

# A high-resolution picture of kinship practices in an Early Neolithic tomb

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To explore kinship practices at chambered tombs in Early Neolithic Britain, here we combined archaeological and genetic analyses of 35 individuals who lived about 5,700 years ago and were entombed at Hazleton North long cairn<sup>1</sup>. Twenty-seven individuals are part of the first extended pedigree reconstructed from ancient DNA, a five-generation family whose many interrelationships provide statistical power to document kinship practices that were invisible without direct genetic data. Patrilineal descent was key in determining who was buried in the tomb, as all 15 intergenerational transmissions were through men. The presence of women who had reproduced with lineage men and the absence of adult lineage daughters suggest virilocal burial and female exogamy. We demonstrate that one male progenitor reproduced with four women: the descendants of two of those women were buried in the same half of the tomb over all generations. This suggests that maternal sub-lineages were grouped into branches whose distinctiveness was recognized during the construction of the tomb. Four men descended from non-lineage fathers and mothers who also reproduced with lineage male individuals, suggesting that some men adopted the children of their reproductive partners by other men into their patriline. Eight individuals were not close biological relatives of the main lineage, raising the possibility that kinship also encompassed social bonds independent of biological relatedness.

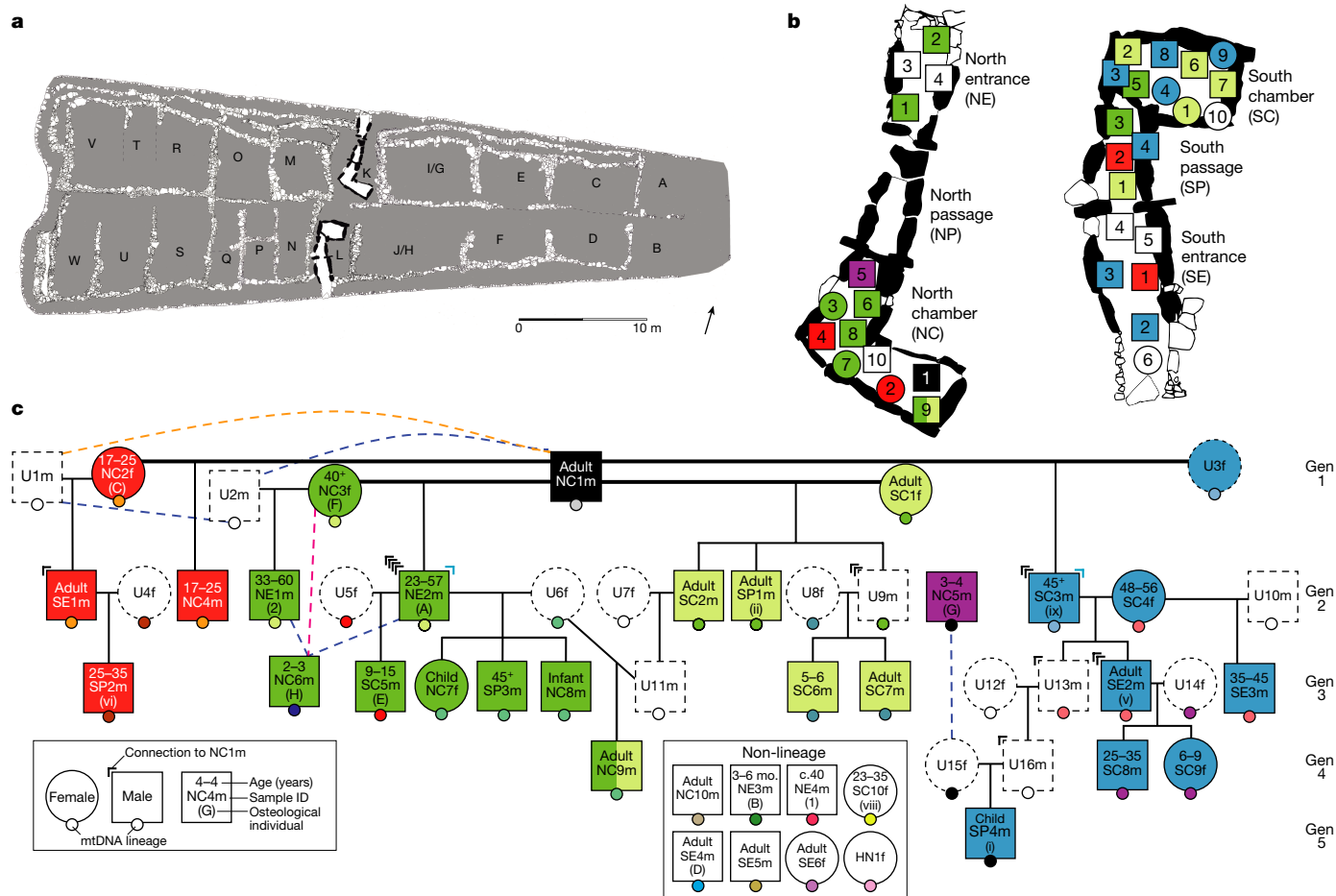
Genome-wide ancient DNA analysis has emerged as a transformative tool for understanding how people in the past related to each other and to people today. To date, these studies have mostly focused on changes in deep ancestry proportions over time, which can be accurately characterized with only a handful of individuals per population<sup>2,3</sup>. Ancient DNA has been increasingly applied to provide insight into social phenomena<sup>4–7</sup>. Yet, while more than a thousand pairs of first-degree to fourth-degree relatives have been documented in the ancient DNA literature, there have been almost no multigenerational families<sup>5,7</sup> where the exact relationships of all the individuals have been uniquely characterized. In studies of Neolithic chambered tombs in Britain and Ireland, relatedness patterns documented to date include cases of first-degree or second-degree relative pairs within or across tombs<sup>8</sup>, persistence of particular Y chromosome lineages in the same tombs<sup>5</sup>, two brothers in the same chamber in England<sup>9</sup>, and an absence of biological kin within the third degree among 11 and 15 sampled individuals at two tombs in Ireland<sup>4</sup>. Our genome-wide data on 35 individuals from the same tomb and reconstruction of a five-generation family including 27 individuals, which we co-analysed with contextual archaeological information, thus offers an unprecedented opportunity to understand social relations within the communities that built and used these tombs. Such comprehensive reconstructions not only provide insight into the

genealogical aspects of kinship in past societies but can also be used to identify kinship practices that extend beyond genealogical descent. Anthropological studies have made it clear that kinship—the relationships of family connection and belonging that have a central role in organizing societies—varies markedly across cultures. Biological relatedness may be of greater or lesser importance in determining kinship; kin need not be biological relatives (or even human), and child rearing is not always centred on the relationship between biological father and mother<sup>10–12</sup>. Funerary practices often have an important role in the social negotiation of connections and divisions between kin, and here we use this insight, along with the ability of ancient DNA to document relatedness, to provide a window into the role of biology in determining kinship among people who buried their dead in Neolithic chambered tombs.

Hazleton North (Gloucestershire, UK), an Early Neolithic Cotswold-Severn chambered long cairn, contained well-preserved human remains and was excavated in its entirety<sup>1</sup>. The tomb was constructed in the thirty-seventh century BC<sup>13</sup>, at least 100 years after cattle and cereal cultivation had been introduced to Britain along with the construction of megalithic monuments<sup>14</sup>; before that, the overwhelming majority of the biological ancestors of those buried at Hazleton North lived in continental Europe<sup>2,3</sup>. There are many other long cairns or long barrows in the region, at least nine of which share a bilateral

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**Fig. 1 | The Hazleton North pedigree in the context of the physical structure of the tomb.** **a**, Plan of the Hazleton North long cairn (grey) showing the L-shaped north and south chambered areas in the middle. The schematic in **a** is adapted from ref. 1, original figure © Historic England. **b**, Burial locations for individuals, with squares for male and circles for female individuals. Individuals are coloured according to the female sub-lineage that they belong to. The relative position of each individual within each compartment does not reflect the exact location where the corpse or remains were placed. **c**, Reconstruction of the pedigree, using the same colour scheme and indicating the locations of

individuals in the tomb, osteological information including age estimates, and different mitochondrial DNA haplogroups as small circles with different colours. Individuals with a dotted outline are unsampled (U) and their existence is inferred. Pink, blue and orange dashed lines indicate probable second-degree, third-degree and fourth-degree relationships, respectively. Marks at the top corners of individuals indicate how many genealogical connections linking individuals in the third through fifth generations to male NC1m traverse through that individual (in blue, connections through stepfathers). Gen, generation; mo., months; mtDNA, mitochondrial DNA.

arrangement of chambers with Hazleton North, although no two sites are identical and others have different chamber arrangements. Hazleton North incorporates two opposed L-shaped chambered areas mirrored around the ‘spine’ of the cairn; these roofed chambered areas were flanked by rectangular cells of masonry on either side of the axial line and the whole cairn was enclosed by a retaining wall<sup>1</sup> (Fig. 1a). The two chambered areas, north and south, each had three compartments: a chamber (innermost), a passage and an entrance (Fig. 1b, Extended Data Fig. 1). Osteological analysis has identified a minimum of 41 individuals within the tomb, including 22 adults<sup>15,16</sup>. The treatment of human remains differs somewhat between the north and the south chambers (Supplementary Information Section 1): bones from more than five individuals in the north chambered area had been gnawed by scavengers<sup>15</sup>, suggesting exposure before deposition (Extended Data Fig. 2); cremated remains from three individuals were placed in the north entrance (one infant, one child and one adult); and the remains in the south chambered area were more commingled and dispersed among neighbouring compartments than in the north chambered area. The individuals buried at Hazleton North exhibit a similar range of pathologies as those from contemporary tombs in southern Britain, such as osteoarthritis and conditions suggesting nutritional stress in childhood<sup>15</sup> (such as cribra orbitalia)

(Supplementary Information Section 1). Isotopic analysis indicates a diet rich in animal proteins<sup>17</sup>, while proteomic analysis confirms that this included dairy products<sup>18</sup>, which is also typical for the region. Bayesian modelling of 44 radiocarbon dates suggested that the monument was built over the course of a decade between 3,695 and 3,650 BC, with the stonework of the north passage collapsing and sealing off the north chamber around 3,660–3,630 BC, and the deposition of the individuals in this study probably ceasing around 3,620 BC<sup>13</sup>. A study of strontium and oxygen stable isotopes on teeth suggested that most of the 22 individuals sampled had spent some of their childhood on geology at least 40 km away<sup>19</sup>. Here we interpret new ancient DNA data alongside the archaeological evidence to reconstruct kinship practices among the community who buried their dead at Hazleton North.

To generate ancient DNA data, we obtained powder from 74 samples, largely petrous bones and teeth. We extracted DNA, generated double-stranded and single-stranded libraries, enriched for molecules overlapping approximately 1.2 million polymorphic positions in the nuclear human genome as well as mitochondrial DNA, and sequenced these libraries (Methods). We obtained data passing standard metrics for DNA authenticity for 156 libraries deriving from 66 samples (Supplementary Table 1). After detecting samples that derived from the

same individual and merging the data, we had genome-wide data from 35 distinct individuals (Extended Data Table 1) with a median coverage of 2.9-fold (range of 0.018–9.75-fold; Supplementary Table 1).

We estimated mismatch rates on the autosomes (Supplementary Tables 4, 5) for each pair of individuals, randomly sampling one DNA sequence at each position on chromosomes 1–22, and computed relatedness coefficients  $r$  (Supplementary Table 5; Methods). We also determined the type of first-degree relationships based on uniparental markers (mitochondrial DNA and Y chromosome) (Supplementary Table 1) and based on the spatial pattern of mismatches along the chromosomes (Supplementary Tables 5, 6, Extended Data Fig. 3). We manually built family trees (Supplementary Information Section 2, Fig. 1c, Extended Data Fig. 4) consistent with the pairwise genetic degrees of relatedness (Extended Data Fig. 2); maternal (mitochondrial DNA) and paternal (Y chromosome) haplogroups; genetic sex (Supplementary Table 1); genetic inbreeding (Extended Data Fig. 9) and age at death. After leveraging the distribution of recombination events (Extended Data Fig. 5), we obtained a unique pedigree that fit the data for 27 individuals (Fig. 1c). We determined that the inferred pedigree (Supplementary Information Section 2) was entirely consistent with independent information from the X chromosome (Extended Data Fig. 6a), the number of shared DNA segments (Extended Data Fig. 6b) and a different methodology for kinship estimation (Extended Data Fig. 7). We introduce a nomenclature to refer to individuals that first specifies the location within the tomb (north chamber (NC), north passage (NP), north entrance (NE), south chamber (SC), south passage (SP), south entrance (SE), unsampled individuals who may not even have been buried in the tomb but who we know must have existed on the basis of their genetic relationship to other individuals (U), and uncertain location within the tomb (HN)); then specifies an arbitrary number to distinguish each individual from the others; and finally gives a letter to indicate their chromosomal sex. In this study, we use ‘m/male/man’ to indicate an individual with an X chromosome and a Y chromosome, and ‘f/female/woman’ to indicate an individual with two X chromosomes, while recognizing that chromosomal sex is only one element in how sex and gender are contextually and culturally defined. In Extended Data Table 1, Supplementary Tables 1, 2, we provide translations between this nomenclature and genetic and osteological identifiers.

The reconstructed pedigree consists of a five-generation lineage descended from one male (NC1m) and four female individuals with whom he reproduced (SC1f, NC2f, NC3f and unsampled female U3f); also interred as part of this family are adult female reproductive partners of lineage male individuals and male line descendants of these women and non-lineage male individuals. The pedigree includes 27 individuals—three times as many individuals as the largest pedigrees reconstructed from ancient DNA<sup>5,7</sup>—and provides the first direct evidence that at least some Neolithic tombs were organized around kinship practices. Eight other individuals are not close biological relatives of these 27 individuals. The reconstructed pedigree includes a sufficiently rich network of relationships to identify kinship practices that would be invisible in smaller datasets (Extended Data Table 2, Supplementary Table 7), while the inclusion in the tomb of eight individuals without evidence of close biological relationships or reproductive partnerships with others in the pedigree suggests either that kinship did not always depend on such relations or that kinship may not have been the only criterion for inclusion in the tomb throughout its use.

Mortuary treatment varied according to chromosomal sex in several ways. First, each third-generation, fourth-generation or fifth-generation individual whose lineage we can trace through the second generation to the first is connected to NC1m entirely through male individuals. Specifically, all 15 of the genealogical connections are through fathers (13 cases) or stepfathers (2 cases) ( $P=0.000061$  from a two-side binomial test; Fig. 1c), providing the first direct evidence that patrilineal descent was a primary determinant of who was interred with whom in a Neolithic tomb. These observations are consistent with the inference that the

persistence of rare Y chromosome haplotypes over time among individuals from the same Neolithic tombs indicates patrilineal practices in these communities<sup>4,8</sup>. Second, 26 of 35 individuals with genetic data are biologically male ( $P=0.00599$  from a two-sided binomial test), consistent with osteological<sup>20</sup> and genetic evidence<sup>8</sup> that chambered tombs in England and Ireland preferentially included biological male individuals (for example, male individuals outnumber female individuals about 1.6 to 1 in Cotswold monuments)<sup>20</sup>. This suggests that the remains of some women were treated in another way (for example, exposure of remains to the elements or scattering of cremated remains away from the tomb). Third, four women among those sampled had reproduced with lineage male individuals, and their presence suggests virilocal burial, that is, burial with a male partner’s lineage rather than their father’s lineage. This, combined with the lack of adult lineage daughters among those sampled (0 adult daughters versus 14 adult sons;  $P=0.00012$  from a two-sided binomial test) and the presence of two lineage daughters who died in childhood, suggests that women generally joined the lineage of their mate. While we do not know the social or geographical distance involved in this patrilocal exogamy, the lack of long runs of homozygosity (which measures how closely two parents of an individual are related to each other) for all but one individual, indicates that inbreeding was effectively avoided (Extended Data Fig. 9). These results show that patrilineal descent had an important role in shaping social relations, a finding that may provide some insight into the nature of the community at Hazleton North (especially given the associations between patrilineal descent, virilocality, polygyny and cattle husbandry documented in ethnographically diverse cultures<sup>21</sup>). However, as we show below, the spatial organization of the dead and the inclusion of individuals who were not part of the biological patriline indicate that other considerations also had an important influence on burial patterns.

We observed six instances of multiple reproductive partners (Fig. 1c), most notably male NC1m who reproduced with four female individuals. We cannot determine whether the latter was an instance of serial monogamy or polygyny, and we cannot exclude the possibility of progeny from unions that were not socially sanctioned in any of the six instances. Where men had multiple reproductive partners, those women were not closely related to one another (Extended Data Fig. 8). However, multiple reproductive partners of female individuals were related in most cases, such as two male individuals in the patriline, NE2m and unsampled male U11m, who are inferred to be third-degree relatives and who both produced offspring with female U6f. Another case is NC3f, who reproduced with male NC1m and also with a different male individual who, although not descending from NC1m, was probably his close relative. Such women may have formed important connections between parallel lineages of related male individuals.

Our data prove that the arrangement of chambers at this Neolithic tomb was centrally determined by notions of kinship, a matter long debated for such monuments<sup>22</sup>. While determination of who could be buried at Hazleton North was primarily patrilineal, we observed a significant spatial patterning in the placement of individuals from different maternal sub-lineages, with all 12 individuals belonging to the sub-lineages of SC1f and U3f buried in the south, and 9 out of 13 belonging to the sub-lineages of NC2f and NC3f buried in the north, including the first-generation mothers in the 3 out of 4 cases where we have been able to locate them ( $P=0.0011$  from a Fisher’s exact test for a difference in the spatial placement of these four sub-lineages) (Fig. 1b). We can therefore describe the pedigree as divided into a ‘southern branch’ and a ‘northern branch’, each consisting of two maternal lines. The fact that this duality is fundamental to the architecture of the tomb suggests that the builders anticipated this division. We infer that the collapse of walling that blocked the junction of the north passage and entrance<sup>1</sup> led to the deposition of longer-lived second-generation and third-generation descendants of NC2f and NC3f outside the north chamber, disrupting this duality and perhaps contributing to the abandonment of the tomb by the northern branch ( $P=0.00408$  from a one-sided Fisher’s exact

test for the individuals in these two sub-lineages with a likely later date of death being buried outside the north chamber). The fact that these branches were based on maternal descent provides evidence that the women who founded each sub-lineage were socially significant in the memories of these communities. The interplay between patrilineal and maternal descent also has implications for interpreting the constitution of personhood and gender in this Neolithic community<sup>23</sup>.

Our genetic analyses of individuals from Hazleton North reveal kinship practices that while consistent with patrilineality cannot all be explained by biological descent. Thus, NE1m, SE1m and SE3m are not descendants of NC1m but instead are sons of women who had other children with him or his male-line genetic descendants; SP2m is the biological son of one of these individuals, SE1m. These four individuals represent cases of incorporation of male individuals into a patriline when their mothers reproduced with a man born into the lineage: this could indicate adoptive kinship, although in two cases the fathers of these male individuals were also third-degree or fourth-degree biological relatives of NC1m (Extended Data Fig. 8). Social fatherhood in this Neolithic community could be as important as biological fatherhood, a pattern observed ethnographically in societies such as the patrilineal and polygynous Nuer<sup>24</sup>. The presence of eight individuals who are not close biological relatives of any member of the lineage could be interpreted in several ways. Three were women; it is possible they were mates of lineage male individuals but did not reproduce, or that we have not sampled their offspring (who probably would not have been buried in the tomb if they were adult daughters). Some or all of these eight may have been considered kin by association or co-residence, or by adoption, raising the possibility of a meaningful role for completely non-biological kinship within the community; however, it is possible that reasons other than kinship were a factor in their inclusion in the tomb and the presence of unrelated individuals is noted at tombs from the same period in Ireland<sup>4</sup>. Overall, however, it is clear that biological relationships and kin membership were critical to the placement of many of the dead in this tomb: two pairs of sub-lineages within a single patriline were core to the layout of the tomb, and most of those buried in the chambers were lineage members. We therefore infer that the patriline and maternal sub-lineages grounded in the first generation both had anchoring roles in how kinship was negotiated at a tomb designed to both bring together and subdivide the community.

This analysis provides additional archaeological insights. Bayesian modelling of radiocarbon dates suggested Hazleton North was probably only in use for up to three generations, but the ancient DNA data document five generations in the southern chamber (Supplementary Information Section 4). Osteological identification of the minimum number of individuals in a tomb has the potential to greatly underestimate the numbers present<sup>25</sup>, yet the 66 skeletal samples that produced genome-wide data included 31 cases of genetic duplicates despite selecting bones and teeth that were not attributed to the same individuals. This suggests that our sampling is well on its way to capturing a good fraction of the individuals whose remains were recovered from the tomb and adds strength to the osteological inference that Hazleton North accommodated tens rather than hundreds of individuals (Supplementary Information Section 1). Approximately 100 long cairns are known within 50 km of Hazleton North; one only 80 m away. Further excavation, radiocarbon dating and ancient DNA analyses are needed to assess how many of these exhibit similar contemporary kinship practices, but it is possible that a high proportion of the local contemporary kin groups built and used such tombs. We have too few measurements of stable isotopes on the individuals that we analysed to be able to study correlations to cross-geology mobility<sup>19</sup>, but isotopic analyses of additional individuals with genetic data could reveal undetected patterns.

This study illustrates how ancient DNA analysis can be combined with archaeological evidence to draw inferences about kinship practices invisible to other methods. In particular, our ability to reconstruct a family tree spanning five continuous generations reveals the first direct

evidence for a central role for patrilineal descent in Neolithic mortuary practices<sup>5</sup>, the acceptance of 'stepsons' into the patriline, and a key role for maternal sub-lineages. Adoption or kinship by association may also have had a role in the inclusion of biologically unrelated individuals. Hazleton North cannot be considered a template for all Neolithic chambered tombs since the layout of such monuments varied and kinship practices could have varied between (and within) the different regions where such tombs were built<sup>22</sup>. Nonetheless, this analysis advances our understanding of kinship and chambered tomb construction in Neolithic Britain. Future research carrying out similar studies in additional tombs both in a Neolithic context in northern Europe and in other cultural contexts has the potential to test alternative theories about kinship in past societies.

## Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41586-021-04241-4>.

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## Methods

### Sampling and ancient DNA data generation

We obtained permission from the Corinium Museum to sample 8 post-cranial bones, 17 petrous bones and 49 teeth from Hazleton North. Processing into powder was carried out in dedicated clean rooms. DNA was extracted from powder using an automated protocol with silica-coated magnetic beads and 'Dabney binding buffer'<sup>26</sup>. DNA extracts equivalent to between 6 and 8 mg of powder were converted into either single-stranded or double-stranded libraries (Supplementary Table 1) following automated library preparation. For some samples, we built multiple libraries. USER treatment was applied before single-stranded library preparation<sup>27</sup> and partial UDG treatment before double-stranded library preparation<sup>28</sup>. Amplified libraries were enriched using two rounds of consecutive hybridization capture enrichment ('1240k' strategy<sup>29,30</sup>) targeting 1,233,013 SNPs and the mitochondrial genome or, 'Twist Ancient DNA' (Supplementary Table 1), a custom probe panel synthesized by Twist Biosciences. The Twist Biosciences custom panel targets the very same 1,233,013 SNPs as well as additional SNPs and tiling regions (Twist probes targeting the mitochondrial genome were spiked in) and was performed for only one round of enrichment using reagents and buffers provided by Twist Biosciences. Captured libraries were sequenced either on an Illumina NextSeq500 instrument with  $2 \times 76$  cycles ( $2 \times 7$  cycles for the indices) or on an Illumina HiSeq X10 with  $2 \times 101$  cycles ( $2 \times 7$  for the indices) (Supplementary Table 1). For this study, we restricted all our analysis to the 1,233,013 SNPs in common between 1240k and Twist Ancient DNA, as well as the mitochondrial genome.

Following the same procedure as in Olalde et al.<sup>31</sup>, we trimmed adapter sequences, merged paired-end sequences, aligned to both the human reference genome (hg19) and the mitochondrial genome (RSRS) using BWA v.0.6.1<sup>32</sup>, and removed PCR duplicate sequences. The computational pipelines are available on GitHub (<https://github.com/DReichLab/ADNA-Tools>, <https://github.com/DReichLab/adna-workflow>).

We evaluated ancient DNA authenticity using several criteria: a rate of cytosine deamination at the terminal nucleotide above 3%; a ratio of Y to combined X + Y chromosome sequences below 0.03 or above 0.35 (intermediate values are indicative of the presence of DNA from at least two individuals of different sex); for male individuals with sufficient coverage, an X chromosome contamination estimate<sup>33</sup> whose lower bound of the 95% confidence interval is below 1.1% (all but one are below 0.5%); and an upper-bound rate for the 95% confidence interval for the rate to the consensus mitochondrial sequence that exceeds 95%, as computed using contamMix-1.0.10<sup>34</sup>.

Out of a total of 74 samples, 8 did not have any library passing these criteria and were discarded, keeping 156 libraries from 66 samples for further analysis (Supplementary Table 1). We retained for analysis one sample (I30332) with 42,000 SNPs recovered that did not have enough data to test for mitochondrial or X chromosome contamination. Given that it did not display evidence of contamination according to the other two authenticity criteria, we decided to include this sample in the kinship analyses but to be cautious in the interpretation of results.

### Genetic sex, mitochondrial and Y chromosome haplogroup determination

To determine genetic sex, we looked for the presence or absence of the Y chromosome by computing the ratio of the number of Y-chromosomal 1240k positions with available data divided by the number of X-chromosomal and Y-chromosomal 1240k positions with available data. Individuals with a ratio of more than 0.35 were considered genetic males and individuals with a ratio of less than 0.03 were considered genetic females (Supplementary Table 1). To check for sex chromosome aneuploidies, we computed the mean coverage on X-chromosomal and Y-chromosomal 1240k positions, and normalized these values by the autosomal coverage on 1240k positions for

each individual. We did not find any evidence for sex chromosome aneuploidies in any individual.

To determine mitochondrial haplogroups (Supplementary Table 1), we constructed a consensus sequence with samtools and bcftools<sup>32</sup>, restricting to sequences with a mapping quality of more than 30 and a base quality of more than 30. We then called haplogroups with Haplogrep2<sup>35</sup>.

We determined Y chromosome haplogroups (Supplementary Table 1) based on the nomenclature of the International Society of Genetic Genealogy (<http://www.isogg.org>) version 14.76 (25 April 2019), restricting to sequences with a mapping quality of 30 or more and a base quality of 30 or more.

### Biological kinship estimation

We estimated pairwise allelic mismatch rates in the autosomes<sup>31,36,37</sup> for each pair of libraries ( $n = 156$ ) deriving from 66 different samples, randomly sampling one DNA sequence at each '1240k' polymorphic position and masking the two terminal nucleotides of each sequence to reduce the effects of post-mortem deamination. We then computed relatedness coefficients  $r$  for each pair (Supplementary Table 4):

$$r = 1 - (2 \times (x - (b/2))/b)$$

with  $x$  being the mismatch rate of the pair under analysis and  $b$  the mismatch rate expected for two unrelated individuals from Neolithic Britain (0.2504; Supplementary Information Section 2.2). We also computed 95% confidence intervals using block jackknife standard errors over 5-Mb blocks<sup>38</sup>.

A total of 105 pairs of libraries stemming from 44 pairs of samples had relatedness coefficients larger than 0.85, indicating that they share their entire genome and that they derived from the same individual. To increase resolution in the kinship analysis, we merged the data from samples deriving from the same individual and from libraries deriving from the same sample, keeping 35 unique individuals for further analysis. We gave a unique identifier to each of these 35 individuals (Supplementary Table 1) based on their burial location and genetic sex (for example, NC1m = male individual 1 from the north chamber).

We recomputed the mismatch rates and relatedness coefficients  $r$  on the merged dataset and annotated degrees of relationship (Supplementary Table 5, Extended Data Fig. 2). We used cut-offs lying halfway between the expected relatedness coefficients for different degrees of genetic relationships<sup>39</sup>: 1 for identical twins or samples deriving from the same individuals, 0.5 for first-degree relationships (parent-offspring and siblings), 0.25 for second-degree relationships (grandparent-grandchild, uncle/aunt-nephew/niece, half-siblings, and double cousins), 0.125 for third-degree relatives (first cousins, great-grandparent-great-grandchild, half uncle/aunt-nephew/niece, among others) and 0.0625 for fourth-degree relationships.

In addition, we determined the type of relationship (siblings or parent-offspring) connecting first-degree relatives based on uniparental markers (mtDNA and Y chromosome) and the DNA sharing along the chromosomes. To analyse DNA sharing patterns along the chromosomes, we computed allelic mismatch rate patterns across sliding windows of 20 Mb, moving by 1 Mb each step (Supplementary Table 6), and visually identified the presence (indicative of a sibling relationship) or absence (indicative of a parent-offspring relationship) of regions with zero or two chromosomes sharing for each first-degree relative pair with sufficient coverage. We illustrate this approach in Extended Data Fig. 3a and annotate the type of relationship for each first-degree pair (Supplementary Table 5).

### Family tree reconstruction

We attempted to reconstruct the family tree relating 27 close biological relatives using the pairwise degrees of genetic relatedness (Extended Data Fig. 2) through a process of triangulation that allowed us to discard

most tree topologies relating these individuals (Supplementary Information Section 2.3). To aid this process, we also incorporated information regarding: the types of first-degree relationships (Supplementary Table 5); the mtDNA and Y chromosome lineages transmitted through maternal and paternal lines (Supplementary Table 1); genetic sex (Supplementary Table 1); the presence or absence of runs of homozygosity (ROH) indicative of inbreeding (Extended Data Fig. 9b); and age at death as determined through osteological analysis (Supplementary Table 1).

After this procedure, we kept two possible tree topologies differing on whether NC1m is the father (Fig. 1c) or the son of SC3m (Extended Data Fig. 4). To disambiguate between these two scenarios, we studied the colocalization of break points of shared DNA segments between individual SC3m and each of his second-degree relatives NC4m, NE2m, SC2m and SP1m (Supplementary Information Section 2.4, Extended Data Fig. 5). This allowed us to obtain a unique family pedigree relating most of the Hazleton North individuals (Fig. 1c).

## Testing the validity of the proposed family tree

We validated the family tree in Fig. 1c using three independent lines of evidence (Supplementary Information Section 2.5): (1) we computed pairwise mismatch rates and relatedness coefficients on the X chromosome (Supplementary Table 5) following the same formula:  $r = 1 - (2 \times (x - (b/2))/b)$ . For male–male comparisons, we adjusted the formula as follows to account for the fact that male individuals have only one X chromosome:  $r = 1 - (x/b)$ . We plotted relatedness coefficients on the X chromosome for first-degree and second-degree pairs (Extended Data Fig. 6a), grouping these pairs based on whether they are expected to share X chromosome DNA according to the tree structure proposed in Fig. 1c. We found that X chromosome sharing patterns perfectly fit the proposed tree structure. (2) For each first-degree or second-degree pair with more than 100,000 overlapping SNPs, we computed allelic mismatch rate values across sliding windows of 20 Mb, moving by 1 Mb each step (Supplementary Table 6). We plotted these values along the chromosomes and visually identified contiguous regions where the allelic mismatch rate is consistent with one chromosome that is identical between the two individuals due to recent descent from a shared ancestor (identical by descent (IBD)) (Extended Data Fig. 3b). In Supplementary Table 5, we annotated the number of such IBD segments identified for each first-degree and second-degree relative pair. We next plotted the number of IBD segments for first-degree and second-degree relationships (Extended Data Fig. 6b), again grouping the pairs according to type of relationship in the proposed tree (Fig. 1c). We recovered the expected pattern<sup>40,41</sup> of a higher number of IBD segments in avuncular and maternal half-sibling pairs than in grandparent–grandchild and paternal half-sibling pairs, adding further support to the proposed tree structure. (3) We replicated our results using the software NgsRelate v.2<sup>42</sup> that uses genotype likelihoods and population allele frequencies to estimate Cotterman coefficients  $k_0$ ,  $k_1$  and  $k_2$ , which correspond to the probability of sharing 0, 1 and 2 alleles in identity by descent. From these coefficients, the software computes the Theta coefficient ( $\theta$ ), which is equivalent to the relatedness coefficient  $r$ . To run NgsRelate, we first created genotype likelihoods directly from the bam alignment files using ANGSD v0.923<sup>33</sup>. We included Hazleton North individuals as well as the set of 53 Neolithic individuals from other sites in Britain. We then ran NgsRelate providing as input the genotype likelihood file and allele frequencies estimated only on the Neolithic set from Britain, to avoid possible bias in allele frequencies stemming from the presence of a high number of closely related individuals at Hazleton North. We observed a strong correlation between both methodologies (Extended Data Fig 7).

## Principal component analysis

To obtain an overview of the ancestry of the Hazleton North individuals, we ran a principal component analysis using the ‘smartpca’ program in EIGENSOFT<sup>43</sup>. We merged the genomic data from the Hazleton North

individuals with other ancient Neolithic and Bronze Age individuals from Britain and Ireland reported in previous publications<sup>2–4,8,9,44</sup>, as well as with 1,109 present-day West Eurasian individuals genotyped on the Affymetrix Human Origins Array<sup>43,45,46</sup>, restricting to 591,642 SNPs that overlapped between the 1240k capture and the Human Origins Array. We projected ancient individuals onto the components computed on present-day individuals with Isqproject:YES and shrinkmode:YES, and plotted the first two principal components (PCs) (Extended Data Fig. 9a). The Hazleton individuals form a homogeneous cluster within the genomic diversity of contemporaneous Neolithic individuals from England, Scotland and Ireland, indicating that they derived from a very similar pool of ancestors as other Neolithic groups across Britain. We did not detect individuals shifted towards smaller values on PC1 that would suggest recent admixture with Mesolithic hunter–gatherers.

## Genetic inbreeding analysis

To study the presence of inbreeding in the Hazleton North group, we used the software hapROH<sup>47</sup> that detects ROH in ancient individuals. ROH are regions of an individual’s genome where the maternal and paternal chromosomes are identical because they derive from a recent common ancestor. The number and length of these segments in a given individual inform about the degree of biological relationship between the parents. We ran hapROH using standard parameters on the Hazleton individuals with data for more than 400,000 SNPs covered (Supplementary Table 1, Extended Data Fig. 9b). The software also computes the ROH expected for offspring of close relatives in outbred populations, and for individuals from populations with a small effective population size<sup>47</sup>. The lack of long ROH in all but one individual (Extended Data Fig. 9b) indicates that the Hazleton community effectively avoided reproductive unions between close relatives. Only one individual (SE6f) had a long ROH of 31 cM, which could be compatible with offspring of second or third cousins. This individual does not belong to the family pedigree.

## Reporting summary

Further information on research design is available in the Nature Research Reporting Summary linked to this paper.

## Data availability

The aligned sequences are available through the European Nucleotide Archive, accession PRJEB46958; the genotype dataset is available as a Supplementary Data file.

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**Author contributions** N.R., O.C., S.C., R.P. and D.R. performed or supervised laboratory work. C.F., V.C., I.A., L.B. and S.C. assembled or contextualized archaeological material. C.F. and I.O. analysed data. C.F., I.O. and D.R. wrote the manuscript. C.F., I.O. and S.C. wrote the supplement.

**Competing interests** The authors declare no competing interests.

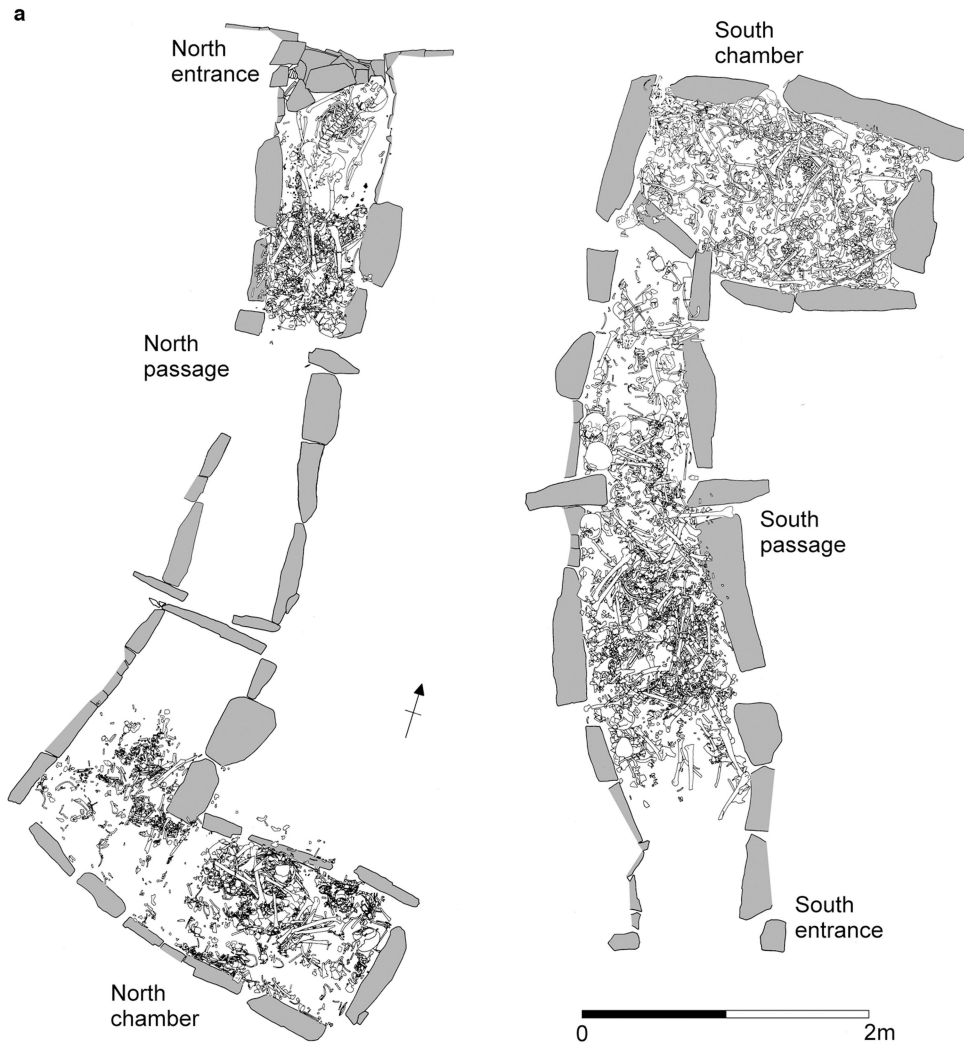
**Additional information**

**Supplementary information** The online version contains supplementary material available at <https://doi.org/10.1038/s41586-021-04241-4>.

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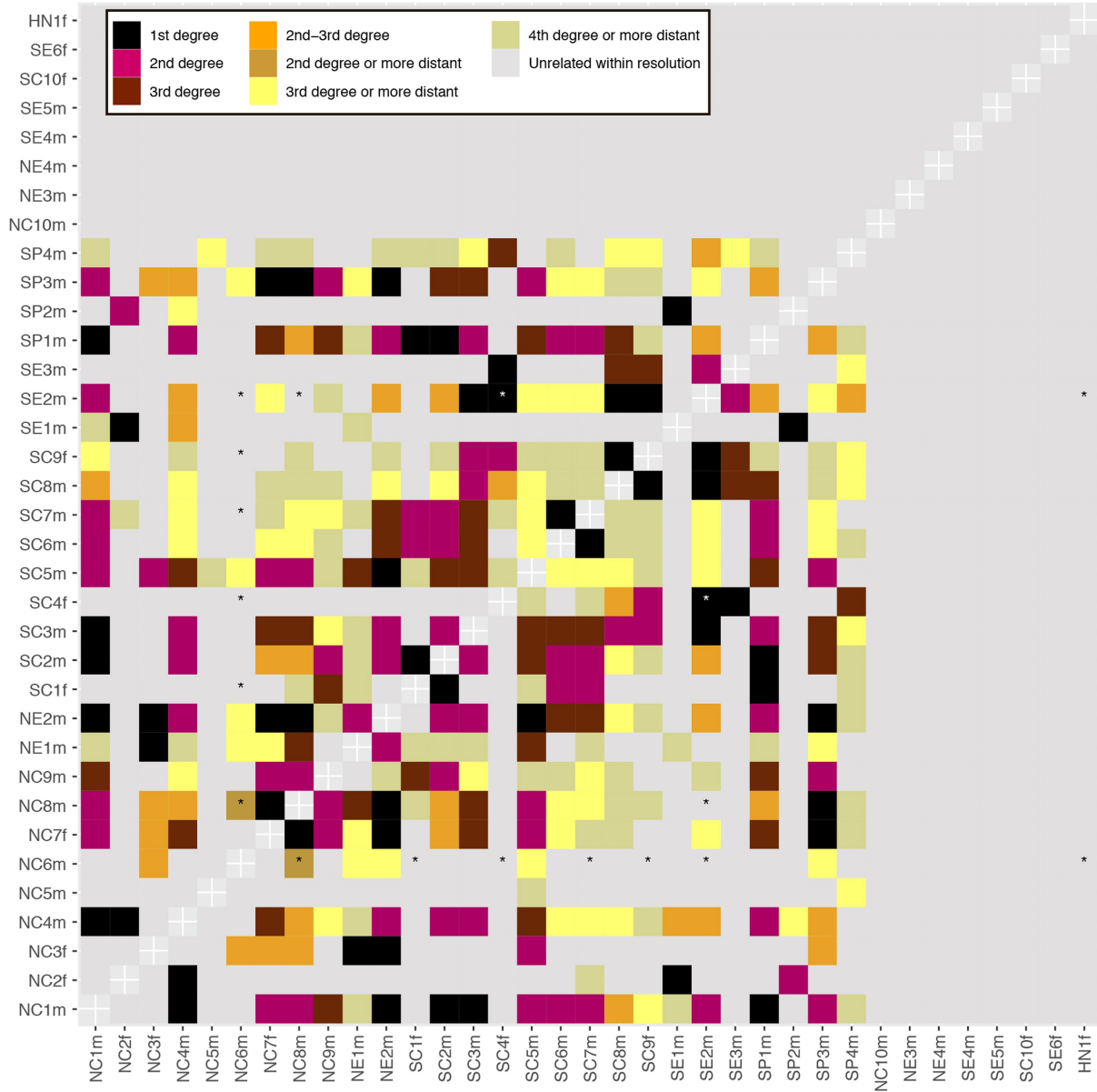
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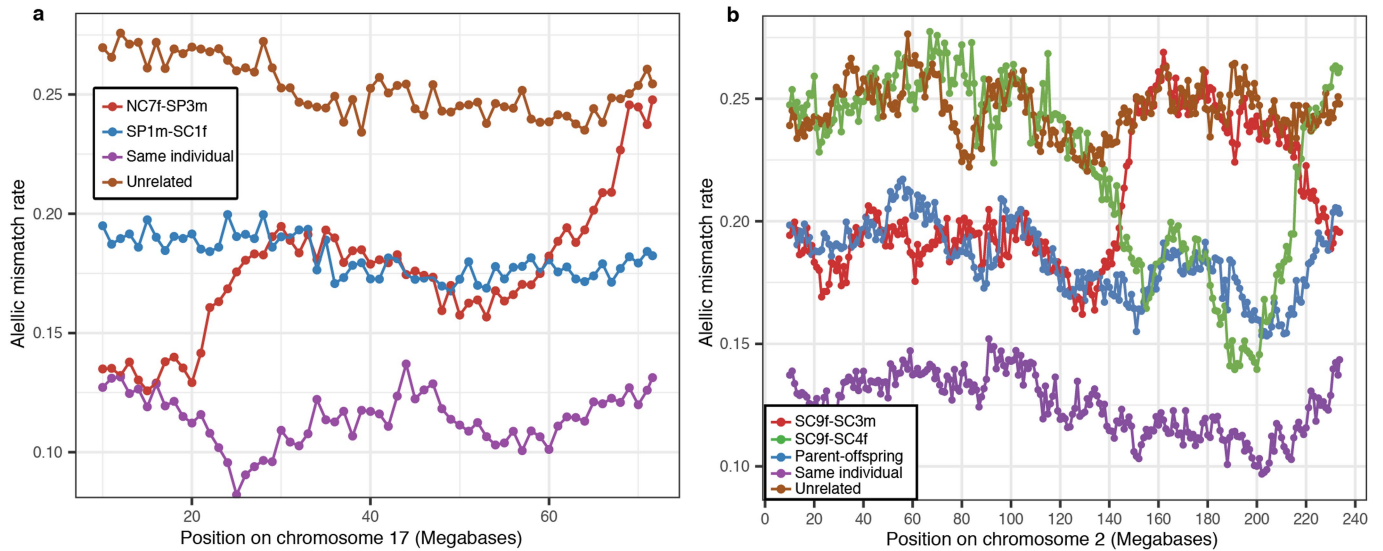
**Extended Data Fig. 1 | The Hazleton North chambered tomb. a**, Distribution of human remains in both chambers. The schematics in **a** are adapted from ref. <sup>1</sup>, original figures © Historic England. **b**, Right humerus from Individual C

showing helical fracture (red arrow), tooth marks (yellow arrow) and gnawed proximal and distal ends (white arrows).



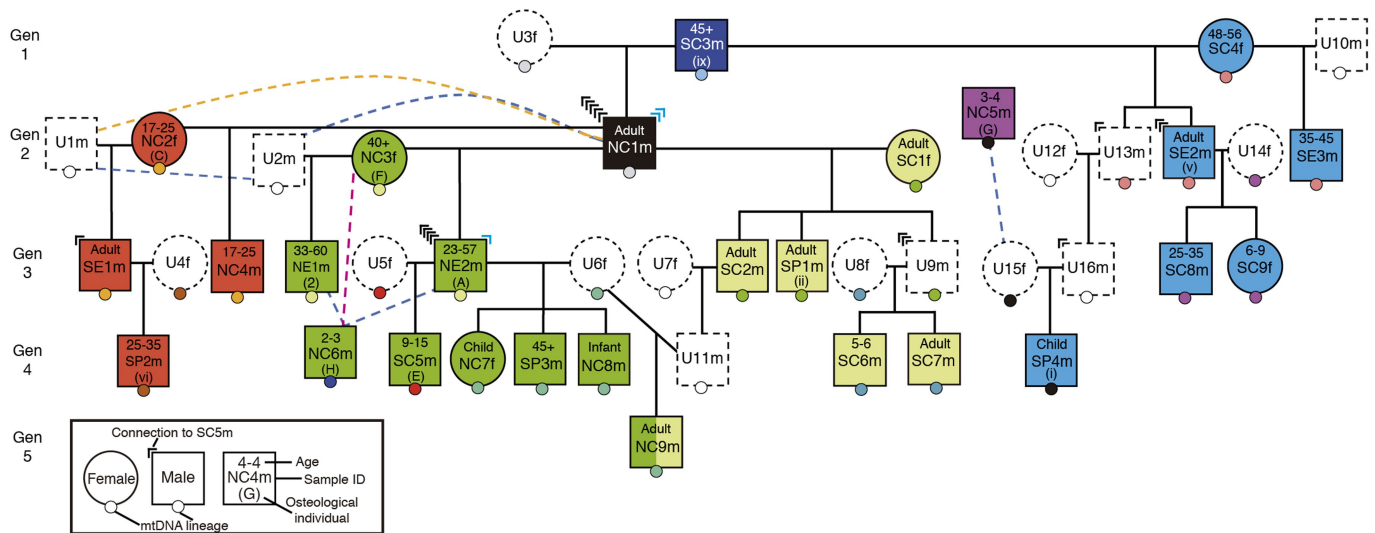


**Extended Data Fig. 2 | Degrees of biological relatedness among individuals at Hazleton North.** (Supplementary Information Section 2.2). Pairs with fewer than 15,000 overlapping SNPs are indicated with an asterisk.



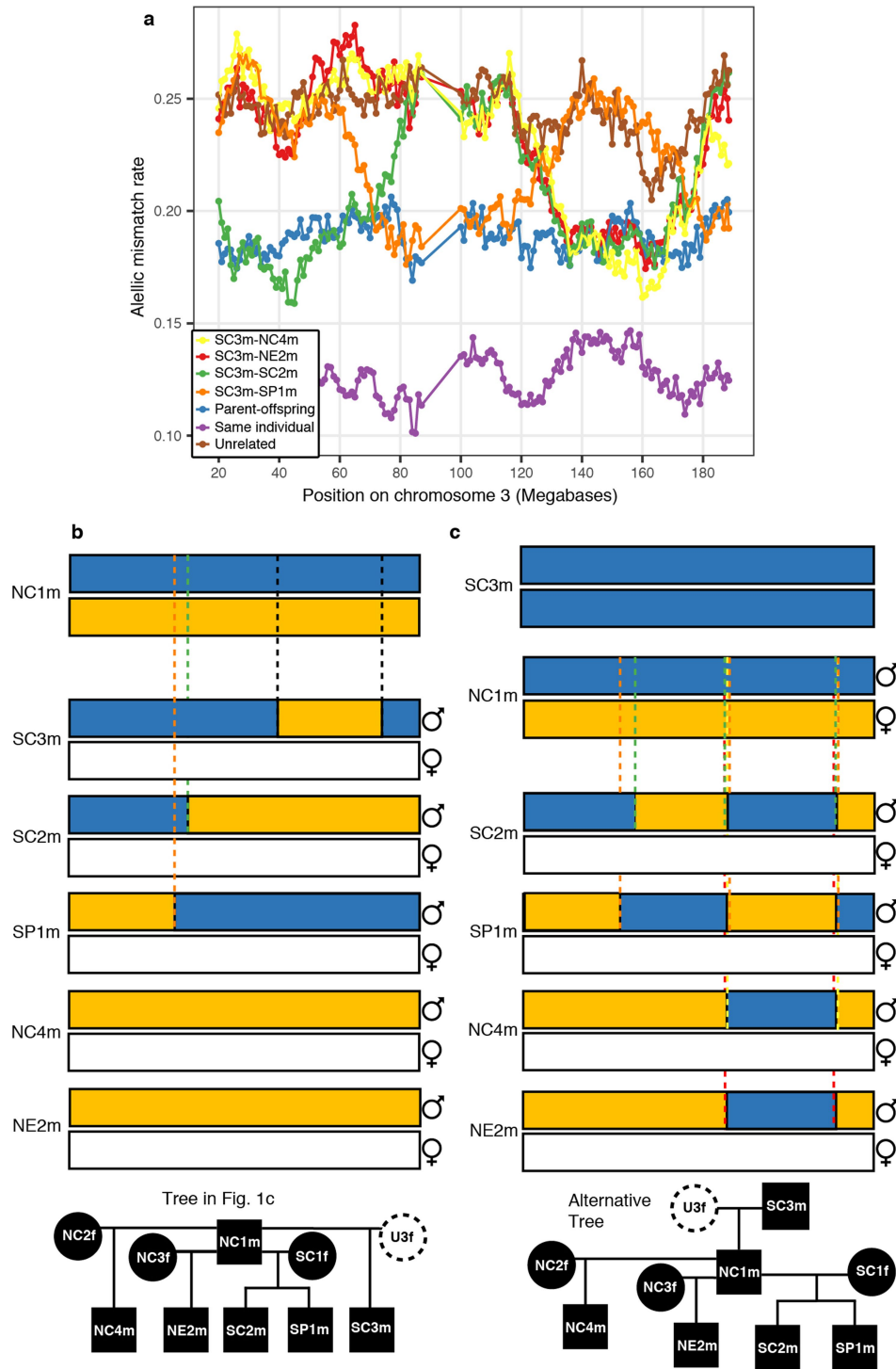
**Extended Data Fig. 3 | Using allelic mismatch rate patterns along the chromosomes to differentiate types of relationships for individuals sharing the same amount of DNA.** **a**, Differentiating between parent-offspring and sibling relationships. Allelic mismatch rate values across sliding windows of 20 Mb, moving by 1 Mb each step. As an example, we show values at chromosome 17 and include for reference a comparison between two unrelated Neolithic individuals from Britain (in brown), and a comparison between one individual and himself (in purple) to show how mismatch rates behave when two chromosomes are shared. The mismatch rate pattern for SP1m-SC1f is compatible with one chromosome shared along the entire chromosome 14 (in fact, along all autosomal chromosomes (Supplementary Table 6)), indicating a parent-offspring relationship. In contrast, the NC7f-SP3m comparison shows regions on chromosome 17 where no chromosome is shared (-65–70 Mb), other regions where two chromosomes are shared (-0–25 Mb) and

other regions where one chromosome is shared (-25–60 Mb), compatible with a sibling relationship. **b**, Comparing DNA sharing patterns between SC9f and her paternal grandparents. We show mismatch rate values at chromosome 2 and include for reference a parent-offspring comparison (SE1m-SP2m; in blue) to show how mismatch rates behave when one chromosome is shared. Two recombination events (one at -145 Mb and other at -220 Mb) in SC9f's father's gamete result in SC9f sharing one chromosome with SC3m from the start of the chromosome to -145 Mb, one chromosome with SC4f from 145 to 220 Mb and one chromosome with SC3m from 220 Mb to the end of the chromosome. This pattern of sharing one chromosome with either SC3m or SC4f (but never both) at every location of the genome is characteristic of comparisons between a grandchild and his/her two grandparents and is also observed in the other autosomal chromosomes.



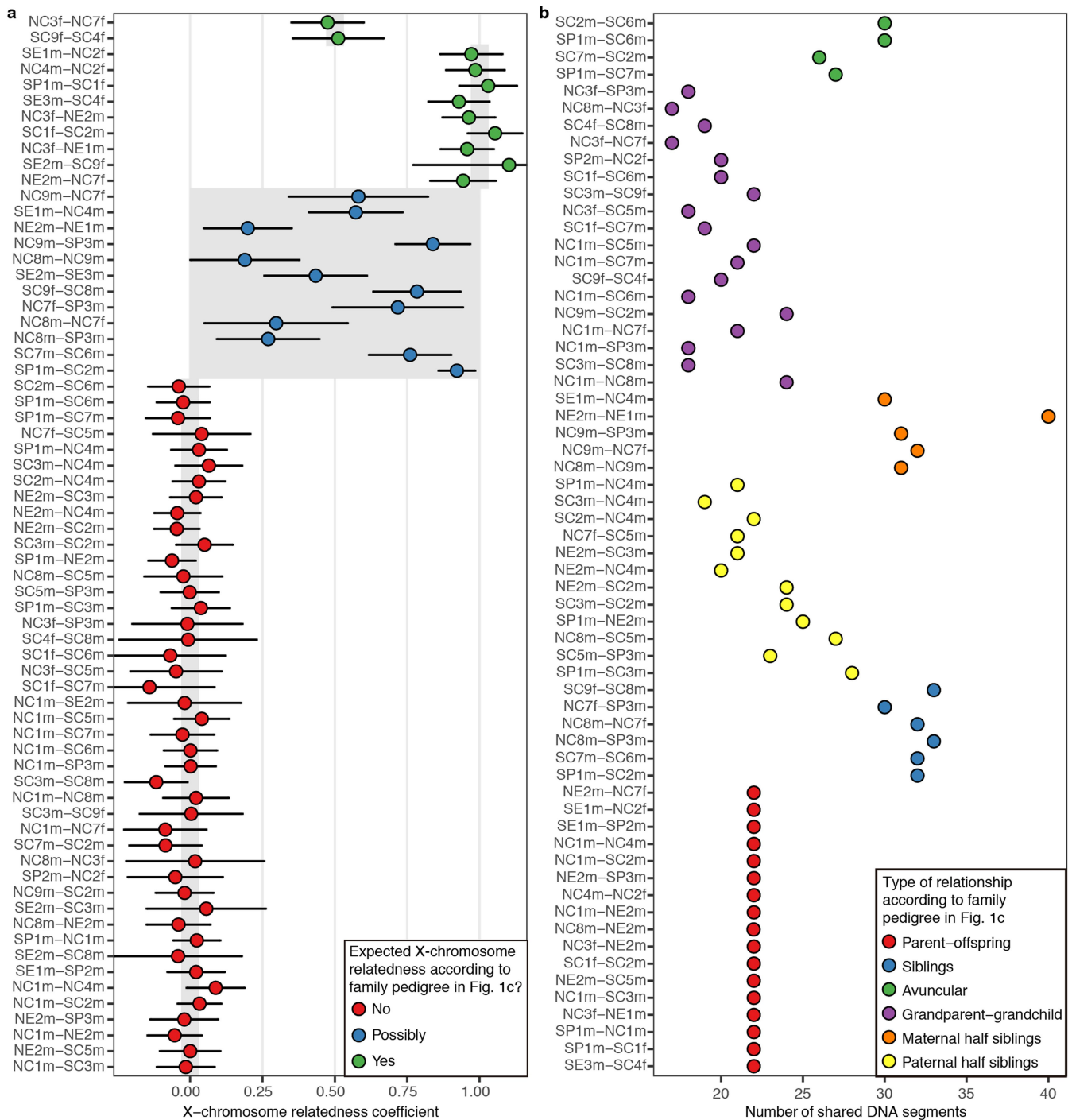
**Extended Data Fig. 4 | Alternative family tree fitting all the genetic evidence except the IBD breakpoints co-localization analysis.** (Supplementary Section 2.4, Extended Data Figure 5). Individuals are coloured

according to the female sub-lineage they belong to (NC1m and NC5m do not belong to any of the four major sub-lineages and are thus given a different color).



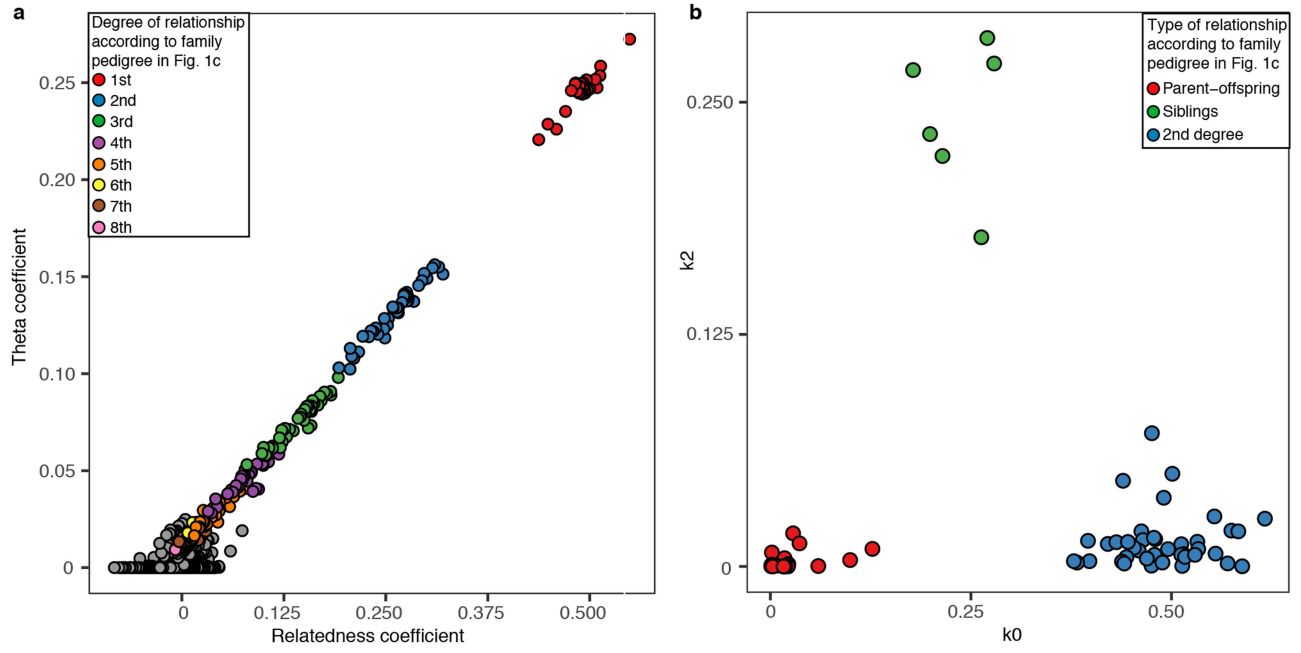
**Extended Data Fig. 5 | Using co-localization of IBD breakpoints to disambiguate between family tree in Fig. 1c and family tree in Extended Data Fig. 4. a.** We show mismatch rate values across sliding windows of 20 Mb on chromosome 3, moving by 1 Mb each step, for comparisons between SC3m and his four second-degree relatives. **b, c.** Recombination events on

chromosome 3 needed to explain the observed mismatch rate patterns under **b**, the scenario of tree in Fig. 1c where 4 recombination events are required, or **c**, the scenario of the tree in Extended Data Fig. 4 where 10 recombination events are required including the extremely implausible occurrence of two recombination events at the same genomic locations in four different gametes.



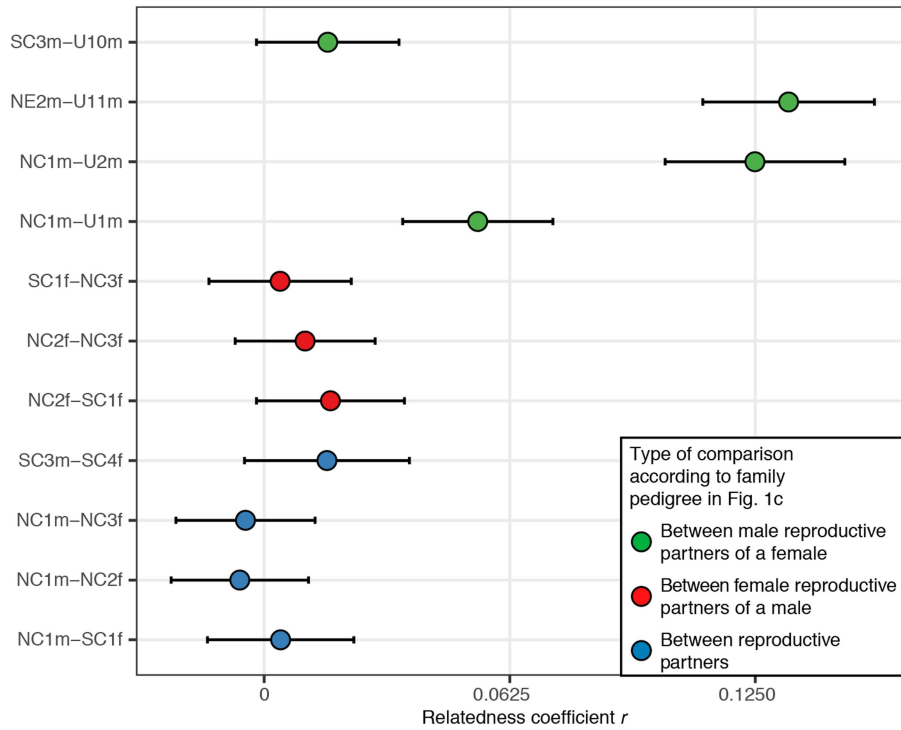
**Extended Data Fig. 6 | Testing the validity of the family pedigree in Fig. 1c using X-chromosome relatedness and number of shared IBD segments.**  
**a**, Relatedness coefficients in the X-chromosome for first- and second-degree relationships with more than 300 overlapping SNPs. For each comparison, expected values according to the type of relationship in the family tree in Fig. 1c

are shown in grey boxes. Bars represent 95% confidence intervals. **b**, Number of shared IBD segments on chromosomes 1-22 for first- and second-degree relationships. Pairs are grouped according to their type of relationship in the family tree in Fig. 1c.



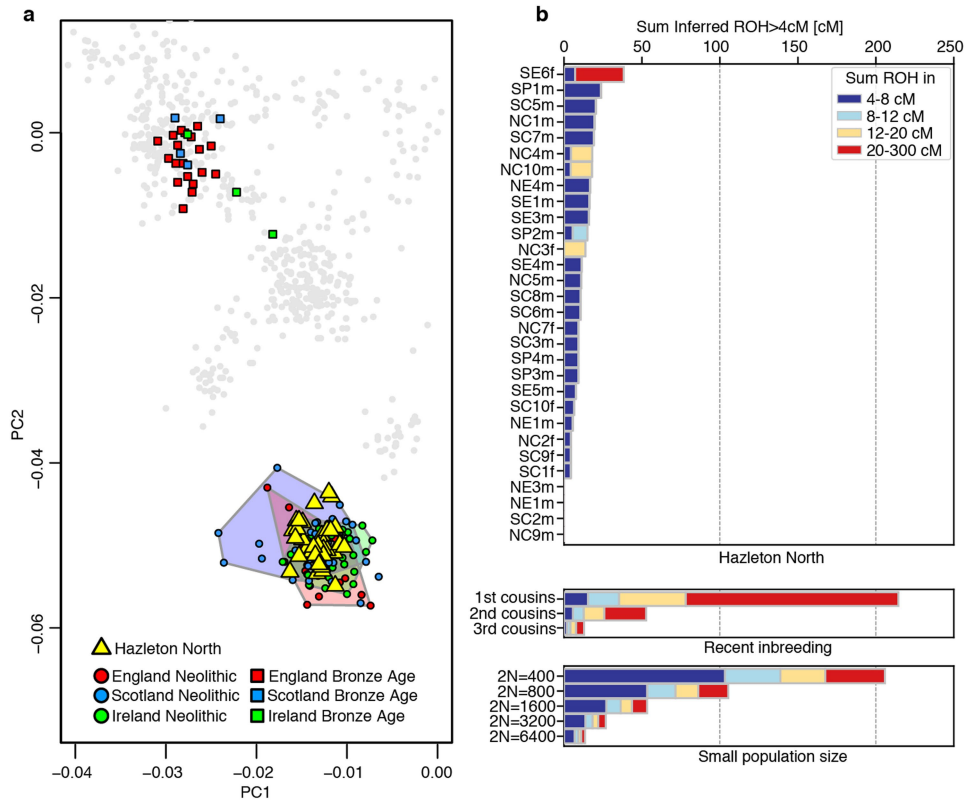
**Extended Data Fig. 7 | Testing the consistency of the kinship results using *NgsRelate*<sup>42</sup>.** **a**, Correlation between the relatedness coefficient  $r$  and the Theta coefficient computed with *NgsRelate*, restricting to comparisons with more

than 15,000 overlapping SNPs. **b**, Cotterman coefficients  $k_0$  and  $k_2$  for first- and second-degree relationships, as computed with *NgsRelate*.



**Extended Data Fig. 8 | Comparing autosomal relatedness between reproductive partners, different male reproductive partners of a female and different female reproductive partners of a male.** To estimate relatedness coefficients between unsampled and sampled male reproductive

partners of a female, we doubled the relatedness coefficient obtained between the son of the unsampled male and the sampled male, to account for the fact that a son is one degree of relationship further away from their father's relatives as compared to his father. Bars represent 95% confidence intervals.



**Extended Data Fig. 9 | Principal Component Analysis and inbreeding analysis. a.** Principal component analysis of Hazleton North individuals and other ancient individuals from Britain and Ireland. Ancient individuals were projected onto the principal components computed on a set of present-day West Eurasians genotyped on the Human Origins Array (not shown in the figure). Individuals with fewer than 15,000 SNPs on the Human Origins dataset

were excluded for this analysis. **b.** Runs of homozygosity (ROH) in different length categories for the Hazleton North individuals with more than 400,000 SNPs covered. ROH were computed using *hapROH*<sup>47</sup>. Below, we plot the expected ROH length distribution for the offspring of closely related parents in outbred populations and for individuals from populations with small effective population size<sup>47</sup>.



**Extended Data Table 1 | Key details for sampled individuals**

Unique identifier	Age-at-death	MtDNA haplogroup	Taphonomy & pathology	Main lineage key relationships
NC1m	Adult	N1b1b	-	Father of SC2m & SP1m by SC1f; father of NC4m with NC2f(C); father of NE2m(A) with NC3f(F); father of SC3m(ix) by U3f.
NC2f(C)	17–25	U8b1b	Gn; CO, PH	Mother of NC4m by NC1m; mother of SE1m by U1m.
NC3f(F)	40+	K1a3a1	OA, ankle trauma	Mother of NE2m(A) by NC1m & NE1m(2) by U2m.
NC4m	17–25	U8b1b	-	Son of NC2f(C) & NC1m; half-brother of SE1m.
NC5m(G)	3–4	T2e1	Gn	Fourth degree relative of SP4m.
NC6m(H)	2–3	J2b1a	Gn; DA	Second-degree relative of NC3f(F); third degree relative of NE2m(A) & NE1m(2)
NC7f	Child	K1b1a	-	Daughter of NE2m(A) & U6f; sister of SP3m & NC8m; half-sister of SC5m(E).
NC8m	Infant, 18–24 months	K1b1a	-	Son of NE2m(A) & U6f; brother of NC7f & SP3m; half-brother of SC5m(E).
NC9m	Adult	K1b1a	-	Son of U6f & U11m.
NC10m	Adult	U3a1	-	None
NE1m(2)	33–60	K1a3a1	AMTL, OA, OD, TB?, DA	Son of NC3f(F) and U2m; half-brother of NE2m(A); 'stepson' of NC1m.
NE2m(A)	23–57	K1a3a1	Gn; DISH, SA, OA	Son of NC1m & NC3f(F); half-brother of NE1m(2); father of SC5m by U5f & of NC7f, SP3m & NC8m by U6f.
NE3m(B)	3-6 mths	V	-	None
NE4m(1)	c. 40	K1a4	Fr. L fibula, OA, DA, AMTL	None
SC1f	Adult	K2b1	-	Mother of SC2m and SP1m by NC1m.
SC2m	Adult	K2b1	-	Son of NC1m & SC1f, brother of SP1m.
SC3m(ix)	45+	W5	Skull fr., OA, PD, DA, AMTL	Son of NC1m & U3f; father of U13m & SE2m(v) by SC4f.
SC4f	48–56	K1d	-	Mother of SE2m(v) & U13m by SC3m(ix); mother of SE3m by U10m.
SC5m(E)	9–15	H1	-	Son of NE2m(A) & U5f; half-brother of SP3m, NC8m & NC7f.
SC6m	5–6	U5b1+16189+@16192	Scurvy	Son of U8f & U9m, brother of SC7m.
SC7m	Adult	U5b1+16189+@16192	-	Son of U8f & U9m; brother of SC6m.
SC8m	25-35	J1c1b1	PD	Son of U14f and SE2m; sister of SC9f.
SC9f	6–9	J1c1b1	Scurvy	Daughter of U14f and SE2m; sister of SC8m.
SC10f(viii)	23-35	K1b1a1d	CO, PH, AMTL	None
SE1m	Older adult	U8b1b	-	NC1m reproduced with his mother NC2f(C); father of SP2m(vi) by U4f; 'stepson' of NC1m.
SE2m(v)	Adult	Too little data	-	Son of SC4f & SC3m(ix); father of SC8m & SC9f.
SE3m	35-45	K1d	-	SC3m(ix) reproduced with his mother SC4f; 'stepson' of SC3m.
SE4m(D)	Adult	J1c1	Fr. R ulna; polio?, twisted spine	None
SE5m	Adult	U5a2d	-	None
SE6f	Adult	U5b1+16189	-	None
SP1m(ii)	33-45	K2b1	-	Son of NC1m & SC1f, brother of SC2m.
SP2m(vi)	25-35	H5	-	Son of SE1m (who was 'stepson' of NC1m).
SP3m	45+	K1b1a	PD, AMTL, DA	Son of NE2m(A) & U6f; brother of NC7f & NC8m; half-brother of SC5m(E).
SP4m(i)	Child	T2e1	-	Son of U16m & U15f.
HN1f	-	K1b1a1	-	None

The individual code consists of the location of the remains, then a number for the individual within that location, and finally their sex. For those with an osteological code, this value is provided in parentheses. Full details, including bone element numbers, radiocarbon dates and stable isotope data are provided in Supplementary Table 1. Fr.=fracture; Gn.=gnawed by canids; AMTL=ante-mortem tooth loss; DA=dental abscess; DISH=Diffuse Idiopathic Skeletal Hyperostosis; CO=cribra orbitalia; OA=osteoarthritis; OD=osteochondritis dissecans; PD=periodontal disease; PH=porotic hyperostosis; SA=septic arthritis.

# Article

## Extended Data Table 2 | Statistically significant patterns in genetic data

Finding	Falsified null hypothesis	Data	P-value
Bias against female burial	Number of females and males buried at Hazleton not significantly different	9 vs 26	0.00599
Strict patrilineality	Number of matrilineal and patrilineal transmissions not significantly different	0 vs 15	0.000061
Strict bias against burial of adult daughters	Number of female and male adult offspring of unions in pedigree not significantly different	0 vs 14	0.00012
Differences in south/north placement in tomb of the four maternal sub-lineages	No association of tomb half and female sub-lineage	NC2f (2 south; 2 north) NC3f (2 south; 7 north) SC1f (5 south; 0 north) U3f (7 south; 0 north)	0.00110
Members of NC2f's and NC3f's sub-lineages stopped being buried in the north chamber later in the tomb's use-life, possibly due to the wall collapse blocking access to the north chamber	No decrease in north chamber burials among members of NC2f's and NC3f's sub-lineages with a likely later death	Earlier death (6 north chamber; 0 other) Later death (1 north chamber; 6 other)	0.00408

Two-sided binomial tests for rows 1–3, Fisher's exact tests for row 4 (two-sided) and row 5 (one-sided). See Supplementary Section 3 and Supplementary Table 7 for details.

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- The exact sample size ( $n$ ) for each experimental group/condition, given as a discrete number and unit of measurement
  - A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
  - The statistical test(s) used AND whether they are one- or two-sided  
*Only common tests should be described solely by name; describe more complex techniques in the Methods section.*
  - A description of all covariates tested
  - A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
  - A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
  - For null hypothesis testing, the test statistic (e.g.  $F$ ,  $t$ ,  $r$ ) with confidence intervals, effect sizes, degrees of freedom and  $P$  value noted  
*Give  $P$  values as exact values whenever suitable.*
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### Software and code

Policy information about [availability of computer code](#)

Data collection BWA v0.6.1, HaploGrep2 v2.1.19, contamMix v1.0-10, ANGSD v0.923, NgsRelate, Samtools, bcftools, other bioinformatics tools and data workflows (<https://github.com/DReichLab/ADNA-Tools> and <https://github.com/DReichLab/adna-workflow>)

Data analysis hapROH v0.3

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio [guidelines for submitting code & software](#) for further information.

### Data

Policy information about [availability of data](#)

All manuscripts must include a [data availability statement](#). This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our [policy](#)

The aligned sequences are available through the European Nucleotide Archive, accession PRJEB46958 [to go live upon publication]; the genotype dataset is available as supplementary material.

## Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

Life sciences  Behavioural & social sciences  Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see [nature.com/documents/nr-reporting-summary-flat.pdf](https://nature.com/documents/nr-reporting-summary-flat.pdf)

## Ecological, evolutionary & environmental sciences study design

All studies must disclose on these points even when the disclosure is negative.

Study description	This study investigates whether or not a large number of individuals buried in the same Neolithic tomb were close biological relatives, and, if they were, how they were related. By comparing these results with archaeological contextual information, such as where each individual was placed in the tomb, we are able to draw detailed inferences about kinship practices. The study collected 74 samples from human skeletal remains for DNA analysis, assessed the biological relationships between those individuals, and situated that alongside the available archaeological data for the site in reaching its conclusions
Research sample	The study collected 74 aDNA samples, largely taken from petrous bones and teeth but also including some post-cranial elements and not known to be from the same individuals, in an attempt to perform as complete as possible as a survey of ancient DNA of individuals buried in Hazleton North chambered cairn. Of these samples, 66 produced viable results. Our genetic analysis indicates that the samples derive from 35 distinct individuals whose remains were found within the tomb; the high rate at which we are obtaining DNA from multiple samples from the same individuals suggests that our sampling is well on its way to achieving its goal of sampling a large fraction of the individuals buried in the tomb. The 35 distinct individuals we identify from DNA can be compared with the osteological estimate of the presence of a minimum number of 41 individuals from the tomb. Archaeological and osteological data is drawn from the original excavation report (Saville 1990) and a PhD study involving a re-examination of the human remains (Cuthbert 2019), both of which are cited in the article. A full catalogue of each skeletal element is available from Corinium Museum on request.
Sampling strategy	The study took a sample from every petrous portion of the temporal bone that was available during the two times the museum was visited (17 in total), and samples from teeth (49 in total), and post-cranial bones (8 in total) designed to cover as many of the individuals buried in the tomb as possible. Petrous bones and teeth were prioritized as they contain the densest concentrations of surviving DNA and so are most likely to yield a result. We never knowingly took multiple samples from the same individual, as our goal was to minimize destruction of skeletal elements while obtaining data from as high a fraction as possible of the individuals in the tomb. Our goal was to sample enough so that we began to saturate the total number of individuals buried in the tomb, as we in fact did by observing that the 66 samples with viable results came from only 35 distinct individuals.
Data collection	<p>The first round of skeletal sampling was performed by a team from the University of Vienna (Olivia Cheronet) in the presence of staff at the Corinium museum. The second round of skeletal sampling was performed entirely by the staff at the Corinium museum.</p> <p>DNA was extracted from using an automated protocol with silica coated magnetic beads and 'Dabney binding buffer'. DNA extracts equivalent to between 6 and 8 mg of powder were converted into either single-stranded or double stranded libraries following automated library preparation. For some samples we built multiple libraries. USER treatment was applied before single-stranded library preparation and partial UDG treatment before double-stranded library preparation. Amplified libraries were enriched using two rounds of consecutive hybridization capture enrichment ('1240K' strategy<sup>32,33</sup>) targeting 1,233,013 SNPs and the mitochondrial genome or, 'Twist Ancient DNA', a custom probe panel synthesized by Twist Biosciences. This custom panel targets the very same 1,233,013 SNPs as well as additional SNPs and tiling regions (Twist probes targeting the mitochondrial genome were spiked in) and was performed for only one round of enrichment using reagents and buffers provided by Twist Biosciences. Captured libraries were sequenced either on an Illumina NextSeq500 instrument with 2x76 cycles (2x7 cycles for the indices) or on an Illumina HiSeq X10 with 2x101 cycles (2x7 for the indices). For this study, we restricted all our analysis to the 1,233,013 SNPs in common between '1240K' and 'Twist Ancient DNA' and the mitochondrial genome.</p> <p>Following the same procedure as in Olalde et al. 2019<sup>36</sup>, we trimmed adapter sequences, merged paired-end sequences, aligned to both the human reference genome (hg19) and the mitochondrial genome (RSRS) using BWA v0.6.146 and removed PCR duplicate sequences. The computational pipelines are available on github (<a href="https://github.com/DReichLab/ADNA-Tools">https://github.com/DReichLab/ADNA-Tools</a>, <a href="https://github.com/DReichLab/adna-workflow">https://github.com/DReichLab/adna-workflow</a>).</p>
Timing and spatial scale	The samples were taken during two sampling visits: the first in 2018 and the second in 2021.
Data exclusions	The only excluded sample was an individual that yield less than 5000 single nucleotide polymorphisms (SNPs) covered at least once, which is too little data for meaningful analysis.
Reproducibility	Only a single library can be made from each extract aliquot so no replication from the same extract is possible. However, the data from the 35 individuals came from 156 distinct libraries (range of 1-15 per individual). For the 29 individuals with more than one library, we had internal replication confirming that the libraries were from the same individuals.
Randomization	The sampling attempted to obtain DNA from every individual in the tomb, and so we aimed to analyze a sample from every individual that appeared osteologically distinct as well as additional samples that potentially could be osteologically distinct.

Blinding

Co-analysis of the genetic and archaeological data was central to the study, so we could not be blind to the sample identity.

Did the study involve field work?  Yes  No

## Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

### Materials & experimental systems

- |                                     |   |
|-------------------------------------|---|
| n/a                                 | Included in the study   |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Antibodies                               |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Eukaryotic cell lines                    |
| <input type="checkbox"/>            | <input checked="" type="checkbox"/> Palaeontology and archaeology |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Animals and other organisms              |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Human research participants              |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Clinical data                            |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Dual use research of concern             |

### Methods

- |                                     |   |
|-------------------------------------|---|
| n/a                                 | Included in the study                           |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> ChIP-seq               |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> Flow cytometry         |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> MRI-based neuroimaging |

## Palaeontology and Archaeology

- Specimen provenance The genetic samples were taken from human remains excavated in the 1981-2 excavations of the long barrow at Hazleton North, England, and subsequently curated at Corinium Museum, Cirencester, England.
- Specimen deposition The human remains are all still curated by Corinium Museum, and are either (a) currently in the Museum or (b) currently on loan to Harvard Medical School and scheduled for return to the Museum within 6 months of the completion of their analysis.
- Dating methods No new dates are provided.
- Tick this box to confirm that the raw and calibrated dates are available in the paper or in Supplementary Information.
- Ethics oversight Permission to study the samples was carried out with official permission from the Corinium Museum which curates the samples for scientific study.

Note that full information on the approval of the study protocol must also be provided in the manuscript.