A complete high-quality MinION nanopore assembly of an extensively drug-resistant *Mycobacterium tuberculosis* Beijing lineage strain identifies novel variation in repetitive PE/PPE gene regions

Arnold Bainomugisa, Tania Duarte, Evelyn Lavu, Sushil Pandey, Chris Coulter, Ben J. Marais and Lachlan M. Coin

**Abstract**

A better understanding of the genomic changes that facilitate the emergence and spread of drug-resistant *Mycobacterium tuberculosis* strains is currently required. Here, we report the use of the MinION nanopore sequencer (Oxford Nanopore Technologies) to sequence and assemble an extensively drug-resistant (XDR) isolate, which is part of a modern Beijing sub-lineage strain, prevalent in Western Province, Papua New Guinea. Using 238-fold coverage obtained from a single flow-cell, *de novo* assembly of nanopore reads resulted into one contiguous assembly with 99.92 % assembly accuracy. Incorporation of complementary short read sequences (Illumina) as part of consensus error correction resulted in a 4,404,064 bp genome with 99.98 % assembly accuracy. This assembly had an average nucleotide identity of 99.7 % relative to the reference genome, H37Rv. We assembled nearly all GC-rich repetitive PE/PPE family genes and identified variants within these genes. With an estimated genotypic error rate of 5.3 % from MinION data, we demonstrated identification of variants to include the conventional drug resistance mutations, and those that contribute to the resistance phenotype (efflux pumps/transpoter) and virulence. Reference-based alignment of the assembly allowed detection of deletions and insertions. MinION sequencing provided a fully annotated assembly of a transmissible XDR strain from an endemic setting and showed its utility to provide further understanding of genomic processes within *Mycobacterium tuberculosis*.

**DATA SUMMARY**


2. The assembled complete genome and its annotations are available in NCBI under accession number CP022704.2 (https://www.ncbi.nlm.nih.gov/sra/?term=CP022704.2).

**INTRODUCTION**

Globally, the tuberculosis (TB) incidence rate has shown a slow decline over the last two decades, although absolute case numbers continue to rise due to population growth, with an estimated 10.4 million new cases occurring in 2016 [1]. TB control gains are threatened by the growing number of drug-resistant strains recorded around the world [2]. An estimated 490,000 incident cases of multi-drug-resistant (MDR) TB, i.e. resistance to at least isoniazid and rifampicin, occurred in 2016 [1]. The incidence of extensively drug-resistant (XDR) strains, i.e. MDR strains with additional resistance to at least one fluoroquinolone and second-line injectable, is also on the rise [1]. Further multiplication of drug resistance in strains that are already highly drug-resistant could lead to programmatically incurable TB, where construction of a curative regimen might be impossible with existing treatment options [3, 4]. Management of drug-resistant TB places a major financial burden on health systems, which may be overwhelmed in settings with high disease burdens [3].

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**Keywords:** *Mycobacterium tuberculosis*; Oxford Nanopore Technologies.

**Abbreviations:** ANI, average nucleotide identity; MDR, multi-drug resistance; XDR, extensive drug resistance; PNG, Papua New Guinea; TB, tuberculosis; ONT, Oxford Nanopore Technologies; PE, proline-glutamate; PPE, proline-proline-glutamate; SMRT, small molecule real-time.

**Data statement:** All supporting data, code and protocols have been provided within the article or through supplementary data files. Six supplementary tables and five supplementary figures are available with the online version of this article.
In the absence of lateral gene transfer [5], drug resistance in *Mycobacterium tuberculosis* (MTB) arises mainly from chromosomal mutations that are selected by chemotherapeutic pressure, which drives drug resistance multiplication and the ongoing evolution of drug-resistant strains [6–8]. Successful transmission of drug-resistant strains results in clonal expansion and potential epidemic spread [9–11]. The acquisition of resistance-conferring mutations has potential to lead to epidemic spread if these drug-resistant strains are readily transmissible [12, 13]. The mechanisms underlying the development of highly transmissible XDR strains are not fully elucidated. One such mechanism is the induction of efflux pumps, which may lead to high-level resistance in mycobacteria [14], without any metabolic compromise. While previous studies described efflux pump genes and identified mutations in some of these genes [15, 16], efflux pumps in a transmissible XDR strain have not been described.

Whole genome sequencing using short reads has elucidated a large number of mutations associated with drug resistance, as well as compensatory mutations, but has limited capacity to resolve large structural variations, gene duplications or variations in repetitive regions [10, 17, 18]. Long-read sequencing could provide a more comprehensive understanding of the evolutionary mechanisms underlying the emergence of highly transmissible drug-resistant strains [17]. In principle, Oxford Nanopore Technologies MinION sequencing technology offers read lengths that are only limited by the length of DNA presented and produces data in real time [19]. The small size, ease of use and cheap unit cost of the MinION nanopore sequencer facilitates successful deployment in resource-limited settings, as has been achieved during the Ebola outbreak in West Africa [20]. Although the potential of MinION to detect drug resistance mutations in *M. tuberculosis* has been demonstrated [21], its application for complete MTB genome assembly has not been reported.

Papua New Guinea (PNG) has a high rate of drug-resistant TB in its Western Province [22]. We recently characterized a drug-resistant TB outbreak on Daru island, which is driven by a modern Beijing sub-lineage 2.2.1.1 strain [23]. Whilst some genetic markers within the strain have been identified [23], the molecular mechanisms responsible for pathogenesis and virulence are not fully elucidated. Genomic regions such as proline-glutamate (PE, 99 loci) and proline-proline-glutamate (PPE, 69 genes) are thought to contribute to virulence and pathogenicity of a strain [24, 25] and yet they are routinely excluded from data analysis after whole genome sequencing [9, 23, 26]. These gene families constitute signature motifs near the N terminus of their gene products and are sub-classified according to sequence features on the C terminus. PE genes are divided into PE_PGRS (polymorphic GC-rich sequence, 65 genes) and PE (no distinctive feature, 34 genes) while PPE genes are divided into PPE_MPTR (major polymorphic tandem repeats, 23 genes), PPE_SVP (Gxx-SVPxxW motif, 24 genes), PPE_PPW (PxxPxxW motif, 10 genes) and PPE (no distinctive feature, 12 genes) [24]. The existence of these subgroups highlights the diversity in their roles towards immune invasion and virulence [27].

Because of the repetitive nature of nucleotides and GC-rich content within the PE/PPE gene regions, they are a source of sequencing error from short-read sequencing such as Illumina. Previous studies have utilized Illumina sequencing to provide understanding of the role of these gene families but some of the reported results are variable and inconsistent with no complete assembly of these genes [27–29]. A study by Elghraoui and colleagues utilized small molecule real-time (SMRT) sequencing, which produces long reads, to assemble a complete genome of MTB H37Ra that included all the PE/PPE gene families [30]. However, this was performed on an attenuated strain, and there remains a need to use cheap long-read sequencing technology on drug-resistant strains to provide insight into the underlying mechanisms of drug resistance and identify key features associated with virulence and transmissibility. We utilized MinION to assemble a comprehensive genome of an XDR strain including variable and repetitive regions such as PE/PPE genes. The assembly will serve as an ideal reference for

**IMPACT STATEMENT**

We recently characterized a modern Beijing lineage strain of *Mycobacterium tuberculosis* responsible for the drug resistance outbreaks in Western Province, Papua New Guinea. As some of the genomic markers responsible for drug resistance and transmissibility among this strain remain unknown, there is a need to elucidate all molecular mechanisms that account for the resistance phenotype, virulence and transmission. Whole genome sequencing using short reads has been widely utilized to study the *M. tuberculosis* genome but it does not generally capture long repetitive regions as variants in these regions are eliminated during analysis. Illumina instruments are known to have a GC bias so that regions that are GC- or AT-rich are under-sampled and this effect is exacerbated in *Mycobacterium tuberculosis*, which has approximately 65 % GC content. In this study, we utilized Oxford Nanopore Technologies (ONT) MinION sequencing to assemble a high-quality complete genome of an extensively drug-resistant strain of a modern Beijing lineage. We were able to assemble all PE/PPE (proline-glutamate/proline-proline-glutamate) gene families that are GC-rich and repetitive in nature. We show the utility of ONT in offering a more comprehensive understanding of the genetic mechanisms that contribute to resistance, virulence and transmission. This is important for setting up predictive analytics platforms and services to support diagnostics and treatment in endemic settings with drug-resistant tuberculosis.
ongoing MDR/XDR outbreak surveillance in Western Province, PNG, and the far north of Queensland.

METHODS

Phenotypic susceptibility testing

We selected an XDR isolate originating from the Western Province of PNG, which we refer to as the WP-XDR strain. This strain was tested for resistance to first- and second-line drugs including: rifampicin (1.0 µg ml⁻¹), isoniazid (0.1 µg ml⁻¹, low-level; and 0.4 µg ml⁻¹, high-level), streptomycin (1.0 µg ml⁻¹), ethambutol (5.0 µg ml⁻¹), pyrazinamide (100 µg ml⁻¹) and second-line drugs amikacin (1.0 µg ml⁻¹), capreomycin (2.5 µg ml⁻¹), kanamycin (2.5 µg ml⁻¹), ethionamide (5.0 µg ml⁻¹), ofloxacin (2.0 µg ml⁻¹), p-aminosalicylic acid (4.0 µg ml⁻¹) and cycloserine (50 µg ml⁻¹). The automated Bactec Mycobacterial Growth Indicator Tube (MGIT) 960 system (Becton Dickinson) was used for susceptibility testing offirst- and second-line drugs. A minimal inhibitory concentration assay for second-line drugs was performed using the Sensititre (TREK diagnostic system) system.

DNA extraction and quantification

DNA was extracted from Lowenstein–Jensen (LI) slopes cultured for 4 weeks at 37 °C. DNA isolation was performed using mechanical and chemical methods. Briefly, five glass beads (0.7 mm; Sigma) were added to 200 µl of PrepMan using mechanical and chemical methods. Briefly, five glass beads (0.7 mm; Sigma) were added to 200 µl of PrepMan using mechanical and chemical methods. DNA was extracted from Lowenstein–Jensen (LI) slopes cultured for 4 weeks at 37 °C. DNA isolation was performed using mechanical and chemical methods. Briefly, five glass beads (0.7 mm; Sigma) were added to 200 µl of PrepMan using mechanical and chemical methods. The solution was dry heat incubated for 10 min and re-suspended with 40 µl of nuclease-free water. The DNA was quantified using a Nanodrop 8000 and a Qubit dsDNA HS assay kit (both from Thermo Fisher Scientific).

MinION library preparation and sequencing

DNA was purified using 0.4 × Agencourt AMPure XP beads (Beckman Coulter) and fragment distribution size was assessed using an Agilent 4200 TapeStation (Agilent). Preparation of a one-dimensional genomic DNA library was performed using the SQQ-LSK108 system (Oxford Nanopore Technologies, Oxford, UK) according to the manufacturer’s instructions. We performed dA-tailing and end-repair using the NEBNext Ultra II End-repair/dA-tail module with two step incubation periods, of 20 min each. A purification step using 0.7 × Agencourt AMPure XP beads (Beckman Coulter) was then performed according to the manufacturer’s instructions. A ligation step was performed using New England Biolabs Blunt/ligase master mix module according to the manufacturer’s instructions and the reaction was incubated at room temperature for 20 min. Adaptor-ligated DNA was purified using 0.4 × Agencourt AMPure XP beads (Beckman Coulter) following the manufacturer’s instructions but using Oxford Nanopore supplied buffers (adaptor bead binding and elution buffers). The library was then ready for MinION sequencing.

With the MinION MK1B device connected to the computer via a USB3, MinKNOW software (v.1.4.3) was started to perform quality control checks on pore activity and equilibrate the flow cell (FLOW-N106, version R9.4). The library was combined with reagents supplied by Oxford Nanopore and loaded onto the flow cell following the manufacturer’s instructions, choosing a 48 h sequencing procedure. Illumina data for the strain were available from our previous study [23].

MinION and Illumina data analysis

Raw files generated by MinKNOW were base called using Albacore (v2.0) to return Oxford Nanopore Technologies (ONT) fastq files. De novo genome assembly was performed using Canu [31] and the assembly was improved using nanopilish (methylation aware option) [32]. The sequence depth of coverage of ONT reads versus the assembly was then assessed using SAMTools [33]. The assembly was further improved through two different error correction steps: Racon (utilizing raw ONT reads) and Pilon (utilizing Illumina reads) [34]. CheckM was used to assess the quality (completeness and heterogeneity) of the assemblies [35]. To determine the accuracy of the assemblies, Illumina reads of the study strain were first de novo assembled using SPAdes 3.11.1 [36]. The resultant Illumina assembly was used to determine the accuracy of ONT assemblies by using statistics from dnaidiff (MUMmer version 3) [37]. The assembly with the highest quality was circularized using Circulator v1.5.1 [38], and similarity with the reference genome H37Rv (NC_000962.3) was calculated using nucmer (MUMmer version 3) [37] and an average nucleotide identity (ANI) calculator [39]. Genome annotation was performed using the NCBI pipeline [40] and circular representation of the genome was viewed using Circos [41]. To evaluate the assembly of PE/PPE gene families, all the PE/PPE gene sequences within MTB H37Rv were searched from Mycobrowser [42] and used to create a customized PE/PPE database. A BLASTN search of the assembly against the database was performed. Percentage breadth of coverage and read depth for PE/PPE gene families from ONT and Illumina reads was assessed using DepthOfCoverage (GATK version 3.7) [43] after mapping to H37Rv using BWA-MEM [44]. The resultant depths were plotted using ggplot (R statistical package). Biodiff (https://www.gitlab.com/LPCDRP/biodiff) was used to call structural variants from the assembly against H37Rv as a reference and as a query. Structural variants were further evaluated by mapping Illumina raw data of lineage-representative genomes including Indo-Oceanic, East-Asian (including Beijing
sequence depth of 238 read length of 5073 bp, longest read 330 509 bp and average XDR strain

Analysis of MinION-derived assembly of the WP-XDR strain

In total, 373952 ONT reads passed base calling with N50 read length of 5073 bp, longest read 330 509 bp and average sequence depth of 238× (Fig. 1). The size of the assembled genome varied among the different consensus tools used: Racon, 4 404 947 bp; Pilon, 4 405 333 bp; and Pilon plus Racon, 4 404 064 bp. Use of raw ONT reads for error correction of the assembly resulted in a 99.92 % accurateassembly (Racon), while use of Illumina reads improved it further to 99.97 % (Pilon). According to CheckM results, the assembly from a combination of Pilon plus Racon tools had better quality scores (Table 1), and we referred to this as WP-XDR assembly. The single contig from this assembly of 4 404 064 bp had a GC content of 65.5 % and had an ANI score of 99.7 % to H37Rv (Fig. S2). NCBI annotation of the genome yielded a total of 3697 coding DNA sequences (CDS), 45 tRNAs, three rRNAs (5S, 16S, 23S) and three non-coding RNAs (Fig. 2). A BLASTN analysis of a PE/PPE customized database against the assembly genome showed 166/168 PE/PPE genes had >90 % nucleotide sequence similarity at an E-score ≤10^{-6} (Fig. S3). Two genes, wag22 and PE_PGRS57, had nucleotide sequence similarity of 58.8 and 78 %, respectively. Evaluation of PE/PPE genes when ONT reads when mapped to H37Rv revealed nearly all PE/PPE genes (166/168) had 100 % breadth of coverage at an average read depth of 299.87 (IQR 285.91–311.36) (Fig. S3). Again wag22 and PE_PGRS57 had breadth of coverage of 26 and 62 %, respectively. From Illumina reads of the study strain, only 54.3 % (92/168) of the PE/PPE genes had 100 % breadth of coverage at an average sequence depth of 46.3 (Fig. S4) with wag22 and PE_PGRS57 having zero coverage.

Genotypic error analysis using MinION reads

We assessed the accuracy of MinION reads to generate accurate SNP calls in the non-repetitive part of the reference genome by excluding repetitive PE/PPE regions. A total of 1254 SNPs and 122 small indels were called from ONT reads while 1098 SNPs and 105 small indels were called from Illumina reads when mapped to H37Rv. Of these, 1095 SNPs (574 non-synonymous and 402 synonymous) and 87 small indels concurred over the technologies. In total, 118 of 159 SNPs and 23 of 35 small indels identified from ONT but not in Illumina reads were from regions of low Illumina coverage (<30× coverage). The remaining 41 SNPs (3.2 %, 41/1254) and 12 small indels (9.8 %, 12/122) were identified in ONT but missing from Illumina (high coverage regions). These might potentially be due to systematic base-calling errors that could not be corrected via consensus calling and polishing. Three (0.2 %) SNPs and 18 (14.7 %) indels were identified by Illumina but not by MinION sequencing. Considering all 194 ONT-only calls (159 SNPs and 35 indels) as false positives and 21 Illumina-only calls (three SNPs and 18 indels) as false negatives, the genotypic error rate from all ONT calls was estimated at 15.6 % [215/(1254+122)*100]. If we ignore inconsistent calls where Illumina coverage was low then the estimated genotypic error rate for base calling from ONT was reduced to 5.3 % [74/(1254+122)*100]. Counts of SNP calling among the non-consensus SNPs from ONT reads did not show any bias to a particular base (Table S2).

ONT and Illumina reads were assessed for SNP identification in PE/PPE gene families. From ONT reads, 158 SNPs were identified from 70 PE/PPE genes (42 PE and 28 PPE) with 88 SNPs (55.6 %) identified from the PE_PGRS subfamily (Table S3). From the Illumina reads, 124 SNPs from

![Fig. 1. Bar graph showing the unimodal sequence length distribution from ONT.](image)
45 PE/PPE genes (25 PE and 20 PPE) were identified, of which 31 SNPs (25%) were from the PE_PGRS sub-family. There were 81 SNPs (from 42 PE/PPE genes) overlapping between ONT and Illumina with PPE54 having the highest number of overlapped SNPs [9] identified within one gene.

**Mutations in known resistance genes, efflux pumps and virulent genes**

Phenotypic drug susceptibility results revealed the WP-XDR isolate to be extensively drug-resistant with susceptibility to only amikacin, kanamycin, \( p \)-aminosalicyclic acid and cycloserine. Table 2 provides details on phenotypic resistance, as well as mutations in genes known to confer drug resistance to first- and second-line drugs and recognized compensatory mutations. Genotypic drug resistance profiles concurred with phenotypic results. Ten mutations were identified in seven genes that encode transmembrane efflux pumps and transporter proteins (Table 3). Table S4 shows 16 SNPs identified in genes that encode virulence proteins; eight (50%) were from the mce gene family, and a mutation within mycP1 (p.Thr238Ala) was also noted. In addition, 27 SNPs were identified in three gene families involved in cell wall synthesis, with 17 in fadD, four in pks and three in mmp gene families (Table S5).

**Structural variants identified from assembly of the WP-XDR strain**

Two reference-based approaches were utilized using biodiff script. First, using H37Rv as a reference to the assembly, we identified known regions of difference for the Beijing sub-lineage [50] and three unknown deletions (Table 4). On evaluating the three unknown deletions using Illumina data of lineage-representative genomes mapped to the reference, Sniffles identified the three unknown deletions still missing among Beijing lineage-representative genomes. Secondly, using the assembly as a reference to H37Rv, we identified two insertions within the WP-XDR assembly of size 4490 bp (2 207 139–2 211 629) and 390 bp (3 488 211–3 488 601). On assessing Illumina data of lineage-representative genomes mapped on the WP-XDR assembly using Sniffles, the 4490 bp insertion with 64.6% GC content was identified among Indo-Oceanic, ancient Beijing, modern Beijing and East-African-Indian lineages but absent from the Euro-American lineage (Fig. 3). As in previous studies [51–53], BLASTN and annotation results of the 4490 bp region was revealed to span seven complete genes that encode proteins that include NADP-dependent oxidoreductase, iron-regulated elongation factor (Tu), PE-family protein and four genes that encode uncharacterized proteins. The second smaller insertion, 390 bp with 66.7% GC content (3 488 211–3 488 601), was identified as being unique among the Beijing lineage but varied in size as it was 835 bp with respect to Indo-Oceanic and East-African-Indian lineages (Fig. 3). Review of the NCBI annotation of the assembly indicated the insertion to be part of a 654 bp gene (CBG40_17120; 3 487 881–3 488 534) that spans 323 bp (3 488 211–3 488 534). A BLAST search of the insertion sequence revealed 50% sequence identity to four M.

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**Table 1.** CheckM quality evaluation and assembly accuracy of draft genomes obtained from the three consensus tools, relative to the SPAdes assembly

<table>
<thead>
<tr>
<th>Tool</th>
<th>Genome size (bp)</th>
<th>Completeness (%)</th>
<th>Contamination</th>
<th>Strain heterogeneity</th>
<th>No. of indels</th>
<th>No. of SNPs</th>
<th>Total errors</th>
<th>Accuracy (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Racon</td>
<td>4 405 333</td>
<td>83.2</td>
<td>0.00</td>
<td>0.00</td>
<td>2421</td>
<td>1026</td>
<td>3447</td>
<td>99.92</td>
</tr>
<tr>
<td>Pilon</td>
<td>4 404 947</td>
<td>97.74</td>
<td>0.00</td>
<td>0.00</td>
<td>1048</td>
<td>236</td>
<td>1284</td>
<td>99.97</td>
</tr>
<tr>
<td>Pilon+Racon</td>
<td>4 404 064</td>
<td>100</td>
<td>0.00</td>
<td>0.00</td>
<td>807</td>
<td>217</td>
<td>1024</td>
<td>99.98</td>
</tr>
</tbody>
</table>

**Fig. 2.** Circular representation of a WP-XDR genome (modern Beijing sub-lineage 2.2.1.1 strain) with gene annotations. From the inner to outer ring: blue, GC content of coding sequence; scattered grey, SNPs relative to H37Rv; grey bars, reverse and forward rRNA; green, reverse coding sequences; red, forward coding sequences; yellow bars, unique regions compared to H37Rv; grey outside ring, assembled contig. Mb, million base pairs.
Table 2. Mutations in candidate drug resistance genes identified from the WP-XDR assembly

<table>
<thead>
<tr>
<th>Drug</th>
<th>In vitro phenotype</th>
<th>Investigated genes</th>
<th>Genes (mutation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rifampicin</td>
<td>Resistant</td>
<td>rpoB, rpoC, rpoA, rpoD</td>
<td>rpoB (p.Ser450Leu) rpoC (p.Val483Gly)</td>
</tr>
<tr>
<td>Pyrazinamide</td>
<td>Resistant</td>
<td>pncA, rpsA, panD</td>
<td>pncA (p.Tyr103Asp) rpsL (p.Lys43Arg) gldB (p.Leu91Pro)</td>
</tr>
<tr>
<td>Streptomycin</td>
<td>Resistant</td>
<td>rpsL, gldB, rrs</td>
<td></td>
</tr>
<tr>
<td>Ethionamide</td>
<td>Resistant</td>
<td>fabG1-inhA, ethA, ethR</td>
<td>fabG1-inhA (C-1ST) gryA (p.Asp94Gly) Nil</td>
</tr>
<tr>
<td>Fluoroquinolone</td>
<td>Resistant</td>
<td>gyrA, gyrB</td>
<td></td>
</tr>
<tr>
<td>Amikacin</td>
<td>Susceptible</td>
<td>rrs, whiB7, gidB</td>
<td>Nil</td>
</tr>
<tr>
<td>Kanamycin</td>
<td>Susceptible</td>
<td>Rv2417c-eis, whiB7, rrs, gidB</td>
<td>Nil</td>
</tr>
<tr>
<td>Capreomycin</td>
<td>Resistant</td>
<td>tlyA, whiB7, rrs, gidB</td>
<td>tlyA (ins397C)</td>
</tr>
<tr>
<td>PAS</td>
<td>Susceptible</td>
<td>ribD, thyA, dfuA, folC</td>
<td>Nil</td>
</tr>
<tr>
<td>Cycloserine</td>
<td>Susceptible</td>
<td>abs, ddl, cycA</td>
<td>Nil</td>
</tr>
</tbody>
</table>

PAS, p-aminosalicylic acid.

tuberculosis H37Rv genomes (E-score=1.1E-175) and 100% sequence identity to 55 M. tuberculosis Lineage 2 genomes (E-score=0) (Table S6). Phyre2 protein modelling [48] of the 390 bp insertion sequence predicted it to be a PPE-like protein with PE8-PPE15 protein used as the best template (79% sequence modelled, 100% confidence) consisting of 73% alpha helices (Fig. S5).

**DISCUSSION**

Long-read sequencing increases the potential for improved understanding of genomic factors that contribute to the M. tuberculosis phenotype because it has inherent power to assemble the entire strain genome. We have shown that a WP-XDR strain can be de novo assembled into one contig using 238× read depth data from one flow-cell of the ONT platform. These results highlighted some of the advantages of using ONT. We were able to resolve the GC-rich and highly repetitive PE/PPE gene families which were poorly resolved when Illumina reads were used. Of the two PE_PGRS subfamily genes (PE_PGRS57 and wag22) that had poor assembly, wag22 has previously been shown to be deleted from Beijing lineage strains [50]. There were twice as many SNPs in PE_PGRS genes based on ONT over Illumina sequencing (Table S2). PE_PGRS gene mutations were under-represented in short-read sequencing, possibly due to their extra GC-rich motifs that could impact on sequencing. Previous studies have identified a greater number of mutations within the PE_PGRS subfamily compared to other PE subfamilies, and attribute this to their involvement in antigenic variation and immune evasion from exposure to the host immune system [27, 58]. PE_PGRS54, which is a member of a larger subfamily, PPE_MPTR (major polymorphic tandem repeat) known to have more amino-acid variations at the C-terminal site [59], had the highest number of mutations (Table S6). Previous studies have identified this gene to be involved in the arrest of phagosome maturation, allowing survival of the bacteria in the macrophages as a result of its long amino-acid length at the C terminus [25, 60]. It has been postulated that PPE54 gene mutations may using long reads in conjunction with Illumina reads to improve on the assembly [54, 55]. Although other long-read sequencing technologies such as SMRT have demonstrated their value in producing high-quality M. tuberculosis genome assemblies [30, 56], there is ongoing research aimed to improve the quality of reads produced by ONT platforms. However, ONT platforms have been shown to be cheaper and user-friendly [57], indicating its future value in TB burdened settings such as Western Province, PNG.

**Table 3.** Mutations in putative efflux pump/transporter genes identified from the WP-XDR assembly

<table>
<thead>
<tr>
<th>Genes investigated</th>
<th>Gene (mutation) identified</th>
</tr>
</thead>
</table>
also play a role in development of isoniazid, rifampicin and ethambutol resistance [61], but we were unable to verify this given the presence of well-characterized drug resistance mutations.

Another advantage of ONT was the resolution of structural variants which are essential in understanding strain-related phenotypes [62]. The identification of the three deleted gene regions and the 4490 bp insertion within the assembly show the limitation of H37Rv when used as a reference genome in genomic studies [51–53]. This could impact on the clear understanding of pathogenicity, virulence and transmissibility. The WP-XDR assembly has potential to be utilized as a reference genome for future investigation of XDR outbreaks in PNG. The second smaller insertion (390 bp) within the assembly had variable sizes among the reference genomes, suggesting independent structural rearrangement among the different lineages. The applied comparative approach highlights the opportunity to study lineage- and strain-specific differences, especially with the advent of long-read sequencing, which can provide complete assemblies.

Although ONT had a greater number of differing variants when compared to Illumina, it had a good estimated genotypic error rate of 5.3 %, identifying all the known drug resistance and compensatory mutations in concordance with in vitro susceptibility testing. This was accompanied

<table>
<thead>
<tr>
<th>Start</th>
<th>Stop</th>
<th>Size (bp)</th>
<th>Region of difference</th>
<th>Genes deleted</th>
<th>RNA deleted</th>
</tr>
</thead>
<tbody>
<tr>
<td>79557</td>
<td>83036</td>
<td>3479</td>
<td>105</td>
<td>Rv0071, Rv0072, Rv0073</td>
<td>ncRv10071, ncRv10071c</td>
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<tr>
<td>1779278</td>
<td>1788514</td>
<td>9236</td>
<td>149</td>
<td>Rv1573, Rv1574, Rv1575, Rv1576c, Rv1577c, Rv1578c, Rv1579c, Rv1580c, Rv1581c, Rv1582c, Rv1583c, Rv1584c, Rv1585c, Rv1586c</td>
<td>Nil</td>
</tr>
<tr>
<td>1986636</td>
<td>1998623</td>
<td>11987</td>
<td>152</td>
<td>pkcD, Rv1576c, Rv1577c, cut1, Rv1759c, Rv1760, Rv1761c, Rv1762c, Rv1763, Rv1764, Rv1765c</td>
<td>Nil</td>
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<tr>
<td>2365414</td>
<td>2366769</td>
<td>1355</td>
<td>Unknown*</td>
<td>Rv2105, Rv2106</td>
<td>Nil</td>
</tr>
<tr>
<td>2535431</td>
<td>2536142</td>
<td>711</td>
<td>181</td>
<td>Rv2262c, Rv2263</td>
<td>Nil</td>
</tr>
<tr>
<td>3122553</td>
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<td>Unknown*</td>
<td>Rv3186, Rv3187</td>
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</table>

*Unknown deletions in the study strain.

Fig. 3. Integrative Genomic Viewer (IGV) of Illumina reads from different M. tuberculosis lineages (A, M. tuberculosis H37Rv; B, Indo-Oceanic; C, ancient Beijing; D, modern Beijing; E, East African Indian; F and G, Euro American) mapped on the ONT assembled genome highlighting the large insertions within the assembled genome. On the left: 4490 bp insertion (2 207 042–2 211 532) spanning seven annotated genes (blue checked arrows, NADH-dependent oxidoreductase, iron-regulated elongation factor-tu, PE family protein and hypothetical/uncharacterized proteins). On the right: 390 bp insertion relative to H37Rv (3 488 211–3 488 601) spanning 323 bp (checked) at the end of a 654 bp coding sequence. ONT, Oxford Nanopore Technologies.
by characterization of mutations within efflux pumps/transporter, biosynthesis pathway encoding genes (ABC, MFS, mmpL, pks and fadD) and virulence genes (mce genes, mtpc1) which are thought to contribute to the resistance phenotype and pathogenesis of the strain [63, 64]. Even with the ongoing improvement of ONT sequencing, we show that ONT data can confidently be utilized for further investigation of genomic variants that influence virulence, resistance and pathogenesis. Accompanied by the portability of ONT devices, this technology has potential for clinical utility as a point-of-care diagnostic tool because it generates reads that can be analysed in real time. This aspect was not explored in the current study but other studies have demonstrated the duration needed for speciation and identification of resistance-conferring genes [20, 65]. This is an ongoing area of research whose future application in the diagnostics of M. tuberculosis will greatly impact on improved management and control of infection.

Some of the limitations of the study included the use of one isolate to identify various structural variants and mutations which could be ‘background’ variations that may not impact on the strain phenotype. Use of a polished consensus assembly for analysis of structural variants could have affected detection of assembly breaks although a polishing step improved the quality of the assembly. Another limitation was the failure to confirm the presence of the identified structural variants using targeted sequencing, although the variants were consistent with Illumina data that had a high Phred score (>90).

In conclusion, we report the value of a portable small sequencer that can sequence an XDR strain, assemble a complete genome including GC-rich repetitive regions and analyse variants with high certainty. Future application of this technology in high-burden settings will have positive impacts on TB control and case management.

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Ethical statement
The isolate was selected from a previous study [23] with ethics clearance from the University of Queensland and the Papua New Guinea Medical Research Advisory Committee to perform detailed whole genome sequencing.

Conflicts of interest
The authors declare that there was no conflict of interest.

Data bibliography
1. NCBI project accession number PRJNA386696 (2018).

References


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