The multifunctional enzyme S-adenosylhomocysteine/methylthioadenosine nucleosidase is a key metabolic enzyme in the virulence of Salmonella enterica var Typhimurium

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Key physiological differences between bacterial and mammalian metabolism provide opportunities for the development of novel antimicrobials. We examined the role of the multifunctional enzyme S-adenosylhomocysteine/Methylthioadenosine (SAH/MTA) nucleosidase (Pfs) in the virulence of S. enterica var Typhimurium (S. Typhimurium) in mice, using a defined Pfs deletion mutant (i.e. Δpfs). Pfs was essential for growth of S. Typhimurium in M9 minimal medium, in tissue cultured cells, and in mice. Studies to resolve which of the three known functions of Pfs were key to murine virulence suggested that downstream production of autoinducer-2, spermidine and methylthioribose were non-essential for Salmonella virulence in a highly sensitive murine model. Mass spectrometry revealed the accumulation of SAH in S. Typhimurium Δpfs and complementation of the Pfs mutant with the specific SAH hydrolase from Legionella pneumophila reduced SAH levels, fully restored growth ex vivo and the virulence of S. Typhimurium Δpfs for mice. The data suggest that Pfs may be a legitimate target for antimicrobial development, and that the key role of Pfs in bacterial virulence may be in reducing the toxic accumulation of SAH which, in turn, suppresses an undefined methyltransferase.

Introduction

Increasing antibiotic resistance is driving the search for novel antimicrobial drug targets and the metabolic differences between prokaryotes and eukaryotes are seen as attractive areas for target discovery [1–5]. While there is considerable overlap between eukaryotes and prokaryotes in core metabolism, there are some important differences [6–8]. The bacterial quorum-sensing systems appear to be unique and the production of the quorum-sensing molecule autoinducer-2 (AI-2) is fed by the activated methyl cycle which metabolises methionine (Met) to produce the key methyl donor S-adenosylmethionine (SAM) [9–15]. The activated methyl cycle recycles Met from SAM (Figure 1). SAM is produced from Met by the SAM synthetase (MetK) in a reaction that is essential to the bacterial cell [16–19]; enteric bacteria such as E. coli and Salmonella enterica var Typhimurium (S. Typhimurium) have no transporters for SAM, and null mutants of MetK are lethal in the bacteria [16,17,20]. As the methyl group is donated from SAM, demethylated SAM is produced, known as S-adenosylhomocysteine (SAH). SAH is
immediately converted to homocysteine and is finally recycled to Met to complete the methyl cycle. SAM is the predominant methyl donor in bacteria and is responsible for methylation of a variety of macromolecules, including DNA, by different SAM-dependent methyltransferases [18,21,22].

In S. Typhimurium and E. coli, the conversion of SAH to homocysteine is a two-step process. SAH is converted to S-ribosylhomocysteine (SRH) by SAH nucleosidase (abbreviated to Pfs) and then from SRH to homocysteine using S-ribosylhomocysteine lyase (LuxS) [23]. AI-2, an integral component of quorum sensing, is produced during the conversion of SRH to homocysteine [9–15]. AI-2 appears to function in intraspecies and interspecies communication, driven by cell density, and controls phenomena including bacterial swarming [9–15]. In eukaryotes and some bacteria, SAH is converted to homocysteine directly through a one-step process by S-adenosylhomocysteine hydrolase (SahH) [10,24,25].

Pfs has two other well-characterised enzymatic activities: it is required for the conversion of methylthioadenosine (MTA) to methylthioribose (MTR) [23], and is indirectly involved in the biosynthesis of biotin, a key vitamin, through the conversion of 5'-deoxyadenosine to 5'-deoxyribose and adenine [26,27]. The nucleoside 5'-deoxyadenosine is an inhibitor of biotin synthase (BioB), and therefore Pfs activity indirectly affects biotin synthesis (Figure 1) [26,27].

The aim of this study was to investigate the ‘essentiality’ of de-methylated SAM salvage and more specifically Pfs in the virulence of S. Typhimurium, and to define the reason(s) why bacteria lacking Pfs were grossly attenuated in mice. Given the multiple reactions mediated by Pfs, its veracity as an antibiotic target will be defined by redundancy and the potential ‘work arounds’ that bacteria typically deploy to effect resistance.

**Experimental procedures**

**Bacterial strains and growth conditions**

The bacterial strains and plasmids used in this study are listed in Table 1. All mutant strains were constructed on the S. Typhimurium SL1344 genetic background and SL1344 was used as the wild-type strain in all
The streptomycin-resistant SL1344 strain has been described previously [28]; its virulence is well-defined. Appropriate antibiotics, including streptomycin (50 μg/ml), chloramphenicol (30 μg/ml), kanamycin (50 μg/ml) and ampicillin (100 μg/ml) were added to growth media as required. To obtain mid-exponential growth phase, S. Typhimurium and E. coli strains were grown in 10 ml Luria-Bertani (LB) broth (BD Difco) overnight, and 100 ml of the overnight culture was sub-cultured into 10 ml fresh LB broth, and grown with shaking (180 rpm) at 37°C for 3–4 h until the optical density reading at 600 nm (OD600) reached 0.6–0.8. Growth phenotypes were characterised in LB broth or M9 minimal medium (2 mM MgSO4, 0.1 mM CaCl2, 0.4% glucose, 8.5 mM NaCl, 42 mM Na2HPO4, 22 mM KH2PO4, 18.6 mM NH4Cl and 100 μM histidine) with or without Met or biotin supplemented as required.

For the quantitation of SAH, bacteria from overnight LB cultures were harvested by centrifugation at 4°C, and the pellets washed twice in 1× PBS before being resuspended in 1× PBS to an equivalent cell density based on OD600 absorbance values. A 1 in 10 dilution was used to inoculate 20 ml M9 minimal medium (supplemented with 100 μM histidine, 50 μg/ml streptomycin and, for strains carrying the pACYC184 plasmid, 30 μg/ml chloramphenicol) prior to incubation with shaking (200 rpm) at 37°C for 24 h. Five replicates of each strain were set up. The following day, the cell density of each culture was measured at OD600 before harvesting the bacteria as described above, cells were washed in 1× PBS and resuspended to the same cell density. Aliquots were centrifuged and the cell pellets snap-frozen in liquid nitrogen before LC–MS analysis.

For growth curves, 200 μl of M9 minimal medium (supplemented with 100 μM histidine, 100 μM Met, 50 μg/ml streptomycin and, for strains carrying the pACYC184 plasmid, 30 μg/ml chloramphenicol) was dispensed into multiple wells of a flat-bottom 96-well plate and inoculated 1 in 100 with bacterial strains of equivalent cell density that had been harvested from overnight LB cultures and washed with 1× PBS before resuspending in M9 medium. Growth was monitored for 22 h at 37°C with intermittent shaking using a FLUOstar Omega microplate reader (BMG LABTECH) set to detect absorbance at 600 nm wavelength.

### Construction of strains and plasmids

Defined deletions of the sequence encoding the pfs, speD, speE, luxS and bioB genes were generated in S. Typhimurium SL1344. The mutant Δpfs was complemented by introducing the pfs from S. Typhimurium and sah gene from Legionella pneumophila into Δpfs strain in trans. These strains are named as Δpfs + pACYC184::pfs and Δpfs + pACYC184::sahH, respectively. Gene deletions and concomitant insertions of an

### Table 1 Strains and plasmids used for this study

<table>
<thead>
<tr>
<th>Strain or plasmid</th>
<th>Relevant phenotypes and genotypes</th>
<th>Source</th>
</tr>
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<tr>
<td><strong>Strains</strong></td>
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<tr>
<td>Salmonella enterica Typhimurium SL1344</td>
<td>Wild-type strain rpsL hisG46; StrR</td>
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<td>Salmonella enterica Typhimurium BRD666</td>
<td>Restriction negative modification positive (r m+) SL1344; StrR</td>
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<td>Escherichia coli DH5α</td>
<td>Cloning strain</td>
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<td><strong>Plasmids</strong></td>
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<tr>
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<td>pACBSR</td>
<td>Medium-copy number, mutagenesis plasmid; p15A ori; Ara-inducible I-Scel and λ Red recombinase; ChlR</td>
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<td>pACYC184</td>
<td>Medium-copy number cloning vector, p15A ori; TefR, ChlR</td>
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<td>This study</td>
</tr>
<tr>
<td>pACYC184::sahH</td>
<td>L. pneumophila sahH cloned into pACYC184; ChlR</td>
<td>This study</td>
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<tr>
<td>pCP20</td>
<td>FLP recombinase, temperature-sensitive replicon; ApR, ChlR</td>
<td>[74]</td>
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<tr>
<td>pKD4</td>
<td>FRT-flanked KmR cassette; ApR, KmR</td>
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Str, streptomycin; Ap, ampicillin; Chl, chloramphenicol; Km, kanamycin; Ara, arabinose; Flp, flippase; FRT, flippase recombinase target.

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antibiotic resistance cassette were constructed using a Lambda Red-mediated ‘gene gorging’ method [29]. All constructs were verified by PCR and moved to a SL1344 or relevant background via P22 phage transduction [28,30]. Primers used to construct mutants are listed in Table 2. Genetic complementation in the complemented Δpfs+pACYC184::pfs strain was achieved by cloning pfs into pACYC184 via the BamHI and SphI sites and then introducing pACYC184::pfs into the Δpfs mutant. Genetic complementation in the complemented Δpfs+pACYC184::sahH strain was achieved by attaching a 3× FLAG-tag sequence at the 5’ end of sahH, then cloning sahH into pACYC184 via the BamHI and SphI sites and then introducing pACYC184::sahH into the Δpfs mutant. All constructs were verified by restriction analysis and DNA sequencing. Key mutants were also sequenced using Illumina whole genome sequencing and analysed using the pipeline at the Wellcome Sanger Institute, Hinxton, U.K.

**Infection of epithelial cells**

Infection of HeLa cells (sourced from American Type Culture Collection (ATCC)) was conducted using established gentamicin protection assay protocols [31,32]. Briefly, HeLa cells were grown in DMEM supplemented with 10% heat-inactivated fetal calf serum (FCS) and 2 mM l-glutamax (Life Technologies), in a humidified 37°C, 5% CO2 incubator. One day prior to infection, HeLa cells were seeded in 24-well plates at 2 × 10^5 cells per well. S. Typhimurium strains were grown to mid-exponential phase prior and frozen in LB with 10% glycerol at −80°C. Bacteria were thawed immediately before infection, washed in antibiotic-free tissue culture medium, and resuspended to an OD600 of 0.1 in medium. HeLa cells were infected at a multiplicity of infection of 100:1, and 1 h post-infection, the multwell plates were washed and incubated for 4 h to allow for intracellular replication. The plates were then washed and overlaid with agar to detect surviving bacteria.

<table>
<thead>
<tr>
<th>Mutants</th>
<th>Sequence (5’−3’)</th>
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<tr>
<td>Δpfs-IsceI-F</td>
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<tr>
<td>Δpfs-IsceI-R</td>
<td>TGGGATAACAGGGTAATGGTAAACAGAGGATTCATGCC</td>
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<tr>
<td>Δpfs-Kan-F</td>
<td>GAAGCAGCTCCAGCTACACA CTGTTCAACTGCCAGTG</td>
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<tr>
<td>Δpfs-Kan-R</td>
<td>GAAGCAGCTCAGCTACACA GAAGCAGCTCAGCTACACA</td>
</tr>
<tr>
<td>ΔspeD-IsceI-F</td>
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<tr>
<td>ΔspeD-IsceI-R</td>
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<tr>
<td>ΔspeD-Kan-F</td>
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</tr>
<tr>
<td>ΔspeD-Kan-R</td>
<td>GAAGCAGCTCCAGCTACACA GAAGCAGCTCAGCTACACA</td>
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<tr>
<td>ΔluxS-IsceI-F</td>
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<td>ΔluxS-IsceI-R</td>
<td>TGGGATAACAGGGTAATGAAACACACACGATCATG</td>
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<tr>
<td>ΔluxS-Kan-F</td>
<td>GAAGCAGCTCCAGCTACACA CTGTTCAACTGCCAGTG</td>
</tr>
<tr>
<td>ΔluxS-Kan-R</td>
<td>GAAGCAGCTCCAGCTACACA GAAGCAGCTCAGCTACACA</td>
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<tr>
<td>ΔbioB-IsceI-F</td>
<td>TGGGATAACAGGGTAATGAAACACACACGATCATG</td>
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<tr>
<td>ΔbioB-Kan-R</td>
<td>GAAGCAGCTCCAGCTACACA GAAGCAGCTCAGCTACACA</td>
</tr>
</tbody>
</table>

Endonuclease restriction sites are underlined; kanamycin cassette specific sequences are in bold; F, forward (5’) primer; R, reverse (3’) primer.
media and diluted in DMEM media with or without L-Met, then added to HeLa cell monolayers at a multiplicity of infection (MOI) of 5–10. The colony-forming units (cfu) in the inoculum were estimated by plating on LB agar plates. Infected HeLa cells were centrifuged at 600×g for 5 min immediately after the addition of bacteria and then incubated for 1 h at 37°C. After 1 h, the tissue culture media was replaced with DMEM media with or without Met and containing 100 μg/ml of gentamicin to kill extracellular bacteria. The concentration of gentamicin was reduced to 10 μg/ml at 2 h post-infection and maintained for the remainder of the experiment. To enumerate intracellular bacteria, cells were washed twice with PBS, and lysed with 1% Triton X-100 (Sigma) for 15 min to release the intracellular bacteria. The bacteria were enumerated by plating appropriate dilutions on LB agar plates.

**Ethics statement**

All animal research conducted in this study was approved by the Animal Ethics Committee (Biochemistry & Molecular Biology, Dental Science, Medicine, Microbiology & Immunology) at The University of Melbourne, under project number 1413141. All experiments were conducted in accordance with the Australian Code of Practice for the Care and Use of Animals for Scientific Purposes, 8th edition 2013.

**Mouse infections**

All animal experiments were carried out at the Biological Research Facility (BRF) at the Department of Microbiology and Immunology, The University of Melbourne, at Peter Doherty Institute for Infection and Immunity. Age- and sex-matched C57BL/6 mice were used between 6–8 weeks of age and all 144 mice used as sources of data in this manuscript were euthanased prior to isolation of infected tissues. To assess the virulence of mutant strains, either the intravenous or oral route of infection was used. For intravenous infections, 200 cfu of each bacterial strain was prepared in 200 μl PBS and injected into the lateral tail vein. For oral infections, mice were orally gavaged with 100 μl 10% sodium bicarbonate immediately before oral gavage of 100 μl containing ~5×107 cfu of S. Typhimurium strains. To prepare the inoculum, all strains of S. Typhimurium were grown shanking at 180 rpm in M9 minimal medium (for S. Typhimurium Δψfs mutant the M9 medium was supplemented with 100 μM Met and 100 μM biotin) at 37°C for 24 h, and stored in 10% glycerol at −80°C until use. Immediately prior to infection, the stored aliquots were thawed and diluted in PBS to the required concentration. At designated time points post-infection, the spleen and liver were removed aseptically, homogenised using the Stomacher 80 Biomaster paddle blender (Seward), and serial dilutions were plated on LB agar plates with 50 μg/ml streptomycin to determine the bacterial load.

**Western blot analysis**

The expression of 3×FLAG:SahH (from pACYC184::sahH) was detected by Western blot analysis using a monoclonal mouse Anti-FLAG antibody (Sigma) at a concentration of 1:2000. S. Typhimurium whole-cell lysates were prepared from overnight cultures. Samples were separated by sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis (PAGE) and transferred to iBlot2 NC Regular stack nitrocellulose (SDS)-polyacrylamide gel electrophoresis (PAGE) and transferred to iBlot2 NC Regular stack nitrocellulose (MF-ChemiBis 3.2).

**Mass spectrometry analysis**

Cell pellets were thawed and metabolites were extracted with 125 μl of 80% acetonitrile and briefly vortexed. Samples were centrifuged for 10 min at 14 000×g and the supernatant transferred to glass vials for LC-MS analysis. SAM was detected on an Agilent 6545b using a zic-pHILIC column (150×4.6 mm, Merck) as previously described [33]. Briefly, data was acquired in full scan mode (60–1200 m/z), and the retention time of SAM was
confirmed with a synthetic standard (Sigma). All features were extracted in the MAVEN software package [34] prior to file conversion with MSConvert. Samples were median normalised and the SAM integrated peak was plotted using GraphPad Prism.

Sequence similarity network analysis
Sequence similarity network analysis: representative protein sequences for MTA/SAH nucleosidase (InterPro family IPR010049 – 3240 sequences) and Adenosylhomocysteinase (InterPro family IPR034373 – 4657 sequences) were extracted from the uniref90 database. All sequences were used as a database to perform an all-against-all pairwise BLAST and a sequence similarity network was generated and assessed with CLANS [35] with E-value cut off as described in the figure legend.

DNA methylation analysis
Strains were grown for 24 h in M9 minimal medium supplemented with Met and appropriate antibiotics. Cells were incubated in lysis buffer (10 mM Tris pH 8.0 containing 0.38% SDS, 2 mg/ml lysozyme, 0.9 mg/ml Proteinase K, and 200 μg/ml RNase A) for 30 min at 37°C, followed by 30 min at 55°C. Genomic DNA was extracted with phenol/chloroform using 5PRIME Phase Lock Gel light tubes (Quintabio), precipitated with 3 M sodium acetate and isopropanol, then washed with 70% (v/v) ethanol. DNA concentration was measured using a NanoDrop 2000 spectrophotometer and visualised after gel electrophoresis through a 0.6% (w/v) agarose gel stained with SYBR Safe DNA gel stain (Invitrogen). Detection of DNA methylation in the form of 5-methylcytosine (5-mC) was performed using the MethylFlash Global DNA Methylation (5-mC) ELISA colorimetric kit as per the manufacturer’s instructions (Epigentek). Duplicate samples (100 ng/well) of each biological replicate were screened in the array and the absorbance read at OD450 using a FLUOstar Omega microplate reader. Percentage of 5-mC DNA in total DNA was calculated after determining the slope of the standard curve using linear regression and the formula: (sample OD − neg control OD)/(slope × 100 ng) × 100%. The presence of dam methylation was assessed by digestion of 1 μg genomic DNA overnight at 37°C with restriction enzymes DpnI, DpnII and MboI, and methylation non-sensitive restriction enzyme Sau3A as per manufacturer’s instructions (NEB). Digested DNA was visualised using gel electrophoresis, as described above, and imaged with the Syngene G:BOX gel documentation system.

Results
S. Typhimurium Δpfs mutant is highly attenuated in mice
A Pfs mutant of virulent S. Typhimurium SL1344 was constructed by allelic exchange and the mutant was sequenced. The role of Pfs in the virulence of S. Typhimurium was examined by testing the growth of the mutant in C57BL/6 mice, and growth was compared with that of the wild-type SL1344 (Figure 2). There was 10 000-fold less S. Typhimurium Δpfs recovered from the spleen or liver of infected mice, compared with wild-type SL1344; complementation with pfs on pACYC184 restored in vivo growth of the bacterium to wild-type levels. C57BL/6 mice were also orally infected with 5 × 107 cfu of SL1344 or the Δpfs mutant (Supplementary Figure S1). The mice were euthanised on day 6 post-infection, the spleen and liver was removed, and the number of bacteria was enumerated by viable count. Again, 10 000-fold less S. Typhimurium Δpfs were recovered compared with SL1344.

These data indicate that MTA/SAH nucleosidase (Pfs) is essential for the replication of S. Typhimurium in the C57BL/6 mouse, which is known to be highly sensitive to infection [36]. To explore why this attenuation occurred, S. Typhimurium Δpfs was investigated for in vitro growth characteristics.

S. Typhimurium Δpfs is an auxotroph requiring biotin or Met for growth under minimal nutrient conditions
The metabolic phenotype of the S. Typhimurium Δpfs mutant was determined in M9 minimal medium. S. Typhimurium Δpfs was unable to grow in M9 minimal medium (Figure 3A), but replicated in nutrient-rich LB medium (Figure 3B), suggesting that the Δpfs mutant is an auxotroph that requires additional nutrients to grow under the nutrient restrictions imposed by M9 medium. To confirm that the deleted pfs gene was responsible for auxotrophy, the replication of S. Typhimurium Δpfs+pACYC184:pfs in M9 minimal medium and LB medium was assessed (Figure 3A,B). This complemented strain grew in M9 minimal medium at levels
comparable to the wild-type, demonstrating that the growth restored in the mutant was dependent on the presence of a functional Pfs.

To further characterise the specificity of auxotrophy in *S. Typhimurium Δpfs*, the bacterium was grown in M9 media supplemented with either 100 μM Met or 100 μM biotin. Supplementation of either Met or biotin supported the growth of the Δpfs mutant at near wild-type levels in M9 (Figure 3C). The study was expanded by titrating the effects of biotin and Met on growth. *S. Typhimurium Δpfs* grew to similar levels as SL1344, albeit at a slower growth rate over 60 h, in M9 minimal medium that contained reducing levels of biotin and Met, and down to 0.1 μM of either (Figure 3C). To test whether the attenuation seen in mice was reflected by reduced intracellular growth in tissue culture cells, the Pfs mutant was used to infect HeLa cells (Figure 3D). Again, growth attenuation was observed, which was restored when the Pfs mutant was complemented with pACYC184 carrying pfs. Given that Met could reverse the auxotrophy observed in M9 medium, the growth in HeLa cells of *S. Typhimurium Δpfs* was compared with mutants unable to both synthesise Met *de novo*, through deletion of *metB*, or by mutating the two Met synthases (MetE and MetH), and transport Met through the high-affinity transporter which consists of MetN, MetI and MetQ [37]. These bacteria that were deficient in Met import and *de novo* synthesis grew intracellularly in HeLa cells (Figure 3D), unlike *S. Typhimurium Δpfs*, which replicated only very slowly inside cells (Figure 3). These data suggest that the growth attenuation observed for *S. Typhimurium Δpfs* in HeLa cells was not due to a Met auxotrophy, but might have been due to suppression of biotin synthesis by accumulating 5′-deoxyadenosine.

To test whether a BioB mutation phenocopied *S. Typhimurium Δpfs*, a defined mutant was constructed and tested in M9 minimal medium, for growth in HeLa cells, and for virulence in mice. *S. Typhimurium ΔbioB* was an auxotroph in M9 (Figure 4A), that was reversed by the addition of biotin (Figure 4B). *S. Typhimurium ΔbioB* was able to grow in HeLa cells (Figure 4C) and was at least as virulent as *S. Typhimurium SL1344* when mice were infected i.v., measured by growth of the *S. Typhimurium ΔbioB* in the liver or spleen of infected mice, examined 5 days post-infection (Figure 4D,E, respectively).

Figure 2. *S. Typhimurium* lacking Pfs is severely attenuated in mice.
C57BL/6 mice were i.v. infected with 200 cfu of wild-type *S. Typhimurium* SL1344 or the Δpfs mutant. On day 5 post-infection, (A) the liver and (B) spleen were harvested to determine the bacterial load in the organs. Symbols represent data from individual animals, and horizontal lines represent the geometric mean of each group. Error bars represent the standard error of the mean (SEM). Data are pooled from three independent experiments. One-way ANOVA with Bonferroni post-test was used for statistical tests; ns = not significant (P > 0.05).

*S. Typhimurium ΔspeD, ΔspeE* and *ΔluxS* mutants are not attenuated for growth in HeLa cells or in mice
We next aimed to determine why *S. Typhimurium Δpfs* was attenuated *in vivo*. Given *S. Typhimurium* with mutations in *de novo* Met biosynthetic pathways showed normal growth and virulence in mice [37], the
Figure 3. S. Typhimurium Δpfs mutant is unable to grow in M9 minimal medium and is attenuated for intracellular growth in HeLa cells.

(A) M9 minimal medium, (B) LB nutrient medium, or (C) M9 minimal media supplemented with indicated concentration of biotin or Met was inoculated with 200 cfu of S. Typhimurium wild-type SL1344 or Δpfs mutant, and grown aerobically at 37°C.
Figure 3. S. Typhimurium Δpfs mutant is unable to grow in M9 minimal medium and is attenuated for intracellular growth in HeLa cells.

The viable bacterial count was determined at indicated time points. Horizontal bars represent the geometric mean of each group, and error bars represent the standard deviation (SD). Dotted lines represent the detection limit for bacterial concentration. Data are pooled from two independent experiments. D) S. Typhimurium wild-type and mutant strains were tested for intracellular replication in HeLa cells. HeLa cells were grown to a monolayer and infected with S. Typhimurium wild-type or mutant strains at a multiplicity of infection (MOI) of 5–10, in complete DMEM. The intracellular bacterial load at 2 h post-infection is expressed as ‘1’ and used as the reference point to calculate fold-change of intracellular bacterial number at subsequent time points. Data are pooled from three independent experiments. Bars represent the mean fold-change and error bars show the SD. Unpaired t-tests were used to compare the intracellular load of wild-type and mutant strains at 10 hr post-infection, and P-values shown were corrected for multiple comparisons using the Bonferroni-Dunn method; ** indicates P < 0.01, and P-values greater than 0.05 were deemed as not significant (ns).

apparent auxotrophy for Met, as was observed in M9 minimal medium, does not readily explain why S. Typhimurium Δpfs was highly attenuated in mice. Therefore, further mutants were constructed to determine if the attenuation in Δpfs was due to either deficiency of downstream metabolites, such as the polyamine, spermidine, or AI-2. To test this, we generated mutants carrying defined deletions in speD and speE, responsible for spermidine biosynthesis, and luxS, the S-ribosylhomocysteine lyase that produces the quorum sensing ligand, AI-2 (Figure 1).

The three mutants were grown in M9 minimal medium and none displayed growth characteristics that were different from the wild-type (Figure 5A). The Δpfs mutant was also tested for swarming and the swarming phenotype was severely reduced with the deletion of Pfs, to levels equal to that observed in a non-flagellated mutant (Supplementary Figure S2). As expected, the ΔluxS mutant also showed a swarming deficiency (data not shown). To determine whether S. Typhimurium ΔspeD, ΔspeE and ΔluxS mutants grew in mammalian cells, HeLa cells were infected with bacterial strains and the number of intracellular bacteria determined over a 10-hr period. All three mutant strains (ΔspeD, ΔspeE and ΔluxS) grew at similar rates to the wild-type in HeLa cells (Figure 5B). We then examined the role of SpeD, SpeE and LuxS enzymes in the virulence of S. Typhimurium in C57BL/6 mice. Mice were i.v. infected with 200 cfu of S. Typhimurium wild-type or mutant strains, and the bacterial load in the spleen and liver was determined at day 5 post-infection. There was no significant difference in the number of bacteria recovered from the spleen or liver of mice infected with ΔspeD, ΔspeE and ΔluxS when compared with the bacterial burden found in the organs of mice infected with wild-type SL1344 (Figure 5C,D). This result indicates that the enzymes SpeD, SpeE and LuxS are not essential for growth of S. Typhimurium in C57BL/6 mice, and that the in vivo growth attenuation observed in S. Typhimurium Δpfs is unlikely to be due to loss of AI-2 (LuxS) or spermidine, the precursor of spermidine, decarboxylated SAM, or MTA. These data do not rule out growth attenuation through the accumulation of MTA or SAH.

The removal of S-adenosylhomocysteine (SAH) from the S. Typhimurium Δpfs mutant restores virulence

In E. coli and Salmonella, the conversion of SAH to homocysteine is a two-step process that occurs via S-ribosylhomocysteine (SRH) [23]. SAH, through Pfs, is converted to SRH, and SRH is converted into homocysteine via LuxS, liberating the quorum sensing AI-2 [9–15,23] (Figure 1). The activated methyl cycle is reinitiated when Met is synthesised from homocysteine by one or both Met synthases [23]. In many eukaryotes and some prokaryotes, such as Legionella pneumophila, this cycle is different, and SAH is converted directly to homocysteine through S-adenosylhomocysteine hydrolase (SahH) [10,24,25]. Previous reports suggest that SAH is a potent inhibitor of SAM-dependent methyltransferase [38–40]. Hence, we hypothesised that the heterologous expression of an exogenous SAH hydrolase (i.e. SahH) in S. Typhimurium Δpfs, might resolve whether the accumulation of the putatively inhibitory metabolite (i.e. SAH) was important in the attenuation of the Δpfs mutant, as SahH restores only one of the three enzymatic functions of Pfs. To test this hypothesis, S. Typhimurium Δpfs + pACYC184::sahH was constructed expressing SahH from Legionella pneumophila and expression of the Legionella enzyme confirmed by Western blot (Figure 6A). Legionella SahH does not have significant protein sequence homology with the enterobacterial Pfs and is of a different length, and similar to the human enzyme (Supplementary Figure S3).
Figure 4. The *S. Typhimurium* ΔbioB mutant is a biotin auxotroph that shows normal growth in HeLa cells, and is fully virulent in mice.

(A) *S. Typhimurium* ΔbioB was unable to grow in M9 minimal medium without biotin supplementation. (B) The growth of *S. Typhimurium* in M9 was restored with addition of 100 μM biotin. Geometric mean ± SD is shown. Dotted lines represent the detection limit for bacterial concentration. Data are pooled from two independent experiments. (C) *S. Typhimurium* ΔbioB grew to similar levels as wild-type SL1344 over a 10-h period in HeLa cells, as described for Figure 3. Data are pooled from three independent experiments. Bars represent the mean fold-change and error bars show the SD. Unpaired *t*-test was used to compare the intracellular load of wild-type and mutant strains at 10 h post-infection; *P*-values greater than 0.05 were deemed as not significant (ns). For assessing virulence *in vivo*, age- and sex-matched C57BL/6 mice were i.v. infected with 200 cfu of indicated strains of *S. Typhimurium*, and the bacterial load in the (D) liver and (E) spleen was determined at day 5 post-infection. Symbols represent data from individual animals, and the geometric mean ± SEM is shown. Data are pooled from two independent experiments. The unpaired *t*-test was used for statistical analyses.
Figure 5. The *S. Typhimurium* ΔspeD, ΔspeE and ΔluxS mutants show no auxotrophy in M9 minimal medium, grow normally in HeLa cells, and are fully virulent in mice.

(A) *S. Typhimurium* ΔspeD, ΔspeE and ΔluxS mutants grew in M9 minimal medium without additional supplementation. Geometric mean ± SD is shown. Dotted lines represent the detection limit for bacterial concentration. Data were pooled from two independent experiments.

(B) *S. Typhimurium* ΔspeD, ΔspeE and ΔluxS grew to similar levels as wild-type SL1344 over a 10-h period in HeLa cells, as described for Figure 3. Data are pooled from three independent experiments. Bars represent the mean fold-change and error bars show the SD. Unpaired *t*-tests were used to compare the intracellular load of wild-type and mutant strains at 10 h post-infection; *P*-values greater than 0.05 were deemed as not significant (ns). For assessing virulence *in vivo*, age- and sex-matched C57BL/6 mice were i.v. infected with 200 cfu of indicated strains of *S. Typhimurium*, and the bacterial load in the (C) liver and (D) spleen was determined at day 5 post-infection. Symbols represent data from individual animals, and the geometric mean ± SEM is shown. Data are pooled from two independent experiments.
The gene encoding SahH from *Legionella pneumophila* was isolated by PCR and ligated to pACYC184 to generate pACYC184::sahH. The SahH produced carried a FLAG tag at the amino-terminus. This plasmid was used to complement *S. Typhimurium* Δpfs mutant and complementation was compared with that achieved by pACYC184::pfs, i.e. the *pfs* from *S. Typhimurium*. (A) Western blot revealed the heterologous expression of SahH in *S. Typhimurium* Δpfs mutant complemented with the sahH gene (Δpfs + pACYC184::sahH) from *Legionella pneumophila*. Cell pellets were collected from 1ml of bacterial
The growth of S. Typhimurium Δpfs in M9 minimal medium was compared with that of the Δpfs mutant, the homologous complemented strain Δpfs+pACYC184∷sahH and the wild-type SL1344 in M9 medium. Growth by S. Typhimurium Δpfs in M9 medium was restored by the heterologous complementation and the Δpfs mutant expressing Legionella SahH grew to a similar level to the wild-type SL1344 (Figure 6B). Similarly in HeLa cells, the growth reduction seen in S. Typhimurium Δpfs (e.g. Figure 3D) was abolished by

Figure 6. The SahH from Legionella pneumophila reverts the M9 auxotrophy of the S. Typhimurium Δpfs mutant and growth in HeLa cells, and restores virulence in mice. Part 2 of 2

The growth of S. Typhimurium Δpfs+pACYC184∷sahH in M9 minimal medium was compared with that of the Δpfs mutant, the homologous complemented strain Δpfs+pACYC184∷pfs and wild-type SL1344 in M9 medium. Growth by S. Typhimurium Δpfs in M9 medium was restored by the heterologous complementation and the Δpfs mutant expressing Legionella SahH grew to a similar level to the wild-type SL1344 (Figure 6B). Similarly in HeLa cells, the growth reduction seen in S. Typhimurium Δpfs (e.g. Figure 3D) was abolished by

Figure 7. LC–MS/MS analysis of SAH levels show SahH complementation reduces SAH accumulation in the absence of Pfs.

S. Typhimurium strains were grown overnight in LB broth, washed then sub-cultured into M9 minimal medium for 24 h. The cells were pelleted, then snap frozen before analysis by mass spectrometry for levels of SAH, determined by ion count at the relevant mass. A SAH standard was used to validate the mass (data not shown). The statistical analysis was conducted using a one-way ANOVA with Bonferroni’s post-test.
complementation with pACYC184::sahH, which restored growth of the Pfs mutant in HeLa cells to wild-type levels (Figure 6C). Lastly, the capacity of S. Typhimurium Δpfs + pACYC184::sahH to grow in mice was examined. On day 5 post-infection, the bacterial burden in the spleen and liver showed that complementation with the sahH gene in S. Typhimurium Δpfs fully restored growth in the mice (Figure 6D,E). This result suggests that the accumulation of SAH was likely the cause of in vivo attenuation in the Δpfs mutant.

The levels of SAH in the different bacterial strains were analysed by mass spectrometry (Figure 7). Wild-type S. Typhimurium SL1344 contained undetectable levels of SAH, in contrast with S. Typhimurium Δpfs which had much higher levels of SAH, measured as ion count by LC–MS. There was no accumulation of SAH Δpfs + pACYC184::pfs, and the levels of SAH were reduced by 60% in the Δpfs strain complemented with pACYC184::sahH. To determine whether this accumulation of SAH was coincident with changes in DNA methylation, the level of DNA methylation in each of the mutants was examined using two methods. Firstly, genomic DNA was extracted from the strains and analysed using antibody specific for methylated DNA (Supplementary Table S1). In addition, DNA samples were subjected to digestion by nucleases that are either methylation dependent, or inhibited by methylation (Supplementary Figure S4). The DNA methylation patterns were not different in the Pfs mutant compared with SL1344, when tested by either method.

Lastly, in order to analyse the relatedness of the bacterial Pfs and eukaryotic SahH enzymes, a Similarity Network Analysis was undertaken using libraries of prokaryotic and eukaryotic protein sequences. The output of the analysis is shown in Figure 8. The analysis reveals that, amongst the Pfs enzymes (MTA/SAH nucleosidases), there are 3–4 subgroupings, but these sequences are inherently different from the SahH enzymes that include some bacteria (e.g. Legionella pneumophila) as well as Archaea and indeed human SahH. These quite different clusterings of sequences suggest separate evolution of the Pfs family of enzymes from the SahH family.

**Discussion**

It has been posited that Pfs, a multifunctional bacterial enzyme, might provide a target for novel antimicrobial development [22]. Pfs has multiple enzymatic roles which include, directly or indirectly, controlling the levels
of a key vitamin, biotin, the detoxification of SAH and the production of a substrate for production of A1-2, and Met and adenine recycling [9–15,22,23,26,27]. We examined the virulence of a S. Typhimurium deletion mutant of pfs, and demonstrated auxotrophy in M9 minimal medium, a lack of intracellular growth in HeLa cells, and a very high degree of growth attenuation in C57BL/6 mice, a highly stringent murine challenge model.

As the enzyme description suggests, the MTA/SAH nucleosidase encoded by pfs, which is present in many bacteria including E. coli and Salmonella, is multifunctional. Pfs catalyses the conversion of SAH to SRH, mediates the hydrolysis of 5-MTA to 5-MTR and adenine, and converts 5′-deoxyadenosine to 5′-deoxyribose and adenine [23,41,42]. Previous studies have shown that the deletion of Pfs causes a growth defect in E. coli and E. coli Δpfs was shown to phenocopy a biotin synthase (bioB gene product) mutant [26,27]. The minimal medium M9, which consists of glucose and salts, requires S. Typhimurium to generate all the necessary cellular substrates including proteins, carbohydrates and lipids for growth to occur. The S. Typhimurium Δpfs mutant was an auxotroph in M9 minimal medium and growth was restored by the provision of Met or biotin. Previous reports have shown that the E. coli Δpfs is growth defective in minimal medium and that the addition of Met or biotin partially restored the growth of the E. coli Δpfs [26]. Recent findings in other bacteria including Neisseria meningitides, Borrelia burgdorferi and Staphylococcus aureus suggested that Pfs is essential for their virulence [43–45], though the specific mechanism(s) through which attenuation occurs is less clear.

In S. Typhimurium Δpfs, the auxotrophic requirement for biotin can be explained by the action of Pfs on 5′-deoxyadenosine, which is both a substrate for Pfs, and a known inhibitor of the biotin synthesis by BioB [27,46]. Why M9 Met auxotrophy was demonstrated by S. Typhimurium Δpfs is less clear; subsequent studies revealed that the conversion of SRH to homocysteine, via LuxS, was dispensable suggesting that the activated methyl cycle per se, as it is currently modelled, is not essential for the provision of substrates for growth in either minimal medium or in mammalian cells. It may be that the accumulation of SAH is inhibitory for MetK and this inhibition is relieved by increased concentrations of Met. The role of Met in the growth attenuation in HeLa cells was explored through comparing the virulence of the S. Typhimurium Δpfs mutant with mutants lacking the major Met importer (MetNIQ), and in de novo synthesis through the loss of MetB or MetE/MetH [37], the partially redundant Met synthases. These combined Met mutants did not phenocopy the attenuation observed in the S. Typhimurium Δpfs mutant in HeLa cells, suggesting that the inability to produce or scavenge Met was not responsible for the observed in vivo growth attenuation. Similarly, since the BioB mutant did not phenocopy the S. Typhimurium Δpfs, and the mutants grew like wild-type in HeLa cells and C57BL/6 mice, the attenuation observed in S. Typhimurium Δpfs cannot be attributed to a reduction in, or loss of, biotin synthesis.

In addition to methylation, SAM provides a substrate for the synthesis of the polyamines spermidine and spermine [18,23,47–49]. Spermidine synthesis proceeds with the decarboxylation of SAM and is catalysed by SAM decarboxylase (SpeD). Decarboxylated SAM donates an aminopropyl group to putrescine to produce spermidine and MTA, and this reaction is catalysed by SpeE [23,49]. MTA is further converted to MTR by Pfs and this inhibition is relieved by increased concentrations of Met. The role of Met in the growth attenuation in HeLa cells was explored through comparing the virulence of the S. Typhimurium Δpfs mutant with mutants lacking the major Met importer (MetNIQ), and in de novo synthesis through the loss of MetB or MetE/MetH [37], the partially redundant Met synthases. These combined Met mutants did not phenocopy the attenuation observed in the S. Typhimurium Δpfs mutant in HeLa cells, suggesting that the inability to produce or scavenge Met was not responsible for the observed in vivo growth attenuation. Similarly, since the BioB mutant did not phenocopy the S. Typhimurium Δpfs, and the mutants grew like wild-type in HeLa cells and C57BL/6 mice, the attenuation observed in S. Typhimurium Δpfs cannot be attributed to a reduction in, or loss of, biotin synthesis.

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LuxS was not essential for the growth of S. Typhimurium in M9 minimal medium, nor for growth in HeLa cells, or in C57BL/6 mice. This observation supports previous studies where E. coli and Salmonella lacking LuxS did not show any growth defect in vitro [56–58]. LuxS plays a potential role in two different aspects of S. Typhimurium pathogenesis: in the recycling of Met, and in quorum sensing [12]. Many pathogenic bacteria synthesise LuxS and, in some bacteria, quorum sensing is involved in the control of virulence gene expression, e.g. Shigella flexneri, Vibrio cholerae, Clostridium perfringens and Streptococcus pyogenes [59–62]. LuxS is also required for meningococcal bacteraemia in mice [63]. In models of the activated methyl cycle in E. coli and Salmonella, LuxS is required to complete the cycle which can replenish Met from homocysteine. In our study,
Met recycling and Met replenishment through LuxS was not essential for S. Typhimurium virulence in vivo. Also, it was concluded that AI-2 mediated quorum sensing was dispensable for S. Typhimurium growth and virulence during systemic infection in highly sensitive mice. These data do not mean that AI-2-mediated quorum sensing is irrelevant to S. Typhimurium virulence and it is possible, through effects on community behaviours like swarming motility, that AI-2 provides S. Typhimurium a fitness advantage ex vivo, which increases transmission efficiency. Indeed, the Pfs mutant lost the swarming phenotype when plated onto soft agar (Supplementary Figure S2).

According to the pathway model (Figure 1), deletion of Pfs should lead to the accumulation of SAH and MTA. Elevated levels of SAH are inhibitory for SAM-dependent methyltransferases [38–40]; similarly, increased concentrations of MTA are inhibitory for polynucleotide synthases and SAM-dependent methyltransferases [64–67]. Having shown that SpeD, SpeE, LuxS and BioB are dispensable, the hypothesis that SAH accumulation is attenuating was tested through the heterologous expression of SahH, from Legionella pneumophila, a SAH hydrolase that lacks MTA nucleosidase activity.

Heterologous expression of L. pneumophila SahH (a specific S-adenosylhomocysteine hydrolase) by S. Typhimurium Δpfs restored the growth and virulence of this Δpfs mutant in M9 minimal medium, HeLa cells and the murine challenge model, and the complemented mutants reduced the levels of SAH that accumulated in the SL1344 Pfs mutant. The degree of reduction was different but the enzyme kinetics of the two enzymes (Salmonella Pfs and Legionella SahH) are unknown though in Met-replete M9 medium, there was growth attenuation over the first 18 hr in the Pfs mutant complemented with SahH, compared with Pfs complementation (Supplementary Figure S5). Further Clustal analysis revealed that the Legionella SahH and human SahH shared homology, suggesting that Legionella may have scavenged SahH from an eukaryotic e.g. the amoebic niche, a theory that has been discussed elsewhere [68,69]. Complementation of growth and virulence by Legionella SahH suggests that suppression of the SAM-dependent methyltransferases by SAH may be responsible for the attenuation observed in S. Typhimurium Δpfs. However, analysis of the DNA from S. Typhimurium Δpfs suggested that DNA methylation was unaffected at the gross level. This could mean that essential methyltransferases targeting other functions were affected by accumulated SAH. S. Typhimurium carries a number of proteins with potential SAM-binding motifs [70], such as PmtA, a phospholipid N-methyl transferase required for membrane synthesis, the expression of which may be repressed by higher levels of SAH.

In summary, these studies suggest that the multifunctional enzyme Pfs may be a legitimate antibiotic target. While the auxotrophy in minimal medium can be alleviated by biotin or Met, the relationship between the availability of these substrates, and the severe attenuation observed in murine infections, is yet to be resolved. Clustal analysis of the E. coli and Salmonella spp. Pfs revealed significant homology, which was absent from the L. pneumophila SahH. S. Typhimurium lacking Pfs shows severe growth attenuation in highly sensitive mice, and this growth inhibition appears to be due to the accumulation of the metabolite SAH. While the SahH enzyme from Legionella pneumophila shares significant homology with the human enzyme, Similarity Network Analysis revealed that many bacterial Pfs genes form a separate and distinct family of proteins, suggesting that it may be possible to generate inhibitors specific for bacterial enzymes, which leave mammalian enzymes unaffected and to effect selective toxicity.

Abbreviations
AI-2, autoinducer-2; cfu, colony-forming units; FCS, fetal calf serum; Met, methionine; MTA, S-adenosylhomocysteine; MTR, methylthioribose; S. Typhimurium, Salmonella enterica var Typhimurium; SAH, S-adenosylhomocysteine; SAH/MTA, S-adenosylhomocysteine/Methylthioadenosine; SAM, S-adenosylmethionine.

Author Contributions
A.U.H. perfomed experiments, generated mutants and helped write the paper, N.W. performed animal experiments and helped write the paper, J.J.W. constructed some key mutants and helped write the paper, H.J. N. designed and helped interpret key in vitro experiments, S.A.C. and D.M.H. provided key data and helped write the paper, I.D.H. generated key figures explaining the relationships between the various enzymes, M.R.D. performed bioinformatic analysis of key genes, M.J.M. helped design experiments and provided key insights into metabolic processes and helped write the paper, T.L. provided key insights into enzyme family relationships and helped write the paper, R.A.S. designed key experiments and helped write the paper.
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Competing Interests
The authors declare that there are no competing interests associated with the manuscript.

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