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Gennaro

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(54) **PROTEINS EXPRESSED BY
MYCOBACTERIUM TUBERCULOSIS AND
NOT BY BCG AND THEIR USE AS
DIAGNOSTIC REAGENTS AND VACCINES**

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patent is extended or adjusted under 35
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8,021,832, which is a continuation of application No.
11/677,502, filed on Feb. 21, 2007, now Pat. No.
7,579,141, which is a division of application No.
10/009,383, filed as application No. PCT/US00/12257
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G01N 33/50 (2006.01)
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(2013.01); **A61K 39/00** (2013.01); **A61K**
2039/53 (2013.01); **C07K 14/35** (2013.01);
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USPC **424/248.1**; 424/184.1; 424/185.1;
424/234.1; 435/863

(58) **Field of Classification Search**
USPC 424/184.1, 185.1, 234.1, 248.1;
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See application file for complete search history.

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Gerard P. Norton; Jianming Jimmy Hao

(57) **ABSTRACT**

The present invention is directed to reagents useful for generating immune responses to *Mycobacterium tuberculosis* and for diagnosing infection and disease in a subject that has been exposed to *M. tuberculosis*.

7 Claims, 8 Drawing Sheets

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MTBN1

MTABPEVRTLREVVLDQLGTAESRAYKMWLPPLTNPVPLNELIARDRRQPLRFALGIMDE
PRRHLDQDVWGVVDVSGAGGNIIGGAPQTGKSTLLQTMVMSAAATHSPRNVQFYCIDLGGG
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QPVASDPYCDVFLIIDGWFGFVGEFPDLEGQVODLAAQGLAFGVHVIISTPRWTELKSRV
RDYLGTKIEFRLGDVNETQIDRITREIPANRPGRAVSMEXHLMIGVPRFDGVHSADNLV
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IVGAAGGMPMPMAFLAPLLPAAADIGLHIIVTCQMSQAYKATMDKPFVGAAPFGSGAPTMLFS
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MTBN2

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MTBN3

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MTBN4

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MTBN5

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IAGPEPAPPKPPPTPPMPIAGPAPTPTESQLAPRRPPTPQTPTGAPQQPESAPHVPSHGP
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GKVATGPSIQARLRAEEASGAQLAPGTEPSFAPLGQPRSYLAPFTRPAPTEPPPSPPQR
NSGRRAEERRVHPDLAAQHAAQPDSTATAATTGGRRRKRAAPDLATQKSLRPAAGPKVK
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ERAGRR

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MTBN7

MAEPLAVDPTGLSAAAAKLAGLVFPQPPAPIAVSGTDSVVAAINETMPSIESLVSDGLPG
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MTBN8

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PQPVTPTATPGPSGPATPTGTPGGEPAHPVKPAALAEQPGVPGQHAGGGTQSGPAHADESAA
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LDRALAAAC

FIG 1B

mtbn1

1 atgactgctg aaccggaagt acggacgctg cgcgaggttg tgctggacca
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1651 ttcaaggcca agcggcgccc ccctggccag gcattttctc tctcgccaga
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mtbn2

1 atggaaaaaa tgtcacatga tccgatcgct gccgacattg gcacgcaagt
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151 gcgacggcgt tcacatcgga gggcatccaa ttgctggctt ccaatgcac
201 ggcccaagac cagctccacc gtgcgggcga agcgggtccag gacgtcgccc
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FIG 2A

mtbn3

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1 atgctgtggc acgcaatgcc accggagcta aataccgcac ggctgatggc
51 cggcgcgggg cgggtcccaa tgcttgcggc ggccgcggga tggcagacgc
101 tttcggcggc tctggacgct caggccgtcg agttgaccgc gcgcctgaac
151 tctctgggag aagcctggac tggaggtggc agcgacaagg cgcttgcggc
201 tgcaacgccg atggtggtct ggctacaaac cgcgtcaaca caggccaaga
251 cccgtgcgat gcaggcgacg gcgcaagccg cggcatacac ccaggccatg
301 gccacgacgc cgtcgctgcc ggagatcgcc gcccaaccaca tcaccagggc
351 cgtccttacg gccaccaact tcttcggtat caacacgac ccgatcgcg
401 tgaccgagat ggattatttc atccgtatgt ggaaccaggc agccctggca
451 atggaggtct accaggccga gaccgcgggt aacacgcttt tcgagaagct
501 cgagcgatg gcgtcgatcc ttgatcccg cgcgagccag agcacgacga
551 acccgatctt cggaatgcc tccctggca gctcaacacc ggttggccag
601 ttgcgcggcg cggctaccca gaccctcggc caactgggtg agatgagcgg
651 cccgatgcag cagctgaccc agccgctgca gcaggtgacg tcgttggtca
701 gccaggtggg cggcacccgg gcgcgcaacc cagccgacga ggaagccgcg
751 cagatgggcc tgctcggcac cagtccgctg tcgaaccatc cgctggctgg
801 tggatcaggc cccagcgcgg gcgcggggct gctgcgcgcg gactcgctac
851 ctggcgaggg tgggtcggtg acccgcaacg cgtgatgtc tcagctgac
901 gaaaagccgg ttgccccctc ggtgatgccg gcggctgctg ccggatcgct
951 ggcgacgggt ggcgcgcgtc cgggtgggtg gggagcgatg ggccaggggtg
1001 cgcaatccgg cggctccacc aggcggggtc tggtcgcgcc ggcaccgctc
1051 gcgcaggagc gtgaagaaga cgacgaggac gactgggacg aagaggacga
1101 ctggtga
```

mtbn4

```
1 atggcagaga tgaagaccga tgccgctacc ctgcgcagg aggcaggtaa
51 tttcgagcgg atctccggcg acctgaaaac ccagatcgac cagggtggagt
101 cgacggcagg ttcgttgcag ggccagtggc gcggcgcggc ggggacggcc
151 gcccaggccg cgggtggtcg cttccaagaa gcagccaata agcagaagca
201 ggaactcgac gagatctcga cgaatattcg tcaggccggc gtccaatact
251 cgagggccga cgaggagcag cagcaggcgc tgcctcgca aatgggcttc
301 tga
```

mtbn5

```
1 atggcgggcg actacgacaa gctcttcgg ccgcacgaag gtatggaagc
51 tccggacgat atggcagcgc agccgttctt cgaccccagt gcttcgtttc
101 cgccggcgcc cgcacggcga aacctaccga agcccaacgg ccagactccg
151 ccccgacgt ccgacgacct gtcggagcgg ttcgtgtcgg ccccgccggc
201 gccaccccca ccccaacct cgcctccgc aactccgatg ccgatcgccg
251 caggagagcc gccctcgccg gaaccggccg catctaaacc acccacacc
301 cccatgcccc tcgccggacc cgaaccggcc ccacccaaac caccacacc
351 ccccatgccc atcgccggac ccgaaccggc cccacccaaa ccaccacac
401 ctccgatgcc catcgccgga cctgcaccca ccccaaccga atcccagttg
```

FIG 2B

451 ggcggggcca gaccacggac accacaaacg ccaaccggag cgccgcagca
501 accggaatca ccggcgcccc acgtaccctc gcacggggcca catcaacccc
551 ggcgcaccgc accagcaccg ccctgggcaa agatgccaat cggcgaaacc
601 ccgcccgtc cgtccagacc gtctgcgtcc ccggccgaac caccgaccog
651 gcctgcccc caacaactcc gacgtgcgag ccggggtcac cgctatcgca
701 cagacaccga acgaaacgtc gggaaggtag caactggtcc atccatccag
751 gcgcggctgc gggcagagga agcatccggc gcgcagctcg ccccggaac
801 ggagccctcg ccagcgccgt tgggccaacc gagatcgat ctggctccgc
851 ccacccgccc cgcgcgcaca gaacctcccc ccagcccctc gccgcagcgc
901 aactccggtc ggcgtgcga gcgacgcgtc caccocgatt tagccgcca
951 acatgcccg gcgcaacctg attcaattac ggccgcaacc actggcggtc
1001 gtcgcgcga ggcgtgcagc ccgatctcg acgcgacaca gaaatcctta
1051 aggcggcgcc ccaagggggc gaagggtgaag aagggtgaag ccagaaacc
1101 gaaggccacg aagccgccc aagtgggtgc gcagcgcgcc tggcgacatt
1151 ggggtgcatg gttgacgca atcaacctgg gcctgtcacc cgacgagaag
1201 tacgagctgg acctgcacgc tcgagtcgc cgcaatcccc gcgggtcgta
1251 tcagatcgcc gtcgtcggtc tcaaagggtg ggctggcaaa accacgctga
1301 cagcagcgtt ggggtcgacg ttggctcagg tgcgggcga ccggatcctg
1351 gctctagacg cggatccagg cgcggaaac ctgcgcgatc gggtagggcg
1401 acaatcgggc gcgaccatcg ctgatgtgct tgcagaaaaa gagctgtcgc
1451 actacaacga catccgcgca cacactagcg tcaatgcggt caatctggaa
1501 gtgctgccgg caccggaata cagctcggcg cagcgcgcg tcagcgacgc
1551 cgactggcat ttcacgcgcg atcctgcgtc gaggttttac aacctcgtct
1601 tggctgattg tggggccggc ttcttcgacc cgctgacccg cggcgtgctg
1651 tccacggtgt ccggtgtcgt ggtcgtggca agtgtctcaa tcgacggcgc
1701 acaacaggcg tcggtcgcgt tggactggtt gcgcaacaac ggttaccaag
1751 atttggcgag cgcgcgatgc gttggtcatca atcacatcat gccgcagaa
1801 cccaatgtcg cagttaaaga cctgggtcgg catttcgaac agcaagttca
1851 acccgccgg gtcgtggtca tgccgtggga caggcacatt gcggccggaa
1901 ccgagatttc actcgacttg ctcgacccta tctacaagcg caaggctctc
1951 gaattggccg cagcgtatc cgacgatttc gagagggtg gacgtcgttg
2001 a

mtbn6

1 ttgagcgcac ctgctgttgc tgctggctct accgcgcgg gggcaaccgc
51 tgcgcggcct gccaccacc gggtagcgat cctgaccggc agacggatga
101 ccgatttggg actgccagcg gcggtgcga tggaaactta tattgacgac
151 accgtcgcgg tgctttccga ggtgttgaa gacacgcgg ctgatgtact
201 cggcggtctt gactttaccg cgcaaggcgt gtggcggttc gctcgtcccg
251 gatcgccgce gctgaagctc gaccagtcac tcgatgacgc cggggtggtc
301 gacgggtcac tgctgactct ggtgtcagtc agtcgcaccg agcgtaccg
351 accgttgggt gaggatgtca tcgacgcgat cgcggtgctt gacgagtcac
401 ctgagttcga ccgcacggca ttgaatcgct ttgtgggggc ggcgatcccg
451 cttttgaccg cggccgtcat cgggatggcg atgcggcggt ggtgggaaac
501 tgggcgtagc ttgtgggtggc cgttggcgat tggcatcctg gggatcgctg

FIG 2C

551 tgcctggtagg cagcttcgtc gcgaacaggt tctaccagag cggccacctg
601 gccgagtgcc tactggtcac gacgtatctg ctgatcgcaa ccgccgcagc
651 gctggccgtg ccgttgccgc gcgggggtcaa ctggttgggg gcgccacaag
701 ttgccggcgc cgctacggcc gtgctgtttt tgaccttgat gacgcggggc
751 ggccctcgga agcgtcatga gttggcgtcg tttgccgtga tcaccgetat
801 cgcggtcatc gcggccgcgc ctgccttcgg ctatggatac caggactggg
851 tccccgcggg ggggatcgca ttccggctgt tcattgtgac gaatgcggcc
901 aagctgaccg tcgcggtcgc gcggatcgcg ctgccgccga ttccggtacc
951 cggcgaaacc gtggacaacg aggagtgtct cgatcccgtc gcgaccccg
1001 aggcctaccg cgaagaaacc ccgacctggc aggccatcat cgcgtcgggtg
1051 cccgcgtccg cgggtccggct caccgagcgc agcaaactgg ccaagcaact
1101 tctgatcgga tacgtcacgt cgggcaccct gattctggct gccggtgcca
1151 tcgcggtcgt ggtgcgcggg cacttctttg tacacagcct ggtggtcgcg
1201 ggtttgatca cgaccgtctg cggatttcgc tcgcggcttt acgccgagcg
1251 ctggtgtgcg tgggcgttgc tggcggcgac ggtcgcgatt ccgacgggtc
1301 tgacggccaa actcatcatc tggtagccgc actatgcctg gctgttgttg
1351 agcgtctacc tcacggtagc cctggttgcg ctggtggtgg tcgggtcgat
1401 ggctcacgtc cggcgcgttt caccggtcgt aaaacgaact ctggaattga
1451 tcgacggcgc catgatcgct gccatcattc ccatgctgct gtggatcacc
1501 ggggtgtacg acacgggtccg caatatccgg ttctga

mtbn7

1 atggctgaac cgttggccgt cgatcccacc ggcttgagcg cagcggccgc
51 gaaattggcc ggcctcgttt ttccgcagcc tccggcgccg atcgcggtca
101 gcggaacgga ttccgtggta gcagcaatca acgagaccat gccaaagcatc
151 gaatcgctgg tcagtgaagg gctgcccggc gtgaaagccg ccctgactcg
201 aacagcatcc aacatgaacg cggcggcgga cgtctatgcg aagaccgac
251 agtcactggg aaccagtttg agccagtatg cattcggctc gtcgggagaa
301 ggccctggctg gcgtcgcctc ggtcggtggt cagccaagtc aggcctacca
351 gctgctgagc acaccgtgt cacaggtcac gaccagctc ggcgagacgg
401 ccgctgagct ggcaccccgt gttgttgca cggtgccgca actcgttcag
451 ctggtccgc acgcggttca gatgtcgaa aacgcaccc ccatcgctca
501 gacgatcagt caaacggccc aacaggccgc ccagagcgcg cagggcgga
551 gcggcccaat gcccgcacag cttgccagcg ctgaaaaacc ggccaccgag
601 caagcggagc cgggtccacga agtgacaaac gacgatcagg gcgaccaggg
651 cgacgtgcag ccggccgagg tcgttgccgc ggcacgtgac gaaggcgccg
701 gcgcatcacc gggccagcag cccggcgggg gcgttcccgc gcaagccatg
751 gataccggag ccggtgcccg cccagcggcg agtccgctgg cggccccgt
801 cgatccgtcg actccggcac cctcaacaac cacaacgtg tag

FIG 2D

mtbn8
1 atgagtatta ccaggccgac gggcagctat gccagacaga tgctggatcc
51 gggcggctgg gtggaagccg atgaagacac tttctatgac cgggcccagg
101 aatatagcca ggttttgcaa agggtcaccg atgtattgga cacctgccgc
151 cagcagaaag gccacgtctt cgaaggcggc ctatgggtccg gcggcgccgc
201 caatgctgcc aacggcgccc tgggtgcaaa catcaatcaa ttgatgacgc
251 tgcaggatta tctcgccacg gtgattacct ggcacaggca tattgccggg
301 ttgattgagc aagctaaatc cgatatcggc aataatgtgg atggcgctca
351 acgggagatc gatatcctgg agaatgaccc tagcctggat gctgatgagc
401 gccataccgc catcaattca ttgggtcacgg cgacgcattg ggccaatgtc
451 agtctggctg ccgagaccgc tgagcgggtg ctggaatcca agaattggaa
501 acctccgaag aacgcactcg aggatttgtc tcagcagaag tcgccgccac
551 cccagacgt gccctacctg gtctgacct cccggggcac accgggcaca
601 ccgggaaccc cgatcacccc gggaacccc atcaccccgg gaacccaat
651 cacacccatc ccgggagcgc cggtaactcc gatcacacca acgcccggca
701 ctcccgctac gccgggtgacc ccgggcaagc cggtcacccc ggtgaccccg
751 gtcaaaccgg gcacaccagg cgagccaacc ccgatcacgc cggtcacccc
801 cccgggtgcc ccggccacac cggcaacccc ggccacgccc gttaccccag
851 ctcccgctcc acacccgcag ccggctccgg caccggcgcc atcgctggg
901 cccagaccgg ttacaccggc cactcccggc ccgtctggtc cagcaacacc
951 gggcacccca gggggcgagc cggcgccgca cgtcaaacc gcggcgcttg
1001 cggagcaacc tgggtgtgcc ggccagcatg cgggcggggg gacgcagtcg
1051 gggcctgccc atgctggacg atccgcgcgc tcggtgacgc cggctgcggc
1101 gtccgggtgt ccgggcgcac gggcgccggc cgcgcgcgcg agcgggtaccg
1151 ccgtggggagc gggcgcgctg tcgagcgtgg gtacggccgc ggcctcgggc
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1251 ggcggcgcca ccgagcacgc gggcggcctc ggcgcggacg gcacctcctg
1301 cccgcccgcg gtccgaccgat cacatcgaca aacccgatcg cagcgagtct
1351 gcagatgacg gtacgcgggt gtcgatgac ccggtgtcgg cggctcgggc
1401 ggcacgcgac gccgccactg cagctgccag cgcgcgccag cgtggccgcg
1451 gtgatgcgct gcggttggcg cgacgcacgc cggcgcgct caacgcgtcc
1501 gacaacaacg cgggcgacta cgggttcttc tggatcaccc cggtgaccac
1551 cgacgggtcc atcgtcgtgg ccaacagcta tgggtggcc tacatacccg
1601 acgggatgga attgccgaat aaggtgtact tggccagcgc ggatcacgca
1651 atcccggttg acgaaattgc acgctgtgcc acctaccgg ttttggccgt
1701 gcaagcctgg gcggtttcc acgacatgac gctgcgggcg gtgatcggta
1751 ccgcgagca gttggccagt tcggatcccg gtgtggccaa gattgtgctg
1801 gagccagatg acattccgga gagcgcaaaa atgacgggccc ggtcgcgct
1851 ggaggtcgtc gacccctcgg cggcggtca gctggccgac actaccgatc
1901 agcgtttgct cgacttggtg ccgcgcgcgc cgggtggatgt caatccaccg
1951 ggcgatgagc ggcacatgct gtgggttcgag ctgatgaagc ccatgaccag
2001 caccgctacc ggccgcgagg ccgctcatct gcgggcgttc cgggcctacg
2051 ctgcccactc acaggagatt gccctgcacc aagcgcacac tgcgactgac
2101 gcggccgtcc agcgtgtggc cgtcgcgga tggctgtact ggcaatacgt
2151 caccgggttg ctcgaccggg ccctggccgc cgcattgctga

FIG 2E

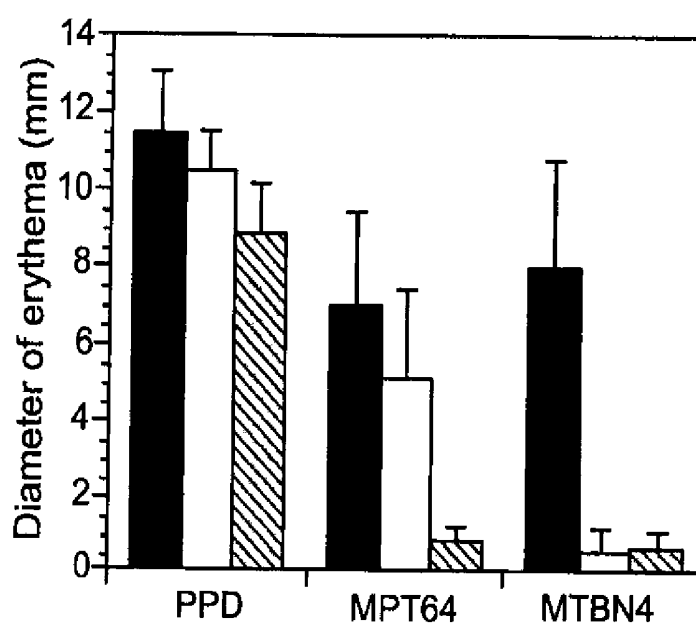


FIG 3

**PROTEINS EXPRESSED BY
MYCOBACTERIUM TUBERCULOSIS AND
NOT BY BCG AND THEIR USE AS
DIAGNOSTIC REAGENTS AND VACCINES**

This application is a divisional of, and claims priority to, U.S. application Ser. No. 13/198,108, filed Aug. 4, 2011 and now pending, which is a continuation of, and claims priority to, U.S. application Ser. No. 12/503,717, filed Jul. 15, 2009 and now issued as U.S. Pat. No. 8,021,832, which is a continuation of, and claims priority to, U.S. application Ser. No. 11/677,502, filed Feb. 21, 2007, now U.S. Pat. No. 7,579,141, which is a divisional of, and claims priority to, U.S. application Ser. No. 10/009,383, filed Mar. 4, 2002 and now issued as U.S. Pat. No. 7,932,373, which claims priority to International Application No. PCT/US00/12257, filed May 4, 2000, which claims priority to U.S. Provisional Application Ser. No. 60/132,505, filed May 4, 1999, the disclosures of each of which are hereby incorporated by reference in their entireties.

BACKGROUND OF THE INVENTION

Tuberculosis infection continues to be a world-wide health problem. This situation has recently been greatly exacerbated by the emergence of multi-drug resistant strains of *M. tuberculosis* and the international AIDS epidemic. It has thus become increasingly important that effective vaccines against and reliable diagnostic reagents for *M. tuberculosis* be produced.

The disclosure of U.S. Pat. No. 6,087,163 is incorporated herein by reference in its entirety.

SUMMARY OF THE INVENTION

The invention is based on the inventor's discovery that a polypeptide encoded by an open reading frame (ORF) in the genome of *M. tuberculosis* that is absent from the genome of the Bacille Calmette Guérin (BCG) strain of *M. bovis* elicited a delayed-type hypersensitivity response in animals infected with *M. tuberculosis* but not in animals sensitized with BCG. Thus proteins encoded by ORFs present in the genome of *M. tuberculosis* but absent from the genome of BCG represent reagents that are useful in discriminating between *M. tuberculosis* and BCG and, in particular, for diagnostic methods (e.g., skin tests and in vitro assays for *M. tuberculosis*-specific antibodies and lymphocyte responsiveness) which discriminate between exposure of a subject to *M. tuberculosis* and vaccination with BCG. The invention features these polypeptides, functional segments thereof, DNA molecules encoding either the polypeptides or the functional segments, vectors containing the DNA molecules, cells transformed by the vectors, compositions containing one or more of any of the above polypeptides, functional segments, or DNA molecules, and a variety of diagnostic, therapeutic, and prophylactic (vaccine) methodologies utilizing the foregoing.

Specifically, the invention features an isolated DNA molecule containing a DNA sequence encoding a polypeptide with a first amino acid sequence that can be the amino acid sequence of the polypeptide MTBN1, MTBN2, MTBN3, MTBN4, MTBN5, MTBN6, MTBN7 or MTBN8, as depicted in FIGS. 1A and 1B, or a second amino acid sequence identical to the first amino acid sequence with conservative substitutions; the polypeptide has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Also included in the invention is an isolated portion of the above DNA molecule. The portion of the DNA molecule encodes a segment of the polypeptide shorter than the full-

length polypeptide, and the segment has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Other embodiments of the invention are vectors containing the above DNA molecules and transcriptional and translational regulatory sequences operationally linked to the DNA sequence; the regulatory sequences allow for expression of the polypeptide or functional segment encoded by the DNA sequence in a cell. The invention encompasses cells (e.g., eukaryotic and prokaryotic cells) transformed with the above vectors.

The invention encompasses compositions containing any of the above vectors and a pharmaceutically acceptable diluent or filler. Other compositions (to be used, for example, as DNA vaccines) can contain at least two (e.g., three, four, five, six, seven, eight, nine, ten, twelve, fifteen, or twenty) DNA sequences, each encoding a polypeptide of the *Mycobacterium tuberculosis* complex or a functional segment thereof, with the DNA sequences being operationally linked to transcriptional and translational regulatory sequences which allow for expression of each of the polypeptides in a cell of a vertebrate. In such compositions, at least one (e.g., two, three, four, five, six, seven, or eight) of the DNA sequences is one of the above DNA molecules of the invention. The encoded polypeptides will preferably be those not encoded by the genome of cells of the BCG strain of *M. bovis*.

The invention also features an isolated polypeptide with a first amino acid sequence that can be the sequence of the polypeptide MTBN1, MTBN2, MTBN3, MTBN4, MTBN5, MTBN6, MTBN7 or MTBN8 as depicted in FIGS. 1A and 1B, or a second amino acid sequence identical to the first amino acid sequence with conservative substitutions. The polypeptide has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Also included in the invention is an isolated segment of this polypeptide, the segment being shorter than the full-length polypeptide and having *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Other embodiments are compositions containing the polypeptide, or functional segment, and a pharmaceutically acceptable diluent or filler. Compositions of the invention can also contain at least two (e.g., three, four, five, six, seven, eight, nine, ten, twelve, fifteen, or twenty) polypeptides of the *Mycobacterium tuberculosis* complex, or functional segments thereof, with at least one of the at least two (e.g., two, three, four, five, six, seven, or eight) polypeptides having the sequence of one of the above described polypeptides of the invention. The polypeptides will preferably be those not encoded by the genome of cells of the BCG strain of *M. bovis*.

The invention also features methods of diagnosis. One embodiment is a method involving: (a) administration of one of the above polypeptide compositions to a subject suspected of having or being susceptible to *Mycobacterium tuberculosis* infection; and (b) detecting an immune response in the subject to the composition, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection. An example of such a method is a skin test in which the test substance (e.g., compositions containing one or more of MTBN1-MTBN8) is injected intradermally into the subject and in which a skin delayed-type hypersensitivity response is tested for. Another embodiment is a method that involves: (a) providing a population of cells containing CD4 T lymphocytes from a subject; (b) providing a population of cells containing antigen presenting cells (APC) expressing a major histocompatibility complex (MHC) class II molecule expressed by the subject; (c) contacting the CD4 lymphocytes of (a) with the APC of (b) in the presence of one or more of the polypeptides, functional segments, and or polypeptide com-

positions of the invention; and (d) determining the ability of the CD4 lymphocytes to respond to the polypeptide, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection. Another diagnostic method of the invention involves: (a) contacting a polypeptide, a functional segment, or a polypeptide/functional segment composition of the invention with a bodily fluid of a subject; (b) detecting the presence of binding of antibody to the polypeptide, functional segment, or polypeptide/functional segment composition, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection.

Also encompassed by the invention are methods of vaccination. These methods involve administration of any of the above polypeptides, functional segments, or DNA compositions to a subject. The compositions can be administered alone or with one or more of the other compositions.

As used herein, an "isolated DNA molecule" is a DNA which is one or both of: not immediately contiguous with one or both of the coding sequences with which it is immediately contiguous (i.e., one at the 5' end and one at the 3' end) in the naturally-occurring genome of the organism from which the DNA is derived; or which is substantially free of DNA sequence with which it occurs in the organism from which the DNA is derived. The term includes, for example, a recombinant DNA which incorporated into a vector, e.g., into an autonomously replicating plasmid or virus, or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule (e.g., a cDNA or a genomic fragment produced by PCR or restriction endonuclease treatment) independent of other DNA sequences. Isolated DNA also includes a recombinant DNA which is part of a hybrid DNA encoding additional *M. tuberculosis* polypeptide sequences.

"DNA molecules" include cDNA, genomic DNA, and synthetic (e.g., chemically synthesized) DNA. Where single-stranded, the DNA molecule may be a sense strand or an antisense strand.

An "isolated polypeptide" of the invention is a polypeptide which either has no naturally-occurring counterpart, or has been separated or purified from components which naturally accompany it, e.g., in *M. tuberculosis* bacteria. Typically, the polypeptide is considered "isolated" when it is at least 70%, by dry weight, free from the proteins and naturally-occurring organic molecules with which it is naturally associated.

Preferably, a preparation of a polypeptide of the invention is at least 80%, more preferably at least 90%, and most preferably at least 99%, by dry weight, the peptide of the invention. Since a polypeptide that is chemically synthesized is, by its nature, separated from the components that naturally accompany it, the synthetic polypeptide is "isolated."

An isolated polypeptide of the invention can be obtained, for example, by extraction from a natural source (e.g., *M. tuberculosis* bacteria); by expression of a recombinant nucleic acid encoding the polypeptide; or by chemical synthesis. A polypeptide that is produced in a cellular system different from the source from which it naturally originates is "isolated," because it will be separated from components which naturally accompany it. The extent of isolation or purity can be measured by any appropriate method, e.g., column chromatography, polyacrylamide gel electrophoresis, or HPLC analysis.

The polypeptides may contain a primary amino acid sequence that has been modified from those disclosed herein. Preferably these modifications consist of conservative amino acid substitutions. Conservative substitutions typically include substitutions within the following groups: glycine and alanine; valine, isoleucine, and leucine; aspartic acid and

glutamic acid; asparagine and glutamine; serine and threonine; lysine and arginine; and phenylalanine and tyrosine.

The terms "protein" and "polypeptide" are used herein to describe any chain of amino acids, regardless of length or post-translational modification (for example, glycosylation or phosphorylation). Thus, the term "*Mycobacterium tuberculosis* polypeptide" includes full-length, naturally occurring *Mycobacterium tuberculosis* protein, as well a recombinantly or synthetically produced polypeptide that corresponds to a full-length naturally occurring *Mycobacterium tuberculosis* protein or to particular domains or portions of a naturally occurring protein. The term also encompasses a mature *Mycobacterium tuberculosis* polypeptide which has an added amino-terminal methionine (useful for expression in prokaryotic cells) or any short amino acid sequences useful for protein purification by affinity chromatography, e.g., polyhistidine for purification by metal chelate chromatography.

As used herein, "immunogenic" means capable of activating a primary or memory immune response. Immune responses include responses of CD4+ and CD8+ T lymphocytes and B-lymphocytes. In the case of T lymphocytes, such responses can be proliferative, and/or cytokine (e.g., interleukin (IL)-2, IL-3, IL-4, IL-5, IL-6, IL-12, IL-13, IL-15, tumor necrosis factor- α (TNF- α), or interferon- γ (IFN- γ))-producing, or they can result in generation of cytotoxic T-lymphocytes (CTL). B-lymphocyte responses can be those resulting in antibody production by the responding B lymphocytes.

As used herein, "antigenic" means capable of being recognized by either antibody molecules or antigen-specific T cell receptors (TCR) on activated effector T cells (e.g., cytokine-producing T cells or CTL).

Thus, polypeptides that have "*Mycobacterium tuberculosis* specific antigenic properties" are polypeptides that: (a) can be recognized by and bind to antibodies elicited in response to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e.g., polypeptides); or (b) contain subsequences which, subsequent to processing of the polypeptide by appropriate antigen presenting cells (APC) and bound to appropriate major histocompatibility complex (MHC) molecules, are recognized by and bind to TCR on effector T cells elicited in response to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e.g., polypeptides).

As used herein, polypeptides that have "*Mycobacterium tuberculosis* specific immunogenic properties" are polypeptides that: (a) can elicit the production of antibodies that recognize and bind to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e.g., polypeptides); or (b) contain subsequences which, subsequent to processing of the polypeptide by appropriate antigen presenting cells (APC) and bound to appropriate major histocompatibility complex (MHC) molecules on the surface of the APC, activate T cells with TCR that recognize and bind to peptide fragments derived by processing by APC of *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e.g., polypeptides) and bound to MHC molecules on the surface of the APC. The immune responses elicited in response to the immunogenic polypeptides are preferably protective. As used herein, "protective" means preventing establishment of an infection or onset of a disease or lessening the severity of a disease existing in a subject. "Preventing" can include delaying onset, as well as partially or completely blocking progress of the disease.

As used herein, a "functional segment of a *Mycobacterium tuberculosis* polypeptide" is a segment of the polypeptide that has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties.

Where a polypeptide, functional segment of a polypeptide, or a mixture of polypeptides and/or functional segments have been administered (e.g., by intradermal injection) to a subject for the purpose of testing for a *M. tuberculosis* infection or susceptibility to such an infection, "detecting an immune response" means examining the subject for signs of an immunological reaction to the administered material, e.g., reddening or swelling of the skin at the site of an intradermal injection. Where the subject has antibodies to the administered material, the response will generally be rapid, e.g., 1 minute to 24 hours. On the other hand, a memory or activated T cell reaction of pre-immunized T lymphocytes in the subject is generally slower, appearing only after 24 hours and being maximal at 24-96 hours.

As used herein, a "subject" can be a human subject or a non-human mammal such as a non-human primate, a horse, a bovine animal, a pig, a sheep, a goat, a dog, a cat, a rabbit, a guinea pig, a hamster, a rat, or a mouse.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention pertains. In case of conflict, the present document, including definitions, will control. Preferred methods and materials are described below, although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention. Unless otherwise indicated, these materials and methods are illustrative only and are not intended to be limiting.

All publications, patent applications, patents and other references mentioned herein are illustrative only and not intended to be limiting.

Other features and advantages of the invention, e.g., methods of diagnosing *M. tuberculosis* infection, will be apparent from the following description, from the drawings and from the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1A and 1B are a depiction of the amino acid sequences of *M. tuberculosis* polypeptides MTBN1-MTBN8 (SEQ ID NOS:1-8, respectively).

FIGS. 2A-2E are a depiction of the nucleotide sequences of the coding regions (mtbn1-mtbn8) encoding MTBN1-MTBN8 (SEQ ID NOS:9-16, respectively).

FIG. 3 is a bar graph showing the delayed-type hypersensitivity responses induced by intradermal injection of 3 different test reagents in female guinea pigs that had been either infected with *M. tuberculosis* cells or sensitized with BCG or *M. avium* cells.

DETAILED DESCRIPTION

The genome of *M. tuberculosis* [Cole et al. (1998) Nature 393: 537-544] contains open reading frames (ORFs) that have been deleted from the avirulent BCG strain.

The polypeptides encoded by these ORFs are designated herein "*M. tuberculosis* BCG Negative" polypeptides ("MTBN") and the ORFs are designated "mtbn." The invention is based on the discovery that a MTBN polypeptide (MTBN4) elicited a skin response in animals infected with *M. tuberculosis*, but not in animals sensitized to either BCG or *M. avium*, a non-*M. tuberculosis*-complex strain of mycobacteria (see Example 1 below). These findings indicate that MTBN (e.g., MTBN1-MTBN8) can be used in diagnostic tests that discriminate infection of a subject by *M. tuberculosis* from exposure to both mycobacteria other than the *M. tuberculosis*-complex and BCG. The *M. tuberculosis*-com-

plex includes *M. tuberculosis*, *M. bovis*, *M. microti*, and *M. africanum*. Thus they can be used to discriminate subjects exposed to *M. tuberculosis*, and thus potentially having or being in danger of having tuberculosis, from subjects that have been vaccinated with BCG, the most widely used tuberculosis vaccine. Diagnostic assays that are capable of such discrimination represent a major advance that will greatly reduce wasted effort and consequent costs resulting from further diagnostic tests and/or therapeutic procedures in subjects that have given positive results in less discriminatory diagnostic tests.

Furthermore, the results in Example 1 show that MTBN4, as expressed by whole viable *M. tuberculosis* organisms, is capable of inducing a strong immune response in subjects infected with the organisms and thus has the potential to be a vaccine.

The MTBN polypeptides of the invention include, for example, polypeptides encoded within the RD1, RD2, and RD3 regions of the *M. tuberculosis* genome [Mahairas et al. (1996) J. Bacteriol. 178: 1274-1282]. Of particular interest are polypeptides encoded by ORFs within the RD1 region of the *M. tuberculosis* genome. However, the invention is not restricted to the RD 1, RD2, and RD3 region encoded polypeptides and includes any polypeptides encoded by ORFs contained in the genome of one or more members of the *M. tuberculosis* genome and not contained in the genome of BCG. The amino acid sequences of MTBN1-MTBN8 are shown in FIGS. 1A and 1B and the nucleotide sequences of mtbn1-mtbn8 are shown in FIGS. 2A-2E.

The invention encompasses: (a) isolated DNA molecules containing mtbn sequences (e.g., mtbn1-mtbn8) encoding MTBN polypeptides (e.g., MTBN1-MTBN8) and isolated portions of such DNA molecules that encode polypeptide segments having antigenic and immunogenic properties (i.e., functional segments); (b) the MTBN polypeptides themselves (e.g., MTBN1-MTBN8) and functional segments of them; (c) antibodies (including antigen binding fragments, e.g., F(ab')₂, Fab, Fv, and single chain Fv fragments of such antibodies) that bind to the MTBN polypeptides (e.g., MTBN1-MTBN8) and functional segments; (d) nucleic acid molecules (e.g., vectors) containing and capable of expressing one or more of the mtbn (e.g., mtbn1-mtbn8) sequences and portions of DNA molecules; (e) cells (e.g., bacterial, yeast, insect, or mammalian cells) transformed by such vectors; (f) compositions containing vectors encoding one or more *M. tuberculosis* polypeptides (or functional segments) including both the MTBN (e.g., MTBN1-MTBN8) polypeptides (or functional segments thereof) and previously described *M. tuberculosis* polypeptides such as ESAT-6, 14 kDa antigen, MPT63, 19 kDa antigen, MPT64, MPT51, MTC28, 38 kDa antigen, 45/47 kDa antigen, MPB70, Ag85 complex, MPT53, and KatG (see also U.S. Pat. No. 6,087, 163); (g) compositions containing one or more *M. tuberculosis* polypeptides (or functional segments), including both the polypeptides of the invention and previously described *M. tuberculosis* polypeptides such as those described above; (h) compositions containing one or more of the antibodies described in (c); (i) methods of diagnosis involving either (1) administration (e.g., intradermal injection) of any of the above polypeptide compositions to a subject suspected of having or being susceptible to *M. tuberculosis* infection, (2) in vitro testing of lymphocytes (B-lymphocytes, CD4 T lymphocytes, and CD8 T lymphocytes) from such a subject for responsiveness (e.g., by measuring cell proliferation, antibody production, cytokine production, or CTL activity) to any of the above polypeptide compositions, (3) testing of a bodily fluid (e.g., blood, saliva, plasma, serum, urine, or

semen or a lavage such as a bronchoalveolar lavage, a vaginal lavage, or lower gastrointestinal lavage) for antibodies to the MTBN polypeptides (e.g., MTBN1-MTBN8) or functional segments thereof, or the above-described polypeptide compositions; (4) testing of a bodily fluid (e.g., as above) for the presence of *M. tuberculosis*, MTBN (e.g., MTBN1-MTBN8) polypeptides or functional segments thereof, or the above-described polypeptide compositions in assays using the antibodies described in (c); and (5) testing of a tissue (e.g. lung or bronchial tissue) or a body fluid (e.g., as above) for the presence of nucleic acid molecules (e.g., DNA or RNA) encoding MTBN polypeptides (e.g., MTBN1-MTBN8) (or portions of such a nucleic acid molecules) using nucleic acid probes or primers having nucleotide sequences of the nucleic molecules, portions of the nucleic molecules, or the complements of such molecules; and (j) methods of vaccination involving administration to a subject of the compositions of either (f), (g), (h) or a combination of any two or even all 3 compositions.

With respect to diagnosis, purified MTBN proteins, functional segments of such proteins, or mixtures of proteins and/or the functional fragments have the above-described advantages of discriminating infection by *M. tuberculosis* from either infection by other bacteria, and in particular, non-pathogenic mycobacteria, or from exposure (by, for example, vaccination) to BCG.

Furthermore, compositions containing the proteins, functional segments of the proteins, or mixtures of the proteins and/or the functional segments allows for improved quality control since "batch-to-batch" variability is greatly reduced in comparison to complex mixtures such as purified protein derivative (PPD) of tuberculin.

The use of the above-described polypeptide and nucleic acid reagents for vaccination also provides for highly specific and effective immunization. Since the virulent *M. tuberculosis* polypeptides encoded by genes absent from avirulent BCG are likely to be mediators of virulence, immunity directed to them can be especially potent in terms of protective capacity. Where vaccination is performed with nucleic acids both in vivo and ex vivo methods can be used. In vivo methods involve administration of the nucleic acids themselves to the subject and ex vivo methods involve obtaining cells (e.g., bone marrow cells or fibroblasts) from the subject, transducing the cells with the nucleic acids, preferably selecting or enriching for successfully transduced cells, and administering the transduced cells to the subject. Alternatively, the cells that are transduced and administered to the subject can be derived from another subject. Methods of vaccination and diagnosis are described in greater detail in U.S. Pat. No. 6,087,163, the disclosure of which is incorporated herein by reference in its entirety.

The following example is meant to illustrate, not limit the invention.

Example 1

MTBN4 Elicits a Specific Skin Reaction in Guinea Pigs Infected with *M. tuberculosis*

Four groups of outbred female guinea pigs (18 per group) were used to test the usefulness of the MTBN4 polypeptide as a *M. tuberculosis*-specific diagnostic reagent. The four groups were treated as follows.

Group 1 animals were infected by aerosol with approximately 100 *M. tuberculosis* strain H37Rv cells.

Group 2 animals were sensitized intradermally with 106 live *M. bovis* BCG Japanese cells.

Group 3 animals were sensitized intradermally with 106 live *M. avium* cells.

Group 4 animals were mock-sensitized by intradermal injection with saline.

Seven weeks after infection or sensitization, the animals were injected intradermally with 1 µg of PPD (6 animals from each group), 2 µg of purified recombinant MPT64 (6 animals from each group), or 2 µg of MTBN4 (6 animals from each group). The diameter of the resulting erythema was measured 24 hours later. Data are expressed as mean diameter of erythema (in mm) and standard deviations are indicated (FIG. 3).

No erythema was detected in the group 4 animals with any test substance and thus no data are shown for this group. On the other hand, group 1 animals (solid bars) showed a significant response with all three test substances. Group 2 animals (open bars) showed a significant response to PPD and MPT64 but not MTBN4.

Group 3 animals showed a significant response to PPD only (hatched bars).

Thus, PPD which contains antigenic/immunogenic molecules common to the *M. tuberculosis*-complex as well as other mycobacterial strains, gave the least discriminatory results in that it induced responses in animals infected with or sensitized to mycobacteria of the *M. tuberculosis*-complex (*M. tuberculosis* and BCG) as well as another non-pathogenic mycobacterium (*M. avium*).

While MPT64, which is encoded and expressed by both *M. tuberculosis* and BCG, did not elicit a response in animals infected with *M. avium*, it did elicit responses in both the *M. tuberculosis* infected and the BCG sensitized animals. Finally, MTBN4 elicited a response in only the *M. tuberculosis* animals. Thus it induced the most specific response and, most importantly, allowed for discrimination between animals infected with *M. tuberculosis* and those sensitized to BCG.

Although the invention has been described with reference to the presently preferred embodiment, it should be understood that various modifications can be made without departing from the spirit of the invention. Accordingly, the invention is limited only by the following claims.

SEQUENCE LISTING

<160> NUMBER OF SEQ ID NOS: 16

<210> SEQ ID NO 1

<211> LENGTH: 591

<212> TYPE: PRT

<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 1

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-continued

1	5	10	15
Gln Leu Gly Thr Ala Glu Ser Arg Ala Tyr Lys Met Trp Leu Pro Pro	20	25	30
Leu Thr Asn Pro Val Pro Leu Asn Glu Leu Ile Ala Arg Asp Arg Arg	35	40	45
Gln Pro Leu Arg Phe Ala Leu Gly Ile Met Asp Glu Pro Arg Arg His	50	55	60
Leu Gln Asp Val Trp Gly Val Asp Val Ser Gly Ala Gly Gly Asn Ile	65	70	75
Gly Ile Gly Gly Ala Pro Gln Thr Gly Lys Ser Thr Leu Leu Gln Thr	85	90	95
Met Val Met Ser Ala Ala Ala Thr His Ser Pro Arg Asn Val Gln Phe	100	105	110
Tyr Cys Ile Asp Leu Gly Gly Gly Gly Leu Ile Tyr Leu Glu Asn Leu	115	120	125
Pro His Val Gly Gly Val Ala Asn Arg Ser Glu Pro Asp Lys Val Asn	130	135	140
Arg Val Val Ala Glu Met Gln Ala Val Met Arg Gln Arg Glu Thr Thr	145	150	155
Phe Lys Glu His Arg Val Gly Ser Ile Gly Met Tyr Arg Gln Leu Arg	165	170	175
Asp Asp Pro Ser Gln Pro Val Ala Ser Asp Pro Tyr Gly Asp Val Phe	180	185	190
Leu Ile Ile Asp Gly Trp Pro Gly Phe Val Gly Glu Phe Pro Asp Leu	195	200	205
Glu Gly Gln Val Gln Asp Leu Ala Ala Gln Gly Leu Ala Phe Gly Val	210	215	220
His Val Ile Ile Ser Thr Pro Arg Trp Thr Glu Leu Lys Ser Arg Val	225	230	235
Arg Asp Tyr Leu Gly Thr Lys Ile Glu Phe Arg Leu Gly Asp Val Asn	245	250	255
Glu Thr Gln Ile Asp Arg Ile Thr Arg Glu Ile Pro Ala Asn Arg Pro	260	265	270
Gly Arg Ala Val Ser Met Glu Lys His His Leu Met Ile Gly Val Pro	275	280	285
Arg Phe Asp Gly Val His Ser Ala Asp Asn Leu Val Glu Ala Ile Thr	290	295	300
Ala Gly Val Thr Gln Ile Ala Ser Gln His Thr Glu Gln Ala Pro Pro	305	310	315
Val Arg Val Leu Pro Glu Arg Ile His Leu His Glu Leu Asp Pro Asn	325	330	335
Pro Pro Gly Pro Glu Ser Asp Tyr Arg Thr Arg Trp Glu Ile Pro Ile	340	345	350
Gly Leu Arg Glu Thr Asp Leu Thr Pro Ala His Cys His Met His Thr	355	360	365
Asn Pro His Leu Leu Ile Phe Gly Ala Ala Lys Ser Gly Lys Thr Thr	370	375	380
Ile Ala His Ala Ile Ala Arg Ala Ile Cys Ala Arg Asn Ser Pro Gln	385	390	395
Gln Val Arg Phe Met Leu Ala Asp Tyr Arg Ser Gly Leu Leu Asp Ala	405	410	415
Val Pro Asp Thr His Leu Leu Gly Ala Gly Ala Ile Asn Arg Asn Ser	420	425	430

-continued

Ala Ser Leu Asp Glu Ala Val Gln Ala Leu Ala Val Asn Leu Lys Lys
 435 440 445

Arg Leu Pro Pro Thr Asp Leu Thr Thr Ala Gln Leu Arg Ser Arg Ser
 450 455 460

Trp Trp Ser Gly Phe Asp Val Val Leu Leu Val Asp Asp Trp His Met
 465 470 475 480

Ile Val Gly Ala Ala Gly Gly Met Pro Pro Met Ala Pro Leu Ala Pro
 485 490 495

Leu Leu Pro Ala Ala Ala Asp Ile Gly Leu His Ile Ile Val Thr Cys
 500 505 510

Gln Met Ser Gln Ala Tyr Lys Ala Thr Met Asp Lys Phe Val Gly Ala
 515 520 525

Ala Phe Gly Ser Gly Ala Pro Thr Met Phe Leu Ser Gly Glu Lys Gln
 530 535 540

Glu Phe Pro Ser Ser Glu Phe Lys Val Lys Arg Arg Pro Pro Gly Gln
 545 550 555 560

Ala Phe Leu Val Ser Pro Asp Gly Lys Glu Val Ile Gln Ala Pro Tyr
 565 570 575

Ile Glu Pro Pro Glu Glu Val Phe Ala Ala Pro Pro Ser Ala Gly
 580 585 590

<210> SEQ ID NO 2
 <211> LENGTH: 99
 <212> TYPE: PRT
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 2

Met Glu Lys Met Ser His Asp Pro Ile Ala Ala Asp Ile Gly Thr Gln
 1 5 10 15

Val Ser Asp Asn Ala Leu His Gly Val Thr Ala Gly Ser Thr Ala Leu
 20 25 30

Thr Ser Val Thr Gly Leu Val Pro Ala Gly Ala Asp Glu Val Ser Ala
 35 40 45

Gln Ala Ala Thr Ala Phe Thr Ser Glu Gly Ile Gln Leu Leu Ala Ser
 50 55 60

Asn Ala Ser Ala Gln Asp Gln Leu His Arg Ala Gly Glu Ala Val Gln
 65 70 75 80

Asp Val Ala Arg Thr Tyr Ser Gln Ile Asp Asp Gly Ala Ala Gly Val
 85 90 95

Phe Ala Glu

<210> SEQ ID NO 3
 <211> LENGTH: 368
 <212> TYPE: PRT
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 3

Met Leu Trp His Ala Met Pro Pro Glu Leu Asn Thr Ala Arg Leu Met
 1 5 10 15

Ala Gly Ala Gly Pro Ala Pro Met Leu Ala Ala Ala Ala Gly Trp Gln
 20 25 30

Thr Leu Ser Ala Ala Leu Asp Ala Gln Ala Val Glu Leu Thr Ala Arg
 35 40 45

Leu Asn Ser Leu Gly Glu Ala Trp Thr Gly Gly Gly Ser Asp Lys Ala
 50 55 60

-continued

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Leu Ala Ala Ala Thr Pro Met Val Val Trp Leu Gln Thr Ala Ser Thr
65              70              75              80

Gln Ala Lys Thr Arg Ala Met Gln Ala Thr Ala Gln Ala Ala Ala Tyr
            85              90              95

Thr Gln Ala Met Ala Thr Thr Pro Ser Leu Pro Glu Ile Ala Ala Asn
            100             105             110

His Ile Thr Gln Ala Val Leu Thr Ala Thr Asn Phe Phe Gly Ile Asn
            115             120             125

Thr Ile Pro Ile Ala Leu Thr Glu Met Asp Tyr Phe Ile Arg Met Trp
            130             135             140

Asn Gln Ala Ala Leu Ala Met Glu Val Tyr Gln Ala Glu Thr Ala Val
145             150             155             160

Asn Thr Leu Phe Glu Lys Leu Glu Pro Met Ala Ser Ile Leu Asp Pro
            165             170             175

Gly Ala Ser Gln Ser Thr Thr Asn Pro Ile Phe Gly Met Pro Ser Pro
            180             185             190

Gly Ser Ser Thr Pro Val Gly Gln Leu Pro Pro Ala Ala Thr Gln Thr
            195             200             205

Leu Gly Gln Leu Gly Glu Met Ser Gly Pro Met Gln Gln Leu Thr Gln
210             215             220

Pro Leu Gln Gln Val Thr Ser Leu Phe Ser Gln Val Gly Gly Thr Gly
225             230             235             240

Gly Gly Asn Pro Ala Asp Glu Glu Ala Ala Gln Met Gly Leu Leu Gly
            245             250             255

Thr Ser Pro Leu Ser Asn His Pro Leu Ala Gly Gly Ser Gly Pro Ser
            260             265             270

Ala Gly Ala Gly Leu Leu Arg Ala Glu Ser Leu Pro Gly Ala Gly Gly
275             280             285

Ser Leu Thr Arg Thr Pro Leu Met Ser Gln Leu Ile Glu Lys Pro Val
290             295             300

Ala Pro Ser Val Met Pro Ala Ala Ala Ala Gly Ser Ser Ala Thr Gly
305             310             315             320

Gly Ala Ala Pro Val Gly Ala Gly Ala Met Gly Gln Gly Ala Gln Ser
            325             330             335

Gly Gly Ser Thr Arg Pro Gly Leu Val Ala Pro Ala Pro Leu Ala Gln
            340             345             350

Glu Arg Glu Glu Asp Asp Glu Asp Asp Trp Asp Glu Glu Asp Asp Trp
355             360             365

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<210> SEQ ID NO 4
<211> LENGTH: 100
<212> TYPE: PRT
<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 4

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Met Ala Glu Met Lys Thr Asp Ala Ala Thr Leu Ala Gln Glu Ala Gly
1              5              10             15

Asn Phe Glu Arg Ile Ser Gly Asp Leu Lys Thr Gln Ile Asp Gln Val
            20             25             30

Glu Ser Thr Ala Gly Ser Leu Gln Gly Gln Trp Arg Gly Ala Ala Gly
            35             40             45

Thr Ala Ala Gln Ala Ala Val Arg Phe Gln Glu Ala Ala Asn Lys
            50             55             60

Gln Lys Gln Glu Leu Asp Glu Ile Ser Thr Asn Ile Arg Gln Ala Gly
65             70             75             80

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-continued

Val Gln Tyr Ser Arg Ala Asp Glu Glu Gln Gln Gln Ala Leu Ser Ser
85 90 95

Gln Met Gly Phe
100

<210> SEQ ID NO 5

<211> LENGTH: 666

<212> TYPE: PRT

<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 5

Met Ala Ala Asp Tyr Asp Lys Leu Phe Arg Pro His Glu Gly Met Glu
1 5 10 15

Ala Pro Asp Asp Met Ala Ala Gln Pro Phe Phe Asp Pro Ser Ala Ser
20 25 30

Phe Pro Pro Ala Pro Ala Ser Ala Asn Leu Pro Lys Pro Asn Gly Gln
35 40 45

Thr Pro Pro Pro Thr Ser Asp Asp Leu Ser Glu Arg Phe Val Ser Ala
50 55 60

Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Thr Pro Met
65 70 75 80

Pro Ile Ala Ala Gly Glu Pro Pro Ser Pro Glu Pro Ala Ala Ser Lys
85 90 95

Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro Pro
100 105 110

Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro
115 120 125

Pro Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Ala Pro Thr
130 135 140

Pro Thr Glu Ser Gln Leu Ala Pro Pro Arg Pro Pro Thr Pro Gln Thr
145 150 155 160

Pro Thr Gly Ala Pro Gln Gln Pro Glu Ser Pro Ala Pro His Val Pro
165 170 175

Ser His Gly Pro His Gln Pro Arg Arg Thr Ala Pro Ala Pro Pro Trp
180 185 190

Ala Lys Met Pro Ile Gly Glu Pro Pro Pro Ala Pro Ser Arg Pro Ser
195 200 205

Ala Ser Pro Ala Glu Pro Pro Thr Arg Pro Ala Pro Gln His Ser Arg
210 215 220

Arg Ala Arg Arg Gly His Arg Tyr Arg Thr Asp Thr Glu Arg Asn Val
225 230 235 240

Gly Lys Val Ala Thr Gly Pro Ser Ile Gln Ala Arg Leu Arg Ala Glu
245 250 255

Glu Ala Ser Gly Ala Gln Leu Ala Pro Gly Thr Glu Pro Ser Pro Ala
260 265 270

Pro Leu Gly Gln Pro Arg Ser Tyr Leu Ala Pro Pro Thr Arg Pro Ala
275 280 285

Pro Thr Glu Pro Pro Pro Ser Pro Ser Pro Gln Arg Asn Ser Gly Arg
290 295 300

Arg Ala Glu Arg Arg Val His Pro Asp Leu Ala Ala Gln His Ala Ala
305 310 315 320

Ala Gln Pro Asp Ser Ile Thr Ala Ala Thr Thr Gly Gly Arg Arg Arg
325 330 335

Lys Arg Ala Ala Pro Asp Leu Asp Ala Thr Gln Lys Ser Leu Arg Pro

-continued

340	345	350
Ala Ala Lys Gly Pro Lys Val Lys Lys Val Lys Pro Gln Lys Pro Lys		
355	360	365
Ala Thr Lys Pro Pro Lys Val Val Ser Gln Arg Gly Trp Arg His Trp		
370	375	380
Val His Ala Leu Thr Arg Ile Asn Leu Gly Leu Ser Pro Asp Glu Lys		
385	390	395
Tyr Glu Leu Asp Leu His Ala Arg Val Arg Arg Asn Pro Arg Gly Ser		
405	410	415
Tyr Gln Ile Ala Val Val Gly Leu Lys Gly Gly Ala Gly Lys Thr Thr		
420	425	430
Leu Thr Ala Ala Leu Gly Ser Thr Leu Ala Gln Val Arg Ala Asp Arg		
435	440	445
Ile Leu Ala Leu Asp Ala Asp Pro Gly Ala Gly Asn Leu Ala Asp Arg		
450	455	460
Val Gly Arg Gln Ser Gly Ala Thr Ile Ala Asp Val Leu Ala Glu Lys		
465	470	475
Glu Leu Ser His Tyr Asn Asp Ile Arg Ala His Thr Ser Val Asn Ala		
485	490	495
Val Asn Leu Glu Val Leu Pro Ala Pro Glu Tyr Ser Ser Ala Gln Arg		
500	505	510
Ala Leu Ser Asp Ala Asp Trp His Phe Ile Ala Asp Pro Ala Ser Arg		
515	520	525
Phe Tyr Asn Leu Val Leu Ala Asp Cys Gly Ala Gly Phe Phe Asp Pro		
530	535	540
Leu Thr Arg Gly Val Leu Ser Thr Val Ser Gly Val Val Val Val Ala		
545	550	555
Ser Val Ser Ile Asp Gly Ala Gln Gln Ala Ser Val Ala Leu Asp Trp		
565	570	575
Leu Arg Asn Asn Gly Tyr Gln Asp Leu Ala Ser Arg Ala Cys Val Val		
580	585	590
Ile Asn His Ile Met Pro Gly Glu Pro Asn Val Ala Val Lys Asp Leu		
595	600	605
Val Arg His Phe Glu Gln Gln Val Gln Pro Gly Arg Val Val Val Met		
610	615	620
Pro Trp Asp Arg His Ile Ala Ala Gly Thr Glu Ile Ser Leu Asp Leu		
625	630	635
Leu Asp Pro Ile Tyr Lys Arg Lys Val Leu Glu Leu Ala Ala Ala Leu		
645	650	655
Ser Asp Asp Phe Glu Arg Ala Gly Arg Arg		
660	665	

<210> SEQ ID NO 6

<211> LENGTH: 511

<212> TYPE: PRT

<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 6

Leu Ser Ala Pro Ala Val Ala Ala Gly Pro Thr Ala Ala Gly Ala Thr		
1	5	10
Ala Ala Arg Pro Ala Thr Thr Arg Val Thr Ile Leu Thr Gly Arg Arg		
20	25	30
Met Thr Asp Leu Val Leu Pro Ala Ala Val Pro Met Glu Thr Tyr Ile		
35	40	45

-continued

Asp	Asp	Thr	Val	Ala	Val	Leu	Ser	Glu	Val	Leu	Glu	Asp	Thr	Pro	Ala	
50						55					60					
Asp	Val	Leu	Gly	Gly	Phe	Asp	Phe	Thr	Ala	Gln	Gly	Val	Trp	Ala	Phe	
65					70					75					80	
Ala	Arg	Pro	Gly	Ser	Pro	Pro	Leu	Lys	Leu	Asp	Gln	Ser	Leu	Asp	Asp	
				85					90					95		
Ala	Gly	Val	Val	Asp	Gly	Ser	Leu	Leu	Thr	Leu	Val	Ser	Val	Ser	Arg	
			100						105					110		
Thr	Glu	Arg	Tyr	Arg	Pro	Leu	Val	Glu	Asp	Val	Ile	Asp	Ala	Ile	Ala	
			115						120				125			
Val	Leu	Asp	Glu	Ser	Pro	Glu	Phe	Asp	Arg	Thr	Ala	Leu	Asn	Arg	Phe	
	130					135						140				
Val	Gly	Ala	Ala	Ile	Pro	Leu	Leu	Thr	Ala	Pro	Val	Ile	Gly	Met	Ala	
145					150					155					160	
Met	Arg	Ala	Trp	Trp	Glu	Thr	Gly	Arg	Ser	Leu	Trp	Trp	Pro	Leu	Ala	
			165						170					175		
Ile	Gly	Ile	Leu	Gly	Ile	Ala	Val	Leu	Val	Gly	Ser	Phe	Val	Ala	Asn	
			180						185					190		
Arg	Phe	Tyr	Gln	Ser	Gly	His	Leu	Ala	Glu	Cys	Leu	Leu	Val	Thr	Thr	
		195					200						205			
Tyr	Leu	Leu	Ile	Ala	Thr	Ala	Ala	Ala	Leu	Ala	Val	Pro	Leu	Pro	Arg	
	210					215					220					
Gly	Val	Asn	Ser	Leu	Gly	Ala	Pro	Gln	Val	Ala	Gly	Ala	Ala	Thr	Ala	
225					230					235					240	
Val	Leu	Phe	Leu	Thr	Leu	Met	Thr	Arg	Gly	Gly	Pro	Arg	Lys	Arg	His	
			245						250					255		
Glu	Leu	Ala	Ser	Phe	Ala	Val	Ile	Thr	Ala	Ile	Ala	Val	Ile	Ala	Ala	
			260						265					270		
Ala	Ala	Ala	Phe	Gly	Tyr	Gly	Tyr	Gln	Asp	Trp	Val	Pro	Ala	Gly	Gly	
		275					280					285				
Ile	Ala	Phe	Gly	Leu	Phe	Ile	Val	Thr	Asn	Ala	Ala	Lys	Leu	Thr	Val	
	290					295						300				
Ala	Val	Ala	Arg	Ile	Ala	Leu	Pro	Pro	Ile	Pro	Val	Pro	Gly	Glu	Thr	
305					310					315					320	
Val	Asp	Asn	Glu	Glu	Leu	Leu	Asp	Pro	Val	Ala	Thr	Pro	Glu	Ala	Thr	
			325						330					335		
Ser	Glu	Glu	Thr	Pro	Thr	Trp	Gln	Ala	Ile	Ile	Ala	Ser	Val	Pro	Ala	
			340					345					350			
Ser	Ala	Val	Arg	Leu	Thr	Glu	Arg	Ser	Lys	Leu	Ala	Lys	Gln	Leu	Leu	
		355					360					365				
Ile	Gly	Tyr	Val	Thr	Ser	Gly	Thr	Leu	Ile	Leu	Ala	Ala	Gly	Ala	Ile	
	370					375						380				
Ala	Val	Val	Val	Arg	Gly	His	Phe	Phe	Val	His	Ser	Leu	Val	Val	Ala	
385					390					395					400	
Gly	Leu	Ile	Thr	Thr	Val	Cys	Gly	Phe	Arg	Ser	Arg	Leu	Tyr	Ala	Glu	
			405					410						415		
Arg	Trp	Cys	Ala	Trp	Ala	Leu	Leu	Ala	Ala	Thr	Val	Ala	Ile	Pro	Thr	
			420					425					430			
Gly	Leu	Thr	Ala	Lys	Leu	Ile	Ile	Trp	Tyr	Pro	His	Tyr	Ala	Trp	Leu	
		435				440						445				
Leu	Leu	Ser	Val	Tyr	Leu	Thr	Val	Ala	Leu	Val	Ala	Leu	Val	Val	Val	
	450					455					460					
Gly	Ser	Met	Ala	His	Val	Arg	Arg	Val	Ser	Pro	Val	Val	Lys	Arg	Thr	

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465	470	475	480
Leu Glu Leu Ile Asp Gly Ala Met Ile Ala Ala Ile Ile Pro Met Leu			
	485	490	495
Leu Trp Ile Thr Gly Val Tyr Asp Thr Val Arg Asn Ile Arg Phe			
	500	505	510

<210> SEQ ID NO 7
 <211> LENGTH: 280
 <212> TYPE: PRT
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 7

Met Ala Glu Pro Leu Ala Val Asp Pro Thr Gly Leu Ser Ala Ala Ala			
1	5	10	15
Ala Lys Leu Ala Gly Leu Val Phe Pro Gln Pro Pro Ala Pro Ile Ala			
	20	25	30
Val Ser Gly Thr Asp Ser Val Val Ala Ala Ile Asn Glu Thr Met Pro			
	35	40	45
Ser Ile Glu Ser Leu Val Ser Asp Gly Leu Pro Gly Val Lys Ala Ala			
	50	55	60
Leu Thr Arg Thr Ala Ser Asn Met Asn Ala Ala Ala Asp Val Tyr Ala			
65	70	75	80
Lys Thr Asp Gln Ser Leu Gly Thr Ser Leu Ser Gln Tyr Ala Phe Gly			
	85	90	95
Ser Ser Gly Glu Gly Leu Ala Gly Val Ala Ser Val Gly Gly Gln Pro			
	100	105	110
Ser Gln Ala Thr Gln Leu Leu Ser Thr Pro Val Ser Gln Val Thr Thr			
	115	120	125
Gln Leu Gly Glu Thr Ala Ala Glu Leu Ala Pro Arg Val Val Ala Thr			
	130	135	140
Val Pro Gln Leu Val Gln Leu Ala Pro His Ala Val Gln Met Ser Gln			
145	150	155	160
Asn Ala Ser Pro Ile Ala Gln Thr Ile Ser Gln Thr Ala Gln Gln Ala			
	165	170	175
Ala Gln Ser Ala Gln Gly Gly Ser Gly Pro Met Pro Ala Gln Leu Ala			
	180	185	190
Ser Ala Glu Lys Pro Ala Thr Glu Gln Ala Glu Pro Val His Glu Val			
	195	200	205
Thr Asn Asp Asp Gln Gly Asp Gln Gly Asp Val Gln Pro Ala Glu Val			
	210	215	220
Val Ala Ala Ala Arg Asp Glu Gly Ala Gly Ala Ser Pro Gly Gln Gln			
225	230	235	240
Pro Gly Gly Gly Val Pro Ala Gln Ala Met Asp Thr Gly Ala Gly Ala			
	245	250	255
Arg Pro Ala Ala Ser Pro Leu Ala Ala Pro Val Asp Pro Ser Thr Pro			
	260	265	270
Ala Pro Ser Thr Thr Thr Thr Leu			
	275	280	

<210> SEQ ID NO 8
 <211> LENGTH: 729
 <212> TYPE: PRT
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 8

Met Ser Ile Thr Arg Pro Thr Gly Ser Tyr Ala Arg Gln Met Leu Asp

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1	5	10	15
Pro Gly Gly Trp Val Glu Ala Asp Glu Asp Thr Phe Tyr Asp Arg Ala	20	25	30
Gln Glu Tyr Ser Gln Val Leu Gln Arg Val Thr Asp Val Leu Asp Thr	35	40	45
Cys Arg Gln Gln Lys Gly His Val Phe Glu Gly Gly Leu Trp Ser Gly	50	55	60
Gly Ala Ala Asn Ala Ala Asn Gly Ala Leu Gly Ala Asn Ile Asn Gln	65	70	75
Leu Met Thr Leu Gln Asp Tyr Leu Ala Thr Val Ile Thr Trp His Arg	85	90	95
His Ile Ala Gly Leu Ile Glu Gln Ala Lys Ser Asp Ile Gly Asn Asn	100	105	110
Val Asp Gly Ala Gln Arg Glu Ile Asp Ile Leu Glu Asn Asp Pro Ser	115	120	125
Leu Asp Ala Asp Glu Arg His Thr Ala Ile Asn Ser Leu Val Thr Ala	130	135	140
Thr His Gly Ala Asn Val Ser Leu Val Ala Glu Thr Ala Glu Arg Val	145	150	155
Leu Glu Ser Lys Asn Trp Lys Pro Pro Lys Asn Ala Leu Glu Asp Leu	165	170	175
Leu Gln Gln Lys Ser Pro Pro Pro Pro Asp Val Pro Thr Leu Val Val	180	185	190
Pro Ser Pro Gly Thr Pro Gly Thr Pro Gly Thr Pro Ile Thr Pro Gly	195	200	205
Thr Pro Ile Thr Pro Gly Thr Pro Ile Thr Pro Ile Pro Gly Ala Pro	210	215	220
Val Thr Pro Ile Thr Pro Thr Pro Gly Thr Pro Val Thr Pro Val Thr	225	230	235
Pro Gly Lys Pro Val Thr Pro Val Thr Pro Val Lys Pro Gly Thr Pro	245	250	255
Gly Glu Pro Thr Pro Ile Thr Pro Val Thr Pro Pro Val Ala Pro Ala	260	265	270
Thr Pro Ala Thr Pro Ala Thr Pro Val Thr Pro Ala Pro Ala Pro His	275	280	285
Pro Gln Pro Ala Pro Ala Pro Ala Pro Ser Pro Gly Pro Gln Pro Val	290	295	300
Thr Pro Ala Thr Pro Gly Pro Ser Gly Pro Ala Thr Pro Gly Thr Pro	305	310	315
Gly Gly Glu Pro Ala Pro His Val Lys Pro Ala Ala Leu Ala Glu Gln	325	330	335
Pro Gly Val Pro Gly Gln His Ala Gly Gly Gly Thr Gln Ser Gly Pro	340	345	350
Ala His Ala Asp Glu Ser Ala Ala Ser Val Thr Pro Ala Ala Ala Ser	355	360	365
Gly Val Pro Gly Ala Arg Ala Ala Ala Ala Ala Pro Ser Gly Thr Ala	370	375	380
Val Gly Ala Gly Ala Arg Ser Ser Val Gly Thr Ala Ala Ala Ser Gly	385	390	395
Ala Gly Ser His Ala Ala Thr Gly Arg Ala Pro Val Ala Thr Ser Asp	405	410	415
Lys Ala Ala Ala Pro Ser Thr Arg Ala Ala Ser Ala Arg Thr Ala Pro	420	425	430

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Pro Ala Arg Pro Pro Ser Thr Asp His Ile Asp Lys Pro Asp Arg Ser
 435 440 445
 Glu Ser Ala Asp Asp Gly Thr Pro Val Ser Met Ile Pro Val Ser Ala
 450 455 460
 Ala Arg Ala Ala Arg Asp Ala Ala Thr Ala Ala Ala Ser Ala Arg Gln
 465 470 475 480
 Arg Gly Arg Gly Asp Ala Leu Arg Leu Ala Arg Arg Ile Ala Ala Ala
 485 490 495
 Leu Asn Ala Ser Asp Asn Asn Ala Gly Asp Tyr Gly Phe Phe Trp Ile
 500 505 510
 Thr Ala Val Thr Thr Asp Gly Ser Ile Val Val Ala Asn Ser Tyr Gly
 515 520 525
 Leu Ala Tyr Ile Pro Asp Gly Met Glu Leu Pro Asn Lys Val Tyr Leu
 530 535 540
 Ala Ser Ala Asp His Ala Ile Pro Val Asp Glu Ile Ala Arg Cys Ala
 545 550 555 560
 Thr Tyr Pro Val Leu Ala Val Gln Ala Trp Ala Ala Phe His Asp Met
 565 570 575
 Thr Leu Arg Ala Val Ile Gly Thr Ala Glu Gln Leu Ala Ser Ser Asp
 580 585 590
 Pro Gly Val Ala Lys Ile Val Leu Glu Pro Asp Asp Ile Pro Glu Ser
 595 600 605
 Gly Lys Met Thr Gly Arg Ser Arg Leu Glu Val Val Asp Pro Ser Ala
 610 615 620
 Ala Ala Gln Leu Ala Asp Thr Thr Asp Gln Arg Leu Leu Asp Leu Leu
 625 630 635 640
 Pro Pro Ala Pro Val Asp Val Asn Pro Pro Gly Asp Glu Arg His Met
 645 650 655
 Leu Trp Phe Glu Leu Met Lys Pro Met Thr Ser Thr Ala Thr Gly Arg
 660 665 670
 Glu Ala Ala His Leu Arg Ala Phe Arg Ala Tyr Ala Ala His Ser Gln
 675 680 685
 Glu Ile Ala Leu His Gln Ala His Thr Ala Thr Asp Ala Ala Val Gln
 690 695 700
 Arg Val Ala Val Ala Asp Trp Leu Tyr Trp Gln Tyr Val Thr Gly Leu
 705 710 715 720
 Leu Asp Arg Ala Leu Ala Ala Ala Cys
 725

<210> SEQ ID NO 9
 <211> LENGTH: 1776
 <212> TYPE: DNA
 <213> ORGANISM: Mycobacterium tuberculosis
 <220> FEATURE:
 <221> NAME/KEY: CDS
 <222> LOCATION: (1)...(1773)

<400> SEQUENCE: 9

atg act gct gaa ccg gaa gta cgg acg ctg cgc gag gtt gtg ctg gac	48
Met Thr Ala Glu Pro Glu Val Arg Thr Leu Arg Glu Val Val Leu Asp	
1 5 10 15	
cag ctc ggc act gct gaa tcg cgt gcg tac aag atg tgg ctg ccg ccg	96
Gln Leu Gly Thr Ala Glu Ser Arg Ala Tyr Lys Met Trp Leu Pro Pro	
20 25 30	
ttg acc aat ccg gtc ccg ctc aac gag ctc atc gcc cgt gat cgg cga	144
Leu Thr Asn Pro Val Pro Leu Asn Glu Leu Ile Ala Arg Asp Arg Arg	

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35	40	45	
caa ccc ctg cga ttt gcc ctg ggg atc atg gat gaa ccg cgc cgc cat Gln Pro Leu Arg Phe Ala Leu Gly Ile Met Asp Glu Pro Arg Arg His 50 55 60			192
cta cag gat gtg tgg ggc gta gac gtt tcc ggg gcc ggc ggc aac atc Leu Gln Asp Val Trp Gly Val Asp Val Ser Gly Ala Gly Gly Asn Ile 65 70 75 80			240
ggg att ggg ggc gca cct caa acc ggg aag tcg acg cta ctg cag acg Gly Ile Gly Gly Ala Pro Gln Thr Gly Lys Ser Thr Leu Leu Gln Thr 85 90 95			288
atg gtg atg tcg gcc gcc gcc aca cac tca ccg cgc aac gtt cag ttc Met Val Met Ser Ala Ala Ala Thr His Ser Pro Arg Asn Val Gln Phe 100 105 110			336
tat tgc atc gac cta ggt ggc ggc ggg ctg atc tat ctc gaa aac ctt Tyr Cys Ile Asp Leu Gly Gly Gly Gly Leu Ile Tyr Leu Glu Asn Leu 115 120 125			384
cca cac gtc ggt ggg gta gcc aat cgg tcc gag ccc gac aag gtc aac Pro His Val Gly Gly Val Ala Asn Arg Ser Glu Pro Asp Lys Val Asn 130 135 140			432
cgg gtg gtc gca gag atg caa gcc gtc atg cgg caa cgg gaa acc acc Arg Val Val Ala Glu Met Gln Ala Val Met Arg Gln Arg Glu Thr Thr 145 150 155 160			480
ttc aag gaa cac cga gtg ggc tcg atc ggg atg tac cgg cag ctg cgt Phe Lys Glu His Arg Val Gly Ser Ile Gly Met Tyr Arg Gln Leu Arg 165 170 175			528
gac gat cca agt caa ccc gtt gcg tcc gat cca tac ggc gac gtc ttt Asp Asp Pro Ser Gln Pro Val Ala Ser Asp Pro Tyr Gly Asp Val Phe 180 185 190			576
ctg atc atc gac gga tgg ccc ggt ttt gtc ggc gag ttc ccc gac ctt Leu Ile Ile Asp Gly Trp Pro Gly Phe Val Gly Glu Phe Pro Asp Leu 195 200 205			624
gag ggg cag gtt caa gat ctg gcc gcc cag ggg ctg gcg ttc ggc gtc Glu Gly Gln Val Gln Asp Leu Ala Ala Gln Gly Leu Ala Phe Gly Val 210 215 220			672
cac gtc atc atc tcc acg cca cgc tgg aca gag ctg aag tcg cgt gtt His Val Ile Ile Ser Thr Pro Arg Trp Thr Glu Leu Lys Ser Arg Val 225 230 235 240			720
cgc gac tac ctc ggc acc aag atc gag ttc cgg ctt ggt gac gtc aat Arg Asp Tyr Leu Gly Thr Lys Ile Glu Phe Arg Leu Gly Asp Val Asn 245 250 255			768
gaa acc cag atc gac cgg att acc cgc gag atc ccg gcg aat cgt ccg Glu Thr Gln Ile Asp Arg Ile Thr Arg Glu Ile Pro Ala Asn Arg Pro 260 265 270			816
ggg cgg gca gtg tcg atg gaa aag cac cat ctg atg atc ggc gtg ccc Gly Arg Ala Val Ser Met Glu Lys His His Leu Met Ile Gly Val Pro 275 280 285			864
agg ttc gac ggc gtg cac agc gcc gat aac ctg gtg gag gcg atc acc Arg Phe Asp Gly Val His Ser Ala Asp Asn Leu Val Glu Ala Ile Thr 290 295 300			912
gcg ggg gtg acg cag atc gct tcc cag cac acc gaa cag gca cct ccg Ala Gly Val Thr Gln Ile Ala Ser Gln His Thr Glu Gln Ala Pro Pro 305 310 315 320			960
gtg cgg gtc ctg ccg gag cgt atc cac ctg cac gaa ctc gac ccg aac Val Arg Val Leu Pro Glu Arg Ile His Leu His Glu Leu Asp Pro Asn 325 330 335			1008
ccg ccg gga cca gag tcc gac tac cgc act cgc tgg gag att ccg atc Pro Pro Gly Pro Glu Ser Asp Tyr Arg Thr Arg Trp Glu Ile Pro Ile 340 345 350			1056
ggc ttg cgc gag acg gac ctg acg ccg gct cac tgc cac atg cac acg			1104

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Gly	Leu	Arg	Glu	Thr	Asp	Leu	Thr	Pro	Ala	His	Cys	His	Met	His	Thr	
	355						360					365				
aac	ccg	cac	cta	ctg	atc	ttc	ggg	gcg	gcc	aaa	tcg	ggc	aag	acg	acc	1152
Asn	Pro	His	Leu	Leu	Ile	Phe	Gly	Ala	Ala	Lys	Ser	Gly	Lys	Thr	Thr	
	370					375					380					
att	gcc	cac	gcg	atc	gcg	cgc	gcc	att	tgt	gcc	cga	aac	agt	ccc	cag	1200
Ile	Ala	His	Ala	Ile	Ala	Arg	Ala	Ile	Cys	Ala	Arg	Asn	Ser	Pro	Gln	
	385				390				395					400		
cag	gtg	cgg	ttc	atg	ctc	gcg	gac	tac	cgc	tcg	ggc	ctg	ctg	gac	gcg	1248
Gln	Val	Arg	Phe	Met	Leu	Ala	Asp	Tyr	Arg	Ser	Gly	Leu	Leu	Asp	Ala	
			405					410						415		
gtg	ccg	gac	acc	cat	ctg	ctg	ggc	gcc	ggc	gcg	atc	aac	cgc	aac	agc	1296
Val	Pro	Asp	Thr	His	Leu	Leu	Gly	Ala	Gly	Ala	Ile	Asn	Arg	Asn	Ser	
			420				425						430			
gcg	tcg	cta	gac	gag	gcc	gtt	caa	gca	ctg	gcg	gtc	aac	ctg	aag	aag	1344
Ala	Ser	Leu	Asp	Glu	Ala	Val	Gln	Ala	Leu	Ala	Val	Asn	Leu	Lys	Lys	
		435				440					445					
cgg	ttg	ccg	ccg	acc	gac	ctg	acg	acg	gcg	cag	cta	cgc	tcg	cgt	tcg	1392
Arg	Leu	Pro	Pro	Thr	Asp	Leu	Thr	Thr	Ala	Gln	Leu	Arg	Ser	Arg	Ser	
	450					455					460					
tgg	tgg	agc	gga	ttt	gac	gtc	gtg	ctt	ctg	gtc	gac	gat	tgg	cac	atg	1440
Trp	Trp	Ser	Gly	Phe	Asp	Val	Val	Leu	Leu	Val	Asp	Asp	Trp	His	Met	
	465			470				475						480		
atc	gtg	ggg	gcc	gcc	ggg	ggg	atg	ccg	ccg	atg	gca	ccg	ctg	gcc	ccg	1488
Ile	Val	Gly	Ala	Ala	Gly	Gly	Met	Pro	Pro	Met	Ala	Pro	Leu	Ala	Pro	
			485				490							495		
tta	ttg	ccg	gcg	gcg	gca	gat	atc	ggg	ttg	cac	atc	att	gtc	acc	tgt	1536
Leu	Leu	Pro	Ala	Ala	Ala	Asp	Ile	Gly	Leu	His	Ile	Ile	Val	Thr	Cys	
			500				505						510			
cag	atg	agc	cag	gct	tac	aag	gca	acc	atg	gac	aag	ttc	gtc	ggc	gcc	1584
Gln	Met	Ser	Gln	Ala	Tyr	Lys	Ala	Thr	Met	Asp	Lys	Phe	Val	Gly	Ala	
		515					520					525				
gca	ttc	ggg	tcg	ggc	gct	ccg	aca	atg	ttc	ctt	tcg	ggc	gag	aag	cag	1632
Ala	Phe	Gly	Ser	Gly	Ala	Pro	Thr	Met	Phe	Leu	Ser	Gly	Glu	Lys	Gln	
	530					535					540					
gaa	ttc	cca	tcc	agt	gag	ttc	aag	gtc	aag	cgg	cgc	ccc	cct	ggc	cag	1680
Glu	Phe	Pro	Ser	Ser	Glu	Phe	Lys	Val	Lys	Arg	Arg	Pro	Pro	Gly	Gln	
	545				550				555					560		
gca	ttt	ctc	gtc	tcg	cca	gac	ggc	aaa	gag	gtc	atc	cag	gcc	ccc	tac	1728
Ala	Phe	Leu	Val	Ser	Pro	Asp	Gly	Lys	Glu	Val	Ile	Gln	Ala	Pro	Tyr	
			565				570							575		
atc	gag	cct	cca	gaa	gaa	gtg	ttc	gca	gca	ccc	cca	agc	gcc	ggg		1773
Ile	Glu	Pro	Pro	Glu	Glu	Val	Phe	Ala	Ala	Pro	Pro	Ser	Ala	Gly		
			580				585					590				
taa																1776
<210> SEQ ID NO 10																
<211> LENGTH: 300																
<212> TYPE: DNA																
<213> ORGANISM: Mycobacterium tuberculosis																
<220> FEATURE:																
<221> NAME/KEY: CDS																
<222> LOCATION: (1)...(297)																
<400> SEQUENCE: 10																
atg	gaa	aaa	atg	tca	cat	gat	ccg	atc	gct	gcc	gac	att	ggc	acg	caa	48
Met	Glu	Lys	Met	Ser	His	Asp	Pro	Ile	Ala	Ala	Asp	Ile	Gly	Thr	Gln	
1			5					10					15			
gtg	agc	gac	aac	gct	ctg	cac	ggc	gtg	acg	gcc	ggc	tcg	acg	gcg	ctg	96
Val	Ser	Asp	Asn	Ala	Leu	His	Gly	Val	Thr	Ala	Gly	Ser	Thr	Ala	Leu	
		20					25					30				

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acg tcg gtg acc ggg ctg gtt ccc gcg ggg gcc gat gag gtc tcc gcc Thr Ser Val Thr Gly Leu Val Pro Ala Gly Ala Asp Glu Val Ser Ala 35 40 45	144
caa gcg gcg acg gcg ttc aca tcg gag ggc atc caa ttg ctg gct tcc Gln Ala Ala Thr Ala Phe Thr Ser Glu Gly Ile Gln Leu Leu Ala Ser 50 55 60	192
aat gca tcg gcc caa gac cag ctc cac cgt gcg ggc gaa gcg gtc cag Asn Ala Ser Ala Gln Asp Gln Leu His Arg Ala Gly Glu Ala Val Gln 65 70 75 80	240
gac gtc gcc cgc acc tat tcg caa atc gac gac ggc gcc gcc ggc gtc Asp Val Ala Arg Thr Tyr Ser Gln Ile Asp Asp Gly Ala Ala Gly Val 85 90 95	288
ttc gcc gaa tag Phe Ala Glu	300
 <210> SEQ ID NO 11 <211> LENGTH: 1107 <212> TYPE: DNA <213> ORGANISM: Mycobacterium tuberculosis <220> FEATURE: <221> NAME/KEY: CDS <222> LOCATION: (1)...(1104) <400> SEQUENCE: 11	
atg ctg tgg cac gca atg cca ccg gag cta aat acc gca cgg ctg atg Met Leu Trp His Ala Met Pro Pro Glu Leu Asn Thr Ala Arg Leu Met 1 5 10 15	48
gcc ggc gcg ggt ccg gct cca atg ctt gcg gcg gcc gcg gga tgg cag Ala Gly Ala Gly Pro Ala Pro Met Leu Ala Ala Ala Ala Gly Trp Gln 20 25 30	96
acg ctt tcg gcg gct ctg gac gct cag gcc gtc gag ttg acc gcg cgc Thr Leu Ser Ala Ala Leu Asp Ala Gln Ala Val Glu Leu Thr Ala Arg 35 40 45	144
ctg aac tct ctg gga gaa gcc tgg act gga ggt ggc agc gac aag gcg Leu Asn Ser Leu Gly Glu Ala Trp Thr Gly Gly Gly Ser Asp Lys Ala 50 55 60	192
ctt gcg gct gca acg ccg atg gtg gtc tgg cta caa acc gcg tca aca Leu Ala Ala Ala Thr Pro Met Val Val Trp Leu Gln Thr Ala Ser Thr 65 70 75 80	240
cag gcc aag acc cgt gcg atg cag gcg acg gcg caa gcc gcg gca tac Gln Ala Lys Thr Arg Ala Met Gln Ala Thr Ala Gln Ala Ala Ala Tyr 85 90 95	288
acc cag gcc atg gcc acg acg ccg tcg ctg ccg gag atc gcc gcc aac Thr Gln Ala Met Ala Thr Thr Pro Ser Leu Pro Glu Ile Ala Ala Asn 100 105 110	336
cac atc acc cag gcc gtc ctt acg gcc acc aac ttc ttc ggt atc aac His Ile Thr Gln Ala Val Leu Thr Ala Thr Asn Phe Phe Gly Ile Asn 115 120 125	384
acg atc ccg atc gcg ttg acc gag atg gat tat ttc atc cgt atg tgg Thr Ile Pro Ile Ala Leu Thr Glu Met Asp Tyr Phe Ile Arg Met Trp 130 135 140	432
aac cag gca gcc ctg gca atg gag gtc tac cag gcc gag acc gcg gtt Asn Gln Ala Ala Leu Ala Met Glu Val Tyr Gln Ala Glu Thr Ala Val 145 150 155 160	480
aac acg ctt ttc gag aag ctc gag ccg atg gcg tcg atc ctt gat ccc Asn Thr Leu Phe Glu Lys Leu Glu Pro Met Ala Ser Ile Leu Asp Pro 165 170 175	528
ggc gcg agc cag agc acg acg aac ccg atc ttc gga atg ccc tcc cct Gly Ala Ser Gln Ser Thr Thr Asn Pro Ile Phe Gly Met Pro Ser Pro 180 185 190	576

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ggc agc tca aca ccg gtt ggc cag ttg ccg ccg gcg gct acc cag acc Gly Ser Ser Thr Pro Val Gly Gln Leu Pro Pro Ala Ala Thr Gln Thr 195 200 205	624
ctc ggc caa ctg ggt gag atg agc ggc ccg atg cag cag ctg acc cag Leu Gly Gln Leu Gly Glu Met Ser Gly Pro Met Gln Gln Leu Thr Gln 210 215 220	672
ccg ctg cag cag gtg acg tcg ttg ttc agc cag gtg ggc ggc acc ggc Pro Leu Gln Gln Val Thr Ser Leu Phe Ser Gln Val Gly Gly Thr Gly 225 230 235 240	720
ggc ggc aac cca gcc gac gag gaa gcc gcg cag atg ggc ctg ctc ggc Gly Gly Asn Pro Ala Asp Glu Glu Ala Ala Gln Met Gly Leu Leu Gly 245 250 255	768
acc agt ccg ctg tcg aac cat ccg ctg gct ggt gga tca ggc ccc agc Thr Ser Pro Leu Ser Asn His Pro Leu Ala Gly Gly Ser Gly Pro Ser 260 265 270	816
gcg ggc gcg ggc ctg ctg cgc gcg gag tcg cta cct ggc gca ggt ggg Ala Gly Ala Gly Leu Leu Arg Ala Glu Ser Leu Pro Gly Ala Gly Gly 275 280 285	864
tcg ttg acc cgc acg ccg ctg atg tct cag ctg atc gaa aag ccg gtt Ser Leu Thr Arg Thr Pro Leu Met Ser Gln Leu Ile Glu Lys Pro Val 290 295 300	912
gcc ccc tcg gtg atg ccg gcg gct gct gcc gga tcg tcg gcg acg ggt Ala Pro Ser Val Met Pro Ala Ala Ala Gly Ser Ser Ala Thr Gly 305 310 315 320	960
ggc gcc gct ccg gtg ggt gcg gga gcg atg ggc cag ggt gcg caa tcc Gly Ala Ala Pro Val Gly Ala Gly Ala Met Gly Gln Gly Ala Gln Ser 325 330 335	1008
ggc ggc tcc acc agg ccg ggt ctg gtc gcg ccg gca ccg ctc gcg cag Gly Gly Ser Thr Arg Pro Gly Leu Val Ala Pro Ala Pro Leu Ala Gln 340 345 350	1056
gag cgt gaa gaa gac gac gag gac gac tgg gac gaa gag gac gac tgg Glu Arg Glu Glu Asp Asp Glu Asp Asp Trp Asp Glu Glu Asp Asp Trp 355 360 365	1104
tga	1107
<210> SEQ ID NO 12 <211> LENGTH: 303 <212> TYPE: DNA <213> ORGANISM: Mycobacterium tuberculosis <220> FEATURE: <221> NAME/KEY: CDS <222> LOCATION: (1)...(300) <400> SEQUENCE: 12	
atg gca gag atg aag acc gat gcc gct acc ctc gcg cag gag gca ggt Met Ala Glu Met Lys Thr Asp Ala Ala Thr Leu Ala Gln Glu Ala Gly 1 5 10 15	48
aat ttc gag ccg atc tcc ggc gac ctg aaa acc cag atc gac cag gtg Asn Phe Glu Arg Ile Ser Gly Asp Leu Lys Thr Gln Ile Asp Gln Val 20 25 30	96
gag tcg acg gca ggt tcg ttg cag ggc cag tgg cgc ggc gcg gcg ggg Glu Ser Thr Ala Gly Ser Leu Gln Gly Gln Trp Arg Gly Ala Ala Gly 35 40 45	144
acg gcc gcc cag gcc gcg gtg gtg cgc ttc caa gaa gca gcc aat aag Thr Ala Ala Gln Ala Ala Val Val Arg Phe Gln Glu Ala Ala Asn Lys 50 55 60	192
cag aag cag gaa ctc gac gag atc tcg acg aat att cgt cag gcc ggc Gln Lys Gln Glu Leu Asp Glu Ile Ser Thr Asn Ile Arg Gln Ala Gly 65 70 75 80	240
gtc caa tac tcg agg gcc gac gag gag cag cag cag gcg ctg tcc tcg Val Gln Tyr Ser Arg Ala Asp Glu Glu Gln Gln Gln Ala Leu Ser Ser	288

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85	90	95	
caa atg ggc ttc tga			303
Gln Met Gly Phe			
100			
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<212> TYPE: DNA			
<213> ORGANISM: Mycobacterium tuberculosis			
<220> FEATURE:			
<221> NAME/KEY: CDS			
<222> LOCATION: (1)...(1998)			
<400> SEQUENCE: 13			
atg gcg gcc gac tac gac aag ctc ttc cgg ccg cac gaa ggt atg gaa			48
Met Ala Ala Asp Tyr Asp Lys Leu Phe Arg Pro His Glu Gly Met Glu			
1 5 10 15			
gct ccg gac gat atg gca gcg cag ccg ttc ttc gac ccc agt gct tcg			96
Ala Pro Asp Asp Met Ala Ala Gln Pro Phe Phe Asp Pro Ser Ala Ser			
20 25 30			
ttt ccg ccg gcg ccc gca tcg gca aac cta ccg aag ccc aac ggc cag			144
Phe Pro Pro Ala Pro Ala Ser Ala Asn Leu Pro Lys Pro Asn Gly Gln			
35 40 45			
act ccg ccc ccg acg tcc gac gac ctg tcg gag ccg ttc gtg tcg gcc			192
Thr Pro Pro Pro Thr Ser Asp Leu Ser Glu Arg Phe Val Ser Ala			
50 55 60			
ccg ccg ccg cca ccc cca ccc cca cct ccg cct ccg cca act ccg atg			240
Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Thr Pro Met			
65 70 75 80			
ccg atc gcc gca gga gag ccg ccc tcg ccg gaa ccg gcc gca tct aaa			288
Pro Ile Ala Ala Gly Glu Pro Pro Ser Pro Glu Pro Ala Ala Ser Lys			
85 90 95			
cca ccc aca ccc ccc atg ccc atc gcc gga ccc gaa ccg gcc cca ccc			336
Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro Pro			
100 105 110			
aaa cca ccc aca ccc ccc atg ccc atc gcc gga ccc gaa ccg gcc cca			384
Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro			
115 120 125			
ccc aaa cca ccc aca cct ccg atg ccc atc gcc gga cct gca ccc acc			432
Pro Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Ala Pro Thr			
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Pro Thr Glu Ser Gln Leu Ala Pro Pro Arg Pro Pro Thr Pro Gln Thr			
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Pro Thr Gly Ala Pro Gln Gln Pro Glu Ser Pro Ala Pro His Val Pro			
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Ser His Gly Pro His Gln Pro Arg Arg Thr Ala Pro Ala Pro Pro Trp			
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Ala Lys Met Pro Ile Gly Glu Pro Pro Pro Ala Pro Ser Arg Pro Ser			
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Ala Ser Pro Ala Glu Pro Pro Thr Arg Pro Ala Pro Gln His Ser Arg			
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Arg Ala Arg Arg Gly His Arg Tyr Arg Thr Asp Thr Glu Arg Asn Val			
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Gly Lys Val Ala Thr Gly Pro Ser Ile Gln Ala Arg Leu Arg Ala Glu			

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gcg gcc aag ggg ccg aag gtg aag aag gtg aag ccc cag aaa ccg aag Ala Ala Lys Gly Pro Lys Val Lys Lys Val Lys Pro Gln Lys Pro Lys 355 360 365			1104
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Ile Asn His Ile Met Pro Gly Glu Pro Asn Val Ala Val Lys Asp Leu	
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Val Arg His Phe Glu Gln Gln Val Gln Pro Gly Arg Val Val Val Met	
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Pro Trp Asp Arg His Ile Ala Ala Gly Thr Glu Ile Ser Leu Asp Leu	
625 630 635 640	
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Leu Asp Pro Ile Tyr Lys Arg Lys Val Leu Glu Leu Ala Ala Ala Leu	
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Met Thr Asp Leu Val Leu Pro Ala Ala Val Pro Met Glu Thr Tyr Ile	
35 40 45	
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Asp Asp Thr Val Ala Val Leu Ser Glu Val Leu Glu Asp Thr Pro Ala	
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Ala Arg Pro Gly Ser Pro Pro Leu Lys Leu Asp Gln Ser Leu Asp Asp	
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Ala Gly Val Val Asp Gly Ser Leu Leu Thr Leu Val Ser Val Ser Arg	
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Thr Glu Arg Tyr Arg Pro Leu Val Glu Asp Val Ile Asp Ala Ile Ala	
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Val Leu Asp Glu Ser Pro Glu Phe Asp Arg Thr Ala Leu Asn Arg Phe	
130 135 140	
gtg ggg gcg gcg atc ccg ctt ttg acc gcg ccc gtc atc ggg atg gcg	480
Val Gly Ala Ala Ile Pro Leu Leu Thr Ala Pro Val Ile Gly Met Ala	
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Ile	Gly	Ile	Leu	Gly	Ile	Ala	Val	Leu	Val	Gly	Ser	Phe	Val	Ala	Asn		
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Arg	Phe	Tyr	Gln	Ser	Gly	His	Leu	Ala	Glu	Cys	Leu	Leu	Val	Thr	Thr		
		195					200					205					
tat	ctg	ctg	atc	gca	acc	gcc	gca	gcg	ctg	gcc	gtg	ccg	ttg	ccg	cgc	672	
Tyr	Leu	Leu	Ile	Ala	Thr	Ala	Ala	Ala	Leu	Ala	Val	Pro	Leu	Pro	Arg		
	210					215					220						
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Gly	Val	Asn	Ser	Leu	Gly	Ala	Pro	Gln	Val	Ala	Gly	Ala	Ala	Thr	Ala		
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gtg	ctg	ttt	ttg	acc	ttg	atg	acg	cgg	ggc	ggc	cct	cgg	aag	cgt	cat	768	
Val	Leu	Phe	Leu	Thr	Leu	Met	Thr	Arg	Gly	Gly	Pro	Arg	Lys	Arg	His		
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Glu	Leu	Ala	Ser	Phe	Ala	Val	Ile	Thr	Ala	Ile	Ala	Val	Ile	Ala	Ala		
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gcc	gct	gcc	ttc	ggc	tat	gga	tac	cag	gac	tgg	gtc	ccc	gcg	ggg	ggg	864	
Ala	Ala	Ala	Phe	Gly	Tyr	Gly	Tyr	Gln	Asp	Trp	Val	Pro	Ala	Gly	Gly		
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Ile	Ala	Phe	Gly	Leu	Phe	Ile	Val	Thr	Asn	Ala	Ala	Lys	Leu	Thr	Val		
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Val	Asp	Asn	Glu	Glu	Leu	Leu	Asp	Pro	Val	Ala	Thr	Pro	Glu	Ala	Thr		
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agc	gaa	gaa	acc	ccg	acc	tgg	cag	gcc	atc	atc	gcg	tcg	gtg	ccc	gcg	1056	
Ser	Glu	Glu	Thr	Pro	Thr	Trp	Gln	Ala	Ile	Ile	Ala	Ser	Val	Pro	Ala		
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Ser	Ala	Val	Arg	Leu	Thr	Glu	Arg	Ser	Lys	Leu	Ala	Lys	Gln	Leu	Leu		
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Ile	Gly	Tyr	Val	Thr	Ser	Gly	Thr	Leu	Ile	Leu	Ala	Ala	Gly	Ala	Ile		
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Ala	Val	Val	Val	Arg	Gly	His	Phe	Phe	Val	His	Ser	Leu	Val	Val	Ala		
385				390						395				400			
ggc	ttg	atc	acg	acc	gtc	tgc	gga	ttt	cgc	tcg	cgg	ctt	tac	gcc	gag	1248	
Gly	Leu	Ile	Thr	Thr	Val	Cys	Gly	Phe	Arg	Ser	Arg	Leu	Tyr	Ala	Glu		
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cgc	tgg	tgt	gcg	tgg	gcg	ttg	ctg	gcg	gcg	acg	gtc	gcg	att	ccg	acg	1296	
Arg	Trp	Cys	Ala	Trp	Ala	Leu	Leu	Ala	Ala	Thr	Val	Ala	Ile	Pro	Thr		
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Gly	Leu	Thr	Ala	Lys	Leu	Ile	Ile	Trp	Tyr	Pro	His	Tyr	Ala	Trp	Leu		
			435			440						445					
ttg	ttg	agc	gtc	tac	ctc	acg	gta	gcc	ctg	gtt	gcg	ctc	gtg	gtg	gtc	1392	
Leu	Leu	Ser	Val	Tyr	Leu	Thr	Val	Ala	Leu	Val	Ala	Leu	Val	Val	Val		
	450					455					460						
ggg	tcg	atg	gct	cac	gtc	cgg	cgc	gtt	tca	ccg	gtc	gta	aaa	cga	act	1440	
Gly	Ser	Met	Ala	His	Val	Arg	Arg	Val	Ser	Pro	Val	Val	Lys	Arg	Thr		
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ctg tgg atc acc ggg gtg tac gac acg gtc cgc aat atc cgg ttc	1533
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Ala Lys Leu Ala Gly Leu Val Phe Pro Gln Pro Pro Ala Pro Ile Ala	
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gtc agc gga acg gat tcg gtg gta gca gca atc aac gag acc atg cca	144
Val Ser Gly Thr Asp Ser Val Val Ala Ala Ile Asn Glu Thr Met Pro	
35 40 45	
agc atc gaa tcg ctg gtc agt gac ggg ctg ccc ggc gtg aaa gcc gcc	192
Ser Ile Glu Ser Leu Val Ser Asp Gly Leu Pro Gly Val Lys Ala Ala	
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ctg act cga aca gca tcc aac atg aac gcg gcg gcg gac gtc tat gcg	240
Leu Thr Arg Thr Ala Ser Asn Met Asn Ala Ala Ala Asp Val Tyr Ala	
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Lys Thr Asp Gln Ser Leu Gly Thr Ser Leu Ser Gln Tyr Ala Phe Gly	
85 90 95	
tcg tcg ggc gaa ggc ctg gct ggc gtc gcc tcg gtc ggt ggt cag cca	336
Ser Ser Gly Glu Gly Leu Ala Gly Val Ala Ser Val Gly Gly Gln Pro	
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agt cag gct acc cag ctg ctg agc aca ccc gtg tca cag gtc acg acc	384
Ser Gln Ala Thr Gln Leu Leu Ser Thr Pro Val Ser Gln Val Thr Thr	
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Gln Leu Gly Glu Thr Ala Ala Glu Leu Ala Pro Arg Val Val Ala Thr	
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Val Pro Gln Leu Val Gln Leu Ala Pro His Ala Val Gln Met Ser Gln	
145 150 155 160	
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Asn Ala Ser Pro Ile Ala Gln Thr Ile Ser Gln Thr Ala Gln Gln Ala	
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Ala Gln Ser Ala Gln Gly Gly Ser Gly Pro Met Pro Ala Gln Leu Ala	
180 185 190	
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Ser Ala Glu Lys Pro Ala Thr Glu Gln Ala Glu Pro Val His Glu Val	
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aca aac gac gat cag ggc gac cag ggc gac gtg cag ccg gcc gag gtc	672
Thr Asn Asp Asp Gln Gly Asp Gln Gly Asp Val Gln Pro Ala Glu Val	
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Val Ala Ala Ala Arg Asp Glu Gly Ala Gly Ala Ser Pro Gly Gln Gln	

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Arg Pro Ala Ala Ser Pro Leu Ala Ala Pro Val Asp Pro Ser Thr Pro				
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Pro Gly Gly Trp Val Glu Ala Asp Glu Asp Thr Phe Tyr Asp Arg Ala				
20	25	30		
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Gln Glu Tyr Ser Gln Val Leu Gln Arg Val Thr Asp Val Leu Asp Thr				
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Cys Arg Gln Gln Lys Gly His Val Phe Glu Gly Gly Leu Trp Ser Gly				
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ggc gcc gcc aat gct gcc aac ggc gcc ctg ggt gca aac atc aat caa				240
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Leu Gln Gln Lys Ser Pro Pro Pro Pro Asp Val Pro Thr Leu Val Val				
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cca tcc ccg ggc aca ccg ggc aca ccg gga acc ccg atc acc ccg gga				624
Pro Ser Pro Gly Thr Pro Gly Thr Pro Gly Thr Pro Ile Thr Pro Gly				
195	200	205		
acc ccg atc acc ccg gga acc cca atc aca ccc atc ccg gga gcg ccg				672
Thr Pro Ile Thr Pro Gly Thr Pro Ile Thr Pro Ile Pro Gly Ala Pro				

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cgt ggc cgc ggt gat gcg ctg ccg ttg gcg cga cgc atc gcg gcg gcg Arg Gly Arg Gly Asp Ala Leu Arg Leu Ala Arg Arg Ile Ala Ala Ala 485 490 495			1488
ctc aac gcg tcc gac aac aac gcg ggc gac tac ggg ttc ttc tgg atc Leu Asn Ala Ser Asp Asn Asn Ala Gly Asp Tyr Gly Phe Phe Trp Ile 500 505 510			1536
acc gcg gtg acc acc gac ggt tcc atc gtc gtg gcc aac agc tat ggg Thr Ala Val Thr Thr Asp Gly Ser Ile Val Val Ala Asn Ser Tyr Gly 515 520 525			1584
ctg gcc tac ata ccc gac ggg atg gaa ttg ccg aat aag gtg tac ttg			1632

-continued

Leu	Ala	Tyr	Ile	Pro	Asp	Gly	Met	Glu	Leu	Pro	Asn	Lys	Val	Tyr	Leu	
530						535					540					
gcc	agc	gcg	gat	cac	gca	atc	ccg	gtt	gac	gaa	att	gca	cgc	tgt	gcc	1680
Ala	Ser	Ala	Asp	His	Ala	Ile	Pro	Val	Asp	Glu	Ile	Ala	Arg	Cys	Ala	
545					550					555					560	
acc	tac	ccg	gtt	ttg	gcc	gtg	caa	gcc	tgg	gcg	gct	ttc	cac	gac	atg	1728
Thr	Tyr	Pro	Val	Leu	Ala	Val	Gln	Ala	Trp	Ala	Ala	Phe	His	Asp	Met	
				565					570					575		
acg	ctg	cgg	gcg	gtg	atc	ggc	acc	gcg	gag	cag	ttg	gcc	agt	tcg	gat	1776
Thr	Leu	Arg	Ala	Val	Ile	Gly	Thr	Ala	Glu	Gln	Leu	Ala	Ser	Ser	Asp	
			580					585					590			
ccc	ggt	gtg	gcc	aag	att	gtg	ctg	gag	cca	gat	gac	att	ccg	gag	agc	1824
Pro	Gly	Val	Ala	Lys	Ile	Val	Leu	Glu	Pro	Asp	Asp	Ile	Pro	Glu	Ser	
		595					600					605				
ggc	aaa	atg	acg	ggc	cgg	tcg	cgg	ctg	gag	gtc	gtc	gac	ccc	tcg	gcg	1872
Gly	Lys	Met	Thr	Gly	Arg	Ser	Arg	Leu	Glu	Val	Val	Asp	Pro	Ser	Ala	
	610					615					620					
gcg	gct	cag	ctg	gcc	gac	act	acc	gat	cag	cgt	ttg	ctc	gac	ttg	ttg	1920
Ala	Ala	Gln	Leu	Ala	Asp	Thr	Thr	Asp	Gln	Arg	Leu	Leu	Asp	Leu	Leu	
625					630					635					640	
ccg	ccg	gcg	ccg	gtg	gat	gtc	aat	cca	ccg	ggc	gat	gag	cgg	cac	atg	1968
Pro	Pro	Ala	Pro	Val	Asp	Val	Asn	Pro	Pro	Gly	Asp	Glu	Arg	His	Met	
				645					650					655		
ctg	tgg	ttc	gag	ctg	atg	aag	ccc	atg	acc	agc	acc	gct	acc	ggc	cgc	2016
Leu	Trp	Phe	Glu	Leu	Met	Lys	Pro	Met	Thr	Ser	Thr	Ala	Thr	Gly	Arg	
			660				665						670			
gag	gcc	gct	cat	ctg	cgg	gcg	ttc	cgg	gcc	tac	gct	gcc	cac	tca	cag	2064
Glu	Ala	Ala	His	Leu	Arg	Ala	Phe	Arg	Ala	Tyr	Ala	Ala	His	Ser	Gln	
		675					680				685					
gag	att	gcc	ctg	cac	caa	gcg	cac	act	gcg	act	gac	gcg	gcc	gtc	cag	2112
Glu	Ile	Ala	Leu	His	Gln	Ala	His	Thr	Ala	Thr	Asp	Ala	Ala	Val	Gln	
	690					695					700					
cgt	gtg	gcc	gtc	gcg	gac	tgg	ctg	tac	tgg	caa	tac	gtc	acc	ggg	ttg	2160
Arg	Val	Ala	Val	Ala	Asp	Trp	Leu	Tyr	Trp	Gln	Tyr	Val	Thr	Gly	Leu	
705					710					715					720	
ctc	gac	cgg	gcc	ctg	gcc	gcc	gca	tgc	tga							2190
Leu	Asp	Arg	Ala	Leu	Ala	Ala	Ala	Cys								
				725												

What is claimed is:

1. A diagnostic composition that discriminates between infection by *Mycobacterium tuberculosis* and vaccination by Bacille Calmette Guerin (BCG) strain of *Mycobacterium bovis*, said composition comprising antigens, all antigens in said composition consisting of at least three different polypeptides of the *Mycobacterium tuberculosis* complex that are not encoded by BCG, and said polypeptides including at least one isolated polypeptide from the group consisting of (i) a first amino acid sequence consisting of the sequence of MTBN4 (SEQ ID NO: 4), (ii) a second amino acid sequence that is an antigenic segment of MTBN4 that has *Mycobacterium tuberculosis* specific antigenic or immunogenic properties and (iii) a third amino acid sequence that is identical to said first or second amino acid sequence but has conservative

substitutions and has *Mycobacterium tuberculosis* specific antigenic or immunogenic properties.

2. The composition of claim 1, wherein said at least one isolated polypeptide comprises consists of said first or second amino acid sequence.

3. The composition of claim 2 further including a pharmaceutically acceptable diluent or filler.

4. The composition of claim 1 further including a pharmaceutically acceptable diluent or filler.

5. A mixture of a composition of claim 1 and a bodily fluid.

6. The mixture of claim 5, wherein the bodily fluid is blood, saliva, plasma, serum, urine, semen, or a lavage.

7. The composition mixture of claim 6, wherein the lavage is a bronchoalveolar lavage, a vaginal lavage, or lower gastrointestinal lavage.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 8,974,800 B2
APPLICATION NO. : 13/893659
DATED : March 10, 2015
INVENTOR(S) : Maria L. Gennaro

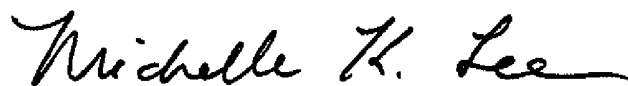
Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In the Claims:

At column 50, claim number 2, line number 48, delete “comprises”

Signed and Sealed this
Nineteenth Day of July, 2016

A handwritten signature in black ink, reading "Michelle K. Lee". The signature is written in a cursive, flowing style.

Michelle K. Lee
Director of the United States Patent and Trademark Office

专利名称(译)	由结核分枝杆菌而不是BCG表达的蛋白质及其作为诊断试剂和疫苗的用途		
公开(公告)号	US8974800	公开(公告)日	2015-03-10
申请号	US13/893659	申请日	2013-05-14
[标]申请(专利权)人(译)	新泽西内科与牙科大学		
申请(专利权)人(译)	医药口腔新泽西理工大学		
当前申请(专利权)人(译)	罗格斯新泽西州立大学		
[标]发明人	GENNARO MARIA LAURA		
发明人	GENNARO, MARIA LAURA		
IPC分类号	A61K39/04 A61K39/00 A61K39/02 G01N33/53 A61K31/711 A61K38/00 A61K48/00 A61P31/04 A61P31/06 C07K14/35 C12N1/15 C12N1/19 C12N1/21 C12N5/10 C12N15/09 C12Q1/02 G01N33/50 G01N33/569		
CPC分类号	A61K39/04 C07K14/35 G01N33/5695 G01N33/5091 A61K38/00 A61K39/00 A61K2039/53 Y10S435/863 A61P31/04 A61P31/06 A61K49/0006 G01N2333/35 G01N2333/57 G01N2800/26		
优先权	12/503717 2011-09-20 US 11/677502 2009-08-25 US 10/009383 2011-04-26 US PCT/US2000/012257 2000-05-04 WO 60/132505 1999-05-04 US		
其他公开文献	US20130251739A1		
外部链接	Espacenet USPTO		

摘要(译)

本发明涉及用于产生针对结核分枝杆菌的免疫应答和用于诊断已经暴露于M的受试者的感染和疾病的试剂。结核。的

MTBN1
MTAIEFVRLTLREVVLDDQLGTAEERAYKMLPPLTNVPLNEELIARDRRQPLRFALGIMDE
PERHLDQVWGVNBAAGRIIGIQAFOYCKSTLLQTMVMEAAATKSPRNVOFYCIDLGGG
QLVLENLPHVGGVANNRSEEDKVRNVAEMDAVMKREITTFEERHVSIGIOMTQCLNDDPS
QVVAESPYGDFLLIUDMPFVGEFFELKGGVQDLAAAGLAFGVHVTSTFENLTKSEV
RDYLGTKIETRLGDVHETQIDKITREIETANRFGRAVSHKELMIQVPRFDCVHSDNLY
EALTAGVQLASQITEQAFVWVLPKELHLRELDNMFPGESDYEETREIFGLKRETDLT
PAHCHMNTNPHLLIFQAARSGKTIJAJAJARACANRSPQQVRFMLADYESGLLDVAFDT
HLGARAIRNSASLEBAVQALAVHLKELLPTDILTAAQLESERHWSDFVVLVDWHM
IVGAAGGMPNAPLAPLLPAAADTGLNITVTQMSQAYKATMDKPVGAAPGSGAPTMLF
GERQEFSESEPKVGRPPQCAFVSPKKEVLAQYTESPPEVFAAPFSAQ

MTBN2
HEKMSHDPAAADIGTGVSDNALHGVTAGSTALTSTVTGLVPAGADEVSAQAATPTSGIO
LLASNAAQCQLHRAAGAVQDVARTYSGIDGGAAGVFAE

MTBN3
MLWHANFPELNTARLMAGAGPAPMLAAAGWOTLSAALDAQAVELTARLNSLGEANTGG
SDKALAAATPHVVVLQZASTGAKIRANGATAQAAVYQAHATTSPSLPIAANKITQAVIT
ATNFFGINTIPIALTENDVPIRMWQAALAMEVQAETAVNTLPKLEPMASILDPGAQ
STNMTFGNPEQSSTFVQCLPMAVOTIGOLGENSHMQLDPLQVTSLSQWGGTG
COMPADESAQKCLTSELSHPLAGSGSPAGSLAASSLPQAGSLTETPLASGL
EKFVAPSVMPAAAAAGSSATGGAAPVGAGAGGGAQSGGSTRFGLVAPAPLAQEREDDED
DWEDDW

MTBN4
NAEKTKGAATLAQEAQNFERISGDLKTCQIDQVESTAGSLQGWGGAAGTLAQAAVVRQ
AANKQKQZLDEISTNIRQAGVQYRADERQQQALSSOMF

MTBN5
KAADYDKLFRPHSGMBAPODDMAAQFFPFPFSAFPAFAGANLFXFNGOTFPPTSDDLSE
FVSAFFPPFPFPFPFPFPFPFIAAGPEPSPAPASKPPTPMPFIAQPEPAPPEKPTPMP
IAGFEPAPKFPPTPMPFIAFADPTTESGLAPRRPPTPQTPTGAPOQPEPAPHVPSHP
HOPRITAPAPPWAKMPIGEFFPEAPGRPSASPAEPTRFAPDHSEERARRGHRYRTDTERNV
GVATGHEICARLBAERASAGLAPCTEPSEPALDQGRETLATPTREPFEPSEPSOR
NSGRARRRVHPLAQHAAGQVDSITAAITGGRHRKRAAPDLDAIQKSLRPAARGPVVK
KVPKPKKATKPKVYVQGGGRHWBALTEINLQLESDEXTLELHAKVRRHGEVQIA
VVLKGGAGKTTCAALGSTLAQVRADRILALDADPGAGNLADRVGRQSGATIAVLAKE
ELHSDIIRAITVIAVHSELPAFETSAORALSDADHIFTADPASSEFNVLACDAG
FFDLITKGLSTVSGVVVAVSVSIDAQQAVALDWLRNNGYQELASRACVVIHIMHGE
PNVAVKDLVHPSGQVCPGRVVMWDRHIAAGTIESLGLLDPITYKKVLELAALSDDF
ERAGRR