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A Mycobacterial Enzyme Essential for Cell Division Synergizes with Resuscitation-Promoting Factor

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**Abstract**

The final stage of bacterial cell division requires the activity of one or more enzymes capable of degrading the layers of peptidoglycan connecting two recently developed daughter cells. Although this is a key step in cell division and is required by all peptidoglycan-containing bacteria, little is known about how these potentially lethal enzymes are regulated. It is likely that regulation is mediated, at least partly, through protein–protein interactions. Two lytic transglycosylases of mycobacteria, known as resuscitation-promoting factor B and E (RpfB and RpfE), have previously been shown to interact with the peptidoglycan-hydrolyzing endopeptidase, Rpf-interacting protein A (RipA). These proteins may form a complex at the septum of dividing bacteria. To investigate the function of this potential complex, we generated depletion strains in *M. smegmatis*. Here we show that, while depletion of *rpfB* has no effect on viability or morphology, *ripA* depletion results in a marked decrease in growth and formation of long, branched chains. These growth and morphological defects could be functionally complemented by the *M. tuberculosis* *ripA* orthologue (*rv1477*), but not by another *ripA*-like orthologue (*rv1478*). Depletion of *ripA* also resulted in increased susceptibility to the cell wall–targeting β-lactams. Furthermore, we demonstrate that RipA has hydrolytic activity towards several cell wall substrates and synergizes with RpfB. These data reveal the unusual essentiality of a peptidoglycan hydrolase and suggest a novel protein–protein interaction as one way of regulating its activity.

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**Introduction**

Though not formally considered virulence factors, genes required for bacterial cell division clearly are necessary for the growth, and thus, pathogenesis, of bacteria. The distinction between homeostatic and virulence genes is blurred when nonessential genes involved in vegetative cell division become essential under specific stressful conditions encountered inside a host. Such an example is seen with the resuscitation-promoting factors (Rpf) encoded by many different bacteria, including *Mycobacterium*. These proteins are named for their ability to resuscitate nonreplicating dormant bacteria. The single Rpf-encoding gene in *Micrococcus luteus* [1] is essential, but as many as three of the five genes encoding RpFA-E can be deleted in *Mycobacterium tuberculosis* without markedly affecting *in vitro* growth. However, one single deletion (*rpFB*) and several of the triple combinations yielded strains unable to grow or divide in stressful conditions in vitro and in vivo [2,3]. This suggests that certain potential cell division proteins that appear to play nonessential roles in homeostatic processes can become vital in conditions of stress.

The vital processes of cell growth and division involve the temporal and spatial coordination of events such as peptidoglycan and cell wall extension, DNA replication, chromosomal partitioning, Z-ring assembly, septum formation, and cytokinesis. Much of the mechanism behind this coordination involves inhibiting and stabilizing proteins that regulate the eventual assembly of a Z-ring at the midcell of bacteria [4,5]. This Z-ring consists primarily of a polymerized ring of tubulin-like FtsZ on the cytoplasmic side of the plasma membrane, stabilized by membrane-associated and integral membrane proteins. Assembly occurs in an ordered fashion that is not entirely linear, with some components assembling before joining the Z-ring [6,7,8]. Some of the last proteins to be recruited to the Z-ring are thought to be the peptidoglycan hydrolizing enzymes, such as AmIC [9] and EnvC [10] in *E. coli*. These enzymes digest the peptidoglycan layers connecting two recently developed daughter cells in the final stage of cell division [11]. While crucial to cell division, the regulation of these potentially lethal enzymes is poorly understood. It is thought that protein-protein interactions play a role in regulating activity and localization [12]. Studying cell wall hydrolases has proven difficult due to the large number encoded in most bacterial genomes and the high degree of functional redundancy. For example, seven hydrolases can be deleted from a strain of *E. coli* without loss of viability [13].

Little is known about the hydrolases involved in mycobacterial cell division. CwlM and Rv2719c were both shown to be mycobacterial cell wall hydrolases [14,15]. Two recently identified hydrolases, RpfB and RpfE, were shown to interact with rpf-interacting protein (RipA), a peptidoglycan endopeptidase [16].
Author Summary

Mycobacteria, like all peptidoglycan-containing bacteria, must extend and cleave the surrounding structurally rigid layer of peptidoglycan to grow and divide. The peptidoglycan hydrolases responsible for this cleavage often have redundant functions, both revealing their importance and making them difficult to study. Furthermore, such hydrolases must be tightly regulated, due to their potentially lytic nature. We recently demonstrated the interaction between a lytic transglycosylase (Rpf) and an endopeptidase (RipA) at the septum of dividing bacteria. To investigate the role of these two hydrolases, we generated conditional mutants of each and were surprised to find that depletion of ripA resulted in long chains of cells. This phenotype was reversed upon induction of ripA, indicating that cell wall expansion and septum formation can be decoupled from the process of septum resolution. In addition, we present data showing that the combination of Rpf and RipA results in enhanced hydrolysis of peptidoglycan in an in vitro assay, suggesting protein–protein interactions as one potential mechanism of regulation.

Rpf proteins constitute a family of lytic transglycosylase enzymes capable of hydrolyzing the glycosidic bonds in the essential stress-bearing, shape-maintaining peptidoglycan layer [17]. RpfB has a structure similar to the E. coli soluble lytic transglycosylase 70 (Sh70) [18] and is known to hydrolyze the beta-1,4-glycosidic bond between N-acetyl muramic acid and N-acetyl glucosamine [19]. RipA has been shown to be a peptidoglycan hydrolase [16]. It is predicted to function as an L,D-endopeptidase, capable of hydrolyzing D-glutamyl-meso-diaminopimelic acid [20], two amino acids that are part of the crosslinking peptides vital for keeping peptidoglycan rigid and stable [21]. Both RpfB and RipA localize to the septa of dividing bacteria [16] and thus may play a role in the late stages of mycobacterial cell division, possibly during regrowth from a stressed state.

Here we show that depletion of ripA in a strain of M. smegmatis results in a significant decrease in growth, formation of long, branched chains, and increased sensitivity to a cell wall–targeting antibiotic. These defects can be functionally complemented with the M. tuberculosis allele of ripA. We demonstrate that the peptidoglycan hydrolytic activity of RipA synergizes with RpfB. Thus, this protein is an unusual example of an essential peptidoglycan hydrolase whose activity may be partially regulated through protein–protein interactions.

Results

Depletion of RipA blocks normal cell division

RipA interacts with RpfB, a lytic transglycosylase, and colocalizes at the septum [16]. We hypothesized that the RipA-RpfB complex may be involved in degrading peptidoglycan at the septum during cell division. To further investigate the function of the individual components of this complex, we attempted to make deletion strains of ripA and rpfB in M. smegmatis. Though disruption of rpfB was successful, we were unable to disrupt the ripA gene in M. smegmatis. We have previously reported that the ripA gene in M. tuberculosis appears to be essential for in vitro growth [22], suggesting that it might also be essential in M. smegmatis.

To test this possibility we constructed depletion strains of M. smegmatis in which ripA (MSMEG3153) or rpfB (MSMEG5439) are transcribed from an inducible tetracycline promoter (Ptet, Figure 1A). We found that the ripA depletion strain had dramatically reduced growth in media lacking inducer (Figure 1B), while the rpfB depletion strain had normal growth in the absence of inducer. Both ripA and rpfB depletion strains grew normally in the presence of inducer. The growth phenotype seen with ripA depletion, as measured by optical density, was dose-dependent. While the optical density of ripA depleted cultures failed to increase in the absence of inducer, we did observe that bacteria formed visible clumps that increased in size during incubation. These clumps (due to filamentation) failed to suspend well and, therefore, were poorly measured using spectrophotometry. Remarkably, this phenotype is reversible, as addition of inducer to growth-arrested, ripA depleted cells resulted in the resumption of normal growth (Figure 1C). This indicates that the frequency of septum resolution can be uncoupled from septum formation and cell elongation. High levels of inducer did not result in gross morphological changes or lysis.

To further confirm the requirement of ripA for growth, we plated ripA-depletion or rpfB-depletion strains of M. smegmatis on plates containing a gradient of inducer. The ripA-depletion strain grew in a Tet-dependent manner, with the highest growth in the center corresponding to the highest concentration of inducer and no growth near the edges of the plate where inducer was the lowest, confirming the requirement of ripA for growth. Conversely, the rpfB-depletion strain grew similarly throughout the plate in a Tet-independent manner, indicating that rpfB is not required for growth under these conditions.

Depletion of RipA results in abnormal morphology

While rpfB depleted cells had normal morphology, ripA depleted cells had markedly abnormal shape (Figures 1C and 2A). They grew in long, branched chains that account for the clumps seen grossly. Staining with the fluorescent membrane dye, TMA-DPH, revealed periodic septa along the chains of bacteria. DNA staining with SYTO 9 revealed nucleoids along the length of the chained bacteria, separated by septa, indicating that DNA segregation and cell elongation. High levels of inducer did not result in gross morphological changes or lysis.

Branches, not observed in wild-type cells, were seen in almost all ripA-depleted bacteria visualized (>95%). Interestingly, 91% (203/222) of the branches visualized originated directly adjacent to septa.

M. tuberculosis ripA allele functionally complements M. smegmatis ripA

The M. tuberculosis ripA gene encodes a 472 amino acid protein that has been shown to degrade peptidoglycan [23]. Its C-terminal 105 amino acids contain a putative endopeptidase domain, which has 40% identity with the Listeria monocytogenes p60 protein (Figure 3A). p60 has been shown to be a cell wall endopeptidase by its ability to degrade cell wall [24,25]. ripA is the first gene in a bicistronic operon, r1478, the downstream gene, encodes a 241-amino acid protein, consisting of a signal sequence followed by a sequence homologous to the C-terminal half of RipA (70% putative hydrolase domain identity and 27% overall amino acid identity). Both genes in the apparent ripA operon encode predicted endopeptidase domains similar to a known p60 hydrolase [24] (Figure 3A). Because the inserted tetracycline-inducible promoter lies upstream of the operon, presumably transcription of both is dependent on the presence of inducer. Thus, either gene could be
responsible for the observed phenotype. There are at least five p60-like genes in many of the mycobacterial species, including *M. tuberculosis*, *M. smegmatis*, and *M. bovis* BCG. These include *rv0024*, *rv1477* (*ripA*), *rv1478*, *rv1566c*, and *rv2190c*. These genes lie in various genomic contexts and there is no function discernable from genomic synteny (Figure 3B).

To identify the responsible gene we tested if the *M. tuberculosis* allele of *ripA* (*ripA-mtb*) was able to complement the *M. smegmatis* with diminished native *ripA* (*ripA-smeg*) production. We expressed *ripA-mtb* from a zeocin-marked episomal plasmid. A *ripA-smeg* depletion strain containing the *ripA-mtb* construct grew similarly to wildtype in the absence of inducer, while a strain carrying an empty plasmid formed chains when *ripA-smeg* was depleted (Figure 3C). In contrast, the *M. tuberculosis* allele of the *ripA* parologue, *rv1478*, was not able to complement an *M. smegmatis* *ripA* depletion. These results confirm that *ripA* is sufficient for complementing the strain depleted of the *ripA* (*ripA-MSMEG3154*) operon and show that *ripA-mtb* is functionally similar to *ripA-smeg*.

Depletion of RipA increases susceptibility to a cell wall-targeting antibiotic

Because depletion of *ripA* may have a marked effect on cell wall structure, we reasoned that strains with diminished expression of *ripA* might have altered susceptibility to antibiotics that target the cell wall. To test this, we grew the *M. Smegmatis* regulated *ripA* strain in the presence of inducer, then washed and spread on plates containing various amounts of inducer (0, 10, 100 ng/ml Tet). A sterilized disc of Whatman paper was placed in the middle of the plate, and antibiotics were added to the disc. After incubating for 3–4 days, the size of the zone of inhibition was measured (Figure 4). Complete depletion of *ripA* resulted in a remarkably high level of susceptibility to β-lactams (carbenicillin). It is unclear why depletion of *ripA* results in such an increase in susceptibility to β-lactams, which target the transpeptidase reaction required for cross-linking peptidoglycan during cell elongation and division. Susceptibility to cycloserine, an analog of D-alanine that inhibits the formation of the cytoplasmic peptidoglycan.
pentapeptide that is eventually transported across the cell membrane and used to cross-link PG strands, was independent of induction of ripA. While increased permeability is often attributed to an observed increase in susceptibility to an antibiotic, depletion of ripA did not affect susceptibility to cycloserine, suggesting a more specific defect.

RipA degrades multiple cell wall substrates

RipA has been predicted and shown to degrade M. luteus cell wall material [25,26]. To test if RipA hydrolyzes peptidoglycan and cell wall material from other species of bacteria, we determined the enzymatic activity of RipA using a variety of FITC-labeled, cell wall–derived substrates. We expressed RipA as a fusion protein with GST in E. coli and purified the fusion protein using affinity chromatography. We found that GST-RipA, but not GST alone, was able to hydrolyze cell wall derived from M. smegmatis as well as peptidoglycan purified from Streptomyces, and had minimal activity against M. luteus cell wall (Figure 5B–D). Therefore, RipA is capable of hydrolyzing cell wall material from several bacterial species.

The interaction of RipA and RpfB results in synergistic hydrolysis of peptidoglycan

The predicted activity of RipA is to cleave the peptide cross-linkages in peptidoglycan and is distinct from Rpf, which is predicted to cleave glycosidic bonds in peptidoglycan (Figure 5A). Given the close proximity of these predicted cleavage sites, we
hypothesized that the interaction of RpfB with RipA may result in enhanced hydrolytic activity. To test this possibility we expressed a portion of RpfB as a fusion protein with GST in E. coli and purified it using affinity chromatography.

Using the same assays described above, we found that GST-RpfB alone had minimal ability to degrade cell wall extracts or purified peptidoglycan. However, when GST-RpfB was combined with GST-RipA, activity was more than the sum of individual enzyme activities. The same result was found with all substrates tested (Figure 5B–D). No increase was detected when GST was combined with GST-RpfB or GST-RipA (data not shown). Addition of twice as much Rpf yielded no increase in hydrolysis, while twice as much RipA yielded twice the hydrolysis (data not shown) indicating the assay is in the linear range.

Discussion

In this work, we demonstrate that RipA is essential for normal cell division in M. smegmatis, with its depletion resulting in long, branched filaments and increased susceptibility to a specific cell wall targeting antibiotic. Furthermore, RipA cleaves peptidoglycan and synergizes with RpfB. Taken together, these data support a model where RipA is 1) required for the final stage of cell division, where daughter cells are separated and 2) has peptidoglycan hydrolytic activity that may be modulated by RpfB under certain conditions.

It is unusual that RipA is essential for normal cell division in M. smegmatis and, apparently, M. tuberculosis [22]. Because bacteria encode a number of hydrolytic enzymes that are, at least in part, functionally redundant, strains carrying deletions of single hydrolase genes are generally viable, though combinations of mutations can result in lack of viability [13]. In M. smegmatis and M. tuberculosis, ripA does not appear to be redundant. Conversely, while M. marinum strains carrying mutations in the homologous gene, ripA, do have abnormal morphology, they are still able to divide. Mutations in the hydrolytic domain of RipA abolished complementation of the defect, confirming the importance of the...
hydrolytic activity of IipA [27]. In *M. marinum*, different *rip* paralogues might be able to complement for loss of *iipA*.

None of the *rpf* genes appears to be essential in *M. tuberculosis* and combinations of at least three *rpf* genes can be deleted in *M. tuberculosis* strains while still maintaining normal in vitro vegetative growth [2]. We demonstrate that RpfB is also not essential in *M. smegmatis*. It logically follows that the interaction between RpfB and RipA must not be essential for RipA function during vegetative growth. Of course, it is possible that another Rpf protein is able to compensate for the absence of RpfB, resulting in increased RipA-dependent activity. For example, RpfE is able to interact with RipA [16]. It is also possible that the RipA-RpfB interaction, and subsequent enhanced hydrolytic activity, is required only under special circumstances, such as growth under specific conditions of stress. As noted, RpfB is required for resuscitation of *M. tuberculosis* in a reactivation mouse model [3]. Likewise, deletion of several combinations of three *rpf* genes results in viable bacteria that are unable to resuscitate from in vitro and in vivo resuscitation assays [2]. Thus, the RipA-RpfB interaction may be necessary under certain conditions.

Figure 4. An *M. smegmatis* strain depleted of RipA is sensitive to a cell wall–targeting antibiotic. The ripA depletion strain of *M. smegmatis* was spread on LB agar plates containing different amounts of anhydrotetracycline inducer (ng/ml concentrations) to regulate the amount of *ripA* expressed. A filter disc with 10 μl antibiotic was placed in the center of plate and the diameter of inhibition of growth was measured after 4 days of growth. Antibiotic concentration on disc: carbenicillin (100 mg/ml) and cycloserine (100 mg/ml).

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Figure 5. Recombinant RpfB and RipA combine to synergistically hydrolyze peptidoglycan. (A) Diagram of the structure of peptidoglycan, indicating where RpfB and RipA are predicted to have hydrolytic activity. One unit of the peptidoglycan is magnified to show the structure of NAG and NAM as well as the amino acids that are part of the DAP-DAP crosslinkage. Lines connecting NAG to NAM represent β-1,4-glycosidic bonds, while residues connecting NAM to NAM depict peptide cross-linkages. NAG: N-acetylglucosamine, NAM: N-acetylmuramic acid. (B–D) N-terminal GST fusion proteins were expressed and purified from *E. coli*. Equal molar amounts of individual or combinations of proteins were incubated with insoluble FITC-labeled substrate: *M. smegmatis* cell wall (B), *Streptomyces* peptidoglycan (C), or *M. luteus* cell wall (D). The extent of hydrolysis was determined by measuring the amount of soluble FITC remaining after centrifugation, and thus released during hydrolysis of the insoluble substrate. GST alone, as well as buffer alone, were used to determine background release of FITC and were subtracted from final values. Data are from representative experiments, each done in triplicate. Data are represented as mean +/− SEM.

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Materials and Methods

Strains and culture conditions

*E. coli* XL-1 (Stratagene) strains were used for cloning and *E. coli* BL21 (DE3) (Stratagene) was used for expression of recombinant proteins from the pET41a (Novagen) or pMal (NEB). *Mycobacterium smegmatis* (mc²155) and *Mycobacterium tuberculosis* (H37Rv) strains were grown at 37°C in Middlebrook 7H9 broth supplemented with ADC and Tween80 and antibiotic when appropriate.

Recombinant protein production

The *E. coli* expression strain, BL21(DE3) was used to synthesize each protein following the Novagen manual protocol. Protein concentrations were measured using the Bradford assay, normalized, and confirmed by coomassie-stained polyacrylamide gels.

Western blotting

Protein samples were combined with 4× Laemmli’s SDS PAGE buffer and boiled at 100°C for 10 minutes. Proteins were separated on 10% Tris-tricine polyacrylamide gels by electrophoresis, transferred to nitrocellulose, and probed with specific antibodies using standard techniques.

Preparation and FITC-labeling of cell wall material

*M. smegmatis* cell wall was prepared as previously described [14]. *Streptomyces* peptidoglycan and lyophilized *M. luteus* cell wall were both obtained from Sigma. The fluorescein isothiocyanate (FITC)-labeled bacterial cell wall was prepared by covalently linking FITC to amine groups in the cell wall. 10 μg FITC (Molecular Probes) was used to label 10 mg of insoluble peptidoglycan or cell wall material following the protocol from Molecular Probes.

Enzyme assay

Recombinant *M. tuberculosis* proteins were incubated with several FITC-labeled cell wall substrates and assayed for activity by measuring FITC release. 25 μg of Rpf or RlpA alone or 25 μg of Rpf and 25 μg RlpA combined, were added to 25 μl of 2 mg/ml substrate and 25 μl 4× reaction buffer (50 mM Tris, 10 mM MgCl₂, 50 mM KCl, 2 mM MnCl₂, 0.01% Chaps, 100 mM KH₂PO₄, pH 5.75). The final volume was brought to 100 μl with H₂O. As a control, 50 μg of lysozyme was added to *M. smegmatis* cell wall. Similar combinations with GST were also tested. GST alone, as well as buffer alone, were used to determine background release of FITC. After incubating at 30°C with enzyme and buffer for 3–5 days, the insoluble substrate was centrifuged (18,000×g) and soluble FITC was measured with filters for excitation 485 nm and emission 530 nm.

Generation of depletion strains

Depletion strains were generated as previously described [28]. Briefly, *M. smegmatis*, with the tetracycline repressor gene integrated into the attB site, was transformed with a suicide vector containing the first 600 nucleotides of *M. smegmatis ripA* gene under control of the tetracycline operator/promoter system (Ptet). Transformants were selected for hygromycin resistance. Appropriate recombination was confirmed using forward primers to Ptet and Prip (native ripA promoter) paired with a reverse primer to the 3’ end of ripA. The presence of a product of appropriate size for the former and lacking in the latter, confirmed the desired strain. Attempts to disrupt the ripA gene in *M. smegmatis* using a nonreplicating suicide vector designed to recombine into the middle of the gene were unsuccessful (though control knockouts, such as rpfB, were successful).

Depletion strain growth and complementation

The ripA and rpfB depletion strains were initially grown in 7H9 media containing kanamycin (selecting for TetR) and hygromycin (selecting for inserted pTet) as well as anhydrotetracycline (Tet). Once cultures reached late log-phase or stationary phase, they were centrifuged (2500 × g for 5 minutes), washed once with PBS, and resuspended in media with varying amounts of Tet. To test recovery of ripA depleted cells, Tet was either added directly to cultures grown without Tet or to fresh media inoculated with cells depleted of ripA. To test complementation, the ripA gene and its native promoter from *M. tuberculosis* was amplified and cloned into an episomal plasmid containing the zacin gene as a marker. This construct, or the isogenic empty vector, was transformed into the ripA depletion strain of *M. smegmatis*. Strains were grown in the presence of tetracycline inducer, washed and inoculated into media lacking inducer. Cultures were monitored by OD₆₀₀ and microscopy. To confirm the essentiality of ripA, depletion strains of *M. smegmatis* were grown on a plate with a gradient of inducer containing 50 nM TMA-DPH or 5 μM SYTO 9 for staining membranes or DNA, respectively. Samples were imaged with a Nikon TE-200 100× (NA 1.4) objective and captured with an Orca-II ER cooled CCD camera (Hamamatsu). Final images were prepared using Adobe Photoshop 7.0.

Susceptibility to antibiotics assay

The ripA depletion strain of *M. smegmatis* was spread on LB agar plates containing different amounts of anhydrotetracycline inducer (ng/ml concentrations) to regulate the amount of ripA expressed. A filter disc with 10 μl of carbencillin (100 mg/ml), iodoniazid (10 mg/ml), or cycloserine (100 mg/ml) was placed in the center of the plate and the diameter of inhibition of growth was measured after 4 days of growth.

Microscopy and imaging

*M. smegmatis* strains were centrifuged at 2500 × g for 2 minutes, washed with 1ml PBS, and resuspended in 20 μl of PBS containing 30 nM TMA-DPH or 5 μM SYTO 9 for staining membranes or DNA, respectively. Samples were imaged with a Nikon TE-200 100× (NA 1.4) objective and captured with an Orca-II ER cooled CCD camera (Hamamatsu).
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Author Contributions

Conceived and designed the experiments: EH ER. Performed the experiments: EH MC. Analyzed the data: EH. Contributed reagents/materials/analysis tools: EH MC LD ER. Wrote the paper: EH.