## STAR METHODS

### KEY RESOURCES TABLE

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CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, David Reich (reich@genetics.med.harvard.edu).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

We generated new genome-wide data from skeletal remains of 15 prehistoric individuals: 5 from eastern Africa, 7 from south-central Africa, and 3 from southern Africa (Tables 1, S1, and S2). One of these individuals, from St. Helena Bay and directly dated...
to ~2100 BP, previously yielded a complete mitochondrial genome (Morris et al., 2014). We directly dated a second South African individual buried in a hunter-gatherer context from Faroaskop to ~2000 BP, and a third individual buried in a pastoralist context from Kasteelberg to ~1,200 BP. We also directly dated and used in-solution enrichment to obtain genome-wide DNA from four individuals from coastal eastern Africa: From the cave site of Panga ya Saidi in the coastal region of southeastern Kenya (~400 BP), Kuumbi Cave in the southeast of Zanzibar Island (Tanzania; ~1,400 BP), and Makangale Cave in the northwest of Pemba Island (Tanzania; ~1,400 BP and ~600 BP). We also obtained genome-wide data from a ~3100 BP individual from a pastoralist context in north-central Tanzania, and ~8100–2500 year old individuals from Malawi.

**Terminology**

There is no widely accepted term with neutral connotations for indigenous communities in southern Africa (Schlebusch, 2010). In this manuscript, we follow San council recommendations in using population-specific terms whenever possible, and alternatively the terms San for Tuu and K’xaa speaking hunter-gatherer groups and Khoe for Khoe-khoe speakers. When necessary we collectively refer to groups with southern Africa-specific ancestry as Khoe-San, or as having San-related ancestry.

**Panga ya Saidi Cave, Kilifi County, Kenya (n = 1)**

Panga ya Saidi is a large limestone cave complex formed within an escarpment c. 15 km from the Indian Ocean coast in southern Kenya. Excavated in multiple Sealinks Project campaigns, the cave’s long and complex depositional sequence spans discontinuously, more than 76,000 years, and contains mainly Later Stone Age (LSA) deposits, overlain by Middle Iron Age (MIA) and Later Iron Age (LIA) deposits dating to the last two thousand years (Helm et al., 2012). The sampled specimen (I0595, Kenya_400BP) is a phalanx recovered from an in situ burial (context 403) and directly AMS radiocarbon dated to 496-322 BP (388 ± 27 uncalibrated BP, OxA-30803). The individual was a tall, robust young adult male. He was buried in a shallow grave in a crouched position with two hands and one foot in the small of the back and the skull disarticulated and placed by the knees. The individual was buried by sediment containing marine shell beads, small knapped stone tools, and Tana Tradition potsherds. The associated faunal remains are exclusively wild, with the exception of a single possible caprine bone. Large numbers of remains of birds, rodents, and other microfauna suggest that the cave may have only been sporadically occupied when the human remains were deposited. We infer from the material culture and fauna that the cave was occupied by foragers during the time the individual was buried, although food producers were present at nearby settlements such as Mtsengo and Mbuyuni (Helm, 2000). Crop remains of African sorghum, pearl millet and finger millet found at the site suggest these foragers had access to agricultural resources.

**Makangale Cave, Pemba, Tanzania (n = 2)**

This limestone cave at the northern end of Pemba Island in the Zanzibar archipelago has been excavated in multiple campaigns, the most recent two seasons conducted by the ERC-funded Sealinks Project at Oxford University in 2012 and then the Max Planck Institute for the Science of Human History in 2016 (unpublished; see also Chami et al., 2009)). The sequence shows clear evidence of human occupation beginning around 1400 BP with an escargotière layer of giant African land snail shells, pottery, and disarticulated human remains. Above this layer, the sequence shows regular human use of the cave into the last thousand years. The first sampled specimen (I0589, Tanzania_Pemba_1400BP) is a sacral vertebra from context 204 (Sealinks Project faunal catalog no. 15336), directly dated to 1421-1307 BP (1520 ± 30 uncalibrated BP, Beta-434912). The second specimen (I2298, Tanzania_Pemba_600BP) is a lower molar from context 301 (Sealinks Project faunal catalog no. 15624), which lies just below the surface and was dated to 639-544 BP (623 ± 20 uncalibrated BP, Wk-43308). Both specimens are associated with a highly unusual faunal assemblage, dominated by fragmented crocodile (Crocodylus cf. niloticus) remains and diverse microfauna, including Rattus rattus (Asian black rat), a nonnative rodent that must have arrived to the area via maritime exchange routes. There are no taphonomic indicators in the faunal assemblage of hunting by humans, nor of crocodile predation on humans. During both of the occupational phases targeted in this study, there were nearby settlements occupied by farmers whose ancestors likely came from the mainland, for example at the sites of Tumbe (c. 1400-1000 BP) and Chwaka (1000-400 BP) (Fleisher and LaViolette, 2013).

**Kuumbi Cave, Zanzibar, Tanzania (n = 1)**

Kuumbi Cave is a limestone solutional cave excavated in multiple campaigns (Chami, 2009; Sinclair et al., 2006). Sealinks Project excavations in 2012 documented a complex depositional sequence stretching discontinuously over 20,000 years, containing evidence of LSA and MIA occupations in five discernible phases (Shipton et al., 2016). The analyzed specimen (I0589, Tanzania_Pemba_1400BP) is a complete second phalanx of an adult (Sealinks Project faunal catalog no. 4353). It was recovered from context 403 and directly AMS radiocarbon dated to 496-322 BP (388 ± 27 uncalibrated BP, OxA-30803). The second specimen (I2298, Tanzania_Pemba_600BP) is a lower molar from context 301 (Sealinks Project faunal catalog no. 15624), which lies just below the surface and was dated to 639-544 BP (623 ± 20 uncalibrated BP, Wk-43308). Both specimens are associated with a highly unusual faunal assemblage, dominated by fragmented crocodile (Crocodylus cf. niloticus) remains and diverse microfauna, including Rattus rattus (Asian black rat), a nonnative rodent that must have arrived to the area via maritime exchange routes. There are no taphonomic indicators in the faunal assemblage of hunting by humans, nor of crocodile predation on humans. During both of the occupational phases targeted in this study, there were nearby settlements occupied by farmers whose ancestors likely came from the mainland, for example at the sites of Tumbe (c. 1400-1000 BP) and Chwaka (1000-400 BP) (Fleisher and LaViolette, 2013).
Luxmanda, Babati District, Tanzania (n = 1)
Luxmanda is an open-air settlement sitting atop the Rift Valley escarpment (1878 m above sea level) at the southern edge of the Mbulu Plateau, just north of Lake Balangida and Mount Hanang. Excavations in 2012, 2013, and 2015 by the RAPT project (Research on the Archaeology of Pastoralism in Tanzania) have shown Luxmanda to be the largest and southernmost known settlement site of the Pastoral Neolithic (PN), the era corresponding to Spread of mobile livestock herding in eastern Africa (Prendergast et al., 2013) (K.M.G. and M.E.P., unpublished data). Despite its isolated location, Luxmanda shows strong material culture affinities to sites of southern Kenya classified as Savanna Pastoral Neolithic (SPN), in particular the Narosura type-site (Odner, 1972); Luxmanda’s ties to SPN sites are further supported by sourcing of obsidian stone tools to the Naivasha Basin in the Central Rift Valley of Kenya. Faunal remains from Luxmanda indicate a diet almost exclusively focused on sheep, goat, and cattle; botanical remains are currently under study. A suite of eleven radiocarbon dates provides a tightly constrained window of occupation c. 3000-2900 BP, which is at the early end of the range for SPN sites. The analyzed specimen (Tanzania_Luxmanda_3100BP) is a petrous bone from a perinatal infant. The infant was found complete and buried just to the west of, and c. 35 cm below, a burnt earth feature, interpreted as a hearth. The burnt earth feature was then overlain by domestic refuse. Collagen from the same petrous bone was AMS radiocarbon dated to 3141-2890 BP (2925 ± 20 uncalibrated BP, ISGS-A3806), a date nearly identical to those of charcoal samples taken from the overlying burned earth feature and domestic refuse.

Hora, Malawi (n = 2)
The Hora 1 and Hora 2 skeletons from Malawi were excavated from the Hora 1 site in the Mzimba District of Malawi, located on the northeastern side of Mount Hora. Mount Hora lies south of the Nyika Plateau and west of the Viphya Mountains, where miombo vegetation grades southwest into edaphic grasslands (DeBusk, 1997). The region divides the Luangwa River Basin in Zambia from Lake Malawi ~130km to the east, which receives all water from the district via the South Rukuru River (Figure S1). The South Rukuru flows south-to-north along the Zambian border before turning east to intersect the Kasitu, a major interior waterway that flows past Mount Malawi. The Hora 1 and Hora 2 skeletons contain paintings that include white stars and abstract red ''gridirons'' or ''nets,'' both of which are motifs replicated at other sites on Mount Hora and nearby localities (Clark, 1956; Cole-King, 1973). Hora 1 was excavated by Clark in 1950, and produced a 2.2 m cultural sequence containing pottery and iron slag at the top, faunal remains and LSA lithic assemblages with mollusk shell beads below this, and two human burials approximately 76cm below the surface (Clark, 1956). The first burial to be revealed was Hora 1 (UCT-242), a short-statured male in his thirties or forties at death, who may have been buried with a flaked stone axe (Morris and Ribot, 2006). The skeleton was left partially in situ during the original excavation and later exhumed by Rangeley (Clark, 1956). About 3 m to the south, the skeleton of a female was recovered (Hora 2, UCT-243); she was of similar stature to the male and died in her early twenties (Morris and Ribot, 2006). The female was found in a flexed position on her left side, and most of the bones of the hands and feet were missing or had been displaced – suggesting a degree of exposure prior to burial (Clark, 1956).

A previous study (Clark, 1956) reports three major stratigraphic units at Hora 1, and places the burials in the upper part of the second unit in association with the “Nachikufan II” – an industry with a type site ~300 km to the east in Zambia. Clark does not consider the burials intrusive, noting that they are overlain exclusively by hunter-gatherer material culture. The earliest occupation layer at the site is “reddish brown earth” that begins at about 1.7 m depth, and contains a lithic assemblage assigned by Clark to the Nachikufu I. On the basis of typology, the earliest deposits at the site were inferred to date to between 16000-11000 BP, the burials to between 10000-5000 BP, and materials in the uppermost unit are likely as recent as the last few hundred years (Miller, 1971). This is supported by a direct AMS age on the Hora 2 skeleton of 8173-7957 BP (7230±60 uncalibrated BP, PSUAMS-2535). Although insufficient collagen was present in the Hora 1 skeleton for a direct age, its archaeological context and genetic similarity to the directly-dated Hora 2 skeleton place it most probably in the early Holocene. New excavations at the Hora 1 site were conducted in 2016 and 2017 within 2-5m of the original burial locations (based on archival photographs), achieving a maximum depth of 1m. A sample of 6842 plotted objects recovered from 3m3 confirmed that below ~30cm depth there is no Iron Age or historical material culture (e.g. slag, pottery, or glass), further establishing that the two skeletons are from a resident hunter-gatherer population.

Fingira, Malawi (n = 3)
The three Fingira samples derived from ex situ human remains (2 adult femora and one subadult femur) recovered in 2016 from Fingira Rock, a large shelter located within the boundaries of Nyika National Park in northern Malawi. The climate is cool and moist compared to the surrounding regions (~1,600 mm annual rainfall, and average daily highs 10 – 20°C). Fingira Rock is an isolated inselberg at ~2100 m above sea level, near the upper limit of the miombo woodland. Within this inselberg is set a single deep rock shelter with 160 m² of deposit (Figure S1). First excavated in 1966, Fingira yielded fragmentary remains of at least 15 human individuals (subadults and adults), plus one more complete burial near the front, in association with rich lithic, archaeofaunal, and archaeobotanical assemblages. All human remains previously recovered from Fingira were studied by Brothwell, and reported in in a previous study (Sandelskow, 1972). They are curated at the Natural History Museum in London.

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In addition to human remains, the Fingira deposits contained bone tools, pigments, and ornaments (Robinson and Sandelowsky, 1968; Sandelowsky, 1972). Two conventional $^{14}$C ages obtained near the base of the ca. 50cm-deep excavation were returned in stratigraphic sequence as 3,260 ± 80 BP and 3,430 ± 80 BP (Sandelowsky 1972). Geometric rock paintings at the site exhibit white overpainting on red, which has been interpreted elsewhere in Malawi as re-use by food producers (Smith, 1985; Zubieta, 2016). Ceramics are rare but present at the site, where the large deposit exhibits predominately Later Stone Age material culture. This was confirmed during 2016 pilot excavations, permitted by the Departments of Antiquities and National Parks and Wildlife.

The deposits at Fingira had not been backfilled after the 1966 excavation, and had undergone extensive erosion and slumping. As the site is accessible to the public, piles of materials had been pulled from the section and placed on fallen rocks. Two of those specimens comprised the two adult femoral specimens reported here. Using original site plans, we identified the extent of the 1966 excavations prior to slumping, including the relative positions of originally-recovered human remains. A large part of this area had been covered with a central termite mound, and it was within this that the partial skeleton of a neonate was recovered. This comprised the subadult sample from Fingira. Direct $^{14}$C AMS ages on these three specimens are reported in Table S2, and show that LSA people were using the site as a cemetery for at least 3,700 years, from 6177-5923 BP (5290±25 uncalibrated BP, UCIAMS-186347) to 2676-2330 BP [2425±20 uncalibrated BP, PSUAMS-1734], 2483-2330 BP (2400±20 uncalibrated BP, PSUAMS-1881)]. The two adult specimens therefore significantly pre-date the earliest known age in Malawi for the Bantu expansion, which derives from the Kasitu Valley (containing Hora Mountain) at 1,750 ± 60 BP (Robinson, 1982). If the Bantu expansion into Malawi began more than ~700 years before what is currently known, then the neonate from Fingira could potentially overlap in time with it. However, the overall antiquity of these specimens makes admixture attributable to the Bantu expansion highly unlikely in light of current knowledge.

**Chencherere II, Malawi (n = 2)**

Mwana wa Chencherere II is a painted rock shelter set in a granitic inselberg at ca. 1,700 m above sea level in the Chongoni Rock Art Region (Smith, 1995). Its relatively high altitude results in cool, moist year-round conditions between 10–20°C. It was excavated by Clark in 1972 (Clark, 1972, 1973), and the faunal assemblage published in detail by Crader (Crader, 1984a), who also reports one adult male burial and the fragmentary remains of seven other individuals (adult and subadult). The site contained large lithic and faunal assemblages, bone and shell tools and ornaments, and increasing abundances of pottery and other evidence of interaction with food-producers over time. The youngest reported date from the site is a conventional $^{14}$C age on charcoal from the top of Level 3, at 800 ± 50 BP. The oldest date is from near the base of Level 4, and is 2,480 ± 200 BP.

All human remains are reported from Levels 2, 3, and 4 – with most in Level 3. The original description of human remains was by Brothwell, and Crader (1984b:Appendix 2) later listed several more individuals that had been discovered within the faunal material. All human remains recovered from Chencherere II were thought to be held at the Natural History Museum in London, but during a 2016 visit to the Malawi National Repository in Nguludi (near Blantyre), six additional specimens were identified. Five of these derived from a cluster in square A3 (Figure S1): right ilium, left femur (sampled), and 3 partial ribs. These were inferred to belong to the same individual, a subadult aged 3-5. An upper right incisor (sampled) was labeled as deriving from square E2, and therefore although of similar ontogenetic age it was deemed likely to be from a different individual.

Although the genetic analysis confirms that these are two different individuals, insufficient material remained from the incisor root of the second individual for a direct age. The first individual returned a direct $^{14}$C AMS age of 5293-4979 BP (4525 ± 25 uncalibrated BP, UCIAMS-186348). As with Fingira, the direct AMS ages on the human remains are substantially older than the conventional charcoal ages suggested, indicating either intrusive charcoal or problems with the original dates.

**St. Helena Bay, South Africa (n = 1)**

In June 2010, an intact skeleton was excavated by Andrew B Smith along the southwest coastal region of South Africa at St. Helena Bay. The skeleton is stored in the Department of Human Biology at the University of Cape Town under the accession number UCT-606. The body had been placed on an impermeable consolidated dune surface, on its right side in a fully flexed position. The bones originate from a single male who stood no more than 1.5 m in height. Dental wear and significant areas of osteoarthritis suggest that he was at least 50 years of age at time of death. Lack of any evidence of tooth decay and excessive occlusal wear suggests a diet typical of hunter-gatherer subsistence. The presence of abnormal bone growths in the right auditory meatus (ear canal opening) caused a condition known as “surfer’s ear” (auditory exostosis) and provides evidence that this individual most likely spent considerable time in the cold coastal waters sourcing food. No obvious cause of death was evident. The results of carbon-14 to stable carbon-13 isotope ratio analysis of a rib provided a date of 2241-1965 BP (2330 ± 20 uncalibrated BP, UGAMS-7255) years before present (UGAMS-7255) with a $d_{13}C$ value of −14.6%. Although the minimum date falls right on the edge of the arrival of pastoralism in the Western Cape, anatomical and archaeological analysis of this skeleton and the associated buried site clearly defines this individual as an indigenous Southern African, predating pastoral arrival into the region. This individual has previously been sampled for mtDNA analysis (Morris et al., 2014).

**Faraoskop, South Africa (n = 1)**

The site of Faraoskop is a rock shelter about 30 km inland from Elands Bay on the west coast of the Western Cape Province of South Africa. The shelter is on the highest ridge of a small hill at an altitude of 300 m above the surrounding plain. Seven skeletons were collected by a local farmer in 1984, but the site was subsequently excavated under controlled conditions in 1987 and 1988 (Manhire,
and five more skeletons were collected including the one referred to here as UCT-386. The shelter has no rock paintings but there is a rich assemblage of Later Stone Age artifacts including stone tools, ostrich eggshell beads, worked marine shell, leather and twine (Manhire, 1993). No pottery is associated with the site. Six skeletons have been C14 dated with resulting dates ranging from 2300 years BP to 1900 years BP (Manhire, 1993; Sealy et al., 1992) but all the dates overlap at the 2σ standard deviation. UCT-386 is the skeleton of a female about 40-50 years at death. A bone sample provided a date of 2017-1748 BP (2000 ± 50 uncalibrated BP, Pta-5283) with a d13C value of −16.8% (Manhire, 1993). A recent re-evaluation of the site indicates the possibility that all of the individuals died in one event. Not only do all of the dates overlap, but the excavation data suggest no separate grave shafts and at least two of individuals show signs of perimortem injury and violent death (Parkington and Dlamini, 2015). The Faraoskop human skeletons are stored in the Department of Human Biology at the University of Cape Town.

Kasteelberg, South Africa (n = 1)
The site of Kasteelberg is on a granite hill on the Vredenberg Peninsula about 4 km from the coastal village of Paternoster, approximately 150 km north of Cape Town (Smith, 1992a). There are several sites on the hill including a small rockshelter, but the human skeleton was excavated from square 22 extension at KBB on the eastern side of the base of the hill. UCT-437 is the nearly complete skeleton of a child of about 4 years of age excavated by Lita Webley in 1986. The body was in a shallow pit about 1.5 m deep and with no stone cover. The specimen has a date of 1282-1069 BP (1310 ± 50 uncalibrated BP, Pta-4373). There were no grave goods in association with the skeleton. The earliest sites on top of the hill provide evidence of domestic sheep at around 2100 years ago. The KBB site at the base of the hill is dated to the latter part of the occupation but present the first appearance of cattle in the region (Smith, 1992b). Overall, the Later Stone Age occupation of the Kasteelberg indicates the presence of herder-foragers who practiced seasonal economic systems, sometimes relying on domestic stock while at other times hunting seals (Sadr et al., 2003). The human skeleton from Kasteelberg is stored in the Department of Human Biology at the University of Cape Town.

METHOD DETAILS

Direct AMS 14C Bone Dates
We report new direct AMS 14C bone dates in this study from multiple laboratories. In general, bone samples were manually cleaned and demineralized in weak HCl and, in most cases (PSU, UCIAMS, OxA, ISGS), soaked in an alkali bath (NaOH) at room temperature to remove contaminating soil humates. Samples were then rinsed to neutrality in Nanopure H2O and gelatinized in HCL (Longin, 1971). The resulting gelatin was lyophilized and weighed to determine percent yield as a measure of collagen preservation (% crude gelatin yield).

Collagen was then directly AMS 14C dated (ISGS, Pta) or further purified using ultrafiltration (PSUAMS, UCIAMS, OxA, Wk, Beta) (Brown et al., 1988; Kennett et al., 2017) or a modified XAD method (Lohse et al., 2014; Stafford et al., 1991). It is standard in some laboratories (PSUAMS, UCIAMS, Wk, OxA) to use stable carbon and nitrogen isotopes as an additional quality control measure. For these samples, the %C, %N and C:N ratios were evaluated before AMS 14C dating. C/N ratios for well-preserved samples fall between 2.9 and 3.6, indicating good collagen preservation (Van Klinken, 1999). Additional quality control work was carried out on the samples from Malawi using FTIR spectra (Figure 1E).

All 14C ages were δ13C-corrected for mass dependent fractionation with measured δ13C/12C values (Stuiver and Polach, 1977) and calibrated with OxCal version 4.3 using the southern hemisphere calibration curve (SHCal13). Given their proximity to the equator, AMS 14C dates for sites in coastal Kenya and Tanzania were calibrated using OxCal v. 4.3 (Bronk Ramsey, 2009) at 95.4% probability employing a mixed curve that combines the SHCal13 (Hogg et al., 2013) and IntCal13 (Reimer et al., 2013) curves at ratios of 70:30 to account for the differential effects of the intertropical convergence zone.

Ancient DNA sample processing in Leipzig: St. Helena Bay sample

DNA extraction and library preparation
30.4 mg of bone powder was removed from the internal root canal of the tooth (SP2809) using a sterile dentistry drill in the clean room facilities of the Max Planck Institute for Evolutionary Anthropology in Leipzig, Germany. A DNA extract (E649) was prepared with a silica-based method, described in detail previously (Rohland and Hofreiter, 2007). 15 μL (15% of the total volume) of the extract was converted into a single-stranded DNA library (AS354) using a modified version of the single-stranded DNA library preparation protocol (Gansauge and Meyer, 2013; Korlević et al., 2015). Library positive and negative controls were carried throughout the library preparation process. Library AS354 was pre-treated with the USER enzyme, a mixture of uracil-DNA glycosylase (UDG) and endonuclease VIII, in order to remove uracils from the internal parts of ancient DNA molecules (Briggs et al., 2010; Meyer et al., 2012). The number of DNA molecules in the library (Table S1) was determined by digital droplet PCR (Bio-Rad OX200), using 1 μL of a 5,000-fold dilution of the library in EBT buffer (10 mM Tris-HCl pH 8.0, 0.05% Tween 20) as template in an Eva Green assay (Bio-Rad) with primers IS7 and IS8 (Meyer and Kircher, 2010). The library was amplified into the PCR plateau in a 100 μL reaction with AccuPrime Pfx DNA polymerase (Dabney and Meyer, 2012) using a pair of primers with two unique index sequences according to a double indexing scheme described in detail elsewhere (Kircher et al., 2011). 50 μL of amplification products were purified using the MinElute PCR Purification Kit (QIAGEN) and eluted in 30 μL TE buffer (10 mM Tris-HCl pH 8.0, 1 mM EDTA). The DNA concentration of the indexed library (AS369) was determined using a NanoDrop 1000 Spectrophotometer.
**Size fractionation for shotgun sequencing**

From the amplified library A5369, 1 μL was taken as template for a second round of amplification in a 100 μL PCR reaction using primers ISS and IS6 (Meyer and Kircher, 2010) with Herculase II Fusion DNA polymerase (Agilent Technologies) under the conditions described in detail previously (Dabney and Meyer, 2012). The concentration of the final library was determined on a Bioanalyzer 2100 instrument (Agilent Technologies) using a DNA-1000 chip. Library A5369 was pooled and sequenced with libraries from another experiment, occupying 33% of one lane of a flow cell on the Illumina HiSeq 2500 platform in rapid mode, using a double index configuration (2 × 76 + 2 × 76) (Kircher et al., 2011).

For a more effective use of sequencing capacity, 10 μL of the library A5369 was additionally separated on a Criterion Precast polyacrylamide gel (10% TBE, BioRad), and the fraction of library molecules with insert sizes larger than 40 bp was gel-excised as described in detail previously (Meyer et al., 2012). Gel-excised library molecules were subjected to a second round of amplification and the concentration of the final library (A5368) was determined using a DNA 1000 chip on the Bioanalyzer 2100.

**Ancient DNA sample processing in Tübingen: Faraoskop and Kasteelberg samples**

**Sampling and extraction**

Sampling took place in the clean room facilities of the Institute for Archaeological Sciences at the University of Tübingen. Both samples were irradiated with UV light for 10 min from all sides to remove surface contamination. The tooth from the South African forager from Faraoskop (UCT386) was sawed apart transversally at the border of crown and root, and dentine from inside the crown was sampled using a sterile dentistry drill, resulting in 56 mg dentine powder. For the femur fragment from the South African pastoralist from Kasteelberg (UCT437), the surface layer from the sampling area was removed with a dentistry drill prior to obtaining four aliquots sampled using a sterile dentistry drill, resulting in 56 mg dentine powder. For the femur fragment from the South African pastoralist from Faraoskop (UCT386) was sawed apart transversally at the border of crown and root, and dentine from inside the crown was sampled using a sterile dentistry drill, resulting in 56 mg dentine powder. For the femur fragment from the South African pastoralist from Kasteelberg (UCT437), the surface layer from the sampling area was removed with a dentistry drill prior to obtaining four aliquots between 51 and 80 mg of bone powder from the inside of the crown.

Extraction was performed following a protocol optimized for the recovery of small ancient DNA molecules (Dabney et al., 2013), resulting in 100 μL of DNA extract per sample. Three of the bone powder aliquots from UCT437 underwent a 10 min pre-digestion step after which the extraction buffer was removed (pre-digest) and replaced by fresh extraction buffer followed by over-night digestion (ON-digest), the powder from UCT386 and one aliquot of UCT437 were extracted without the pre-digestion step (full-digest). All eight resulting extracts were taken along for further library preparation. Negative controls were included in the extraction and taken along for all further processing steps.

**Screening**

Two double-indexed libraries were produced from an aliquot of 20 μL of the full-digest extractions of UCT386 and UCT437 (Kircher et al., 2011; Meyer and Kircher, 2010). Positive and negative controls were included in library preparation and taken along into sequencing. Libraries were enriched for human mitochondrial DNA (Marcic et al., 2010) and both enriched and shotgun libraries were sequenced on a HiSeq2500 with 2x101+8 cycles. Processing by the EAGER pipeline (Peltzer et al., 2016) and schmutzi (Renaud et al., 2013) resulted in an endogenous DNA content of 39% and 8% and an estimated mitochondrial contamination of 0%–2% and 1%–3% for UCT386 and UCT437, respectively.

**Library preparation for shotgun sequencing**

For UCT386 four more libraries were produced from an aliquot of 20ul of full-digest extract each, including a DNA repair step with UDG and endonuclease VIII to remove deaminated bases (Briggs and Heyn, 2012). For UCT437, six additional UDG-treated libraries were produced from 20ul each of the three pre-digest and the three ON-digest extracts. After indexing PCR (Kircher et al., 2011), aliquots of the UDG-treated libraries were size selected on a PAGE gel to remove fragments of sizes below 35 and above 80 bp (Meyer et al., 2012).

**Ancient DNA sample processing in Dublin: Malawi samples**

Sampling took place in ancient DNA-dedicated clean room facilities at University College Dublin. The petrous part of the temporal bone was selected for analysis from each individual from Hora, and other skeletal parts for the two other sites. Each complete petrous was UV irradiated for 10 min on each side prior to processing to reduce surface contamination. Any remaining sediment was removed using a Renfert Basic Classic Sandblaster (Renfert GmbH) at low power. Bone powder was retrieved from the petrous of UCT242 (Hora 1) by drilling a small hole on the superior surface of the petrous with a 4.8mm High Speed Cutter (Dremel) until the cochlea was accessible. Bone powder was then collected directly from the cochlea using a 3.2mm Tungsten Carbide Cutter (Dremel). The petrous from UCT243 (Hora 2) was cut from anterior to posterior using a Dremel drill at a location that transected the cochlea. The powder was collected directly from the cochlea using a 3.2mm Tungsten Carbide Cutter (Dremel). Powder aliquots from both samples were then UV irradiated for 5 min and placed in 2.0mL Eppendorf tubes.

**Ancient DNA sample processing in Boston: Tanzania samples, Kenya samples, and Malawi sample powder**

Sampling and DNA extraction

In a dedicated ancient DNA facility at Harvard Medical School, samples were UV-irradiated for 10 min in a UVP crosslinker. At the chosen part of each sample (root for teeth and compact part for bones) the surface was removed with a sanding disk. About 75 mg (±10 mg) of fine powder was obtained by drilling into the physically cleaned part with a sterile dentist drill bit and collected for DNA extraction (Table S1). In the case of KC-10-1011(4353) (I0589), additional bone powder was collected for a second DNA extraction attempt. The seven Malawi Holocene samples arrived in the Boston laboratory as powders prepared in Dublin, Ireland.
Starting from the sample powder, we followed the Dabney et al., 2013 extraction protocol for all samples, but replaced the funnel/MinElute assemblage with the pre-assembled Roche columns (Korlević et al., 2015) and eluted two times in 45 μL for a total of 90 μL DNA extract.

**Initial library preparation**

One initial barcoded library (L1) was prepared from 30 μL DNA extracts for all but two samples (Hora1 and Hora2), which were discolored and we reduced the volume to 3 μL (reducing the volume seems to mitigate library preparation inhibition, which we often find to be associated with discolored DNA extracts) following protocols published previously (Rohland et al., 2015) (Table S1). For three samples (I0589, I0595, I1048) the initial library was UDG-treated (Briggs et al., 2010) following a modification from Rohland et al., 2015 (partial UDG treatment) that is tailored to inefficiently remove terminal Uracils therefore leaving the aDNA authenticity signal in the terminal bases while efficiently removing miscoding damage within the molecules. The initial libraries for the other samples (I2298, I2966, I2967) were not UDG-treated. The last step of the library preparation, the amplification with universal primers, was set up in the cleanroom, but the thermal cycling happened in another laboratory physically separated from the cleanroom. The final products of our barcoded libraries cannot be sequenced right away; an additional PCR step is needed to finalize the adaptor sites. This is advantageous in that we can incorporate a second set of barcodes through dual indexing to differentiate two or more experiments done with the same barcoded library within the same sequencing run (see below).

**Screening**

Each initial library underwent screening that consisted, first, of shallow shotgun sequencing after an indexing PCR (that adds dual indices to each library, (Kircher et al., 2011), and second, target capture enrichment for mitochondrial DNA and a varying number of nuclear loci to assess mitochondrial haplogroup, mitochondrial contamination, aDNA authenticity and nuclear complexity (Meyer et al., 2014; Rohland et al., 2015). This experiment is finished by adding unique index combinations to each captured library, which is then subsequently pooled with the shotgun indexing PCR product for sequencing. Sequencing was done on an Illumina NextSeq500 with 2x76cycles + 2x7cycles.

We demultiplexed reads to be sample-specific requiring that the that the 7bp P5 and P7 indices matched (allowing one mismatch). Sample identification was further ensured by requiring that additional 7bp internal barcodes matched, again allowing one mismatch. We merged with a modified form of SeqPrep (https://github.com/jstjohn/SeqPrep) (the modification ensures that the highest quality base is retained in the overlap region), requiring at least 15bp overlap between forward and reverse reads, allowing one mismatch, retaining only reads of length greater than or equal to 30 base pairs, generating single ended reads.

Reads were then aligned using the same algorithm of BWA (version 0.6.1) (Li and Durbin, 2009) using ancient parameters to allow an increased mismatch rate, and to disable seeding (“-n 0.01 –o 2 –l 16500”). Multiple sequencing runs were run to increase coverage, and merged together. Duplicates were then removed by identifying clusters of reads which have the same start and stop position, and the same mapped orientation. The highest base quality representative of each set is used to represent the cluster. The mix of mitochondrial and nuclear loci necessitates two different references for the alignment process: for mitochondrial analysis, we use the RSRS (Behar et al., 2012) mitochondrial genome; for nuclear analysis we use the hg19/GRCh37, 1000 Genomes release reference genome.

**Additional libraries and processing**

To collect more nuclear data for a subset of the samples, we prepared additional barcoded libraries (I0589, I0595, I1048, I3726) without UDG-treatment from existing DNA extracts (L2-L5). For sample I0589 we collected more bone powder than necessary for one extraction, and therefore prepared three additional libraries from a newly prepared DNA extract (E2). Four additional libraries for one sample (I3726) were prepared on an Agilent Bravo Workstation using an automated protocol based on the partial UDG protocol that replaced the MinElute cleanups with magnetic bead cleanups.

All libraries underwent the same procedure as outlined, screening (see above) and nuclear target enrichment (see below), with the exception that up to 4 libraries from the same sample were pooled in equimolar concentrations before screening and nuclear target capture (Table S1). Preprocessing and alignment for nuclear data used the same procedure as performed in screening, without requiring the mitochondrial alignments.

**Shotgun genome sequencing**

Shotgun sequencing of the ancient South African from St. Helena Bay was performed at the Max Planck Institute in Leipzig, Germany, on four lanes of the Illumina HiSeq 2500 platform in rapid mode, using a double index configuration (2 × 76 + 2 × 76) (Kircher et al., 2011). An indexed X174 control library was spiked in prior to sequencing. Base calling was done with the machine-learning algorithm freeIBIS (Renaud et al., 2013). Overlapping pair-end reads were merged (Kircher, 2012) and mapped to the human reference genome (hg19/GRCh37, 1000 Genomes release) using the Burrows-Wheeler Aligner (BWA) (Li and Durbin, 2009). BWA parameters were adjusted for ancient DNA sequences (“-n 0.01 –o 2 –l 16500”), to allow for more mismatches and indels and to turn off the seeding (Meyer et al., 2012). A total of 64,128,220 raw sequences were obtained from the first shotgun sequencing of the library A5369. Another 800,205,849 raw sequences were generated from the size-selected library using four lanes of the Illumina HiSeq 2500. Only mapped sequences longer than 35 bp were retained and duplicates removed (bam-rmdup; https://bitbucket.org/ustenzel/biohazard), leaving 9,880,908 sequences from the first shotgun run and 52,551,348 sequences from sequencing the size-fractionated library. Duplication rates were 1.02 and 1.06, respectively, indicating that both libraries were not sequenced to exhaustion. The proportion of sequences ≥ 35bp that mapped to the human reference genome was ~28%.
Shotgun sequencing of the ancient South African from St. Helena Bay (UCT386) and the ancient South African from Kasteelberg (UCT437) was performed at the University of Tuebingen using two lanes of an Illumina HiSeq2500 instrument for 2x101+8 cycles (UCT386 non-UDG-treated library), on 50% of two lanes of an Illumina NextSeq500 instrument for 2x151+8 cycles (UCT437 non-UDG-treated library), on a complete run of an Illumina HiSeq2500 instrument for 2x125+8 cycles (four UCT386 UDG-treated and size-selected libraries and six UCT437 UDG-treated and size-selected libraries), and on 5.5 lanes of an Illumina HiSeq2500 instrument for 2x125+8 cycles (four UCT386 UDG-treated libraries without size-selection and three UCT437 UDG-treated full-digest libraries without size-selection). The samples were processed using the EAGER pipeline (Peltzer et al., 2016), clipping adapters and merging reads subsequently with an overlap of 10bp. Resulting reads were then mapped within the pipeline against the human reference genome GrCh37 and using BWA 0.7.5 (Li and Durbin, 2009) for further downstream analysis.

Shotgun sequencing of the Malawi_Hora_8100BP samples was performed at Harvard Medical School using an Illumina NextSeq500 instrument. Preprocessing and alignment used the same procedure as performed in screening.

In-solution nuclear target enrichment
After libraries passed screening QC (that is, there was evidence of authentic ancient DNA), we performed nuclear target enrichment of the short (but barcoded) libraries following (Fu et al., 2015) aiming to enrich for about 1.24M SNPs in total (Fu et al., 2015; Haak et al., 2015; Mathieson et al., 2015) using a semi-automated approach on a Perkin Elmer Evolution P3 liquid handler. For two libraries (S0589.E1.L1 and S2595.E1.L1) the desired 1.24M targeted SNPs were captured in two independent reactions by enriching, first, for about 0.39M SNPs, and second, for 0.84M SNPs. The other first libraries (L1) were enriched in one single reaction (1240k). After indexing PCRs with dual indices and equimolar pooling sequencing was performed on an Illumina NextSeq500 with 2x76cycles + 2x7cycles. Preprocessing and alignment used the same procedure as described in screening, without requiring the mitochondrial alignments.

Genotyping and initial processing of 34 present-day individuals from Malawi
 Genome-wide data was collected from 34 present-day individuals from Malawi using the Affymetrix Human Origins SNP array (Paterson et al., 2012), under a protocol for analysis of de-identified DNA samples approved by the Harvard Human Research Protection Program (Protocol 11681, most recently reviewed on 12 July 2016). Quality control of the data prior to merging involved screening for outlier individuals, excess missingness, as well as deviations from Hardy-Weinberg equilibrium, and was performed in a manner similar to what has previously been described (Lazaridis et al., 2014).

Data processing and preparation
We extracted genotypes from the ancient genomes by drawing a random sequence read at each position, ignoring the first and last 3 bp of every read and any read containing insertions or deletions in their alignment to the human reference genome. If the randomly drawn haploid genotype of an ancient individual did not match either of the alleles of the biallelic SNP in the reference panel, we set the genotype of the ancient individual as missing.

We added these pseudohaploid genotypes to 17 million dinucleotide transversion SNPs identified between present-day genomes from the Simons Genome Diversity Panel (which includes human-fixed differences to chimpanzee). We also added the ancient genotypes to 550 individuals from 56 African populations genotyped on the Affymetrix Human Origins array (Lazaridis et al., 2014; Paterson et al., 2012; Pickrell et al., 2012; Pickrell et al., 2014; Skoglund et al., 2015). To all these datasets we added diploid genotypes from two archaic human genomes — a Neanderthal and a Denisovan (Meyer et al., 2012; Prüfer et al., 2014). The populations shown in Figure S2 are individuals from the Affymetrix Human Origins array, when we in the text refer to Khomani_San, they are the individuals from the Simons Genome Diversity panel and so are not shown in the legend in Figure S2.

QUANTIFICATION AND STATISTICAL ANALYSIS
Population genetic approaches that quantify shared genetic drift, such as f-statistics and admixture graph fitting, are maximally robust when ascertainment of SNPs are performed in an outgroup (Paterson et al., 2012; Wang and Nielsen, 2012), such that there is no bias in allele frequencies between the analyzed populations and the polymorphism that appeared by mutation in the ancestral population of all analyzed populations. Whereas the Human Origins Array that we use for most analyses comprises 13 different panels ascertained in modern humans (Paterson et al., 2012), none of these can be regarded as outgroup-ascertained for the purpose of African populations. To obtain an outgroup-ascertained set of SNPs for African populations for our modeling of deep phylogenetic relationships, we identified 814,242 transversion SNPs polymorphic between the archaic Denisovan (Meyer et al., 2012) and Neanderthal (Prüfer et al., 2014) genomes (together labeled as ‘Archaic’ here). Since the ancestors of Denisovans and Neanderthals are consistent with having diverged from sub-Saharan lineages before those lineages separated from each other (Green et al., 2010; Mallick et al., 2016; Meyer et al., 2012; Prüfer et al., 2014; Reich et al., 2010), the ascertained SNPs that were also present as polymorphisms in sub-Saharan Africa were highly likely to have been polymorphic before the African populations diversified. We extracted these positions from the 1000 genomes project MSL (Mende from Sierra Leone: 81 unrelated individuals), and YRI (Yoruba from Ibadan, Nigeria; 107 unrelated individuals), to increase power. These 1000 genomes project sequences were processed by sampling a random sequence at each position as for the ancient data, setting the genotype as missing if it did not match either of the two alleles in the ascertained SNP set.
Principal component analysis and ADMIXTURE clustering analyses

We used smartpca (Patterson et al., 2006) to compute principal components using all transversion and transitions SNPs, and the present-day populations shown in Figure 1 and Figure S2. We projected the ancient individuals the option lsaproject: YES, on eigenvectors computed using the present-day populations on the Affymetrix Human Origins array. To deal with the confounder factor of recent admixture with western Eurasian-related populations on the PCA, we removed all northern Africans, eastern African Cushitic speakers, Nama, and 8 Khomani individuals that had 5% or more cluster membership in the shared with Europeans in an ADMIXTURE analysis.

For our main ADMIXTURE clustering analysis (Alexander et al., 2009) (Figure 1B; Figure S3) we excluded 166,439 SNPs that were in a CpG context and thus retain postmortem damage, and used 431,134 SNPs and 208 selected ancient- and present-day individuals genotyped on the Human Origins Array. In Figure 1B, we show only eight representative individuals for the non-African Japanese (originally n = 29), and Sardinian (n = 27) populations. For the authentication analysis investigating evidence of contamination, we used PMDtools (Skoglund et al., 2014a) to isolate sequences from each sample that had clear evidence of contamination according the postmortem damage score (PMD score > 3, using only based with phred-scaled quality of at least 30 to compute the score), and performed clustering analysis only on 111,208 transversion SNPs (Figure S3). The exclusion of transition SNPs is due to the PMD score approach enriching for C>T and G>A substitutions indicative of ancient DNA.

Symmetry statistics and admixture tests

D-statistics, f4-statistics, and f2-statistics (Patterson et al., 2012; Reich et al., 2009) were computed with POPSTATS (Skoglund et al., 2015). f2-statistics test two pairs of populations are symmetric with respect to one another, and quantify any asymmetry arising from admixture. More specifically, if p1, p2, p3, and p4 are the derived allele frequencies at a biallelic SNP locus in population 1, population 2, population 3, and population 4, we can estimate f2 as a sum over all SNP loci f2 = \sum (p1 - p2)(p3 - p4) (Reich et al., 2009). D-statistics (Green et al., 2010), which are also used in this study, are a version of f2-statistics with a denominator to normalize for heterozygosity, but in practice both statistics have similar power to detect deviations from the null model, and f2-statistics have the additional advantage of being directly informative about admixture proportions and shared genetic drift (Patterson et al., 2012).

For the statistics in Figure 3B and Figure 3C, we used 814,242 transversion SNPs polymorphic between the archaic Denisovan (Meyer et al., 2012) and Neanderthal (Prüfer et al., 2014) genomes (together labeled as ‘Archaic’ here). We extracted these loci from the 1000 genomes project MSL (Mende from Sierra Leone; 81 unrelated individuals), and YRI (Yoruba from Ibadan, Nigeria; 107 unrelated individuals). The statistics in Figure S2C used either complete genome sequences from the Simons Genome Diversity Panel (Mallick et al., 2016), or panel 5 of the Human Origins Array, which comprises 119,413 SNPs that were originally ascertained as polymorphic positions in a single Yoruba individual. We used this set to test whether the ancient individuals were closer to one of two San groups because some of the SNPs on the Human Origins array were ascertained in one of the San groups, potentially affecting the statistic. In Table S6 we report multiple D-statistics for different configurations of populations using transversion SNPs in complete genomes from the Simons Genome Diversity Project and ancient shotgun sequences.

Y chromosome and mitochondrial haplogroups

For Y chromosome haplogroup calling, we filtered reads with mapping quality < 30 and bases with base quality < 30, and determined the most derived mutation for each sample using the tree of the International Society of Genetic Genealogy (http://www.isogg.org) version 11.110 (21 April 2016). We also used Yfitter (Jostins et al., 2014) to confirm the haplogroups of the male Faraoskop and St. Helena Bay individuals using the entire shotgun sequence data, with identical haplogroup calls as the other approach.

For mitochondrial DNA haplogroups, we used Haplogrep2 (Weissensteiner et al., 2016) with PhyloTree 17 (van Oven and Kayser, 2009), restricting analysis to sites with base quality 10, and depth 1. These relatively permissive thresholds were used to maximize coverage on the mitogenome. For sample I2966, which was found to have mitochondrial contamination, we first restricted to damaged reads using a PMD score threshold of 3 (Skoglund et al., 2014a).

Ancestry model and estimates with qpAdm

Clustering analyses and PCA are sensitive to genetic drift, such as the genetic drift that occurs in a population after the time ancient individuals lived (Skoglund et al., 2014b), and may thus not provide an accurate view of shared ancestry between ancient and present-day individuals. We employed a framework for estimating ancestry proportions that is based on f4-symmetry statistics, taking advantage of the fact that f4-statistics are proportional to admixture proportions and genetic drift. In the well-documented case of Neanderthal admixture into non-African populations, for example, the statistic f4(chimpanzee, Neanderthal; African, non-African) is proportional to \alpha x, where \alpha is the proportion of Neanderthal-related ancestry (approximately 2%) and x is proportional to the amount of genetic drift that occurred from the divergence of Neanderthal ancestors and African ancestors, to the divergence of the sampled Neanderthal genome and the Neanderthal population that admixed with non-Africans. By analyzing many such f4-statistics, (Lazaridis et al., 2014) and (Tilak et al., 2015) showed that it is possible to estimate admixture proportions for a target population without detailed assumptions about population phylogeny, and also to perform hypothesis tests for whether a particular mixture model fits the data (Reich et al., 2012) and to estimate standard errors for admixture proportions with a weighted block jackknife procedure over large segments over the genome (in this study 5 cM). This has been implemented as the qpAdm algorithm in the ADMIXTOOLS package and requires the proposal of a set of source populations as well as a set of outgroups that are proposed to not
share drift with the target population more recently than the source populations. In other words, appropriate source populations do not need to be the true source populations but instead, need only be more closely related to the true source populations than they are to any of the outgroups. Violations of these assumptions can be detected as an increase in rank in the matrix of $f_4$-statistics computed (Reich et al., 2012). We also analyze a statistic using fitted allele frequencies predicted using the estimated mixture proportions, $f_4$ (Target population, Fitted Target population; Mbuti, Test). Deviations of this statistic from 0 are informative about whether some outgroups have an excess, or deficiency, of shared drift with the Target population under the fitted mixture proportions.

Here, we used a model with 19 populations (Mbuti, Dinka, Mende, South_Africa_2000BP, Tanzania_Luxmanda_3100BP, Ethiopia_4500BP, Levant_Neolithic (PPNB), Anatolia_Neolithic, Iran_Neolithic, Denisova, Loschbour, Ust_Ishim, Georgian, Iranian, Greek, Punjabi, Orcadian, Ami, and Mixe), using previously published complete genomes (Fu et al., 2014; Lazaridis et al., 2014; Mallick et al., 2016; Meyer et al., 2012) and ancient DNA data enriched using the 1240k SNP set (Lazaridis et al., 2016; Mathieson et al., 2015) to maximize the power to infer admixture proportions for the ancient African populations. These populations, and in particular the ones from Africa, were chosen to capture major strands of ancestry and extremes in population differentiation found in sub-Saharan Africa (Figure 1).

We then successively moved a set of candidate source populations (Mende, Dinka, Mbuti, South_Africa_2000BP, Tanzania_Luxmanda_3100BP, Ethiopia_4500BP, PPNB Anatolia_Neolithic, Iran_Neolithic) from the outgroup set to test if they fit as sources in admixture models. Using these 9 candidate sources populations, for each target population we thus tested 9 one-source ancestry models, $\binom{9}{2} = 36$ two-source admixture models, and $\binom{9}{3} = 84$ three source admixture models, for a total of 9+36+84 = 129 models. In Figure S2 and Table S5, we show admixture proportions for the model with the lowest chi-square score (or highest p value), if that model had a p value > 0.01. If a one-source model did not fulfill this criterion, we considered two-source models, and then subsequently three-source models if no two-source model fulfilled the criteria.

We successfully obtained mixture models for 55 Target populations, comprising both ancient populations (we excluded Malawi_Chencherere_5200BP due to low SNP coverage) and populations genotyped on the Affymetrix Human Origins array, all shown in Table S5. In one analysis, Tanzania_Luxmanda_3100BP was also used as a target population, and in these analyses it was dropped from the outgroup set. We highlight some notable mixture models inferred here:

- Kenya_400BP, Tanzania_Pemb_1400BP and Hadza1 are all fitted as having ~100% Ethiopia_4500BP-related ancestry. The other group of Hadza samples are fitted as having 19 ± 8% Dinka-related ancestry (the remainder being Ethiopia_4500BP-related).
- Tanzania_Pemb_600BP, Malawi_Chewa, Malawi_Ngoni, Malawi_Tumbuka, Malawi_Yao, Yoruba, Esan, Gambian, Luo, BantuKenya, BantuSA_Ovambo, Himba, Wambo, BantuSA_Herero are all fitted as consistent with having ~100% Mende-related western African-related ancestry. The Mandenka, from the western African coast, are fitted as having 2.8% ± 0.6% Levant Neolithic-related ancestry (PPNB).
- The Luhyia, an eastern Bantu-speaking group, are fitted as having 40% ± 6% Dinka-related ancestry, with the remainder being western African Mende-related ancestry.
- The Biaka, a western rainforest hunter-gatherer group in Cameroon, is fitted as having 72% ± 2% Mbuti-related ancestry (the Mbuti are an eastern rainforest hunter-gatherer group), with the remainder being western African Mende-related ancestry.
- The minimum indigenous southern African ancestry observed in Khoe-groups and Bantu-speakers in southern Africa is ~8% ± 2% in the Damara, and the remainder is western African-related.
- The maximum indigenous southern African ancestry observed in the present-day populations is the 91% ± 1% inferred for the Ju_hoan_North, with the remainder being related to Tanzania_Luxmanda_3100BP.
- Some populations in northern and eastern Africa are fitted as having large proportions of Tanzania_Luxmanda_3100BP-related ancestry. This includes the Maasai (49% ± 2%) and Datog (66% ± 3%) who have ancestry also related to the Dinka; the Kikuyu (63% ± 2%) who also have ancestry related to the Mende; and finally, the Afar (79% ± 3%) and Somali (62% ± 6%) who have large amounts of inferred Tanzania_Luxmanda_3100BP-related ancestry in addition to ancestry related to the Iran Neolithic.

**Maximum likelihood tree model**

We used the four ancient African shotgun genomes together with complete genomes from African populations in the Simons Genome Diversity project (Mallick et al., 2016), excluding populations with evidence of asymmetrical allele sharing with non-Africans indicative of gene flow (Table S6), to reconstruct a maximum likelihood tree using Treemix v1.12 (Pickrell and Pritchard, 2012). We performed 100 bootstrap replicates to assess the uncertainty of the fitted model. While this tree is not an adequate representation of human population history in Africa, we found 100% bootstrap support for the Ethiopian_4500BP Mota being most closely related to the ancestral population of all non-Africans (Figure 3A).

**Testing a tree-like model of African population history**

The maximum-likelihood tree based on allele frequency covariance between the ancient African genomes and complete genomes from the SGDP panel (Figure 3A) recapitulates many aspects of previous analyses of African populations (Pickrell et al., 2012;
Schlebusch et al., 2012). When African populations are forced into a tree (not allowing for mixture), southern African ancestors diverge first, followed by Central African rainforest hunter-gatherers (e.g., Mbuti), western Africans, and present-day eastern Africans (Dinka, Hadza), then non-Africans.

To scrutinize this tree-like model in more detail, we computed all 35 D-statistics that include the outgroup and have an expected value of 0 if a tree-like model (chimpanzee, (South_Africa_2000BP, (Mbuti, (Mende, (Dinka, Ethiopia_4500BP)))) is true. We find that most statistics computed are inconsistent with the null model (Figure S4A). Notably, we reject the hypothesis that the ancient South Africans are an outgroup to other African populations for several pairs of present-day populations (Table S6; Figure S4A). For instance, with genome sequence data we find that the Ethiopian_4500BP Mota genome shares more derived alleles with the ancient South Africans than with present-day western Africans ([D[chimpanzee, South_Africa_2000BP; Yoruba, Ethiopia_4500BP] > 0). This is also true when West Africans are compared to Mbuti ([D[chimpanzee, South_Africa_2000BP; Yoruba, Mbuti] > 0). Similar excess of shared derived alleles is observed for the eastern Pygmies (Mbuti) compared to western Pygmies, and even when contrasting western African populations such as the Yoruba and Mende (Figure 3B; Figure 3C; Table S6; Figure S4). This could be explained in two ways: 1) there has been gene flow between ancient southern Africans and a broad set of other populations that has resulted in a gradient of southern African relatedness, or 2) there is a gradient of ancestry in western Africans that is basal to southern Africans, causing an attraction to the outgroup (chimpanzee in this case) (see below).

Testing admixture graph models of African population history
To reconstruct admixture graph models relating the histories of African populations, we used South_Africa_2000BP (South Africa), Mende (MSL; West Africa), and Ethiopia_4500BP (East Africa) to represent major lineages contributing to present-day Africans. In addition, we sought to explain one of the most surprising observations in our data, that the Mende and Yoruba West African populations are not symmetrically related to South_Africa_2000BP, and so we also included the Yoruba (YRI) in these analyses. For all admixture graph analyses, we used 814,242 transversion SNPs polymorphic between the archaic Denisovan (Meyer et al., 2012) and Neanderthal (Prüfer et al., 2014) genomes (together labeled as ‘Archaic’ here).

A tree-like model does not fit the data
We first tested a strict tree-like model with no admixture edges, hypothesizing that the topology obtained from basic tree reconstructions where southern Africans are the earliest diverging lineage, and Yoruba and Mende are a clade to the exclusion of the Ethiopia_4500BP, is true (Figure S4B). We found that this model is strongly rejected by the data with 19 predicted $f_4$-statistics deviating from the empirically observed data by $|Z| > 3$. The most deviating $f_4$-statistic was $f_4$ (Archaic, Ethiopia_4500BP; MSL, YRI), which is predicted to be 0 in the tree-like model but is empirically observed to be $f_4 = 0.000427$, $Z = 9.157$. Insight into the imperfect fit can be obtained by inspecting all four significant $f_4$-statistics that are predicted to be zero in the model:

- $f_4$(Archaic, Ethiopia_4500BP; MSL, YRI) = 0.000427, $Z = 9.157$
- $f_4$(Archaic, South_Africa_2000BP; MSL, YRI) = 0.000193, $Z = 4.742$
- $f_4$(South_Africa_2000BP, Ethiopia_4500BP; MSL, YRI) = 0.000235, $Z = 4.583$
- $f_4$(Archaic, South_Africa_2000BP; Ethiopia_4500BP, MSL) = −0.000728, $Z = −3.068$

The three most significant deviations all test the (MSL, YRI) clade. The deviations thus indicate shared history either between MSL and the outgroup Archais, or between YRI and South_Africa_2000BP/Ethiopia_4500BP. These observations could be parsimoniously explained by any of the following gene flow events:

- Gene flow from a basal human lineage (separating from the ancestors of all sub-Saharan Africans before their separation from each other) into the ancestors of MSL to a greater extent than YRI (there is no evidence of specifically Neanderthal/Denisovan gene flow since $f_4$(chimpanzee, Archais; MSL, YRI) = −1.6 in this data)
- Gene flow related to YRI into the ancestors of Ethiopia_4500BP and South_Africa_2000BP
- Gene flow related to Ethiopia_4500BP into the ancestors of YRI more than MSL
- Gene flow most closely related to MSL into the common ancestors of the archaic Neanderthals and Denisovans (we exclude this as implausible)

The fourth deviating statistic $f_4$(Archaic, South_Africa_2000BP; Ethiopia_4500BP, MSL) ($Z = 3.068$) suggests that Ethiopia_4500BP and MSL are not a clade with respect to South_Africa_2000BP. This weakens the case for gene flow into Yoruba alone as a sufficient explanation, since the Yoruba do not enter into this statistic. Instead, either excess basal ancestry in the Mende and gene flow between South_Africa_2000BP and Ethiopia_4500BP could explain this particular statistic.

Admixture models with gene flow events
We proceeded by testing models with one gene flow event positing that YRI have mixture related to either South_Africa_2000BP or Ethiopia_4500BP, or that MSL have ancestry from a basal lineage (Figure S4C; Figure S4D; Figure S4E). We found that neither of these fit the data, with between 4 and 13 $f_4$-statistic outliers with $|Z| > 3$.

Testing admixture graphs with two admixture events, we found that a model where both YRI and MSL have ancestry from a basal African lineage had its single outlier in an $f_4$ statistic $f_4$(Archaic, Ethiopia_4500BP; Ethiopia_4500BP, YRI) that is more negative in the
data than the model (Figure S4F). This $f_2$-statistic has Ethiopia_4500BP appearing twice, and can thus be rearranged to be an $f_4$ statistic $f_4$(Archaic, YRI; Ethiopia_4500BP) that is not positive enough ($Z = 3.157$), and can thus be interpreted as the model under-representing the external drift in the Ethiopia_4500BP genome. We do not consider this outlier to compromise the model, as the processing of the Ethiopia_4500BP (Mota) genome that we use is pseudo-haploid (single random sequence read), and thus there is no real information about the external drift of the Mota lineage. This outlier may thus reflect a difficulty in modeling external drift for pseudo-haploid samples.

In addition, we found that an admixture graph where both Ethiopia_4500BP and YRI are mixed between MSL- and South_Africa_2000BP lineages does not fit the data (Figure S4G), with 3 outliers, and the worst being $Z = -4.835$. However, a model where the YRI has ~2% ancestry from a population that is mixed between South_Africa_2000BP and Ethiopia_4500BP fits the data with 2 outliers that are not too surprising after correcting for multiple hypothesis testing (Figure S4H). These outliers are $f_2$(Archaic, South_Africa_2000BP; Ethiopia_4500BP, MSL) ($Z = 3.068$) which was also significant for the tree model without admixture, and $f_4$(Archaic, South_Africa_2000BP; Ethiopia_4500BP, YRI) ($Z = 3.018$). Both these outliers could be consistent with the presence of basal African ancestry in YRI and MSL, or unmodeled gene flow between the ancestors of South_Africa_2000BP and Ethiopia_4500BP.

We thereby conclude that the most parsimonious admixture graph models identified here posit either the presence of basal African ancestry in Mende and Yoruba (Figure 3D; Figure S4F), or alternatively admixture from a source related to both South_Africa_2000BP and Ethiopia_4500BP in the Yoruba but not the Mende (with evidence also for gene flow between South_Africa_2000BP and Ethiopia) (Figure 3F; Figure S4G).

Automated grafting of populations onto a skeleton admixture graph

Using the admixture graph model in which the Yoruba and Mende both carry ancestry from a basal western African population, we automatically added additional populations to each possible node in the graph. We evaluated the fit in terms of the number and deviation of outlying $f_2$-statistics, as well as whether the added branches had zero drift length. We show the results of this procedure in Table S7, with the key to the node labels used shown in Figure S5A. We highlight topologies that we consider optimal fits in Figure S5B-5G.

We find that the Malawi_Hora_8100BP shotgun sequence data can be best fitted as mixed between the lineage leading to South_Africa_2000BP, and the lineage related to Ethiopia_4500BP that also is fitted as forming part of the ancestry of YRI and MSL (Figure S5B). Similarly, the Mbuti can be best fitted as mixed between the lineage related to Ethiopia_4500BP that also is fitted as forming part of the ancestry of YRI and MSL, and second a lineage that diverged prior to South_Africa_2000B but after the basal western African lineage (Figure S5C). Non-Africans (Japanese) are fitted as having part Archaic ancestry (Green et al., 2010), with the remainder of their ancestry again being derived from the lineage related to Ethiopia_4500BP that also is fitted as forming part of the ancestry of YRI and MSL (Figure S5D). This analysis is consistent with the possibility that the same human lineage contributed ancestry both to the source of non-Africans and many African populations today.

Support for a single out-of-Africa founding population

Simple tree models suggest that non-African variation represented by Sardinian, English, Han Chinese and Japanese falls within the variation of African populations. To test whether non-Africans are indeed consistent with being descended from a homogeneous population that separated earlier from a subset of African populations – beyond the known effects of archaic admixture in non-Africans – we used African populations with little or no known West Eurasian mixture (South_Africa_2000BP, Mbuti, Biaka, Mende, Ethiopia_4500BP, Dinka) and tested whether they are consistent with being an unrooted clade with respect to a diverse set of non-Africans (Orcadian, Onge, Mixe, Motala_Mesolithic, Japanese, Anatolia_Neolithic) using qpWave (Patterson et al., 2012; Reich et al., 2012). We found that this model was consistent with the data ($p = 0.53$) (transition SNPs excluded to a final set of 110,507 transition SNPs). Even when we add New Guinean highlanders to the set of non-Africans, the single-source model for the out-of-Africa founders is not rejected ($p = 0.11$).

Date of admixture between expanding agriculturalists and previously established foragers

We estimated the date of admixture between expanding agriculturalists related to western Africans (probably Bantu-speakers) and the previously established foragers using the ALDER software (Loh et al., 2013), which uses the rate of decay per generation of the linkage disequilibrium that is created in admixed populations and that can be detected using signed linkage disequilibrium weighted by allele frequency differences between populations taken as proxies for the ancestral populations (Moorjani et al., 2011). To maximize statistical power, we used the full 1240k data merged with 1000 genomes phase 3 genotype data. We then estimated weighted linkage disequilibrium in 99 unrelated LWK individuals (Luhya from Webuye, Kenya) a Bantu-speaking group, using 85 unrelated West African MSL individuals (Mende from Sierra Leone) and 4 ancient eastern African hunter-gatherer individuals (Ethiopia_4500BP, Tanzania_Pemba_1400BP, Tanzania Zanzibar_1400BP, Kenya_400BP). The analysis used a total of 1,070,197 SNPs. We obtained significant evidence for one-reference weighted LD decay for both putative source populations ($Z \sim 6$), as well as for the two-reference weighted LD decay ($Z = 4.89, p = 10^{-4}$). The estimated date of mixture was 16.8 generations ago, with a standard error of 3.4. Assuming a generation time of 30 years, this suggests that admixture occurred on average 380 to 760 years ago (95% confidence interval). We note that a previous study (Pickrell et al., 2014) did not obtain high-confidence support for West African related admixture in the Luhya, and we
hypothesize that our clear demonstration of this is due to both the availability of a more accurate ancestral population in the form of the ancient eastern Africans, as well as to the increased leverage from the large samples size 1000 Genomes data.

**Evidence for selective sweeps in the ancestry of present-day San**

We performed a genome-wide scan for large genomic regions with excessive allele frequency between present-day San (Khomani and Ju_hoan_North) and the two ancient South_Africa_2000BP, with the Mbuti as a second outgroup. Previous statistics such as the Locus-Specific Branch length (LSBL) (Shriver et al., 2004) and the derivative Population Branch Statistic (PBS) (Yi et al., 2010) also estimate a branch length in a three-population phylogeny, but use $F_{ST}$ as their base, which can be undefined when used for loci with fixed differences. Thus, we computed the statistic $f_3 = (p_{San} - p_{South_Africa_2000BP})(p_{San} - p_{Mbuti})$ for windows of 500kb, separated by 10kb. We only retained windows with at least 50 SNPs, resulting in a total of $l = 262,047$ autosomal loci. We approximated the neutral genome-wide average $\mu$ and its standard deviation $\sigma$ by subsampling 546 of the windows, requiring that these were separated by at least 5 million base pairs (Mb) and thus approximately independent. We then standardized the distribution of the test statistic $f_w$ in each window as $Z(f_w) = (f_w - \mu) / \sigma$. We show the most deviating windows in Table 2.

**Evidence for polygenic selection**

Investigating evidence for polygenic selection in African populations is complicated by the fact that most information on the genomic basis of human phenotypic variation has been based on analysis of highly differentiated populations from Africans such as Europeans. In the absence of rich phenotypic information about sub-Saharan African populations, we used gene ontology (GO) information, which draws on information from a wide array of studies on humans and non-human model organisms. We focused on 208 GO categories that contained at least 50 genes each, which allows us to compute genome-wide block jackknife standard errors using the entire genic regions. We focus on $f_3$-statistics that measure the length of one of the branches in a hypothetical three-way tree-like population history (assuming no admixture). The three populations we focused on were the ancient South_Africa_2000BP ($n = 2$ pseudohaploid draft genomes), the present-day San ($n = 6$ complete genomes), and the Mbuti ($n = 4$ complete genomes). The availability of the two ancient South_Africa_2000BP genomes can in principle inform us about selection in the last ~2000 years since these individuals lived, or starting further back in time in case they are not from a direct ancestral population of the present-day San (our data suggest that they are more closely related to the Khomani San than to the Ju'_hoan). We thus computed the statistic $f_3(Mbuti, South_Africa_2000BP; San)$ which is proportional to allele frequency differentiation in the present-day San compared to the other two populations.

We find that the “RESPONSE_TO_RADIATION” GO category is an outlier that shows the greatest degree of differentiation in this analysis. However, this could also be due to genes in this category constantly being under rapid evolution or having other differences compared to other categories. To test this, we computed the statistic $f_3(San, South_Africa_2000BP; Mbuti)$, and found no strong signal in the response to radiation category. Instead, the category with the strongest evidence of differentiation in the Mbuti lineage since the divergence from the San groups is “REGULATION_OF_GROWTH,” suggesting the possibility of relatively recent evolution of the shorter stature of present-day rainforest hunter-gatherer populations.

**DATA AVAILABILITY**

Raw sequence data (bam files) from the 15 newly reported ancient individuals is available from the European Nucleotide Archive. The accession number for the sequence data reported in this paper is ENA: PRJEB21878. The newly reported SNP genotyping data is available to researchers who send a signed letter to D.R. containing the following text: “(a) I will not distribute the data outside my collaboration; (b) I will not post the data publicly; (c) I will make no attempt to connect the genetic data to personal identifiers for the samples; (d) I will use the data only for studies of population history; (e) I will not use the data for any selection studies; (f) I will not use the data for medical or disease-related analyses; (g) I will not use the data for commercial purposes.”
Figure S1. Archaeological Context and Dating of Newly Reported Human Remains from Malawi, Related to Figure 1

(A) Locations of the three sites in Malawi.

(B) Plan of the Hora rock shelter showing locations of the original Hora 1 (male) and Hora 2 (female) burials relative to new excavations from 2016 and 2017.

(C) Plan of Fingira Cave showing locations of human remains recovered in 1966 (green dots) and 2016 (green stars).

(D) Plan of Chencherere II rock shelter redrawn from Crader (1984a), showing approximate locations where human remains were recovered in 1972 (red and green stars), and the approximate locations of the remains sampled from Nguludi for this study (blue stars).

(E) Ultrafiltered collagen FTIR spectra for three individuals from Malawi (MW2 = UCIAMS-186346, MW3 = UCIAMS-18647, MW4 = UCIAMS-186348) compared to well-preserved standards and samples (Modern Caribou, BU4, RP01, POR1). These samples show the typical features for well-preserved collagen. The top of the figure shows the FTIR spectra for a single XAD purified sample (MW1 = PSUAMS-1734) compared against a well preserved standard (Beauford Whale). These data, along with the stable isotopes and C:N ratios, indicate that these samples are well preserved and that the AMS $^{14}$C dates are reliable.
Figure S2. Ancient Individuals and African Population Structure, Related to Figure 1
(A) Full legend and PC1-PC2 scatterplot (same analysis as Figure 1) with all present-day populations indicated with separate symbols.
(B) PC1-PC2 computed only with southern African populations (and west African Yoruba).
(C) Symmetry statistics confirming that ancient individuals from the western Cape region in South Africa are more closely related to southern San such as the Khomani rather than more northern San populations such as the Ju hoan_North.
Figure S3. Admixture Clustering Analysis of K = 2–K = 7 Clusters, Related to Figure 1

Each bar shows inferred cluster membership in each individual for a given number of clusters. This analysis used 431,134 SNPs that were not in a CpG dinucleotide context. We also show an inset with cluster membership inferred for damage-restricted ancient DNA sequences (PMD-score > 3) in a parallel run with only 111,208 transversion SNPs.

Figure S4. Admixture Models of African Population History, Related to Figure 3

(A) A tree-like model. Tests of all possible triplet topologies predicted to be symmetric by the tree obtained for major African population lineages in the maximum likelihood model in Figure 3A.

(B–H) Under each graph we show the outlier $f_4$ statistic with the greatest Z-score between model and empirical values. When the same group appears twice in an $f_4$ statistic (e.g., $f_4$(Archaic, South_Africa_HGs; South_Africa_HGs,YRI)), this corresponds to an $f_3$ statistic. Graphs F and H are the most parsimonious fits with only two admixture events.
Figure S5. Automatically Adding Additional Populations to a Skeleton Graph, Related to Figure 3

(A) Skeleton graph invoking widespread admixture from a hypothesized ancestral northeast African population.
(B–D) Best fitting insertion points for Malawi_Hora_8100BP shotgun sequence data (B), Mbuti (C), and Japanese non-Africans (D), in the skeleton graph displayed in (A). For details see Table S7.
(E–G) Best three graph solutions, all without outliers, for grafting Malawi_Hora_8100BP (Malawi_HGs) onto the skeleton graph in (A). For details see Table S6.