49 Quaternary fluvial records provide information on terrestrial palaeoclimate (e.g., Pigati *et al.*, 2014),
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16 50 neotectonics (e.g., Pinter and Keller, 1995), archaeological context (Mishra *et al.*, 2007), and a wide
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18 51 variety of other areas (see Bridgland & Westaway, 2014). These sediments offer unique challenges, and
19 52 failure to consider these may hamper, or worse, lead to erroneous palaeoenvironmental interpretations.
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21 53 The Younger Dryas Stadial, which corresponds to Greenland Stadial-1 (GS-1; ~12.9-11.7 ka BP;
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23 54 (Rasmussen *et al.*, 2006) has been well described and has been recognised in proxy records from
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25 55 California (e.g., Hendy *et al.*, 2002). The “Younger Dryas Impact Hypothesis” (YDIH) is the relatively
26 56 new suggestion (Firestone *et al.*, 2007) that global events approximately 12.9 kyBP – including climatic
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28 57 cooling, extinction of North American megafauna, demise of the Clovis archaeological culture, and other
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30 58 changes worldwide – resulted from the impact of a 5-km diameter comet into the southern margin of the
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32 59 Laurentide ice sheet. The YDIH is controversial and has been heavily contested (e.g., Pinter and Ishman,
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34 60 2007; Pinter *et al.*, 2011; Boslough *et al.*, 2013; van Hoesel *et al.*, 2014; Holliday *et al.*, 2014; Meltzer *et al.*,
35 61 2014; Daulton *et al.*, 2016). Although many sites globally have been put forward as containing
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37 62 evidence for the YDIH (e.g., LeCompte *et al.*, 2012; Bunch *et al.*, 2012; Wittke *et al.*, 2013; Petaev *et al.*,
38
39 63 2013; etc.) one key sedimentary section from Arlington Canyon, Santa Rosa Island, in the Northern
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41 64 Channel Islands of California (Fig. 1) has played a particularly important role in the ongoing development
42 65 of the YDIH (AC003 in Kennett *et al.*, 2008 *et seq*; site III in this study). Several key papers have
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44 66 focused on this locality with the interpretation of “intense biomass burning” and associated rapid
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46 67 landscape change (Kennett *et al.*, 2008), the presence of nanodiamonds (Kennett *et al.*, 2009a) and shock-
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48 68 synthesized hexagonal diamonds (Kennett *et al.*, 2009b). More recently, dating evidence from the site was
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50 69 a key component used in a Bayesian chronological analysis which found a synchronous age for the start
51 70 of the Younger Dryas boundary or ‘impact’ layer (Kennett *et al.*, 2015).
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53 71 The purpose of this study was to systematically sample locality AC003 (Kennett *et al.*, 2008 *et seq*;
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55 72 site III in this study) in Arlington Canyon for its evidence of palaeofire and relate this single site
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57 73 stratigraphy to other multiple sites along the canyon that we also investigated for fire history (see Pinter *et al.*,
58 74 2011; Hardiman *et al.*, 2016). We demonstrate that evidence in Arlington Canyon is inconsistent with
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1 75 the catastrophic extraterrestrial impact and the associated local manifestations that have been proposed.
2
3 76 More broadly, the widely divergent interpretations of events preserved in Arlington Canyon illustrate
4
5 77 general challenges in using palaeobotanical and charcoal records from fluvial sequences. We present
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7 78 recommendations and protocols for analysis of Quaternary fluvial deposits, particularly for the collection
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9 79 of macro-charcoal (defined here as >125 µm) and interpretation of palaeofire from these more complex
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11 80 sedimentary sequences.

12 81 13 14 82 **Material and Methods**

15 83
16 84 Arlington Canyon is one of a series of north-south oriented drainages along the northern flank of Santa
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18 85 Rosa Island, carrying discharge from the island's interior northward to the coast (Fig 1). Santa Rosa
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20 86 Island has been slowly uplifting through the Quaternary (Pinter *et al.*, 2001), resulting in rugged
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22 87 topography and streams within deeply incised canyons (Schumann *et al.*, 2016). At the base of Arlington
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24 88 and selected neighbouring canyons, one aggradational terrace level forms a morphological bench up to
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26 89 ~25 m above the modern stream. This terrace sedimentary fill consists of fluvial and localized colluvial
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28 90 deposits that aggraded from the canyon base during the latest Pleistocene until the mid-to late Holocene
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30 91 (Pinter *et al.*, 2001; Schumann *et al.*, 2014). This was followed by a cessation of deposition and
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32 92 reinitiation of incision that cut base level to the bottom of the canyon and exposed the Pleistocene to
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34 93 Holocene fill deposits in a narrow "slot canyon" through the terrace (Schumann *et al.*, 2016). The terrace
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36 94 fill sequence consists of several fluvial cut-and-fill packages, consisting of channel and floodplain
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38 95 deposits that pinch out laterally or grade into colluvial deposits at their margins. Distinguishable
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40 96 stratigraphic units can be traced laterally over distances of metres to tens of metres, but these fluvial units
41
42 97 change in texture and character both vertically and horizontally. Sandy point bars and silt-dominated
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44 98 overbank deposits are punctuated by conglomeratic channel fills. Distinguishable depositional units range
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46 99 in thickness from less than 1 metre to more than 10 m. Between cut-and-fill packages, several
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50 100 depositional hiatuses and erosional unconformities are marked by weakly developed palaeosols,
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49 101 characterized by darker colour (Pinter *et al.*, 2011; Schumann *et al.*, 2014), enriched clay content, and
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51 102 weak soil structure developed on these undulating palaeo-topographic surfaces.

52 103 Our Locality I (see Scott *et al.*, 2010 supplementary data) is an exposure more than 10 m high and
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54 104 100 m long (UTM). Our Localities II and III are located just 110 m and 190 m north, respectively, of
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56 105 Locality I, but the lateral variability in the fluvial architecture makes it impossible to correlate sections at
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58 106 the scale of individual units. Locality III is a 4-m high exposure on the western side of the canyon
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60 107 (33° 59' 25.526"N 120° 9' 32.208"W) (See Supplementary Materials, Fig. S1). Wittke *et al.* (2013) claim
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62 108 that "coordinates, photographs, stratigraphic descriptions, and radiocarbon ages presented in their papers
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1
2 109 (e.g. Scott *et al.*, 2010 and Pinter *et al.*, 2011) conclusively demonstrate that none of their samples
3 110 collected were taken from the same stratigraphic section studied by Kennett *et al.* (2008).” On the
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5 111 contrary, our Locality III is identical to their locality AC003 (See Supplementary Materials, Fig. S2).
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7 112 Furthermore, material from AC003 was sent to the senior author in March 2007 by G. James West (via
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9 113 John Johnson) with a request to report on the charcoal. Lithological logs of other Arlington sections and
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11 114 radiocarbon data are given in Hardiman *et al.* (2016).

12 115 13 14 116 *Sampling procedures*

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16 117 The large changes in depositional facies over short distances within the Arlington Canyon fluvial
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18 118 sequence, combined with high vertical-relief and cut-and-fill sedimentary packages require extensive
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20 119 detail in the stratigraphic descriptions (Fig. 2) and a large number of dated samples in order to correlate
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22 120 packages of sedimentary aggradation through the full sequence. Our sampling goals included: (1) to
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24 121 collect organic material in the sediments, with particular interest in charcoaled plants (macrocharcoal,
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26 122 >125 μ m), and (2) to obtain material for radiocarbon dating. Thus we sampled every horizon with visible
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28 123 charcoal. At some intervals where continuous sampling was necessary, we used a core box that could be
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30 124 hammered into the section and removed for later sub-sampling. All samples were photographed *in situ*
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32 125 before removal.

33 126 34 127 *Sample processing and radiocarbon dating*

35 128 In order to separate charcoal or macroscopic plant material from bulk sediment, we first removed any
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37 129 large rock clasts. Sediment was then soaked in warm water for disaggregation; if needed we used 10%
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39 130 hydrogen peroxide (Rhodes, 1998). It should be noted that the charcoal in such water baths generally does
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41 131 not float off, as suggested by Firestone *et al.* (2007). The samples were then wet sieved to produce
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43 132 residues of below 62 μ m, below 125 μ m, and above 125 μ m. Charcoal was picked from the >125 μ m
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45 133 residues. We note that in all samples, charcoal pieces are liable to fragment, so counts of number of
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47 134 fragments are not meaningful, particularly in fluvial sediments. Some of the charcoal residues were
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49 135 cleaned by dissolving the sediment in 40% HF (see Scott, 2010).

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51 136 Particularly for fluvial deposits, a pervasive issue for radiocarbon dating is the potential for "old
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53 137 wood" charcoal dates (Schiffer, 1986; Gavin, 2001; Bird, 2013). Because charcoal is chemically inert and
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55 138 mechanically robust, it can sometimes survive erosion from a pre-existing deposit, transport through the
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57 139 fluvial system, and redeposition. In order to minimize the danger of dating secondary, re-deposited
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59 140 charcoal, we identified organic material before submission for radiocarbon analysis and selected only
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61 141 fragile but well preserved charred plant parts, rather than more robust charcoal fragments. Picked

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2 142 samples of charcoalfied wood, seeds, carbonaceous spherules and coprolites were sent for radiocarbon
3 143 dating by two different laboratories: the Keck Carbon Cycle AMS Laboratory at University of California
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5 144 (UC) Irvine and the Oxford Radiocarbon Accelerator Unit, RLAHA, University of Oxford (see Hardiman
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7 145 *et al.*, 2016).

8
9 146 There are several methods to separate charcoalfied plant material from disaggregated sediment
10 147 samples (see Scott, 2010). Samples picked from sediments were studied by light microscopy or mounted
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12 148 on aluminium stubs for scanning electron microscopy. Some charcoal was embedded into resin blocks
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14 149 and polished for examination under oil reflective microscopy. We attempted to use the protocol outlined
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16 150 in Firestone *et al.* (2007) and Kennett *et al.* (2008, 2009b) for specimen isolation, but following these we
17 151 were not successful. We found that none of the charcoal separation techniques cited in Firestone *et al.*
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19 152 (2007) worked for the Arlington samples, so it is uncertain how these were collected, processed or picked.
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21 153 Sampling protocols provided in “Separation of YD Event Markers (8/10/2007),” a guide provided by one
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23 154 of its authors (Allen West, GeoScience Consulting), will break up charcoal fragments in to a large number
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25 155 of smaller fragments.

26 156 27 28 157 *Microscopy of palaeobotanical samples*

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30 158 Samples were identified under water by reflected light under a low-power binocular microscope. Some
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32 159 samples were picked using dark-field lighting (see Glasspool and Scott, 2013) that facilitated the
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34 160 separation of charred and un-charred plant fragments. Some specimens were gold-coated using a Poloron
35 161 sputter coater. Uncoated specimens were studied using a Hitachi S3000N variable pressure SEM under
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37 162 low vacuum and in backscatter electron mode. Coated samples were studied using secondary electron
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39 163 mode. Specimens were also gold coated and examined using a Philips Environmental SEM.

40 164 Uncharred fungal sclerotia, charred sclerotia, carbonaceous spherules, and wood charcoals were
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42 165 embedded in polyester resin, cut, and polished. Reflectance was measured using a Leica DM2500
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44 166 microscope linked to a MSP200 photometer reflectance system. The specimens were measured under oil
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46 167 of refractive index 1.518, using light filtered to 546 nm. Mean random reflectance (Ro %) was measured,
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48 168 and temperature conversion was achieved by comparison with wood and fungus charcoal experimental
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50 169 charcoalfication curves. Full charcoal reflectance methodology and background are presented in Scott
51 170 and Glasspool (2005, 2007), McParland *et al.* (2009), and Scott (2010).
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53 171 54 55 172 *Organic Geochemistry (Analytical Pyrolysis)*

56 173 Analytical pyrolysis was carried out using an SGE Pyrojector pressurised with helium at 15 psi
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58 174 and fitted to an HP5890 Series II gas chromatograph (GC) interfaced to an HP5972 MSD mass
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1 spectrometer at Royal Holloway University of London. Samples (~1 mg) were loaded into and introduced
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spectrometer at Royal Holloway University of London. Samples (~1 mg) were loaded into and introduced with a P-3 pelletizer, and pyrolysis was carried out at 650°C. Pyrolysate was transferred to the chromatography column with a constant flow of helium of 0.7 cm³/min. into the GC inlet kept at 280°C. The column (J&W DB5, 30 m x 0.25 mm x 0.25 µm film thickness) was initially at 50°C for 2 min., then heated at 7.5°C/min to a final temperature of 330°C. Splitless injection was applied with a delay time of 1.5 min., and the GC-MS interface temperature was set at 300°C.

Sedimentological , straigraphical and biological description

Along much of the Arlington Canyon study area, the basal 1-2 m of Quaternary fill consist of horizontal to sub-horizontally bedded silt-dominated strata, with dispersed sand-size grains. We sampled and measured as low in the section as possible, sometimes hand-excavating several decimetres below groundwater level. Because the basal sediments were wet in outcrop, they gave the impression of being darker in colour and, seemingly, more organic-rich (Kennett *et al.*, 2008, 2009b). This was not the case; the samples lightened to a grey-brown colour upon drying (Fig. 2; Supplementary Materials, Fig. S3).

Within these fine-grained basal facies are isolated sand- and gravel-rich laminae that occur as lenses, bar forms, and thin channels (Fig. 2, S3). This coarser clastic fraction includes small rounded granules and pebbles and a few, isolated more angular and larger rock fragments. Some of the horizons contain charcoal, but the charred fragments were not uniformly distributed within them. Conglomeratic units occurred as lenses or as distinct channel fills. The base of section IIIc, for example, comprises a >1 m-thick gravel layer. Less than 8 m to the north, this horizon has thinned and is no longer present (Log IIIa). Log IIId is located identical to the section described by Kennett *et al.* (2008, 2009b), and the photograph showing the position of their recorded section is shown in Wittke *et al.* (2013, Supplementary Information) and here in Fig. S2.

Overlying the basal, predominantly fine-grained deposits in Arlington Canyon is a sand-dominated package, consisting predominately of laminated and cross-bedded sands. Charcoal and charred plant fragments are widespread, ranging in size up to >1 cm in diameter (See Supplementary Materials, Fig. S4); some horizons also contain un-charred and partially charred plant material. Within the coarse sands, there are abundant coarser granule lenses and isolated pebbles (Supplementary Materials, Fig. S5). At Locality III at the ~2 m level, there is a thin clay-rich band, dark but not organic-rich, that is clearly identifiable on the log and photos of Kennett *et al.* (2008, 2009b; Wittke *et al.*, 2013) (Fig. 2). The next metre higher in the section at Locality III is predominantly fine sand with some cross beds, scattered charcoal fragments (Fig. 2), and some coarse sand that often fills small channels (Fig. 2). This unit is

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2 208 cross-cut by an erosional ravinement surface that is widespread in Arlington Canyon, locally high in relief
3 209 and down-cutting through the underlying units by >10 m in some locations.
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5 210 6 7 211 *Charcoal distribution and identification*

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9 212 Charcoal (Figs. 3, 4; Supplementary Materials, Fig. S6) in the Arlington Canyon sequence, especially
10 213 wood charcoal is concentrated in the basal ~3 m of the sections (Table 1). Charcoal becomes less common
11
12 214 higher in the sequence. Charcoal occurs as thin discontinuous layers, lenses and as scattered fragments
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14 215 (Fig. S4). In cross-bedded units, charcoal is concentrated in foreset cross-beds (Fig. S4a). In sample
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16 216 AC003, we have noted abundant charcoal, often up to 5 mm in size. Secondary wood charcoal from
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18 217 Arlington Canyon samples tends to dominate (Table 1). However there is an equal proportion of conifer
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20 218 (Fig. 4a-c) and angiosperm (Fig. 4f-i) wood charcoal throughout the sequence (Fig. 4). In addition, small
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22 219 herbaceous angiosperm axes (Figure 4j) are common in some samples, but leaf (Fig. 4d,e), bark charcoal
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24 220 and seeds are relatively rare.
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26 221 27 222 *Carbonaceous spherules and "elongate" forms*

28 223 Firestone *et al.* (2007) coined the term "carbon spherules," referring to "highly vesicular,
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30 224 subspherical-to-spherical objects 0.15–2.5 mm in diameter, with cracked and patterned surfaces, a thin
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32 225 rind, and honeycombed (spongy) interiors." According to Firestone *et al.* (2007), these particles were
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34 226 formed during high-temperature ignition associated with the Younger Dryas extraterrestrial impact event.
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36 227 Kennett *et al.* (2008) identified "carbon elongates," which were described as similar in size, context, and
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38 228 origin, but ellipsoidal in shape and with "a much coarser interior cellular structure." In our Arlington
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40 229 samples, carbonaceous spherular forms occur throughout the section but are more common in the basal 2
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42 230 m (Table 1)(Fig. 3; Supplementary Materials, Fig. S7). They range in size from 250µm-1.5 mm in
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44 231 diameter. In cross section, they often show a thin surface rind and internally a spongy internal texture
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46 232 (Fig. S7). The internal anatomy of these spherules is very diverse. Most of the spherules are black in
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48 233 colour. Our sediment sample AC003 from West contains common carbonaceous spherules (Fig. S5f).

49 234 Carbonaceous particles that match the description of 'carbon elongates' occur throughout Locality III
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51 235 and other Arlington sections and are very abundant within several samples (Supplementary Materials, Fig.
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53 236 S8)(e.g., SRI-10-56; Table 1). Some 'elongate' forms show hexagonal morphology (Fig. S8f). In most
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55 237 samples they are black, but in SRI-10-55 they show a range of colours from brown to black (Fig. S8b).
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57 238 Sample AC003 contains a few 'carbon elongates'.

58 239 59 240 *Glassy carbon* 60

1
2 241 Firestone *et al.* (2007) also identified glass-like carbon, consisting of angular fragments up to several
3 242 cm in size, with glassy texture “suggest[ing] melting during formation” purportedly recording impact-
4 generated, high-intensity fire. Material that could be described as “glassy carbon” occurs throughout the
5 243 Arlington section but is rarely abundant (Table 1). It occurs as small pieces usually a few mm in size Fig.
6 244 3l). It is common in sample SRI-10-55 from Log IIIa. Sample AC003 from West contains a few
7 245 specimens of glassy carbon.
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12 247 Three samples of glassy carbon from Arlington Canyon were examined by analytical pyrolysis/gas
13 chromatography/mass spectrometry and compared with samples of charcoal prepared by treatment of
14 248 *Sequoia* at 350°, 450° and 600°C (Scott and Glasspool, 2005) and with a sample of synthetic glassy
15 249 carbon (Alfa-Aesar 42130, Type 1, 200-400 µm spherical). While we do not believe that there is any
16 250 similarity between glassy carbon as recorded in sediments and true commercially produced glassy carbon,
17 251 we nevertheless examined both materials. As anticipated, the synthetic glassy carbon, which is specified
18 252 to be stable up to 1100°C, gave no chromatographic peaks. The chromatograms of the *Sequoia* and Santa
19 253 Rosa samples are shown in Fig. S9-f, and compared in a bar chart showing the relative percentage peak
20 254 areas of the 16 most prominent compounds present (Table 2).
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30 257 *Nanodiamonds*

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32 258 We examined three different specimen sets of carbonaceous spherules for the presence of
33 259 nanodiamonds: 1) five spherules/fragments from SRI 09-28A; 2) eight spherules/fragments from AC003;
34 and 3) 13 acid-washed spherules/fragments from AC003. For a detailed discussion on the interpretation
35 260 of this evidence please refer to Daulton *et al.* (2016).
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43 264 **Data interpretation**

44 265 *Charcoal*

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46 267 The majority, but not all, of charcoal found in Quaternary terrestrial sediments come from wildfires
47 (Glasspool and Scott, 2013). Most modern charcoal accumulations within fire areas are produced by the
48 268 charring of surface litter from low-temperature surface fires (Scott, 2010; Scott *et al.*, 2014). Higher
49 temperature crown fires often totally combust the plant material and leave no macroscopic charcoal
50 269 residue. Charcoal in fluvial settings may indicate not only fire occurrence but also, in some
51 270 circumstances, deposition during post-fire erosion (Brown *et al.*, 2013). Charcoal type also may indicate
52 burning of trees, shrubs or herbs (Scott, 2010). In this study, charcoal from Arlington Canyon was derived
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2 274 from conifer trees, angiosperm trees and shrubs and herbaceous angiosperms. This suggests that the fire
3 275 was probably predominantly a surface fire (Scott *et al.*, 2000; Scott, 2010).
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5 276 6 7 277 *Carbonaceous Spherular Forms*

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9 278 Two carbonaceous forms – widely known within palaeobotanical circles, but perhaps less so
10
11 279 elsewhere – have been reported in samples from Arlington Canyon and have created much confusion.
12
13 280 Carbonaceous spherular forms (so-called ‘carbon spherules’) ranging in size from less than 100 µm to
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15 281 over 1 mm occur frequently in charcoal residues from most wildfires. Such material is particularly
16
17 282 common in charred litter from surface fires. Even in the case of a hot crown fire, most charcoal comes
18
19 283 from the charring of surface-dwelling plants and litter (Scott, 2010).

20
21 284 One of the most common spherular types found in the Arlington Canyon samples are fungal sclerotia
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23 285 (Fig. S7). Sclerotia are common both in the soil and attached to living and dead plant debris. The sclerotia
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25 286 are resting cysts (Fig. S5) that often form during periods of water stress (Amasya *et al.*, 2015). Their
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27 287 occurrence in charcoal residues is not unexpected. The genus *Sclerotium* is common, but in both modern
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29 288 and Quaternary sediments, *Cennococcum* is also widespread (Ferdinandson *et al.*, 1925; Sakagami and
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31 289 Watanabe, 2009; Benedict, 2011). Sclerotia have a distinctive morphology: in cross section they have a
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33 290 thin crust, and the interior may be foam-like (Fig. S7). Their texture can be modified by fire, and the level
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35 291 of modification is a function of temperature (Scott *et al.*, 2010). Just as with wood and other fungal
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37 292 material, the reflectance of charred sclerotia increases with increasing temperatures (Scott *et al.*, 2010;
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39 293 Scott and Glasspool, 2007). The number of sclerotia in a sediment sample will be controlled by their
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41 294 abundance in the source area and by sedimentological processes. Many fluvial processes concentrate
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43 295 organic matter, including sclerotia (Malloch *et al.*, 1986).

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45 296 Carbonaceous spherular forms are found throughout the Arlington sequence but are more common
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47 297 near the base of the section. This concentration may be due to either external factors (greater
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49 298 concentration of the presumed source material) or internal processes such as sedimentary concentration
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51 299 (in the low-energy, fine-grained deposits that predominate near the base of the Arlington sequence). It is
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53 300 possible that carbonaceous spherular forms have multiple origins, but most ‘carbon spherules’ that we
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55 301 have examined can be confidently identified as fungal sclerotia (see also discussion in Daulton *et al.*,
56
57 302 2016).

58 303 59 304 *‘Carbonaceous elongates’/ Coprolites*

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61 305 The elongate forms described by Kennett *et al.* (2008) also may have a range of origins. Some may
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63 306 represent fungal sclerotia (Sakagami and Watanabe, 2009). However, by far the most common origin is
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2 307 arthropod fecal pellets (coprolites) (Scott, 1992). Arthropod coprolites are abundant in fluvial and indeed
3 308 all terrestrial sediments since the Devonian (e.g., Scott, 1977; Chaloner *et al.*, 1991; Scott *et al.*, 1992;
4
5 309 Habgood *et al.*, 2004; Edwards *et al.*, 2012). They may be produced by a wide range of arthropods, the
6
7 310 smallest (<50 μm) from mites, to collembolan and termites, and the largest coprolites (>1 mm) from
8
9 311 millepedes (Scott, 1992). These particles have a range of shapes and contents. Many of the coprolites
10
11 312 from the sediments at Locality III in Arlington are cylindrical with rounded ends (Fig. S8). These are
12
13 313 uncharred, partially charred, or occur as charcoal (Fig S8b). When charred, coprolites may shrink and the
14
15 314 inside preferentially combust, leaving hollow shells. A significant number of the Arlington coprolites
16
17 315 have a hexagonal cross section, which is typical of termite frass (Light, 1930; Lance, 1946; Scott, 1992;
18
19 316 Collinson, 1999b; Colin *et al.*, 2011) (Fig. S8d). Such frass is abundant in archaeological deposits
20
21 317 (Adams, 1984) and has been identified at other California sites (Light, 1930; Lance, 1946; Anderson and
22
23 318 Stillick, 2013). We have experimentally charred termite frass at a range of temperatures. We found that
24
25 319 the outer shape is retained and the reflectance increases with temperature (Scott and Glasspool, 2007;
26
27 320 McParland *et al.*, 2007).

28 321 29 322 *Glassy Carbon*

30 323 Some carbonaceous materials found in sediments have been termed “glassy carbon” because they
31
32 324 exhibited a glassiness or vitreous appearance (Scheel-Ybert, 1998). Material of the same name – but
33
34 325 structurally and chemically distinct – was also synthesized by carbonization of polymer precursors
35
36 326 starting in the mid-1950s. True glassy or vitreous morphology in carbonaceous materials does not result
37
38 327 exclusively from high temperatures (Marguerie and Hunot, 2007; Fabre 1996), but can also result from
39
40 328 the fine-grained homogenous nature of the material. McParland *et al.* (2010) showed that neither the
41
42 329 charcoals associated with glassy carbon, nor the glassy carbon itself in the sediments exhibited features of
43
44 330 high-temperature formation. Another explanation for the origin of glassy carbon comes from the
45
46 331 charcoalification process itself, which involves pyrolysis in the absence of oxygen (Scott, 2010;
47
48 332 Beaumont, 1985, section 2.5).

49 333 The chromatogram of the pyrolysate of sample AC003 (Fig. S9d) shows a composition similar to
50
51 334 those obtained from samples of *Sequoia* experimentally charred at 350 and 450°C (Fig. S9bc). In addition
52
53 335 to aromatic hydrocarbons, oxygen and nitrogen-containing compounds, viz. pyridine, phenol,
54
55 336 benzonitrile, benzofuran, methylphenols and dibenzofuran, are present. The chromatograms of the 350°
56
57 337 and 450° experimental *Sequoia* samples and AC003 are similar to those obtained by Kaal *et al.* (2009)
58
59 338 from 6200 year-old Fabaceae-derived charcoal from Campo Lamiero, northwest Spain. The implication is
60
61 339 that the charcoal sample AC0003 was formed at a temperature < 600°C. Chromatograms produced from

1 340 samples 10-36 and 10-57 (Fig. S9ef) resemble those of the 600°C *Sequoia* charcoal (Table 2). The
2
3 341 implication is that these charcoals were formed at a higher temperature than that experienced by sample
4
5 342 AC003, but there is no evidence from this analysis of their formation at >1000°C.

6
7 343 Based on the chromatographic and combustion results from the Arlington Canyon samples, we
8
9 344 conclude that much of this glassy carbon was likely produced as solidified tar. Tar is produced during
10
11 345 charcoalification, mostly at temperatures below 500°C (Beaumont, 1985), and this represents the typical
12
13 346 temperatures of many surface fires (Scott *et al.*, 2014). The chemistry of tars produced during this process
14
15 347 is well understood (e.g., Ku and Mun, 2006).

16 348
17 349 *What we can and cannot say about charcoal in fluvial sediments at Arlington Canyon.*

18
19 350 *Quantity of charcoal* - The quantity of charcoal in any one sample from fluvial sediments is not
20
21 351 indicative of the size of a fire. The amount of charcoal depends on the amount of charred litter, as most
22
23 352 macroscopic charcoal comes from the charring of surface-dwelling plants and litter from low-temperature
24
25 353 surface fires (Scott, 2010). In addition, charcoal can be locally concentrated in some facies (Glasspool and
26
27 354 Scott, 2013). After the Hayman fire in Colorado in 2002, charcoal was transported out of the fire-affected
28
29 355 area by flooding rivers. One downstream channel was filled with several metres of charcoal (see Fig. 9c
30
31 356 of Scott, 2010), which was not indicative of the size of the fire but rather of taphonomic processes.

32 357 *Local or regional fire* – Large charcoal fragments may be transported a considerable distance. Large
33
34 358 >1 cm pieces of charcoal may be transported down rivers and into marine sediments (e.g., Nichols *et al.*,
35
36 359 2000; Scott, 2010). Un-charred and charred plants have different hydrodynamic qualities, as do different
37
38 360 plant organs and charcoal formed at different temperatures (e.g., Nichols *et al.*, 2000; Scott, 2010; Scott *et*
39
40 361 *al.*, 2014). It is reasonable to infer that a fire was local if there is charcoal from a variety of plants, of a
41
42 362 range of sizes and varying from charred to un-charred.

43 363 *Intensity, severity or type of fire* - There has been much confusion of the terms “fire intensity”, “fire
44
45 364 severity” and “burn severity” (Keeley, 2009). Fire intensity refers to the total energy released by a fire and
46
47 365 not the energy release rate. Fire intensity data do not provide information on the temperature of the fires
48
49 366 or surface fire conditions. It is not possible to determine fire intensity simply from the amount of charcoal.
50
51 367 Fire severity refers to the extent of loss or damage to vegetation, which again cannot be determined from
52
53 368 charcoal assemblages. It is possible to obtain some temperature data from the measurement of the
54
55 369 reflectance of charcoal (Scott, 2010), and charcoal temperature profiles may help distinguish the
56
57 370 occurrence of ground, surface, or crown fires (Scott *et al.*, 2014; McParland *et al.*, 2009; Hudspith *et al.*,
58
59 371 2014). Ground fires, as opposed to surface fires tend to destroy the vegetation, with little charcoal
60
372 remaining. Crown fires can reach higher temperatures than most surface fires.

1
2 373 *Vegetation affected by wildfire* – an important feature of charcoal is that it retains anatomical
3 374 information that allows taxonomic identification (Scott, 2010). The charcoal from the Arlington section is
4
5 375 mainly from coniferous and angiosperm secondary wood and indicates that a forested ecosystem was
6
7 376 affected by wildfire. However, small axes of herbaceous angiosperms and shrubs suggest that fire on this
8
9 377 landscape included mainly surface fire. The reduction of charcoal at higher levels in the Arlington
10
11 378 sequence likely results from the documented loss of most large conifers from the Northern Channel
12
13 379 Islands by the end of the Pleistocene (Anderson *et al.*, 2010). Grasslands produce much smaller inputs of
14
15 380 charcoal (Bond, 2015).

16 17 381 18 382 *What can and cannot be interpreted from organic fractions*

19 383 The occurrence of fungal sclerotia tells us little about the environment of deposition, and less about
20
21 384 fire regime. They are common in many soils and especially those of temperate and arctic-alpine climatic
22
23 385 zones (Sakagami and Watanabe, 2009). However, more sclerotia are formed during periods of water
24
25 386 stress, so there may be some indication of rainfall variability (Benedict, 2011; Fernandez and Koide,
26
27 387 2013). The sizes of coprolites that are composed of plant material may also indicate the occurrence of
28
29 388 mites, springtails and millipedes, all found in decaying plant litter (e.g., Chaloner *et al.*, 1991; Scott *et al.*,
30
31 389 1992), or of termites, which tend to be found in somewhat drier environments (Harris, 1971).

32 390 33 391 34 35 392 *Dating*

36
37 393 Eleven radiocarbon dates were obtained from site III primarily from charcoal fragments and also from a
38
39 394 piece of uncharred wood (see Table 3). All new dates are shown calibrated using the IntCal13 calibration
40
41 395 curve (Reimer *et al.*, 2013) using OxCal v4.2.4 (Bronk and Lee, 2013) (see Table 3). The oldest age
42
43 396 returned was 14,080-14,500 cal BP, and the youngest age 12,710-12,850 cal BP (see Fig. S2c). These new
44
45 397 chronological data are consistent with the radiocarbon dates presented in Kennett *et al.* (2008) from the
46
47 398 same locality. However radiocarbon dates on charcoal fragments from elsewhere in Arlington Canyon and
48
49 399 from similar deposits in neighbouring canyons shows deposition and fire activity as early as 29,222–
50
51 400 28,394 cal BP (Pinter *et al.*, 2011), with charcoal diminishing in quantity higher in the Arlington
52
53 401 sequence, but on-going into the Holocene (Anderson *et al.*, 2010). Indeed the distribution of charcoal
54
55 402 through Arlington Canyon clearly indicates a record of more than one fire event, as shown in both the
56
57 403 wider chronological and sedimentological evidence (Hardiman *et al.*, 2016). These data are inconsistent
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59 404 with the single, catastrophic impact-induced ignition interpreted by Firestone *et al.* (2007), Kennett *et al.*
60
61 405 (2008), and other YDIH proponents.

Discussion

Like many Quaternary deposits, the fluvial sequence in Arlington Canyon contains a significant quantity and range of organic material, much of which has been charred. Abundant charcoal implies the occurrence of fire, but whether these fires were started by lightning, humans, or extraterrestrial impact requires additional lines of evidence (Hardiman *et al.*, 2016; Scott *et al.*, 2016).

Arlington Canyon has featured centrally in results suggesting a global-scale impact drove broad changes at the onset of the Younger Dryas (the YDIIH). Wittke *et al.* (2013) assert that we did not study the same section as theirs (AC003). This is not true. While Kennett *et al.* (2008, 2009b) gave UTM coordinates without specifying which datum or map projection was used, we were able to navigate to their published location using the North American Datum 1983 (NAD83) and found there the largest, best exposed, and most accessible outcrop in Arlington Canyon. Later we surmised that Kennett *et al.* (2008, 2009b) had used NAD27 (confirmed in Wittke *et al.*, 2013). We subsequently measured, sampled, and dated the small section at that location.

We have described, analyzed, and sampled sequences in Arlington and in other canyons on Santa Rosa Island, which include material ranging in age from ~29,000 cal a BP to ~5,000 a BP (Scott *et al.*, 2010; Pinter *et al.*, 2011; Hardiman *et al.*, 2016). We continue to be puzzled why YDIIH proponents have focused extraordinary attention on one single age horizon in one <5 m section, when such a broad range of deposits and ages are represented in the surrounding area (see Hardiman *et al.*, 2016). We show from our lithological logging and analysis that there was not an ‘impact horizon’ as claimed.

Carbonaceous materials from Arlington Canyon do not require extraterrestrial input or ignition, or in some cases preclude such an event. Carbonaceous spherular forms (‘carbon spherules’) and coprolites (‘carbon elongates’) occur in multiple samples from multiple horizons on Santa Rosa Island and on neighboring islands and from sites throughout the world. They occur in sediments of a wide range of ages, from well before the Younger Dryas to well after (e.g. Anderson *et al.*, 2010; Scott *et al.*, 2010) (Table 1). Many of the carbonaceous spherular forms have features identical to those of fungal sclerotia. None of the samples or morphologies observed to date require catastrophic high-temperature combustion or other extraterrestrial influence. Many of the ‘carbon elongates’ are demonstrated to be arthropod faecal pellets (Fig. S8); those with hexagonal morphology are identified as termite frass (see Scott, 1992).

Many YDIIH proponents repeatedly use glassy carbon as an indicator of high-temperature fires (Firestone *et al.*, 2007; Kennett *et al.* 2008; Bunch *et al.* 2012; Wittke *et al.*, 2013; Kinzie *et al.*, 2014). Most glassy carbon is in fact produced as solidified tars from a low- to medium-temperature charring process, as shown here, being common in fires of those temperatures. This has also been referred to as vitreous

1
2 440 charcoal, glassy charcoal, etc. by numerous authors and was demonstrated by McParland *et al.* (2010) to
3 441 be of low-temperature origin. None of the carbon forms from Arlington Canyon yield evidence of higher-
4
5 442 than-normal burning temperatures.

6
7 443 Wood charcoal is abundant in lower portions of the Arlington Canyon sequence, including from
8
9 444 deposits both older and younger than the Younger Dryas. Charcoal distribution in fluvial sediments is
10 445 strongly influenced by taphonomic processes, so that the type and quantity of charcoal varies both
11
12 446 laterally and vertically. The number of charcoal particles per unit volume or weight of sediment samples
13
14 447 cannot be interpreted in terms of “fire frequency” or “fire intensity”.

15
16 448 Kennett *et al.* (2008, 2009b) repeat the narrative from Firestone *et al.* (2007) that the purported
17 449 Younger Dryas impact created intense wildfires across much of the planet, including in particular, Santa
18
19 450 Rosa Island. Marlon *et al.* (2009) found no evidence of regionally synchronous fires across North
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21 451 America, and the current study finds no evidence of high-temperature fires in Arlington Canyon. The
22
23 452 occurrence of 'carbon spherules' does not indicate high temperature. Spherules and charcoal from AC003
24 453 had low reflectance, typical of low-temperature surface fires. Wittke *et al.* (2013) claim to have produced
25
26 454 spheres from high-temperature experiments involving combusting wood (their Fig. 8). However, these are
27
28 455 not carbon spheres but rather are inorganic in composition, comprising aluminium and silica and are not
29
30 456 relevant to the origin of the carbonaceous spherules.

31
32 457 The occurrence of nanodiamonds, particularly the hexagonal 2H polytype lonsdaleite, in Younger
33 458 Dryas boundary sediments is considered by YDIH proponents as among the strongest evidence of impact
34
35 459 shock processing of the crust. We have demonstrated elsewhere (Daulton *et al.*, 2016) that the
36
37 460 observations and interpretations were erroneous.

38
39 461 We conclude that YDIH proponents fail to explain the broad discrepancies between their
40 462 interpretations and the findings of independent researchers. Contrary evidence is ignored, and a broad
41
42 463 range of evidence is twisted to fit the YDIH. On Santa Rosa Island (Pinter *et al.*, 2011) as well as other
43
44 464 California Channel Islands (Pigati *et al.*, 2014), widespread and frequent fires occurred both before and
45
46 465 after the onset of the Younger Dryas, recording predominantly low-temperature surface fires.
47
48 466 Stratigraphic concentrations of charcoal are related to the nature of the original fires but also to how much
49 467 litter there was to char and a wide range of other taphonomic as well as transportation and depositional
50
51 468 processes. The sediments in Arlington Canyon lack evidence for meteoritic/cometary material from an
52
53 469 impact in North America, evidence of associated impact processes, and evidence of impact generated fires
54
55 470 (see also comments by Boslough *et al.*, 2013).

56 57 58 471 **Conclusions**

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60

1
2 473 Fluvial deposits in Arlington Canyon, Santa Rosa Island, and material in those deposits document a
3 474 long-term and mostly gradual evolution of the Arlington palaeo-landscape since the latest Pleistocene.
4
5 475 This was driven by some combination of climate change, post-glacial sea-level rise, climate-driven
6
7 476 vegetation changes, extinction of the local megafauna (*Mammuthus exilis*), and the arrival and subsequent
8
9 477 expansion of human activities (e.g., Rick *et al.*, 2014). These changes have driven a long-term shift in fire
10
11 478 regimes. The size range of the charcoal fragments in the latest Pleistocene to Holocene sediments from
12
13 479 Arlington Canyon, as well as the presence of charred and non-charred plant material, suggests a surface
14
15 480 fire regime, with charcoal moved to the stream by overland flow and subsequent fluvial transport. This
16
17 481 range of material, together with SEM and reflectance analyses, indicate low-temperature surface-fire
18
19 482 regimes of coniferous and mixed coniferous/angiosperm forests. The distribution of charcoal in the
20
21 483 sequence suggests multiple fire events through the record. We find no evidence for a single, high-intensity
22
23 484 crown fire, nor any evidence of the kind of catastrophic, transformative fire event proposed in the YDIH.

24
25 485 Carbonaceous spherules recorded by Kennett *et al.* (2008) are predominantly fungal sclerotia, and
26
27 486 ‘carbon elongates’ are predominantly arthropod coprolites; those with hexagonal cross sections probably
28
29 487 are termite frass. Glassy carbon present in these deposits formed from the precipitation of tars during the
30
31 488 charcoalification process. None of these materials indicate high temperatures. The presence of
32
33 489 nanodiamonds in Arlington Canyon spherules has not been confirmed by independent studies, and we
34
35 490 find no evidence of nanodiamonds. Material identified as lonsdaleite at Arlington Canyon by Kennett *et*
36
37 491 *al.* (2009b) is inconsistent with the lonsdaleite structure and more consistent with polycrystalline
38
39 492 aggregates of graphene and graphane (see Daulton *et al.*, 2010, 2016). None of the evidence supports the
40
41 493 contention that there is an impact horizon in the Arlington sequence. By extension, our research suggests
42
43 494 that similar problems may exist at other sites supporting the purported Younger Dryas impact.

44
45 495 *Acknowledgements.* We thank staff with Channel Islands National Park. ACS thanks Sharon Gibbons and
46
47 496 Neil Holloway for technical support. We also thank Tom Higham of the Oxford Accelerator Unit for
48
49 dates and for his comments on the manuscript. This research was supported by grants from the National
50
51 Geographic Society (8321-07) to NP, from the National Science Foundation (EAR-0746015) to NP and
52
53 RSA, and a Leverhulme Emeritus Fellowship (EM-2012-054) to ACS.
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Figure Captions

Figure 1. Map of Santa Rosa showing location of Section III (AC 003 of Kennett *et al.*, 2008).

Figure 2. Detailed lithological logs of site III, Arlington Canyon, showing site in 2010 (above) and 2013 (below).

Figure 3. Organic fractions from sieved samples, Site III Arlington Canyon. Images (a,c,e) are reflected light under water; images (b,d,f) are dark field images of same samples highlighting charred and uncharred plant material. (a,b) Sample 10-56 Section IIIa mid section. The image shows large uncharred wood fragments (brown) with wood charcoal (black) and coprolites. (c,d) Sample 10-56 Section IIIa mid section. (e,f) Sample 10-56 Section IIIa mid section. (g) Large charcoal fragments, sample SRI-13-19, section IIIf below mid section. (h) Specimen of wood charcoal shown in g and put into water showing fragmentation. (i) Glassy carbon, sample SRI-10-56, section IIIa. (j) Carbonaceous spherules, sample SRI-10-56, section IIIj. (k) Carbon elongates (coprolites). Sample SRI-10-56, section IIIa

Figure 4. Charcoal from sediments from Site III, Arlington Canyon. (a) Scanning Electron Microscopy of conifer secondary wood, SRI-10-65, section IIIc. (b) Detail of image a showing rays and ray pits. (c) Detail of image a showing growth ring. (d) Conifer leafy shoot, cf *Cupressus* sp., SRI-13-11, section IIIc. (e) Conifer needle, *Pinus* sp., SRI-13-21. (f) Angiosperm secondary wood, SRI-13 core IIID, 37 cm from base. (g) Detail of image f showing vessels. (h) Angiosperm secondary wood, SRI-13-21. (i) Detail of image h showing multiseriate rays. (j) Small angiosperm axis, SRI-13 core IIID, 37 cm from base. (k) Detail of image j showing vessels.

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Table 3. Radiocarbon dates obtained from Site III, Arlington Canyon, Santa Rosa Island, CA and used in this study.

Table 1. Distribution of charcoal and other organic materials from site III Arlington Canyon.
A=Abundant; C=Common; F=Frequent, R=Rare, P=Present

Sample No.	Section	Height above base (cm)	Charcoal									Uncharred	
			>1 cm	5mm-1cm	<5mm	<1mm	1-2mm axes	Leaves	Spherules	Coprolites	Glassy carbon	Wood	
SRI-10-47	IIIA	10-12		C		C	R	P		C			
SRI-10-48	IIIA	20-22			A	C	R	R		C			
SRI-10-49	IIIA	27				R							
SRI-10-50	IIIA	69				R							
SRI-10-51	IIIA	53			C		R		R	R	R		
SRI-10-52	IIIA	62	P				P			A			
SRI-10-53	IIIA	72					P			R	R		
SRI-10-54	IIIA	88	C				R			R			
SRI-10-55	IIIA	95	A				R		R	A	P		
SRI-10-56	IIIA	118-120	A	C	C	F				A		C	
SRI-10-57	IIIA	131	C	C	F	F		R		R	C	C	
SRI-10-58	IIIA	147-148	A	F	F	F			R	R		C	
SRI-10-59	IIIA	197-198				F		R		R			
SRI-10-60	IIIA	260											
SRI-10-61	IIIA	131		F					R			F	
SRI-10-62	IIIA	131	R				R			R			
SRI-10-63	IIIA	215	P	A			F	P		C			
SRI-10-65	IIIB	55		A	F	F				F			
SRI-10-66	IIIB	83				R				R			
SRI-10-67	IIIB	110			R					R			
SRI-13-01	IIID	118-120			A	A			R	C			
SRI-13-02	IIID	135-140		A	A		C						
SRI-13-03	IIID	154-156				F	R						
SRI-13-04	IIID	200-202			A		F	R	R	R			
SRI-13-05	IIID	210-212		C	C		C			A	R		
SRI-13-06	IIID	270-242			C					C			
SRI-13-07	IIID	15-17		C			F	P	R	C			
SRI-13-08	IIID	28-30		C			R						
SRI-13-09	IIID	40-42	C				F			C	R	F	
SRI-13-10	IIID	60-62		C			F		R	C			
SRI-13-11	IIIF	0-4			A			P		C		F	
SRI-13-12	IIIF	50-52	A	F	F		C		R	F		P	
SRI-13-13	IIIF	115-117			R								
SRI-13-14	IIIF	125-127			C				R	R			
SRI-13-15	IIIF	150-152		A			C			C	P		
SRI-13-16	IIIF	190-192		A			F		R	C	R		
SRI-13-17	IIIF	200-202			C		C			F			
SRI-13-18	IIIF	220-222				A	R			F	R		
SRI-13-19	IIIF	140-142	A	C								A	
SRI-13-20	IIIE	100-102	C	C			C		C	F			
SRI-13-21	IIIE	153-155		A	A					C	R		
SRI-13-22	IIIE	192-194		P	A		C			F			
SRI-13-23	IIIE	218-220			C		F			F	R		
SRI-13-24	IIIC	12-15	A	A	C				C	C			

Table 2. List of compounds detected by pyrolysis-gas chromatography-mass spectrometry of glassy carbon and charcoals. "Ret Time" is retention time and reflects size of molecules going through the mass spectrometer.

	RetTime	compound	SEQUOIA350		SEQUOIA450		SEQUOIA600		AC003		SRI-10-36		SRI-10-57	
			PeakArea	%	PeakArea	%	PeakArea	%	PeakArea	%	PeakArea	%	PeakArea	%
1	3.67	benzene	3870640	3.16	7580848	15.19	40095737	61.92	3269558	13.35	5264142	48.17	10828341	47.92
2	4.82	toluene	7961682	6.51	11875708	23.79	9754648	15.06	6938243	28.33	1952563	17.87	3464002	15.33
3	6.36	ethylbenzene	221024	0.18	466479	0.93	1496316	2.31	601975	2.46	190718	1.75		0.00
4	6.53	xylene	1531400	1.25	2302114	4.61	704280	1.09	1350791	5.52	291081	2.66	93810	0.42
5	6.94	styrene	975975	0.80	1184472	2.37	4603983	7.11	1218587	4.98	390031	3.57	324752	1.44
6	8.62	phenol	9613598	7.86	8787116	17.60		0.00	2488814	10.16	430317	3.94	497775	2.20
7	8.90	benzotrile		0.00	101814	0.20		0.00	954222	3.90		0.00	646202	2.86
8	9.09	benzofuran	1187128	0.97	1338316	2.68		0.00	991371	4.05		0.00		0.00
9	10.17	2-methylphenol	2085471	1.70	1878744	3.76		0.00	1838438	7.51		0.00		0.00
10	10.63	4-methylphenol	4177201	3.41	3869322	7.75		0.00	2038107	8.32		0.00		0.00
11	13.05	naphthalene	1902648	1.55	4387767	8.79	5362078	8.28	848406	3.46	1386444	12.69	3971877	17.58
12	15.19	methylnaphthalene	820391	0.67	1924983	3.86	334955	0.52	478329	1.95	114184	1.04	320448	1.42
13	15.48	methylnaphthalene	300671	0.25	815107	1.63	286347	0.44	349751	1.43	99494	0.91	172470	0.76
14	16.69	acenaphthene		0.00	438952	0.88		0.00	178534	0.73	248839	2.28	718862	3.18
15	19.10	dibenzofuran	849154	0.69	1381920	2.77	276435	0.43	262903	1.07	70450	0.64	435839	1.93
16	23.22	phenanthrene	355260	0.29	921184	1.85	399201	0.62	113065	0.46	256888	2.35	1124154	4.97
17	23.51	anthracene	94274	0.08	2747425	5.50			42551	0.17	36582	0.33	0	0.00

Table 3 – New radiocarbon dates obtained from Site III, Arlington Canyon, Santa Rosa Island, CA in this study. (References, 1= Hardiman *et al.*, 2016)

¹⁴ C publication code	Site signifier	Sample number	Dated material	δ ¹³ C	¹⁴ C age (yr BP)	¹⁴ C age error (1σ)	Reference
UCIAMS-84951	IIIa	SRI-10-63	Charcoal	-	11005	25	1
UCIAMS-84950	IIIa	SRI-10-63	Charred coprolites	-	11755	30	1
UCIAMS-84949	IIIa	SRI-10-62	Charred twigs	-	11030	30	1
UCIAMS-84948	IIIa	SRI-10-61	Uncharred wood	-	10935	30	<i>This Study</i>
UCIAMS-84947	IIIa	SRI-10-56	Charred coprolites	-	11095	30	1
UCIAMS-84946	IIIa	SRI-10-56	Charred twigs	-	11035	30	1
UCIAMS-84945	IIIa	SRI-10-52	Charred twigs	-	11000	25	1
UCIAMS-84944	IIIa	SRI-10-47	Charred twigs	-	12310	30	1
UCIAMS-84943	IIIa	SRI-10-47	Charred coprolites	-	11885	30	1
OxA-29224	III f	SRI-13-11	Small charred axis	-24.55	11130	50	1
OxA-29225	III f	SRI-13-11	Small charred axis	-24.62	11085	50	1

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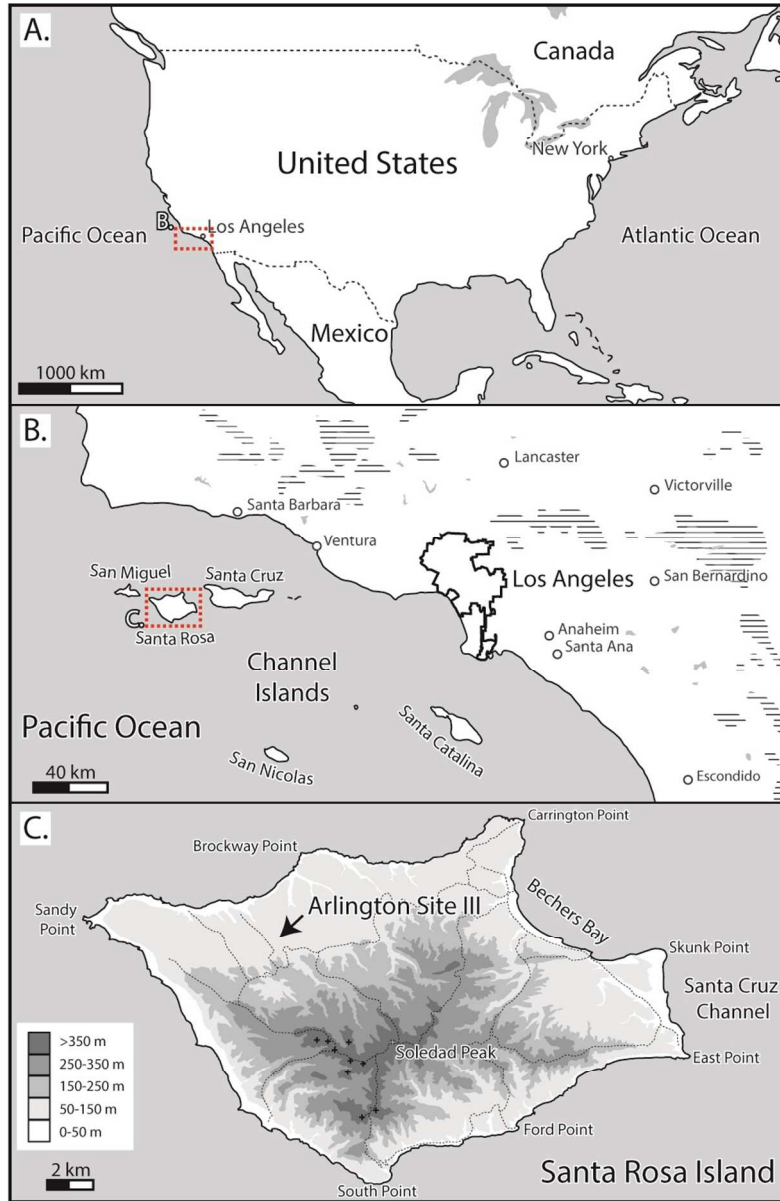


Figure 1. Map of Santa Rosa showing location of Section III (AC 003 of Kennett et al., 2008).
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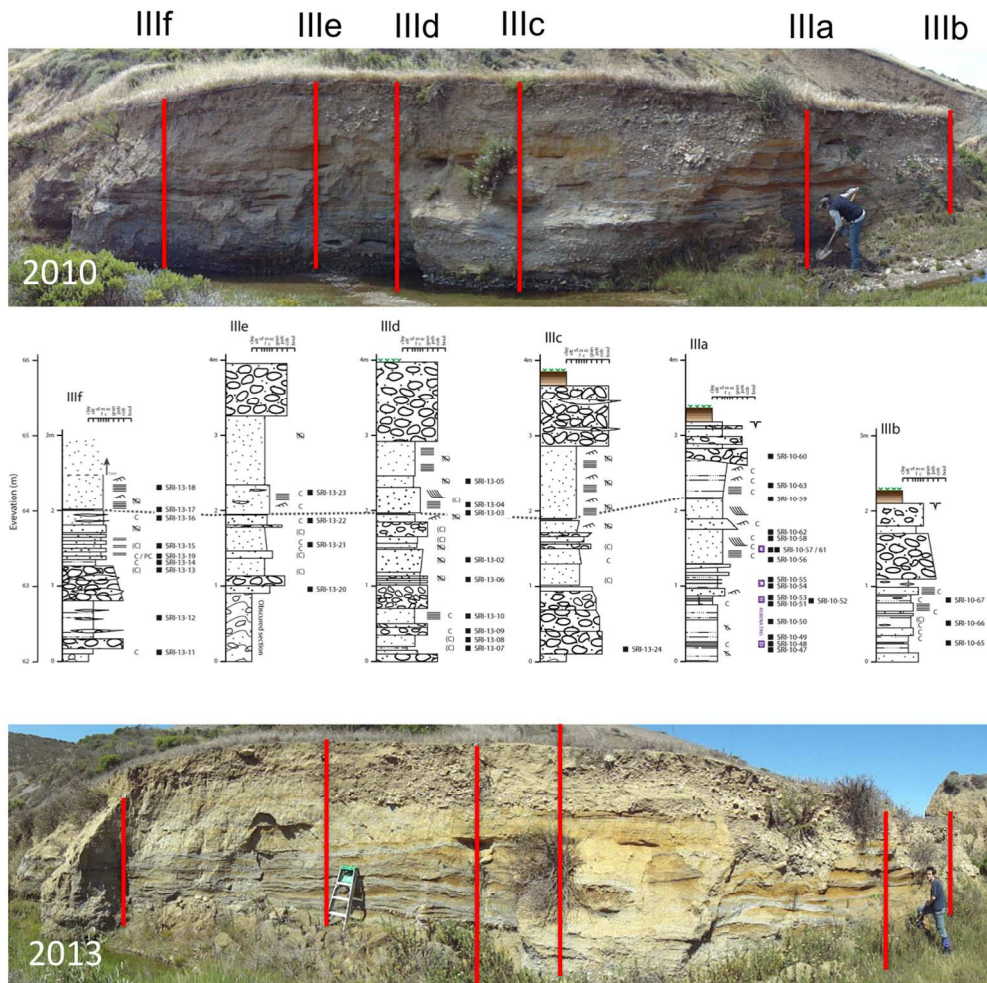


Figure 2. Detailed lithological logs of site III, Arlington Canyon, showing site in 2010 (above) and 2013 detail in the stratigraphic de 528x521mm (72 x 72 DPI)

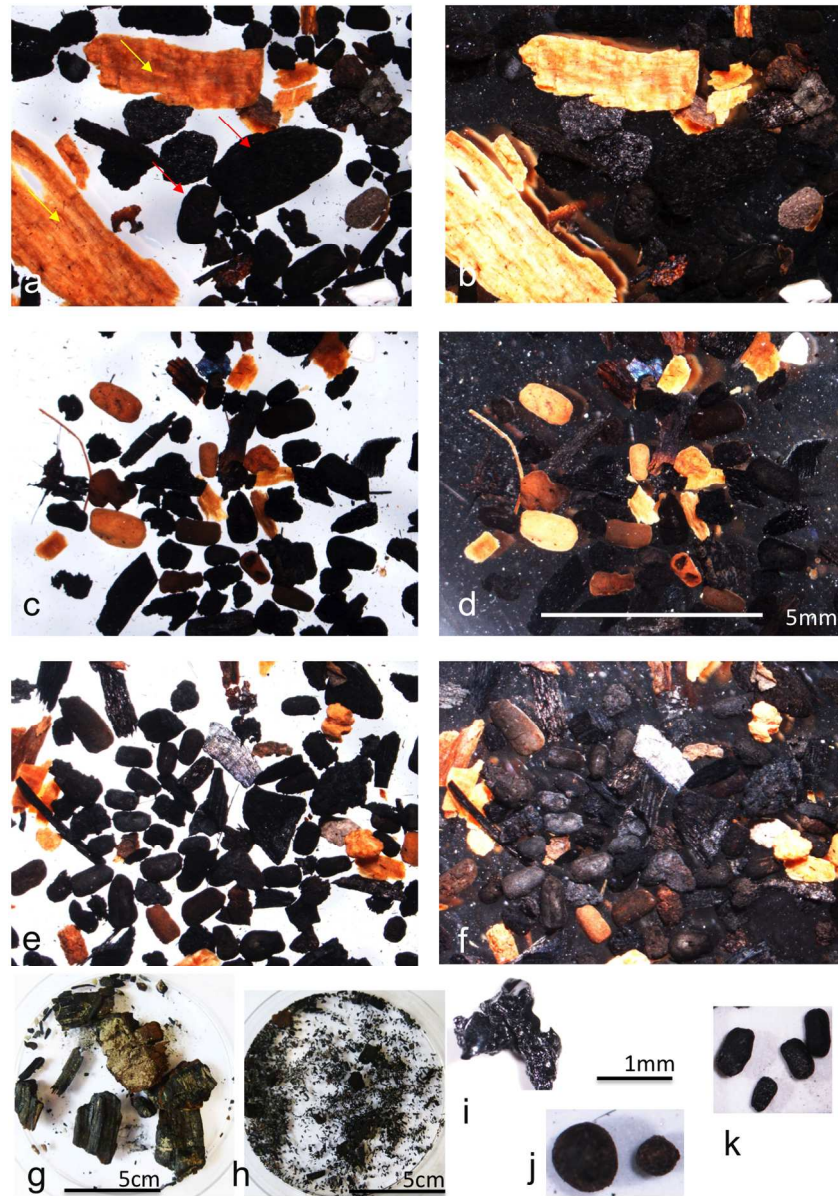


Figure 3. Organic fractions from sieved samples, Site III Arlington Canyon. Images (a,c,e) are reflected light under water; images (b,d,f) are dark field images of same samples highlighting charred and uncharred plant material. (a,b) Sample 10-56 Section IIIa mid section. The image shows large uncharred wood fragments (brown) with wood charcoal (black) and coprolites. (c,d) Sample 10-56 Section IIIa mid section. (e,f) Sample 10-56 Section IIIa mid section. (g) Large charcoal fragments, sample SRI-13-19, section IIIf below mid section. (h) Specimen of wood charcoal shown in g and put into water showing fragmentation. (i) Glassy carbon, sample SRI-10-56, section IIIa. (j) Carbonaceous spherules, sample SRI-10-56, section IIIj. (k) Carbon elongates (coprolites). Sample SRI-10-56, section IIIa
Charcoal (Figs. 3, 4; Suppleme
527x752mm (72 x 72 DPI)

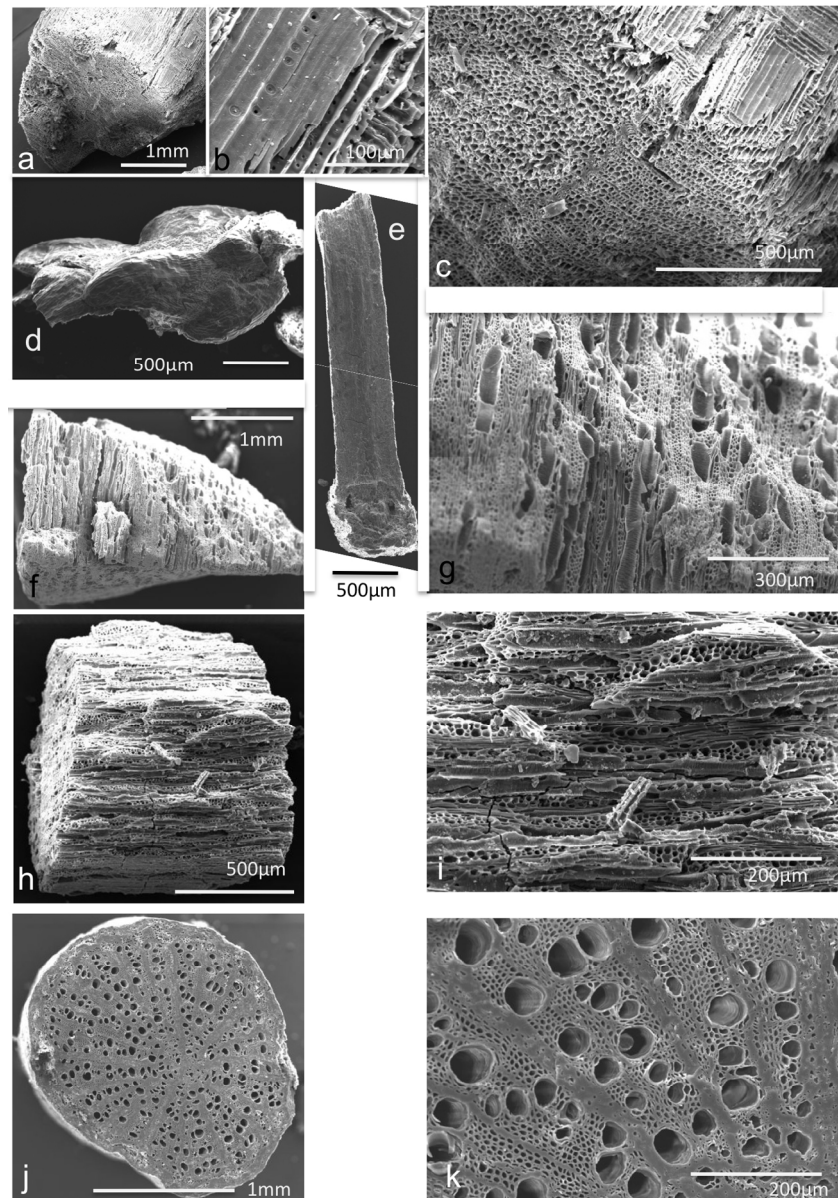


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Charcoal (Figs. 3, 4; Supplemental Fig. S1)
526x753mm (72 x 72 DPI)

Supplemental Materials

Supplemental Figure Captions

Figure S1 Photograph of outcrop of Site III, Arlington Canyon. (a) Site in 2010. (b) Detail of position of logs IIIa and IIIb. (c) Detailed log IIIa with dated samples. (d) Key to lithological logs.

Figure S2. The central part of the Site III section (AC003) of Kennett *et al.* (2008, 2009b; Wittke *et al.*, 2013) at Arlington Canyon. (a) Lithological log showing dated horizons (from Kennett *et al.*, 2009b). Note the dark layer labelled 2m from the surface. Note also the YD 'impact horizon' has been identified between 3.8 m and 5 m below surface. (b) Photograph from Wittke *et al.* (2013) supporting materials of the site identified as AC003. This clearly shows sample points and 4 of which are denoted with arrows. It also shows position of sampled section and a scale. The distance from the top to the black layer is 2m. However, the distance to the base of the section shown is just over 4m. The YD 'impact horizon' is indicated by these authors with yellow lines. However this corresponds to a depth of 3.8-4.1m. It is unclear where the units below recorded in Kennett *et al.* (2008, 2009b) are situated. (c) Photograph of the same section taken by Scott in 2010 clearly showing the Kennett section with four sample positions indicated by red arrows, as in B. The position of the claimed YD 'impact horizon' as reported by Wittke *et al.* (2013) is indicated with the yellow dots. The water in the stream is approximately 4 m from the top of the section. There is no evidence of material being accessed below the water surface. It is unclear, therefore, where the section 4-5-5.0 m occurs and it is these samples that contain most of their 'impact' markers and dated material.

Figure S3. Lateral variation of facies at the base of the section at Site III, Arlington Canyon. Section IIIId is the position of the section described in Kennett *et al.* (2008). Note the dark bed below the thin conglomerate (3-4m below surface; it is dark because it is wet. This is part of a channel complex with silts, sands and gravels. 2m laterally (section IIIc) these beds are represented by coarse gravel conglomerates that thin out only a few metres to the north (section IIIa).

Figure S4. Sediments and carbonaceous fossils from our locality III in Arlington Canyon, which is the same locality as AC003 of Kennett *et al.*, (2008, 2009b) and Wittke *et al.* (2013). (a) Typical distribution of charcoal in the mid sandy layers in the section. Note the charcoal is strongly related to sedimentary structures and not evenly distributed. Note samples taken from each of the three lines

1 would give quantitatively different results. (b) Range of charcoal sizes from very small < 1mm
2 fragments to pieces >1cm in fine sands. Note also the larger pieces break on extraction. (c) Taking
3 and recording sample of charcoal-rich sand from section IIIa. (d) Taking sediment core from base of
4 section IIIf.
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10 Figure S5. Sieved samples (>125 μ m) from Site III Arlington Canyon. (a) Sample SRI-13-11. Base of
11 section IIIf. The predominantly silty rocks also contain a significant amount of sand and pebbles.
12 The pebbles are both angular and rounded and are of a variety of rock types. (b) Pebbles from
13 sample SRI-13-17 showing rounded dark organic rich silt pebbles. Mid section IIIf. This horizon is
14 dark and can be traced across the outcrop half way up the section. (c) Range of pebble types from
15 sample SRI-13-05 showing they comprise both rounded and angular pebbles. Section III d upper part
16 of section. (d) Iron stained sand with charcoal fragments from upper sandy layer, sample SRI-13-23,
17 Section IIIe. (e) Sand, pebbles with charcoal fragments, including coprolites, and bone (brown),
18 sample SRI-13-05, section III d upper part. (f) Charcoal residue with wood fragments, small axes,
19 carbonaceous spherules and glassy carbon, sample SRI-13-11, base of section IIIf. (g) Sample 2007
20 AC003 from G. James West, sieved in water. Note brown iron staining of some of the charcoal
21 fragments and sand grains. (h) Charcoal-rich sample after dissolving mineral matter in 40% HF.
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35 Figure S6. Sediments and charcoal from Site III, Arlington Canyon. (a) Sediment sample from the base of
36 section IIIC showing fine-medium silty-sand that becomes lighter as the sediment dries. (b) Lower
37 surface of sediment sample from the base of section IIIC showing fine-medium silty-sand that
38 becomes lighter as the sediment dries. (c) Top of specimen b showing a single large charcoal
39 fragment. (d) Specimen shown in c that has been gently sieved in water through 125 μ m sieve. Note
40 that one piece can break in to many hundreds of fragments.
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48 Figure S7. Carbonaceous spherules from Arlington Canyon, Site III. (a) Carbonaceous spherules from
49 sample from G. James West collection AC003. (b) Specimen from Arlington Canyon, sample
50 2007AC-003 (collected by J. West and J.J. Johnson) (12.8-13.1 Ka) Section through spherule. (c)
51 Detail of image b showing outer rind and 'cellular' interior. (d) Carbonaceous spherules (fungal
52 sclerotia). from charcoal residue after low temperature surface fire, Thursley, Common, Surrey,
53 England. (e,f) Scanning Electron Micrographs of carbonaceous spherules from charcoal residue after
54 low temperature surface fire, Thursley Common, Surrey, England, 2006 showing features
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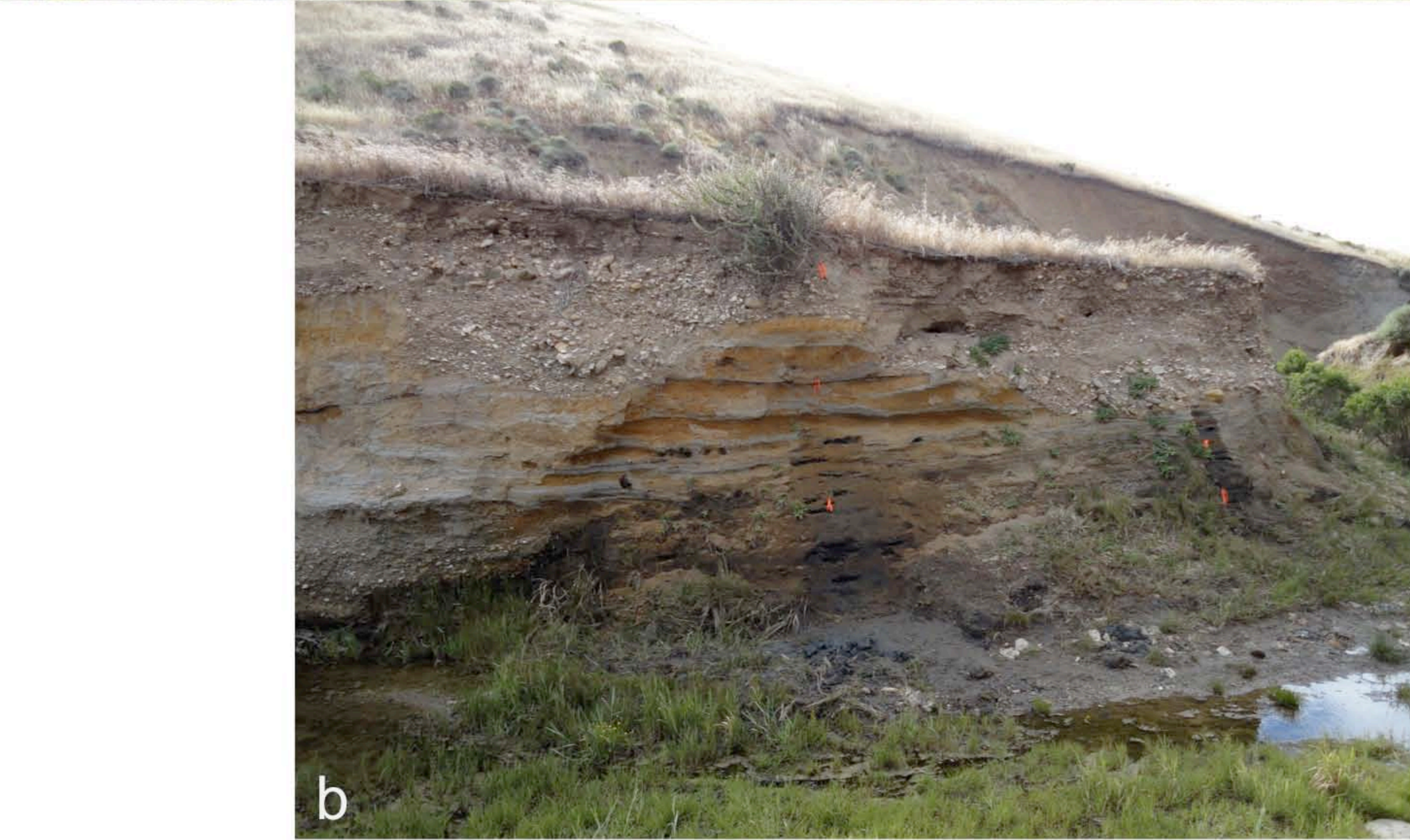
1 comparable to the Spherules in images a-c. (e) Broken spherules. (f) Detail of image e showing
2 rind, radially arranged outer cells and equi-dimensional inner 'cortical' cells and fused medullary
3 hyphae. (g-i) Scanning Electron Micrographs of sclerotia from the fungus *Conococcum geophilum*
4 Fr. In mycorrhiza with young growth of *Picea glauca* after fire event, Peace River, Canada.
5 Collected by T. Längle and photographed by A.G.Heiss. (g) Whole sclerotium. (h) Broken
6 sclerotium. (i) Broken sclerotium showing rind and fused medullary hyphae. Figures S7b and S7c
7 from Scott *et al.*, (2011). Figures S7g-i from Auxillary Material from Scott *et al.*, (2010). Note
8 Figure S7e was also published in Pinter *et al.*, (2011) and origin stated clearly.
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18 Figure S8. Coprolites ('Carbon elongates') from Arlington Canyon, Site III. (a) Charcoal fragments, carbon
19 spherule and carbon elongates (coprolites) from SRI-10-56. (b) Carbon elongates (coprolites) from
20 SRI-10-56 showing a range from uncharred (brown) to charred (black) forms. Some of the coprolites
21 show hexagonal morphology indicating their origin as termite frass. (c) Cluster of carbon elongates,
22 probable charred termite frass, sample SRI-13, core section IIID, 72-74 cm from base. (d) Scanning
23 Electron Micrograph of cluster of modern charred termite frass. (e) Scanning Electron Micrograph
24 of single termite coprolite. (f) Scanning Electron Micrograph of broken termite coprolite showing
25 hexagonal morphology. (g) Scanning Electron Micrograph of cluster of termite frass.
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33 Figure S9. Chromatograms of charcoals pyrolysed at 650C. (a) *Sequoia* charcoal made at 350C. (b)
34 *Sequoia* charcoal made at 450C. (c) *Sequoia* charcoal made at 600C. (d) Santa Rosa glassy carbon
35 sample AC003. (f) Arlington glassy carbon sample SRI-10-36. (g) Arlington glassy carbon sample
36 SRI-10-57.
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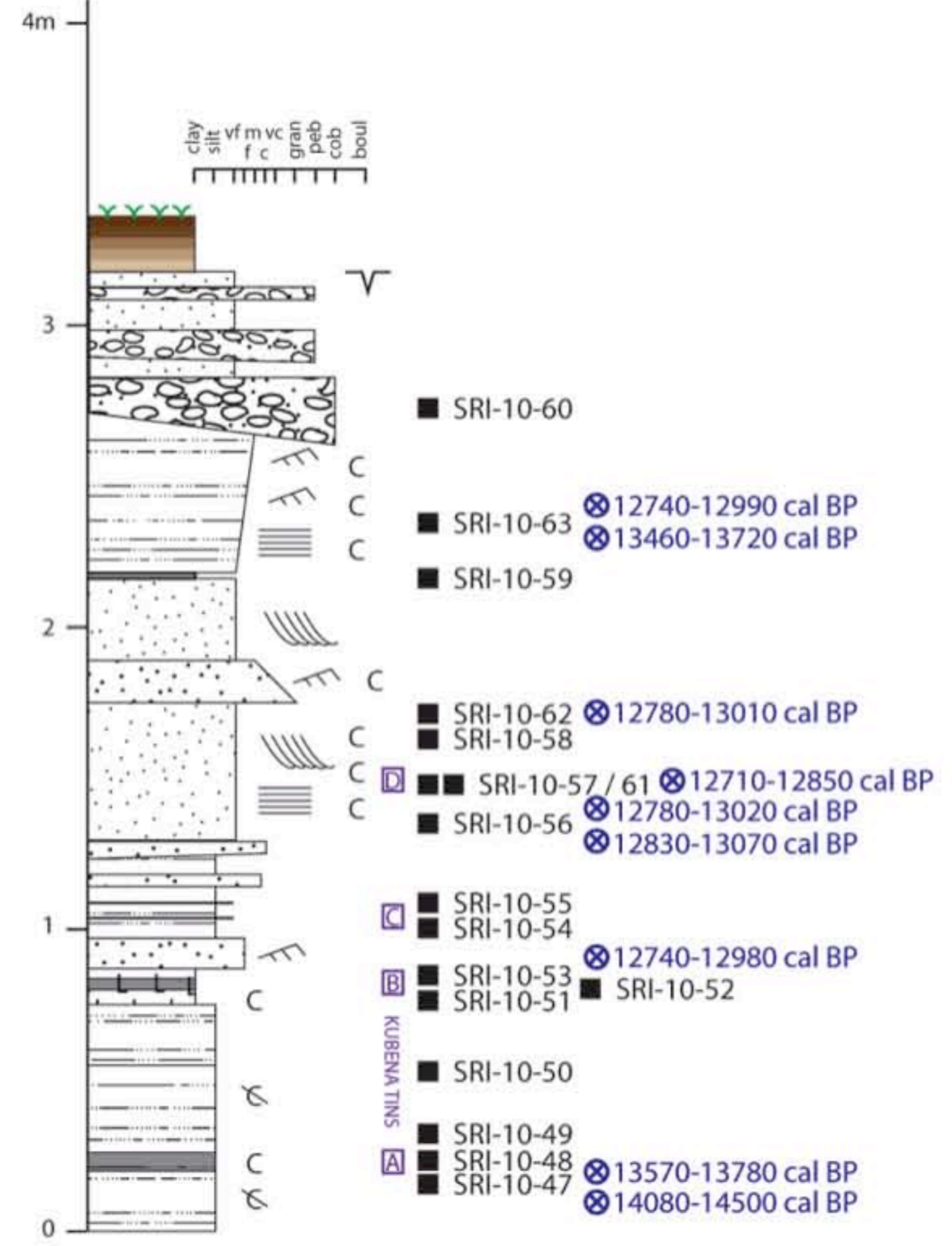
a



b



IIIa

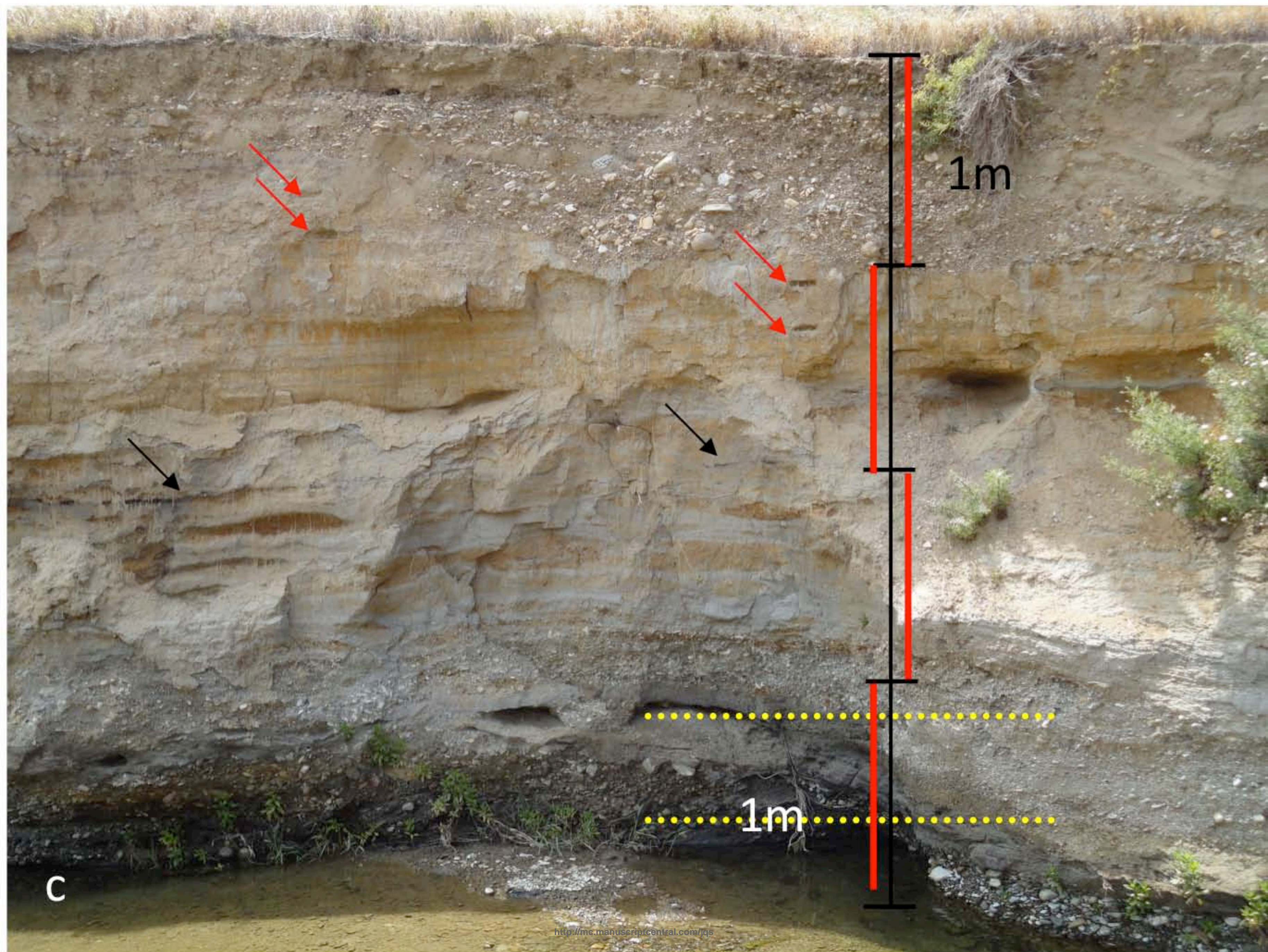
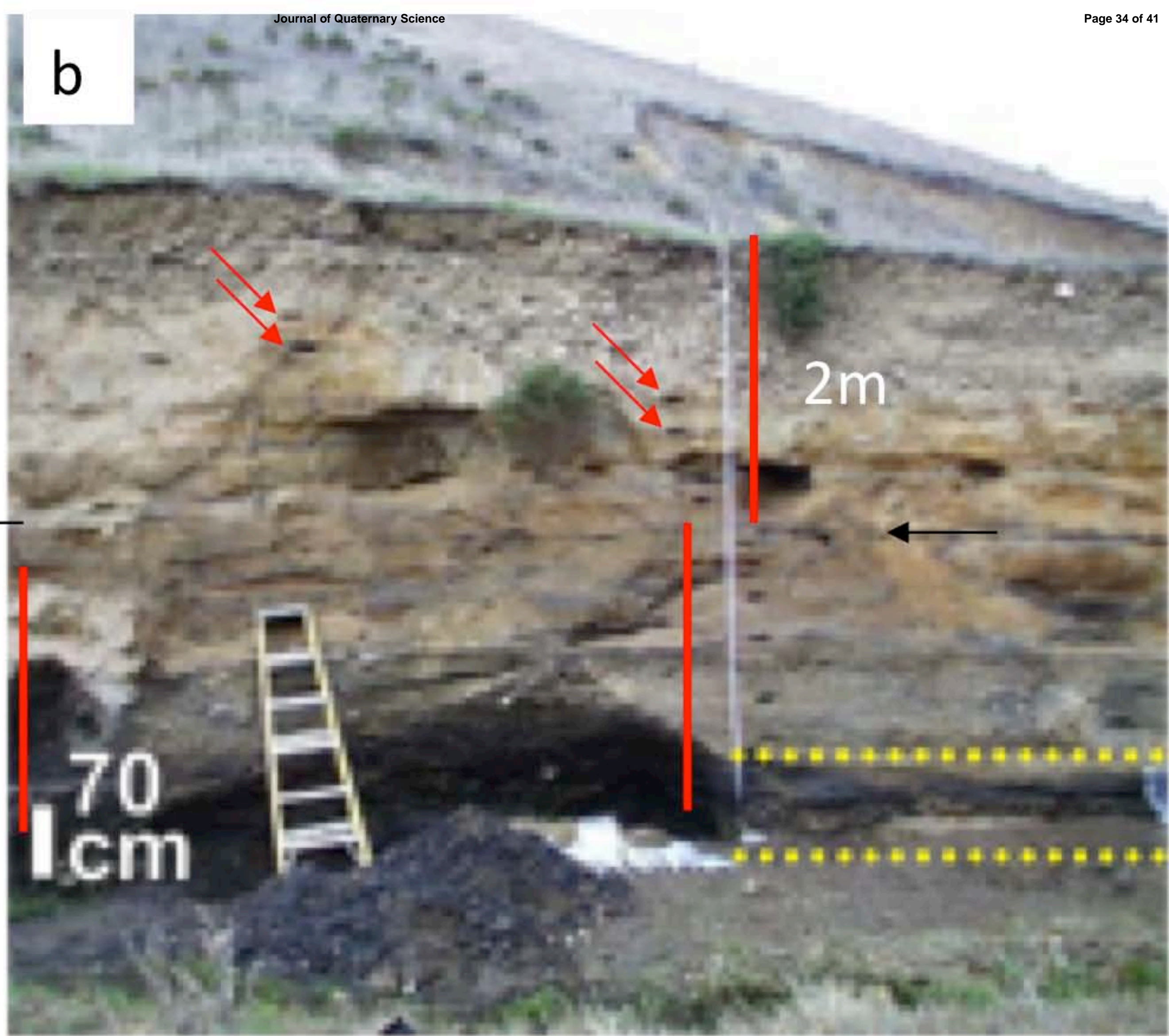
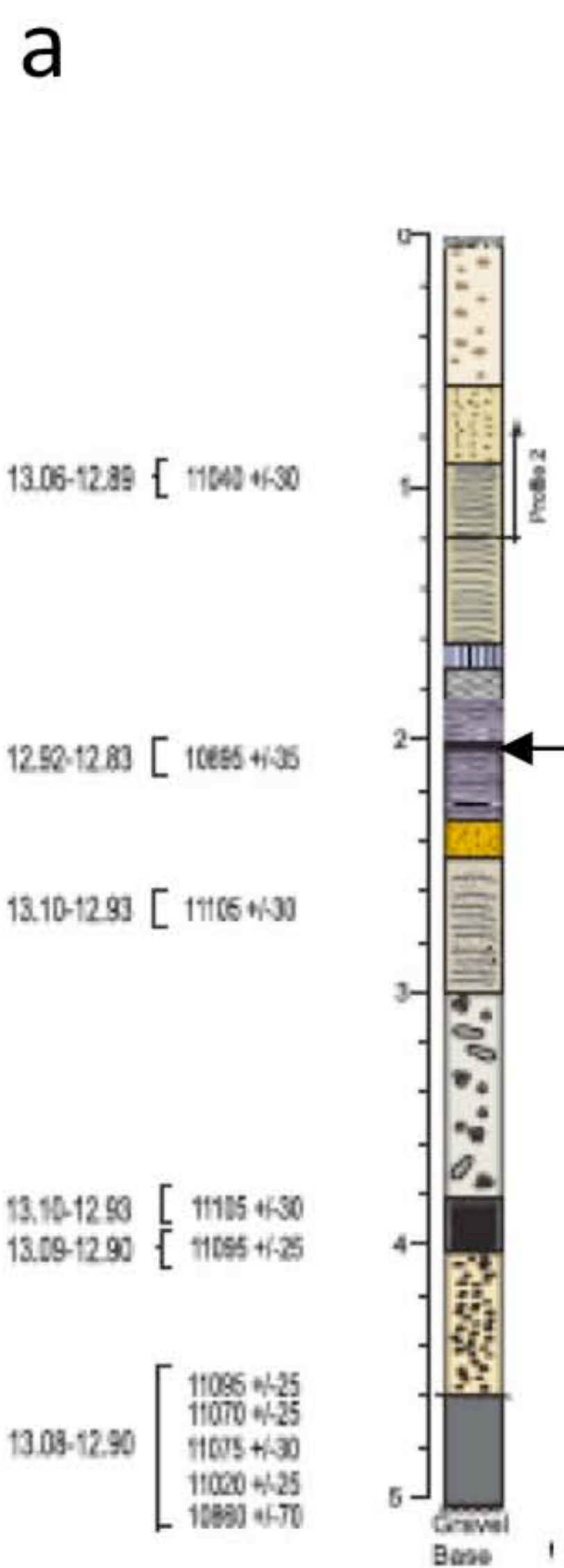


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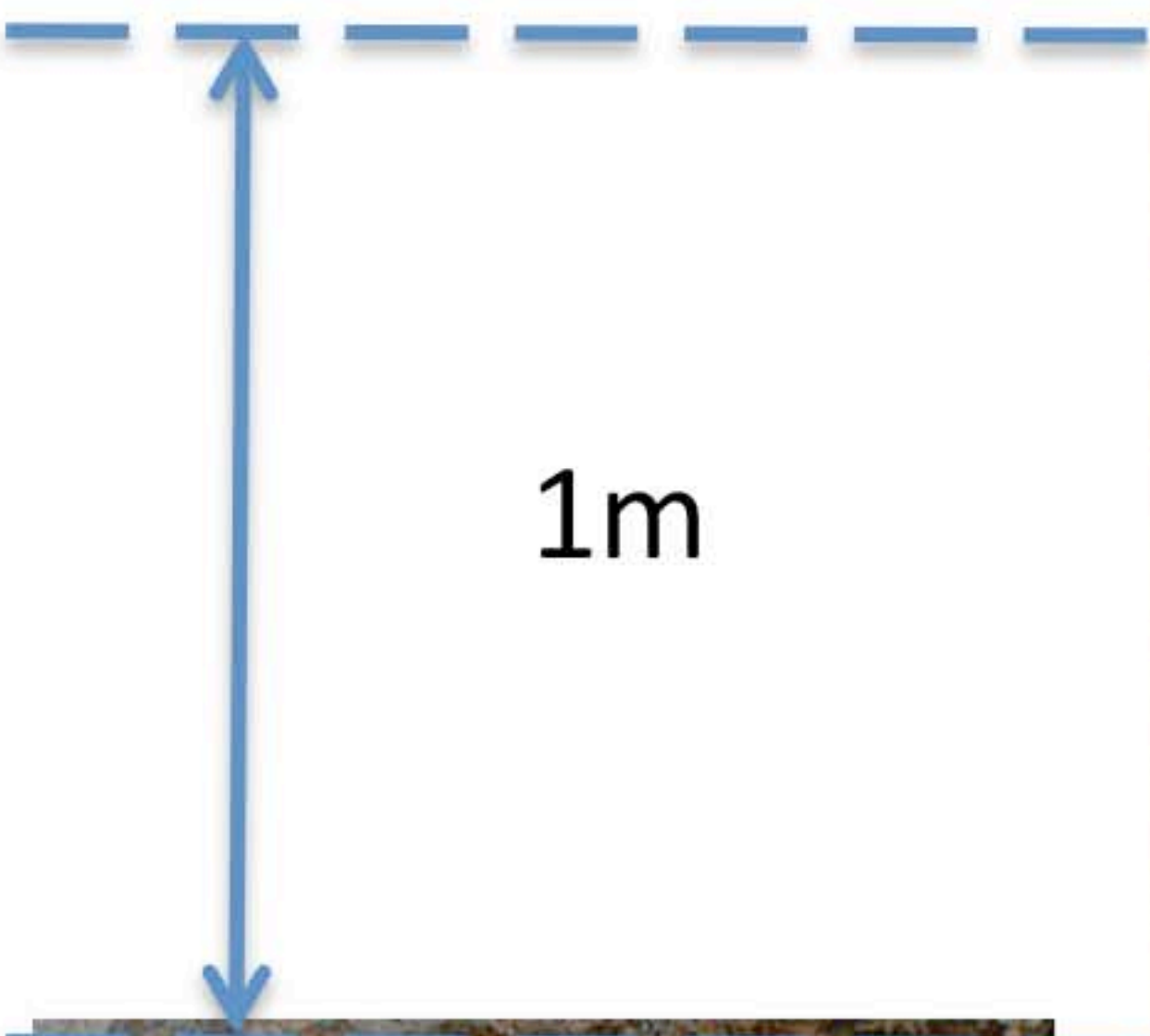
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|--|-----------|--|----------------------------------|--|---------------------------|
| | Clay | | Darker Horizons | | Laminations |
| | Silts | | Large Charcoal fragments | | Tree Roots <i>in Situ</i> |
| | Fine Sand | | Small charcoal fragments | | Radiocarbon date (cal BP) |
| | Sand | | v.rare charcoal fragments | | Sample |
| | Granules | | Mudcracks | | |
| | Pebbles | | Current ripple cross laminations | | |
| | Cobbles | | Planar cross bedding | | |
| | Boulders | | wave ripple cross laminations | | |

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III d

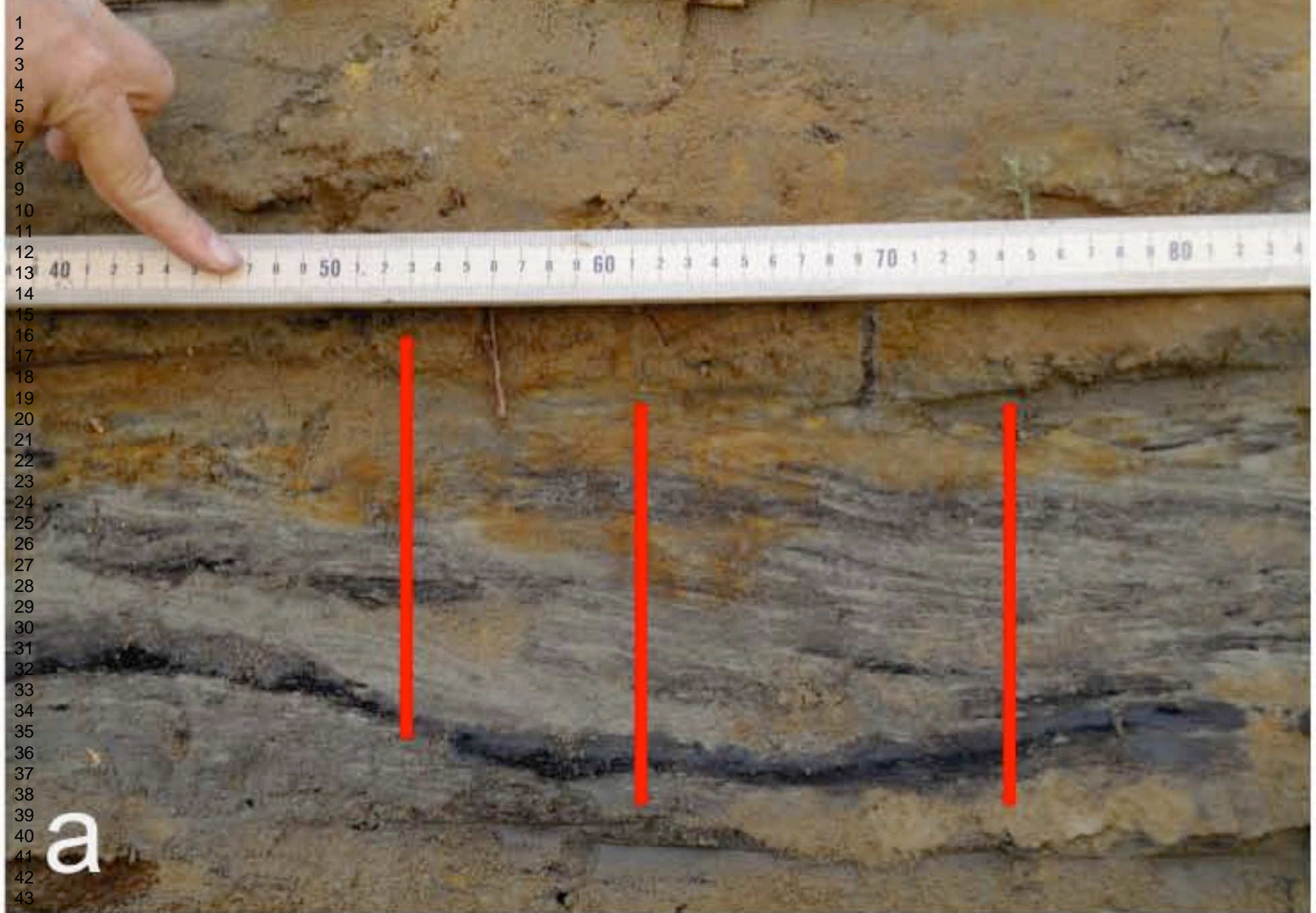
III c

III a

2m

6m



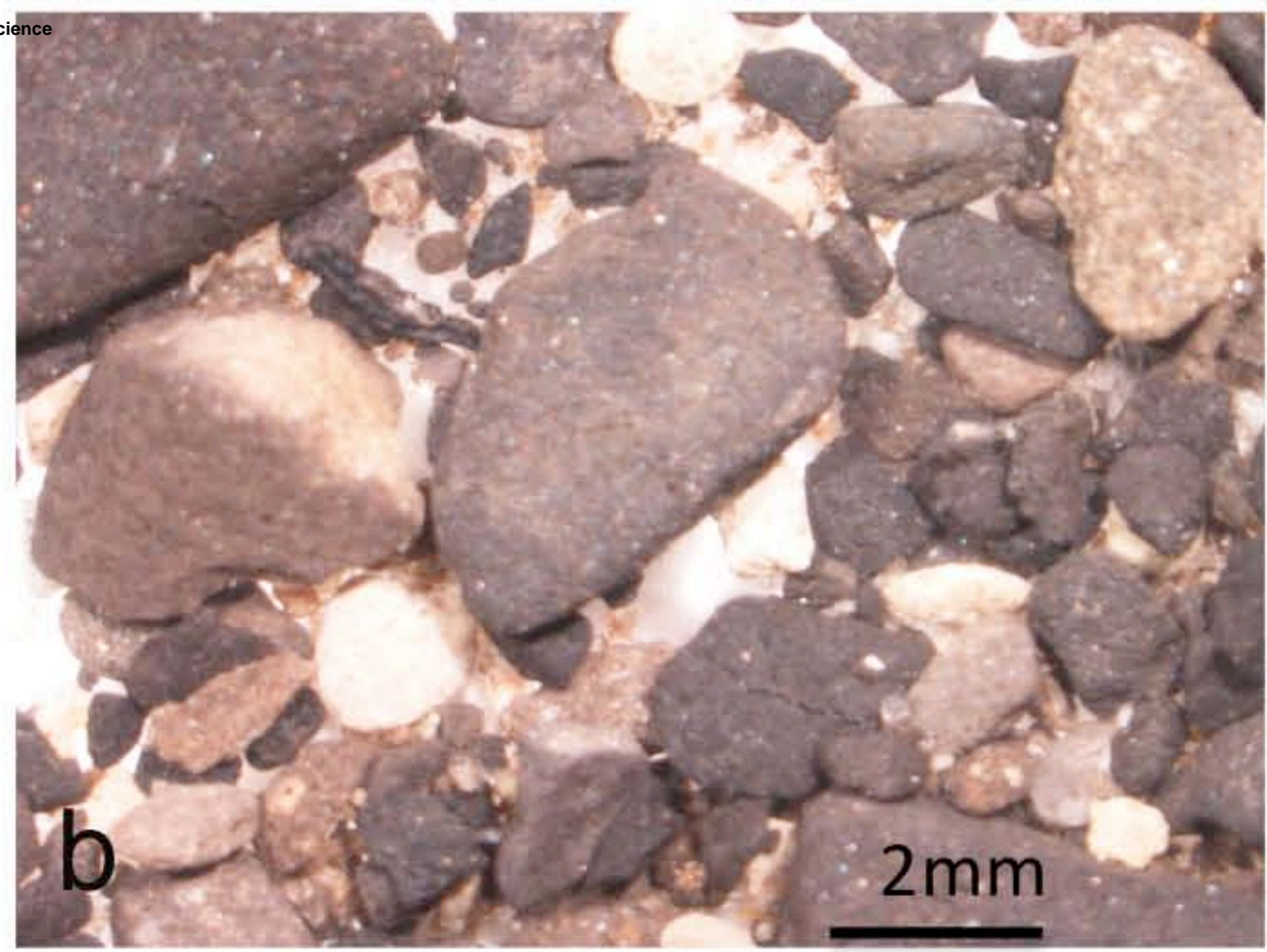


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a

5cm



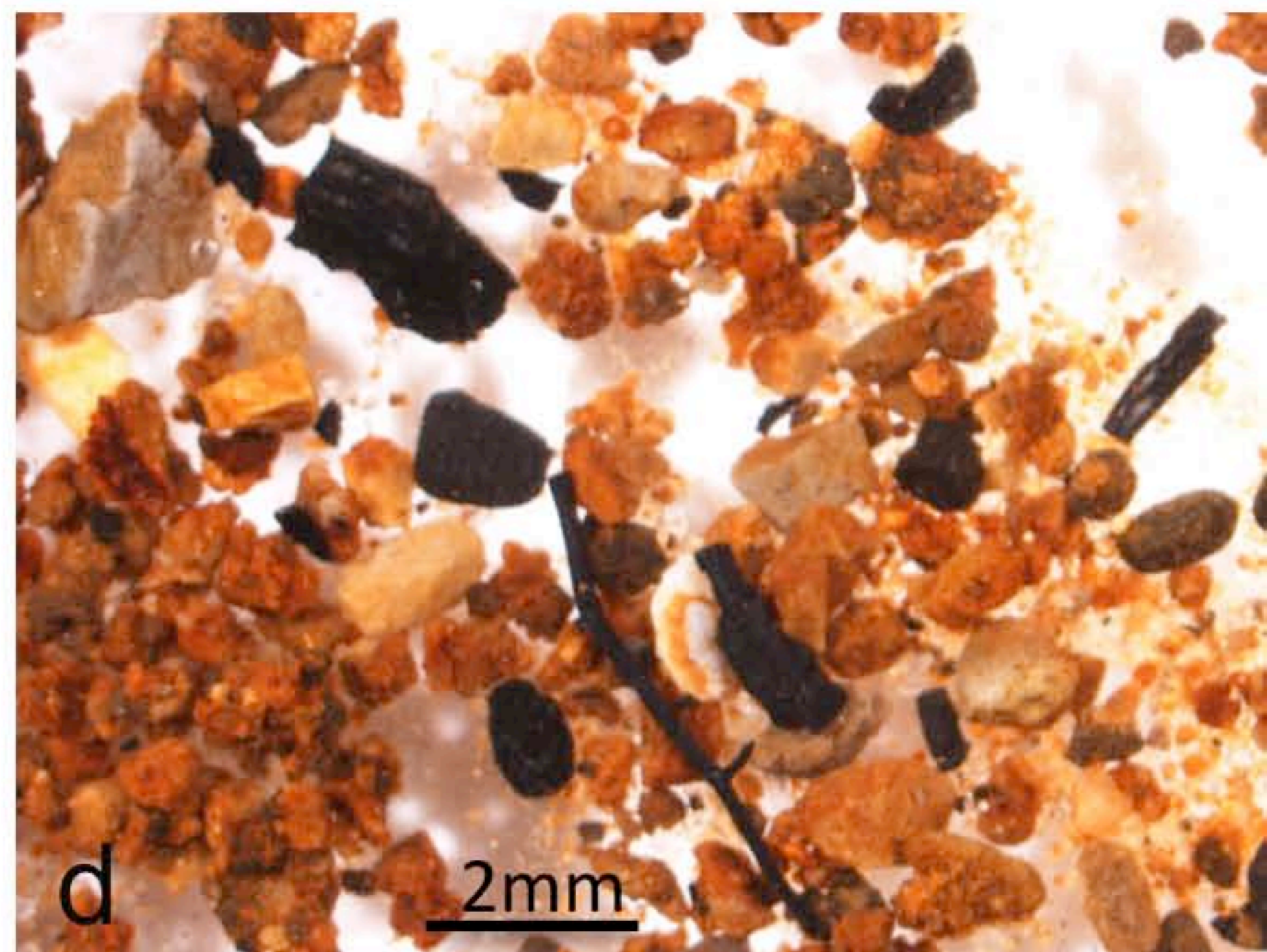
b

2mm



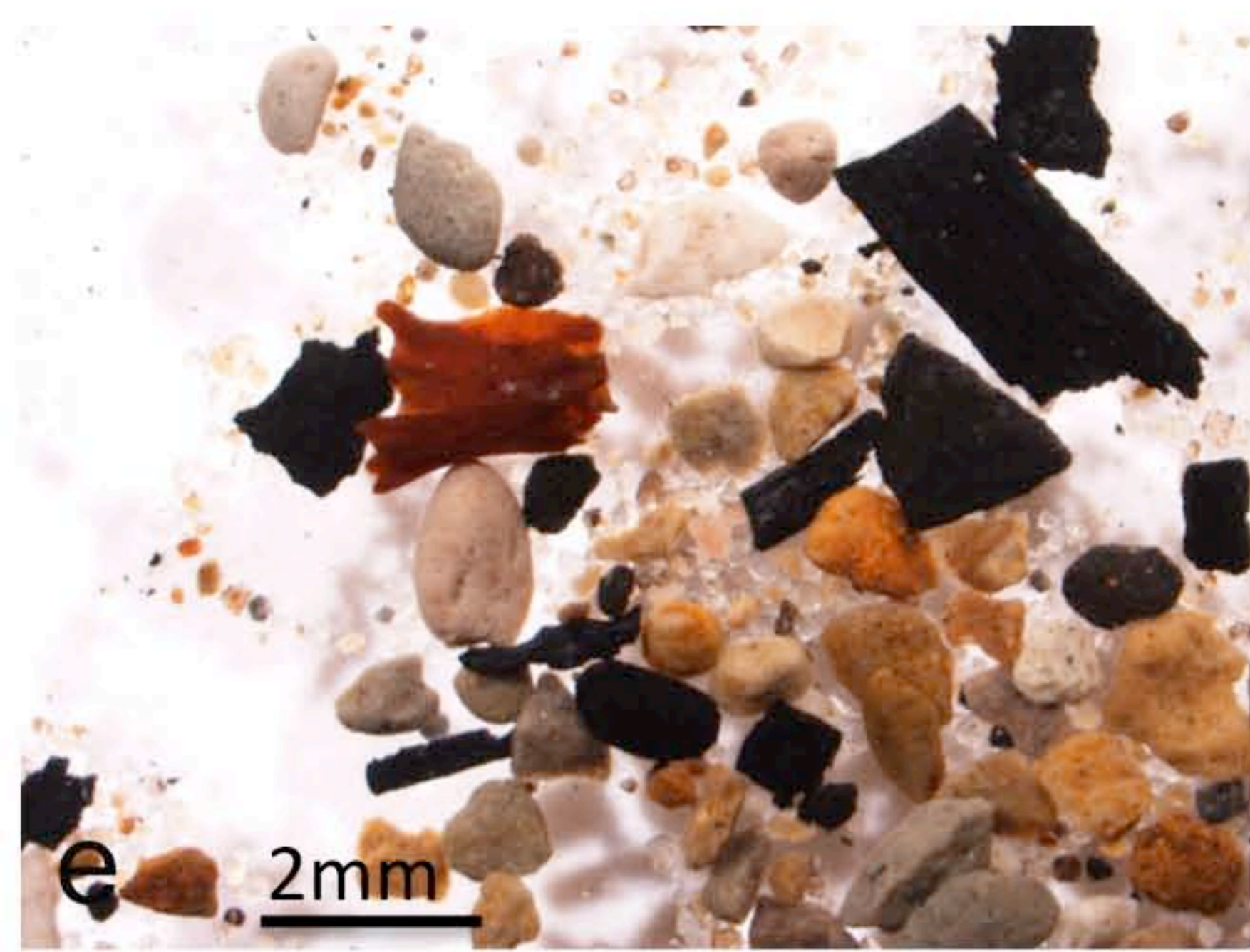
c

2mm



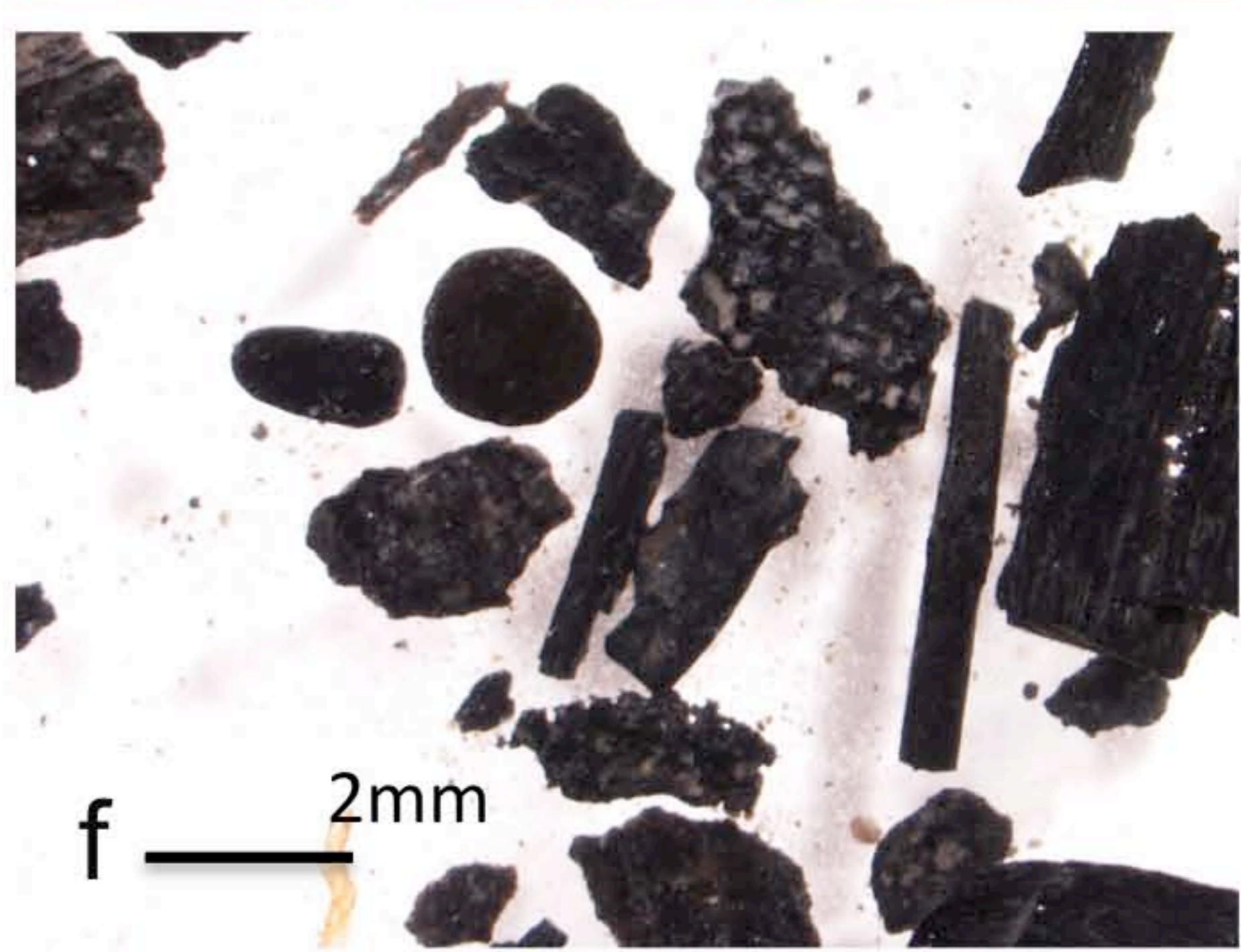
d

2mm



e

2mm



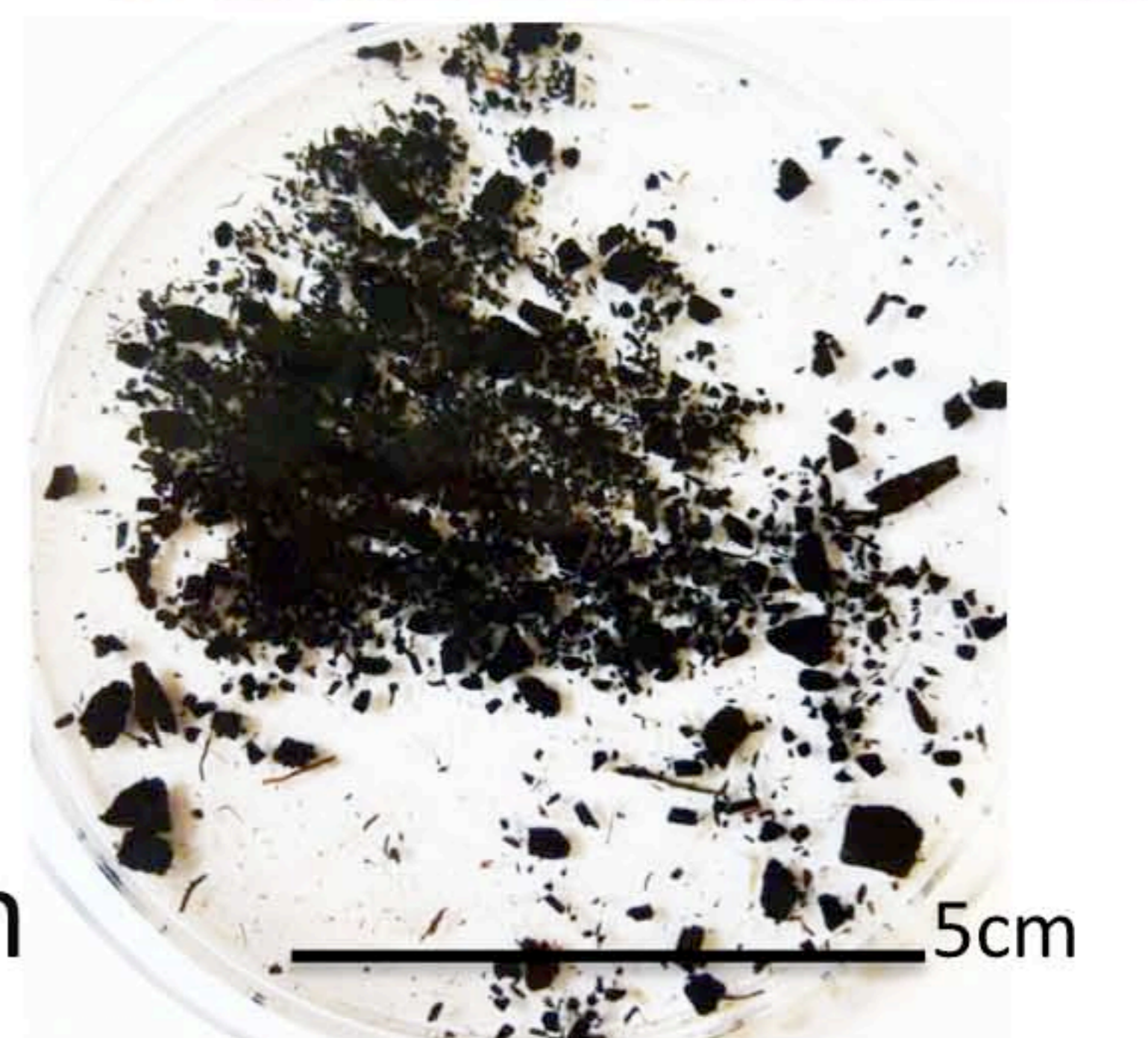
f

2mm



g

2mm



h

5cm

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a

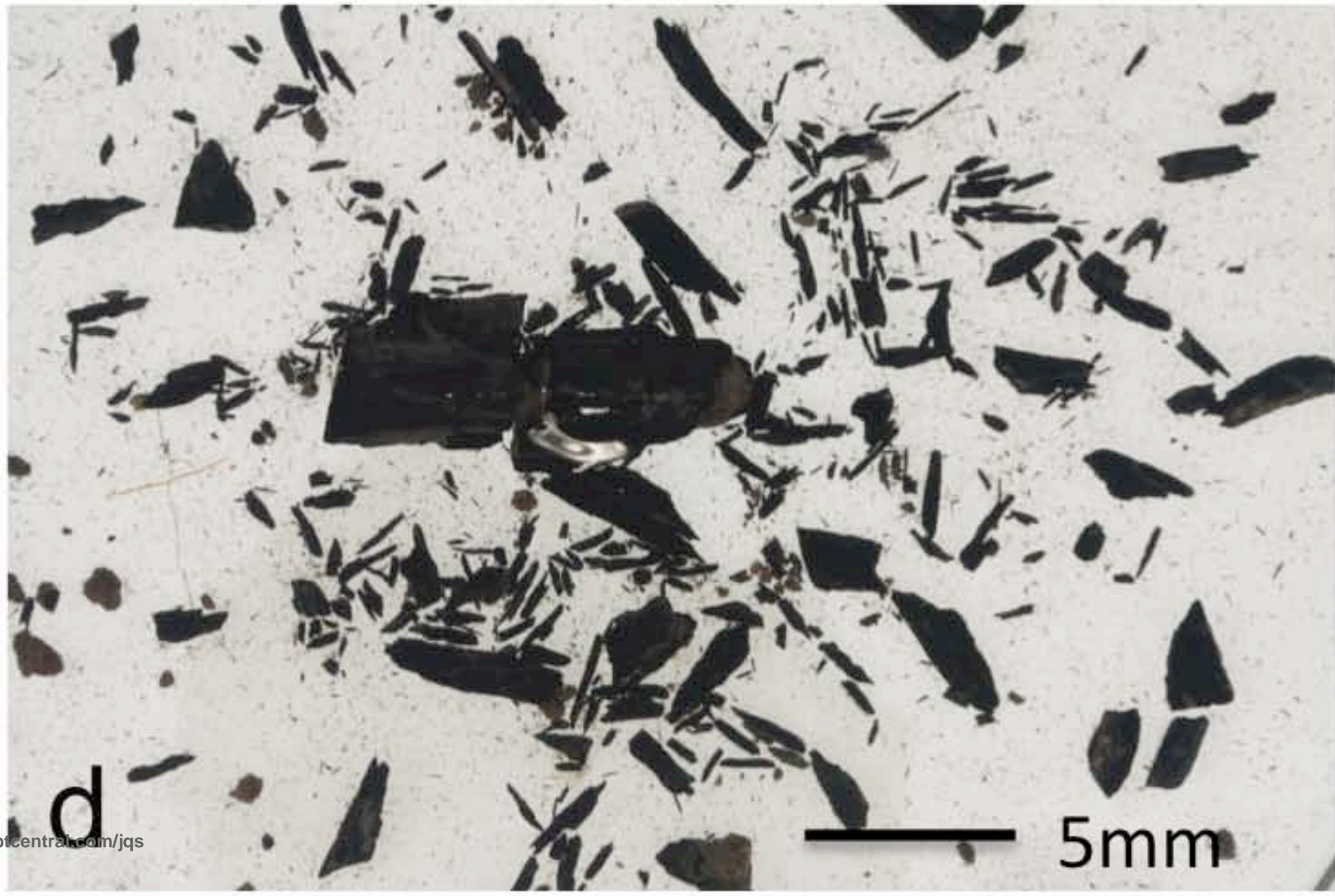


b

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c



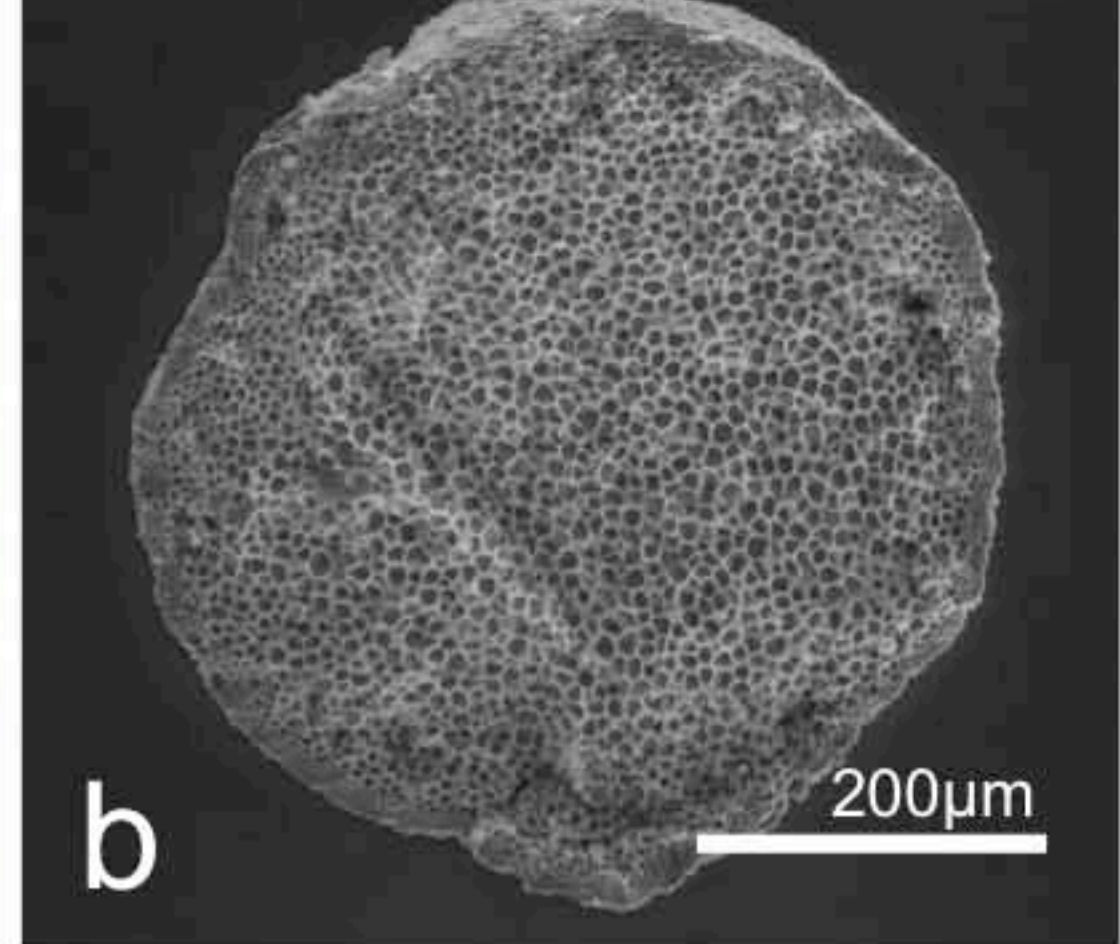
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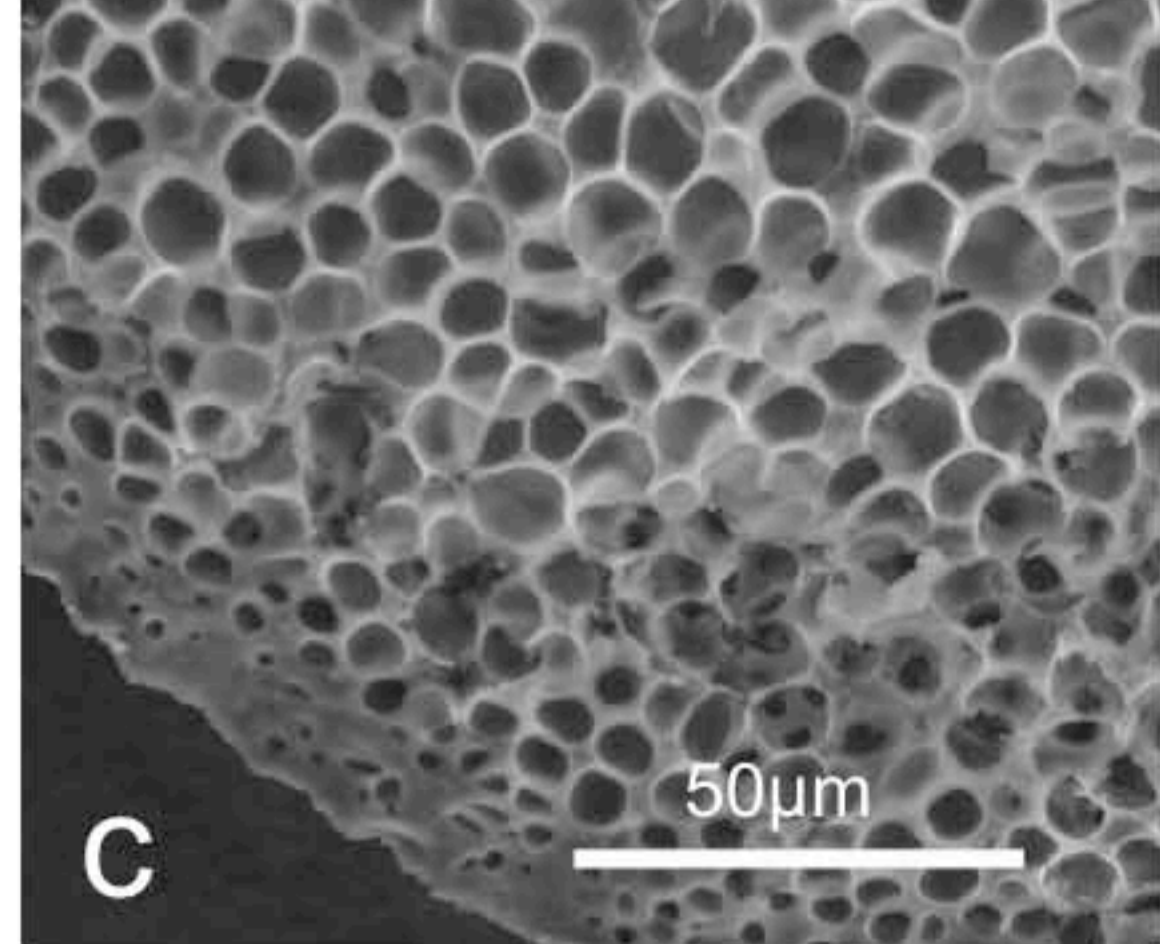
a

200µm



b

200µm



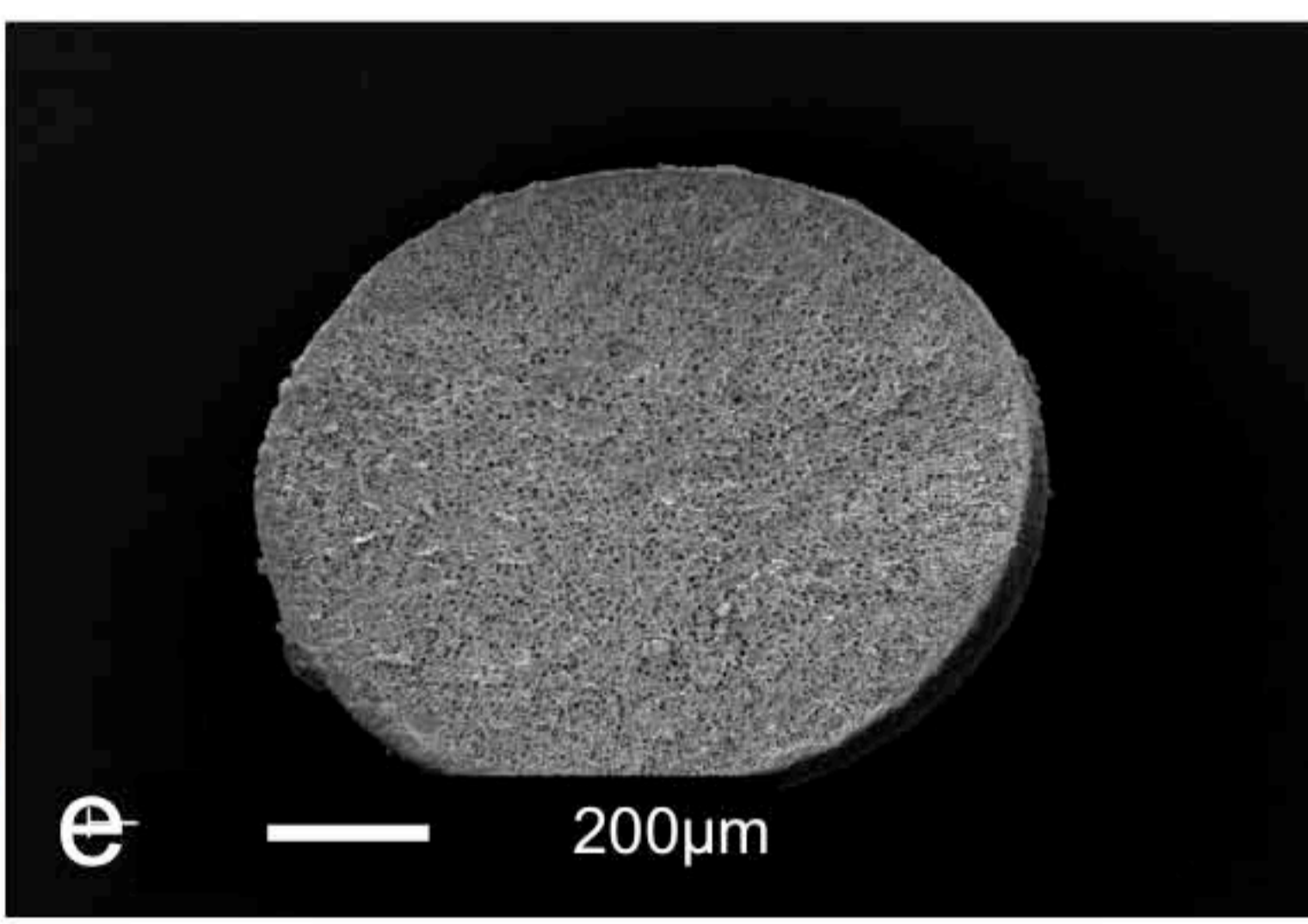
c

50µm



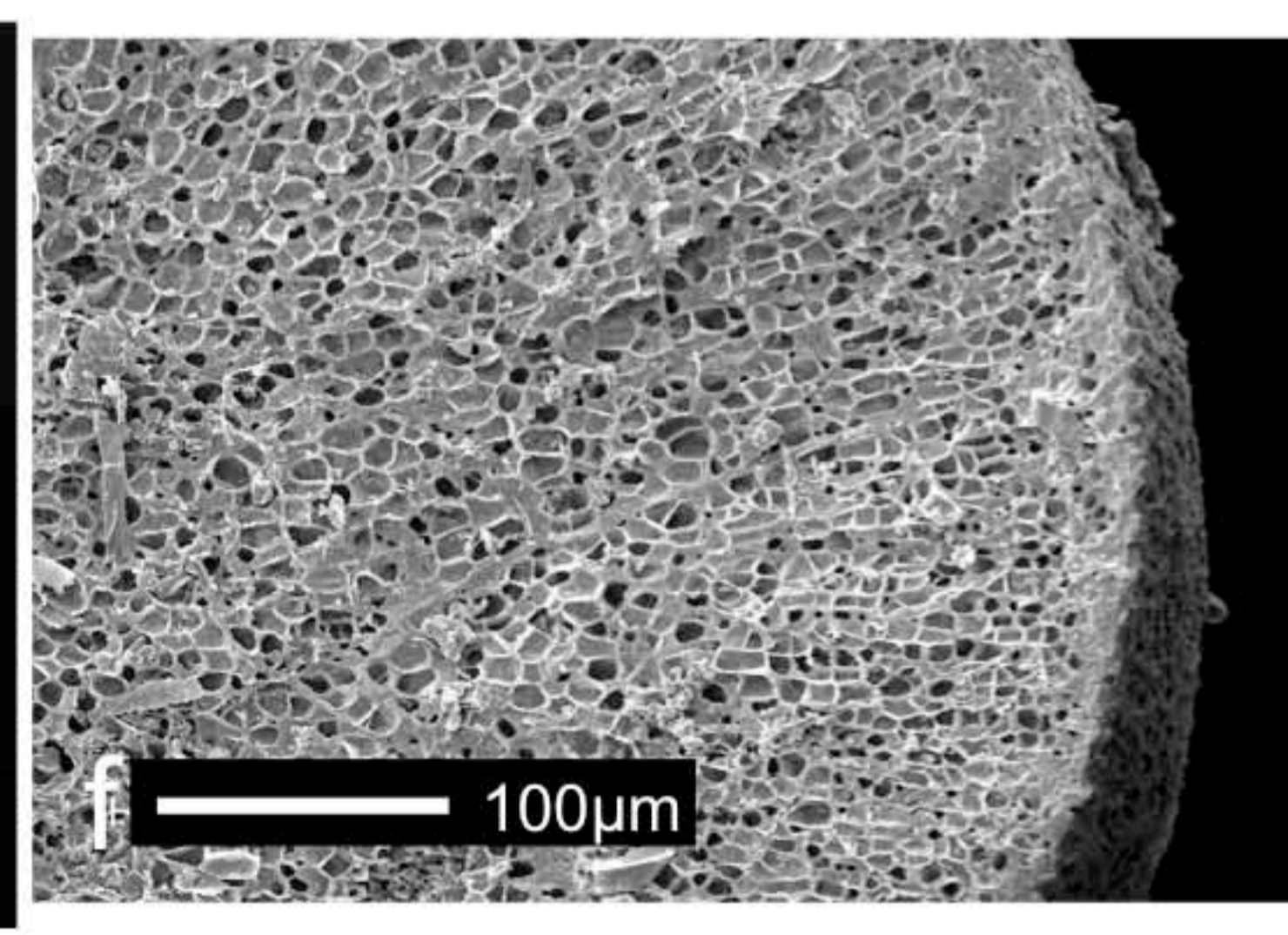
d

200µm



e

200µm



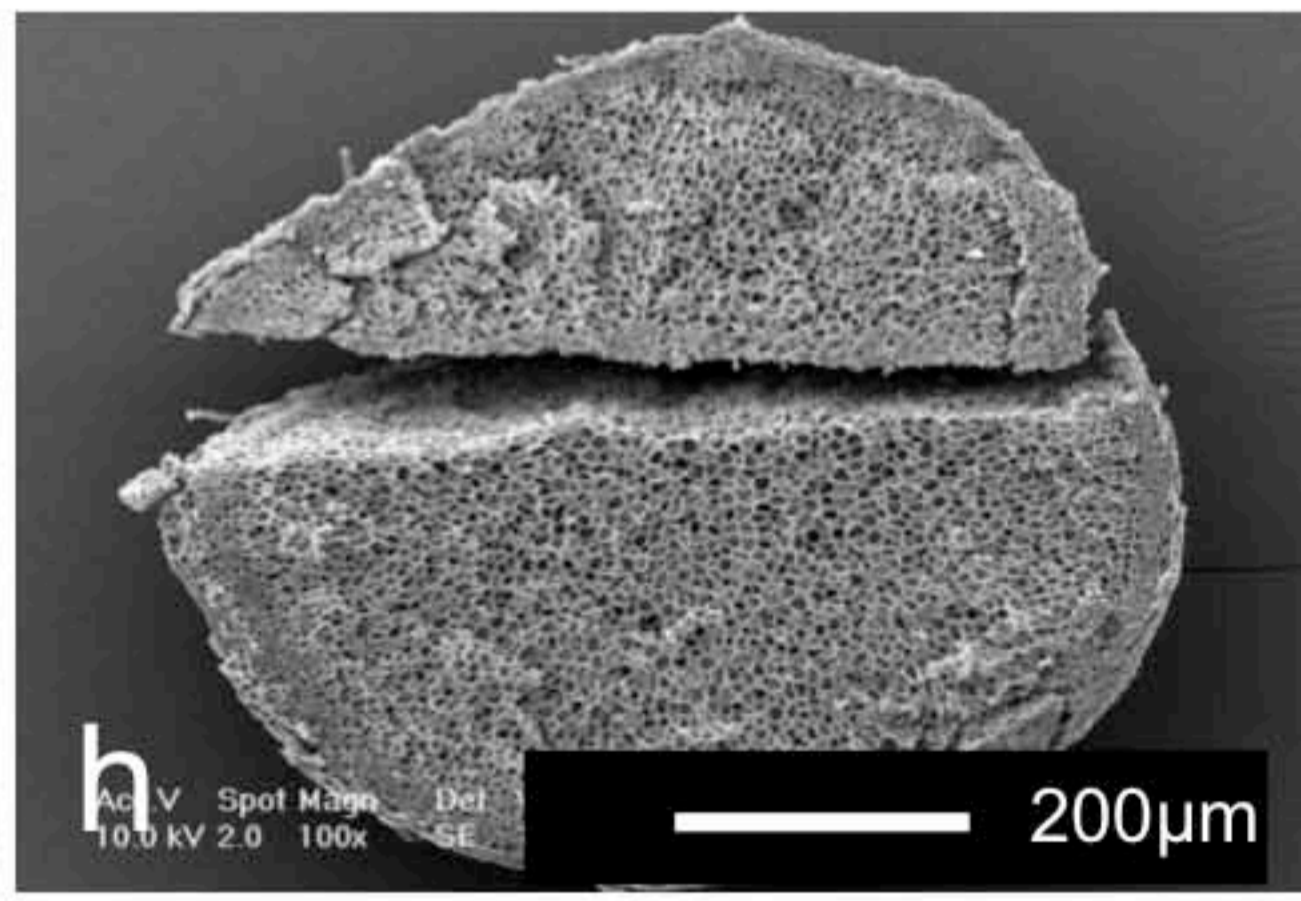
f

100µm



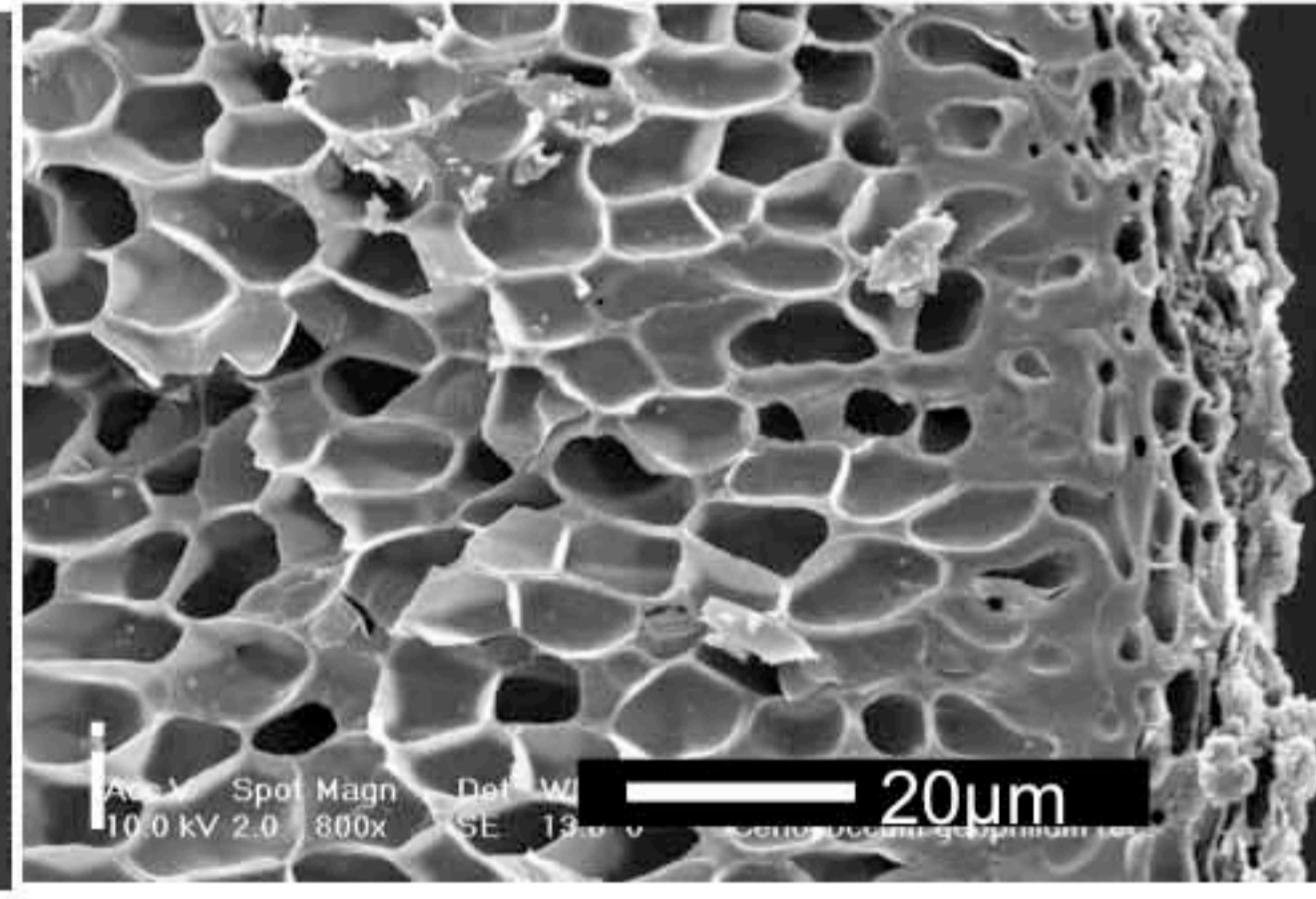
g

200µm



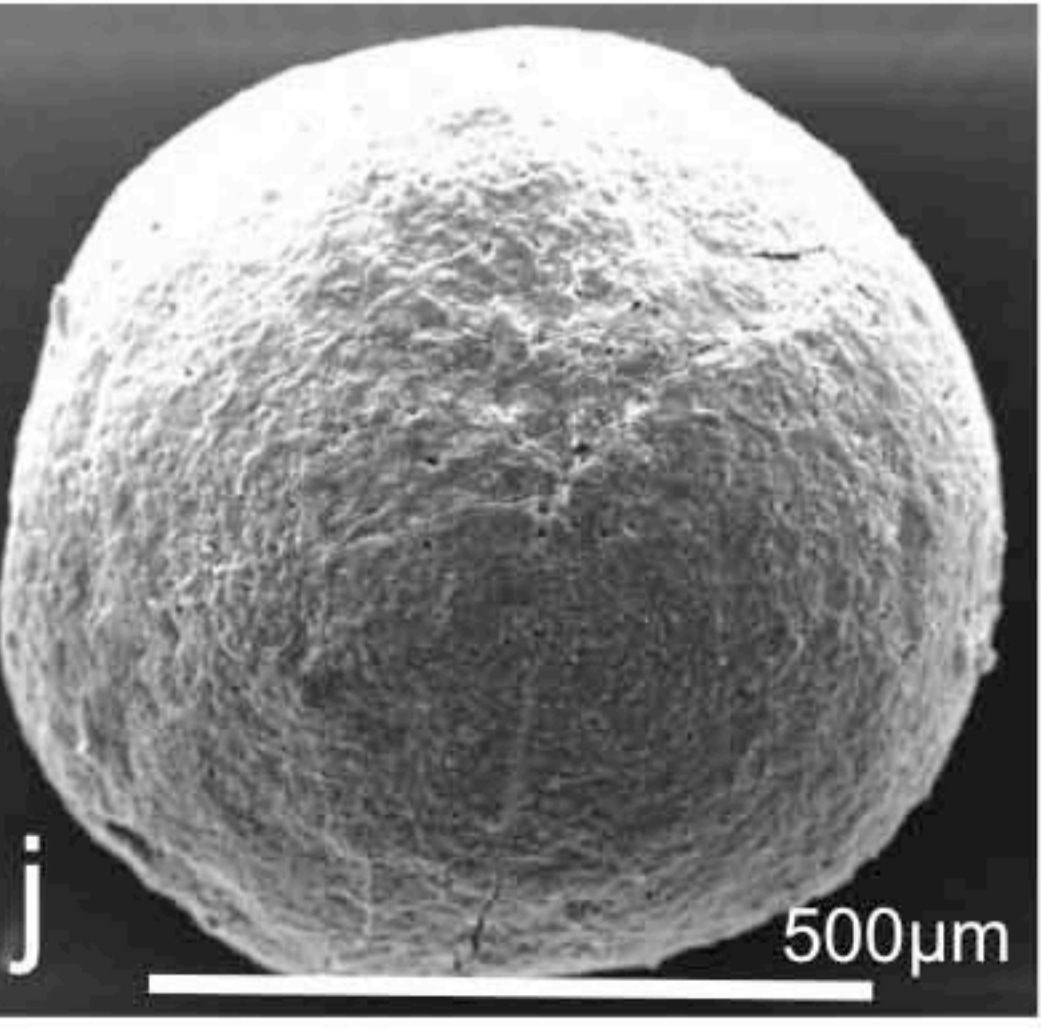
h

200µm



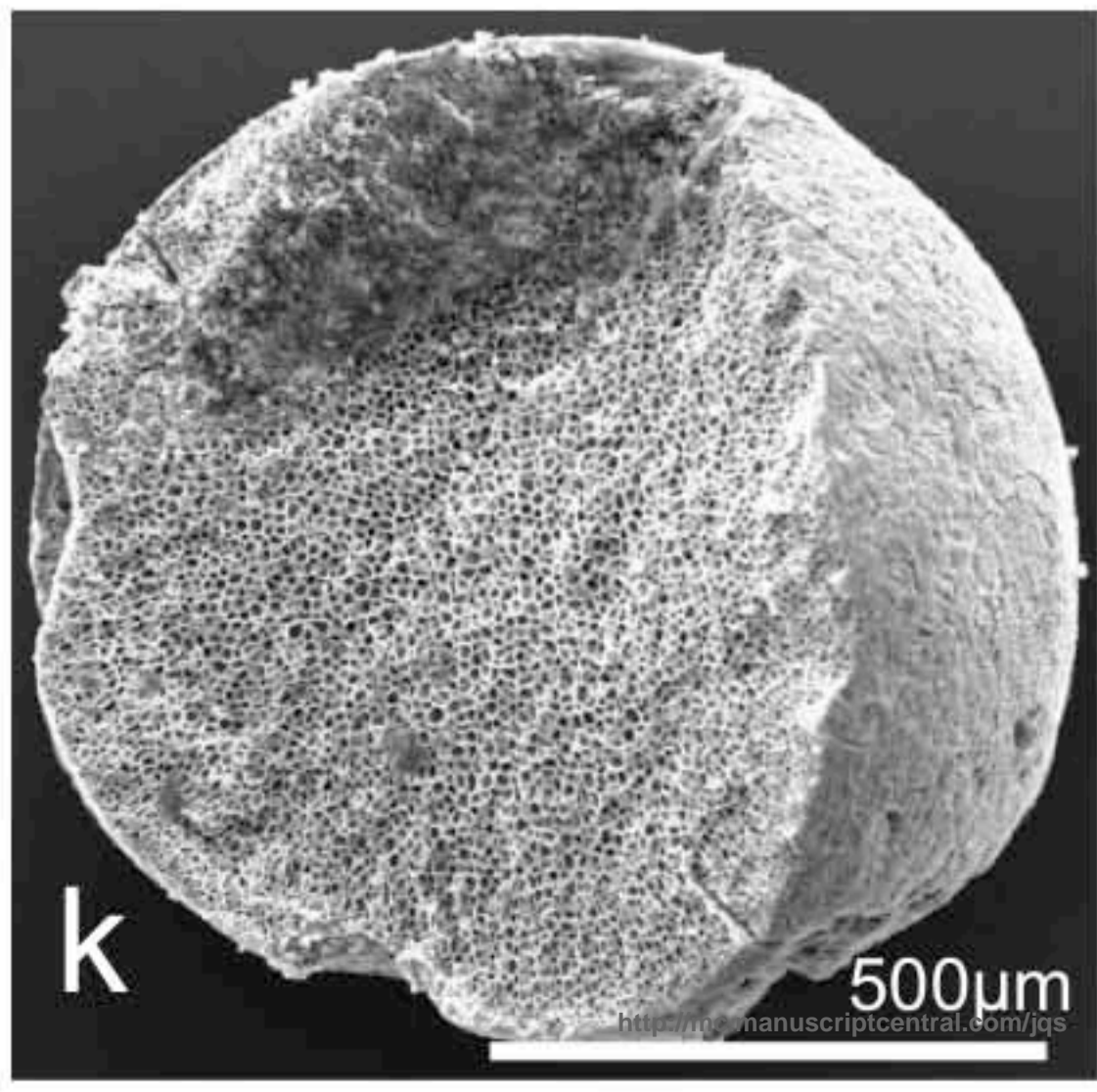
i

20µm



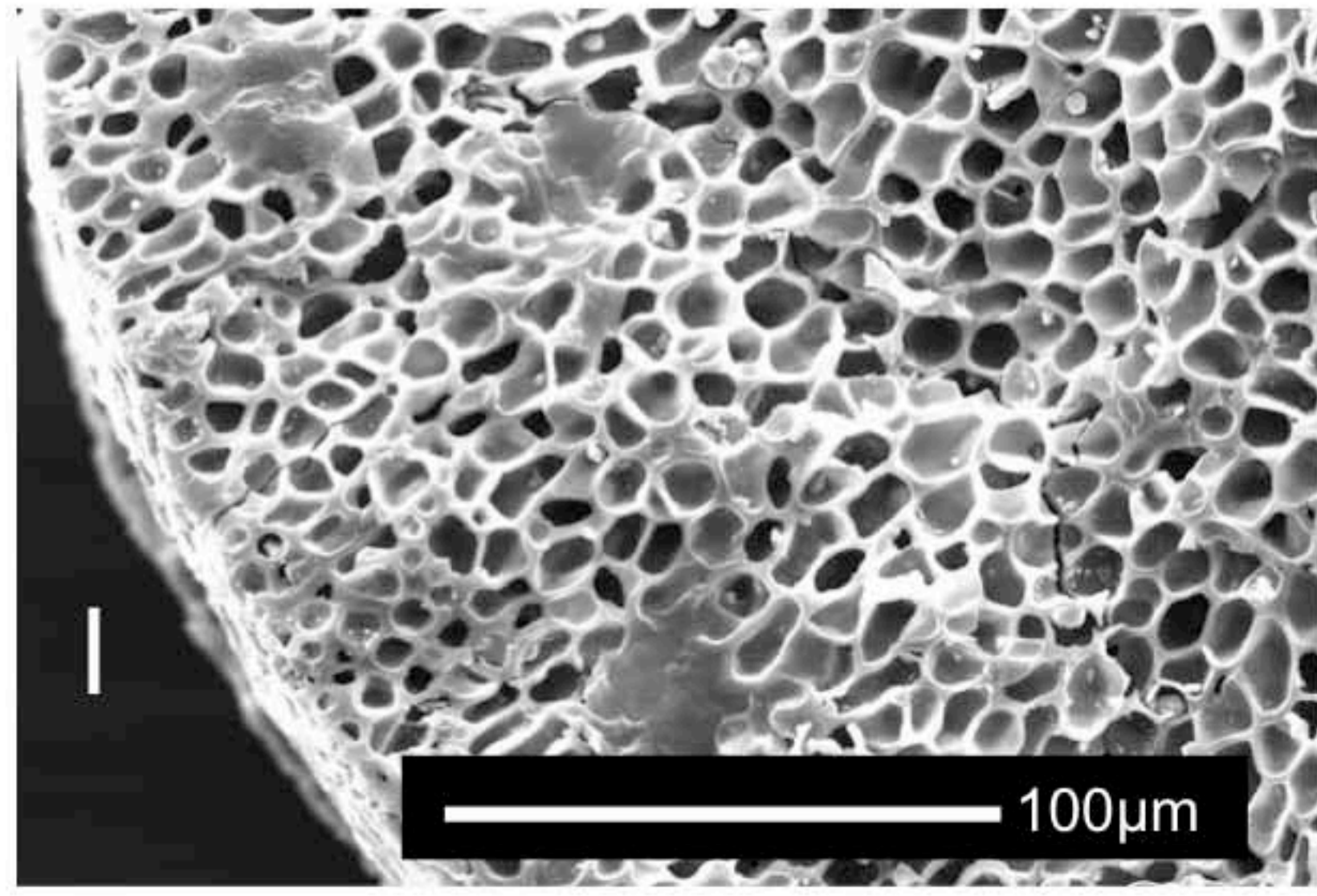
j

500µm



k

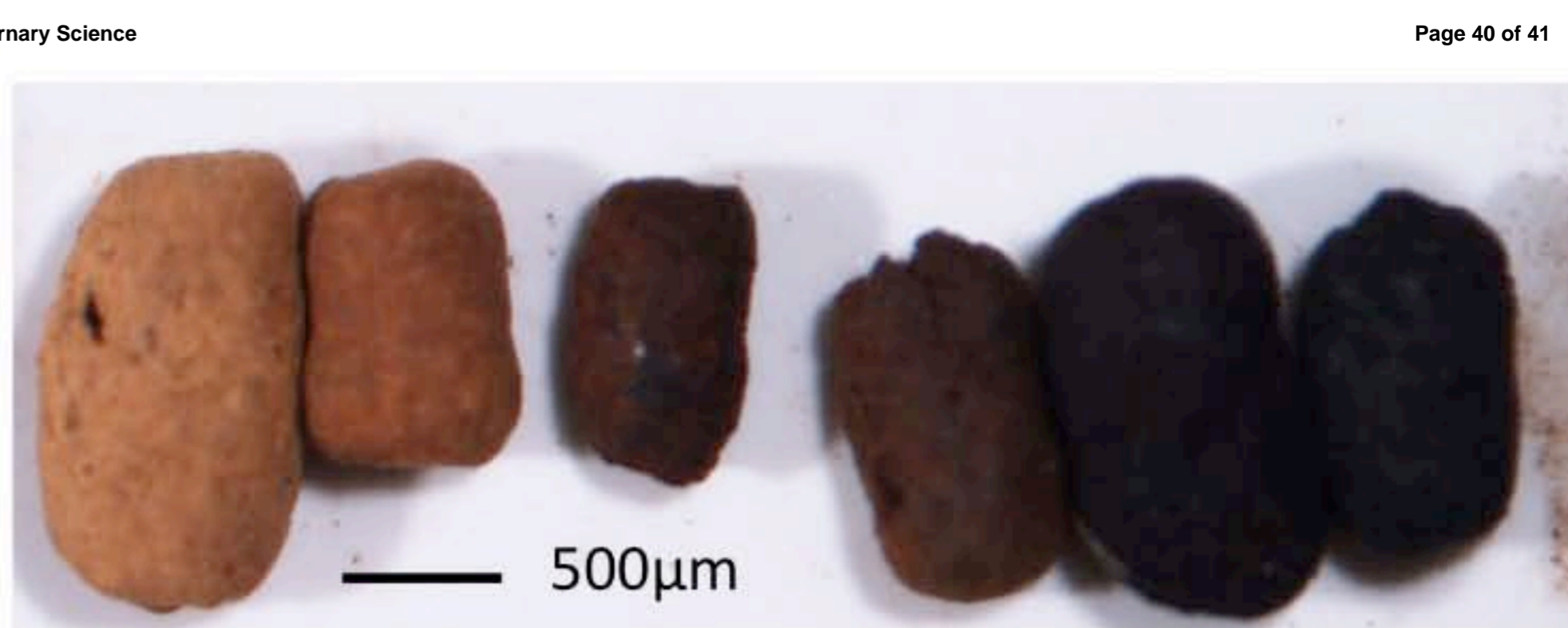
500µm



l

100µm

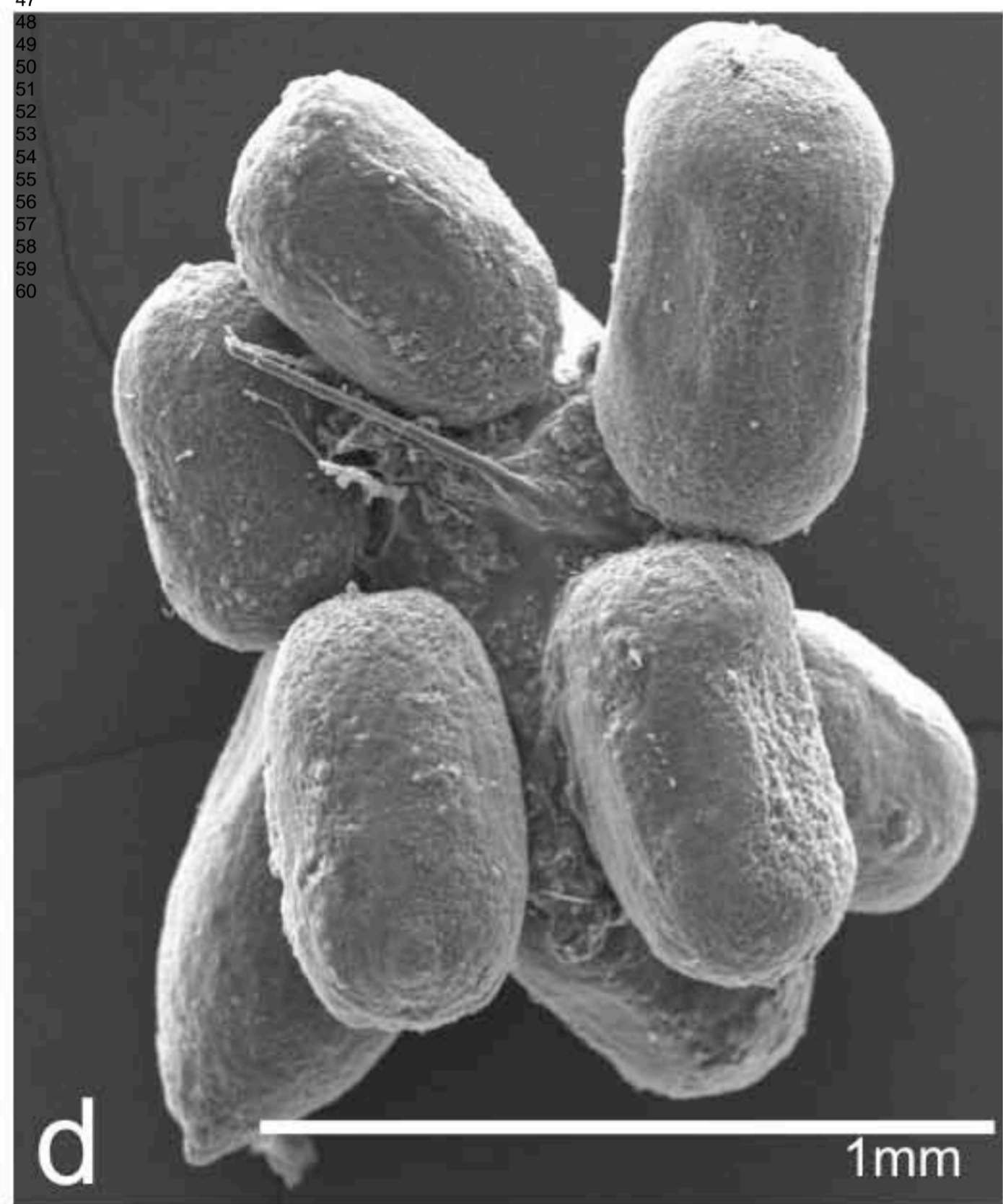
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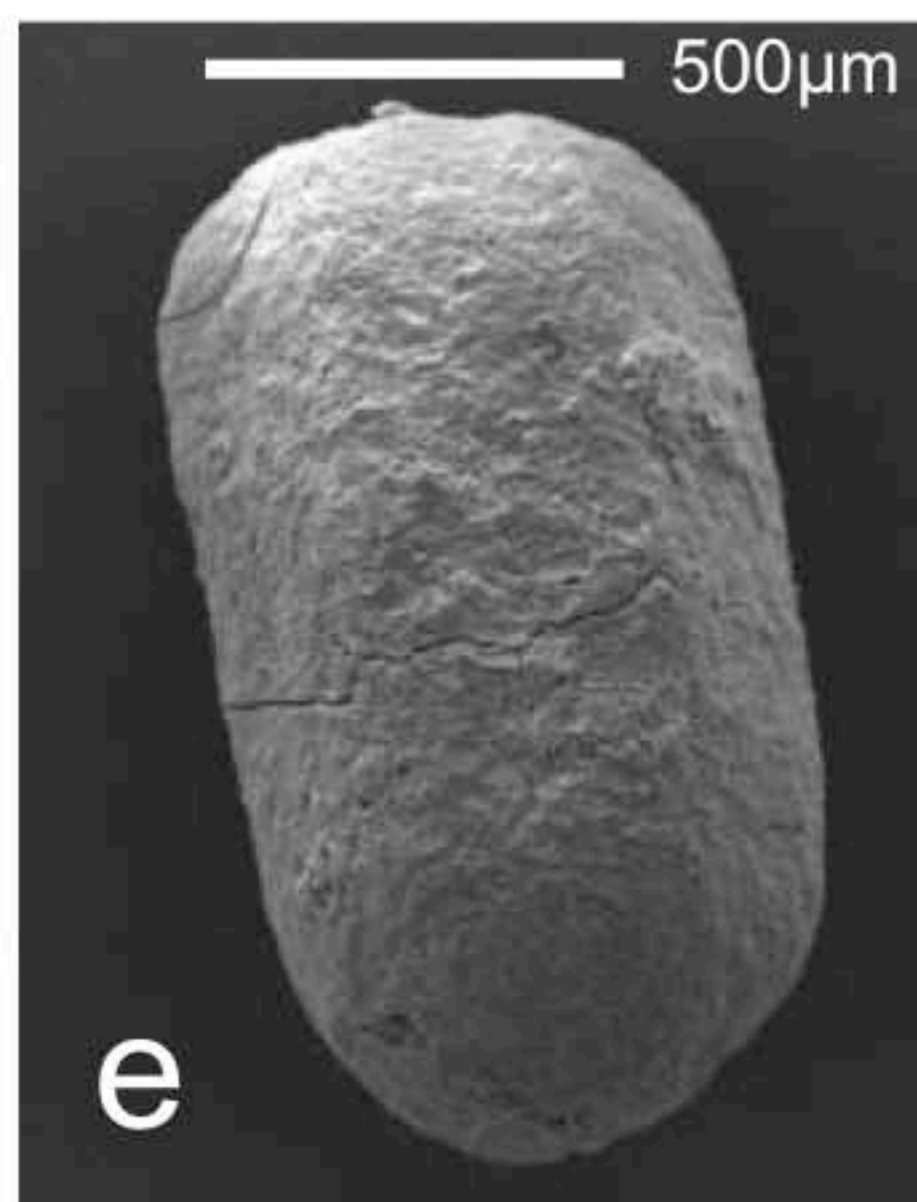
b



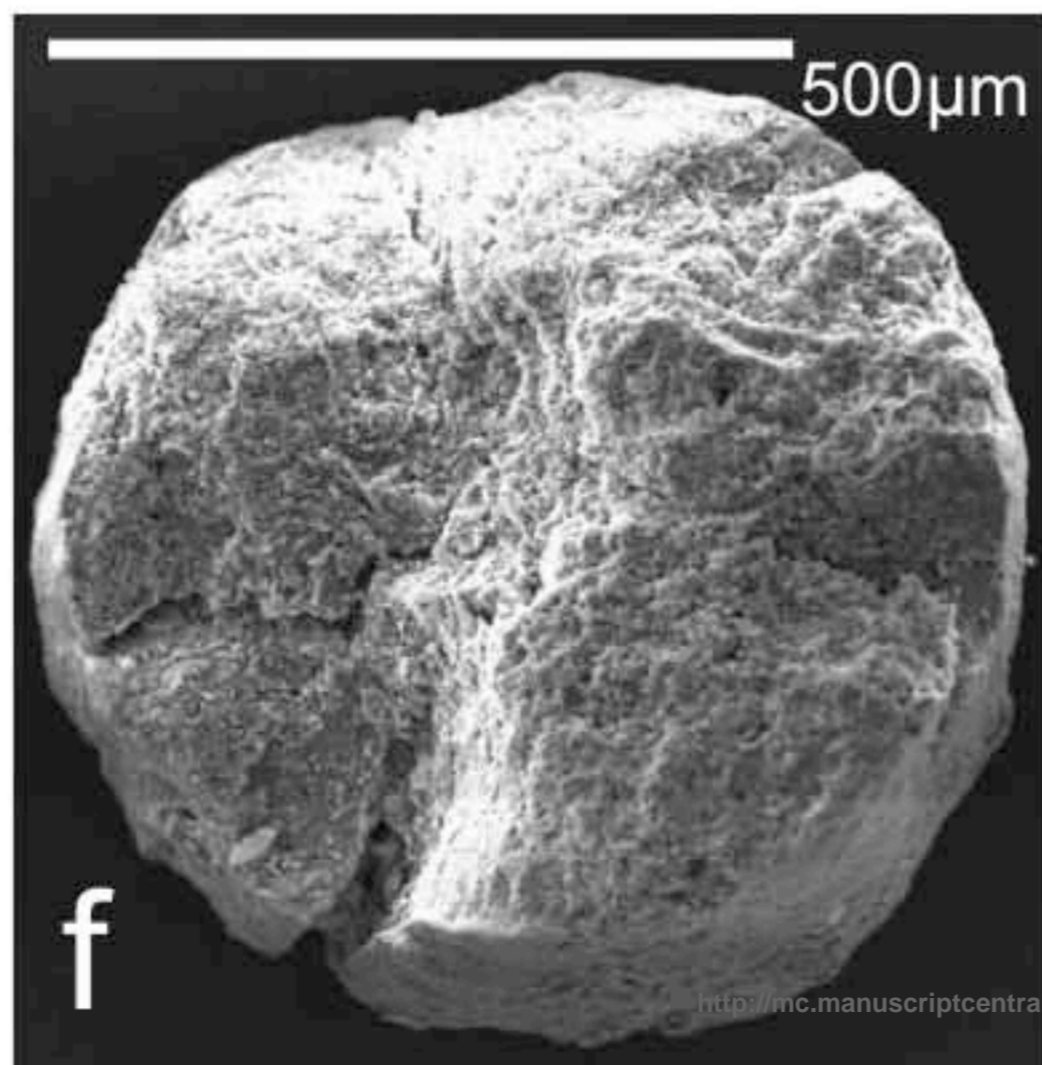
c



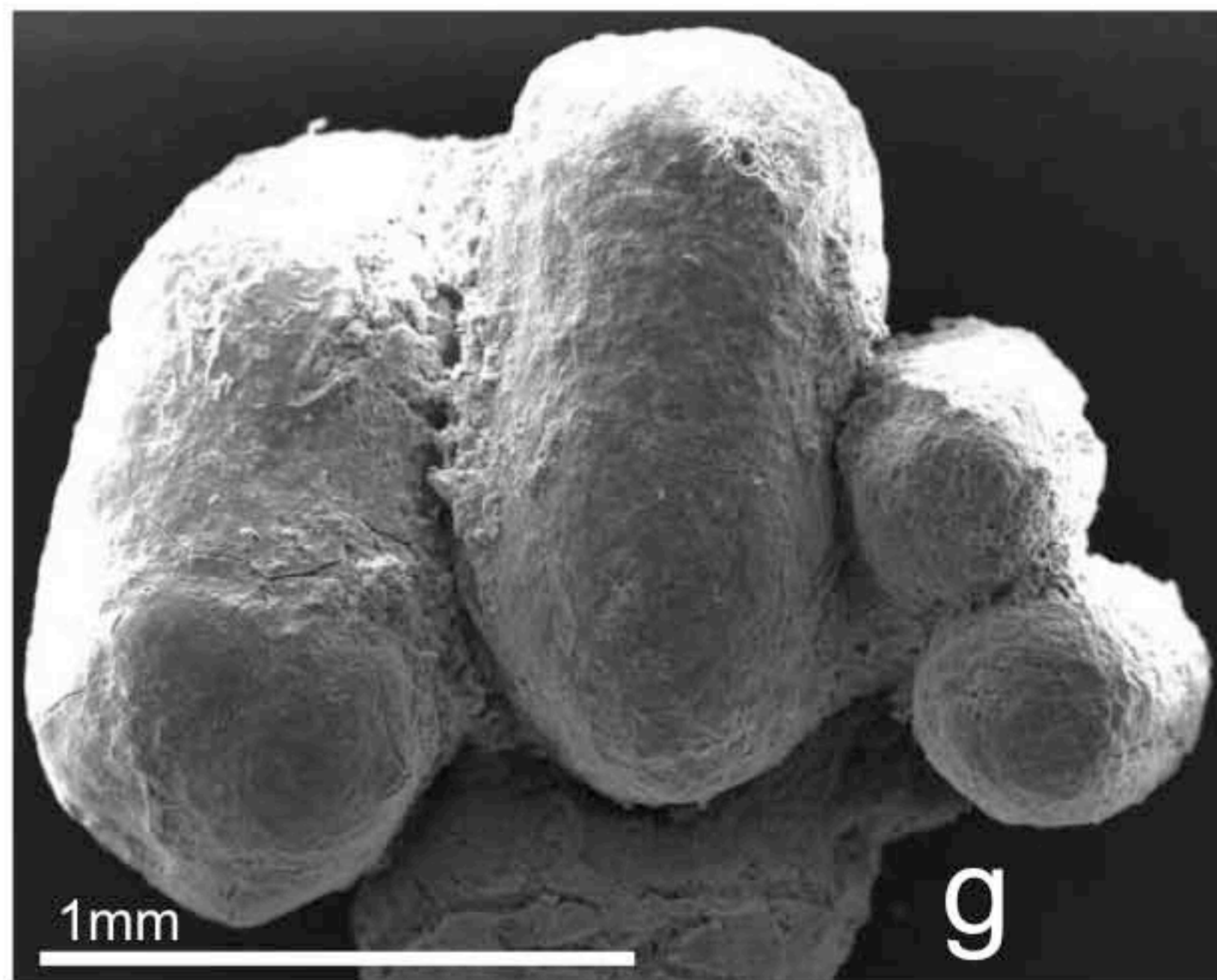
1mm



500µm



500µm



1mm

g

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