Genome-wide estimation of recombination, mutation and positive selection enlightens diversification drivers of *Mycobacterium bovis*

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Genome sequencing has reinvigorated the infectious disease research field, shedding light on disease epidemiology, pathogenesis, host–pathogen interactions and also evolutionary processes exerted upon pathogens. *Mycobacterium tuberculosis* complex (MTBC), enclosing *M. bovis* as one of its animal-adapted members causing tuberculosis (TB) in terrestrial mammals, is a paradigmatic model of bacterial evolution. As other MTBC members, *M. bovis* is postulated as a strictly clonal, slowly evolving pathogen, with apparently no signs of recombination or horizontal gene transfer. In this work, we applied comparative genomics to a whole genome sequence (WGS) dataset composed by 70 *M. bovis* from different lineages (European and African) to gain insights into the evolutionary forces that shape genetic diversification in *M. bovis*. Three distinct approaches were used to estimate signs of recombination. Globally, a small number of recombinant events was identified and confirmed by two independent methods with solid support. Still, recombination reveals a weaker effect on *M. bovis* diversity compared with mutation (overall r/m = 0.037). The differential r/m average values obtained across the clonal complexes of *M. bovis* in our dataset are consistent with the general notion that the extent of recombination may vary widely among lineages assigned to the same taxonomical species. Based on this work, recombination in *M. bovis* cannot be excluded and should thus be a topic of further effort in future comparative genomics studies for which WGS of large datasets from different epidemiological scenarios across the world is crucial. A smaller *M. bovis* dataset (*n* = 42) from a multi-host TB endemic scenario was then subjected to additional analyses, with the identification of more than 1,800 sites wherein at least one strain showed a single nucleotide polymorphism (SNP). The majority (87.1%) was located in coding regions, with the global ratio of non-synonymous upon synonymous alterations (dN/dS) exceeding 1.5, suggesting that positive selection is an important evolutionary force exerted upon *M. bovis*. A higher percentage of SNPs was detected in genes enriched into “lipid metabolism”, “cell wall and cell processes” and “intermediary metabolism and respiration” functional categories, revealing their underlying importance in *M. bovis* biology and evolution. A closer look on genes prone to horizontal gene transfer in the MTBC ancestor and included in the 3R (DNA repair, replication and recombination) system revealed a global average negative value for Tajima’s D neutrality test, suggesting that past selective sweeps and population expansion after a recent bottleneck remain as major evolutionary drivers of the obligatory pathogen *M. bovis* in its struggle with the host.

The *Mycobacterium tuberculosis* complex (MTBC) is one of the most successful taxon of bacterial pathogens and a paradigmatic case in bacterial evolution, revealing a strikingly high nucleotide identity at the genome level (> 99%) among its members1,2. The different MTBC ecotypes cause tuberculosis (TB), an infectious granulomatous disease, in a broad group of host species, ranging from micro-mammals to humans3–1. Currently, the...
complex encompasses human [M. tuberculosis (Mtb), M. africanum] and animal-adapted pathogens (M. bovis, M. caprae, M. pinnipedii, M. microti, M. mungi, M. orygis, M. suricattae, “chimpanzee bacillus” and “dassie bacillus”)5,6. M. canettii (also known as “smooth tubercle bacilli”) has an average nucleotide identity of 98% with the aforementioned mycobacteria and comparative genomic works suggest that M. canettii and the rest of MTBC have diverged very recently from a common ancestor7. Considering this notion, several authors refer to M. canettii as an MTBC member8.

The MTBC has been systematically described as a strictly clonal complex, with population structure being apparently dominated by reductions in diversity, bottlenecks, selective sweeps and genetic drift9,10. Assuming the strictly clonal evolution of the complex, polymorphisms such as deletions cannot be restored by recombination11. Based on this premise, the successive events of genomic deletions of the regions of difference (RD) and TbD1 (Mtbc specific deletion 1 region) have been proposed as molecular markers of MTBC evolution12,13. Comparative genomics and whole genome sequencing (WGS) works support the division of human-adapted members into nine lineages (M. tuberculosis L1 to L4, L7 and L8; and M. africanum L5, L6 and L9), with lineages L2 to L4 sharing the deletion of TbD1 region12,13. Moreover, animal-adapted members have been proposed to share a common ancestor and are defined by clade-specific deletions in the RD7, RD8, RD9 and RD1014–16.

Events of horizontal gene transfer (HGT) and recombination are assumed to be rare and to have occurred in the ancestors of MTBC, rather than throughout the diverging history of MTBC members17–19. Two early reports by Hughes and collaborators (2002) and Gutacker and collaborators (2006) suggested that recombination events might have helped to shape the polymorphisms marking specific loci of M. tuberculosis strains9,10. The apparent absence of recombination in MTBC has been attributed to: (1) loss of mechanistic processes and ability for HGT; (2) rareness of HGT events; and (3) no opportunity for recombination events within MTBC ecological niches17,18. More recently, a few Whole Genome Sequencing (WGS) studies applied to MTBC strains20 and M. bovis21 provided evidences of recombination, with the first suggesting that strains frequently exchange small DNA fragments, but because of the limited nucleotide sequence variation, these events remain unnoticed.

Mycobacterium bovis is the MTBC member most frequently recovered from livestock, mainly cattle, although it can also be isolated from free-ranging and fenced wildlife22–24. M. bovis evolved to five main clonal complexes [European 1 (Eu1), European 2 (Eu2), European 3 (Eu3), African 1 (Af1) and African 2 (Af2)], defined based on spoligotyping profile, specific deletions and single nucleotide polymorphisms (SNPs) in specific genes25–29. These clonal complexes evidence the diversity structure of M. bovis population and association with geographic regions. Furthermore, a recent WGS work by Zimpel and collaborators (2020) devised an M. bovis SNP-based phylogeny with over 1900 genomes, which suggested the existence of at least four distinct lineages in the world (named Lb1 to Lb4), that are not entirely concordant with the previous defined clonal complexes, although geographic specificities may also be confirmed30. These authors performed phylogenetic and molecular dating divergence analyses but did not investigate recombination30.

Previous works employing different molecular techniques such as spoligotyping, MIRU-VNTR (Mycobacterial Interspersed Repetitive Unit-Variable Number of Tandem Repeat) and, more recently, SNP typing, revealed a certain level of genetic diversity among M. bovis strains31–35. The differentiation of genetic variants has become a crucial tool to study disease epidemiology, contributing to gain insights into pathogenesis, virulence and disease transmission. The arrival of WGS methodologies opened the possibility to shed light into the evolutionary drivers exerted upon M. bovis genomes during adaptation and persistence to different hosts and epidemiological scenarios.

In this work, we take advantage of a comparative genomic analysis of a diverse M. bovis dataset (n = 70), including isolates from different clonal complexes to gain insights into the evolutionary processes of M. bovis, specifically addressing phylogenetic relationships and recombination events. Complementary to this analysis, the sub-dataset of M. bovis isolates (n = 42) obtained from a well characterized multi-host TB endemic region in Portugal31,36 was further explored to infer the balance between the relative rates of nonsynonymous (dN) to synonymous (dS) nucleotide substitution, and the evolutionary contribution of specific groups of genes referred to in the literature as having been acquired though HGT by the MTBC ancestor37–39, as well as genes encoding 3R (DNA repair, replication and recombination) system components39. The genes proposed to be acquired through HGT were selected since they may represent ancient polymorphisms, and so it is expected that they might contain a higher fraction of synonymous alterations. The genes included in the 3R system were selected since previous work performed with M. tuberculosis strains suggest a general negative/purifying selection acting upon these genes and that they might play an important role in evolution39. Another objective of the work was to infer the presence of recombination events. For this purpose, and considering that our dataset from Portugal only had genomes included in European clonal complex 2 and strains without a clonal complex assigned, we decided to include publicly available genomic data to end up with representatives from all clonal complexes and to increase robustness and breadth of results.

Methodology

Mycobacterium bovis isolates dataset. Forty-two newly sequenced M. bovis genomes from an endemic multi-host TB scenario in Portugal (details below), previously characterized from an epidemiological point of view30, were at the centre of this work. Considering that the dataset from Portugal only has representatives of European 2 clonal complex and strains without complex assigned, publicly available whole genome sequencing data was added in order to enlarge the dataset with representatives from all M. bovis clonal complexes. Therefore, three sources of whole genome sequencing data were used in this work: complete/draft genome assemblies up to a maximum of 10 scaffolds deposited at NCBI (National Center for Biotechnology Information) (n = 15 isolates); Illumina fastq files deposited at SRA (Sequence Read Archive) representative of M. bovis clonal complex diversity (n = 12 isolates)30; and 42 newly sequenced genomes from Portugal. Mycobacterium bovis BCG (bacil-
lus Calmette-Guérin) was excluded from the NCBI search. *M. bovis* AF2122/97 commonly used as reference genome was included in the dataset. Due to the public unavailability of whole genome sequences from representatives of African 1 clonal complex, and the low numbers of genomes from representative strains of AF2 and Eu1, raw sequencing data available at SRA was used in those cases. The work of Zimpel and collaborators (2020) helped in the identification of genomes from the aforementioned clonal complexes and in the selection process of *M. bovis* to include in the dataset. For Eu3, only one type genome is described (Branger et al., 2020), thus the genome that we included is the solo representative of the Eu3 complex.

Globally, the dataset included 70 *M. bovis* isolated from eight host species, distributed by 12 countries between 1985 and 2016. Thirty-six were assigned as Eu2, seven as Eu1, one as Eu3, three as Af1, four as Af2 and 19 were not attributed to any clonal complex (details below). Detailed information about the *M. bovis* used in this study (including accession numbers) can be found in Table 1 and Supplementary Table 1.

**Newly sequenced genomes (dataset from Portugal).** Forty-two newly sequenced *M. bovis* whole genomes originating from animal TB hotspots in Portugal and scattering a period of over 12 years were at the centre of this study, as the underlying wildlife-livestock disease system has been monitored regularly\(^1-3\) (Supplementary Fig. 1). These strains were isolated from cattle (*n* = 14), red deer (*n* = 16) and wild boar (*n* = 12) from 2003 to 2015, according to the ensuing procedure: animal tissue samples were pooled and processed following the protocol guidelines recommended in the OIE Manual for Terrestrial Animals and inoculated onto Stonebrink and Löwenstein-Jensen pyruvate solid media and liquid medium. Cultures were incubated at 37 °C and inspected weekly for growth for a minimum period of 12 weeks. Colonies were directly stored at glycerol solution at -80°C. The DNA for the WGS procedure was obtained after a single in vitro passage of original archived samples in mycobacteria selective medium (Middlebrook 7H9, BD Diagnostics). For that purpose, frozen culture stocks were re-cultured on Middlebrook 7H9 supplemented with 5% sodium pyruvate and 10% ADS enrichment (50 g albumin, 20 g glucose, 8.5 g sodium chloride in 1 L water) at 37 °C. After four weeks’ growth, the culture medium was renewed, and the cultures were monitored regularly until growth was observed. Cells were harvested by centrifugation, the pellet was resuspended in 500 μL phosphate buffer saline (PBS), heat-killed at 99 °C during 30 min, centrifuged, and the supernatant stored at -20 °C until WGS. All procedures were performed on a level 3 biosafety facility.

WGS paired-end genomic libraries were prepared with unique indexing of each DNA sample and sequenced using Illumina MiSeq (2 × 250 pb) (40 samples) and HiSeq (2 × 150 pb) (two isolates) technology (Eurofins Genomics, Germany). The genomic DNA was sequenced using the Illumina Genome Analyzer with the paired-end module attachment and libraries were constructed with Nextera XT DNA Library Prep Kit from Illumina, according to the manufacturer's specifications.

**Clonal complex assignment.** Considering the data recovered from SRA (*n* = 12), the clonal complex identification was available as metadata of the corresponding publications\(^5,6,10,41,43\). When considering complete genomes, with the exception of *M. bovis* AF2122/97 and *M. bovis* 3601 that are recognized members of Eu1 and Eu3 clonal complexes, respectively\(^25,29\), whole genome alignment with *M. tuberculosis* H37Rv (NCBI accession NC_000962.3) was performed using MAFFT (Multiple alignment program for amino acid or nucleotide sequences, version 7.458) with parameter--addfragments\(^49\). Then, the presence of the deletions and/or SNP characteristic of the different clonal complexes was searched.

The newly sequenced *M. bovis* (*n* = 42) and raw reads from draft assembly genomes (*n* = 3) were aligned with reference genome *M. tuberculosis* H37Rv via vSNP pipeline and the presence of the deletions and/or SNP characteristic of the different clonal complexes was searched.

Information from the presence/absence of characteristic deletions and/or SNP and spoligotyping profile were gathered to assign the genomic data to the corresponding clonal complex. For four draft assemblies it was not possible to infer the spoligotyping profile, so and they were included in the “without complex” group.

**Bioinformatics analysis.** The bioinformatics workflow followed in this work started from de novo assembly and map to reference strategies, with the purpose to explore recombination events and the polymorphisms of specific gene groups. Figure 1 provides a flowchart of the steps followed. For the recombination analysis, all the genomes were used to increment the robustness of inferences and the associated metrics.

**De novo genome assembly.** In order to mitigate errors in the generation of genome consensus sequences, we first obtained de novo assemblies and, then, the core multi-alignment. The Unicycler pipeline, currently available at https://github.com/rrwick/Unicycler\(^49\), was implemented to perform de novo assembly for 54 sequenced genomes (42 newly sequenced and 12 fastq files recovered from SRA). Briefly, before de novo assembly, reads quality analysis was performed in FastQC version 0.11.7 (https://github.com/s-andrews/FastQC), and whenever necessary cleaned with Trimmomatic version 0.36 (options “cut adapter and other illumina-specific sequences from the read” and “cut bases off the end of a read, if bellow a threshold quality of 20” were applied) (http://www.usadellab.org/cms/?page=trimmomatic)\(^50\). Then, SPAdes optimiser\(^49\) was used for genome assembly and Pilon version 1.18\(^18\) for post-assembly optimization. A conservative bridging mode was selected to avoid misassembly and the k-mer size was searched and selected between 20 and 95% of read length. Following SPAdes guidelines and considering reads’ size, contigs with less than 300 bp were removed and a 20 read depth coverage cut-off was established\(^52\). In the de novo assembly strategy, no genome regions, such as the highly repetitive Proline-Glutamate (PE) and Proline-Proline Glutamate (PPE) paralogous genes, were removed.
| M. bovis ID | Clonal complex | Country | Year | Host species | References | Type of sequence |
|------------|---------------|---------|------|--------------|------------|----------------|
| Mb0220     | w/o CC        | Portugal| 2003 | Cattle       |            | Newly sequenced |
| Mb0261     | Eu2           | Portugal| 2006 | Red deer     |            | Newly sequenced |
| Mb0601     | Eu2           | Portugal| 2007 | Cattle       |            | Newly sequenced |
| Mb0769     | Eu2           | Portugal| 2008 | Cattle       |            | Newly sequenced |
| Mb0783     | Eu2           | Portugal| 2008 | Wild boar    |            | Newly sequenced |
| Mb0865     | Eu2           | Portugal| 2008 | Cattle       |            | Newly sequenced |
| Mb0891     | Eu2           | Portugal| 2009 | Red deer     |            | Newly sequenced |
| Mb0895     | Eu2           | Portugal| 2008 | Wild boar    |            | Newly sequenced |
| Mb1317     | Eu2           | Portugal| 2010 | Cattle       |            | Newly sequenced |
| Mb1339     | Eu2           | Portugal| 2010 | Cattle       |            | Newly sequenced |
| Mb1458     | w/o CC        | Portugal| 2010 | Wild boar    |            | Newly sequenced |
| Mb1480     | w/o CC        | Portugal| 2010 | Cattle       |            | Newly sequenced |
| Mb1654     | Eu2           | Portugal| 2011 | Cattle       |            | Newly sequenced |
| Mb1670     | w/o CC        | Portugal| 2011 | Red deer     |            | Newly sequenced |
| Mb1711     | Eu2           | Portugal| 2011 | Red deer     |            | Newly sequenced |
| Mb1712     | Eu2           | Portugal| 2011 | Red deer     |            | Newly sequenced |
| Mb1714     | Eu2           | Portugal| 2011 | Cattle       |            | Newly sequenced |
| Mb1744     | w/o CC        | Portugal| 2012 | Wild boar    |            | Newly sequenced |
| Mb1746     | Eu2           | Portugal| 2012 | Red deer     |            | Newly sequenced |
| Mb1758     | Eu2           | Portugal| 2012 | Cattle       |            | Newly sequenced |
| Mb1769     | Eu2           | Portugal| 2012 | Wild boar    |            | Newly sequenced |
| Mb1785     | Eu2           | Portugal| 2012 | Red deer     |            | Newly sequenced |
| Mb1789     | Eu2           | Portugal| 2012 | Cattle       |            | Newly sequenced |
| Mb1841     | Eu2           | Portugal| 2012 | Cattle       |            | Newly sequenced |
| Mb1870     | Eu2           | Portugal| 2012 | Wild boar    |            | Newly sequenced |
| Mb1915     | Eu2           | Portugal| 2013 | Red deer     |            | Newly sequenced |
| Mb1948     | w/o CC        | Portugal| 2013 | Red deer     |            | Newly sequenced |
| Mb1960     | Eu2           | Portugal| 2013 | Red deer     |            | Newly sequenced |
| Mb2026     | Eu2           | Portugal| 2013 | Cattle       |            | Newly sequenced |
| Mb2043     | Eu2           | Portugal| 2013 | Red deer     |            | Newly sequenced |
| Mb2067     | Eu2           | Portugal| 2013 | Wild boar    |            | Newly sequenced |
| Mb2206     | Eu2           | Portugal| 2014 | Cattle       |            | Newly sequenced |
| Mb2235     | w/o CC        | Portugal| 2014 | Red deer     |            | Newly sequenced |
| Mb2277     | w/o CC        | Portugal| 2014 | Red deer     |            | Newly sequenced |
| Mb2300     | Eu2           | Portugal| 2014 | Wild boar    |            | Newly sequenced |
| Mb2310     | Eu2           | Portugal| 2015 | Red deer     |            | Newly sequenced |
| Mb2313     | Eu2           | Portugal| 2015 | Wild boar    |            | Newly sequenced |
| Mb2325     | Eu2           | Portugal| 2015 | Red deer     |            | Newly sequenced |
| Mb2328     | Eu2           | Portugal| 2015 | Red deer     |            | Newly sequenced |
| Mb2347     | w/o CC        | Portugal| 2015 | Wild boar    |            | Newly sequenced |
| Mb2395     | Eu2           | Portugal| 2015 | Wild boar    |            | Newly sequenced |
| Mb2397     | Eu2           | Portugal| 2015 | Wild boar    |            | Newly sequenced |
| Mb502499   | Af1           | Ghana   | NA   | Human        |            | SRA deposited  |
| Mb502526   | Af1           | Ghana   | NA   | Human        |            | SRA deposited  |
| Mb1203064  | Af1           | Ghana   | NA   | Human        |            | SRA deposited  |
| Mb1177155  | Af2           | France  | NA   | Wild boar    |            | SRA deposited  |
| Mb1791710  | Af2           | Tanzania| NA   | Chimpanzee   |            | SRA deposited  |
| Mb1791712  | Af2           | Tanzania| NA   | Chimpanzee   |            | SRA deposited  |
| Mb1792006  | Eu1           | USA     | 2006 | Cattle       |            | SRA deposited  |
| Mb1792127  | Eu1           | USA     | 2008 | Cattle       |            | SRA deposited  |
| Mb1792361  | Eu1           | USA     | 2013 | Cattle       |            | SRA deposited  |
| Mb7240242  | Eu1           | USA     | 2016 | Cattle       |            | SRA deposited  |
| Mb7240415  | Eu1           | USA     | 2014 | Cattle       |            | SRA deposited  |
| Mb1791984  | Eu1           | USA     | 2005 | Cattle       |            | SRA deposited  |
| MBE1       | w/o CC        | Egypt   | 2014 | Cattle       |            | assemble/draft genomes NCBI |
| MBE3       | w/o CC        | Egypt   | 2014 | Cattle       |            | assemble/draft genomes NCBI |

Continued
| M. bovis ID | Clonal complex(a) | Country | Year | Host species | References | Type of sequence |
|-------------|-------------------|---------|------|--------------|------------|-----------------|
| MBE4        | w/o CC            | Egypt   | 2014 | Cattle       | NA         | assemble/draft genomes NCBI |
| MBE10       | w/o CC            | Egypt   | 2015 | Cattle       | NA         | assemble/draft genomes NCBI |
| Mb0077      | w/o CC            | Canada  | 2006 | Elk          | NA         | assemble/draft genomes NCBI |
| Mb0565      | w/o CC            | Canada  | 2011 | Cattle       | NA         | assemble/draft genomes NCBI |
| BMR25       | w/o CC            | Canada  | 1985 | Buon         | NA         | assemble/draft genomes NCBI |
| Mb3601      | Eu3               | France  | 2014 | Cattle       | NA         | assemble/draft genomes NCBI |
| Mb0476      | Eu2               | Canada  | 2002 | Cattle       | NA         | assemble/draft genomes NCBI |
| MbSP38      | Eu2               | Brazil  | 2010 | Cattle       | NA         | assemble/draft genomes NCBI |
| Mb1595      | w/o CC            | Korea   | 2012 | Cattle       | NA         | assemble/draft genomes NCBI |
| Mb0030      | w/o CC            | China   | NA   | NA           | NA         | assemble/draft genomes NCBI |
| Mb0001      | Eu2               | Brazil  | 2015 | Tapirus terrestris | NA | assemble/draft genomes NCBI |
| Mb0003      | w/o CC            | India   | 1986 | Cattle       | NA         | assemble/draft genomes NCBI |
| Mb31150     | Af2               | Uganda  | NA   | Chimpanzee   | NA         | assemble/draft genomes NCBI |

Table 1. Characteristics of *Mycobacterium bovis* genomes used in this work. Eu1: European 1, Eu2: European 2, Eu3: European 3, Af1: African 1, Af2: African 2, and w/o CC: without clonal complex. NA: non-available information.

Figure 1. Bioinformatics workflow followed in this study.
The quality of de novo assemblies was assessed by QUAST pipeline (http://quast.sourceforge.net/quast.html), which promotes the remapping of contigs with *M. bovis* AF2122/97 reference genome (NCBI accession number LT708304.1) (quality parameters presented in Supplementary Table 1).

**Genome map to reference.** The FASTQ files from the newly sequenced *M. bovis* obtained from Illumina sequencing were aligned with *M. bovis* AF2122/97 reference genome (LT708304.1) with the help of vSNP pipeline (https://github.com/USDA-VS/vSNP). The standard filtering parameters or variant quality score recalibration were applied according to Genome Analysis Toolkit (GATK)’s Best Practices recommendations[33-35]. Results were filtered using a minimum SAMtools quality score of 150 and AC = 2. Reads were also examined using Kraken (http://ccb.jhu.edu/software/kraken/) to exclude contamination. The vSNP pipeline used for the map to sequence strategy in our work examines a series of defining SNPs and targets also to exclude mixed infection scenarios. Genome coverage by reads was superior to 99% (Supplementary Table 1).

To avoid mapping errors and false SNPs, a variant was filtered out if: (1) it was supported by less than 20 reads, (2) it was found in a frequency of less than 0.9, (3) it was registered in at least one strain but also with a gap in at least another strain. SNPs and positions with mapping issues or alignment problems were visually validated with Integrated Genomics Viewer (IGV) version 2.4.19 (http://software.broadinstitute.org/software/igv)[36]. Since Proline-Glutamate (PE) and Proline-Proline Glutamate (PPE) genes are highly repetitive and part of multi-gene families, they are prone to misreading by Illumina sequencing and mis-mapping and so are preferentially removed from the bioinformatics workflow of *Mycobacterium tuberculosis* complex members when a strategy of map to sequence is used to confirm SNPs. We thus filtered PE/PPE genes out from the analysis, as well as indels. All SNPs were grouped into functional categories according to Bovilist (http://genolist.pasteur.fr/Bovilist/).

The SnpEff pipeline (https://pcingola.github.io/SnpEff/) was employed to infer SNP consequences (synonymous or non-synonymous alterations). A new database for *M. bovis* AF2122/97 genome (LT708304.1) was created.

**Global core genome multi-alignment.** The core genome multi-alignment was performed with Parsnp v1.2, currently available at https://github.com/marbl/parsnp[37], using the 69 complete genomes/draft assemblies (with option -c) and *M. bovis* AF2122/97 (LT708304.1) as reference. Four core multi-alignment were performed: including only members of Eu2 clonal complex (n = 37), including all members of European clonal complexes (n = 44), including a junction of European and African clonal complexes (n = 51), and including all *M. bovis* from this study (n = 70).

The core alignments generated by Parsnp were used to infer maximum-likelihood (ML) phylogenetic trees using RAxML, via CIPRES Science Gateway v3.3 (http://www.phylo.org/)[38], with 1000 bootstrap replications.

**Estimation of recombination events.** The presence of recombination events was examined using three different algorithms and bioinformatics tools in parallel: SplitsTree4 software, Gubbins (Genealogies Unbiased By recomBinations In Nucleotide Sequences) pipeline and RDP4 (Recombination Detection Program, version beta 4.101) software.

The split decomposition method implemented in SplitsTree4 v4.15.1 (http://www.splitstree.org/) was implemented to compute unrooted phylogenetic networks, which were validated statistically using the Phi test, with a significance threshold of p = 0.05. The core multi-alignments from Parsnp analysis were used as input and the split decomposition as network criteria was implemented.

Gubbins pipeline v2.3.1 (https://github.com/sanger-pathogens/gubbins[39]) was run using default parameters, as another way to assess the impact of recombination on *M. bovis*. The algorithm implemented in the pipeline reconstructs the clonal genealogy relating the complete genomes/draft assemblies of our dataset and the reference genome (*M. bovis* AF2122/97, LT708304.1) to each other; and scans the positions of SNPs across each branch of the tree in order to detect clusters of SNPs that would indicate recombination events. The null hypothesis for branch assumes the absence of any recombination events, therefore implying that the SNPs occurring on the branch should be evenly distributed. The core multi-alignments from Parsnp and the best scoring ML tree from RAxML were used as input files.

Finally, to confirm the recombination events suggested by the Gubbins pipeline, six algorithms (RDP[41], GENECONV[42], Bootscan[43], Maxchi[44], Chimaera[45], and SIScan[46]) implemented in RDP4[47] were applied to the core multi-alignments from Parsnp under default settings. We established that at least three of the algorithms implemented in RDP4 had to concordantly evidence a significant signal to validate each recombination event.

Considering that both Gubbins and RDP4 software seek recombination signals by inspecting the complete multi-alignment in windows of 500 bp maximum, and to confirm that the inclusion of PE/PPE genes in the de novo assembly process did not interfere with the recombination signals found, the neighbourhood of genes in which recombination events were identified were further inspected through a synteny analysis. Synteny maps, using complete genomes, were constructed with MAUVE—multi-genome alignment (http://darlinglab.org/mauve/mauve.html) to exclude local genome translocations or inversions. Furthermore, a synteny analysis with aminoacidic sequences was performed via SyntTax webserver (https://archaea.i2bc.paris-saclay.fr/SyntTax/) using complete genomes.

**Gene diversity analyses.** The genome dataset obtained from a multi-host TB system in Portugal was subjected to deeper analyses with the objective to examine the polymorphisms in the genes referred in the literature as having been acquired through HGT by the MTBC ancestor[37,38] and in the genes encoding 3R (DNA repair, replication and recombination) system components[39]. Gene sequences of the 42 *M. bovis*, together with gene sequence from the reference genome (*M. bovis* AF2122/97, NC_002945.4), were aligned using ClustalX v2.1
Results and discussion

Global phylogenetic analysis. A Maximum Likelihood (ML) phylogenetic tree based on the 69 M. bovis isolates and reference genome was obtained (Fig. 2A). This strategy allows the generation of a more robust tree, when comparing with single gene based trees or multi-locus based trees, that do not capture the variability across the entire genome and consequently present low inter-specific discriminatory power\(^\text{68,69}\). The resulting topology of the ML tree generally agrees with clonal complex classification, with genomes of Eu2 clustering in one tree branch and genomes of Af1 also clustering together (Fig. 2A). Results are also in agreement with the known M. bovis evolutionary relationships that present a large division between Eu1 members and a group composed by all the other clonal complexes and genomes without assigned clonal complex\(^\text{30}\). Small inconsistencies between clonal complex and the relationships observed at the phylogenetic tree can be explained by the fact that clonal complexes are described based on specific genomic regions, while the phylogenetic tree is based on core genome multi-alignment representing the whole genomes.

Evidences of recombination in Mycobacterium bovis. Mycobacterium tuberculosis complex is described to have clonally evolved, and most evidences accumulated over the years support the idea that ongoing HGT and recombination events do not occur at detectable levels in the MTBC\(^\text{15,17,18}\). Previous works have suggested that there might be limited recombination among MTBC strains\(^\text{20,21}\), while others were not successful to identify measurable recombination events\(^\text{70,71}\). To revisit this issue with focus on M. bovis, and unlike previous works that only accounted for M. tuberculosis\(^\text{70,71}\); or that accounted MTBC as a whole, with few M. bovis representatives\(^\text{20}\); or that only considered a restrict M. bovis dataset\(^\text{21}\), in this work a total of 70 strains, with representatives from all clonal complexes, was used to screen for recombination. The dataset was scaled in four cumulative levels: (1) Eu2 members, (2) all European clonal complexes members (i.e. European), (3) both European and African clonal complexes (Eu + Af) and (4) the entire dataset (encompassing the genomes that are not included in any of the clonal complexes already described).

To investigate this postulate further, a split decomposition network was performed to assess for the absence of recombination events between genomes, since this method enables the visualization of ancestral relationships between individuals and displays conflicting phylogenetic signals. The presence of cycles in the network (i.e. regions that do not converge into a single tree), was confirmed in all four datasets under analysis, however none was supported statistically by the Phi test (Eu2, \(p = 0.0956\); European, \(p = 0.1637\); Eu + Af, \(p = 0.2774\); entire dataset \(p = 0.2451\)), providing poor evidence for the presence of recombination events (Fig. 3A-D).

Following this analysis, and considering the observation of cycles in all networks, the reconstruction algorithm implemented in Gubbins pipeline was applied in order to reconstruct the clonal genealogy and to perform a complementary estimation of the impact of recombination in M. bovis genomes. A cumulative number of recombination events was inferred with the majority occurring in terminal branches (i.e. occurring in a single genome) (Table 2). The metrics showed consistency across the datasets and revealed that recombination events occurred two hundred to three hundred times less frequently than mutations, once the rho/theta parameter...
that represents the relative rates of recombination and point mutation on a branch presented an average value between 0.0037 and 0.0056 (Table 3). Recently, a published work with 38 *M. bovis* strains evidenced a higher rho/theta value (rho/theta = 0.1) than the one obtained for this dataset, however the work by Patané and co-workers used reference-based assemblies to infer recombination parameters, a procedure detail that was already associated with enrichment of putative recombination events at terminal branches due to the assembly procedure.

Following, the r/m parameter, which represents the ratio of diversity introduced by recombination and mutation, revealed an average value between 0.025 and 0.037, pointing that recombination has a lower overall
Recombination event | Identification | Core-alignment positions | Genome positions(a) | Gene name | Mb gene name | Classification of gene function | M. bovis isolate ID
--- | --- | --- | --- | --- | --- | --- | ---
#1 | Gubbins | 945,923–945,950 | 1,220,297–1,220,324 | PE PGRS22 | Mb1121 | PE-PGRS family protein | Mb2026
#2 | Gubbins; RDP4 | 1,176,674–1,177,221 | 1,475,305–1,475,975 | rrs | Mb5019 | Ribosomal RNA 16S | Mb0003
#3 | Gubbins; RDP4 | 1,532,736–1,532,787 | 1,953,495–1,953,548 | narX | Mb1765c | Probable nitrate reductase NarX | Mb1792361
#4 | Gubbins | 1,532,751–1,532,781 | 1,953,840–1,953,870 | narX | Mb1765c | Probable nitrate reductase NarX | Mb1792361
#5 | Gubbins; RDP4 | 1,794,609–1,794,714 | 2,283,200–2,283,315 | pkS12 | Mb2074c | Probable polyketide synthase pkS12 | Mb0003
#6 | Gubbins | 1,794,627–1,794,780 | 2,283,713–2,285,136 | pkS12 | Mb2074c | Probable polyketide synthase pkS12 | Mb0003
#7 | Gubbins | 2,242,002–2,242,098 | 2,839,474–2,839,570 | tatA | Mb2121 | Probable Sec-independent protein translocase membrane-bound protein tatA | Mb0565
#8 | Gubbins | 3,244,551–3,244,556 | 4,003,420–4,003,425 | espA | Mb3646c | Conserved hypothetical alanine and glycine rich protein | Mb2043

Table 4. Detailed information concerning the recombination events identified by Gubbins and RDP4 in the entire dataset. Genome positions according to M. bovis AF2122/97.

| Recombination event | Alignment positions | RDP (p-value) | GENECONV (p-value) | Bootscan (p-value) | MaxChi (p-value) | Chimaera (p-value) |
|---|---|---|---|---|---|---|
| #2 | 1,176,674–1,177,221 | 7.524×10⁻²² | 1.871×10⁻²⁰ | 1.004×10⁻¹⁵ | 9.926×10⁻⁰⁵ | 9.753×10⁻⁰⁵ |
| #3 | 1,532,736–1,532,787 | 3.771×10⁻¹⁰ | 5.216×10⁻⁰⁸ | 5.634×10⁻¹² | – | – |
| #5 | 1,794,609–1,794,714 | 1.388×10⁻¹¹ | 2.324×10⁻¹⁰ | 6.200×10⁻¹³ | – | – |

Table 5. Statistical values associated with different algorithms implemented in RDP4 for the confirmed recombination events.

Effect on M. bovis genetic diversity when comparing with mutation (Table 3). To make a broad comparison, the r/m parameter was estimated using a similar methodology for an MTBC dataset composed by 23 genomes, revealing a mean value of 0.486, while for the 38 M. bovis dataset of Patané and co-workers it evidenced a mean value of 0.98. In the first study there were only two M. bovis (M. bovis BCG and reference strain) within the 23 genomes included in the work, so the obtained value might be biased by the overrepresentation of M. tuberculosis genomes. In the second report, the M. bovis population under analysis was mainly recovered from American countries and livestock hosts. In contrast, in our dataset, a higher number of geographic locations and host species is represented, and genomes grouped into different clonal complexes with distinct population genetic signatures were also used, enabling a deeper and wider population knowledge. The differential r/m average values obtained with our dataset are consistent with the notion that the extent of recombination varies widely among lineages assigned to the same taxonomical species, so these results suggest that M. bovis clonal complexes might exhibit a differential impact of recombination, as also suggested by Didelot & Maiden. Nevertheless, enlarging significantly this dataset with the inclusion of a higher number of M. bovis genomes would allow further clarification of this point. Both r/m and rho/theta parameters present variability among the tree branches, a result that is in agreement with reports concerning other bacterial species.

Finally, to confirm the recombination events identified by Gubbins pipeline, the different core multi-alignments were also independently tested in RDP4 software with six different algorithms. Globally, less than half of the events identified by Gubbins were confirmed by RDP4 (Tables 4, 5). Considering the entire dataset, three recombination events were confirmed, two involving internal nodes and another one involving a single genome in a terminal branch and for which a clonal complex could not be assigned (Tables 4, 5). The identification of events in terminal branches might be a sign that recombination is still ongoing in contemporary M. bovis strains or the result of misalignment. In this putative recombination region, circa 20% of positions have an undefined nucleotide (N), which can therefore influence the recombination signal (Supplementary Fig. 2). Moreover, this region affects the rrs gene, encoding the 16S ribosomal RNA that is expected to be highly conserved, so this putative recombination signal could be the result of a sequencing error or wrong alignment. Whole genome alignment between Mb0003 and M. bovis AF2122/97 was thus then performed and the presence of undefined nucleotides and of SNPs was confirmed, so the likely issues related to wrong alignment did not arise as a consequence of the bioinformatics procedure implemented in this work.

No gaps or undefined nucleotides were identified in the recombination regions of internal nodes (Figs. 4, 5). With respect to these events, one encompasses exclusively Eu2 genomes, affecting the pkS12 gene that encodes a probable polyketide synthase; while the other one is registered across Eu1 genomes and affects narX gene.
encoding a probable nitrate reductase (Table 4). Overall, the recombination analysis suggested the presence of a limited number of recombination segments with statistical support, and the inferred metrics indicate a lower effect of recombination on *M. bovis* genealogy. The recombination signal was expected to be low, however it is important to distinguish true evolutionary signals from background noise, which is a challenging task. In order to decrease the noise signal proposed to be introduced by reference-based assemblies and misalignment issues,

Figure 4. Detailed visualization of alignment in the recombination region of *M. bovis* dataset affecting the *narX* gene encoding a probable nitrate reductase. No gaps or undefined nucleotides were identified in the recombination region of internal nodes. This specific event was registered across Eu1 genomes. The quality of sequencing of *narX* gene was evaluated by read mapping against *M. bovis* AF2122/97. The SNP positions suggested in the recombination region were confirmed by applying the criteria referred to in the methods section (at least 20 reads and 0.9 frequency of alteration). The polymorphisms at *narX* gene were fully confirmed in genomes Mb1792361 and Mb7240415 (2.3%).

Figure 5. Detailed visualization of alignment in the recombination region of *M. bovis* dataset affecting the *pks12* gene. No gaps or undefined nucleotides were identified in the recombination region of internal nodes. With respect to this event affecting the *pks12* gene that encodes a probable polyketide synthase, it encompasses exclusively Eu2 genomes. The quality of sequencing of *pks12* was evaluated by read mapping against *M. bovis* AF2122/97. The SNP positions suggested in the recombination region were confirmed by applying the criteria referred to in the methods section (at least 20 reads and 0.9 frequency of alteration). The polymorphisms were fully confirmed for genomes Mb0891, Mb1711, Mb1789, Mb1870, Mb1758, Mb2043, Mb1960.
with the exception of complete genomes, all the remaining ones were de novo assembled and the quality of assemblies was checked and secured via QUAST pipeline analysis (Supplementary Table 1). Moreover, a series of complementary analyses was performed to provide robustness and accurateness to the overall investigation. Thus, the quality of sequencing of narX and pks12 genes was evaluated by read mapping against M. bovis AF2122/97. The SNP positions suggested in the recombination region were confirmed by applying the criteria referred in the methods section (at least 20 reads and 0.9 frequency of alteration). The polymorphisms at narX gene were fully confirmed in two genomes (Mb1729361 and Mb7240415: 2.3%), as well as in the case of pks12 gene for genomes Mb0891, Mb1711, Mb1789, Mb1870, Mb1758, Mb2043, Mb1960. However, for genome Mb2043, six out of eight positions did not meet the read depth criteria because the SNPs were supported by a maximum of 17 reads that was below the established cut-off of 20. Recombination at this genome spot could thus be confirmed for six genomes (8.6%) (Figs. 4, 5).

PE and PPE genes have repetitive regions prone to misreading by Illumina sequencing and mis-mapping and so are commonly removed from the bioinformatics workflow of Mycobacterium tuberculosis members only when a strategy of map to sequence is used. The inference of recombination events applied in this work was based on de novo assemblies for which PE/PPE were not filtered out. We believe that the strategy applied, with the implementation of three different, complementary approaches and algorithms by SplitsTree, Gubbins pipeline and RDP4 software, is robust to deal and filter recombination regions arising from false signals. Nevertheless, to exclude the interference of PE/PPE genes on the identification of SNP clusters by Gubbins and RDP4 software, and consequently on the identification of the recombination regions proposed to affect narX and pks12 genes, the neighbourhood of these genes was inspected (Supplementary Fig. 3–5). In M. bovis AF2122/97, the narX gene is delimited by narX2 and Mb1764c, while pks12 is surrounded by Mb2075c e Mb2073c (Supplementary Fig. 3–5). Synteny maps with MAUVE using complete genomes yielded plots providing information about gene order conservation and rearrangements, showing four colinear blocks, without signs of genome translocations or inversions. Furthermore, a complementary analysis with aminoacidic sequences evidenced synteny in all complete genomes and no PE/PPE were identified in the neighbouring regions of narX or pks12. For narX, one genome (Mb0030) had a lower synteny score, since narX gene is identified in two segments (segment 1891 and 1890). For pks12, Mb0030 and Mb003 present lower synteny scores due to a similar situation, whereas pks12 is identified in two and three segments, respectively, representing different domains of the protein (Supplementary Fig. 3–5). Considering this information and that both Gubbins and RDP4 software perform an analysis inspecting the core multi-alignment in windows with a maximum of 500 bp, we confirmed that the PE/PPE genes did not interfere with the recombination signals affecting narX and pks12.

Although the recombination signals detected in this dataset may be considered residual, recombination in M. bovis cannot indeed be excluded and should thus continue to be the subject of further analyses for which sequencing of whole genomes from different epidemiological scenarios is crucial.

Comparing the obtained ML phylogenetic trees before and after the recombination correction (Fig. 2A,B) did not lead to significant changes in the inferred phylogenetic relationships, with M. bovis strains being gathered within the same groups.

**An evolutionary scenario for M. bovis from a multi-host TB system in Portugal.** A SNP alignment containing 1816 polymorphic positions was obtained after mapping reads of 42 newly sequenced M. bovis against the reference genome of M. bovis AF2122/97. The majority of SNPs (87.1%) was located in coding regions and the affected genes were characterized according to functional categories displayed in Bovilist (Fig. 6A,B). After accounting for the total number of genes per functional category, the genes encompassed in “Lipid metabolism” category presented the higher number of SNPs, followed by “Cell wall and cell process” and “Regulatory from the ancestral state and representing positive (diversifying or directional) and/or relaxed purifying selection evolution. M. bovis” (Fig. 6A,B). After accounting for the total number of genes per functional category, the genes encompassed in “Lipid metabolism” category presented the higher number of SNPs, followed by “Cell wall and cell process” and “Regulatory from the ancestral state and representing positive (diversifying or directional) and/or relaxed purifying selection evolution. M. bovis” (Fig. 6A,B). After accounting for the total number of genes per functional category, the genes encompassed in “Lipid metabolism” category presented the higher number of SNPs, followed by “Cell wall and cell process” and “Regulatory from the ancestral state and representing positive (diversifying or directional) and/or relaxed purifying selection evolution. M. bovis” (Fig. 6A,B). 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Globally, the average dN/dS ratio is superior to 1.5, which suggests a global evolutionary pressure to escape from the ancestral state and representing positive (diversifying or directional) and/or relaxed purifying selection scenarios. In the categories “Virulence, detoxification, adaptation”, “Insertion seqs and phages” and “Regulatory proteins”, over two-thirds of SNPs were non-synonymous (Fig. 6B).

In all categories, there were genes with more than one SNP, leading to an average rate of mutation (i.e. the mean value of SNPs per gene) greater than one (Fig. 6A). The higher mutation values were harboured by pks12 (Mb2074c) with 15 SNPs and fas (Mb2553c) with 8 SNPs. Both genes are involved in fatty acid metabolism. The pks genes encode polypeptide syntheses (PKS) which are multifunctional enzymes involved in the biosynthesis of mycobacterial cell wall lipids. The pks gene encodes a multifunctional polypeptide that is involved in the synthesis of mycoketides. The fas gene is involved in the synthesis of mycolic acids. Both genes play an import role in the biosynthesis of the cell wall that is at the interface with the host.

**SNP-detailed analysis of HGT and 3R genes.** To further study the evolutionary processes within M. bovis, two specific groups of genes were analysed. Previous published works using sequence composition and phylogenetic methods identified genes that were acquired through HGT by the MTBC ancestor before diversification. Those genes are listed in Supplementary Table 2. The SNP distribution was analysed in a total of 77 genes presumably involved in HGT, and 26 polymorphic sites were identified, leading, in the majority of cases (78%), to a non-synonymous (NS) change (Supplementary Table 2). Previous work conducted with MTBC genomes evidenced that putative HGT regions present a higher ratio of NS SNPs when comparing with the rest of the genome. If one considers that these recombination tracts were acquired by the MTBC ancestor and, thus, they over-represent ancient polymorphisms, then it would be expected a higher fraction of synonymous alterations, since NS substitutions are expected to be eliminated by negative selection, as the changes in amino acid
might modify protein function. So, our results suggest that functional consequences may arise from substitutions in HGT-like genes, which remits to their importance on valuable adaptive genetic diversity.

In parallel with this analysis, the genes encoding 3R (DNA repair, replication and recombination) system components were thoroughly examined, following the previous published list by dos Vultos and collaborators (2008)\textsuperscript{39}. The exchanges of identical DNA fragments cannot be directly observed, although it might be a frequent process when involving closely related bacteria, such as in the case of this dataset; plus, this process might be crucial as a DNA repair method\textsuperscript{72} and thus play a role in homologous recombination. A total of 26 polymorphic positions distributed by 54 genes were identified (Supplementary Table 3). In this group of genes, NS changes account for about 65% of the consequences, which is in agreement with a previous report for \textit{M. tuberculosis} strains\textsuperscript{39}.

Gene and nucleotide diversity (\(\pi\)) were evaluated for the genes presenting polymorphisms. Gene diversity is a measure of the uniqueness of a particular gene sequence in a population. Average values of 0.256 and 0.226 were obtained for HGT and 3R group genes, respectively. When the value of gene diversity index is zero, all the sequences under analysis are equal. Therefore, the values obtained in this work reveal that there is limited genetic diversity within the selected panel of genes. The nucleotide diversity (\(\pi\)) compares the similarity per site between two nucleotide sequences. When \(\pi\) is superior to 0.003 it can be considered that the group of sequences under analysis is highly diverse. In our analysis, both gene groups reveal an average value inferior to 0.003, with HGT registering 0.00034 and the 3R circa. 0.00021. No gene had a \(\pi\) value higher than 0.003, thus also confirming limited nucleotide diversity within the selected gene panels.

The Tajima’s D test of neutrality was also evaluated, and in both groups there were genes with positive and negative values, evidenced by an average value inferior to zero. The selection against deleterious mutations, past selective sweeps and population expansion after a recent bottleneck are pointed as possible causes to decrease the result from Tajima’s D test.

**Balance of forces in \textit{M. bovis} evolution.** Natural selection is a mechanism of evolution and has been associated with MTBC evolution\textsuperscript{7}. Selective sweeps (i.e. positive selection that leads to the fixation of a new
beneficial mutation) and background selection (i.e. selection against a deleterious mutation that leads to the elimination of any mutation linked to the target of selection) are both linked to the action of natural selection.

In this work, several evidences support the importance of natural selection: (1) SNP distribution is not random, with genes included in the “lipid metabolism”, “cell wall and cell processes” and “intermediary metabolism and respiration” categories presenting a higher SNP rate; (2) regions proposed to be transferred from holism ancestor also accumulate an excess of SNPs; and (3) the HGT and 3R groups evidenced a global average value inferior to zero in the neutrality tests, indicating a past selective sweep or expansion after bottleneck. Furthermore, the high proportion of low-frequency genetic variants, particularly singletons, is one of the features associated with MTBC population genetics, and proposed to reflect the influence of background selection, an effect that is also confirmed in this work, as 372 (20.5%) of the 1816 considered SNPs are strain-specific.

The global elevated value of dN/dS ratio is commonly associated with a positive selection force, likely due to diversifying selection and local selective sweep. However, such patterns in effective population size might have contributed, partially, to this unusual rate of NS per synonymous mutations, once mutations that might have been deleterious in a population with a large effective population size can drift to a high frequency in a small population and, in that way, reflecting reduction in the efficacy of purifying selection as a consequence of increased genetic drift.

The affected genes could confer important adaptive advantages through NS substitutions, however functional studies would be necessary to understand the consequences arising from those SNPs and to infer what would be the benefits for mycobacteria. Recent work performed by Yang and collaborators with M. tuberculosis strains suggested that this evolutionary pressure could allow accessory genes (i.e. genes that are not present in all strains or strain-specific genes) to gradually dominate and eventually become core genes (i.e. present in all strains). This could provide important adaptive and resistance capacities, if considering that accessory genes might be involved in virulence, immune system evasion or antibiotic resistance.

Therefore, a deeper understanding of the role of these evolutionary forces is required to determine which genes have contributed significantly to M. bovis evolution in its trajectory of interaction with different hosts in specific disease systems.

Final conclusions and future work

The study of genetic relatedness and structure of obligatory pathogen populations might provide important insights into their intraspecific genomic diversity and evolution arising upon the interaction with the host. In recent years, many technological advances have shed light onto the biology of M. bovis, however the use of high-throughput technologies such as WGS to understand evolutionary steps is still infrequent, with most works in the TB field being focused on M. tuberculosis or in the molecular epidemiology of M. bovis.

In the current work, a diverse M. bovis dataset, with representatives of all described clonal complexes, was used to assess how different evolutionary forces impact and shape the genetic diversity of a population. Altogether, we ended up with a dataset composed of 70 M. bovis strains, representing the most diverse dataset available to infer recombination, when comparing with other publicly available works. Furthermore, we used isolates obtained from multiple hosts, including humans. Although we may speculate that the inclusion of more genomes might have an impact on the identification of recombination events and recombination metrics, this pilot work is already significant in the context of present knowledge. More complete analyses may be conducted in the future with larger M. bovis datasets to confirm our findings.

The impact of recombination in our dataset was assessed through three complementary strategies. Moreover, efforts to avoid unreliable alignments and to guarantee data quality were made, so that the assessment of recombination signals would be as accurate as possible. Although residual, two approaches support a number of recombination events in the examined dataset, which argue against the paradigm that MTBC is strictly clonal. Despite the limited effects on M. bovis diversity when comparing with mutation, recombination events need to be considered in future evolutionary research works in order to further understand their true impact on biological processes, once they may be an important force generating diversification that may translate into virulence, immune evasion and/or antibiotic resistance phenotypes.

Indeed, previous WGS works support recombination in M. canetti, showing that strains are highly recombinogenic and evolutionary early–branching, with larger genome sizes, 25-fold more SNPs relative to MTBC members. Those works also provide experimental evidence of how pks5-recombination-mediated bacterial surface remodelling in M. canetti increased virulence, driving evolution from smooth to rough morphology and from generalist mycobacteria (M. canetti) towards professional pathogens of mammalian hosts (MTBC). Moreover, a recent work performed by Chiner-Oms and collaborators (2019) found evidences of recombination between the MTBC ancestor and M. canetti ancestor (before diverging to M. canetti), thus proposing the existence of recombination potential before the diversification of MTBC into different ecotypes. So, efforts to expand this topic across all MTBC ecotypes should continue in the future. In this work, we excluded recombination in genomes from the African clonal complexes, nevertheless, a broader sample dataset would be necessary to accurately address the differences amongst clonal complexes members.

Following, the comparative genomic analyses performed in a smaller group of genomes representative of M. bovis population from an endemic TB scenario in Portugal suggested that genes included in the “lipid metabolism”, “cell wall and cell processes” and “intermediary metabolism and respiration” categories have a superior importance in M. bovis evolution and a global positive selection force was suggested to be acting upon this population, as informed by the elevated dN/dS ratio.

Finally, this work reinforces the value of WGS as a high-resolution tool for the analysis of M. bovis genomic diversity and provides insights into the role of recombination and positive selection as evolutionary driving forces in a pathogen affecting a large range of host species, with economical and biodiversity impacts across the world.
Data availability
The newly sequencing data included in this work is deposited under the following Biosample accession numbers: SAMN17004141-SAMN17004143, SAMN17004145-SAMN17004174, SAMN17004176-SAMN17004184 and under the Bioproject accession number PRJNA682618 at a public domain server in National Centre for Biotechnology Information (NCBI) SRA database.

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M.V.C. conceived this work and secured resources and funding. A.C.R. performed the bioinformatic analyses under the guidance of M.V.C. and explored the data under MVC supervision. A.C.R. wrote the first draft of the manuscript and M.V.C. critically revised all versions. Both authors approved the final version.

Competing interests
The authors declare no competing interests.

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