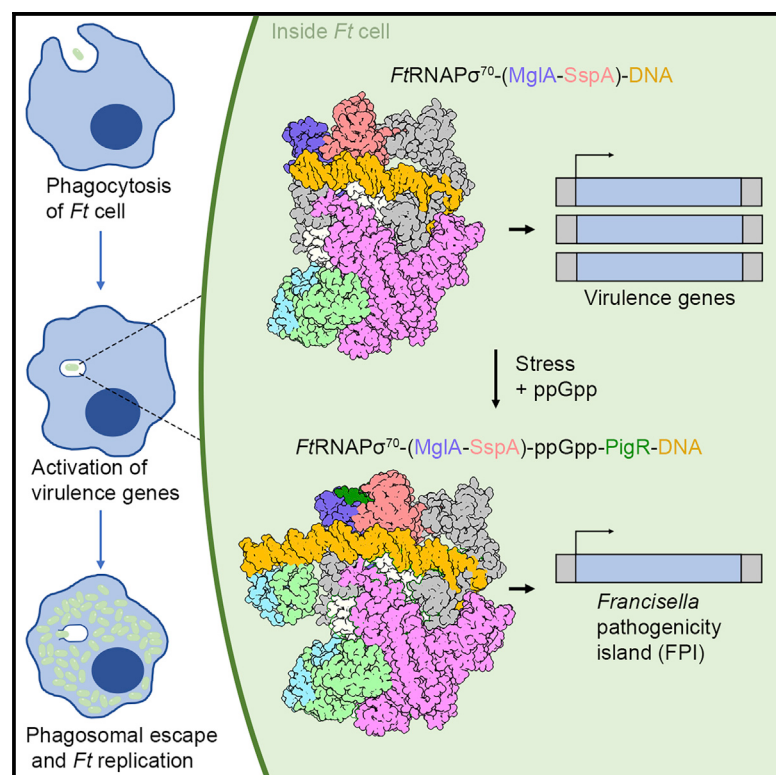


Structural Basis for Virulence Activation of *Francisella tularensis*

Graphical Abstract



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In Brief

Francisella tularensis pathogenicity is driven by unique regulators, the MglA-SspA heterodimer, PigR and the alarmone ppGpp that combine to activate *Francisella* pathogenicity island transcription by an unknown mechanism. Using cryo-EM, crystallography, and cellular studies, Travis et al. reveal how MglA-SspA binds RNAP and employs two strategies to activate virulence genes.

Highlights

- *Ft*RNAP σ^{70} -(MglA-SspA)-DNA cryo-EM structure reveals MglA-SspA enables σ^{70} -DNA binding
- *Ec*RNAP σ^{70} -(SspA)₂-DNA complex shows conserved mode of SspA-family binding to RNAP σ^{70}
- PigR binding to *Ft*RNAP σ^{70} -(MglA-SspA)-DNA via ppGpp reveals α CTD-promoter recruitment
- Combined data show two (MglA-SspA)-based strategies for *Ft* virulence activation



Article

Structural Basis for Virulence Activation of *Francisella tularensis*

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SUMMARY

The bacterium *Francisella tularensis* (*Ft*) is one of the most infectious agents known. *Ft* virulence is controlled by a unique combination of transcription regulators: the MglA-SspA heterodimer, PigR, and the stress signal, ppGpp. MglA-SspA assembles with the σ^{70} -associated RNAP holoenzyme (RNAP σ^{70}), forming a virulence-specialized polymerase. These factors activate *Francisella* pathogenicity island (FPI) gene expression, which is required for virulence, but the mechanism is unknown. Here we report *Ft* RNAP σ^{70} -promoter-DNA, *Ft* RNAP σ^{70} -(MglA-SspA)-promoter DNA, and *Ft* RNAP σ^{70} -(MglA-SspA)-ppGpp-PigR-promoter DNA cryo-EM structures. Structural and genetic analyses show MglA-SspA facilitates σ^{70} binding to DNA to regulate virulence and virulence-enhancing genes. Our *Escherichia coli* RNAP σ^{70} -homodimeric EcSspA structure suggests this is a general SspA-transcription regulation mechanism. Strikingly, our *Ft* RNAP σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA structure reveals ppGpp binding to MglA-SspA tethers PigR to promoters. PigR in turn recruits *Ft* RNAP α CTDs to DNA UP elements. Thus, these studies unveil a unique mechanism for *Ft* pathogenesis involving a virulence-specialized RNAP that employs two (MglA-SspA)-based strategies to activate virulence genes.

INTRODUCTION

Francisella tularensis (*Ft*), the causative agent of tularemia, is a Gram-negative γ -proteobacterium that is one of the most infectious pathogens known, with as few as 10 organisms constituting an infectious dose (Dennis et al., 2001; Maurin, 2015). Due to its high infectivity and ease of aerosolization, several countries have developed the bacterium as a bioweapon and the CDC has listed it as a category A bioterrorism agent (Dennis et al., 2001; Maurin, 2015). Multidrug-resistant forms of *Ft* can be readily generated in the laboratory, further heightening concern of this pathogen (Gestin et al., 2010; Loveless et al., 2010; Suter et al., 2014). The factors and molecular mechanisms that control *Ft* pathogenicity are not fully understood. However, a cluster of genes present on the *Francisella* pathogenicity island (FPI) has been shown to be necessary for phagosomal escape by this bacterium into the cytosol of infected macrophages, a key step in *Ft* replication and pathogenesis (Lauriano et al., 2004; Nano et al., 2004; Larsson et al., 2005; Weiss et al., 2007;

Barker et al., 2009; Bröms et al., 2010; Eshraghi et al., 2016). The FPI cluster, which encodes a type VI secretion system (Nano et al., 2004; Barker et al., 2009; Bröms et al., 2010; Russell et al., 2014), is present in two chromosomal copies in the most virulent *Ft* subspecies, *F. tularensis tularensis* and *F. tularensis holarctica*.

Recent studies have revealed the *Ft* transcription regulatory system that mediates FPI activation. This system is composed of the stringent starvation protein A (SspA), the macrophage growth locus protein A (MglA) and the pathogenicity island gene regulator (PigR) (Lauriano et al., 2004; Charity et al., 2007, 2009; Brotcke and Monack, 2008; Rohlfing and Dove, 2014; Cuthbert et al., 2015, 2017). Insight into how these proteins sense infection to activate the FPI was revealed by studies showing that the stress alarmone, ppGpp, produced upon infection binds directly to the MglA-SspA complex (Cuthbert et al., 2017). The only component of the *Ft* virulence circuitry with homologs in other Gram-negative bacteria is SspA, which is homodimeric in these bacteria and is involved

in stress and virulence responses (Hansen et al., 2005a). Interestingly, our recent data showed that MglA is an SspA-like protein that heterodimerizes with the *Ft*SspA (Cuthbert et al., 2015, 2017), and studies revealed that MglA and SspA, which are expressed during *Ft* infection, are constitutively associated with the *Ft*RNA polymerase (RNAP) (Ramsey et al., 2015). Bacterial RNAP core enzymes consist of large β and β' subunits that combine to form the active site. Their assembly is facilitated by the homodimeric N-terminal domains of the α subunit (α NTDs) (Murakami and Darst, 2003; Feklistov et al., 2014; Murakami, 2015). Bacterial σ factors are dissociable RNAP subunits that dictate DNA promoter specificity (Feklistov et al., 2014).

Unlike most bacteria, *Ft* encodes only two σ factors, the general “housekeeping” σ^{70} and σ^{32} . Notably, *Ft* utilizes the σ^{70} containing holoenzyme (*Ft*RNAP σ^{70}) for activation of its virulence genes. Intriguingly, *Ft* also encodes two distinct α RNAP subunits, which heterodimerize (Charity et al., 2007; Mukhamedyarov et al., 2011). Thus, *Ft* utilizes a regulatory circuitry to activate pathogenesis that includes a unique RNAP and unique combination of regulatory factors. How MglA-SspA interacts with the *Ft*RNAP σ^{70} is unknown. Moreover, while other SspA-like proteins are well-known regulators of virulence genes, their mechanisms have remained enigmatic. The molecular basis by which Pigr, *Ft*RNAP σ^{70} - (MglA-SspA), and the stress signal ppGpp collaborate to activate transcription of FPI genes is also not known. Here we address these key issues by employing cryo-electron microscopy (cryo-EM), X-ray crystallography, and cellular analyses. Our studies unveil specific molecular mechanisms for activation of *Ft* virulence genes while also revealing a conserved interaction mechanism of SspA proteins for RNAP σ^{70} holoenzymes.

RESULTS

Overall *Ft*RNAP σ^{70} -(MglA-SspA)-DNA Structure

To elucidate the *Ft*RNAP σ^{70} structure and determine how it interacts with MglA-SspA, we used single-particle cryo-EM. *Ft*RNAP σ^{70} -(MglA-SspA) complexes, consisting of ($\beta\beta'\alpha_1\alpha_2\omega$) σ^{70} -(MglA-SspA), were isolated from cells of the live vaccine strain (LVS) (Fortier et al., 1991), an attenuated form of *Ft* subspecies *holarctica*, which contained an RNAP β' subunit with an engineered TAP-tag (Figure S1A). Previous studies showed the resultant RNAP is fully active and that MglA and SspA are isolated in stoichiometric amounts with the cellular *Ft*RNAP σ^{70} (Figure S1A) (Charity et al., 2009). Notably, the *Ft*RNAP σ^{70} -(MglA-SspA) complex can also be pulled down if only MglA is tagged (Figure S1A). Using the *Ft*RNAP σ^{70} -(MglA-SspA) complex purified with the β' -TAP tag, we determined the 3.46 Å resolution cryo-EM structure of *Ft*RNAP σ^{70} -(MglA-SspA) bound to a 53-mer DNA site that encompassed base pairs (bps) –51 to 1 bp beyond the transcription start site (+1) of the FPI-housed *iglA* promoter (Figure 1A) (Ramsey et al., 2015). The structure revealed unambiguous density for most of the protein subunits. The promoter DNA, from bps –35 to +1, was also visible while the DNA upstream of the –35 element was disordered (Figure 1A; Table S1). The MglA-SspA heterodimer is well resolved in the

structure and makes extensive interactions with the *Ft*RNAP σ^{70} holoenzyme.

*Ft*RNAP Contains a Distinct α Heterodimer

The *Ft*RNAP core enzyme harbors a similar overall organization as observed in *E. coli* and other structurally characterized bacterial RNAPs (Murakami and Darst, 2003; Murakami, 2015). The C α atoms of the individual *Ft* β , β' , and ω subunits in the structure can be superimposed onto those of the *E. coli* RNAP with root mean squared deviations (RMSD) of 1.4 Å, 1.3 Å, and 1.4 Å, respectively. But while the *Ft* α subunits also superimpose well with the *E. coli* α subunit (RMSD = 1.4 Å), instead of an α homodimer, the *Ft*RNAP core contains an α heterodimer that is composed of two subunits, which share only 32% sequence identity. The presence of two different *Ft* α subunits suggested the possibility that four *Ft*RNAP core enzymes, (α_1) $_2$ $\beta\beta'$, (α_2) $_2$ $\beta\beta'$, ($\alpha_1\alpha_2$) $\beta\beta'$, and ($\alpha_2\alpha_1$) $\beta\beta'$, might form, with the latter enzyme species differing in which α contacts β and which α contacts β' (Charity et al., 2007; Mukhamedyarov et al., 2011). RNAP α subunits consist of N-terminal and C-terminal domains connected by a long linker. Density is not visible for the linkers or the α C-terminal domains (α CTDs) in the *Ft*RNAP σ^{70} -(MglA-SspA)-DNA structure, but the α_1 and α_2 N-terminal domains (α NTDs) are well resolved and the data clearly show only one form of the *Ft*RNAP core, which contains a specific α_1 - α_2 heterodimer with α_1 contacting β and α_2 interfacing with β' (Figures S1B and S1C).

The *Ft* α NTDs from each α subunit make distinct contacts important for formation of the RNAP core, burying 2,670 Å² of protein surface from solvent. The α_1 subunit uses residues that are not conserved in α_2 to form a network of selective hydrogen bonds with the β subunit (Figures S1B and S1C). Specific contacts to β are provided from α_1 residues Asn137, Lys76, and Asp174 to β residues Glu729, Val773, and Arg823 (Figure S1C). Residues in α_2 , Arg183, Thr188, Phe174, and Tyr47 which are not found in α_1 , contact β' residues Glu411, Glu441, the C β of Glu529, and the carbonyl atom of Arg533, respectively (Figure S1B). Interestingly, α_2 also contributes a few contacts to the β subunit; residues Arg31, Tyr35, Phe39, and Asn43 interact with β residues Glu822, Arg1228, Thr1229, and Lys1231, respectively (Figure S1B). Thus, the two α subunits in *Ft* are not interchangeable and form a selective heterodimer that makes specific interactions with β and β' , critical for correct *Ft* RNAP core assembly.

MglA-SspA Contacts with *Ft*RNAP σ^{70}

The mechanism by which MglA-SspA interacts with *Ft*RNAP σ^{70} has been unknown. One hypothesis has been that because MglA-SspA is a heterodimer, it would interact with the distinct subunits of the *Ft*RNAP α heterodimer (Rohlfing and Dove, 2014; Cuthbert et al., 2017). However, in the *Ft*RNAP σ^{70} -(MglA-SspA)-DNA structure, the MglA-SspA heterodimer docks opposite the α heterodimer and near the promoter DNA binding region (Figures 1A and 1B; Figures S2A–S2G). The extensive interactions between MglA-SspA and RNAP are consistent with it existing as a constitutive subunit of the polymerase (Ramsey et al., 2015). MglA-SspA interacts with the RNAP β' subunit but primarily interfaces with σ^{70} (which dissociates during

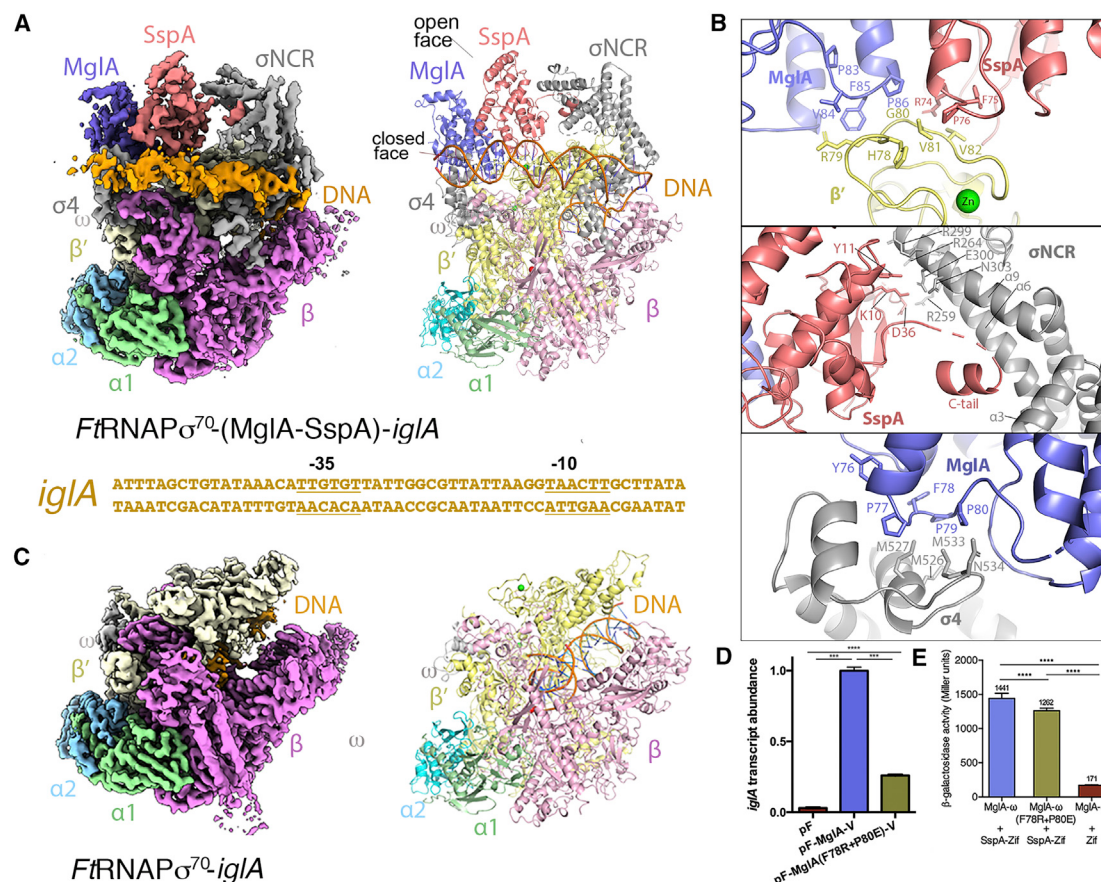


Figure 1. Structures of *FtrNAP* σ^{70} -(MgIA-SspA)-DNA and *FtrNAP* σ^{70} -DNA Complexes

(A) *FtrNAP* σ^{70} -(MgIA-SspA)-*igIA* promoter DNA cryo-EM structure. Left, density map with protein subunits and DNA labeled. Right, ribbon diagram of the complex (obtained in the absence of ppGpp). Below the structure is the *igIA* promoter sequence used in the structure with -35 and -10 elements labeled.

(B) Close-up views of the three main MgIA-SspA and *FtrNAP* σ^{70} contact points.

(C) *FtrNAP* σ^{70} -*igIA* promoter DNA complex. Left is the density map, and right is the corresponding ribbon diagram; density is absent for promoter DNA and σ^{70} in this structure.

(D) qRT-PCR data showing the relative abilities of VSV-G tagged WT and mutant MgIA, supplied by vectors pF-MgIA-V and pF-MgIA(F78R-P80E)-V respectively, to promote *igIA* expression in Δ *mglA* mutant cells. Transcripts were normalized to *tul4*, whose expression is independent of MgIA. pF is an empty vector control. Error bars represent standard deviations of the mean. Statistical significance was assessed in Prism using a two-tailed t test assuming equal variance; ****p* < 0.001, *****p* < 0.0001. Experiments were performed with biological triplicate samples at least twice, and data from a representative experiment are shown.

(E) Bacterial two-hybrid assay of the ability of MgIA(F78R-P80E)- ω to interact with the SspA-Zif fusion protein. Assays were performed with cells of the *E. coli* reporter strain KDZif1 Δ Z containing compatible plasmids directing the IPTG-controlled synthesis of the indicated proteins. Cells were grown in LB supplemented with IPTG (50 μ M) and then assayed for β -galactosidase activity. Statistical significance was assessed in Prism using one-way ANOVA with Tukey's multiple comparisons; *****p* < 0.0001. Experiments were performed with biological triplicate samples at least twice, and data from a representative experiment are shown.

transcription elongation), consistent with the heterodimer specifically forming a complex with the *FtrNAP* σ^{70} holoenzyme. Indeed, the structure shows that the function of the MgIA-SspA complex as a transcription activator does not involve DNA binding. Rather, the heterodimer stabilizes σ^{70} contacts with the RNAP core and promoter DNA, permitting the σ^2 and σ^4 domains, which are conserved with the corresponding *E. coli* σ^{70} domains, to bind the -10 and -35 promoter elements, respectively. The *igIA* promoter bound in this complex adopts an open conformation as ascertained by the presence of a transcription bubble (Figure 1A).

The *FtrNAP* σ^{70} -(MgIA-SspA)-DNA structure reveals three contact points between MgIA-SspA and the *FtrNAP* σ^{70} holoen-

zyme. The interface involving the β' subunit includes interactions from both MgIA and SspA. In this interface, the so-designated "closed face" of the MgIA-SspA heterodimer, which includes regions 83-PVFP-86 and 74-RFP-76 of MgIA and SspA, respectively, interacts with β' residues 77-KHRGVV-82 (Figure 1B, top panel; Video S1). The remaining contact surfaces are between MgIA-SspA and σ^{70} . Each subunit of the MgIA-SspA heterodimer interacts with a different region of σ^{70} , thereby stabilizing the extended form of σ^{70} on the DNA. SspA interacts with the σ^{70} nonconserved region (σ NCR) and MgIA contacts the σ^4 domain (Figure 1A). The σ NCR, which is located between σ^{70} regions 1 and 2, is found in larger, type 1 σ proteins such as σ^{70} (Feklistov et al., 2014). SspA contacts two regions of the

σ NCR. One interaction interface includes SspA residues Lys10 and Tyr11 contacting σ NCR residues Glu300, Asn303, and Arg299 and SspA residue Asp36 making electrostatic interactions with σ NCR residue Arg259 (Figure 1B, middle panel). The second, more extensive contact region involves the SspA C-terminal tail (C-tail), residues 202–209 (Video S1). These SspA residues, which are disordered in MglA-SspA crystal structures (Cuthbert et al., 2017), adopt a helical structure in the RNAP complex that docks into a cavity between σ NCR helices α 3, α 6, and α 9 (Figure 1B, middle panel). With the exception of the SspA C-tail, there are no large structural changes in MglA-SspA upon *Ft* RNAP σ^{70} binding.

The third contact surface between MglA-SspA and *Ft* RNAP σ^{70} involves interactions between MglA and σ^{70} . In this interface, MglA makes multiple contacts to the σ 4 region, which recognizes the –35 element of the promoter via a helix-turn-helix (HTH) motif. MglA residues 76–YPFPP–80 form a network of contacts with σ 4 helices α 2 and α 3 and the loop connecting the two helices. In particular, MglA residue Phe78 binds near a shallow hydrophobic cavity within this long loop composed of Met526, Met527, and Met533. MglA residue Pro80 also inserts into this hydrophobic cleft and interacts with residues Met533, Thr535, and the C β of Asn534 (Figure 1B, bottom panel; Video S1). The structure explains why MglA-SspA might not associate with the *Ft* RNAP σ^{32} holoenzyme, as the smaller *Ft* σ^{32} protein does not contain an NCR, and the σ 4 loop region contacted by MglA differs in sequence and length between the two *Ft* σ factors.

PigR-Independent Virulence Role for MglA-SspA

Our *Ft* RNAP σ^{70} -(MglA-SspA)-promoter DNA structure indicates that MglA-SspA functions to facilitate the interaction of σ^{70} with both the core RNAP and promoter DNA, leading to open complex formation. Notably, the promoter DNA used to obtain the structure did not contain a pre-formed bubble (from non-complementary DNA) (Figure 1A), and thus promoter opening was mediated *de novo* by the *Ft* RNAP σ^{70} -(MglA-SspA) complex. This finding suggests the possibility that MglA-SspA might activate *Ft* transcription in a PigR-independent manner. To investigate this possibility further, we next obtained a 3.0 Å cryo-EM structure of *Ft* RNAP σ^{70} with *iglA* promoter DNA in the absence of MglA-SspA (Figures S3A–S3G; Table S1). Strikingly, in this structure, density was absent for both σ^{70} and the promoter DNA, consistent with data showing FPI transcription is dependent on MglA-SspA (Charity et al., 2007; Rohlfing and Dove, 2014) (Figure 1C). Interestingly, DNA and some RNA density were present in the active site channel of the RNAP, presumably carried over from the two-step purification of the holoenzyme from *Francisella* cells. Hence, the *Ft* RNAP σ^{70} purified in this manner captures the RNAP in complex with non-specific DNA, and the addition of *iglA* promoter DNA is insufficient to allow promoter binding or formation of a stable open promoter complex by the *Ft* RNAP σ^{70} without MglA-SspA.

Examination of the *iglA* promoter shows that while it harbors a strong –10 element, required for binding the σ 2 domain, the –35 element is not optimal, which suggests a role for the MglA- σ 4 interaction in the formation of a stable transcription initiation complex. To test further the importance of MglA contacts to

σ 4 in transcription, we constructed an MglA mutant with substitutions F78R and P80E, as the structure shows these substitutions should impede MglA-SspA binding to *Ft* σ 4. qRT-PCR analyses revealed that the VSV-G tagged MglA(F78R-P80E) mutant in LVS Δ *mglA* mutant cells significantly impaired transcription activation of the *iglA* gene compared to wild-type (WT) VSV-G tagged MglA (Figure 1D). Notably, the MglA(F78R-P80E) mutant interacts with SspA as well as the WT MglA protein in a bacterial two-hybrid assay indicating that the impact on expression was not due to impaired MglA-SspA heterodimer formation (Figure 1E). MglA-SspA binding to *Ft* RNAP σ^{70} does not induce structural changes in polymerase subunits, nor does the heterodimer directly interact with σ 2 to influence melting of the –10 element. Therefore, the mechanism by which MglA-SspA favors open promoter complex appears similar to that of the *Thermus thermophilus* TAP activator, which promotes the generation of the *Tt* RNAP open complex by forming stabilizing adhesive protein-protein interactions with the polymerase (Feng et al., 2016). Unlike *Tt*TAP, MglA-SspA appears to act as a σ^{70} activator by making direct contacts to σ^{70} and the RNAP β' subunit as well as facilitating the interaction of σ^{70} with promoter DNA.

The structural data showing that the presence of MglA-SspA at the *iglA* promoter is required to drive open complex formation suggested that the heterodimer alone might impact *Ft* transcription of virulence genes even in the absence of PigR (Ramsey et al., 2015). To test this hypothesis and obtain information on genes that may be regulated by MglA-SspA but not PigR, we performed RNA-seq studies (STAR Methods). In these experiments, the transcriptomes of LVS WT cells, LVS Δ *mglA* mutant cells, and LVS Δ *pigR* mutant cells were analyzed using biological triplicate samples. These studies revealed that the MglA-SspA regulon (as defined by genes whose expression changes by a factor of 2 or more in Δ *mglA* mutant cells compared to WT with an adjusted p value < 0.05) is larger than previously described (Figures 2A and 2B; Table S2). They also revealed that although most of the PigR-regulated genes (those whose expression changes by a factor of 2 or more in Δ *pigR* mutant cells compared to WT with an adjusted p value < 0.05) are also controlled by MglA-SspA, many MglA-SspA regulated genes are not regulated by PigR (Figure 2A–B).

Conserved SspA-Binding Mode with RNAP σ^{70}

The finding that MglA-SspA alone controls transcription of multiple genes led us to question whether homodimeric SspA proteins might utilize a similar mechanism. SspA proteins play key roles in starvation responses and virulence in multiple bacteria, including *E. coli*, *Yersinia enterocolitica*, and *Neisseria gonorrhoeae* (Ishihama and Saitoh, 1979; Williams et al., 1994; De Reuse and Taha, 1997; Badger and Miller, 1998; Hansen et al., 2003). Interestingly, like MglA-SspA, the transcription regulation functions of *Ec*SspA transcription are mediated through the RNAP σ^{70} holoenzyme (Hansen et al., 2003). In addition, the region encompassing *Ec*SspA residues 83–FPHPP–87, which has been shown to be key for its transcription function (Hansen et al., 2005a, 2005b), corresponds to MglA residues 76–YPFPP–80 and *Ft*SspA residues 74–RFPAP–78, which our cryo-EM structure revealed contact σ 4 and β' (Figure 1A). As

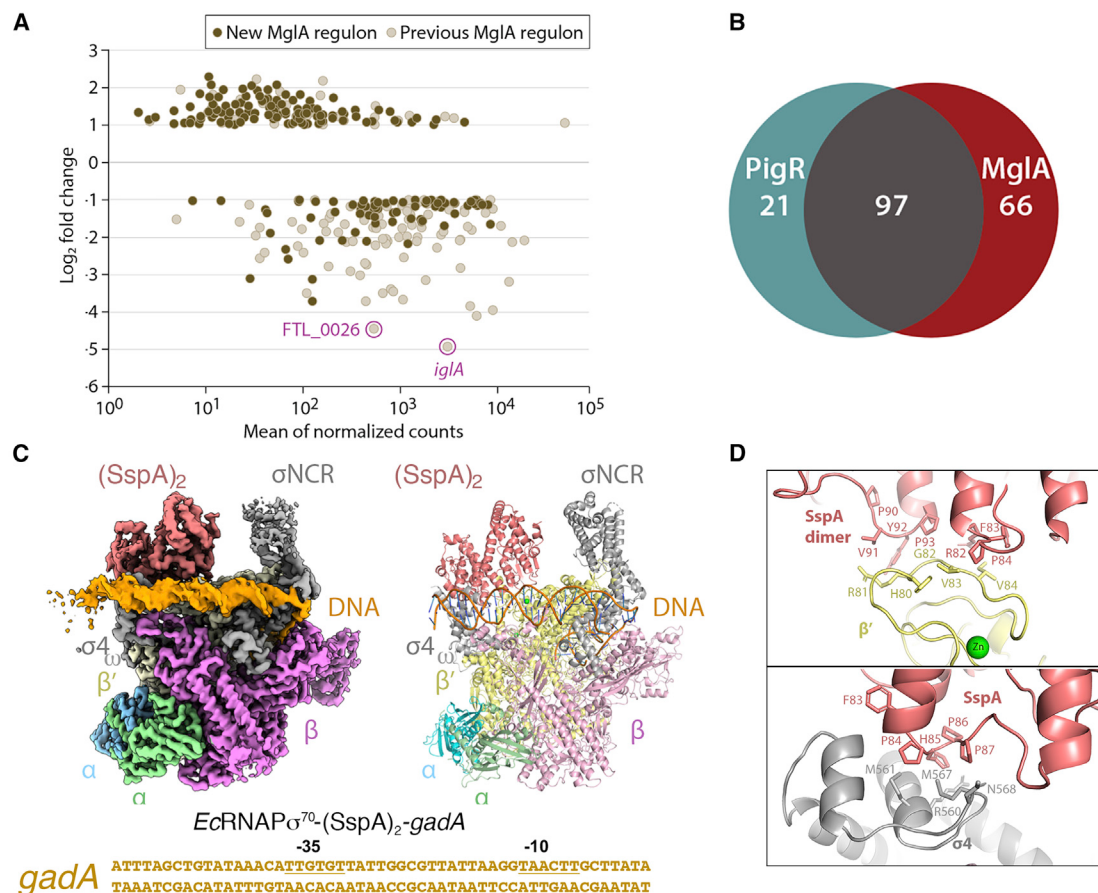


Figure 2. MglA-SspA Mediated-PigR Independent Transcription and General Binding Mode of SspA Proteins for RNAP_{σ70}

(A) DESeq2 was used to conduct differential gene expression analysis between WT and Δ mglA mutant RNA-seq libraries. The graph shows log₂ fold change in transcript abundance in Δ mglA compared to WT cells. Previously described MglA-controlled transcripts (such as *iglA* and *FTL_0026*) are in beige (with the *iglA* and *FTL_0026* transcripts highlighted). Newly identified MglA-transcripts are in brown. Analyses were done using biological triplicates.

(B) Venn diagram indicating MglA-regulated genes that are controlled by PigR and those that are not. Most genes regulated by PigR are also regulated by MglA-SspA but MglA-SspA regulates multiple genes independently of PigR.

(C) Cryo-EM EcRNAP_{σ70}-(SspA)₂-gadA DNA structure. Left is cryo-EM map, right is ribbon diagram, and below is the sequence of the *gadA* promoter used in the structure, with -35 and -10 elements labeled. The complex is shown in the same orientation as Figure 1A.

(D) Close-up views of contact points between EcSspA and EcRNAP_{σ70}.

the *E. coli* protein is the best studied of the SspA homologs, we turned to cryo-EM studies to examine the interaction of the EcSspA homodimer with EcRNAP_{σ70}.

EcSspA was shown to be important in *E. coli* acid stress resistance during stationary phase; in a microarray study using an SspA mutant, acid-resistance glutamate-dependent (*gad*) genes were downregulated, with the greatest effect observed for *gadA* (Hansen et al., 2005a, 2005b). Thus, we obtained a 3.2 Å cryo-EM structure of the *E. coli* RNAP_{σ70}-(SspA)₂ complex bound to the *gadA* promoter. The structure shows that the EcSspA homodimer forms essentially the same complex with EcRNAP_{σ70} as MglA-SspA does with FtRNAP_{σ70}. Specifically, EcSspA contacts the σ4 domain of σ₇₀ and the RNAP β' subunit using corresponding surfaces, and the structure adopts an open promoter complex as in the FtRNAP_{σ70}-(MglA-SspA)-DNA structure (Figure 2C; Figures S4A–S4G; Table S1). The SspA dimer is also juxtaposed near the σNCR but

unlike FtMglA-SspA does not contact the σNCR in this complex. Similar to the MglA-SspA complex with FtRNAP_{σ70}, residues 74-PHP-76 of one SspA subunit contacts the β' subunit, while these residues in the other subunit of the EcSspA homodimer interact with the open loop in σ4 (Figure 2D). The σ4 loop of the Ecσ₇₀ is different from other Ecσ factors, explaining the preference of SspA proteins for the RNAP_{σ70} holoenzyme. Residues 90-PVYP-93 of one SspA subunit also interact with Ecβ', analogous to the interaction between MglA residues 83-PVFP-86 and Ftβ'. The N-terminal helix of Ecβ' contributes additional contacts to EcSspA, as does the C-terminal helix of ω, which is disordered in most RNAP structures (Figure S5). Similar to the Ft *iglA* promoter, Ec *gadA* lacks an optimal -35 element and EcSspA appears to stabilize the RNAP σ4 domain onto promoter DNA. Notably, the phage P1 *P*_{late} promoter, which requires EcSspA for transcription activation, lacks a recognizable RNAP_{σ70} -35 element (Hansen et al., 2003).

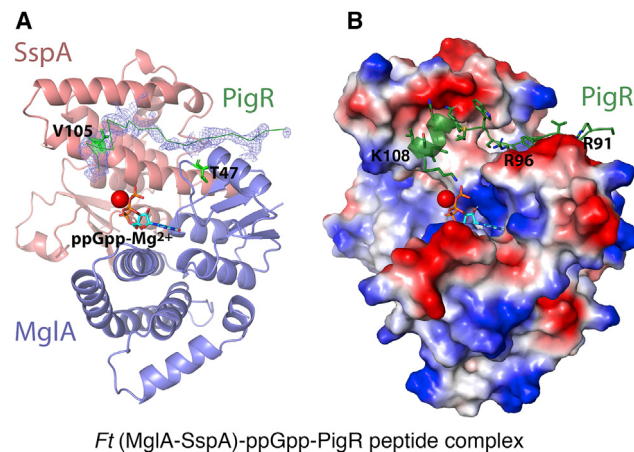


Figure 3. X-Ray Structure of (MglA-SspA)-ppGpp-PigR Peptide Complex

(A) Ribbon diagram of MglA-SspA with $2F_o - F_c$ electron density (0.65σ) shown before the PigR peptide was added. Shown also are the locations of SspA V105 and MglA T47 (green sticks). The PigR peptide is shown as a green ribbon, ppGpp as sticks, and Mg^{2+} as a red sphere.

(B) Electrostatic surface of MglA-SspA with the PigR peptide depicted as a green ribbon. Positive and negative regions are blue and red, respectively.

Ft(MglA-SspA)-ppGpp-PigR Crystal Structure

A pivotal step in *Ft* pathogenicity is escape of *Francisella* cells from the phagosome into the cytosol. This step depends on transcription of FPI encoded genes, the full activation of which requires $RNAP^{\sigma^{70}}$ -(MglA-SspA), the *Francisella*-specific protein PigR and the stress alarmone ppGpp (Lauriano et al., 2004; Brotcke et al., 2006; Charity et al., 2007, 2009; Brotcke and Monack, 2008; Rohlfing and Dove, 2014; Cuthbert et al., 2017). MglA-SspA binds ppGpp in the absence of PigR, but ppGpp is required for high affinity binding of the C-terminal 22 residues of PigR to MglA-SspA (Cuthbert et al., 2017). Although unknown, the structure of the 111 residue PigR protein is predicted to contain a MerR-like winged HTH (wHTH) (Heldwein and Brennan, 2001; Brown et al., 2003) followed by the C-terminal tail, composed of residues 90–111. To delineate the details of the PigR interaction with (MglA-SspA)-ppGpp, we solved the (MglA-SspA)-ppGpp-PigR(90–111) crystal structure to 2.95 Å resolution (Figure 3A; Table S3; STAR Methods). In the structure, the vast majority of contacts to the ppGpp are from the *Ft*-specific MglA protein, which shares only ~25 sequence identity with SspA homodimeric proteins. Consistent with this, previous studies showed that MglA-SspA binds ppGpp with a K_d of 12 μ M, while the *Ec*SspA homodimer showed very weak binding to ppGpp (Cuthbert et al., 2017). The (MglA-SspA)-ppGpp-PigR structure reveals that the PigR C-tail binds within the ppGpp binding pocket or “open face” (Figure 1A). The N-terminal region of the PigR fragment forms a strand that extends into the (MglA-SspA)-ppGpp binding pocket, and the remaining PigR residues fold into an amphipathic helix (Figures 3A and 3B).

Previous data showed that mutations in SspA and MglA residues that directly contact ppGpp impaired PigR binding, in line with data showing that ppGpp is required for high affinity PigR binding (Rohlfing and Dove, 2014; Cuthbert et al., 2017).

However, SspA(V105E) and MglA(T47A) substitutions do not impact ppGpp binding but significantly impede the interaction of PigR with MglA-SspA (Rohlfing and Dove, 2014). Consistent with these data, our structure reveals that SspA residue Val105 and MglA residue Thr47 line the PigR binding pocket, with the hydrophobic face of the PigR C-terminal helix making extensive interactions with SspA residue Val105 (Figure 3A). Complementary electrostatic interactions also occur between PigR residues Arg91 and Arg96 and the negatively charged surface of the $\alpha 2$ helix of MglA (Figure 3B). The ppGpp binding pocket of MglA-SspA and the C terminus of the PigR helix (sequence: KAKS) docked in this pocket are both highly positively charged; PigR helix residue Lys108 is positioned to contact ppGpp. Thus, ppGpp appears to facilitate PigR binding to MglA-SspA via direct contacts to ppGpp as well as by preventing unfavorable charge-charge clashes between PigR C-terminal residues and the basic ppGpp binding pocket in MglA-SspA.

FtRNAP σ^{70} -(MglA-SspA)-ppGpp-PigR Complex

Our (MglA-SspA)-ppGpp-PigR crystal structure revealed that PigR is anchored to the (MglA-SspA)-ppGpp complex via its C-terminal tail. To understand how PigR enhances FPI transcription in the context of the full *Ft* RNAP complex, we obtained a 4.5 Å resolution cryo-EM structure of the *Ft* RNAP σ^{70} -(MglA-SspA)-ppGpp-PigR-*igI*A promoter DNA complex (Figure 4A; Figures S6A–S6H). In this complex, MglA-SspA is bound in the identical location as in the *Ft* RNAP σ^{70} -(MglA-SspA)-DNA structure. Density for ppGpp in the MglA-SspA pocket was evident. Unlike the *Ft* RNAP σ^{70} -(MglA-SspA)-DNA complex, in this structure density is also present for the upstream DNA, including the DNA binding site for PigR, the PigR response element (PRE), and DNA upstream and downstream of the PRE (Figure 4A). Density for the PigR wHTH near the PRE site was too weak to be readily modeled. But there are ~10 residues between the MglA-SspA binding region of PigR and the wHTH-PRE binding site, which would allow the connection of the two PigR regions (Video S2). These data indicate that PigR binds the (MglA-SspA)-ppGpp complex as a monomer, bringing its wHTH motif within proximity of the PRE.

Unexpectedly, in the *Ft* RNAP σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA structure we also observed two globular regions of density contacting the DNA upstream and downstream of the PRE. Both these PRE-flanking DNA sites are very AT rich (Figure 4A). Strikingly, examination of the DNA sequences of promoters known to be regulated by PigR, such as the *igI*A, *pdp*A, and *FTL_0026* promoters, revealed that all contain AT-rich regions flanking their PRE sites (Figure 4B). Studies in model bacteria have revealed the importance of AT-rich regions located upstream of the -35 element (Newlands et al., 1991; Ross et al., 1993; Estrem et al., 1998; Ross and Gourse, 2009), termed Upstream (UP) elements, as contact sites for the C-terminal domains of the α subunits (α CTDs) of RNAP (Newlands et al., 1991; Ross et al., 1993; Estrem et al., 1998; Ross and Gourse, 2009). Consistent with the assignment of these densities in our structure as *Ft* α CTDs, lower contoured density maps revealed a connection between the densities bound to the AT-rich DNA sites and the *Ft* RNAP α NTDs, and the *Ec* α CTD structure could be readily docked into these densities (Figure 4C). At the current resolution,

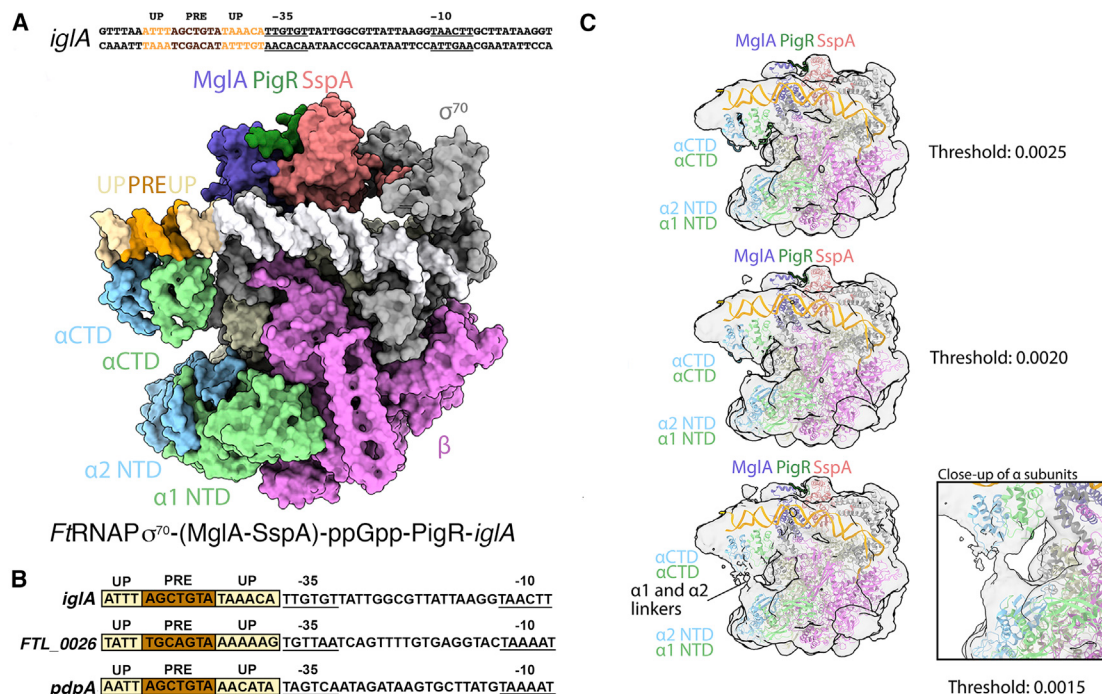


Figure 4. Cryo-EM Structure of the *FtrNAP* σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA Complex

(A) Cryo-EM structure of the *FtrNAP* σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA complex. The PigR C-tail is modeled from the crystal structure. Above the structure is the full promoter sequence used in the structure, with promoter elements labeled. (B) DNA sequences of PigR controlled genes showing just the promoter regions. -35 and -10 elements are indicated, PRE elements are colored brown, and AT-rich UP elements are light yellow. (C) Images of the *FtrNAP* σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA density map at various contour levels. Right shows a close up of the linker region, which leaves ambiguous which NTD is linked to which CTD.

however, we were unable to assign which α CTD, α 1 or α 2, is bound to which UP element (upstream or downstream of the PRE). But both α CTDs are positioned on the DNA as in previously observed α CTD-UP structures with conserved DNA-binding residues proximal to the DNA (Benoff et al., 2002; Ross and Gourse, 2009). Notably, density was not visible for the α CTDs in our *FtrNAP* σ^{70} -*iglA*, *EcrNAP* σ^{70} -(SspA)₂-*gadA*, or *FtrNAP* σ^{70} -(MglA-SspA)-*iglA* structures indicating that PigR recruits the α CTDs to the DNA. Precisely how PigR facilitates UP element binding by the *Ft* α CTDs is currently unclear due to the weak density of the PigR wHTH. However, the proximity of the PRE to the UP elements suggests that PigR may recruit the α CTDs through direct protein-protein contacts. The possibility that distinct α CTDs may bind separate surfaces of PigR and be directed to different UP elements would provide another rationale for the presence of two different *Ft* α subunits, in addition to the importance of the *Ft* α NTDs in RNAP core assembly. Alternatively, PigR binding to the PRE may help order and stabilize the upstream DNA, damping its thermal motions, thereby favoring α CTD binding.

Our structural data suggest that in the presence of ppGpp, PigR binds to MglA-SspA and PRE DNA to enable the recruitment to and/or stabilization of the α CTDs on the PRE-adjacent UP elements, forming stable initiation complexes (Figure 4A; Video S2). To test this hypothesis, we generated reporter con-

structs in which the *iglA* and *FTL_0026* promoters were transcriptionally fused to *lacZ* on a chromosomal integration vector, thus allowing us to assess the effects of mutations within the proposed UP promoter elements in their native contexts (Ramsey et al., 2015). Using these constructs, we showed that substitutions of bps within the PRE as well as within the proposed UP elements downstream of the PRE in these promoters significantly impaired or abrogated PigR-dependent transcription (Figures 5A–5D). Thus, these data support the structural model demonstrating the importance of UP elements in PigR-mediated *Ft* virulence gene activation.

DISCUSSION

Francisella tularensis is the causative agent of the zoonotic disease tularemia (Barker and Klose, 2007). Due to its low infectious dose, ability to be transmitted to humans via multiple routes, and the potential to cause life-threatening infections, *Ft* has been developed as a bioterrorist agent. Indeed, the potential of *Ft* to be exploited deliberately to cause disease was recognized by the WHO as early as the 1970s (Dennis et al., 2001; Maurin, 2015). In addition, there is growing concern over the increase in reported outbreaks of tularemia in Europe over the past two decades (Eliasson and Bäck, 2007; Faber et al., 2018; Schroll et al., 2018). While *Ft* can infect more than 200 different

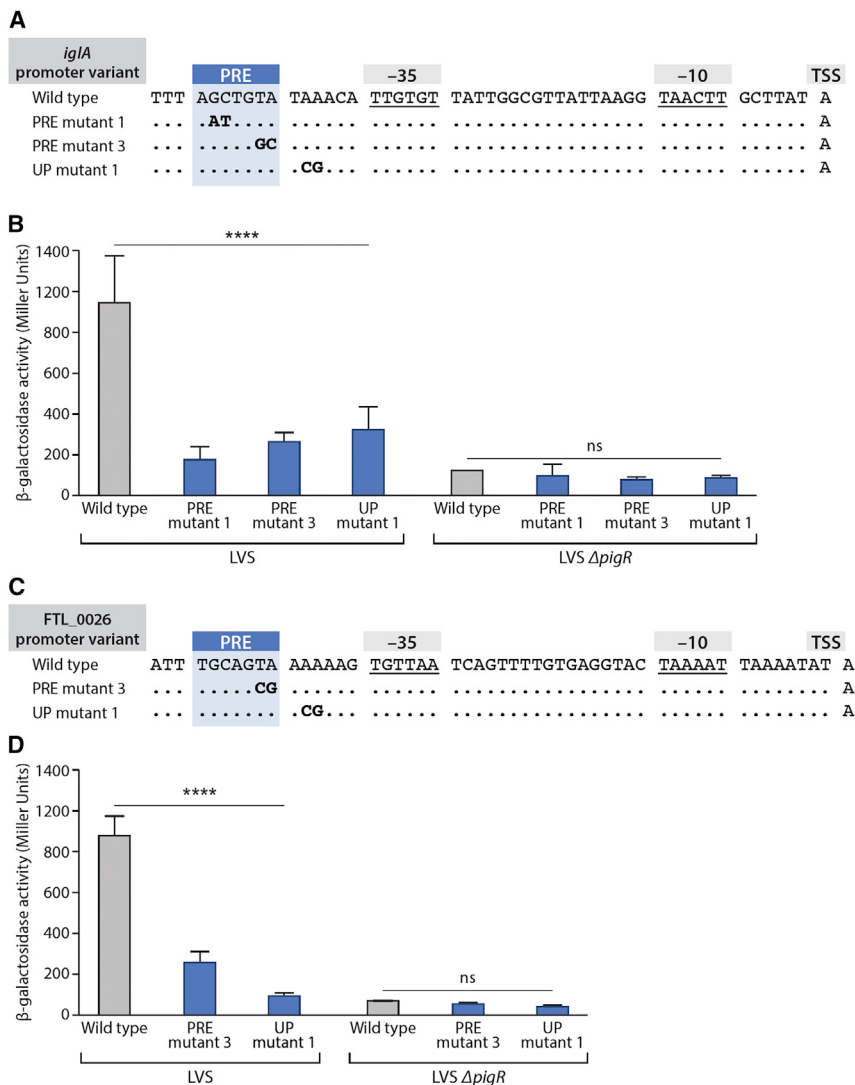


Figure 5. Test of Importance of UP Elements in Activity of PigR-Regulated Promoters

(A) Mutations were made in the PRE (PRE mutant 1 and PRE mutant 3 in bold) as well as in the UP element downstream of the PRE (UP mutant 1, in bold). Wild-type is also labeled. In PRE mutant 3, the TA bps of the PRE at positions 6 and 7 are mutated to CG. In UP mutant 1, bps 2 and 3 in the UP element downstream of the PRE were changed from AA to GC. All mutations impaired PigR-dependent regulation.

(B) Quantification of *igIA-lacZ* expression in LVS wild-type (LVS) and $\Delta pigR$ mutant (LVS $\Delta pigR$) cells containing the indicated *igIA* promoter variants (x axis) by β -galactosidase assay (Miller units). Promoter variants linked to a *lacZ* reporter gene were integrated into the *FTL_0111* locus. Statistical significance was assessed in Prism using one-way ANOVA with Tukey's multiple comparisons; ****p < 0.0001. Experiments were performed with biological triplicate samples at least twice, and data from a representative experiment are provided.

(C) Double mutations were made in the PRE (PRE mutant 3, as indicated in bold) and in the UP element downstream of the PRE (UP mutant 1, as indicated in bold). Wild-type is labeled. In PRE mutant 3, the TA bps of the PRE at positions 6 and 7 are mutated to CG. In UP mutant 1, bps 2 and 3 in the *FTL_0026* promoter downstream UP element were changed from AA to GC. All mutations impaired PigR-dependent regulation.

(D) Quantification of *FTL_0026-lacZ* expression in LVS wild-type (LVS) and $\Delta pigR$ mutant (LVS $\Delta pigR$) cells containing the indicated *FTL_0026* promoter variants (x axis) by β -galactosidase assay (Miller units). Promoter variants linked to a *lacZ* reporter gene were integrated into the *FTL_0026* locus. Statistical significance was assessed in Prism using one-way ANOVA with Tukey's multiple comparisons; ****p < 0.0001. Experiments were performed with biological triplicate samples at least twice, and data from a representative experiment are shown.

organisms and multiple cell types, it primarily invades macrophages, where it is initially contained within a phagocytic vacuole. A necessary step in *Ft* virulence is escape from the phagosome into the cytosol, where the bacterium is able to rapidly divide and spread to surrounding tissues. What controls this step was unclear until a spontaneous *Ft* mutant with altered colony formation and inability to replicate in macrophages was identified. The *mgIA* gene was subsequently identified as a mutated locus key for this step (Baron and Nano, 1998). Later studies revealed that, in addition to the MglA protein, the *Ft* SspA and PigR proteins and the stress molecule ppGpp are essential for robust transcription of *Francisella* pathogenicity island (FPI) genes (Lauriano et al., 2004; Charity et al., 2007, 2009; Brotcke and Monack, 2008; Rohlfing and Dove, 2014; Cuthbert et al., 2017).

Unlike many bacteria, which employ specialized σ factors for activation of virulence genes, *Ft* utilizes the *Ft* RNAP σ^{70} holoenzyme, which contains the housekeeping σ^{70} protein, to transcribe FPI genes (Kazmierczak et al., 2005; Ramsey et al.,

2015). As noted, MglA was established early on as a central activator of *Ft* virulence, and our studies have revealed that MglA harbors an SspA-like fold and specifically heterodimerizes with *Ft*SspA. MglA-SspA interacts with *Ft* RNAP σ^{70} to form a virulence-specialized RNAP holoenzyme. Here we undertook cryo-EM, X-ray crystallography, and genetic studies to elucidate the molecular underpinnings of this *Ft*-specialized virulence activating transcription machinery. Our *Ft* RNAP σ^{70} -(MglA-SspA)-DNA cryo-EM structure revealed that the MglA-SspA heterodimer does not interact with DNA, as typical transcription activators do, but instead makes extensive contacts with the RNAP σ^{70} holoenzyme that fastens the enzyme to the promoter (Lee et al., 2012; Browning and Busby, 2016). Specifically, the MglA-SspA heterodimer primarily interacts with σ^{70} and makes additional contacts to the core RNAP β' subunit. Through contacts to the σ NCR, located at the σ^{70} N-terminal region of the σ^{70} protein, and the $\sigma 4$ domain, which is located at the C terminus of σ^{70} , MglA-SspA properly orients the malleable σ^{70} to key DNA promoter elements, i.e., the -10 and -35 elements, as well as

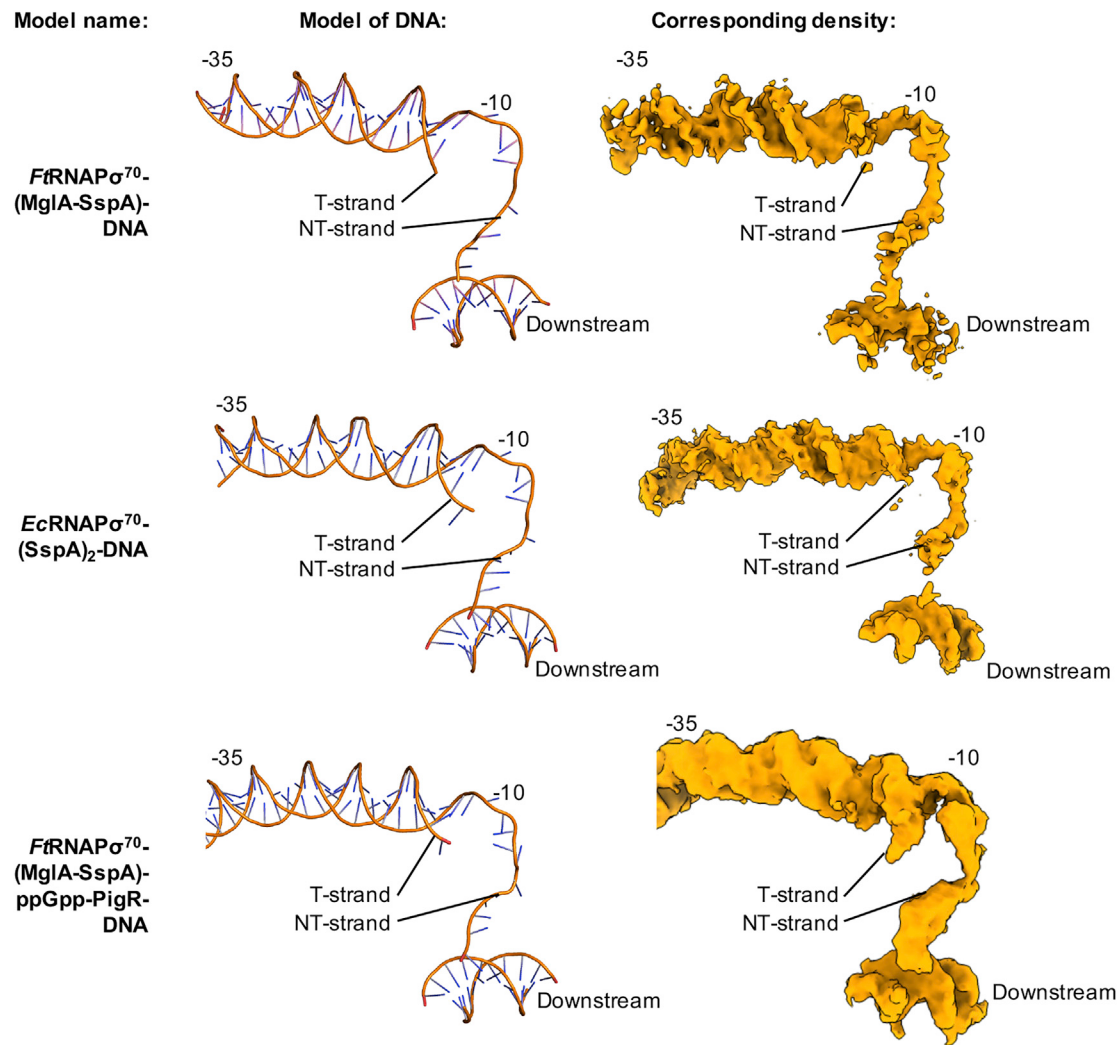


Figure 6. Open-Promoter Complexes Are Formed in *Ft*(MglA-SspA)- and *Ec*(SspA)₂-Containing Transcription Complexes
Protein subunits were removed from each model for clarity. Left shows ribbon diagrams of the DNA and right shows the corresponding DNA density.

affixes it to the RNAP core, leading to the formation of an open promoter complex (Bae et al., 2015; Murakami, 2015).

Interestingly, our structures showed that the MglA-SspA heterodimer is similar in its overall oligomeric organization and subunit structure to SspA homodimers (Cuthbert et al., 2017). Our cryo-EM analyses of an *E. coli* RNAP σ^{70} -(SspA)₂-promoter DNA structure revealed the same overall organization of SspA with the *Ec*RNAP. Moreover, both *Ft*RNAP σ^{70} -(MglA-SspA)-DNA and *Ec*RNAP σ^{70} -(SspA)₂-DNA structures form open promoter complexes (Figure 6). Promoters of the genes activated by MglA-SspA in *Ft* and by the SspA homodimer in *E. coli* are not ideal. Indeed, while they contain near-optimal -10 motifs (both contain key bps A₋₁₁ and T₋₇ found in the T₋₁₂A₋₁₁T₋₁₀A₋₉A₋₈T₋₇ consensus motif), their -35 motifs poorly match the ideal -35 consensus element. In particular, the *Ps* promoter activated by *Ec*SspA lacks any recognizable -35 element. Previous structural studies revealed that key base contacts to the -35 element are made by residues Gln414, Glu410, and Arg409 from the *Taq*

σ^A (Campbell et al., 2002). These residues, which are conserved in the *Ec* and *Ft* σ^{70} proteins, interact with bases within the T₋₃₅T₋₃₄G₋₃₃ and the C₋₃₁ bps. The -35 element of the *iglA* promoter, T₋₃₅T₋₃₄G₋₃₃T₋₃₂G₋₃₁T₋₃₀, departs from the optimal -35 motif (T₋₃₅T₋₃₄G₋₃₃A₋₃₂C₋₃₁A₋₃₀) and notably lacks the guanine of the C₋₃₁-G₋₃₁ bp, which is specifically read by Arg409 (Campbell et al., 2002). While phosphate contacts to the -35 motif are made by residues in the σ^4 domain in the *Ft*RNAP σ^{70} -(MglA-SspA)-DNA complex, base specific contacts are missing. Further, the *iglA* lacks an extended -10 motif (T₋₁₇G₋₁₆T₋₁₅G₋₁₄), which data have shown can substitute for weak -35 elements (Kumar et al., 1993; Barne et al., 1997). Analyses of promoters of genes shown to be activated by MglA-SspA in a PigR-independent manner by RNA-seq revealed that all contained poor -35 elements with a few (*FTL_1012*, *FTL_1511*, and *FTL_0361*) possibly also harboring poor matches to the -10 element; we note the -35 and -10 elements of these genes are predictions based on our annotated transcription start

site (Figure S7A). Thus, our structural data as well as genetic studies indicate MglA-SspA functions as an activator of weak promoters. This was also supported by our reporter assays, which demonstrated that the interactions between MglA-SspA and σ^4 that enable the latter domain to contact the -35 element were necessary for activation of the *iglA* promoter (Figure 1D). Future transcription studies on homodimeric SspA proteins will be needed to assess their roles in transcription at specific promoters.

Thus, based on our findings the MglA-SspA heterodimer appears to function as a σ activator, a term recently coined by Cartagena et al. to describe the *Salmonella enterica* Crl protein (Cartagena et al., 2019). The authors proposed that Crl, which is a small arc-shaped monomeric protein composed of β strands, acts as a σ^S activator through its interactions with σ^S region 2 and the RNAP β' subunit, which enable σ^S -RNAP complex formation (Cartagena et al., 2019). MglA-SspA stabilizes primarily the σ^4 region of σ^{70} along with β' , enabling σ^{70} binding to both the *Ft* RNAP core and promoter DNA. These data indicate that σ activators may comprise multiple structural families and employ distinct interactions to aid in the assembly or stabilization of σ subunits with RNAP core enzymes and promoter DNA. Given that the *Ft* RNAP σ^{70} -(MglA-SspA)-DNA structure assumed an open promoter complex, we hypothesized that this complex alone might be capable of transcription activation in a manner independent of PigR. We thus performed RNA-seq analyses to delineate and compare the MglA-SspA and PigR regulons. Although these studies indicate that MglA-SspA and PigR activate many of the same genes, they also revealed that MglA-SspA activates the expression of a number of genes independently of PigR, the roles of which in virulence remain to be elucidated.

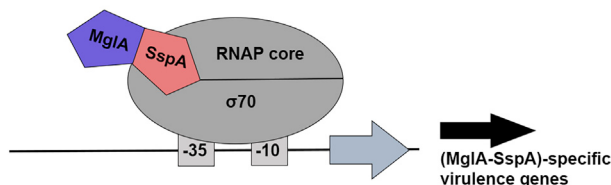
Collectively, our structural and genetic analyses indicate that the MglA-SspA heterodimer contributes to *Ft* virulence in a PigR-independent manner. Regardless, optimal activation of the FPI requires both the *Ft*-specific PigR protein and ppGpp. Unlike MglA-SspA, PigR has a conventional DNA-binding, winged HTH motif. While the wHTH of PigR places it in the MerR family of proteins, which are known to act as dimers (Heldwein and Brennan, 2001; Brown et al., 2003), our studies indicate that PigR functions as a monomer. Indeed, our (MglA-SspA)-ppGpp-PigR peptide crystal structure demonstrated that only one C-terminal tail of PigR inserts into the ppGpp binding pocket of MglA-SspA. Binding of ppGpp to MglA-SspA facilitates its interaction with the C-tail of PigR, thus revealing a second function of MglA-SspA: recruitment of an additional transcription activator. This interaction depends on the presence of ppGpp. ppGpp has emerged as a central second messenger in bacteria and shown to bind a variety of protein targets (Zhang et al., 2018; Wang et al., 2019) and crucially, to play an important role in reprogramming transcription (Kanjee et al., 2012; Krásný and Gourse, 2004; Liu et al., 2015). This latter process has been extensively studied in *E. coli*. In *E. coli* RNAP ppGpp binds to two sites: site 1, which is at the interface of the ω and β' subunits, and site 2, which is a pocket formed by the interaction of β' and the transcription factor DksA (Ross et al., 2013, 2016; Zuo et al., 2013). Binding of ppGpp to either RNAP site 1 or site 2 can inhibit transcription, but only binding of ppGpp to site 2 can stimulate

transcription (Ross et al., 2013, 2016). Notably, the amino acid residues that form this *Ec* RNAP pocket are not found in *Ft* RNAP, and DksA does not appear to be present in *Ft*. Thus, *Ft* employs ppGpp as a potentiator of virulence through its interaction with the MglA-SspA heterodimer. The ppGpp-binding pocket of MglA-SspA is also distinct from the ppGpp binding sites that have been characterized in multiple proteins from other bacteria (Kanjee et al., 2012).

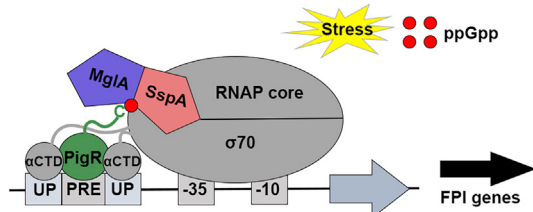
To understand how PigR collaborates with *Ft* RNAP σ^{70} -(MglA-SspA) to drive FPI activation, we solved the structure of the *Ft* RNAP σ^{70} -(MglA-SspA)-ppGpp-PigR-*iglA* DNA complex (Figure 4). Because PigR functions as a monomer, its specific binding to the PRE, on its own, would likely contribute only weakly to the affinity of the RNAP complex for regulated promoters. However, the structure uncovered an unexpected role for PigR: the recruitment of the *Ft* RNAP α CTDs to DNA sites that directly flank the PRE. Examination of these DNA sites revealed they are AT rich. Such AT-rich sites upstream of the -35 element have been termed upstream (UP) elements and were discovered by Ross and Gourse as key contributors to transcription activation in *E. coli* through their interactions with the α CTDs of RNAP (Ross et al., 1993, 2003). *Ft* is unusual in encoding two distinct α RNAP subunit genes. To our knowledge, the only other bacterium with two α subunits that has been characterized experimentally is *Streptomyces granaticolor*, where the presence of two α subunits with molecular weights of 40 kDa and 43 kDa was reported (Najmanová et al., 2000). However, it was revealed that the smaller α subunit is produced from proteolytic cleavage that occurred during sporulation and was not encoded by a separate gene (Najmanová et al., 2000). Our structures have revealed that a distinct *Ft* α NTD heterodimer is critical for proper assembly of β and β' into a functional RNAP core (Figure S1B). The presence of two α CTDs in *Ft* also presents the opportunity for their different surfaces to be utilized in interactions with, for example, distinct regions of PigR, or potentially other transcriptional activators, thus endowing additional versatility in *Ft* α CTD function.

Early studies suggested that α CTD interactions with UP elements function to not only enhance binding of RNAP to promoters but also to stimulate isomerization from the closed to the open complex (Rao et al., 1994; Saecker et al., 2011). However, recent single molecule experiments from the Gelles lab examining *E. coli* RNAP transcription from the *rrnB* P1 promoter revealed that α CTD-UP interactions stimulated RNAP binding, with no effect on open promoter formation (Mumm et al., 2020). Our *Ft* RNAP σ^{70} -(MglA-SspA)-DNA complex revealed that MglA-SspA alone facilitated open promoter complex formation when in complex with *Ft* RNAP σ^{70} and thus is consistent with the Gelles study in that the α CTD-UP element interaction does not appear to be required for open promoter complex formation. The addition of the *Ft* α CTD-UP contacts, which occurs only in the presence of PigR, thus likely functions to enhance RNAP binding. Notably, these are the first data revealing the utilization of UP elements in *Ft* transcription. Underscoring the importance of this interaction, the known PigR regulated promoters all contain AT-rich sites flanking their PRE sites (Figure 4B). Comparison of our RNA-seq data of additional genes regulated in a PigR-dependent manner showed

MgIA-SspA (PigR-independent) regulated promoters



(MgIA-SspA)- and PigR-dependent promoters

**Figure 7. Two Roles of MgIA-SspA in *Ft* Virulence Activation**

Schematic model for *Ft* virulence gene activation. Upper, PigR-independent genes are activated by MgIA-SspA stabilizing σ^{70} and DNA interactions with *Ft* RNAP. Lower, PigR-dependent genes involve PigR binding to (MgIA-SspA)-ppGpp to recruit α CTDs to UP elements.

that, again, all have relatively weak predicted -35 elements (Figure S7B). However, the most striking findings from MEME analyses of these PigR regulated promoters were that not only is the PRE conserved, but also that the AT-rich motifs that directly flank the PRE are remarkably conserved (Figure S7C). These data indicate that α CTD recruitment to the *Ft* UP elements is a key and previously unrecognized mechanism for PigR-mediated activation of virulence genes.

In conclusion, our studies have elucidated the molecular basis for virulence activation of the dangerous pathogen *Ft*. Specifically, these collective studies have unveiled MgIA-SspA as the linchpin in *Ft* virulence, whereby this heterodimer employs two roles in the activation of virulence genes: one involving a σ^{70} activation function and a second mechanism whereby binding of a small molecule alarmone, ppGpp, produced during infection, to the MgIA-SspA heterodimer anchors a unique transcription activator, PigR, to virulence promoters, which in turn allows the recruitment and stable binding of the two *Ft* α CTDs to previously uncharacterized UP elements (Figure 7). These data thus provide the foundation for the future development of specific therapeutics that directly target virulence activation in *Francisella tularensis*.

During the review process of this manuscript, Wang et al., (2020) published a 3.68 Å resolution cryo-EM structure of *E. coli* SspA bound to *E. coli* RNAP and a generic promoter, one not characterized as being regulated by *E. coli* SspA, which is structurally similar to our SspA-regulated *Ec*RNAP(SspA)₂-*gadA* promoter complex solved at 3.2 Å resolution.

Limitations of Study

Our work provides structures of key *Francisella tularensis* virulence transcription complexes. The resolution of the *Ft*RNAP σ^{70} -(MgIA-SspA)-ppGpp-PigR-DNA structure limits the details that can be discerned from this complex, and the location

of PigR in this structure is not clearly resolved. An additional limitation of the *Ft*RNAP σ^{70} -(MgIA-SspA)-ppGpp-PigR-DNA structure is the inability to assign which α CTD is bound to which UP element.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.molcel.2020.10.035>.

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AUTHOR CONTRIBUTIONS

M.A.S. and R.G.B. conceived the structural experiments and S.L.D. the genetic studies; B.A.T., M.A.S., K.M.R., S.M.P., T.T., J.M.W., and A.B. performed experiments or analyses; M.B., A.H., and A.B. provided cryo-EM consulting and experimental input; M.A.S., B.A.T., and R.G.B. wrote the manuscript with significant input from S.L.D. and K.M.R.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-Digoxigenin-AP, Fab fragments	Millipore Sigma	Cat# 11093274910; RRID: AB_2734716
Bacterial and Virus Strains		
<i>E. coli</i> C41(DE3)	Lucigen	Cat# 60442-1
<i>E. coli</i> BL21(DE3)	NEB	Cat# C25271
<i>E. coli</i> XL1 Blue	Agilent	Cat# 200130
<i>E. coli</i> DH5 α F' I ^q	NEB	Cat# C29921
<i>E. coli</i> BL21-Gold(DE3)pLysS	Agilent	Cat#230134
<i>E. coli</i> KDZif1 Δ Z	Vallet-Gely et al., 2005	N/A
<i>F. tularensis</i> subsp. <i>holarctica</i> LVS β -TAP	Charity et al., 2007	N/A
<i>F. tularensis</i> subsp. <i>holarctica</i> LVS MglA-TAP	Charity et al., 2007	N/A
<i>F. tularensis</i> subsp. <i>holarctica</i> LVS Δ mglA β -TAP	Rohlfing and Dove, 2014	N/A
LVS	Fortier et al., 1991	N/A
LVS Δ mglA	Charity et al., 2007	N/A
LVS Δ pigR	Charity et al., 2009	N/A
LVS P _{iglA} -lacZ	Ramsey et al., 2015	N/A
LVS Δ pigR P _{iglA} -lacZ	Ramsey et al., 2015	N/A
LVS P _{iglA} -M1-lacZ	Ramsey et al., 2015	N/A
LVS Δ pigR P _{iglA} -M1-lacZ	Ramsey et al., 2015	N/A
LVS P _{iglA} -M3-lacZ	This paper	N/A
LVS Δ pigR P _{iglA} -M3-lacZ	This paper	N/A
LVS P _{iglA} -UPM1-lacZ	This paper	N/A
LVS Δ pigR P _{iglA} -UPM1-lacZ	This paper	N/A
LVS P _{FTL_0026} -lacZ	This paper	N/A
LVS Δ pigR P _{FTL_0026} -lacZ	This paper	N/A
Chemicals, Peptides, and Recombinant Proteins		
C-terminal PigR Peptide: KRNVFSRCWINMNLYSVIAKAS	Genscript	N/A
Guanosine-3',5-bisdiphosphate (ppGpp)	TriLink Biotechnologies	Cat# N-6001
Sarkosyl	Sigma	Cat# L5125-50G
Phenylmethylsulfonyl fluoride (PMSF)	Sigma	Cat# 10837091001
Lysozyme	Worthington Biochem	Cat# LS002933
Sodium deoxycholate	Alfa Aesar	Cat# B20759
EDTA	Sigma	Cat# EDS-100G
1,4-Dithiothreitol (DTT)	Thermo Fisher	Cat# BP172-25
Polyethyleneimine (PEI)	Spectrum	Cat# 46AT97
Polysorbate 20 (Tween 20)	Thermo Fisher	Cat# BT337-500
CHAPSO	Sigma	Cat# C3649-1G
Difco Mueller Hinton Broth	BD Diagnostic Systems	Cat# 275730
IsoVitalX Enrichment	BD Diagnostic Systems	Cat# 211876
Cystine Heart Agar	BD Diagnostic Systems	Cat# 247100
Bovine Hemoglobin	VWR	Cat# 90000-410
BugBuster 10X Protein Extraction Reagent	Millipore Sigma	Cat# 70921
AcTEV™ Protease	Life Technologies	Cat# 12575015
Rnase-free DNase I	Lucigen Corporation	Cat# D9905K

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
cOmplete Mini EDTA-free Protease Inhibitor Cocktail Tablets	Millipore Sigma	Cat# 11836170001
Millipore Millex Sterile Syringe Filters: PES Membrane NP-40 Alternative	Millipore Sigma	Cat# SLGP033RS
NP-40 Alternative	Millipore Sigma	Cat# 492018
FastStart Essential DNA Green Master	Roche	Cat# 06402712001
Superscript III Reverse Transcriptase	Life Technologies	Cat# 18080093
RQ1 DNase, Promega	Fisher Scientific	Cat# PR-M6101
CDP-Star	Millipore Sigma	Cat# 11759051001
ULTRAhyb Ultrasensitive Hybridization Buffer	Life Technologies	Cat# AM8670
Kanamycin A	Research Products International	Cat# K22000
Carbenicillin disodium salt	VWR	Cat# IC19509205
Tetracycline hydrochloride	Fisher Scientific	Cat# BP912-100
O-nitrophenyl- β -D-galactoside	Millipore Sigma	Cat# N1127
Isopropyl β -D-1-thiogalactopyranoside (IPTG)	Millipore Sigma	Cat# 420322
Ferric pyrophosphate	Millipore Sigma	Cat# P6526
Critical Commercial Assays		
Direct-zol Miniprep kit	Zymo Research	Cat# R2070
Ribo-Zero Magnetic Kit (Bacteria)	Illumina	Cat# MRZB12424
KAPA Stranded RNA-Seq Kit	Kapa Biosystems	Cat# KK8400
KAPA Single-Indexed Adaptor Kit	Kapa Biosystems	Cat# KK8710
PCR DIG Probe Synthesis Kit	Millipore Sigma	Cat# 11636090910
DIG Wash and Block Buffer Set	Millipore Sigma	Cat# 11585762001
Deposited Data		
Coordinates/StructureFactors for the <i>Ft</i> (MglA-SspA)-ppGpp-PigR crystal structure	This paper	PDB: 6WEG
Cryo-EM map of <i>Ft</i> RNA σ^{70} -(MglA-SspA)- <i>iglA</i>	This paper	EMD-21851
Cryo-EM map of <i>Ft</i> RNA σ^{70} -(MglA-SspA)-ppGpp-PigR- <i>iglA</i>	This paper	EMD-21852
Cryo-EM map of <i>Ec</i> RNA σ^{70} -(SspA) $_2$ - <i>gadA</i>	This paper	EMD-21853
Cryo-EM map of <i>Ft</i> RNA σ^{70} - <i>iglA</i>	This paper	EMD-21850
<i>Ft</i> RNA σ^{70} -(MglA-SspA)- <i>iglA</i> coordinates	This paper	PDB: 6WMR
<i>Ft</i> RNA σ^{70} -(MglA-SspA)-ppGpp-PigR- <i>iglA</i> coordinates	This paper	PDB: 6WMT
<i>Ec</i> RNA σ^{70} -(SspA) $_2$ - <i>gadA</i> coordinates	This paper	PDB: 6WMU
<i>Ft</i> RNA σ^{70} - <i>iglA</i> coordinates	This paper	PDB: 6WMP
RNA sequencing	This paper	GEO: GSE150932
Oligonucleotides		
<i>iglA</i> -1 (top): ATTTAGCTGTATAAACATTGTGTTAT TGCGGTTATTAAGGTAAGTCTGCTTATA	IDT	N/A
<i>iglA</i> -1 (bottom): TATAAGCAAGTTACCTTAATAA CGCCAATAACACAATGTTTATACAGCTAAAT	IDT	N/A
<i>iglA</i> -2 (top): GTTTAAATTTAGCTGTATAAACATTGT GTTATTGGCGTTATTAAGGTAAGTCTGCTTATAAGGT	IDT	N/A
<i>iglA</i> -2 (bottom): ACCTTATAAGCAAGTTACCTTAATAA CGCCAATAACACAATGTTTATACAGCTAAATTTAAAC	IDT	N/A
<i>gadA</i> (top): CTGTAATGCCTTGCTTCCATTGCGGATA AATCCTACTTTTTTATTGCCTTC	IDT	N/A
<i>gadA</i> (bottom): GAAGGCAATAAAAAAGTAGGATTT ATCCGCAATGGAAGCAAGGCATTACAG	IDT	N/A
Recombinant DNA		
pET-15b vector	Novagen	Cat# 69661-3
pIA900 vector	Svetlov & Artsimovitch, 2015	Addgene plasmid #104401

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
LA4 vector	Borukhov and Goldfarb, 1993	N/A
pMCSG21 vector	DNASU	N/A
pET-15b-PigR	Genscript	This paper
pET-28-a-His6MBP-MglA	Cuthbert et al., 2015	N/A
pMCSG21-SspA	Cuthbert et al., 2015	N/A
pF	Rohlfing and Dove, 2014	N/A
pF-MglA-V	Rohlfing and Dove, 2014	N/A
pF-MglA(F78R+P80E)-V	This paper	N/A
pSP109	This paper	N/A
pSP111	This paper	N/A
pSP113	This paper	N/A
pSP125	This paper	N/A
pBR-MglA- ω	Rohlfing and Dove, 2014	N/A
pACTR-SspA-Zif	Rohlfing and Dove, 2014	N/A
pACTR-Ap-Zif	Rohlfing and Dove, 2014	N/A
pBR-MglA(F78R+P80E)- ω	This paper	N/A
Software and Algorithms		
Coot v0.9	Emsley et al., 2010	https://www2.mrc-lmb.cam.ac.uk/personal/pemsley/coot
MotionCor2 v1.2.2	Zheng et al., 2017	https://emcore.ucsf.edu/ucsf-software
Phenix v1.17	Afonine et al., 2012	https://www.phenix-online.org/download/
RELION-3.0	Zivanov et al., 2018	https://www3.mrc-lmb.cam.ac.uk/relion/index.php/Main_Page
UCSF Chimera v1.13.1	Pettersen et al., 2004	https://www.cgl.ucsf.edu/chimera
UCSF ChimeraX v0.91	Goddard et al., 2018	https://www.rbvi.ucsf.edu/chimerax/
HT-Seq v0.6.1	Anders et al., 2015	https://github.com/simon-anders/htseq
Bowtie2 v2.2.6	Langmead and Salzberg, 2012	http://bowtie-bio.sourceforge.net/bowtie2/index.shtml
DESeq2 v1.14.1	Love et al., 2014	https://bioconductor.org/packages/release/bioc/html/DESeq2.html
GCTF v1.06	Zhang, 2016	https://www2.mrc-lmb.cam.ac.uk/research/locally-developed-software/zhang-software/#gctf
cisTEM v1.0.0	Grant et al., 2018	https://cistem.org/
crYOLO v1.5.6	Wagner et al., 2019	window:cryolo">https://sphire.mpg.de/wiki/doku.php?id=pipeline>window:cryolo
Trimmomatic v 0.36	Bolger et al., 2014	http://www.usadellab.org/cms/?page=trimmomatic
Prism	GraphPad	https://www.graphpad.com/scientific-software/prism/
XDS	Kabsch, 2010	http://xds.mpimf-heidelberg.mpg.de
3DFSC	Tan et al., 2017	https://3dfsc.salk.edu/
Other		
Ni-NTA agarose	ThermoFisher	Cat# R90115
HiLoad 26/600 Superdex 200 pg	GE Healthcare	Cat# GE28-9893-36
HiLoad 26/600 Superdex 75 pg	GE Healthcare	Cat# GE28-9893-34
DE53 diethylaminoethyl cellulose	Whatman	Cat# 4058200
Heparin Sepharose 6 Fast Flow	GE Healthcare	Cat# GE17-0998-01
Quantifoil R1.2/1.3 Cu 300 holey carbon grids	EM Sciences	Cat# Q350-CR1.3

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
UltrAufoil R1.2/1.3 Au 300 holey gold grids	EM Sciences	Cat# Q350AR1.3A
Calmodulin Affinity Resin, Agilent Technologies	Fisher Scientific	Cat# 50-125-262
IgG Sepharose® 6 Fast Flow	Millipore Sigma	Cat# GE17-0969-01
KAPA Pure Beads	Kapa Biosystems	Cat# KK8000
Poly-Prep Chromatography Columns	Bio-Rad	Cat# 7311550
LightCycler 96 Instrument	Roche	Cat# 05815916001
Nylon Membrane, positively charged	Millipore Sigma	Cat# 11209299001

RESOURCE AVAILABILITY**Lead Contact**

Further information for resources and requests should be directed to and will be fulfilled by the Lead Contact, Maria A. Schumacher (maria.schumacher@duke.edu).

Materials Availability

All reagents generated in this study are available upon request from the Lead Contact without restriction.

Data and Code Availability

The (MglA-SspA)-ppGpp-PigR peptide crystal structure coordinates and structure factors have been deposited in the Protein Data Bank under the accession number, PDB: 6WEG. Cryo-EM density maps have been deposited in the Protein Data Bank with the accession numbers PDB: 6WMM, PDB: 6WMP, PDB: 6WMU and PDB: 6WMT and EMD with accession codes EMD-21851, EMD-21850, EMD-21853 and EMD-21852. RNA-Seq data have been deposited in the National Center for Biotechnology Information Gene Expression Omnibus (GEO) under accession number GEO: GSE150932.

EXPERIMENTAL MODEL AND SUBJECT DETAILS**Growth conditions**

F. tularensis subsp. *holarctica* LVS and its derivatives were grown aerobically at 37°C in either Mueller Hinton broth (BD Diagnostic Systems), supplemented with 0.1% glucose, 0.025% ferric pyrophosphate, and 2% Isovitalex (BD Diagnostic Systems) or on cysteine heart agar (BD Diagnostic Systems) supplemented with 1% hemoglobin solution (VWR). When appropriate, kanamycin (Research Products International) was used for selection at either 5 µg/mL or 10 µg/mL. *Escherichia coli* strains XL1 Blue and DH5αF'1^q were used for routine cloning. *E. coli* strains were grown aerobically at 37°C in LB medium.

METHOD DETAILS**Purification of *Ft*RNA σ^{70} complexes and PigR**

*Ft*RNA σ^{70} and *Ft*RNA σ^{70} -(MglA-SspA): *Ft*RNA σ^{70} -(MglA-SspA) used for cryo-EM studies was purified directly from cells of the *Francisella tularensis* live vaccine strain (LVS) that synthesized the β' -subunit of RNAP with a tandem affinity purification (TAP)-tag at its C terminus (LVS β' -TAP) (Charity et al., 2007). *Ft*RNA σ^{70} -(MglA-SspA) was also purified from cells of LVS that synthesized MglA with a TAP-tag at its C terminus (LVS MglA-TAP) (Charity et al., 2007). For Cryo-EM studies *Ft*RNA σ^{70} but lacking the MglA-SspA complex (*Ft*RNA σ^{70}) was purified from LVS Δ mglA mutant cells that synthesize β' -TAP (LVS Δ mglA β' -TAP) (Rohlfing and Dove, 2014). In all cases, cells were grown with aeration at 37°C in 400 mL supplemented Mueller-Hinton Broth to mid-exponential phase (OD₆₀₀ ~0.35) and harvested by centrifugation at 4°C. Cell pellets were resuspended in 5 mL lysis buffer [20 mM K-HEPES, pH 7.9, 50 mM KCl, 0.5 mM dithiothreitol (DTT), 1x BugBuster (Millipore Sigma), 10% glycerol, with one protease inhibitor tablet (Millipore Sigma) per 10 mL] and 10 U DNase I (Lucigen) was added to the cell lysis buffer. Cells were lysed by incubation at 37°C for 30 min and insoluble material was removed by centrifugation. Lysates were passed through a 0.22 micron filter and the buffer concentration was adjusted to 10 mM Tris-HCl pH 8, 150 mM NaCl, 0.1% NP-40 (Millipore Sigma). The lysates were incubated with rocking for 2 hr at 4°C in chromatography columns (Bio-Rad) with 200 µL IgG-Sepharose 6 Fast Flow beads (Millipore Sigma) that had been washed twice and resuspended in IPP150 buffer (10 mM Tris-HCl pH 8, 150 mM NaCl, 0.1% NP-40). Lysates were allowed to flow through the column and beads were washed three times with 10 mL IPP150 buffer and once with 10 mL Tobacco Etch Virus (TEV) cleavage buffer (10 mM Tris-HCl pH8, 150 mM NaCl, 0.1% NP-40, 0.5 mM EDTA, 1 mM DTT). Beads were incubated with rocking overnight at 4°C with 1 mL TEV cleavage buffer and 100 units AcTEV Protease (Life Technologies). Samples were eluted from the beads and combined with the eluate from a wash with 200 µL TEV cleavage buffer and the buffer concentration was adjusted to include 3 mM CaCl₂.

Samples were incubated with rocking for one hr at 4°C in clean chromatography columns with 200 μ L calmodulin binding beads (Fisher Scientific) that had been washed twice and resuspended in calmodulin binding buffer (10 mM K-HEPES pH 7.9, 150 mM NaCl, 10 mM β -mercaptoethanol, 1 mM Mg-acetate, 1 mM imidazole, 2 mM CaCl_2). Unbound samples were allowed to flow through and beads were washed three times with 10 mL calmodulin binding buffer and once with 200 μ L calmodulin elution buffer (10 mM K-HEPES pH 7.9, 150 mM NaCl, 10 mM β -mercaptoethanol, 1 mM Mg-acetate, 1 mM imidazole, 2 mM EGTA). Samples were eluted from beads using 1 mL calmodulin elution buffer. $\text{Fts}^{\sigma 70}$ copurified with the core enzyme in stoichiometric amounts in both cases as did MglA-SspA in the non ΔmglA cells.

PigR: An artificial gene, codon optimized for *E. coli* expression, encoding *F. tularensis* PigR was purchased from Genscript. The gene was cloned into the pET15b plasmid at the BamHI and NdeI restriction sites, providing a hexahistidine tag at the N terminus of the expressed protein. *E. coli* C41(DE3) cells were transformed with the plasmid for protein expression. Cells were induced with 0.5 mM isopropyl β -D-1-thiogalactopyranoside (IPTG) when they reached an OD_{600} of ~ 0.6 . After induction at 37°C for 3 h, cells were pelleted and resuspended in Buffer A [20 mM Tris-HCl pH 7.5, 300 mM NaCl, 5% glycerol, 7.5% sodium lauroyl sarcosinate (sarkosyl), 5 mM β -mercaptoethanol (BME)]. Cells were lysed by sonication and the soluble fraction was subjected to affinity chromatography using nickel-nitrilotriacetic acid (Ni-NTA) resin. Wash and elution steps were performed with Buffer B (20 mM Tris-HCl pH 7.5, 300 mM NaCl, 5% glycerol, 0.05% sarkosyl, 5 BME) supplemented with increasing concentrations of imidazole. Several milligrams of pure (> 95%) PigR per liter of bacterial culture were obtained using this method.

Purification of $\text{EcRNAP}^{\sigma 70}$ and EcSspA

$\text{EcRNAP}^{\sigma 70}$ holoenzyme: The EcRNAP core enzyme (α dimer, β , β' , ω) and σ^{70} expression strains were a gift from Drs. Wilma Ross and Richard Gourse (UW-Madison). The core enzyme was expressed from a pIA900 vector in BL21(DE3) cells and purified by polyethylenimine (PEI) precipitation, ammonium sulfate precipitation, Ni-NTA affinity chromatography, and heparin-Sepharose chromatography (Ross et al., 2003). For EcRNAP core expression, the cells were grown at 37°C until the OD_{600} reached 0.6 and were then induced with 1 mM IPTG for 3 hr at 30°C. Cells were pelleted and resuspended in Buffer C (50 mM Tris-HCl pH 8.0, 5% glycerol, 2 mM EDTA, 233 mM NaCl, 1 mM β -mercaptoethanol, 0.1 mM DTT, 0.26 mM PMSF). Lysozyme and deoxycholate were added to the mixture to a final concentration of 0.25 mg/mL and 0.07% respectively. Next, the cells were lysed by sonication and diluted with an equal volume of 0.2 mM NaCl + TGED (10 mM Tris-HCl pH 8.0, 5% glycerol, 0.1 mM EDTA, 0.1 mM DTT). After centrifugation to remove cell debris, the supernatant was cooled to 4°C. While stirring, PEI was added dropwise to a final concentration 0.6%. The resulting white precipitate was collected by centrifugation at 7500 rpm at 4°C and the supernatant was discarded. The pellet was extracted twice with 0.5 M NaCl + TGED by dissolving the pellet and repeating the centrifugation step. RNAP was eluted from the pellet with 1.0 M NaCl + TGED and the centrifugation step was repeated again. Next, 0.35 mg of ammonium sulfate was added per mL of supernatant and the mixture was stirred on ice for 30 min. The precipitate was collected by centrifugation and the pellet was stored in 0.2 M NaCl + TGED overnight at 4°C. The next day, the pellet was resuspended in Buffer D (20 mM Tris-HCl pH 8.0, 500 mM NaCl, 5% glycerol, 0.1% Tween, 10 mM imidazole). The resuspension was loaded onto a Ni-NTA column and washed with an equal volume of Buffer D. EcRNAP core was eluted from the column with Buffer F (20 mM Tris-HCl pH 8.0, 500 mM NaCl, 5% glycerol) supplemented with increasing concentrations of imidazole. To remove nucleic acids that may have copurified with the EcRNAP core, the EcRNAP -containing elutions were pooled and subjected to heparin Sepharose chromatography. This was performed by bringing the EcRNAP solution to 0.2 mM NaCl and 2 mM DTT, loading onto the heparin Sepharose column pre-equilibrated with 0.2 M NaCl + TGED, washing with two column volumes of 0.2 M NaCl + TGED, and eluting with 15 mL 0.6 M NaCl + TGED. The purified EcRNAP -containing fractions were dialyzed against 0.1 mM NaCl + TGED, flash frozen, and stored at -80°C until use. The σ^{70} subunit was expressed without an affinity tag from the LA4 expression vector in BL21 Gold (DE3)pLysS cells and purified from inclusion bodies (Anthony et al., 2003). Cells expressing σ^{70} were grown at 37°C until reaching an OD_{600} of ~ 0.6 . The cells were induced with 0.5 mM IPTG for 3 h at 37°C. The names and components of the buffers used for the following purification steps are the same as in Borukhov and Goldfarb (Borukhov and Goldfarb, 1993). σ^{70} containing cell pellets were suspended in 100 mL of Lysis Buffer (40 mM Tris pH 7.9, 0.3 M KCl, 10 mM EDTA, 0.1 mM PMSF), 0.2 mg/mL lysozyme, and 0.2% sodium deoxycholate. The cells were then lysed by sonication and centrifuged to pellet the insoluble inclusion bodies. The pellet was resuspended in 50 mL Lysis Buffer with 0.5% Triton X-100, sonicated, and pelleted two additional times. The inclusion bodies were solubilized by dissolving the pellet in 20 mL Denaturing Buffer (50 mM Tris pH 7.9, 6 M guanidine HCl, 10% glycerol, 10 mM MgCl_2 , 10 μ M ZnCl_2 , 1 mM EDTA, 10 mM DTT) and centrifuged to remove insoluble particles. The supernatant was dialyzed against 2 L of Reconstitution Buffer (50 mM Tris pH 7.9, 20% glycerol, 10 mM MgCl_2 , 10 μ M ZnCl_2 , 1 mM EDTA, 1 mM DTT) overnight. The next day, precipitate was removed by centrifugation and the supernatant was diluted 5-fold with Buffer E (40 mM Tris pH 8.3, 5% glycerol, 1 mM EDTA, 0.1 mM DTT) (Ross et al., 2003). Lastly, anion-exchange chromatography was performed using DE53 diethylaminoethyl cellulose (Whatman) pre-equilibrated in Buffer E. The protein was loaded onto the column, which was then washed with 15 mL Buffer E supplemented with 50 mM NaCl. The protein was eluted with 15 mL of Buffer E supplemented with 300 mM NaCl. Pure σ^{70} was obtained from this step. To form the $\text{EcRNAP}^{\sigma 70}$ holoenzyme, EcRNAP was mixed 1:3 with σ^{70} and incubated at room temperature (rt) in Buffer G (20 mM Tris-HCl pH 8.0, 5 mM MgCl_2 , 5% glycerol, 100 mM NaCl, 10 mM BME) for one hr followed by incubation at 4°C overnight. The holoenzyme was further purified by size exclusion chromatography using a HiLoad 26/600 Superdex 200 pg column (GE Healthcare) with Buffer G as the elution buffer. Fractions corresponding to purified holoenzyme were combined, flash frozen in liquid N_2 , and stored at -80°C until use.

EcSspA: An artificial gene encoding full-length *E. coli* SspA inserted into the pET15b plasmid at the BamHI and NdeI restriction sites was purchased from Genscript. C41(DE3) cells were transformed with the construct. For protein expression, the cells were grown and induced with 0.5 mM IPTG for 3 h at 37°C. For purification, the cells were pelleted and resuspended in 100 mL of Buffer H (20 mM Tris-HCl pH 7.5, 300 mM NaCl, 5% glycerol, 1 mM BME). Cells were lysed by sonication and the cleared lysate was loaded onto a Ni-NTA column pre-equilibrated with Buffer H. SspA was washed and eluted with Buffer H supplemented with increasing concentrations of imidazole, yielding purified SspA. SspA was buffer exchanged with Buffer H to remove imidazole and was flash frozen and stored at –80°C until use.

Cryo-EM specimen preparation

***FtRNAP* σ^{70} -(MgIA-SspA)-DNA and *FtRNAP* σ^{70} -DNA:** 0.4 mg/mL *FtRNAP* σ^{70} $\pm$ (MgIA-SspA) was mixed 1:2 with *iglA* promoter DNA and incubated for one hr at 22°C. The *iglA* promoter DNA site (top strand: ATTTAGCTGTATAAACATTGTGTTATTGGCGTTATTAAGGTAACCTTGCTTATA. Bottom strand: TATAAGCAAGTTACCTTAATAACGCCAATAACACAATGTTTATACAGCTAAAT) (IDT) were annealed by heating at 95°C for two min followed by slowly cooling to rt. For grid preparation, 3 μ L of sample was applied to an UltrAufoil R1.2/1.3 Au 300 (Quantifoil) holey gold grid at 95% humidity and 22°C. The grid had been glow discharged for 30 s prior to sample application. After a 15.0 s incubation period, the grid was blotted for 2.0 s and plunge frozen into liquid ethane using a Leica EM GP2 (Leica Microsystems). All grids were stored in liquid nitrogen until data acquisition.

***FtRNAP* σ^{70} -(MgIA-SspA)-ppGpp-PigR-DNA:** To prepare this complex, 0.4 mg/mL *FtRNAP* σ^{70} -(MgIA-SspA) was mixed 1:2 with *iglA* DNA containing 6 additional upstream nucleotides and 4 additional downstream nucleotides (*iglA* top strand: GTTTAAATTAGCTGTATAAACATTGTGTTATTGGCGTTATTAAGGTAACCTTGCTTATAAGGT. Bottom strand: ACCTTATAAGCAAGTTACCTTAAACGCCAATAACACAATGTTTATACAGCTAAATTTAAAC). Then, 1 mM ppGpp and 10 μ M PigR were added and the mixture was incubated at 22°C for 1 h. For grid preparation, 3 μ L of complex was applied to a glow discharged Quantifoil R1.2/1.3 Cu 300 holey carbon grid, incubated at 95% humidity and 22°C for 15.0 s, blotted for 2.5 s, and plunge frozen into liquid ethane using a Leica EM GP2.

***EcRNAP* σ^{70} -(SspA)₂-DNA and *EcRNAP* σ^{70} -DNA:** Purified *EcRNAP* σ^{70} holoenzyme was thawed, concentrated to 6 mg/mL, and mixed 1:3 with SspA and 1:2 with *gadA* promoter DNA. The *gadA* promoter DNA site (top strand: CTGTAATGCCTTGCTTCATTGCGGATAAATCCTACTTTTTATTGCCTTC. Bottom strand: GAAGGCAATAAAAAAGTAGGATTTATCCGCAATGGAAGCAAGGCATTACAG) was annealed by heating at 95°C for two min followed by slowly cooling to rt. The complex was incubated at rt for 1 h. CHAPSO was added immediately prior to freezing at a concentration of 8 mM to alleviate orientation bias. Grids were prepared on Quantifoil R1.2/1.3 Cu 300 holey carbon grids as described for *FtRNAP* σ^{70} -(MgIA-SspA)-ppGpp-PigR-DNA.

Cryo-EM data acquisition and processing

***FtRNAP* σ^{70} -(MgIA-SspA)-DNA:** A Titan Krios G3i (ThermoFisher Scientific) with Falcon 3EC direct electron detector at Duke University was used to collect 1,672 30-frame movies on grids of the *FtRNAP* σ^{70} -(MgIA-SspA)-DNA complex in counting mode. EPU software was used for automated data acquisition, the magnification was 75,000 $\times$ to give a pixel size of 1.06 Å, and the dose rate and total exposure for each movie was 0.8 e[–]/pix/s and 42 e[–] Å^{–2}, respectively. The defocus range was set from –1.0 to –2.25 μ m. After manual curation of the raw data, 1,291 movies were imported into RELION-3.0 for processing (Zivanov et al., 2018). The wrapper to MotionCor2 was used for motion correction and dose weighting (Zheng et al., 2017). For frame alignment, 5 $\times$ 5 patches were used, and the applied B-factor was 150 Å². GCTF was used for CTF estimation on unweighted summed images (Zhang, 2016). Three 2D templates generated from $\sim$ 1,000 manually picked particles were used to autopick a total of 618,526 particles. The particles were extracted with a box size of 352 pixels and rescaled to 176 pixels (2.12 Å/pix). After sorting the particles by Z-score with the 2D templates as references, the 578,236 particles with the highest Z-scores were input to reference-free 2D classification. From this, 21 classes containing 478,764 particles showed clear structural features. These classes were combined and input to a 2nd round of reference-free 2D classification. Next, 29 classes containing 242,263 particles were used for 3D classification with an *E. coli* RNAP holoenzyme (PDB: 6CA0) (Narayanan et al., 2018) low-pass filtered to 60 Å as the initial model. One 3D class composed of 50,316 particles showed clear density that was unaccounted for after fitting an *E. coli* holoenzyme structure (Narayanan et al., 2018) that was well fit with the MgIA-SspA heterodimer structure (PDB: 5U56) (Cuthbert et al., 2017). The particles in this class were then re-extracted without scaling and subjected to 3D refinement. After CTF refinement and Bayesian polishing, the global resolution improved to 3.46 Å. To improve the local resolution of MgIA-SspA, a masked 3D classification was performed using a mask only around MgIA-SspA. Two classes (21,457 particles) showing well-resolved MgIA-SspA density were combined and refined. While the global resolution for the reconstruction was unchanged, the local resolution for MgIA-SspA was improved. Lastly, this final polished particle set was exported to cisTEM (Grant et al., 2018) for a final round of 3D-refinement, which showed improved density for MgIA-SspA. All local resolution calculations were performed in RELION-3.0 (Zivanov et al., 2018) and viewed in UCSF Chimera (Pettersen et al., 2004).

***FtRNAP* σ^{70} -DNA:** For this structure, data were collected at Duke University with a Titan Krios G3i operating at 300 keV and equipped with a K3 direct electron detector (Gatan, Inc.) and Latitude S automated data acquisition software. A total of 3,490 movies were collected in counting mode with a magnification of 22,500 $\times$ to give a pixel size of 1.07 Å and a defocus range of –1.0 to –2.0 μ m. The dose rate was 15 e[–]/pix/s and the total exposure for each movie was 60 e[–] Å^{–2}. Manual curation resulted in the removal of 974 movies from the data. The remaining 2,516 movies were imported into RELION-3.0 for processing (Zivanov et al., 2018). MotionCor2 (Zheng

et al., 2017) was used for motion correction and dose-weighting with a 1×1 patch and an applied B factor of $150 \text{ \AA}^{-2}$. CTF estimation was performed with GCTF (Zhang, 2016) on the unweighted summed images. Before autopicking, ~ 1000 particles were manually picked and subjected to reference-free 2D classification to generate templates. Two templates were used to autopick 1,503,586 particles from the summed images. The particles were extracted with a box size of 352 pixels and scaled to 88 pixels ($4.28 \text{ \AA}/\text{pix}$) for faster processing. Particles were sorted based on their similarity to the autopicking references and a set of the best 1,455,290 particles were input to reference-free 2D classification. From this, 30 classes with clear structural features were combined to form a particle set of 745,802 particles to use for 3D classification. The 3D classification was performed with a $60 \text{ \AA}$ map from the *FtRNAP* σ^{70} -(MglA-SspA)-DNA data as the initial model. Class 7 was composed of 137,682 particles and showed the most well-resolved structural features. These particles were re-extracted without scaling ($1.07 \text{ \AA}/\text{pix}$) and input to 3D refinement, resulting in a $3.25 \text{ \AA}$ reconstruction. After two rounds of CTF refinement and Bayesian polishing, the global resolution of the refined map improved to $3.0 \text{ \AA}$. Three-dimensional FSC calculations using 3DFSC showed that the best and worst directional resolutions are ~ 2.9 and $\sim 3.3 \text{ \AA}$ (FSC 0.143), respectively, which is reflective of some particle orientation bias in the data.

***FtRNAP* σ^{70} -(MglA-SspA)-ppGpp-PigR-DNA:** For this structure, 7,362 movies (60 frames per movie) were collected at Duke University with a Titan Krios G3i operating at 300 keV and equipped with a K3 direct electron detector. The data was acquired using Latitude S software. Movies were collected in counting mode at a magnification 22,500x with a pixel size of $1.07 \text{ \AA}$. The defocus range was set to -1.25 to $-2.75 \text{ \mu m}$ and the dose rate for each movie was $15 \text{ e}^-/\text{pix}/\text{s}$ to give a total electron exposure of $60 \text{ e}^- \text{ \AA}^{-2}$. After manual curation, 7,081 movies were imported into RELION-3.0 for processing (Zivanov et al., 2018). MotionCor2 was used for beam-induced motion correction (Zheng et al., 2017) with dose-weighting for each dose-fractionated movie stack. The applied B-factor for frame alignment was 150 and 1×1 patches were used. The contrast transfer function was estimated for each unweighted summed image using a wrapper to GCTF (Zhang, 2016) in RELION-3.0 (Zivanov et al., 2018). $\sim 1,000$ particles were manually selected and classified using reference-free 2D classification. Subsequently, auto-picking was performed with 4 templates that each represented a different particle orientation to give a total particle set of 3,485,423 particles. Particles were extracted with a box size of 352 pixels, rescaled to 88 pixels ($4.28 \text{ \AA}/\text{pix}$), and were sorted by similarity to the 2D templates using the Particle Sort job. The 258,463 particles with the worst Z-scores (least similarity) were removed from the data after sorting. Next, the remaining particles were classified by reference-free 2D classification. After 2D classification, high-resolution classes representing 1,901,718 particles were combined and used for 3D classification. The initial model for 3D classification was created by generating a $60 \text{ \AA}$ density map from a *EcRNAP* σ^{70} holoenzyme structure (PDB: 6CA0) (Narayanan et al., 2018; Kang et al., 2017). Two holoenzyme classes were combined and re-extracted with a pixel size of $2.14 \text{ \AA}$. Refinement of this particle set yielded a $5.0 \text{ \AA}$ reconstruction with extra density upstream of the -35 region of DNA. A focused classification of this upstream region revealed density for 2 RNAP α C-terminal domains. A focused classification of this upstream region revealed density for 2 RNAP α C-terminal domains and spurious density for PigR. To perform the focused classification, a spherical mask was created in RELION-3.0 to encompass the DNA upstream of -35 . For a view of the mask, see the data processing flowchart (Figure S6C). The focused classification was performed with partial signal subtraction. One class composed of 22.9% of the particles revealed density for 2 α CTDs. After re-extraction, CTF refinement, and Bayesian polishing, a final map with a global resolution of $4.5 \text{ \AA}$ was obtained. Additional focused classifications were performed to further resolve the spurious PigR density but were unsuccessful likely due to heterogeneity.

***EcRNAP* σ^{70} -(SspA) $_2$ -DNA:** Remote data collection was performed at the Pacific Northwest Cryo-EM Center (PNCC) using a Titan Krios G3i operating at 300 keV and equipped with a K3 direct electron detector. A total of 7,363 50-frame movies were collected in counting mode with a magnification of 81,000x, a pixel size of $0.53 \text{ \AA}$, a dose per frame of $1 \text{ e}^- \text{ \AA}^{-2}$, and a defocus range of -1.0 to $3.0 \text{ \mu m}$. Following manual curation, 5,708 movies were imported to RELION-3.0 for processing. During beam-induced motion correction with MotionCor2, the movies were binned by a factor of 2. CTF estimation was performed using GCTF. For particle picking, $\sim 1,000$ particles were picked and exported to crYOLO (Wagner et al., 2019) to train a general model which was used to pick 672,043 particles from the full dataset. The particles were binned 4x and subjected to one round of reference-free 2D classification. Next, 40 2D classes comprising 393,711 particles were selected and input to 3D classification. All 4 classes represented holoenzymes with clear density for SspA. The class with the best occupancy for SspA contained 111,519 particles and was 3D auto-refined to $8.8 \text{ \AA}$. Lastly, a focused 3D-classification of SspA with partial signal subtraction of the RNAP core and σ^{70} subunit was performed. Class 2 comprised 49,560 particles and had the best resolution for SspA. A final 3D refinement was performed on this class to give a map with a global resolution of $3.2 \text{ \AA}$.

Model building and refinement

To build the *FtRNAP* core and σ^{70} subunits, a model of *EcRNAP* σ^{70} holoenzyme (PDB: 6CA0)⁴ was fit into the density map using Chimera (Pettersen et al., 2004). The *FtRNAP* β , β' , ω , and σ^{70} subunits share 59%, 62%, 42% and 45% sequence identity with the corresponding subunits of *EcRNAP* and the two *Ft* α subunits share 39% and 41% sequence identity with the *EcRNAP* α subunit. After fitting, the RNAP core subunits and σ^{70} subunits were replaced by the *Ft* sequences and re-modeled to fit the map using Coot (Emsley et al., 2010). In the *FtRNAP* σ^{70} -(MglA-SspA)-DNA structure, Chimera was also used to fit a model of MglA-SspA (PDB: 5U56) (Cuthbert et al., 2017) into the density map. The C-terminal helix of SspA, not visible in our previous crystal structure (Cuthbert et al., 2017) was built *de novo* to fit the electron density. Density was not present for DNA bps -52 to -36 and was not modeled. In *FtRNAP* σ^{70} -DNA, nonspecifically bound DNA and RNA in the active site were fit into the map using Chimera using the corresponding DNA and RNA from PDB: 6ALH (Kang et al., 2017). Density was absent for the *igIA* promoter DNA in this structure. To build the

*Ec*RNA σ^{70} -(SspA) $_2$ -DNA structure, an *Ec*RNA σ^{70} holoenzyme (PDB: 6CA0) (Narayanan et al., 2018) was manually fit into the density map using Chimera (Pettersen et al., 2004). Next, a crystal structure of *Y. pestis* SspA (PDB: 1YY7) (Hansen et al., 2005a) was fit into the density corresponding to SspA. Using Coot, the *Y. pestis* SspA sequence was replaced by the *E. coli* SspA sequence and all RNAP holoenzyme and SspA residues were inspected and remodeled as necessary. Some parts of the RNAP model, such as the N terminus of β' , and the C terminus of ω , were built *de novo*. Density was not present for nucleotides –45 to –38 in the upstream promoter DNA region and was not modeled. In the *Ft*RNA σ^{70} (MglA-SspA)-ppGpp-PigR-DNA structure, density was evident for a ppGpp bound to MglA-SspA as previously observed (Cuthbert et al., 2017) and in our (MglA-SspA)-ppGpp-PigR crystal structure (below). Density was evident for most of the promoter in this structure except –58 to –53. In addition, density for a ppGpp molecule bound in the *Ft*RNA σ^{70} between the β' and ω subunits, termed the site 1 ppGpp binding site in the *E. coli* RNAP, was observed. The role of this ppGpp binding site is unclear. Indeed, binding to site 2, which is between ppGpp, DksA and the RNAP channel in *E. coli* RNAP, results in the majority of ppGpp's effects on transcription (Ross et al., 2013). There is no DksA in *Ft* and the residues involved in site 2 binding are also not present in *Ft*RNA σ^{70} . While the density for the *Ft* α CTDs was evident in this structure, the side chains could not be fitted and hence the α CTDs were constructed as polyalanine models.

For all deposited cryo-EM coordinates, flexible regions and unstructured loops were removed from the model if the electron density was not well resolved. Real-space refinement was performed in Phenix (Afonine et al., 2012) using Ramachandran and secondary structure restraints. Refined coordinates were inspected and modified in Coot (Emsley et al., 2010), as needed.

***Ft*(MglA-SspA)-ppGpp-PigR determination**

A *his₆-sspA*-(*his₆-mbp-mglA*) coexpression system was generated by cotransforming C41(DE3) cells with plasmids encoding *F. tularensis* *sspA* (cloned into the pMCSG21 vector using ligation-independent cloning) and the *his₆-mbp-mglA* fusion (cloned into pET28A). C41(DE3) cells were transformed with both expression plasmids and cells grown to an OD₆₀₀ of 0.5 at 37°C. The cells were then induced by addition of 0.5 mM IPTG overnight at 15°C. Cells were lysed in 20 mM Tris, pH 7.5, 200 mM NaCl, 10% glycerol, and 7.5 mM imidazole, 1 mM β ME, 2 mg/L DNase I, and 1 mM PMSF with a microfluidizer. Cell debris was removed by centrifugation at 15,000 rpm and the resultant supernatant was loaded onto a Ni-NTA column and washed with increasing concentrations of imidazole in a buffer consisting of 20 mM Tris, pH 7.5, 200 mM NaCl, 10% glycerol, and 1 mM β ME. The protein was eluted using the buffer containing 0.25 to 2.0 M imidazole. The MBP-tag was removed overnight at rt using his-tagged TEV protease. The treated protein sample was next applied to a Ni-NTA column, which removed the His₆-TEV tag and any uncleaved fusion protein. The MglA-SspA was collected in the flow through and the protein further purified via size exclusion chromatography using a Superdex S75 column (GE Healthcare). The buffer used for size exclusion chromatography was 20 mM Tris, pH 7.5, 200 mM NaCl, 10% glycerol, and 1 mM DTT. Crystals were grown using the hanging drop vapor diffusion method by mixing the protein complex at 24 mg/mL with ppGpp (TriLink) and 22-mer PigR C-terminal peptide (KRNVSRCWINMNLVSVIKAKS) at final concentrations of 1 mM and 0.5 mM, respectively. This solution was mixed 1:1 with a crystallization reagent consisting of 0.1 M Tris-HCl pH 8.0 and 24% PEG 4000. Data collection revealed no peptide density, likely due to low peptide solubility. Thus, excess PigR peptide was added to the drops, which were then allowed to sit for 2 months prior to data collection. To collect data, the crystals were cryoprotected by dipping them for several seconds in a 1 μ L drop containing the crystallization solution supplemented with 18% ethylene glycol. X-ray intensity data were collected at the Advanced Light Source (ALS) beamline 8.3.1. The data were processed and scaled using XDS. Phaser in CCP4 was used to solve the structure by molecular replacement using an MglA-SspA heterodimer from our previously solved structure (Cuthbert et al., 2017). The crystallographic ASU contained two MglA-SspA dimers. Each dimer has a bound ppGpp but packing only permits the interaction of the PigR peptide with one dimer. Density was found for all but the last three residues of the peptide in this dimer (though a low contour level of the map was required) and PigR peptide side chain density was also weak. After multiple rounds of rebuilding and refinement in Phenix (Emsley et al., 2010; Afonine et al., 2012), the model converged to $R_{\text{work}}/R_{\text{free}}$ values of 21.8%/28.7% at 2.95 Å resolution.

Plasmids for qRT-PCR analyses

Plasmid pF-MglA-V directs the synthesis of LVS MglA with a vesicular stomatitis virus-glycoprotein (VSV-G) epitope-tag fused to its C terminus (MglA-V) under the control of the *groEL* promoter and has been described previously (Rohlfing and Dove, 2014). Plasmid pF-MglA(F78R+P80E)-V directs the synthesis of mutant MglA-V containing amino acid substitutions F78R and P80E in MglA. This plasmid was made by introducing DNA specifying the corresponding mutations into pF-MglA-V. Plasmid pF was the empty vector control (Rohlfing and Dove, 2014).

qRT-PCR analyses

RNA was isolated from cells of the LVS Δ *mglA* mutant strain that contained plasmid pF, pF-MglA-V, or pF-MglA(F78R+P80E) that were grown with aeration at 37°C in supplemented Mueller-Hinton Broth to mid-log (OD₆₀₀ = 0.35–0.4). Nucleic acids were purified using the Zymo Direct-zol RNA Miniprep Plus kit (Zymo Research) according to kit instructions and eluted in 80 μ L RNase-free water. Purified nucleic acids were treated with 10 units of RQ1 DNase (Fisher Scientific) in 1x RQ1 DNase buffer at 37°C for one hr. RNA was purified from DNase-treated samples using the Zymo Direct-zol RNA Miniprep Plus kit according to kit instructions. cDNA was generated from RNA using Superscript III reverse transcriptase (Life Technologies). Relative transcript abundance for *iglA* was determined relative to the amount of *tul4* transcript by qPCR using FastStart Essential DNA Green Master mix (Roche)

with a Roche LightCycler 96 detection system and relative expression values were calculated by using the comparative threshold cycle (C_T) method ($2^{-\Delta\Delta C_T}$) (Livak and Schmittgen, 2001). The reported fold enrichment values are the means from three biological replicates, and error bars represent the standard deviation of the mean. The data shown are from one representative experiment.

Plasmids for *Francisella* reporter strains

Mutations in the *igIA* promoter were generated in plasmid pMO1, which harbors a transcriptional fusion between the *igIA* promoter and the *lacZ* gene (Ramsey et al., 2015). Plasmid pSP109 contains DNA specifying the *igIA* promoter containing the PRE 3 mutation and was used to generate strains LVS P_{igIA} -M3-*lacZ* and LVS $\Delta pigR$ P_{igIA} -M3-*lacZ*. Plasmid pSP111 contains DNA specifying the *igIA* promoter containing the UP 1 mutation and was used to generate strains LVS P_{igIA} -UPM1-*lacZ* and LVS $\Delta pigR$ P_{igIA} -UPM1-*lacZ*. Plasmid pSP113 harbors a transcriptional fusion between the wild-type *FTL_0026* promoter and the *lacZ* gene and was made by replacing the *igIA* promoter region in plasmid pMO1 with 1100 bp of DNA upstream of the *FTL_0026* gene by isothermal assembly. Plasmid pSP113 was used to generate strains LVS P_{FTL_0026} -*lacZ* and LVS $\Delta pigR$ P_{FTL_0026} -*lacZ*. Mutations in the *FTL_0026* promoter in plasmid pSP113 were generated by site-directed mutagenesis. Plasmid pSP123 contains DNA specifying the *FTL_0026* promoter containing the PRE 3 mutation and was used to generate strains LVS P_{FTL_0026} -M3-*lacZ* and LVS $\Delta pigR$ P_{FTL_0026} -M3-*lacZ*. Plasmid pSP125 contains DNA specifying the *FTL_0026* promoter containing the UP 1 mutation and was used to generate strains LVS P_{FTL_0026} -UPM1-*lacZ* and LVS $\Delta pigR$ P_{FTL_0026} -UPM1-*lacZ*.

Francisella reporter strains

Reporter strains LVS P_{igIA} -*lacZ* and LVS $\Delta pigR$ P_{igIA} -*lacZ* contain *lacZ* integrated downstream of the wild-type *igIA* promoter at the *FTL_0111* locus (Ramsey et al., 2015). Strains LVS P_{igIA} -M1-*lacZ* and LVS $\Delta pigR$ P_{igIA} -M1-*lacZ* contain *lacZ* integrated downstream of the *igIA* promoter containing the PRE 1 mutation at the *FTL_0111* locus (Ramsey et al., 2015). Reporter strains described in this manuscript were created by electroporation of integration plasmids, described above, into LVS and selection of single homologous recombination events between the integration plasmid and the chromosome. Electroporations were performed using LVS cells washed three times with 10% sucrose and approximately 0.5–2.5 μ g plasmid DNA in a 2 mm cuvette at 2.5 kV. Cells were allowed to recover, shaking aerobically at 37°C for 4–8 hr in supplemented Mueller-Hinton Broth and cells containing an integration event were selected on cysteine heart agar with 1% hemoglobin and 10 μ g/mL kanamycin. Reporter strains containing the *igIA-lacZ* transcriptional fusion were integrated at the *FTL_0111* *igIA* locus as determined by Southern blotting (Ramsey et al., 2015). Digested chromosomal DNA was transferred to a positively charged nylon membrane (Millipore Sigma). The blot was blocked and washed using the DIG Wash and Block buffer set (Millipore Sigma) following manufacturer's instructions. The Southern blot probe was synthesized using the PCR DIG Probe Synthesis Kit (Millipore Sigma) and hybridized to the blot overnight in ULTRAhyb hybridization buffer (Life Technologies). The probe was detected after incubation with anti-digoxigenin-AP antibody (Millipore Sigma) and CDP-Star (Millipore Sigma). Reporter strains containing the *FTL_0026* promoter-*lacZ* transcriptional fusion and its derivatives were integrated at the *FTL_0026* locus as determined by PCR.

Plasmids for bacterial two-hybrid assays

Plasmids pBR-MglA- ω , pACTR-SspA-Zif, and pACTR-Ap-Zif were used for two-hybrid studies (Rohlfing and Dove, 2014). Plasmid pBR-MglA(F78R+P80E)- ω is identical to pBR-MglA- ω except that the MglA moiety of the MglA- ω fusion protein contains amino acid substitutions F78R and P80E. Plasmid pBR-MglA(F78R+P80E)- ω was made by introducing DNA specifying the F78R and P80E substitutions in MglA into plasmid pBR-MglA- ω .

β -Galactosidase assays

β -Galactosidase assays were performed with cells of the LVS and LVS $\Delta pigR$ reporter strains (Ramsey et al., 2015). Specifically, for β -galactosidase assays with LVS and LVS $\Delta pigR$ reporter strains, cells were grown with aeration at 37°C in supplemented Mueller-Hinton Broth to mid-log ($OD_{600} \sim 0.3$). Cells were lysed in Z-buffer (60 mM Na_2HPO_4 , 40 mM NaH_2PO_4 , 10 mM KCl, 1 mM $MgSO_4$, 50 mM β ME) with SDS-chloroform and assayed for β -galactosidase using ONPG (O-nitrophenyl- β -D-galactoside) as substrate at 28°C. Bacterial two-hybrid assays were performed using *E. coli* reporter strain KDZif1 Δ Z (Rohlfing and Dove, 2014). In detail, for bacterial two-hybrid assays, *E. coli* reporter strain KDZif1 Δ Z was transformed with plasmid pBR-MglA- ω together with pACTR-SspA-Zif, plasmid pBR-MglA(F78R+P80E)- ω together with pACTR-SspA-Zif, and plasmid pBR-MglA- ω together with pACTR-Ap-Zif. Transformants were first grown overnight with aeration at 37°C in LB containing carbenicillin (100 μ g/mL), tetracycline (10 μ g/mL) and IPTG (50 μ M). Cells were then back-diluted into LB containing carbenicillin (100 μ g/mL), tetracycline (10 μ g/mL) and IPTG (50 μ M) and grown with aeration at 37°C to mid-log ($OD_{600} \sim 0.3$ –0.5). Cells were lysed in Z-buffer with SDS-chloroform and assayed for β -galactosidase using ONPG as substrate at 28°C.

RNA-Seq

For RNA-Seq experiments, RNA was isolated from cells of strains LVS, LVS $\Delta mglA$, and LVS $\Delta pigR$ (Charity et al., 2007, 2009). The cells were grown with aeration at 37°C in supplemented Mueller-Hinton Broth to mid-log (OD_{600} 0.35–0.4) using the Direct-zol Miniprep kit (Zymo Research). RNA (5 μ g) was treated with the Ribo-Zero Magnetic Kit (Bacteria) (Epicenter) and sequencing libraries were constructed using the KAPA stranded RNA-Seq library preparation kit (Kapa Biosystems) following the manufacturer's

instructions and sequenced on an HiSeq2500. Reads were trimmed with Trimmomatic (v 0.36) and aligned to the *F. tularensis* LVS genome (NC_007880) using bowtie2 (v 2.2.6) (Langmead and Salzberg, 2012). HTSeq (v 0.6.1) (Anders et al., 2015) was used to count reads that map to annotated genes and DESeq2 (v 1.14.1) (Love et al., 2014) was used for differential gene expression analysis.

QUANTIFICATION AND STATISTICAL ANALYSIS

qRT-PCR

In Figure 1D, experiments were performed with biological triplicate samples at least twice and data from a representative experiment are shown. Statistical significance was assessed in Prism using a two-tailed t test assuming equal variance.

LVS reporter strain β -galactosidase assays

In Figure 5, experiments were performed with biological triplicate samples at least twice and data from a representative experiment are shown. Statistical significance was assessed in Prism using one-way ANOVA with Tukey's multiple comparisons.

Two-hybrid β -galactosidase assays

In Figure 1E, experiments were performed with biological triplicate samples at least twice and data from a representative experiment are shown. Statistical significance was assessed in Prism using one-way ANOVA with Tukey's multiple comparisons.

RNA-Seq

Transcriptomes were analyzed using biological triplicate samples and DESeq2 was used for differential gene expression analysis. Genes are considered differentially expressed if transcript abundance changed 2-fold or greater in mutant cells compared to WT with an adjusted p value of less than 0.05.