Two Coregulated Efflux Transporters Modulate Intracellular Heme and Protoporphyrin IX Availability in *Streptococcus agalactiae*

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Abstract

*Streptococcus agalactiae* is a major neonatal pathogen whose infectious route involves sepsisemia. This pathogen does not synthesize heme, but scavenges it from blood to activate a respiration metabolism, which increases bacterial cell density and is required for full virulence. Factors that regulate heme pools in *S. agalactiae* are unknown. Here we report that one main strategy of heme and protoporphyrin IX (PPIX) homeostasis in *S. agalactiae* is based on a regulated system of efflux using two newly characterized operons, gbs1753 gbs1752 (called *pefA pefB*), and gbs1402 gbs1401 gbs1400 (called *pefR pefC pefD*), where *pef* stands for ‘porphyrin-regulated efflux’. *In vitro* and *in vivo* data show that *PefR*, a MarR-superfamily protein, is a repressor of both operons. Heme or PPIX both alleviate *PefR*-mediated repression. We show that bacteria inactivated for both *Pef* efflux systems display accrued sensitivity to these porphyrins, and give evidence that they accumulate intracellularly. The Δ*pefR* mutant, in which both *pef* operons are up-regulated, is defective for heme-dependent respiration, and attenuated for virulence. This new *Pef* regulon controls intracellular heme and PPIX availability in *S. agalactiae*, and is needed for its capacity to undergo respiration metabolism, and to infect the host.

Introduction

Heme (iron protoporphyrin IX) is a redox-active molecule, and a cofactor for numerous cell functions used in oxygen sensing and signal transmission, metabolism, and metal homeostasis [1]–[3]. In addition to its varied activities as a cofactor, heme promotes toxic oxygen radical production [4]. The duality between heme as a multifunctional cofactor, and a potentially toxic molecule, suggests the need for strict limitation of its intracellular levels. Metal-free protoporphyrin IX (PPIX) is also found intracellularly, as a heme precursor in bacteria that synthesize heme, and as an intermediate during iron recovery from heme, as shown in *Escherichia coli* [5]. Thus, cells might have to deal with both heme and PPIX intracellular pools.

While most studied organisms synthesize heme to ensure activity of hemoproteins under the appropriate conditions, numerous bacteria lack heme biosynthesis genes, making heme-catalyzed processes fully dependent upon external heme supplies. For example, *Hemophilus influenzae*, *Bacteroides* sp., and several Firmicutes, including *Lactococcus lactis*, *Enterococcus faecalis*, *Streptococcus agalactiae*, and numerous *Lactobacillus* sp., require heme to activate a respiration metabolic pathway [6]–[18]. Some of these bacteria (e.g., *H. influenzae*, *E. faecalis*, *Bacteroides* sp., *Lactobacillus brevis* and *Lactobacillus plantarum*) also encode heme-dependent catalases, which rely on exogenous heme for both their stability and activity [19]. Thus, heme supplied by the environment can have a determining effect on the metabolic and enzymatic capacities in organisms lacking the heme biosynthetic pathway.

All bacteria, regardless of their heme biosynthesis capacities, need to manage their intracellular heme pools. Regulation may be exerted at the levels of biosynthesis, uptake, degradation, and possibly efflux, which is in some cases coordinated with the cell iron status [20]–[25]. The use of efflux as a means of heme homeostasis remains in question. The sole candidate, HtrAB, a heme-regulated transporter, was first reported as having a role in heme toxicity in *Staphylococcus aureus*; orthologs of this system were also described in *L. lactis* and *Bacillus anthracis* [12], [26]–[28]. HtrAB in *S. aureus*, *B. anthracis*, and likely several Gram-positive pathogens responds to an extracytoplasmic heme sensor, HssS (part of HssSR two-component system) to activate expression [27], [28].

*S. agalactiae* is an important human pathogen that does not synthesize its own heme. Nevertheless, infection by this bacterium involves compulsory passage through the bloodstream, causing sepsisemia and subsequent meningitis [29], [30]. We showed previously that although *S. agalactiae* generally grows by a fermentation metabolism, it can also use heme, present in blood, to activate the terminal cytochrome *bd* quinol oxidase for respiration metabolism.
Author Summary

The infectious route of numerous bacterial pathogens includes sepsis, where bacteria are exposed to hemo-rich blood. Heme (iron protoporphyrin IX) is generally considered a bacterial iron source. However, while some pathogens do not biosynthesize heme, they use environmental heme to activate key functions. For example, incorporation of heme by the major neonatal pathogen Streptococcus agalactiae activates a latent respiration chain. Respiration metabolism stimulates S. agalactiae growth and survival in blood, and is needed for virulence. While the importance of heme in S. agalactiae behavior is documented, how it manages its intracellular heme pools remains unknown. We discovered a novel regulon, called Pef for “porphyrin-regulated efflux”, that modulates S. agalactiae intracellular availability of heme and protoporphyrin IX. A single transcriptional regulator, PefR, represses two distinct efflux transport operons. Regulator-mediated repression is alleviated by heme or protoporphyrin IX. Importantly, over-expression of Pef efflux transporters led to intracellular heme insufficiency, and consequent respiration and virulence defects. Inversely, when Pef efflux transporters were inactivated, results indicated an increased intracellular accumulation of heme and protoporphyrin IX. These studies point to the important role of regulated efflux transport systems in bacterial pathogens for maintaining intracellular heme at levels sufficient to stimulate growth and promote infection.

Thus, beyond the potential use of heme uptake to acquire iron, the use of heme to activate the energetically favorable respiration metabolism confers a significant gain in cell density, and is required for full bacterial virulence in a septicemia model [17], [18]. 

Despite the importance of growth in heme-rich blood as an initial step of S. agalactiae infection, the functions involved in regulating intracellular heme levels are unknown. Homology of HrtAB and HsaSR exist in S. agalactiae, and might participate in modulating heme toxicity upon sensing of extracellular heme. However, the mechanism by which HrtAB modulates heme toxicity is unknown, and the question remains whether other systems are needed to regulate heme pools. Here we report the existence of a novel regulon comprising two efflux operons and a single repressor that regulates heme pools. Here we report the existence of a novel regulon comprising two efflux operons and a single repressor that regulates heme pools.

Results

Identification of gbs1753 gbs1752 as a putative heme- and PPIX-induced locus in S. agalactiae

Transcriptome studies, comparing S. agalactiae gene expression under respiration (i.e., aerobic growth in the presence of exogenous heme and menaquinone) versus aerobic fermentation (i.e., no addition) conditions, initially revealed that gbs1753 gbs1752 was 2.4 to 5.1 fold higher under respiration conditions (AD, PG, EC, and P. Gleser [Pasteur Institute]; Table S1). The gbs1753 ORF encodes an integral membrane protein of the drugH+ antiporter family, belonging to the major facilitator superfamily (MFS) [31]. The gbs1752 ORF encodes an unknown protein with 2 transmembrane domains. We chose this locus for study, as it was conserved among Gram-positive bacteria that lack a complete heme biosynthesis pathway, but encode cytochrome bd quinol oxidases, indicating a capacity for respiration metabolism (Fig. 1 and Table S2).

The +1 transcriptional start of gbs1753 gbs1752 was located by 5’-RACE PCR mapping at 134 nucleotides upstream of the gbs1753 start codon at a cytosine (Fig. 2A). The gbs1753 promoter region was fused to a lacZ reporter, referred to as P_{gbs1753-lacZ} and expressed on a low copy number plasmid (Fig. 2A) [32]. In the absence of added heme, P_{gbs1753-lacZ} displayed basal level expression in static and aerobic growth conditions (possibly reflecting trace amounts of heme in BHI medium). Cultures grown with added heme (from 0.1 μM to 10 μM) displayed up to 9-fold higher P_{gbs1753-lacZ} expression compared to controls without heme, even in non-respiration-permissive conditions (Fig. 2B), indicating that heme, and not the state of respiration, was the inducing factor for the gbs1753 promoter. Significant P_{gbs1753-lacZ} induction was observed at heme concentrations of 0.3 μM and above. Similarly, PPIX, gallium protoporphyrin (GaPPIX), and zinc mesoporphyrin (ZnMPIX) also induced P_{gbs1753-lacZ}, while free iron had no inducing effect. These initial results indicate that gbs1753 expression is induced by different porphyrin molecules, regardless of their metal core status.

Identification of Gbs1402 as a putative regulator of gbs1753 gbs1752 expression

A random mutagenesis approach was used to identify genetic factors that affect gbs1753 gbs1752 expression. A transposon-generated

![Figure 1](image-url)
mutant library was screened for clones in which expression of the
P_{gbs1753}-lacZ fusion was up-regulated, as detected by deep blue
colony color. Several mutations mapped in a single ORF, gbs1402.
The gbs1402 gene encodes a multiple antibiotic resistance MarR-
like regulator (for review see [33]), and is located just upstream
of gbs1401 gbs1400, encoding a putative ABC-type multidrug transport
complex (Fig. 1B). Sequence analysis predicted that the Gbs1401
and Gbs1400 proteins both contain a transmembrane domain and
an ATPase signature. These observations led us to formulate a
working hypothesis that both Gbs1401 and Gbs1400 proteins
are involved in PPIX and metalloporphyrin efflux, and are
regulated just upstream of pefAB genes, for porphyrin-regulated
efflux. gbs1753 gbs1752 are renamed pefA pefB; gbs1401 gbs1400 are renamed pefC pefD; and the
gbs1402 gene for the potential regulator is renamed pefR (Fig. 1).

**PefR binds specifically to pefAB and pefRCD promoter regions and represses their transcription**

Proteins of the MarR family characteristically bind to DNA
inverted repeat motifs (IR) [33]. Sequence analysis revealed the
presence of a near-perfect 18-nucleotide IR within a 23 bp
consensus, (5′-TAAAATAGTTCTCAGGTAAGT4′) present one upstream of pefA pefR, and twice (one identical sequence, and
one inexact copy with 3 nucleotide substitutions in italics) upstream of pefC pefD (Fig. 3A). The IR is not present
elsewhere on the S. agalactiae genome. This 23-nucleotide sequence
contains the −10 region (TAAAAT) of a putative promoter and
constituted a candidate target site for PefR binding to pefAB and
pefRCD promoter regions. Mobility shift assays were performed
using a purified PefR His-fusion protein, in combination with
DNA fragments comprising the IRs of pefAB or pefRCD promoter
regions, plus a fragment without an IR as negative internal control
(Fig. 3B). PefR caused a mobility shift of both the pefAB and
pefRCD fragments in a protein concentration-dependent manner.
No shift was observed with the control DNA fragment.

To determine whether PefR impacts on pefA pefB and pefR pefC
pefD expression in *vivo*, Northern blot experiments were performed
on the WT strain and an in-frame ΔpefR mutant, using pefAB and
pefRCD as probes (Fig. 3C). The deduced transcript sizes were
compatible with organization of the pefA and pefB genes as one
operon, and of pefR, pefC, and pefD genes as another. Compared to
the WT strain, expression of both pefAB and pefRCD operons
was strongly increased in the ΔpefR mutant. We also evaluated
expression of the P_{pefA-lacZ} transcriptional fusion in the WT
and ΔpefR mutant strains. β-galactosidase expression in the ΔpefR
background was 12 times higher than in the WT strain (Miller
Units were 62.9 ± 9.2 in ΔpefR versus 5.1 ± 1.7 in the WT strain).
The above *in vitro* and *in vivo* results indicate that PefR is a
transcriptional repressor of both the pefAB and the pefRCD
operons, by binding directly to the operator regions.

PefR is an intracellular heme and PPIX sensor and
regulator of pef expression

A characteristic of MarR family regulators is their binding to
effector molecules, which leads to induction or repression of their
target genes [34]–[36]. Transcriptional fusions showed that
the pefAB operon was induced in the presence of metalloporphyrins or
PPIX (Fig. 2B). As PefR binds the two pef promoter regions, and
appears to repress transcription, we hypothesized that PefR
binding and repression are modulated by heme and PPIX. To test
Figure 3. PefR is a repressor of pefAB and pefRCD loci. A. A conserved 23-nucleotide motif is present once upstream of pefAB, and twice upstream of pefRCD and is highlighted. The IR present in each of the motifs is marked by arrows. The −10 and −35 motifs, and the mRNA pefAB and putative pefRCD start sites are in blue. Start codons and RBS sequence are in italics. One motif upstream of pefRCD (dotted arrows) differs from the other motifs by 3 nucleotides, which are in gray italics. B. PefR binds the pefAB and pefRCD promoter regions in gel shift assays. Two pmoles of pefAB or pefRCD promoter fragment were incubated in the presence of 0, 4, 17, 34, 63 pmoles of PefR (lanes 1 to 5) and 4.6 pmoles control fragment, corresponding to an 116-bp fragment of the pefA gene. C. Northern blot analyses of pefAB and pefRCD mRNA in the WT and ΔpefR strains, using locus-specific probes (see Materials and Methods). As probe efficiencies and times of exposure differ for each target RNA, differences between pefAB and pefRCD levels are not comparable. ldhL mRNA (ctrl) was used to control for RNA quantity.

doi:10.1371/journal.ppat.1000860.g003

Expression of hrtAB as a function of heme and PPIX levels in S. agalactiae

Genomic studies of S. agalactiae revealed the existence of an analog of HrtAB, a heme-regulated transport system initially characterized in S. aureus and shown to be involved in heme toxicity [37]. Gbs0119 and Gbs0120 showed 45% and 29% identity with HrtB and HrtA of S. aureus respectively. We used Northern blot experiments (Fig. 5A) and a lacZ promoter fusion, PpefCD-lacZ (Fig. 5B), to assess the heme and PPIX concentrations needed to induce the hrtAB locus. Interestingly, hrtAB expression was low at heme concentrations below 1 μM, with strong induction at 10 μM; PPIX did not induce its expression, as was observed in S. aureus. Importantly, no hrtAB induction was observed at heme levels where pefAB was induced, i.e., between 0.1 and 0.5 μM heme (Fig. 2B). We conclude that the pef regulon is induced at heme concentrations below those needed for hrtAB induction, indicating that these functions are active under different conditions.

Role of pefAB and pefRCD operons in porphyrin sensitivity

The above results led us to ask whether pefAB and pefRCD loci are involved in porphyrin efflux. Several in vivo approaches were developed to explore this question. We constructed in-frame pefA (encoding a putative drug:H+/H2 efflux transporter) and pefB (encoding a membrane protein of unknown function) deletion mutants; as the two components of pefCD are predicted to encode a single ATP-dependent transporter, we generated a deletion removing both ORFs. Mutations were combined to inactivate both pef operons. Tests were also performed with the ΔpefR deletion mutant, in which expression of both pef loci were highly induced (Fig. 3C).

Sensitivity of ΔpefR, ΔpefA ΔpefCD, and ΔpefB ΔpefCD mutants to metalloporphyrins was evaluated by plate inhibition tests (Fig. S2). Both ΔpefA ΔpefCD and ΔpefB ΔpefCD mutants showed greater sensitivity than the WT strain to 1 nmole heme (Fig. S2A). Similar results were obtained using Gallium PPIX (GaPPIX; tested at 50 nmole, Fig. S2B). We noted that ΔpefA ΔpefCD was more sensitive than ΔpefB ΔpefCD; this phenotypic difference might suggest an accessory role of PeF in PeFp PeFb function. The single ΔpefA, ΔpefB, ΔpefC or ΔpefD mutants gave little or no inhibition by these porphyrins (data not shown), suggesting a functional redundancy between the efflux systems encoded by these loci. Sensitivity of the ΔpefIR mutant to heme and GaPPIX did not differ significantly from the WT. We further tested heme-mediated inhibition in liquid medium by growing WT and pef single and double mutants, and the ΔpefR mutant, in 0 or 1 μM heme (Fig. 6C).
and data not shown). The ΔpefA ΔpefCD and ΔpefB ΔpefCD mutants, but not the other tested strains, displayed a slight growth inhibition in the presence of heme. Addition of 2 μM or 4 μM heme exacerbates growth retardation of these mutants (data not shown). These results indicate the need for at least one of the Pef efflux systems to avoid heme toxicity.

To evaluate PPIX accumulation in S. agalactiae WT and mutant cells, we exploited its reactivity upon exposure to visible light; when excited by light, PPIX generates reactive oxygen species [38],[39]. The WT strain, and ΔpefAB ΔpefCD single and combined mutants, and the ΔpefR mutant, were grown with 10 μM PPIX and exposed to visible light for 0, 10 or 50 minutes. Viability of cells grown without PPIX and exposed to light was equivalent for all strains (Fig. 6B). Strikingly, inactivation of both putative pumps led to total mortality upon short light exposure (Fig. 6B). These results suggested that PPIX accumulates in S. agalactiae when pefAB and pefCD systems are inactivated. There were also marked differences between WT and ΔpefR sensitivity to PPIX after a longer (50 minute) light treatment. The ΔpefR mutant, which is up-regulated for pefAB and pefCD expression, showed essentially full viability, compared to a >10-fold drop for the WT strain. These data, showing heme and PPIX sensitivities of ΔpefAB ΔpefCD mutants, and lower PPIX sensitivity of the ΔpefR mutant give strong evidence for a role of the pef regulon in porphyrin efflux and intracellular homeostasis.

Intracellular porphyrin availability in S. agalactiae WT and ΔpefA ΔpefCD strains

We used two approaches to evaluate differences in intracellular porphyrin levels in the WT versus ΔpefA ΔpefCD strain. First, we exploited the fact that heme- and PPIX- induce P_{pefA-lacZ} (same as P_{gbs1753-lacZ} above), to compare induction levels in the WT and ΔpefA ΔpefCD double mutant strains (Fig. 7A). In response to heme and PPIX addition, β-galactosidase activity was respectively about 2-fold and 3-fold increased in the ΔpefA ΔpefCD mutant compared to WT, suggesting that more heme and PPIX are available to activate the pefA (gbs1753) promoter in this mutant.

Second, we used an in vivo “heme and PPIX sensor”: the E. faecalis catalase KatA is degraded if not bound to a porphyrin molecule [19]. Intracellular availability of heme or PPIX was evaluated by comparing the relative stability of the E. faecalis KatA expressed in the WT strain and ΔpefA ΔpefCD and ΔpefR mutants...
Figure 5. The \textit{hrtAB} locus is induced by higher heme concentrations than \textit{pefAB} or \textit{pefRCD}, and is not induced by PPIX. A. Northern blot analyses of \textit{gbs0119} expression in the presence of heme or PPIX in WT NEM316. Cultures were grown in the presence of 0, 1, 5 and 10 \textmu M heme or PPIX, and harvested for total RNA extraction in early stationary phase. \textit{idhL} (\textit{gbs0947}; ‘ctrl’) mRNA was used as RNA quantity control. The hybridization was performed on the same membrane as that used in Fig. 4B. B. Expression analysis of \textit{P}_{\text{pefA1}}/\text{lacZ} by determination of \beta-galactosidase activity (Miller Units). Measurements were performed three or more times. doi:10.1371/journal.ppat.1000860.g005

(Fig. 7B and data not shown). Cells were grown without porphyrins or in 1 \textmu M heme or PPIX, and KatA levels were followed in Western blots on cell lysates. Amounts of KatA in the WT and \Delta\textit{pefR} mutant strains were not significantly different (data not shown), which might reflect the limits of this reporter system. However, in the \Delta\textit{pefA} \Delta\textit{pefCD} mutant to which heme or PPIX was added, KatA showed pronounced stabilization, as expected for higher intracellular porphyrin levels in that strain. Results of both in vivo systems used above point to greater availability of porphyrins in the \Delta\textit{pefA} \Delta\textit{pefCD} mutant.

Physiological impact of \textit{pefAB} and \textit{pefRCD} activities on \textit{S. agalactiae} respiration and virulence

\textit{S. agalactiae} takes up exogenous heme, which activates its membrane cytochrome \textit{bd} quinol oxidase, and is needed for respiration metabolism; the shift to respiration increases cell density by at least 20% compared to aerobic fermentation growth [17]. We compared WT, \Delta\textit{pefR}, \Delta\textit{pefA} \Delta\textit{pefCD}, \Delta\textit{pefB} \Delta\textit{pefCD}, and the \Delta\textit{pefR} strain complemented by a plasmid-carried \textit{pefR} gene (\textit{pefRCD}) for their capacity to grow in respiration conditions (Fig 8A). The WT, \Delta\textit{pefA} \Delta\textit{pefCD}, \Delta\textit{pefB} \Delta\textit{pefCD}, and the complemented \Delta\textit{pefR} strain showed increased growth densities, indicative of respiration growth in these conditions. In contrast, the \Delta\textit{pefR} mutant was not augmented, suggesting a respiratory defect of this strain. The \textit{pefR}:\textit{IS51} insertional mutant gave the same results as the \Delta\textit{pefR} deletion strain (data not shown).

To confirm that the growth difference observed above was due to respiration metabolism, we performed the same growth experiments in the presence of 2-n-heptyl-4-hydroxyquinoline N-oxide (HQNO), which inhibits respiration but allows fermentation growth [data not shown; [17]]. All tested strains except \Delta\textit{pefR} displayed lower growth densities in HQNO, which were comparable to those in aerobic fermentation; HQNO had no effect on cell density of the \Delta\textit{pefR} mutant. These results confirm that all strains tested above (Fig. 8A), except \Delta\textit{pefR}, activated respiration growth.

Respiration growth provokes an increase in oxygen consumption [17]. This property was examined to confirm the respiration-defect of \Delta\textit{pefR}. Oxygen consumption in aerobic fermentation growth by the WT, \Delta\textit{pefR}, and \Delta\textit{pefR} [\textit{pefRCD}] strains was measured as 20.3, 21.0, and 31.1 \textmu M.min$^{-1}$ per OD$_{600}$=1 cells (Table 1), which in these conditions reflects cytoplasmic NADH oxidase activity [17]. In respiration-permissive conditions, oxygen consumption by the WT and \Delta\textit{pefR} [\textit{pefRCD}] strain was markedly higher (44.2 and 51.6 \textmu M.min$^{-1}$ respectively), but only moderately higher for the \Delta\textit{pefR} mutant (30.2 \textmu M.min$^{-1}$) (Table 1), further indicating the respiration-defect in the \Delta\textit{pefR} mutant. These results, together with the above studies, indicate that high PefAB and PefCD efflux activities impact on respiration by diminishing heme availability.

WT, \Delta\textit{pefAB} \Delta\textit{pefCD} and \Delta\textit{pefR} strains were tested in a mouse infection model (Fig. 8B). Virulence of the parental strain, and the \Delta\textit{pefA} \Delta\textit{pefCD} or \Delta\textit{pefB} \Delta\textit{pefCD} mutants was essentially the same. In contrast, the \Delta\textit{pefR} mutant (as well as the \textit{pefR}:\textit{IS51} insertional mutant; data not shown) showed markedly attenuated virulence. Earlier findings showed that a \textit{S. agalactiae} respiration-defective mutant is attenuated for virulence [17]. Reduced intracellular heme of \textit{pefR} leads to a respiration deficiency, which can explain virulence attenuation of this mutant.

Discussion

The present study uncovers a previously unknown system of porphyrin homeostasis in \textit{S. agalactiae}, comprising two operons encoding distinct efflux pumps, controlled by a single intracellular regulator. Our results indicate that PefR, like numerous MarR-type proteins, acts as an intracellular sensor to regulate efflux of its ligands [40],[33], in this case, porphyrins. The range of PefR regulation is confined to the \textit{pefAB} and \textit{pefRCD} operons, as deduced from the presence of an IR within a 23 bp sequence uniquely upstream of these operons. This configuration is typical of MarR binding sites [33],[41],[42]. Expression of the \textit{pefAB} and \textit{pefRCD} efflux loci limits \textit{S. agalactiae} intracellular porphyrin availability; at high \textit{pef} expression levels, \textit{S. agalactiae} may be unable to activate respiration metabolism, and to cause septicemic infection in the animal host. The management of intracellular porphyrin levels by efflux pumps, and regulation by an intracellular sensor in \textit{S. agalactiae} constitutes new information on heme and PPIX homeostasis strategies.
complete pef regulon might conceivably arise from a genetic transfer event. However, this possibility is unlikely, as the identity between their pef ORFs (73% to 49%) is not higher than the 72% identity between these species at the genome level (Average Nucleotide Identity; calculated as in [46]). Conservation of the pef regulon may indicate a role in survival and/or infection in the particular biotopes of both these species.

Results of this study add heme and PPIX to the inventory of MarR interactants. Several MarR proteins were previously shown to bind to, and regulate efflux of lipophilic and planar molecules, such as antibiotics, aromatic aldehydes or fatty acids, usually as a means to limit cellular toxicity [47]–[51]. PefR activity would expectedly be tuned to retain sufficient intracellular heme for functions such as respiration. Indeed, pefAB expression in cells grown with porphyrins is induced to only half the levels compared to a fully induced ΔpefR mutant (data not shown). This, and the fact that the ΔpefR mutant is respiration defective, supports the need for basal levels of intracellular heme. Moreover, the presence of two putative PefR binding sites upstream of pefRCD (versus one upstream of pefAB) might influence the differential regulation of the two loci; possibly, lower levels of pefCD expression could participate in limiting intracellular heme depletion.

PefAB and PefCD add to a handful of other systems described as being involved in porphyrin efflux. The PefAB and PefCD systems share common features with two efflux systems in eukaryotes that control porphyrin pools: Feline leukemia virus receptor C, FLVRC, is an MFS-family protein (like PefA) that exports cytoplasmic metalloporphyrins across an ion-proton gradient [52], [53]. The breast cancer resistance protein Bcrp/ABCG2 in humans is an ABC family transporter (like PefCD) that regulates eukaryotic intracellular heme levels under hypoxic conditions [54], [55]. Just two possible porphyrin efflux systems were described in bacteria, neither of which is an analog of PefAB or PefCD: In E. coli, TolC effluxes PPIX, possibly to facilitate its turnover after iron extraction from heme; neither its regulation, nor its capacity to efflux heme have been reported [5], [56]. In S. aureus, the hrtAB locus is regulated by heme, but not PPIX, and its inactivation provokes accrued heme sensitivity [28], [27]. The S. agalactiae hrtAB homologous system (gbs0120 gbs0119) is likely to have a similar role in limiting heme toxicity. In contrast, the pef regulon is activated by PPIX, and at heme concentrations that do not induce hrtAB.

As PefAB efflux relies on proton motive force (PMF) to export substrates, we speculate that activity would be stimulated by the greater PMF generated under respiration conditions, as shown in L. lactis [57], [17], [58]. Indeed, the PefAB system is conserved in streptococci and lactobacilli having the capacity to activate a respiration metabolism. As PefCD requires ATPase activity, its activity likely varies according to growth conditions [59], [60]. The need for both PefAB and PefCD activities in S. agalactiae may accommodate the different metabolic states encountered in vivo, e.g., anaerobic fermentation in abscess, aerobic fermentation in lungs, and respiration in sepsis.

A model for heme and PPIX homeostasis in S. agalactiae, based on the newly characterized Pef operons, is proposed (Fig. 9): PefR represses expression of pefAB and pefRCD operons. Heme and PPIX are assimilated by unknown mechanisms. Once intracellular, these molecules interact with PefR, which detaches from its binding sites to activate PefAB and PefCD efflux pumps. Activities of the two Pef systems may assure rapid adjustment of intracellular...
Figure 7. PPIX intracellular accumulation in vivo. 

A. Expression analysis of PpefA-lacZ by β-galactosidase activity determinations was performed in early stationary phase cells of WT and ΔpefA ΔpefCD strains. S. agalactiae and derivatives were grown in BHI liquid medium. Results represent the mean ± standard deviation from triplicate experiments. Asterisks denote statistically significant differences as determined by Student’s t-test (p<0.05).

B. PPIX- or heme-dependent production of the E. faecalis catalase KatA in S. agalactiae WT, and ΔpefA ΔpefCD mutant strains. Cells were grown in M17G supplemented or not with 1 μM of PPIX or heme. The loading of equivalent amounts of protein was verified by Coomassie stained gels performed in parallel. KatA was detected in total S. agalactiae protein extracts by immunoblot assays. Results shown are representative of 3 experiments.

doi:10.1371/journal.ppat.1000860.g007
heme levels under different cell physiological conditions. Whether PPX efflux by \textit{pefAB pefCD} has a biological role in \textit{S. agalactiae} as an enzymatic byproduct of iron capture as in \textit{E. coli} [5] remains to be determined. At higher heme concentrations, HrtAB homologs Gbs0120 and Gbs0119 could be activated \textit{via} the putative two-component system Gbs0122 (heme sensor) and Gbs0121 (regulator), as was demonstrated for HsRS in \textit{S. aureus} [27]. This system would have a dominant role in protecting cells in a heme-rich environment, e.g., in conditions of massive degradation of host red blood cells; such a role is consistent with the strong induction of \textit{hrtAB} (>200-fold) when heme levels are high (Fig. 5). Inactivation of \textit{pefAB} and \textit{pefCD} loci results in increased intracellular heme and PPX pools, with consequences on growth and on heme-dependent enzyme activities; constitutive activity of these operons by \textit{pefR} inactivation depletes these intracellular pools, resulting in respiration and virulence defects.

A role for respiration functions in virulence has been reported for diverse bacterial pathogens, including \textit{Brucella abortus}, \textit{Shigella flexneri}, \textit{S. aureus} and \textit{Mycobacterium tuberculosis}; unlike \textit{S. agalactiae}, these bacteria biosynthesize heme and are autonomous for respiration [61]–[64]. In \textit{S. agalactiae}, respiration is part of the metabolic reprogramming that occurs during the requisite septicemic phase of infection, in heme-rich blood [17]. We suggest that a correlation between unavailability of heme due to high \textit{pefAB} and \textit{pefCD} activity, and the respiration defect explains virulence attenuation in the \textit{ΔpefR} mutant. The use of efflux to regulate metabolic activity of a major bacterial pathogen has not been previously reported, and will lead to a better understanding of how bacteria deal with heme and other porphyrins in the variable conditions they encounter during infection.

### Materials and Methods

**Ethics statement**

All animal experiments were performed in strict accordance with INRA institutional guidelines of good animal practice (Jouy-en-Josas, France), and approved by the Direction des Services Vétérinaires (accreditation number 78-63).

***Bacterial strains, growth conditions and plasmids***

Strains and plasmids used in this study are described in Tables 2 and 3. NEM316, a \textit{S. agalactiae} capsular serotype III strain was used as wild type (referred to as WT) and was isolated from a fatal case of septicemia [65]. \textit{S. agalactiae} and its derivatives were cultivated at 37°C in M17 medium (Oxoid) supplemented with glucose (0.2% for pre-cultures and 1% for test cultures [referred to as M17G]). For β-galactosidase assays, \textit{S. agalactiae} was grown in BHI (brain heart infusion, Difco Laboratories) supplemented with 0.8% glucose. Aerated cultures were grown under agitation at

| Table 1. Measurement of oxygen consumption according to metabolism in WT and \textit{ΔpefR} strains. |
|---------------------------------|
| **Strains** | **Aerobic fermentation (μM.min⁻¹)** | **Respiration** |
| WT | 20.3±1.6 | 44.2±5.7 |
| \textit{ΔpefR} | 21.0±6.6 | 30.2±4.8 |
| \textit{ΔpefR} + \textit{ppefR} | 31.1±3.2 | 51.6±0.3 |

\textit{S. agalactiae} WT, \textit{ΔpefR}, and \textit{ΔpefR} complemented by a plasmid carrying a wild-type copy of \textit{pefR} (Δ\textit{pefR} + \textit{ppefR}) were grown to late exponential phase. Oxygen consumption was measured on whole cells with a Clark-type oxygen electrode. Results represent the means ± SD of two (for \textit{ΔpefR} + \textit{ppefR}) or three experiments.

doi:10.1371/journal.ppat.1000860.t001
200 rpm in a ratio of air space to liquid of approximately 5/1. Respiring cultures were grown under agitation in the presence of 1 μM heme (from a stock solution of 10 mM hemin chloride, Fluka) and 10 μM vitamin K2 (a menaquinone; Sigma). *E. coli* was cultivated in LB medium (Luria-Bertani, Difco Laboratories) at 37°C with aeration by shaking at 180 rpm. Antibiotics were used as needed at the following concentrations: for *S. agalactiae*, 500 μg/ml erythromycin, 1000 μg/ml kanamycin, 4 μg/ml chloramphenicol, 5 μg/ml tetracycline; for *E. coli*, 100 μg/ml erythromycin, 20 μg/ml chloramphenicol.

**RNA extraction, Northern blot and RACE PCR**

RNA was extracted from *S. agalactiae* cultures as described [66]. Cells from 20 ml culture with an OD_{600} of 0.5~0.8 (exponential growth phase) or OD_{600}~1.5 were harvested by centrifugation at 6000×g during 10 min. Total RNA was extracted with a guanidine isothiocyanate and phenol-chloroform step [67], using the TRIZOL Reagent (Invitrogen).

For Northern blot analysis, RNA samples (30 μg) were mixed with an equal volume of glyoxal load dye (Ambion) and were electrophoretically separated on a 0.9% agarose glyoxal gel [68]. RNA samples were transferred to a Biodyne B membrane (Pall) according to manufacturer’s instructions. Hybridization and detection were performed using ECL direct nucleic acid labeling and detection systems (Amersham). A ~500 base pair (bp) DNA labeled fragment of each ORF tested was used as probe. Primers used for probe generation are shown in Table S3.

The pefa transcriptional start was mapped using a 5′/3′ rapid amplification cDNA ends (RACE) kit, second generation (Roche Applied Science) according to supplier’s instructions. Mapping was realized with RNA extracted from respiring cells, using primers:

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**Figure 9. Model for pefa and pefCD functions regulated by an intracellular sensor in *S. agalactiae*.** Extracellular heme and PPIX may be internalized by as yet unknown transporters. By homology with a recently described system in *S. aureus* [27], heme, but not PPIX, may bind to a putative external two-component receptor protein Gbs0122, the homolog of *S. aureus* HssS, resulting in gbs0119 gbs0120 (gmkAB) induction. Once inside the cell, free porphyrin molecules may encounter different binding proteins, including PefR. Apo-PefR binds to pefa and pefCD promoter regions to repress their expression. PefR-porphyrin binding releases PefR from DNA, to enable pefa and pefCD expression. The PefA and PefCD loci mediate efflux of free heme and PPIX to avoid toxicity. Overexpression of PefA and PefCD leads to heme depletion and respiration and virulence defects. Phenotypes of ΔpefR versus Δpefa ΔpefCD mutants are shown in the Table below.

doi:10.1371/journal.ppat.1000860.g009
5’TAGATGTAGGTGCTAACGTCG3’ and 5’CTTGTGTAGGCGCTTACGTCG3’.

β-galactosidase assays

Plasmid pTCV-lac is a low copy number plasmid used to evaluate promoter activities in *S. agalactiae* [32]. A DNA fragment containing *gbs1753* promoter was PCR-amplified with primers 5’GCGTAGAATTCAATATGGAG3’, containing an *EcoRI* site (underlined) and 5’TATCTGCGGATCCATTTCTGAT3’, containing a *BamHI* site (underlined). After digestion of the amplified fragment by *EcoRI* and *BamHI*, the *gbs1753* promoter was cloned into plasmid pTCV-lac, resulting in plasmid pTCV-J11 carrying the *Pgbs1753-lacZ* (or *PpefA-lacZ*) fusion. The plasmids pTCV-lac and pTCV-J11 were subsequently transformed into *S. agalactiae* by electroporation and recombinant clones were obtained by erythromycin selection. The same strategy was performed to generate a *PpefA-lacZ* fusion using the primers 5’ATAGCGCCAAATTTCTGGAG3’ and 5’CTTGGATTCTTTGCAGAC3’, giving rise to plasmid pTCV-J21.

Mutagenesis and screening conditions

Random insertional mutagenesis was performed on *S. agalactiae* with pG+host8::ISS1 basically as described [69]. Both *PpefA-lacZ* (reporter for *gbs1753* transcription) and pG+host8::ISS1 plasmids were established in *S. agalactiae*. The strain was grown at 30°C with kanamycin for 2.5 h. The temperature was then shifted to 37°C for 2.5 h and plated on M17G with erythromycin, kanamycin and 80 μg/ml X-gal (5-bromo-4-chloro-3-indolyl-beta-D-galactopyranoside, BioMerieux), to obtain ~100 CFU per plate. After 24 h incubation at 37°C, deep blue colonies were selected. Stable ISS1 mutants were isolated by growth at 30°C without erythromycin as described [70]. The flanking chromosomal DNA was sequenced using primer 5’AGGGCATGAAAAGATTCGAG3’. From the 5000 mutants we screened, 7 were in the pefR locus. Stable ISS1 mutants were then retested for *gbs1753* up-regulation.

**Table 2. Strains used in this study.**

| Strain name | Main characteristics | Reference |
|-------------|----------------------|-----------|
| E. coli TOP10 | F- mcrA Δ(mmr-hsdRMS-mcrBC) ΔBloxZ ΔDlacX74 recA1 araD139 Δ(ara-leu)7697 galU galK rpsL (StrR) endA1 supE44 | Invitrogen |
| GBS1753 | supE hsdR5 thdR5 thi Δ(lac-proAB)F’(traD36 lacI5 prophage07 lacZΔM15) repA9 | [71] |
| BL21 Star (DE3) | F- ompT hsdS2(rB maker) gal dcm (DE3) | Invitrogen |
| *S. agalactiae* NEM316 | Serotype III isolated from neonatal blood culture | [65] |
| NEM11 | NEM316 ΔpefA In-frame deletion of amino acid (aa) positions 36–433 of PefA. This study |
| NEM12 | NEM316 ΔpefB In-frame deletion of aa positions 26–211 of PefB. This study |
| NEM13 | NEM316 ΔpefC ΔpefD. Double deletion of aa positions 133–605 of PefC and positions 1–419 of PefD. This study |
| NEM14 | NEM316 ΔpefR In-frame deletion of aa positions 24–145 of PefR. This study |
| NEM15 | NEM316 ΔpefA ΔpefCD This study |
| NEM008 | NEM316 gbs1402::ISS1. ISS1 insertion between codon positions 81 and 82 aa of the gbs1402 ORF. This study |

**Table 3. Plasmids used in this study.**

| Plasmid name | Main characteristics | Reference |
|--------------|----------------------|-----------|
| pTCV-lac | Conjugative E. coli-Gram-positive bacteria shuttle plasmid with β-galactosidase reporter construct. EmR KmR. | [32] |
| pTCV-J11 | *PpefA-lacZ* or *PpefR-lacZ* gbs1753 (pefA) promoter region cloned into pTCV-lac. EmR KmR. This study |
| pTCV-J21 | *PpefA-lacZ* gbs0119 promoter region cloned into pTCV-lac. EmR KmR. This study |
| pG+host8::ISS1 | Vector for insertional mutagenesis. TetR. | [69] |
| pG+host5 | Temperature sensitive vector used in Gram-positive bacteria. EmR. | [70] |
| pRC2 | Cloning vector for complementation. CmR. | [72] |
| pRC2-J14 | pRC2 plasmid containing pefR gene, referred as ppefR. CmR. This study |
| pET100D-TOPO | Cloning vector. AmpR | Invitrogen |
| pET100-J14 | Expression N-terminal His tagged PefR. AmpR. This study |
| pLUMB5 | *PpefA-lacZ* KatA-His6. Expression E. faecalis KatA. Kanamycin resistance promoter. CmR | [19] |

doi:10.1371/journal.ppat.1000860.t002
centrifugation at 6000 rpm for 10 min and resuspended in 10 ml binding buffer (Tris HCl 50 mM pH 8, NaCl 300 mM, Imidazole 50 mM). Cells were broken by shaking with glass beads in a Fast Prep apparatus (MP Biomedicals). The lysate was centrifuged at 14000 rpm for 15 min at 4°C. The recombinant His-tagged fusion protein was purified by nickel affinity chromatography using His-Select affinity (Sigma) equilibrated with binding buffer according to supplier’s instructions. The elution was carried out with binding buffer containing 300 mM Imidazole. All fractions were collected and analyzed by SDS-PAGE. Purified proteins were dialyzed against 50 mM Tris HCl pH 8, 80 mM NaCl and 6.25% glycerol. Protein concentrations were determined by the Lowry assay (Bio-Rad).

Immunoblotting analysis

Strains harboring P_apla-o-katA-His6 plasmid were incubated overnight under static conditions with or without 1 μM heme or PPIX. Cell growth was harvested by centrifugation at 6000×g during 10 min. Cells were broken by shaking with glass beads in a Fast Prep apparatus (MP Biomedicals). Immunoblotting analysis was done with rabbit anti-KatA antiserum as described, and kindly provided by Dr. L. Hederstedt [19]. Detection was performed using ECL Plus Western blotting detection reagents (GE Healthcare).

Electrophoretic mobility shift assay (EMSA)

A 281 bp DNA fragment containing the gbs1753 promoter region and a 248 bp DNA fragment containing the gbs1402 promoter region were generated by PCR from genomic DNA of S. agalactiae and purified using the PureLink PCR Purification Kit (Invitrogen). Binding assays with PefR were carried out using a binding buffer (5% glycerol, 20 mM Tris HCl, pH 8, 200 mM MgCl2, 0.2 mM MgG6P, 1 mM EDTA, 0.2 mM DTT, 0.3 mM BSA) in a final volume of 15 μl at 37°C. Reaction mixtures were incubated at 37°C for 20 min, after which they were analyzed by gel electrophoresis on an 8% polyacrylamide gel in TBE 1× buffer, stained with ethidium bromide. Reaction mixtures were incubated at 37°C for 20 min, after which they were analyzed by gel electrophoresis on an 8% polyacrylamide gel in TBE 1× buffer, stained with ethidium bromide. Reactions were carried out in the presence of a 116 bp non-specific competitor DNA (internal coding gbs1753 sequence) where specified. Where tested, heme or PPIX were added 20 min after protein and DNA components. Gel shift experiments were performed with two independent batches of purified PefR.

Construction of Δgbs1753 (ΔpefA), Δgbs1752 (ΔpefB), Δgbs1401 Δgbs1400 (ΔpefCD) and Δgbs1402 (ΔpefR)

mutants and pefR plasmid construction

In-frame pefA, pefB, pefCD, and pefR deletion mutants were constructed using the strategy described below. Briefly, the two regions flanking the locus to be deleted were independently amplified by PCR (see Table S3 for primers). Amplified fragments were digested by HindIII or EcoRV enzyme and ligated to each other. The resulting fragments were amplified by PCR, digested by EcoRV enzyme and cloned into the thermosensitive shuttle plasmid pG+host5. Electroporation of S. agalactiae strains and allelic exchange were done as described [70]. In-frame deletions were confirmed by PCR and sequence analysis. The pefR gene was amplified (Table S3 for primers), and cloned using EcoRI and BamHI on fragment ends, onto plasmid pRC2 cut with the same enzymes; the resulting plasmid, pRC2::J14, is referred to as ppefR in Results.

PPIX and metalloporphyrin sensitivity tests

Stock solutions (10 mM) of PPIX, GaPPIX, ZnPPIX, and ZnMPPIX (Frontier Scientific) were prepared in DMSO. PPIX sensitivity of S. agalactiae NEM316 and its derivatives was examined on early stationary growing cells (OD600 = 1.6). For PPIX light sensitivity tests, serial dilutions were spotted on plates and exposed to visible light for 0, 10 or 50 minutes. A 500 W halogen lamp with a glass lid placed at a distance of 55 cm was used as light source. Metalloporphin sensitivity tests on plates were performed as described [12]. Briefly, 1:100 dilutions of stationary-phase cultures were prepared in M17G soft (0.7%) agar, and poured over M17G plates. Metalloporphin (heme, 5 μl of a 0.2 mM solution; GaPPIX, 5 μl of a 10 mM solution) was spotted onto the plates. All plates were incubated for 20 h at 37°C and then photographed.

Oxygen consumption measurements

Oxygen consumption was determined as described [17]. Briefly, S. agalactiae WT and ΔpefR late exponential phase cultures, prepared in aerobic fermentation or respiration conditions, were washed twice with PBS at 4°C and resuspended in the same buffer to obtain a 1 ml bacterial suspension at OD600 = 1.0. Oxygen consumption was followed with a Clark-type oxygen electrode (Liquif-Phase Oxygen electrode unit DW1, Hansatech instruments). The maximum oxygen consumption rate was measured following the addition of 10 mM glucose.

Mouse virulence assay

Pathogen-free 6 week-old Swiss CD1 mice (Charles River laboratories) were infected intravenously with S. agalactiae NEM316 or derivatives. Groups of 7–8 mice were anesthetized with ketamine (100 μg/g, Merial) and xylazine (12 μg/g, Bayer) and were inoculated in the eye vein with 107 CFU. Mortality was observed over a 10-day period. Animal experiments were performed in triplicate.

Bioinformatics

To find the conserved sequence between pef promoter regions, sequence were aligned with ClustalW and manually refined. A search for the 23 bp consensus in S. agalactiae NEM316 genome was done by visual inspection, and using BlastN program with EXPECT threshold set at 100. Pef protein homologs were identified using Genome Region Comparison, http://cmr.jcvi.org/cgi-bin/CMR/CmrHomePage.cgi, and/or were retrieved from genomic Blast databases, http://blast.ncbi.nlm.nih.gov/Blast.cgi/; using BlastP program with an E-value threshold set at 0.01.

Supporting Information

Figure S1 A. PPIX displaces PeR from its DNA target. Gel mobility shift analysis of the effect of PPIX on PeR binding to the pefAB (right) and pefRCD (left) promoter regions. Samples contained 2 pmoles of P_pefAB or P_pefRCD fragment and 30 pmoles of PeR, to which were added either sample buffer, 120 pmoles (left) or 240 pmoles (right) of PPIX. On the two top panels, lanes 1–2 were juxtaposed to lane 3–4, which were separated on the initial same gel. B. Heme, but not iron, displaces PefR from its DNA target. Samples contained 2 pmoles of P_pefRCD fragment and 32 pmoles of PeR, to which were added either sample buffer, 160 pmoles heme, or 160 pmoles iron. Lanes 1–2 were juxtaposed to lane 3–4, which were separated on the initial same gel.

Found at: doi:10.1371/journal.ppat.1000860.s001 (1.51 MB EPS)

Figure S2 Differential heme (FePPIX) and GaPPIX sensitivity of the pef mutants. Stationary phase cultures of WT, ΔpefR, ΔpefA

Intracellular Heme Control by an Efflux Regulon
ΔpfeFCD or ΔpfeBΔpfeFCD strains were spread on M17G plates and 1 n mole of heme (Δ) or 50 n moles of GaPPIX (B) were pipetted directly onto plates. Plates were incubated 24 hours, and then photographed. Representative results of at least 3 experiments are shown. 

Found at: doi:10.1371/journal.ppat.1000860.s001 (1.40 MB EPS)

**Table S1** Transcriptome analysis in respiration versus aerobic fermentation conditions. Cells were grown in M17 medium with 1% glucose, supplemented or not with a mixture of 10 K2 (a menaquinone) and 48 mK vitamin K2 (a menaquinone) and 48 µg/ml hemoglobin as heme source. Cells were harvested at OD600 = 0.3 for RNA extraction. Total RNA was extracted and analyzed by hybridization on NEM316 derived whole genome DNA macroarray as described [66]. Fold change in expression is the mean of two independent macroarray experiments. Shown are results of *S. agalactiae* genes whose expression was induced or repressed at least 2-fold in respiration compared to aeration conditions. Genes further studied in this work are in bold. Gene assignments are according to the Sagalist web site (http://genolist.pasteur.fr/SagaList/) and BLAST searches (http://blast.ncbi.nlm.nih.gov/Blast.cgi/). 

Found at: doi:10.1371/journal.ppat.1000860.s002 (1.40 MB EPS)

**Table S2** Distribution of pfe regulon and cytochrome genes among Lactobacillales. * Reorganization of gene or domain order; # frameshift mutation in pfeA homologous gene. In bold, species containing the complete pfe regulon; ** Imperfect repeats, character-

istic of MarR DNA binding sites, are present in respective pfeAB and pfeB/CD promoter regions of this strain; *** gbs1401 and gbs1400 ORFs are missing in *S. agalactiae* 2603V/R [43].

Found at: doi:10.1371/journal.ppat.1000860.s004 (0.06 MB DOC)

**Table S3** Primers used in this study.

Found at: doi:10.1371/journal.ppat.1000860.s005 (0.05 MB DOC)

**Acknowledgments**

We are grateful to IERP for technical assistance in animal studies. We thank Lars Hederstedt (Lund University, Sweden) for the generous gift of plasmid and anti-KatA antibody. We are grateful to Marie-Agnès Petit for application of ANI program and for critical and thoughtful reading of the manuscript, Gilles Lambertet for precious help in oxygen uptake determinations and Olivier Son and Isabelle Poquet for advice in gel shift assays (Micalis). We thank Cécile Wandsman, Patrick Trieu-Cuoit, and Philippe Glaser (Institut Pasteur), Claire Poyart (Hôpital Cochin), and Philippe Bouloc (Institut de Généétique et Microbiologie) for stimulating discussions, and the members of the Micalis ‘Bâtiment 222’ lab for their support in the course of this study.

**Author Contributions**

Conceived and designed the experiments: AF DL PG AG. Performed the experiments: AF DL ADB EC. Analyzed the data: AF DL PG AG. Wrote the paper: AF AG.

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