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## ***Bacillus anthracis* HssRS signaling to HrtAB regulates heme resistance during infection**

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### **Abstract**

*Bacillus anthracis* proliferates to high levels within vertebrate tissues during the pathogenesis of anthrax. This growth is facilitated by the acquisition of nutrient iron from host heme. However, heme acquisition can lead to the accumulation of toxic amounts of heme within *B. anthracis*. Here, we show that *B. anthracis* resists heme toxicity by sensing heme through the HssRS two-component system, which regulates expression of the heme-detoxifying transporter HrtAB. In addition, we demonstrate that *B. anthracis* exhibits elevated HssRS function compared to its evolutionary relative *S. aureus*. Elevated heme sensing is likely required by *B. anthracis* due to the significant heme sensitivity exhibited by members of the genus *Bacilli*. We also demonstrate that *B. anthracis* depends on conserved residues within the previously uncharacterized sensing domain of the histidine kinase HssS for HssS function. Finally, we show that the heme- and HssRS-regulated *hrtAB* promoter is activated in a murine model of anthrax. These results demonstrate the evolutionary conservation of heme sensing among multiple Gram-positive bacteria and begin to provide a mechanistic explanation for the heme resistance of *B. anthracis*. Further, these data suggest that heme stress is experienced by bacterial pathogens during infection.

### **Keywords**

Heme; two-component system; Staphylococcus; Bacillus; anthrax

## **INTRODUCTION**

*Bacillus anthracis* is capable of causing devastating disease upon breaching host epithelial barriers (Dixon *et al.*, 1999; Mock and Fouet, 2001). A spore-forming Gram-positive pathogen that primarily infects livestock, *B. anthracis* can cause a rare zoonotic disease in humans known as anthrax (Dixon *et al.*, 1999; Mock and Fouet, 2001). Anthrax arises when spores germinate within host tissues and can progress to systemic bacteremia followed by sudden death (Dixon *et al.*, 1999; Mock and Fouet, 2001). Like all bacterial pathogens, *B. anthracis* must circumvent mechanisms of innate immunity to replicate within host tissues and cause disease.

One mechanism of innate immunity that represents a challenge to bacterial pathogens is nutritional immunity, which refers to the host-mediated sequestration of nutrients such as metal ions required for bacterial growth (Bezkorovainy, 1981; Corbin *et al.*, 2008). In particular, free iron is maintained at a concentration within host tissues that is too low to support the growth of most bacteria, including *B. anthracis* (Bullen, 1999; Reniere *et al.*,

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2007). However, iron in the form of the porphyrin heme is an abundant molecule within host erythrocytes as the cofactor of hemoglobin and represents a source of nutrient iron for invading bacteria (Reniere *et al.*, 2007). *B. anthracis* and related pathogens express a set of proteins known collectively as the iron-regulated surface determinant (Isd) system devoted to the acquisition and degradation of host heme for nutrient iron needs (Gat *et al.*, 2008; Maresso *et al.*, 2006; Maresso *et al.*, 2008; Mazmanian *et al.*, 2003; Skaar and Schneewind, 2004). The importance of heme acquisition to pathogenesis is highlighted by the decreased virulence observed for pathogens inactivated for members of the Isd system (Reniere and Skaar, 2008; Skaar *et al.*, 2004; Torres *et al.*, 2006).

Although heme is a valuable source of iron for *B. anthracis* during infection, heme and hemin (the free, Fe(III) form of heme found in solution) can catalyze the formation of reactive oxygen species, leading to protein and lipid damage that cause bacterial cell death (Everse and Hsia, 1997; Reniere *et al.*, 2007; Torres *et al.*, 2007). *S. aureus*, an evolutionary relative of *B. anthracis*, is susceptible to hemin toxicity; however, staphylococci are able to adapt to growth in the presence of hemin (Torres *et al.*, 2007). In *S. aureus*, adaptation to hemin toxicity depends on expression of the heme-regulated ABC transporter (HrtAB), which is up-regulated upon exposure of staphylococci to hemin and alleviates hemin toxicity (Friedman *et al.*, 2006; Torres *et al.*, 2007). It has been hypothesized that HrtAB may directly export excess heme from the bacterial cytoplasm or may detoxify heme stress by exporting a toxic metabolite that accumulates in cells exposed to excess heme (Stauff and Skaar, 2008). Although the mechanism of heme detoxification by HrtAB remains unclear, the ability of *S. aureus* to express HrtAB upon heme exposure demonstrates that staphylococci sense and respond to heme in a way that prepares bacterial cells for growth in high heme concentrations.

In *S. aureus*, HrtAB expression is regulated by a two-component system (TCS) designated the heme sensor system (HssRS) (Stauff *et al.*, 2007; Torres *et al.*, 2007). HssS is a membrane-localized histidine kinase that senses heme through an unknown mechanism and undergoes histidine phosphorylation (Stauff *et al.*, 2007). HssR is a cytoplasmic response regulator that recognizes phosphorylated HssS and catalyzes phosphotransfer to a conserved HssR aspartic acid residue (Stauff *et al.*, 2007). When phosphorylated, HssR binds to a direct repeat (DR) within the *hrtAB* promoter, inducing expression of HrtAB (Stauff *et al.*, 2007). Importantly, interfering with proper heme metabolism through mutation of *hrtAB* or *hssRS* alters the virulence properties of *S. aureus*, highlighting the importance of heme sensing to the physiology and pathogenesis of staphylococci (Stauff *et al.*, 2008; Stauff and Skaar, 2008; Torres *et al.*, 2007).

It is currently unclear whether heme sensing and detoxification is a general feature of Gram-positive pathogens that associate with host tissues rich in heme. It is likely that a pathogen such as *B. anthracis*, which is capable of replicating to a high density within the bloodstream of its host (Dixon *et al.*, 1999), inducing hemolysis (Shannon *et al.*, 2003), and acquiring heme (Gat *et al.*, 2008; Maresso *et al.*, 2006; Maresso *et al.*, 2008), may require systems devoted to the avoidance of heme toxicity. Consistent with this supposition, Van Heyningen demonstrated in 1948 that *B. anthracis* is capable of proliferating on plates containing high concentrations of hemin (Heyningen, 1948). In contrast, *B. subtilis* (a bacterium not commonly associated with vertebrate tissues containing heme) is sensitive to hemin toxicity (Heyningen, 1948). Accordingly, *B. anthracis* encodes putative orthologues of *S. aureus* *hssRS/hrtAB* (Torres *et al.*, 2007), whereas *hss/hrt* genes are absent from the genome of *B. subtilis* (Stauff and Skaar, 2008).

Here, we demonstrate that *B. anthracis* encodes functional HssRS and HrtAB systems required for protection from hemin toxicity. We show that differential hemin resistance

within the *Bacilli* observed by Van Heyningen correlates with the presence of *hssRS/hrtAB* genes. Furthermore, *B. anthracis* HssRS exhibits an elevated autophosphorylation and phosphotransfer rate that correlates with the increased *in vivo* activity of this TCS. More specifically, we report that domains of the histidine kinase HssS are likely responsible for the elevated function of *B. anthracis* HssS. *B. anthracis* is highly sensitive to heme toxicity in the absence of HrtAB function, and may therefore require an increased ability to mount a heme detoxification response through heme sensing by HssS and signal transduction through HssR. We also demonstrate that mutation of residues within the previously uncharacterized signal recognition domain of HssS eliminates heme sensing by this histidine kinase. Last, we show that the *hrtAB* promoter is activated in a cutaneous model of anthrax, suggesting a role for heme sensing during *B. anthracis* infection. These results demonstrate that heme sensing by HssRS and subsequent heme detoxification by HrtAB are conserved in *B. anthracis* and that heme stress may be encountered by a wide range of pathogenic or saprotrophic Gram-positive bacteria.

## RESULTS

### Phylogenetically related *Bacilli* display differential heme resistance

To test whether encoding *hss/hrt* correlates with resistance to heme toxicity, we assessed the heme resistance of four *Bacilli* that contain these genes (*B. anthracis*, *B. cereus*, *B. thuringiensis*, and *B. israelensis*) and two with no identifiable *hss/hrt* orthologues (*B. licheniformis* and *B. subtilis*) (Fig. S1). We found that each of the four *Bacilli* encoding putative Hss/Hrt systems are able to proliferate in the presence of heme, while both *Bacilli* lacking Hss/Hrt systems succumb to heme toxicity (Fig. S1). These data link the presence of *hss/hrt* genes with resistance to heme toxicity.

### *B. anthracis* encodes *hss/hrt* genes required for heme resistance

We chose to investigate Hss/Hrt function in *B. anthracis* based on the fact that this organism is a mammalian pathogen and one that displays an intimate association with host blood during infection. *B. anthracis* HrtAB and HssRS share significant sequence identity with the corresponding proteins in *S. aureus* (HrtA, 49.3%; HrtB, 30.9%; HssS, 35.3%; HssR, 48.4%) (Stauff and Skaar, 2008; Torres *et al.*, 2007). The *hrtAB* and *hssRS* operons of *B. anthracis* are located less than 3 kb apart on the chromosome and are separated by two open reading frames, one of which encodes an uncharacterized transcription factor (BAS3024, Fig. 1A). An HssR consensus binding site can be found within the promoter of *B. anthracis* *hrtAB* (GTTTCATATT(N<sub>2</sub>)GTTTCATATT) (de Been *et al.*, 2008; Stauff *et al.*, 2007), suggesting that HssRS may regulate HrtAB expression in *B. anthracis* (Fig. 1A). Furthermore, we found that *B. anthracis* is capable of adapting to heme toxicity (Fig. S2), a phenotype that is likely due to expression of HrtAB in heme-treated cells prior to exposure to toxic levels of heme (Torres *et al.*, 2007). These observations are consistent with the proposition that *B. anthracis* adapts to heme through HssRS-dependent expression of HrtAB.

Accordingly, we assessed the ability of *B. anthracis* mutants lacking *hrtA* and *hssRS* to resist heme toxicity. While wildtype *B. anthracis* is able to proliferate in 10  $\mu$ M heme, *B. anthracis*  $\Delta$ *hrtA* and  $\Delta$ *hssRS* are unable to proliferate in this heme concentration, phenotypes that can be at least partially complemented by providing a full-length copy of the corresponding gene *in trans* (Fig. 1B; see *Heme preparation and concentrations* in Experimental Procedures for choice of heme concentration here and throughout). Importantly, *B. anthracis*  $\Delta$ *hrtA* and  $\Delta$ *hssRS* display no growth defect in the absence of heme (Fig. S3). This demonstrates that *B. anthracis* requires functional Hss/Hrt systems for heme resistance.

## The *B. anthracis* *hrtAB* promoter displays enhanced constitutive and hemin-induced activity

Although *S. aureus* and *B. anthracis* both require HrtAB for hemin resistance, it is unclear whether these systems display overlapping function. We therefore performed interspecies complementation experiments and found that *B. anthracis* or *S. aureus hrtAB* restore hemin resistance to *S. aureus*  $\Delta hrtAB$ , suggesting orthology at the level of transporter function between HrtAB from these organisms (Fig. 2A). In addition, we found that in *S. aureus* both the *S. aureus* and *B. anthracis hrtAB* promoters respond to hemin in a similar manner (Fig. 2B). These results confirm that *S. aureus* HssR is capable of activating the *S. aureus* and *B. anthracis hrtAB* promoters, consistent with the presence of an HssR consensus site within these promoters (Fig. 1A).

However, in *B. anthracis* the *hrtAB* promoter exhibits elevated levels of activation in response to hemin (Fig. 2C). Furthermore, the *B. anthracis hrtAB* promoter displays increased activity in the absence of hemin when compared with the *S. aureus hrtAB* promoter (Fig. 2C). In keeping with this, we tested whether the elevated constitutive activity of the *B. anthracis hrtAB* promoter depends on HssRS signaling and binding of HssR to the *hrtAB* promoter direct repeat (DR). Mutation of the DR or disruption of *hssRS* eliminates the hemin responsiveness and reduces the constitutive activity of the *B. anthracis hrtAB* promoter (Fig. 2D). This demonstrates that *B. anthracis* HssRS responds to hemin and activates the *hrtAB* promoter in a DR-dependent manner. Furthermore, these data suggest that in *B. anthracis*, HssRS constitutively induces HrtAB expression even in the absence of hemin, an activity not exhibited by *S. aureus* HssRS (Stauff *et al.*, 2007; Torres *et al.*, 2007).

The fact that *hrtAB* promoters from both species respond to hemin in a similar manner in *S. aureus* (Fig. 2B) but not in *B. anthracis* (Fig. 2C), despite similar DR sequences, indicates that bases outside of the *hrtAB* promoter DR may be important for promoter activity in *B. anthracis*. Accordingly, we found that single-base DR mutations have similar effects on the ability of *S. aureus* and *B. anthracis* HssR to activate the *hrtAB* promoter. Specifically, we found that mutation of four conserved DR residues, or removal or addition of spacer residues between the repeats eliminates the ability of *S. aureus* and *B. anthracis* HssR to activate the *S. aureus hrtAB* promoter in response to hemin (Fig. 2E, constructs 3, 8 – 10). Furthermore, mutation of T3 of the first repeat or T2 of the second repeat decreases hemin responsiveness in both species to the same extent (Fig. 2E, constructs 5 and 7). While *B. anthracis* HssR is more sensitive to mutation of A5 of repeat 2 and *S. aureus* is more sensitive to mutation of T9 of repeat 1 (Fig. 2E, constructs 4 and 6), both mutations decrease *hrtAB* promoter activity in response to hemin. This provides the first indication that HssR is exquisitely sensitive to alterations in the distance between each repeat and mutation of single DR residues. Overall, this suggests that HssR-*hrtAB* promoter interactions display similarity in *S. aureus* and *B. anthracis*, pointing to sequences outside of the DR to explain the hemin-independent, HssRS-dependent *hrtAB* promoter activity observed in *B. anthracis*.

### *S. aureus* and *B. anthracis* HssRS provide differential hemin resistance

We next tested whether *S. aureus* and *B. anthracis* HssRS display differential activity *in vivo* by assessing the ability of HssRS from *S. aureus* and *B. anthracis* to complement *S. aureus*  $\Delta hssRS$ . To account for potential differences in protein abundance, we tagged each *hssS* in the context of an *hssRS*-encoding plasmid, creating *hssR-hssSmyc* plasmids for the expression of HssR and HssSmyc from either species. We found that *S. aureus* and *B. anthracis* HssSmyc are expressed equivalently in these strains and that HssR-HssSmyc from either species is able to complement *S. aureus*  $\Delta hssRS$  (Fig. 3A).

In these experiments, we noticed that *B. anthracis* HssR-HssSmyc provides significantly greater complementation to *S. aureus*  $\Delta hssRS$  than HssR-HssSmyc from *S. aureus* (Fig. 3A). We tested whether differential complementation is conserved in *B. anthracis*  $\Delta hssRS$  and observed a similar trend (Fig. 3B). These results indicate that the molecular function of *B. anthracis* HssRS may differ from that of *S. aureus* HssRS in a way that provides for an increased ability to sense and respond to heme.

### **S. aureus and B. anthracis HssS exhibit differential function in vivo**

The ability of *B. anthracis* HssR-HssSmyc to impart elevated hemin resistance may be due to differences in HssR, HssS, or both. We reasoned that HssS differences are likely to contribute to differential activity of this TCS, as *B. anthracis*/*S. aureus* HssS share significantly less sequence identity than HssR (Stauff and Skaar, 2008). We performed complementation experiments with HssSmyc alone and found that *B. anthracis* HssSmyc provides significantly greater hemin resistance to *S. aureus*  $\Delta hssS$  than HssSmyc from *S. aureus* (Fig. 3C). We next attempted to repeat these observations in *B. anthracis*. Notably, HssSmyc from *B. anthracis* complements *B. anthracis*  $\Delta hssS$ , while HssSmyc from *S. aureus* fails to complement this strain in spite of similar expression of both proteins (Fig. 3D). These results suggest that, although *S. aureus* and *B. anthracis* HssRS are functionally orthologous TCS, *B. anthracis* HssRS provides elevated hemin resistance to bacteria due to differences in HssS function.

### **HssS chimeras reveal a complex domain interplay leading to elevated B. anthracis HssS activity**

The ability of *B. anthracis* HssS to provide greater hemin resistance than *S. aureus* HssS may be due to differences in the function of any of the domains comprising HssS. A membrane-localized histidine kinase, HssS is predicted to contain four domains: an N-terminal signal recognition domain flanked by two transmembrane helices (Mascher *et al.*, 2006), and a tripartite signaling domain consisting of a HAMP linker domain (Aravind and Ponting, 1999), a phosphate-accepting and response regulator recognition HisKA domain (Grebe and Stock, 1999), and a C-terminal ATPase domain (Grebe and Stock, 1999).

To define HssS domains required for the increased *in vivo* activity of *B. anthracis* HssS, we constructed chimeric forms of HssSmyc with either the sensing, HAMP, HisKA, or ATPase domain of *S. aureus* HssS in place of the respective *B. anthracis* HssS domain (Fig. 4; for detailed description of HssS chimeras, see Table S1). We reasoned that if a single *B. anthracis* HssS domain is responsible for the increased *in vivo* activity of this histidine kinase, replacing this domain with the corresponding *S. aureus* domain would decrease the function of the resulting chimera. The function of each chimera was tested in *S. aureus* instead of *B. anthracis* due to the inability of *S. aureus* HssS to rescue *B. anthracis*  $\Delta hssS$  (Fig. 3D) or to signal to *B. anthracis* HssR (see Fig. 6F).

Surprisingly, all chimeras which express in *S. aureus* and contain a single *S. aureus* domain allow this organism to proliferate to similar levels as *S. aureus* expressing full length *B. anthracis* HssSmyc (Fig. 4). In contrast, all chimeras containing two or more *S. aureus* HssS domains (with the exception of a chimera containing the entire *S. aureus* signaling domain, which is non-functional) function similar to *S. aureus* HssS *in vivo*, requiring over 8 hours to proliferate in 20  $\mu$ M hemin (Fig. 4). These data indicate that *B. anthracis* HssS is able to withstand single domain swapping with *S. aureus* HssS and retain its elevated function. It is likely that a complex interplay between HssS domains is responsible for the differential activity between *S. aureus* and *B. anthracis* HssS, an interplay that is perturbed by the insertion of more than one *S. aureus* HssS domain into *B. anthracis* HssS.

## B. anthracis HssS requires conserved residues in the predicted extracytoplasmic sensing domain for function in vivo

In HssS, signal recognition is predicted to occur within the putative extracytoplasmic domain, which is flanked by transmembrane helices (see Fig. 5A for schematic). However, no studies have addressed the importance of this input domain in HssS function. In this regard, we subjected the predicted sensing domain of *B. anthracis* HssS to mutational analysis to test whether it is required for HssS activity.

To identify residues within the predicted sensing domain that may be required for HssS activity, we aligned HssS sequences from *Firmicutes* predicted to encode *hss/hrt*. This analysis revealed a number of conserved residues within the predicted sensing domain (Fig. 5B). We selected 15 perfectly or highly conserved residues and a single non-conserved residue in *B. anthracis* HssS for alanine substitution mutational analysis, focusing on two patches of conserved sequence (T124-N129 and F149-F165, Fig. 5B). We also mutated the predicted phosphorylated histidine residue (H248) of HssS. We then expressed these point mutants in *B. anthracis*  $\Delta hssS$  and analyzed their expression and function.

By immunoblotting against the C-terminal C-Myc epitope tag added to HssS, we found that all point mutants with the exception of HssS:F149A are expressed to levels similar to wildtype HssS (Fig. 5C). As expected, HssS:H248A is unable to rescue the hemin sensitivity of *B. anthracis*  $\Delta hssS$ , presumably due to an inability to signal to HssR (Fig. 5D). Furthermore, HssS:Q145A, a mutant in a non-conserved HssS residue, complements  $\Delta hssS$  at levels similar to wildtype HssS (Fig. 5D). However, many of the point mutants are unable to complement *B. anthracis*  $\Delta hssS$  (Fig. 5D). Specifically, *B. anthracis* HssS mutants at residues K42, T124, F127, R151, E160, or R162 are unable to complement  $\Delta hssS$  to varying degrees. These data suggest that conserved residues within the predicted HssS sensing domain are required for HssS function.

## B. anthracis HssS-HssR undergoes rapid autophosphorylation and phosphotransfer

Because autophosphorylation is the major catalytic activity of histidine kinases (Mascher *et al.*, 2006), we hypothesized that differences in autophosphorylation between *S. aureus* and *B. anthracis* HssS may correlate with the increased ability of *B. anthracis* HssS to provide hemin resistance (Fig. 3C, Fig. 4). *In vitro*, *B. anthracis* HssS undergoes autophosphorylation that is dependent on conserved histidine 248 (Fig. 6B), correlating with its lack of functionality *in vivo* (Fig. 5D). Significantly, we found that the autophosphorylation rate of *B. anthracis* HssS is 208 (+/- 50) times greater than that of *S. aureus* HssS (Fig. 6A, 6B). This correlates with the ability of *B. anthracis* HssS to provide enhanced hemin resistance to *S. aureus*  $\Delta hssS$  (Fig. 3C, Fig. 4). Furthermore, these data also correlate with the increased *in vivo* activity of an HssS chimera consisting of the *S. aureus* HssS sensing domain fused to the *B. anthracis* HssS signaling domain (Fig. 4, chimera 1).

We next tested HssS-HssR phosphotransfer and found that both *S. aureus* and *B. anthracis* HssRS are capable of engaging in intraspecies phosphotransfer dependent on a conserved aspartic acid residue of HssR (Fig. 6C) (Stauff *et al.*, 2007). We also analyzed interspecies phosphotransfer using *S. aureus* HssS and found that *B. anthracis* HssR fails to catalyze phosphotransfer from *S. aureus* HssS (Fig. 6D, 6F), providing a potential explanation for the inability of *S. aureus* HssS to complement *B. anthracis*  $\Delta hssS$  (Fig. 3D). This indicates that evolutionary divergence has occurred between *S. aureus* and *B. anthracis* HssRS. We also tested interspecies phosphotransfer with *B. anthracis* HssS and found that HssR from both species are capable of catalyzing phosphotransfer from *B. anthracis* HssS (Fig. 6C, 6E, 6G). Notably, when phosphorylated *B. anthracis* HssS is incubated with *S. aureus* HssR,

phosphorylated *B. anthracis* HssS levels do not decrease because *B. anthracis* HssS continues to undergo autophosphorylation even after HssR is added (Fig. 3E).

In these experiments, we observed differences in the rate of *B. anthracis* and *S. aureus* HssS-HssR phosphotransfer. Specifically, we found that *B. anthracis* HssS-HssR phosphotransfer occurs rapidly (peaking in under 10 seconds, Fig. 6G) while *S. aureus* HssS-HssR phosphotransfer occurs at a relatively slow rate (peaking at about 10 minutes, Fig. 6F). The rate of phosphotransfer appears to be dictated by HssR rather than HssS, as slow phosphotransfer on the part of *S. aureus* HssR also occurs in a phosphotransfer reaction containing *B. anthracis* HssS (Fig. 6G). These results indicate evolutionary divergence between *S. aureus* and *B. anthracis* HssR that favors rapid phosphorylation of *B. anthracis* HssR. Importantly, the increased rate of *B. anthracis* HssS phosphorylation and HssS-HssR phosphotransfer may in part explain the ability of this TCS to provide elevated hemin resistance to *S. aureus* (Fig. 3A).

### ***B. anthracis* $\Delta$ hrtA is more sensitive to hemin toxicity than *S. aureus* $\Delta$ hrtA**

We reasoned that the enhanced function of *B. anthracis* HssRS may confer elevated hemin resistance to *B. anthracis* when compared with *S. aureus*. However, we found that both species display similar hemin resistance by growth curve analysis (Fig. 7A). We next reasoned that in the absence of Hss/Hrt function, *B. anthracis* may display enhanced sensitivity to hemin toxicity and may therefore require elevated HssRS activity to compensate. We therefore tested the relative hemin sensitivity of *S. aureus*  $\Delta$ hrtA and *B. anthracis*  $\Delta$ hrtA and found that after 4 hours in 10  $\mu$ M hemin, *S. aureus*  $\Delta$ hrtA exhibits a slight growth inhibition, while *B. anthracis*  $\Delta$ hrtA exhibits a dramatic decrease in surviving bacteria (Fig. 7B). Together with previous data, these results suggest that *B. anthracis* may have evolved a highly active HssRS system to cope with the fact that this organism is extremely susceptible to hemin toxicity.

### ***B. anthracis* experiences heme stress during growth within vertebrates**

Given the capacity of *B. anthracis* to acquire heme (Gat *et al.*, 2008; Maresso *et al.*, 2006; Maresso *et al.*, 2008; Skaar *et al.*, 2006) and to avoid hemin toxicity through HssRS-regulated expression of HrtAB (Fig. 1 and Fig. 2), we wondered whether HrtAB is expressed during systemic anthrax infection. To this end, we constructed a luminescence-based *hrtAB* promoter reporter by fusing the *S. aureus* *hrtAB* promoter to the *lux* genes of *Photobacterium luminescens* in the plasmid pXen-1 (Francis *et al.*, 2000). We utilized the *S. aureus* *hrtAB* promoter instead of the *B. anthracis* *hrtAB* promoter because the latter displays significant activity in the absence of hemin (Fig. 2C), making hemin-dependent induction and constitutive promoter activity difficult to differentiate. When *B. anthracis* is transformed with this construct, bacilli undergo luminescence in response to hemin (Fig. 8A).

We next infected mice subcutaneously (Chand *et al.*, 2009; Glomski *et al.*, 2007; Lamanna and Jones, 1963; Watts *et al.*, 2009) with spores prepared from *B. anthracis* harboring either pXen-1 or *phrt*.Lux and tracked bioluminescence of germinating *B. anthracis* cells over time. We found that *B. anthracis* harboring pXen-1 do not undergo detectable bioluminescence during infection (Fig. 8B, top), but that *B. anthracis* harboring *phrt*.Lux induces detectable levels of bioluminescence 24–36 hours post infection (Fig. 8B, bottom). Based on the strict hemin responsiveness of the *S. aureus* *hrtAB* promoter in *B. anthracis* (Fig. 2B, Fig. 8A), these data suggest that during infection, *B. anthracis* activates HssRS to alleviate heme toxicity.

## DISCUSSION

*Bacillus anthracis* is capable of invading host tissues to cause disease. During infection, *B. anthracis* acquires host heme to satisfy nutrient iron needs (Gat *et al.*, 2008; Maresso *et al.*, 2006; Maresso *et al.*, 2008; Reniere *et al.*, 2007). It appears that for *B. anthracis*, amassing of exogenous heme as a nutrient must be balanced with the need to overcome the toxicity of the heme molecule.

We have demonstrated a high level of orthology between the *S. aureus* and *B. anthracis* Hss/Hrt systems. *S. aureus* and *B. anthracis* mutants lacking members of these operons display a similar hemin-sensitive phenotype (Fig. 1) (Torres *et al.*, 2007). Also, *B. anthracis* *hssRS* and *hrtAB* are able to complement *S. aureus* mutants lacking these genes, indicating orthologous function of both systems (Fig. 2, Fig. 3). Furthermore, our data demonstrate functional overlap between the *S. aureus* and *B. anthracis* heme sensor systems at the levels of signal sensing, signal transduction, autophosphorylation by HssS, phosphotransfer to HssR, interaction between HssR and the *hrtAB* promoter, and activation of the *hrtAB* promoter by HssR (Fig. 2, Fig. 3, Fig. 6) (Stauff *et al.*, 2007). These similarities point to the overall evolutionary conservation of Hss/Hrt function between *S. aureus* and *B. anthracis*, a further indication that these systems are important for the fitness of both bacteria at some point throughout their life cycles.

Nonetheless, we have observed significant functional differences between HssRS/HrtAB from *S. aureus* and *B. anthracis*. First, the *B. anthracis* *hrtAB* promoter displays significant activity in the absence of hemin (Fig. 2C, 2D). This activity is dependent on HssRS and the DR to which HssR binds, suggesting that *B. anthracis* HssRS engages in constitutive signaling that induces low-level activation of the *hrtAB* promoter. The fact that this occurs only in *B. anthracis* and that most DR mutations are tolerated to a similar extent by both species points to sequences outside of the DR that may be important for HrtAB expression. It is possible that other unidentified factors expressed only by *B. anthracis* (such as the putative transcriptional regulator located between *hssRS* and *hrtAB*, Fig. 1A) may interact with the *hrtAB* promoter outside of the DR, functioning with HssR to increase expression of HrtAB. Alternatively, another *B. anthracis*-specific two-component systems may utilize the *hrtAB* promoter DR, overlapping with HssRS to induce expression of HrtAB (de Been *et al.*, 2008).

Another key functional difference between the *S. aureus* and *B. anthracis* heme sensor systems occurs at the levels of HssS autophosphorylation and phosphotransfer to HssR (Fig. 6). The signaling domain of *B. anthracis* HssS displays a significantly elevated rate of autophosphorylation *in vitro* compared to the corresponding *S. aureus* HssS domain. Although the molecular basis for the increased autophosphorylation rate of the *B. anthracis* HssS signaling domain is unclear, it is likely that it involves interplay between the three cytoplasmic sub-domains. This is indicated by data demonstrating that the increased *in vivo* activity of full-length *B. anthracis* HssS can be reversed by insertion of multiple HssS domains from *S. aureus* HssS (Fig. 4). We have also found that *B. anthracis* HssS-HssR phosphotransfer occurs more rapidly than *S. aureus* HssS-HssR phosphotransfer. These activities are likely to allow the rapid induction of HrtAB expression in *B. anthracis*, alleviating heme toxicity. Accordingly, *S. aureus* expressing *B. anthracis* HssRS is more resistant to hemin toxicity than wildtype *S. aureus*. These results suggest that a certain degree of functional divergence has occurred throughout the course of evolution between *S. aureus* and *B. anthracis* HssRS.

In the absence of the ability to respond to growth in the presence of hemin, *B. anthracis* is hypersensitive to hemin toxicity (Fig. 7). It is possible that *B. anthracis* may amass hemin



more rapidly than *S. aureus* or that *B. anthracis* may be more sensitive to heme-induced perturbation than *S. aureus*. It should be noted that the Isd heme iron acquisition system is not expressed in the conditions used in these experiments (data not shown), indicating that heme toxicity can occur independently of Isd-mediated heme uptake. Regardless of the reason for increased heme susceptibility in the absence of a functional heme detoxification response, the heme resistance of wildtype *B. anthracis* is similar to that of wildtype *S. aureus*. This is an indication that in *B. anthracis*, HssRS and/or HrtAB function more efficiently than the corresponding systems in *S. aureus*. Throughout the course of evolution, *B. anthracis* may have accumulated amino acid substitutions in HssRS that lead to constitutive activity of this TCS, increased autophosphorylation of HssS, and rapid HssS-HssR phosphotransfer. It is possible that these activities compensate for the heme sensitivity of *B. anthracis* and that such compensatory mutations may have been necessitated by the association between *B. anthracis* and host tissues rich in heme.

As the first component in the HssRS/HrtAB-mediated process of heme detoxification, the histidine kinase HssS controls heme resistance by detecting exogenous heme and transducing signals through the membrane. However, the mechanism of heme sensing by this histidine kinase remains elusive (Stauff and Skaar, 2008). Here, we present data showing that mutation of any of a number of conserved residues in the predicted HssS sensing domain eliminates the ability of *B. anthracis* to overcome heme toxicity. These residues cluster within two conserved patches, a possible indication that these regions of the protein interface with the HssS ligand. Although these studies do not identify the signal directly recognized by HssS, they do suggest that the previously uncharacterized sensing domain of HssS is involved in detection of heme. The fact that this sensing domain is flanked by transmembrane helices and is predicted to reside in the extracytoplasmic space indicates that HssS detects a signal located either between the membrane and the cell wall, or within the membrane itself. We hypothesize that the HssS ligand may be heme or a molecule that is either modified or induced by heme exposure, and that this molecule is recognized by conserved HssS sensing domain residues shown here to be required for HssS activity.

The fact that *B. anthracis* activates the *hrtAB* promoter during infection of vertebrates (Fig. 8) and heme detoxification appears to be involved in *S. aureus* virulence (Torres *et al.*, 2007) points to a role for heme sensing during the life cycles of two Gram-positive pathogens. Furthermore, a number of pathogenic and saprotrophic relatives of *S. aureus* and *B. anthracis* including *B. cereus* and *Listeria monocytogenes* are predicted to express HssRS/HrtAB (Stauff and Skaar, 2008; Torres *et al.*, 2007). It is therefore likely that heme sensing and detoxification are processes engaged in by a number of related Gram-positive organisms. This is supported by the observation made by Van Heyningen of differential heme resistance among phylogenetically related *Bacilli* (Heyningen, 1948). Van Heyningen's method may have differentiated organisms that express functional Hss/Hrt systems from those that do not based solely upon the heme resistance of the organism. Importantly, the fact that HssRS/HrtAB orthologues are only found in those *Bacilli* that exhibit pathogenic or saprotrophic lifestyles indicates that these systems are not required for the fitness of bacteria that do not encounter exogenous heme throughout their life cycles.

It has been suggested that heme sensing and detoxification evolved in dense microbial communities rich in heme-synthesizing bacteria (Miller *et al.*, 2007; Pedersen *et al.*, 2008). This may explain the recent discovery of a heme-regulated HrtAB orthologue required for heme resistance in the lactic acid bacterium *Lactococcus lactis* (Pedersen *et al.*, 2008). Although *L. lactis* is not a pathogen, it may encounter environmental heme synthesized by nearby bacteria (Tatsumi and Wachi, 2008), necessitating HrtAB function (Pedersen *et al.*, 2008). Notably, *L. lactis* does not possess *hssRS* genes and may regulate expression of *hrtAB*

through a mechanism other than TCS-based signal sensing (Pedersen *et al.*, 2008). This indicates evolutionary conservation of HrtAB function but divergence of HrtAB regulation between the *Lactobacillales* (which include the *Streptococci* as well as *L. lactis*) and the *Bacillales* (including *S. aureus* and *B. anthracis*).

Throughout the course of evolution, the pathogenic or saprotrophic *Firmicutes* may have acquired heme detoxification systems from bacteria such as *L. lactis*, thereby being provided with a means to facilitate survival within host tissues rich in heme. Within the *Firmicutes*, selective pressure to maintain a heme sensor system and a heme detoxification system may only be exerted on bacteria that commonly associate with exogenous heme. The evolutionary origin of the bacterial heme detoxification response will become clearer as the mechanism of heme sensing and detoxification are elucidated and as more organisms are identified that are capable of sensing and responding to heme toxicity.

## EXPERIMENTAL PROCEDURES

### Bacterial strains and growth conditions

*S. aureus* strain Newman (Duthie and Lorenz, 1952),  $\Delta hrtAB$  (Friedman *et al.*, 2006),  $\Delta hrtA$  and  $\Delta hssS$  (Torres *et al.*, 2007) have been described previously. *B. cereus*, *B. thuringiensis*, *B. israelensis*, *B. licheniformis*, and *B. subtilis* were acquired from the Bacillus Genetic Stock Center (College of Biological Sciences, The Ohio State University, www.bgsc.org). *B. anthracis* Sterne was used for all experiments (Sterne, 1937). *S. aureus* was grown in tryptic soy broth (TSB) or brain-heart infusion broth (BHI); *Bacilli* were grown in BHI or Luria broth (LB). All plasmid construction was performed using *Escherichia coli* DH5 $\alpha$ . Plasmid DNA to be introduced into *B. anthracis* was propagated in the *E. coli* strain K1077 (Kim *et al.*, 2004). Plasmid selection was performed in *S. aureus* and *B. anthracis* by supplementing media with 10  $\mu$ g/ml chloramphenicol (complementation and reporter plasmids) or 20  $\mu$ g/ml kanamycin (knockout constructs).

### Genetic manipulation of *S. aureus*

Genetic manipulation of *S. aureus* and construction of *S. aureus*  $\Delta hssRS$  was performed as described previously (Bae and Schneewind, 2006; Stauff *et al.*, 2007).

### Genetic manipulation of *B. anthracis*

Electroporations were performed as previously described (Kim *et al.*, 2004), with modifications. *E. coli* K1077 harboring plasmids to be introduced into *B. anthracis* were grown overnight in 15 ml of LB in 50 ml conical tubes. DNA was miniprep from K1077 and dialyzed against water to remove traces of salt prior to electroporation. Electrocompetent *B. anthracis* cells were prepared by growing bacteria in BHI + 0.5 M D-sorbitol to OD<sub>600</sub> = 0.2 and washing cells three times in SMG (0.5 M D-sorbitol, 0.5 M D-mannitol, 10% glycerol) without proteinase K treatment. Four  $\mu$ l of DNA was added to 50  $\mu$ l competent cells and mixtures were incubated on ice for 10 minutes. Electroporations were carried out in 1 mm gap cuvettes with a Bio-Rad Gene Pulser using an exponential decay protocol set to 2500 V, 1 ms time constant. After electroporation, cells were recovered in BHI + 0.5 M D-sorbitol for 2 hours at 30 °C with shaking and plated onto BHI + antibiotic. For generation of *B. anthracis* *hrtA*, *hssRS*, and *hssS* knockout constructs for insertion/deletion mutagenesis, the erythromycin resistance gene *ermC* was inserted between 1 kilobase regions flanking *hrtA*, *hssRS*, or *hssS* in the thermosensitive mutagenesis plasmid pLM4 (Kern and Schneewind, 2008). Mutagenesis was performed as described previously (Kern and Schneewind, 2008). The incorporation of mutations, loss of pLM4, and maintenance of pXO1 were confirmed by PCR.

## Epitope tagging of hssS

A C-terminal *c-myc* epitope tag was incorporated into *S. aureus* and *B. anthracis* *hssS* alone and *hssS* within the *hssRS* operon using a PCR-based approach as described previously (Stauff *et al.*, 2007). Briefly, PCR reactions were assembled containing a 5' primer annealing to the 5' end of *hssR* (for amplification of *hssR-hssSmyc*) or *hssS* (for *hssSmyc*), dilutions of a 3' primer annealing to the 3' end of *hssSmyc* and encoding the *c-myc* epitope tag (EQKLISEEDL), and excess of a 3' primer annealing to the *c-myc* coding region and incorporating a restriction endonuclease site. Tagged *hssS* is designated *hssSmyc* and tagged *hssRS* is designated *hssR-hssSmyc*.

## Construction of complementation plasmids

All complementation and reporter plasmids were constructed in a pOS1 (Schneewind *et al.*, 1992) or pOS1*plgt* (Bubeck Wardenburg *et al.*, 2006) background. A *B. anthracis* *hrtA* complementation construct was generated by PCR-SOE (Horton *et al.*, 1990) using a method described previously (Torres *et al.*, 2007). This resulted in a seamless fusion between the *B. anthracis* *hrtAB* promoter and the *B. anthracis* *hrtA* open reading frame. PCR-SOE DNA was digested with EcoRI and BamHI and inserted into pOS1*plgt*. For generation of a complementation plasmid containing *B. anthracis* *hssRS* under the control of its native promoter, *hssRS* was amplified by PCR and inserted into pOS1 between the PstI and BamHI sites. For generation of complementation plasmids containing *S. aureus*/*B. anthracis* *hssR-hssSmyc* under the control of the *S. aureus* *hssRS* promoter, the *S. aureus* *hssRS* promoter was first inserted between the EcoRI and NdeI sites of pOS1*plgt* to make pOS1*phssRS*. Promoterless *S. aureus* and *B. anthracis* *hssR-hssSmyc* were then amplified by PCR and inserted between the NdeI and BamHI sites of pOS1*phssRS*. For generation of complementation plasmids containing *S. aureus*/*B. anthracis* *hrtAB* under the control of their native promoters, *hrtAB* was amplified by PCR and inserted into pOS1*plgt* between the XhoI and EcoRI sites. To construct plasmids encoding *hssSmyc* under the control of the *S. aureus* *hssRS* promoter, *hssSmyc* was PCR amplified and inserted into pOS1*phssRS* between the NdeI and BamHI sites. Chimeric forms of *hssSmyc* were constructed by cloning *S. aureus* and *B. anthracis* *hssSmyc* in tandem in pCR2.1 (Invitrogen) and joining HssS domains together by inverse PCR followed by blunt-end ligation (see *Construction of Xyle reporter plasmids* below). Chimeric forms of *hssSmyc* were then inserted into pOS1*phssRS*. HssSmyc chimeras are described in Table S1.

## B. anthracis HssS alignment and mutagenesis

An alignment of conserved residues in HssS was created by assembling HssS amino acid sequences from all available sequenced *Firmicute* genomes (with the exception of *Exiguobacterium sibiricum*, which clusters outside of the *Staphylococcus*, *Listeria*, and *Bacillus* clusters) and aligning by Clustal V method using the Lasergene 6 software package.

## HssSmyc immunoblotting

For *S. aureus* HssSmyc immunoblots, cell lysates were prepared from bacteria grown in 5 ml TSB in 15 ml conical tubes for 15 hours at 37 °C with shaking. For *B. anthracis* HssSmyc immunoblots, cell lysates were prepared from bacteria grown in 5 ml BHI in 15 ml conical tubes for 12 hours at 30 °C with shaking. Bacteria were pelleted, washed in 5 ml TBS (50 mM tris, pH 7.5, 150 mM NaCl), digested in 0.5 ml TSM (100 mM tris, pH 7.0, 500 mM sucrose, 10 mM MgCl<sub>2</sub>) with 12.5 µg/ml lysostaphin (*S. aureus*) or 2 mg/ml lysozyme (*B. anthracis*), and protoplasts were sonicated in 0.5 ml TBS. Equal amounts of lysate and 2X SDS loading buffer were mixed, boiled and loaded onto 10% acrylamide gels, subjected to SDS-PAGE, and transferred to nitrocellulose membranes. Membranes were probed with 9E10 anti-C-Myc monoclonal primary and AlexaFluor-680-conjugated anti-

mouse secondary (Invitrogen) antibodies. Membranes were dried and scanned using an Odyssey Infrared Imaging System (LI-COR Biosciences).

### Construction of XylE reporter plasmids

The *B. anthracis hrtAB* promoter-*xylE* fusion was created by PCR-SOE (Horton *et al.*, 1990) as described for the *S. aureus hrtAB* promoter-*xylE* fusion (Torres *et al.*, 2007). Mutagenesis of the *S. aureus hrtAB* promoter-*xylE* fusion was carried out by inverse PCR-based mutagenesis. Briefly, a non-overlapping non-mutagenic primer and a primer introducing the desired mutation were designed which amplify in opposite directions on the plasmid template. Inverse PCR was carried out using *S. aureus hrtAB* promoter-*xylE* fusion as template and 5' phosphorylated oligos. Amplified DNA was blunt end ligated, transformed into *E. coli*, minipreped, and mutagenized constructs were validated by DNA sequencing.

### Construction of Lux reporter plasmids

To generate a construct in which the *lux* genes of pXen-1 (Xenogen) (Francis *et al.*, 2000) are under the control of the *S. aureus hrtAB* promoter (selected for lack of activity in the absence of hemin, Fig. 2C), primers containing a 5' EcoRI site and a 3' BamHI site were used to amplify the *hrtAB* promoter from *S. aureus* genomic DNA. PCR-amplified DNA was inserted into pXen-1 to create the plasmid *phrt.Lux*.

### Construction of plasmids for expression of recombinant proteins

Construction of protein expression constructs for *S. aureus* HssS, HssS:H249A, and HssR has been described previously (Stauff *et al.*, 2007). Construction of expression constructs for *S. aureus* HssR:D52N and *B. anthracis* HssR, HssS, HssS:H248A and HssR:D54N was performed as described previously (Stauff *et al.*, 2007). Briefly, *B. anthracis hssR* or *hssS* were amplified by PCR and inserted between the NdeI and BamHI sites of pET15b (Invitrogen). Constructs were validated by sequencing and were then mutagenized by inverse PCR (see *Construction of XylE reporter plasmids* above). For protein expression, pET15 constructs were introduced into *E. coli* BL21 (DE3). Bacteria were grown overnight in terrific broth (TB) and subcultured 1:100 into fresh TB. Bacteria were incubated at 37 °C with shaking to OD<sub>600</sub> = 0.4, and were then transferred to a 16 °C incubator with shaking. Cells were grown for another hour and then IPTG was added to a final concentration of 0.2 mM. Cells were grown overnight and hexahistidine-tagged proteins were purified as described previously (Stauff *et al.*, 2007). To validate the integrity of recombinant HssS, *S. aureus* and *B. anthracis* HssS were analyzed by mass spectrometry (Vanderbilt University Proteomics Core Facility). Intact masses were determined for both proteins and N- and C-termini were validated by MS/MS following digestion with trypsin protease.

### XylE reporter assays

XylE assays were performed as previously reported (Stauff *et al.*, 2007) with modifications. Bacteria harboring reporter plasmids were grown overnight in BHI (*S. aureus* and *B. anthracis* when analyzing hemin induction of *hrtAB* promoter) or LB (*B. anthracis* when analyzing hemin-independent *hrtAB* promoter activity) and subcultured 1:50 into 2 ml fresh medium in 15 ml conical tubes. Bacteria were grown for 5 hours at 37 °C and cytoplasmic fractions were prepared and analyzed as previously reported (Stauff *et al.*, 2007) except that for *B. anthracis*, lysozyme (2 mg/ml) was used instead of lysostaphin. Absolute XylE activities were determined for *B. anthracis* reporters due to lysozyme interference during protein quantification.

### Bacterial growth curve analyses

Triplicate cultures of bacteria were grown 12 hours at 30 °C with shaking in 5 ml of medium in 15 ml tubes. Each culture was subcultured 1:50 into 150 µl fresh medium in a 96-well round-bottom plate. Plates were wrapped to avoid evaporation and were grown at 37 °C with shaking at 180 rpm. 600 nm absorbance readings were taken on a Cary 50 MPR microplate reader coupled to a Cary 50 Bio UV-visible spectrophotometer (Varian, Inc.). For all experiments, absorbance readings of the triplicate subcultures were averaged and one standard deviation from the mean was plotted as error bars.

### Hemin preparation and concentrations

Hemin (Fluka) stocks were made by dissolving hemin in 0.1 M NaOH to a concentration of 10 mM. Hemin concentrations were chosen based on the following considerations. For all experiments attempting to assess *hrtAB* promoter activity without any detectable interference of hemin toxicity with strain growth (such as in hemin-sensitive mutants), promoter induction, or Xyle synthesis, 2 µM hemin was utilized. Five µM hemin fully induces the *hrtAB* promoter in *S. aureus* with no detectable hemin toxicity and was therefore utilized to pre-adapt *B. anthracis* to hemin toxicity. Ten µM hemin was selected for growth curve analysis of *B. anthracis hss/hrt* mutants as a hemin concentration known to completely inhibit growth of *S. aureus hss/hrt* mutants within 12 hours of subculture, and as a hemin concentration known to partially inhibit growth of wildtype *S. aureus* in TSB. Twenty µM hemin is known to inhibit growth of *S. aureus* in BHI and was therefore used to inhibit growth of *Bacilli*; 20 µM hemin also completely inhibits growth of *S. aureus* and *B. anthracis ΔhssS* up to 24 hours and was therefore used in experiments assessing the hemin resistance of complemented derivatives of *ΔhssS*. Thirty µM hemin was used to inhibit the growth of *S. aureus* and *B. anthracis ΔhssRS* due to the significant hemin resistance of strains harboring *hssRS* complementation plasmids. Fifty µM hemin was chosen in experiments analyzing the native or pre-adapted hemin resistance of wildtype strains in BHI due to the inherent hemin resistance of these strains.

### HssS-HssR autophosphorylation and phosphotransfer assays

To analyze autophosphorylation of *S. aureus* and *B. anthracis* HssS signaling domain, 25 µl reaction mixtures were prepared containing 288 µg/ml HssS, 50 mM tris, pH 8.0, 5 mM MgCl<sub>2</sub>, 200 mM KCl, 0.2 mM DTT, 10 % glycerol, 20 µM unlabeled ATP, 5 µCi [ $\gamma^{32}$ P]ATP. 5 µl samples were mixed with 2X SDS-PAGE loading buffer at various time points after the addition of radiolabeled ATP, samples were resolved by SDS-PAGE, and gels were dried and analyzed using a PhosphorImager. To analyze transphosphorylation from HssS to HssR, *B. anthracis* and *S. aureus* HssS were labeled for 15 and 50 minutes, respectively, in reaction mixtures described above. HssR or HssR mutant was added to this reaction to a final concentration of 890 µg/ml. Samples were taken and analyzed as described above.

### Bacterial hemin killing assays

Five ml cultures of *S. aureus* and *B. anthracis* wildtype and *ΔhrtA* were grown in 15 ml conical tubes for 12 hours at 30 °C with shaking at 180 rpm. Bacteria were subcultured 1:100 into BHI or BHI + 10 µM hemin and incubated at 37 °C for 4 hours. At the time of subculture and following incubation, bacterial colony forming units (CFU) were enumerated by performing serial dilutions in PBS and plating cells on BHI agar.

### *B. anthracis* infections and IVIS

Spores for infection of mice were prepared as described previously (Marraffini and Schneewind, 2006) with modifications. Briefly, *B. anthracis* harboring pXen-1 or *phrt.Lux*

were grown for 12 hours in 5 ml 2X SG medium (Leighton and Doi, 1971) containing 10 µg/ml chloramphenicol in 15 ml conical tubes at 30 °C with shaking. Cells were subcultured 1:100 into 250 ml flasks containing 30 ml 2X SG with 10 µg/ml chloramphenicol. Flasks were shaken at 37 °C for 4 days. Spore-containing cultures were then pelleted and washed three times with water, heated to 70 °C for 30 minutes, and then washed again with water. Purified spores were serially diluted and plated onto BHI to obtain a viable spore count. Spore preparations were stained using malachite green to microscopically confirm the presence of spores and the absence of vegetative bacilli. Spores harboring Lux plasmids were diluted to a concentration of 200 spores/µl in sterile PBS. One-hundred µl of spore dilutions (20000 spores/mouse) were injected subcutaneously into the inguinal region of 6-week-old A/J mice (two groups of 5 mice). Following spore inoculation, mice were visualized at 12-hour intervals using a Xenogen IVIS 200. Mice were monitored for signs of disease and were sacrificed when found to be moribund.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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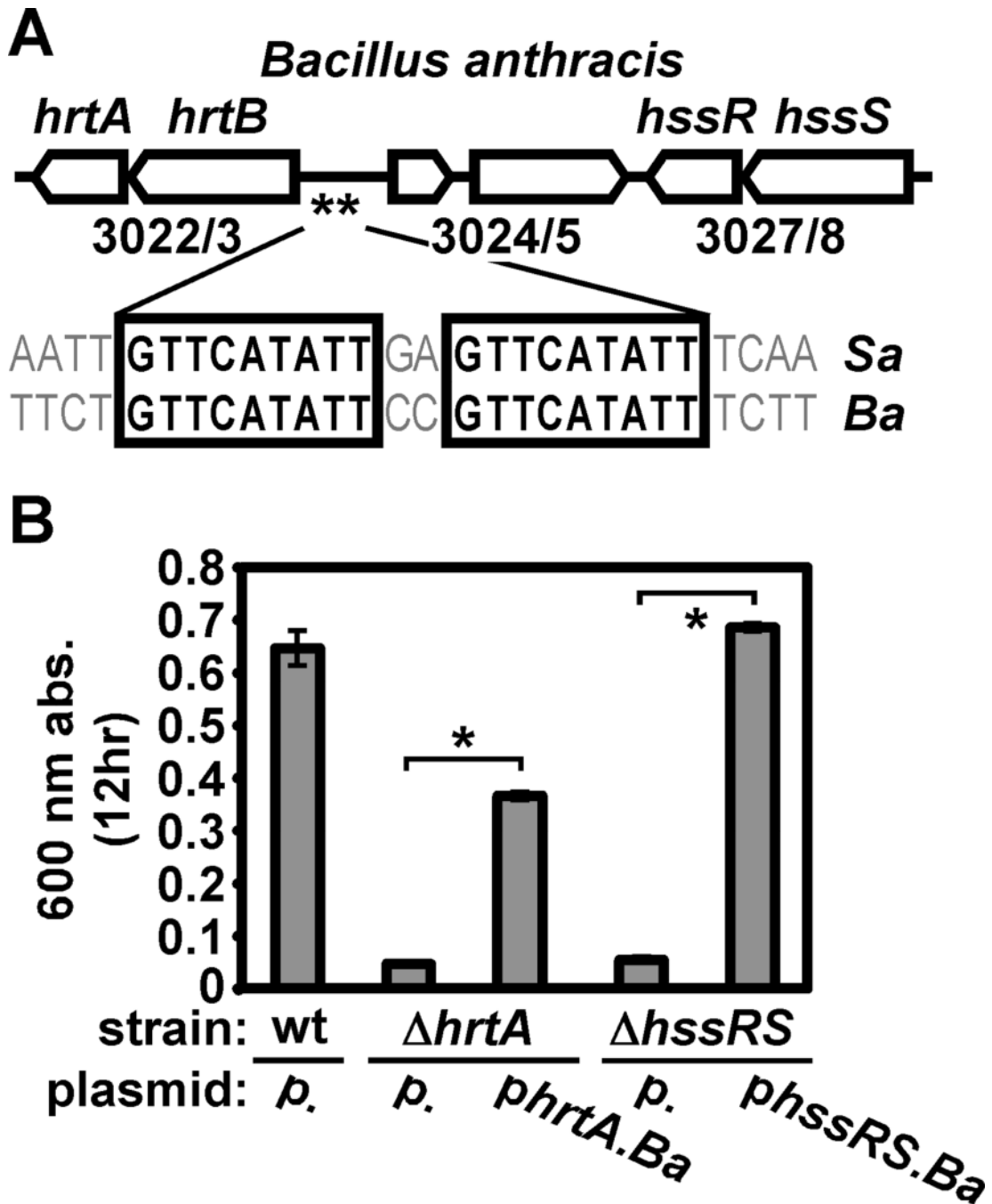
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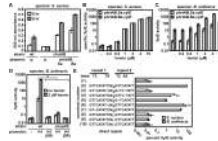
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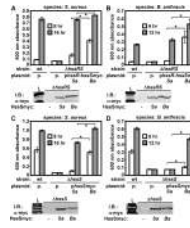
**Figure 1. *B. anthracis* requires HrtAB and HssRS for hemin resistance**  
**A.** Schematic of the *B. anthracis* *hss/hrt* locus. Open arrows represent open reading frames (ORF). Designations above ORFs denote predicted gene function (*hrtA*: ATPase; *hrtB*: permease; *hssS*: histidine kinase; *hssR*: response regulator). Numbers below ORFs correspond to gene numbers in the *B. anthracis* Sterne genome. Double asterisks signify a putative HssR binding site (consensus sequence GTTCATATT(N<sub>2</sub>)GTTCATATT) upstream of *hrtAB*; alignment shows the known *S. aureus* (*Sa*) and predicted *B. anthracis* (*Ba*) HssR binding sites. **B.** Contribution of *hrt* and *hss* to hemin resistance of *B. anthracis*. *B. anthracis* wildtype (wt) harboring plasmid pOS1 (p.),  $\Delta$ *hrtA* with either pOS1 or pOS1 encoding *hrtA* (*phrtA.Ba*), and  $\Delta$ *hssRS* with either pOS1 or pOS1 encoding *hssRS* (*phssRS.Ba*) were grown

for 12 hours in BHI. Bacteria were sub-cultured 1:100 into BHI containing 10  $\mu$ M hemin (a hemin concentration known to inhibit *S. aureus hss/hrt* mutants) in triplicate, grown for 12 hours at 37 °C, and culture densities were determined and averaged. Error bars correspond to one standard deviation from the mean; asterisks denote statistically significant complementation by Student's *t* test ( $p < 0.05$ ).



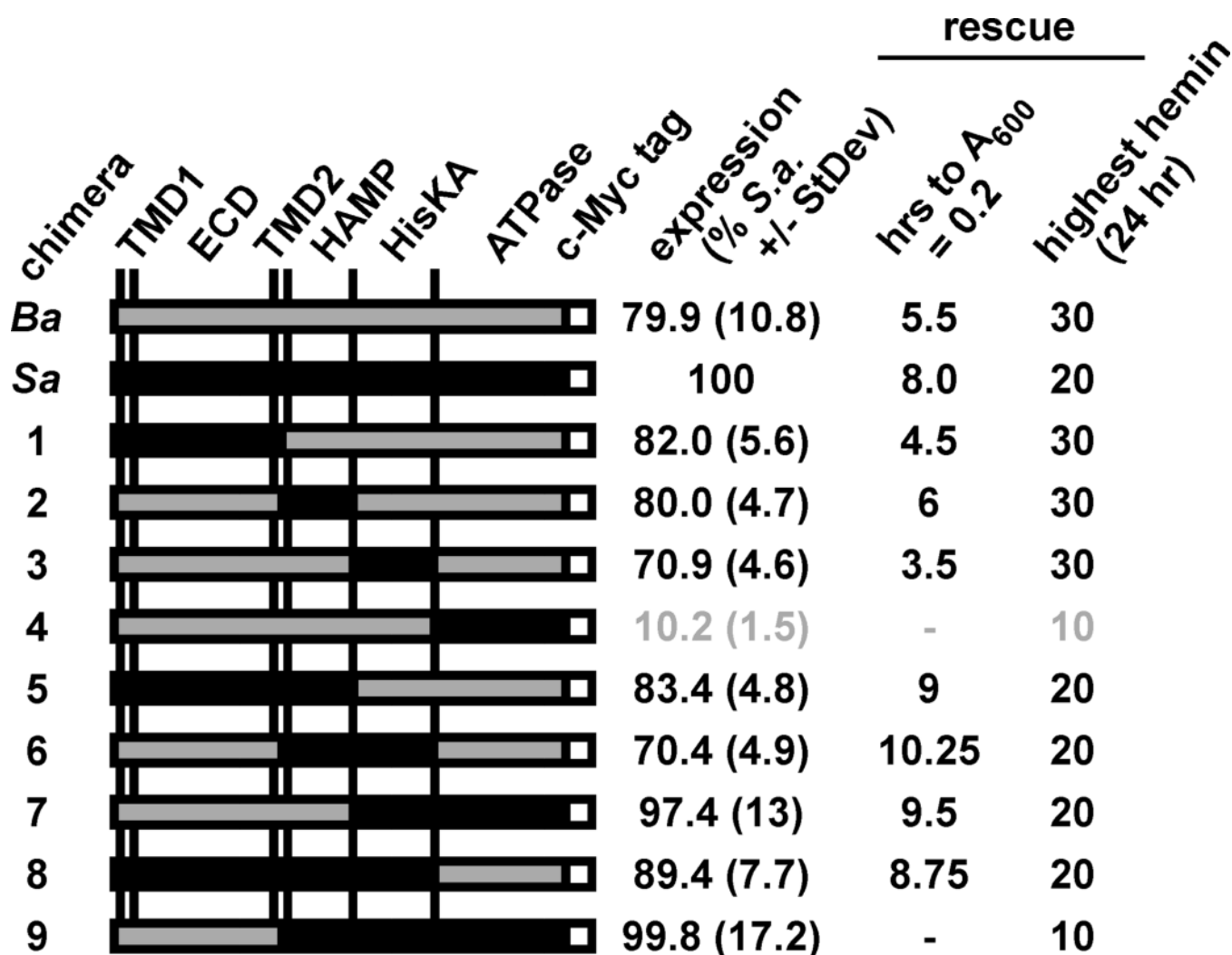
**Figure 2. Elevated activity of the *B. anthracis hrtAB* promoter**

**A.** Complementation of *S. aureus*  $\Delta hrtAB$  with *S. aureus* or *B. anthracis hrtAB*. *S. aureus* wildtype (wt) transformed with pOS1 (p.),  $\Delta hrtAB$  with pOS1,  $\Delta hrtAB$  with pOS1 encoding *S. aureus hrtAB* (*phrtAB.Sa*), and  $\Delta hrtAB$  with pOS1 encoding *B. anthracis hrtAB* (*phrtAB.Ba*) were grown in TSB + 10  $\mu$ M hemin (a hemin concentration known to inhibit growth of *S. aureus hss/hrt* mutants). Proliferation was tracked by growth curve analysis. Culture densities at 6 hour (white bars) and 8 hour (grey bars) time points are shown. **B.** Reporter assay monitoring *S. aureus* and *B. anthracis hrtAB* promoter activity in *S. aureus*. *S. aureus* harboring either an *S. aureus* (*phrtAB.Sa.xylE*, white bars) or *B. anthracis* (*phrtAB.Ba.xylE*, grey bars) *hrtAB* promoter XylE reporter were grown without hemin (–) or in the indicated concentration of hemin and reporter activity was measured. **C.** Reporter assay monitoring *S. aureus* and *B. anthracis hrtAB* promoter activity in *B. anthracis*. Strains were grown in LB to avoid contaminating hemin in BHI. **D.** The dependence on HssRS and the *hrtAB* promoter direct repeat (DR) for constitutive and hemin-induced *hrtAB* promoter activity in *B. anthracis*. *B. anthracis* wildtype (wt) or  $\Delta hssRS$  harboring a plasmid containing either a promoterless *xylE* (–, wildtype only), *xylE* under the control of the *B. anthracis hrtAB* promoter (*hrt*), or *xylE* under the control of the *hrtAB* promoter mutated at four conserved DR bases (*hrt*(DR), shown in schematic form in *E*, construct #3) were grown in LB (white bars) or LB + 2  $\mu$ M hemin (a hemin concentration that activates the *hrtAB* promoter without any detectable growth inhibition of *B. anthracis*  $\Delta hssRS$ ; grey bars) and XylE activity was measured. **E.** Reporter assay measuring sensitivity of *S. aureus* and *B. anthracis* HssRS to perturbations in the *hrtAB* promoter DR. *S. aureus* (white bars) or *B. anthracis* (grey bars) harboring a promoterless *xylE* (construct number 1), *xylE* under the control of the *S. aureus hrtAB* promoter (2), or *xylE* under the control of the *S. aureus hrtAB* promoter containing the indicated DR mutations (3–10; mutations are in grey) were grown in BHI or BHI + 2  $\mu$ M hemin. Reporter activity was measured and is expressed as the percent of hemin-induced activity observed for the wildtype *hrtAB* promoter. All results are representative of at least three independent experiments. Error bars correspond to one standard deviation from the mean of triplicate samples within the same experiment; asterisks denote statistically significant differences by Student's *t* test ( $p < 0.05$ ).

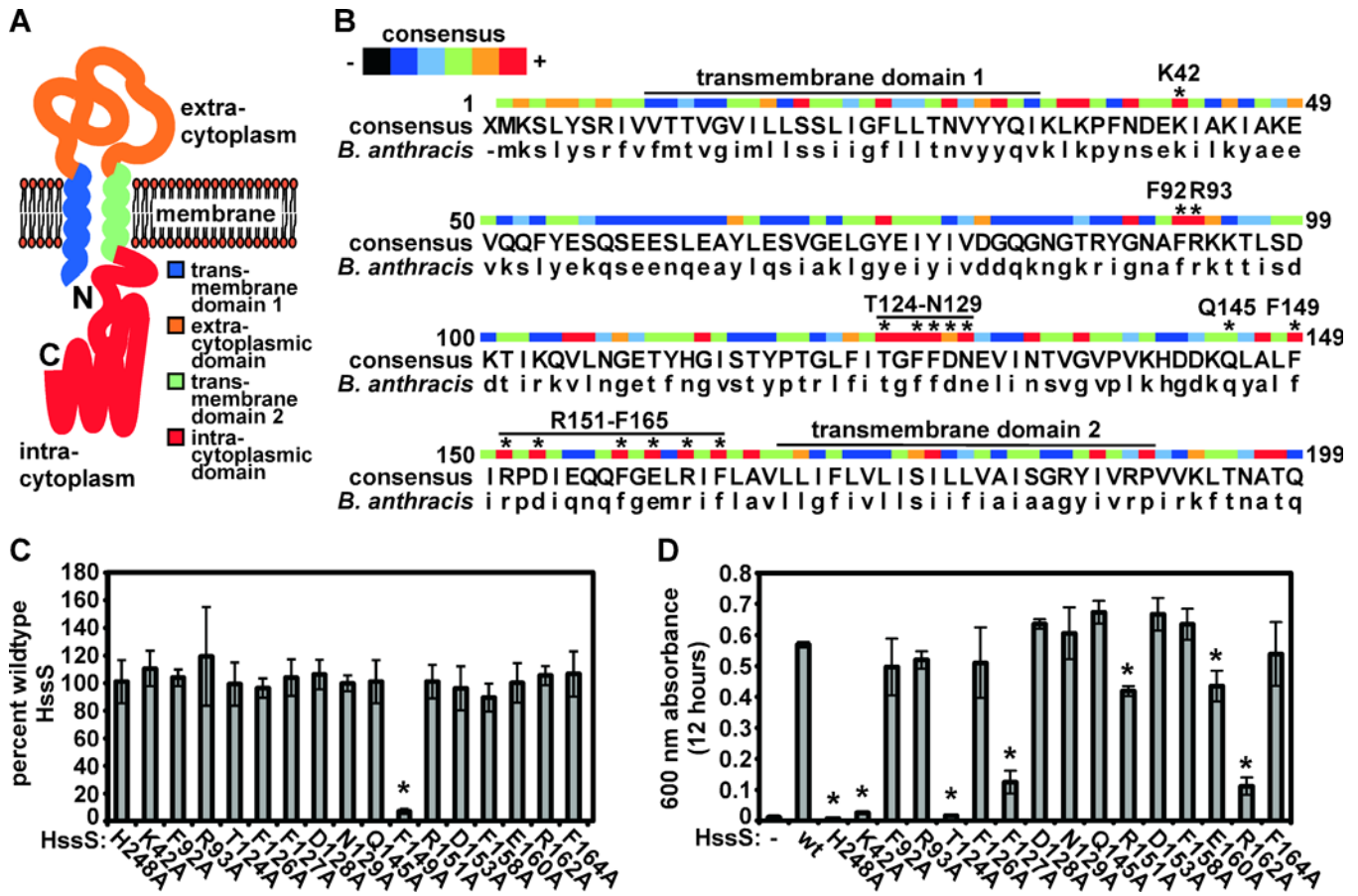


**Figure 3. *B. anthracis* HssRS displays elevated hemin sensing activity *in vivo***

**A.** Complementation of *S. aureus*  $\Delta hssRS$  by *S. aureus* and *B. anthracis* HssRS. Top: triplicate cultures of *S. aureus* wildtype (wt) harboring pOS1 (p.),  $\Delta hssRS$  with pOS1,  $\Delta hssRS$  with pOS1 encoding *S. aureus* HssR-HssSmyc (*phssR-hssSmyc.Sa*), and  $\Delta hssRS$  with pOS1 encoding *B. anthracis* HssR-HssSmyc (*phssR-hssSmyc.Ba*) were grown in TSB + 30  $\mu$ M hemin (a hemin concentration chosen due to the exceptional hemin resistance of strains containing the *phssR-hssSmyc.Ba* plasmid). Proliferation was tracked by growth curve analysis. Average culture density at 8 hour (white bars) and 16 hour (grey bars) time points are shown. Bottom: anti-C-Myc immunoblot showing expression of *S. aureus* and *B. anthracis* HssSmyc in *S. aureus*  $\Delta hssRS$ . Lane 1: molecular weight ladder (L). Lanes 2–4: *S. aureus*  $\Delta hssRS$  transformed with pOS1 (–), *phssR-hssSmyc.Sa* (*Sa*), or *phssR-hssSmyc.Ba* (*Ba*). **B.** Top: Complementation of *B. anthracis*  $\Delta hssRS$  by *S. aureus* and *B. anthracis* HssRS (same plasmids described in A). Experiment was carried out as described in A, except that earlier time points (6 hour (white) and 12 hour (grey)) are shown due to sporulation and autolysis observed at late time points in *B. anthracis* cultures. Bottom: anti-C-Myc immunoblot showing expression of *S. aureus* and *B. anthracis* HssSmyc in *B. anthracis*  $\Delta hssRS$ . Lanes are the same as those described in A. **C.** Top: Complementation of *S. aureus*  $\Delta hssS$  by *S. aureus* and *B. anthracis* HssS. Top: *S. aureus* wildtype harboring plasmid pOS1,  $\Delta hssS$  with pOS1,  $\Delta hssS$  with pOS1 encoding *S. aureus* HssSmyc (*phssSmyc.Sa*), and  $\Delta hssS$  with pOS1 encoding *B. anthracis* HssSmyc (*phssSmyc.Ba*) were analyzed as described in A, except that subcultures were performed in 20  $\mu$ M hemin (chosen due to lower hemin resistance of complemented  $\Delta hssS$  strains compared to strains described in A). Bottom: anti-C-Myc immunoblot showing expression of *S. aureus* and *B. anthracis* HssSmyc in *S. aureus*  $\Delta hssS$ . Lane 1: molecular weight ladder (L). Lanes 2–4: *S. aureus*  $\Delta hssS$  transformed with empty plasmid (–), *phssSmyc.Sa* (*Sa*), or *phssSmyc.Ba* (*Ba*). **D.** Top: Complementation of *B. anthracis*  $\Delta hssS$  by *S. aureus* and *B. anthracis* HssS. Experiment was performed as described in C. Bottom: anti-C-Myc immunoblot showing expression of *S. aureus* and *B. anthracis* HssSmyc in *B. anthracis*  $\Delta hssS$ . Lanes are the same as those described in C. All results are representative of at least three independent experiments. Error bars correspond to one standard deviation from the mean of triplicate samples within the same experiment. Asterisks denote statistically significant differences in complementation by Student's *t*-test ( $p < 0.05$ ).

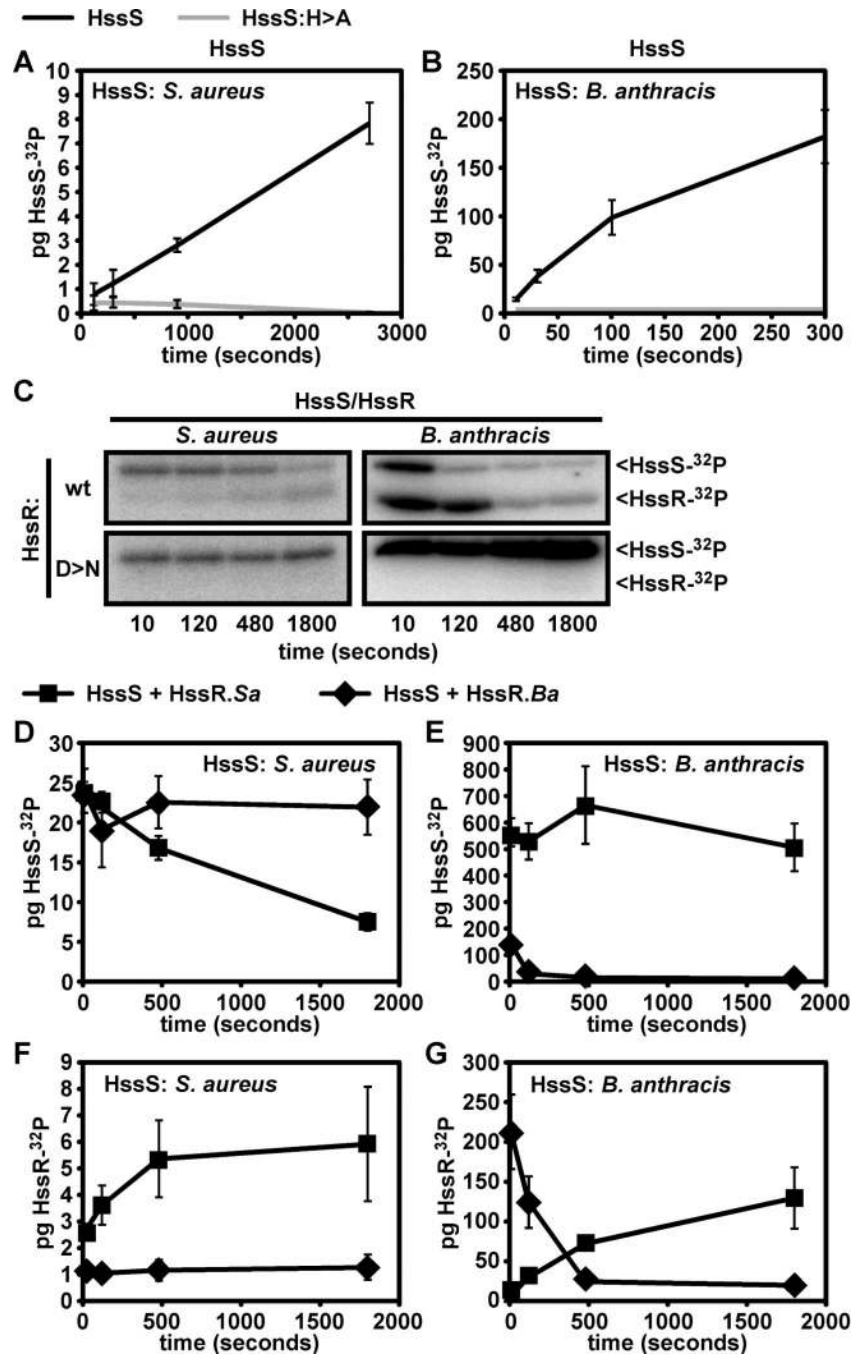


**Figure 4. Rescue of *S. aureus*  $\Delta hssS$  by *S. aureus*/*B. anthracis* HssS chimeras**  
 Left: schematic of HssS chimeras. Black: *S. aureus* HssS; grey: *B. anthracis* HssS; white: C-terminal C-Myc epitope tag. TMD1/2, predicted transmembrane domains; ECD, predicted extracytoplasmic sensing domain; HAMP, intracytoplasmic linker domain; HisKA, phosphate acceptor domain; ATPase, ATP hydrolysis domain. Chimera junctions are defined in Table S1. Right side: chimera functional data. Column 1: Expression of HssS chimeras. Triplicate cultures of *S. aureus*  $\Delta hssS$  expressing each shown HssS variant were prepared and HssS expression was analyzed by immunoblot against the C-Myc epitope tag of HssS followed by LiCor-based quantification. Chimera levels relative to abundance of *S. aureus* HssS were determined and averaged. Standard deviations are shown. Data corresponding to chimeras with less than 50% expression are highlighted in grey. Columns 2 and 3: rescue of  $\Delta hssS$  by HssS chimeras. Column 2: strains were inoculated into medium containing 20  $\mu$ M hemin (a hemin concentration that partially inhibits wt *S. aureus* and completely inhibits *S. aureus*  $\Delta hssS$ ) and proliferation was tracked by growth curve analysis. The length of time required to initiate growth (OD<sub>600</sub> = 0.2) was determined. Column 3: strains were inoculated into medium containing 10, 20, or 30  $\mu$ M hemin. The highest hemin concentration each strain was able to grown in by 24 hours is indicated. Results are representative of at least three experiments.



**Figure 5. Residues within the predicted *B. anthracis* HssS sensing domain required for HssS function**

**A.** Schematic of *B. anthracis* HssS. *B. anthracis* HssS is predicted to consist of an N-terminal signal peptide/transmembrane domain (blue), an extracytoplasmic sensing domain (orange), a second transmembrane domain (green), and an intracytoplasmic signaling domain (red). **B.** Alignment of conserved HssS sensing domain residues. An HssS consensus sequence was generated from an alignment performed using data from all species with complete genome sequences that are predicted to contain *hss/hrt* (capital letters). Color code indicates level of conservation (blue, little conservation; red, absolute conservation). Asterisks above residues designate amino acids subjected to alanine substitution mutagenesis. The signaling domain autophosphorylated histidine residue (H248) was also selected for mutagenesis. **C.** Expression of *B. anthracis* HssSmyc point mutants. Triplicate cultures of *B. anthracis*  $\Delta hssS$  expressing each HssSmyc point mutant were prepared and HssS expression was analyzed by immunoblot against the C-Myc epitope tag of HssSmyc followed by LiCor-based quantification. Mutant levels relative to abundance of wildtype *B. anthracis* HssSmyc were determined and averaged. Error bars correspond to standard deviations; asterisk denotes point mutant with expression significantly lower than wild type. **D.** Rescue of *B. anthracis*  $\Delta hssS$  by HssS point mutants. Strains were inoculated into BHI containing 20  $\mu$ M hemin (a hemin concentration that does not significantly inhibit wt *B. anthracis* in BHI but completely inhibits *B. anthracis*  $\Delta hssS$ ) and grown for 12 hours. The mean optical density of triplicate cultures was determined. Error bars correspond to standard deviations; asterisks denote strains with culture density significantly lower than those of  $\Delta hssS$  expressing wildtype HssSmyc.

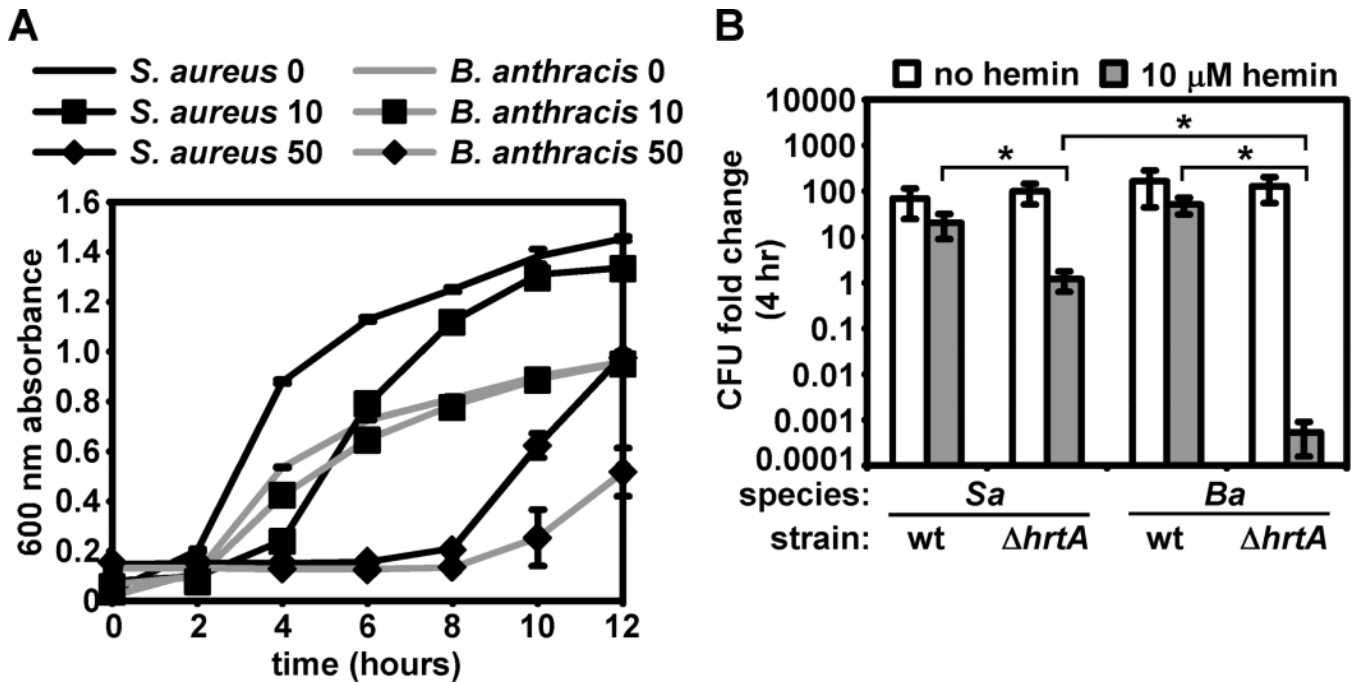


**Figure 6. *B. anthracis* HssS–HssR exhibit enhanced autophosphorylation and phosphotransfer *in vitro***

**Symbols for proteins used in A and B.** Black lines: wildtype HssS signaling domain (*S. aureus* HssS in A, *B. anthracis* HssS in B); grey lines: mutant HssS with alanine substituted for predicted phosphorylated histidine residue (HssS:H>A). **A.** Autophosphorylation of *S. aureus* HssS and phosphorylation site mutant. The described *S. aureus* HssS variants were incubated with  $^{32}\text{P}$ -ATP for the indicated time and samples were taken and analyzed by SDS-PAGE followed by quantification via phosphorimager analysis. **B.** Autophosphorylation of the described *B. anthracis* HssS variants. Experiments were performed as described in A. **C.** Intraspecies HssS-HssR phosphotransfer using *S. aureus*

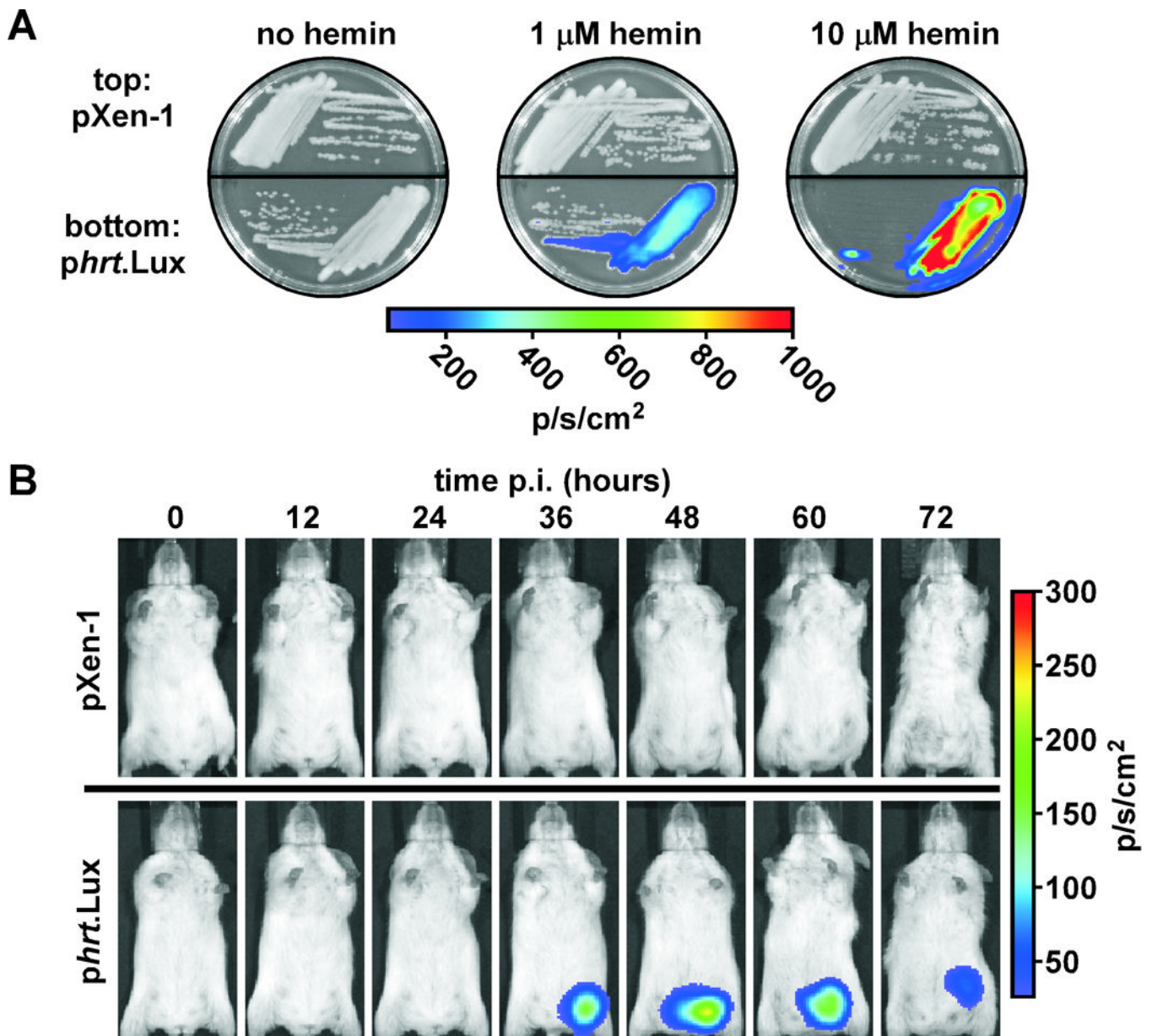
and *B. anthracis* HssRS. Phosphorylated *S. aureus* (left panels) or *B. anthracis* (right panels) HssS was added to wildtype HssR (wt, top panels) or HssR mutated at the predicted phosphorylated residue (D>N, bottom panels). Samples were analyzed as described in A. ***D through G: interspecies phosphotransfer. Symbols for reactions analyzed in D through G:*** Lines with squares, reactions containing HssS (*S. aureus* HssS in *D* and *F*, *B. anthracis* HssS in *E* and *G*) and wildtype *S. aureus* HssR (HssR.Sa); lines with diamonds, reactions containing HssS and wildtype *B. anthracis* HssR (HssR.Ba). ***D.*** Amount of phosphorylated HssS in phosphotransfer reactions containing *S. aureus* HssS. Panel shows levels of phosphorylated HssS after the addition of HssR from the indicated species. ***E.*** Amount of phosphorylated HssS in phosphotransfer reactions containing *B. anthracis* HssS. Experiment was performed as in *D* using *B. anthracis* HssS. ***F.*** HssR phosphorylation in phosphotransfer reactions containing *S. aureus* HssS. ***G.*** HssR phosphorylation in phosphotransfer reactions containing *B. anthracis* HssS. Quadruplicate reactions were performed and averaged for all experiments. Error bars correspond to one standard deviation from the mean of data from all four experiments.





**Figure 7. Differential hemin sensitivity of *S. aureus* and *B. anthracis*  $\Delta hrtA$**

**A.** Proliferation of wildtype *S. aureus* and *B. anthracis* in hemin. Growth curve analysis was performed on *S. aureus* (black lines) and *B. anthracis* (grey lines) grown in BHI without hemin (no symbols), BHI + 10  $\mu$ M hemin (a hemin concentration that inhibits *hss/hrt* mutants; boxes), or BHI + 50  $\mu$ M hemin (a hemin concentration that inhibits wt *S. aureus* and *B. anthracis*; diamonds). **B.** Resistance of wildtype and  $\Delta hrtA$  *S. aureus* and *B. anthracis* to hemin. *S. aureus* (*Sa*) and *B. anthracis* (*Ba*) wildtype (wt) and  $\Delta hrtA$  were grown overnight and subcultured into BHI or BHI + 10  $\mu$ M hemin. At the time of subculture and after a four-hour incubation, serial dilutions were performed and plated and colony forming units (CFU) were enumerated. Data is presented as the fold change in CFU following incubation in the presence or absence of hemin (log scale). Asterisks denote statistically significant differences by Student's *t*-test ( $p < 0.05$ ). All results are representative of triplicate independent experiments. Error bars correspond to one standard deviation from the mean of triplicate samples within the same experiment; asterisks denote statistically significant differences by Student's *t* test ( $p < 0.05$ ).



**Figure 8. HrtAB expression in an animal model of cutaneous anthrax**

**A.** Hemin-dependent luminescence of *B. anthracis* harboring *phrt.Lux*. *B. anthracis* harboring promoterless pXen-1 or pXen-1 with the Lux genes under the control of the *hrtAB* promoter (*phrt.Lux*) were grown on BHI agar containing no hemin, BHI + 1  $\mu$ M hemin, or BHI + 10  $\mu$ M hemin. Plates were visualized by IVIS. **B.** Activation of the *hrtAB* promoter during anthrax infection. Spores of *B. anthracis* harboring pXen-1 or *phrt.Lux* were used to infect groups of five A/J mice via a sub-dermal injection in the left inguinal region. Luminescence of *B. anthracis* was tracked every 12 hours post infection (p.i.) by IVIS. Shown is one mouse representative of each group.