Systematic bone tool production at 1.5 million years ago

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Recent evidence indicates that the emergence of stone tool technology occurred before the appearance of the genus *Homo*¹ and may potentially be traced back deep into the primate evolutionary line². Conversely, osseous technologies are apparently exclusive of later hominins from approximately 2 million years ago (Ma)^{3,4}, whereas the earliest systematic production of bone tools is currently restricted to European Acheulean sites 400–250 thousand years ago^{5,6}. Here we document an assemblage of bone tools shaped by knapping found within a single stratigraphic horizon at Olduvai Gorge dated to 1.5 Ma. Large mammal limb bone fragments, mostly from hippopotamus and elephant, were shaped to produce various tools, including massive elongated implements. Before our discovery, bone artefact production in pre-Middle Stone Age African contexts was widely considered as episodic, expedient and unrepresentative of early *Homo* toolkits. However, our results demonstrate that at the transition between the Oldowan and the early Acheulean, East African hominins developed an original cultural innovation that entailed a transfer and adaptation of knapping skills from stone to bone. By producing technologically and morphologically standardized bone tools, early Acheulean toolmakers unravelled technological repertoires that were previously thought to have appeared routinely more than 1 million years later.

The incorporation of animal resources into the diet of hominins was a milestone in the evolution of our species⁷, with recent evidence confirming that, by 2.6 Ma, hominins modified mammal carcases with stone tools to access meat⁸. As a consequence of the encroachment of technology-driven hominins in the carnivore guild, one of the questions that follow is when the utility of animal skeletal elements expanded to include artefact manufacture in addition to their primary role for consumption^{4,9}. This innovation entailed specific knowledge—technological and also anatomical—to be transmitted through social learning in parallel with lithic technology, opening opportunities for new flexible adaptations that contributed to shaping the evolution of Pleistocene hominin cultures.

The Lower Pleistocene bone tool evidence is sparse; limb shaft fragments and horn cores were used—but not intentionally shaped by knapping—in digging and termite foraging activities at several southern African sites dated between 2.4 and 0.8 Ma^{3,10-13}. At Olduvai Gorge in Tanzania, only a few fossils previously purported as bone tools¹⁴ have stood scrutiny^{4,15,16} and include surface finds (that is, out of their original sedimentary context) and isolated specimens dispersed across a stratigraphic interval spanning over 1 million years¹⁴. Konso in Ethiopia has also yielded some scattered knapped bone artefacts, including the discovery of a bone handaxe found on the surface but attributed to 1.4 Ma deposits¹⁷.

Bone tools shaped by knapping become more frequent in the Eurasian Middle Pleistocene after 500 thousand years ago (ka)⁵. Bone bifaces

are found at several sites from the Levant at Revadim Quarry¹⁸, Central Europe at Vértesszőlős¹⁹ and Bilzingsleben²⁰, and southern Europe at Fontana Ranuccio and Castel di Guido⁵, as well as at the Bashiyi Quarry, Chongging, China²¹. Possible expedient bone shaft fragments bearing one or more edges with flake removal scars on the cortical or the medullar surface, some of which have been interpreted as wedges or intermediate tools, are present in Europe as early as marine isotope stage 9 at Gran Dolina, Spain²², Schöningen, Germany^{23,24}, and in Italy⁵ at Castel di Guido, Bucobello, La Polledrara di Cecanibbio and Rebibbia-Casal de' Pazzi. In East Asia, similar tools are reported throughout the Pleistocene²⁵. Bone tools shaped with techniques such as scraping, grinding and gouging, which allowed the production of diversified and specialized tool types such as spear and arrow points, barbed points, awls and needles, only appear during the Upper Pleistocene in the African Middle Stone Age after 90 ka²⁶⁻²⁹ and in Eurasia after 45 ka³⁰⁻³⁴.

The sparse nature of the early knapped bone tool record has prevented researchers from identifying behavioural consistencies in their production and use, and establish the role they had in early hominin subsistence and cognition. Here we describe a bone tool assemblage from a single horizon securely dated to 1.5 Ma at Olduvai Gorge Bed II, Tanzania, which precedes other evidence of systematic bone tool production by more than 1 million years and sheds new light on the almost unknown world of early hominin bone technology.

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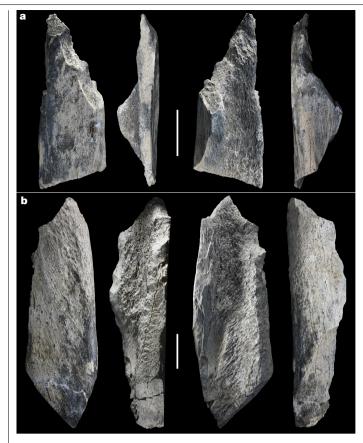


Fig. 1 | Tools made on long bone diaphysis of very large mammals. a, Indeterminable taxon larger than two tonnes (accession number T69L20-3009). b, Elephant (accession number T79L10-2511). Scale bars, 5 cm. See also Extended Data Fig. 6 and Supplementary Videos 4 and 5.

The T69 Complex archaeological site

The T69 Complex is located in the Frida Leakey Korongo (FLK) West Gully at Olduvai Gorge, in northern Tanzania (Extended Data Fig. 1). This site is positioned in the stratigraphic interval between Middle and Upper Bed II, in which early Acheulean assemblages are reported. Radiometric and chronostratigraphic data firmly position the T69 Complex site at 1.5 Ma (Methods).

The T69 Complex includes seven trenches excavated between 2015 and 2022 (Methods; Supplementary Information Videos 1 and 2). The archaeological assemblage that yielded the bone tools was deposited within interlayering silts and sands indicative of an alluvial plain environment with alternating decantation and water flow facies. It contains over 10,900 stone tools that are 2 cm or longer, mostly made on locally available quartzite, and numerous smaller lithic artefacts (more than 41,000). The lithic assemblage includes cores (n = 747) and split cobbles (n = 50), pounded (n = 273) and retouched (n = 230)tools, flakes (n = 2,804) and flake fragments (n = 8,358), and is attributed to the Acheulean³⁵ owing to the presence of large cutting stone tools (LCT; n = 37). The assemblage also contains over 9,419 identifiable vertebrate fossils and 13,413 unidentified bone fragments. Abundant fish, crocodile and hippopotamus remains are consistent with lithological data indicating the proximity of water sources. The large mammal assemblage is dominated by bovids and hippopotamus, the latter represented by relatively complete carcasses. Equids, suidae, elephants, rhinoceros and other taxa are also present. Hippopotamus is the most abundant genus, and bones frequently exhibit evidence of anthropogenic manipulation, suggesting that hominins were attracted to the T69 Complex area by the availability of hippopotamus carcasses (Methods; Supplementary Data 1).

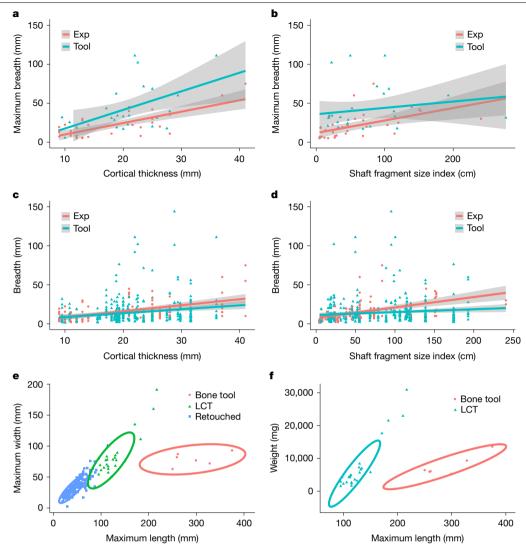
The bone tool assemblage

The T69 Complex faunal assemblage, including the bone tools, presents an excellent state of preservation that allows documentation of natural and anthropogenic modifications in detail. Taphonomic and morphometric analyses rule out natural processes to account for the modifications recorded on the 27 specimens identified as bone tools (Methods). Several factors are known to cause flake removals on bone that mimic intentional flaking⁴, including carnivores gnawing, crocodiles biting, trampling and fracturing to access marrow. Several reasons rule out these factors: carnivore remains amount to less than 1% of the identified specimens. With the exception of two possible tooth marks (Supplementary Data 2), no modifications produced by carnivores are recorded on the T69 Complex bone tools, whereas they are abundant in bone assemblages modified by these agents and yielding fragments bearing flake removals 36,37. The fresh appearance of the edges of the tools and the rarity of striations attributable to trampling indicate that this process had minimal effect on the state of preservation of the bone artefacts and cannot be the cause for the numerous invasive flake removals present on them. Experimental breakage of large mammal limb bones for marrow extraction, including elephant bones, has demonstrated that flake removal scars resulting from this activity rarely amount to more than four (Extended Data Fig. 4f-h), they are mostly isolated or, when contiguous, rarely exceed three, and they preferentially occur on the ends rather than the lateral sides of the splinters^{4,9}. T69 Complex faunal fragments display an average of 2.1 flake scars that occur as isolated removals in 84.3% of the specimens (Extended Data Fig. 4b-d), conforming with a pattern typical of limb bone breaking to access marrow. By contrast, the T69 Complex bone tools exhibit an average of 12.9 flake scars per specimen (Extended Data Fig. 4i-k) and are always arranged contiguously and preferentially-but not exclusively-on their lateral edges.

All 27 bone tools were found in situ during excavation (Supplementary Video 3). Eighteen (66.7%) come from bones attributed to mammals heavier than two tonnes (Fig. 1, Extended Data Fig. 5 and Supplementary Data 2). Among the 16 taxonomically identifiable specimens, eight are from elephant, six are from hippopotamus and two from bovids (Supplementary Data 2). Thus, although the mammal assemblage is dominated by bovids (41.1%), a minimum of 50% of bone tools are made from elephant, which only makes 1.1% of the T69 Complex taxa composition. With the exception of a single proximal portion of a large boyid radius that preserves part of its epiphysis, all tools are exclusively made on limb bone shaft fragments, particularly femora, tibiae and humeri.

The bone tools display morphological, technological and dimensional characters identifying behavioural patterns previously undocumented at early hominin sites. The surface of flake removals is fresh, which is indicative of the short period of subaerial exposure elapsed between bone tool production and burial. All tools made of hippopotamus limb fragments come from bones broken while they were in a nutritive state. For those made of elephant, both fresh and partially weathered bones were used (Supplementary Data 2), which suggests that hominins accessed fresh carcasses but also defleshed bones. The tools are considerably longer than most of the faunal assemblage and fall within the size range of experimentally broken elephant long bones⁴ (Extended Data Fig. 4a,e,i). Bone tools made from elephant bones are the largest, ranging between approximately 22 and 38 cm in length, and approximately 8 and 15 cm in width. Tools made from hippo bones are slightly smaller with ranges between approximately 18 and 30 cm in length and approximately 6 and 8 cm in width (Extended Data Fig. 5).

Elephant bone tools bear on average more flake removal scars $(\mu = 17.3)$ than those made on hippopotamus bone $(\mu = 13.3)$. Among the taxonomically unidentifiable specimens, seven bone tools show sizes and number of flake removal scars that are compatible with those made on elephant or hippopotamus bones. Eight tools are smaller in size and bear on average six flake removal scars. When shaping bone



 $Fig.~2 \,|\, Metric comparison of the breadth of flake removal scars relative to the dimension of the bone tools, and the size and weight of the notched bone tools compared with the lithic artefacts found in the same assemblage.$

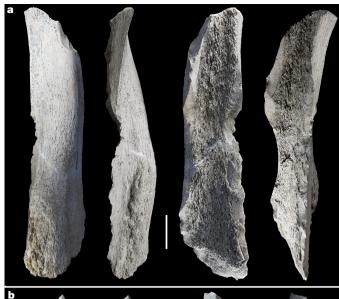
 \mathbf{a} - \mathbf{d} , Scatterplot of the breadth for the most invasive (\mathbf{a}, \mathbf{b}) and all (\mathbf{c}, \mathbf{d}) removal scars present on the FLK T69 Complex bone tools and on bone fragments produced during experimental (Exp) bone fracturing to expose the marrow of horse and elephant limb bones compared with the cortical thickness (\mathbf{a}, \mathbf{c}) and the shaft fragment size index (\mathbf{b}, \mathbf{d}) . The maximum breadth of the largest flake removal scars on the bone tools and on fragments of limb bones fractured for marrow extraction demonstrates that the former are considerably larger

tools, the T69 Complex knappers invested substantial effort in first producing invasive flake removals to shape the tool and, subsequently, regularizing the resulting edges via trimming. These two steps are documented by comparing the breadth of the flake removal scars with the cortical thickness and the shaft fragment size index (Fig. 2a-d). The most invasive flake removal scars on the bone tools are significantly longer (F = 21.17, d.f. = 2 and 461, P < 0.0001) than those on fragments of limb bones fractured experimentally for marrow extraction (Fig. 2a,b). The trimming phase, by contrast, produced numerous small contiguous flake removal scars comparable in size-but not in their arrangementwith those on bone fractured experimentally for marrow extraction, which are generally isolated (Fig. 2c,d). Production of blanks for the largest artefacts required striking heavy and sizeable stone percussors against stationary bones⁴, whereas morphology of flake removals in all bone tools is compatible with the use of handheld hammerstones during the shaping stage.

irrespective of the cortical thickness or the shaft fragment size index. The difference in trend observed when considering all flake removal scars is probably due to the production of numerous small contiguous retouch flakes to shape the bone tool edges. **e.f.** Length and width (**e**) and length and weight (**f**) comparisons between the T69 Complex bone tools and the lithic assemblage found in the same horizon are shown. Bone tools are more elongated than the LCT, but their current weight falls within the range of variation recorded for the LCT found at the T69 Complex. Trend lines and grey bands indicate the linear regression method and the 90% confidence interval, respectively (**a**–**d**), and the ellipses denote the 90% confidence interval (**e.f**).

Six tools made on more than 2-tonne mammal bones show a recurrent modification pattern to produce a particular shape (Fig. 3 and Extended Data Figs. 7 and 9). With an average of 16.8 removals per specimen, they feature one crescent and one pointed end, coupled with a large invasive notch produced by contiguous removals in the mesial part of the object. Most removal scars making the notch are present on the medular surface. The pointed end of these tools systematically corresponds to the robust mid-portion of the diaphysis, and the rounded end to the metaphysis. The large notch may have facilitated the prehension of the tool while preserving its sturdy shape and heavy weight.

A function as heavy-duty tools for the larger bone artefacts is suggested by comparison with LCT and smaller retouched lithics from the T69 Complex assemblage. The notched bone tools are larger, more elongated and more intensively shaped than the stone tools—LCT average of 10.1 flake scars per artefact—whereas their current weight falls within the range of recorded variation for LCT found at the site



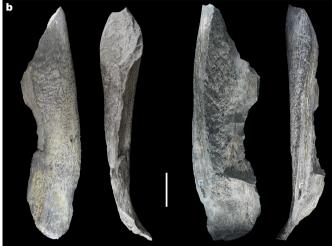


Fig. 3 | Large bone tools made on diaphysis fragments. a, Elephant humerus (accession number T79L10-9047). b, Hippopotamus femur (accession number T79L10-18461). Scale bars, 5 cm. See also Extended Data Fig. 7 and Supplementary Videos 6 and 7.

(Fig. 2e,f). Furthermore, distal fracture patterns on some of the specimens evoke their use in percussive and compressive activities³⁸⁻⁴¹ (Methods; Extended Data Fig. 11).

In summary, to produce bone artefacts at the T69 Complex, hominins selected bones from large mammals and preferentially elephant. Precise anatomical knowledge and understanding of bone morphology and structure are suggested by preference given to thick limb bones as tool blanks. Excellent understanding of bone fracture mechanics is shown by the preferential use of large mammal fresh bones and the application of recurrent flaking procedures. Mental templates are suggested by the production of morphologically similar, elongated, pointed and notched bone tools.

Implications for early bone tool technology

Systematic bone tool production from a single horizon dated to 1.5 Ma at the T69 Complex prompts a reconsideration of the emergence and evolution of bone technology among early hominins. Previous evidence on bone tool production in the Early Stone Age concerned either used unmodified bone fragments^{3,11-13}, isolated discoveries of possible knapped tools preventing firm conclusions on early hominin behaviours^{4,14-17}, or bone tools shaped by knapping only occurring

after 500 ka^{5,6,9,18-21,42}. By contrast, the T69 Complex contains, within the same archaeological assemblage, a set of bone tools that share consistent technological features and signal to the existence of a patterned behaviour.

The T69 Complex bone tools demonstrate that.1 million years before the production of fully formatted bone bifaces, East African hominins complexified their technology by systematically integrating large, knapped bone tools into it. Such integration occurred at a pivotal time in the evolution of African cultural adaptations—namely, the transition between the late Oldowan and the early Acheulean-and may have had a profound effect on the complexification of behavioural repertoires observed in the latter period⁴³, including enhancements in cognition and mental templates⁴⁴, artefact curation and raw material procurement⁴⁵. This innovation took place at a moment in which large bifacial stone tools had a minor role in their technical systems 46 and had yet to acquire the large size, refined technology and typical symmetrical morphology known for bifaces from subsequent Acheulean assemblages⁴⁷. Particularly at butchery sites of large mammals where suitable raw material was readily available, large heavy bone tools may have fulfilled functions that were later achieved by large bifacial stone tools, which afforded both heavy weight and effective cutting edges. This hypothesis may explain why after the advent of systematic lithic handaxe production, heavy-duty bone tools such as those found at the T69 Complex, potentially less efficient in cutting tasks, may have become rare. In this view, Middle Pleistocene Acheulean bone handaxes⁵ could be interpreted as a local transfer to bone of a knapping technology in contexts where scarce availability of optimal lithic raw materials hindered the production of stone LCT.

The alternative scenarios entail that bone technologies appeared and disappeared across more than 1 million years, or were more common in the Early Stone Age than previously thought. In both cases, evidence of such technology is yet to be reported adequately. Preservation bias against organic materials and the limited availability of studies targeting the identification of knapped bone may be masking the widespread and systematic use of osseous technologies, until now thought to appear much later in our evolutionary history. Future research needs to investigate whether similar bone tools were already produced in earlier times, persisted during the Acheulean and eventually evolved into Middle Pleistocene bone bifaces similar in shape, size and technology to their stone counterparts.

Online content

Any methods, additional references. Nature Portfolio reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41586-025-08652-5.

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Methods

Site context

Location and age. The T69 Complex (WGS84 UTM Zone 36S, x = 760990; y = 9669250) is located in the Frida Leakey Korongo (FLK) West outcrop, within the Main Gorge at Olduvai Gorge, in northern Tanzania. Excavations were conducted annually through eight field seasons between 2015 and 2022. Seven trenches (T69, T71, T74, T75, T77, T78 and T79) were dug at the site, of which three (T71, T74 and T75) yielded no archaeological materials (T71 was placed above and T74–T75 below the fossil-bearing horizons). Separate names were given to the trenches as test pits progressed around the main outcrop, but eventually they were all linked up into one single excavation area (Supplementary Video 1) named T69 Complex, after the first trench we dug at the FLK West locality.

Stratigraphically, the site is positioned in Olduvai Bed II, which overlays Bed I and is capped by Bed III (Extended Data Fig. 1e). The top of Bed I (Tuff IF) is dated to 1.803 ± 0.002 Ma 48 , whereas the base of Bed III is estimated at 1.14 ± 0.05 Ma 49 . Within Bed II, the T69 Complex is placed within a sandstone bed in the Tuff IIC stratigraphic interval 50 that marks the transition between Middle and Upper Bed III and sits 7 m above the Bird Print Tuff and 4 m below Tuff IID 35 . A tuff located 20 cm below the Bird Print Tuff 51 has been dated at 1.664 ± 0.019 Ma 52 , and an average age of 1.339 ± 0.024 Ma has been estimated for Tuff IID 53 . The bone tool level in the T69 Complex has been dated at 1.48 ± 0.2 (1 σ) Ma using direct cosmogenic nuclide isochron burial dating 35 , which is consistent with the constraining ages available for Beds I, II and III (Extended Data Fig. 1e).

During the Tuff IIC interval, the Olduvai perennial palaeolake was constrained by the graben between the Fifth and FLK faults. The FLK locality was positioned in the southeastern lake-margin zone, with drainage inputs towards the north-northeast⁵⁰. The T69 Complex deposits are dominated by the fluvial and lacustrine processes occurring at the eastern margin of the Bed II palaeolake⁵⁰. The lower interval consists of alluvial massive sands and silty decantation facies (Extended Data Fig. 1d). These deposits were partially eroded by water flows, which led to the sedimentation of the sandstone bed containing the bone tool horizon. This sandstone has a sheet-like geometry and contains mainly tuffaceous facies with a massive or poorly bedded structure. The sandstone bed includes local lenses of augitic cross-laminated facies more abundant towards the base, and discontinuous patches of cemented very fine sandstone towards the top. Thin sections of the sandstone indicate a clay matrix and subrounded to rounded grains, from very coarse to fine in size, that locally include claystone rip-up subrounded clasts (Extended Data Fig. 1f). These sedimentological features suggest that cross-laminated facies were generated by local phases of traction and sedimentation under low regime flow conditions, whereas predominant massive or weakly bedded facies were formed by relatively rapid deposition of suspended load during decreasing flood. The sandy facies were buried by progradation of lacustrine brown clays (Extended Data Fig. 1d).

Excavation and sample preparation. Sedimentary units with no archaeological remains overburdening fossiliferous beds, which in some parts of the outcrop were more than 6 m thick, were removed manually with larger picks. Excavation of archaeological units initiated with small picks, and when remains were found, these were uncovered sequentially with screwdrivers, dental tools and wooden sticks. Each item found in situ, regardless of its size, was given a unique ID, positioned three-dimensionally with a total station, and recorded following the protocols outlined by de la Torre et al.⁵⁴. Azimuth (dip direction, $0-360^{\circ}$) and dip angle $(0-90^{\circ})$ of the major axis of suitable elongated items⁵⁵ were recorded with compass and clinometer during the excavation process. All sediment collected from the archaeological units was dry-screened with 2-mm-mesh sieves. All items with an individual ID were hand-labelled, and a QR code was attached and added to a geographical information system that integrates all archaeological information for the FLK site.

Following conservation protocols detailed elsewhere ⁵⁶, some bone tools were stabilized by temporary consolidation with cyclododecane (a cyclic alkane hydrocarbon) before being lifted from the excavation. Solutions of paraloid B-72 (ethyl methacrylate—methyl acrylate co-polymer) or B-44 (methylmethacrylate—ethylacrylate) on acetone were also used. Mechanical removal of sedimentary accretions avoided contact with bone surfaces. When necessary, excessive consolidant was removed, breaks aligned and further consolidation carried out. Gaps or cracks that undermined structural condition of fossils were filled with B-44 or B-72 mixed with acetone and microscopic glass spheres.

Site formation. The T69 Complex bone horizon spans an average thickness of approximately 50 cm and was excavated across a surface of 295 m² (Extended Data Fig. 2 and Supplementary Video 2). Using ArcGIS Pro (v3.3), the horizontal distribution pattern was tested through an average nearest neighbour analysis that compared the actual distance between items to a hypothetical random sample 57 . The complete assemblage and the lithic and fossil assemblages showed a significantly clustered distribution (P = 0.000; Extended Data Fig. 3-1), whereas the bone tools were randomly distributed across the clusters (P = 0.6785).

Orientation^{58,59} and fabric analyses^{55,60} were conducted using Oriana software (v3.13) to investigate potential rearrangements of the assemblage. Orientation angular histograms show a circular-shaped distribution (Extended Data Fig. 3-2a), which becomes more homogeneous when axial values are projected (Extended Data Fig. 3-2b). When considering orientation as axial data (0-180°), statistical tests do not allow rejecting the null hypothesis of uniformity. By contrast, all azimuthal data statistical tests reject the uniform distribution against unimodal and plurimodal distributions, indicating a preferential orientation pattern with a mean vector at N22° E (see statistical results in $table 1 in Supplementary \, Data \, 1). \, Stereographic \, projection \, of \, azimuth \,$ and dip values showed that most of the data concentrated around the external area and dominated by low dip angles (Extended Data Fig. 3-2c). Fabrics indicate a gridle or planar shape (Extended Data Fig. 3-2d,e), with eigenvector 1 pointing towards N27° E and thus coinciding with the mean vector direction (table 2 in Supplementary Data 1).

In summary, horizontal distribution patterns do not indicate a differential distribution between fossils and lithics, with both groups appearing together in clusters and bone tools scattered randomly across the excavation surface. The planar or girdle fabric with low dip angles suggests that archaeological items were accumulated on a subhorizontal surface and did not experience post-depositional vertical dispersion. Nonetheless, azimuthal data in the horizontal dimension indicate a notable influence of the dip direction, characterized by a mean vector that coincides with the north-northeast palaeo-drainage trend estimated by Hay⁵⁰ in the lake-margin zone near the FLK locality. This suggests that the archaeological assemblage may have experienced some rearrangements by the alluvial flows depositing the sandstone bed. Given the generally excellent preservation of the archaeological items, the sheer abundance of less than 2 cm lithics, the sedimentological features of the sandstone bed (see the 'Location and age' section), and the orientated but scarcely clustered angular pattern, this gentle reorganization would be produced by low-energy flood-sheets with a poor sorting capacity, in which the item direction was determined by the depositional surface.

Faunal assemblage. The T69 Complex fossil assemblage (n = 22,832; bones of 2 cm or more = 45.3%; bones of less than 2 cm = 54.7%) contains abundant mammal (more than 11,627), reptile (n = 1,453), fish (n = 1,084) and bird (n = 286) specimens. Mammal bones of 2 cm or more (n = 8,930) were sampled for full zooarchaeological analysis following refs. 56,61,62. This zooarchaeological control sample (ZCS; n = 2,075) included (1) all items identified as bone tools, (2) all proboscidean fossils (given their relevance among the bone tool assemblage, the entire T69 Complex fossil collection was inspected to ensure all elephant specimens were analysed), and (3) a randomly selected sample from all other

2 cm or more mammal bone specimens. Whenever possible, the taxa and skeletal element were recorded. Otherwise, the specimens were attributed to a mammal size class based on their cortical thickness following Bunn⁶³. Fresh and weathered bone fractures were distinguished based on criteria available in the literature⁶⁴⁻⁷¹.

The ZCS skeletal part profile is dominated by axial bones (ribs and vertebrae), followed by appendicular elements (particularly tibia, femur, humerus and unidentifiable long bone fragments; tables 3 and 4 in Supplementary Data 1). Taxonomically, the ZCS assemblage is dominated by Bovidae (n = 691; 33.3%) and Hippopotamidae (n = 233; 11.2%), with considerably lower proportions of Equidae (n = 49; 2.3%) and Elephantidae (n = 22; 1.1%), and residual presence of other taxa such as Rhinocerotidae (n = 8; 0.4%), Suidae (n = 7; 0.3%) and others (n = 2; less than 0.1%); table 5 in Supplementary Data 1). The assemblage contains at least 16 different individuals from 10 distinct species (table 6 in Supplementary Data 1): Hippopotamus cf. gorgops (minimum number of individuals (MNI) = 4), Equus sp. (MNI = 2), Elephas cf. recki (MNI = 2), Diceros cf. bicornis (MNI = 2), Damaliscus cf. lunatus (MNI = 1), Connochaetes aff. gentryi (MNI = 1), Eudorcas cf. thomsonii (MNI = 1), Hippotragus gigas (MNI = 1), Kobus sp. (MNI = 1) and Tragelaphus aff. imberbis (MNI = 1). Overall, taxonomic composition and animal size distribution (table 7 in Supplementary Data 1) of the T69 Complex site suggests similar ecological conditions to the modern-day Serengeti ecosystem, with open and seasonal grassland habitats dominated by large communities of size 2 and 3 bovids near rivers and at the margin of an alkaline lake that accommodated hippopotamids, fish and crocodilians^{72,73}.

Cut marks (present in more than 5.5% of the ZCS specimens) and percussion marks (more than 3.1%) indicate that early humans butchered carcasses at the site (tables 8 and 9 in Supplementary Data 1). According to the tooth-marked bone proportion (more than 2.4%), carnivores had a lesser role in the formation of the assemblage (table 10 in Supplementary Data 1). A short period of subaerial exposure is documented in 39.8% of specimens, although stage 2 (ref. 74) bones dominate the sample and more weathered bones are also present (table 11 in Supplementary Data 1). All these taphonomic signatures indicate separate assemblage formation episodes and are consistent with the alluvial dynamics that characterize the T69 Complex depositional environment.

Site function. Taxa represented, skeletal part profiles and taphonomic patterns indicate that hominins occupied a floodplain environment where they accessed relatively complete carcasses of hippopotamus. Other mammal body parts that show anthropogenic modifications—including bovids (cut-marked bones) and elephants (flaked bones)—were potentially transported by hominins to the site. Hominins also imported large quantities of quartzite from the nearby hill of Naibor Soit oproduce stone tools. The input of other biotic as well as abiotic agents to the formation of the assemblage is attested by the presence of tooth-marked bones—indicating carnivore action—birds, reptiles and fish—unrelated to the formation of the anthropogenic assemblage—and the sedimentary evidence—pointing at gentle rearrangements of parts of the deposit.

The large size of both the stone tool and bone assemblages suggests that hominins visited the area repeatedly, probably attracted by the availability of dead hippopotamus. Exploitation of hippopotamus carcasses was not limited to meat procurement, but also to the production of bone tools. Cut marks produced by lithics clearly indicate that stone tools imported from Naibor Soit had a role in the butchery activities, but we argue that hominins were also importing bone raw materials; elephant fossils are rare in the T69 Complex faunal composition, whereas they are overrepresented in the bone tool assemblage. We hypothesize that bone-tool makers accessed elephant limbs elsewhere and shaped large diaphyseal fragments that were then imported to the T69 Complex and used in percussive and compressive activities, potentially related to butchery of the hippopotamus carcasses that attracted hominins to the area.

Bone tools

Taphonomy. Bone tools (n=27) with most conspicuous technological features were recognized onsite during the excavation process (Supplementary Video 2), whereas others were identified during the inspection of the entire sample of mammal bones of 2 cm or longer. Each specimen was first examined with a magnifying glass with incident light. Anthropogenic modifications were distinguished from natural modifications based on published criteria $^{41,74-85}$, with a particular attention on the natural and anthropogenic processes that could produce flaking scars on faunal remains, such as trampling $^{86-88}$, carnivore alterations 36,37,83,8990 and marrow extraction $^{4,9,91-95}$ (see below and Supplementary Data 3).

Bone tool analysis and comparison with archaeological and experimental samples. The archaeological material considered in the present study comprised: (1) the 27 bone specimens bearing features, that is, flake scars, impact and morphology, that led us to consider them as bone tools (henceforth, BTS; Figs. 1 and 3 and Extended Data Figs. 6–10). (2) A sample of 4,507 long bone fragments from the same excavation (henceforth, random control sample (RCS)). The RCS was selected to verify whether the bone tools stood out morphologically when compared with the rest of the fauna or instead represented an extreme in size variation, that is, in maximum length, width and thickness of the bone fragments. (3) A subsample of 250 long bone fragments from RCS (henceforth, random control subsample (RCSS)). The RCSS was selected to verify whether the number of flake removal scars observed on the BTS stood out from other specimens in the faunal assemblage or instead represented an extreme in variation.

To establish whether marrow extraction activities could have produced a flaking pattern akin to that observed on the RCSS and BTS, we compared the T69 Complex specimens with two published experimental samples of large mammal long bones fractured to expose marrow. The first, henceforth, experimental sample 1 (ES1), consisted of nine elephant long bones fractured by knapping that produced 107 diaphyseal fragments⁴. The second, ES2, derives from the fracturing of six horse long bones by knapping, which resulted in 33 diaphyseal fragments and 284 flakes and splinters⁹. The maximum length, width, thickness and cortical thickness of the specimens included in BTS, ES1 and ES2 were collected using a digital calliper. The following variables were recorded for specimens with flake removal scars: number of scars, the location of each scar (cortical or medullar surface, distal. proximal, right or left lateral edge), their arrangement (isolated, contiguous or series of contiguous flakes) and the breadth of each scar. The contiguous category includes both adjacent and overlapping flake removal scars. Series of contiguous flake scars imply two or more groups of contiguous scars separated by an unmodified edge. The breadth of the flake removal scars was compared with the cortical thickness of the diaphyseal fragment and with the shaft fragment size index, that is, the product of the maximum length and width (in centimetres) divided by two⁴, to detect anomalously invasive flake removal scars on BTS compared with those recorded for the ES1 and ES2 specimens.

Recurrent patterns in the anthropogenic modification of bone fragments were sought in the frequency, size and arrangement of flake removal scars as well as on fractures resulting from potential use. Technological features were described following conventions proposed by Inizan et al. ⁹⁶ and adapted to bone tools analysis by Pante et al. ¹⁶. Size comparison was done between the heavy-duty bone tools, that is, a subsample of the BTS, and the lithic LCT and retouched flakes found in the T69 Complex assemblage. Finally, the weight of the heavy-duty bone tools and lithic LCT was also compared.

All bone tools were photographed with a Nikon D7500 camera (Nikon DX 18–140 mm 1:3.5–5.6 and Nikon N 105 mm 1:2.8 lenses), and 3D scanned with an Artec Space Spider 3D scanner. Relevant bone surface modifications were analysed with a Leica DVM6 digital microscope

(Planapo FOV 12.55 and 43.75 objectives, up to \times 675). The quantitative analyses and graphs were done in R-CRAN (v4.4.1)⁹⁷.

Reporting summary

Further information on research design is available in the Nature Portfolio Reporting Summary linked to this article.

Data availability

All data generated or analysed during this study are included in this published article and its Supplementary Information files.

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Author contributions I.d.I.T. and J.K.N. directed the Olduvai Gorge Archaeology Project fieldwork. I.d.I.T. and R.M. supervised excavations. A.B.-C. conducted the geological study. R.F.P. supervised conservation of the faunal assemblage. I.M. conducted the zooarchaeological analysis. I.d.I.T., J.K.N., F.d'E. and L.D. analysed the bone tools. Quantitative analysis was produced by L.D. (bone tools), A.B.-C. (orientation and fabrics) and I.M. (zooarchaeology). A.T. produced the imagery and collected the 3D scan data. I.d.I.T., F.d'E. and L.D. wrote the first draft. All authors discussed the results and contributed to the manuscript.

Competing interests The authors declare no competing interests.

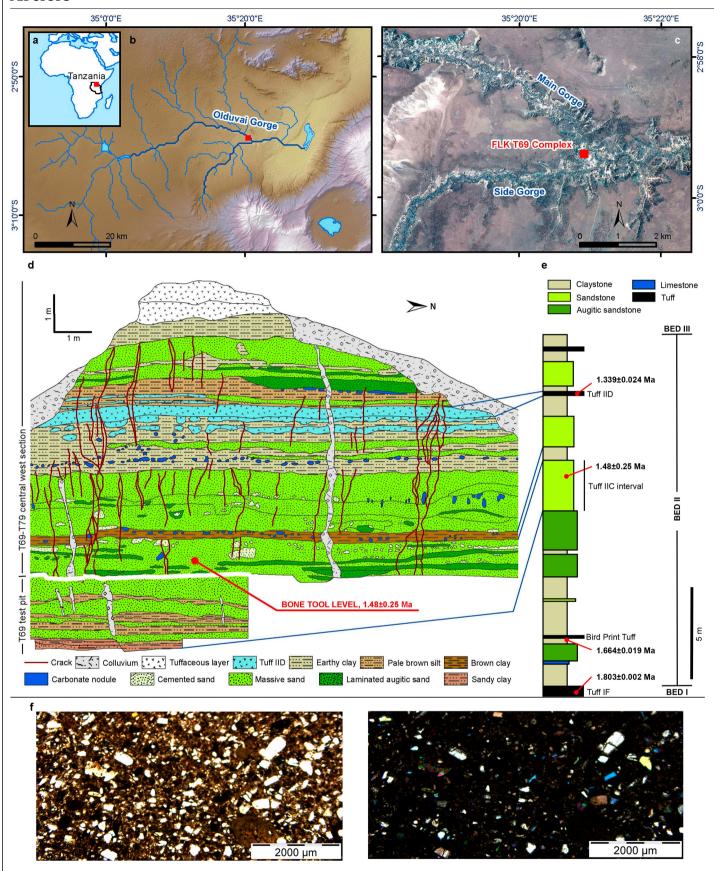
Additional information

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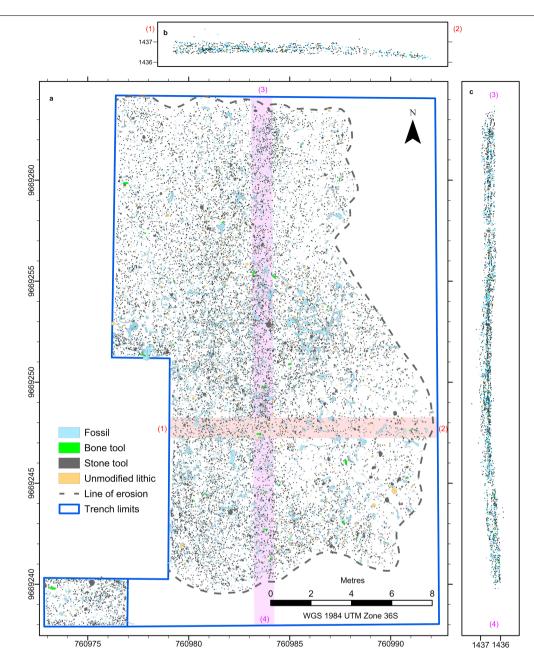
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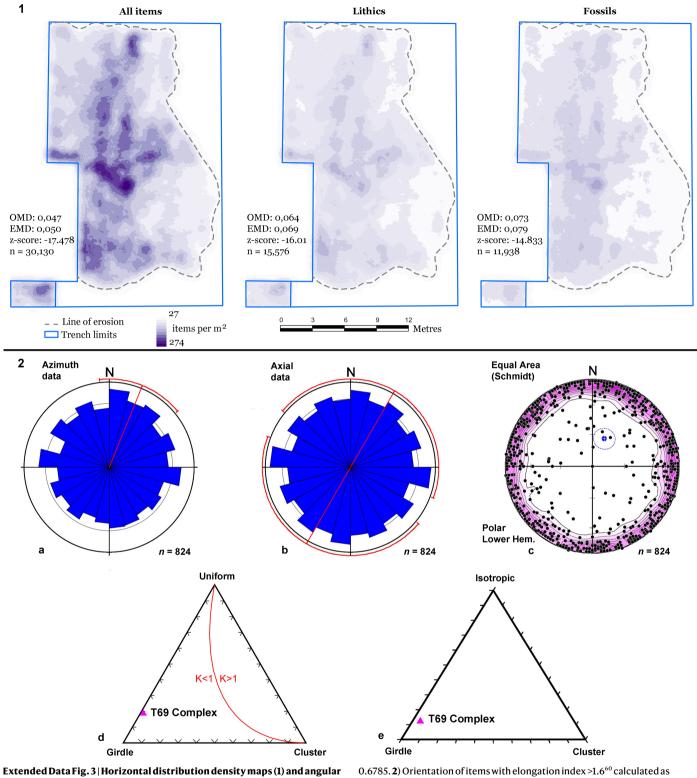
 $\textbf{Extended Data Fig. 1} | See \ next \ page \ for \ caption.$

Extended Data Fig. 1 | Geographic and stratigraphic context of the FLK T69 Complex site. a, Location of Olduvai Gorge in Tanzania. b, Main geographical features of the Olduvai Basin. The geographical image was adapted from NASA (https://lpdaac.usgs.gov/products/astgtmv003/), USGS. c, Position of the T69 Complex in the Main Gorge at Olduvai. The geographical image was adapted with permission from ref. 98, Elsevier. d, Stratigraphic position of the bone tool

level shown on a composite section of the deepest part of trench T69 and the central area of the T69-T79 west section. \mathbf{e} , Position of the T69 Complex within the general chrono-stratigraphic sequence of Bed II in the FLK area. Dates by 35,48,52,53 . \mathbf{f} , Normal (left) and polarized (right) thin section photographs of the sandstone containing the bone tool horizon.

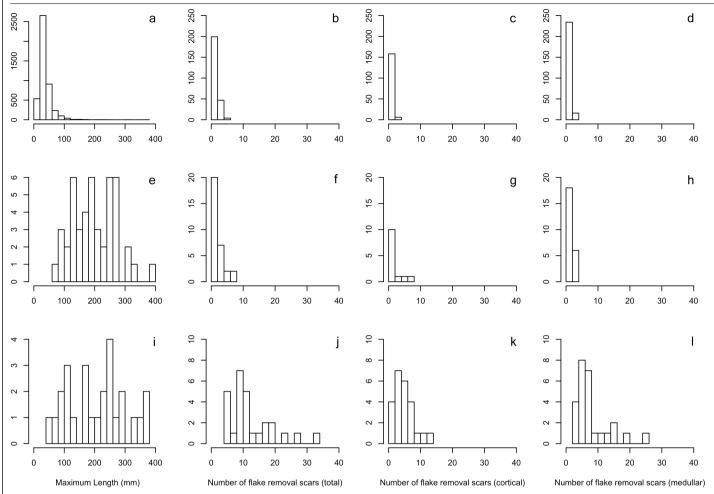


Extended Data Fig. 2 | Distribution map of the T69 Complex bone tool horizon. a, Plan view. b, East-West cross section within a 1-metre-wide area between (1) and (2). c, North-South cross section within a 1-metre-wide area between (3) and (4). See also Supplementary Video 2.



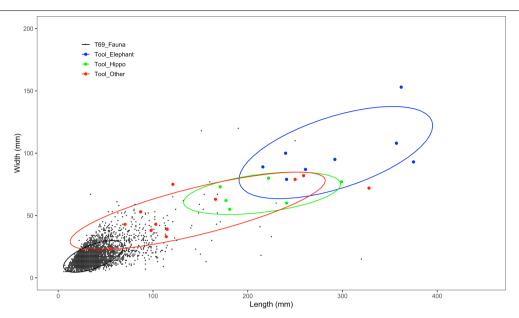
histograms and stereographic projections (2) of lithics and fossils from the T69 Complex bone tool horizon. 1) Observed Mean Distance (OMD) and Expected Mean Distance (EMD) are expressed in metres. *P*-values of complete, lithic and bone assemblages = 0.000. *P*-value of bone tool distribution =

0.6785.2) Orientation of items with elongation index >1.6% calculated as azimuthal (a) and axial data (b). c, Stereographic projection of azimuth and dip values. d, Vollmer's trivariate fabric shape triangle% e, Benn's fabric shape triangle.



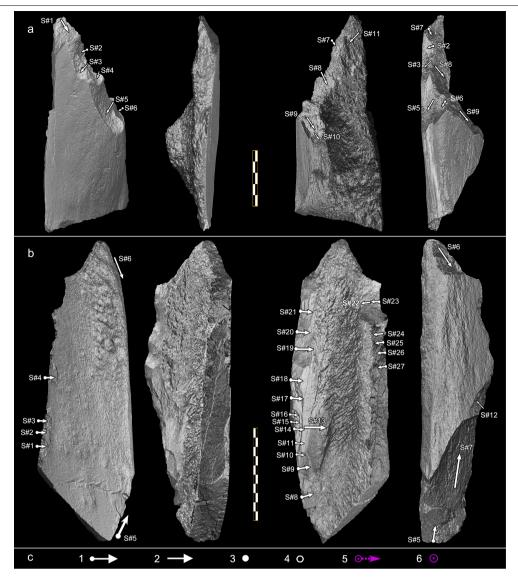
Extended Data Fig. 4 | Metric comparison (a,e,j) and number of flake removal scars (b-d, f-h,j-l) per specimens. Comparison of the maximum length between a, the bone fragments from the T69 Complex faunal assemblage, e, the diaphyseal fragment produced when fracturing elephant limb bones⁴, and i, the bone fragments bearing flake removal scars interpreted as bone tools from the T69 Complex. The tools are considerably longer than the remainder of the faunal assemblage and fall within the size range of experimentally broken elephant long bones. Comparison of the number of flake removal scars per specimen between (b-d), a randomly selected sample of diaphyseal fragments from the T69 Complex faunal assemblage, (f-h) bone fragments produced during experimental bone

fracturing to expose the marrow of horse 9 and elephant 4 limb bones, and (j-l) bone fragments bearing flake removal scars interpreted as tools from the T69 Complex. The total number of flake removal scars rarely exceed four in experimental settings (f) whether they are present on the cortical (g) or medullar aspect (h), which broadly matches the pattern observed for the comparative sample from the T69 faunal assemblage (f-h). Conversely, the bone tools from the T69 Complex display instead an average of 12.9 flake scars per specimen (j), preferentially arranged contiguously on both the cortical (k) and medullar (l) aspects.



Extended Data Fig. 5 | **Size comparison between the bone tools and the faunal assemblage from the T69 Complex.** Scatterplot of the length and width of the bone fragments from the T69 Complex faunal assemblage (black) compared to those recorded for the bone tools made of elephant bones (blue), hippo bones (green), and bones from other or unidentified species (red). Part

of that latter sample falls within the range of variability of both elephant and hippo as their cortical thickness suggests they originate from size 5 or 6 mammals (i.e., mammals weighting more than two tonnes). Confidence ellipses at 90%.



Extended Data Fig. 6 | **3D surface model images of bone tools.** Illustrated in Fig. 1. **a**, Bifacially-flaked tool (accession number T69L20-3009; dimensions $166 \times 63 \times 44$ mm). The left part (cortical aspect) is fractured, almost certainly after the distal end had already been shaped (as indicated by the truncated morphology of S#1). Order of flaking can be reconstructed in part (S#5 is offset by posterior S#6 and S#4; S#3 is partially erased by S#2), and indicates a recurrent sequence of shaping of the edges near the tip. **b**, Tool made of a proboscidean long bone fragment (accession number T79L10-2511; dimensions $292 \times 95 \times 66$ mm). The scars on the medullar aspect are organized in two series of contiguous removals: the left edge series are struck from the cortical side and penetrate invasively into the cancellous tissue. In the right edge, S#24-27 were removed from a pre-existing fracture plane, thus indicating that the

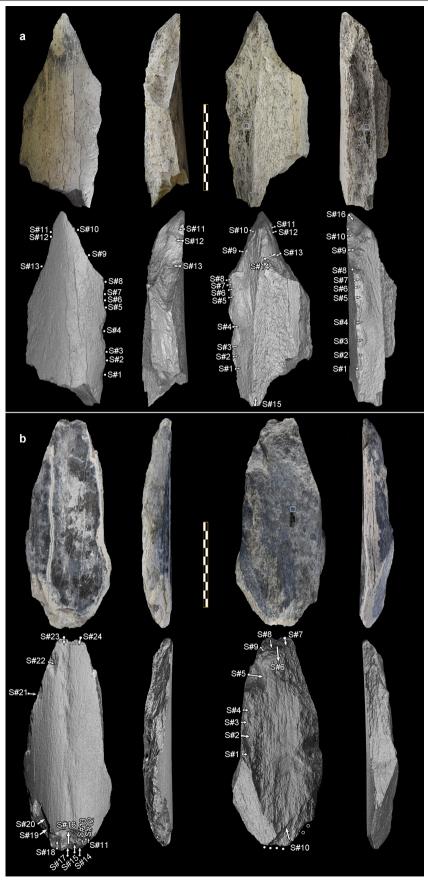
shaping took place after the bone shaft had been split. The distal end was fractured after shaping had occurred, as part of S#22 was removed by the tip fracture, and the edge where S#21 is located was also truncated by the fracture. The V-shaped angle of such fracture runs parallel to the tool axis, which is compatible with a percussive and compressive motion likely causing the tip breakage. Each scale bar segment, 1 cm. See also Supplementary Videos 4 and 5. **c**, Symbology (based on 101): 1-Direction of flaking on scar with percussion point. 2-Direction of flaking on scar where percussion point is missing. 3- Percussion point of removal on the adjacent flaking surface. 4- Position of removal with absent percussion point on the adjacent flaking surface. 5- Striking direction on undetached flakes. 6- Percussion point of undetached flake on the adjacent flaking surface.



$Illustrated in Fig.\,3.\,These\,artefacts\,share\,similar\,techno-morphological$ features, in which the portion of long limb diaphysis closer to the epiphysis

(where bones are thicker) is shaped into the tip of the tool, the base of the artefact is shaped in the portion near the metaphysis (thinner and with a more oval natural shape), and one lateral edge in the medial portion of the shaft is shaped into a deep notch. a, Massive tool made from the humerus of an elephant (accession number T79L10-9047; dimensions 375 × 93 × 105 mm). Series 1 (S#1-4): flake removals on the cortical aspect of the tip. Series 2 (S#5-8): sequence of alternating removals that indicates consecutive flipping of the blank. Series 3 (S#9-10): the cortical surface is used to strike flakes that are removed sequentially and clockwise. This series is truncated by (and hence is anterior to) the series starting with S#11. Series 4 (S#11-19): shaping of a $crescent\ bottom\ at\ the\ base\ of\ the\ tool.\ Instead\ of\ wide\ and\ invasive\ removals$

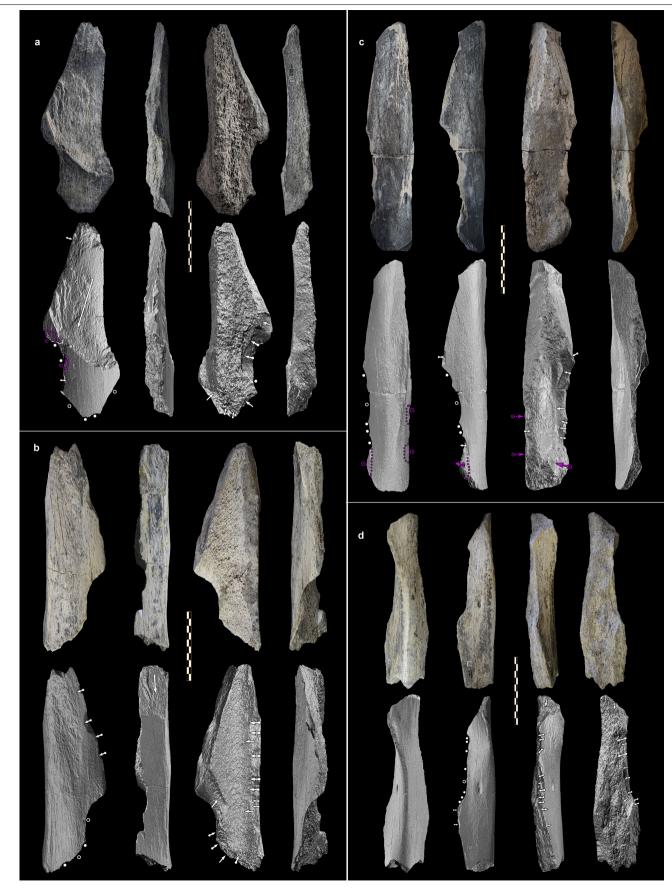
such as those from the previous series, scars in Series 4 trim the edge to create a $semicircular \, shape. \, Series \, 5 \, (S\#20\text{-}29): unifacial \, removals \, from \, the \, cortical$ surface into the medullar side remove cancellous tissue and create a denticulate edge. Series 6 (S#30-34): a deep notch with several extractions within is created in the medial part of the edge. **b**, Unifacially-shaped bone tool (accession number T79L10-18461; dimensions 299 × 77 × 54 mm). Removals S#8-12 use scar S#1 as their percussion platform. Most of flaking is concentrated on one edge, where deep and invasive scars (S#3 and S#7) were succeeded by a contiguous sequence of flake removals aimed at creating a large notch (S#8-12) and shape the tip (S#13-16). Two V-shape scars (S#5-6) forming a dihedral angle at the tip run parallel to the tool axis are interpreted as fractures probably caused by percussive and compressive motions during tool use. Each scale bar segment, 1 cm. See additional details in Supplementary Videos 6 and 7.



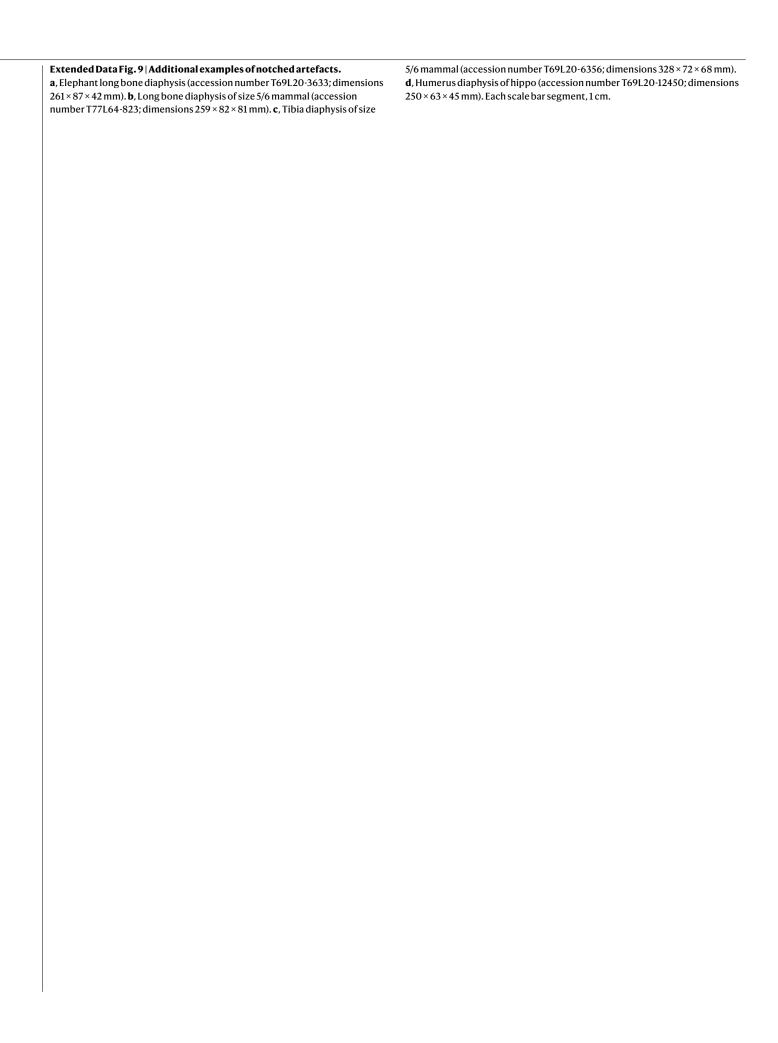
 $\textbf{Extended Data Fig. 8} | See \ next \ page \ for \ caption.$

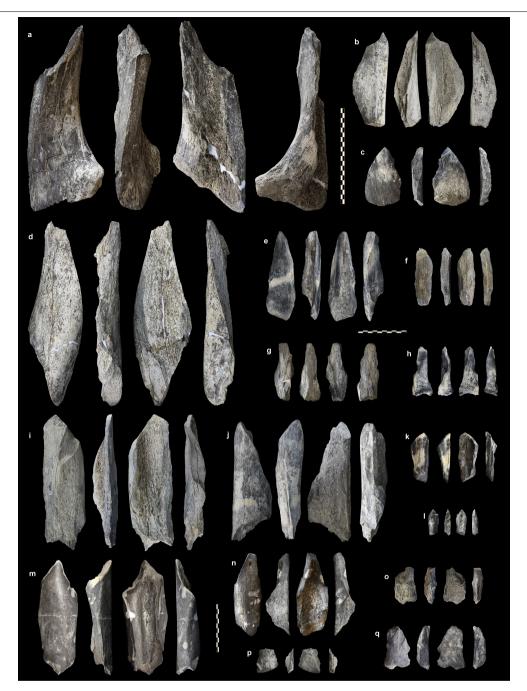
 $\label{lem:extended} \textbf{ Data Fig. 8} \ | \ \textbf{ Additional examples of elephant bone artefacts. a,} \\ \textbf{ Unifacially-flaked bone tool (accession number T69L20-13007; dimensions 216} \\ \times 89 \times 60 \ \text{mm}) \ \text{on an indeterminable limb bone diaphyseal fragment. Bilateral removals on the distal part - S#9-10 on the right and S#11-13 on the left sides (cortical view) - shape a pointed end. Flake removals S#1-13 were struck from the cortical surface into pre-existing large fracture planes, and therefore occurred after the initial splitting of the diaphysis. Percussion points of most scars penetrate deep into the edge, producing large notches (e.g., S#1, S#4,$

S#6). **b**, Bifacially-flaked tool made on indeterminable diaphyseal fragment of limb bone (accession number T79L10-13631; dimensions $240 \times 100 \times 40$ mm). Most scars on the medullar surface (S#1-5) used a pre-existing large fracture plane as the striking platform. Scar S#10 removes part of another pre-existing fracture plane, and is in turn largely obliterated by the S#11-20 flaking series. Removals on the proximal end (S#10-20) shape a crescent via recurrent flaking of the same area. Each scale bar segment, 1 cm.

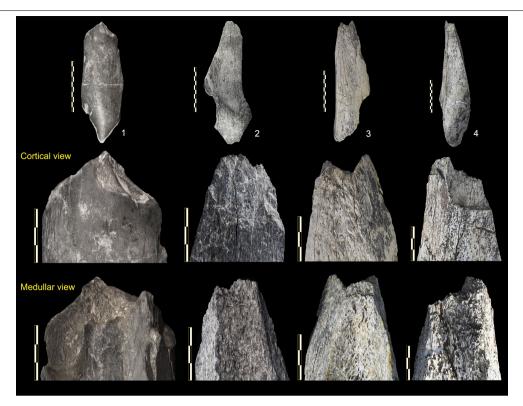


 $\textbf{Extended Data Fig. 9} | See \ next \ page \ for \ caption.$





 $\textbf{Extended Data Fig. 10} | \textbf{Remaining specimens in the T69 Complex bone tool assemblage.} \\ \textbf{See accession numbers, skeletal, taxonomic and dimension data in Supplementary Data 2, and annotated flaking patterns in Supplementary Video 8. Each scale bar segment, 1 cm.} \\$



Extended Data Fig. 11 | Recurrent distal fracture patterns on additional bone tools. 1, Tibia diaphysis of cf. *Hippopotamus* (accession number T69L20-1872; dimensions $222 \times 80 \times 54$ mm). 2, Accession number T69L20-3633

 $(see \, Extended \, Data \, Fig. \, 9a). \, 3, Accession \, number \, T77L64-823 \, (see \, Extended \, Data \, Fig. \, 9b). \, 4, Elephant \, long \, bone \, diaphysis \, (accession \, number \, T78L84-75; \, dimensions \, 353 \times 103 \times 53 \, mm). \, Each \, scale \, bar \, segment, 1 \, cm.$

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All data generated or analysed during this study are included in this published article and its supplementary information files.

- A description of any restrictions on data availability

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| For Bayes | ian analysis, information on the choice of priors and Markov chain Monte Carlo settings | | | | | | |
| For hierar | For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes | | | | | | |
| Estimates | of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated | | | | | | |
| ' | Our web collection on <u>statistics for biologists</u> contains articles on many of the points above. | | | | | | |
| Software an | d code | | | | | | |
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| Data collection | ArcGIS Pro v. 3.3, R-Cran v. 4.4.1, Oriana v. 3.13 | | | | | | |
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| Sample size | This archaeolog tools larger tha zooarchaeologi among the bon The archaeolog archaeological | one tool assemblage reported here contains 27 artefacts, derived from an archaeological horizon excavated across 295 square meters. archaeological horizon also yielded abundant fossils (over 22,800 remains) from mammals, reptiles and fish, as well as over 10,900 stone larger than 2 cm. Mammal bones ≥2 cm (n = 8,930) were sampled for full zooarchaeological analysis. The sample selected for full rechaeological analysis (n = 2,075) included all items identified as bone tools, all proboscidian fossils (as these are particularly significant and the bone assemblage), and a sample from all other ≥2 cm mammal bone specimens. Inchaeological bone tools were compared to: 1- Dimensions of a control sample (n = 4,507) of long bone fragments from the same declogical excavation. 2- Features of flake removals in a sample (n = 250) of long bone fragments. 3-Published experimental samples of the Coopen et al., ref 9) and elephant (Backwell and d'Errico, ref. 4) long bones fractured to expose marrow. | |
| fossils. This selection included all items identified as bone tools plus all | | e size of the mammal assemblage, full zooarchaeological analysis was conducted in a sample of the determinable mammal ection included all items identified as bone tools plus all proboscidean fossils, as well as a sample from all other determinable the comparison of dimensions and flaking patterns between bone tools and the rest of the mammal assemblage from the T69 trol sample of the latter was selected (see details in Methods). | |
| Replication | All bones tools were first-hand analysed by two pairs of co-authors (IdIT-JKN and LD-Fd'E), and then each artefact was discussed in group meetings during two separate sessions of several days each, approximately 1 year apart. Results are available in Supplementary Information Dataset 2, which contains all data collected from each item interpreted as a bone tool. Each item is labelled with a unique ID that will enable reproducibility of data collection for any researcher re-studying the assemblage. | | |
| Randomization | Horizontal spatial distribution of all archaeological was tested through an Average Nearest Neighbour Analysis that compared the actual distance between items to a hypothetical random sample, following Ebdon (ref. 57). | | |
| Blinding | | e independently analysed by IdIT and LD-Fd'E, after which specimen features were discussed as a group and recorded in | |

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| Palaeontology and archaeology | | MRI-based neuroimaging | |
| Animals and other organisms | | | |
| Clinical data | | | |
| Dual use research o | of concern | | |
| | | | |
| ' | | | |
| Palaeontology an | d Archaeology | | |
| | | | |
| Specimen provenance | All specimens are derived from excavations directed by at the FLK West T69 Complex in Olduvai Gorge. Fieldwork permits were renewed annually by the Tanzanian Commission for Science and Technology (COSTECH), the Ngorongoro Conservation Authority Area (NCAA), and the Department of Antiquities, Tanzanian Ministry of Natural Resources and Tourism, and an export permit for analysis was granted by the Department of Antiquities. | | |
| The state of the s | | nanently housed at the Olduvai Research Laboratory in the Leakey Camp, a research and storage facility Ministry of Natural Resources and Tourism. | |
| Dating methods No new dates are provided | | | |
| Tick this box to confi | rm that the raw and calibra | ated dates are available in the paper or in Supplementary Information. | |
| Ethics oversight Fieldwork and analysis prot Ministry of Natural Resource | | ocols abided by guidance provided by COSTECH, NCAA, and the Department of Antiquities, Tanzanian es and Tourism. | |
| Note that full information on | the approval of the study prot | tocol must also be provided in the manuscript. | |
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was applied. Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosiacism, off-target gene editing) were examined.