FAD-sequestering proteins protect mycobacteria against hypoxic and oxidative stress

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Liam K. Harold†§, James Antoney†||, F. Hafna Ahmed†, Kiel Hards†, Paul D. Carr*, Trevor Rapson‖, Chris Greening*‡**, Colin J. Jackson*‡, and Gregory M. Cook†∥‡

From the †Department of Microbiology and Immunology, School of Biomedical Sciences, University of Otago, Dunedin 9054, New Zealand, ‡Maurice Wilkins Centre for Molecular Biodiscovery, The University of Auckland, Private Bag 92019, Auckland 1042, New Zealand, ¶Research School of Chemistry, The Australian National University, Canberra, Australia, ■The Commonwealth Scientific and Industrial Research Organisation, Land and Water Flagship, Canberra, Australian Capital Territory, Australia, and **School of Biological Sciences, Monash University, Melbourne, Australia

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The ability to persist in the absence of growth triggered by low oxygen levels is a critical process for the survival of mycobacterial species in many environmental niches. MSMEG_5243 (fsq), a gene of unknown function in Mycobacterium smegmatis, is up-regulated in response to hypoxia and regulated by DosRDosS/DosT, an oxygen- and redox-sensing two-component system that is highly conserved in mycobacteria. In this communication, we demonstrate that MSMEG_5243 is a flavin-sequestering protein and henceforth refer to it as Fsq. Using an array of biochemical and structural analyses, we show that Fsq is a member of the diverse superfamily of flavin- and deazaflavin-dependent oxidoreductases (FDORs) and is widely distributed in mycobacterial species. We created a markerless deletion mutant of fsq and demonstrate that fsq is required for cell survival during hypoxia. Using fsq deletion and overexpression, we found that fsq enhances cellular resistance to hydrogen peroxide treatment. The X-ray crystal structure of Fsq, solved to 2.7 Å, revealed a homodimeric organization with FAD bound covalently. The Fsq structure also uncovered no potential substrate-binding cavities, as the FAD is fully enclosed, and biochemical studies indicated that the Fsq:FAD complex is relatively inert and does not share common properties with electron-transfer proteins. Taken together, our results suggest that Fsq reduces the formation of reactive oxygen species (ROS) by sequestering free FAD during recovery from hypoxia, thereby protecting the cofactor from undergoing autoxidation to produce ROS. This finding represents a new paradigm in mycobacterial adaptation to hypoxia.

Mycobacteria are obligately aerobic bacteria that are able to survive prolonged periods of hypoxia and nutrient limitation. One of the key challenges for mycobacterial persistence during hypoxia is combating the generation of reactive oxygen species (ROS) during re-entry into aerobic conditions and ensuing growth. The majority of ROS production in mycobacterial cells is generated through the autoxidation of molecules with sufficient electrochemical potential to form superoxide (1). Flavins, which include FAD, are one class of molecules that have sufficient potential to autoxidize (in the presence of light or metal ions) (1, 2). In mycobacteria, autoxidation of flavins generates superoxide, which is subsequently converted into hydrogen peroxide by superoxide dismutase. Catalase then in turn catalyzes the decomposition of hydrogen peroxide to water and oxygen (3, 4). Mycobacteria also have a range of other strategies to detoxify ROS, for example using a phenolic glycolipid present in their cell wall to scavenge ROS and mycothiol to reduce disulfide bonds (5–7).

In mycobacteria, a conserved mechanism in the adaptation to low oxygen is the activation of the two-component regulatory system DosRDosS/DosT. The histidine kinases DosS and DosT use a heme group to sense the redox state and oxygen concentration of the cell (8, 9). Both kinases in turn phosphorylate DosR, a LuxR-type helix-turn-helix transcriptional activator that binds 18-bp palindromic sequences in promoter regions (10–12). In the fast-growing soil bacterium Mycobacterium smegmatis, the dos system controls 11 operons and 49 genes in response to oxygen deprivation (13). Many of the genes under Dos control remain uncharacterized.

Fsq is among the mycobacterial proteins of the dos regulon and is highly up-regulated under hypoxia (13). Fsq has recently been identified as belonging to a diverse and largely uncharacterized superfamily of flavin/deazaflavin oxidoreductases (FDORs) that are abundant in mycobacteria and other Actinobacteria (14, 15). This superfamily is characterized by a split β barrel structural fold that generally binds flavin/deazaflavin cofactors. The FDOR superfamily can be further split into two families:

4The abbreviations used are: ROS, reactive oxygen species; FDOR, flavin/deazaflavin oxidoreductase, SSN, sequence similarity network; qPCR, quantitative PCR; VC, vector control; FMN, flavin mononucleotide; LBT, lysogeny broth with 0.05% Tween 80; HdB, Hartmans de Bont; TEV, tobacco etch virus; SEC, size-exclusion chromatography.
the FDOR-A family in which each has $F_{420}$ as a cofactor and the more diverse FDOR-B family that can bind $F_{420}$, FAD, or FMN as cofactors, or heme as a substrate. The FDOR-B family exhibits diverse functions, including pyridoxine 5’-phosphate oxidases (PnPOxs), biliverdin reductases, and heme oxygenases (14).

Understanding of physiological roles of the uncharacterized genes of the dos regulon remains an important goal as they may provide insight into the unique mechanisms of adaptation these bacteria have evolved to survive both the stress of entry into hypoxia and re-entry into aerobic growth conditions during their normal life cycle. In this communication, we determined the role of Fsq in adaptation to hypoxia through a combination of bioinformatic, genetic, biochemical, and physiological approaches.

Results and discussion

Homologues of Fsq are highly conserved among mycobacteria

A previous bioinformatics analysis of the diverse superfamily of FDORs placed Fsq within a discrete FAD-binding subfamily alongside MSMEG_6368 and rv3129 from *Mycobacterium tuberculosis* (14). We compared the genetic organization, expression levels, and primary sequences of Fsq and MSMEG_6368 to gain insight into their possible cellular roles. Based on pairwise alignment, Fsq shares 64% amino acid identity with rv3129 and 54% identity with MSMEG_6368. The fsg and rv3129 genes are both part of the dosR regulon and share similar genomic contexts, with both being in the same region as dosR (Fig. 1a). Conversely, MSMEG_6368 is not part of the dosR regulon and does not share the same genomic context as fsg (13). Quantitative RT-PCR con-

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**Figure 1.** Organization, regulation, and abundance of Fsq. a, genomic context of fsg in *M. smegmatis* and its homolog rv3129 in *M. tuberculosis*. Matching colors between *M. smegmatis* and *M. tuberculosis* represent homologous genes. Gene sizes are not to scale. b, comparison of expression ratio of MSMEG_6368 and fsg, with tgs1 as a positive control, from cells harvested at 6 days’ growth in aerobic and hypoxic conditions. The gene sigA was used as the reference. Error bars represent S.D. of biological triplicates. Statistical testing was performed using an unpaired Student’s t test, where ** = $p < 0.01$. c-f, SSNs of Fsq-related proteins are shown with different cutoffs. In an SSN, protein sequences are represented as nodes (dots) that are joined by edges (lines) which represent a similarity metric such as Basic Local Alignment Search Tool (BLAST) E-value (46). Nodes are colored according to taxonomic order. c, SSN with no logE cutoff applied. d, SSN of Fsq-related proteins with a logE cutoff of $-36$. A large heterogeneous cluster containing Fsq, MSMEG_6368, and rv3129 separates from several small clusters. e, the large heterogeneous cluster examined with a cutoff of $-53$. A large compact cluster of mycobacterial sequences separates from other clusters. f, the large cluster from previous panel with a cutoff of $-67$. A cluster containing MSMEG_6368 separates from one containing Fsq and rv3129, suggesting that they may have distinct functions.
fsq contributes to the survival of Mycobacterium smegmatis under hypoxic growth conditions

To determine the function of fsq, we constructed a markerless deletion mutant strain Δfsq (Fig. 2a) (Table S1). The Δfsq deletion mutant was confirmed by PCR and whole genome sequencing. Strain Δfsq and the isogenic WT parent strain were grown under aerobic conditions on a range of different carbon sources (fermentable and nonfermentable) and no differences in growth rate or final cell yield were observed between the strains (Fig. S1). However, when strains were grown in sealed serum vials that became hypoxic as cells grew (∼37 h), we observed a difference in the final optical density (600 nm) between the Δfsq mutant and the WT (Fig. 2b). To investigate this effect in more detail, we compared the cell viability of the Δfsq mutant to the isogenic WT and a ΔdosR mutant during prolonged incubation under hypoxia (Fig. 2c). The Δfsq mutant exhibited a survival defect compared with WT strain from day 10 post inoculation onwards, with the maximum difference observed being a 2-log difference at 22 days post inoculation (Fig. 2c). Consistent with previous findings (13), the ΔdosR mutant showed a 3-log decrease in viability after 10 days compared with WT. We were able to complement the Δfsq defect under hypoxia using complementing vector (pLH2, fsq+) induced with 20 ng/ml tetracycline (Fig. 2d). After 18 days, there was 0.24% survival for the WT with an empty vector, 0.22% survival for complemented Δfsq, and 0.02% survival for Δfsq mutant with an empty vector (Fig. 2d).

The loss of viability observed for Δfsq mutant cells was initially hypothesized to be caused by Fsq playing a role in the complete activation of DosR given the two genes are in close proximity (Fig. 1a). To address this hypothesis, we determined the effect of the Δfsq mutation on the expression of a dos-regulated gene (hyd3-lacZ) in M. smegmatis (16). When the expression of hyd3-lacZ was measured in the WT background, hyd3-lacZ was up-regulated in response to hypoxia (Fig. 2e). No up-regulation of hyd3-lacZ was observed in the ΔdosR mutant, but WT levels of hyd3-lacZ expression were observed in the Δfsq mutant, confirming that fsq did not play a role in DosR activation.

MSMEG_6368 shares 54% identity with Fsq but is not part of the dosR regulon, suggesting a distinct functional role. To investigate the functional role of MSMEG_6368 and rule out MSMEG_6368 compensating for the loss of fsq in the Δfsq mutant, we created a markerless deletion mutant of MSMEG_6368 (strain ΔMSMEG_6368) (Table S1). We then grew the ΔMSMEG_6368 mutant and the isogenic WT strain on a range of carbon sources (fermentable and nonfermentable) under aerobic conditions and found that there was no detectable difference in the growth rate or final cell yield (Fig. S2, a and b). No phenotypic differences in growth or cell viability between the ΔMSMEG_6368 mutant and the isogenic WT strain were observed under hypoxia (Fig. S2, c and d). We then used qPCR to monitor the expression of MSMEG_6368 in the WT and Δfsq mutant (Fig. S2e). During hypoxia, the expression of MSMEG_6368 was not different in the Δfsq mutant compared with the WT. These data suggest that MSMEG_6368 does not provide compensation for fsq in the Δfsq mutant during hypoxia.

Biochemical, structural, and electrochemical analysis suggests Fsq is an FAD-binding protein

To determine the biochemical function of Fsq, we expressed Fsq in Escherichia coli and purified the protein to homogeneity. Fsq had previously been demonstrated to preferentially bind to FAD over other structurally similar cofactors FMN and F420P with a Kd value of 3.6 μM (14). We set about testing whether Fsq could function as a flavin-sequestering protein by solving and analyzing its X-ray crystal structure and testing its electrochemical properties. Fsq heterologously produced in E. coli copurified with a cofactor that has a UV-visible absorbance spectrum with peaks at 374 nm and 456 nm, which is consistent with FAD (Fig. 3a). To confirm the identity of this cofactor, we crystallized the purified protein (Fig. 3b), obtaining 2.7 Å resolution diffraction (Table 1). The crystal contained 20 molecules in the asymmetric unit, arranged as homodimers (Fig. 3c). Omit electron density (mFo − DFo) consistent with FAD was observed at the dimer interfaces/cofactor-binding sites for 17 of these chains (Fig. 4, a and b). The remaining three chains had no FAD bound at this site, although they were part of dimers in which the other monomer was bound to FAD. There was no electron density to support covalent attachment of FAD to the protein, which is consistent with previous work showing that bound
Figure 2. Creation and hypoxic survival of Δfsq markerless deletion mutant. a, diagram of fsq markerless deletion construction. b and c, long-term survival of M. smegmatis WT (blue) compared with ΔdosR (black) and Δfsq (red) mutants under hypoxic conditions. Growth was in HdB minimal medium supplemented with 22 mM glycerol. Entry of the vials into a hypoxic state was estimated by the decolorization of methylene blue (1.5 g/ml) which occurred at 27 h. b, growth measured using optical density at 600 nm wavelength with an average of three biological replicates shown with error bars representing S.D. Unpaired Student’s t test of WT compared with Δfsq and WT compared with ΔdosR at each time point. *. p < 0.05; **, p < 0.01; ***, p < 0.001; ****, p < 0.0001. d, complementation of survival defect of Δfsq mutant using pLH2 (fsq+) (Δfsq comp) (red) compared with an empty vector control in wild type (WT VC) (blue) and an empty vector in Δfsq mutant (Δfsq VC) (black). Vectors were induced using 20 ng/ml tetracycline just prior to entry into hypoxia. Survival percentage of day 18 with an average of three biological replicates shown with error bars representing S.D. Unpaired Student’s t test of WT compared with Δfsq comp and WT compared with Δfsq VC. *, p < 0.05. e, measurement of DosR activity through hyd3 LacZ expression in M. smegmatis WT compared with ΔdosR and Δfsq mutants under hypoxic conditions. Strains were grown into hypoxia in HdB with 22 mM glycerol. Entry into hypoxia was indicated by methylene blue (1.5 g/ml) control vials. Samples (2 ml) were taken at 27, 48, 72, and 144 h and used to preform β-gal assays to assess the level of dosR activity. Key: WT, blue circles; ΔdosR mutant, black triangles; Δfsq mutant, red squares; β-gal activity, open symbols; optical density (600 nm), closed symbols. Each sample shows average of a biological triplicate with error bars representing S.D.
FAD can be readily removed from Fsq through partial unfolding-refolding (14).

Comparison of the apo- and holo- forms of Fsq revealed a large conformational change in the holo-structure that allows for the accommodation of FAD (Fig. 4, a and b). Most notably, a loop region in the apo-protein forms/H9251-helix-6 in the holo-protein, which interacts with the phosphate groups of FAD (Fig. 4, c and d). Although the apo-protein contains both/H9251-helix-9 and/H9252-helix-10, the position of/H9251-helix-9 is shifted in the holo-protein, causing/H9251-helix-10 to unravel. This lengthens loop-a between/H9251-helix-9 and β-sheet-7, allowing it to accommodate the isoalloxazine group of FAD (Fig. 4, b and d). This substantial change upon cofactor binding is unusual for proteins of the FDOR family, as structurally characterized FMN- and F420-binding proteins show only slight topological differences between their apo- and holo-forms (Fig. S3) (14, 15, 17, 18). This structural rearrangement results in essentially complete enclosure of the cofactor within the interior of the protein. From the pairwise alignment of Fsq and rv3129 we found that the majority of the key residues were maintained except for a proline 45 and leucine 30, which were deleted in rv3129.

Flavin/deazaflavin-binding proteins can typically carry out three broad roles: enzymes that catalyze chemical reactions (19), electron transport proteins that transfer electrons to partner enzymes for catalysis (20), or cofactor-binding proteins that bind to and sequester or transport cofactors (21–23). We therefore sought to test each of these hypotheses. In FDORs with known enzymatic activity, catalysis occurs on the Re-face of the isoalloxazine rings of their flavin cofactors, which is solvent accessible to allow substrate binding (15, 17). In contrast, for Fsq, we found that the isoalloxazine ring of FAD is fully enclosed by the protein, making the Re-face completely inaccessible as a substrate-binding site (Fig. 4e). This is achieved by an offset parallel stacking interaction between the xylene and pyrazine rings of the isoalloxazine and Phe-136 of the secondary chain in the dimer, and loop-a (which undergoes a conformational change upon FAD binding) covers the pyrimidine ring with hydrophobic interactions from Trp-115 from the primary chain (Fig. 4f). The lack of any substrate-binding site will prevent binding or catalysis of small molecule substrates, making it unlikely that Fsq is an enzyme.

Protein film voltammetry was employed to determine the effect of Fsq on the electrochemical properties of the cofactor, which could also provide insight into its physiological role. Both Fsq and FAD displayed a single reversible redox peak at all pH values that we tested, with no evidence of one-electron reductions (Fig. 5a), as is seen in some flavin-dependent enzymes such as fumarate reductase (24). At pH 7.3, the redox potential of FAD was found to be −260 ± 3 mV (versus normal hydrogen electrode), whereas that of Fsq:FAD was −246 ± 9 mV (Fig. 5a). Thus, the redox potential of FAD is not substantially changed by Fsq. This is in contrast to many enzymes and electron-transfer proteins that can alter the redox potentials more than 100 mV in either direction to carry out catalysis (25, 26). There was relatively little difference in the effect of pH on the redox potential of free and complexed FAD (−45 mV/pH unit and −39
mV/pH unit, respectively) with no redox-linked $pK_a$ observed in the range of 3.6–7.3. This contrasts again with electron-transfer proteins, such as flavodoxins, that have a redox-linked $pK_a$ between pH 4.6 and 6.8 (25, 27) (Fig. 5b). The values for free FAD are consistent with previous studies of free FAD on carbon nanotube–modified electrodes (28). In summary, FAD appears to be relatively stable and unreactive when bound to Fsq, making it unlikely that this protein:flavin complex is involved in catalysis or electron-transfer reactions, leaving the possibility that it is involved in FAD sequestration.

**Fsq promotes resistance to oxidative stress**

Based on the structural and biochemical data that suggest Fsq is unlikely to have either a catalytic or electron-transfer role in mycobacterial physiology, and the reduced fitness of the $Δfsq$ strain in hypoxic conditions, we hypothesized one benefit of FAD sequestration by Fsq could be the prevention of FAD autooxidation to avert superoxide formation and cellular damage. One of the key challenges that mycobacterial species must overcome to survive hypoxia is mitigating oxidative damage upon entry to and re-aeration from hypoxia. To test this hypothesis, we challenged the $Δfsq$ mutant and isogenic WT grown in either hypoxia (inducing conditions or aerobic conditions (noninducing) with hydrogen peroxide (Fig. 6). Under aerobic conditions, there was no significant difference in cell viability between the WT and $Δfsq$ mutant in response to hydrogen peroxide challenge (Fig 6a). However, under hypoxic conditions, the $Δfsq$ mutant was significantly ($p = 0.0082$) more sensitive to hydrogen peroxide challenge compared with the WT (Fig 6b). When Fsq was overexpressed in cells using the tetracycline-inducible vector pLH2 ($fsq^+$), we could protect cells from killing by hydrogen peroxide stress compared with the empty vector control (VC) (Fig. 6c). These results establish the molecular process by which Fsq confers a fitness advantage to *M. smegmatis* during recovery from hypoxia. However, they do not explain the fitness advantage of Fsq to *M. smegmatis* during long-term hypoxic growth. Thus, although at least one physiological function of Fsq is now defined, other specific functions related to hypoxic survival could also exist.

**Comparison to other cofactor-binding proteins in mycobacteria**

Having confirmed that Fsq protects *M. smegmatis* against oxidative damage when emerging from hypoxic conditions by sequestering free FAD, we examined whether other examples of mycobacterial cofactor-sequestrer proteins have been reported in the literature. *M. tuberculosis* and *M. smegmatis* have both been shown to harbor a nonfunctional nitroreductase designated Acg that has been shown to function as a flavin mononucleotide (FMN) storage protein or used to sequester FMN from other biochemical pathways (29). Acg is not related in sequence or structure to Fsq. However, as we have observed with Fsq, the Acg structure encloses (“caps”) FMN in a way to sterically prevent reactivity with NAD(P)H. Thus, like Fsq, Acg also appears to have evolved from a superfAMILY in which most proteins are enzymes and, under selective pressure, have lost the ability to bind substrate to develop a cofactor-sequestration function. Notably, the $acg$ gene is also regulated as part of the dosR regulon (29). The observation that mycobacteria has evolved two different proteins that each sequester flavin cofactors to prevent deleterious interactions, both of which are under control of the DosR regulon, highlights the biochemical challenges imposed by the descent into, and emergence from, hypoxia. These examples provide us with deeper understanding of the mechanisms that mycobacteria have evolved to adapt to the pressures associated with hypoxic conditions.

**Conclusions**

Mycobacteria utilize a diverse array of flavins in a wide variety of essential electron-transfer reactions, including FAD, FMN, and F$_{A2O}$ (30). Previously, it has been shown that mycobacteria possess proteins such as Acg to sequester FMN in hypoxic conditions. In this study we infer, using a combination of biochemical, structural, and electrochemical data, that Fsq from *M. smegmatis* functions to sequester FAD under hypoxia. The expression of *fsq* is controlled by DosR and mutants defective in *fsq* are hypersensitive to oxidative stress when recovering from hypoxia. These data suggest that the ability of Fsq to

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**Table 1**

Crystallography statistics for Fsq

| Data collection | Space group | P2 21 21 |
|----------------|-------------|----------|
| Unit cell parameters | a (Å) | 80.27 |
| | b (Å) | 186.04 |
| | c (Å) | 395.39 |
| | α, β, γ (°) | 90, 90, 90 |
| Wavelength (Å) | 1.5418 |
| Resolution range (Å)* | 29.6–2.69 (2.74–2.69) |
| Unique reflections | 82,043 |
| Completeness (%) | 99.9 (99.9) |
| $R_{merge}$* | 0.215 (1.444) |
| $R_{free}$* | 0.084 (0.565) |
| Mean (I/σ(I)) | 10.5 (1.7) |
| CC$_{1/	ext{merge,all}}$ | 0.993 (0.551) |
| Solvent content (% v/v) | 45.7 |
| Molecules per asymmetric unit | 20 |

- $R_{merge} = \langle \delta F_h^2 \rangle \langle \delta F \rangle^2$ where $\langle \delta F \rangle$ is the average intensity of i symmetry-related observations of the unique reflection $h$.
- $R_{free} = \langle \delta F_h^2 \rangle / \langle \delta F \rangle^2$ where $\langle \delta F \rangle$ is the average intensity of a set of reflections randomly chosen from the data set.
- $CC_{1,	ext{merge,all}} = \langle F_1 \rangle - \langle F_2 \rangle / \langle F_1 \rangle \langle F_2 \rangle$ and 5% of the data that were excluded from the refinement were used to calculate $R_{free}$. 

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**Fig. 5**

**Fig. 6**

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sequester FAD protects it from autoxidation and the production of ROS under hypoxia.

Experimental procedures

Strains and growth conditions

*M. smegmatis* mc²155 was grown in either lysogeny broth with 0.05% Tween 80 (LBT) or Hartmans de Bont (HdB) minimal medium supplemented with 0.05% tyloxapol and either 22 mM glycerol or 30 mM succinate as the sole carbon and energy source. When required, media were supplemented with 20 µg/ml kanamycin, 50 µg/ml hygromycin, or 5 µg/ml gentamicin. If solid media were required, 1.5% agar LBT was used. Aerobic growth was carried out in flasks of 125 ml with 25 ml medium or 250 ml with 50 ml medium in rotary incubator shaking at 200 rpm, 37 °C. E. coli strain DH10β was grown in lysogeny broth or M9 minimal medium and, when required, 50 µg/ml kanamycin, 50 µg/ml hygromycin, or 20 µg/ml gentamicin. *M. smegmatis* cells were grown into hypoxia using 30 ml of medium in 120-ml serum vials using the Berney-Cook model of hypoxia where depletion of oxygen was confirmed using a control vial that contained methylene blue (1.5 g/ml) that becomes colorless when low oxygen is present. To count colony-forming units (cfu/ml), each culture was serially diluted in PBS (pH 7.0) and spotted onto agar plates. All survival studies were performed with biological triplicates, and plates were spotted in technical triplicate. Optical density to monitor growth of bacteria were done at 600 nm wavelength using a Jenway 6300 spectrophotometer. Once optical density of cells was above 0.5, the cells were diluted 1 in 10 in 0.85% saline buffer to bring the optical density (600 nm) below 0.5. β-gal assays were performed as described previously.
FAD-sequestering proteins protect mycobacteria

Figure 6. Fsq heightens resistance to hydrogen peroxide. a, challenge of WT compared with ∆fsq with 20 or 50 mM hydrogen peroxide for 3 h. Cells were grown aerobically in HdB with 22 mM glycerol for 10 days before harvest and resuspension in phosphate buffer saline with 0.5% tyloxapol. Cells were normalized to 1 optical density (600 nm) before treatment. b, challenge of WT compared with ∆fsq with 20 or 50 mM hydrogen peroxide for 3 h. Cells were grown hypoxically in HdB with 22 mM glycerol for 10 days before harvest and resuspension in phosphate buffer saline with 0.5% tyloxapol. Cells were normalized to 1 optical density (600 nm) before treatment. c, challenge of WT either expressing fsq (fsq+) (WT pLH2) or a vector control (VC) (WT pMind) control induced with 20 ng µl−1 tetracycline, challenged with 50 or 75 mM hydrogen peroxide for 3 h. Cells were grown aerobically to exponential phase n HdB with 22 mM glycerol before harvest and resuspension in phosphate buffer saline with 0.5% tyloxapol. Cells were normalized to 1 optical density (600 nm) before treatment. a−c, error bars represent S.D. of three biological replicates. Unpaired Student’s t test compared with WT for each treatment, where *p < 0.05; **p < 0.01.

Sequence similarity network

Sequences were obtained via BLAST searches using MSMEG_5243 as the query and were curated with CD-Hit (34) to remove sequences with more than 90% identity. Sequence similarity networks were obtained using an all-versus-all BLAST of the curated sequences as described previously (14) using BLAST + (35). Networks were visualized in Cytoscape 3.6.1 (36) with nodes colored according to taxonomic order and BLAST E-values used as the edges.

Markerless deletion mutant construction

In brief, to generate the deletion of fsq, two homologous regions either side of target gene, we generated and fused together using overlap extension PCR (Primers LH1, LH2, LH3, LH4) (Table S2) and cloned into pX33 using the SpeI restriction enzyme (New England Biolabs). For the deletion of MSMEG_6368, following the initial PCR that generated the two homologous regions either side of target gene (Primers LH7, LH8, LH9, LH10) (Table S2), we used Gibson cloning (New England Biolabs) as per the manufacturer’s instructions to join the homologous regions to pX33. The pX33 constructs with homologous regions were then electroporated into M. smegmatis and grown at 28 °C to ensure permissibility of the pX33 temperature-sensitive plasmid. Confirmation of the plasmid in the cell was achieved by treating cell colonies with 250 mM catechol, which causes any colonies containing the plasmid to turn yellow because of the catechol dioxygenase produced by pX33. Following confirmation of the plasmid presence, the cells were grown on liquid media at 28 °C and then plated 40 °C to force integration based on the homology of the gene flanking regions. Integrants were confirmed through catechol screening (as above). Integrants were then spread-plated on sucrose (10% w/v) containing media to force recombination to either the WT genotype or the markerless deletion mutant. All mutants were confirmed using whole genome sequencing (WGS) and aligned using Geneious to an isogenic WT to confirm that the gene was deleted.

Expression and purification of Fsq

The vector encoding Fsq includes an N-terminal His-tag followed by a TEV protease cleavage site, as described previously (14). E. coli BL21(DE3) (Invitrogen) cells were transformed with the pETMCSIII vector containing the Fsq construct using electroporation. Colones were picked and grown in 5-ml starter cultures before being transferred to 1 liter modified autoinduction medium (5 g of yeast extract, 20 g of tryptone, 85.5 mM NaCl, 22 mM KH_2PO_4, 42 mM NaH_2PO_4, 0.6% glycerol, 0.05% glucose, 0.2% lactose, and 100 µg/ml of ampicillin) and grown for 16 h at 30 °C before centrifugation at 4000 × g for 15 min and removal of supernatant. Cell pellets were frozen and resuspended in 40 ml of buffer A (50 mM NaH_2PO_4, pH 8.0, 300 mM NaCl, 25 mM imidazole) before being ruptured by sonication on ice. Turbonuclease (0.1 µl) (Sigma-Aldrich) was added prior to pelleting cell debris by centrifugation at 20,000 × g for 60 min. The resulting supernatant was then loaded onto a 5-ml HisTrap nickel affinity column (GE Healthcare) pre-equilibrated with buffer A. The column was washed with 25 ml of buffer A containing 25 mM imidazole, 25 ml of 8% buffer B (43 mM imidazole) and eluted in 100% buffer B (250 mM imidazole). Buffer B has the same composition as buffer A except for the addition of 250 mM imidazole. The eluted protein was then passed over a GE HiPrep 26/10 desalting column to buffer exchange into 50 mM Tris-HCl, 200 mM NaCl, pH 8, and His-tag was cleaved by incubating the eluent overnight with TEV protease (purified in-house) (37), 1 mM DTT, and 0.5 mM EDTA. The cleaved His-tag and TEV protease were then removed by passing the reaction mixture over a 5-ml HisTrap nickel affinity column and collecting the flow-through. The sample was concentrated by centrifugation in an Amicon Ultra-15 10 MWCWO spin filter at 4000 × g to a final volume of 2 ml and further purified by size-exclusion chromatography (SEC) using a GE HiLoad 16/600 Superdex 75-ml column with buffer containing 20 mM HEPES, 150 mM NaCl, pH 7.5. The final sample was diluted to adjust buffer concentration to 20 mM HEPES, 50 mM NaCl, pH 7.5, and concentrated to 1.3 ml for storage at 4 °C.

X-ray crystallography

High-throughput screens from Hampton Research were used to identify initial crystal-forming conditions for Fsq using the 1.3 Å protein solution. Crystals grew in 20% PEG 1500, 4% 2-methyl-2,4-pentanediol, and 0.1 M citric acid, pH 3.5. Using the same conditions as a cryo-buffer, data were collected in-
house with a Xenocs GeniX 3D Cu HF microbeam X-ray generator and a MAR345 plate detector. The diffraction data were integrated using XDS (38) and AIMLESS (39) as part of the CCP4 suite (40) was used for scaling. This was followed by molecular replacement with Phaser (41) using the structure of a related protein of unknown function from Corynebacterium glutamicum (PDB ID: 3FKH), which has 45% amino acid sequence identity to Fsq. Initial model building was performed using Buccaneer (42), followed by manual model building in Coot (43). Refinement was performed with Refmac 5.5 (44) and PHENIX 1.13_2998 (45). Omit m_F - DF, electron density was used to model the position of bound FAD molecules.

**DNA manipulation and cloning**

All molecular biology techniques were carried out according to standard procedures. All restriction enzymes and DNA modifying enzymes used were from New England Biolabs and were in their appropriate buffers. All constructs were confirmed using either PCR or restriction digests followed by sequencing. PCR for construct creation was done using Thermo Fisher Phusion polymerase following the recommended conditions with appropriate annealing temperatures and extension time. The construct for *fsq* tetracycline (pLH2–)-inducible expression was made using the pMIND vector backbone as described previously (46) with an artificial ribosome-binding site (47) using the primers LH5 and LH6 to amplify *fsq* (Table S2).

**Quantitative RT-PCR**

Total RNA was extracted using TRIzol (Ambion) as per the manufacturer’s instructions and then purified using the RNA Clean and Concentrator kit (Zymo) as per the manufacturer’s instructions. The cells were lysed by three cycles in a mini-Beadbeater (Biospec Products) at 4800 rpm for 30 s and on ice between each of the cycles. DNA was removed by treatment with 3 units RNase-free DNase using the TURBO DNA-free kit (Ambion) as per the manufacturer’s instructions. The RNA concentration was determined using a NanoDrop One spectrophotometer. Reverse transcriptase PCR was performed using SuperScript III (Invitrogen), according to the manufacturer’s instructions for cDNA synthesis. After cDNA synthesis, qPCR was carried out using Platinum SYBR Green qPCR SuperMix-UDG with ROX (Invitrogen). The qPCR reactions were carried out on a QuantStudio 6 (Applied Biosystems). As an internal control and for the normalization of results the gene *siga* was used. Error bars are representative of biological triplicate. Primers used are listed in Table S2 (MSMEG_6368 = LH11, LH12; *fsq* = LH13, LH14; *tgs1* = tgsfw, tgsrev).

**Hydrogen peroxide challenges**

Cells were grown in HdB minimal media with glycerol as the sole carbon and energy source. Cell were then harvested and resuspended in PBS (pH 7.5) with 0.05% tyloxapol normalized to an *A_{600}* of 1.0 and then treated with 20, 50, or 75 mM hydrogen peroxide and left shaking at 180 rpm at 37°C for 3 h. Cells were then plated using serial dilutions 5 μl spots on LBT solid media to determine cell viability.

**Electrochemistry**

Cyclic voltammetry was carried out using a BASi Epsilon Potentiostat with a C3 cell stand. A three-electrode system was employed comprising a modified glassy carbon electrode as working electrode, Pt wire counter and Ag/AgCl-saturated NaCl reference electrode. All potentials are quoted versus the normal hydrogen electrode, calculated as Ag/AgCl + 196 mV. The working electrode was prepared by polishing according to the manufacturer’s instructions using 0.05 mm alumina polish, followed by sonication for 5–10 min in Milli-Q water. Carbon nanotubes (single-walled COOH-functionalized 1–2 nm, nanostructured and amorphous materials, outer diameter 1–2 nm) were dispersed in Milli-Q water using sonication to give a concentration of 1 mg/ml. SEC-purified Fsq (~1 μg) was mixed with an equal volume of nanotube suspension and applied directly to the freshly polished electrode surface and allowed to dry at ambient temperature. Films prepared with protein-free FAD were used as controls. Oxygen was removed from the buffer through purging with argon. The accession number for MSMEG_5243 is PDB ID: 6ECI.

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