ARTICLE

The phylogenetic and geographic structure of Y-chromosome haplogroup R1a

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R1a-M420 is one of the most widely spread Y-chromosome haplogroups; however, its substructure within Europe and Asia has remained poorly characterized. Using a panel of 16 244 male subjects from 126 populations sampled across Eurasia, we identified 2923 R1a-M420 Y-chromosomes and analyzed them to a highly granular phylogeographic resolution. Whole Y-chromosome sequence analysis of eight R1a and five R1b individuals suggests a divergence time of \sim 25 000 (95% CI: 21 300–29 000) years ago and a coalescence time within R1a-M417 of \sim 5800 (95% CI: 4800–6800) years. The spatial frequency distributions of R1a sub-haplogroups conclusively indicate two major groups, one found primarily in Europe and the other confined to Central and South Asia. Beyond the major European versus Asian dichotomy, we describe several younger sub-haplogroups. Based on spatial distributions and diversity patterns within the R1a-M420 clade, particularly rare basal branches detected primarily within Iran and eastern Turkey, we conclude that the initial episodes of haplogroup R1a diversification likely occurred in the vicinity of present-day Iran.

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INTRODUCTION

High-throughput resequencing efforts have uncovered thousands of Y-chromosome variants that have enhanced our understanding of this most informative locus' phylogeny, both through the resolution of topological ambiguities and by enabling unbiased estimation of branch lengths, which, in turn, permit timing estimates.^{1–5} The International Society of Genetic Genealogy¹⁰ has aggregated these variants and those discovered with previous technologies into a public resource that population surveys can leverage to further elucidate the geographic origins of and structure within haplogroups.^{6–13}

Y-chromosome haplogroup R (hg R) is one of 20 that comprise the standardized global phylogeny.¹⁴ It consists of two main components: R1-M173 and R2-M479¹⁵ (Figure 1). Within R1-M173, most variation extant in Eurasia is confined to R1a-M420 and R1b-M343.¹⁶ In Europe, R1a is most frequent in the east, and R1b predominates in the west.¹⁷ It has been suggested that this division reflects episodic population expansions during the post-glacial period,

including those associated with the establishment of agricultural/ pastoral economies. $^{3,18-21}_{\ }$

More than 10% of men in a region extending from South Asia to Scandinavia share a common ancestor in hg R1a-M420, and the vast majority fall within the R1a1-M17/M198 subclade.²² Although the phylogeography of R1b-M343 has been described, especially in Western and Central Europe,^{15,23–25} R1a1 has remained poorly characterized. Previous work has been limited to a Europeanspecific subgroup defined by the single-nucleotide polymorphism (SNP) called M458.^{22,26–30} However, with the discovery of the Z280 and Z93 substitutions within Phase 1 1000 Genomes Project data¹ and subsequent genotyping of these SNPs in \sim 200 samples, a schism between European and Asian R1a chromosomes has emerged.³¹ We have evaluated this division in a larger panel of populations, estimated the split time, and mapped the distributions of downstream sub-hgs within seven regions: Western/Northern Europe, Eastern Europe, Central/South Europe, the Near/Middle East, the Caucasus, South Asia, and Central Asia/Southern Siberia.

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Figure 1 Haplogroup (hg) R1a-M420 topology, shown within the context of hg R-M207. Common names of the SNPs discussed in this study are shown along the branches, with those genotyped presented in color and those for which phylogenetic placement was previously unknown in orange. Hg labels are assigned according to YCC nomenclature principles with an asterisk (*) denoting a paragroup.⁶³ Dashed lines indicate lineages not observed in our sample. The marker Z280 was not used as it maps to duplicated ampliconic tracts.

MATERIALS AND METHODS

Population samples

We assembled a genotyping panel of 16244 males from 126 Eurasian populations, some of which we report upon for the first time herein and others that we have combined from earlier studies,^{22,29–30,32–45} and updated to a higher level of phylogenetic resolution. All samples were obtained using locally approved informed consent and were de-identified.

Whole Y-chromosome sequencing

We analyzed 13 whole R1 Y-chromosome sequences: 8 novel, 2 previously published,⁴ and 3 from the 1000 Genomes Project.² All sequences were generated on the Illumina HiSeq platform (Illumina, San Diego, CA, USA) using libraries prepared either from genomic DNA or from flow sorted Y-chromosomes drawn from lymphoblast cell line cultures induced to the metaphase cell division stage (Supplementary Table 1). We used Bowtie⁴⁶ to

map 101-bp sequencing reads to the hg19 human reference, and we called genotypes across 9.99 Mb and estimated coalescence times using a rate of 1 SNP accumulating per 122 years as described in Poznik *et al.*⁴

SNP analysis

We selected binary markers (Supplementary Table 2) from the International Society of Genetic Genealogy¹⁰ database and from whole Y-chromosome sequencing and genotyped samples either by direct Sanger sequencing or RFLP assays. Within the full panel, 2923 individuals were found to be members of hg R1a-M420. These M420 carriers were then genotyped in a hierarchical manner (Figure 1) for the following downstream markers with known placement on the tree: SRY10831.2, M17, M198, M417, Page7, Z282/S198, Z284/S221, M458, M558/CTS3607, Z93/S202, Z95, Z2125, M434, M560, M780, M582, and for three SNPs whose placement within the R1a topology was previously unknown: M746, M204, and L657.

We generated spatial frequency maps for the R1a subgroups that we determined to occur at 10% frequency or greater within a studied region. To do so, we used Surfer (v.8, Golden Software, Inc, Golden, CO, USA) with the *kriging* algorithm and the option to use bodies of water as break-lines. We carried out spatial autocorrelation analysis to detect clines by calculating Moran's I coefficient using PASSAGE v.1.1 (www.passagesoftware.net) with a binary weight matrix, 10 distance classes, and the assumption of a random distribution. Haplogroup diversities were calculated using the method of Nei.⁴⁷ To investigate the genetic affinities among populations, we used the freeware popSTR program (http://harpending.humanevo.utah.edu/popstr/) to perform a principal component analysis (PCA) based on R1a subgroup frequencies.

Short tandem repeat (STR) analysis

We genotyped 1355 samples for 10–19 STRs (Y-STRs; Supplementary Table 3) and calculated haplotype diversities, also using the method of Nei.⁴⁷ Coalescence times (T_d) of R1a subhaplogroups were estimated using the ASD₀ methodology described by Zhivotovsky *et al*⁴⁸ and modified according to Sengupta *et al.*⁴¹ Given the uncertainty associated with Y-STR mutation rates,²⁴ we used both the evolutionary effective mutation rate of 6.9×10^{-4} per 25 years⁴⁸ and, for comparison, the pedigree mutation rate of 2.5×10^{-3} per generation.⁴⁹

RESULTS AND DISCUSSION

Refinement of hg R1a topology

Figure 1 shows, in context, the phylogenetic relationships of the markers we genotyped in this study. These include three for which phylogenetic placement was previously unknown: M746, M204, and L657. We localized the rare M204 SNP based on a single Iranian sample confirmed by Sanger sequencing to carry the derived allele.^{37,50}

Phylogeography

We measured R1a haplogroup frequency by population (Supplementary Table 4). Of the 2923 hg R1a-M420 samples, 2893 were derived for the M417/Page7 mutations (1693 non-Roma Europeans and 1200 pan-Asians), whereas the more basal subgroups were rare. We observed just 24 R1a*-M420(xSRY10831.2), 6 R1a1*-SRY10831.2(xM198), and 12 R1a1a1-M417/Page7*(xZ282,Z93). We did not observe a single instance of R1a1a-M198*(xM417,Page7), but we cannot exclude the possibility of its existence.

Of the 1693 European R1a-M417/Page7 samples, more than 96% were assigned to R1a-Z282 (Figure 2), whereas 98.4% of the 490 Central and South Asian R1a lineages belonged to hg R1a-Z93 (Figure 3), consistent with the previously proposed trend.³¹ Both of these haplogroups were found among Near/Middle East and Caucasus populations comprising 560 samples.

Subgroups of both R1a-Z282 and R1a-Z93 exhibit geographic localization within the broad distribution zone of R1a-M417/Page7.

Among R1a-Z282 subgroups (Figure 2), the highest frequencies (\sim 20%) of paragroup R1a-Z282* chromosomes occur in northern Ukraine, Belarus, and Russia (Figure 2b). The R1a-Z284 subgroup (Figure 2c) is confined to Northwest Europe and peaks at $\sim 20\%$ in Norway, where the majority of R1a chromosomes (24/26) belong to this clade. We found R1a-Z284 to be extremely rare outside Scandinavia. R1a-M458 (Figure 2d) and R1a-M558 (Figure 2e) have similar distributions, with the highest frequencies observed in Central and Eastern Europe. R1a-M458 exceeds 20% in the Czech Republic, Slovakia, Poland, and Western Belarus. The lineage averages 11-15% across Russia and Ukraine and occurs at 7% or less elsewhere (Figure 2d). Unlike hg R1a-M458, the R1a-M558 clade is also common in the Volga-Uralic populations. R1a-M558 occurs at 10-33% in parts of Russia, exceeds 26% in Poland and Western Belarus, and varies between 10 and 23% in the Ukraine, whereas it drops 10-fold lower in Western Europe. In general, both R1a-M458 and R1a-M558 occur at low but informative frequencies in Balkan populations with known Slavonic heritage. The rarity of R1a-M458 and R1a-M558 among Central Asian and South Siberian R1a samples (4/301; Supplementary Table 4) suggests low levels of historic Slavic gene flow.

In the complementary R1a-Z93 haplogroup, the paragroup R1a-Z93* (Figure 3b) is most common (>30%) in the South Siberian Altai region of Russia, but it also occurs in Kyrgyzstan (6%) and in all Iranian populations (1–8%). R1a-Z2125 (Figure 3c) occurs at highest frequencies in Kyrgyzstan and in Afghan Pashtuns (>40%). We also observed it at greater than 10% frequency in other Afghan ethnic groups and in some populations in the Caucasus and Iran. Notably, R1a-M780 (Figure 3d) occurs at high frequency in South Asia: India, Pakistan, Afghanistan, and the Himalayas. The group also occurs at >3% in some Iranian populations and is present at >30% in Roma from Croatia and Hungary, consistent with previous studies reporting the presence of R1a-Z93 in Roma.^{31,51} Finally, the rare R1a-M560 was only observed in four samples: two Burushaski speakers from north Pakistan, one Hazara from Afghanistan, and one Iranian Azeri.

Y-STR haplotype networks and diversity

We genotyped a subset of 1355 R1a samples for 10–19 Y-chromosome STR loci (Supplementary Table 3) and constructed networks for both hg R1a-Z282 and hg R1a-Z93 (Supplementary Figure 1 and Supplementary Figure 2). Although we could assign haplotypes to various haplogroups, power to identify substructure within hg R1a-M198 was limited, consistent with previous work.^{22,52} Although haplotype diversity is generally very high (H>0.95) in all haplogroups (Supplementary Table 3), lower diversities occur in south Siberian paragroup R1a-Z93* (H=0.921), in Jewish R1a-M582 (H=0.844) and in Roma R1a-M780 (H=0.759), consistent with founder effects that are evident in the network patterns for these populations (Supplementary Figure 2).

Origin of hg R1a

To infer the geographic origin of hg R1a-M420, we identified populations harboring at least one of the two most basal haplogroups and possessing high haplogroup diversity. Among the 120 populations with sample sizes of at least 50 individuals and with at least 10% occurrence of R1a, just 6 met these criteria, and 5 of these 6 populations reside in modern-day Iran. Haplogroup diversities among the six populations ranged from 0.78 to 0.86 (Supplementary Table 4). Of the 24 R1a-M420*(xSRY10831.2) chromosomes in our data set, 18 were sampled in Iran and 3 were from eastern Turkey. Similarly, five of the six observed R1a1-SRY10831.2*(xM417/Page7) chromosomes



Figure 2 (a–e) Spatial frequency distributions of Z282 affiliated haplogroups. Each map was generated using the frequencies from Supplementary Table 4 among 14461 individuals, distributed across 119 population samples (references listed in Supplementary Table 4). Because of the known difference between the origin and present distribution of the Roma and Jewish populations, they were excluded from the plots. Additional populations from literature²⁷ were used for the M458 map.

were also from Iran, with the sixth occurring in a Kabardin individual from the Caucasus. Owing to the prevalence of basal lineages and the high levels of haplogroup diversities in the region, we find a compelling case for the Middle East, possibly near present-day Iran, as the geographic origin of hg R1a.

Spatial dynamics of R1a lineage frequencies

We conducted a spatial autocorrelation analysis of the two primary subgroups of R1a (Z282 and Z93) and of each of their subgroups independently (Supplementary Figure 3). Each correlogram was statistically significant. We observed clinal distributions (continually decreasing frequency with increasing geographic distance) across a large geographic area in the two macrogroups and in M558 and M780 as well. One group (Z2125) did not reveal any discernible pattern, and the analysis of four groups (Z282*, Z284, M458, and Z93*) indicated potential clinal distributions that do not extend across the full geographic range under study. Therefore, we also analyzed partial ranges for Z282* and M458 in Europe, the Caucasus, and the Middle East, and for Z284 in Europe, but these partial range analyses also failed to yield evidence of clinal distributions.

We also conducted PCA of R1a subgroups (Figure 4). The first principal component explains 21% of the variation and separates European populations at one extreme from those of South Asia at the other. The second explains 14.7% of the variation and is driven almost exclusively by the high presence of M582 among some Jewish populations, particularly the Ashkenazi Jews. PC2 separates them from all other populations. When we consider haplogroups rather than populations (Supplementary Figure 4), we see that the clustering of European populations is due to their high frequencies of M558, M458, and Z282*, whereas the M780 and Z2125* lineages account for the South Asian character of the other extreme.

To put our frequency distribution maps, PCA analyses, and autocorrelation results in archaeological context, we note that the earliest R1a lineages (genotyped at just SRY10381.2) found thus far in European ancient DNA date to 4600 years before present (YBP), a time corresponding to the Corded Ware Culture,⁵³ whereas three



Figure 3 (a-d) Spatial frequency distributions of Z93 affiliated haplogroups. Maps were generated as described in Figure 2.

DNA sample extracts from the earlier Neolithic Linear Pottery Culture (7500–6500 YBP) period were reported as G2a-P15 and F-M89(xP-M45) lineages.⁵⁴ This raises the possibility of a wide and rapid spread of R1a-Z282-related lineages being associated with prevalent Copper and Early Bronze Age societies that ranged from the Rhine River in the west to the Volga River in the east⁵⁵ including the Bronze Age Proto-Slavic culture that arose in Central Europe near the Vistula River.⁵⁶ It may have been in this cultural context that hg R1a-Z282 diversified in Central and Eastern Europe. The corresponding diversification in the Middle East and South Asia is more obscure. However, early urbanization within the Indus Valley also occurred at this time⁵⁷ and the geographic distribution of R1a-M780 (Figure 3d) may reflect this.

To evaluate the potential role of R1a diversification in these post-Neolithic events, we took two approaches toward estimating the time to the most recent common ancestor (TMRCA). The first was a Y-STR-based coalescent time estimation, the results of which (Supplementary Table 5) demonstrate the unsuitability of the pedigree mutation rate, as supported also by the evidence in Wei *et al*,³ the ages being severely underestimated. Alternatively times based on the evolutionary mutation rate,⁴⁸ which is prone to overestimation, should be regarded as the upper bounds on the sub-hg dispersals. The second approach was TMRCA estimation based on whole Y-chromosome sequencing data.

Whole Y-chromosome sequences from R1a and R1b: TMRCA estimates

The SNPs that we genotyped across 126 populations reveal considerable information about the topology of the haplogroup tree, but they were ascertained in a biased manner, and they are too few in number to convey any meaningful branch-length information. Hence, our SNP genotyping results are devoid of temporal information. To obtain unbiased branch lengths to estimate TMRCA, we analyzed whole Y-chromosome sequences (9.99 Mb of which were usable) of 13 individuals: 8 R1a and 5 R1b. We used MEGA⁵⁷ to construct a bootstrap consensus maximum likelihood tree (Figure 5) based on 928 R1 SNPs (Supplementary Data File 1), of which 462 were previously named.¹⁰ To define the ancestral and derived states of SNPs corresponding to the roots of the R1a and R1b subtrees (branches 23 and 8 in Figure 5, respectively), we called genotypes and constructed the tree jointly with previously published hg E sequences,⁴ which constituted an outgroup.



Figure 4 Principal component analysis of hg R1a subclades. The plot was obtained by collapsing the 126 populations into 49 regionally/culturally defined groups and calculating R1a subclade frequencies relative to R1a-M198. We excluded one population with small overall sample size and all populations in which fewer than 5 R1a Y-chromosomes were observed.



Figure 5 Y-chromosome phylogeny inferred from 13 \sim 10-Mb sequences of hg R individuals. Branches are drawn proportional to the number of derived variants. Each of the 24 branches is labeled by an index, and the number of SNPs assigned to the branch is shown in brackets. The tips of the tree are labeled with sequencing coverage, population, ID, and the most derived commonly known SNP observed in the corresponding sample.

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A consensus has not yet been reached on the rate at which Y-chromosome SNPs accumulate within this 9.99 Mb sequence. Recent estimates include one SNP per: ~100 years,⁵⁸ 122 years,⁴ 151 years⁵ (deep sequencing reanalysis rate), and 162 years.⁵⁹ Using a rate of one SNP per 122 years, and based on an average branch length of 206 SNPs from the common ancestor of the 13 sequences, we estimate the bifurcation of R1 into R1a and R1b to have occurred ~25 100 ago (95% CI: 21 300–29 000). Using the 8 R1a lineages, with an average length of 48 SNPs accumulated since the common ancestor, we estimate the splintering of R1a-M417 to have occurred rather recently, ~5800 years ago (95% CI: 4800–6800). The slowest mutation rate estimate would inflate these time estimates by one-third, and the fastest would deflate them by 17%.

With reference to Figure 1, all fully sequenced R1a individuals share SNPs from M420 to M417. Below branch 23 in Figure 5, we see a split between Europeans, defined by Z282 (branch 22), and Asians, defined by Z93 and M746 (branch 19; Z95, which was used in the population survey, would also map to branch 19, but it falls just outside an inclusion boundary for the sequencing data⁴). Star-like branching near the root of the Asian subtree suggests rapid growth and dispersal. The four subhaplogroups of Z93 (branches 9-M582, 10-M560, 12-Z2125, and 17-M780, L657) constitute a multifurcation unresolved by 10 Mb of sequencing; it is likely that no further resolution of this part of the tree will be possible with current technology. Similarly, the shared European branch has just three SNPs.

We caution against ascribing findings from a contemporary phylogenetic cluster of a single genetic locus to a particular pre-historic demographic event, population migration, or cultural transformation. The R1a TMRCA estimates we report have wide confidence intervals and should be viewed as preliminary; one must sequence tens of additional R1a samples to high coverage to uncover additional informative substructure and to bolster the accuracy of the branch lengths associated with the more terminal portions of the phylogeny. Although some of the SNPs on the lineages we have defined by single SNPs are undoubtedly rare (eg, the Z2125 sub-hg M434, Figure 1, Supplementary Table 4), it remains possible that future genotyping effort using the SNPs in Supplementary Data File 1 may expose other substructure at substantial frequency, commensurate with more recent episodes of population growth and movement. In addition, high coverage sequences using multiple male pedigrees sampled across various haplogroups in the global Y phylogeny will be needed to more accurately estimate the Y-chromosome mutation rate. Nonetheless, despite the limitations of our small sample of R1a sequences, the relative shortness of the branches and their geographic distributions are consistent with a model of recent R1a diversification coincident with range expansions and population growth across Eurasia.

CONCLUSION

Our phylogeographic data lead us to conclude that the initial episodes of R1a-M420 diversification occurred in the vicinity of Iran and Eastern Turkey, and we estimate that diversification downstream of M417/Page7 occurred ~5800 years ago. This suggests the possibility that R1a lineages accompanied demic expansions initiated during the Copper, Bronze, and Iron ages, partially replacing previous Y-chromosome strata, an interpretation consistent with albeit limited ancient DNA evidence.^{54,60} However, our data do not enable us to directly ascribe the patterns of R1a geographic spread to specific prehistoric cultures or more recent demographic events. Highthroughput sequencing studies of more R1a lineages will lead to further insight into the structure of the underlying tree, and ancient

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DNA specimens will help adjudicate the molecular clock calibration. Together these advancements will yield more refined inferences about pre-historic dispersals of peoples, their material cultures, and languages.^{57,61,62}

CONFLICT OF INTEREST

PAU consulted for and has stock in, and CDB is on the advisory board of a project at 23andMe. CDB is on the scientific advisory boards of Personalis, Inc.; InVitae (formerly Locus Development, Inc.); and Ancestry DNA. The remaining authors declare no conflict of interest.

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