

1 **Earliest human burial in Africa**

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70 **The origin and evolution of mortuary practices are topics of intense interest and**
71 **debate. Human burials dated to the African Middle Stone Age (MSA) are**
72 **exceedingly rare, and unknown in East Africa. Here, we describe the partial**
73 **skeleton of a c. 2.5-3.0 year-old child dating to 78.3 ± 4.1 ka, recovered in the MSA**
74 **layers of Panga ya Saidi (PYS), an archeological site in the tropical upland coast**
75 **of Kenya. Recent excavations revealed a pit feature containing a child in a flexed**
76 **position. Taphonomic, geochemical, granulometric, and micromorphological**
77 **analyses of the burial pit content and encasing archaeological layers indicate that**
78 **the pit was deliberately excavated. Strict articulation or good anatomical**
79 **association of the skeletal elements and histological evidence of putrefaction**
80 **support the in-place decomposition of the fresh body. Absent to minimal**

81 **displacement of the unstable joints during decomposition points to an interment**
82 **in a filled space (grave earth) making PYS the oldest human burial in Africa. The**
83 **morphological assessment of the partial skeleton is consistent with its assignment**
84 **to *H. sapiens*, although the preservation of some primitive features in the dentition**
85 **supports increasing evidence for non-gradual accretion of modern traits during**
86 **the emergence of our species. The burial sheds new light on how MSA populations**
87 **interacted with the dead.**

88

89 Increasing scrutiny is being placed on the interplay between biological and cultural
90 factors in the evolution of our lineage and the emergence of our species in Africa¹⁻².
91 Mortuary practices play a key role in this debate as growing evidence supports an
92 ancient origin and possible long-term evolution of these behaviours. Formal burials,
93 defined as the interment of a dead body in an excavated grave, may have been preceded
94 by more elusive practices and only performed by the latest representatives of our genus.
95 Testing this scenario is made difficult, particularly in Africa, by the scarcity of sites
96 with clear and well-dated evidence of dead body treatment.

97 Panga ya Saidi (PYS) has emerged as one of the key Middle Stone Age (MSA) and
98 Later Stone Age (LSA) sites of Africa given its excellent preservation of environmental
99 proxies which demonstrate its unique long-term position in a coastal tropical forest-
100 grassland ecotone^{5,6}, as well as its distinctive and continuous sequence of technological
101 innovations and symbolic traits^{6,7}. The excavated cave sequence is ~3 metres deep and
102 encompasses 19 layers (Fig. 1). A series of 20 stratigraphically ordered radiocarbon
103 and optically stimulated luminescence (OSL) ages, when included in a Bayesian model,
104 indicate human occupation from ~78 ka to 500 years ago⁶, including in each of the last
105 five marine isotope stages.

106 The 2013 excavations at PYS revealed a partial pit feature in profile, markedly
107 contrasting in texture and colour in comparison to the surrounding matrix (Fig. 1,
108 Supplementary Information A). An OSL tube (OSL4) and a micromorphology sample
109 (PYS 13_1) were placed in the feature in 2013. Upon removal, the micromorphology
110 sample revealed the presence of heavily degraded bones inside the section. Excavations
111 were expanded in 2017 to expose the top of the feature, which was positioned at the
112 bottom of MSA Layer 18. The plan view of the pit was subcircular, measuring 36.7 cm
113 (north-south) x 39.8 cm (west-east) x 12.5 cm (depth) (Fig. 1b, Supplementary
114 Information A). Limited hand excavation at the top of the pit indicated that the feature
115 contained a concentration of fragile and degraded bones in association with MSA lithic
116 artefacts (Extended Data Fig. 1, Supplementary Information B), embedded in a matrix
117 different from the surrounding sediments of Layer 19 (Supplementary Information A).
118 The excavation surface indicated the presence of decomposed bones, in what was later
119 shown to be the base of a skull and an articulated spine of a child (Fig. 1b).

120 Several small fragments of unidentified bone were recovered during the 2017
121 excavation but due to their poor state of preservation, it was decided to plaster the whole
122 feature and transport it for careful laboratory excavation. The plastered remains were
123 first taken to the National Museums of Kenya (NMK) in Nairobi, and then to the
124 Conservation and Restoration laboratories at CENIEH in Burgos, Spain. The low
125 density and extreme fragility of the bones necessitated a combination of mechanical
126 and digital cleaning (Supplementary Information A).

127 Careful examination revealed the articulated partial skeleton of an immature human
128 (Fig. 2). Sample OSL4, taken directly from the feature in 2013, was processed at Royal
129 Holloway, London, resulting in a **stratigraphically coherent age of $\sim 76.0 \pm 7.4$ ka**
130 **Incorporation of this OSL age into the Bayesian model yielded an estimated age of 78.3**

131 ± 4.1 ka **for the pit** (Extended Data Fig. 2, Supplementary Information C). Hand
132 excavation of the sediment block encasing the skeletal elements revealed the presence
133 of lithics and fauna (Supplementary Information B, Extended Data Fig. 1) consistent
134 with the surrounding layers. These layers (18 and 19) produced a large assemblage of
135 artefacts (n=2194) that have been shown to be both distinct from the LSA layers above
136 (16-1) (Shipton et al. 2018) and consistent with other MSA assemblages in East Africa
137 (Blinkhorn and Grove 2018; Grove and Blinkhorn 2020), thereby providing additional
138 and robust proof that the pit was associated with MSA occupations. Screening of the
139 sediment and skeletal elements for ancient DNA proved inconclusive (Supplementary
140 Information D).

141 As explained below, the field and laboratory evidence strongly indicates that
142 this is a primary and intentional burial, and that the child's body was deliberately placed
143 in an excavated pit that was posteriorly and rapidly backfilled with sediment.

144

145 **Primary and intentional deposit**

146 The skeletal remains consist of a considerable part of the basicranium, a
147 fragment of the left hemi-mandible with a complete ramus, five teeth (right M₁, right
148 M¹, left dm² and the *in situ* and un-erupted left M₁ and left M¹), the cervical and thoracic
149 spine with associated ribs, the right clavicle, and the left humerus (Fig. 2). Beyond these
150 better-preserved bones, there were several fragments corresponding to the cranial,
151 facial, pectoral, pelvic and limb areas, although anatomical identification is difficult
152 due to the severe diagenesis (bioerosion, recrystallisation) of the skeleton (Extended
153 Data Figs. 3-4, Extended Data Table 1, Supplementary Information E). Fragments of
154 the left radius and ulna (in anatomical connection through a lump of sediment), and a
155 deformed fragment of the left parietal, were also recovered from the burial fill, although

156 detached from the main block (Supplementary Information A). Several unidentifiable
157 small, white and flat bone fragments recovered in the field, prior to plastering the
158 feature, probably correspond to the crushed and severely distorted upper part of the
159 cranial vault. The advanced diagenesis of the bones prevented the preservation and/or
160 physical recovery of the remaining skeletal elements. Photographs taken after
161 micromorphology sampling in 2013 show the proximal portion of the right femora
162 inside the section (see Supplementary Information A Fig. 2) and photographs of the
163 plan view in 2017 show the proximal end of the left femur (Fig. 1 b) . Based on dental
164 development we estimate that the child, which we named 'Mtoto' ('child' in Swahili),
165 died at the age of 2.5-3.0 years (Supplementary Information F).

166 According to Duday (2006), the identification of a primary burial (the fresh
167 body was placed in a location where the entire process of decomposition took place)
168 is based on four criteria: i) the macroscopic anatomical integrity of the body, specially
169 of unstable articulations; ii) the minimal displacement of the bones that can be
170 explained within the course of decomposition; iii) abundance of terrestrial gastropods
171 that feed on earthworms in close proximity to the corpse and iv) geochemical and
172 histological analyses in favour of an ins situ-decomposition and putrefaction. All these
173 criteria are met with the PYS finding.

174 The majority of the bones appear in either strict articulation or in good
175 anatomical association and minor displacements can be explained as a consequence of
176 decomposition and subsequent formation of secondary spaces. The combination of the
177 photographic, microtomographic and surface scanner data, together with the total
178 station coordinates of the feature, confirm that the body was deposited in a flexed right
179 lateral decubitus position with the thighs flexed towards the torso at an angle less than
180 90° (Fig. 3, Extended Data Fig. 5). The vertebral column forms an arc stretching from

181 the cervical to the distal thoracic area, and this, together with the relative position of
182 the lower limbs, denotes a tightly flexed position of the body. The body is not lying
183 flat, but the spine it is at an angle of approximately 12° above the horizontal axis. The
184 thorax is laterally compressed (Fig.3, Extended Data Fig. 5). The ribs on the right side
185 are flattened and those on the left side are at a higher angulation. There is a gap between
186 the anterior ends of the right and the left ribs from the same vertebral level, consistent
187 with the interpretation that the child was lying on its right side. Although the
188 mechanical pressure of the sediment flattened the thorax, the rib cage did not collapse,
189 preserving the original spatial relationship and curvature of the ribs, pointing to
190 decomposition in a filled space (Supplementary Information A). Indeed, the
191 preservation of most of the thorax articulations and its volume indicates that the
192 destruction of the soft tissues and viscera did not produce a large, temporary empty
193 space. This phenomenon tends to occur in contexts characterised by particularly fluid
194 sediments that infiltrate by percolation and it is indirect but solid evidence of a deposit
195 made in bare earth¹⁰. The particle size analysis confirms that the sediment inside the
196 burial presents a higher proportion of both silt and sand and a lower proportion of clay
197 in comparison to Layers 17-19 (except for two samples from top of Layer 18; Fig. 4
198 and Supplementary Information G). This would favour a progressive infilling of the
199 internal space as the cadaver decomposed and reinforces the hypothesis of an in-situ
200 decomposition of the cadaver.

201 The right clavicle displays an oblique orientation, with a descent of the sternal
202 extremity of almost 90° (Supplementary Information A). Similarly, the first and second
203 right ribs are also distally displaced, and rotated medially about 90°, but they preserve
204 the intercostal space, arguing in favour of a minimal displacement of the sternal
205 articulation of the pectoral girdle. The depression of the clavicle and the adoption of an

206 oblique orientation are typical of tightly shrouded burial¹¹. This is consistent with the
207 upper part of the body being wrapped in a perishable cloth/material, or alternatively,
208 the body being densely packed within its pit structure. In either case, such a deliberate
209 treatment of the body would explain the exceptional in-place preservation of the scapula
210 and humerus of the over-hanging arm, and the intact articulation of the vertebral column
211 and ribs, which would otherwise likely collapse as the decomposition advances ¹⁰.

212 Rotation of the head is common in burials, as a result of the gravity force and the
213 weight of the cranium when the decay of the cranio-vertebral attachments places it in
214 an unstable position (Extended Data Fig. 5). For Mtoto, the cranium and first three
215 cervical vertebrae are disarticulated as a unit and partially dislocated from the column.
216 The movement of the head points to the existence of some empty space around it and
217 it contrasts with the progressive infilling and minimal displacement of the rest of the
218 body. In a fresh cadaver, this type of head dislocation involving the cervical vertebrae
219 couldn't indicate collapse due to the decay of a perishable support placed beneath the
220 head^{12,13}. Mtoto's head dislocation, together with the depression of the clavicle and the
221 first two ribs, would be compatible with the upper part of the body being wrapped and
222 the head supported with a perishable material. The differential preservation of the upper
223 versus the lower part of the body could be additional evidence for this protective
224 treatment. These evidences would be supportive of a more elaborated involvement of
225 the community in the funerary rite versus the structured abandonment (Pettitt, 2010) or
226 the happenstance burial (Deffleur, 1993).

227 Despite extreme bone fragility, the anatomical integrity and strict articulation of
228 some of the so-called unstable or labile articulations such as those between the
229 vertebrae, between the vertebrae and the ribs, and between the left scapulothoracic
230 articulation and the left temporo-mandibular joint indicate this is a primary and

231 undisturbed deposit^{10,11} and that the child was covered with sediment rapidly after its
232 placement. Additionally, taphonomical, histological and geochemical analyses support
233 the in-situ decomposition and putrefaction of 'Mtoto'. The anatomical alignment and
234 advanced diagenesis of Mtoto contrasts with the highly fragmented status and more
235 variable diagenetic conditions of non-human faunal remains recovered from Layers 17-
236 19 (Extended Data Table 2, Supplementary Information H). Taken together, all
237 available evidence argues in favour of rapid burial after death of the individual,
238 protecting the skeleton from intense post-depositional breakage as experienced by the
239 faunal remains in the surrounding layers.

240 Optical microscopy of bone fragments (probably upper limb) demonstrates that
241 human and non-human bone followed different taphonomic trajectories. Human bone
242 underwent predominantly non-Wedl bioerosion (attributed to bacteria^{17,18}) and
243 enlargement of osteocyte lacunae and canaliculi (probably due to fungal, or
244 fungal+bacterial action) (Extended Data Figs. 3-4). Although variable across the
245 sample, bioerosion of the larger human bone fragments was arrested (General Histology
246 Index: 2-3). Pervasive recrystallisation of the bone hydroxyapatite into a Ca-enriched,
247 amorphous or cryptocrystalline calcium phosphate - more advanced in endosteal
248 regions - may be associated with 0.1-1 μ m-scale bacterial microtunnelling¹⁹. Fe and Mn
249 oxide deposition - probably bacterially mediated - in bone (and adhering sediment)
250 likely resulted from episodes of waterlogging. The most parsimonious interpretation of
251 human bone diagenesis is that the child's body decomposed as a buried fleshed cadaver,
252 in an episodically waterlogged burial environment (Supplementary Information E).
253 Additionally, the skull of 'Mtoto' presents several star-shaped marks and bore holes
254 indicative of insect and gastropod's activity, compatible with with in-situ
255 decomposition.

256 Higher concentrations of MnO and CaO in the pit fill are consistent with *in situ*
257 decomposition of the body^{15,16}. Micromorphological and histological evidence suggests
258 that, while both Ca, Mn (and Fe) mobilisation were diachronous, calcite and Mn oxide
259 deposition in and around the bone may have been mediated to a significant extent by
260 putrefactive bacteria (Supplementary Information E and Extended Data Figs. 3-4,
261 Extended Data Table 1).

262 The sediment matrix is devoid of microcharcoal, ash or other (putative) microscopic
263 human inputs. Five land snail shell fragments (*Achatina* sp.) were found in close
264 association with the skeleton, around the occipital area. Although one of the shell
265 fragments bear lines incised by a point, not seen in fragments from the encasing layers
266 (Extended Data Fig. 6, Supplementary Information I), fragments of this genus are also
267 abundant in penecontemporaneous Layer 18 and bear traces of heating and
268 consumption. Thus, the evidence is not sufficient to suggest deliberate placement of
269 shell fragments in the pit. Nevertheless, the *Achatina* shells from the pit are
270 significantly larger in comparison to those from penecontemporaneous Layer 18,
271 indicating that they did not experience intensive breakage from processes such as
272 trampling (Extended Data Fig. 6, Supplementary Information I). Analysis of reddish
273 agglomerates spotted during the excavation of the child's remains showed that they
274 were not anthropogenic in origin and cannot therefore be interpreted as evidence that
275 the child's body or the wrapping that protected it was covered in ochre (Supplementary
276 Information G).

277

278 **Burial versus funerary caching**

279 Additional to the placement of the body, the recognition of a burial requires the
280 identification of purportedly excavated burial pit and the posterior intentional covering

281 of the corpse (Pettitt, 2010). The distinction of a new stratum is key to discern a burial
282 from the accommodation of a body in a natural place such as cave fissures or hollows,
283 also known as funerary caching (Gargett, 1999). The excavation of Trench 4 exposed
284 a distinct feature, a well demarcated pit with different colour and density matrix with
285 no comparable counterpart in the remainder of the sequence Supplementary
286 Information A Fig. 1 and 3) which can only have resulted from intentional digging into
287 layer 19. The sedimentological analyses provides evidence of a clear burial cutting. **The**
288 **burial fill is a ferruginous mix of silt and sand, compositionally similar to the top of**
289 **Layer 18 and the base of Layer 17, and different from Layer 19 in which the pit appears**
290 **to have been excavated (Fig. 4, Supplementary Information G). The fine –grained**
291 **texture of the intraskeletal matrix may be representative of the original composition of**
292 **the burying sediment, or it may have resulted from infiltration of the sediment between**
293 **the bones as the cadaver skeletonised. (Supplementary Information E and Extended**
294 **Data Figs. 3-4, Extended Data Table 1) and compatible with the evidence of a**
295 **progressive infilling in a deposit made in bare earth The lack of diagnostic flood**
296 **features of mass flow deposits in the burying sediment (Supplementary Information E)**
297 **makes unlikely that possibility of the sediment being washed into the pit during a flood**
298 **event shortly after the deposition of the corpse. Based on the bone microscopical study,**
299 **the most parsimonious explanation is to interpret that Mtoto’s body was deliberately**
300 **covered using as backfill sediment** scooped from the colluvial deposits that made up
301 the Layer 18 cave floor.

302 In sum, the interpretation for an intentional burial^{20,21} of Mtoto is based on: a) the
303 identification of a clear pit feature dugged into layer 19, b) the geochemical and
304 granulometric evidence discriminating the burial fill from the surrounding layers and
305 suggesting that sediment gradually filled empty spaces created by the body putrefaction

306 and insect activity, c) the overall completeness and anatomical integrity of the skeleton
307 and the alignment of the body in a tightly flexed position in the pit consistent with a
308 rapid covering after the body's deposition. The taphonomic reconstruction, the
309 sedimentary evidence, and the anatomical positioning are consistent with the burial of
310 a 'fresh' body and its subsequent decomposition in the pit.

311

312 **Taxonomic assessment**

313 To investigate Mtoto's taxonomic affiliation, the PYS teeth were compared against
314 a large sample of *H. neanderthalensis* and recent and fossil *H. sapiens*. The expression
315 of several crenulations and mesial accessory tubercles make the PYS dm² more
316 primitive than recent counterparts (Fig. 5, Extended Data Fig. 7, Supplementary
317 Information F). The morphology of the PYS M¹ samples fall within the range of
318 variation of *H. sapiens*, though it resembles the morphologically more complex Aterian
319 individuals in its pronounced Carabelli expression, the non-reduction and even in the
320 exceptionally large and subdivided hypocones²². In both the dm² and the M¹, the
321 occlusal polygon is more rhomboidal than in fossil and recent *H. sapiens*, but not as
322 skewed as in Neanderthals (Supplementary Information F). The size cusp sequence of
323 the M¹ falls between recent and Upper Palaeolithic *H. sapiens* on the one hand, and
324 Neanderthals and Qafzeh on the other. Although the morphology of both of the PYS
325 M₁ samples are compatible with that of *H. sapiens*, the profuse crenulation of their
326 enamel makes this morphology more complex than that of recent *H. sapiens* and some
327 fossil *H. sapiens* specimens such as those from Die Kelders in South Africa, Dolni
328 Vestonice and Pavlov in Europe, Qafzeh and Misliya in the Levant, and Daoxian and
329 Tubo in China^{23,24}. The size cusp sequences in the M₁s are intermediate between that
330 found in early *H. sapiens* specimens (Qafzeh H4, Les Rois, Abri Pataud, Dolni

331 Vestonice DV14) and in Neanderthals and recent *H. sapiens*²⁵. The geometric-
332 morphometric shape analysis of the occlusal outline of the EDJ reveals that all teeth
333 cluster with *H. sapiens* except for the M₁ that is closer to the Neanderthal distribution
334 in the inward location of the metconid tip and the more rounded lingual profile (Fig. 5).

335 The dimensions of Mtoto's teeth fall within the range of variation of recent *H.*
336 *sapiens* and are smaller than those of Neanderthals and some fossil *H. sapiens* such as
337 Qafzeh and Mesolithic teeth. The enamel of both deciduous and permanent molars of
338 PYS is thick, a primitive condition shared with *H. sapiens* and the majority of
339 specimens in the hominin fossil record except for Neanderthals (Supplementary
340 Information F).

341 Overall, Mtoto's dentition is consistent with an assignment to *H. sapiens*, although
342 it preserves some primitive features indicating that the child was less morphologically
343 derived than other broadly contemporaneous populations (Supplementary Information
344 F). The mandibular ramus shows a symmetric mandibular notch, where the condylar
345 and coronoid processes are levelled. This, together with the very arched temporal
346 squama, also align Mtoto with *H. sapiens*.

347

348 **Implications of the PYS burial for human evolution**

349

350 Despite Africa's alleged centrality for the emergence of 'modern human behaviour' in
351 the Pleistocene^{3,26}, evidence for mortuary practices in the continent is scarce and often
352 ambiguous. Intentional defleshing has been inferred from the orientation and location of
353 cut marks on the 600 kyr old Bodo skull, attributed to *Homo heidelbergensis/H.*
354 *rhodesiensis*. Defleshing and curation, supported by the presence of both diagnostically
355 located cutmarks and polishing are recorded on the skull of the juvenile individual from

356 Herto. Furthermore, evidence pointing to *funerary caching* - - the deposition of corps
357 in an existing natural feature– has been proposed for the Atapuerca-Sima de los Huesos
358 hominins and *H. naledi*. Until now, one of the earliest possible burials in Africa was
359 the infant hominin (BC3) found in 1941 at Border Cave, South Africa, thought to date
360 to $\sim 74 \pm 4$ ka²⁷ and apparently associated with a perforated and ochred *Conus* shell.
361 Although recent reappraisal of the evidence confirms the presence of a pit,
362 documentation on this burial is limited. No information is available on the degree of
363 articulation and the position of the remains^{28,29} within the pit and its age is inferred from
364 a stratigraphic correlation with a section, dated by ESR, located more than 10 m far
365 from the pit. The chronological and stratigraphic data for the BC3 specimen is coherent
366 overall with an age not younger than 58 ka²⁷, and possibly as old as 74 ka, but a more
367 constrained chronology for the skeleton is not available.

368 In contrast to the situation at Border Cave, the contextual, chronological and
369 taphonomic information at PYS are all supportive of a primary burial and meet the
370 criteria for simple, early inhumations of hominins in the Late Pleistocene²¹. On the basis
371 of multiple, and stratigraphically coherent, OSL dates, PYS represents the earliest
372 unequivocal evidence of an intentional burial in Africa at 78.3 ± 4.1 ka, clearly
373 demonstrating that complex treatment of the dead was practiced by *H. sapiens* by late
374 MIS 5. Moreover, the PYS burial reveals a clear and direct association between *H.*
375 *sapiens* and MSA technology, providing further confirmation of the link between this
376 technology and our early members of our species (Righter et al. 2017).

377 The PYS child in Kenya, in combination with the infant burial from Border
378 Cave and the funerary caching or interment of a juvenile at Taramsa in Egypt ~ 69 ka³⁰,
379 suggests that *H. sapiens* were intentionally preserving the corps of young members of
380 their groups at this time. Prior to ~ 74.6 ka, however, there are no unambiguous burials

381 of *H. sapiens* in Africa, despite the fact that earlier MSA populations demonstrate
382 sophisticated forms of symbolic expression³¹⁻³⁴. The absence of burials from the onset
383 of the MSA by 305 ka³⁵, and the rare occurrence of burials after 74 ka, may be a
384 consequence of cultural choice, such as the disposal of bodies away from residential
385 sites and in ways that are not consistent with their preservation. It may also indicate a
386 shift, sometime between 150 ka and 80 ka, from defleshing and curation, recorded at
387 Herto, to funerary caching and burials, observed at more recent sites from East Africa.

388 Interestingly, this provides a point of contrast with Neanderthals and early *H.*
389 *sapiens* outside of Africa, who buried their dead in residential sites by ~120 ka³⁶, about
390 45,000 years before PYS, suggesting that intentional burial may have emerged in
391 multiple hominin lineages. Infant and child burials are ubiquitous in Neanderthal and
392 early *H. sapiens* sites in the Levant and Europe, comprising 55-35% of all known
393 interments after 120 ka³⁶. Burial in residential localities, such as at PYS, has been
394 suggested to reflect mourning behaviour and the intention to keep the dead nearby³⁶.
395 Despite being the cradle of *H. sapiens*³⁷, Africa demonstrates a scarcity of mortuary
396 practices over most of the MSA that provides little current support for modern-like
397 conceptions of the afterlife and/or treatment of the dead. Nonetheless, cross-cultural
398 evidence in *H. sapiens* clearly emphasises that the absence of a behaviour does not
399 necessarily imply that capacity for such behaviour was lacking. Evidence for advanced
400 planning and symbolism from ~320 ka and particularly after 100 ka at East African
401 sites³⁸ suggests that modern cultural traits gradually permeated through those societies.
402 Limited evidence for mortuary behaviours in Africa may be due to practices leaving
403 elusive archaeological traces, taphonomy, or limited investigation. The highly fragile
404 and taphonomically-altered nature of the PYS skeletal remains highlight the
405 problematic conservation issues at even a well-preserved site. However, in the light of

406 the available evidence, the possibility of a non-African origin for the burial culture
407 cannot be discarded.

408 Beyond providing evidence for the earliest clear *H. sapiens* burial in Africa, the
409 ~74.6 ka skeletal remains are also of interest for the insight they provide into the
410 evolution of our species. While the mandibular and dental assessment of Mtoto are
411 consistent with its assignment to *H. sapiens*, the preservation of some primitive dental
412 features in comparison to other penecontemporaneous populations, suggests that our
413 species may have evolved in subdivided and regionally distinct populations and in a
414 variety of paleoecological settings (Supplementary Information J). Our study suggests
415 that the biological and socio-cultural evolution of *H. sapiens* was as a complex, likely
416 nonlinear and regionally diverse process.

417

418

419

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421

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519

520 **Online content**

521 Any methods, additional references, Nature Research reporting summaries, source
522 data, extended data, supplementary information, acknowledgements, peer review
523 information; details of author contributions and competing interests; and statements
524 of data and code availability are available at <https://doi.org/10.1038/s41586-020-2018-2>

525

526 **Methods**

527 **Dating**

528 **Optically stimulated luminescence dating.** A suite of seven single-grain optically
529 stimulated luminescence (OSL) ages for PYS were published⁶, including a detailed
530 description of the luminescence measurement and data analysis techniques used (their
531 Supplemental Note 3). In the present study, we report an additional OSL age, produced
532 using the method previously described⁶ for a sample from the burial infill (sample
533 PYS13-OSL4). Data used to calculate the age for OSL4 are presented in bold in
534 Supplementary Information C Tables 1-3, alongside data from the main publication⁶
535 for comparison. Since all quantities reported here were measured using the same
536 instruments and methods as those reported previously⁶, it is reasonable to treat them as
537 internally consistent and therefore comparable. Data for OSL4 are consistent with those
538 for previously reported PYS samples, particularly those for OSL3 and OSL5, which
539 yield contemporaneous ages. Particularly noteworthy is the similarity between the ages
540 for OSL4 (burial infill) and OSL3 and OSL5. If the burial pit had been dug into much
541 older sediments, and then infilled using those sediments, some grains would probably
542 have had their luminescence signal reset during the process. This might be expected to
543 yield a younger age for the burial infill than the surrounding sediments, which is not
544 the case. Similarly, the overdispersion for OSL4 is consistent with that for OSL5, where
545 no evidence for post-depositional disturbance has been found. The similarity in age and
546 overdispersion between samples OSL3-5 does not unequivocally demonstrate that the
547 burial infill is contemporaneous with Layers 17 and 18, since if the pit was backfilled
548 with spoil this might have occurred under subdued light conditions leading to poor
549 luminescence signal resetting. Also, given the ~7% uncertainties on individual ages,

550 archaeologically important differences in age could go unnoticed. Nonetheless, the
551 absence of a discrepancy between the ages and overdispersion of the burial infill and
552 surrounding sediments suggests that the burial is contemporaneous with Layers 17/18,
553 at least at the temporal resolution achievable using luminescence techniques.

554 **Bayesian Modelling and age of the PYS burial.** Bayesian modelling enables the
555 relative stratigraphic information recorded during excavation to be formally
556 incorporated into posterior age estimates deriving from chronometric data expressed as
557 probability distributions (in this case calibrated radiocarbon and OSL probability
558 distributions) and ‘prior’ information, i.e. observations on the data we collect. In
559 archaeology, this is often the stratigraphic and other relative information. Hence
560 ‘posterior’ data, is a probability function that reflects the level of confidence associated
561 with the values of the unknown parameters, in this case chronometric measurements,
562 after the observation of the prior information. Details about the data and structure of
563 comprehensive model for the dating of the PYS sequence was described previously⁶.

564 Here we update the model by adding the new OSL age obtain for the burial infill
565 (OSL4) and re-running the new model. We use OxCal 4.3.2⁴¹, and the code is shown
566 in Supplementary Information C. By inserting a “Date” command in the OxCal
567 structure we allow the model to find the best fit for the age of the burial based on all
568 OSL ages obtained for relevant Layers 17 and 18 (Supplementary Information C Table
569 4). Because of uncertainties in the bleaching history of sediment grains deriving from
570 disturbed contexts, such as a burial, we do not tie the “Date” command to OSL4; instead
571 we treat the burial infill sediment as another independent age for Layer 18.

572 Given that Layers 17 and 18 are statistically indistinguishable from each other and from
573 OSL4, we may assume that both layers are quasi-contemporaneous, at least in the

574 precision offered by the luminescence methods, for the studied period. The new model
575 is shown in Extended Data Figure 1.

576 **Screening for Ancient DNA**

577 **DNA extraction and library preparation.** We screened for ancient DNA 8
578 undiagnosed skeletal fragments and 12 sediment samples associated with the PYS
579 burial pit (Supplementary Information D Table 1). DNA was extracted from ~50mg of
580 bone or sediment using a silica-based method developed for the retrieval of short DNA
581 molecules⁴² on an automated liquid handling platform⁴³. 15% of each extract were
582 converted into a single-stranded DNA library⁴⁴, and barcoded with a pair of unique
583 indices⁴⁵, following the modifications described in Korlević et al. (2015)⁴⁶. The number
584 of DNA molecules incorporated into each library was assessed by quantitative PCR as
585 described elsewhere⁴⁷. Extraction and library negative controls were carried through all
586 steps of the experiments. Libraries were pooled and shotgun sequenced on a HiSeq
587 platform (Illumina).

588

589 **Mitochondrial capture and sequencing.** 1 µg of each amplified DNA library was
590 enriched whether for mammalian mitochondrial DNA (mtDNA) using a probe set of
591 242 taxa⁴⁸ or for human mtDNA with a probe set covering the full human mitochondrial
592 genome^{49,50}. The enriched libraries were pooled in two sets according to the capture
593 probe used and sequenced on a MiSeq platform (Illumina).

594

595 **Sequence processing and mapping.** The reads obtained from the sequencing of the
596 mtDNA enriched libraries were trimmed to remove adapter sequences and overlapping
597 paired-end reads were merged using leeHom⁵¹. Sequences from the mammalian

598 mtDNA enriched libraries were aligned to a non-redundant database of 796 mammalian
599 mitochondrial genomes from the NCBI Reference Sequence database (RefSeq) using
600 nucleotide BLAST (BlastN)⁵² with default parameters. Sequences were then assigned
601 to different taxa using MEGAN⁵³ and each ancient taxon is deemed present in the
602 dataset according to the requirement described earlier⁵⁴. The sequences from the human
603 mtDNA enriched libraries were aligned to the revised Cambridge Reference
604 mitochondrial genome (rCRS, NC_0120920) using the Burrows-Wheeler Aligner
605 (BWA)⁵⁵ with optimized parameters for ancient DNA «-n 0.01 -o 2 -l 16500»⁵⁶.
606 Aligned sequences shorter than 35 bases with a mapping quality lower than 25 were
607 filtered out. PCR duplicates were removed by merging sequences with identical
608 alignment start and end coordinates using bam-rmdup
609 (<https://github.com/mpieva/biohazard-tools>).

610

611 **Virtual reconstruction**

612 PYS was recovered encased in a block and had to be manually and mechanically
613 prepared for study. Given the delicate preservation and the infantile age stage of the
614 specimen, a microCT scan was performed to extract digital 3D models of the teeth and
615 bones, which were still embedded within the sediment. Because of the delicate state of
616 preservation and the need of preserving the information regarding the position of the
617 body a combination of mechanical and virtual isolation of the skeleton was decided.
618 However, the low density of the bones prevented a proper virtual isolation of the
619 elements, and it was necessary to carefully isolate the cranial from the postcranial
620 elements (Supplementary Information A) and to combine microCT and surface
621 scanners to reconstruct the original position of the child inside the block. The microCT
622 scans were performed at the Laboratory of Microscopy of the Centro Nacional de

623 Investigacion sobre la Evolucion Humana–Unique Scientific & Technical
624 Infrastructures (CENIEH–ICTS, Burgos, Spain) with a Phoenix v|tome|x s (GE
625 Measurement & Control). First, the entire block was scanned at 140 kV and 400 μ A,
626 with a 0.2 mm Cu filter and an integration time of 333 ms, resulting in an isometric
627 voxel size of 0.1227 mm. As the remains were carefully manually cleaned, we
628 performed a microCT of the block containing the cranial remains at 200 Kv and 400
629 μ A, with the same filter and integration time, resulting in an isometric voxel size of
630 0.0769 mm, and another scan of the block containing the postcranial elements at 140kV
631 and 250 μ A, with the same filter and integration time, resulting in an isometric voxel
632 size of 0.0479 mm. Each isolated tooth as well each smaller block resulting from the
633 mechanical excavation were also microCT scanned. Additionally, surface scanners
634 with a hand scanner (Artec Space Spider) under conditions of artificial light were
635 performed of each excavated element and processed with Artec Studio 11 software to
636 create surface models. Digital cleaning and segmentation was performed with Mimics
637 18.0 (Materialise, Be) and Avizo 7.0 (Visualization Sciences Group, 2012) using a
638 combination of automatic and manual segmentation, generating STL files of all PYS
639 remains. Digital renderings of the specimen in several views were done with Avizo,
640 Mimics and MeshLab, respectively. With the digital renders, it was possible to identify
641 several anatomical elements, such as the first vertebrae in anatomical connection. The
642 identification of two teeth and several shells in the first microCT of the main block was
643 useful as reference landmarks to later orient the microCT of the skull and the CT and
644 surface scanner of the thorax in their original position. Furthermore, and to understand
645 the position of the child within the block, the skull of a *Homo sapiens* child in a similar
646 stage of development was microCT scanned and virtually reconstructed. Using Mimics
647 and MeshLab software, both PYS and *Homo sapiens* crania were superimposed with

648 the registration and align tool. A best-fit was made to reference them in the best position
649 and see which parts were deformed or displaced. Furthermore, both crania were placed
650 again in the burial pit using as a reference the 2 teeth identified in the first microCT
651 scan of the main block. In addition, a comparative human skeleton of *H. sapiens* of
652 similar age was scanned using the same surface scanner to obtain the digital models of
653 all the long bones, hips and ribs, to help reconstruct the original position of the child
654 inside the burial. Using Mimics software, all the models were scaled to obtain a length
655 similar to that of a 2.5-3 year old (following Scheuer and Black, 2000⁵⁷). To align the
656 comparative skeleton with the PYS remains, the STL models of the vertebral column,
657 ribs, clavicle, scapula and humerus were aligned using both the CT of the entire block
658 and the CT of the PYS vertebral column.

659 Once all the remains and comparative models were aligned, 3ds Max 2020 (Autodesk)
660 was employed combining the topographic information available from the field,
661 photographs of the excavation plan, all the STL and OBJ models of both Mtoto and the
662 comparative skeleton to produce the closest reconstruction of the original position of
663 the child when it was found at the site. Each anatomical element was reoriented
664 following the topographic field information and the orthophotos of the site. When all
665 the remains were located in space, we proceeded to create textures, shadows and lights,
666 to visualize the preserved skeletal elements and to integrate them in their original
667 position in the comparative skeleton. Some renderings have been made with the semi-
668 transparent comparative skeleton, to understand and visualize the position of the PYS
669 into the burial, and understand its anatomical connection.

670

671 **Thin section micromorphology and bone microscopy**

672 A micromorphology sample collected at the end of the 2013 field season (PYS
673 2013_M1:19/18) intercepted soft, degraded bone. This sample was not processed for
674 micromorphological analysis: it was stored until 2019, when it became apparent that it
675 may have contained parts of Mtoto's skeleton (probably fragments of lower limbs and
676 ribs). The box was then excavated at CENIEH (Burgos). Centimetre-sized bone
677 fragments and lumps of the sediment matrix were collected for microscopic
678 investigation, the results of which are reported here. The 2013 micromorphology
679 sample was taken in a 6x10cm polyurethane box. Prior to its excavation at CENIEH, it
680 had been stored at room temperature for 5.5 years. Human bone fragments and sediment
681 lumps between the bones were collected on excavation of the sample box in 2019. Bone
682 and sediment were air dried and impregnated with polyester resin under desiccation
683 vacuum. One uncovered petrographic thin section (30µm thick) was produced from the
684 impregnated block. The section was examined under a polarizing microscope (x10 to
685 x400) at plane polarized (PPL), cross-polarized (XPL) and oblique incident light (OIL).
686 Description of sediment features follows Bullock et al. (1985)⁵⁸, Stoops (2003)⁵⁹ and
687 Stoops et al. (2018)⁶⁰. Estimates and measurements of sediment inclusions and
688 histological attributes of the bone fragments were made with the aid of standard semi-
689 quantitative estimation charts and the *analySIS pro5* image analysis software. The
690 degree of diagenetic alteration of bone was estimated mainly through optical
691 microscopy, using the General Histology Index (GHI: Hollund et al. 2012⁶¹) – an
692 estimate of microstructural alteration similar to the more commonly used Oxford
693 Histology Index (OHI: Hedges et al. 1995⁶²; Millard 2001⁶³), but taking into account
694 other types of structural and compositional alteration (generalised destruction, staining,
695 accumulation of authigenic deposits, fissuring) besides bioerosion. Where identifiable,
696 microscopic bioerosion (“*microscopical focal destruction*” – MFD: Hackett (1981)⁶⁴

697 was recorded following the typology proposed by Hackett (1981)⁶⁴ and Jans (2008)¹⁷.
698 SEM-EDS analysis (on a Zeiss EVO-MA15, with an Oxford Instruments InCA Max 80
699 mm EDS) was carried out on the polished, uncoated thin section, to observe bone
700 microstructure at higher magnifications and determine elemental composition of
701 selected features. Low-vacuum conditions were used (60 Pa) to prevent charging of the
702 sample surface; strict operating conditions of 50 μ A filament current, 2.525 A gun
703 current, 20 kV accelerating voltage, and an 8.5 mm working distance to achieve an
704 acquisition rate of 15 kcps, were applied to standardise the analyses. A polished Co
705 standard was analysed to adjust for beam current drift, and a polished dolomite standard
706 was used to confirm the accuracy of the calculated absolute element concentrations.
707 Navigation on the sample was aided by section scans. Data are reported as non-
708 normalised percentage weights.

709

710 **Dental analysis**

711 **Comparative metrical and morphological study of teeth.** The evaluation of
712 morphological features was made on the original fossils and the virtual images obtained
713 by microtomography. The descriptive terminology used in this report derives from the
714 following sources: Carlsen 1987⁶⁵, Tobias 1991⁶⁶, Turner et al. 1991⁶⁷, Scott and Turner
715 1997⁶⁸, Martín-Torres et al. 2007⁶⁹, Martín-Torres et al. 2008⁷⁰, Martínez de
716 Pinillos et al. 2017⁷¹, and Table 4 in Martín-Torres et al. 2012⁷², which includes a
717 modified version of the Arizona State University Dental Anthropological System
718 (ASUDAS) of scoring. Occlusal wear was recorded following Molnar 1971⁷³. The
719 comparison was focused in the Late Pleistocene samples, *H. sapiens* and *H.*
720 *neanderthalensis*, although several samples from the Middle to late Pleistocene Africa
721 were included in order to assess the variability of the African fossil record. The

722 mesiodistal (MD) and buccolingual (BL) dimensions of the PYS child were measured
723 by JMBC to the nearest 0.1 mm, following the methods of Flechier, Lefèvre, and
724 Verdéne (1973)⁷⁴. Apart from the MD and the BL, we have also calculated the
725 computed crown area (CCA: MD X BL) and the measured crown area (CI: [BL/MD] x
726 100). In addition, the cusp areas of the M1s were measured following Bermúdez de
727 Castro et al. (2001)⁷⁵ and using the criteria outlined by Bailey (2004)⁷⁶. The areas were
728 measured three times and the average of the three values was used. The total crown
729 base area (TCBA) was calculated as the sum of all the individual cusp areas. The PYS
730 values were compared against a large hominin sample of *H. sapiens*, *H.*
731 *neanderthalensis* and some relevant Middle to Late Pleistocene fossils from Africa.

732 **Enamel Thickness.** Virtual sectioning of the molars was performed following the
733 protocol described in Olejniczak and colleagues (2008)⁷⁷. The mCT image stack was
734 imported into Amira (6.3.0, FEI Inc.) and rotated into anatomical position. Then, the
735 tip of three dentine horns (protoconid, metaconid and hypoconid in the mandibular
736 molars and protocone, paracone and hypocone in the maxillary molars) were identified
737 and the image stack was adjusted to intersect these three points of interest. A new plane
738 perpendicular to the plane containing the three dentine horns was rotated to pass
739 through the mesial dentine horns (protoconid and metaconid in the mandibular molars
740 and protocone and paracone in the maxillary molars). We assessed enamel thickness
741 from virtual 2D mesial cross-section planes in each PYS molar as described in Martin
742 (1985)⁷⁸ using Amira (6.2, FEI Inc.) and ImageJ (1.51, NIH). In each mesial plane, we
743 measured the enamel (c) and dentine cap (b, including the pulp) areas (in mm²), adding
744 up into the total crown area (a, in mm²), and the enamel-dentine junction (EDJ) length
745 (d, in mm). We calculated the average enamel thickness (AET=c/d), the relative enamel
746 thickness (RET= 100*AET/(b^{1/2})) and the percentage of dentine and pulp in the molar

747 crown ($b/a=100*b/a$ in %). We assessed volume enamel thickness of the molar caps in
748 the complete sample from PYS. Using Amira (6.3.0, FEI Inc.) we performed the
749 segmentation of the dental tissues (enamel, dentine and pulp). We used the
750 semiautomatic tool, threshold-based segmentation, and manual corrections. We
751 employed the protocol of Olejniczak et al. (2008)⁷⁷ for the definition of the cervical
752 plane. That is, the plane halfway between the most apical continuous ring of enamel
753 and the plane containing the last hint of enamel. The following variables were measured
754 and/or calculated: volume of the enamel (V_e in mm^3); volume of the coronal dentine
755 including the pulp enclosed in the crown (V_{cdp} in mm^3); total volume of the crown,
756 including the enamel, dentine and pulp (V_c in mm^3); surface of the EDJ (SEDJ in
757 mm^2); percentage of dentine and pulp in the total crown volume (V_{cdp}/V_c
758 $=100*V_{cdp}/V_c$ in %); 3D average enamel thickness (3D AET= $V_e/SEDJ$ in mm) and,
759 3D relative enamel thickness (3D RET= $100*3D AET/(V_{cdp}^{1/3})$ a scale-free
760 measurement)^{77,79}. In order to extract the largest amount of information of the PYS
761 specimens and the comparative sample, including the occlusal worn molars, we
762 assessed lateral (non-occlusal) enamel thickness in the complete sample. In Amira
763 (6.3.0, FEI Inc.) we defined the occlusal basin plane, a plane parallel to the cervical
764 plane and tangent to the lowest enamel point of the occlusal basin. All material above
765 the occlusal basin plane was removed and only the enamel, dentine and pulp between
766 these two planes were measured^{80,81}. The following variables were measured and/or
767 calculated: lateral volume of the enamel (L V_e in mm^3); lateral volume of the coronal
768 dentine including the pulp enclosed in the crown (LV $_{cdp}$ in mm^3); total lateral volume
769 of the crown, including the lateral enamel, dentine and pulp (LV $_c$ in mm^3); lateral
770 surface of the EDJ (LSEDJ in mm^2); percentage of dentine and pulp in the lateral crown
771 volume (LV $_{cdp}/LV_c=100*LV_{cdp}/LV_c$ in %); 3D average enamel thickness (3D

772 LAET=LVe/LSEDJ in mm) and, 3D lateral relative enamel thickness (3D
773 LRET=100*3D LAET/(LVcdp1/3) a scale-free measurement)⁸². The results of the 2D
774 and 3D measurements in PYS specimens were compared with two populations,
775 Neanderthals and modern humans (MH). Adjusted Z-scores^{83,84} of the three variables
776 accounting for tissue proportions (percentage of dentine, AET and RET) were
777 computed to compare 2D and 3D values of the PYS specimens to the means and
778 standard deviations of the Neanderthal and MH groups. This statistical method allows
779 the comparison of unbalanced samples by using Student's inverse t distribution. In
780 these Z-scores the -1.0 to +1.0 interval comprises the 95% of the variation in the
781 reference sample. In addition, standard box and whisker plots were computed to
782 represent three set of variables of crown volume and lateral volume (including 3D
783 Vcdp/Vc, 3D AET, 3D RET and 3D LVcdp/Vc, 3D LAET, 3D LRET) in the PYS
784 sample and the complete comparative specimens and/or groups.

785 **Tissue distribution (cartographic maps).** In order to visualize enamel thickness
786 topographic distribution in PYS specimens, 3D chromatic maps were generated in
787 Amira (6.3.0, FEI Inc.). The defined chromatic scale is from thinnest (blue) to thickest
788 (red)^{85,86}. For comparative purposes, we generated the chromatic maps of a selected
789 sample of specimens, including: Neanderthals from Roc de Marsal (lower and upper
790 deciduous m2), La Quina (upper permanent M1) and Abri Suard (lower permanent
791 M1). Fossil *H. sapiens* from La Madaleine (lower deciduous m2), Qafzeh (upper
792 deciduous m2 and upper and lower permanent M1). And, modern humans from
793 European origin (upper and lower deciduous m2, and upper and lower permanent M1).

794 **Geometric morphometric of the EDJ.** We performed the Geometric morphometric
795 (GM) analysis of the EDJ morphology on the virtual surfaces of PYS specimens and a
796 comparative sample that included Neanderthals, fossil *H. sapiens* and modern humans

797 (Supplementary Information F Table 4). We reconstructed the slightly worn dentine
798 horns of the comparative sample in Geomagic Studio (version 2012;
799 www.geomagic.com) with the fill-holes tool. When necessary we mirrored the
800 comparative specimens according to the PYS molar type. Using landmark tool in
801 Amira, we placed one landmark on the dentine horn tip of each main cusp (protocone,
802 paracone, metacone and hypocone on maxillary molars, and protoconid, metaconid,
803 hypoconid, entoconid and hypocunulid on mandibular molars). Following, we placed a
804 set of semilandmarks (96 and 95 points in maxillary and mandibular molars
805 respectively) along the marginal crests. In each molar, configuration segments were
806 saved independently. Using the package R, we generated the document containing the
807 coordinate configuration for each molar type. Following, using R package we
808 performed the weighted between-group principal component analysis (bgPCA) based
809 on the Procrustes and deformation-based shape residuals⁸⁷. Finally, we tested for
810 allometry on the landmark-based analyses using the coefficient of determination (R²)
811 of a multiple regression⁸⁸, in which the explicative variable is the centroid size and the
812 dependent variables are the bgPC scores⁸⁹.

813

814 **Faunal analysis**

815 **Zooarchaeology**

816 Two separate zooarchaeological studies were conducted: N.A. analysed remains from
817 burial fill at the Max Planck Institute for the Science of Human History in Jena, and
818 M.E.P. analysed remains from Trenches 3-4 at the National Museums of Kenya in
819 Nairobi. Access to comparative skeletal material for eastern Africa was distinct in each
820 study, and analytical protocols differed somewhat.

821 **Fauna, burial fill.** For the burial fill (context 809 in Trench 8), all bone fragments were
822 sorted, counted and measured (length, width and thickness) using a digital caliper
823 (Mitutoyo 500–463), regardless of identifiability. The specimens were recorded in
824 detail using codes following a zonation system that allow for the description of
825 fragmentation patterns. All identified fragments were examined for bone surface
826 modifications under a Nikon C-PCN stereomicroscope.

827 **Fauna, Trenches 3-4.** For Trenches 3-4, faunal remains are reported from the
828 following layers and contexts: Layer 17 (contexts 316X, 317Y-Z, 420B-F), Layer 18
829 (422A-C), Layer 19 (423A-D). These remains were weighed and sorted into those
830 identifiable at least to a minimal level (e.g., mammal limb bone), and those not easily
831 identifiable at any level. For each identified specimen, recorded variables include taxon,
832 element, portion, side, weathering, breakage, and bone surface modifications. Cortical
833 surfaces of all identified specimens were examined with a 20x hand lens under strong
834 oblique light.

835

836 **Sediment analysis**

837 Two sets of samples were analysed in the framework of this study. The first set
838 comprises sediment samples from archaeological layers and the burial pit. The second
839 set includes reddish micro-agglomerates associated with the child remains.

840 *Sediment analysis from archaeological layers and the burial pit.* Two groups of
841 sediment samples were analysed. The first group comprises twenty-four samples of
842 sediment collected during the excavation of Trenches 3 and 8 in 2017 (PYS-2017-
843 200127 to PYS-2017-200153). They come from layers 17, 18 and 19, and were
844 collected at 2 cm depth interval. They do not include sediment from the burial
845 (Supplementary Information G Table 1).

846 The second group comprises five sediment samples collected close to the human
847 remains (Supplementary Information G Table 2). Three samples, reddish-brown in
848 colour, were retrieved close to the maxilla, the face, and inside the cranial vault. The
849 other two, brownish in colour, were collected close to the occipital bone and the
850 postcranial perimeter. These five samples were retrieved during the excavation of the
851 skeleton at the CENIEH's Conservation and Restoration Laboratory.

852 The sediments samples were studied at the UMR 5199 PACEA and UMR 5060
853 IRAMAT/CRP2A laboratories in Bordeaux, France. The samples were examined and
854 photographed with a motorised Leica Z6 APOA microscope equipped with a DFC420
855 digital camera. Uploaded images were treated with Leica Application Suite (LAS)
856 equipped with the Multifocus module. Samples were then prepared for grain-size
857 analysis with a Horiba LA-950 laser particle size analyser. The sample pre-treatment
858 included suspension in sodium hexametaphosphate (5 g/l) and hydrogen peroxide
859 (35%) at room temperature for 12 h. The resulting compound was subjected to 60 s
860 ultrasonification to achieve optimal dispersion. The Mie solution to Maxwell's equation
861 provided the basis for calculating the particle size^{90,91}, using a refractive index of 1.333
862 for water and 1.55 – 0.01i for the particles. The grain-size distributions expressed in ϕ
863 units were decomposed in different Gaussian populations (parametric curve fitting
864 method) using the mixdist R package⁹² to identify the main modes and their relative
865 proportions. The limits used for grain-size classes are as follows: <7 μm (clays), 7–
866 63 μm (silts), 63-2000 μm (sands). They are based on studies showing that the amount
867 of clay particles measured by laser diffraction is usually underestimated (e.g. Konert
868 and Vandenberghe, 1997⁹³).

869 Elemental analysis was carried out by using a hand-held SPECTRO xSORT energy
870 dispersive X-ray fluorescence (EDXRF) spectrometer from Ametek. This instrument is

871 equipped with a silicon drift detector (SDD), a low power W X-ray tube with an
872 excitation source of 40 kV, and an X-ray beam of 8 mm. Spectra acquisition times were
873 set to 300 s. Measurements were performed with a constant working distance by using
874 a positioning device consisting of a lead receptacle to which the spectrometer was fixed.
875 Samples were previously ground and homogenized with an agate mortar and then
876 placed into a plastic cup covered with a Prolene® thin film. Three measurements were
877 taken on each sample. Element contents were calculated as the average of these
878 acquisitions. In order to precisely quantify the elemental composition of the samples a
879 dedicated calibration for sediment samples was applied (see Sitzia et al., 2019⁹⁴ for
880 details). This calibration allows quantifying Si, K, Ca, Ti, V, Cr, Mn, Fe, Ni, Zn, Ga,
881 As, Rb, Sr, Y, Zr, and Ba. Centred log ratio (*clr*) was used to analyse geochemical data
882 as an alternative to the raw percentages, following Aitchison (1986)⁹⁵. Prior to
883 statistical analysis, a non-parametric replacement was made for the null values below
884 the limit of detection, according to the method proposed by Martín-Fernández et al.
885 (2003)⁹⁶. Some elements were excluded when they displayed more than one third of
886 missing values (Cr, Ni, Sr) or showed a heterogeneous distribution (V). We performed
887 principal component analysis (PCA) of EDXRF concentrations for the thirteen major,
888 minor, and trace elements more frequently detected (Si, K, Ca, Ti, Mn, Fe, Zn, Ga, As,
889 Rb, Y, Zr, Ba). All data analyses were done with the *ade4* package⁹⁷ for R software.

890 We performed XRD analyses on powder samples by using a Bruker D8 Advance
891 diffractometer (Bragg-Brentano Theta-Theta geometry, fixed sample in the horizontal
892 plane, movable tube and detector), goniometer diameter 600mm, Cu anti-cathode X
893 source, Cu-K α incident X incident doublet radiation. Acquisitions were conducted with
894 an angular range of 3-60° at 2 Θ (reticular distances 1.54-29 Å), measurement step
895 0.01°, a 181px linear detector type Bruker LynxEye covering simultaneously 2.6 ° in

896 2θ , an analysis time per point of 550 s (approx. 9 min/px), an angular reproducibility
897 of 0.017° , supporting samples holders of 25 mm in diameter and 1.5 mm thickness,
898 made of PolyMethyl MethAcrylate (PMMA) holders. Samples were previously ground
899 and homogenized with an agate mortar. The time of analysis was 5 h. Mineralogical
900 phases and the semi-quantitative analysis were achieved by using the routine
901 DIFFRAC.SUITE™ EVA software package (Bruker AXS GmbH, Germany) which
902 allows mathematical deletion of bottom noise and Cu-K α 2 component, combined with
903 the specific powder diffraction file (PDF2) database (International Centre for
904 Diffraction Data—ICDD, Pennsylvania, USA).

905 *Reddish micro-agglomerates associated with the child remains*

906 The reddish micro-agglomerates, possibly fragments of iron-rich rocks commonly
907 called ochre, were detected in the sediment coating the child remains during their
908 excavation and consolidation at the CENIEH. An analytical pipeline involving
909 observation and sampling under low-magnification optical microscope, Raman
910 spectroscopy, X-ray Diffraction (XRD) and Scanning Electron Microscopy coupled
911 with Energy Dispersive X-Ray Spectroscopy (SEM-EDS), was applied to establish the
912 elemental and mineralogical composition of these micro-agglomerates. The aim was to
913 investigate whether ochre may have been involved in the mortuary practices that have
914 led to the deposition and preservation of the child remains. All analyses were conducted
915 at the CENIEH Archaeometry and Microscopy laboratories.

916 Two groups of reddish micro-agglomerates were analysed. The first set consists of 55
917 micro-agglomerates contained in samples of reddish-brown clay sediment collected
918 close to the maxillary, the face, and inside the cranial vault. The micro-agglomerates
919 typically correspond to $\sim 900 \times 600 \mu\text{m}$ associations of grains featuring distinct reddish
920 shades (Supplementary Information G Figure 1). The second group is composed of two

921 larger ~2-10 mm– red fragments collected on the ribs and vertebral bodies
922 (Supplementary Information G Figure 2). Fifteen agglomerates from the first group and
923 the two composing the second group were submitted to analytical microscopy (Raman
924 spectroscopy, SEM-EDS) and XRD analysis (Supplementary Information G Table 3).

925 A FEI Quanta 600 SEM fitted with EDS, and Oxford Instruments INCA software to
926 interface with the EDS, was used to obtain an estimation of the bulk elemental
927 composition of the agglomerates. Operating conditions were 15kV of acceleration
928 voltage, working distance of 10 mm, and acquisition time of 90 seconds, resulting in
929 typical dead times between 10 and 17 %. Unless otherwise specified, results are given
930 as compound weight %, oxygen calculated by stoichiometry and normalised to 100%.

931 All SEM micrographs are Low Vacuum Secondary Electron (SE) images.

932 The samples were first observed at low magnification – typically ranging 100-200x –
933 to capture the entire particle (Supplementary Information G Figure 3). Three
934 representative locations were then selected to investigate the samples microstructure at
935 1200x where surface features and mineral phases were measured. Three areas were
936 analysed at each location to estimate the average bulk chemical composition. Finally, a
937 mapping of the matrix between crystals was conducted at 2400x and compared to the
938 bulk composition obtained at 1200x (Supplementary Information G Figure 4).
939 Occasionally, other magnifications were used to better investigate texture, grain
940 morphology and their relation with elemental composition. Analyses at high
941 magnifications were conducted to avoid potential biases due to surface irregularities
942 and voids. The analytical procedure was adapted from the analysis of similar mineral
943 particles described in Pitarch Martí et al. (2017)⁹⁸.

944 The mineralogical composition of the sample was established by XRD. Due to the
945 destructive nature of this technique, these analyses were not performed on agglomerates

946 analysed by SEM-EDS and RAMAN but on residues coming from the same samples.
947 Sediment was hand grounded in an agate mortar with a pestle. The resulting powder
948 was tapped on silicon discs and mounted into stainless steel sample holders. Patterns
949 were collected with a PANalytical X'Pert PRO MPD diffractometer with CuK α
950 radiation ($\lambda=1.5406 \text{ \AA}$) and a solid-state multichannel detector. Each sample was
951 scanned over the 2-theta range 5- 80° with a step size of 0.03. The working tension and
952 intensity were 45kV and 40mA respectively, and the time of analysis ranged from 3 to
953 16 h. Semi-quantitative analysis was performed on the XRD data using the X'Pert High
954 Score software. Due to the small amount of material analysed the results should be
955 regarded as qualitative.

956 Raman spectra were acquired using a DRX Thermo Scientific Raman dispersive
957 spectrometer, with a laser emitting at 780 nm. The power radiation measured under the
958 Olympus x50 microscope objective was about 0.5 mW-0.8 mW. Acquisitions of about
959 25 seconds and multiple additions were used. The spectrometer worked in a spectral
960 range from 55 to 3350 cm⁻¹. The calibration of the spectrometer was done with a
961 polystyrene standard (main band: 1000 cm⁻¹). The recorded mineral spectra were
962 contrasted against the RRUFF database library for phase identification⁹⁹ as well as
963 Julien et al. (2004)¹⁰⁰, Bellot-Gurlet et al. (2009)¹⁰¹, Hanesch (2009)¹⁰² and Babay et al.
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1146 E.S., L.M.F analysed the hominin fossil; P.F. conducted the mechanical restoration and
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1152 **Competing interests** The authors declare no competing interests.

1153 **Additional information**

1154 **Supplementary Information** is available in the online version:

1155 **Correspondence and requests for materials** should be addressed to M.M.-T., N.B.
1156 or M.D.P.

1157 **Extended Data Fig. 1. PYS MSA lithics.** Above: The size distribution of flakes
1158 through Layers 18-17 (MSA) and Layer 16 (early LSA). Notice the decrease in size
1159 across Layers 17-16. The small sample of flakes recovered from the burial (n=14),
1160 shown here with a boxplot, falls within the variation in product weight for the MSA.
1161 Below: Facetted limestone flakes from MSA layers of PYS. A: limestone flake with
1162 facetted dihedral platform from burial context (809). B: Retouched limestone flake
1163 with large facetted platform from Layer 17.

1164

1165 **Extended Data Fig. 2. Bayesian model for the age estimation of PYS.** Left: Bayesian
1166 model of all available age determinations from PYS, produced using OxCal v4.4.2 and

1167 IntCal20. Right: Age estimate of the burial determined using Bayesian model. A full
1168 description (OxCal code) of the age model is provided in Supplemental Information C.
1169

1170 **Extended Data Fig. 3. Micromorphological and histological analysis. a,**
1171 Ferruginous microfacies (MF-fer: here a coarser-grained variant) adhering on
1172 cancellous bone (plane polarised light: PPL). **b,** Sharp boundary between ferruginous
1173 (MF-fer) and carbonate microfacies (MF-carb) in sediment between Mtoto's bones
1174 (cross-polarised light: XPL). Angular ferruginous nodules and/or MF-fer intraclasts are
1175 present within the carbonate microfacies. **c,** Mtoto's bone fragment. A Fe oxide-stained
1176 calcite crust (Ca) covers the bone surface. A thin zone of non-clouded bone ('HAP' –
1177 conventionally 'hydroxyapatite') immediately beneath the bone surface is underlain
1178 with clouded, Ca-enriched bone of the mesosteum (CLb). Note the sharp boundary
1179 between HAP and CLb (especially at the left side), and dense bioerosion foci within
1180 CLb, such as the linear longitudinal tunnels; budding tunnels (dark spots) and enlarged
1181 osteocyte lacunae (smaller dark spots). Double arrow shows two possible Wedl tunnels.
1182 PPL. **d,** As in (c), but in XPL. Note that the calcite crust (Ca) comprises two layers: a
1183 Fe oxide-stained, microcrystalline layer, and a latter one, of clear, coarser-crystalline
1184 calcite (grey arrow). Birefringent areas within clouded bone (CLb) mark osteons. Note
1185 the loss of birefringence in bioerosion foci (e.g. lower right corner). **e,** Enlarged
1186 osteocyte canaliculi and lacunae in clouded, Ca-enriched bone. PPL. **f,** Advanced
1187 alteration of putative human bone (GHI: 2), with small areas of preserved histology,
1188 pervasive 'clouding', fissuring (e.g. blue arrow), dissolution pores (e.g. green arrow)
1189 and Fe and Mn impregnation (black spots). Note the spatial patterning of bioerosion,
1190 with domains of larger, circular coalescent non-Wedl MFD (e.g. red arrow) and smaller,
1191 more typical tunnels (budding and linear longitudinal: e.g. white arrow). A crust of

1192 calcite speleothem (grey arrows) encrusts a transverse fracture across the bone (PPL).
1193 **g**, Budding and linear longitudinal tunnels in highly altered bone (area marked with
1194 white arrow in (f). Some smaller-scale, spongiform bioerosion is also shown,
1195 surrounded with permineralised rims of redeposited 'hypdoxyapatite' (white bands).
1196 SEM image. **h**, Periosteum of clouded bone (Cb), encrusted with carbonate deposits
1197 (MF-carb). Note the irregular, pitted microrelief of the bone surface, indicative of
1198 dissolution. Larger circular-elliptical pores (blue/turquoise) are haversian canals.
1199 Circles show foci of fine-scale bacterial bioerosion within clouded bone. SEM image,
1200 with colour temperature filter to enhance resolution.

1201

1202 **Extended Data Fig. 4. Histological analysis of Mtoto's bone.** Elemental composition
1203 of clouded bone (Cb) and encrusting carbonate sediment (Cc) (SEM-EDS image and
1204 spectra). The pictured area corresponds to that of Extended Data Figure 3 c,d. Diffuse
1205 lighter grey areas within the clouded bone may be perimineralised rims around
1206 bioerosion foci. Note the variable enrichment in Ca (especially in Spectrum 21) and the
1207 low concentrations of Fe, Al and Mg in the authigenic Ca-P phase that makes up
1208 clouded bone.

1209

1210 **Extended Data Fig. 5. Reconstruction of key taphonomic events of Mtoto's burial.**

1211 The 3D sequence (a,b,c) illustrates the reconstruction of the key taphonomic events
1212 affecting the shape and relationship of the head and the spine. **a**, Original right lateral
1213 decubitus position of the child in the burial pit. **b**, Lateral compression of the thorax
1214 because of the sediment weight; the ribs are flattened but the rib cage does not collapse
1215 as it is common in decomposition in filled space (earth grave). **c**, The head dislocates

1216 as it is typical in the case of burials with perishable head support. **d**, Ideal reconstruction
1217 of Mtoto's original position at the moment of its discovery at the site.

1218

1219 **Extended Data Fig. 6. PYS Shell analysis. a**, Fragments of *Achatina* cf. *fulica* found
1220 in close association with the child skeleton. Observation of anatomical features allows
1221 for a precise identification of the provenance of the shell fragments. Fragments PYS-
1222 2017-200407 and PYS-2017-200404 come from the area of the body whorl adjacent to
1223 the middle portion of the parietal callus. Fragment PYS-2017-200405 comes from a
1224 portion of the shell close to that of the previous fragment, and may derive from the
1225 same individual. Fragment PYS-2017-200406 comes from the middle of the body
1226 whorl, on its dorsal aspect. Although anatomically it is compatible with a provenance
1227 from the same individual, it is very dark in colour, suggestive of a higher Mn intake;
1228 additionally, the concretion coating its outer surface has a different texture, indicating
1229 that it had a different taphonomic history, possibly indicating it derives from a different
1230 shell. Fragment PYS-2017-200086.D comes from the middle of the body whorl, on its
1231 ventral aspect. **b**, Refitting of fragments PYS-2017-200407 and PYS-2017-200404.
1232 The two large fragments refit along an ancient fracture perpendicularly intercepting the
1233 shell growth lines. **c**, Modern striations on the inner surface of specimen PYS-2017-
1234 200406, probably produced during excavation or cleaning of the fragments. The
1235 modern origin of the striations is shown by their random orientation and absence of the
1236 thin manganese patina adhering on the inner surface of this specimen. **d**, Micrographs
1237 and 3D reconstruction of an area of fragment PYS-2017-200404 outer surface showing
1238 two grooves obliquely crossing the decussated sculpture of the outer shell surface. The
1239 grooves internal morphology and outline indicate that they were made by a pointed
1240 instrument, possibly a stone tool, following the irregular morphology of the shell

1241 natural surface and slightly changing direction when running into concave areas. The
1242 antiquity of the lines is demonstrated by the red sediment coating the specimen, which
1243 fills the striations and almost completely buries them when they run into natural
1244 grooves of the shell. **e**, Fragments of *Achatina cf. fulica* found in feature 809 (bottom)
1245 and their anatomical origin (top). The twelve fragments mostly come from the *Achatina*
1246 snail's body whorl and the last whorls of the spire; only two come from the parietal
1247 wall and the apex. They present a similar state of preservation, colour, taphonomic
1248 modifications and type of associated concretions with those observed on the five
1249 fragments found in direct association with the skeleton. None of these fragments bear
1250 incisions similar to those recorded on specimen PYS-2017-200404. Fragments
1251 comprising the control sample from Layer 18 are, in general, less frequently encrusted
1252 with concretions in comparison to those from the skeleton and the burial pit (809). **f**,
1253 Biplot correlating the length and width of *Achatina cf. fulica* fragments from the burial
1254 pit and associated skeleton in comparison to those from Layer 18 (L18) (top), and the
1255 length and width distributions of *Achatina cf. fulica* fragments from these two contexts
1256 (bottom). The fragments from the burial pit are significantly larger in size ($p= 0,001$)
1257 while displaying the same length/width ratio. Incorporation in the grave fill preserved
1258 *Achatina* fragments from greater levels of fragmentation in comparison to those in
1259 Layer 18, which may have been subject to trampling.

1260

1261 **Extended Data Fig. 7. PYS human dental tissue comparison. a**, Adjusted Z-scores
1262 of 2D and 3D, complete crown and lateral enamel, variables. Z-score graphs for the
1263 crown average enamel thickness (2D and 3D AET and 3D LAET), relative enamel
1264 thickness (2D and 3D RET and 3D LRET), and percentage of dentine and pulp (2D
1265 Acdp/Ac, 3D Vcdp/Vc and 3D LVcdp/LVc) in the three PYS molars in comparison

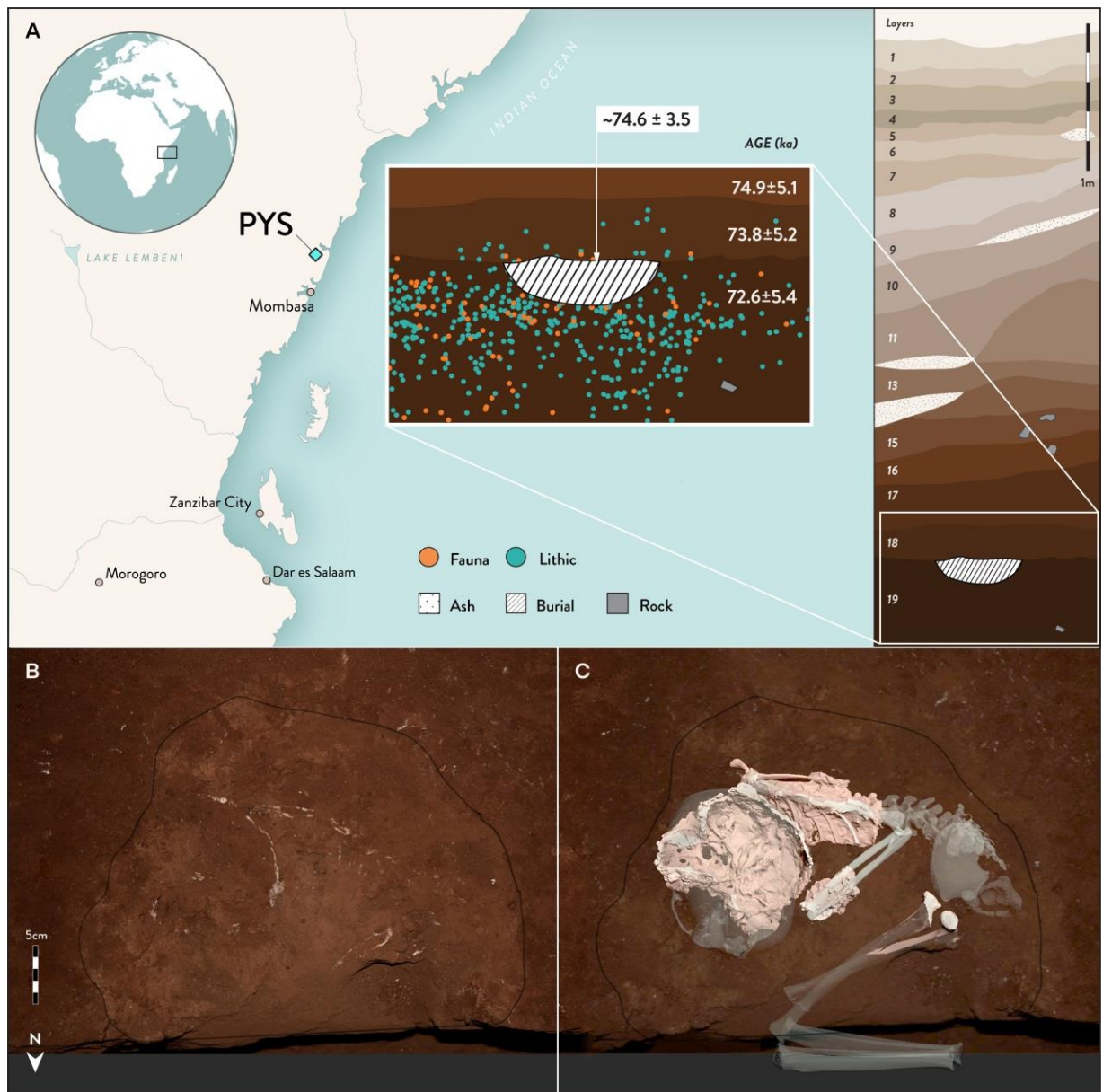
1266 with the variation expressed in Neanderthals (blue triangles) and recent modern humans
1267 (red circles). The solid line passing through zero represents the mean, and the -1 and 1
1268 values correspond to the 95% limit of variation of the two comparative samples. **b**,
1269 Enamel thickness cartographies of the PYS upper and lower molars compared with
1270 those of extinct and extant specimens. Topographic thickness variation is rendered by
1271 a pseudo-colour scale ranging from thinner (dark-blue) to thicker (red). (O=occlusal,
1272 B=buccal). Ldm2 (first row): PYS, NEA=Neanderthal (Roc de Marsal), FHS=fossil *H.*
1273 *sapiens* (Qafzeh 10) and MH=modern human of European origin. RM1 (central row):
1274 PYS, NEA=Neanderthal (La Quina), FHS=fossil *H. sapiens* (Qafzeh 15) and
1275 MH=modern human of European origin. RM1 (central row): PYS, NEA=Neanderthal
1276 (Abri Suard), FHS=fossil *H. sapiens* (Qafzeh 15), and MH=modern human of European
1277 origin.

1278

1279 **Extended Data Table 1. Diagenesis of identifiable and putative human bone in**
1280 **Mtoto's section.** lld: linear-longitudinal MFD; bd: budded MFD; lam: lamellate MFD;
1281 HAP: hydroxyapatite; Cc: calcite.

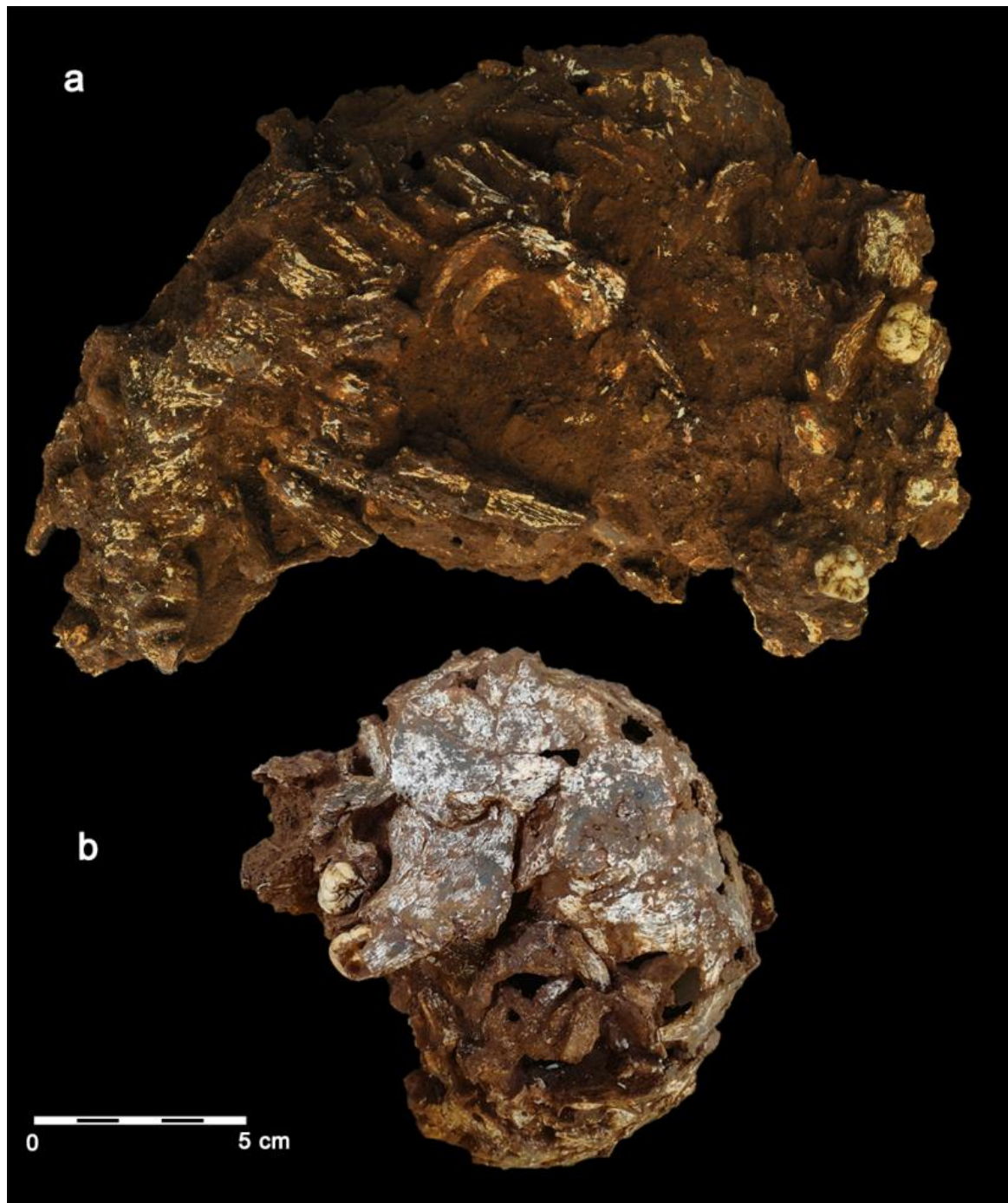
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1283 **Extended Data Table 2. PYS faunal remains.** Fragmentation of the appendicular
1284 skeletal elements recorded from the fill close to the burial in Trench 8. The high level
1285 of fragmentation of the faunal remains contrasts with the relative completeness but
1286 advanced degradation of Mtoto's skeletal remains, indicating different taphonomic
1287 histories for faunal and human remains.



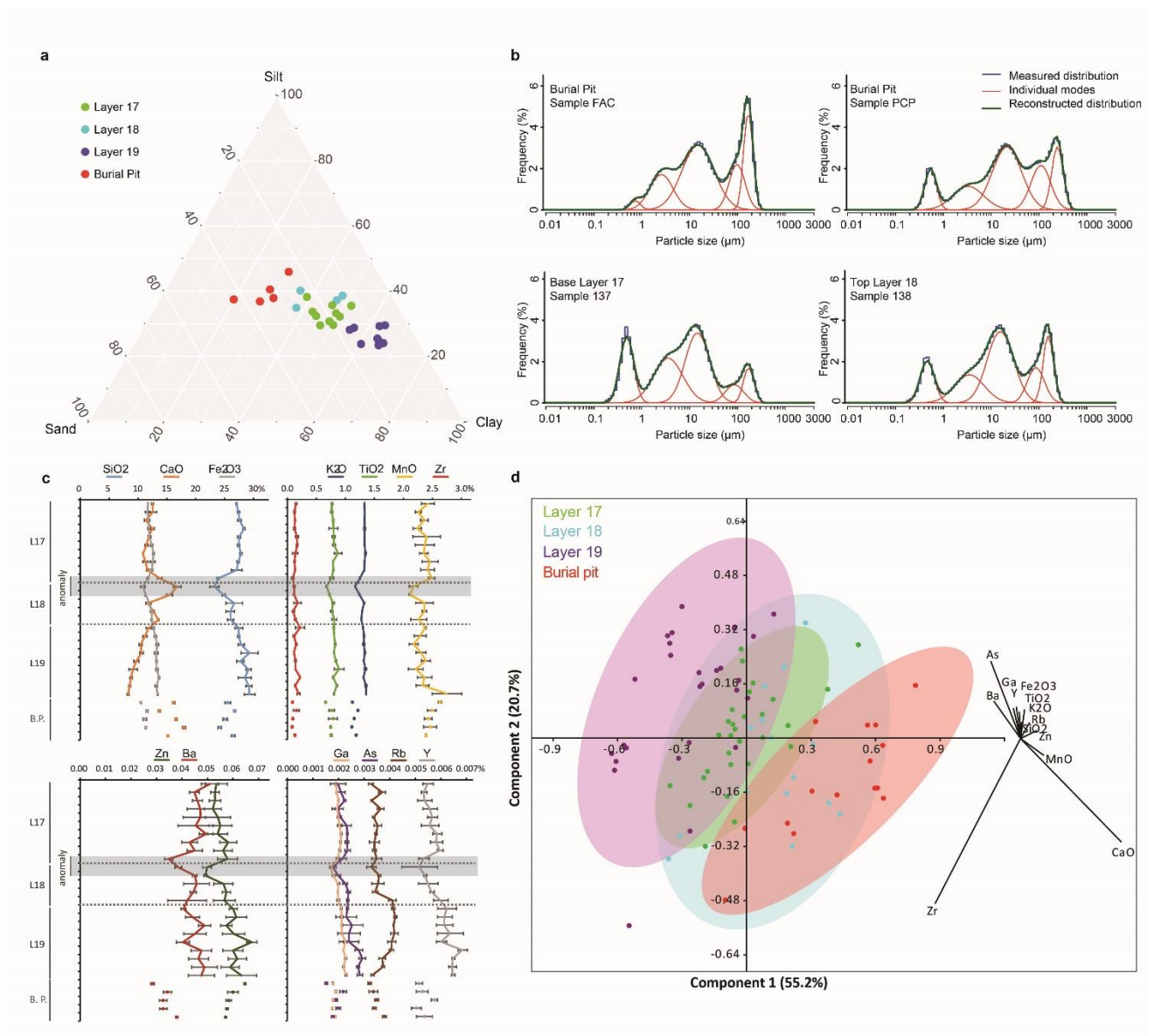
1289

1290 **Fig. 1: Location of PYS and stratigraphic context of burial.** a, PYS is located in the uplands of
 1291 Kenya's coastal plain. Panel on right side shows the 19 stratigraphic layers with the location of burial pit
 1292 in MSA Layer 19. Detailed inset of burial pit shows piece-plotted objects, including MSA lithics and
 1293 fauna and three associated OSL dates in Layers 17-19. The Bayesian age is $\sim 74.6 \pm 3.5$ ka. b, Plan view
 1294 of the 2017 excavation. The black line delimits the pit feature and the change in texture from the
 1295 surrounding matrix. The faint white outline of the decomposed skull, spine, femur and other severely
 1296 degraded bones can be observed on the surface of the pit. c, Same view with the superimposition of the
 1297 virtual reconstruction of the CT and surface scans of Mtoto. The preserved parts (solid) are superimposed
 1298 over a semi-transparent comparative skeleton to better depict the position of the child.



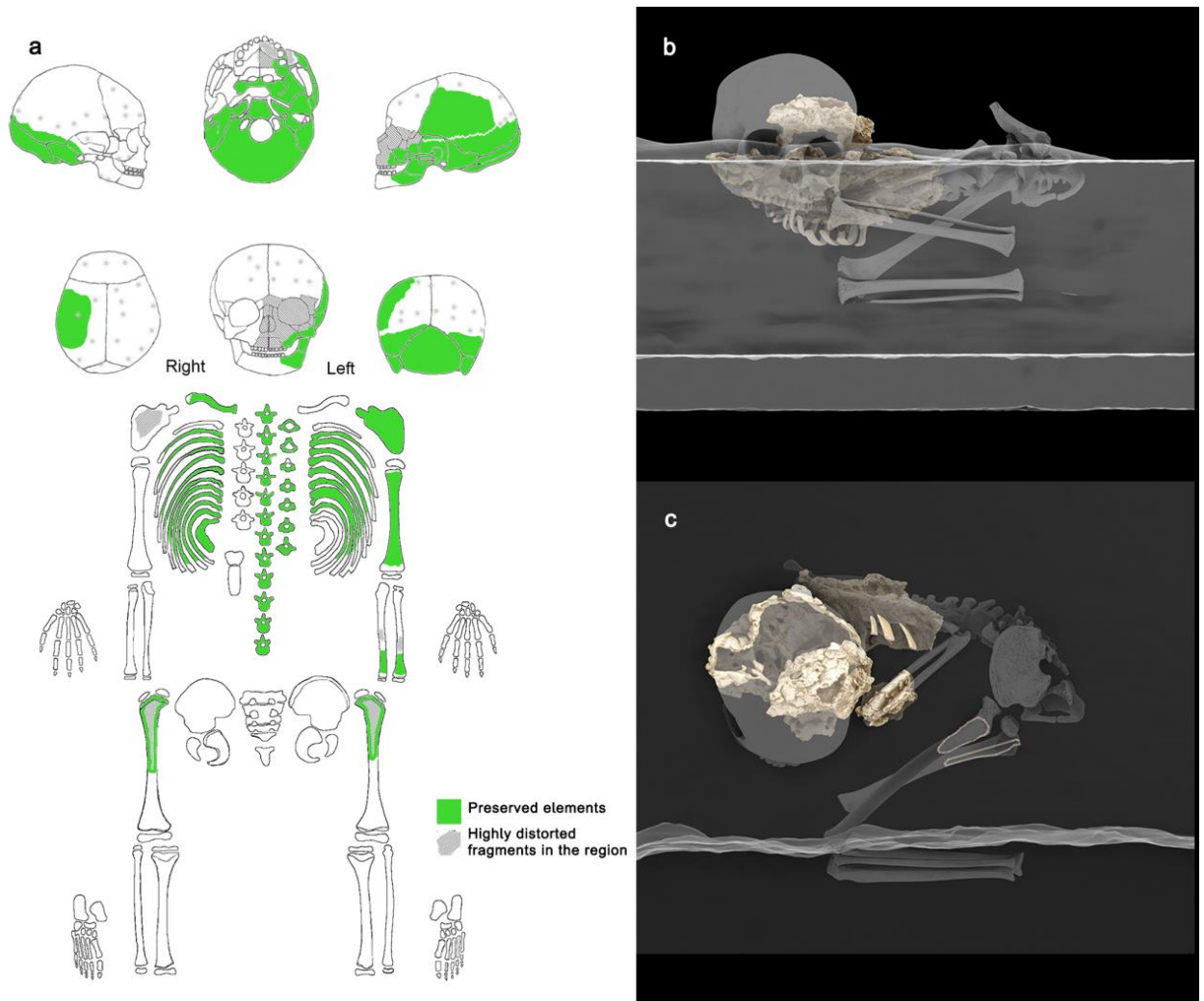
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1300 **Fig. 2: PYS human fossil.** **a**, External view of the PYS main block where the flexed spine with
 1301 articulated vertebrae and ribs, as well as some teeth are partially exposed on the surface. The photograph
 1302 was taken after the initial cleaning and removal of three right thoracic ribs, which revealed how the first
 1303 and second ribs descended and rotated into the thoracic cavity while preserving the intercostal space.
 1304 This argues in favour of minimal displacement of this part of the body as a unit given the sternal
 1305 articulation of the pectoral girdle. **b**, External view of the left side of Mtoto's skull and left hemimandible
 1306 showing the strict and intact temporo-mandibular articulation. The photograph was taken after the skull
 1307 was cleaned and separated from the postcranial elements. The unerupted M_1 and M^1 are held in place
 1308 despite the fact that their roots were not developed, supporting the interpretation of an undisturbed
 1309 deposit. The first three cervical vertebrae are rotated but in place, and connected to the foramen magnum.



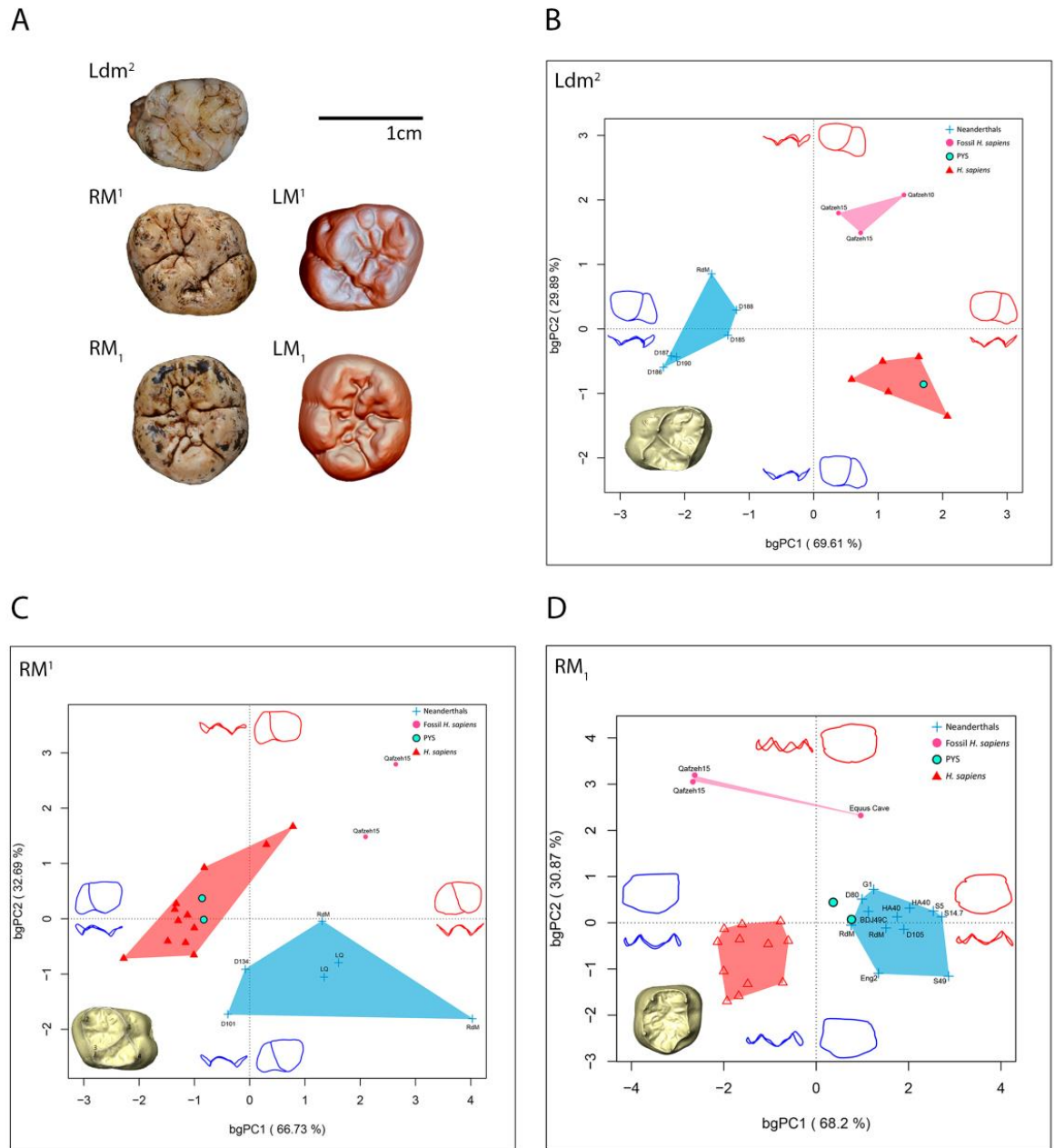
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1311 **Fig. 3: Analysis of sediment from the stratigraphic sequence and the burial pit.** a, Particle size
 1312 ternary diagram indicating the higher content of sand and silt in the burial pit in comparison to the
 1313 encasing archaeological layers, and in particular Layer 19. b, Examples of particle size distribution and
 1314 multimodal decomposition showing the similarities of sand and silt modes between the burial samples
 1315 and samples 137 (base of Layer 17) and 138 (top of Layer 18). c, Elemental profiles of sediment from
 1316 Layers 17, 18, 19 and the burial pit. Element concentrations are expressed in percentages. Sediment
 1317 samples from the burial pit display an elemental composition remarkably similar to the three samples
 1318 identified as an anomaly at the top of Layer 18 and the base of Layer 17. d, Results of PCA of the centre
 1319 log ratio data (selected elements: SiO₂, K₂O, TiO₂, MnO, Fe₂O₃, Zn, Ga, As, Rb, Y, Zr and Ba).
 1320 Confidence ellipses at 95%. The burial pit samples markedly differ from Layer 19.



1321

1322 **Fig. 4: Mtoto's preservation and position in the pit.** a, Preserved parts of the PYS skeleton. Front b
 1323 and top c view of the virtual reconstruction of the preserved skeletal parts (solid) superimposed over a
 1324 transparent comparative skeleton. The fragments of both the left and right femora could not be recovered,
 1325 but their outline was identified on the plan view and in the wall profile, marked as a solid line over the
 1326 transparent skeleton. The position of the right radius and ulna fragments and the right parietal are
 1327 approximate, since they were found detached from the original block.



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Fig. 5: PYS human dental remains. **a**, PYS dental remains: isolated teeth (left column) and mCT 3D reconstruction of the two molars included in the maxillary and mandibular bones (right column). All molars are positioned with the mesial surface towards the top and the distal surface towards the bottom. L (left); R (right); dm2 (second deciduous molar), M¹ (permanent upper first molar), M₁ (permanent lower first molar). **b**, Between-group principal component analysis (bgPCA) of the Procrustes shape coordinates of the PYS Ldm2 enamel dentine-junction surface (EDJ) compared with to those of Neanderthals (n=6), fossil *H. sapiens* (n=3) and modern humans (n=5). **c**, Between-group principal component analysis (bgPCA) of the Procrustes shape coordinates of the PYS RM1 EDJ compared with to those of Neanderthals (n=6), fossil *H. sapiens* (n=2) and modern humans (n=12). **d**, Between-group principal component analysis (bgPCA) of the Procrustes shape coordinates of the PYS RM1 EDJ compared with to those of Neanderthals (n=12), fossil *H. sapiens* (n=3) and modern humans (n=12).