Comparative genomic and phylogeographic analysis of *Mycobacterium leprae*

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Reductive evolution and massive pseudogene formation have shaped the 3.31-Mb genome of *Mycobacterium leprae*, an unculturable obligate pathogen that causes leprosy in humans. The complete genome sequence of *M. leprae* strain Br4923 from Brazil was obtained by conventional methods (6× coverage), and Illumina resequencing technology was used to obtain the sequences of strains Thai53 (38× coverage) and NHDP63 (46× coverage) from Thailand and the United States, respectively. Whole-genome comparisons with the previously sequenced TN strain from India revealed that the four strains share 99.995% sequence identity and differ only in 215 polymorphic sites, mainly SNPs, and by 5 pseudogenes. Sixteen interrelated SNP subtypes were defined by genotyping both extant and extinct strains of *M. leprae* from around the world. The 16 SNP subtypes showed a strong geographical association that reflects the migration patterns of early humans and trade routes, with the Silk Road linking Europe to China having contributed to the spread of leprosy.

Leprosy is a chronic dermatological and neurological disease that results from infection with the unculturable pathogen *M. leprae*¹ and causes nerve damage that can lead to severe disabilities. There is no known reservoir for *M. leprae* other than human beings. New opportunities for understanding the transmission of the leprosy bacillus and its phylogeny have arisen following the determination of the complete 3.3-Mb genome sequence of the TN strain, from Tamil Nadu, India².

A notable feature of the *M. leprae* genome is the exceptionally large number of pseudogenes, which occupy almost half of the TN chromosome². The resulting loss of function most likely accounts for the exceptionally slow growth rate of the bacillus and for researchers' failure to culture it *in vitro*. Given this extensive genome decay, one might expect to find more genetic variability between different isolates of *M. leprae*, but initial analysis of SNPs demonstrated that these were very rare, occurring roughly once every 28 kb. Furthermore, all

extant isolates of *M. leprae* were nearly indistinguishable, belonging to one of only four SNP types, and are derived from a single clone³. Variable number tandem repeats (VNTRs) have also been investigated in *M. leprae*^{3–8} and, in some cases, have proved useful for countrywide epidemiological surveys⁹. However, owing to variability of VNTR profiles in samples taken from different sites on the same patient, their utility may be limited^{7,10,11}.

The emerging discipline of microbial phylogeography is a powerful means of monitoring not only the spread of microbes but also the movement of their hosts. For instance, compelling associations were found between the genotypes of *Helicobacter pylori* strains and their places of origin, and the migration and ethnicity of their human hosts^{12–14}. *M. leprae* is also proving useful in this respect, with its spread reflecting the migrations of early humans³, and similar studies with tuberculosis patients suggest that *Mycobacterium tuberculosis* lineages have also adapted to particular human populations¹⁵.

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Received 19 June; accepted 1 September; published online 1 November 2009; corrected after print 5 February 2010; doi:10.1038/ng.477

In the present work, the complete genome sequence of Br4923, a Brazilian strain of *M. leprae*, has been determined and compared with that of the TN isolate, leading to the discovery of a total of only 155 SNPs, of which 78 are informative. To deepen the comparison and avoid possible ascertainment bias¹⁶, the genomes of strains from North America and Thailand were resequenced using Illumina (Solexa) technology, revealing comparably low levels of diversity. For phylogeographic purposes, the presence of the 78 informative SNPs was subsequently surveyed in ~400 isolates, enabling classification of *M. leprae* into 16 SNP subtypes of limited geographic distribution that correlate with the patterns of human migrations and trade routes.

RESULTS

Complete genome sequence of Br4923

The Br4923 strain of *M. leprae* was chosen for complete genome analysis because it was originally isolated from a patient in Brazil, the country with the second highest leprosy burden, and because Brazil is geographically remote from India^{1,17}. The genome comprises 3,268,071 bp and is thus 141 bp smaller than that of the TN strain². No evidence was found for DNA inversions, translocations or

duplications, and no transposition or amplification of either the defective insertion sequence elements or the four families of dispersed repeats was detected¹⁸, consistent with the findings of an earlier quantitative PCR study³.

On alignment of the two genomes, 194 polymorphic sites were uncovered: these correspond to 155 SNPs, 31 VNTR regions and eight insertion or deletion (indel) events. On verification of the original sequence data, nine SNPs were found to stem from sequencing errors². The distribution of the true SNPs were as follows: 52 in genes encoding proteins, 39 in pseudogenes, 26 in noncoding regions (or as-yet-unidentified pseudogenes) and 38 in dispersed repeats. The majority of the 31 VNTR regions were found to affect homopolymeric tracts (HPT), di- and trinucleotide repeats, as well as some longer sequences, the longest of which is 52 bp in length and present in two and three copies in the TN and Br4923 strains, respectively. Variation in the di- and trinucleotide repeats is now well documented^{4,5,8–11}. It is noteworthy that, at the VNTR loci, the repeat copy number was consistently higher in the TN strain than in Br4923, as 23 of the 31 Brazilian VNTR were shorter than their Indian counterparts. Six of the eight indels occurred in repetitive elements and the others were in an intergenic region and the gene *ML0825c*.

Figure 1 SNP in dispersed repeats. The sites of SNP located within the RLEP, LEPREP, REPLEP and LEPRPT elements present in the genomes of the TN and Br4923 strains are shown and color coded. The position of the SNP is given above and the repeat identifier at left. When occupied by the same nucleotide in both strains, the coloring is continuous: for example, light blue for G. If the same change occurs within a repeat family, a different continuous color is used: for example, light orange for A. When a site is polymorphic between strains, discontinuous darker colors are used. Deletions are identified by *.

of the *M. leprae* strains Thai53 (38×) and NHDP63 (46×) from Thailand and the United States, respectively, and the reads were mapped onto the TN consensus sequence. The entire genome was covered by the assembly except for the dispersed repeats that could not be distinguished owing to the short read length. This analysis led to the identification of a combined total of 201 SNPs and 14 single-base indels, many of which were shared with the TN or Br4923 strains (Supplementary Table 1), and it uncovered five pseudogenes (Fig. 2a). A distance matrix was then established (Fig. 2b) and used to construct a phylogenetic tree by neighbor joining²⁰. The topology of the tree (Fig. 2c) is fully consistent with the previous scheme derived by SNP typing³.

Synonymous and nonsynonymous substitutions

Determining the ratio of nonsynonymous to synonymous SNPs (dN/dS) is a popular method used to predict whether any purifying selection is operating²¹. When the four genomes were compared, we found 43 (42%) synonymous and 59 (58%) nonsynonymous changes in the protein-coding genes (Supplementary Table 1); these are values similar to those reported for the *M. tuberculosis* complex (38% and 62%, respectively), in which SNPs are >10-fold more abundant²². The dN/dS ratios were calculated within the *M. leprae* genes through pairwise comparisons revealing an average value of 0.70. A ratio

close to 1 is indicative of no, or weak, selection against nonsynonymous SNPs as recently described for *Salmonella typhi* (with a dN/dS = 0.66)²³. Like *S. typhi*, another obligate human pathogen, *M. leprae* also seems to show little genetic drift, probably because of its small effective population size²¹. Phylogenetic trees were constructed using the nonsynonymous and synonymous SNPs²⁴, and these had similar topology to that shown in Figure 2c.

Phylogeographic survey of extant *M. leprae*

In our previous study, we had genotyped 175 strains of *M. leprae* from 21 different geographic origins but found only four phylogenetic groups: SNP-types 1–4 (ref. 3). A close relationship was established between the geographical origin of the strain and its genotype, giving a first impression of how leprosy may have disseminated around the globe as humans migrated. However, certain parts of the world were unrepresented, particularly the Middle East and Europe. Because these regions were of historical importance in the migration of human populations—notably the Middle East region, where early humans are thought to have arrived from Africa and from whence the European and Asian migrations began²⁵—we made intensive efforts to obtain samples from these settings. Samples were obtained from Iran and Turkey in the Middle East and from China, Korea and Japan in the Far East.

The 84 informative polymorphic sites were used in our survey (Fig. 3a, Supplementary Table 2), and these sites enabled us to reconstruct the movement of leprosy between peoples and countries. A total of 400 samples from 28 regions were successfully screened, resulting in the definition of 16 different *M. leprae* subtypes, as compared to the four originally described³; these are referred to as A–P (where A corresponds to the TN isolate and Br4923 is P, Fig. 3a). SNP type 1 is now subdivided into four subtypes (A–D), SNP type 2 into four subtypes (E–H), SNP type 3 into five subtypes (I–M) and SNP type 4 into three subtypes (N–P). For instance, subtypes A and B differ from each other at one HPT and 21 SNP. Likewise, although there are no SNP differences within members of SNP type 4, these can be subdivided based on differences in their indel or HPT profiles.

To exclude the possible effects of ascertainment bias¹⁶, we challenged this evolutionary model by phylogenetic analysis using other actinobacteria, such as *M. tuberculosis*, as an

Figure 4 Maximum likelihood analysis and geographical distribution. The tree was rooted between genotypes 2 and 3, and a single member of each genotype present per country was included. For each replicate, tree leaves were attributed to 1 of 11 broad geographical locations and a maximum likelihood inference of ancestral (geographical) states is shown. Branch lengths are arbitrary, and nodes with bootstrap support values above 60 are indicated together with the distribution of ancestral states at each internal node. Vertical bars indicate major geographical locations for each cluster.

Figure 5 Dissemination of leprosy throughout the world. Pillars are located on the country of origin of the *M. leprae* sample and color coded according to the scheme for the 16 SNP subtypes shown in **Figure 3** (note that this differs from the color scheme in **Fig. 4**). The thickness of the pillar corresponds to the number of samples (1–5, thin; 6–29, intermediate; >30, broad). The gray arrows indicate the migration routes of humans, with the estimated time of migration in years shown^{25,46}. The red dots indicate the location of the Silk Road in the first century, and * denotes results obtained from a-DNA.

outgroup. Inspection of the positions corresponding to the *M. leprae* SNPs in other genome sequences allowed the likely ancestral base to be deduced (**Fig. 3b**, **Supplementary Table 2**). This was successful for 13 of the 15 groups of markers and enabled us to establish a consensus ancestral sequence, which was then used in either neighbor-joining or maximum likelihood analysis to produce a phylogenetic tree (**Fig. 3c**). The topology of the trees was fully consistent with that of the scheme and phylogenetic tree for the four fully sequenced *M. leprae* strains (**Figs. 2c** and **3a**), thereby confirming its robustness.

An additional phylogenetic analysis was performed using maximum likelihood analysis^{26,27} to probe the relationship between the genotype of the *M. leprae* isolates and the affected individual's country of origin. For completeness, an example of each genotype found within a country was included, resulting in a sample size of 61. Overall, there was a consistent trend in the relationship between genotype and the geographic region (**Fig. 4**), and this trend gives an indication of the likely direction of gene flow and dissemination of the leprosy bacillus.

The global distribution of the 16 SNP subtypes was then plotted to show the frequency of each genotype per country. Again, with

the exception of islands³, this plot showed a reasonably tight correlation between the country of origin of an individual, the *M. leprae* genotype of the strain infecting the individual, and the known pattern of human migrations (**Fig. 5**). It is of interest that the *M. leprae* strains from both Turkey and Iran fall into the same two groups, F and K. This is consistent with F being a precursor to strains that migrated eastward with human populations, later giving rise to SNP type A found in India and Southeast Asia, whereas group K may have been an ancestor for *M. leprae* associated with westward migrations of humans that led to genotype M, associated with Europe and the Americas, and genotype P, characteristic of South America. Findings with Chinese isolates of *M. leprae*⁹ are also important, as, unlike many samples from Southeast Asia, these are not genotype 1 but genotype 3, subtype K.

Phylogeography of ancient *M. leprae*

Studying ancient DNA (a-DNA) is a valuable yet challenging approach, as this not only enables us to obtain samples from countries where leprosy is extinct but also provides information about different time periods and past epidemics. Skeletal remains were obtained from leprosy graveyards in Croatia, Denmark, Egypt, England, Hungary and Turkey, and all showed clear osteological evidence of lepromatous leprosy (**Table 1**). These remains were first shown to be positive for *M. leprae* DNA by single-round RLEP PCR²⁸ and then were used to generate PCR products for sequence analysis of all three SNP typing loci and, when possible, the appropriate subtypes. In all 13 cases, the a-DNA samples were found to be of SNP type 3, and 7 of these were successfully subtyped. The a-DNA analysis revealed that *M. leprae* from the UK belonged to SNP subtype 3I, whereas samples from Hungary were of SNP subtypes 3K or 3M (**Table 1**). Both extinct and extant samples of *M. leprae* from Turkey belonged to type 3K, whereas the 1,500-year-old Egyptian sample was found to be SNP type 3. All of the archaeological cases exhibited different VNTR profiles, which assisted in authentication of the a-DNA analysis (data not shown).

Table 1 Ancient *M. leprae* DNA, details of cases studied and bone samples taken from archaeological sites

Country (site)	Burial no. or marking	Age at death	Sex	Samples	Period (century AD)	SNP type	SNP subtype	Ref.
Croatia	2A	50–60	M	Both rhino-	8th–9th	3	— ^a	— ^b
	3A	20–25	F	Maxillary		3	— ^a	
Denmark	G483	25–30	F	Palatine	11th–15th	3	I/J	— ^b
Egypt (Dakhleh Oasis)	K2-B116	~23	M		4th–5th	3	K/L/M	38
England (Blackfriars, Ipswich)	1914	35–40	M	Maxillary palatine process	13th–16th	3	I	28
England	11287	Child	M	Tibia	9th–11th	3	— ^a	— ^b
	11503	30–40	M	Foot bones	9th–11th	3	— ^a	
	11784	30–40	M	Palatine	9th–11th	3	— ^a	
Hungary	503	30–35	F	Nasal region	11th	3	M	47
(Püspökladány)	222	45–50	M	Nasal region	10th	3	K	
Hungary	KD271	50–60	M	Skull and lower limbs	7th	3	K	48
(Kiskundorozsma)								
Turkey (Kovuklukaya)	KK 20/1	Mature	F	Nasal region	8th–9th	3	K	49

^aNot yet established. ^bC. Watson, D. Lockwood, personal communication.

DISCUSSION

Here, we describe the complete genome sequence and comparative analysis of Br4923, a Brazilian strain of *M. leprae*, and its use for the discovery of SNPs and other polymorphic markers with phylogeographic potential. This finding was then complemented by genome resequencing of strains from Thailand and the United States. When the four genome sequences were compared, remarkably little genomic diversity was uncovered, consistent with the hypothesis that leprosy has arisen from infection with a single clone that has passed through a recent evolutionary bottleneck^{2,3}. The four strains, which came from widely separated countries, have genomes that are 99.995% identical. In terms of the diagnosis, treatment and prevention of leprosy, this is extremely encouraging, as it means that antigenic drift in *M. leprae* should be negligible and the sequences of drug targets will not vary. Indeed, only 49 of the estimated 1,614 proteins in *M. leprae* show any amino acid change (Supplementary Table 1).

In a recent comparative study with *M. tuberculosis* and *Streptomyces coelicolor*, it was estimated that most of the mutations (56%) associated with pseudogenes or noncoding regions in *M. leprae* could be attributed to transitions, mainly of the C→T or G→A types²⁹. When the noncoding or pseudogene-containing regions were compared in the two completely sequenced *M. leprae* genomes, the frequency of these transitional mutations (59%) was found to be very close to that value, whereas in the coding sequences, the frequency was 63.5%. As outlined above, the same mutations are also dominant in the repetitive elements, where they account for 97% of the changes. However, as these elements appear to undergo recombination (Fig. 1), this particular value may be misleading. Nonetheless, a decrease in the G+C content of a genome is thought to be a hallmark of reductive evolution^{30,31}, and at 57.8% *M. leprae* clearly conforms to this rule, as the G+C content of most sequenced mycobacterial genomes is ~66%.

Five new pseudogenes were uncovered by this work (Fig. 2a), and their orthologs in *M. tuberculosis* are all known or predicted to be nonessential³². The most noteworthy of these pseudogenes was found in Br4923, where the counterpart of *ML0825c* in the TN strain has acquired a frameshift mutation as the result of a thymidine insertion in codon 16 (TN position 978629). *ML0825c* encodes a transcriptional regulator belonging to the ArsR family that is predicted to repress target gene expression in the absence of heavy metal cations. In *M. tuberculosis*, the *ML0825c* ortholog *rv2358* is the proximal gene in a bicistronic operon with *zur* (*furB*), whose expression is repressed by *rv2358* in a zinc-dependent manner^{33,34}. In an *M. tuberculosis zur* mutant, expression of 32 genes was found to be upregulated³⁵. It is not known whether loss of *ML0825c* has any phenotypic consequences, but it is conceivable that, as a result of polarity from the frameshift mutation, *Zur* may not be made, thus leading to derepression of the *Zur* regulon. This mutation is found only in *M. leprae* strains belonging to SNP subtypes 4O and 4P. The present study has also uncovered another useful marker for SNP subtype 4P, Del-1476519, a single-base deletion restricted to *M. leprae* strains from South America or the Caribbean. We can conclude that loss of the corresponding thymidine must have occurred within the last 500 years, since the introduction to Brazil of the ancestral strain from West Africa during the slave trade.

Through genome comparisons, some authors have attempted to reconstruct the evolution of *M. leprae* and to predict whether pseudogene formation was a gradual or stochastic event^{36,37}. Using a new approach based on the number of nonsynonymous substitutions per site in pseudogenes, one group estimated that *M. leprae* and *M. tuberculosis* diverged 66 million years ago and that a single pseudogenization event must have occurred in the leprosy bacillus

in the last 10–20 million years³⁶. Although we cannot predict the precise date of pseudogene formation here, our data indicate that this process still occurs.

The main goal of this study was to discover new polymorphic markers in order to improve the resolution of epidemiological and phylogeographic studies of *M. leprae*. This was successfully achieved, and interrogation of the 84 informative sites has led to the 4 SNP types³ being resolved into 16 subtypes. Together with an increase in the number of countries surveyed, this study has resulted in a much deeper level of understanding of the global spread of leprosy, with confirmation of the relationship between SNP type and geographical origin of the strains.

Paleomicrobiology was particularly helpful in this respect, and the analysis of ancient *M. leprae* DNA, present in skeletal remains from countries where leprosy has been eradicated, not only enabled us to expand our geographical coverage but also provided insight into the genotypes of strains circulating in Europe, Turkey and Egypt as long as 1,500 years ago. From the initial scheme for the evolution of the different *M. leprae* genotypes, it was predicted that European isolates should belong to SNP type 3, and this was indeed found to be the case (Table 1). Thanks to the new tools developed here, it was also possible to subtype the isolates in some cases, despite the limiting amounts and extensive fragmentation of the a-DNA. Examination of B116 (Table 1), the Egyptian skeleton from the Kellis-2 site in the Dakhleh Oasis³⁸, deserves some comment, as this specimen comes from a region close to the proposed origin of both *M. leprae* and *Homo sapiens*. Based on genotype data from East African strains of *M. leprae*, one might expect to find SNP type 2 in Egypt, whereas in reality we found type 3. B116, who has a calibrated radiocarbon date of 445 ± 50 years AD, in the Roman period, was buried in close proximity to another individual with leprosy, termed B6. Dietary studies using ¹⁵N stable isotopic analysis showed that both burials were outliers compared to other inhumations at Kellis 2 and may therefore not have been of Egyptian origin. Alternatively, from the phylogeny shown in Figure 3c, it is possible that when the ancestor of *M. leprae* separated from the other mycobacteria, its genotype was between SNP types 2 and 3.

The present phylogeographic scheme, based on results obtained with ~400 samples of *M. leprae* from 28 different locations, extends considerably the hypothesis proposed earlier in which the progenitor strain may have originated in East Africa and was SNP type 2 (ref. 3). This then gave rise to SNP type 1, which spread eastward with humans into Asia and SNP type 3, which disseminated westward into the Middle East and Europe before spawning the type 4 strains that are found in West Africa and countries linked to West Africa by the slave trade. Two new conclusions can be drawn from interpretation of the refined scheme (Figs. 3a, 4 and 5).

First, leprosy appears to have been introduced into Asia by two different routes: a southern route, associated with the SNP type 1 strains encountered in the Indian subcontinent, Indonesia and the Philippines, and a more northerly route starting in the Eastern Mediterranean region and extending via Turkey and Iran to China and from there to Korea and Japan. Here, strains with the 3K SNP subtype are preponderant, and the trade route between Europe and Asia known as the Silk Road appears likely to have been a means of transport and disease transmission (Fig. 5). The Black Death, caused by *Yersinia pestis*³⁹, is thought to have reached Europe in the fourteenth century from China via the Silk Road, carried by humans and their fleas. For leprosy (Fig. 3a) the opposite route of transmission may have operated, with the disease originating in Europe or the Middle East and then spreading to the Far East. Leprosy was thought

to have reached China from India, in about 500 BC, and then to have spread from China to Japan⁴⁰. This proposition is incompatible with the data presented here.

Second, it seems unlikely that leprosy was introduced into the Americas by early humans via the Bering straits; rather, it appears more probable that it was brought by immigrants from Europe, as most of the *M. leprae* strains found in North, Central and South America have the 3I genotype found in European leprosy cases. This interpretation is consistent with paleological findings because skeletons with signs of leprosy are limited to the postcolonial period⁴¹.

Finally, it is worth discussing the enormous discrepancy between the period at which pseudogene formation is thought to have arisen and the origin of early humans. It has been estimated recently that the bulk of the pseudogenes in *M. leprae* arose no earlier than 9 million years ago³⁶. Pseudogene formation is an indicator of radical change in the lifestyle of the host bacterium, such as from the free-living to pathogenic state or of adaptation to life within a particular tissue or cell type^{30,31}. In the case of *M. leprae*, obligate parasitism of humans or another primate species would represent such a change. Although modern humans represented by *H. sapiens* have existed only since approximately 250,000 years ago and left Africa within the last 100,000 years to settle other regions, earlier hominids are thought to have diverged from chimpanzees over 5 million years ago⁴². Reconciliation of the estimated time of pseudogene formation with human evolution could be achieved if an ancestor of *M. leprae* infected an early primate and then underwent genome decay and was subsequently transmitted vertically—although this seems unlikely, given that more genetic diversity among *M. leprae* isolates would be expected if this were true. Alternatively, the genome decay could well be ancient, but *M. leprae* may only recently have become a human pathogen. For instance, it is conceivable that an ancestral form of *M. leprae* infected an invertebrate host such as an insect, which later acted as a vector for transmitting the bacillus to humans. Support for the latter scenario is provided by studies of the related pathogen *Mycobacterium ulcerans*, which is at an early stage of reductive evolution⁴³ and appears to be transmitted to humans by water bugs and/or mosquitoes^{44,45}. Further insight into the timing of pseudogene formation in *M. leprae* will be provided by microbiology and paleomicrobiology and by deeper genome sequence analysis.

METHODS

Methods and any associated references are available in the online version of the paper at <http://www.nature.com/naturegenetics/>.

Accession codes. Br4923 genome sequence, GenBank FM211192; reads from Thai53 and NHDP63 resequencing analysis, SRP001064 (trace depository).

Note: Supplementary information is available on the Nature Genetics website.

ACKNOWLEDGMENTS

We thank the study participants and staff from the various leprosy clinics for their invaluable participation and support; J. Molto, A. Marcsik, Y. Selim Erdal, E. Popescu, M. Slaus and J. Boldsen for their encouragement and help with a-DNA samples; and L. Frangeul, A. Kapopoulou and S. Uplekar for assistance with bioinformatics. This work received the financial support of the Fondation Raoul Follereau, the Génopole programme, and the US National Institutes of Health, National Institute of Allergy and Infectious Diseases (grant RO1-AI47197-01A1 and contract NO1-AI25469).

AUTHOR CONTRIBUTIONS

M.M., N.H., P.J.B. and S.T.C. designed the study; A.P.-M., M. Matsuoka, A.K., Y.D., S.J., T.H.R., L.V.-C., M.M.S., S.B., M. Macdonald, B.R.S. and J.S.S. contributed sources of *M. leprae* DNA and demographic information; N.H., M. Monot, N.Z.,

D.S., T.G., J.T., K.H., A.G., J.R., P.S. and P.B. performed DNA sequencing and bioinformatics; G.M.T., H.D.D., A.B., S.M., C.W. and D.L. provided, sequenced and analyzed a-DNA and contributed archaeological insight; and S.T.C. wrote the manuscript with contributions from M. Monot, N.H. and G.M.T. plus comments from all authors.

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ONLINE METHODS

Complete sequence of Br4923 and informatics. Br4923 was originally isolated in 1996 from the skin biopsy of a Brazilian patient, inoculated into nude mice, and its whole genome sequence obtained from a pcDNA2.1 shotgun library. Briefly, 5 µg of purified DNA was sheared by nebulization and cloned in the pcDNA2.1 vector using adapters as described previously⁵⁰. The sequencing template was obtained with a Templiphi kit (GE Healthcare), subjected to BigDye Terminator version 3.1 cycle sequencing method and run in an ABI3730 DNA sequencer (Applied Biosystems). The sequence was assembled from 21,300 reads, analyzed and annotated using the Staden package (Trev, Gap4), Blast, Act and Artemis^{51–55}, as described previously^{2,56}.

Genome resequencing of Thai53 and NHDP63 and informatics. The strains Thai53 and NHDP63 were originally from an individual from Thailand and a native-born American from Louisiana, United States, respectively. Genomic DNA fragment sequencing libraries were prepared using the DNA Sample Prep Kit (Illumina) according to the protocol supplied with the reagents and using 5 µg of genomic DNA. DNA fragment libraries were loaded into one (Thai53) or two lanes (NHDP63) of a flow cell and sequenced on the Genome Analyzer II (Illumina) using the 36 Cycle Sequencing Kit version 1. Data were processed using the Illumina Pipeline Software package version 1.0 and reads either mapped onto the TN consensus sequence using MAQ⁵⁷ or assembled using Edena⁵⁸ and aligned to the reference genome using blastn⁵⁵. Data were managed using GAP4 and variations reported by direct comparisons.

Phylogeny. Multiple alignments were generated using ClustalW⁵⁹ and phylogenetic trees calculated by the neighbor-joining or maximum likelihood methods using web-based software (see URL). To calculate dN/dS ratios, we used the START 2 package⁶⁰ with the Nei-Gojobori method and the Jukes-Cantor correction⁶¹. For the phylogeographic analysis, we selected one SNP per linkage disequilibrium block ($n = 25$) for each combination of geographical location and genotype ($n = 61$), excluding samples from islands. These sequences were input to the RaxML software^{26,27}, which produced a maximum likelihood tree and 200 bootstrap replicates. Trees were rooted between genotypes 2 and 3. For each replicate, the tree leaves were attributed 1 of 11 broad geographical locations and a maximum likelihood inference of ancestral (geographical) states was performed using the ape package under R⁶². The inferred transition rates were averaged over all replicates and applied to the maximum likelihood tree (Fig. 4).

Sample preparation from extant *M. leprae*. Details of the specimens examined and their origin may be found in **Supplementary Table 3**. In this study, we performed PCR amplification with >600 samples of *M. leprae* comprising purified DNA, fresh skin biopsies, paraffin-embedded biopsy samples and slit-skin smears from microscope slides. *M. leprae* cells and/or DNA were prepared differently according to the sample source. Fresh skin biopsies were treated as described previously and DNA released by 'freeze boiling'⁶³. Paraffin-embedded specimens were first heated at 60 °C for 3 h, then the paraffin was removed by two extractions with 1 ml of xylene for 15 min, which was followed by one extraction with 100% ethanol for 15 min. Tissue was progressively rehydrated by soaking in 30% ethanol and, finally, in water. Then biopsies were minced with fine scissors followed by shearing in a Qiagen lyser with 3 mm glass beads (10 min, maximum power). The supernatant was transferred to an Eppendorf tube and 'freeze boiled'. After removal of debris, the supernatant was used directly in PCR. For slit-skin samples we used the Qiaamp DNA micro kit (Qiagen, Inc.) to recover DNA from stained microscope slides.

Standard PCR, SNP analysis and sequence reactions. The seven well-characterized strains of *M. leprae* used as sources of DNA for initial work were TN, India 2 and Thai53 (all SNP type 1), Africa (SNP type 2), NHDP63 and NHDP98 (both SNP type 3) and Br4923 (SNP type 4)³. Details of the primers used for genotyping may be found in **Supplementary Table 4**. Reactions (20 µl) typically contained *M. leprae* DNA from different samples, 10 mM Tris-HCl (pH 9.0), 50 mM KCl, 1.5 mM MgCl₂, 1.25 U *Taq* DNA polymerase (Q-Biogene) and 200 nM of forward and reverse primers. PCR was carried out for 45 cycles consisting of denaturation at 94 °C for 1 min, annealing at 55 °C for 1 min and extension at 72 °C for 2 min, with a final extension at 72 °C for 10 min in

a thermocycler (PTC-100, MJ Research, Inc.). If amplification failed, the PCR reaction was repeated after addition of 1 µl of T4GP32 (Q-Biogene). After enzymatic treatment with Exonuclease I (USB, Corp.) and Shrimp Alkaline Phosphatase (USB, Corp.), PCR products were submitted to BigDye Terminator version 3.1 cycle sequencing and analyzed using an ABI3100 DNA sequencer (Applied Biosystems). Sequence data were analyzed as above^{51,54}.

Ancient DNA (a-DNA) studies. Details of samples taken for the bioarchaeological part of the study are shown in **Table 1**. Measures to prevent cross-over contamination²⁶ were followed from the time of sampling. The strategy followed included (i) the use of multiple extraction and template blanks, (ii) reproducibility, (iii) appropriate molecular behavior, (iv) confirmation of product identity with sequencing and (v) replication of key findings at a separate center. One of the samples, burial 1914 from Ipswich, was sampled twice to provide sufficient material for analysis at center 2, Manchester University. Gloves were worn and changed between handling different skeletal components. Samples of bone were removed from the skeleton using sterile disposable scalpel blades and transferred into sterile plastic containers for transport to the laboratories. The work surface was cleaned between samples using a proprietary multisurface cleaner containing bleach.

a-DNA extraction, PCR and sequencing. At center 1, University College London, samples of bone or scrapings were finely ground in autoclaved pestles and mortars and DNA from bone powder extracted using the NucliSens kit from bioMérieux, as previously described²⁸. DNA extracts were stored at –20 °C until assayed. At center 2, Manchester University, DNA from burial 1914 was extracted independently using a modification of the method of Yang⁶⁴, which has previously been shown to be the most efficient of five tested methods for extraction of a-DNA from bones⁶⁵.

The presence of residual *M. leprae* DNA was first confirmed using a sensitive PCR method, which amplifies the 37-copy repetitive element RLEP^{18,19,28}. SNP typing methods were developed to amplify fragmented DNA likely to persist in skeletal remains and applied to the extracts. The oligonucleotide primers used for this and the key PCR conditions are listed in **Supplementary Table 5**.

The Excite core kit (BioGene) was used for all PCR amplifications. This is a uracil-N-glycosylase-ready kit suitable for real-time and routine hot-start PCR applications. SYBR green was included in the PCR master mixes at a final dilution of 1/55,000, and reactions were performed and monitored on the Corbett RotorGene 3000 real-time PCR platform (Corbett Research) in a final volume of 25 µl. Forty-five cycles of amplification were performed for all methods. Melt analysis was performed using the RotorGene software and, additionally, all products were run on 3% agarose gels. Template blanks containing water in place of bone extract were alternated with samples in the RotorGene chamber and monitored to ensure absence of contamination. Positive control samples were not amplified during the course of the bioarchaeological study.

PCR products were separated on 3% (wt/vol) low-melting-point agarose (Invitrogen), and bands were excised with a sterile scalpel blade and purified using a GeneClean III DNA isolation kit (Fisher Life Sciences), then sequenced using BigDye Terminators, as outlined above.

URLs. Mobyle@Pasteur, <http://mobyle.pasteur.fr>; NCBI short read archive, http://www.ncbi.nlm.nih.gov/Traces/sra_sub/sub.cgi.

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Corrigendum: Comparative genomic and phylogeographic analysis of *Mycobacterium leprae*

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Nat. Genet. 41, 1282–1289 (2009); published online 1 November 2009; corrected after print 5 February 2010.

In the version of this article initially published 1 November 2009, Ali Khamesipour was misspelled in the author list. The error has been corrected in the HTML and PDF versions of the article.