

Genomic Characterization of IS6110 Insertions in *Mycobacterium orygis*

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ABSTRACT: *Mycobacterium orygis*, a subspecies of the *Mycobacterium tuberculosis* complex (MTBC), has emerged as a significant concern in the context of One Health, with implications for zoonosis or zoonoanthroposis or both. MTBC strains are characterized by the unique insertion element IS6110, which is widely used as a diagnostic marker. IS6110 transposition drives genetic modifications in MTBC, imparting genome plasticity and profound biological consequences. While IS6110 insertions are customarily found in the MTBC genomes, the evolutionary trajectory of strains seems to correlate with the number of IS6110 copies, indicating enhanced adaptability with increasing copy numbers. Here, we present a comprehensive analysis of IS6110 insertions in the *M. orygis* genome, utilizing ISMapper, and elucidate their genetic consequences in promoting successful host adaptation. Our study encompasses a panel of 67 paired-end reads, comprising 11 isolates from our laboratory and 56 sequences downloaded from public databases. Among these sequences, 91% exhibited high-copy, 4.5% low-copy, and 4.5% lacked IS6110 insertions. We identified 255 insertion loci, including 141 intragenic and 114 intergenic insertions. Most of these loci were either unique or shared among a limited number of isolates, potentially influencing strain behavior. Furthermore, we conducted gene ontology and pathway analysis, using eggNOG-mapper 5.0, on the protein sequences disrupted by IS6110 insertions, revealing 63 genes involved in diverse functions of Gene Ontology and 45 genes participating in various KEGG pathways. Our findings offer novel insights into IS6110 insertions, their preferential insertion regions, and their impact on metabolic processes and pathways, providing valuable knowledge on the genetic changes underpinning IS6110 transposition in *M. orygis*.

KEYWORDS: *Mycobacterium orygis*, IS6110, bovine tuberculosis, zoonosis, *Mycobacterium tuberculosis* complex, One Health

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Introduction

The insertion sequence IS6110 is found explicitly in *Mycobacterium tuberculosis* complex (MTBC), the causative agent of tuberculosis (TB) in humans and other mammals, including wildlife and farm animals, which are responsible for zoonotic or zoonoanthropotic TB transmission. This unique trait made IS6110 a historical epidemiological marker for diagnosing MTBC in biological samples, enabling the detection of TB outbreaks and transmission chains, although currently being replaced by robust sequencing methods.^{1–3} The transposition of IS6110 nurtures various genetic modifications in MTBC strains, conferring genomic plasticity and significant biological implications.⁴ Insertions of IS6110 amid the coding region might render the gene inactive or act as a mobile promoter controlling gene activation, leading to unusual consequences in the host-pathogen evolution.^{5,6} The members of the MTBC are classified into high (>6) and low (<7) copy strains based on the content of IS6110, although no clear evidence accounts for its phenotypic consequences in bacterial physiology and pathogenesis.⁷ However, *M. tuberculosis* Beijing/W lineage (L2) possessing a high content of 15 IS6110 insertions is linked with high virulence and colossal spread of drug-resistant strains.⁸ It has

been demonstrated that a reasonable number of IS6110 might render strain-specific phenotypes catering to selective advantages during its course of infection rather than excessive accumulation resulting in inactivation or deletion of essential genetic regions, being detrimental to the bacterium.⁵ Among the animal-adapted MTBC, *Mycobacterium bovis* (*M. bovis*) is known to possess only 1 or few copies of IS6110,⁴ whereas *M. orygis* possess high copy numbers ranging from 17 to 20, however, the exact position of insertions are yet unknown⁹ and not much information is available on other members which possess a zoonotic risk. *M. orygis*, has been isolated from various species, including oryx, waterbucks, cattle, deer, rhinoceros, monkeys, and humans and the exact extent of its host range still remains unknown.^{9–16} A recent study in India identified 7 *M. orygis* isolates among 940 clinical isolates from humans.¹⁷ Our laboratory has frequently been isolating *M. orygis* from cattle and wild ungulates recently, and various reports on this species are also being reported in South Asia.^{13,14}

The advent of Whole Genome Sequencing (WGS) has facilitated to localize the IS6110 position in the genomes of MTBC and has been recently used to identify its role in bovine adaptation.^{18,19} This study investigates the distribution of



IS6110 among *M. orygis* strains and its effect on the genes by *in silico* characterization. We have manifested the use of WGS to localize the IS6110 copies and their chromosomal distribution using the sequences of 67 *M. orygis* strains, comprising 56 WGS paired reads downloaded from the National Centre for Biotechnology Information–Sequence Read Archive (NCBI-SRA) and 11 WGS paired reads sequenced from our lab and to evaluate the level of IS6110 sequence stability and their orthology evolving in a multi-host system. Ever since the publication of the complete genome of *M. orygis* 51145,²⁰ it has been used well as a reference genome (GenBank accession No. CP063804) for analysis of *M. orygis* strains.¹³ Although the complete genome has been sequenced, the genes are not functionally annotated, and the mycobrowser (<https://mycobrowser.epfl.ch/>) identifies these genes as unknown, thereby demanding the use of orthologous comparative methods to identify its function. Hence, to overcome this, we additionally used *M. tuberculosis* H37Rv (NC000962.3) as the reference genome.

Materials and Methods

Data acquisition and analysis

To compare the sequence of our study isolates with existing sequences available in the database, NCBI SRA was searched with the term “*Mycobacterium orygis*.” All the sequences branded as *M. orygis* or *M. tuberculosis* var. *orygis* were downloaded from SRA with the fasterq-dump tool from the SRA toolkit (version 2.9.6) available at <https://ncbi.github.io/sra-tools/>. Only the sequences which downloaded as paired reads, and the sequences which possessed the species-specific Region of Difference (RDs) and single nucleotide polymorphisms (SNPs) were used for further analysis (Table S6). A total of 56 paired-end sequences of *M. orygis* strains isolated from human and animal sources were used, along with 11 study isolates. The raw sequences were filtered using trimomatic (version 0.39) with a framework of minimum phred quality score and read length set to 30 and 80, respectively. The filtered sequences were further analyzed through vSNP <https://github.com/USDA-VS/vSNP> using *M. orygis* 51145 (CP063804) as the reference genome to generate SNPs. The phylogenetic tree was constructed based on the aligned whole-genome SNP sequences under a GTRCATI model of substitution and maximum-likelihood algorithm with a bootstrap replication of 1000²¹ and visualized using the interactive tree of life (iTOL).²²

Analysis of IS6110 distribution

ISMMapper (version 2.0)²³ pipeline https://github.com/jhawkey/IS_mapper was used to localize IS6110 on short-read sequences with *M. tuberculosis* H37Rv (NC000962.3) and *M. orygis* 51145 (CP063804) as reference genomes. The presence of IS6110 was visualized manually with Integrative Genomics Viewer^{24,25} using the bam file generated by ISMapper 2.0.

Analysis of orthologous genes

The genomic positions of the IS6110 were deduced through ISMapper analysis. The genes surrounding the insertion sites (intergenic) and genes possessing the insertion sites within them (intragenic) were identified based on the respective reference genomes to determine orthologous genomic sites of the IS.

Gene ontology enrichment and KEGG pathway analysis

The protein sequences of the upstream and downstream genes of IS6110 and the protein sequence of the genes which possessed the IS6110 insertions were used to perform functional annotation using eggNOG-mapper (version emapper—2.1.9)²⁶ based on eggNOG orthology data.²⁷ Sequence searches were performed using DIAMOND (version 2.0.11)²⁸ to mine the orthologs.

Results

Distribution of IS6110 among *M. orygis* strains

In our analysis of IS6110 localization among the 67 sequences from both reference genomes, we observed that 4.5% (3/67) of the strains in the human cluster did not possess any IS6110 insertions and were classified as no-copy strains. We found that 91% (61/67) of the strains were high-copy, while 4.5% (3/67) were classified as low-copy. The number of IS6110 loci varied within the samples. In the human cluster, the loci ranged from 2 to 30 (with a median of 25) using the reference CP063804. In the animal cluster, the loci ranged from 20 to 42 (with a median of 26.5) using the same reference. However, when using the reference NC000962.3, the number of loci ranged from 2 to 43 (with a median of 32) in the human cluster and from 26 to 54 (with a median of 35) in the animal cluster. Overall, we identified a total of 24 common insertion loci. The specific regions of these insertion sites are tabulated and schematically represented, along with the phylogenetic tree generated using the maximum likelihood approach with all 67 *M. orygis* sequences (Figure 1 and Table 1).

Unique and novel IS6110 insertions in *M. orygis* strains

In addition to the previously mentioned typical and known IS6110 insertions, we also identified novel IS6110 insertions using ISMapper. When mapped against CP063804, we identified 107 novel insertion loci, of which, 61.7% (66/107) were intragenic and 38.3% (41/107) were intergenic. Approximately 58% (62/107) of these insertion loci were unique or shared between 2 and 3 strains, while 15 loci were shared in at least 4 strains, with only one being intergenic. Of the 15 common insertion sites, 11 were exclusively found in the animal cluster, 3 in both the human and animal clusters, and 1 insertion site

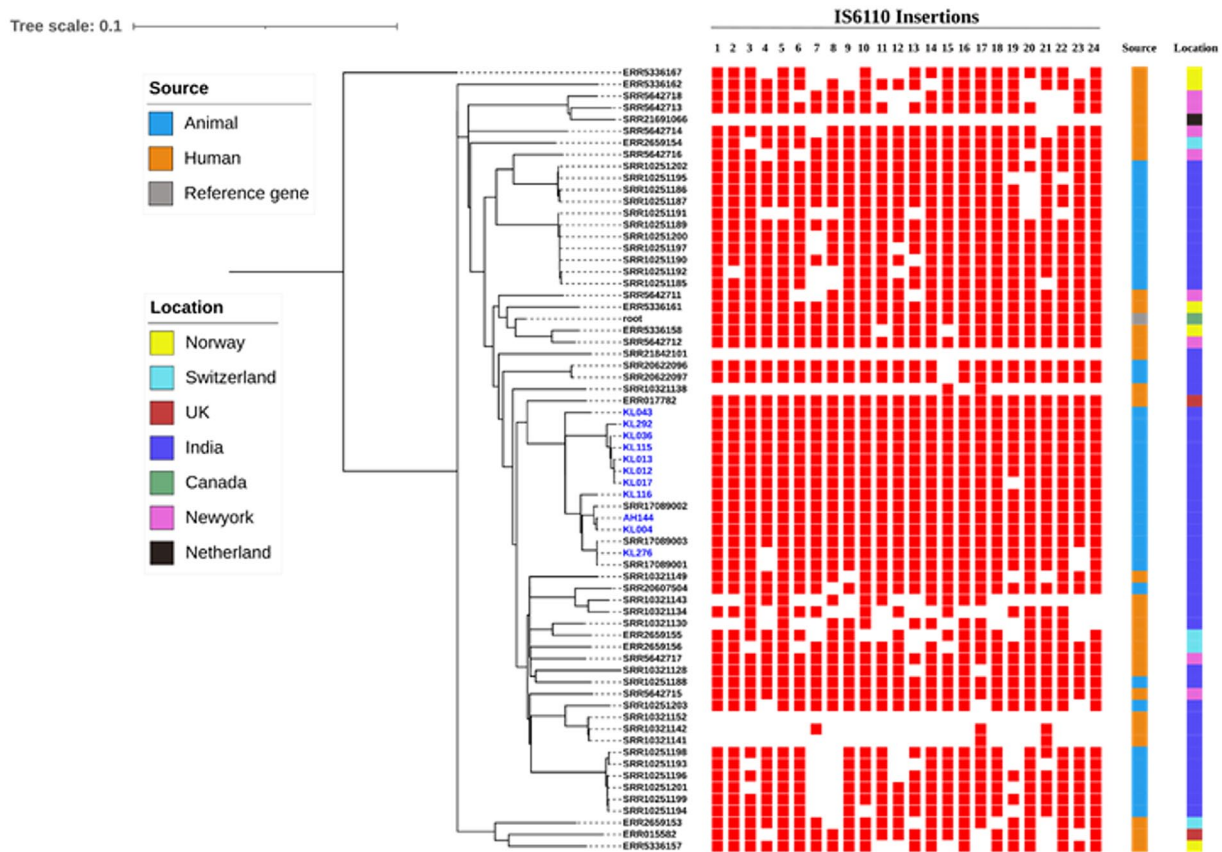


Figure 1. Contextual Phylogenetic tree combining the 11 study isolates along with 56 *M. orygis* isolates around the globe: Constructed using whole genome SNPs obtained after alignment with *M. orygis* reference genome CP063804. The heatmap shows the presence or absence of IS6110 in 67 *M. orygis* genomes. Red squares indicate the presence of IS6110 in a specific site. Descriptions of each IS6110 site are described in Table 1. The source and the geographic location of the isolates are represented in the respective bands. The study isolates are labeled in blue.

was found only in the human cluster. Almost 124 novel insertion loci were identified when mapped against NC000962.3, out of which 39.5% (49/124) were intergenic and 60.5% (75/124) intragenic. Only 14 insertion sites were identified in both clusters, of which only 2 were intergenic. Notably, we observed variations in the localization and orientation of IS6110 insertions among the strains despite being present in the same genomic region. These findings shed light on the diversity and distribution of novel IS6110 insertions, providing valuable insights into the genetic landscape of the studied clusters. The genes on either side of IS6110 and those that contained IS6110 in both the H37Rv and 51145 reference genomes were combined. We identified 227 genes, among which 85 were from *M. tuberculosis*, 64 from *M. orygis* and 79 genes were found to be common in both *M. orygis* and *M. tuberculosis* (Supplemental Table S1). Using eggNOG Mapper, the protein sequences of these 227 genes were subjected to functional annotation.

Genetic significances of IS6110 insertions

Approximately 204/227 genes were scanned by eggNOG 5.0, and 82 were identified to be functionally involved in Gene Ontology (GO) and KEGG pathway (KP) enrichment

analysis. Almost 37/82 genes were exclusively linked to various processes in GO and 19/82 genes were mapped in numerous KP. Additionally, we also identified 26 genes that were involved in both GO and KP. A total of 63 genes were encompassed in 669 GOs comprising 455 Biological processes (BP), 161 Molecular Functions (MF) and 53 Cellular components (CC), and the distribution of the genes in various functions are represented in Figure 2. Among the 63 genes involved in GO, 32 were found only in the human cluster, 15 in the animal cluster, and 16 in both clusters. The intergenic insertion sequence upstream of Rv0742 (Hypothetical protein) and Rv2965c/Rjtmp_003059 (*kdtB/coaD*) were found in almost 61 and 60 isolates, respectively. The intragenic insertion sequences present within the genes Rv1266/Rjtmp_001332 (*pknH*), Rv2565 (NTE family protein), Rv1753 (*PPE24*), Rv0402/Rjtmp_000420/Rjtmp_000422 (*mmpL1*) and Rv1367c/Rjtmp_001448 (Hypothetical protein) were also found in almost 55 to 61 isolates. All these genes were involved in various GO processes (Supplemental Tables S2 and S3). Other insertion regions in the human cluster were unique and found in only 1 strain or at least 2 strains. A similar pattern was observed in the animal cluster except for 1 intergenic region between Rv2338c/Rjtmp_002419 (*moeW/thiF* family adenylationase) and Rv2339/Rjtmp_002420 (*mmpL9*) and an

Table 1. Details of IS6110 insertions among all *M. orygis* isolates.

| IS NO. | ORIENTATION | LOCI | POSITION | NO. OF ISOLATES |
|--------|-------------|----------------------------|-----------------|-----------------|
| 1 | R | RJtmp_000364: RJtmp_000365 | 418095-419370 | 61 |
| 2 | R | RJtmp_000420: RJtmp_000421 | 481559-482854 | 59 |
| 3 | R | RJtmp_000782: RJtmp_000783 | 835711-837062 | 60 |
| 4 | R | RJtmp_000841: RJtmp_000842 | 892501-893790 | 52 |
| 5 | R | RJtmp_001232: RJtmp_001233 | 1304672-1305974 | 60 |
| 6 | F | RJtmp_001332: RJtmp_001333 | 1423001-1424355 | 55 |
| 7 | R | RJtmp_001435: RJtmp_001436 | 1536637-1537898 | 44 |
| 8 | R | RJtmp_001448: RJtmp_001449 | 1550641-1551955 | 58 |
| 9 | F | RJtmp_001738: RJtmp_001739 | 1891878-1893148 | 56 |
| 10 | R | RJtmp_001757: RJtmp_001758 | 1909881-1911200 | 59 |
| 11 | F | RJtmp_001833: RJtmp_001834 | 198611-1987961 | 55 |
| 12 | R | RJtmp_001837: RJtmp_001838 | 1994072-1995329 | 44 |
| 13 | F | RJtmp_001872: RJtmp_001873 | 2035374-2036722 | 50 |
| 14 | F | RJtmp_001988: RJtmp_001989 | 2151738-2152992 | 58 |
| 15 | F | RJtmp_002433: RJtmp_002434 | 2602192-2603492 | 56 |
| 16 | R | RJtmp_002654: RJtmp_002655 | 2842275-2843648 | 60 |
| 17 | F | RJtmp_002901: RJtmp_002902 | 3068456-3069736 | 59 |
| 18 | F | RJtmp_003056: RJtmp_003057 | 3264524-3265824 | 57 |
| 19 | R | RJtmp_003106: RJtmp_003107 | 3314904-3316281 | 51 |
| 20 | R | RJtmp_003217: RJtmp_003218 | 3429234-3430489 | 52 |
| 21 | F | RJtmp_003255: RJtmp_003256 | 3470593-3471871 | 53 |
| 22 | F | RJtmp_003278: RJtmp_003279 | 3493266-3494534 | 55 |
| 23 | R | RJtmp_003430: RJtmp_003431 | 3653243-3654495 | 52 |
| 24 | R | RJtmp_003855: RJtmp_003856 | 4137024-4138299 | 59 |

intragenic insertion in Rv2430/RJtmp_002513 (*PPE41*) which were found in at least 6 and 7 strains, respectively.

Almost 62 KPs were identified in which nearly 45 genes were involved, including 20 from the human cluster, 14 from the animal cluster and 11 genes from both human and animal clusters. Here too, we found that the genes from both clusters mainly were unique or shared between a couple of strains. The majority of the genes (23/45) were involved in metabolic pathways (map01100) and 13/45 genes were involved in the biosynthesis of secondary metabolites (map01110). In addition to metabolic pathways, genes such as Rv0742, Rv2964/RJtmp_003056 (*purU*), Rv2965c/RJtmp_003059 (*kdtB/coaD*) were found in almost 60 strains and these genes were involved in glycerolipid metabolism (map00561), Glycolate and dicarboxylate metabolism (map00630) & Pantothenate and CoA biosynthesis (map00770) respectively. The gene Rv1368/RJtmp_001450 (*lprF*), found in 7 isolates (3 in human and 4 in

animal cluster) was involved in the TB pathway (map05152), and RJtmp_000891 (*pip*) found only in 14 isolates from the animal cluster was involved in arginine and proline metabolism (map00330). Most of the genes *dnaN*, *gap*, *icd*, *lpdB*, *proA*, *trpE*, *glgC*, *plcC*, and RJtmp_002367 were involved in at least 5 or more pathways but were found only in 1 or 2 isolates (Supplemental Tables S4 and S5).

IS6110 insertions and genetic significance among our study isolates

All our study isolates belonged to the animal cluster and were high-copy strains possessing more than 25 IS6110 insertions. In addition to the previously mentioned insertion regions, we also identified an IS6110 intergenic insertion between Rv1470/RJtmp_001552 (*trxA*) and Rv1471/RJtmp_001553 (*trxB1*) in 2 of our study isolates (KL013, KL017). *TrxA* was involved in

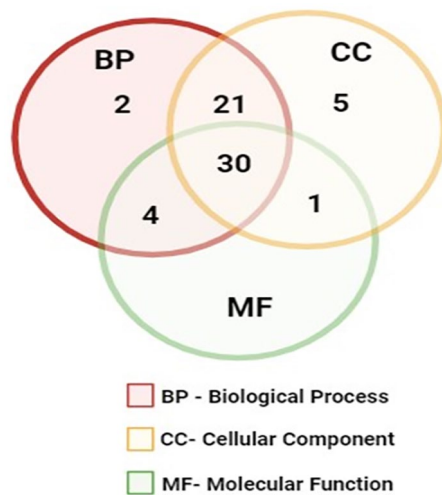


Figure 2. Distribution of 63 genes in various GO process.

seleno compound metabolism (map00450), NOD-like receptor signaling pathway (map04621), and Fluid shear stress and atherosclerosis pathway (map05418). Another intragenic insertion within Rjtmp_001745 was exclusively found in 6 of our isolates (KL012, KL013, KL017, KL115, KL036, and KL292). Strain KL012 and KL292 possess a unique insertion in the genes Rjtmp_000047 (*marR* transcriptional regulator) and Rjtmp_002096 (*csm5*), respectively and the gene *csm5* was identified to be involved in 11 GOs. A total of 7 unique insertions were found in strain KL036, among which 3 were intergenic and 4 intragenic. The intragenic insertions were found in genes Rv1364/Rjtmp_001464 (*carB*), Rv1631/Rjtmp_001705 (dephospho-coA kinase), Rv2379/Rjtmp_002451 (*mbtF*), and Rv1609/Rjtmp_001681 (anthranilate synthase component). Among these genes, *carB* was involved in almost 50 GOs and *mbtF* is involved in the Biosynthesis of siderophore group non-ribosomal peptides (map01053). Intergenic insertions were found between Rv1187/Rjtmp_001253 (*rocA/pruA*) & Rv1188/Rjtmp_001254 (proline dehydrogenase/*putA*), Rv1523/Rjtmp_001610 (methyltransferase) and Rv1524/Rjtmp_001611 (glycosyltransferase), and Rv3800/Rjtmp_003912 (*pks13*) and Rv3801/Rjtmp_003913 (*fadD32*). Among these genes, *pruA* and *putA* are involved in arginine & proline metabolism along with the *pip* gene and *fadD32* is involved in nearly 47 different GOs.

Discussion

The fact that members of the MTBC exhibit non-specific host preferences have now been well established, as demonstrated by the isolation of *M. tuberculosis* from cattle and the identification of *M. bovis* and *M. orygis* in humans. The concept of host tropism within MTBC species reflects the interplay between host-driven and pathogen-driven processes, where variations in the host immune response and alterations in MTBC phenotypes contribute to differences in infection, persistence, disease development, and transmission.²⁹ The adaptation of the pathogen to the host is a highly complex process, as the mechanisms underlying the transition from latent TB infection (LTBI) to active TB

disease are still not fully understood. In this context, the MTBC species possess a distinctive insertion sequence called *IS6110*, which belongs to the IS3 family and plays a significant role in transposition. The transposition of *IS6110* follows a copy-out-paste-in mechanism.³⁰⁻³² This implicit trait offers an outstanding chromosomal polymorphism to study TB outbreaks and its unquestionable role as a clinical epidemiological marker.

The perseverance of this insertion element in MTBC might also steer phenotypic changes and affect the fitness of the strain. Various reports claim these transposition events are instigated by external stress conditions leading to genetic variability.^{33,34} *IS6110* insertions are well known to disturb gene expression by intruding protein-coding genes, by facilitating recombination events resulting in deletions and inversions, or by up-regulating the expression of adjacent genes due to its presence within the promoter region.^{18,33,35,36} Recently Charles et al¹⁹ have made use of the WGS approach to study the abundance and chromosomal distribution of *IS6110* copy on *M. bovis* genomic data of French animal field strains and stated that these insertions seem to be stable within specific genotypes over time and between host species, signifying that the transposition of *IS6110* is not an evolutionary driver for modern French *M. bovis* strains at least over a 15-year period. The effect of *IS6110* on genome function prompted us to take a deeper look at the distribution and patterns of *IS6110* insertions among the recently circulating strain *M. orygis*, the emerging pathogen in India.

M. tuberculosis as a reference genome has been well utilized and is also well known to possess 16 *IS6110* insertions and all the possible regions of insertions have been vastly explored.³⁷ Meanwhile, in the case of *M. orygis*, van Ingen et al had previously identified these strains as high copy strains possessing 17 to 20 *IS6110* copies. However, the exact regions of insertions remained unknown.⁹ Here, we identified 25 *IS6110* insertions in the reference strain and any additional insertions other than the ones found in the reference were identified as novel insertions.

We found that most of the strains in this study were high copy strains, but also identified strains with no copy (SRR10321152, SRR21691066, SRR21842108) and low copy (SRR10321141, SRR10321142, SRR10321138) *IS6110* insertions although insignificant. No copy and low copy strains were found only in the human cluster, whereas the animal cluster consisted of only high copy strains. Among all the isolates, KL036 possessed the highest number of *IS6110* insertions with 42 insertions against CP063804 and 54 against NC000962.3. Most novel insertions were highly confined to a single strain or found in 2 or 3 strains in both human and animal clusters, indicating the potential role of *IS6110* transposition.

The 25 known *IS6110* were present in all the high-copy strains but were found to be localized in different positions and orientations within the same region and these insertions were mostly intergenic. Nearly all the high copy strains possessed *IS6110* insertions downstream to the genes *mosR*, *lipx*, *pknH*, *EccA5*, *MgtC/sapB* family protein, and ExeA family protein. A few other

genes include oxidoreductase, phospholipase, transposase and hypothetical proteins. We found 2 genes for each of *mmpL1* (RJtmp_000420 (912bp), RJtmp_000422 (1971bp)), *lipX* (RJtmp_001232 (177bp), RJtmp_001234 (129bp)) and *mmpL10* (RJtmp_001247 (1743bp), RJtmp_001249 (1269bp)) in *M. orygis* genome compared to Rv0402c (2877bp), Rv1169c (303bp), and Rv1183 (3009bp), respectively in *M. tuberculosis* genome. A closer observation showed the presence of IS6110 insertion sequence RJtmp_000421, RJtmp_001233, and RJtmp_001248 between these genes, which leads to the speculation that the existence of 2 genes might be due to the transposition of the insertion element. Most novel and unique insertion sequences were found within the hypothetical protein, PPE/PE family proteins or PPE-domain-containing proteins. Few transcriptional regulators, such as *marR*, *gntR*, *LuxR*, and *TetR* had IS6110 insertions upstream of the gene but were confined to a single strain. We found no insertion sequences upstream of the previously reported genes *PhoP*, *essS*, or *ctpD*. However, we found an insertion element in the upstream region of *dnaN* in 1 strain from the human cluster.¹⁸

Functional annotation by Gene ontology and KEGG pathway analysis identified numerous functions and pathways in which the genes affected by IS6110 were involved. However, only 7 insertion sequences were found in all these isolates, whereas the rest were only present in a single or a couple of strains. The effect of the insertion sequence between *pknH* and *embR* is unknown. However, the genes *pknH* and *embR* are non-essential regulatory proteins found to be involved in various processes, including biological processes, cellular components and molecular function by GO analysis. Similarly, another insertion sequence existing between *purU* and *coaD/kdtB* was also observed in all the isolates, and the gene *kdtB* is involved in the cell wall and cell process.

In the case of the novel and unique insertions, none of the loci were found among our isolates except for Rv2357 (*glyS*) and Rv1368/RJtmp_001450 (*lprF*). The former was found in 6 isolates, 3 from each cluster including 2 study isolates (KL043 and KL036) and the latter was found in 7 isolates in the animal cluster, including 3 study isolates (KL004, KL115, and AH144). Another insertion loci within Rv0804/RJtmp_000891 (*pip*) was found in all 11 of our study isolates and 3 more in the animal cluster. The gene *glyS* known as glycyl-tRNA synthetase, catalyzes the synthesis of glycyl-tRNA, which is required to insert glycine into protein.³⁸ The role of *LprF* in the tuberculosis pathway is unclear, and we speculate its possibility to display TLR2 agonist activity as most of the Lipoproteins like *LprG*, *LprA* and *LpqH* are well known for the same.³⁹⁻⁴¹ Similarly, in the arginine and proline metabolism, proline iminopeptidase (*Pip*) explicitly catalyzes the removal of N-terminal proline residues from peptides.⁴² Gamma-glutamyl phosphate reductase (*proA*) catalyzes the NADPH-specific reduction of L-gamma-glutamyl phosphate into L-glutamate-5 semialdehyde, which spontaneously cyclizes to form pyrroline-5-carboxylate.⁴³ This intermediate product of proline biosynthesis is converted into glutamate by pyrroline-5-carboxylate dehydrogenase (*PruA/RocA*).⁴⁴

To the best of our knowledge, this is the first study to report the identification and localization of IS6110 insertion sites in the *M. orygis* genome. Some of the insertion sites identified in this study have previously been recognized as insertion hotspots of MTBC. Only a few isolates in this study had IS6110 insertions in the hotspots region like Rv1758 (*cut1*), Rv1777 (*cyp144*), Rv3183, Rv3327, Rv1371, Rv1765c.^{45,46} Nearly all the isolates had an insertion element in Rv1682 and Rv1169, the hotspot regions for L5 and L6 lineages of MTBC, respectively.⁴⁶ Gene interruption or regulation of genes in these hotspot regions might differ depending on the orientations and/or insertion sites of IS6110 in the same genomic region. Fewer studies have suggested that the location of IS6110 in the same transcriptional orientation and not more than 400 bp upstream of a gene may lead to upregulation of the gene.^{6,45} Although we have also detected similar insertions in our analysis, it warrants an additional *in vivo* or *in vitro* study to assess their role in virulence and other survival traits. The study is the beginning, and we presume that the genetic changes observed due to IS6110 transposition could be partly responsible for the fitness of *M. orygis* in establishing infection in various hosts. At the same time, other explanations could also support this success. The sequences used in this study were limited to fewer numbers and most of the sequences obtained were from India. An in-depth analysis involving more sequences from various other geographic locations will provide a clear understanding regarding the fitness of the organism. Further studies are warranted to correlate the effect of IS6110 transposition events to the successful adaptation of the pathogen.

Conclusion

In conclusion, our findings provide novel information about IS6110 insertions and the preferential insertion regions in the *M. orygis* genome. We have also elucidated the genomic impact of these insertions and their effect on various metabolic processes and pathways suggestive of their pertinence. Henceforward, as and when MTBC strains are being vastly sequenced, it is ideal to locate the IS6110 insertion sites, which will provide a better understanding regarding its biological role in the organism's adaptation, evolution, and fitness.

Author Contributions

Ahmed Kabir Refaya: Methodology, Software, Formal analysis, Investigation, Data curation, Writing – original draft, Writing – review & editing, Visualization. **Umashankar Vetrivel:** Investigation, Resources, Writing – review & editing. **Kannan Palaniyandi:** Conceptualization, Methodology, Formal analysis, Investigation, Resources, Writing – review & editing, Visualization, Supervision, Project administration, Funding acquisition. The authors have read and approved the manuscript.

Supplemental Material

Supplemental material for this article is available online.

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