SUPPLEMENTARY DATA LEGENDS

Fig. S1: Rv1813 is detected in M. tuberculosis culture filtrate but not in cellular fractions. Lysate obtained

from M. tuberculosis H37Rv strain grown in Sauton's medium to logarithmic phase (OD580~0.7) were frac-

tionated and probed with anti-Rv1813 polyclonal antibodies. Culture filtrates were obtained from the

same culture. Anti-FtsZ (FtsZ is a cytoplasmic protein), anti-GlnA (GlnA is a membrane protein) and

anti-RpfB (RpfB is a membrane and cell-wall anchored protein) antibodies were used to confirm the purity

of mycobacterial fractions.

Fig. S2: <sup>1</sup>H-<sup>15</sup>N HSQC spectrum of Rv1813c

This spectrum was obtained at 800 MHz, 20°C and pH 6,8 with 0.3 mM <sup>15</sup>N-uniformly labeled sample.

Cross peak assignments are indicated using the one-letter amino acid code and number following the full-

length protein sequence numbering.

Fig. S3: Confocal microscopy analysis of Rv1813c family members localization in Dictyostelium.

Dictyostelium cells expressing the indicated constructs were fixed, processed for immunofluorescence,

and analyzed by confocal microscopy (Airyscan).

(A) Cellular localization of Nt-GFP-tagged Rv1813c. (B) Mitochondrial localization of untagged Rv1813c

expressed in Dictyostelium. Rv1813c was labeled with a rabbit pAb anti-Rv1813c antibody. Rv1813c col-

ocalizes with Mitoporin, a mitochondrial specific protein. (C) Mitoporin localization in untransfected pa-

rental Ax2 cells. Cells were labeled with a mouse mAb to mitoporin revealing characteristic ring shaped

structures. A maximum projection of Z confocal sections is shown on the right panel. (D) Localization of

different Rv1813c family members of M. tuberculosis and M. marinum expressed in Dictyostelium and

revealed by anti-myc labelling. White arrows indicate so mitochondria with affected shapes. Bar, 5 μm.

Fig. S4: Rv1813c family localization in HeLa cells

HeLa cells expressing the indicated constructs were fixed, processed for immunofluorescence, and ana-

lyzed by confocal microscopy (Airyscan). Cells were colabeled either with rabbit polyclonal anti-Rv1813c,

mouse mAb anti-Cytochrome c, and mitotracker deep red (upper panel) or anti-myc, rabbit anti-grp75

(mitochondria marker) and mitotracker deep red (lower panels). Bar, 10 µm.

Supplementary table 1: NMR and refinement statistics for RV1813 protein structures

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