Repression by the H-NS/YmoA histone-like protein complex enables IscR dependent regulation of the *Yersinia* T3SS

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Abstract

The type III secretion system (T3SS) is an appendage used by many bacterial pathogens, such as pathogenic *Yersinia*, to subvert host defenses. However, because the T3SS is energetically costly and immunogenic, it must be tightly regulated in response to environmental cues to enable survival in the host. Here we show that expression of the *Yersinia* Ysc T3SS master regulator, LcrF, is orchestrated by the opposing activities of the repressive H-NS/YmoA histone-like protein complex and induction by the iron and oxygen-regulated IscR transcription factor. While deletion of *iscR* or *ymoA* has been shown to decrease and increase LcrF expression and type III secretion, respectively, the role of H-NS in this system has not been definitively established because *hns* is an essential gene in *Yersinia*. Using CRISPRi knockdown of *hns*, we show that *hns* depletion causes derepression of *lcrF*. Furthermore, we find that while YmoA is dispensable for H-NS binding to the *lcrF* promoter, YmoA binding to H-NS is important for H-NS repression activity. We bioinformatically identified three H-NS binding regions within the *lcrF* promoter and demonstrate binding of H-NS to these sites in vivo using chromatin immunoprecipitation. Using promoter truncation and binding site mutation analysis, we show that two of these H-NS binding regions are important for H-NS/YmoA-mediated repression of the *lcrF* promoter. Surprisingly, we find that IscR is dispensable for *lcrF* transcription in the absence of H-NS/YmoA. Indeed, IscR-dependent regulation of LcrF and type III secretion in response to changes in oxygen, such as those *Yersinia* is predicted to experience during host infection, only occurs in the presence of an H-NS/YmoA complex. These data suggest that, in the presence of host tissue cues that drive sufficient IscR expression, IscR can act as a roadblock to H-NS/YmoA-dependent repression of RNA polymerase at the *lcrF* promoter to turn on T3SS expression.
Author summary

In facultative pathogens, horizontally-acquired virulence factors must be integrated into existing regulatory networks that respond to environmental cues. The Ysc type III secretion system (T3SS) is encoded on a virulence plasmid carried by the plague agent *Yersinia pestis* and the closely related enteropathogen *Yersinia pseudotuberculosis*. H-NS is a histone-like protein suggested to repress horizontally-transferred virulence genes. Here we show that in *Y. pseudotuberculosis*, H-NS and its binding partner YmoA facilitate oxygen-dependent regulation of the Ysc T3SS by the transcription factor IscR. As oxygen availability is low in the intestine where the T3SS is not required for *Yersinia* colonization, and higher in extraintestinal tissue where the T3SS is critical for virulence, these data suggest that the Ysc T3SS was integrated into the pre-existing H-NS and IscR regulatory networks to optimize bacterial pathogenesis.

Introduction

Virulence factors are critical components that allow pathogens to establish or sustain infections within a given host. One common bacterial virulence factor is a needle-like apparatus, known as the type III secretion system (T3SS) [1,2]. Enteropathogenic *Yersinia pseudotuberculosis* is one of three human pathogenic *Yersinia* spp. that use the T3SS to inject effector proteins into host cells that dampen host immune responses, facilitating extracellular growth [3–6]. Members of human pathogenic *Yersinia* spp. include *Yersinia pestis*, the causative agent of plague, and the enteropathogens *Yersinia enterocolitica* and *Yersinia pseudotuberculosis*. While the T3SS is critical for infection, this apparatus appears to be metabolically burdensome since T3SS activity leads to growth arrest [7,8]. In addition, the Ysc T3SS is associated with pathogen-associated molecular patterns (PAMPs) recognized by several innate immune receptors, and some of these T3SS-associated PAMPS have evolved under selective evolutionary pressure by the ensuing immune response [5,9]. Without tight regulation of T3SS expression and deployment, these metabolic and immunological burdens would decrease the chance of *Yersinia* survival in the host.

The Ysc T3SS is encoded on a 70 kb plasmid for *Yersinia* Virulence, known as pYV or pCD1 [10]. Transcriptional regulation of T3SS genes is maintained by a master regulator called LcrF/VirF [11–14]. LcrF itself is also encoded on pYV, within the YscW-lcrF operon, and is highly conserved among all three human pathogenic *Yersinia* spp. LcrF is part of a larger family of AraC-like transcriptional regulators, and orthologs exist in other T3SS-encoding pathogens, such as ExsA in the nosocomial pathogen *Pseudomonas aeruginosa* [15]. The YscW-lcrF operon is regulated by various mechanisms in response to different environmental stimuli, including temperature, oxygen, and iron availability [16,17]. For example, an RNA thermometer blocks the ribosome binding site of lcrF at room temperature, but melts at mammalian body temperature, allowing lcrF translation [16].

Temperature-dependent transcriptional control of yscW-lcrF has been previously linked to the Histone-like Nucleoid structuring protein, H-NS, and an H-NS-binding protein called YmoA (*Yersinia* modulator) [16]. H-NS contains an N-terminal oligomerization domain and a C-terminal DNA minor-groove binding domain separated by a flexible linker [18,19]. H-NS preferentially binds AT rich regions of DNA [19,20]. Once H-NS binds a high-affinity site, H-NS oligomerizes on the DNA [21,22]. H-NS oligomers can either form a nucleoprotein filament on a contiguous stretch of DNA, or H-NS can form DNA bridges when multiple discrete H-NS binding regions are brought together, either way leading to transcriptional silencing of
that particular gene [23]. Interestingly, H-NS in multiple bacterial pathogens has been shown to silence certain gene targets during growth outside of the mammalian host (20–30°C), but fails to silence these targets to the same magnitude when exposed to mammalian body temperature (37°C) [24–26]. This suggests H-NS may play a role in repressing virulence factors outside host organisms in facultative pathogens. However, H-NS is an essential gene in pathogenic Yersinia [27–30], making it challenging to definitively test the role of H-NS in regulating gene expression in these organisms.

YmoA, an E. coli Hha (“high hemolysin activity”) ortholog, has been suggested to modulate H-NS repression of a subset of promoters and deletion of ymoA in Yersinia leads to changes in gene expression of putative H-NS targets [16,31–33]. YmoA and Hha lack a DNA binding domain; instead, these proteins form a heterocomplex with H-NS or H-NS paralogs [34–37]. Recent data has suggested that Hha contributes to H-NS silencing by aiding in H-NS bridging [38]. While YmoA alone cannot bind the yscW-lcrF promoter, H-NS alone or the H-NS/YmoA complex can [16]. In Y. pestis, YmoA is suggested to have a higher turnover rate at 37°C compared to environmental temperatures [32]. Current models suggest that degradation of YmoA, and therefore a reduction in the H-NS/YmoA complex at 37°C, relieves repression of yscW-lcrF [32]. Yet, ymoA deletion mutants exhibit higher levels of T3SS expression at 37°C compared to a parental strain in all three pathogenic Yersinia species [16,31,32], suggesting that some YmoA is present at 37°C during mammalian infection, and that H-NS retains repressive activity.

The Iron Sulfur Cluster Regulator IscR is a critical positive regulator of lcrF [17,39]. IscR belongs to the Rrf2 family of winged helix-turn-helix transcription factors [40,41]. IscR was first characterized in E. coli where it exists in two forms: holo-IscR bound to a [2Fe-2S] cluster, and cluster-less apo-IscR [42–45]. Both forms of IscR bind DNA, but while both apo-IscR and holo-IscR bind to so-called type II motif sequences, only holo-IscR binds type I motifs [43,44]. Holo-IscR represses its own expression through binding two type I motifs in the isc promoter [46]. Thus, conditions that increase iron-sulfur cluster demand, such as iron starvation or oxidative stress, lead to a lower holo- to apo-IscR ratio and higher overall IscR levels. We have previously shown that low iron and oxidative stress lead to upregulation of IscR in Yersinia, and subsequently upregulation of lcrF transcription and T3SS expression at 37°C [17,39].

In this study, we further analyze the role of H-NS and YmoA in regulation of the yscW-lcrF promoter at 37°C and integrate this into our current understanding of positive regulation by IscR. Promoter truncations and site mutations were used to probe the cis-acting sequences in the yscW-lcrF promoter region that are required for H-NS/YmoA repressive activity. Direct DNA binding of H-NS and IscR to this upstream region was investigated by ChIP-qPCR. A requirement for H-NS in controlling lcrF expression was examined using a CRISPRi mediated hns knockdown strain. The results revealed a specific role of H-NS in repression of lcrF, identified two H-NS binding regions in the yscW-lcrF promoter required for H-NS/YmoA repression, and showed that disruption of the H-NS/YmoA complex eliminates repression. Furthermore, our results reveal a surprising finding that the positive impact of IscR is eliminated in strains defective in formation of the H-NS/YmoA repressive complex. Taken together, our data suggest that an H-NS/YmoA complex is critical for proper IscR-dependent regulation of LcrF and the T3SS in response to changes in oxygen tension.

Results

Knockdown of H-NS leads to derepression of LcrF

H-NS is essential in human pathogenic Yersinia [27–30], complicating analysis of its involvement in Ysc T3SS expression. Therefore, to test the involvement of H-NS in lcrF regulation,
we used CRISPRi to knock down H-NS expression in *Y. pseudotuberculosis* and measured the effect on *lcrF* expression levels at 37˚C. For this CRISPRi system pioneered in *Y. pestis* [47], target gene guide RNAs and dCas9 can be induced in the presence of anhydrotetracycline (aTC). CRISPRi knockdown led to a ~6-fold decrease in H-NS transcription when exposed to aTC (Fig 1A). Importantly, this reduction of H-NS expression led to a ~31-fold increase in *lcrF* mRNA, demonstrating that H-NS represses LcrF transcription (Fig 1B). Knockdown of H-NS did not affect expression of *gyrA*, a housekeeping gene not predicted to be regulated by H-NS (Fig 1C). These data provide the first direct evidence that H-NS negatively influences *Yersinia* LcrF expression at 37˚C.

**H-NS occupies three DNA sites within the yscW-*lcrF* promoter**

H-NS and H-NS/YmoA complexes have been shown *in vitro* to bind a *yscW-*lcrF* promoter fragment containing sequences between the -2 to +272 nucleotides relative to the transcriptional start site [16]. However, the specific H-NS binding regions were not identified. We used FIMO-MEME suite tools to predict putative H-NS binding sites upstream of *yscW-*lcrF and identified three predicted H-NS binding sites, referred to as Sites I, II, and III. Site III is contained within the -2 to +272 nucleotide sequence originally shown to interact with H-NS, and Site I and II are located further upstream (Fig 2A). To test whether H-NS binds to these predicted sites, we carried out ChIP-qPCR analysis to assess H-NS occupancy *in vivo*. In order to immunoprecipitate H-NS-DNA complexes, we used a chromosomally-encoded 3xFLAG tagged H-NS allele. This FLAG tag did not affect the ability of H-NS to repress LcrF expression (Fig A in S1 Text). Consistent with this, H-NS enrichment was readily detectable at all three predicted sites in the *yscW-*lcrF promoter when bacteria were cultured at 26˚C. However, H-NS binding at these predicted sites was greatly diminished at 37˚C (Fig 2B), despite the fact that H-NS was required for repression of *yscW-*lcrF at this temperature. Interestingly, previous
reports have shown that H-NS in other facultative pathogens represses expression of certain virulence genes at environmental temperatures (<30˚C) but exhibits decreased binding at mammalian body temperature (37˚C) [24,26,48], in line with our findings. No enrichment of H-NS was seen at a control pYV-encoded promoter that was not predicted to bind H-NS at either temperature (DN756_21750). These data suggest that H-NS occupies the yscW-lcrF promoter at high levels under environmental temperatures at which the T3SS is known to be efficiently repressed [11,16]. Nevertheless, it is likely that some H-NS binds to the yscW-lcrF promoter at 37˚C, but is below the limit of detection of ChIP-qPCR.

YmoA is predicted to enhance the repressive ability of H-NS on the yscW-lcrF promoter at both 25˚C and 37˚C, but previous in vitro data suggests that this effect is not mediated by increasing H-NS binding to the yscW-lcrF promoter [16]. Consistent with this, no difference
in H-NS enrichment at the *yscW-lcrF* promoter was observed *in vivo* by ChIP-qPCR in the *ymoA* mutant compared to the parental strain at 26˚C, suggesting that YmoA does not affect H-NS occupancy at the *yscW-lcrF* promoter at this temperature (Fig 2C). In addition, deletion of *ymoA* did not change our ability to detect H-NS binding to the *yscW-lcrF* promoter at 37˚C. Since previous data showed that IscR exerts its positive effect at 37˚C, we also assayed H-NS enrichment in a strain lacking IscR and found no difference between an *iscR* mutant and the wildtype strain at 26˚C (Fig 2C) or 37˚C (Fig 2D). Taken together, these data indicate that H-NS binds to the three predicted DNA sites in the *yscW-lcrF* promoter *in vivo* and occupancy of these sites is higher at environmental temperatures compared to 37˚C but is independent of IscR or YmoA.

**Two H-NS binding regions are required to repress *yscW-lcrF* promoter activity**

To determine whether any of the three identified H-NS binding Sites I, II, and III function in H-NS-dependent regulation of the *yscW-lcrF* promoter at 37˚C, we systematically truncated the *yscW-lcrF* promoter and tested promoter activity using a *lacZ* reporter (Fig 3A). We also compared expression in a mutant lacking YmoA as a close mimic for strains lacking H-NS since *ymoA* is not an essential gene, whereas *hns* cannot be deleted. Eliminating the region containing H-NS binding Site I did not significantly affect promoter activity (Fig 3B; promoter 1 compared to promoter 2). However, additional truncation of H-NS binding Site II led to a significant increase in promoter activity that was less sensitive to YmoA, suggesting that some of the H-NS repressive effect had been lost by Site II truncation (Fig 3B; promoter 2 compared to promoter 3). Importantly, further truncation to eliminate the IscR binding site led to deregulated promoter activity (Fig 3B; promoter 4). Lastly, truncation to eliminate the -35 and -10 promoter elements led to a complete lack of promoter activity (Fig 3B; promoter 5), as expected. Taken together, these data suggest that H-NS binding Sites II and III are required for repression of *yscW-lcrF* promoter activity.

Since Site III is downstream of the transcription initiation site, we could not assess its contribution through promoter truncation. Thus, we mutated Site III in the *yscW-lcrF* promoter 2 *lacZ* construct by switching the TA-rich sequences to CG-rich sequences to perturb H-NS binding and compared it to a similarly mutated Site II. Mutation of Site III resulted in significantly increased promoter activity in the WT genetic background (Fig 4A). Mutation of Site II did not lead to a statistically significant increase (Fig 4A, WT background), although mutation of both Site II and Site III resulted in promoter activity that trended higher than mutation of Site III alone. Importantly, mutation of Site II and/or III did not significantly change promoter activity in the absence of YmoA (Fig 4A, Δ*ymoA* background), suggesting YmoA mediated repression was eliminated. Thus, these data suggest that H-NS binding to at least two of the three identified binding regions plays a major role in H-NS-mediated repression of the *yscW-lcrF* promoter.

Given the location of the IscR binding site between H-NS Sites II and III, we also determined whether IscR regulation of the *yscW-lcrF* promoter is impacted by H-NS/YmoA-mediated repression. While deletion of *iscR* did not affect YmoA or H-NS expression (Fig B in S1 Text), we found that deletion of *iscR* led to a decrease in *yscW-lcrF* promoter activity compared to the wildtype strain for all promoter constructs containing the known IscR binding site (Fig 3; promoters 1–3), as expected from our previous studies [17,39]. Surprisingly, in the absence of *ymoA*, deletion of *iscR* had no effect on promoter activity (Fig 3B), suggesting that IscR is dispensable for *yscW-lcrF* promoter activity in the absence of an H-NS/YmoA-repressive complex.
Since IscR had no effect on \( yscW-lcrF \) promoter activity in the absence of YmoA, we tested whether IscR enrichment at the \( yscW-lcrF \) promoter in vivo was impacted by YmoA. We used a chromosomal 3xFLAG tagged IscR allele previously shown not to affect IscR activity [49]. IscR enrichment at the \( yscW-lcrF \) promoter was not influenced by the presence of YmoA since the levels were the same between the wildtype strain and the \( ymoA \) mutant at either 26˚ or 37˚C (Fig 2E). However, IscR enrichment at the \( yscW-lcrF \) promoter was ~3-fold higher at 37˚C compared to 26˚C, while IscR enrichment at the promoter of another known IscR target, the \( suf \) operon, did not significantly differ with temperature (Fig C in S1 Text). This increase in IscR enrichment at the \( yscW-lcrF \) promoter at 37˚C is not due to increased IscR levels since we did not observe higher levels of IscR protein when \( Yersinia \) was cultured at 37˚C compared to 26˚C (Fig B in S1 Text). Thus, a lack of IscR binding cannot explain the absence of IscR regulation of the \( yscW-lcrF \) promoter in strains deleted for \( ymoA \). Rather, based on the results establishing a repressive effect of an H-NS/YmoA complex acting through H-NS binding to

![Fig 3. The identified H-NS binding Sites II and III are important for regulation of \( yscW-lcrF \) promoter activity. (A) Schematic of \( PyscW-lcrF::lacZ \) fusions. Five constructs (p1-p5) were used to assess which regions of \( p\yscW-lcrF \) allows for H-NS-YmoA repression and IscR activation. (B) \( Yersinia \) harboring the various \( p\yscW-lcrF::lacZ \) plasmids were grown aerobically under T3SS-inducing conditions (low calcium LB at 37˚C) for 1.5 hrs and assayed for β-galactosidase (Miller units). The average of at least three biological replicates are shown ± standard deviation. Statistical analysis was performed using an unpaired Student’s t-test (* \( p < .05 \), ** \( p < .01 \), *** \( p < .001 \), **** \( p < .0001 \), and n.s. non-significant).

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\( \text{H-NS/YmoA complex enables IscR regulation of } Yersinia \text{ LcrF} \)
regions flanking the IscR binding site, we considered the notion that IscR acts by interfering with H-NS/YmoA repression of the yscW-lcrF promoter.

To test whether IscR potentiation of yscW-lcrF promoter activity was affected by the H-NS/YmoA complex, we mutated the characterized IscR binding site in the yscW-lcrF promoter 2 lacZ construct (pNull) and introduced into WT and ΔymoA Y. pseudotuberculosis genetic backgrounds. All these strains were grown aerobically under T3SS-inducing conditions (low calcium LB at 37˚C) for 1.5 hrs and assayed for β-galactosidase (Miller units). The different promoter constructs are listed on the x-axis. The average of at least three biological replicates are shown ± standard deviation. Statistical analysis was performed using (A-B) a one-way ANOVA with Bonferroni’s multiple comparisons test on either (yellow bars) all WT genetic backgrounds carrying different reporter constructs or (blue, green, or red bars) each individual reporter construct expressed in WT, ΔiscR, ΔymoA, or ΔiscR/ΔymoA; or (B) an unpaired t test (purple bars). (*) p < .05, (**) p < .01, (***) p < .001, **** p < .0001, and n.s. non-significant.

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IscR is not required for LcrF expression or type III secretion in the absence of YmoA

Since we found that loss of iscR did not decrease yscW-lcrF promoter activity in the absence of YmoA, we assessed T3SS activity of Y. pseudotuberculosis expressing or lacking iscR and/or
ymoA. Consistent with previous studies, we observed ~18-fold decrease in secretion of the T3SS effector protein YopE upon iscR deletion, while ymoA deletion led to ~6-fold increase in YopE secretion (Fig 5A). As expected, the effect of YmoA on T3SS activity required LcrF (Fig D in S1 Text). Importantly, YopE secretion in the ΔiscR/ΔymoA double mutant was similar to the ΔymoA mutant, indicating that IscR is dispensable for T3SS activity in the absence of YmoA (Fig 5A). As proteins of the YmoA family lack a DNA binding domain and are thought to affect transcription by interacting with H-NS [33,38], we tested whether the ability of YmoA to bind H-NS was required for its ability to control lcrF expression and, therefore, T3SS activity. Previous studies showed that a YmoA D43N mutant cannot interact with H-NS in vitro [50]. A Y. pseudotuberculosis ymoA Δ43N mutant was able to express YmoA (Fig E in S1 Text), but exhibited ~6-fold increase in YopE secretion, similar to ymoA deletion (Fig 5A). This suggests that YmoA represses yscW-lcrF through its interaction with H-NS. Furthermore, there was no difference in YopE secretion between the ymoA Δ43N mutant and the iscR/ymoA Δ43N double mutant. These effects on YopE secretion are most easily explained by changes in LcrF levels. As expected, the ΔiscR mutant had a ~5-fold reduction in lcrF mRNA compared to wild-type, while the ΔymoA and ymoA Δ43N mutants displayed ~10-fold elevated lcrF mRNA (Fig 5B and 5C). In contrast, we observed no difference in LcrF expression between the ΔymoA and ΔiscR/ΔymoA mutants or between the ymoA Δ43N and ΔiscR/ymoA Δ43N mutants. Likewise, mutation of the IscR binding site mutant in the yscW-lcrF promoter (lcrF pNull) reduced LcrF
expression and T3SS activity in the presence of YmoA, but not in the absence of YmoA (Fig F in S1 Text). Collectively, these data suggest that YmoA requires H-NS binding to inhibit lcrF transcription, and that IscR regulation of the yscW-lcrF promoter and T3SS expression is only important in the presence of the H-NS/YmoA complex.

**IscR-dependent regulation of LcrF and the T3SS in response to oxygen availability requires YmoA**

We previously showed that low iron and high oxidative stress lead to elevated IscR levels, which then activate T3SS expression through upregulation of LcrF [17]. The data shown here suggest that IscR is dispensable in the absence of H-NS/YmoA activity at the yscW-lcrF promoter. To test how YmoA affects the ability of IscR to regulate LcrF in response to environmental cues, we measured lcrF mRNA levels in ΔiscR and ΔymoA mutants under aerobic or anaerobic conditions at 37˚C. As expected, under aerobic conditions iscR mRNA levels were increased ~4-fold compared to anaerobic conditions, and this upregulation of iscR levels led to a ~12-fold induction in lcrF levels in the wildtype strain (Fig 6A and 6B). In contrast, lcrF

![Fig 6](https://doi.org/10.1371/journal.pgen.1010321.g006)
mRNA and protein levels were not affected by oxygen in the ΔymoA and ΔiscR/ΔymoA mutants (Fig 6A and 6C). Unlike what we observed using rich media (Fig B in S1 Text), deletion of ymoA reduced expression of IscR mRNA and protein expression under these minimal media conditions (Fig 6B and 6C), although this does not explain elevated LcrF/T3SS expression in the ymoA mutant. Taken together, these data suggest that the H-NS/YmoA complex facilitates IscR regulation of LcrF and the T3SS in response to environmental cues.

Discussion

The histone-like protein H-NS has been suggested to repress expression of horizontally transferred genes, but the hns gene is essential in pathogenic Yersinia [27–30]. In this study, CRISPRi knockdown in Y. pseudotuberculosis revealed that hns is required for repression of the LcrF T3SS master regulator. Furthermore, genetic analysis revealed that YmoA must interact with H-NS to repress lcrF transcription. We identified two relevant H-NS binding sites in the yscW-lcrF promoter that contribute to promoter repression in a manner dependent on YmoA. Interestingly, we found that the transcription factor IscR promotes lcrF expression and T3SS activity only in the presence of H-NS/YmoA. Y. pseudotuberculosis IscR levels are low under anaerobic conditions, such as those in the intestinal lumen. As Yersinia cross the intestinal barrier, oxygen tension increases. Oxidative stress has been shown to stimulate elevated IscR levels that drive LcrF expression and type III secretion, which is critical for extraintestinal infection [17,51–54]. Importantly, oxygen- and IscR-dependent regulation of LcrF and the T3SS required an H-NS/YmoA complex. These data suggest that following acquisition of the pYV virulence plasmid encoding the Ysc T3SS during Yersinia evolution, Ysc T3SS genes may have come under general repression by H-NS and its modulator YmoA but evolved to respond specifically to IscR-responsive host tissue cues to enable T3SS-mediated target host cell immunomodulation.

YmoA was previously shown to bind H-NS and the H-NS/YmoA complex was proposed to regulate LcrF expression [16,50]. Indeed, a YmoA point mutation that eliminates H-NS binding phenocopied the ymoA deficient strain, suggesting YmoA must interact with H-NS to repress LcrF and T3SS expression. However, YmoA was not shown to affect H-NS binding to the yscW-lcrF promoter in vitro [16], and our ChIP-qPCR analysis did not find a change in H-NS yscW-lcrF promoter occupancy in the presence or absence of YmoA at 26˚C. At 37˚C, H-NS occupancy was below the limit of detection by ChIP-qPCR, so we could not rule out H-NS binding to the yscW-lcrF promoter at this temperature, nor the effect of YmoA on residual H-NS binding. However, deletion of ymoA or knockdown of hns both caused elevated LcrF expression at 37˚C, indicating that both proteins are needed to repress the yscW-lcrF promoter at mammalian body temperature. Furthermore, the fact that mutation of the Site II/III H-NS binding sites in an lcrF transcriptional reporter construct led to an increase in promoter activity at 37˚C suggests that H-NS repression of lcrF activity is at least partially due to direct binding. However, it is possible that H-NS may have additional indirect effects on lcrF transcription. A recent study showed that deletion of ymoA impacts expression of almost 300 genes, including known regulators [55], raising the possibility that deletion of ymoA may affect the yscW-lcrF promoter both directly and indirectly. However, the fact that deletion of YmoA did not affect promoter activity of the yscW-lcrF Site II/III mutant promoter construct suggests that YmoA does not affect yscW-lcrF promoter activity independently of the H-NS binding sites. Taken together, these data suggest that YmoA binding to H-NS does not alter H-NS occupancy at the lcrF promoter but, rather, potentiates H-NS repressive activity. How YmoA affects H-NS activity will need to be addressed by future studies. Importantly, a Y. pseudotuberculosis strain lacking ymoA is attenuated in colonization of intestinal lymph tissue and vital
organs following oral infection [55]. While derepression of the T3SS may play a role in this virulence defect, the effect of ymoA deletion on expression of Yersinia adhesins and other virulence factors likely contributes [55]. Indeed, YmoA modulates the carbon storage regulator (Csr) global regulatory system known to regulate several virulence factors. We observed that ymoA deletion led to lower IscR levels following growth in minimal media (Fig 6) but not in rich media (Fig B in S1 Text), suggesting that perhaps YmoA contributes to iscR expression under nutrient limiting conditions.

The pYV plasmid in the region of the lcrF gene has been reported to take on different architecture depending on temperature [56]. A “bent” pYV architecture was suggested to be stabilized by H-NS at lower temperatures while this bending was not detected at 37˚C. H-NS oligomers can either form a nucleoprotein filament on a contiguous stretch of DNA or can form DNA bridges when multiple discrete H-NS binding regions are brought together [57,58]. It is possible that at lower temperatures, H-NS forms nucleoprotein filaments at the lcrF gene locus on pYV, perhaps nucleated from the identified binding sites, and thus H-NS enrichment via ChIP-qPCR is readily detectable. In contrast, it is possible that at 37˚C, fewer H-NS molecules are bound and the sensitivity of ChIP-qPCR is not sufficient to detect this binding. However, the fact that mutation of the H-NS binding Sites II and III in the yscW-lcrF promoter leads to derepression of promoter activity at 37˚C, and that this is independent of YmoA, strongly suggest that H-NS/YmoA is still active at the wildtype yscW-lcrF promoter at 37˚C.

Previous reports have suggested that H-NS represses some target genes under environmental conditions, but to a lesser extent at mammalian body temperature. For example, the Shigella flexneri T3SS is co-regulated by the AraC transcriptional regulator VirF and H-NS [59,60]. VirF promotes VirB, which ultimately activates the Shigella T3SS [61]. The Shigella T3SS is only expressed at mammalian body temperature and this is controlled by preventing expression of VirF at environmental temperatures. H-NS binds to two distinct sites upstream of virF leading to the formation of a DNA bridge [62], repressing virF transcription [48]. Furthermore, more H-NS binding to the virF promoter is observed at lower temperatures (<30˚C) compared to mammalian body temperature (37˚C). Similarly, we observed H-NS enrichment at the yscW-lcrF promoter robustly at environmental temperature while it was below the limit of detection by ChIP-qPCR at 37˚C. However, Yersinia H-NS repression of LcrF still plays an important role at 37˚C, as CRISPRi knockdown at this temperature leads to lcrF derepression.

IscR-regulated genes in other bacterial pathogens have also been shown to be repressed by H-NS, and H-NS has been shown to repress T3SS genes in many bacterial species. In Vibrio species, for example, nitrosative stress and iron starvation drive V. vulnificus IscR to compete with H-NS binding to the promoter of the vvhBA operon, which encodes an extracellular pore-forming toxin essential for its hemolytic activity [63–66]. Furthermore, in V. parahaemolyticus, H-NS represses expression of the T3SS-1 regulator ExsA, while exsA is induced by the transcription factor HlyU [67]. Like IscR, HlyU may also sense oxygen levels [68]. In Salmonella enterica, H-NS has been suggested to bind to a high-affinity nucleation site in the regulatory region of the Salmonella pathogenicity island (SPI)-1 T3SS regulator HilD, leading to repression of hilD [69]. In addition, genetic evidence suggests that this H-NS repression of hilD can be counteracted by HilD itself [69]. Interestingly, IscR has been shown to repress the Salmonella pathogenicity island (SPI)-1 T3SS regulator HilD in response to iron depletion, possibly facilitating expression of the SPI-1 T3SS in the intestinal lumen where it is needed to enter intestinal epithelial cells [70]. Likewise, in Pseudomonas aeruginosa, the H-NS family members MvaT and MvaU repress the promoter of the T3SS master regulator ExsA, an AraC family transcription factor with similarity to Yersinia LcrF [11,71], while the cAMP-responsive Vfr protein directly activates exsA transcription [72]. Collectively, these data are consistent with the idea that T3SS genes must come under H-NS family repression following horizontal
gene transfer, necessitating a positive regulatory factor that must overcome H-NS repression to induce virulence gene expression in response to an appropriate signal. In the case of the Yersinia, H-NS repression of the T3SS master regulator LcrF plays an active role in allowing T3SS genes to be responsive to relevant host tissue cues through IscR.

Materials and methods

Bacterial strains and growth conditions

Bacterial strains used in this paper are listed in Table A in S1 Text. Y. pseudotuberculosis were grown, unless otherwise specified, in LB (Luria Broth) at 26˚C shaking overnight. To induce the T3SS, overnight cultures were diluted into low calcium LB medium (LB plus 20 mM sodium oxalate and 20 mM MgCl₂) to an optical density (OD₆₀₀) of 0.2 and grown for 1.5 h at 26˚C shaking followed by 1.5 h at 37˚C to induce Yop synthesis, as previously described [73].

For growing Yersinia under varying oxygen conditions, casamino acid-supplemented M9 media, referred to as M9 minimal media in this study, was used [74]. Growth of cultures to vary oxygen tension was achieved by first diluting 26˚C overnight aerobic cultures of Y. pseudotuberculosis to an OD₆₀₀ of 0.1 in fresh M9 minimal media supplemented with 0.9% glucose to maximize growth rate and energy production under anaerobic conditions, and incubating for 12 hrs under either aerobic or anaerobic conditions at 26˚C. Both aerobic and anaerobic cultures were then diluted to an OD₆₀₀ of 0.1, grown for 2 hrs at 26˚C, and shifted to 37˚C for 4 hrs.

Construction of Yersinia mutant strains

The Yersinia mutants were generated as described in [49]. H-NS was tagged with a C-terminal 3xFLAG affinity tag at the native locus through splicing by overlap extension [75], using primer pair Fhns_cds/Rhns_cds (Table B in S1 Text) to amplify ~500bp upstream of hns plus the hns coding region excluding the stop codon, F3xFLAG/R3xFLAG to amplify the 3xFLAG tag, and F3'hns/R3'hns to amplify the ~500 bp downstream region of hns including the stop codon. For the AymoA mutant, primer pairs F5/R5ΔymoA were used to amplify ~1000 bp 5' of ymoA and F3/R3ΔymoA to amplify ~1000 bp 3' of ymoA. To generate the ymoAD43N mutant, primer pairs pUC19_YmoA_F and pUC19_YmoA_R were used to amplify 250 bp upstream of ymoA to 250 downstream of the ymoA start codon and the amplified product cloned into a BamHI and SacI digested pUC19 plasmid. Q5 site directed mutagenesis was performed using primer pairs ymoAD43N_F and ymoAD43N_R. The resulting plasmid, pUC19 ymoAD43N, was digested with BamHI and SacI and the resulting fragment was ligated into the suicide plasmid pSR47s. Mutant strains were generated as described above.

In order to generate lacZ promoter constructs of ymoBA and hns, primer pairs pFU99a_y-moA_F/pFU99a_ymoA_R and pFU99a_hns_F/pFU99a_hns_R were used to amplify ~500 bp upstream of ymoA and hns, respectively, which included the first ten amino acids of ymoA and hns. These promoters and first ten amino acids of YmoA and H-NS were fused in frame to lacZ and cloned into a BamHI- and Sall-digested pFU99a using the NEBuilder HiFi DNA Assembly kit (New England Biolabs, Inc) and electroporated into Y. pseudotuberculosis.

In order to generate lacZ promoter constructs of yscW-lcrF, the reverse primer pFU99a_yscWlcrF_R was used with the following forward primers: pFU99a_yscWlcrF_p1 (promoter construct 1/ -505 to +294 of yscW), pFU99a_yscWlcrF_p2 (promoter construct 2/ -309 to +294 of yscW), pFU99a_yscWlcrF_p3 (promoter construct 3/ -166 to +294 of yscW), pFU99a_yscWlcrF_p4 (promoter construct 4/ -47 to +294 of yscW), or pFU99a_yscWlcrF_p5 (promoter construct 5/ +101 to +294 of yscW). These promoter fragments were cloned into a BamHI- and Sall-digested pFU99a and electroporated into Y. pseudotuberculosis. To generate
the Site 2, Site 3, Site 2/Site3, pNull, pNull/Site2, pNull/Site3, and pNull/Site2/Site3 mutant lacZ promoter constructs, Q5 mutagenesis was used with primers listed in Table B in S1 Text. The single mutant plasmids were generated first, which were then used as templates to generate the double mutant plasmids, and the double mutant plasmid was used as a template to generate the triple mutant plasmid.

**Type III secretion system secretion assay**
Visualization of T3SS cargo secreted in broth culture was performed as previously described [76]. Briefly, *Y. pseudotuberculosis* in LB low calcium media (LB plus 20 mM sodium oxalate and 20 mM MgCl2) was grown for 1.5 h at 26˚C followed by growth at 37˚C for 1.5 h. Alternatively for the ±O2 cultures, *Y. pseudotuberculosis* was grown in M9 minimal media with 0.9% glucose as described above. Cultures were normalized to OD600 and pelleted at 13,200 rpm for 10 min at room temperature. Supernatants were removed, BSA spiked into the supernatant to serve as a control, and proteins precipitated by addition of trichloroacetic acid (TCA) at a final concentration of 10%. Samples were incubated on ice for at least 1 hr and pelleted at 13,200 rpm for 15 min at 4˚C. Resulting pellets were washed twice with ice-cold 100% acetone and resuspended in final sample buffer (FSB) containing 0.2 M dithiothreitol (DTT). Samples were boiled for 5 min prior to separating on a 12.5% SDS-PAGE gel. Coomassie stained gels were imaged using Bio-Rad Image Lab Software Quantity and Analysis tools. YopE bands were quantified using this software and normalized to the BSA protein precipitation control.

**Western blot analysis**
Cell pellets were collected, resuspended in FSB plus 0.2 M DTT, and boiled for fifteen minutes. At the time of loading, supernatants and cell pellets were normalized to the same number of cells. After separation on a 12.5% SDS-PAGE gel, proteins were transferred onto a blotting membrane (Immobilon-P) with a wet mini trans-blot cell (Bio-Rad). Blots were blocked for an hour in Tris-buffered saline with Tween 20 and 5% skim milk, and probed with the rabbit anti-RpoA (gift from Melanie Marketon), rabbit anti-LcrF (gift from Gregory Plano), rabbit anti-IscR [44], rabbit anti-YmoA (gift from Gregory Plano), rabbit anti H-NS (gift from Robert Landick), mouse M2 anti-FLAG (Sigma), goat anti-YopE (Santa Cruz Biotech), and horse-radish peroxidase-conjugated secondary antibodies (Santa Cruz Biotech). Following visualization, quantification of the bands was performed with Image Lab software (Bio-Rad).

**Quantitative RT-PCR**
RT-qPCR was carried out as previously described [49] using the primers in Table B in S1 Text. The expression levels of each target gene were normalized to that of 16S rRNA present in each sample and calculated by utilization of a standard curve. At least three independent biological replicates were analyzed for each condition.

**β-galactosidase assays**
*Y. pseudotuberculosis* harboring promoter-lacZ fusion plasmids were grown in LB low calcium media (LB plus 20 mM sodium oxalate and 20 mM MgCl2) for 1.5 h at 26˚C followed by growth at 37˚C for 1.5 h. Protein expression was stopped by incubating cells on ice for 20 minutes. Cultures were spun down and resuspended in Z Buffer [77]. Samples were permeabilized using chloroform and 0.1% sodium dodecyl sulfate, incubated with 0.8 mg/mL ONPG, and β-galactosidase enzymatic activity was terminated by the addition of 1M sodium bicarbonate. β-galactosidase activity is reported as Miller units.
CRISPRi knockdown

Knockdown of H-NS via CRISPRi methods was adapted from [47]. In order to generate the pgRNA-tetO-JTetR-H-NS plasmid, a protospacer-adjacent motif (PAM) was located near the promoter of hns [78]. Two oligonucleotides (hns_gRNA_F and hns_gRNA_R) consisting of 20-nt targeting the hns promoter region with BbsI cohesive ends were synthesized and annealed before being cloned into pgRNA-tetO-JTetR (Addgene) by Golden Gate assembly. The plasmids pdCas9-bacteria (Addgene) and pgRNA-tetO-JTetR-H-NS were transformed into wildtype Y. pseudotuberculosis sequentially. These plasmids induce expression of dCas9 and gRNA-H-NS when exposed to anhydrotetracycline. Y. pseudotuberculosis cultures carrying these plasmids were sub-cultured to OD_{600} 0.2 and incubated at 26˚C for 3 hrs in the presence or absence of 1μg/mL anhydrotetracycline, and then transferred to 37˚C for 1.5 hrs to induce the T3SS. Samples were collected, and RNA was isolated for qRT-PCR analysis.

Bioinformatic prediction of H-NS/YmoA binding sites

A training set of known H-NS binding sites in E. coli K-12 substr. MG1655 from RegulonDB was used to generate an H-NS binding motif using MEME-suite 5.1.1 tools [79,80]. FIMO was then used to scan for an H-NS binding site near the regulatory region of the yscW-lcrF promoter.

ChIP-qPCR

Cells were grown for 3 hrs at 26˚C or 37˚C with shaking at 250 rpm and protein/nucleic acids were crosslinked using 1% formaldehyde at 26˚C or 37˚C for 10 min. Crosslinking was quenched with the addition of ice cold 0.1 M glycine and incubated at 4˚C for 30 min. 32 x 1 [OD_{600}] cells were harvested for each replicate and cell pellets were stored at −80˚C. DNA was fragmented by resuspending samples using IP buffer (100mM Tris-HCl, pH 8, 300mM NaCl, 1% Triton X-100, 1 mM PMSF) and sonicated at 25% Amplitude 15s on/59s off for a total of 8 cycles per sample. After sonication, lysates were treated with micrococcal nuclease and RNase-A for 1hr at 4˚C. Lysates were clarified via centrifugation at 13,000 rpm for 15 min at 4˚C. Lysates were pre-cleared using Dynabeads Protein A/G for 3hr at 4˚C. Immunoprecipitation was performed by adding Sigma monoclonal mouse anti-FLAG M2 antibody to samples and incubated overnight at 4˚C. Dynabeads Protein A/G were added to samples and washes were performed to remove non-specific binding. After H-NS-DNA or IscR-DNA complexes were eluted, samples were placed at 65˚C for 5 hr to reverse crosslinks. DNA was then purified using Qiangen PCR purification kit and input samples were diluted 1:100 while samples treated with antibody or control samples not treated with the antibody were diluted 1:5 and qPCR was performed to assess IscR/H-NS binding to promoters of interest. Percent input was calculated by the following equation: 100^\left(\frac{CT_{\text{input}} - CT_{\text{input+AB}}}{\text{CT}_{\text{input}}}\right).

Supporting information

S1 Text. Fig A in S1 Text. 3xFLAG tag allows for detection of H-NS using FLAG antibody and does not affect H-NS ability to repress LcrF. Fig B in S1 Text. IscR does not regulate YmoA or H-NS expression. Fig C in S1 Text. IscR enrichment at the suf promoter is not influenced by temperature. Fig D in S1 Text. YmoA affects LcrF dependent type III secretion activity. Fig E in S1 Text. The YmoA^{D43N} mutant protein is expressed. Fig F in S1 Text. IscR binding to the yscW-lcrF promoter is dispensable in the absence of ymoA. Table A in S1 Text. Strains used in this study. Table B in S1 Text. Y. pseudotuberculosis primers used in this study.
Table C in S1 Text. Plasmids used in this study.

(AOCX)

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