Two-Component System Cross-Regulation Integrates Bacillus anthracis Response to Heme and Cell Envelope Stress

Laura A. Mike, Jacob E. Choby, Paul R. Brinkman, Lorenzo Q. Olive, Brendan F. Dutter, Samuel J. Ivan, Christopher M. Gibbs, Gary A. Sulikowski, Devin L. Stauff, Eric P. Skaar

Abstract

Two-component signaling systems (TCSs) are one of the mechanisms that bacteria employ to sense and adapt to changes in the environment. A prototypical TCS functions as a phosphorelay from a membrane-bound sensor histidine kinase (HK) to a cytoplasmic response regulator (RR) that controls target gene expression. Despite significant homology in the signaling domains of HKs and RRs, TCSs are thought to typically function as linear systems with little to no cross-talk between non-cognate HK-RR pairs. Here we have identified several cell envelope acting compounds that stimulate a previously uncharacterized Bacillus anthracis TCS. Furthermore, this TCS cross-signals with the heme sensing TCS HssRS; therefore, we have named it HssRS interfacing TCS (HitRS). HssRS reciprocates cross-talk to HitRS, suggesting a link between heme toxicity and cell envelope stress. The signaling between HssRS and HitRS occurs in the parental B. anthracis strain; therefore, we classify HssRS-HitRS interactions as cross-regulation. Cross-talk between HssRS and HitRS occurs at both HK-RR and post-RR signaling junctions. Finally, HitRS also regulates a previously unstudied ABC transporter implicating this transporter in the response to cell envelope stress. This chemical biology approach to probing TCS signaling provides a new model for understanding how bacterial signaling networks are integrated to enable adaptation to complex environments such as those encountered during colonization of the vertebrate host.

Introduction

One mechanism by which bacteria sense and adapt to their environment is through the use of two-component signaling systems (TCSs). A prototypical TCS consists of a membrane-bound histidine kinase (HK) and cytoplasmic response regulator (RR). In the presence of a specific signal, the HK autophosphorylates at a conserved intracellular histidine residue and then transfers the phosphate to an aspartate on the cognate RR. This phosphorylation event activates the RR, which binds to target promoter regions and subsequently regulates gene expression [1]. In this manner the classic bacterial TCS is thought to function in a linear fashion, in that each HK has a defined input that results in a specific output from the RR.

TCSs are present in nearly all sequenced bacterial genomes as well as some fungal, archaeal, and plant species; most species encode 20-30 TCSs, while others carry upwards of 200-300 putative TCSs [2]. The conserved nature of the TCS signaling pathway means that core structural elements are similar amongst most HKs and RRs. The appreciation that interactions, both beneficial and detrimental, occur between TCSs has become more prevalent over the past two decades. This ‘cross-talk’ has been previously defined as the “communication between two pathways that, if eliminated, would leave intact two distinct, functioning pathways” [3]. When cross-talk between two signaling systems is of physiological significance to an organism this is referred to as ‘cross-regulation’ [3]. An activating signal is required in order to observe cross-regulation, but not cross-talk.

The current understanding of HK-RR signaling purity implicates molecular recognition, HK phosphatase activity, and substrate competition as the primary mechanisms for ensuring the proper transcriptional response [4]. Molecular recognition dictates HK-RR specificity as HK-RR pairs have co-evolved amino acid residues that drive a strong preference for phospho-transfer from the HK to its cognate RR. This phenomenon is observable in vitro as most HKs have a strong kinetic preference for...
Author Summary

The vertebrate host is a hostile environment to microbes, where large variations in temperature, oxygen, and nutrient availability challenge microbial replication. In order to successfully infect vertebrates, pathogens must sense that they have entered their host and alter gene expression accordingly. One way bacteria adapt to their environment is by using two-component signaling systems (TCSs), which are comprised of a signal sensor and a transcriptional regulator. The core signaling components of TCSs are conserved across the bacterial kingdom; however, examples of interactions between two distinct TCSs are extremely rare. Here we have probed how Bacillus anthracis, the causative agent of anthrax, adapts to life in the vertebrate host. The B. anthracis heme sensing TCS, HssRS, senses heme as a component of vertebrate blood and protects the bacteria from heme toxicity. We have identified a new TCS (HitRS), which is activated by compounds that alter the integrity of the cell envelope, and interacts with HssRS to coordinate a simultaneous response to both heme and cell envelope stress. This reciprocal HssRS-HitRS signaling is an unusual demonstration of bacterial signal cross-regulation and suggests that coordinating the response to heme and cell envelope stress is important for B. anthracis adaptation to the vertebrate host.

their cognate RR over other RRs encoded in the same genome [5]. Furthermore, if an HK is a bi-functional kinase/phosphatase, its phosphatase activity is specific for its cognate RR [6]. This buffers against non-specific phosphorylation of the cognate RR and limits the activation of the pathway [4]. TCS signaling purity is further maintained by a high ratio of RR:HK, which allows the cognate RR to outcompete other RRs for binding to its cognate HK [4,7]. Specificity may also be influenced by the temporal expression and spatial localization of TCSs [3].

Much focus has been placed on the maintenance of specificity at the HK-RR level and the general conclusion has been that cross-talk between non-cognate HK-RR pairs is possible, but rare. Most instances in which non-cognate HK-RR interactions have been described involve genetic manipulation of the system [6,8,9,10,11]. In all of these examples cross-talk is unresponsive to stimuli. The unresponsiveness to stimuli may be overcome by evolving the non-cognate HK-RR pair to increase phosphotransfer efficiency. In one example of this strategy, mutation of CpxA from Escherichia coli resulted in increased phosphotransfer to the non-cognate RR OmpR, as well as stimulii-responsive [12].

Only a few examples of natural cross-regulation, which are responsive to stimuli, have been reported. These include the integration of anaerobic nitrate and nitrite respiration in E. coli, PhoPR and YycGF interaction in Bacillus subtilis, BecRS and YycPQ in B. subtilis, ArcBA and EnvZ-OmpR in E. coli, and PmrAB and QseBC in E. coli [13,14,15,16,17]. In these instances, both HK-RR cross-phosphorylation and differential regulation of promoters by RRs are mechanisms by which signal integration is achieved.

Signal integration from multiple TCSs may be achieved at points in the signaling cascade other than at the HK-RR level. In fact, data suggest that TCSs naturally intersect at points in the signal transduction pathway other than at the HK-RR juncture. Even before an HK phosphorylates its cognate RR, interactions between HKS of individual TCSs can impact HK phosphorylation status and the downstream signaling cascade [18]. At the transcriptional level, multiple RRs may influence overlapping sets of genes [3]. This is typically manifested as two unique RRs regulating dissimilar, but overlapping DNA consensus sequences in the target promoter region [19,20,21,22]. Nonetheless, there is at least one example of cross-regulation in E. coli that demonstrates that homologous RRs may differentially target similar DNA consensus sequences [23]. Alternatively, a protein product regulated by a RR may itself be a transcriptional regulator or phosphatase that impinges on the regulatory outputs of other TCSs [3,24,25,26]. These examples of HK-RR and post-RR TCS interactions illustrate that bacterial signaling networks are not purely insulated linear systems that have a singular response to a particular input. Rather, interactions between TCSs at both the HK-RR and post-RR regulatory levels may allow bacteria to integrate multiple signals in order to flourish in diverse environments.

Bacillus anthracis is a Gram positive, spore-forming pathogen that is the causative agent of anthrax. The lifecycle of B. anthracis spans diverse environments. B. anthracis spores can be found in the soil; when the spores encounter a favorable environment, such as a mammalian host, they germinate and enter the vegetative part of the B. anthracis lifecycle. Reflective of the complex environmental conditions to which B. anthracis must sense and respond this pathogen encodes approximately 45 TCSs. Upon gaining entry to the blood, the concentration of B. anthracis can reach levels as high as 10^9 bacteria/ml [27]. This rapid proliferation presumably necessitates efficient nutrient utilization and metabolic adaptation.

Upon entering the blood, B. anthracis encounters an environment devoid of free iron, but replete with red blood cells (RBCs) which contain an abundant source of iron in the form of heme-bound hemoglobin. B. anthracis lyses RBCs and imports heme from the hemoglobin released into the microenvironment in order to satisfy its need for nutrient iron [28]. Paradoxically, too much heme is toxic to B. anthracis. One characteristic that enables B. anthracis to flourish in the blood is its ability to sense and detoxify heme. To adapt to the iron-poor, heme-rich environment B. anthracis detects heme using the heme sensing system (HssRS) TCS [29]. HssRS activates the expression of the heme regulatory transporter (hrtAB) which is thought to protect the bacteria from heme toxicity by exporting heme [29,30]. B. anthracis experiencing heme stress during infection has been observed in vivo as the hrt promoter (Pht) is activated in a murine model of cutaneous anthrax [29]. Notably, HssRS is conserved in many Gram positive bacteria suggesting that this is a common mechanism by which these organisms deal with heme toxicity [31].

Previously, a collection of small molecule activators of HssRS was identified in a high-throughput screen [32]. Here, we report that one of these compounds, VU0120925 (‘205), up-regulates Pht independent of HssRS. Exploiting this atypical stimulation of Pht by ‘205 led to the identification of a TCS that cross-talks with HssRS. This TCS is annotated as BAS1816-17 in the B. anthracis Sterne genome and due to the cross-talk between BAS1816-17 and HssRS observed here, we have named it HssRS interfacing TCS (HitRS). This TCS is comprised of the HitS HK and its cognate RR HitR, and targets a direct repeat (DR) in the hrtP promoter (Pht). Heme-stimulated HssRS cross-regulates Pht at both the HK-RR and RR-DR level, while HitRS cross-regulates Pht only at the RR-DR level. Since cross-talk is signal responsive and observed in parental B. anthracis Sterne, we assign the HssRS-HitRS interactions described here as cross-regulation. These results are summarized in Table 1, Table 2, and Figure 1. To define the native activity of HitRS, a phenotypic microarray resulted in the identification of five other small molecule activators of this TCS, many of which are implicated in cell envelope stress. This suggests that the bacterial responses to heme and cell
envelope stress are integrated and provides a platform for examining the intersection of heme toxicity with the integrity of the cell envelope. Altogether, this chemical biology approach to studying TCS biology has allowed us to dissect a complex signaling network in *B. anthracis* which may serve as a model for natural TCS cross-regulation during growth in diverse environments, including the vertebrate host.

**Results**

**VU0120205 (‘205) activates the Phrt direct repeat independent of HssR**

Previous work identified a series of small molecule activators of HssRS in *Staphylococcus aureus* [32]. These compounds were evaluated for HssRS stimulation in *B. anthracis* by quantifying the activation of the *hit* promoter (*Phrt*) fused to the *xylE* reporter gene (*phit*). From this collection, compound VU0120205 (‘205) was resynthesized and determined to have negligible activation of Phrt in the parental *B. anthracis* strain, Sterne; although, some activation may be masked by the high basal Phrt expression in this strain (Figure 2A and 2C) [29]. While all activation of Phrt by heme is lost in ΔhssRS, Phrt is strongly activated upon ‘205 exposure in the absence of the HssS histidine kinase (HK) and its cognate response regulator (RR) HssR (Figure 2A and 2B). Four conserved nucleotides in the Phrt direct repeat (DR) that are required for heme-induced activation of Phrt are also required for ‘205-mediated activation of Phrt (Figure 2A and 2D) [29]. These data suggest that another regulator in *B. anthracis* stimulates *hit* expression through the *hit* DR and that this regulator is activated by ‘205. This model is supported by the observation that there is a second DR in a distinct *B. anthracis* promoter that is closely related to the Phrt DR, but located in a different region of the chromosome [33]. Based on these observations, we hypothesized that the factor that recognizes this alternative DR may also target the Phrt DR and activate *hit* expression independent of HssR.

This alternative DR retains the four nucleotides required for HssR binding to Phrt, but differs from Phrt by four other nucleotides (Figure 2D) [29]. This DR variant is located in the predicted promoter region upstream of four previously uncharacterized genes including a TCS and an ABC transporter. Based on similarities between these DRs, and data described below, we have named these genes *hit*RS interfacing DR (*hit*RS) and *hit*RS interfacing transporter (*hit*PQ (Figure 2D)). In keeping with this, we will refer to the candidate promoter upstream of these genes as Phrt. HitP and HitQ are conserved in the *Bacilli* genus and are predicted to function together as an ABC transporter. The *hit*R and *hit*s genes encode a putative TCS in *B. anthracis* where HitS is the histidine kinase (HK) and HitR is the RR that are predicted to regulate Phrt [33]. Furthermore, BLAST analysis of HitR identifies HssR as the most similar RR in *B. anthracis* with 87% coverage and 52.5% identity between the two proteins. Consistent with this, HitS is most similar to HssS in *B. anthracis* with 81.8% coverage and 40.7% identity. Therefore, HssRS is the most closely related TCS to HitRS in *B. anthracis* and these systems likely recognize a similar DR. Taken together, these observations lead to the hypothesis that HitRS is a functional TCS that responds to ‘205 and regulates both Phrt and Phit.

**‘205-dependent activation of Phrt is mediated through HitRS**

‘205 activates Phrt in a manner independent of HssR but dependent on the Phrt DR. We hypothesized that this activation of Phrt is due to HitRS activity. Therefore, we generated ΔhitRS and ΔhssRSΔhitRS strains to determine if ‘205-dependent activation of Phrt is mediated through HitRS. The *phrt* reporter plasmid was transformed into these strains and the responsiveness of Phrt to ‘205 was quantified. The high basal activation of Phrt in ΔhitRS was greatly reduced in ΔhssRSΔhitRS (Figure 2A). These data reveal that the ‘205-induced activation of Phrt requires HitRS.

**B. anthracis** HssRS activates Phrt in response to heme treatment [29]. To determine if HitRS responds to heme and regulates the *hit* promoter, the activation of Phrt in ΔhitRS upon treatment with 2 μM heme was quantified and found to be similar to the parental strain (Figure 2B). This indicates that heme does not trigger HitRS to activate *hit* expression.

**HitRS targets the Phit DR**

To test the hypothesis that HitRS targets Phit in addition to Phrt, Phit was fused to the *xylE* reporter gene (*phit*) and transformed into the parental strain, ΔhssRS, ΔhitRS, and ΔhssRSΔhitRS. These strains were grown in the presence of vehicle or 50 μM ‘205 and then XylE activity was quantified. *Phit* is activated in the presence of ‘205 in a manner that requires HitRS, but not HssRS (Figure 3A).

The Phit DR contains the four conserved nucleotides previously shown to be required for heme-mediated activation of Phrt (Figure 2D) [29]. These four conserved nucleotides were mutated on the Phit reporter plasmid (*phitDR*) and the resulting construct was transformed into *B. anthracis* Sterne. Mutation of these four conserved nucleotides ablated ‘205-induced activation of Phit (Figure 3A). Together these data reveal that ‘205 activates HitRS, which regulates both Phit and Phrt in a manner that requires four conserved nucleotides in the DR of both of these promoters.

**HitRS is a two-component system that cross-phosphorylates with HssRS**

If HitRS functions as a prototypical TCS, HitS should phosphorylate HitR at a conserved aspartate. To test this hypothesis, the intracellular domain of HitS and the full length HitR were each N-terminally tagged with a hexa-histidine sequence, heterologously expressed in *E. coli*, and purified over a Ni-NTA column (Figure S1). The full length HitR predicted from the annotated *B. anthracis* Genome is highly insoluble when heterologously expressed. There are three candidate ATG translation start sites within the first 81 nucleotides of the predicted HitR open reading frame. When HitR was aligned with homologs from other Bacilli it became apparent that the third ATG is most likely the true translation start site. Expressing HitR with the N-terminus beginning at the third predicted translation start site and therefore lacking the first predicted 26 amino acids resulted in soluble RR (Figure S1B). The conserved residues predicted to be required for phosphorelay were identified in HitR and HitS. The conserved aspartate at amino acid position 56 in the receiver domain of HitR was mutated to an asparagine and the conserved histidine at amino acid position 137 in the dimerization and histidine phosphotransfer (DHp) domain of HitS was mutated to an alanine, and these variant forms of HitR and HitS were also purified (Figure S1). Since HitR and HitS share high sequence homology to HssR and HssS, we hypothesized that cross-talk between HitRS and HssRS could be mediated by non-cognate phosphate-transfer. To determine if TCS cross-phosphorylation occurs in *vivo*, HssR, HssS:H248A, HssR, and HssR:D54N proteins were also purified as described previously (Figure S1) [29].

First, the ability of the histidine kinases to autophosphorylate was determined by incubating the purified HKs in the presence of [γ-32P]-ATP and then resolving the proteins using SDS-PAGE.
Both HKs autophosphorylate in the presence of ATP and these activities require the conserved phosphate-accepting histidines (Figure 3B). Next the capacity of HsS and HsS to phosphotransfer to both HitR and HsS was investigated. HsS and HsR were first autophosphorylated and then co-incubated with each RR. The resulting phosphorylation state of each protein was visualized and the percent of phosphotransfer from each HK to each RR quantified (Figure 3C and 3D). Each HK preferentially phosphorylates its cognate RR, although there is significant cross-transfer to both HitR and HssR as the HK and HitR RR deletions (Figure S3). These observations suggest that the similarity between HssR and HitR may permit cross-phosphorylation by the HKs and that HitR functions as a TCS.

To further delineate the relative preference each HK has for both RRs, each RR was diluted over a concentration range of 2-logs. The HitR and HssR dilutions were then incubated with auto-phosphorylated HssS or HitS and the percent of phosphotransfer quantified (Figure S3). Plotting the percent of phosphotransfer against the RR concentration identifies a clear hierarchy in the preference for phosphorylation in vitro, such that the phosphotransfer of HitS to HitR is most efficient, followed by HssS to HssR, then HitS to HssR, and finally HssS to HitR (Figure S3). These results are consistent with previous reports that HKs preferentially phosphorylate their cognate RR over the non-cognate RR in vitro [3].

The direct repeat determines the response of each promoter region

The in vitro data presented here highlight the possibility that cross-signaling may occur due to phosphorylation of a non-cognate RR by a HK. However it is also possible that the similarity of Hit and HitR as well as the homology of HitR and HssR may permit cross-talk between these two TCSs at the RR-DRIV level. The four nucleotides of the HitR DR previously

---

**Table 1. Strains and plasmids.**

| Strain or Plasmid       | Description                                                                 | Source |
|-------------------------|-----------------------------------------------------------------------------|--------|
| Strains                 |                                                                             |        |
| _B. anthracis_          |                                                                             |        |
| Sterne                  | Non-encapsulated _B. anthracis_; parental strain for all mutants; obtained from Dr. Olaf Schneewind | [50]   |
| ΔhssRS                  | _hssRS_ TCS deletion                                                        |        |
| ΔhitRS                  | _hitRS_ (BAS1816-17) TCS deletion                                           |        |
| ΔhssRSΔhitRS            | _hssRS_ and _hitRS_ deletions                                               |        |
| ΔhssSΔhitR              | _hssS_ HK and hitR RR deletions                                             |        |
| ΔhssR                   | _hssSRR_ deletion                                                           |        |
| ΔhitR                   | _hssSRR_ and hitR RR deletions                                             |        |
| ΔhitSΔhssS              | _hitS_ HK and _hssS_ RR deletions                                           |        |
| ΔhitSΔhssS-hitRΔD56N    | _hitS_ HK and _hssS_ RR deletions; chromosomal _hitR_ containing D56N mutation |        |
| ΔhitR                   | _hitR_ RR deletion                                                          |        |
| Strains                 | _E. coli_                                                                   |        |
| DH5α                    | Strain used for molecular cloning                                           |        |
| Top10                   | Strain used for molecular cloning                                           |        |
| K1077                   | Methylase-deficient primary recipient for _B. anthracis_ DNA                | [51]   |
| BL21 (DE3)              | Recombinant protein expression                                              | Invitrogen |
| Plasmids                |                                                                             |        |
| phit                     | _Phit_ driven _XyIE_ reporter                                               |        |
| phtΔDR                  | _Phit_ driven _XyIE_ reporter with 4 conserved nucleotides mutated (Fig. 1D) |        |
| phitΔDR′                 | _Phit_ driven _XyIE_ reporter with 4 conserved nucleotides mutated (Fig. 1D) |        |
| pht-hitΔDR              | _Phit_ driven _XyIE_ reporter with 4 nucleotides unique to _Phit_ DR (Fig. 1D) |        |
| pht-hitΔDR′             | _Phit_ driven _XyIE_ reporter with 4 nucleotides unique to _Phit_ DR (Fig. 1D) |        |
| pET15b-hssR             | _HssR_ expression construct with N-terminal his(6)-tag                     |        |
| pET15b-hssRD54N         | _HssRD54N_ expression construct with N-terminal his(6)-tag                 |        |
| pET15b-hssS             | _HssS_ intracellular domain expression construct with N-terminal his(6)-tag |        |
| pET15b-hssSH248A         | _HssS_H248A_ intracellular domain expression construct with N-terminal his(6)-tag |        |
| pET15b-SacIΔ-hitR        | _HitR_ expression construct with N-terminal his(6)-tag; pET15b with SacI restriction site engineered into MCS |        |
| pET15b-SacIΔ-hitRD56N    | _HitRD56N_ expression construct with N-terminal his(6)-tag; pET15b with SacI restriction site engineered into MCS |        |
| pET15b-hitS              | _HitS_ intracellular domain expression construct with N-terminal his(6)-tag |        |
| pET15b-hitS-H137A        | _HitSH137A_ intracellular domain expression construct with N-terminal his(6)-tag |        |

**DOI:** 10.1371/journal.ppat.1004044.t001

---

Two-Component System Cross-Regulation

---

PLOS Pathogens | www.plospathogens.org 4 March 2014 | Volume 10 | Issue 3 | e1004044
shown to be required for \textit{HssR-P}\textit{hrt} interactions are conserved in the \textit{Phit} DR. However, four other nucleotides within the DR discriminate the \textit{Phit} DR from the \textit{Phrt} DR (Figure 2D). This suggests a model whereby the four nucleotides unique to \textit{Phit} are sufficient to differentially regulate \textit{Phit} as compared to \textit{Phrt}. To test this hypothesis, the four nucleotides unique to the \textit{Phit} DR were incorporated into the \textit{Phrt} reporter construct to make \textit{Phrt-hitDR} and the four nucleotides unique to the \textit{Phrt} DR were switched into the \textit{Phit} reporter construct to generate \textit{phrt-hitDR}. These constructs were transformed into \textit{B. anthracis} Sterne and the strains were exposed to either heme or '205. The response of each promoter to heme and '205 was quantified by measuring the XylE activity in each strain. As predicted, swapping the four unique nucleotides in each DR switches the responsiveness of each promoter to heme

\begin{table}[h]
\centering
\begin{tabular}{|l|c|c|c|c|c|c|c|c|c|}
\hline
\textbf{Strain} & \textbf{phrt} & \textbf{phit} & \textbf{phrtDR}\textsuperscript{*} & \textbf{phitDR}\textsuperscript{*} & \textbf{phrt-hitDR} & \textbf{phit-hrtDR} \\
\hline
\textit{B. anthracis} Sterne & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 \\
\textit{DhssRS} & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 \\
\textit{DhitRS} & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 & \textsuperscript{†} 0 \\
\textit{DhssS\_hitR} & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 \\
\textit{DhitS\_hitR} & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 \\
\textit{DhitS\_hitR\_hitR\_D56N} & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 & nd 0 \\
\hline
\end{tabular}
\caption{Summary of XylE reporter activities.}
\end{table}

See Table 1 for descriptions of strains and plasmids. \textsuperscript{†} = XylE reporter activity is elevated relative to vehicle. 0 = XylE reporter activity is similar to vehicle. Results reported are for \textit{B. anthracis} exposed to 1 \textmu M heme, 50 \textmu M '205, and 20 \textmu M NDGA, except results marked with an * denote exposure to 5 \textmu M heme. nd = not determined.
doi:10.1371/journal.ppat.1004044.t002

Figure 1. A summary of cross-talk observed between \textit{HssRS} and \textit{HitRS} in \textit{B. anthracis}. (A-C) \textit{HssRS} was previously shown to be stimulated by heme to activate \textit{Phrt} [29]. (D-F) Here, a second TCS \textit{HitRS} has been identified to be stimulated best by '205 and NDGA to activate \textit{Phit}. (G-H) At the RR-DR level cross-talk occurs as \textit{HitR} regulates \textit{Phrt} and \textit{HssR} regulates \textit{Phit}. (I) At the HK-RR level cross-talk occurs when \textit{HssS} cross-phosphorylates \textit{HitR}, this has been observed both in \textit{vitro} and in \textit{vivo}. (J) Cross-phosphorylation from \textit{HitS} to \textit{HssR} has only been observed in \textit{vitro}, so it is denoted with a dashed, gray line.
doi:10.1371/journal.ppat.1004044.g001
Two-Component System Cross-Regulation

and ‘205 (Figure 4). Therefore, these four unique nucleotides in the Phrt DR are sufficient to determine the response of the promoter to environmental stimuli.

It is also notable that while the four divergent nucleotides determine the response of the promoter to heme and ‘205, the high basal activation of Phrt is not transferred to Phrt when the Phrt DR was placed in Phrt. This result, when considered in conjunction with the observation that the high Phrt background is also lost upon hssR deletion and Phrt DR mutation (Figure 2A), suggests that the high basal activation of Phrt is due to the interaction of HssR with the Phrt DR in the context of Phrt.

Finally, the use of parental B. anthracis Sterne in this experiment enabled a concentration of heme to be administered (5 μM) which the ΔhssRS mutant cannot tolerate. This concentration of heme activates Phrt and Phrt-hitRDR in the parental strain. Since we did not mutate any TCSs in this B. anthracis strain, the activation of Phrt by heme indicates that cross-talk from HssRS to Phrt may occur naturally.

Mechanisms of HitRS cross-signaling to Phrt

Our observations that HitS phosphorylates HssR in vitro (Figure 3C and 3D) and HitRS regulates Phrt in ΔhssRS (Figure 2A) led us to hypothesize that cross-talk from HitRS to Phrt occurs in vivo at either the HK-RR or RR-DR level. To delineate cross-talk at the HK-RR level in vivo, a ΔhssSΔhitR strain was generated to eliminate any influence HssS or HitR may have on phosphotransfer from HitS to HssR. The ΔhssSΔhitR double knock-out strain was transformed with the phit reporter plasmid. If phosphotransfer from HitS to HssR occurs in vivo, then activation of Phrt would be observed in ΔhssSΔhitR following ‘205 stimulation of HitS; this is not the case (Figure 5A). Despite observing HitS to HssR cross-phosphorylation in vitro, phosphotransfer from HitS to HssR is not a source of cross-talk in vivo.

Next, the influence of HitR on the Phrt DR was examined to determine if cross-talk occurs at the RR-DR level. The Phrt reporter construct was transformed into the parental strain, ΔhssR, and ΔhssRΔhitR strains. As expected, when HitS was stimulated with ‘205, activation of Phrt was observed in ΔhssS but not in ΔhssSΔhitR (Figure 5A). These data signify that HitR can regulate the Phrt DR; however, this activity may be masked by the high basal activity of HssR in the parental strain. In summary, cross-talk from HitRS at Phrt in vivo is mediated by HitR regulation of Phrt, and not cross-phosphorylation of HssR by HitS.
Mechanisms of HssRS cross-signaling to Phit

Since Phit is activated in the presence of heme (Figure 4), we hypothesized that cross-talk from HssRS to Phit occurs at either the HK-RR or RR-DR level. The in vivo contribution of cross-phosphorylation from HssS to HitR was investigated by the same strategy used in the previous data section. A ΔhitSΔhssR double knockout strain was generated and transformed with the hitR reporter plasmid. If phosphotransfer from HssS to HitR occurs in vivo, then Phit is predicted to be activated in ΔhitSΔhssR in the presence of heme (Figure 5B). This does indeed occur. Furthermore, the activation of Phit upon heme treatment in ΔhitSΔhssR is absent when the HitR phosphate-accepting aspartate is mutated in this genetic background (Figure 5B). Together, these data indicate that HssS can cross-phosphorylate HitR to activate Phit in vivo.

Next, the ability of HssR to regulate Phit at the RR-DR level was probed. Using an analogous strategy, the Phit reporter construct was transformed into the parental strain, ΔhitR, and ΔhssRΔhitR. HssS was stimulated with 1 μM heme since ΔhssRΔhitR cannot tolerate higher concentrations of heme; no Phit activation was observed in the parental strain under these experimental conditions (Figure 5B). However, treating parental B. anthracis and ΔhitR with 5 μM heme resulted in Phit stimulation in both strains, implicating HssS-Phit interactions as a source of heme-dependent Phit activation (Figure 5C). These data reveal that in vivo HssS influences Phit upon heme exposure at both the RR-DR and HK-RR level associating HitPQRS with B. anthracis heme homeostasis.

The identification of HitRS activators

The cross-talk we have observed between HssRS and HitRS points to a natural function for HitRS that may be coordinated with the bacterial response to heme toxicity. Identifying the native signal of HitRS would not only inform the function of the putative HitPQ ABC transporter, but may also explain the physiological function of HssRS and HitRS cross-talk. Therefore we sought to identify other signals that activate HitRS by performing a Biolog Phenotypic Microarray.

Twenty 96 well plates (PM1-20) were purchased from Biolog’s Phenotypic Microarray Library for Microbial Cells. Each well contains a different compound that may differentially impact the growth and behavior of bacteria. Parental B. anthracis Sterne transformed with the Phit reporter plasmid was grown in each of these plates and activation of Phit was monitored by XylE assay. The screen was repeated twice and twelve molecules that activated Phit were repurchased from commercial vendors. The specific activation of HitRS was tested by exposing both parental B. anthracis Sterne and ΔhitRS, transformed with the Phit reporter plasmid, to each compound in a concentration range that spanned at least 2-logs and did not inhibit bacterial growth. This screen identified four compounds that specifically activate Phit in a manner that requires HitRS (Figure 6).

The compounds identified in the phenotypic microarray lead us to believe that the integrity of the cell envelope is involved in HitRS activation. The most potent activator identified is nordihydroguaiaretic acid (NDGA), which is more potent than nordihydroguaiaretic acid (NDGA), which is more potent than...
‘205 (Figure 6A). NDGA is an antioxidant isolated from the creosote bush with anti-inflammatory and antiviral activities [34]. The impact of NDGA on bacterial physiology is not well-understood, therefore we interrogated other HitRS activators in an effort to ascribe function to this system.

The second best activator of HitRS identified in the phenotypic microarray is chlorpromazine (Figure 6B). Chlorpromazine is used to treat a range of human pathologies ranging from allergies to psychoses [35]. Nonetheless some studies have probed the effect of chlorpromazine on bacteria. Most of this work has focused on the chemorepellent activity of chlorpromazine to B. subtilis, and one study notes that chlorpromazine inhibits cell wall biogenesis in Bacillus megaterium [36,37].

Both low concentrations of vancomycin and high concentrations of sodium phosphate (pH 7) were found to weakly activate HitRS (Figure 6D and 6E). The concentration range of vancomycin is limited due to the antibiotic nature of the compound. Vancomycin is best known for inhibiting Gram positive cell wall biosynthesis, but at low concentrations it has also been shown to disrupt membrane integrity [38]. The sodium phosphate salt only showed weak activity at high concentrations, beyond which significant growth defects were observed.

The description of the biological activity of each of these compounds in the literature led to the hypothesis that HitRS may be activated by cell envelope stressors. The detergent Tween 80 weakly activates HitRS at high concentrations (0.5–2.5%); however, other membrane disrupting compounds such as SDS, daptomycin, gramicidin, and Triton X-100 do not activate HitRS (data not shown). This suggests that general membrane stress does not activate HitRS. Recently, a small molecule inhibitor of S. aureus wall teichoic acid (WTA) synthesis called targocil was identified [39]. Targocil inhibits the transport of the nascent WTA from inside the membrane to the outer surface by TarGH in S. aureus. A microarray performed on targocil-treated S. aureus revealed that hitAB was the second most up-regulated operon next to cweA, a small peptide of unknown function that is a reporter of cell wall stress [40]. The up-regulation of hitAB in response to targocil treatment in S. aureus prompted the investigation of whether targocil activates HitRS in B. anthracis [41]. Targocil does specifically activate Phit in a manner that requires HitRS (Figure 6C).

In summary, five additional activators of HitRS have been identified, including one compound, NDGA, which is a more potent activator of HitRS than ‘205. The mechanism by which these molecules activate HitRS remains to be determined. Deciphering the common impact each of the HitRS activators has on B. anthracis physiology may uncover the native function of this new TCS and elucidate why HssRS and HitRS cross-talk, providing insight into the mechanisms of heme toxicity.

Cross-regulation between HssRS and HitRS

The cross-talk from HssRS to Phit occurs in response to heme in B. anthracis Sterne. This indicates that this interaction is not just cross-talk, but more specifically cross-regulation. We hypothesized that the HitRS-dependent activation of Phit is also an example of cross-regulation, despite only observing up-regulation in a genetically manipulated organism. To circumvent the need to use a B. anthracis mutant to test cross-talk from HitRS to Phit, we used the more potent HitRS activator NDGA, which was identified in the phenotypic microarray. When parental B. anthracis Sterne carrying the phit reporter plasmid was treated with NDGA, Phit is significantly up-regulated (Figure 7A). Activation of Phit in response to NDGA is not due to cross-phosphorylation from HitS to HssR, since Phit activation is lost in ΔhitDRΔhssS (Figure 7A). This result suggests that cross-regulation is mediated by HitR regulation of Phit. To address whether cross-talk between HssRS and HitRS occurs natively, RNA was purified from B. anthracis treated with heme, 200 μM NDGA, chlorpromazine, or targocil for 10 min. Quantitative PCR on the cDNA generated from these experimental growth conditions reveals that all tested activating molecules up-regulate both hitA and hitP, further solidifying that cross-regulation between HssRS and HitRS occurs naturally in vivo (Figure 7B).

Discussion

Here, a small molecule activator (‘205) of S. aureus HssRS was used to identify a previously uncharacterized TCS, HitRS, in B. anthracis and dissect the complex signaling interactions between HitRS and HssRS. The interactions between HssRS and HitRS, which we have documented here, are summarized in Table 1, Table 2 and Figure 1. HitRS recognizes a DR in Phit, while HssRS impacts activation at Phit (Figure 7A). DGAS hitsR is an example of a DR that operates in a manner that cross-regulates PhitDR, but the PhitDR is not sufficient to induce high basal activation of the Phit promoter. In total, the high basal activity of Phit requires the Phit DR, but the Phit DR is not sufficient to induce high basal activation of the Phit promoter. It implicates that other cis- or trans-acting cellular factors influence the high basal level of B. anthracis Phit activation.

Despite the high basal activity of Phit, it is clear that HitRS cross-regulates Phit upon NDGA treatment in B. anthracis Sterne (Figure 7A). Pairing genetics with chemical biology has revealed that HitRS cross-regulates Phit at the RR-DR level (Figure 5A), while HssRS impacts activation at Phit through HIK-RR and RR-DR interactions (Figure 5B and 5C). The observation that in vivo HitS does not cross-phosphorylate HssR, while HssS does cross-phosphorylate HitR may be explained by differential kinase and phosphatase activities by HitS and HssS on each of the RRs. Identifying the factors that influence cross-phosphorylation in vivo

Figure 4. The direct repeat in each promoter determines specificity in vivo. B. anthracis Sterne (WT) was transformed with the phit and phit xyIE reporter plasmids. Furthermore, the DR of each promoter was swapped into the other reporter plasmid (phit-hitDR and hit-hrtDR) and transformed into WT. Each strain was grown in triplicate and exposed to DMSO, 5 μM heme, or 50 μM ‘205 for 24 h and XyIE activity was quantified. Error bars represent ±2SD of data averaged from four independent experiments. Significance was calculated by an unpaired Student’s t-test, where ** = P<0.0001.

doi:10.1371/journal.ppat.1004044.g004
will further our understanding of mechanisms by which bacterial TCSs are insulated against cross-talk at the HK-RR level. Furthermore, it is important to recognize that the cross-talk between HitRS and HssRS occurs in parental *B. anthracis* Sterne in response to heme and NDGA, suggesting that the cross-talk described here occurs naturally and is an example of TCS cross-regulation. There are few documented instances of cross-regulation in bacterial signaling networks; therefore, using HitRS and HssRS as a model of TCS cross-regulation will expand our understanding of the mechanisms by which bacteria integrate TCS signaling pathways.

HitRS and HssRS are more closely related to one another than any of the 45 other TCSs in *B. anthracis*. Determining the native function of HitRS is essential for understanding if the cross-regulation between HssRS and HitRS occurs in parental *B. anthracis* Sterne in response to heme and NDGA, suggesting that the cross-talk described here occurs naturally and is an example of TCS cross-regulation. There are few documented instances of cross-regulation in bacterial signaling networks; therefore, using HitRS and HssRS as a model of TCS cross-regulation will expand our understanding of the mechanisms by which bacteria integrate TCS signaling pathways.

HitRS and HssRS are more closely related to one another than any of the 45 other TCSs in *B. anthracis*. Determining the native function of HitRS is essential for understanding if the cross-regulation between HssRS and HitRS occurs in parental *B. anthracis* Sterne in response to heme and NDGA, suggesting that the cross-talk described here occurs naturally and is an example of TCS cross-regulation. There are few documented instances of cross-regulation in bacterial signaling networks; therefore, using HitRS and HssRS as a model of TCS cross-regulation will expand our understanding of the mechanisms by which bacteria integrate TCS signaling pathways.

**Figure 5. In vivo cross-signaling between HssRS and HitRS.** (A) Cross-talk from HitRS to Phrt. *B. anthracis* Sterne (WT), ΔhssR, ΔhssRΔhitR, and ΔhssSΔhitR were transformed with the *xyle* reporter plasmid, phrt. Each strain was treated with either 50 μM 205 or an equivalent volume of vehicle (DMSO). (B) Cross-talk from HssRS to Phit at low heme concentrations. WT, ΔhitR, ΔhssRΔhitR, ΔhitSΔhssR, and ΔhssSΔhitRΔhitR:D56N were transformed with the *xyle* reporter plasmid, phit. Each strain was treated with either 1 μM heme or an equivalent volume of vehicle (0.1 M NaOH). (C) Cross-talk from HssRS to Phit. WT and ΔhitR phit were treated with either 5 μM heme or an equivalent volume of vehicle (0.1 M NaOH). In all instances, cultures were grown for 24 h and *xyle* activity was quantified. Error bars represent ±SEM of data averaged from two independent experiments performed with biological triplicates. Significance was calculated by an unpaired Student’s *t*-test, where ** = *P* ≤ 0.001 and *** = *P* ≤ 0.0001.

doi:10.1371/journal.ppat.1004044.g005

The intercalation of heme into membranes and the resulting disruption of the osmotic gradient is the mechanism by which free heme causes erythrocyte lysis [44,45,46]. The genetic architecture of the HssRS-HitRS regulatory network results in the up-regulation of HitRS by HssRS in response to heme toxicity. However, the analogous HitRS cross-regulation of Phrt does not increase expression of HssRS as HssRS is not under the control of Phrt. This differential expression of TCSs in signaling networks has been identified in other systems and allows bacteria to integrate signals from multiple pathways [24,25,26]. It is possible that the activation of HrtAB protects the bacteria from heme toxicity specifically, and HitPQ buffers *B. anthracis* against more general cell envelope stresses. The potential up-regulation of HitRS during heme stress supports the model that HitRS may sense more general cell envelope damage, which may also enable *B. anthracis* to better cope with heme-induced stress.

Alternatively, the cross-signaling between HssRS and HitRS may occur to supplement HssRS-mediated heme tolerance. In 1948, van Heyningen noted that *Bacillus* species exhibit differential sensitivity to heme [47]. Both *B. anthracis* and *B. cereus* are resistant to heme toxicity and encode HssRS-HitRS, while *B. subtilis* and *B. licheniformis* are sensitive to heme toxicity and do not encode HssRS-HitRS [data not shown][29,47]. Moreover, *B. anthracis* HssRS is more active than *S. aureus* HssRS and ΔhitA *B. anthracis* is more sensitive to heme toxicity than ΔhitA Δ*S. aureus* [29]. These data suggest that *B. anthracis* has differential sensitivities and responses to heme toxicity and it is possible that hitPQR contributes to *Bacillus* adaptation to heme stress under the conditions tested here.

More generally, the interactions between HssRS and HitRS suggest that *B. anthracis* may integrate multiple signals to better adapt and survive in diverse environments. The *Bacillus* have evolved complex signal transduction pathways, perhaps as a means to coordinate sporulation with signals from diverse niches spanning soil, water, plant surfaces, and various insect and mammalian hosts. The complexity of *Bacillus* signaling networks is highlighted by the reports of cross-regulation between the PhoPR and YycG TCSs as well as the many-to-one signaling cascade that integrates multiple HKs on a single RR to coordinate
Two-Component System Cross-Regulation

Figure 6. Five novel activators of HitRS. B. anthracis Sterne (WT) and ΔhitRS were transformed with the xylE reporter plasmid, phit. The strains were treated with either 50 μM 205 or a range of concentrations of the following compounds: (A) nordihydroguaiaretic acid (NDGA), (B) chlorpromazine, (C) targocil, (D) vancomycin, and (E) sodium phosphate (NaH₂PO₄) pH 7. Cultures were grown for 6 h and harvested. Cells were lysed and XylE activity was quantified. In all instances, error bars represent ±SEM of data averaged from two independent experiments performed with biological triplicates. Significance was calculated by an unpaired Student’s t-test, where * = P<0.01, ** = P<0.001, and *** = P<0.0001.
doi:10.1371/journal.ppat.1004044.g006

sporulation [15,48,49]. The capacity of the Bacilli to integrate signaling cascades may have facilitated the evolution of cross-regulation between HssRS and HitRS.

In summary, these studies have provided an array of insights into TCS biology and bacterial signaling networks. First, investigating the RR-DR interactions between two closely related TCSs reveals that HitR may recognize other DRs, but other cellular factors may insulate this cross-talk at the RR-DR level. For example, HitR may regulate HssR, but HssR does not seem to regulate HitR. At non-toxic concentrations of heme, HssR does not seem to regulate P₃hit. Conversely, P₃hit in the parental B. anthracis strain is activated only at higher concentrations of heme either through cross-phosphorylation from HssS to HitR or activation of Phit by HssR (Figure 5B and 5C). At non-toxic concentrations of heme, HssR does not seem to regulate Phit (Figure 5B), suggesting that other cis- or trans-acting factors are required to regulate HssR activity specifically at Phit and therefore limit HssR activation of the Phit DR at sub-toxic concentrations of heme. In E. coli nitrate and nitrite metabolism are coordinated by NarPQ and NarXL, which also employs differential regulation of similar DNA consensus sequences by two homologous RRs [3,23]. The determinants that drive differential regulation at shared consensus sequences are not well understood. Identifying factors that limit cross-regulation could reveal additional levels at which cross-talk is limited in bacterial signaling networks. At the HK-RR level, it is intriguing that the ‘least preferred’ cross-phosphorylation from HssS to HitR in vitro is observed in vivo, but the parallel cross-phosphorylation from HitS to HssR is not. Dissecting signaling determinants, such as HK phosphatase activity, which impact in vivo cross-phosphorylation will further add to the understanding of specificity determinants in TCS biology. The combination of chemical and genetic approaches employed here have enabled the dissection of a novel B. anthracis TCS signaling network, providing a unique, natural model for defining the mechanisms by which bacterial pathogens sense and respond to the complex host environment.

Materials and Methods

Bacterial strains and growth conditions

B. anthracis strain Sterne was used for all experiments [50]. Freezer stocks were streaked onto either LB- or BHI-agar plates and grown at 37°C for 24 h. Colonies from these plates were used to inoculate either LB or BHI and the cultures were grown with shaking at 180 rpm and 37°C; if the incubation was longer than 8 h the cultures were grown at 30°C, 180 rpm. All plasmid construction was performed in E. coli DH5α or TOP10. To move plasmids from E. coli to B. anthracis they were first transformed into E. coli K1077 [51]. Proteins were expressed in E. coli BL21DE3 pREL. Antibiotic concentrations used were ampicillin 100 μg/ml in E. coli (pET15b constructs), chloramphenicol 10 μg/ml in B. anthracis (XylE reporter assays) and 34 μg/ml in E. coli (protein expression), and kanamycin 20 μg/ml in B. anthracis and 40 μg/ml in E. coli (knockout generation).

Preparation of heme and other compound stocks

10 mM heme stocks (Fluka) were prepared daily in 0.1 M NaOH. A 50 mM stock of 205 was prepared in DMSO and stored at -20°C. A 20 mM stock of NDGA was prepared in ethanol and stored at -20°C. Stacks of 250 μM vancomycin and 100 mM chlorpromazine were prepared in water and stored at
−20°C. A 1 M stock of sodium phosphate monobasic was prepared in water, brought to a pH of 7, and stored at room temperature. A stock of 20 mg/ml of targocil was prepared in DMSO and stored in the dark at 4°C.

Genetic manipulation of B. anthracis

Electroporations and mutant generation were performed as previously described [29]. Electroporations were also performed with the following modifications. DNA purified for K1077 E. coli was eluted in 10 mM Tris-HCl, pH 8.5 and this solution was used for electroporations. After the B. anthracis competent cells were prepared, the optical density at 600 nm (OD_{600}) was determined in phosphate buffered saline (PBS, 137 mM NaCl, 2.7 mM KCl, Na_2HPO_4 10 mM, KH_2PO_4 1.8 mM, pH 7.4). Cells were flash frozen in liquid nitrogen and thawed on ice for electroporations. For electroporation the competent cells were diluted in SMG (0.5 M D-sorbitol, 0.5 M D-mannitol, 10% glycerol) to a final OD_{600} of 8.0. To 50 µl of cells 0.5 – 5 µl of DNA was added (approximately 3 µg total). Cells were incubated with DNA in a pre-chilled 1 mM cuvette for 1 min on ice and then electroporated using conditions described previously [29]. Cells were recovered in 1 ml BH +0.5 M D-sorbitol for 3 h at 30°C, 200 rpm in aeration tubes. All genomic lesions were generated without an antibiotic cassette. To generate an unmarked, deletion strain, 1,000 bp (including 100 bp of the target gene) from the 5’ and 3’ flanking regions of the gene targeted for deletion were cloned into pLM4. To generate a targeted lesion on the chromosome, 2,000 bp surrounding the site targeted for mutation were cloned into pLM4 and then lesions were introduced by inverse PCR. The pLM4 constructs were transformed into B. anthracis for mutagenesis as described previously [52].

Generation of xylE reporter plasmids

Previously, hit promoter activity was monitored using a pOS1 based xylE reporter gene system [29]. To maintain continuity with previous studies, the hit promoter xylE reporter construct was also generated in a pOS1 background [53]. Notably phit, phitDR, phit-hitDR, and phit-hitIDR did not have high basal activity, suggesting that the high basal activity of phit is not due to the copy number of pOS1. The hit promoter-xylE fusions was generated by PCR-SOE [54]. This resulted in a seamless fusion between the hit promoter and the xylE reporter gene. PCR-SOE DNA was digested with BamHI and EcoRI and ligated into pOS1. Mutagenesis of the hit DR was carried out by inverse PCR-based mutagenesis. The selected mutations were incorporated on one primer and a second non-overlapping, non-mutagenic primer was designed to amplify in the opposite direction. The primers were 5’ phosphorylated by T4 polynucleotide kinase (NEB) and the parental plasmid amplified using the high-fidelity DNA polymerase Pfu Turbo. The amplified DNA was blunt end ligated and transformed into E. coli. Mutagenized constructs were verified by DNA sequencing. XylE assays - XylE assays were performed as described previously with the following exceptions [29]. All cultures were grown overnight in 5 ml LB/chloramphenicol at 30°C, 180 rpm. Bacteria were sub-cultured into 4 ml LB/chloramphenicol and grown as indicated: either 1:100 sub-culture and growth for 6 h at 37°C, 180 rpm or 1:1,000 sub-culture and growth for 24 h at 37°C, 180 rpm.

Generation of protein purification constructs

The hitR gene was amplified from genomic DNA by Pfu Turbo using the third ATG as the start site. Inverse PCR was used to insert a SacII site in the multiple cloning site (MCS). Briefly, two complementary, overlapping primers that amplify in the opposite direction were designed, which included the addition of the SacII site after the BamHI restriction. The PCR product was transformed into E. coli and successful incorporation of the SacII site was confirmed by sequencing. This inverse PCR generated pET15b_SacII and resulted in the MCS changing from: CATATGCGTGGATCCCGGTGCT to CATATGCGTGGATCCCGGTGCT. The hitR PCR product was digested with SacII and BamHI-HF (NEB) and ligated into pET15b_SacII. The intracellular domain (ICD) (bp 221 to the end) of hitS was amplified using Pfu Turbo and digested with SmaI. The vector pET15b was digested with XhoI and purified using the Qiagen gel purification kit. Blunt ends were generated on pET15b by treating the cut vector with DNA Polymerase I, Klenow fragment (NEB) and subsequently ligated to the digested hitS ICD.

Figure 7. HssRS and HitRS cross-regulate. (A) Cross-regulation of phit by HitRS. B. anthracis Sterne and ΔhitR hitSS were transformed with the xylE reporter plasmid, phit. The strains were treated with either 20 µM NDGA or vehicle (ethanol). Cultures were grown for 24 h and harvested. Cells were lysed and XylE activity was quantified. Error bars represent ±SEM of data averaged from three independent experiments performed with biological triplicates. The significance was calculated by an unpaired Student’s t-test, where *** = P<0.0001. (B) WT was grown to mid-log and then lesions were introduced by inverse PCR. The pLM4 constructs were transformed into B. anthracis and then lesions were introduced by inverse PCR. The pLM4 constructs were transformed into B. anthracis for mutagenesis as described previously [52].
The mutation of HitR D56N and HitS ICD H137A were generated by inverse PCR as described above. All constructs result in the generation of N-terminal hexahistidine-tagged proteins and were transformed into BL21 DE3 pRLE.

Protein purification

Protein purifications were carried out as described previously with the following modifications [29]. All proteins were induced in 0.5 mM IPTG, cells were lysed for 5 min on an EmulsiFlex C3 (Avestin) at a peak PSI of 20,000 and the lysates were centrifuged at 20,000 x g for 1 h at 4°C. HitS ICD and HssS ICD and their corresponding mutants were purified in TBS (50 mM Tris, pH 8.0, 300 mM KCl) containing 10 mM imidazole and eluted in 75 mM imidazole in TBS, HitR and HssR, and their corresponding mutants were purified in TBS containing 1 mM EDTA, 10% glycerol, and 10 mM imidazole and eluted in 50 mM imidazole in TBS. HitR was induced after 2 h of growth at 16°C in 0.5 mM IPTG and a second time 5 h later with an equal volume of IPTG to bring the final concentration to 1 mM.

Autophosphorylation and phosphotransfer assays

Autophosphorylation assays were performed as described previously with the following modifications [29]. Ten μl autophosphorylation reactions were assembled with HssS and HitS ICDs and their corresponding mutants at a final concentration of approximately 10 μM (640 μg/ml). For HssS 10 μCi of [γ-32P]-ATP was used and due to the high autophosphorylation of HitS, 5 μCi of [γ-32P]-ATP was used. All reactions were incubated at 37°C for 30 min. To analyze the phosphotransfer of each HK to each RR, the HKs were autophosphorylated as described above in a 30 μl reaction volume. After the HKs autophosphorylated for 30 min, 10 μl were sampled into 2X SDS-PAGE loading buffer. To the remaining 20 μl, 3 μl of 100 μM RR was added for a final concentration of approximately 10 μM. HK-RR mixtures were incubated for 5 min and then 10 μl was sampled into 2X SDS-PAGE loading buffer. Samples were analyzed as described previously [29]. To examine the preferential phosphotransfer of each HK for each RR, a 220 μl autophosphorylation reaction was run for each HK. Serial dilutions of each RR were prepared so each HK for each RR, a 220 μl autophosphorylation reaction was assembled with HssS and HitS ICDs and their corresponding mutants at a final concentration of approximately 10 μM (640 μg/ml). For HssS 10 μCi of [γ-32P]-ATP was used and due to the high autophosphorylation of HitS, 5 μCi of [γ-32P]-ATP was used. All reactions were incubated at 37°C for 30 min. To analyze the phosphotransfer of each HK to each RR, the HKs were autophosphorylated as described above in a 30 μl reaction volume. After the HKs autophosphorylated for 30 min, 10 μl were sampled into 2X SDS-PAGE loading buffer. To the remaining 20 μl, 3 μl of 100 μM RR was added for a final concentration of approximately 10 μM. HK-RR mixtures were incubated for 5 min and then 10 μl was sampled into 2X SDS-PAGE loading buffer. Samples were analyzed as described previously [29]. To examine the preferential phosphotransfer of each HK for each RR, a 220 μl autophosphorylation reaction was run for each HK. Serial dilutions of each RR were prepared so when 2 μl of the RR stock was added to 9.5 μl of phosphorylated HK, the final RR concentration was 20, 10, 5, 2.5, 1.25, 0.625, 0.31, 0.16, 0.08, and 0.04 μM. Each RR concentration was incubated with each phosphorylated HK for 30 sec, after which 10 μl of 2X SDS-PAGE buffer was added. The samples were stored on ice and analyzed as described previously [29].

Biolig Phenotypic Microarray screen

Phenotypic Microarray plates (PM 1-20) and IF-0a GN/GP (1.2X) were purchased from Biolog. Preparation of PM inoculating fluids was adapted from the manufacturer’s recommendations for _B. subtilis_. The following solutions were prepared: PM-A = 800 mM tricarballylic acid, pH 7.1; PM-B = 240 mM MgCl2 and 120 mM CaCl2; PM-C = 5 mM L-arginine and 6 mM L-glutamic acid; PM-D = 0.5 mM L-cystine pH 8.5 and 1 mM 5-UMP; PM-E = 0.6% tween-80; PM-F = 300 mM D-glucose and 600 mM pyruvate. In all instances pH was adjusted with NaOH and all solutions were filter sterilized and stored at 4°C. 12X PM additive for PM-1,2,5,7,10,15 ml each of PM-B, PM-C, PM-E, PM-F, and 15 ml each of PM-D and sterile water. 12X PM additive for PM-3,6,8,9,11,12,14,16,17,18,19,20 ml each of PM-A and PM-D. 12X PM additive for PM-4,5,13 ml each of PM-B, PM-C, PM-E, PM-F, PM-G, and 15 ml of PM-A and 10 ml of sterile water. 12X PM additive for PM-5,8 = 5 ml each of PM-B and PM-G and 15 ml of PM-A and 25 ml of sterile water. 12X PM additive for PM-9,10 = 5 ml each of PM-B, PM-E, PM-F, PM-G and 30 ml of sterile water. The inoculating solution for each plate consisted of 20 ml of IF-0A GN/GP (1.2X), 2 ml of 12X PM additive (specific for each plate, described above), and 20 μl of 10 mg/ml chloramphenicol. _B. anthracis_ PM-2 was grown overnight on solid media and the colonies were inoculated into the inoculating solution to a final OD600 of approximately 0.6. The inoculating solution with the bacterial suspension was aliquoted 100 μl into each well of a 96 well PM plate. A single well was also inoculated with 1.67 μl of 50 mM 205 as a positive control; the well was selected based on chemical redundancy with other wells on the same plate to avoid masking a potential hit. Plates were covered with a Kimwipe to allow oxygen exchange and incubated at 37°C, 180 rpm. At 3 h and 5 h, wells with significant liquid loss were supplemented with 50 μl of sterile water. After 6–7 h, each PM plate was centrifuged at 3,200 x g for 10 min. The supernatant was removed by pipetting and the plate frozen at −80°C. To analyze the XylE activity, the PM plates were thawed and cells were lysed by the addition of 50 μl of 2 mg/ml lysozyme in 100 mM potassium phosphate buffer (pH 8.0), 10% (v/v) acetone. Cells were incubated at 37°C for 30 min. To each well 25 μl of 100 mM potassium phosphate (pH 8.0), 0.2 mM pyrocatechol was added immediately before assaying XylE activity [56]. Each PM plate was screened twice, using a different well for a 205 positive control each time.

RNA purification

_B. anthracis_ Sterne was grown overnight in 5 ml of LB at 30°C, 180 rpm. The overnight cultures were sub-cultured 1:100 into 5 ml of LB for 5 h at 37°C, 180 rpm. 5 μl of each compound prepared at 1000× concentration in DMSO was added to each culture and incubated for 10 min at 37°C, 180 rpm. To preserve the cultures 5 ml of ice cold acetonitrile:ethanol was added to each sample and immediately stored at −70°C.

The RNA was purified using a Qiagen RNeasy Prep kit according to the RNeasy lipid tissue directions. Briefly, samples were thawed on ice and then pelleted at 3,200 x g for 10 min. The supernatants were removed and the pellets dried for 10 min on ice. Each pellet was resuspended in 1 ml of Trizol and transferred to a bead beater tube containing Lysing Matrix B (MP). Cells were lysed on a FastPrep-24 bead beater for 45 s at 6 m/s. Samples were cooled on ice and then mixed with 200 μl chloroform. Samples were pelleted at 16,100 x g for 15 min at 4°C. The upper aqueous layer was transferred to a new tube and then RNA was isolated according to the Qiagen RNeasy lipid tissue directions and eluted twice, each time in 50 μl of water.

DNA contamination was removed by adding 8 μl of RQ1 DNase (Promega), 12 μl 10X RQ1 DNase buffer, and 2 ul of RNase inhibitor (Promega) to each sample and incubating at 37°C for 2 h. DNase was removed using the Qiagen RNeasy mini prep RNA clean-up protocol and eluted twice, each time in 50 μl of water.

qRT-PCR

cDNA was generated by adding 4 μl of Random Hexamers (Promega) to 2 μg of RNA template in 28 μl of water. The primers were annealed to the template for 5 min at 70°C and then placed on ice. To 14 μl of primer-annealed template was added 5 μl of 5X M-MLV reverse transcriptase buffer, 1.25 μl of dNTPs, 1.25 ul of RNase inhibitor, and either 1.5 μl of M-MLV reverse transcriptase (Promega) or water. cDNA was generated at 37°C for 1 h and then diluted 1:50 in DNase-/RNase-free water and stored at −20°C. qRT-PCR was performed in 12.5 μl final
volume using SYBR Green Supermix, 16S RNA as a normalizing control, and 57°C as an annealing temperature. Primer sets used are in Table S1.

205 synthesis

All reagents and starting materials were purchased from Sigma Aldrich with the exception of disulfonyl chloride (I) which was purchased from ChemCollect (Remscheid, Germany). Dichloromethane was used as received in a Sure/Seal bottle. Triethylamine was stored over potassium hydroxide. All glassware was flame dried prior to use. Column chromatography was conducted on an Isco Combiflash automated purification system. Melting points were determined using a Stanford Research Systems OptiMelt system. NMR spectra were recorded on a 400 MHz Bruker spectrometer. All chemical shifts were reported relative to residual solvent peaks. LC/MS was conducted and recorded on an Agilent Technologies 6130 Quadrupole instrument.

N2, N7-di-p-tolyl-9H-fluorene-2, 7-disulfonamide (VU0120205). I was added to a solution of 4-toluidine (307 mg, 7.53 mmol) and triethylamine (1.05 mL, 7.53 mmol) in 10 ml dichloromethane at 0°C with stirring (Figure S4). The solution was allowed to warm to room temperature (23°C) and maintained under an atmosphere of argon (5 h). The reaction was concentrated and the residue purified by column chromatography (Hexanes/EtOAc, 100:0 – 0:100 gradient) which gave 500 mg (40%) of VU0120205 as a yellow powder: mp 234–238°C; 1H-NMR (400 MHz, (CD)2CO) δ 8.37 (s, NH), 8.10 (d, J = 8.0, 2H), 7.98 (s, 2H), 7.81 (d, J = 8.4, 2H), 7.10 (d, J = 8.4, 4H), 7.03 (d, J = 8.4, 4H), 4.00 (s, 2H), 2.20 (s, 6H); 13C-NMR (100 MHz, (CD)2CO) δ 145.90, 144.87, 140.49, 136.17, 136.53, 130.59, 127.38, 125.11, 122.5, 122.18, 37.69, 20.84; LRMS m/z calculated for C27H25N2O4S2 [MH]+ 505.1, found 505.1.

Statistics

Where indicated, Student’s t-tests were calculated using either Excel 2007 or GraphPad Prism 5.

Supporting Information

Figure S1 Purified HitR, HitS, HssR, and HssS. (A-B) Each histidine kinase (A) and response regulator (B) was expressed in E. coli and purified by Ni-affinity purification. The dialyzed elutions were run on an SDS-PAGE gel to assess purity. (TIF)

Figure S2 HssS and HitS do not cross-phosphorylate QseB. Each HK was auto-phosphorylated (0.5 μM final concentration) and then mixed with recombinant QseB (1 μM final concentration). The reactions were incubated at 37°C and sampled for SDS-PAGE analysis at 10 s and 10 min. (TIF)

Figure S3 Each histidine kinase preferentially phosphorylates its cognate response regulator in vitro. (A-B) A serial dilution of RR concentrations ranging from 20 μM to 39 nM was prepared. Each HK was autophosphorylated and then mixed with each RR concentration for 30 sec. Phosphotransfer reactions were quenched with SDS-PAGE loading buffer and resolved on a gel. A representative image is shown in (A). Data from at least four replicates were average and plotted in (B). Error bars represent ± SD. (TIF)

Figure S4 Schematic of VU0120205 synthesis. Disulfonyl chloride (I) was added to a solution of 4-toluidine and triethylamine in dichloromethane at 0°C with stirring. The solution was allowed to warm to room temperature and maintained under an atmosphere of argon. The reaction was concentrated and the residue purified by column chromatography which produced VU0120205 as a yellow powder. (TIF)

Acknowledgments

We thank Dr. Maria Hadjifrangiskou for sharing purified QseB and critical reading of the manuscript, Dr. Suzanne Walker for sharing targecl, and Dr. Nicholas Bergman and Dr. Karla Passalacqua for their work on the preliminary B. anthracis microarrays. We acknowledge Amy Tesenrieder for the preparation of VU0120205.

Author Contributions

Conceived and designed the experiments: LAM JEC PRB LQO DLS. Performed the experiments: LAM JEC PRB LQO DLS. Analyzed the data: LAM JEC PRB LQO DLS. Contributed reagents/materials/analysis tools: BFD SJIII CMG GAS. Wrote the paper: LAM EPS.

References

1. Stock AM, Robinson VL, Gouldeau PN (2000) Two-component signal transduction. Annu Rev Chem 69: 183–215.
2. Skerker JM, Prichuk BS, Stryaporn A, Labin EA, Ashenberg O, et al. (2008) Rewiring the specificity of two-component signal transduction systems. Cell 133: 1043–1054.
3. Laub MT, Goulani M (2007) Specificity in two-component signal transduction pathways. Annu Rev Genet 41: 121–145.
4. Podgornaia AI, Laub MT (2013) Determinants of specificity in two-component signal transduction.Curr Opin Microbiol 16: 156–162.
5. Skerker JM, Prasol MS, Perchuk BS, Biondi EG, Laub MT (2005) Two-component signal transduction pathways regulating growth and cell cycle progression in a bacterium: A system-level analysis. PLoS Biol 3: e334.
6. Groban ES, Clarke EJ, Salis HM, Miller SM, Voigt CA (2009) Kinetic buffering of cross talk between bacterial two-component sensors. Journal of Molecular Biology 390: 380–393.
7. Cai SJ, Inouye M (2002) EnvZ-OmpR interaction and osmoregulation in Bacillus subtilis. Mol Microbiol 70: 494–506.
8. Stryaporn A, Perchuk BS, Laub MT, Goulani M (2010) Evolving a robust signal transduction pathway from weak cross-talk. Mol Syst Biol 6: 452.
9. Matsubara M, Kitaoka SI, Takeda SI, Mizuno T (2000) Tuning of the porin expression under anaerobic growth conditions by His-to-Asp cross-phosphorelay through both the EnvZ-oosensor and ArcB-anosensor in Escherichia coli. Genes Cells 5: 555–569.
10. Stewart V (2003) Biochemical Society Special Lecture. Nitrate- and nitrite-responsive sensors NarX and NarQ of proteobacteria. Biochem Soc Trans 31: 1–10.
11. Silvia JC, Haldimann A, Prahalad MK, Walsh CT, Wanner BL (1998) In vivo characterization of the type A and B vancomycin-resistant enterococci (VRE) VanRS two-component systems in Escherichia coli: a non-pathogenic model for studying the VRE signal transduction pathways. Proc Natl Acad Sci U S A 95: 11951–11956.
12. Siryaporn A, Prahalad MK, Walsh CT, Wanner BL (1998) Cross-talk suppression between the CpxA-CpxR and EnvZ-OmpR two-component systems in Escherichia coli. Mol Microbiol 70: 494–506.
13. Matsubara M, Kitaoka SI, Takeda SI, Mizuno T (2000) Tuning of the porin expression under anaerobic growth conditions by His-to-Asp cross-phosphorelay through both the EnvZ-oosensor and ArcB-anosensor in Escherichia coli. Genes Cells 5: 555–569.
14. Stewart V (2003) Biochemical Society Special Lecture. Nitrate- and nitrite-responsive sensors NarX and NarQ of proteobacteria. Biochem Soc Trans 31: 1–10.
15. Howell A, Dubrac S, Noone D, Varaghezei KI, Devine K (2006) Interactions between the YycFG and PhoPR two-component systems in Bacillus subtilis: the PhoR kinase phosphorylates the non-cognate YyfF response regulator upon phosphate limitation. Mol Microbiol 59: 1199–1215.
16. Guckes KR, Kostakioti M, Breland JG, Go AP, Shaffer CL, et al. (2013) Strong cross-system interactions drive the activation of the QseB response regulator in the absence of its cognate sensor. Proc Natl Acad Sci U S A 110: 16592–16597.
17. Rietkotter E, Hoyer D, Mascher T (2008) Bacitracin sensing in Bacillus subtilis. Mol Microbiol 67: 768–785.
18. Goodman AL, Merighi M, Hyedo M, Venter I, Filloux A, et al. (2009) Direct interaction between sensor kinase proteins mediates acute and chronic disease. Proc Natl Acad Sci U S A 106: 162–167.
19. Jubelin G, Vinayae A, Belouc C, Ghigo JM, Lazaroni JC, et al. (2005) CpxR/ompR interplay regulates curli gene expression in response to osmolarity in Escherichia coli. J Bacteriol 187: 2038–2049.
20. Batchelor E, Walthers D, Kenney LJ, Goulian M (2005) The Escherichia coli CpxA-CpxR envelope stress response system regulates expression of the porinsompF and ompG. J Bacteriol 187: 5723–5731.
21. Delgado MA, Moulim G, Grosman EA (2006) The PmrA/PmrB and RcsC/RcsB systems control expression of the Salmo nella O-antigen chain length determinant. Mol Microbiol 56: 39–50.
22. Moulim G, Grosman EA (2005) Control of the Salmonella sfgl gene by three two-component regulatory systems. Mol Microbiol 47: 335–344.
23. Darwin AJ, Tyler KL, Busby SJ, Stewart V (1997) Differential regulation by the homodimeric response regulators NarL and NarP of Escherichia coli K-12 depends on DNA binding site arrangement. J Biol Chem 272: 283–289.
24. Gun J, Miller SL (1996) PhoP-PhoQ activates transcription of pmrAB, encoding a two-component regulatory system involved in Salmonella typhimurium antimicrobial peptide resistance. J Bacteriol 178: 6857–6864.
25. Vianney A, Jubelin G, Renaud S, Dorel C, Lejeune P, et al. (2005) Escherichia coli tol and sci genes participate in the complex network affecting curli synthesis. Microbiology 151: 2467–2479.
26. Birkey SM, Liu W, Zhang X, Duggan MF, Hulett FM (1998) Pho signal transduction network reveals direct transcriptional regulation of one two-component system by another two-component regulator: Bacillus subtilis PhoP directly regulates production of RvD. Mol Microbiol 30: 943–953.
27. Telford 1, Zornetta I (2012) Bacillus anthracis factors for phagosomal escape. Toxins 4: 536–553.
28. Honsa ES, Maresso AW (2011) Mechanisms of iron import in anthrax. Biometals 24: 535–545.
29. Batchelor E, Walthers D, Kenney LJ, Goulian M (2005) The Escherichia coli CpxA-CpxR envelope stress response system regulates expression of the porinsompF and ompG. J Bacteriol 187: 5723–5731.
30. Darwin AJ, Tyler KL, Busby SJ, Stewart V (1997) Differential regulation by the homodimeric response regulators NarL and NarP of Escherichia coli K-12 depends on DNA binding site arrangement. J Biol Chem 272: 283–289.
31. Anzaldi LL, Skaar EP (2010) Overcoming the heme paradox: Heme toxicity and prevention of a transcriptional heme stress response. J Bacteriol 190: 3588–3596.
32. Stauf DL, Bagaley D, Torres VJ, Joyce R, Anderson KL, et al. (2008) Staphylococcus aureus HtrA is an ATPasese required for heme toxicity and prevention of a transcriptional heme stress response. J Bacteriol 190: 3588–3596.
33. Stauf DL, Torres VJ, Skaar EP (2007) Signaling and DNA-binding activities of the Staphylococcus aureus HtrA-HsH two-component system required for heme sensing. J Biol Chem 282: 26111–26121.