Ancient DNAs and the Neolithic Chinese super-grandfather Y haplotypes

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Abstract

Background: Previous studies identified 3 Neolithic Han Chinese super-grandfather Y haplotypes, F5 (O2a2b1a1a), F46 (O2a2b1a2a1), and F11 (O2a1b1a1a1a), but their relationships with the archaeological and written records remain unexplored. **Results:** We here report genome wide DNA data for 11 ancient human samples (0.02x-1.28x) from China ranging from 6500 to 2500 years before present (YBP), which also includes ~11 Mb Y NRY data for one sample. The 11 ancient samples belonged to 4 different genetic groups, designated as Dashanqian (DSQ) of Xiajiadian Culture in the Northeast, Duzhong (DZ) of late Yangshao Culture in the Central Plains, Zhengzhou Xishan (ZX) of Miaodigou Culture in the Central Plains, and Others. Present day F5 samples were closer in autosomal distances to the ZX and DSQ groups while O1, O2, and C samples were closer to the DZ group. We also sequenced the Y chromosome of one of these ancient samples K12 from DSQ and found both K12 and a previously reported ~4000 year old sample MG48 from Northwest China to have the F2137 (O2a2b1a1a1a2a) haplotype, belonging to the most prolific branch F438 immediately under F5. We further found close relationships between ZX and DSQ and between ZX and ancient M117 Tibetans or present day Southwest Dai Chinese carrying the F5 subtype O2a2b1a1a6. **Conclusions:** The results imply radiations of F5 subtypes from the putative place of F5 origin in Henan. These results are remarkably consistent with archaeological and written records.

Background

There are numerous data for human activity in China from the time of the Neolithic period to the beginning of written records. There were the Gaomiao and Pengtoushan Culture of ~5800 BC in the South (Hunan), the Jiahu Culture and Peiligang Culture of 7000-5000 BC in the Central Plains (Henan), the Xinglongwa Culture of 6200-5400 BC and later the Hongshan and Xiajiadian Cultures in the Northeast (Inner Mongolia-Liaoning border), and the Dadiwan Culture of 5800-5400 BC in Gansu and Western Shaanxi [1]. At 5000 BC to 3000 BC, the Yangshao Culture was the most popular and existed extensively along the Yellow River in China and flourished mainly in the provinces of Henan, Shaanxi and Shanxi with the early period of the Culture mostly found in Shaanxi and the late period in Henan [1]. Elements of the middle to late Yangshao Culture, the Miaodigou Culture, have been found widely in China, including the Hongshan Culture in the Northeast (3500 BC), indicating the broad cultural migration and influence of this Culture [2].

By analyzing the Y chromosome haplotype patterns, three Neolithic super-grandfather haplotypes have been discovered that together account for ~40% of present day Han Chinese males [3-5]. The expansion dates are estimated 5400 YBP (years before present) for O3a2c1a-M117-F5 (O2a2b1a1a-F5 or F8, ISOGG 2017), 6500 YBP for O3a2c1-F46 (O2a2b1a2a1-F46), or 6800 YBP for O3a1c-002611-F11 (O2a1b1a1a1a-F11), and these three haplotypes represent 16%, 11%, and 14% of present day Han Chinese males, respectively. Several historiographies on ancient Chinese mention the great leaders/ancestors around the time of 5000 years ago or earlier, including *Yan* Emperor (Yandi), *Huang* Emperor (Huangdi), and *Chi You*.

It remains unclear how the Neolithic Cultures were related to the super-grandfather haplotypes and how the different Neolithic Cultures were interconnected. We here addressed these questions by analyzing ancient DNA samples from 9 different sites in Central and Northern China. We found evidence of F5 associated autosomes in the Miaodigou Culture in Henan and both F5 haplotype and F5 associated autosomes in the Xiajiadian Culture in Inner Mongolia. The results provide a coherent account of information from the relevant fields.

Results

Relationships among aDNAs in Central and Northern China

We collected 11 skeletal remains from 9 sites in Central and Northern China that were 2500-6500 years old (Table 1 and Figure 1). DNAs were extracted and sequenced to different degrees of coverage (0.018x-1.27x). SNPs were called using the Hg19 reference genome. We obtained two sets of SNPs, one slow and the other fast, with the slow set informative for phylogenetics and the fast set representing natural selection and maximum saturation as detailed before [6-12]. Slow evolving SNPs are yet to reach mutation saturation and hence better qualified for phylogenetic inferences. That the slow set and the fast set can give rise to different results or conclusions was strong justification for using the slow set or for distinguishing the two sets. We then merged the SNPs of each of the aDNA samples with those of the 1000 genomes project (1kGP) [13], and obtained pairwise genetic distances, i.e., mismatch in homozygous (hom) sites for slow SNPs or mismatch in all sites for fast SNPs, between each aDNA sample and all individuals in the 1kGP.

Table 1. Information on ancient samples for which we report the nuclear sequence data in this study

Figure 1. Locations of ancient samples from China

Because different ancient samples were sequenced at different coverages, it is unrealistic to use SNP mismatches to infer relationships as there would be few shared SNPs among different samples. As an alternative, we calculated the correlation coefficient R of two genomes in their distance to the East Asian (ASN) samples in 1kGP, assuming that different random sampling of a fraction of the whole set of ~15K slow SNPs are roughly equivalent in representing the whole set. Verification of this R correlation approach has been described previously [6].

We tested the correlation of ancient samples within themselves relative to their correlation with present day Han Chinese in order to better infer the relationships among the ancient samples. We obtained for each aDNA R values with each other and with the 211 Han Chinese samples in 1kGP. We then ranked these R values as shown in Table 2, which then allowed us to infer genetic relationships among these aDNAs. We grouped these 11 aDNAs into 4 groups based on being from the same site and being closely related to each other as indicated by correlation rankings. In Table 2, rank values means the rank of a sample on the column among all 222 samples in values of correlation to a sample listed on the row, e.g., K2 from the column was ranked 7th among all samples in correlation with K12 from the row. The 4 groups were designated as the Dashanqian (DSQ) group including the 2 samples from the Xiajiadian Culture in Dashanqian in Chifeng (K2 and K12) known to be related to the Hongshan Culture [14], one sample S1 from the Xiaoheyuan Culture at the Hala Haigou site in Chifeng related to the Hongshan Culture [15] and one sample FQ17 from the East Zhou dynasty at Fushan Qiaobei Shanxi [16]; the Duzhong (DZ) group including a late Yangshao sample DZ14 from Duzhong Henan [17], one sample SG2046 from Sanguan Hebei known to be influenced by both Yangshao and Xiajiadian [18], one sample ZK22 from Zhukaigou Inner Mongolia [19], and one sample H69 from the early Shang site at Xuecun Henan [20]; the Zhengzhou Xishan (ZX) group including 2 samples from the same Zhengzhou Xishan site of Miaodigou Culture in the Central Plains (ZX167 and ZX107) [21]; and finally the Other group including PAB3002 of Gaotaishan Liaoning [22].

Table 2. Ranks in correlation between each pair of ancient samples among 211 Han Chinese in 1KG and 11 ancient samples. Rank values listed means the rank of a sample on the column among all 223 samples in values of correlation to a sample listed on the row.

These groupings mostly showed expected relationships based on geophysical distances, for example, the close genetic relationship of the 3 Chifeng samples in the DSQ group. Also, FQ17 of Shanxi in the DSQ group was located in the Y shaped region known to be a common migration route linking the Central Plains with the Northeast [23]. A notable exception was ZX167 who had K2 and FQ17 from DSQ group among the most related aDNA samples rather than ZX107 from the same site (K2 and FQ17 ranked higher than all other aDNAs in R correlation to ZX167, being at 69th and 74th, respectively). However, ZX107 did have ZX167 as the most related aDNA (the ranking of 97th was the highest among all aDNAs in R correlation to ZX107), consistent with the geographical location.

ZX167 was the 172th ranked in relation to K2 and the closest to K2 among non DSQ samples. K2 was the 69th ranked in relation to ZX167 and the closest to ZX167 among all aDNAs samples here. This indicated gene flow between DSQ and ZX, which is consistent with the known archaeological findings of a migration of Miaodigou Culture to the Niuheliang site in Chifeng Inner Mongolia [2].

Relationships between aDNAs and present day samples carrying the super-grandfather haplotypes

We first determined to which present day populations the ancient Chinese samples may be closely related. We used the informative slow SNPs to calculate pairwise genetic distances between each ancient DNA and each of the 1kGP samples as described previously [6, 7]. For samples such as K2 with relatively large number of slow SNPs covered, we were able to perform informative distance analyses and principle components analysis (PCA). The results showed K2 to be closest to East Asians in genetic distances (Figure 2A). K2 was closer to Europeans than most East Asians as shown by PCA (Figure 2B), which is consistent with the location of K2 at Chifeng, Inner Mongolia where East and West admixture are known to have occurred [24]. Interestingly, K2 was more related to South East Asians such as KHV (Kihn Vietnamese) and Hunan people and least related to Northern Chinese CHB (Han Chinese in Beijing), consistent with the known migration of ancient Southwest people from Gaomiao Culture in Hunan to the Jiahu Culture in Henan and in turn to the Hongshan Culture in the Northeast as indicated by archaeological records (8 pointed star) [25]. In contrast, PCA plots using fast SNPs clustered all aDNA samples here with PEL (Peruvian in Lima, Peru) and as outliers to the ASN samples (Figure 2C and data not shown), which was reminiscent of the surprising finding with European aDNAs that ancient samples (>2000 years old) were often not the direct ancestors of present day Europeans living in the same area and do not cluster with them on PCA plots [26-29]. These results from the fast SNPs were in direct conflict with all other lines of data and hence most likely wrong, which in turn provided additional support for the theoretical justification for using slow SNPs in demographic inferences [6-12].

Figure 2. Relationship of ancient Chinese genomes with present day populations. A. Pairwise genetic distance between K2 sample and 1kGP samples. Distances were mismatches in homozygous sites. Standard error of the mean (SEM) is shown. B. PCA plots (PC1 and PC2) of K2 merged with 1kGP samples using slow SNPs. Only

ASN, EUR, and AFR samples of 1kGP are selectively shown. C. PCA plots (PC1 and PC2) of K2 merged with 1kGP samples using fast SNPs.

We performed correlation analyses using distances to European samples of 1kGP to determine whether the aDNAs here were more related to East Asians (CHS, Southern Han Chinese) or Europeans (CEU, Utah Residents with Northern and Western European Ancestry). All informative aDNAs with R values greater than expected by random simulations (>0.02) showed greater affinity to CHS than to CEU (Table 3). Some samples had two few SNPs to be informative. Therefore, although the numbers of SNPs here were limited, which would weaken the strength of correlations, most samples were still informative enough to be able to properly identify the correct group affiliations of these aDNAs.

Table 3. Correlation of ancient samples from China with Han Chinese (CHS) or Europeans (CEU) of 1kGP. Shown are Spearman correlation R values in distance to EUR samples in 1kGP. Two samples from Han Chinese of 1kGP were shown as positive controls. Comparisons were deemed non-informative when R values were $0.1 - 0$ or $-0.1 - 0$, which happens to be found for randomly scrambled distance values as shown for CHS_HG00478.

Our previous studies showed that individuals carrying the same Y haplotype were also more related in autosomes, which could only be shown with slow but not fast autosomal SNPs [7]. To study the relationship of the ancient DNAs with the super-grandfathers, we next determined the autosomal relationship of each aDNA to present day people carrying the super-grandfather Y haplotypes. We calculated the average rank in R values of each aDNA to the F5 samples relative to the F11 and F46 samples in the 211 Han Chinese in 1kGP. A ratio <1 in the F5 rank versus the F11 or F46 rank means closer relationship to the F5 samples relative to the F11 or F46 samples. Relative to F11 samples, DSQ and ZX but not DZ were found closer to F5 samples, indicating that the DSQ and ZX groups had more F5 associated autosomes and less F11 associated autosomes whereas the DZ group had the opposite (Figure 3). Relative to the F46 samples, DSQ and DZ groups were more evenly related to F5 and F46 but the ZX group was still more related to F5, indicating that the ZX group had probably the most F5 associated autosomes among the aDNAs examined here. While DSQ and ZX groups were both related to F5, DSQ was more related to F46 than ZX.

We also did the same analysis using total pairwise distance values calculated from the set of fast evolving SNPs, as would normally be done by others in the field. The results showed that all 3 aDNA groups (DSQ, DZ, and ZX) had F5/F11 ratio <1 (0.81-0.84), indicating no sensible correlation of any kind. Such dramatic difference in results obtained from the fast and the slow SNPs was consistent with previous findings [6, 7], confirming again that only slow SNPs could be used for informative analyses in phylogenetic studies or in detecting specific association between autosomes and Y haplotypes.

Figure 3. Autosomal relationships of aDNAs with Han Chinese samples of 1kGP carrying F5, F11, or F46 haplotypes. R correlation values with 211 Han Chinese and 12 aDNAs were obtained and ranked for each aDNA sample. The average rank to F5, F11, or F46 was calculated for each sample listed here and the ratios in the ranks are shown either for each sample (A) or the average of each group (B).

To determine further the possible Y haplotypes associated with the aDNA groups, we studied the autosomal relationship of aDNAs with present day samples grouped by different haplotypes. We determined the relative distance to DSQ vs DZ or to ZX vs DZ for different sets of 1kGP samples with each set associated with a particular haplotype. For each sample, we calculated a rank in the subtraction value of 'the rank to DSQ

subtracting the rank to DZ (DSQ-DZ)' with small values (<120 with 120 being the middle rank corresponding to the subtraction value 0) meaning higher rank in relation to DSQ relative to DZ. We also similarly calculated a rank in the value of 'the rank to ZX subtracting the rank to DZ (ZX-DZ)'. The samples of the M134 clade containing both F5 and F46 samples were significantly closer to ZX than samples carrying O1 and O2 haplotypes (F5/F46 and O1/O2 in Figure 4). Relative to DSQ, samples carrying C3 (3 C3-Z1300 and 1 C3-CTS9677-M217) were the closest to the DZ group although not significant. The results suggest that DZ group had more autosomal ancestry from Southern China as O1, O2, and C3-Z1300 or C3-CTS9677-M217 are known to be common in the South [5].

Figure 4. Relative autosomal distance of aDNAs to Han Chinese in 1kGP grouped by Y haplotypes. Ranks in R to aDNA groups were determined and ranks were shown for the subtraction values of 'rank to DSQ subtracting the rank to DZ (DSQ-DZ)' or 'rank to ZX subtracting the rank to DZ (ZX-DZ)'. Small subtraction ranks (<120) means closer rank to DSQ or ZX, and higher value means closer rank to DZ.

Ancient DNA relationships with Southwest Chinese carrying M117 or the F5 subtype O2a2b1a1a6

Tibetans are known to commonly carry M117 [30] and a subtype of M117 or F5, O2a2b1a1a6-CTS1642, is common in Southwest Chinese and also found in one of 3 ancient (3150-2400 YBP) Tibetan samples Chokhopani-C1 from Nepal [31, 32]. If both DSQ and ZX groups may be candidate group for the origin of F5, it would be important to ask which group is more related to Southwest people carrying M117 or O2a2b1a1a6. We made use of 4 previously published Tibetan genomes of 3150-1250 years old, including 3 M117 (including the C1 sample with the CTS1642 haplotype) and one D haplotype (markers downstream of M117 were not covered for some samples) [30]. We obtained the slow SNPs set and calculated the pairwise genetic distance with each member of 1kGP for each of the ancient Tibetans.

To determine which of the 11 aDNA samples was closely related to the M117 Tibetans, we obtained correlation R ranking of the 3 M117 Tibetans in relation to each aDNA among 227 samples including 211 Han Chinese, 11 aDNAs, and 4 Tibetan aDNAs (Figure 5A). The highest rank for the M117 Tibetans was found in correlation with the ZX group among the 4 aDNA groups. The results indicated possible gene flow between the ZX group and the M117 Tibetans.

The haplotype O2a2b1a1a6 was commonly found in Southwest Chinese such as the Dai, in addition to Tibetans [31, 32]. To further confirm the putative role of ZX in the origin of this haplotype, we studied the CDX samples (Chinese Dai in Xishuangbanna) of the 1kGP. We first confirmed that CDX samples carrying O2a2b1a1a6 were closely related to M117 Tibetans in autosomal genetic distances. We obtained R values of each aDNA with the CDX samples using their genetic distance to East Asian samples in 1kGP. We classified the M117 samples in CDX into 2 different haplogroups (O2a2b1a1a6 and other M117), and calculated the average R of the CDX in relation to the Tibetans. The results showed that CDX samples carrying O2a2b1a1a6 were significantly closer to M117 Tibetans than CDX samples carrying other M117 haplotypes (Figure 5B). CDX samples carrying the C haplotype were significantly closer to the D type Tibetan than to M117 Tibetans, consistent with C and D belonging to the ABCDE clade [7]. The results indicated that the autosomes of ancient M117 Tibetans were more

related to those associated with O2a2b1a1a6, suggesting that ancient Tibetans likely carried the O2a2b1a1a6 haplotype, just like the present day Tibetans [31].

We next divided the CDX samples into 3 Y groups (O2a2b1a1a6, Other M117, and C) and tested their relationships with the 4 groups of aDNAs here. We obtained correlation values of each aDNA to each of 99 CDX, 103 CHB, and 50 CHS samples of 1kGP. Among the 4 aDNA groups, the ZX group was found closest to the O2a2b1a1a6 samples and the only aDNA group that showed significantly closer distance to O2a2b1a1a6 samples than to C samples (Figure 5C). The results suggest a specific association of the ZX group with populations that carried the O2a2b1a1a6 haplotype, indicating either that ZX was enriched with the O2a2b1a1a6 haplotype or that ZX was the ancestral population giving rise to the group carrying the O2a2b1a1a6 haplotype. Given the present day distribution of O2a2b1a1a6 in the Southwest, it is more likely for ZX, being located in the Central Plains, to be the ancestral group to the O2a2b1a1a6 haplotype.

Figure 5. Relationship of the O2a2b1a1a6 haplotype with aDNA samples. A. Relationship between ancient Tibetans and the aDNA groups in this study. B. Relationship of Tibetans with CDX samples carrying different Y haplotypes. C. Relationship with aDNA groups of this study for CDX samples carrying different Y haplotypes.

Analyses of ancient Y chromosomes

Given the autosomal relationships of both the DSQ and ZX group with present day people carrying the F5 haplotype, it may be expected that at least some members of these two groups may carry the F5 haplotype. We performed Y chromosome sequence analysis on the only sample here that was good enough for high coverage sequencing analysis, the ~3000 year old K12 from Upper Xiajiadian Culture in West Liao River Valley, Northeast China [14]. We also analyzed the published sequence of the MG48 individual from the ~4000 year old Mogou site of the Qijia Culture in Northwest China [33]. The results showed both K12 and MG48 to be F438-F2137 with no mutations in sites that define downstream haplotypes under F2137. The results provided additional support for the presence of the F5 lineage in the DSQ group.

Discussion

Our results here showed that aDNAs in Central and Northern China from this study could be separated into 4 groups based on autosomal relationships among them. Two among these, ZX and DSQ, were related to F5 associated autosomes. The DZ group was more related to autosomes associated with the O1, O2, and C3-Z1300 that are commonly found in Southern China while the ZX group was more associated with the F5 and F46 haplotypes common in the Central Plains and the North [5]. This indicates that the DZ group might be migrants from the South. Consistently, analyses of human skulls of the Duzhong sites indicated close relationship with populations from Southern China [34, 35]. Human skulls from the Zhengzhou Xishan site or other Miaodigou sites such as Shanxian Henan however showed mixed features related to both Yangshao and Dawenkou people [36, 37]. Thus, people from different sites in Henan, such as Duzhong and Xishan, appear to have different cranial features, consistent with our DNA findings here. The DNA results confirm the suggestion based on archaeological and historical records that the early Yangshao Culture and its possible predecessor the Peiligang/Jiahu Culture may be associated with migrants under the legendary *Yan* or *Fuxi* Emperor from Southern China such as the Pengtoushan and Gaomiao Culture in Hunan [25, 38].

The DSQ group had one sample K12 carrying F5 while the ZX group was non-informative for Y chromosome. There are at least 7 branches immediately under F5 [32]. Among these, the F438 branch appears to be of high social economic status (SES) based on it having shorter branch length, more descendant branches, and higher fitness (lower risk for autism) [4]. The O2a2b1a1a1a2a2-F2137* haplotype of K12 and MG48 belongs to F438. If the original F5 haplotype had a fitness advantage that might have contributed to its super-grandfather status in the first place, haplotypes with fewer random variations from the original F5 haplotype should be expected to retain the most of the fitness advantage of F5 and as such to be more likely to confer high SES status and produce more descendants. Thus populations in ancient times near the time of the original F5 would be expected to be enriched with haplotypes closest to F5. Thus the finding of two of two informative samples that have high coverage sequence data being of the F438-F2137 haplotype is consistent with *a priori* expectation of higher prevalence of a high SES haplotype in ancient times.

It appears that the ZX group may be more directly linked to the origin of F5 as it was related to both DSQ samples in the Northeast and samples from the Southwest carrying the F5 subtype O2a2b1a1a6. The common presence of F438-F2137 in the 3000-4000 YBP time period in the North indicates unlikely the presence of F5* haplotype in the North in ancient times. The most parsimonious explanation for these observations is the diversification and radiation of F5 sub-branches from a centrally located population such as ZX where F5* might originate. This conclusion is remarkably consistent with previous molecular findings on present day Y haplotypes [39] and archaeological studies [40].

Present day Chinese are thought to be the descendants of *Yan* and *Huang*, two legendary great leaders from ~5000 YBP. Based on archaeological and historical records, scholars have suspected an association of *Yan* with the Yangshao Culture in the Central Plains (but with ancestry from the South such as the Gaomiao Culture in Hunan) and *Huang* with the Hongshan Culture in the Northeast [23, 38, 41]. The finding here of F5 lineage in both the Central Plains and the North at 5000-3000 YBP is in line with the *Yan-Huang* legends.

The Miaodigou Culture was a most popular Culture of its time and known to have impacted the Northeast Hongshan Culture (and the subsequent Xiajiadian Culture to which the DSQ group belonged), more so than any other Culture of the time such as the Dawenkou Culture [2]. Our DNA findings here suggest that there were people in the Central Plains closely related to the super-ancestor F5* lineage and possibly associated with the Miaodigou Culture. These conclusions from DNA studies are consistent with the suggestion from archaeological and historical studies that the Miaodigou Culture, and in particular the first walled town (made of rammed earth) of Xishan, may be linked to the lineage of *Huang* who is known to be the first to have built walled towns in Chinese history [21]. People of this lineage are believed to have also lived in the Northeast (Hongshan and Xiajiadian Culture) including the great Zhuanxu Emperor, a grandson of *Huang*, and to have migrated down to the Central Plains in later times during a cold climate period (~4200 YBP) [23, 40, 41]. Samples from the Niheliang site of Hongshan Culture (~5000 YBP) are 13.7% for haplotype O2a2b1-M134 (downstream sites remain to be determined) and future studies of more samples are needed to determine if F5 was present at this site [42].

Conclusion:

Overall, our study identified the presence of F5 genomes in ancient samples from the Central Plains and the Northeast at 5000-3000 YBP and implicates the origin of the F5 lineage in the Central Plains and subsequent diversification and migration to the Northeast and Southwest. The remarkable unification of ancient DNA results with archaeological and written records can only be found when we used slow SNPs but not fast SNPs. This provides further confirmation of our new molecular methodology in demographic inferences.

Materials And Methods:

Ancient DNA sequencing

Archaeological sites and samples were described in details in supplementary materials.

The post and pre-PCR labs are located in separate buildings to avoid cross contamination. Sample preparation, DNA extraction, PCR set-up and library preparation were completed in the pre-PCR laboratory. PCR and sequencing were carried out in the post-PCR laboratory. To prevent modern sources of contamination in ancient DNA, researchers in the pre-PCR lab followed a strict workflow protocols when doing the laboratory work [43].

All the samples were exclusively taken from teeth in this research. ZX107, ZX167, DZ14, and H69 were stored at the archaeology institute of Henan. Other samples were kept at the State Museum of Research Center for Chinese Frontier Archeology at Jilin University. Sample decontamination, grinding and DNA extraction were performed as described by Zhang et al [43]. The DNA libraries were built using the NEBNext® Ultra™ DNA Library Prep Kit for Illumina® in accordance to manufacturer's instructions. Probes of ~11 Mb targeted regions of the NRY chromosome were designed and used for DNA capture in this study [44]. All the libraries were sequenced on the HiSeq X-Ten platform.

Sequence download

We downloaded ancient and modern human genome sequences from 1000 genomes (<http://www.internationalgenome.org>) and the relevant websites using publically available accession numbers (SRP065070).

Selection and identification of SNPs

Selecting SNPs: The random selection of autosomal 255K SNPs as fast evolving SNPs (none from the X chr) and the selection of the slow evolving SNPs were as described previously [7].

Calling SNPs: Before aligning, Adapterremoval 2.2.1a was used to trim low-quality nucleotides (>25 bp) and remove adapter residues [45]. We then aligned the sequences to the human genome hg37 using BWA 0.7.10 [46]. The mapped reads were filtered for mapping quality 30 and sorted using SAMtools 0.1.19 [47] and Picardtools 1.107 (<http://picard.sourceforge.net>). Duplicates were removed using Picardtools 1.107. The BAM files were re-aligned using GATK-3.2-2 [48]. We calculated the ancient DNA damage and the distribution of read lengths using MapDamage2 in this study [49]. The distribution of read lengths of these samples ranged from ~45 to ~100 bp. The damage of ancient DNA was shown in Supplementary Fig. S1. We used GATK-3.2-2 and

vcftools 0.1.15 to call SNPs[48, 50, 51]. For better accuracy of calling SNPs, the Phred quality score (Q score) was set at 50, DP $\geq$ 10, and alt frequency $\geq$ 0.8 of the modern data. For the ancient Y chromosome DNAs, the filters for calling SNPs included DP $\geq$ 4 and alt frequency $\geq$ 0.8.

Genetic distance analyses and other population genetics methods

Genetic distance: Pairwise genetic distances of either hom distances (mismatch in hom sites) or total distances (mismatch in het and hom sites combined) were calculated using the custom made software “dist” as previously described [7]. This software is freely available at <https://github.com/health1987/dist> and has been described in detail in previous publications [52, 53].

PC analysis: We performed PCA using GCTA and PLINK 1.9 [54, 55].

Other methods: Other common statistical methods used were Student’s t test, chi square test, and Fisher’s exact test, 2 tailed, and Spearman correlation coefficient R analysis using Prism6 of Graphpad.

Declarations

Ethics approval and consent to participate:

The study of all human remains in this research was approved by the State Administration of Cultural Heritage of China.

Consent for publication:

Not applicable.

Availability of data

The raw reads of ancient DNAs reported here have been deposited at NCBI with accession number xxxx.

Competing interests:

The authors declare that they have no competing interests.

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Authors' contributions:

Ye Zhang and Jiawei Li did the sample preparation, DNA extraction, and DNA library preparation. Ye Zhang, Xiaoyun Lei, and Hongyao Chen analyzed the WGS data and the Y-capture data. Hui Zhou and Shi Huang conceived and supervised the study. YZ and SH wrote the manuscript.

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Tables

Table 1. Information on ancient samples for which we report the nuclear sequence data in this study.

Sample name	Ages (BP)	# SNPs slow	Coverage	Culture	Sites
FQ17	2500	1200	0.113	East Zhou	Fushan Qiaobei, Shanxi
K2	3000	8300	1.2796	Xiajiadian Upper	Dashanqian, Chifeng, Inner Mongolia
K12	3000	2000	0.2071	Xiajiadian Upper	Dashanqian, Chifeng, Inner Mongolia
S1	3920	350	0.043	Xiaoheyan	Halahaigou, Chifeng, Inner Mongolia
SG2046	3500	1500	0.1668	Xiajiadian Lower	Sanguan, Hebei
DZ14	5000	1980	0.0712	Late Yangshao	Duzhong, Mianchi Henan
ZX167	5300	3722	0.0674	Miaodigou	Zhengzhou Xishan, Henan
ZX107	5300	161	0.0179	Miaodigou	Zhengzhou Xishan, Henan
PAB3002	3000	2300	0.2361	Gaotaishan	Pinganbao, Liaoning
ZK22	4200	500	0.0485	Zhukaigou	Zhukaigou, Ordos, Inner Mongolia
H69	3500	1692	0.0304	Early Shang	Xuecun, Zhengzhou, Henan

Table 2. Ranks in correlation between each pair of ancient samples among 211 Han Chinese in 1KG and 12 ancient samples. Rank values means the rank of a sample on the column among all 223 samples in values of correlation to a sample listed on the row.

	FQ17	K2	K12	S1	SG2046	DZ14	ZK22	H69	ZX167	ZX107	PAB3002
FQ17		170	173	169	219	125	152	78	74	207	220
K2	20		7	90	216	205	61	169	69	167	192
K12	184	103		5	221	179	174	109	189	192	142
S1	217	215	149		217	187	203	106	210	193	195
SG2046	220	219	221	151		40	164	166	172	179	212
DZ14	209	221	219	220	10		21	5	214	218	214
ZK22	169	187	189	180	198	42		16	220	202	113
H69	102	217	155	7	213	34	18		221	120	190
ZX167	23	172	164	89	161	127	207	195		97	189
ZX107	219	213	198	135	203	181	196	94	159		204
PAB3002	221	214	151	61	214	135	123	133	203	178	

Table 3. Correlation of ancient samples from China with Han Chinese (CHS) or Europeans (CEU) of 1kGP. Shown are Spearman correlation R values in distance to EUR samples in 1kGP. Two samples from Han Chinese of 1kGP were shown as positive controls. Comparisons were deemed non-informative when R values were 0.1 – 0 or -0.1 – 0, which happens to be found for randomly scrambled distance values as shown for CHS_HG00478.

Sample names	R in EUR distance		T test
	CHS	CEU	
FQ17	0.010	0.039	ni
K2	0.423	0.229	<0.01
K12	0.179	0.165	
S1	0.125	0.080	<0.01
SG2046	0.165	0.152	
DZ14	0.136	0.126	
ZK22	0.182	0.073	<0.01
H69	0.158	0.138	<0.05
ZX167	0.306	0.211	<0.01
ZX107	-0.024	0.010	ni
PAB3002	0.114	0.088	<0.01
CHS_HG00478	0.623	0.376	<0.01
CHB_NA18611	0.512	0.380	<0.01
CHS_HG00478 random	-0.008	-0.020	ni

Supplemental Figure Legend

Figure S1. Ancient DNA damage assessment of all samples in this study

Figures

Figure 1

Location of ancient samples in China

Figure 2

Relationship of ancient Chinese genomes with present day populations. A. Pairwise genetic distance between K2 sample and 1kGP samples. Distances were mismatches in homozygous sites. SEMs are shown. B. PCA plots (PC1 and PC2) of K2 merged with 1kGP samples using slow SNPs. Only ASN, EUR, and AFR samples of 1kGP are selectively shown. C. PCA plots (PC1 and PC2) of K2 merged with 1kGP samples using fast SNPs.

Figure 3

Autosomal relationships of aDNAs with Han Chinese samples of 1kGP carrying F5, F11, or F46 haplotypes. R correlation values with 211 Han Chinese and 11 aDNAs were obtained and ranked for each aDNA sample. The average rank to F5, F11, or F46 was calculated for each sample listed here and the ratios in the ranks are shown either for each sample (A) or the average of each group (B).

Figure 4

Relative autosomal distance of aDNAs to Han Chinese in 1kGP grouped by Y haplotypes. Ranks in R to aDNA groups were determined and ranks were shown for the subtraction value of ‘rank to DSQ subtracting the rank to DZ (DSQ-DZ)’ or ‘rank to ZX subtracting the rank to DZ (ZX-DZ)’. Small subtraction ranks means closer rank to DSQ or ZX, and higher value means closer rank to DZ.

Figure 5

Relationship of the O2a2b1a1a6 haplotype with aDNA samples. A. Relationship between ancient Tibetans and the aDNA groups in this study. B. Relationship of Tibetans with CDX samples carrying different Y haplotypes. C. Relationship with aDNA groups of this study for CDX samples carrying different Y haplotypes.

Supplementary Files

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