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Contrasting patterns of Y chromosome and mtDNA variation in Africa: evidence for sex-biased demographic processes

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To investigate associations between genetic, linguistic, and geographic variation in Africa, we type 50 Y chromosome SNPs in 1122 individuals from 40 populations representing African geographic and linguistic diversity. We compare these patterns of variation with those that emerge from a similar analysis of published mtDNA HVS1 sequences from 1918 individuals from 39 African populations. For the Y chromosome, Mantel tests reveal a strong partial correlation between genetic and linguistic distances (r=0.33, P=0.001) and no correlation between genetic and geographic distances (r=-0.08, P>0.10). In contrast, mtDNA variation is weakly correlated with both language (r=0.16, P=0.046) and geography (r=0.17, P=0.035). AMOVA indicates that the amount of paternal among-group variation is much higher when populations are grouped by linguistics ($\Phi_{CT} = 0.21$) than by geography ($\Phi_{CT} = 0.06$). Levels of maternal genetic among-group variation are low for both linguistics and geography ($\Phi_{CT} = 0.03$ and 0.04, respectively). When Bantu speakers are removed from these analyses, the correlation with linguistic variation disappears for the Y chromosome and strengthens for mtDNA. These data suggest that patterns of differentiation and gene flow in Africa have differed for men and women in the recent evolutionary past. We infer that sex-biased rates of admixture and/or language borrowing between expanding Bantu farmers and local hunter-gatherers played an important role in influencing patterns of genetic variation during the spread of African agriculture in the last 4000 years.

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Introduction

An important question in population genetics is identifying the best predictors of genetic relationships among human populations. Several studies indicate strong correlations between genetic and linguistic relationships among globally distributed human populations.^{1,2} At the subcon-

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tinental scale, correlations between genetic variation and linguistic or geographic variation differ substantially. Y chromosome studies have shown that geographic distances correlate with genetic affinities among populations in Europe,³ the Americas,⁴ and Austronesia,⁵ whereas language better explains Y chromosome relationships in Siberia.⁶ Mitochondrial DNA (mtDNA) studies suggest that linguistic relationships are better correlated with genetic affinities among South American populations,⁷ while both geography and language are correlated with maternal variation in Austronesia.⁸ In Africa the question of gene-language relationships remains equivocal; some classical genetic and Y chromosome studies point to language,^{9,10} while other Y chromosome and mtDNA studies identify geography^{11,12} as a better predictor of genetic affinities.

The distribution of linguistic variation has been strongly influenced by the Neolithic Revolution, particularly in Africa. Linguistic, archeological, and ethnographic data suggest that all four African language families arose before agriculture in West Africa (Niger-Congo), Northeastern Africa (Afroasiatic), the middle Nile region (Nilo-Saharans), and East Africa (Khoisan).¹³⁻¹⁷ Early dispersals of Niger-Congo, Afroasiatic, and possibly Nilo-Saharan languages are likely associated with migrating farmers.^{14–17} Diamond and Bellwood¹⁵ hypothesized that early farmers replaced the languages of hunter-gatherers living in their path of expansion and that this replacement would lead to strong correlations between linguistic and genetic variation. Their least equivocal example of an association of a language group with the spread of agriculture are the Bantu expansions. Beginning ~ 4000 years ago, farmers speaking Niger-Congo Bantu languages expanded from a southern Cameroonian homeland over most of subequatorial Africa.^{13,18,19} Evidence for the concordant spread of Bantu genes and languages comes from autosomal,^{9,20} mtDNA,^{12,21} and Y chromosomal^{10,11,22-27} data.

The nonrecombining portion of the Y chromosome (NRY) and mtDNA are both haploid and uni-parentally inherited and, hence, are expected to have a four-fold reduction in effective population size (N_e) relative to the autosomes. In the absence of selection, the reduced N_e leads to an increased rate of genetic drift, which makes these haploid regions sensitive indicators of such demographic processes as bottlenecks, population subdivision, and population size and range expansions. The comparative study of patterns of variation at these loci allows the examination of the relative contribution of males and females in shaping African genetic diversity. In this study, we test for associations between genetic, linguistic and geographic differentiation to (1) identify correlates of genetic diversity in Africa, (2) examine the degree of concordance between the Y chromosome and mtDNA, and (3) assess the effects of sex-specific demographic processes shaping patterns of variation.

Subjects and methods Population samples

Samples include representatives of the four major language families: Khoisan, Afroasiatic, Nilo-Saharan, and Niger-Congo (Table 1; Figure 1). Many of the 40 populations in Table 1 were analyzed in previously published studies;^{22,23,28,29} however, several markers were typed in these samples for the first time in the current study. Differences in the number of samples in this and previous studies reflect differences in the availability of DNA, the inclusion of new samples, and/or the merging or splitting of populations according to language or ethnographic criteria. Sampling protocols were approved by the Human Subject Committee at the University of Arizona and those of collaborating institutions.

Y chromosome markers and terminology

Fifty biallelic Y-linked markers, SNPs and indels, were typed using a hierarchical protocol.^{23,26,27} First, we typed mutations defining major haplogroups (eg, haplogroup A defined by M91) and then we typed all markers within a haplogroup until the most derived mutation in that haplogroup was determined (Figure 2). Thus, not every individual was typed for every marker. Markers were typed using allele-specific PCR, restriction enzyme digest, or direct sequencing. Protocols and primer sequences for these assays were previously published.^{23,30} We follow the terminological conventions recommended by the Y Chromosome Consortium³⁰ for naming NRY lineages.

MtDNA

To compare maternally and paternally inherited patterns of variation, we re-examined 366 bp of mtDNA HVS1 sequence data compiled from a number of previous studies.^{12,31–33} The data set includes 39 populations from the major language groups: Khoisan (!Kung1, !Kung2, Khwe, Hadza), Nilo-Saharan (Kanuri, Songhai, Turkana, Nubian, Sudanese, Mbuti, Datoga), Afroasiatic (Moroccan Berber, non-Berber Moroccan, Egyptian, Algerian Mozabite, Tuareg, Somalian, Amhara, Hausa, Podokwo, Mandara, Uldeme, Iragw), Niger-Congo non-Bantu (Fulbe = Fulfulde, Yoruba, Serer, Wolof, Mandinka, Tupuri), and Niger-Congo Bantu (Bubi, Fang, Biaka, Kikuyu, Mozambique1, Mozambique2, Bakaka, Bassa, Mbenzele, Sukuma) (Figure 1). Some populations represented in the original data sets^{12,31–33} were omitted because they are not found on the African mainland, are Cameroonian populations not represented in the Y chromosome data set,³³ or because linguistic designations could not be inferred.

Mantel tests

The correlation among genetic, linguistic, and geographic distances was assessed by the Mantel test³⁴ employing ARLEQUIN 2.000.³⁵ To test whether statistically significant associations between linguistic and genetic affiliations

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Table I Sampled population	Table 1	Sampled	populations
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Geographic regionEthnicityNFeWest Africa	amily liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo	Sublevel Atlantic Dogon Kwa Kwa Kwa Kwa	Latitude/Longitude 14N 15W 15:5N 15W 14N 3W 6N 1E
West AfricaGambia/SenegalWolof34NGambia/SenegalMandinka39NMaliDogon55NGhanaEwe30NGhanaGa29N	liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo	Atlantic Atlantic Dogon Kwa Kwa Kwa	14N 15W 15:5N 15W 14N 3W 6N 1E
Gambia/SenegalWolof34NGambia/SenegalMandinka39NMaliDogon55NGhanaEwe30NGhanaGa29N	liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo	Atlantic Atlantic Dogon Kwa Kwa Kwa	14N 15W 15:5N 15W 14N 3W 6N 1E
Gambia/SenegalMandinka39NMaliDogon55NGhanaEwe30NGhanaGa29N	liger-Congo liger-Congo liger-Congo liger-Congo liger-Congo	Atlantic Dogon Kwa Kwa Kwa	15:5N 15W 14N 3W 6N 1E
MaliDogon55NGhanaEwe30NGhanaGa29N	liger-Congo liger-Congo liger-Congo liger-Congo	Dogon Kwa Kwa Kwa	14N 3W 6N 1E
Ghana Ewe 30 N Ghana Ga 29 N	liger-Congo liger-Congo liger-Congo	Kwa Kwa Kwa	6N 1E
Ghana Ga 29 N	liger-Congo liger-Congo	Kwa Kwa	5 51 05
Change Fourte 22	liger-Congo	Kwa	5:5N 0E
Gnana Fante 32 N			6:5N 1:5E
Central Africa			
Cameroon, North Podokwo 19 A	froasiatic	Chadic	12:5N 14:5E
Cameroon, North Mandara 28 A	froasiatic	Chadic	12:5N 14:5E
Cameroon, North Uldeme 13 A	froasiatic	Chadic	12:5N 14:5E
Cameroon, North Tupuri 9 N	liger-Congo	Gur	10N 13:5F
Cameroon South Bassa 11 N	liger-Congo	Bantu	3N 10F
Cameroon South Bakaka 17 N	liger-Congo	Bantu	3N 10.5F
Cameroon South Ngoumba 31 N	liger-Congo	Bantu	3N 10F
Cameroon South Bakola Pygmies 33 N	liger-Congo	Bantu	2.5N 10.5F
CAR Riaka Pygnies 31 N	liger-Congo	Bantu	3N 16F
CAR Baka Pygmies 18 N	liger-Congo	Bantu	3N 16E
CAN Baka Tyginies 10 N	iger-congo	banta	SINTOL
East Africa	linen Commo	Dentu	1.61 205
DRC INande 18 IN	liger-Congo	Bantu	
DRC Herna 18 IN	liger-Congo	Bantu	
DRC Alur 9 N	llio-Sanaran	INIIOUC Cualantia	I SIN SUE
DRC Midul Pygmies 47 N	liio-sanaran	Sudanic	ZIN Z9E
Tanzania S. Cusnitic 9 A			45 35E
Kenya Massal 26 N	llio-Sanaran	NIIOTIC	25 3/E
Kenya Luo 9 N	llio-sanaran	NIIOTIC	0:55 34:5E
Kenya Kikuyu & Kamba 42 N	liger-Congo	Bantu	1:55 38:5E
Uganda Ganda 26 N	liger-Congo	Bantu	IN 32E
Ethiopia Amhara 18 A	froasiatic	Semitic	10N 39E
Ethiopia Oromo 9 A	froasiatic	Cushitic	6N 39E
Ethiopia South Semitic 20 A	froasiatic	Semitic	12N 38E
South Africa			
Namibia !Kung/Sekele 32 K	hoisan	Northern	19:0/S 13:39E
Namibia Tsumkwe San 29 K	hoisan	Central	19:13S 17:42E
Namibia Dama 18 K	hoisan	Central	24:50S 17:00E
Namibia Nama 11 K	hoisan	Central	24:00S 17:50E
Namibia Herero 24 N	liger-Congo	Bantu	22:30S 18:58E
Namibia Ambo 22 N	liger-Congo	Bantu	22:30S 18:58E
South Africa Sotho-Tswana 28 N	liger-Congo	Bantu	27S 27E
South Africa Zulu 29 N	liger-Congo	Bantu	31:35S 28:47E
South Africa Xhosa 80 N	liger-Congo	Bantu	33:58S 25:36E
Zimbabwe Shona 49 N	liger-Congo	Bantu	17:50S 31:03E
North Africa			
Egypt Egyptian 92 A	froasiatic	Erythraic	30N 30E
Tunisia Tunisian 28 A	froasiatic	Semitic	35N 10E

reflect the same events in population history or parallel, but separate isolation by distance processes, we performed partial correlations holding geography (or language) constant.³⁶ Genetic distances were based on Slatkin's³⁷ linearized Φ_{ST} values (ie, incorporating molecular distances among haplogroups). Geographic distances between populations were calculated using approximate latitude and longitude data for the sample sites (Table 1). We used a novel approach for classifying linguistic relationships

among populations. One of us (CE) constructed tree relationships among the languages spoken by the study populations using several sources of linguistic, archeological, and ethnographic data. Divergence times between related languages were estimated using archeological dates and glottochronological methods.³⁸ Linguistic relationships among populations in this study, as well as among the populations in the mtDNA data set, are available at http://www.u.arizona.edu/~ewood/data.html. We also

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Figure 1 Map of Africa. The approximate location of 40 populations typed for Y chromosome markers in this study (\bullet) and 39 populations surveyed for HVS1 sequence data^{12,31,32,33} (\bigcirc) are indicated. The distribution of the four African language families was constructed using Greenberg's³⁹ classifications and further refined with data from the ethnologue (http://www.ethnologue.com/). Three shades of gray on map refer to the distribution of language families: Khoisan (light gray, southwest), Afroasiatic (light gray, north), Niger-Congo (medium gray), and Nilo-Saharan (dark gray). The circled geographic regions include North, West, Central, East, and South Africa.

performed Mantel tests with matrices constructed using (1) the method described by Poloni *et al*¹⁰ that uses the tree relationships of the languages defined by Greenberg,³⁹ (2) the tree relationships among languages reported in this study without making use of divergence times, (3) equal distances among populations of different language families, and/or (4) variable distances among populations of different language families. All matrices yielded very similar correlations (both *r* and *P* values) for the entire data set. Results differed slightly among matrices when we removed the Bantu speakers.

AMOVA

Analyses of molecular variance (AMOVA) were also performed using ARLEQUIN 2.000.³⁵ Both haplogroup frequencies and molecular differences among haplogroups were taken into account. We grouped populations by five geographic regions (West, Central, East, South, and North Africa) and by four linguistic groups (Afroasiatic, Nilo-Saharan, Khoisan, and Niger-Congo) (Figure 1). All samples used for the Mantel analysis were also used in the AMOVA: 1122 individuals from 40 populations for the Y chromosome and 1918 individuals from 39 populations for mtDNA.

Given that levels of population differentiation can be influenced by (1) sample composition and the Y chromosomal and mtDNA data sets presented here are sampled differently, (2) the differing rates and modes of evolution characterized by the Y chromosome and mtDNA systems, and (3) the type of polymorphisms examined (eg, preascertained Y chromosome SNPs *versus* mtDNA HVS1 sequence data), direct comparisons between these haploid

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Figure 2 Maximum-parsimony tree of 50 Y chromosome biallelic markers typed in this survey. The root of the tree is denoted by an arrow. Major clades (ie, A–R) are labeled with large capital letters. Subclade labels (eg, A3b) are indicated to the left of the branches. Mutation names are given along the branches. The length of each branch is not proportional to the number of mutations or the age of the mutation. Only the names of the 36 haplogroups observed in the present study are shown to the right of the branches. Haplogroup frequencies are shown on the far right.

genetic systems should be considered with caution. Nevertheless, by comparing linguistic and geographic associations within a locus, we can ask whether mtDNA and the Y chromosome have been influenced by similar demographic processes.

Results

Geographic distribution of Y chromosome haplogroups in African populations

Phylogenetic analysis of the 50 Y biallelic markers used in this study yielded 36 haplogroups (Figure 2) (for appendix please refer to Supplementary Information). The vast majority of these lineages (98.1%) belong to five major haplogroups: A (7.1%), B (10.2%), E (70.2%), J (5.4%), and R (5.2%). Haplogroup A is closest to the root of the tree and is found most frequently in the Khoisan, particularly the A2 and A3b1 lineages (47.7%). Haplogroup B chromosomes are most frequently observed in Pygmies (48.9%), with B2a* and B2b* being nearly exclusive to this group. Haplogroup E is overwhelmingly the most common in this study. Over half of the individuals in our study (51%) are members of the subclade E3a, which is defined by the P1 mutation. Niger-Congo speakers have the highest frequency of E-P1* chromosomes (40.7%) and the largest proportion of E-M191 chromosomes (27.5%), particularly in Bantu speakers (31.5%). The E3b1 (E-M78) lineage is most frequent in Afroasiatics (22.5%). In this study, haplogroup J is concentrated in Afroasiatics (19.5%). While African haplogroup R chromosomes are generally quite rare, R-P25* chromosomes are found at remarkably high frequencies in northern Cameroon (60.7-94.7%). The remaining haplogroups (K, F*, I, and G) account for only 1.9% of the individuals in our data set.

Analysis of molecular variance (AMOVA)

The overall Y chromosome Φ_{ST} for the 40 populations is 0.32 (Table 2), a value that is similar to that found in a previous study of African Y-SNP diversity ($\Phi_{ST} = 0.34$).²⁸ This value is also similar to that obtained when our African sample is grouped into five geographic regions. When

populations are grouped according to language family, the proportion of among-group variance ($\Phi_{\rm CT} = 0.21$) is more than three times higher than when populations are grouped according to geographic location ($\Phi_{\rm CT} = 0.06$) (Table 2). AMOVA results for the mtDNA data are also presented in Table 2. The continental mtDNA $\Phi_{\rm ST}$ is 0.15. MtDNA Φ -statistics are very similar when populations are placed in either linguistic or geographic groups.

These results indicate that Y chromosome variation is significantly partitioned among both geographic and linguistic groups. Therefore, both language and, to a lesser extent, geography are probably important (albeit overlapping) predictors of African genetic structure.

Mantel tests

To test the underlying cause of association between genetic and linguistic versus geographic variation, we performed Mantel tests. These tests ask whether there is a correlation between geographic (or language) distance and genetic distance. Mantel tests reveal a statistically significant positive correlation between Y chromosome variation and linguistics (r = 0.32, P = 0.001) that explains 8.9% of the genetic variance. The correlation between genetic and linguistic variation remains strong when geography is held constant (r = 0.33, P = 0.001). In contrast, there is no correlation between paternal genetics and geographic distances (r = 0.01, P > 0.10) (Table 3). Mantel test results based on the mtDNA HVS1 data are also presented in Table 3. The correlation between maternal genetics and linguistics is significant (r = 0.23, P = 0.016), but weakens when geography is held constant (r = 0.16, P = 0.046). Similarly, a significant correlation between mtDNA and geography (r = 0.23, P = 0.008) weakens when linguistics is held constant (r = 0.17, P = 0.035). It is important to note that a failure to find correlations in Mantel tests does not mean that two variables are not related in some way. Rather, it means that processes that might cause a positive correlation (eg, isolation by distance or directional gene flow in the case of geography, or strict language-gene co-

		Within populations		Among populations within groups		Among groups	
Group	No. of groups	Variance (%)	Φ_{ST}	Variance (%)	Φ_{SC}	Variance (%)	Φ_{CT}
Y chromosome	1	68.2	0.32				
Linguistic groups ^a	4	62.1	0.38	16.6	0.21	21.3	0.21
Geographic groups ^b	5	67.4	0.33	26.2	0.28	6.4	0.06
mtDNA	1	84.7	0.15				
Linguistic groups ^a	4	84.0	0.16	13.5	0.14	2.5	0.03
Geographic groups ^b	5	84.1	0.16	12.3	0.13	3.6	0.04

 Table 2
 Analysis of molecular variance (AMOVA)

All Φ -statistics. *P*-values are less than 0.01.

^aAfroasiatic, Khoisan, Niger-Congo, Nilo-Saharan (see Figure 1).

^bWest, Central, East, South, North Africa (see Figure 1).

Table 3Correlation and partial correlation coefficients, r(P-value), between genetic, linguistic, and geographicdistances

	Ŷ	mtDNA
Genetics and Linguistics Genetics and Linguistics, Geography held constant Genetics and Geography Genetics and Geography, Linguistics held constant	0.32 (0.001) 0.33 (0.001) -0.01 (0.508) -0.08 (0.859)	0.23 (0.016) 0.16 (0.046) 0.23 (0.008) 0.17 (0.035)

evolution) are unlikely to be the only processes operating (Tables 2 and 3).

Discussion

Mantel tests show a statistically significant positive correlation between Y chromosome and linguistic variation, while there is no correlation between Y chromosome and geographic variation. Furthermore, when populations are grouped according to language, the amount of amonggroup paternal differentiation (Φ_{CT}) is substantially higher than when grouped according to geographic location. Correlations with mtDNA show a different pattern. Maternal variation is weakly correlated with both language and geography and maternal among-group differentiation is nearly the same when populations are grouped according to linguistic affiliation or geographic location. These results suggest that patterns of differentiation and gene flow in Africa have been different for men and women in the recent evolutionary past.¹⁰ In the following sections, we discuss (1) the relationships among genetic, linguistic, and geographic differentiation and the population history factors that may underlie these relationships, and (2) the effects of Bantu expansions on the distribution of Y chromosome and mtDNA variation in Africa.

Associations between genetic and linguistic variation

The association of genetic and linguistic variation has been observed at the global level,¹ as well as on the regional scale.^{36,40,41} What are the underlying causes of these associations? Sokal⁴¹ stated that a common language usually reflects a common origin for two populations, and a related language indicates a common origin farther back in time. This is generally thought to be an outcome of common historical processes leading to genetic and linguistic diversification – for example, a founding population may reproduce biologically and linguistically in a new location and replace the genes and languages of previous residents.^{2,15,36} Discrepancies between genetic and linguistic differentiation could arise through a number of processes: genetic admixture can occur without language change, languages can be transmitted horizontally without

significant genetic change, and/or genetic and linguistic evolution may proceed at heterogeneous rates.^{2,15,42,43}

We found a statistically significant association between NRY variation and linguistic differentiation and a marginally significant association between mtDNA variation and linguisitic variation. However, when we performed Mantel tests controlling for geographic distance, the partial correlation between maternal genetic and linguistic variation weakens, while that between paternal genetic and linguistic variation remains statistically significant (Table 3). This suggests that the observed association between Y chromosome and language variation reflects the same co-evolutionary population history events.³⁸ These differing patterns for the Y chromosome and mtDNA could be the result of a greater degree of female than male admixture and/or the adoption of languages by females to a greater extent than males (see below). In either case, the implication is that African languages tend to be passed from father to children.¹⁰

Associations between genetic and geographic distances show the opposite trend than do the aforementioned associations between genetic and linguistic variation; there is no correlation between Y chromosome variation and geographic distance. In contrast, there is a stronger correlation between mtDNA variation and geographic distance, albeit only marginally significant when language is held constant (Table 3). Thus, the genetics–language correlation is stronger for the Y chromosome and the different pattern shown by mtDNA data suggests that men and women did not have identical demographic histories.

Effect of Bantu expansions on Y chromosome and mtDNA variation

Numerous studies suggest that the Bantu expansions have had a substantial impact on the distribution of genetic variation in Africa.^{9,10,12,22,25,27} Is the strong Y chromosome-linguistics correlation we observe across the entire continent primarily the result of the massive migrations of Bantu farmers? We sought to clarify the effect of each language group on influencing the paternal geneticslinguistics association and the genetics-geography association by repeating the Mantel test after removing each language group in turn (ie, Afroasiatic, Khoisan, Nilo-Saharan, Niger-Congo non-Bantu, and Niger-Congo Bantu). If one language group were disproportionately contributing to the overall pattern, then the association is expected to weaken upon removal of this group. There is only a single language group that has this effect: the removal of Bantu-speakers causes the paternal geneticslinguistics correlation to drop from r = 0.33 to 0.08 (Figure 3).We note that the same trend is observed, albeit to a lesser extent, when other language matrices (see Subjects and Methods) are employed in the Mantel tests (data not shown). The lower correlation coefficient when Bantu populations (but not other linguistic groups) are

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Figure 3 Partial correlation coefficient between genetics and linguistics holding geography constant (black bars) and genetics and geography holding linguistics constant (white bars) for the Y chromosome and mtDNA. **P<0.01, *P<0.05.

removed suggests that Bantu are contributing more to the language–Y chromosome relationship than any other language group.

The Y chromosome-geography correlation shows a different pattern. While the removal of the Bantu populations does not produce a correlation, the additional removal of four northern Cameroonian populations results in a statistically significant positive correlation (r = 0.27, P = 0.008). The strong effect of these northern Cameroonian populations on the Y chromosome results can be explained by the very high frequency of derived paternal (but not maternal) lineages that originated in non-African populations.^{27,33} We note that this increased geographical correlation is not entirely attributable to the northern Cameroonian populations because when only these populations are removed, there is no Y chromosome-geography correlation (r = -0.002, P > 0.10). Thus, in the absence of the unique populations from northern Cameroon, the removal of Bantu speakers leads to an association between Y chromosome and geographic differentiation, consistent with a recent dispersal of Bantu Y chromosomes.

On the other hand, the exclusion of Bantu speakers strengthens both the mtDNA–linguistics and mtDNA–geography correlations (Figure 3). Upon further exploration, we discovered that the increase in both of these maternal genetic correlations is due solely to Bantu-speaking Pygmy populations, specifically, the Biaka and Mbenzele (data not shown). The strong effect of these Pygmy populations on the mtDNA–linguistics correlation suggests that the horizontal transfer of languages from Bantu farmers to hunter-gatherer Pygmy females¹⁹ occurred without significant genetic change.^{19,31,36} The stronger mtDNA associations with both linguistic and geographic variation observed when the Biaka and Mbenzele popula-

tions are removed may reflect the fact that these two populations are maternal genetic outliers.³¹

To further investigate the effect of the Bantu expansions on patterns of geographic variation, we grouped populations by their geographic location and removed each language group in a series of four AMOVA runs (data not shown). Unlike the case for any other language family, the removal of Bantu populations results in a higher Y chromosome $\Phi_{\rm CT}$ (0.28) than when they are included (0.06). This supports the hypothesis that Bantu Y chromosomes (eg E-P1*, E-M191) are acting to homogenize geographically differentiated populations. A similar analysis of mtDNA results in slightly higher $\Phi_{\rm CT}$ value when the Bantu populations are excluded (0.07 *versus* 0.04).

If Bantu males and females dispersed equally from their West African homeland, replacing the genes of local hunter-gatherers in their path of expansion (equal sex ratio model), then we would expect similar patterns of association for paternally and maternally inherited loci. If, on the other hand, one sex dispersed more effectively (sexbiased model), we would expect to find differences in the degree of association between genetic and linguistic variation for the two haploid loci. Several explanations have been offered for observed differences in patterns of Y chromosome and mtDNA variation among populations.^{31,44-46} Our results support the sex-biased model whereby the replacement of pre-existing languages by Bantu languages more closely parallels the turnover of Y chromosomes than mtDNA. How can this be explained? One possibility is that Bantu male farmers dispersed over longer distances or in greater numbers than Bantu females. Another possibility is that males and females dispersed equally, but there was a higher 'effective' migration rate for Bantu Y chromosomes than Bantu mtDNA. As Bantu farmers dispersed, they likely intermarried to some extent with the original inhabitants related to modern Pygmies and Khoisan.¹⁵ In present-day African populations, the direction of intermarriage is usually between huntergatherer women and farmer (Bantu) men and the children of these marriages generally become farmers residing in their father's village^{19,47} (ie, patrilocality). If this were typical of practices that existed throughout the Bantu expansions, we would expect Bantu mtDNA to be diluted (with hunter-gatherer mtDNA) to a greater extent than Y chromosomes. It is also possible that if the ancestral Bantuspeaking population were highly polygynous, then indigenous Y chromosomes would have been replaced by a more homogeneous pool of Bantu Y chromosomes, leading to a stronger correlation between linguistic and genetic variation. Indeed, polygyny is known to be substantially higher among food-producers than among hunter-gatherers.³¹ In combination with the above processes, the adoption of Bantu languages by hunter-gatherer Pygmies may weaken maternal genetic-linguistic associations. Although our data cannot address the relative impact of

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While earlier studies of Y chromosome variation have noted a correspondence between high-frequency haplogroups and the distribution of Bantu speakers (ie E-P1* and E-M191),^{11,22–27} this is the first study to demonstrate a statistically significant correlation between Y chromosome SNP haplogroups³⁰ and linguistic differentiation in Africa. The data presented here are consistent with the hypothesis that prehistoric agriculture dispersed hand-in-hand with Bantu languages and Y chromosomes, with languages and Y chromosomes replacing those of hunter-gatherers in the paths of expansion.¹⁵ Not all populations speaking Bantu languages in our study showed the effects of complete paternal genetic replacement (eg, the Bantu-speaking western Pygmies and northern Cameroonians). It is important to note that different mutation rates, as well as methods used to assay variation, on the NRY and mtDNA may contribute to some of the contrasting patterns observed here.46 Future studies that examine Y chromosome and mitochondrial DNA sequence variation in the same samples representing African geographic and linguistic diversity will help to further elucidate the effects of Bantu expansions on the complex genetic landscape of Africa.

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Supplementary Information accompanies the paper on European Journal of Human Genetics website (http://www.nature.com/ejhg).