Proteomic analysis of drug-susceptible and multidrug-resistant nonreplicating Beijing strains of *Mycobacterium tuberculosis* cultured in vitro

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**ABSTRACT**

The existence of latent tuberculosis infection (LTBI) is one of the main obstacles hindering eradication of tuberculosis (TB). To better understand molecular mechanisms and explore biomarkers for the pathogen during LTBI, we cultured strains of *Mycobacterium tuberculosis* (Mtb) under stress conditions, mimicking those in the host granuloma intracellular environment, to induce entry into the non-replicating persistence stage. The stresses included hypoxia, low pH (5.0), iron deprivation (100 μM of 2',2'-dipyridyl) and nutrient starvation (10% M7H9 medium). Three Mtb strains were studied: two clinical isolates (drug-susceptible Beijing (BJ) and multidrug-resistant Beijing (MDR-BJ) strains) and the reference laboratory strain, H37Rv. We investigated the proteomics profiles of these strains cultured in stressful conditions and then validated the findings by transcriptional analysis. NarJ (respiratory nitrate reductase delta chain) was significantly up-regulated at the protein level and the mRNA level in all three Mtb strains. The narJ gene is a member of the narGHJI operon encoding all nitrate reductase subunits, which play a role in nitrate metabolism during the adaptation of Mtb to stressful intracellular environments and the subsequent establishment of latent TB. The identification of up-regulated mRNAs and proteins of Mtb under stress conditions could assist development of biomarkers, drug targets and vaccine antigens.

**1. Introduction**

Tuberculosis (TB), caused by *Mycobacterium tuberculosis* (Mtb), is a major global health problem and one of the leading causes of death worldwide [1–3]. Currently, the major obstacle to control and eradication of TB is the ability of Mtb to persist as a life-long infection in humans in a dormant or non-replicating state (NRP) [3,4]. This condition is asymptomatic and is known as latent TB infection (LTBI) [4]. In a small proportion of LTBI cases (5–10%), the pathogen might be reactivated, leading to active TB [1]. Humans are the only reservoir of this pathogen, which needs to be considered for TB control [5].

Of particular concern are strains belonging to the East Asian or Beijing (BJ) lineages of Mtb. These have caused numerous outbreaks worldwide [6]. The hyper-virulent BJ lineage has been emerging throughout the world associated with disease outbreaks and antibiotic resistance [7,8]. Multidrug resistant BJ strains (MDR-BJ) show resistance to several current anti-tuberculosis drugs.

Metabolic processes play important roles in the pathogenesis of Mtb [9]. However, the metabolic processes that occur on entering dormancy, leading to survival and drug resistance in the host, are poorly understood [9]. Multiple-stress conditions within the host, such as oxygen depletion and immune responses, induce mycobacteria to enter a dormancy stage during which they are phenotypically drug resistant. Study of gene regulation and protein expression of Mtb may clarify the processes by which Mtb achieves dormancy. This adaptation may produce more virulent phenotypes of Mtb that can survive within granulomas of host [4,10].

*Mycobacterium tuberculosis* H37Rv is the most studied strain of TB in research laboratories [11]. However, a laboratory strain might not exhibit the same virulence properties as clinical strains [4]. To better

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understand the NRP stage of *Mtb*, in vitro models have been designed to mimic intracellular stress conditions. In this study, we aimed to mimic multiple stresses in vitro by combining conditions of hypoxia, acidic pH, iron deprivation and nutrient starvation. These conditions induce *Mtb* to enter the NRP stage. We investigated the protein expression patterns (using proteomic approaches and mRNA quantitative analysis) of *Mtb* reference strain H37Rv, and two clinical strains, BJ and MDR-BJ, cultured in this way.

2. Materials and methods

2.1. Bacterial strains, media and growth conditions

*Mycobacterium tuberculosis* H37Rv laboratory strain (NCBI:txid1773) and two clinical isolates, drug-susceptible Beijing (BJ) (NCBI: txid634955) and multidrug-resistant Beijing (MDR-BJ), were used. The strains used in this study cannot be linked to patient information, a condition approved by the Khon Kaen University Ethics Committee (No. HE621448). Three strains of *Mtb* were cultured in 20 ml of Middlebrook 7H9 (M7H9) medium (Sigma) and supplemented with 0.2% of glycerol and 10% of BBL™ Middlebrook OADC Enrichment (BD, US). Bacterial cells were grown for 7 days with shaking at 37 °C. The number of cultivable cells was estimated by plate count technique.

To mimic the multiple stress conditions, these media were prepared as follows: M7H9 medium with supplements was diluted to 10% with sterile water. The iron chelator was added to 100 μM, methylene blue was added to 1.5 mg/ml, 36% HCl was added to adjust the pH to 5.0. Log-phase *Mtb* cells, 10^5 CFU/ml were then inoculated into 20 ml of multiple stress media with parafilm-sealed test tubes, which were incubated at 37 °C without shaking. The decolorization of methylene blue was used to indicate oxygen depletion (3–4 weeks), which causes *Mtb* cells to enter the non-replicating/dormant stage after 4 weeks. Log-phase *Mtb* cells, 10^5 CFU/ml were then inoculated into 20 ml of control (M7H9 media) tubes were not sealed and incubated at 37 °C with shaking for 4 weeks.

2.2. Protein preparation, in-gel digestion and LC–MS/MS

Bacterial cell pellets were harvested by centrifugation after 4 weeks of incubation. Protein was then isolated from the cells by using lysis buffer (0.5 M Na_2HPO_4, 5 M NaCl, 1 M imidazole, 100 mg/ml lysozyme, 1X protease inhibitor cocktail (Amresco, USA), 1 M dithiothreitol) with 0.1 mm zirconia/silica beads (BioSpec Products, Inc.). Protein concentrations were determined by the Bradford protein assay, using Bradford reagent (Bio-Rad Laboratories, Inc.). To separate proteins, 5 μg of total proteins of each sample were electrophoresed through 12.5% SDS-PAGE. Gels were then cut, and in-gel digestion was performed.

To investigate the protein expression patterns of *Mtb*, liquid chromatography tandem mass spectrometry (LC–MS/MS) analysis and associated bioinformatics analysis were performed as previously described [12]. The detailed methods of this part are shown in supplementary materials and methods.

2.3. RNA extraction and cDNA synthesis

Bacterial cells were harvested by centrifugation after 4 weeks of incubation. RNA was isolated using Trizol reagent (Invitrogen, USA) with silica beads, according to the manufacturer’s instructions. Five hundred nanograms of RNA was treated with DNase I (Invitrogen, USA), to remove genomic DNA, according to the manufacturer’s instructions. cDNA was then synthesized using SuperScript III Reverse Transcriptase (Invitrogen, USA) according to the manufacturer’s instructions.

2.4. Quantitative real time PCR (qRT-PCR)

*Mycobacterium tuberculosis* genes that were differentially expressed between the active stage and NRP stage, including *narJ*, *tcrY*, *uvrC*, *Rv1356c* and *Rv3134c*, were selected for further study. The primer sequences used in this study are listed in Table 1. The expression of *Mtb* genes was examined by qRT-PCR. The PCR reaction was performed on a real-time PCR instrument (Applied Biosystems QuantStudio 6 Flex Real-Time PCR System) using SsoFast™ EvaGreen® Supermix (Bio-Rad Laboratory, Inc., USA). The relative expression of genes was examined by 2^ΔΔCt_. The 16S rRNA gene was used as an internal control.

2.5. Construction of protein-protein interaction networks

STRING (search tool for the retrieval of interacting genes/proteins) software was used to construct interaction networks of protein. Predicted functional partners of protein by STRING version 11.0 (https://string-db.org/) with high confidence interaction score (0.700).

2.6. Statistical analysis

GraphPad prism 5.0 was used for all data analysis. All experiments were done in triplicate and values are expressed as mean ± SD. Student’s t-test was used to analyze the difference among two groups. Statistically significant differences between groups are indicated by *p < 0.05; **p < 0.01; ***p < 0.001.

3. Results

3.1. Proteins detected only during the NRP stage or the active stage

To identify proteins found only in *Mtb* cultures under multiple stress conditions (i.e. during the NRP stage), proteomic profiles of the NRP and active stages were compared. During the NRP stage of H37Rv, BJ and MDR-BJ strains, 1,946, 1849 and 1947 proteins were detected, respectively. Of these, 220, 52 and 197 proteins were found only in the NRP stage of H37Rv, BJ and MDR-BJ strains, respectively (Fig. 1A–C). The details and complete lists of these proteins and their relative quantities are given in Table S1.

The same analyses identified 1,750, 1891 and 1799 proteins expressed by H37Rv, BJ and MDR-BJ strains, respectively, during growth in culture. There were 24, 94 and 49 proteins unique to the active stage of H37Rv, BJ and MDR-BJ strains, respectively (Fig. 1A–C).

3.2. Common unique proteins in NRP stage

Ten proteins were found only in the NRP stage in all three *Mtb* strains (Fig. 2). Further analysis revealed only one protein, *narJ*, that was characterized in the reference strains, whereas another 9 proteins were classified as non-characterized proteins. Fifty-seven, 33 and 35 proteins were unique to H37Rv, BJ and MDR-BJ strains, respectively during the

Table 1

| Gene | Direction/position of primer | Sequence |
|------|-----------------------------|----------|
| 16S rRNA | Forward, 469–491 | 5′-TTGACCGGTAGGGTAGGGAAGAAGC-3′ |
|       | Reverse, 909–888 | 5′-CCGTATGTTTTTAGCCTGGG-3′ |
| narJ  | Forward, 439–512 | 5′-CCTTATGACACACGGTGCC-3′ |
|       | Reverse, 583–586 | 5′-GGGAGCGTCAAGGTTAAAGG-3′ |
| tcrY  | Forward, 796–815 | 5′-CGAGGAGGCTTGTAGCTGCC-3′ |
|       | Reverse, 930–911 | 5′-CGGAGAGGTTGTACCCGG-3′ |
| uvrC  | Forward, 114–133 | 5′-GAGGCTGCTACTGCTGCC-3′ |
|       | Reverse, 308–289 | 5′-GAATTCCGCGATACGCTACTCC-3′ |
| Rv1356c | Forward, 531–533 | 5′-CGGCCGCTGTCGCTGATT-3′ |
| Rv3134c | Forward, 472–491 | 5′-TCCAGGTATCGGTTGTTGTT-3′ |

* The 16S rRNA gene was used as a housekeeping gene to normalize the data. Primers from Huile et al. (2002) [13].
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3. Further analysis found only 1, 1 and 2 proteins, respectively, that were well-characterized in the database of the M. tuberculosis H37Rv reference strain and significantly up-regulated. These proteins were Rv3134c (in the H37Rv strain), tcrY (BJ strain) and Rv1356c and uvrC (both in the MDR-BJ strain). The details and complete list of proteins in the three Mtb strains, and their relative quantities are given in Table 2 and Table S2.

3.3. Validation of proteomic analysis by quantitative real-time PCR

The shared and strain-specific proteins found by LC-MS/MS analysis to be over-expressed during the NRP stage in the three different Mtb strains were then further validated by measuring mRNA expression levels using qRT-PCR. The relative mRNA expression level of narJ was up-regulated in all strains. The fold-changes of expression level in H37Rv, BJ and MDR-BJ strains were 2.397 (p > 0.05), 51.536 (p < 0.01) and 16.661 (p < 0.01), respectively. These results were correlated with relative intensities (log2-fold) of proteins analyzed by LC-MS/MS. The intensity (log2-fold) of narJ in H37Rv, BJ and MDR-BJ strains were 17.953 (p < 0.001), 18.730 (p < 0.001) and 14.614 (p < 0.001), respectively (Fig. 3). In addition, the relative mRNA expression levels of Rv3134c, tcrY, Rv1356c and uvrC were significant up-regulated, with fold-changes of 3.344, 2.057, 11.735 and 10.941, respectively (Fig. 4). The results of qRT-PCR for all proteins confirmed the results of the LC-MS/MS.

4. Discussion

There have been few comparisons of proteomics profiles of BJ and MDR-BJ strains in the same experiment. These strains are very similar, and...
differing only by several mutations associated with drug resistance. However, our proteomic profiles revealed a large number of strain-specific proteins—38 (33 + 5) in BJ and 183 (35 + 148) in MDR-BJ (Fig. 2). This suggests that the multidrug-resistant phenotype might be associated with a large number of proteins.

The narJ protein was found only in the NRP stage of BJ and MDR-BJ. This protein is a subunit of nitrate reductase enzyme which bacteria use nitrate as a final electron acceptor instead of oxygen [14]. Nitrate reductase activity occurs at a low level during an aerobic growth of Mtb and significantly increases during the NRP stage under hypoxia [14]. Nitrate reductase activity of Mtb is also correlated with the virulence and is also associated with its growth under anaerobic conditions [15]. NarJ is a promising biomarker for the dormancy stage, especially in the BJ strain.

STRING analysis of narJ, the respiratory nitrate reductase delta chain, revealed the protein association networks involving this gene, including proteins encoded in narGHJ operon family. The narG/Rv1161, narH/Rv1162 and narI/Rv1164 genes encoded respiratory nitrate reductase alpha, beta and gamma chains, respectively. These proteins interact with each other at the same time and place. The other proteins shown in Fig. S1 with high and significant score of association with narJ included narX/Rv1736c, nitrate reductase, narK2/Rv1737c, nitrate/nitrite transporter, narU/Rv0267, integral membrane nitrate extrusion protein (nitrite facilitator), nirD/Rv0253, nitrite reductase [NAD(P)H] small subunit, tybA/Rv1165, GTP-binding translation elongation factor TypA (tyrosine phosphorylated protein A) (GTP-binding protein) and mrdA1/Rv0994, molybdopterin biosynthesis protein. Network nodes represent proteins; spliced isoforms or post-translational modifications are collapsed, i.e. each node represents all the proteins produced by a single protein-coding gene locus. Therefore, Mtb nitrate reductase is also the genes encoded narGHJ operon and is also associated with narK 1, narK2, narK3, narL, narX, and narU [15].

The NarGHJ operon encodes nitrate reductase [14]. To survive in a stress condition, narK2/Rv1737c is up-regulated during anaerobic conditions [15], which allows the transport of nitrate into and nitrite out of the cell [16]. These transportations may generate the ATP, which is necessary to Mtb survival in the absence of oxygen as a terminal electron acceptor [16]. Respiratory reduction of nitrate through NarGHJ operon could provide energy for the latent stage survival of the Mtb [16].

Nitrate reductase within phagosomes in macrophages is due to the anaerobic environment there [14]. The composition of nitrate reductase operon family in mycobacterial: the narG and narH bind to the plasma membrane of phagosome via the interaction between a hydrophobic patch of narH and narL which are bound to the cell membrane [15]. NarJ is a specific ligand that recognizes and binds to narG, and forms a complex with narGH to facilitate narJ, [4Fe–4S] cluster, and molybdenum cofactor successively. NarJ dissociates from the complex before interacting with membrane-binding narL. Nitrate passes through the cell wall into the cytoplasm via transmembrane protein narK2. It is reduced via narG and nitrite is released outside the macrophage via an unknown exporter [15]. Therefore, nitrate can be used as an alternative nitrogen source when the availability of other nitrogen sources is limited [16]. Further studies are needed concerning the availability and capability for utilization of other nitrogen sources during the latent infection stage. Such studies would provide a deeper insight into the possible link between dormancy and nitrogen assimilation and assist in identifying future drug targets.

Rv3134c, found only in H37Rv strain during the NRP stage, was annotated as a universal stress protein which could play an important role in adaptation to hypoxia and its up-regulation by at least 10-fold [17]. Moreover, its participating in the phosphorylation in the two-component regulatory system devRS [17]. A member of the dormancy Rv3132c/Rv3133c/Rv3134c regulon, it was induced in response to hypoxia, a strain lacking Rv3134c gene does not induce most genes of the dormancy regulon, the hypoxic regulation of hspX was eliminated. These results suggest a possible role for Rv3132c/Rv3133c/Rv3134c in latent stage of Mtb [17,18]. The regulon might give insights into the dormant or NRP stage of Mtb infection [17,18]. Rv3134c was significantly correlated with the expected six proteins including devR/Rv3133c, devS/Rv3132c, dorT/Rv2027c, Rv0079, hspX/Rv2031c and Rv1738. The network of direct interconnections between Rv3134c and other proteins (STRING version 11.0) with a high confidence score (0.700) is shown in Fig. S2.

TcrY, a protein seen only in the BJ strain during the NRP stage, was annotated as a two-component sensor kinase. This is associated with a two-component regulatory signal-transduction system, tcrXY, which activates phosphorylation [19,20]. Two component signal transduction systems, its process the signals resulting from the stress environment which developed by bacterium [20]. M. tuberculosis H37Rv cells lacking the tcrXY regulon show an increased virulence with significantly shorter survival times in SCID mice [19,20]. The network of direct high-confidence scoring (0.700) interconnections between TcrY and other proteins, as analyzed by STRING version 11.0, is shown in Fig. S3.

MDR-BJ strain-specific proteins during the NRP stage were Rv1356c and uvrC. These were annotated as a hypothetical protein and excinuclease ABC (subunit C - nuclease), respectively. Rv1356c is significantly up-regulated after 24 h under nutrient starvation in the H37Rv strain [21]. UvrC is a DNA-repair enzyme that catalyzes the excision of UV-damaged nucleotide segments producing oligomers having the modified base(s) [22]. In fact, Mtb is exposed to a variety of environmental, endogenous physical and chemical stresses that could produce genotoxic damage, but it possesses an efficient system to counteract the harmful effects of DNA-damaging assaults [22]. The STRING database in combination between the Rv1356c and uvrC with other proteins (STRING analysis version 11.0, with high confidence scores (0.700)) (Fig. S4). In the right-hand group, we found that the Rv1356c was associated with Rv1353c, Rv1354c and moeY/Rv1355c which is a yca/Rv1625, a central protein interconnecting the two groups with a high confidence score. In the left-hand group, there are three connectors (rpoB/Rv0667, DNA-directed RNA polymerase (beta chain) RpoB (transcriptase beta chain) (RNA polymerase beta subunit), pykA/Rv1617, probable pyruvate kinase PykA and ndkA/Rv2445c, probable nucleoside diphosphate kinase NdkA (NdkA (NDP kinase) (nucleoside-2-P kinase)). However, we want to focus on rpoB, a protein involved in resistance to first-line drugs (rifampicin) [23]. RpoB connects with gyrA/Rv0006, DNA gyrase (subunit A) GyrA (DNA topoisomerase (ATP-hydrolysing)) (DNA topoisomerase II) (type II DNA
topoisomerase) and gyrB/Rv0005, DNA gyrase (subunit B) GyrB (DNA topoisomerase (ATP-hydrolysing)) (DNA topoisomerase II) (type II DNA topoisomerase), all associated with resistance to second-line drugs (fluoroquinolones) [23], and then gyrAB associated between uvrA and uvrB, Uvr ABC system proteins with uvrC also known as excinuclease ABC.

Strains of the *M. tuberculosis* Beijing lineage are globally distributed and are often associated with severely pathogenic and virulent drug-resistant TB. The ability of *Mtb* to enter the NRP stage renders it even more resistant to drugs and allows it to persist in the host without causing symptoms. To deal with the problem of LTBI, understanding of the molecular mechanisms used by the pathogen is necessary. Strain-specific proteins identified in this study, particularly narJ, could be used in the search for new therapeutic targets to prevent and control TB, especially LTBI.

Author statement

Bhanubong Saiboonjan: Methodology, Validation, Investigation, Conceptualization, and Writing – review & editing. Sittiruk Roytrakul: Funding acquisition and Resources. Arumee Sangka: Supervision and Investigation. Wises Namwat: Supervision, Conceptualization, Investigation, and Writing – review & editing.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence any aspect of the research, its conduct, or the manuscript in preparation. The authors declare that they have no known competing financial interests.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.bbrep.2021.100960.

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