

A weakly structured stem for human origins in Africa

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Abstract

While a modern human origin within Africa is now broadly accepted, considerable uncertainty surrounds specific models of divergence and migration across the continent. Progress is hampered by a paucity of fossil and genomic data, as well as variability in dating. Here we use linkage disequilibrium and diversity-based statistics, optimized for rapid, complex demographic inference to discriminate among such models. We infer detailed demographic models for populations across Africa, including representatives from eastern and western groups, as well as 44 newly whole-genome sequenced individuals from the Nama (Khoe-San). Despite the complexity of African population history, present-day population structure dates back to Marine Isotope Stage (MIS) 5. The earliest population divergence among contemporary populations occurs 120-135kya, between the Khoe-San and other groups. Prior to the divergence of contemporary African groups, we infer long-lasting structure between two or more weakly differentiated ancestral *Homo* spp. populations connected by gene flow over hundreds of thousands of years (i.e. a weakly structured stem). We find that weakly structured stem models provide more likely explanations of polymorphism that had previously been attributed to contributions from archaic hominins in Africa. In contrast to models with archaic introgression, we predict that fossil remains from coexisting ancestral populations should be morphologically similar. Despite genetic similarity between these populations, an inferred **SG: 4 to 10%** of random genetic differences among [modern human populations] can be attributed to genetic drift between stems. We show that model misspecification explains variation in previous divergence time estimates and argue that studying a suite of models is key to robust inferences about deep history.

Introduction

Archaeological sites from the Middle Stone Age (approx. 300kya-40kya) are widely distributed across Africa, and are particularly well represented in the northern, eastern and southern parts of the continent. Similarly, fossil crania such as those from the sites of Jebel Irhoud¹, Herto² and Klasies River³ demonstrate that anatomically derived *Homo sapiens* features were also present across the continent

during this period. It has been difficult to reconcile these lines of evidence with evidence from genomics, which have suggested a predominantly tree-like model of recent population divergence from a single ancestral population. It is unclear whether fossil specimens and archaeological sites represent populations which contributed to our ancestors as population precedents, or were local “dead-ends” from which contemporary *Homo sapiens* do not descend. Recently, synthetic attempts to reconcile genetic and paleoanthropological data include proposals for a Pan-African origin of *Homo sapiens* by which populations in many regions of the continent contributed to the formation of *Homo sapiens* beginning at least 300kya^{4,5,6}.

Genetic models have been hampered in their contribution to this discussion because they primarily assume (or, at least, have been tested under) a tree-like model of isolation-with-migration. Alternative theoretical scenarios have been proposed, such as stepping stone models⁷ or population coalescence and fragmentation⁶. These approaches are more challenging to interpret and fit to data. However, new population genetic inference tools now allow for inference involving tens to hundreds of genomes from multiple populations and greater complexity^{8,9,10}. Inspired by evidence for Neanderthal admixture with modern humans in Eurasia, several recent articles have shown that introducing an archaic ghost population contributing to African populations in the period surrounding the Out-of-Africa migration event substantially improves the description of genetic data relative to single-origin models^{11,12,13,14,9,15,16,15}. This has driven speculation about the geographic range of this ghost population, possible links to specific archaic remains, and the possibility of finding ancient DNA evidence (e.g., Hsieh et al.¹³). However, these prior articles share two weaknesses. First, they only contrast a single-origin model with an archaic admixture model, leaving out other plausible models (Figure S2 and Henn et al.¹⁷). Second, they focus on a small subset of African diversity, either because of small sample sizes (2-5 genomes) or because they rely on 1000 Genomes data which only recruited populations of recent West African ancestry. While ancient DNA from Eurasia has helped us understand early human history outside of Africa, there is no comparably ancient DNA to elucidate early history in Africa.

Here, we therefore aim to discriminate among a broader set of demographic models by interrogating the genomes of contemporary populations. We take as our starting point 4 classes of models (single population expansion, single population expansion with regional persistence, archaic admixture, and multi-regional evolution, Figure S2), using 290 genomes from southern, eastern, and western Africa as well as Eurasia. By including geographically and genetically diverse populations across Africa, we infer demographic models that explain more aspects of genetic diversity in more populations than previously reported. These analyses confirm the inadequacy of tree-like models and provide an opportunity to directly evaluate a wide range of alternative models.

Results

We inferred detailed demographic histories using 4x-8x whole-genome sequencing data for four diverse African populations, comprising the Nama (Khoen-San from South Africa, newly presented here), Mende (from Sierra Leone, MSL from the Phase 3 1000 Genomes Project¹⁸), Gumuz (recent descendants of a hunter-gatherer group from Ethiopia^{19,20}), and Eastern African agriculturalists (Amhara and Oromo from Ethiopia¹⁹). The Amhara and Oromo populations, despite speaking distinct Afro-Asiatic languages, are highly genetically similar^{21,20} and thus the two groups were combined for a larger sample size (Figure 1). We also included the British (GBR) from the 1000 Genomes Project in our demographic models as a representative source of back-to-Africa gene flow and recent colonial admixture in South Africa. Finally, we used a high-coverage ancient Neanderthal genome from Vindija Cave, Croatia²² to account for archaic gene flow from Neanderthals in non-Africans and gauge the relative time depth of divergence, assuming Neanderthals diverged 550kya from a common stem. For each population, we computed low-order allele frequency and linkage disequilibrium (LD) statistics that are well suited for both low- and high-coverage genomes^{9,23}. Using a maximum-likelihood inference framework, we then fit to these statistics a family of parameterized demographic models that involve population splits, size changes, continuous and variable migration rates, and punctuated admixture events, to learn about the nature of population structure over the past million years.



Figure 1: **Geography and genetic diversity within Africa.** The combined 1000 Genomes and AGRP datasets include populations from West, East, and South Africa. Both PCA (B, C) and ADMIXTURE (D) illustrate the well-characterized high levels of genetic diversity and signatures of recent migration and gene flow within Africa. We built parameterized models using the newly-sequenced Nama, Mende from Sierra Leone, Gumuz, Oromo and Amhara from Ethiopia, and the British and Vindija Neanderthal individual (bold fonts). **APR: work on this caption**

A Late Middle Stone Age common ancestry for contemporary humans

We began with a model of geographic expansion from a single ancestral, unstructured source followed by migration between populations, without allowing for contribution from an archaic African lineage or population structure prior to the expansion (Figure S2A). As expected⁹, this first model was a poor fit to the data qualitatively (Figure S4) and quantitatively (log-likelihood (LL) $\approx -189,400$, Table S1). We next explored a suite of models in which population structure predates the differentiation of contemporary groups, including models allowing for ancestral reticulation (Figure S2B), archaic admixture (Figure S2C), and African multi-regionalism (Figure S2D).

Regardless of the model choice for early epochs, inference of human demographic history in the last 150kya was remarkably robust. The earliest divergence among contemporary human populations differentiates the southern African Nama from other African groups between 110–135kya, with low to moderate levels of subsequent gene flow (Table 1). In none of the high-likelihood models which we explored did the divergence between Nama and other populations exceed ~ 140 kya. We conclude that, regardless of model specification, geographic patterns of *Homo sapiens* population structure date back to the Late Middle Stone Age in Africa, likely arising during MIS 5. Although we find evidence for earlier population structure in Africa (see below), present-day populations cannot be easily mapped onto these more ancient stem groups as only a small proportion of drift between modern populations can be attributed to drift between stems (Figures 3 and S10–S13).

Given this consistency in inferred recent history and the numerical challenge of optimizing a large number of parameters, we fixed several parameters related to recent population history so as to focus on more ancient events (Supp. Methods). The time of divergence of Western and Eastern African populations was set to 60kya, just prior to the split of Eurasians and East Africans 50kya. All models predict relatively high subsequent gene flow between the two regions ($m \approx 2 \times 10^{-4}$). Admixture from Neanderthals to Europeans directly following the out-of-Africa migration was set to 1.5% at 45kya (Supp. Methods). Back-to-Africa gene flow at the beginning of the Holocene primarily affected the ancestors of the Ethiopian agricultural populations, comprising over half of their genetic ancestry, estimated to be

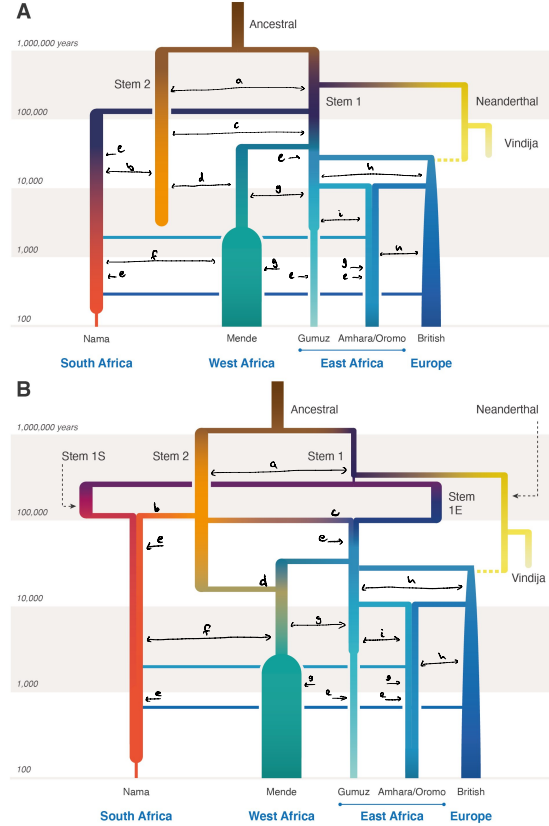


Figure 2: **Two best fit models with a weakly structured stem.** In both the the parameterizations of early population structure, continuous migration (A) and fragmentation with mergers (B), models that include ongoing migration between stem populations outperform those in which stem populations are isolated. Most populations are connected by continuous, reciprocal migration that are indicated by double-headed arrows (labels matched to migration rates and divergence times in Table 1), and which lasts for the duration of concurrence of contemporaneous populations. The merger-with-stem-migration model (B) outperformed the continuous-migration model (A), $LL = -102,600$ vs $-115,500$, respectively, and both models provide a much better fit to diversity and LD statistics than a single-origin model of recent expansion.

64–65%. The past 5,000 years also saw major demographic changes, including strong population growth for Western Africans as they specialized in yam and oil palm agriculture (estimated 3-fold growth). We observe significant gene flow from the Amhara and Oromo into the Nama, a signal which is likely a proxy for the movement of Eastern African caprid and cattle pastoralists^{24,25}, here estimated to constitute a 25% ancestry contribution 2,000 ya. Colonial period admixture from Europeans into the Nama was estimated at 15%, similar to proportions inferred by **ADMIXTURE**²⁶ (Figure 1).

Deep but connected population structure within Africa

To account for population structure prior to 135kya, we allowed for two or more “stem” populations which could diverge either before or after the Neanderthal split. We considered models both with and without migration between these stem populations, and in both cases we tested two different types of gene exchange during the expansion phase. 1) One of the stem population expands (splits into contemporary populations), and the other stem population(s) has continuous symmetric migration with those populations; or 2) one or more of the stem populations expands, with instantaneous pulse (or “merger”) events from the other stem population, so that diversifying branches are formed through the

Likelihood	Label	Population Pair	Divergence Time (kya)	Migration rate per generation	Migration duration (ky)
Continuous Model					
LL= -115,500	a	Stem 1, Stem 2	1,163	6.43e-5	1,028
	b	Stem 2, Nama	NA	5.82e-5	130
	c, d	Stem 2, Other Africans*	NA	3.10e-5, 1.64e-4	130, 55
	e, f	Nama, Other Africans*	135	4.1e-5, 9.8e-6	135, 60
	g	Mende, East Africans	60	2.14e-4	60
	h	East Africans, British	50	4.17e-5	50
	i	Gumuz, Amhara/Oromo	12	3.36e-4	12
Merger Model					
LL= -102,600	a	Stem 1, Stem 2	1,442	1.16e-4	963
	–	Stem 1S, Stem 1E	479	0 (fixed)	–
	b	Stem 2 to Nama	119	0.71	pulse
	c	Stem 2 to Stem 1E	98	0.50	pulse
	d	Stem 2 to Mende	25	0.18	pulse
	e, f	Nama, Other Africans*	119	4.4e-5, 7.1e-6	119, 60
	g,	Mende, East Africans*	60	1.98e-4	60
	h	East Africans, British	50	3.87e-5	50
	i	Gumuz, Amhara/Oromo	12	3.59e-4	12

Table 1: **Migration and divergence parameters from best fit models.** Labeled migration rates correspond to symmetric continuous migration bands shown in Figure 2. Both the continuous-migration and merger models inferred a relatively deep split of human stem branches, though they were connected by ongoing migration that maintained their genetic similarity. *: The ancestors of the Nama diverged from other African groups $\sim 120\text{--}135\text{kya}$, which later branch into West and East African groups 60kya . Migration rates and durations are shown for 1) East Africans and their ancestors, and 2) the Mende, respectively.

coalescence of multiple ancestral populations. Depending on parameter values, this scenario encompasses archaic introgression and fragmentation-and-coalescence models.

Allowing for continuous migration between the stem populations substantially improved the fits relative to zero migration ($LL \approx -102,600$ vs. $-107,700$ in the merger scenario and $LL \approx -115,500$ vs. $-126,600$ with continuous migration). With continuous migration between stems, population structure extends back to $1.1\text{--}1.4\text{Mya}$ (Table 1). Migration between the stems in these models is moderate, with a fraction of migrant lineages each generation estimated as $m = 6.43 \times 10^{-5} - 1.16 \times 10^{-4}$. For comparison, this roughly compares to inferred migration rates between connected contemporary populations over the past 50kya (Table 1). This ongoing (or at least, periodic) gene flow qualitatively distinguishes these models from previously proposed archaic admixture models (Figure S2C) as the early branches remain closely related (Figure 3).

Bootstrap-based confidence intervals are relatively narrow within each specified model (Tables S1–S5), reflecting an informative statistical approach. As expected, model assumptions have a larger impact on parameter estimates. To convey model uncertainty we highlight features of two inferred models with high likelihoods, the “multiple-merger” and the “continuous-migration” model that both allow migration between stem branches. These differ primarily in the timing of the early divergence of stem populations and their relative N_e (Figure 2). The two models also differ in the mode of divergence during the Middle Stone Age.

Under the continuous-migration model, one of the two stems diverges into lineages leading to contemporary populations in western, southern and eastern Africa, and the other (Stem 2) contributes variable ancestry to those populations. This migration from Stem 2 is highest with the Mende ($m = 1.6 \times 10^{-4}$) compared to the Nama and East African populations ($m = 5.8 \times 10^{-5}$ and 3.1×10^{-5} , respectively), with migration allowed to occur until the relatively recently demographic events 5kya . A sampled lineage from the Nama, Mende, and Gumuz have probabilities of being in Stem 2 at the time of Stem 1 expansion (135ka) of approximately 0.145, 0.2, and 0.13, respectively, though these probabilities change over time, precluding the notion of a fixed admixture proportion.

In contrast, under the multiple-merger model, stem populations merge with varying proportions to form the different contemporary groups. We observe a sharp bottleneck in Stem 1 down to $N_e = 117$ after the split of the Neanderthal branch. This represents the lower bound allowed in our optimization (i.e. 1% of the ancestral N_e), although the size of this bottleneck is poorly constrained ($\sigma_{N_e} = 838$). After a long period of exchange with Stem 2, Stem 1 then fractures into “Stem 1E” and “Stem 1S” 479kya .

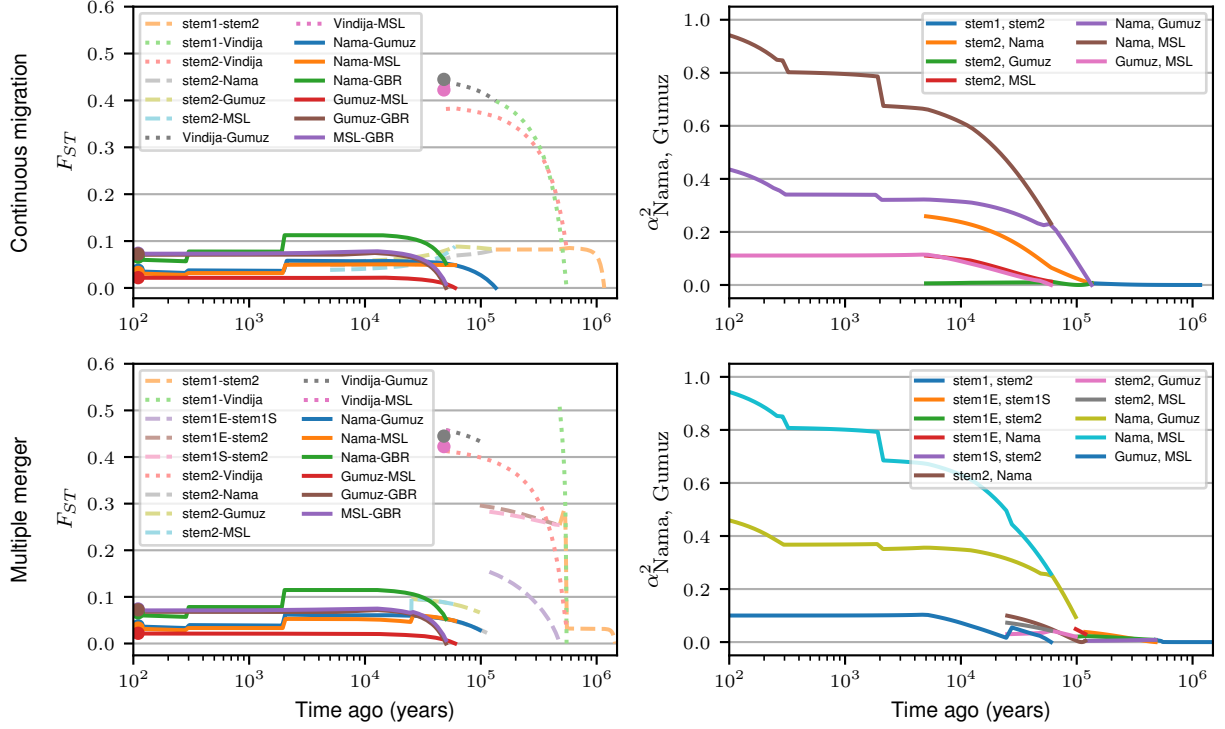


Figure 3: **Predicted population structure over time.** From the best fit models of our two parameterizations (top row: continuous migration, bottom row: multiple merger), we computed predictions of population structure between contemporaneous populations over time (F_{ST}) and the proportion of drift between pairs of sampled populations that aligns with drift between populations in the past (α^2).

The timing of this divergence was also poorly constrained ($\sigma_T = 166\text{kya}$). These populations evolve independently until approximately 119kya when Stem 1S and Stem 2 combine to form the ancestors of the Nama, with proportions 29%, 71% respectively. Similarly, Stem 1E and Stem 2 combine in equal proportions (50% each) to form the ancestors of the Western Africans and Eastern Africans (and thus also all individuals who later disperse during the Out of Africa event). Finally, the Mende receive a large additional pulse of gene flow from Stem 2, replacing 18% of their population 25kya. The later Stem 2 contribution to the Western African Mende resulted in better model fits (LL). This may indicate that an ancestral Stem 2 population occupied Western or Central Africa, broadly speaking. The differing proportions in the Nama and Eastern Africans may also indicate geographic separation of Stem 1S in southern Africa and Stem 1E in eastern Africa.

Reconciling multiple lines of genetic evidence

Previous studies have found support for archaic admixture in Africa using two-locus statistics^{13,9}, the conditional SFS (cSFS)¹⁵, and reconstruction of gene genealogies¹⁰. However, none of these studies considered a weakly structured stem. We validated our inferred models with additional independent approaches. We find that the observed cSFS (conditioned on the derived allele being carried in the Neanderthal sample) is very well-described by the merger model (Figures 4A-C and S14-S17), even though this statistic was not used in the fit. Both models outperform archaic models fit directly to the cSFS (for example, compare with Figure 1 in Durvasula and Sankararaman¹⁵).

We used **Relate**¹⁰ to infer the distribution of coalescence rates over time, which are commonly interpreted as effective population size histories. Trajectories of N_e from prior studies strongly support a bottleneck in Out of Africa populations. African populations also dip in effective size around 100-

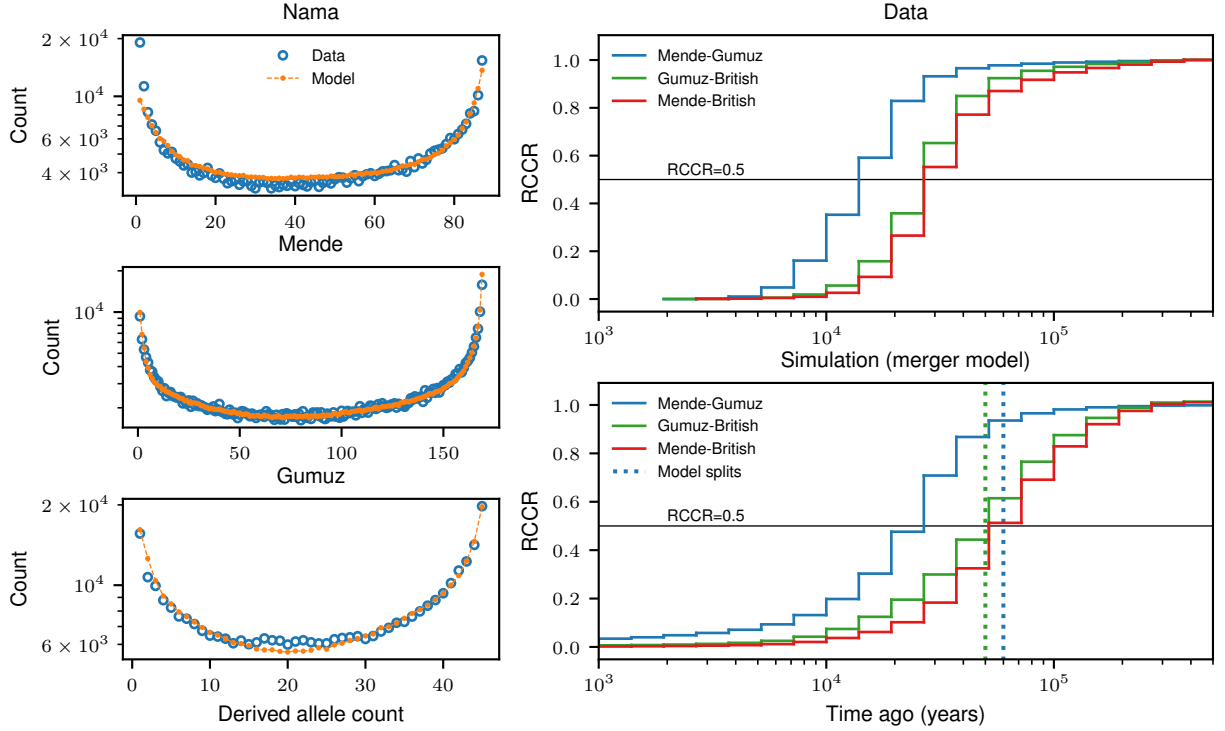


Figure 4: **Model validation using independent statistics.** (A–C) Using our best fit models to the LD and pairwise diversity statistics, we simulated expected conditional site-frequency-spectra (cSFS) and compared to the observed cSFS from the data. Our inferred models provide a good fit to the data, even though these were not used in our inference. Across the three populations, ancestral state misidentification was consistently inferred to be 1.5–1.7% for intergenic loci (Supp. Methods). (D, E) We used Relate¹⁰ to reconstruct genome-wide gene genealogies, which we used to estimate coalescence rate trajectories and cross-coalescence rates between pairs of populations. While coalescence rate distributions are informative statistics about past evolutionary processes, interpretation can be hindered by migration and population structure, and translating relative cross-coalescence rate curves (RCCR) into population divergence times is especially prone to misinterpretation. For example, the Mende-Gumuz comparison shows a more recent increased RCCR than either population with the British, a pattern that is recapitulated under our best-fit model, even though the Mende-Gumuz split occurs prior to the Gumuz-British split.

200kya, although this decrease is more modest than the OOA event²⁷. One possible explanation is that ancestral population structure during the Middle Stone Age inflated N_e and subsequent gene flow among ancestral Africans resulted in a more recent decreased N_e ²⁸. We simulated genomic data under four inferred demographic models to compare with observed coalescence rate histories. All models, including the single-origin model, generate multi-modal curves indicating that the ancestral increase in N_e between 100kya–1Mya is a general feature of the coalescent rate in humans, although none of the models perfectly match the exact timing of the early fluctuations in N_e observed in the data. Each simulated model also underestimated the level of population growth during the Holocene, indicating additional free parameters are necessary to refine these growth rates.

Relative cross-coalescence rates (rCCR) have recently been used to estimate divergences between a pair of populations, as measured by the rate of coalescence between two groups divided by the mean within population coalescence. Simulations of rCCR accuracy, however, focus on a ‘clean split’ between populations whereby groups diverge without subsequent gene flow. Published estimates of the earliest human divergences with rCCR, which range from 150kya–100kya²⁹, may be significantly biased when

compared to more complex models with gene flow as inferred here. We find that midpoint estimates of rCCR are poor estimates for population divergence, often underestimating divergence time by 50% or greater (e.g., Mende vs. Gumuz ~15kya compared to a true divergence of 60kya), and recent migration can lead to the misordering of divergence events (Figure 4E). We suggest that rCCR analyses which do not fit multiple parameters including gene flow should be interpreted with caution.

Discussion

Any attempt at building detailed models of human history is subject to model misspecification. This is true of earlier studies, which often assumed that data inconsistent with a single origin model must be explained by archaic admixture. This is also true of this study. While it remains prohibitive to fully explore the space of plausible models of early human **BMH: we use this phrase a lot. human or hominin? Tim?** population structure, we sought to capture model uncertainty by exploring multiple parameterizations of early human population structure. The best-fit models presented here include reticulation and low-level migration between early human populations rather than archaic admixture. We cannot rule out that more complex models involving additional stems, or hybrid models including both weak structure and archaic admixture may better explain the data. Because parameters related to the split time, migration rates, and relative sizes of the early stems were variable across models, reflecting a degree of confounding among these parameters, we refrained from introducing additional branches (and therefore additional parameters) during that period. **SG: Some statement about hitting the boundary** Rather than interpreting the two stems as representing well-defined and stable populations over hundreds of thousands of years, we interpret the weakly structured stem as consistent, e.g., with a coalescence and fragmentation model⁶. Eventual ancient DNA samples from Africa could further distinguish the archaic admixture model from the weakly structured stem model. **SG: Or models involving more diversity within Africa?**

The Middle Stone Age in Africa

By contrast, our inferred models paint a more consistent picture of the late Middle Stone Age as a critical period of change, assuming that estimates from the recombination clock accurately relate to geological chronologies (Supplement). During the Middle Stone Age, the Multiple Merger indicates three major stem lineages in Africa, tentatively assigned to southern (Stem 1S), eastern (Stem 1E) and western/central Africa (Stem 2). While the length of isolation among the stems is variable across model fits, models with this period of divergence, isolation and then a merger event (i.e. a “reticulation”) out-performed models with bifurcating divergence and continuous gene flow.

A population reticulation involves multiple stems contributing genetically to the formation of a group. One way in which this can happen is through the geographic expansion of one or both stems. For example, if during MIS 5, either Stem 1S from southern Africa moved northward thereby encountering the Stem 2, or Stem 2 moved from central/western Africa southward into Stem 1S – then we could observe disproportionate ancestry contributions from different stems in modern groups. We observed two merger events. The first, between Stem 1S and Stem 2, results in the formation of an ancestral Khoe-San population 120kya. The second 100kya between Stem 1E and Stem 2, results in the formation of the ancestors of East/West Africans as well as later “Out of Africans”. The rapid rise in sea levels and increased precipitation during MIS 5e, following a glacial period of aridity across Africa³⁰, might have triggered migration inland away from the coasts, as has been suggested for the Paleo-Agulhas plain³¹.

Following these merger events, the stems subsequently fracture into subpopulations which then appear to persist over the past 120kya. These subpopulations can be linked to modern-day groups despite subsequent gene flow across the continent; for example, a genetic lineage sampled in the Nama has an XX probability of being in the ancestral ‘southern’ subpopulation 50kya versus XX probability of being in the ‘eastern’ subpopulation (Figure XX). We also find that Stem 2 continued to contribute to western Africans during the Last Glacial Period, indicative that this gene flow likely occurred in western/central Africa (Table 1).

Contrasting archaic admixture and a weakly structured stem

Evidence for archaic admixture in Eurasia has bolstered the plausibility of archaic admixture having also occurred in Africa. For this reason, previous work has focused on archaic admixture to explain patterns of polymorphism inconsistent with a single origin model. Here, we have shown that weakly structured stem models better capture these patterns. They also help explain an ecological riddle posed by the archaic admixture model. Neanderthal populations were separated from early *Homo sapiens* by thousands of kilometers and continental geographic barriers. By contrast, an archaic hominin population in Africa would need to have stayed in relative reproductive isolation from the human lineage over hundreds of thousands of years despite closer geographic proximity and reproductive compatibility. The weakly structured stem model resolves this ecological riddle by accommodating continuous or recurrent contact between two or more groups present in Africa.

There is evidence for both deleterious and adaptive archaic-derived alleles in modern genomes in the form of a depletion of Neanderthal ancestry in regulatory regions³² or an increased frequency of archaic-related haplotypes such as at *EPAS1* among Tibetans (e.g., Zhang et al.³³). **SG: Probably not the best reference** Under an archaic African admixture model, the estimated 8–10% introgression rate is much higher than Neanderthal gene flow, and would have plausibly been fertile ground for dramatic selection for or against archaic-derived haplotypes over a short period³⁴. By contrast, adaptation under a weakly structured stem would have occurred continuously over much longer periods. Patterns of polymorphism that are inconsistent with the single-stem model predictions have been used to infer putative archaic admixed segments^{11,13,34,15}, negative selection against such segments³⁴, and pervasive positive selection³⁵. However, such approaches are subject to high false positives in the presence of population structure with migration³², and their interpretation should be re-examined in light of a weakly structured stem model in Africa.

Multiple studies have shown a correspondence between phenotypic differentiation, usually assessed with measurements of the cranium, and genetic differentiation among human populations and between humans and Neanderthals^{36,37,38}. If we assume that this correspondence also holds for early human ancestral populations in Africa, the weakly structured stem model would predict that the fossils deriving from each of the two stems should be morphologically quite similar (supplementary text X) **APR: Will we still include this?**, even if they are from different parts of Africa, with differences across stems closer to those across contemporary human populations than to human-Neanderthal differences. Differences within stems would be comparable to differences within recent human populations. Identifying the morphological signature of ancient structure in the fossil record will require careful analysis of many broadly similar remains. Our model further predicts that morphologically distinct crania such as XX and YY are unlikely representatives of branches that contributed appreciably to modern human ancestries. **SG: Here I am being somewhat forceful. We could use conditional to emphasize that these conclusion are under the weakly structured stem model.**

Methods

See Supplement, for now.

Acknowledgements

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