New insights into the Tyrolean Iceman’s origin and phenotype as inferred by whole-genome sequencing

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The Tyrolean Iceman, a 5,300-year-old Copper age individual, was discovered in 1991 on the Tisenjoch Pass in the Italian part of the Ötztal Alps. Here we report the complete genome sequence of the Iceman and show 100% concordance between the previously reported mitochondrial genome sequence and the consensus sequence generated from our genomic data. We present indications for recent common ancestry between the Iceman and present-day inhabitants of the Tyrrhenian Sea, that the Iceman probably had brown eyes, belonged to blood group O and was lactose intolerant. His genetic predisposition shows an increased risk for coronary heart disease and may have contributed to the development of previously reported vascular calcifications. Sequences corresponding to ~60% of the genome of Borrelia burgdorferi are indicative of the earliest human case of infection with the pathogen for Lyme borreliosis.
In September 1991, two hikers discovered a human corpse (Fig. 1), partially covered by snow and ice, on the Tisenjoch Pass in the Italian part of the Ötztal Alps. The Tyrolean Iceman, a 5,300-year-old Copper age individual, is now conserved at the Archaeological Museum in Bolzano, Italy, together with an array of accompanying artefacts. An arrowhead lodged within the soft tissue of the left shoulder, having caused substantial damage to the left subclavian artery, indicated a violent death\(^1\). Speculations on his origin, his life habits and the circumstances surrounding his demise initiated a variety of morphological, biochemical and molecular analyses. Studies of the Iceman's mitochondrial genome began 3 years after his discovery with an analysis of the HVSI region\(^2\) and led to the sequence analysis of the entire mitochondrial genome\(^3,4\). Although these mitochondrial DNA (mtDNA) studies yielded conclusive data, no successful amplification of nuclear DNA from the Iceman has been reported to date. As cold environments often provide for good biomolecular preservation and as methods for ancient DNA analysis have recently improved, a whole-genome sequencing study was initiated in 2010. A 0.1-g bone biopsy was taken from the Iceman's left ilium under sterile conditions in the Iceman's preservation cell at the South Tyrol Archaeological Museum in Bolzano, Italy, DNA was extracted at the Institute of Human Genetics in Tübingen, Germany and a sequencing library was generated at febit Gmbh in Heidelberg, Germany. Paired-end high-throughput sequencing on the SOLID 4 platform was performed at Life Technologies facilities, Beverly, MA, USA. We found indication for recent common ancestry between the Iceman and present-day inhabitants of the Tyrrhenian Sea (particularly Corsica and Sardinia), that the Iceman probably had brown eyes, belonged to blood group O and was probably lactose intolerant. We further found a genetic predisposition for an increased risk for coronary heart disease (CHD), which may have contributed to the development of previously reported vascular calcifications. In addition, we found sequences corresponding to ~60% of the genome of \(B.\) burgdorferi that are indicative of the earliest human case of infection with the pathogen for Lyme borreliosis.

**Results**

**Overall sequencing results.** We obtained a total of \(2.93\times10^9\) sequencing reads out of which \(1.11\times10^9\) reads (37.9%) mapped uniquely to the human reference genome (hg18). Paired-end reads covered 96% of the reference genome with small variation among chromosomes (max = 97.74% for chr22, min = 89.61% for chrY) (Supplementary Fig. S1a and Supplementary Table S1). The average depth of coverage for non-redundant reads excluding duplicates is 7.6-fold, ranging from 4.76-fold for chromosome X to 10.8-fold for chromosome Y (Supplementary Fig. S1b). More than 65% of the genome is covered at least five-fold. Regions with more than \(\times100\) coverage were excluded from analysis, as these most likely represent PCR duplication artefacts.

Deamination-based damage, as the most common modification in ancient DNA\(^5,6\), potentially complicates correct identification of base substitutions that result from evolutionary processes. The sequencing of a 4,000-year-old palaeo-eskimo’s DNA sequence demonstrated that exclusion of postmortem, damage-based changes had little impact on the high-throughput sequencing result\(^5\). However, to estimate the extent of miscoding lesions (that is, postmortem changes in DNA), we determined the transition and transversion ratios (Supplementary Table S2 and Supplementary Table S3) and found them to be largely consistent with ratios observed for other genomes recently analysed by high-throughput sequencing.

One limitation to the whole-human genome sequencing has been the ability to provide broad access to the underlying sequence information in a useable format such that individuals can readily query the genome for their own specific research needs. To enable full access to the Iceman's genome to a broader scientific community, we have generated a genome browser (http://IcemanGenome.net)\(^8\).

This browser contains all the aligned sequencing reads to the Human Genome Reference 19, provides a basic level of annotation including individual single-nucleotide polymorphisms (SNPs), insertions and deletions, and gene identification. We have also incorporated a query engine for more complicated analysis routines, such as the comparison of the Iceman's genome to other contemporary genomes (Methods).

**Contamination controls.** Assessment of the percentage of miscoding lesions and potential human contamination was carried out through analysis of the mtDNA. As the mitochondrial genome is naturally at a very high ratio as compared with the nuclear genome (\(\times1160\) coverage versus \(\times7.6\) in this specific case), it provides a good target to identify low-frequency variants that can be caused by contamination. Additionally, the Iceman's mitochondrial genome had previously been sequenced in a different laboratory using a different sample\(^3\), thus serving as an ideal independent comparison. Allele ratios were determined for the previously noted positions at which the Iceman differs from the Cambridge Reference Sequence (rCRS, NC_012920). For each read not consistent with the known Iceman genome, the specific exchange type was examined as to whether it is a typical deamination product or whether it constitutes a known human mitochondrial variant (Supplementary Fig. S2, Supplementary Table S4). The average rate of concordance with the known Iceman genome is over 94%, the percentage of exchanges corresponding to miscoding lesions is under 5% and the rate of exchanges corresponding to other known human sequences is <2.5%. While the latter may be caused by modern contamination (either human or other), it may also suggest heteroplasy of certain positions of the Iceman's mitochondrial genome. Although this phenomenon was thought to be rare, recent studies have shown that heteroplasy is not uncommon in healthy mitochondrial cells\(^6\). This analysis suggests a maximum potential contamination level of 2.5%, an amount that would have little impact on the allele calls for the nuclear genome.

The next step comprised the analysis of DNA damage patterns. As sequences derived from ancient DNA molecules show characteristic nucleotide composition patterns, these can be used to distinguish them from contaminants of modern DNA fragments. In particular, postmortem deamination of cytosine to uracil leads to an increased rate of observed C to T transitions, particularly near the ends of the DNA fragments\(^7\). Furthermore, postmortem DNA fragmentation occurs at an increased rate at the 3′-end of purine

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**Figure 1 | The Tyrolean Iceman.** The mummy of the Tyrolean Iceman in his preservation cell at the Archaeological Museum of Bolzano.
nucleotides due to depurination, leading to a characteristic enrichment in purines at the base preceding the sequencing read.

In order to determine whether the Iceman DNA shows these characteristic patterns, we analysed a random subsample of the aligned sequencing reads for their nucleotide composition and misincorporation patterns. From each of the SAM files containing the aligned read data of the three sequencing slides, we randomly sampled one out of every 1,000 reads, and combined them into an initial analysis file containing a total of 1,156,960 reads. To simplify the subsequent nucleotide substitution analysis, we removed reads that showed insertions or deletions with respect to the reference sequence, as well as reads that mapped closer than 10 bp from the edges of the respective chromosome. This final read set contained 1,148,786 reads, and all subsequent results are based on this set. For each read, we then retrieved the corresponding genomic sequence from the human genome hg18 build, extended to include sequence up to 10 bp upstream (downstream) for reads that map to the forward (reverse) strand.

The average nucleotide composition around the ends of the analysed reads shows the typical pattern expected for ancient DNA fragmentation (Fig. 2a and b). A sharp increase in the frequency of purines (A and G) is observed at the first base upstream of the forward reads, accompanied by a complementary increase of pyrimidines (C and T) just downstream of the reverse reads. Additionally, substitution rates for each of the possible nucleotide mismatches along the sequencing read also show the expected pattern. While substitution rates are generally low (<0.25%) for most mismatch types, C to T transitions show an increased overall rate, in particular towards the end of the reads (up to 3.5% at the first nucleotide). As above, the increase in C to T transitions for the forward reads are accompanied by the complementary increase in G to A transitions observed for the reverse reads. Overall, these results are comparable to other ancient human genomes recently published, such as the Saqqara and the Australian Aboriginal genome8,9, and therefore provide additional indication that the sequences are indeed ancient.

**SNP analysis.** To gain first insight into genetic traits and into physiological and pathological characteristics of the Iceman, we identified 2.2 million SNPs in the Iceman’s genome sequence of which 1.7 million SNPs have previously been reported in dbSNP, with ~900,000 and 800,000 positions being present in a heterozygous and a homozygous state, respectively (Supplementary Table S5). As the 1.000 Genomes Project has demonstrated, using higher sequence coverage should result in the detection of ~3.5 million SNPs per individual; however, due to the limited amount of ancient DNA, lower sequencing coverage was generated, resulting in the detection of fewer SNPs. As shown in Supplementary Table S5, we found a concordance rate of ~92% for homozygous SNPs but only of 55% for heterozygous SNPs indicative of an undercalling of heterozygous SNPs as expected for 7.6-fold coverage.

We compared the number and frequency of homozygous to heterozygous SNPs known from dbSNP on each chromosome. The highest frequency of homozygous SNPs among autosomes was detected for chromosome 13, where 29,400 of the 43,680 SNPs were shown to be homozygous (67.3%). We detected heterozygous SNPs in regions with high similarity to other genomic regions and pseudo-autosomal regions. However, the overall highest rate of homozygous SNPs, namely 96.8% (16,080 of 16,606 SNPs), was detected in the X chromosome. As the Iceman was a male, these results further demonstrate the low potential degree of contamination, consistent with the results derived from the mtDNA analysis.

We furthermore analysed SNPs, including variants in specific genomic regions, of potential clinical or functional relevance. In the 5,300-year-old mummy, our analysis did not identify variations in human accelerated regions10, which are apparently functional sections of the genome showing human-specific evolution over the last few million years.

SNPs of potential clinical relevance with additional direct assessment of further SNPs of phenotypic interest (for example, lactase persistence, pigmentation and blood group) are summarized in Supplementary Table S6. Not all listed SNPs necessarily infer a phenotypic consequence or a high predisposition risk. A number of variants supported by morphological or radiological data (such as risk factors for atherosclerosis), or variants that are likely to lead to a specific phenotype (such as lactase nonpersistence) have been resequenced using the Sanger method. These are listed together with individual coverage depth in Table 1. Sanger sequencing, carried out using extract from a separate sample, confirmed the next-generation sequencing results in all cases where amplification was successfully carried out, further underlining the low degree of contamination. Additionally, some of the characteristic SNPs of the Iceman’s mitochondrial genome were screened using mitochondrial primers to confirm authenticity (Table 2).

As a next step we analysed the genetic ancestry of the Iceman. As the Iceman’s mtDNA haplotype has not yet been detected among thousands of sampled contemporary individuals, it has been difficult to assess his genetic ancestry with high accuracy. The first analysis was to determine if the Iceman’s autosomal DNA shows an affinity to any specific population or if he remains an outlier among contemporary samples. We intersected genotype calls of greater than ×6 coverage from his genome with the population reference sample consisting of more than 1,300 Europeans genotyped for SNPs on the Affymetrix 500K array11,12, 125 individuals from seven North African populations ranging from Egypt to Morocco on the Affymetrix 6.0 array and 20 Qatari samples from the Arabian Peninsula13. When plotting the Iceman’s genotype along the first two major axes of variation in Principal Component space (PC1 versus PC2), PC1 is driven by a north-to-south gradient differentiating North Africans from Europeans, and PC2 aligns individuals along north-to-south gradient within Europe (Fig. 3a). The Iceman clusters nearest to southern European samples, suggesting no greater genetic affinity with the North African or Middle Eastern components of variation than present day southern Europeans (Fig. 3a). When considering only European populations, however, we observe that the Iceman clusters closest with five outlier contemporary samples from south-western Europe. In particular, the Iceman abuts the Italian samples originating from geographically isolated regions such as Sardinia (Fig. 3b). Analysis of a larger set of samples, including
Sardinians from HGDP across a smaller subset of SNPs, further supports the clustering of the Iceman with samples from Sardinia based on autosomal SNPs (Fig. 3f). A potential reason for this may be an error in genotype calls due to low-coverage sequencing. To evaluate this hypothesis, we repeated the principal component analysis (PCA) analysis including

### Table 1 | Verification of nuclear SNPs.

<table>
<thead>
<tr>
<th>dbSNP # (b126)</th>
<th>Association</th>
<th>Forward primer 5’-3’</th>
<th>Reverse primer 5’-3’</th>
<th>Fragment size (bp)</th>
<th>AT (°C)</th>
<th>Independent PCR replication results</th>
<th>NGS data coverage</th>
<th>HapMap frequency of Iceman's genotype (sample size)</th>
<th>Primer reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>rs10757274</td>
<td>Coronary artery disease</td>
<td>CCCCCGTG GTCTAAG TCTAG</td>
<td>AGAATCCC TACCCCTAC ATCCTCTCT</td>
<td>82</td>
<td>55</td>
<td>nsa</td>
<td>8G, 1A</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>rs2383206</td>
<td>Coronary artery disease</td>
<td>TCTATTCT GCTTGGTC CCCCTTTGCT</td>
<td>GGTTCAAGGA TTCAAGGCACT TCGTG</td>
<td>78</td>
<td>55</td>
<td>G/G</td>
<td>8G</td>
<td>G/G=0.246 (130)</td>
<td>NA</td>
</tr>
<tr>
<td>rs5351</td>
<td>Atherosclerosis</td>
<td>CTATCCCTA ATGTTTTTAC AAGCAGCC</td>
<td>ATGGCCAAT GCAAGCAGA</td>
<td>74</td>
<td>55</td>
<td>C/T</td>
<td>20T, 14C</td>
<td>C/T=0.416 (226)</td>
<td>C/T=0.567 (194)</td>
</tr>
<tr>
<td>rs4988235</td>
<td>Lactase non-persistence Y-hg G</td>
<td>AATGCAGGG CTCTAAAGAC ACA</td>
<td>ACGATGCAGGG CATCTACGC TTCAACCAAA CTACAACT</td>
<td>111</td>
<td>53</td>
<td>G/G</td>
<td>14G</td>
<td>G/G=0.088 (226)</td>
<td>G/G=0.833 (204)</td>
</tr>
<tr>
<td>rs2032636</td>
<td>Y-hg G</td>
<td>GTCTATGTGCT AGACAAAGA</td>
<td>GTAAGCTGTA CACATGAAAA AGATG</td>
<td>72</td>
<td>59</td>
<td>T/T</td>
<td>7T</td>
<td>T/T=0.000 (29)</td>
<td>NA</td>
</tr>
<tr>
<td>rs2178500</td>
<td>Y-hg G2a</td>
<td>GTCTATGTGCT AGACAAAGA</td>
<td>GTAAGCTGTA CACATGAAAA AGATG</td>
<td>64</td>
<td>54</td>
<td>nsa</td>
<td>4G</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>rs7892988</td>
<td>Y-hg G2a3</td>
<td>GTCTATGTGCT AGACAAAGA</td>
<td>GTAAGCTGTA CACATGAAAA AGATG</td>
<td>70</td>
<td>54</td>
<td>nsa</td>
<td>7T</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>rs17817449</td>
<td>BMI</td>
<td>AAGAAAGAGT TGCTGAGA GCACCTAAG</td>
<td>AAGAAAGAGT TGCTGAGA GCACCTAAG</td>
<td>81</td>
<td>53</td>
<td>nsa</td>
<td>5C</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>rs1426654</td>
<td>Skin colour (light)</td>
<td>GTATATGTT CAGGCTTGG ATTGTC</td>
<td>CAAGGGATCT TGAACAGGACTG</td>
<td>95</td>
<td>55</td>
<td>A/A</td>
<td>29A</td>
<td>A/A=1,000 (126)</td>
<td>A/A=0.990 (204)</td>
</tr>
</tbody>
</table>

**Abbreviations:** NA, not applicable; nsa, no successful amplification/sequencing; n.n., no name (no RefSNP accession ID).


Independent verification was carried out for several autosomal and Y-chromosomal SNPs. Owing to the strictly limited amount of available sample material, PCR was only carried out once for each SNP, and verification was limited to eight SNPs.

### Table 2 | Mitochondrial primer sequences.

<table>
<thead>
<tr>
<th>Variant position (rCRS/ Iceman)</th>
<th>Forward primer 5’-3’</th>
<th>Reverse primer 5’-3’</th>
<th>Laboratory sample</th>
<th>Fragment length including primers</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>8137 (C/T)</td>
<td>L8105: GTCCCCACATTAGGGCTAAACAGATG</td>
<td>H8241: CCCTAGTATACCCCCGGCTGAG</td>
<td>Bolzano, muscle sample</td>
<td>89 bp</td>
<td>This study</td>
</tr>
<tr>
<td>16224 (T/C)</td>
<td>L16209: CCCCCATGCTTACAAGCAAGT</td>
<td>H16348: ATGGGGAGCAAGGGATTTG</td>
<td>Tübingen, bone sample; Bolzano, muscle sample</td>
<td>179 bp</td>
<td>Handt 1996; Haak 2005</td>
</tr>
<tr>
<td>16311 (T/C)</td>
<td>L16287: CACCCCATGCTTACAAGCAAGT</td>
<td>H16397: GTGAGAGCTGTCGAG</td>
<td>Bolzano, muscle sample</td>
<td>162 bp</td>
<td>Handt 1996</td>
</tr>
<tr>
<td>16326 (T/C)</td>
<td>CACTAGGATGACCAAAACC</td>
<td>H16397: GTGAGAGCTGTCGAG</td>
<td>Bolzano, muscle sample</td>
<td>137 bp</td>
<td>This study</td>
</tr>
<tr>
<td>16311 (T/C)</td>
<td>L16287: CACCCCATGCTTACAAGCAAGT</td>
<td>H16397: GTGAGAGCTGTCGAG</td>
<td>Bolzano, muscle sample</td>
<td>131 bp</td>
<td>Handt 1996</td>
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<td>16326 (T/C)</td>
<td>CACTAGGATGACCAAAACC</td>
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<td>Bolzano, muscle sample</td>
<td>131 bp</td>
<td>Handt 1996; This study</td>
</tr>
</tbody>
</table>

To establish the authenticity of the sample extracts before whole-genome sequencing (and replication attempts) of nuclear SNPs, a selection of the Iceman’s known mitochondrial variants, hitherto unique in this combination*, were tested.
Figure 3 | Autosomal and Y-chromosome evidence of the Iceman’s origin. (a) Principal component analysis (PCA) of the Icman genome and SNP array data for European, North African and Near Eastern populations. Population samples were as follows: Europe S (for example, Italy), Europe SW (Spain/Portugal), Europe W (for example, France), Europe C (for example, Germany), Europe NW (for example, UK), Europe NNE (for example, Sweden), Europe SE (for example, Greece), Europe ESE (Cyprus/Turkey), North Africa (Algeria, Libya, Egypt, Morocco, Sahara and Tunisia) and Near East (Qatar). PCA was performed using 123,425 SNPs. (b) PCA projecting the Icman into genetic clusters within Europe. PCA was performed using 132,981 SNPs after intersecting the Icman genome with 1,387 European samples from the PopRes dataset. Samples clustering close to the Icman were of Sardinian origin. (c) The phylogenetic relationship of Y-chromosome haplogroup G2a4 within haplogroup G was constructed using markers ascertained in the Icman as listed in Supplementary Table S7. Unboxed and boxed marker labels indicate observed ancestral and derived allele status, respectively. (d) Spatial frequency distribution of G2a4-L91 chromosomes in Europe. Appropriate sample locations of 7,797 individuals from 30 populations are indicated with red dots. (e) Spatial distribution of L91 frequency in various regions of Corsica and Sardinia. Marker L91 was genotyped by HaeIII RFLP analysis, which identifies a G to C transversion at position 250 within the 447-bp PCR fragment amplified using primers (F) 5’-cttgccatcctgcaaggg-3’ and (R) 5’-gtggaggtgctgccgctc-3’. The frequency data were converted to spatial-frequency maps using Surfer software (version 7, Golden Software, Inc.), following the Kriging procedure. (f) PCA of the Icman genome combined with SNP array data for 1,387 European samples from PopRes and 28 Sardinians from HGDP. PCA was performed using 28,003 SNPs.

data from five HapMap CEU individuals with high-confidence genotype calls on multiple array platforms that were also sequenced as part of the 1,000 Genomes Project. To establish controls for the PCA analysis and to address any issues that may arise from the low sequencing coverage of the Icman, we randomly chose five CEU individuals from the Pilot 1 data release, which had a similar (or lower) coverage to the Icman, and were sequenced using the same SOLiD technology. The 1,000G CEU sample genotypes were inferred using a single-sample procedure similar to that used for the Icman (Supplementary Methods). SNP array genotype data for the same individuals were obtained from HapMap release 28. The merged HapMap/1,000 Genomes/PopRes/Icman dataset contained 125,729 SNPs. The 1,000 Genomes control samples were projected onto the PC space inferred from the rest of the dataset using both the low-coverage SNP calls and the high-confidence HapMap genotypes. The low-coverage 1,000 Genome control samples fall closer to the middle of the PCA plot, near central Europe, when compared with their corresponding genotype HapMap data (Supplementary Fig. S3a). The 1,000 Genomes control samples illustrate how low-sequence-coverage genomes tend to project towards the averaged centre of the distribution due to the missing genotypes. Thus, the Icman’s position along with the 1,000 Genomes control samples is shifted towards the origin as those missing genotype values are set to 0 after the normalization step in the PCA. When using a ×12 threshold (though with fewer loci) the placement of the Icman shifts from the European continent closer to the Sardinian sample group, as there are no missing genotypes, reflecting his true position more accurately. (Supplementary Figs S3b–d show placement of the Icman on PopRes PCA based on varying degrees of coverage and thresholding for missing genotypes.) To determine the Icman’s paternal ancestry, his Y-chromosome haplogroup allocation was assessed according to the hierarchical
order of markers organized by the International Society of Genetic Genealogy\textsuperscript{14}. The detection of four phylogenetically equivalent SNPs\textsuperscript{15} and an independent verification of the rs2032636 (M201) SNP by Sanger sequencing of a PCR amplicon obtained using genomic DNA isolated from a 2007 muscle biopsy unequivocally assigns the Iceman to haplogroup G, specifically subgroup G2a (Supplementary Table S7). While haplogroup G displays its highest frequency in the Caucasus\textsuperscript{16}, it is also present at ~11% in present day Italy\textsuperscript{17}. Genetic analysis of ancient DNA from 5,000-year-old skeletons from a burial cave site in southern France were mostly assigned to haplogroup G2a-P15 (ref. 18). Haplogroup G2a3 was also reported in an ancient DNA sample of an early Neolithic individual from Saxony-Anhalt, Germany\textsuperscript{19}. We used the phylogenetic relationships of the ancestral and derived allelic states for 13 other detected SNPs within the G hierarchy (Supplementary Table S7, Fig. 3c) to place the Iceman into the G2a4-L91 lineage that is notably divergent from G2a3-U8 lineages typical of modern continental Europeans. Knowledge about the phylogeography of haplogroup G2a4 is unreported. We addressed this issue here by analysing the G2a4-defining L91 SNP in 7,797 chromosomes from 30 regions across Europe. Fig. 3d shows the spatial frequency distribution of G2a4 throughout Europe. The highest frequencies (25 and 9%) occur in southern Corsica and northern Sardinia, respectively, (Fig. 3e) while in mainland Europe the frequencies do not reach 1%.

We also performed a detailed analysis of the Iceman's phenotype and the diseases he may have suffered from. One trait associated with the beginning of agriculture in Europe is lactase persistence (the ability to digest milk after early childhood), which is commonly associated with a polymorphism in the MCM6 gene (\( \sim 13.910^{-1} \))\textsuperscript{20}. Palaeogenetic analyses from various prehistoric sites failed to detect the derived allele in any of the tested Neolithic samples, indicating that lactase persistence was rare in the Neolithic and, due to the substantial selective advantage conferred by this trait, gained in frequency over the next millennia and was widespread in Central Europe by the Middle Ages\textsuperscript{21}. Comparisons between genotype and phenotype (diagnostic methane–hydrogen breath test) in South European individuals have shown that the homozygous ancestral C allele causes clinical symptoms in over 85% of cases\textsuperscript{22}. Although a small number of genetically non-lactase-persistent individuals show no malabsorption problems, this may not least depend on the age variation of the study group: the onset age of lactose malabsorption is at about 50 years in the range of the study\textsuperscript{23}. The most strongly associated variant, rs12913832, shows the homozygous A allele in the Iceman's genome (coverage depth \( \times 31 \)), which is associated with brown eye colour in over 80% of cases even when regarded alone\textsuperscript{30}. Branicki et al.\textsuperscript{31} defined a haplotype of five SNPs as predictor: rs4778138, rs4778241, rs495174, rs12913832 and rs916977. The Iceman's haplotype (for which TTTTC and TTTTT were addressed together as the last SNP is a clear heterozygote) in the Branicki study encompassed 46 individuals, 8 of which were blue-eyed, the residual 38 having a non-blue eye colour. The haplotype defined by Sturm et al.\textsuperscript{30} narrows the field down further: individuals who shared the Iceman's A-AAA haplotype had blue eyes in under 5% of cases, 40% having green or more often hazel eyes, while 55.7% of individuals had brown eyes.

Further, phenotypically relevant variants include a homozygous deletion (coverage depth \( \times 7 \)) at rs8176719, a T allele at rs505922 and lack of significant deletions in the \( RHD \) gene, which are characteristic of blood group O Rh-positive carriers\textsuperscript{32,33}.

**Metagenomic analysis.** Metagenomic analysis was performed using the BlastN algorithm. Despite the presence of an exceptionally high number of human reads (77.6%), only a small proportion of the assignable reads derive from the bacterial domain (0.84%). Most prevalent bacterial species were found to be in the phylum Firmicutes, mostly the spore-forming Clostridia (72%). Within the Spirochaetes phylum, 45.7% of the reads (0.16% of the total bacterial hits) were assigned to sequences of the pathogen \( B. burgdorferi \), which is known to cause Lyme disease in humans (Fig. 5). To confirm this result of the metagenomic approach, the entire number of reads was mapped against the \( B. burgdorferi \) reference genome and the strain-specific \( B. burgdorferi \) plasmids. One million sequence reads covered about 60% of the \( B. burgdorferi \) genome including a high representation of the bacterial epsipeme, which was covered up to 67% at least once. While several reads were specific for Borrelia, the 60% coverage still represents an upper boundary as reads

![Figure 4 | CT image of abdomen and coronal reconstruction. The aorta is visible as a strip-shaped structure to the right of the vertebral column. Two calcifications constituting atherotic plaques are visible at the left contour of the aortic bifurcation.](image)
may map to other species besides *Borrelia*. The coverage of selected *Borrelia* genome and plasmid sequences for our NGS data is presented in Supplementary Table S9. Remarkably, *B. burgdorferi* sensu stricto infections have been linked to vascular calcifications19,23. In summary, our data point to the earliest documented case of a *B. burgdorferi* infection in mankind. To our knowledge, no other case report of borreliosis is available for ancient or historic specimens.

**Discussion**

Next-generation sequencing technology enabled the reconstruction of the nuclear genome of the Tyrolean Iceman. For control purposes, temporally and spatially separate DNA analysis was performed on different samples, yielding consistent data. The mitochondrial consensus generated from our data showed 100% concordance with the previously published Iceman mitochondrial genome, underlining the sequence authenticity. Sequence analysis showed genetic distance from modern mainland European populations, but proximity to the extant populations of Sardinia. Interestingly, the Iceman's Y-haplogroup G2a4 has hitherto only been found at appreciable frequency in Mediterranean islands of the Tyrrhenian Sea (Sardinia and Corsica). Although admixture and demographic history cannot be reconstructed from one individual alone, the Iceman's Y-chromosomal data document the presence of haplogroup G in Italy by the end of the Neolithic and lends further support to the demic diffusion model36. The affinity of the Iceman's genome to modern Sardinian groups may reflect relatively recent common ancestry between the ancient Sardinian and Alpine populations, possibly due to the diffusion of Neolithic peoples. The Iceman's stable isotope constitution37 that localized his residential mobility to the Alpine region, taken together with their initial genomic ancestry profile, suggests differential genetic destinies for populations of mainland Europe and those of the islands of the Tyrrhenian Sea that began to diverge at least 5,000 years ago. The Iceman's genetic signature may at one time have been more frequent in Neolithic South Tyrol although further ancient DNA analyses from these regions will be necessary to fully understand the genetic structure of ancient Alpine communities and migration patterns between the insular and mainland Mediterranean.

Autosomal data also yielded novel insights into the Iceman's phenotype, such as eye colour and the inability to digest lactose. The availability of the complete, excellently preserved body allows comparisons between genetic data and observed morphological and radiological characteristics, such as the vascular calcifications diagnosed in previous CT analyses, which were now shown to have a probable hereditary component. The discovery of predisposition risk factors for atherosclerosis points towards a hereditary component for the vascular calcifications radiologically observed in the mummy. Finally, the detection of *B. burgdorferi* is the oldest documentation of this pathogen in a human to date.

**Methods**

**DNA extraction.** DNA extraction from the bone sample whose DNA was subjected to next-generation sequencing was performed at the Division of Molecular Genetics, Institute of Human Genetics at the Eberhard-Karls-University of Tübingen, Germany. All surfaces, instruments and disposables were treated with bleach (Sodium hypochlorite 3%, Roth), DNA away solution (Roth) and/or irradiated with UV light. Blank controls were carried throughout all steps to monitor possible contamination during laboratory procedures. The sample was ground into fine bone powder using a mortar and pestle after brief nitrogen deep freezing. DNA extraction was carried out according to the protocol described by Scholz and Pusch38.

Aqueous solutions of PCR components were screened for purity with a highly sensitive contamination monitoring protocol (intra Alu-PCR) reported elsewhere39,40.

**Library preparation.** The library was prepared following the SOLID protocol for low-input fragment library preparation. The library was mainly prepared following the manufacturer's instructions. Variations from the protocol are listed in the following.

A total amount of 22.6 ng of unsheared genomic DNA diluted in 10 μl water was end-repaired using 1 μl End Polishing Enzyme 1 (100 μl −1) and 2 μl End Polishing Enzyme 2 (10 μl −1), 4 μl dNTP mix (10 mM), 20 μl ×5 End-Polishing Buffer in a total volume of 100 nl. Following incubation at room temperature for 30 min, DNA was purified using SOLID Library Column Purification Kit. SOLID Adaptors (P1: 5′-GGACCAGCGCCGCCGCTCTCCTCTATGGGCAGTCGGTGGAAGCTGGATT-3′; P2: 5′-AGAGAATGAGGACCGGCAGGGCCAGTT-3′), each at 2.5 μl, were ligated to purified DNA using 40 μl ×5 T4 Ligase Buffer, 10 μl T4 Ligase (5 μl −1) in a total volume of 200 μl by incubation at room temperature for 15 min. After another purification step, ligated DNA was eluted in 40 μl nuclease-free water. No additional size selection was carried out to avoid loss of material. DNA was then incubated with 380 μl Platinum HiFi PCR Amplification Mix and 100 ng of both Library PCR Primers each ( Primer 1: 5′-CCACTAGCGCCGCTCTCCTCTTCCTCATTG-3′, Primer 2: 5′-CTGCCCGGCTCTCCTATCCTT-3′) to repair the gap in the double-stranded DNA molecules introduced during adaptor ligation using the following cycling conditions: ×1 72 °C 20 min; ×1 95 °C 5 min; 2 cycles 95 °C 5 s, 62 °C 15 s, 70 °C 1 min, one cycle 70 °C 5 min and hold at 4 °C. DNA was purified using the PureLink® PCR Purification Kit (Invitrogen). Eluted DNA (40 μl) was again cycled using 100 μl ×2 Phusion HF Master Mix (Finnzyme) and 8 μl each of both Library PCR Primers 1 and 2 in a total volume of 200 μl. This mixture was divided among four wells of a 96-well plate and cycled using the following conditions: 12 cycles 95 °C 1 s, 62 °C 15 s, 70 °C 1 min; and 1 cycle 70 °C 5 min, hold at 4 °C. PCR products were purified as described above. Six different dilutions of the eluate were quantified, yielding an average concentration of the undiluted sample of 104.3 ng μl −1. A 2 ng μl −1 dilution was sequenced using a paired-end sequencing protocol on a SOLID+4 at the Applied Biosystems facility in Beverly, USA.

**Sequencing.** Starting with the 2 ng μl −1 dilution of the fragment library preparation, a 60 pg μl −1 dilution was applied to the emulsion PCR to perform single-molecule amplification of the library and to template small magnetic beads. The emulsion was subsequently broken via organic solvent; beads were washed, enriched for templated beads and isolated using the EZBead system. After a 3′ end modification of the DNA bound to magnetic beads, templated beads were deposited onto chemically modified slides and loaded into three different flow cells of a SOLID+4 instrument. The DNA was sequenced using the SOLID+ TOP Paired-End Sequencing Kit MM500/25; 50 bp were sequenced via ligation-based sequencing, starting at the P1 adaptor (forward direction) and 25 bp starting at the P2 adaptor (reverse direction). The resulting sequence data were mapped to hg18 and the analysis was done off instrument.

**Next-generation sequencing data analysis.** SOLID data were mapped and paired using the mapreads program incorporated into the Bioscope analysis pipeline (Life Technologies). The 50-bp-long F3 tag was mapped with a seed-and-extend approach where the data are first mapped with up to two mismatches in a 25-bp seed and alignment is potentially extended further by an Iceman alignment. The shorter 25-bp F5 tag was mapped as a 25mer allowing up to two mismatches and no extension of any kind was attempted. After mapping, the data were then paired with Bioscope, which is the process of matching up the mapped F3 and F5 reads attempting to find the best possible pair where multiple mappings are found. During pairing, the insert size is calculated for the two tags being paired and this insert size is used to classify the pairs; a pair that is of the expected order, orientation and insert size is classified as a good mate. To assure that the data are
not biased by PCR duplicates and other redundant molecules that may be over-represented in the library, especially considering the age of the sample, the three slides of data were merged together and PCR duplicates were removed using Picard (http://picard.sourceforge.net/). These non-redundant good mates—along with any singletons, which were flagged as primary alignments by mapreads but do not have a mapped mate—were used for SNP calling and the resultant coverage of these reads was >7.6. The SNP caller, diBayes (http://solidsoftwarertools.com/gf/project/dibayes/), is also part of the Bioscope package. Default medium stringency settings were applied.

**Functional SNP analysis.** SNPs of potential clinical relevance were primarily identified through comparison with the Human Gene Mutation Database release 2011.1, with additional direct assessment of further SNPs of phenotypic interest (for example, lactase persistence, pigmentation and blood groups) and are summarized in Supplementary Table S6. Allele frequency was derived from the 1,000 Genomes Project data via dbSNP. To determine coverage depth for each allele in the Iceman’s genome, generated reads were viewed in the Integrative Genomics Viewer (IGV, http://broadinstitute.org/software/igv/), version 1.5.64. Y-chromosomal haplotype allocation was carried out according to the nomenclature of the Y-Chromosome Consortium. In addition, more recently reported subvariants of haplogroup G2a were characterized in the Iceman. Supplementary Table S8 contains a detailed analysis of all identified SNPs as to formation of premature stop codons, altered start codons and readthrough stop codons in genes.

**Principal component analysis.** We used PCA to identify clusters of genetic similarity among samples with European ancestry and projected the inferred genotype for the Iceman sample based on aDNA sequencing to 7.6-fold coverage. We used the same subset of samples and SNPs reported by Novembre et al. In order to reproduce the PCA map of Europe, which includes 1,387 samples from throughout Europe with four grandparents born in a single country and 197,146 autosomal SNPs. As predicted from the aDNA sample coverage, 31% of the SNPs had missing genotypes in the Iceman, so after quality-control filtering for missingness, a total of 132,981 SNPs remained for subsequent analyses. PCA was performed using the GATK unified genotyper to call genotypes, with default parameters for the Iceman sample based on aDNA sequencing to 7.6-fold coverage. We used the GATK unified genotyper to call genotypes, with default parameters for the Iceman sample based on aDNA sequencing to 7.6-fold coverage.

**Artificial SNP analysis.** SNPs of potential clinical relevance were primarily identified through comparison with the Human Gene Mutation Database release 2011.1, with additional direct assessment of further SNPs of phenotypic interest (for example, lactase persistence, pigmentation and blood groups) and are summarized in Supplementary Table S6. Allele frequency was derived from the 1,000 Genomes Project data via dbSNP. To determine coverage depth for each allele in the Iceman’s genome, generated reads were viewed in the Integrative Genomics Viewer (IGV, http://broadinstitute.org/software/igv/), version 1.5.64. Y-chromosomal haplotype allocation was carried out according to the nomenclature of the Y-Chromosome Consortium. In addition, more recently reported subvariants of haplogroup G2a were characterized in the Iceman. Supplementary Table S8 contains a detailed analysis of all identified SNPs as to formation of premature stop codons, altered start codons and readthrough stop codons in genes.

**References**

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Author contributions


Additional information

Accession codes: The sequencing data have been uploaded to the European Nucleotide Archive under accession code ERP001144.

Supplementary Information accompanies this paper at http://www.nature.com/naturecommunications

Competing financial interests: I.S., S.M., M.S., C.L., T.T.H. and A.S. are employees of Life Technologies. M.M., V.B. and A.K were previously employed by Fehlt biomed GmbH.

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