| 1   | Variability in Middle Stone Age symbolic traditions: the marine shell beads from Sibudu               |
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| 2   | Cave, South Africa  |
| 3   |   |
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| 15  |   |
| 16  | Highlights  |
| 17  |   |
| 18  | • We analyse marine gastropods from Sibudu dated to between 70 ka and 46 ka                           |
| 19  | • We study present day gastropod biocoenoses and thanatocoenoses along the KwaZulu Natal              |
| 20  | coast   |
| 21  | • Nassarius kraussianus shells were used as beads at Sibudu 46 ka                                     |
| 22  | • Mancinella capensis and Afrolittorina africana may have been used as beads 70 ka                    |
| 23  | • Observed differences in perforated taxa over time suggests variability in symbolic traditions       |
| ~ 1 |   |

# 25 Abstract

26 Located in the KwaZulu-Natal, 15 km from the coast, Sibudu has vielded twenty-three marine 27 gastropods, nine of which perforated. At  $70.5 \pm 2.0$  ka, in a Still Bay Industry, there is a cluster of 28 perforated Afrolittorina africana shells, one of which has red ochre stains. There is also a 29 perforated Mancinella capensis and some unperforated shells of both A. africana and M. 30 *capensis*. The cluster may represent an area where the shells were processed or where apparel to which shells were attached was lost. In a Howiesons Poort layer,  $64.7 \pm 1.9$  ka, there is a single 31 32 perforated Afrolittorina africana shell. This shell may be from the Still Bay and may have been 33 dislodged by rock fall. Two Nassarius kraussianus shells, one of them perforated and bearing 34 traces of utilisation as a bead, were found in a late Middle Stone Age layer with an OSL age of 35  $46.6 \pm 2.3$  ka. N. kraussianus was not found in older layers, even though this taxon occurs in 36 southern Cape sites during the Middle Stone Age. Where the perforations are undamaged and 37 suitable for microscopy, they appear to have been made by punching the shells with a pointed 38 tool made of bone or hard wood. Such perforations have been replicated experimentally. Unlike 39 the Blombos beads, the Sibudu shells from the Still Bay and Howiesons Poort layers bear no 40 compelling evidence of use wear in their perforations. We therefore cannot be certain that they 41 were suspended. However, if the Sibudu shells were beads, the changing use of taxa through time 42 suggests variability in symbolic traditions.

43

# 44 Keywords

45 Middle Stone Age, marine shell ornaments, microscopy, taphonomy

46

#### 48 **1. Introduction**

49 Early instances of personal decoration (e.g. Deacon, 1995; Ambrose, 1998; Kuhn et al., 2001; Henshilwood et al., 2004; Vanhaeren et al. 2006; Bouzouggar et al., 2007; Bar-Yosef Mayer et 50 51 al., 2009; d'Errico et al. 2009; Conard, 2010; d'Errico and Backwell, 2016; Bicho et al., 2018; 52 see Steele et al. 2019 for a review) have attracted the attention of archaeologists interested in the emergence of modern cultural traits because they are generally considered a convincing hallmark 53 54 of symbolically mediated behaviour and, by extension, a reflection of cultures comparable to those known today (e.g. McBrearty and Brooks, 2000; Wadley, 2003; Kuhn and Stiner, 2007; 55 56 Henshilwood and Dubreuil, 2011; d'Errico et al., 2003; d'Errico and Stringer, 2011; Shipton et 57 al., 2018). The discovery, more than a decade ago, of marine shells used as beads at the southern 58 African Middle Stone Age (MSA) site of Blombos Cave, in layers dated to ca 72 ka 59 (Henshilwood et al., 2004; d'Errico et al., 2005; Vanhaeren et al., 2013) has challenged the 60 longstanding paradigm according to which personal decoration did not occur before 40 ka. Additional marine shell beads dated to between ca 100 ka and 70 ka were identified in 61 62 subsequent years at seven Aterian and Levantine Mousterian sites of North Africa and Western Asia (Vanhaeren et al., 2006; Bouzouggar et al., 2007; d'Errico et al., 2009; Bar-Yosef et al., 63 64 2009) and in the Howiesons Poort of Border Cave in South Africa (d'Errico and Backwell, 65 2016). At each of these sites personal ornaments consist of perforated shells belonging almost 66 exclusively to a single species: Nassarius gibbosulus in Morocco and Algeria, this species and 67 Glycymeris sp. in Israel, and Nassarius kraussianus and Conus ebraeus shells in South Africa. 68 Taphonomic analysis and experimental reproduction of the modifications recorded on the shells 69 have shown that N. kraussianus were collected alive, whereas N. gibbosulus and Glycymeris sp. were gathered dead on the shores. Some of these beads bear traces of pigment and intense use-70 71 wear. Experimental reproduction of use-wear recorded on *Nassarius kraussianus* shell beads 72 from Blombos Cave has recently shown that a clear change in the way of stringing beads and the 73 visual appearance of the resulting beadwork occurred at Blombos Cave between lower and upper 74 Still Bay layers (Vanhaeren et al., 2013). It has also been shown that these changes were 75 coincident with environmental change and the way in which the habitation space was used. When 76 recovered at sites excavated with modern standards these shell beads often bear traces of red 77 pigment. Some of them are blackened by heating either accidentally or for the purpose of 78 changing their colour (d'Errico et al., 2015). 79 While confirming the existence of early bead-using traditions these discoveries have also raised

80 questions about the mechanisms that have led to the emergence and maintenance of such

81 innovation. On the one hand, no personal ornaments have been so far reported from sites securely

82 dated to between 70 ka and 50 ka and the ornaments attested in Africa at ca. 45 ka (Ambrose,

83 1998; d'Errico et al., 2012; Gliganic et al., 2012; Miller and Willoughby, 2014), almost

84 exclusively consisting of ostrich egg shell and stone beads, suggests cultural discontinuity with

85 previous traditions. This contradicts (d'Errico and Stringer, 2011) the scenario according to

86 which the emergence of cultural complexity would be reflected by a process of continuous

87 accretion and implementation of cultural innovations. On the other hand, although convincing

and associated with other striking evidence of cultural complexity, the Blombos and Border Cave

shell beads were, until now, the only reliable instance of personal ornamentation prior to 70 ka

90 for the southern African continent.

91 In a previous paper we reported the presence of possible shell beads in the Still Bay layers at

92 Sibudu Cave, KwaZulu-Natal, South Africa (d'Errico et al., 2008). They consist of six

93 Afrolittorina africana, five of which were recovered in the Still Bay and one in the lowermost

94 Howiesons Poort layer of this site. Three of these shells bear perforations, one has residues of red

95 pigment, and three are blackened by heating. One shell bears a complete and a broken perforation

96 that can correspond to a repair after a first break. Taphonomic analysis of the archaeological

97 specimens based on present day Afrolittorina africana biocoenoses, microscopic examination,

98 experimental perforation of modern shells, and a review of the natural agents that may

99 accumulate marine shells at inland sites such as Sibudu has suggested to us probable human

100 involvement in the collection, transport, modification, and abandonment of Sibudu's

101 Afrolittorina africana. Here we report on the discovery of an extended collection of marine shells

102 from the late Middle Stone Age (MSA), Howiesons Poort (HP), Still Bay (SB), and pre-Still Bay

103 (pre-SB) layers of this site. These shells belong to three species (Afrolittorina africana,

104 Mancinella capensis and Nassarius kraussianus); some are perforated, heated, and bear red

105 pigment residues. Study of archaeological and experimentally perforated shells, and comparison

106 with shells from modern biocoenoses (*i.e.* living assemblages) and thanatocoenoses (*i.e.* dead

107 assemblages) of the Ballito Bay shores, allows us to reinforce the argument for their being

108 purposely collected, modified and used as beads. A detailed description of this material and

109 related behavioural correlates is key to assessing recent claims according to which the production

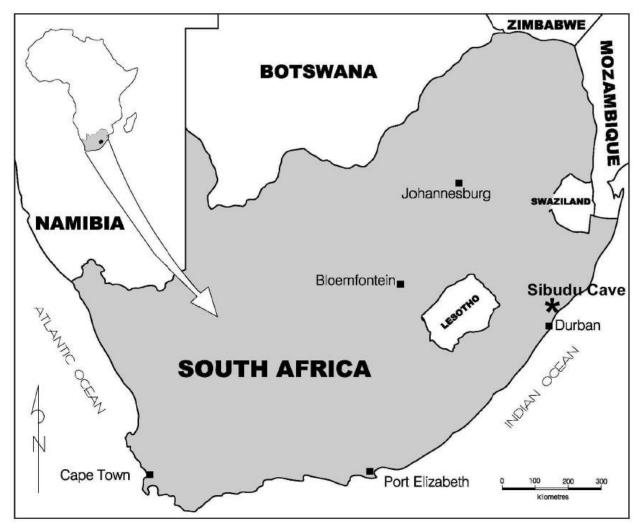
110 of personal ornaments would have little implication for the emergence of complex human

111 cognition (Haidle et al., 2015; Garofoli, 2014).

- 112
- 113

## 114 **2. Background to Sibudu**

- 115 Sibudu is a large rock shelter above the uThongathi River, about 40 km north of Durban in
- 116 KwaZulu-Natal, South Africa (Fig. 1). At present the shelter is about 15 km inland of the coast
- and the maximum distance from the coast during the last glacial episode is likely to have been
- 118 about 20 km.



119

- 120 Figure 1. Location of Sibudu Cave.
- 121

122 The site was excavated under the directorship of Lyn Wadley between 1998 and 2011. Since

- 123 2011 the site has been excavated by Nicholas Conard, University of Tübingen. A long and
- 124 detailed Middle Stone Age (MSA) sequence with good organic preservation occurs within its
- three metre deep sediments. The site's cultural succession (Lombard 2004, 2005, 2006a, 2006b,
- 126 2007, 2011; Wadley, 2004, 2005, 2006, 2007, 2010a,b; Villa et al., 2005; Cochrane, 2006, 2008;
- 127 Delagnes et al., 2006; Villa and Lenoir, 2006; Backwell et al., 2008; d'Errico et al., 2008;
- 128 Wadley and Mohapi, 2008; Wadley et al., 2009, 2011; Wadley and Kempson, 2011; d'Errico et

- 129 al., 2012b; de la Peña et al. 2013; de la Peña and Wadley, 2014a, b, 2017; de la Peña, 2015;
- 130 Soriano et al. 2015; Wozcieszak and Wadley, 2018), chronology and micromorphology ((Schiegl
- 131 et al., 2004; Pickering, 2006; Schiegl and Conard, 2006; Wadley and Jacobs 2006; Jacobs et al.,
- 132 2008a, b; Goldberg et al., 2009) and environmental context (Cain, 2004, 2006; Plug, 2004, 2006;
- Allott, 2006; Glenny, 2006; Herries, 2006; Renaut and Bamford, 2006; Reynolds, 2006; Sievers,
- 134 2006; Wells, 2006; Clark and Plug, 2008; Wadley et al., 2008; Robinson and Wadley, 2018;
- 135 Clark 2019) are already published extensively elsewhere.
- 136
- 137 The Wadley excavation method is described in detail in Wadley and Jacobs (2006). In brief, the 138 sediment was excavated in 50 cm quadrants (a-d) within each metre square. Quadrant a is the 139 north-east quadrant in each metre square and B4a, for example, describes quadrant a in square 140 B4. Twenty-one square metres of deposit were excavated into the ~58 ka occupations and six 141 square metres were excavated as a deep sounding into older occupations with ages between 77.2 142  $\pm$  2.1 and 61.7  $\pm$  1.5 ka (Fig. 2). The ages are derived from single-grain optically stimulated 143 luminescence (OSL) analysis of sedimentary quartz grains (Jacobs et al., 2008a, b). The deepest 144 layers contain assemblages that are informally called pre-Still Bay, while Still Bay, Howiesons 145 Poort and late MSA Industries occur more recently. Occupations at Sibudu were discontinuous, 146 with hiatuses that, in layers more recent than 58 ka, were sometimes as long as 10,000 years. The 147 final Wadley excavations of 2011 reached a layer called BS16 (Brown Sand 16), which is 148 undated, but older than 77 ka. Characteristic finds, constituent stratigraphic units, date intervals 149 of the different cultural assemblages and OSL ages for the stratigraphic units are summarised in 150 Figure 2.

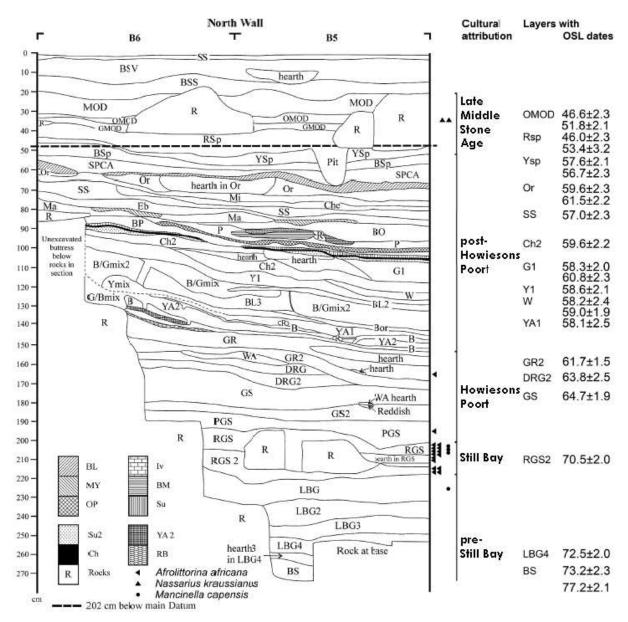


Figure 2. Stratigraphy of Sibudu Cave along the North section with indication of the stratigraphic
layers, their cultural attribution, and optically stimulated luminescence (OSL) dates after Wadley
and Jacobs, 2006 and Jacobs et al., 2008a,b.

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In the lower most stratigraphic unit BS16 BS16 of the Wadley excavation, a long bifacial point was recovered (Wadley, 2012). It is narrow and thin, but has a rounded base, not a pointed one like a 'classic' Still Bay point. The Conard excavations below BS16 recovered more bifacial points, some serrate (Rots et al. 2017). The uppermost BS layer has an OSL age of  $77.2 \pm 2.1$  ka. The BS member varies between 30 and 40 cm in thickness and, apart from the basal layer BS16, contains an informal assemblage with only rare retouched pieces. The LBG (Light Brownish Sand) member above this is about 50 cm thick and it comprises four layers with two available OSL ages,  $73.2 \pm 2.3$  and  $72.5 \pm 2.0$  ka (Jacobs et al., 2008b). The informal lithic assemblage in these layers has only rare examples of retouched pieces and has been designated pre-Still Bay. Bone tools with double bevel working ends come from the pre-Still Bay layers.-Some ochre pieces with groups of lines forming fan-shaped motifs that may have been deliberately scored come from the pre-Still Bay layers (Hodgskiss, 2012, 2013).

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169 The Still Bay Industry at the site is therefore immediately preceded by a deep sequence that 170 extends about 80 cm in the Wadley excavation and even farther in the recent Conard one. The 171 Still Bay occurs in the member Reddish Grey Sand (RGS), a loose, sandy sediment named after 172 the Munsell colour reading 5YR 5/2 Reddish-Grey. Two layers, RGS2 and RGS have been 173 named although the division between the two is not distinct, and the entire member is seldom 174 more than 20 cm thick (Fig. 2). Several separate lenses have been distinguished by subtle colour 175 changes within RGS2 and RGS. RGS, at the top of the sequence, has an OSL age of  $70.5 \pm 2.0$ 176 ka, and the entire Still Bay seems to represent a short pulse of occupation at Sibudu. Since the 177 RGS layer is directly below a Howiesons Poort layer (PGS) with an OSL age of  $64.7 \pm 1.9$  ka, an 178 occupation hiatus of about five thousand years seems to occur at Sibudu between the Still Bay 179 and the Howiesons Poort. Excavation of the Still Bay layers was hampered by extensive rock fall 180 that began in the BS member and continued, perhaps intermittently, until about 65 ka. The rock 181 fall has encroached on the six square metres of the deep sounding and this has resulted in 182 restricted areas of occupation within the SB and HP Industries. Perhaps because of the extensive 183 rockfall, the combustion features in Layer RGS are concentrated in squares B4 and C4 and in a 184 single quadrant in square B5 where fewer rocks occur (Fig. 3). At least three of the ashy features appear to be hearths. All lithics and bone fragments larger than 20 mm were piece-plotted during 185 186 the excavation of the Still Bay layers, as well as shells, bifacial tools and worked ochre. The Still 187 Bay Industry at Sibudu, like that occurring in MSA sites along the South African Cape coast, is 188 marked by the presence of bifacial tools with invasive retouch, including some rare lanceolate 189 points. Whole bifacial points are relatively uncommon at the site, but bifacial point fragments are 190 abundant, as is also the case at other Still Bay sites, such as Blombos (Henshilwood et al., 2001) 191 and Hollow Rock (Evans, 1994). This pattern is probably due to the vulnerability of the long, 192 thin bifacial points. In a preliminary study, bifacially worked tools comprised 44% of all the 193 retouched pieces in the RGS member (Wadley, 2007). A more detailed analysis (Soriano et al., 194 2015) showed that stone knapping in RGS and RGS2 was mostly intended for the production of

195 bifacial foliate points and that most were broken during utilization, sharpening or resharpening, 196 so that many discarded tips and tip flakes were found. Dolerite, quartzite and hornfels were used 197 for the manufacture of the points. Residue and use-trace analysis of a few Sibudu Still Bay 198 bifacial points and point fragments suggests that they were mostly pointed hunting weapons, but 199 at least one of the points was a cutting tool (Lombard, 2006a) and the majority of points studied 200 by Soriano and colleagues seem to have been cutting tools. Raman Spectroscopy and Scanning 201 Electron Microscopy (Wozcieszak and Wadley, 2018) support the original interpretation that the 202 tools were hafted with an adhesive containing ochre powder. Other lithics in the Still Bay 203 assemblage include splintered pieces, a few segments and unifacial points, and scrapers of 204 various types (Wadley, 2007). Worked ochre is less common in the Still Bay than elsewhere in 205 the site (Hodgskiss, 2013). No worked bone has yet been found in the Still Bay layers at Sibudu. 206 In contrast, worked bone is extremely common in the younger Howiesons Poort Industry, 207 including pointed forms, awls, smoothers, pièces esquillées and pressure flakers (d'Errico et al., 208 2012b). The use of bone pressure flakers during the Howiesons Poort at Sibudu has been recently 209 supported by analysis of quartz bifacial points (de la Peña et al., 2013). The layers associated 210 with the Howiesons Poort are (from the base to the top): Pinkish Grey Sand (PGS), Grey Sand 211 (GS, GS2 and GS3), Dark Reddish Grey (DRG) and Grey Rocky (GR and GR2) (Wadley and 212 Jacobs, 2006). The stratigraphy is clear, but the rock fall mentioned earlier has caused some 213 disturbance to the oldest Howiesons Poort layer, PGS, as well as to the Still Bay layers. PGS has 214 an OSL age of  $64.7 \pm 1.9$  ka, while the younger Howiesons Poort layer, GR2, has an OSL age of 215  $61.7 \pm 2$  ka (Jacobs et al., 2008b). GR2 is an artificial spit to divide the light, brownish-grey silt 216 of GR. There are several hearths in GR. The Howiesons Poort layers have rich lithic assemblages 217 with many backed tools and small bladelets (Wadley and Mohapi, 2008; de la Peña and Wadley, 218 2014). Worked ochre is more common in the Howiesons Poort than in the Still Bay (Hodgskiss, 219 2013).

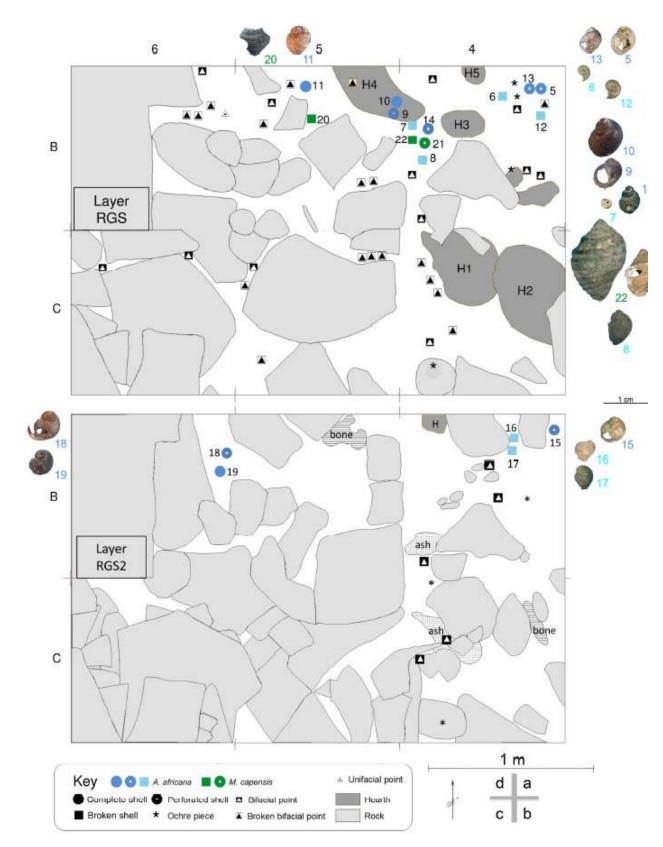


Figure 3. Spatial distribution of selected archaeological finds, rocks and ash lenses in squares BC/4-6 within layer RGS (top) and RGS 2 (bottom). Colors and symbols identify find category,
shell taxa and state of completeness. See Table 1 for descriptive information on the illustrated

- and numbered marine shell finds.
- A long sequence of post-Howiesons Poort (post-HP) layers occurs after the HP; these are about
- 226 100 cm in depth and they contain many strata wherein the inhabitants burnt bedding layers made
- of plant material (Goldberg et al., 2009; Wadley et al., 2011). The weighted mean age of the
- post-HP is  $58.5 \pm 1.4$  ka (Jacobs et al., 2008a). Some post-HP lithics have been described
- 229 (Cochrane, 2006; Villa et al., 2005; Conard et al., 2012; de la Peña and Wadley, 2017), as has the
- rich fauna (Clark and Plug, 2008). Layer BSp is the youngest of the post-HP layers and the late
- 231 MSA succeeds the post-HP. The Late MSA layer OMOD (Orange Mottled Deposit) has an age
- estimate of  $46.6 \pm 2.3$  ka. Lithics in OMOD include bifacial and unifacial points, scrapers and
- convergent flakes.
- 234 The GIS-based Coexistence Approach (CA<sub>GIS</sub>) analysis demonstrates that the late MSA was
- 235 warmer in winter than was previously the case, summer precipitation increased and vegetation
- became more closed (Bruch et al., 2012). *Podocarpus* sp. was consequently present in layer
- 237 OMOD (Hall et al., 2014). Furthermore, as was the case in the earlier SB and HP occupations
- 238 where vegetation was closed, small bovids like *Philantomba monticola* (blue duiker),
- 239 *Cephalophus natalensis* (red duiker) and *Raphicerus campestris* (steenbok) occurred (Wadley et
- al., 2008). Nonetheless, large animals like *Tragelaphus oryx* (eland), *Equus quagga* (zebra) and
- 241 Syncerus caffer (buffalo) were still present. Stable carbon and oxygen isotope data were retrieved
- from faunal tooth enamel throughout the sequence (Robinson and Wadley, 2018). The pre-SB
- seems to have been dominated by more closed environments than at other times, and was likely
- forested and mesic. More open and perhaps drier conditions occurred between  $\sim$ 48–38 ka ago,
- but the remaining periods were not dissimilar from today (Robinson and Wadley, 2018).
- 246 Throughout the Sibudu sequence there is bone preservation, though it is often burnt. People
- collected many suids and a diverse assemblage of small game in the pre-SB (Clark, 2019). High
- frequencies of blue duiker (*Philantomba monticola*) in the SB and before suggest the early use of
- remote capture technology (Clark 2019). In HP layers, and even earlier, there are remains of large
- bovids and bushpig, and small creatures are well-represented (Clark and Plug, 2008; Plug
- 251 personal communication 2009), including *Philantomba monticola* (blue duiker) and *Chlorocebus*
- 252 *pygerythrus* (vervet monkey). In the upper part of the post-HP, there is an increase in plains game
- such as zebra and small bovids become rare (Clark and Plug, 2008). Birds, reptiles, rodents and
- fish occur throughout the sequence, as well as fresh water and marine molluscs (Plug, 2006).
- 255 Marine species include turtles, fish, crustaceans and molluscs (Plug, 2006).
- 256 Since most of the shells that we describe here come from the SB layers of Sibudu, we provide

257 more detail on the broad context of this Industry.

#### **3. The Still Bay**

259 The southern African Still Bay (Goodwin and van Riet Lowe, 1929; Sampson, 1974) has the 260 lanceolate, bifacially shaped, point as its *fossile directeur*, but it also contains thin, long bifacial 261 points of other shapes. The thin Still Bay points tended to break easily both during manufacture 262 and use. The Industry at Blombos Cave on the coast in the Cape, more than 1000 km from 263 Sibudu, has a well-known and remarkable Still Bay sequence with many bifacial points. Heat 264 treatment of silcrete was used at this site to improve its quality for production, sometimes 265 through pressure flaking, of the fine points (Brown et al., 2009; Mourre et al., 2010; Villa et al., 266 2009; Soriano et al., 2015). Bone tools in the form of fully shaped points and awls produced by

267 scraping were found in Still Bay layers at Blombos and Peers Caves (d'Errico and Henshilwood,

268 2007). Modified red ochre is common at Still Bay sites (Henshilwood et al., 2009; Dayet et al.,

269 2013) and, at Blombos, fragments of hematite are engraved with abstract motifs (Henshilwood et

al., 2009). Perforated *Nassarius kraussianus* shells were used as beads by the Still Bay

inhabitants of Blombos Cave during the M1 archaeological phase (Henshilwood et al., 2004;

272 d'Errico et al., 2005; Vanhaeren et al., 2013).

273 Little is known about the origin of the Still Bay. At Sibudu and Blombos archaeological layers

274 immediately below the Still Bay contain relatively simple flake-based assemblages with few

275 retouched tools. In contrast, pre-Still Bay layers at Diepkloof and Klasies River contain unifacial

and bifacial pieces shaped with invasive retouch that appear to announce the technical

277 complexity of the Still Bay (Porraz et al., 2013).

278 To date, four sites – Sibudu (Wadley, 2007) and Umhlatuzana in KwaZulu-Natal Province

279 (Lombard et al., 2010; Mohapi, 2013), Diepkloof in the Western Cape Province (Rigaud et al.,

280 2006; Porraz et al., 2013), and Apollo 11 in the South of Namibia (Wendt, 1974; Vogelsang et

al., 2010; Lombard and Högberg, 2018) have stratigraphic sequences in which Still Bay

assemblages precede Howiesons Poort ones. Chronostratigraphy, Optically stimulated

283 luminescence (OSL) and Thermoluminescence (TL) dating of pre-Still Bay, Still Bay, and

Howiesons Poort layers have, for some years, situated the Still Bay between 72 and 71 ka and,

after a possible archaeological hiatus, the younger Howiesons Poort between 65 and 60 ka

286 (Jacobs et al., 2008b, 2013; Jacobs and Roberts, 2008). This view, i.e. that the Howiesons Poort

would be younger than the Still Bay, has been recently challenged by the OSL and TL dating of

the Diepkloof sequence (Tribolo et al., 2009; 2013) and fired the controversy surrounding

calculation of internal dose rate in OSL dating (Guérin et al., 2013, but see responses by

Galbraith, 2015; Jacobs and Roberts, 2015). Tribolo and colleagues propose a mean age of  $109 \pm$ 

- 10 ka for the Still Bay of Diepkloof, i.e. almost 40 ka older than the age estimated by Jacobs et
- al. (2008b) for the same site, and a duration of 50 ka (from  $105 \pm 10$  to  $65 \pm 8$  ka) for the
- 293 Howiesons Poort. They use these age estimates to support the hypothesis that both the Still Bay
- and the Howiesons Poort emerged during the last interglacial and coexisted during OIS 5 and 4
- in southern Africa. So far, there is no evidence in the KwaZulu-Natal region suggesting that the
- 296 Howiesons Poort was contemporaneous with the Still Bay.
- 297

#### 298 4. Material and Methods

- 299 4.1. Taxonomic identification
- 300 Modern and archaeological marine gastropods were identified at genus and, where possible,
- 301 species level using criteria proposed by Kilburn and Rippey (1982), Branch and Branch (1981),
- 302 Richards (1981), Branch et al. (1994), Marais and Seccombe (2010). Archaeological specimens
- 303 were compared to modern shell reference collections hosted at the KwaZulu-Natali Museum,
- 304 Pietermaritzburg. The taxonomic status of each species was checked and, where needed,
- 305 corrected using the WoRMS Editorial Board (2014).
- 306

# 307 *4.2. Survey of the biocoenoses*

- 308 Present day availability of marine gastropod species in the intertidal zone was checked at Ballito
- 309 Bay (29°32'23.52''S, 31°13'13.22''E) and Shaka's Rock (29°30'49.92''S, 31°13'57.03''E) by
- 310 shore bound surveying and snorkelling at rocky outcrops in February 2012. This led to the
- 311 creation of a list of identified species and determination of their relative abundance.
- 312

#### 313 *4.3. Survey of the thanatocoenoses*

- 314 A marine shell thanatocoenoses composed of 522 remains of marine gastropods was collected by
- the authors of this study in February 2012 at four locations: Shaka's Rock (29°30'49.92''S,
- 316 31°13'57.03''E), Ballito Bay (29°32'23.52''S, 31°13'13.22''E), Emerald Cove (30°47'00.80''S,
- 317 30°25'48.92''E) and Umdloti Beach (29°38'56.67''S, 31°08'00.86''E). These four locations are
- found along the sandy beaches and rocky outcrops of the KwaZulu-Natal coast. Dead shells were
- 319 sampled at each location for two hours during low tide between the upper beach and the
- 320 strandline. Shell fragmentation was described using five grades: 1) complete, 2) 99-75%, 3) 74-
- 321 50%, 4) 49-25%, 5) <24% preserved. Fragmentation properties were also described
- 322 independently of their location, according to genus by using a ternary taphogram (Kowalewski et
- al., 1995) in which the three states are represented by complete or almost complete specimens

- 324 (grades 1 and 2), damaged specimens (grades 3 and 4), and fragments (grade 5). The Software
- 325 PAST was used to produce the ternary taphogram.

# 326 4.4. Malacological reference collection

327 Our reference collection also comprises 470 Afrolittorina africana collected alive in September 328 2006 at two locations - uThongathi River mouth (29°34'22.93''S, 31°11'07.17''E) and Ballito 329 Bay (29°32'23.52''S, 31°13'13.22''E) - on the coast of Kwa-Zulu-Natal (d'Errico et al., 2008), 330 and four Nassarius kraussianus collected dead at the same time inside the river mouth and on the 331 oceanic beach close to the uThongathi River (Fig. 4). An additional five and nine dead N. 332 kraussianus were collected respectively in February 2012 and November 2013 on the oceanic 333 beaches of Ballito Bay (Kwa-Zulu-Natal) and Die Hoop (34°29'6.76''S, 20°29'28.16''E) in the 334 Western Cape). The surface of five N. kraussianus from Goukou (Western Cape) were also 335 analysed: one from the biocoenosis, one from the thantatocoenosis (d'Errico et al., 2005), as well 336 as two experimentally burned specimens and one burned and etched specimen; d'Errico et al., 337 2015). 3D scanning of a 877 x 660  $\mu$ m representative surface of these shells was performed with 338 a Sensofar S neox Confocal Imaging Profiler (Sensofar, Barcelone) after optical examination of 339 the piece. Surfaces were scanned with a 20x objective allowing a lateral sampling interval of 340  $0,645 \ \mu m$  and a vertical resolution of  $0.02 \ \mu m$ .



341

342 Figure 4. Marine shells from the Pleistocene layers of Sibudu Cave. See Table 1 for contextual

343 and descriptive information. 1-2: *Nassarius kraussianus*, 3-19: *Afrolittorina africana*; 20-23:

- 344 *Mancinella capensis*. Scale bar = 1 cm.
- 345

#### 346 4.5. Experimental reference collections

347 Experimental criteria to identify techniques used to perforate A. africana are known in the 348 literature (d'Errico et al., 2008). We have applied a similar approach to the other perforated 349 gastropod species found at Sibudu, Mancinella capensis. Seven types of tools were used to 350 perforate modern specimens of this species: a retouched lithic point, a small crab claw, a small 351 pebble, and bone and wooden awls of two sizes. Shells were perforated by punching the body 352 whorl through the aperture and from the outer dorsal side. The pebble was only used on the outer 353 dorsal side. For all experimentally perforated shells we recorded the location, morphology, size, 354 maximum diameter, and orientation of the perforations, the location of micro-chipping, and the height of the shell aperture. 355

356 Three fresh and two dead *Mancinella* shells were experimentally heated in a fire made from a

357 large sheaf of *Cyperus involucratus* culms. The fire combusted the culms rapidly; the maximum

temperate was in excess of 800°C for five minutes, then the fire died immediately.

359

## 360 4.6. Archaeological shells

361 Both sides of the Sibudu shells were digitised at a resolution of 1200 dpi with an Epson 362 Perfection 1660 Photo scanner. The resulting images served as a base to produce drawings of the 363 two aspects of each shell with Adobe Illustrator. The archaeological specimens were also examined and photographed with a motorised Leica Z6 APOA equipped with a DFC420 digital 364 365 camera and a Leica Application Suite (LAS) equipped with the Multifocus module. The 366 Multifocus module is designed to acquire extended depth of field images from the microscope. 367 Once digital images have been collected at different Z-positions, adapted algorithms combine 368 them into one single sharp composite image that massively extends the depth of focus. 369 Measurements were taken with a digital calliper. The location of spots of red pigment on shells 370 was systematically recorded and selected spots were analysed with a Jobin-Yvon T64000 Raman 371 spectrometer operated in triple subtractive mode. The 514.5 nm line of an argon ion laser was 372 used as the excitation source. Backscattered spectra were collected via an Olympus BX40 373 microscope Raman attachment, and the light dispersed via 1800 lines/mm gratings onto a liquid-374 nitrogen cooled CCD detector. Power for the sample was kept fairly low (1.2mW) to minimize 375 localised heating effects. A narrow bandpass filter was used to remove laser plasma lines from 376 the spectra. 3D scanning with a Confocal Imaging Profiler of a surface of a N. kraussianus from

377 Sibudu was performed as described above.

## 379 **5. Results**

380 5.1. Taxonomic identification

381 Twenty-three marine gastropods were recovered in the Pleistocene layers of Sibudu Cave (Table

382 1, Fig. 4). They belong to three species (Afrolittorina africana, Mancinella capensis and

383 Nassarius kraussianus) living in different habitats and having a different diet. Afrolittorina

384 *africana* (Krauss in Philippi, 1847) lives on exposed rocks located at the top of the intertidal

385 shore along the coast or in the mouth of estuaries; they shelter in little pools and moist crevices,

- 386 where they can be found in vast numbers (Branch and Branch, 1981; McQuaid, 1981). Juvenile
- 387 *A. africana* feed on bacteria and diatoms, adults mainly on lichens and blue–green algae.
- 388 Mancinella capensis (Petit de la Saussaye, 1852; Claremont et al., 2013), previously called

389 *Reishia capensis* (Petit de la Saussaye, 1852), can be found living under rocks and in rock

390 crevices in high energy wave action shores from low neap-tide downward. It feeds on large

391 ascidians and gastropods such as *Burnupena* spp. (Marais and Seccombe, 2010). *Nassarius* 

392 *kraussianus* (Dunker, 1846) is a scavenging gastropod adapted to estuarine environments

393 (Palmer, 1980; Branch et al., 1994). Present day distribution of *A. africana* and *M. capensis* 

394 ranges from Cape Aghulas to Northern Natal. *N. kraussianus* can be found all along the South

395 African coast, except the North-West coast.

Table 1. Taxonomic, contextual and descriptive data on the gastropod shells found in the Middle Stone Age layers of Sibudu Cave.

| Layer       | Cultural<br>attribution | Square     | Number | Date          | Species        | Perforation | Pigment | Heating | Length<br>(mm) | Perforation<br>max. diam.<br>(mm) | Perforation<br>min. diam.<br>(mm) |    |
|-------------|-------------------------|------------|--------|---------------|----------------|-------------|---------|---------|----------------|-----------------------------------|-----------------------------------|----|
| OMOD        | Late MSA                | A5a Hearth | 2404   | 30/08/2005    | N. kraussianus | no          | no      | no      | 6,67           | na                                | na                                | 1  |
| OMOD        | Late MSA                | C3c        | na     | 26/08/1999    | N. kraussianus | yes         | no      | yes     | 8.30           | 6.26*                             | 3.78*                             | 2  |
| GR2         | HP                      | C4b        | na     | 14/02/2009    | A. africana    | no          | no      | no      | 6.20           | na                                | na                                | 3  |
| PGS         | HP                      | B5d        | 2057   | 10/11/2004    | A. africana    | yes         | no      | no      | 5              | 2.4*                              | 2.27*                             | 4  |
| RGS         | SB                      | B4a        | na     | 17/11/2009    | A. africana    | yes         | no      | no      | 5.55           | 1.28**                            | 1.18**                            | 5  |
| RGS         | SB                      | B4a        | na     | 13/11/2009    | A. africana    | na          | no      | no      | na             | na                                | na                                | 6  |
| RGS         | SB                      | B4c        | na     | 15/11/2009    | A. africana    | na          | yes     | no      | na             | na                                | na                                | 7  |
| RGS         | SB                      | B4d        | na     | 15/11/2009    | A. africana    | na          | no      | yes     | 8,5            | na                                | na                                | 8  |
| RGS         | SB                      | B5a Hearth | 1989   | 11/11/2004    | A. africana    | yes         | yes     | yes     | 5.79           | 2.8*                              | 2.46*                             | 9  |
| RGS         | SB                      | B5a Hearth | 1989   | 11/11/2004    | A. africana    | no          | no      | yes     | 7.97           | na                                | na                                | 10 |
| RGS         | SB                      | B5c        | 1991   | 11/11/2004    | A. africana    | no          | no      | no      | 5.33           | na                                | na                                | 11 |
| RGS         | SB                      | B5c        | 1988   | 11/11/2004    | M. capensis    | na          | no      | yes     | na             | na                                | na                                | 20 |
| RGS true 2  | SB                      | B4a        | na     | 17/11/2009    | A. africana    | na          | yes     | no      | na             | na                                | na                                | 12 |
| RGS true 2  | SB                      | B4c        | na     | 17/11/2009    | M. capensis    | yes         | no      | no      | 10,5           | 2.8                               | 1.8                               | 21 |
| RGS brown   | SB                      | B4c        | na     | 23/11/2009    | M. capensis    | na          | yes     | yes     | 18.60          | na                                | na                                | 22 |
| RGS brown 2 | SB                      | B4a        | na     | 24/11/2009    | A. africana    | yes         | no      | no      | 6.20           | 2.91                              | 2.40                              | 13 |
| RGS brown 2 | SB                      | B4c        | na     | 25/11/2009    | A. africana    | yes         | no      | yes     | 6.15           | 1.75                              | 1.22                              | 14 |
| RGS2        | SB                      | B4a        | na     | 27/11/2009    | A. africana    | yes         | no      | no      | 6.11           | 2.23                              | 1.52                              | 15 |
| RGS2        | SB                      | B4a        | na     | 27/11/2009    | A. africana    | no          | yes     | no      | na             | na                                | na                                | 16 |
| RGS2        | SB                      | B4a        | na     | 27/11/2009    | A. africana    | na          | no      | yes     | na             | na                                | na                                | 17 |
| RGS2        | SB                      | B6a        | na     | 16/11/2004    | A. africana    | yes         | no      | yes     | 6.14           | 2.27                              | 1.67                              | 18 |
| RGS2        | SB                      | B6a        | na     | 16/11/2004    | A. africana    | no          | no      | yes     | na             | na                                | na                                | 19 |
| LBG         | pre-SB                  | B5a        | 2062   | 14-18/11/2004 | M. capensis    | na          | yes     | no      | na             | na                                | na                                | 23 |

N.: Nassarius; A: Afrolittorina; M.: Mancinella; na: not applicable; \*perforation with post-depositional damage; \*\*perforation produced by chemical alteration; MSA: Middle Stone Age, HP: Howiesons Poorts, SB: Still Bay, figure numbers in *italics* indicate shells published in d'Errico et al. 2008.

397 Table 1. Taxonomic, contextual and descriptive data on the gastropod shells found in the Middle

398 Stone Age layers of Sibudu Cave.

399

396

400 5.2. Analysis of the biocoenoses and thanatocoenoses

401 The survey of two marine shell biocoenoses along the Dolphin coast allowed for the

402 identification of marine gastropods belonging to 18 genera, among which Afrolittorina, Cypraea,

403 *Littorina*, *Nerita*, *Nodilittorina* and *Turbo* are the most represented (Table 2).

| Table 2. Marine gastropods collected in modern bio- and thanatocoenoses on the Dolphin coast of Kwa | -Zulu-Natal |
|---|-------------|
|---|-------------|

|                | Biocoenosis        |       | Thanatocoenosis |       |            |         |     |      | 32  | Fragmentation |     |     |       |                        | Sibudu   |      |   |     |  |  |
|----------------|--------------------|-------|-----------------|-------|------------|---------|-----|------|-----|---------------|-----|-----|-------|------------------------|----------|------|---|-----|--|--|
| Genus          | Ballito            | Shaka | Ballito         | Shaka | Emerald    | Umdloti | Ν   | %    | 1   | 2             | 3   | 4   | 5     | R                      | PHP      | HP   | SB                                      | PSB |  |  |
| Afrolittorina* |                    |       | -               |       | 5          | -       | 5   | 1.1  | 5   |               |     | -   | -     |                        |          |      | 2                                       |     |  |  |
| Agrobuccinum   | 12                 | -     | 1               | 2     | 12         | 1       | 4   | 0,9  | 122 | 2             | -   | 2   | 2     | 1                      | 72       | 22   | 122                                     | 2   |  |  |
| Buffonaria     | <u>i</u>           | 2     | 4               | 4     | 82         | 5       | 13  | 2,9  | 823 | $\sim$        | 1   | 3   | 9     | 525                    | 12       | 23   | 323                                     | 12  |  |  |
| Bullia         | 3 <u>2</u>         | 3     | 52              | 1     | 12         | 9       | 61  | 13,4 | 38  | 14            | 6   | 3   |       |                        | 82       | 28   | -                                       |     |  |  |
| Burnupena      |                    | 3     | 13              | 26    |            | 10      | 49  | 10,8 |     | 1             | 6   | 8   | 34    |                        | 12       | 74   |   | 3   |  |  |
| Bursa          | •                  | •     | 73              | 2.72  | æ          | 3       | 3   | 0,7  |     | -             |     | 1   | 2     |                        | -        |      |   |     |  |  |
| Charonia       | 35                 |       | -               |       | 2          | 4       | 6   | 1,3  |     | -             | -   | 3   | 3     |                        | 18       | -    | -                                       | -   |  |  |
| Conus          | 84                 |       | 2               | 1     | 14         | 4       | 21  | 4,6  | -   | ~             | -   | 3   | 18    |                        | 28       | -    | -                                       | -   |  |  |
| Crepidula      | 12                 | 2     | 1               | 12    | 12         | 1       | 2   | 0,4  | 1   | 1             |     | -   | 1.0   | 1                      | 12       | 22   | -                                       | 4   |  |  |
| Cronia         |                    | 2     | 2               | 120   | 3 <b>4</b> | -       |     | 12   |     | 12            |     | -   | 1.4   | 1                      | 12       | 22   | 123                                     | 2   |  |  |
| Cymatium       | 32                 | 9     | 15              | 8     | 14         | 6       | 43  | 9,5  | 1   | 2             | 2   | 5   | 33    |                        | 82       | 28   |   | 3   |  |  |
| Cypraea        |                    |       | 2               | 5     | 7          | 10      | 24  | 5,3  | 7   | 02            |     | 5   | 12    |                        | 17       | -    | 0.50                                    | ~   |  |  |
| Epitonium      |                    |       | 1               | 3.00  |            | 1       | 2   | 0,4  |     | 2             |     | -   | -     | -                      |          |      | -                                       |     |  |  |
| Ficus          | 2                  | -     | 2               | 1     |            | 3       | 3   | 0,7  |     | 2             | 1   | 1   | 1     |                        |          |      | -                                       |     |  |  |
| Fusinus        |                    | 2     |                 |       |            | 1       | 1   | 0,2  | -   | 2             | -   | - 2 | 1     | -                      |          | -    | -                                       | 2   |  |  |
| Hydatina       |                    | 2     | 22              | -     | 34         |         |     |      | -   | -             | -   | -   | 0-0   | -                      | 19       | -    | -                                       | -   |  |  |
| Janthina       | 12                 | 12    | 2               | 823   | 3          | 4       | 9   | 2,0  | 823 | 7             | 2   | 2   | 32313 |                        | 12       | 23   | 1.0                                     | 12  |  |  |
| Littorina      |                    |       |                 | 12    | 10 - C     |         |     |      |     |               |     | 8   |       |                        | 82       | 28   |   | 3   |  |  |
| Mancinella***  |                    | -     | 8               | 44    | 6          |         | 58  | 12,7 |     | -             |     | 4   | 54    |                        |          |      | 3                                       | 1?  |  |  |
| Minolia        | -                  | -     | 3               |       | -          |         | 3   | 0,7  | 1   | 1             | 1   |     | -     |                        | -<br>-   |      | -                                       |     |  |  |
| Morula         |                    |       | -               | 2     | 1          | 5-6     | 3   | 0,7  | 1   | -             | -   | 1   | 1     |                        |          | -    |   | -   |  |  |
| Natica         |                    | 2     | 22              | 1     |            | 1.0     | 1   | 0,2  |     |               | -   | 2   | 1     | 12-11                  |          | - 20 | i anna anna anna anna anna anna anna an | 2   |  |  |
| Nassarius**    | 12                 | 2     | 5               |       | 2          | 122     | 5   | 1,1  | 3   | 2             |     | -   | 120   | 1                      | 2        | 28   | 12.2                                    | 2   |  |  |
| Nerita         |                    |       | 6               | 12    | 2          | 120     | 18  | 4,0  | 3   | 2             | 7   | 4   | 2     |                        | <u>_</u> | 23   |   | 2   |  |  |
| Nodilittorina  |                    |       | Ĩ               |       | 22         | 100     | -   |      |     |               |     | 1   |       |                        | 83       | 23   |   |     |  |  |
| Oliva          | 1000<br>1 <b>-</b> | -     | -               | 2     | 1          | 3       | 6   | 1,3  |     | 3             | -   | 1   | 2     | -                      |          | -    |   | -   |  |  |
| Oxystele       |                    | -     | 17              | 13    | -          | 8       | 38  | 8,4  | 24  | 8             | 2   | 1   | 3     |                        |          |      | 2                                       | -   |  |  |
| Peristernea    |                    | -     | -               | 1     | 1          | 1       | 3   | 0,7  | 2   | -             |     | -   | 1     | 1949)<br>19 <b>-</b> 8 | -        | -    | -                                       | -   |  |  |
| Phalium        |                    | 2     | 22              | 3     | 1          | -       | 4   | 0.9  | -   | 2             | -   | -   | 4     | 1.00                   |          | - 20 | i anna anna anna anna anna anna anna an | 2   |  |  |
| Philippia      |                    | 2     | 22              | -     | 1          | 823     | 1   | 0,2  | 122 | 1             |     | 12  | 120   | 1                      | 12       | 28   | 12.1                                    | 2   |  |  |
| Purpura        | 12                 | 2     | 23              | 2     | 2          | 120     | 2   | 0.4  | 823 | 2             | 1   | 25  | 2     |                        | 12       | 23   | 3 <b>.</b>                              | 2   |  |  |
| Ranella        | 32                 | •     | 4               | 2     | 8          | 8       | 20  | 4,4  |     |               | 1   | 12  | 19    |                        | 83       | 23   |   |     |  |  |
| Strombus       |                    |       | - 24            | -     | -          | -       | -   |      | -   | -             | ್   | -   |       |                        |          | -    |   | -   |  |  |
| Thais          |                    | 8     | 2               | 5     | 4          | 1       | 12  | 2,6  | 1   | 2             |     | 2   | 7     | 1999                   |          | 7    |   | -   |  |  |
| Throchus       |                    |       | 2               | -     | 1          | 1       | 4   | 0,9  | 1   | 2             | 1   | 2   | -     |                        |          | 2    |   |     |  |  |
| Tonna          | 52                 | 2     | 1               | 2     | 4          | 5       | 12  | 2,6  | -   | -             | -   | 4   | 8     | 1943)<br>2520          | 38       | 28   | 1                                       | 100 |  |  |
| Tricolia       | 12                 | ÷.    | 12              | 2     | 3          | 2       | 17  | 3,7  | 16  | 1             |     |     | •     | 0.00                   |          | 2    | 3                                       | 1   |  |  |
| Turbo          |                    |       | 12              |       | -          | -       | 1/  | 5,7  | 10  | 1             | -   |     | 100   | 100                    |          | 4    | -                                       | ÷.  |  |  |
| Turritella     |                    |       | 3               | 100   | 10         | 2       | 2   | 0,4  |     | 2             |     | 8   | 100   |                        | 8        | -    | 100                                     | 6   |  |  |
| Total          | 2                  |       | 155             | 134   | 76         | 81      | 455 | 0,1  | 104 | 51            | 30  | 51  | 219   |                        | 2        | 9    | 11                                      | 1   |  |  |
|                | -                  |       |                 |       |            |         |     |      |     |               |     |     |       |                        |          |      |   |     |  |  |
| Indetermined   | 32                 | 9     | 11              | 37    | 10         | 13      | 70  | 13,4 | 1.2 | -             | 1.0 | 2   | 67    | 1                      | 4        | 36   | 1                                       | 7   |  |  |

•: present, ••: abundant, •••: very abundant, R: recent, HP: Howiesons Poort, SB: Still Bay, PHP: Post Howiesons Poort, PSB: Pre-Still Bay

\*: Afrolittorina africana, \*\*: Nassarius kraussianus, \*\*\*: Mancinella capensis, \*\*\*\*: one specimen bears a perforation on the body whorl

406 Table 2. Marin gastropods collected in modern bio- and thanatocoenoses on the Dolphin coast of407 KwaZulu Natal.

408

409 Analysis of four thanatocoenoses along the same coast identified between 17 and 22 genera

410 (Table 2). Conus, Cymathium, Cypraea, Thais and Tonna are found in all samples. Of the three

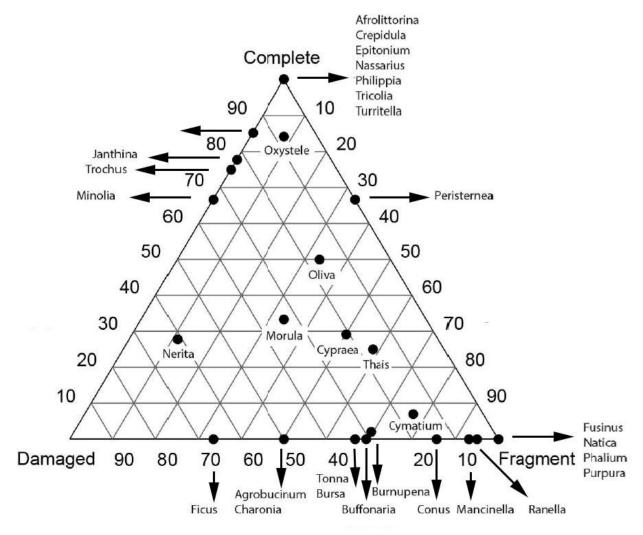
411 shell species found at Sibudu, only two, A. africana and M. capensis, are found alive on present

412 day shores of the Ballito Bay area (Table 2, Fig. 4). Although the biocoenosis of this species was

413 not surveyed by us, *N. kraussianus* is likely living in local estuaries and reported from the

414 Mzingazi in Richards Bay, 120 km up north (Teske et al., 2007). The scarcity of dead A. africana

- 415 and the fragmentary state of *M. capensis* are probably due to the strong wave action, which
- 416 crushes or breaks them as soon as they are dislodged by waves. *N. kraussianus* is apparently
- 417 more resistant to wave action as we found it on the Ballito Bay shores, 5 km away from the
- 418 closest estuary. The degree of fragmentation varies according to genera (Table 2, Fig. 5).



420 Figure 5. Ternary taphogram showing the state of fragmentation of identified marine shells

421 genera found on the Dolphine's coast beaches. See methods and Table 2 for information on how

422 the taphogram was made and on which malacological reference collections were included.

- 423
- 424 Afrolittorina, Bullia, Crepidula, Epitonium, Nassarius, Oxystele, Peristernia, Philippia, Tricolia
- 425 and *Turritella* are most often complete, while *Agrobuccinum*, *Buffonaria*, *Burnupena*, *Bursa*,
- 426 Charonia, Conus, Ficus, Fusinus, Mancinella, Natica, Phalium, Purpura, Ranella, and Tonna,
- 427 are only found as damaged specimens or fragments. The remainder of the genera are represented
- 428 by at least one complete or almost complete specimen.

- 429 Natural perforations only occur on two species, *Nassarius* and *Burnupen*a. Nine out of ten
- 430 *Nassarius* found on Ballito Bay and Thongathi beaches bear perforations, three on the ventral
- 431 and seven on the dorsal side. Perforations on the ventral aspect of the recovered *Nassarius*
- 432 consist of worn holes bored into the shell by *Natica* (Palmer, 1980), those on the dorsal aspect
- 433 result from mechanical abrasion by sand and wave action on the beach (Fig. 6).

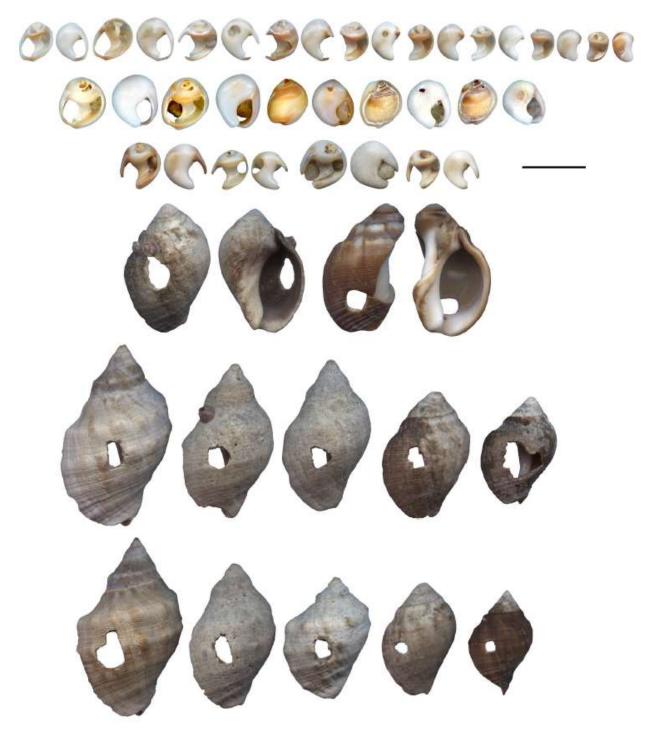


Figure 6. Top: *Nassarius kraussianus* collected dead on the sandy beaches of Die Hoop in the top
row, Ballito Bay in the middle row and close to the uThongathi River in the bottom row. Bottom: *Mancinella capensis* and *Burnupena burnupena* with natural perforations (top row) and
experimental perforations made with a stone (middle row) and a bone tool (bottom row) exerting
pressure on the inside of the shell through its aperture.

440

441 All perforated beached *Nassarius* display granules firmly stuck within the shell's interior.

442 Alterations on *Nassarius* from Ballito and Thongathi thanatocoenoses differ from those recorded

443 on dead *Nassarius* from estuarine environments (d'Errico et al., 2005: 12, fig.4) where gradual

decalcification in calm waters is the main taphonomic agent. The latter are porous and display

445 perforations with thin, crenulated edges resulting from the gradual weakening of the shell wall

446 whereas Ballito and Thongathi Nassarius are not etched, have smooth hole-edges and a shiny

447 appearance (Fig. 6). At microscopic scale, micro-pits due to impacts are observed (Fig. 7b).

448 Abrasion first produces sub-millimetric perforations on the spire, then causes an enlargement of

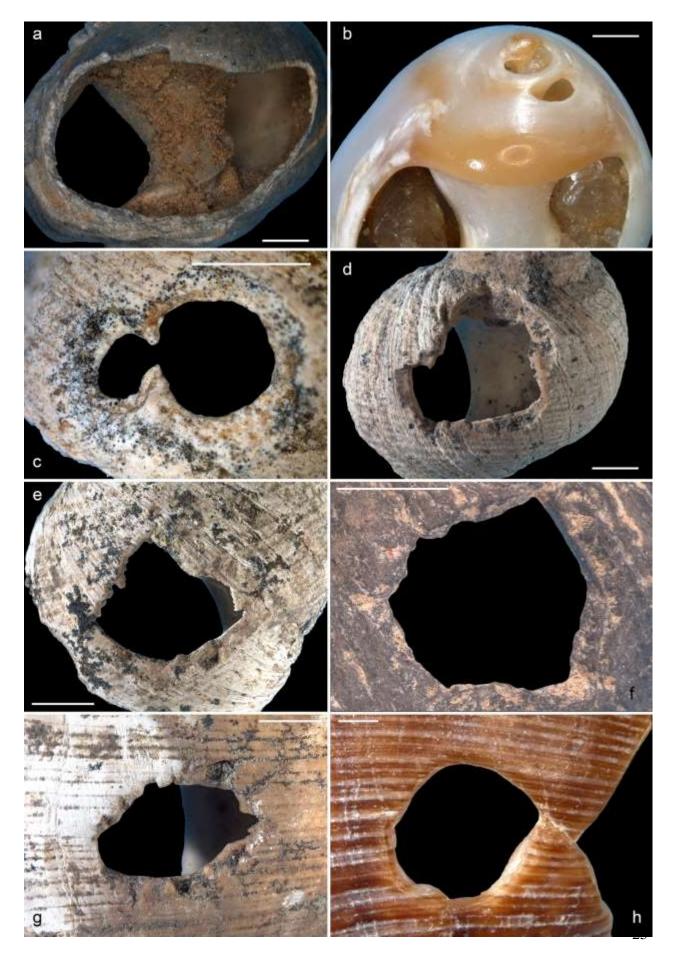
that perforation, the opening of a second millimetric perforation on the spire, the removal of the

450 apex and the opening of a wide perforation on the body whorl that increases its size until partial

451 removal of the shell's lip. The surviving segment of the lip takes a typical pointed morphology

452 characteristic of beach worn Nassarius (Fig. 6). Only one fragment of Burnupena over the 49

453 collected bears a sub-circular worn perforation of unknown origin (Fig. 7h).



- 457 Figure 7. Macro photos of a) the perforated *Nassarius kraussianus* from Sibudu (Fig. 4 n°2), b) a
- 458 beach worn *Nassarius kraussianus*, c) two very small juxtaposed perforations interpreted as
- 459 resulting from a chemical alteration on *Afrolittorina africana* (Fig. 4 n°5), d-e) perforations on
- 460 Afrolittorina africana (Fig. 4 n°15 and 14) that must result from pressure on the inside of the
- shell, g) perforation on *Mancinella capensis* (Fig. 4 n°21), and h) natural perforation on
- 462 Burnupena from Umdloti.
- 463

# 464 *5.3. Stratigraphic and spatial distribution.*

465 Marine gastropods were recovered in the pre-Still Bay, Still Bay, Howiesons Poort and late MSA

466 layers (Table 1). N. kraussianus is only found in the late MSA, A. africana in different

467 Howiesons Poort and Still Bay layers, *M. capensis* in the Still Bay and possibly in the pre-Still

Bay. The discovery of a single *A. africana* in the lowermost Howiesons Poort layer PGS and the

469 nature and thickness of this layer has suggested (d'Errico et al., 2008) that this specimen may

470 have derived from the underlying Still Bay layer RGS, where most of the other shells were found.

471 The enlarged collection of marine gastropods described here includes another *A. africana* from

472 Howiesons Poort layer GR2, which confirms the presence of marine gastropods in Sibudu

473 Howiesons Poort layers.

474 In Still Bay layers two clusters of five and four perforated A. africana shells were found in the 475 same lens subdivided in three sub-units of different colour (RGS, RGS brown, RGS true). The 476 first cluster is associated with two hearths (H3-4); the other is located East of hearths H3 and H5 477 (Fig. 3). Two lumps of ochre occur together with the second cluster in B4a (Fig. 3). Two other 478 clusters of three and two shells were found in the underlying lens RGS2: one in the same location 479 as the second cluster of the overlying lens RGS; the other in sub-square B6a (Fig. 3). Another 480 piece of ochre is found nearby and a fourth one at one meter in square C4 (Fig. 3). The three Still 481 Bay M. capensis were found within the first cluster of lens RGS (Fig. 3). This clustering of the 482 perforated shells contrasts with the dispersed distribution of bifacial points and point fragments

483 484

# 485 5.4. Taphonomic and technological analysis

in the same layer (Fig. 3).

486 The absence of features diagnostic of estuarine (d'Errico et al., 2005) or beach induced alterations

487 (Fig. 6, 7) suggests that the two *N. kraussianus* from the late MSA layer OMOD were either

488 collected alive or within an estuarine thanatocoenosis soon after their death. One is complete

489 (Fig. 4  $n^{\circ}$ 1), the other (Fig. 4  $n^{\circ}$ 2) bears a perforation on the dorsal aspect and traces of heating 490 that caused blackening and weakening of the shell (d'Errico et al., 2015). The latter has probably 491 favoured the micro breakage of most of the perforation edge (Fig. 7a). A portion of the original 492 perforation edge is nevertheless preserved and reveals a smooth appearance comparable to that 493 produced on experimentally worn N. kraussianus (Vanhaeren et al. 2013). Although the 494 smoothing visible on the preserved portion of the original perforation edge is comparable to that 495 observed on highly abraded beach worn N. krausianus, such a taphonomic process can be 496 discarded for this archaeological specimen since it would entail advanced damage, not observed 497 here, on the spire.

498 Of the 17 A. africana, four are complete or only slightly damaged (Fig. 4 n°3,10,11 and 19),

499 seven have a perforation on the body whorl (Fig. 4 n°4,5,9,13-15 and 18), two have their body

500 whorl partially removed by breakage (Fig. 4  $n^{\circ}$  8 and 17), three are apical fragments (Fig. 4  $n^{\circ}$ 6-7

and 12), and one consists of a portion of the body whorl (Fig.  $4 n^{\circ}16$ ). Seven are blackened by

beating and four bear micro-residues of red material (Table 1). Among the seven perforated

503 shells, one displays two conical contiguous perforations with evidence of etching suggesting

504 chemical dissolution (Fig.  $n^{\circ}7c$ ). The perforations on the other *A. africana* (Fig. 7d-f) are similar

to those produced experimentally by punching the inner aspect of the body whorl, i.e. through the

aperture, with a pointed tool (d'Errico et al., 2008, Fig. 7-8). This is suggested by their location

and the presence of micro-chipping around the perforation edge on the outer surface of the shell.

508 In one case (Fig. 4 n°18) this action was performed twice (d'Errico et al., 2008), probably to re-

509 use the shell after breakage of the first perforation. In two cases, post-depositional damage of the

510 original perforation (fig. 4 n°4 et 9) prevents identifying the morphology of the tool-tip used to

511 produce the holes. The hole morphology and location on the other specimens support the use of a

512 thin point made of bone or hard wood. Microscopic analysis of these perforations reveals no

513 obvious traces of use-wear (Fig. 7).

514 Of the three (Fig. 4 n°20-22), possibly four (Fig. 4 n°23), *M. capensis*, one (Fig. 4 n°22) is well

515 preserved, blackened by heating, and misses its outer lip and a part of the adjacent body whorl. A

516 second (Fig. 4 n°21) exhibits a perforation on the body whorl, a broken outer lip with post

517 depositional damage, and misses its apex. A third (Fig.4 n°20) is only represented by a small

518 portion of a body whorl which is blackened by heating and displays crenulated edges identical to

519 those observed on specimens of the same species experimentally broken by intense heating. The

520 fourth specimen, possibly *M. capensis* (Fig. 4 n°23), consists of a fragment which displays, on

521 one side, a crenulated edge similar to the one described for the third specimen. On the other

- sides, it has smoothed edges as well as an orange stain on the dorsal and the ventral aspect which
- 523 contrasts with the light brownish grey colour of the sediment of layer LGB.
- 524 Experimental attempts to perforate *M. capensis* shells reveal that punching the shell from the
- 525 outer surface is ineffective whatever tool-type (bone or lithic point) used. Pressure through the
- 526 aperture with a bone or lithic point resulted in sub-rectangular perforations with edges exposing
- 527 the inner prismatic layer and occasional micro-chipping (Fig. 6). These features are also found on
- 528 the single perforated *M. capensis* from Sibudu (Fig. 4 n°21). As with the *A. africana*, the
- 529 perforation on the *M. capensis* shows no compelling evidence of usewear.
- 530

# 531 6. Discussion

532 In previous papers we reviewed the potential causes for the accumulation of marine gastropods at 533 inland southern African sites (d'Errico et al., 2005, 2008) and concluded that humans were to be 534 considered the more likely agent responsible for the accumulation and modification of six A. 535 africana found at Sibudu (d'Errico et al., 2008). Analyses of 17 supplementary gastropods (A. 536 africana, N. kraussianus and M. capensis) from Sibudu MSA layers, local bio- and 537 thanatocoenoses, and additional perforation experiments allow us to assess this diagnosis anew. 538 The distance of Sibudu from the sea during its Pleistocene occupation rules out the possibility 539 that these marine shells were accumulated by natural processes. The survey and fragmentation 540 analysis of natural gastropod assemblages confirm that the Sibudu gastropod collection is not 541 representative of present day biocoenoses or thanatocoenoses of the Dolphin Coast. Taxa well 542 represented in the biocoenoses such as Burnupena, Littorina, Morula, Nerita, Nodilittorina, and 543 Peristerna, are absent at Sibudu. Similary, Bullia, Burnupena, Cymatium, Oxystele, and Ranella, 544 representing 50% of present day thanatocoenoses, are absent at Sibudu. It is unlikely that the 545 Sibudu gastropods were collected for consumption or to be used as tools. Almost all of them are 546 tiny gastropods with low caloric value (2 kcal for 0.4 g of dry soft tissue extracted from 100 A. 547 africana and 4 kcal for 0.8 g extracted from 100 N. kraussianus; d'Errico et al., 2005, 2008) 548 which makes a subsistence strategy based on transporting them for 15 km ineffective. Although 549 the bivalves, like *Perna perna*, which is present at the site, and a few gastropods found at Sibudu 550 could have been transported for food, their low numbers suggest that the consumption of 551 seafood, if it did occur, was only occasional at the site (Plug, 2006). The small size of the 552 gastropods is also in contradiction with their use as tools (e.g. containers). 553 Sibudu *N. kraussianus* shells bear surface features that are incompatible with a gathering of dead 554 specimens on beaches and rather point to a collection of living specimens or a thanatocoenosis

555 soon after the death of the organisms in their estuarine habitat. N. kraussianus is listed among the 556 64 species of marine shells found in the stomach of a loggerhead sea turtle (*Caretta caretta*) 557 (Hughes, 1974). The identification of remains of this species in the MSA layers of Sibudu (Plug, 558 2006) opens the possibility that N. kraussianus reached the site accidentally in the stomach of 559 prey. Caretta caretta crushes large gastropod shells before ingurgitating them as they are only 560 found in the form of fragments in the turtle's stomach (Hughes, pers. com.). Small gastropods, 561 such as *N. kraussianus*, are probably swallowed whole. However, it is unlikely that *Caretta* 562 *caretta* made the keyhole perforation on one of the two Sibudu *N. kraussianus*. This perforation 563 type is rarely found in estuarine thanatocoenoses and when it does it only occurs on decalcified 564 shells (d'Errico et al., 2005), which is not the case for the Sibudu specimen. The state of 565 preservation of this archaeological specimen is also incompatible with the use-wear observed on 566 beach worn Nassarius. In contrast, even if damaged, the use wear on a well preserved portion of 567 the perforation edge is comparable to that observed on archaeological and experimental N. 568 kraussianus shells used as beads (Vanhaeren et al., 2013). The use of N. kraussianus as beads is 569 documented in the Still Bay of Blombos Cave, dated to ca. 72 ka (Henshilwood et al., 2004) and 570 in the Early Later Stone Age of Border Cave, dated to between 42-44 ka cal BP (d'Errico et al., 571 2012). The perforated Sibudu N. kraussianus comes from a layer dated 46.6  $\pm$  2.3 ka (Jacobs 572 2008a). Transport in the stomach of a turtle cannot be discarded for the complete N. kraussianus. 573 Its occurrence in the same late MSA layer as the perforated specimen makes it however equally 574 possible that it corresponds to a shell lost or discarded before being perforated to be used as bead. 575 No natural agents are known to bring A. africana or M. capensis to inland sites and neither of 576 these two species is reported as being found in the stomach of identified prey at Sibudu. Littorinidae are present in the Still Bay layers of Blombos Cave (d'Errico et al., 2005) and the 577 578 MSA layers of Klasies River (Thackeray, 1988). Due to lack of clear human induced 579 modifications, they have been interpreted as "incidental shells" which came to the site 580 accidentally with seafood (e.g. attached to the byssus of mussels). This is unlikely for the A. 581 africana from Sibudu considering the scarcity of shellfish consumption at this site. Also, none of 582 these Littorinidae from Blombos or Klasies River are perforated, while a third of those from 583 Sibudu are, and another third may have been perforated before post-depositional damage. 584 The spatial distribution of A. africana and M. capensis provides clues for discussing the reasons 585 for their presence at the site. Contrary to the lithic bifacial points, found all over the excavated 586 area, perforated and clearly unperforated gastropods are found in clusters associated with some 587 hearths and are absent around other hearths (Fig. 3). This indicates that different spatial use and

588 patterns of discard applied to the shells and the points. It also suggests that the loss or disposal of 589 the shells is the consequence of an activity conducted close to hearths but that this activity did 590 not systematically occur every time a hearth was created and maintained. This pattern also 591 contradicts the hypothesis according to which the shells found at the site correspond to accidental 592 losses of individual beads during a variety of everyday activities. If this was the case, one would 593 expect to find a more widespread if not random distribution of exclusively perforated or broken 594 shells. The observed spatial distribution is compatible with an occasional subsistence activity or 595 the manufacture or maintenance of beadwork.

596 The perforations observed on the Sibudu *A. africana* are of three types: 1) natural perforation

597 either made by a scavenger or by post-depositional chemical dissolution (Fig. 4 n°5), 2)

598 perforations produced and/or enlarged by post-depositional processes for which an anthropogenic

599 intervention is difficult to prove (Fig. 4 n°4 and 9) and 3) small perforations with features

600 implying that they have been produced by punching the shell through its aperture (Fig. 4 n°13-15

and 18). The only perforated *M. capensis* (Fig. 4  $n^{\circ}21$ ) falls in this last category.

602 Considering the small caloric value of *A. africana*, shells belonging to this last category could

603 well have been intended for beads. Microscopic analysis of their perforation edges identifies no

or undiagnostic use-wear that would unambiguously demonstrate their use as beads. The absence

of use-wear could be explained by the fact that 1) they have been lost or disposed before use-

606 wear could develop, 2) they were attached in such a way that use-wear could not develop, 3) their

607 use as beads produced breakage of the perforation edges rather than identifiable wear, 4) they

have been perforated by humans for an unknown reason, and 5) the perforations were made by an

609 unknown natural agent. The first two hypotheses are consistent with the spatial distribution of

archaeological finds within layer RGS and RGS2. The third hypothesis is contradicted by the

611 presence, among the perforated specimens, of small perforations that can hardly result from

612 enlargement of previously even tinier holes (Fig.4 n°14). The fourth and fifth hypotheses are

613 difficult to test at present as they would require a larger assemblage and, probably, the creation of

an experimental protocol seeking to explore the impact of post-depositional damage on shells

615 incorporated in Southern Africa multi-layered MSA sequences in shelters and cave.

616

#### 618 **7. Conclusions**

619 Three species of perforated marine shell were found in MSA layers of Sibudu: Nassarius 620 kraussianus, Afrolittorina africana and Mancinella capensis. Nassarius kraussianus was found 621 only in the late MSA in OMOD layer dated to ca 47 ka; it did not occur in Howiesons Poort or 622 earlier industries. A N. kraussianus shell bead and an unperforated shell belonging to the same 623 species were recovered from OMOD. The perforated shell had been heated. N. kraussianus was 624 found at Border Cave, north of Sibudu, in early LSA contexts. It was first used as a personal 625 ornament in the Western Cape Still Bay. Sibudu confirms what has been recently highlighted at 626 Border Cave (d'Errico et al., 2012): after an early introduction, for example at Blombos, this 627 species was subsequently used to produce beads at late MSA/early LSA sites located along the 628 eastern flank of South Africa. The absence, at Sibudu, of ostrich eggshell beads, found at Border 629 Cave in association with the *N. kraussianus*, may indicate that ostrich eggshell beads are an 630 intrusive phenomenon in this part of southern Africa and that in the earliest phases of their spread 631 their distribution may have been restricted to north-east KwaZulu-Natal. Ostrich did not, and 632 does not, occur in the Sibudu area, and no ostrich bones or eggshell occur in the MSA layers of 633 the site.

634 Seven perforated A. africana shells were found in Sibudu, as well as ten unperforated A. africana 635 shells. Six perforated A. africana shells were from the Still Bay and one from the Howiesons 636 Poort, though the Howiesons Poort one may be misplaced from the Still Bay as the result of rock 637 fall disturbance. Three of the unperforated and one perforated shell showed traces of pigment, 638 while three perforated and four unperforated shells had been heated. The only perforated M. 639 *capensis* shell was found in the Still Bay and this shell's small perforation has features implying 640 that it was produced by punching the shell through its aperture. Three unperforated M. capensis 641 shells were found in the Still Bay; two had traces of pigment in them and two had been heated. 642 The interpretation of Sibudu's A. africana shells as beads used for a short time only, together 643 with unused shells, or shells intended for perforation represents, at present, the most 644 parsimonious explanation for this evidence. However, we are still far from the degree of certainty 645 attained at other MSA sites from northern and southern Africa where the larger number of 646 specimens found, and the consistency in traces of manufacture and utilisation make the diagnosis 647 more robust. If formally demonstrated, the use of shell beads at Sibudu during the late MSA, 648 Howiesons Poort and Still Bay occupations would represent supporting evidence for shell bead 649 use in the Howiesons Poort and early LSA, already suggested for Border Cave (d'Errico et al. 650 2012; Beaumont and Bednarik, 2013). Furthermore, it would corroborate the evidence for

- 651 comparable behaviour by Still Bay populations, attested at Blombos Cave (Henshilwood et al.,
- 652 2004). The use of different shell species for making beads at Sibudu suggests that the Howiesons
- 653 Poort and the Still Bay are not a homogeneous cultural phenomenon. Instead, these entities are
- associated, in different regions of southern Africa, with distinct bone tool technologies (d'Errico
- et al. 2012) as well as personal ornament traditions (d'Errico and Backwell, 2016). Moreover, the
- 656 presence of *N. kraussianus* beads in Sibudu's final MSA and Border Cave's early LSA points to
- 657 dynamically changing traditions through time.
- 658

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