

Serological Expression Cloning and Immunological Evaluation of MTB48, a Novel *Mycobacterium tuberculosis* Antigen

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Improved diagnostics are needed for the detection of *Mycobacterium tuberculosis*, especially for patients with smear-negative disease. To address this problem, we have screened *M. tuberculosis* (H37Rv and Erdman strains) genomic expression libraries with pooled sera from patients with extrapulmonary disease and with sera from patients with elevated reactivity with *M. tuberculosis* lysate. Both serum pools were reactive with clones expressing a recombinant protein referred to here as MTB48. The genomic sequence of the resulting clones was identical to that of the *M. tuberculosis* H37Rv isolate and showed 99% identity to the *Mycobacterium bovis* and *M. bovis* BCG isolate sequences. The genomic location of this sequence is 826 bp upstream of a region containing the *esat-6* gene that is deleted in the *M. bovis* BCG isolate. The *mtb48* 1,380-bp open reading frame encodes a predicted 47.6-kDa polypeptide with no known function. Southern and Western blot analyses indicate that this sequence is present in a single copy and is conserved in the *M. tuberculosis* and *M. bovis* isolates tested but not in other mycobacterial species tested, including *Mycobacterium leprae* and *Mycobacterium avium*. In addition, the native protein was detected in the cytoplasm, as was a processed form that was also shed into the medium during culture. Serological analysis of recombinant MTB48 and the *M. tuberculosis* 38-kDa antigen with a panel of patient and control sera indicates that the inclusion of recombinant MTB48 in a prototype serodiagnostic test increases assay sensitivity for *M. tuberculosis* infection when it is combined with other known immunodominant antigens, such as the 38-kDa antigen.

Mycobacterium tuberculosis continues to be the major infectious cause of human death in developing countries and has reemerged in industrialized countries (6, 14, 17, 21, 22, 27, 28). This reemergence is due, in part, to infection with the human immunodeficiency virus (HIV), which has led to increased susceptibility to *M. tuberculosis* and active disease (15, 35), development of new strains of multiple-drug-resistant *M. tuberculosis* (10, 30), and reduced resources for treatment and surveillance of patients (3, 22). Control of this infection will depend on new antibiotic therapy, improved early detection for both smear-positive and -negative cases, and the development of an effective vaccine. The development of a sensitive rapid serodiagnostic test would complement present methods of diagnosis, including skin testing, DNA amplification, bacterial culture, and radiological imaging and has become a major research goal (4, 11, 12, 13, 14, 16). A rapid and inexpensive assay could reduce the need for the more expensive tests, possibly reducing the cost of diagnosis, and would be of value in an emergency room setting, where early knowledge of a patient's disease status would be helpful. Finally, an inexpensive diagnostic test is needed in developing countries, where more expensive diagnostic tools are not available. To date, many *M. tuberculosis* antigens, such as the 38-kDa lipoglycoprotein (1), MTC28 (25), MPT32 (31), MTB81 (20), CFP-10 (9), and the 19-kDa (2) and 14-kDa (33) antigens, have been

tested for their utilities in the development of a rapid serodiagnostic test (5, 23, 32). However, the sensitivity needed for a useful test has not been achieved due to the heterogeneous human response to *M. tuberculosis* antigens (23) and to the low seroreactivities of problematic groups, such as HIV-positive patients or those with smear-negative disease (12, 19, 32). Here we describe the cloning of *mtb48* by screening a *M. tuberculosis* genomic expression library with pooled sera from patients with extrapulmonary disease and with sera from patients with elevated reactivity with *M. tuberculosis* lysate. The *mtb48* 1,380-bp open reading frame (ORF) encodes a predicted 47.6-kDa polypeptide with no known function. Southern blot and bioinformatic analyses indicate that this sequence is present as a single copy and is highly conserved in the *M. tuberculosis* and *Mycobacterium bovis* isolates tested but not in other mycobacterial species tested, including *Mycobacterium leprae* and *Mycobacterium avium*. Western blot analysis shows that the native protein is cytoplasmic and not membrane bound. A truncated form of the native protein is found both in the cytoplasm and in the culture filtrate (culture filtrate protein [CFP]), indicating that this form could be shed. Because serological reactivity to MTB48 is complementary to that of other *M. tuberculosis* antigens, this new antigen adds significantly to the antigen pool that will be needed for a highly sensitive and specific serodiagnostic test.

MATERIALS AND METHODS

Mycobacterial strains. Sean Skerritt (Seattle Veterans Affairs Hospital, Seattle, Wash.) provided *M. tuberculosis* strains H37Ra (ATCC 25177), H37Rv, and

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Erdman (ATCC 35801), *M. tuberculosis* strain C is a clinical isolate provided by Lee Riley (University of California, Berkeley). Pelleted samples of *M. bovis* BCG and *M. leprae* were provided by Paul Tan (Genesis Corp., Auckland, New Zealand). Other species of mycobacteria, including *Mycobacterium scrofulaceum* (ATCC 19981), *Mycobacterium smegmatis* (ATCC 19420), *Mycobacterium fortuitum* (ATCC 6841), *Mycobacterium mageritense* (ATCC 29571), and *Mycobacterium goodii* (ATCC 14470), were obtained from the American Type Culture Collection (ATCC; Manassas, Va.).

Library preparation and serological expression screening. *M. tuberculosis* cells were lysed in phenol with a cell disrupter (Mini-Beadbeater; type BX-4; Biospec Products, Bartlesville, Okla.) following the manufacturer's instructions. After the addition of TE buffer (10 mM Tris, 1 mM EDTA [pH 8]), the lysate was extracted two times with phenol, followed by extraction with phenol-chloroform-isoamyl alcohol and chloroform. Twenty micrograms each of *M. tuberculosis* Erdman and H37Rv strain genomic DNA was sheared with a sonicator (B. Braun Biotech, Inc., Allentown, Pa.) to generate fragments of approximately 0.5 to 5.0 kbp. DNA fragments were blunted with T4 DNA polymerase (Gibco BRL, Grand Island, N.Y.) and ligated to *EcoRI* adapters (Stratagene, La Jolla, Calif.). Adapted inserts were then phosphorylated with T4 polynucleotide kinase (Stratagene) and size selected with a Sephacryl S-400 HR column (Sigma Chemical Co., St. Louis, Mo.). Insert DNA was ligated to a Lambda ZAP II, *EcoRI*-digested, calf intestinal alkaline phosphatase-treated vector (Stratagene), and the ligation mixture was packaged with Gigapack II Gold packaging extract (Stratagene).

Immunoreactive proteins were screened from approximately 5×10^5 PFU of recombinant lambda phage from both the Erdman and the H37Rv genomic expression libraries with nitrocellulose filters (Schleicher & Schuell, Keene, N.H.). Reactive plaques were assessed with two pools of *Escherichia coli*-adsorbed patient sera. The first was a pool of three patient serum samples that were strongly reactive with *M. tuberculosis* lysate (20), as assessed by enzyme-linked immunosorbent assay (ELISA) (H37Rv library; TbH clones), and the second was a pool of five serum samples from *M. tuberculosis*-infected patients with extrapulmonary disease (Erdman library; XP clones). Positive plaques were visualized with an alkaline phosphatase-conjugated goat anti-human immunoglobulin G (IgG), IgA, and IgM (heavy and light chain) secondary antibodies (Zymed Laboratories Inc., South San Francisco, Calif.) developed with nitroblue tetrazolium-5-bromo-4-chloro-3-indolylphosphate (Gibco BRL). Excision of phagemid was done by the Lambda ZAP II protocol, and the resulting plasmid DNA was sequenced with an ABI 377 automated sequencer (Perkin-Elmer/ABI, Foster City, Calif.) with M13 forward, reverse, and specific internal DNA sequencing primers. Nucleic acid and protein homology searches were performed with software from DNASTAR, Inc. (Madison, Wis.) against the EMBL and GenBank database and the Swiss and PIR and Translated database. Protein analysis was performed with the PSORT program (National Institute for Basic Biology, Okazaki, Japan) and Protean analysis software from DNASTAR, Inc., including the Kyte-Doolittle hydrophilicity plot, Jameson-Wolf antigenicity index, and Emini surface probability plot.

Expression and purification of recombinant antigens. Four recombinant antigens were generated, including the amino terminus (rTbH4), the carboxy terminus (rXP-1), and the full-length sequence (rMTB48) of MTB48, as well as the mature polypeptide for the *M. tuberculosis* 38-kDa antigen (r38 kDa). Expression of rTbH4, rXP-1, and r38 kDa was accomplished by amplifying a plasmid insert (TbH4 and XP-1) or *M. tuberculosis* Erdman genomic DNA (38 kDa) with *Pfu* polymerase (Stratagene) and clone-specific primers that included a 5' *NdeI* restriction site (in italics in the sequences below), an ATG initiation codon (underlined in the sequences below), and a nucleotide sequence coding for 6 histidines (boldface in the sequences below) and a 3' primer with a stop codon and a *HindIII* or *EcoRI* restriction site (italic in the sequences below). Oligonucleotide primers for the amplification of TbH4 included 5'-ATTAGGTCATATGCACCATCACCATCACCATACGCAGTCGCAGACCGTGACGG and 3'-TATAGGAATCCTTCAATCCTCGGTGTAGAGCGCCTCG. XP-1 was amplified with 5'-CAATTACATATGCATCACCATCACCATCAACGACGGCGAAGGAAGTGTGC and 3'-AACCTGGAATTCGTCATGCTCACTTCGAC, and the 38-kDa antigen was amplified with 5'-CAATTACATATGCATCACCATCACCATCACCATCCTGTGGCTCGAAACCCAGG and 3'-GTACGGAATTCGTTGGTCAAACGAGGCTAGCTGG. Amplification product was digested with the appropriate restriction enzymes, isolated by gel electrophoresis, and ligated into a pET17b plasmid vector (Novagen, Madison, Wis.) previously cut with *NdeI* and *HindIII* or *EcoRI* and dephosphorylated. The ligation mixture was transformed into XL1 Blue MRF' competent cells (Stratagene), and plasmid DNA was prepared for sequencing (Qiagen Inc., Valencia, Calif.). Recombinant protein was expressed by transforming plasmid DNA into BL21 pLys S competent cells (Novagen) and inducing a single-colony cell culture with 2 mM isopropyl- β -D-thiogalactopyr-

anoside (IPTG; Sigma Chemical Co.). Full-length antigen was amplified with primers 5'-CTAGTTAGTACTCAGTCGCAGACCGTG and 3'-GCAGTGACGAATTCACCTCGACTCC by using 200 ng of total genomic *M. tuberculosis* Erdman strain DNA and Accuzyme (Intermountain Scientific Corporation, Inc., Kaysville, Utah) polymerase. The PCR product was digested with *ScaI* and *EcoRI* and ligated into a pET28 vector modified to include an in-frame methionine and a 6-histidine tag at the amino terminus. The full-length construct was sequenced and transformed into the HMS174 pLys S cell line (Novagen), and a single-colony culture was induced with IPTG. Cultures were disrupted by sonication and recombinant protein was recovered from cell lysate with an Ni-nitrilotriacetic acid agarose matrix (Qiagen Inc.). Purified protein was then passed over a Q anion-exchange column (rXP-1 and rMTB48) (Bio-Rad, Hercules, Calif.), following the manufacturer's instructions, and was finally dialyzed in 10 mM Tris (pH 8.0). Recombinant proteins were quality checked for purity by sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis (PAGE), followed by staining with Coomassie blue stain and N-terminal protein sequencing (26), and were quantified by a Micro BCA assay (Pierce, Rockford, Ill.). Recombinants were assayed for endotoxin contamination by the *Limulus* assay (Bio Whittaker, Walkersville, Md.).

Rabbit antiserum. Rabbit antiserum to recombinant antigens was prepared by injecting 200 μ g of protein in 1 ml of incomplete Freund's adjuvant (IFA; Sigma Chemical Co.) and 100 μ g of muramyl dipeptide subcutaneously. Six weeks later, the animals were boosted with 100 μ g of protein in IFA, followed by a final boost with 50 μ g administered intravenously 1 month later. Animals were killed 1 week after the final boost, and sera were stored in aliquots at -20°C .

Study population. Serum samples were obtained from individuals who had pulmonary tuberculosis alone prior to treatment (culture and/or acid-fast bacillus smear positive) or who had documented coinfections with HIV, as evidenced by a positive HIV type 1- and HIV type 2-specific antibody ELISA. These included samples from Brazil (Roberto Badaro, Federal University of Bahia, Salvador, Brazil), Uganda and South Africa (Milton Tam, Program for Appropriate Technology in Health, Seattle, Wash.), and the Philippines (Babi Tan, American Leprosy Foundation, Cebu City, Philippines) and sera from patients with tuberculosis (Nevin Turgay, Bornova-Izmir, Turkey). Samples from patients with lung cancer and pneumonia (China) were obtained from Robert Cole (ICT Diagnostics, Frenchs Forest, Australia). To further evaluate the specificity of the MTB48 antigen, we obtained sera from individuals who were purified protein derivative (PPD) positive (induration, >10 mm) (culture, clinically, and radiographically negative for tuberculosis) and individuals who were PPD negative (King County Tuberculosis Clinic, Seattle, Wash.). Sera from healthy blood donors (from the United States) were obtained from Boston Biomedica (West Bridgewater, Mass.), and sera from individuals who were HIV seropositive alone were obtained from Robert Ackridge (Fred Hutchinson Cancer Research Institute, Seattle, Wash.).

Southern blot analysis. Genomic DNA was prepared by standard techniques and digested with *PstI* (Gibco BRL). One microgram of each digest was run on a 1.0% agarose gel and stained with ethidium bromide to confirm the equivalent loading of each lane prior to overnight transfer to a Nytran nylon membrane (Schleicher & Schuell). Insert DNA isolated by gel electrophoresis was labeled by random hexamer primer extension with [^{32}P]dCTP and hybridized with blots overnight at 65°C . Blots were washed at 65°C twice for 15 min each time with $2\times$ SSC ($1\times$ SSC is 0.15 M NaCl plus 0.015 M sodium citrate) and $0.1\times$ SSC containing 0.1% SDS and exposed to X-ray film (Kodak, XAR-5).

Western blot analysis. Recombinant antigens and native *M. tuberculosis* proteins were subjected to SDS-PAGE analysis with 15% polyacrylamide minigels. A total of 200 ng of rMTB48 per lane was used for screening of patient serum by Western blot analysis. Cellular localization of native protein was determined by loading 10 μ g of an *M. tuberculosis* cytoplasm preparation; 2.5 μ g each of *M. tuberculosis* cell membrane fraction, cell wall fraction, and CFPs (provided by John Belisle, Colorado State University, produced through National Institute of Allergy and Infectious Diseases-National Institutes of Health Tuberculosis Research Materials contract N01-AI-25147); and 25 ng of recombinant protein. The proteins were transferred to nitrocellulose (BA-85; Schleicher & Schuell) and blocked for 1 h at room temperature with phosphate-buffered saline (PBS) containing 1% Tween 20. The blots were then washed three times for 10 min each time in PBS containing 0.1% Tween 20 and 0.5 M sodium chloride (wash buffer). Next, the blots were probed for 1 h at room temperature with serum diluted 1:500 in wash buffer, followed by three washes (10 min each time) in wash buffer. The blots were then incubated for 45 min at room temperature with protein A-horseradish peroxidase diluted 1:20,000 in wash buffer and again washed three times (10 min each time) in wash buffer. Finally, the blots were incubated in chemiluminescent substrate (ECL; Amersham Plc, Little Chalfont, United Kingdom) for ≈ 1 min and were then exposed to X-ray film (XAR5) for

TABLE 1. Comparison of the seroreactivities of recombinant proteins TbH4 and XP-1 with full-length MTB48

Patient identification	ELISA result ^a		rMTB48
	rXP-1	rTbH4	
383004	0.956	1.942	2.349
392004	7.446	4.375	5.287
393004	1.515	3.994	4.379
395004	2.392	1.494	1.591
396004	0.621	1.990	2.158
398004	0.902	1.737	2.241
407004	1.201	1.625	2.145
416004	0.830	1.628	1.917
445004	3.642	3.387	4.076
454004	1.070	1.085	1.005
465004	0.856	2.431	2.796
470004	1.595	2.795	3.455
476004	1.204	3.117	3.662
482004	2.119	4.313	4.386
487004	0.626	1.795	2.080
489004	2.667	4.578	4.405
490004	0.502	1.613	1.491
506004	0.452	1.444	1.524
514004	2.720	4.778	4.881

^a Data are normalized to a cutoff value (based on the mean plus 3 standard deviations) of 1.0 to allow comparison among groups. Positive results are shown in boldface type.

10 to 60 s, as required. The integrity of the protein in *M. tuberculosis* cellular protein fractions was confirmed by probing a Western blot with a rabbit anti-tubulin antiserum.

ELISA. Ninety-six-well microtiter plates (Corning Costar, Cambridge, Mass.) were coated overnight at 4°C with recombinant proteins (200 ng/well). The plates were then aspirated and blocked with PBS containing 1% (wt/vol) bovine serum albumin for 2 h at room temperature. This was followed by washing in PBS containing 0.1% Tween 20 (PBST). A dilution of serum (1/100) in PBS containing 0.1% bovine serum albumin was added to the wells, and the plates were incubated for 30 min at room temperature, followed by washing six times with PBST and then incubation with protein A-horseradish peroxidase conjugate (1/20,000 dilution; Sigma Chemical Co.) for 30 min. The plates were then washed six times in PBST and were then incubated with tetramethylbenzidine substrate (Kirkegaard & Perry Laboratories, Inc., Gaithersburg, Md.) for 15 min. The reaction was stopped by the addition of 1 N sulfuric acid, and the plates were read at 450 nm with an ELISA plate reader (EL311; Biotek Instruments, Hyland Park, Va.). The cutoff for the assays was determined from the mean for the negative population plus 3 standard deviations of the mean. For the data in Table 1, the signal-to-cutoff ratio was determined by dividing the sample ELISA value by the cutoff value.

Nucleotide sequence accession number. The GenBank accession number for *mtb48* is AY029285.

RESULTS

Expression cloning and molecular characterization of *mtb48*.

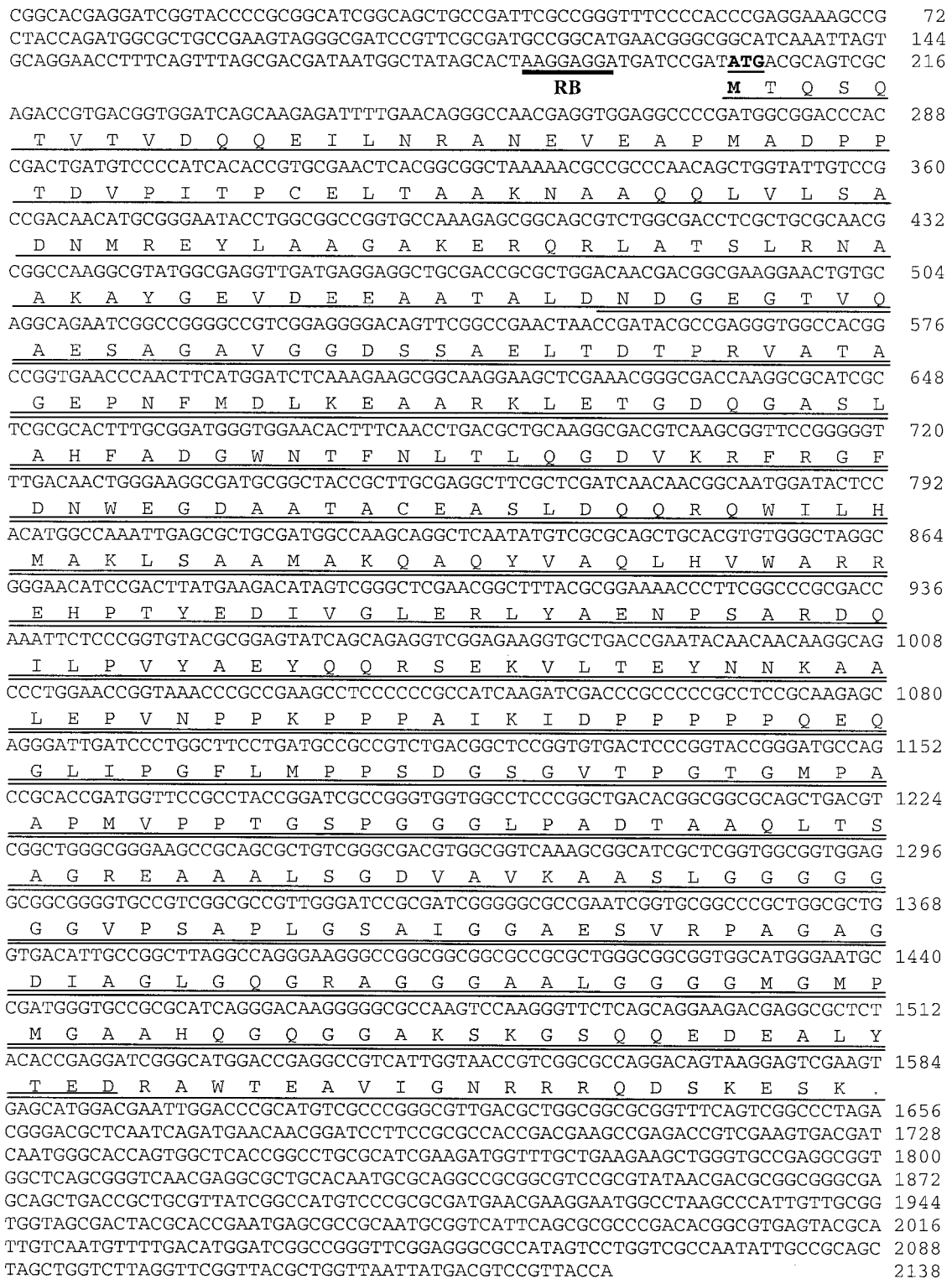
Two genomic expression libraries, constructed with randomly sheared *M. tuberculosis* Erdman strain and H37Rv strain genomic DNA, were screened with two pools of patient sera. The first pool consisted of sera from five patients with extrapulmonary disease (Erdman library), and the second pool consisted of sera from three patients whose sera were strongly reactive with the *M. tuberculosis* lysate (H37Rv library). These screenings resulted in the cloning of *mtb48* in two sections (clone *tbh4* from the H37Rv library and clone *xp-1* from the Erdman library) that overlap with 100% identity by 1,041 bp (Fig. 1). Clone *tbh4* contains the amino-terminal 1,319 bp, while clone *xp-1* contains the carboxy 1,101-bp terminus of a 1,380-bp ORF. The cloned inserts contain a predicted ribo-

some-binding site (Shine-Dalgarno sequence; Fig. 1) 17 bp upstream from the predicted ATG start codon and an ORF that represents a predicted polypeptide of 47.6 kDa. The predicted polypeptide contains no hydrophobic leader sequence, nor does it contain predicted transmembrane domains, as determined by the PSORT program. However, analysis of hydrophilicity indicates three hydrophobic stretches in the carboxy-terminal half of the polypeptide that could serve as transmembrane regions (Fig. 2).

Genomic organization of *mtb48*. Database searches with the nucleotide and predicted protein sequences for *mtb48* resulted in a 100% match with *M. tuberculosis* H37Rv genomic DNA (GenBank accession number AL022120) and hypothetical protein Rv3881c (GenBank accession number G70803), a 99% match with *M. bovis* (GenBank accession number U34848) and *M. bovis* BCG (GenBank accession number U35019) strain DNA, and a 98.7% match with the *M. bovis* and *M. bovis* BCG predicted protein sequences. Of interest, a partial genomic sequence (GenBank accession number AF001303) from an Indian *M. tuberculosis* clinical isolate (isolate NTI 64719) shows at least 98.8 to 99.6% identity to the *mtb48* sequence, given the degenerate bases contained in this sequence. Together, this information indicates that the *mtb48* sequence is well conserved between species (*M. tuberculosis* and *M. bovis*) and between geographically isolated strains from the United States and India. The genomic location of *mtb48* is 826 bp upstream from a 9,504-bp sequence (which contains the *esat-6* [accession number U34848] gene) that is deleted in the BCG strain of *M. bovis* (24). Southern blot analysis was performed with genomic DNA from both clinical and environmental *Mycobacterium* isolates. An *mtb48*-specific probe hybridized to 5.5-kbp reactive *Pst*I fragments in *M. tuberculosis* isolates, including strains H37Ra, H37Rv, Erdman (data not shown), and C (data not shown), and to a 6.5-kbp fragment in the *M. bovis* BCG isolate (Fig. 3). No reactivity was seen with *M. leprae*, *M. smegmatis*, *Mycobacterium vaccae*, *M. goodii*, *Mycobacterium chelonae*, *M. fortuitum*, *M. scrofulaceum*, or *M. avium*. The *Pst*I fragments correspond in size to those of the fragments predicted from the genomic DNA sequence. The difference in fragment size between *M. tuberculosis* isolates and the *M. bovis* BCG isolate can be attributed to a deletion event in BCG which removes the *Pst*I site that is found in the *M. tuberculosis* isolates.

Recombinant protein expression and purification. Four recombinant proteins representing the amino (rTbH4) and carboxy (rXP-1) termini and the full-length sequence (rMTB48) for MTB48 and the mature polypeptide for the *M. tuberculosis* 38-kDa antigen (r38 kDa) were overexpressed and purified by using a 6-histidine tag and metal affinity chromatography. Recombinant constructs were 1,341, 1,125, 1,404, and 1,080 bp, including the ATG start codon, 6-histidine tag, and stop codon, for rTbH4, rXP-1, rMTB48, and the 38-kDa antigen, respectively, and expressed recombinant proteins were 46.0, 38.6, 48.6, and 36.8 kDa for rTbH4, rXP-1, rMTB48, and the 38-kDa antigen, respectively. All recombinant proteins were tested for purity by N-terminal sequencing, Western blotting, and Coomassie blue staining and were determined to be over 95% pure.

Western blot analysis. The cellular localization of native protein was determined by probing a Western blot with rabbit



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FIG. 1. Relationship of full-length MTB48 to partial ORFs from genomic clones *tsh4* and *xp-1*. Partial ORFs for Tsh4 (upper line in the underlines) and XP-1 (lower line in the underlines) are underlined, and a double line indicates their overlap. The indicated polypeptide sequences were used to develop constructs for rTsh4, rXP-1, and rMTB48 protein expression. A potential ribosome-binding site (RB) is indicated by a thick bar, and the predicted ATG initiation codon is shown in bold and underlined. The nucleotide sequence number is shown on the right.

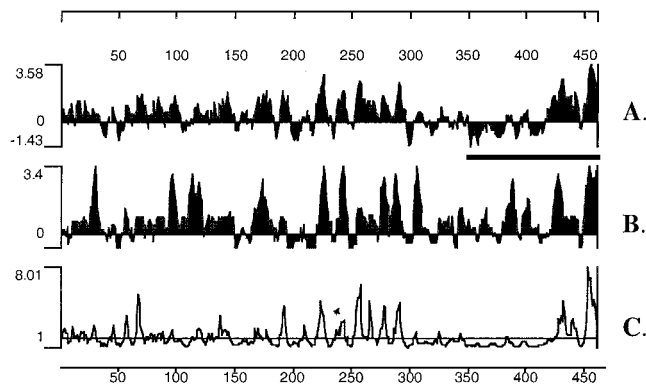


FIG. 2. Protein sequence analysis of MTB48 including hydrophilicity (A), antigenic index (B), and surface probability (C). Analysis was performed with DNA Star protein analysis software (Protean 4.0) by using the Kyte-Doolittle hydrophilicity plot, the Jameson-Wolf antigenic index, and the Emini surface probability plot. A thick bar indicates a region proposed to be cleaved from the native protein to allow release from the cell. Protein amino acid numbers are shown above and below the plots, while the relative peak heights are shown to the left.

rMTB48 antiserum. The blot contained fractionated *M. tuberculosis* proteins, including cytoplasmic, membrane, and cell wall fractions; CFPs; and rMTB48. Two major bands of approximately equal reactivity at 43 and 55 kDa were seen in the cytoplasmic fraction, while bands with little or no reactivity were observed in the membrane or cell wall fractions. A single band of 43 kDa was seen in the CFP lane, while two bands of

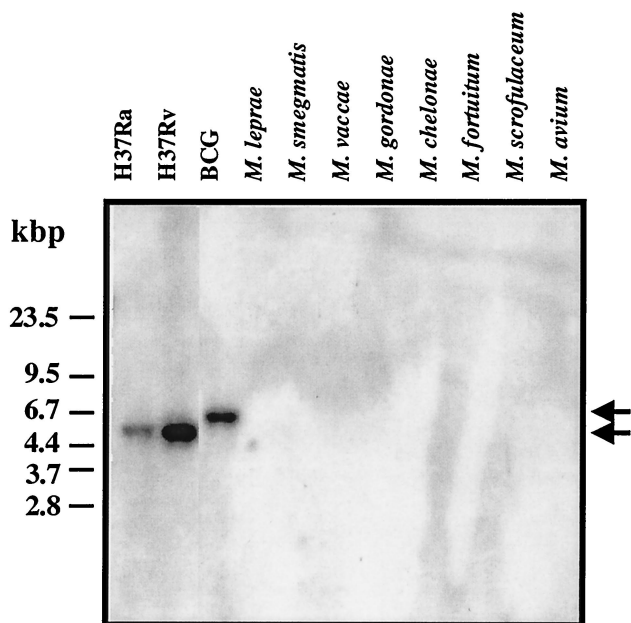


FIG. 3. Southern blot analysis of *mtb48* showing hybridization of a labeled *tbh4* nucleotide probe with *Pst*I-digested genomic DNA from the *M. tuberculosis* and *M. bovis* BCG isolates tested but not from *M. leprae* or the environmental mycobacteria tested. Arrows indicate the migration of a labeled 6.5-kbp fragment from the BCG strain of *M. bovis* (top arrow) and 5.5-kbp DNA fragments from *M. tuberculosis* isolates (bottom arrow).

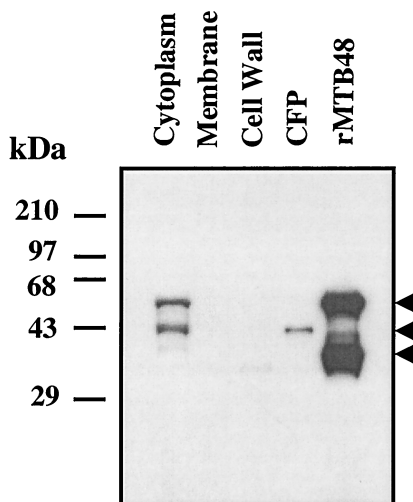


FIG. 4. Western blot analysis indicates that the cellular localization of native MTB48 protein is in the cytoplasm and the culture filtrate (CFP). Ten micrograms of an *M. tuberculosis* cytoplasm preparation, 2.5 μ g each of cell membrane, cell wall, and CFPs, and 25 ng of recombinant MTB48 were separated by SDS-PAGE, transferred to nitrocellulose, and probed with an MTB48 rabbit polyclonal antibody. A 55-kDa band (top arrow) is seen in both the cytoplasm and the recombinant protein lanes, while a 43-kDa band (middle arrow) is seen in both the cytoplasm and the CFP lanes. A second band of reactivity at approximately 38 kDa is shown in the recombinant protein lane (bottom arrow).

38 and 55 kDa were seen in the recombinant protein lane (Fig. 4). The 43-kDa form found in the CFP fraction appears to be identical in size to the 43-kDa truncated or processed form of MTB48 seen in the cytoplasmic fraction. Both the cytoplasmic native protein and the recombinant protein each show a processed form, with the said forms being of approximately equal molar concentrations, indicating that the two proteins might be processed by the same or a similar process.

ELISA evaluation of recombinant proteins XP-1, TbH4, and MTB48. Nineteen MTB48-positive patient serum samples were assayed with rXP-1 and rTbH4 to determine the general location of antigenic epitopes on rMTB48 (Table 1). rTbH4 was reactive with 19 of 19 serum samples, while rXP-1 was reactive with 11 of 19 serum samples. These data indicate that approximately 42% of MTB48-positive patient serum samples are reactive only with an amino-terminal epitope(s), while 58% of the serum samples are reactive with a central epitope(s) and, potentially, an additional amino-terminal epitope(s).

ELISA evaluation of sera from patients with tuberculosis and controls. Sera from individuals with tuberculosis in the presence or absence of HIV infection, PPD-positive individuals with no evidence of disease, and PPD-negative individuals were tested by ELISA with tuberculosis lysate, r38 kDa, and rMTB48 (Table 2). The sensitivities of recombinant proteins with serum samples from HIV-infected individuals in Africa with *M. tuberculosis* coinfections were 44.4 and 66.7% for rMTB48 and r38 kDa, respectively. The sensitivity of rMTB48 and r38 kDa used in combination was 74.1% (a gain of 7.4% over that obtained with r38 kDa only), indicating that the reactivities of the two recombinant antigens were complementary. Overall sensitivities for patients with only *M. tuberculosis*

TABLE 2. ELISA reactivities of *M. tuberculosis* lysate, rMTB48, and r38 kDa with HIV-positive and -negative sera from patients with tuberculosis and controls^a

Serum group and patient status	Origin	No. of serum samples	HIV status	Smear result	No. of samples positive by ELISA			
					Lysate	MTB48	38 kDa	38 kDa + MTB48
Sera from patients with:								
TB, ^b HIV, and <i>M. tuberculosis</i> coinfection	Uganda	17	+	+	7	8	12	13
	Uganda	2 ^c	+	-	1	0	1	1
	South Africa	8	+	+	6	4	5	6
Total		27			14	12	18	20
TB	Brazil	248	-	+	165	81	120	154
		52	-	-	24	9	11	16
	Uganda	15	-	+	10	10	12	12
	South Africa	51	-	+	47	10	15	23
	Turkey	11	-	+	8	3	9	9
	Philippines	31	-	+	24	10	11	17
	11	-	-	7	2	4	5	
Total		419			285	125	182	236
Control sera								
HIV infection	United States	11	+		0	0	0	0
Class II TB (PPD positive)	Africa	3	-		1	0	0	0
	Europe ^d	6	-		0	0	0	0
	Southeast Asia	7	-		1	0	2	2
	United States-Latin America	41	-		4	1	3	4
Total		57			6	1	5	6
Class 0/I TB (PPD negative)	Africa	1	-		0	0	0	0
	Europe ^d	3	-		2	0	0	0
	Southeast Asia	3	-		0	0	0	0
	United States-Latin America	22	-		0	2	1	3
Total		29			2	2	1	3
Other								
Lung cancer	China	13	-		2	0	0	0
Pneumonia	China	18	-		5	0	1	1
Healthy controls	China (Caucasian)	9	-		2	0	0	0
	United States	95	-		1	2	2	4
Total		135			10	2	3	5

^a Control sera were from HIV-positive, PPD-positive, and PPD-negative individuals, healthy controls, and patients with lung cancer and pneumonia.

^b TB, tuberculosis.

^c One individual was positive for tuberculosis by X ray.

^d Primarily from Russia and its satellite regions where tuberculosis is endemic.

infection were 29.8 and 43.4% for rMTB48 and r38 kDa, respectively. The sensitivity for the antigens used in combination, however, was 56.3%, indicating a greater contribution (a gain of 12.9%) of rMTB48 with these serum samples. Among the 419 serum samples from patients with only tuberculosis evaluated by ELISA, rMTB48 was reactive with 32.1% of the serum samples from patients with smear-positive tuberculosis and 17.2% of patients with smear-negative disease; and r38 kDa was reactive with 47 and 23.4% of smear-positive and smear-negative patients, respectively. Comparison of the ELISA values for serum samples from *M. tuberculosis*-infected

patients in Brazil and Africa (smear positive) shows a similar sensitivity. rMTB48 detected *M. tuberculosis* in 32.7% of the Brazilian samples and 30.3% of the African samples, while r38 kDa detected *M. tuberculosis* in 48.4 and 40.9% of these samples, respectively. Also, the gain in sensitivity with rMTB48 over that with r38 kDa was similar for both: 13.7 versus 12.1%, respectively. The specificities for rMTB48 and r38 kDa, including all control sera, were 97.8 and 96.1%, respectively, while the specificity for rMTB48 and r38 kDa in combination was 94%. No antibody reactivity with rMTB48 was seen for sera from the non-*M. tuberculosis*-infected control groups, including

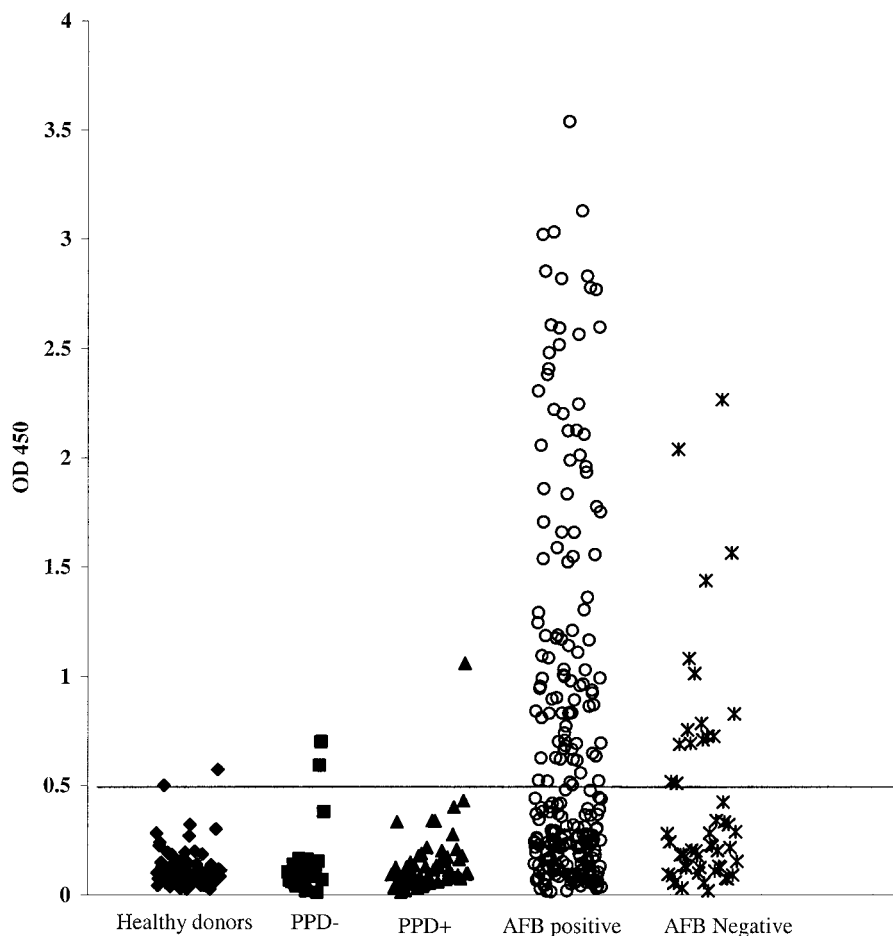


FIG. 5. Distribution of MTB48 ELISA reactivity among sera from healthy blood donors ($n = 95$), PPD-positive individuals ($n = 57$), PPD-negative individuals ($n = 29$), and Brazilian tuberculosis patients who were smear (acid-fast bacillus [AFB]) positive ($n = 248$) and negative ($n = 52$). The assay cutoff value (horizontal bar) was determined from the mean for the healthy donor population plus 3 standard deviations of the mean. OD 450, optical density at 450 nm.

sera from HIV-positive patients, lung cancer patients, and pneumonia patients. Specificities of 98.2 and 93.1% were determined for MTB48 with class II (PPD positive) and class 0/I (PPD negative) *M. tuberculosis*-infected sera, respectively. ELISA results for a subset of 248 smear-positive and 52 smear-negative Brazilian *M. tuberculosis* patient serum samples are displayed graphically in Fig. 5. Taken together, these data indicate that rMTB48 adds to the sensitivity of a 38-kDa antigen-based serological test for *M. tuberculosis* in a variety of diverse clinical and geographic settings.

DISCUSSION

The development of a serodiagnostic assay for *M. tuberculosis* with high degrees of sensitivity and specificity will most likely require a panel of several recombinant antigens or a fusion polyprotein that comprises the critical epitopes. This is due to the highly heterogeneous nature of antigen recognition by infected individuals that results, in part, from diverse genetic backgrounds, exposure to different stage-specific or strain-specific antigens, bacillary loads, patient health, and individual treatments (23). Data presented in this report indicate

that MTB48 is an important component of the antigen cocktail that will be required for diagnostic assay development. MTB48, which was cloned from both the Erdman and the H37Rv strains of *M. tuberculosis* by serological expression screening with *M. tuberculosis*-infected patient serum, encodes a predicted 47.6-kDa polypeptide (approximately 54 kDa under reducing conditions, possibly due to the high proline content [7.6%]) with no known function and is restricted to virulent and avirulent strains of the *M. tuberculosis* complex and *M. bovis*. *mtb48* genomic nucleotide sequences from the Erdman (clone *xp-1*) and H37Rv strains (clone *tbh4*) were identical, and database searches showed that the *mtb48* genomic sequence (Erdman strain) was identical to the H37Rv genomic sequence (GenBank accession number AL022120) and 99% identical to the corresponding *M. bovis* sequence (GenBank accession number U34848). A partial genomic sequence from an *M. tuberculosis* clinical isolate from India (GenBank accession number AF001303) also shows approximately 99% identity to *mtb48*. Taken together, this information indicates that the *mtb48* sequence is well conserved between species and between geographically isolated strains from the United States and India. The native MTB48 protein is cytoplasmic and appears to be

cleaved near the carboxy terminus to form a 43-kDa protein that is then shed. One cannot rule out the possibility, however, that in the culture filtrate the protein is degraded to the smaller fragment by secreted or shed proteases. The carboxy-terminal segment contains hydrophobic regions that could be removed upon processing, thus allowing secretion or shedding of the antigen. Preliminary epitope mapping shows that of the MTB48-positive patient serum samples tested, 42% recognize an amino-terminal epitope(s), which is missing in rXP-1, while 58% recognize one or more central epitopes, indicating the possibility of several B-cell epitopes along the molecule. Protein analysis (Fig. 2) shows that the amino terminus and central region of the protein are hydrophilic and are predicted to be highly antigenic. Surface probability predictions also indicate clusters of surface-exposed regions in the amino-terminal region and in the midregion of the molecule.

The serodiagnostic potential of rMTB48 compared to that of r38 kDa was evaluated by ELISA. The 38-kDa *M. tuberculosis* antigen was chosen for comparison because of its relatively high degrees of sensitivity and specificity, as determined in numerous studies (8, 9, 18, 20, 31, 34). Serological sensitivity for the 38-kDa antigen ranges from 16 to 89%, and specificity ranges from 93 to 100% (5, 8, 18, 22, 29, 34), depending largely upon the selection of patient populations in each study. In the present study, overall sensitivity values for rMTB48 and r38 kDa, including those for samples from patients with HIV coinfection and smear-positive and -negative patients, were 30.7 and 44.8%, respectively, with a sensitivity of 57.4% when both antigens were used in combination. Samples from patients with HIV coinfection alone resulted in higher values (44.4, 66.7, and 74.1%, respectively), as did samples from smear-positive patients (32.1, 47, and 60.6%, respectively). Values for samples from smear-negative patients were 17.2, 23.4, and 32.8%, respectively. Thus, MTB48 is a highly conserved antigen and its serological reactivity complements that of the 38-kDa antigen in the detection of *M. tuberculosis* infection.

Although rMTB48 adds to the overall sensitivity of the 38-kDa antigen, an *M. tuberculosis* serodiagnostic reagent with high degrees of sensitivity and specificity will depend on the use of several antigens, including r38 kDa and rMTB48. Detection of problematic infections, such as smear-negative infections, depends on a combination of techniques, including serology, detection of clinical symptoms, and radiography. Additional effort must be directed toward the serological detection of smear-negative infections. Although patients with smear-negative infections are less infectious and have a lower mortality rate than patients with smear-positive infections, the number of patients with smear-negative infections is estimated to exceed the number of patients with smear-positive infections, and over half of the patients with smear-negative infections will progress to active disease (7). In addition, coinfections with HIV have disproportionately increased the number of smear-negative cases. Our studies continue to be directed toward the development of a rapid serodiagnostic assay that uses an optimized set of recombinant antigens that will be expressed as one or more fusion polyproteins.

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