Structural analysis of the recognition of the -35 promoter element by SigW from *Bacillus subtilis*

Eunju Kwon, Shankar Raj Devkota, Deepak Pathak, Pawan Dahal, Dong Young Kim

College of Pharmacy, Yeungnam University, Gyeongsan, Gyeongbuk, South Korea

* dyokim@ynu.ac.kr

Abstract

Sigma factors are key proteins that mediate the recruitment of RNA polymerase to the promoter regions of genes, for the initiation of bacterial transcription. Multiple sigma factors in a bacterium selectively recognize their cognate promoter sequences, thereby inducing the expression of their own regulons. In this paper, we report the crystal structure of the σ4 domain of *Bacillus subtilis* SigW bound to the -35 promoter element. Purine-specific hydrogen bonds of the -35 promoter element with the recognition helix α9 of the σ4 domain occur at three nucleotides of the consensus sequence (G-35, A-34, and G′-31 in G-35A-34A-33A-32C-31C-30T-29). The hydrogen bonds of the backbone with the α7 and α8 of the σ4 domain occurs at G′-30. These results elucidate the structural basis of the selective recognition of the promoter by SigW. In addition, comparison of SigW structures complexed with the -35 promoter element or with anti-sigma RsiW reveals that DNA recognition and anti-sigma factor binding of SigW are mutually exclusive.

Introduction

Transcription in bacteria is initiated by sigma factors, which recruit the core RNA polymerase to a cognate promoter [1, 2]. Sigma factors selectively recognize promoter elements, -10 and -35 elements, and additional sequences, including extended -10 element and discriminator, which are present upstream of the transcription start site [3, 4]. During transcription initiation, the -10 element is strand-separated to form a transcription bubble [5, 6], whereas the -35 element is recognized by a helix-turn-helix (HTH) motif in the sigma factor, without strand separation [7, 8].

Sigma factors are categorized into two families, based on the sequence homology with *Escherichia coli* sigma factors: housekeeping σ^70^ required for bacterial homeostasis; and σ^54^ activated for nitrogen utilization [9]. The σ^70^ family is further sub-divided into five groups [4, 10]. Group I sigma factors are composed of σ_1.1, σ_2, σ_3, and σ_4 domains, which are responsible for the recognition of the discriminator, -10, extended -10, and -35 elements, respectively. The primary sigma factors, which regulate the transcription of housekeeping genes, belong to group I. Group II-V are classified depending on the presence or absence of the domains and
motifs in group I. Alternative sigma factors, which are activated in response to a range of stress conditions, belong to groups II–V [10].

Group IV sigma factors are known as extracytoplasmic function (ECF) sigma factors, because they are activated in response to extracytoplasmic stresses [3, 10, 11]. They are the most divergent group of sigma factors, and almost all bacteria contain multiple ECF sigma factors. As an extreme case, *Streptomyces coelicolor* contains approximately 50 ECF sigma factors in its genome [10, 12]. The ECF sigma factors are composed of only σ2 and σ4 domains, which bind -10 and -35 elements, respectively [3, 13–15]. In many cases, promoter binding of ECF sigma factors is inhibited by binding to the cytoplasmic domain of transmembrane anti-sigma factors [10, 11, 16], and is activated to initiate the transcription of target genes in response to a specific stress signal by being released from the anti-sigma factor.

*Bacillus subtilis* contains at least seven ECF sigma factors: SigM, SigV, SigW, SigX, SigY, SigZ, and YlaC [17]. Of these sigma factors, SigW induces the transcription of its target regulon to counteract cell envelope stresses caused by antibiotics [18, 19], alkaline pH [20], and high salt concentration [21, 22]. In the absence of these stresses, SigW is downregulated by anti-sigma factor RsiW, which is localized in the plasma membrane [16, 23]. It is released from the anti-sigma factor under the stress conditions [24–27]. Under these conditions, transmembrane RsiW is sequentially cleaved by regulated intramembrane proteolysis by PrsW and RasP proteases; therefore, the cytoplasmic domain of RsiW is released together with SigW from the plasma membrane [25, 27]. Subsequently, the cytoplasmic domain of RsiW is completely degraded by ClpXP protease [28], allowing SigW to bind to the core RNA polymerase, and thereby to activate SigW-dependent transcription.

Even though crystal structures have been reported showing interactions between the σ4 domain and the -35 promoter element in group I and group IV sigma factors [3, 7, 8, 29, 30], structural information from diverse sigma factors is required to understand the selective recognition of the -35 element. In the work reported here, we determined the crystal structure of the \( \sigma^W_4 \)–35 element complex, and analyzed the promoter binding mode of SigW. The structure reveals the unique regulation mode of SigW-dependent transcription, and the recognition specificity of the -35 element by the \( \sigma_4 \) domain.

**Materials and methods**

**Plasmid preparation and protein expression**

DNA encoding the σ4 domain of *B. subtilis* SigW (σ\(^W_4\) residues 126–187) was amplified from the genome of *B. subtilis* 168 strain using polymerase chain reaction and inserted into pET-Duet-1 vector (Merck Millipore, Billerica, MA, USA) expressing the N-terminal 6\(_X\)His tag and TEV protease cleavage site. The plasmid was transformed into *E. coli* strain BL21-star (DE3) (Thermo Fischer Scientific, Waltham, MA, USA) and cells were grown in Luria Broth media at 37°C. When the culture reached an OD\(_{600}\) of 0.6–0.7, the medium was cooled, and 0.4 mM isopropyl β-D-1-thiogalactopyranoside was added to the culture to induce \( \sigma^W_4 \) expression. After overnight incubation at 15°C, the cells were harvested by centrifugation at 3,000 g for 10 min.

**Purification of \( \sigma^W_4 \)–35\(^W\)**

Cells expressing 6\(_X\)His-\( \sigma^W_4 \) were resuspended in buffer A (20 mM HEPES pH 7.5, 1.0 M NaCl, and 10% (v/v) glycerol), lysed by sonication, and clarified by centrifugation at 20,000 g for 30 min, after the addition of DNase I and RNase A at a concentration of 10 μg/ml. \( \sigma^W_4 \) was purified by immobilized metal affinity chromatography (IMAC) and size exclusion chromatography (SEC). Clarified cell lysate was loaded onto a 5 mL HisTrap nickel chelating column (GE Healthcare Bio-sciences, Uppsala, Sweden), and the resin was washed with buffer A,
containing 80 mM imidazole. Proteins bound to the resin were eluted by an imidazole gradient (0.08–1.00 M imidazole). Fractions that contained 6XHis-σW were pooled and treated with TEV protease overnight at 25°C to cleave the His tag. After complete cleavage, the protein solution was dialyzed against buffer A for 3 h and passed through Ni-NTA resin to remove the 6XHis tag (Thermo Fisher Scientific, Rockford, IL, USA). σW was further purified by SEC using Superdex 75 preparatory grade column (GE Healthcare Biosciences) pre-equilibrated with buffer B (20 mM HEPES pH 7.5, 1.0 M NaCl, and 5% (v/v) glycerol).

The -35 element recognized by σW was prepared by mixing two complementary DNA fragments. Two single-stranded DNAs (5’−ATTGAAACCTTT−3’ and 5’−AAAAGGTTCCTCAA−3’ ) were synthesized (Biobasic, Seoul, South Korea), mixed at a 1:1 molar ratio in buffer B, and purified by SEC using a Superdex75 analytical column. The purified -35 element (-35W) was then mixed with σW at a 1:1:1 molar ratio. The mixture was dialyzed in buffer C (20 mM HEPES pH 7.5, 0.2 M NaCl, and 5% (v/v) glycerol) and concentrated to 15 mg/mL for crystal screening.

Crystallization, data collection, and structure determination
Crystallization of the σW/-35W complex was performed using the micro-batch method at 20°C. The drop for crystal screening was prepared by mixing 1 μL of σW/-35W (15 mg/mL) and 1 μL of crystallization solution under a layer of Al’s oil (Hampton Research, Aliso Viejo, Ca, USA). Crystals of σW/-35W grew completely in a month under the conditions of Wizard Precipitant Synergy 127 (0.1M imidazole/hydrochloric acid pH 6.5, 30% (v/v) PEG1500, 10% (v/v) isopropanol, and 0.1 M CaCl2) (Rigaku, Tokyo, Japan). Crystals of σW/-35W were picked using a cryo-loop (Hampton Research) and flash-frozen in a cold nitrogen stream. Diffraction data were collected at PLS-BL7A (Beam line 7A, Pohang Light Source, South Korea) [31] and were indexed, integrated, and scaled using MOSFLM [32].

The crystal structure of σW/-35W was determined by the molecular replacement (MR) method using PHASER [33]. The structure of the E. coli σ4/-35 element (PDB ID: 2H27) was used as a template for MR. MR solution was found from the truncated σ4 (residues 127–186)/-35 element. Cycles of refinement and model building were performed at 3.1 Å resolution using PHENIX.refine [34] and COOT [35]. Final refinement resulted in R / Rfree values of 24.8 / 29.0% without residues in the disallowed region of the Ramachandran plot. The data collection and refinement statistics are summarized in Table 1. The final coordinates and structure factors were deposited in the Protein Data Bank (PDB ID: 6JHE). Structural alignment was performed using the DALI server [36]. Protein-ligand interactions were analyzed with LigPlot+ [37] and PDBPISA [38]. The free energy change (ΔG) of the protein caused by ligand binding was analyzed using PDBPISA [38]. The conversion of ΔG to a dissociation constant (Kd) was calculated using the equation Kd = e(ΔG/RT) (R = 1.987 cal/molK; T = 293K). DNA geometry was analyzed using w3DNA [39]. Surface charge distribution was calculated using APBS [40]. The figures were drawn using PyMOL [41] and ALSCRIPT [42].

Accession number
The final coordinates and structure factors were deposited in the Protein Data Bank (PDB ID: 6JHE for σW/-35W).

Results and discussion
Overall structure
B. subtilis σW (residues 125–187) recognizes a cognate -35 promoter element (Fig 1A and 1B). Its consensus sequence is identified as T₃₆G₃₅A₃₄A₃₃A₃₂C₃₁X₃₀T₂₉T₂₈T₂₇, based on
the promoter sequences of the SigW regulon [43]. σ\textsuperscript{W\textsubscript{4}} was purified under high salt conditions (1 M NaCl) to minimize its instability, and dialyzed in low salt buffer together with the double-stranded DNA of A\textsubscript{38}T\textsubscript{37}T\textsubscript{36}G\textsubscript{35}A\textsubscript{34}A\textsubscript{33}A\textsubscript{32}C\textsubscript{31}C\textsubscript{30}T\textsubscript{29}T\textsubscript{28}T\textsubscript{27}(-35\textsuperscript{W}) to allow σ\textsubscript{W\textsubscript{4}} binding to -35\textsuperscript{W}. Crystals of σ\textsubscript{W\textsubscript{4}}/-35\textsuperscript{W} belonging to a hexagonal space group grew under conditions containing PEG1500 and isopropanol as protein precipitants. Diffraction data were collected at a resolution of 3.1 Å (Table 1), and the structure was determined by molecular replacement using the structure of a truncated E. coli σ\textsubscript{E\textsubscript{4}}/-35\textsuperscript{E} (-35 promoter element for SigE binding) as a template [8].

The crystal structure contains a σ\textsubscript{W\textsubscript{4}} monomer and a double-stranded -35\textsuperscript{W} in the asymmetric unit. Residues 134–186 of SigW and 11 nucleotide pairs of -35\textsuperscript{W} were traced into the electron density (S1 Fig) and the final structure was refined at R / R\textsuperscript{free} values of 24.8 / 29.0% (Table 1). σ\textsubscript{W\textsubscript{4}} is comprised of four α-helices (α6–α9) in the crystal structure of the SigW/RsiW complex [23]. However, residues 125–133, which correspond to α6, are disordered in the crystal structure of σ\textsubscript{W\textsubscript{4}}/-35\textsuperscript{W} (Fig 1A). The residues on α6 are likely to be flexible because they are not bound to DNA directly. α8–α9 of σ\textsubscript{W\textsubscript{4}} forms the HTH motif, and α9 is inserted into the major groove of -35\textsuperscript{W} as a DNA recognition helix (Fig 1C and 1D) [7]. Positively-charged residues are distributed on the DNA binding surface of α8–α9, whereas hydrophobic patches are distributed on the opposite side that interacts with σ\textsubscript{W\textsubscript{2}} in the crystal structure of SigW/RsiW (Fig 1E and 1F) [23].

| Table 1. Data collection and refinement statistics. |
|--------------------------------------------------|
| **Data collection**                               |
| Data set                                          |
| σ\textsubscript{W\textsubscript{4}}/-35\textsuperscript{W} |
| Space group                                      |
| P6-22                                            |
| Unit cell                                        |
| a, b, c (Å)                                      |
| 61.61, 61.61, 119.96                             |
| α, β, γ (˚)                                      |
| 90.00, 90.00, 90.00                              |
| Resolution (Å)                                  |
| 30.0–3.00 (3.18–3.00)                           |
| Wavelength (Å)                                  |
| 0.97933                                         |
| Total/Unique reflections                        |
| 45479/3013                                      |
| Completeness (%)                                |
| 99.4 (100.0)                                    |
| I/σ                                             |
| 62.8 (16.4)                                     |
| R\textsubscript{merge} (%)                      |
| 10.4 (52.4)                                     |
| **Refinement**                                  |
| Resolution                                      |
| 30.0–3.10                                       |
| No. reflections, working/free                   |
| 2715/140                                       |
| R\textsubscript{work}/R\textsubscript{free} (%) |
| 24.8/29.0                                      |
| No. atoms                                       |
| 884                                             |
| Protein                                         |
| 436                                             |
| DNA                                             |
| 448                                             |
| B factors                                       |
| 69.0                                            |
| RMSD                                            |
| Bond length (Å)                                 |
| 0.012                                           |
| Bond angle (˚)                                  |
| 1.438                                           |
| Ramachandran plot (%)                           |
| Favor                                           |
| 94.1                                            |
| Allowed                                         |
| 5.9                                             |
| Disallowed                                      |
| 0.0                                             |

https://doi.org/10.1371/journal.pone.0221666.t001
Interactions between $\sigma^W_4$ and the -35 promoter element

The recognition helix $\alpha_9$ mediates the major interactions between $\sigma^W_4$ and -35$^W$ through hydrogen bonds and hydrophobic interactions. The bases and backbones of three purine nucleotides (G$_{-35}^W$ and A$_{-34}^W$ in the non-template strand and G'$_{-31}^w$ in the template strand; ' indicates the template strand of DNA) form hydrogen bonds with the residues on the N-terminal half of $\alpha_9$ in $\sigma^W_4$ (Fig 2A–2C and S2 Fig). The guanine oxygen (O6) and backbone phosphate (OP2) of G$_{-35}^W$ form hydrogen bonds with side chain amino groups (NH$_2$) of R175 and R172, respectively. The purine nitrogen (N7) and backbone phosphate (OP2) of A$_{-34}^W$ interact with the side chain oxygens (OG1) of T171 and T168. The guanine oxygen (O6) of G'$_{-31}^w$ forms a hydrogen bond with the side chain amino group (NZ) of K170. The electron density map for the side chain of K170 is relatively weak; however, the most-preferred rotamer is at hydrogen bond distance to the O6 of G'$_{-31}^w$ (S2B Fig).

Hydrophobic interactions are observed between K170-G'$_{-31}^w$, T171-A$_{-34}^W$, H174-T$_{-32}^W$, T$_{-33}^W$, and R175-T$_{-36}^W$ (S3 Fig).

In addition to $\alpha_9$, $\alpha_7$-$\alpha_8$ contributes to -35$^W$ binding without base specificity. The backbone atoms (OP1, OP2, and OP2) of G'$_{-30}^w$ form hydrogen bonds with the side chain amino group (NZ) of K148, the side chain oxygen (OG) of S154, and the backbone nitrogen of L155, respectively (Fig 2B and 2C). Altogether, $\alpha_9$ in $\sigma^W_4$ specifically recognizes -35$^W$, and $\alpha_7$-$\alpha_8$ provides additional contacts, leading to a tighter interaction.
**Structural comparison of σW4/-35W and σE4/-35E**

SigE is an *E. coli* ECF sigma factor activated in response to envelope stress and induces transcription of heat shock proteins [45]. Its σ4 domain (σE4) recognizes the -35 element, of which the consensus sequence is G_{-35}G_{-34}A_{-33}A_{-32}C_{-31}T_{-30}T_{-29} (-35E) [46]. The σE4 structure is highly similar to σW4 [8]. σW4 is superimposed on σE4 with a root mean square deviation (RMSD) value of 1.8 Å for 53 Cα atoms (S4A Fig). The overall fold is conserved between σW4 and σE4 and the main differences are observed at the N- and C-termini (S4A and S4B Fig).

https://doi.org/10.1371/journal.pone.0221666.g002
Overall, the interactions of $\sigma^E_4$ and $\sigma^W_4$ with the corresponding -35 elements are conserved. Three nucleotides, $G_{35}$, $G_{34}$, and $G'_{31}$, in -35$E$, which correspond to $G_{35}$, $A_{34}$, and $G'_{31}$ in -35$W$, mediate purine nucleotide-specific interactions with the recognition $\alpha$-helix (Fig 2A). The backbone and base of $G_{35}$ ($G_{35}$ in $\sigma^W_4$) form hydrogen bonds with R173 (R172 in $\sigma^W_4$) and R176 (R175 in $\sigma^W_4$). The backbone and base of $G_{34}$ ($A_{34}$ in -35$W$) form hydrogen bonds with T169 (T168 in $\sigma^W_4$) and S172 (T171 in $\sigma^W_4$). Although the -34 position is not identical between -35$E$ and -35$W$ ($G_{34}$ in -35$E$ and $A_{34}$ in -35$W$), hydrogen bonds mediated by the backbone phosphate and purine N7 are conserved. The base of $G'_{31}$ forms a hydrogen bond with R171 (R170 in $\sigma^W_4$). The backbone phosphate of $G'_{31}$ also forms a hydrogen bond with R178, which is not observed in the $\sigma^W_4/-35W$ structure. $A'_{30}$ in -35$E$ mediates backbone interactions similarly to $G'_{30}$ in -35$W$ (Fig 2). The backbone phosphate of $A'_{30}$ forms hydrogen bonds with the NH1 of R149 (K148 in $\sigma^W_4$) and the backbone nitrogen of Y156 (L155 in $\sigma^W_4$) (Fig 2D and 2E). Interaction between S154 and $A'_{30}$ in $\sigma^W_4/-35W$ is missing in the $\sigma^E_4/-35E$ structure. In summary, purine base-specific hydrogen bonds in the structures of $\sigma^W_4/-35W$ and $\sigma^E_4/-35E$ are conserved, whereas the hydrogen bonds with the nucleotide backbone are slightly different.

Hydrophobic interactions between $\sigma^E_4$ and -35$E$ are mostly conserved in the $\sigma^W_4/-35W$ structure (S3 Fig). Residues R171, F175, and R176 in $\sigma^E_4$ (K170, H174, and R175 in $\sigma^W_4$) interact with T$'_{32}$, T$'_{33}$, and C$'_{36}$ in -35$E$ ($T'_{32}$, $T'_{33}$, and $T_{36}$ in -35$W$), respectively. Hydrophobic interactions between P166/G168/T169 and $G_{34}$ and between Y156 and $A'_{30}$ are observed only in $\sigma^E_4/-35E$, whereas the interaction between K170 and $G'_{31}$ in $\sigma^W_4/-35W$ is observed only in $\sigma^E_4/-35E$. A cation-$\pi$ interaction is observed between R176 and the pyrimidine ring of C$'_{34}$ in the structure of $\sigma^E_4/-35E$. However, the distance between the base (T$'_{36}$) and the corresponding residue (R175) is too far to form a cation-$\pi$ interaction in the structure of $\sigma^W_4/-35W$ (S5 Fig).

### DNA geometry of -35$W$

Nucleotides $A_{33}A_{32}$ in -35$E$ ($G_{35}G_{34}A_{33}A_{32}C_{31}T_{30}T_{29}$) do not form hydrogen bonds with $\sigma^E_4$, although these nucleotides are conserved among the -35 elements of the SigE regulon, and mutating these nucleotides in characteristic oligo(dA)/oligo(dT)-like DNA geometry that is rigid and straight with a narrow minor groove [8, 48]. A previous structural study of the $\sigma^E_4/-35E$ complex suggested that the geometry of the narrowed minor groove is critical for $\sigma^E_4$ recognition [8], and we show that -35$W$ also displays a narrowed minor groove (Fig 3). Like -35$E$, the narrowing of the minor groove of -35$W$ begins at $A_{33}A_{32}$ and is stabilized downstream of the -35 element, even though the downstream sequence of -35$W$ has an insertion of two cytosines ($A_{33}A_{32}C_{31}C_{30}T_{29}$) (Fig 3). In contrast, $G_{33}A_{32}$ in -35$A$ (the -35 element of Thermus aquaticus SigA) has a wider minor groove than normal B-DNA (Fig 3). The crystal structure of $\sigma^W_4/-35W$ supports the suggestion that the $A_{33}A_{32}$ conservation in the -35 element for group IV sigma factors is critical for the formation of the narrow minor groove [8].

### Structural comparison of $\sigma^W_4/-35W$ and SigW/RsiW

The crystal structure of SigW complexed with anti-sigma RsiW was previously reported [23]. The structures of $\sigma^W_4$ under the binding of -35$W$ and RsiW superimpose with an RMSD value of 1.6 Å for 53 $\alpha$-carbon atoms (S4B Fig). Although conformational differences of $\sigma^W_4$ in the structures bound to -35W and RsiW are minor, $\sigma^W_4$ bound to -35W exists in a slightly compact conformation. $\sigma^W_4$ interacts with -35$W$ through the residues K148, S154, L155, T168, K170, T171, R172, and R175, with a surface area of 621.2 Å$^2$ buried at the binding interface (Fig 4A and
4B). σW interacts with RsiW through the residues I150, K170, H174, R177, E178, R181, R185, and L187, these interactions result in an approximately 50% larger burial of surface area (915.2 Å²) (Fig 4C and 4D). The surface area of σW that binds -35W and RsiW partially overlaps with residue K170 on σW (Fig 4A and 4C), indicating that the interactions of -35W and RsiW with σW are mutually exclusive.

In the crystal structure of the SigW/RsiW complex, the -10 element-binding surface of σW is buried in the surface of σW [23], whereas the -35 element-binding surface of σW is directly blocked by RsiW (Fig 4), suggesting that SigW inhibition by RsiW competes with -35 element binding in the promoter. The binding of the -35 element to σW results in a ΔG of -10.5 kcal/mol, which corresponds to a Kd value of 1.47 x 10⁻⁸ M. RsiW binding to SigW results in a surface burial of 1619.6 Å², and a ΔG of -17.85 kcal/mol. The free energy change corresponds to a Kd value of 4.84 x 10⁻¹⁴ M. These observations indicate that DNA binding of SigW is structurally repressed in the presence of RsiW.

**Role of conserved residues in σW**

*B. subtilis* contains multiple ECF sigma factors that respond to diverse environmental stresses. The positions L137, L147, and E157 of σW are highly conserved in *B. subtilis* ECF sigma factors (Fig 5A and 5B). The highly conserved residues are likely to be involved in the intrinsic folding of SigW, but not in DNA binding. L137 and L147 stabilize σW as part of the central innermost hydrophobic cluster (S6A and S6B Fig). E157 is associated with σW binding in the crystal structure of the SigW/RsiW complex (S6C and S6D Fig).

The residues involved in DNA binding are less conserved than those involved in intrinsic folding. Residues K170, T171, and R175, which mediate purine-specific hydrogen bonds in σW, are aligned to K/R, S/T, and K/R in *B. subtilis* ECF sigma factors, as well as *E. coli* SigE (Fig 5A and 5B). The conservation of these residues correlates with the three nucleotides that mediate purine-specific interactions (G₃₅, A₃₄, and G'₃₁ in -35W). For example, A₃₄ in the SigV promoter interacts with T144 and T147, as does A₃₄ in -35W, and T168/T171 in σW, whereas C₃₄ in the SigZ promoter is aligned with G138 and S141. These observations suggest that sequence variation in the DNA-binding interface confers the specificity required to discriminate between -35 elements (Fig 5C). In summary, conserved residues in *B. subtilis* ECF σ₄ are mainly involved in intramolecular stability and interactions with the -35 element. Slight
sequence variations in the residues involved in the binding of the -35 element may contribute to the fine-tuning of the promoter selectivity and binding affinity for individual ECF sigma factors.

![Fig 4. Structural comparison of $\sigma^W_4$ and SigW/RsiW.](https://doi.org/10.1371/journal.pone.0221666.g004)

![Fig 5. Conserved residues in $\sigma^W_4$.](https://doi.org/10.1371/journal.pone.0221666.g005)
Conclusions

Group IV ECF sigma factor is activated in response to environmental stress and initiates transcription of its own regulon to counter that stress. The crystal structure of $\sigma_4$/-35W shows that SigW selectively recognizes cognate -35 promoter elements which have a narrowed minor groove, in a similar manner to the E. coli SigE. Comparison with the SigW/RsiW structure shows that SigW binding to the -35 promoter element and anti-sigma RsiW is mutually exclusive. These results provide the structural basis for the mechanism of SigW activation, and improve our understanding of the selective interactions between $\sigma_4$ domains and their cognate -35 promoter elements.

Supporting information

S1 Fig. Overall electron density map of $\sigma_4$-35W. Red and orange stick models indicate $\sigma_4$ and -35W, respectively. 2Fo-Fc maps are drawn at two different orientations. The contour level is set to 1.0 $\sigma$.
(TIF)

S2 Fig. Map of electron density around the binding interface in the $\sigma_4$-35W structure. 2Fo-Fc maps are drawn at same scale and orientation as those for models in Fig 2C and arranged in the same order as the panels in Fig 2C. The contour level is set to 1.0 $\sigma$.
(TIF)

S3 Fig. Hydrophobic interactions between the $\sigma_4$ domain and the -35 promoter element. (A, B) Hydrophobic interactions between $\sigma_4$ and -35W shown at two different orientations. Residues and nucleotides, which are associated with hydrophobic bonds, are drawn as red and orange stick models. Dotted lines indicate hydrophobic interactions. (C, D) 2Fo-Fc electron density maps in (C) and (D) are shown at same scale and orientation as those for models in S2A and S2B Fig, respectively. The contour level is set to 1.0 $\sigma$. (E, F) Hydrophobic interactions between $\sigma_4$ and -35E. Residues and nucleotides which are associated with hydrophobic bonds are drawn as purple and green stick models.
(TIF)

S4 Fig. Structural comparison of $\sigma_4$ domains. (A) Superposition of $\sigma_4$-35E and $\sigma_4$-35W structures. N- and C-termini of $\sigma_4$ are labeled. (B) Distribution of root-mean-square values between Cα positions of superimposed $\sigma_4$ and $\sigma_4$ structures. (C) Superposition of $\sigma_4$ domains from the structures of $\sigma_4$-35W and SigW/RsiW.
(TIF)

S5 Fig. Cation-π interaction. (A) The green dotted line indicates the cation-π interaction between R176 of $\sigma_4$ (cyan model) and C-36 of 35E (green). The corresponding residue in $\sigma_4$ (R175) and base in -35W (T-36) are drawn as a stick model. The distance between the Arg and pyrimidine is labeled.
(TIF)

S6 Fig. Conserved residues of $\sigma_4$ domains. The residues that participate in intramolecular interactions with L137 (A), L147 (B), or E157 (C) are drawn as stick models and the residue number is labeled. The helix and loop in $\sigma_4$ are colored pink and light brown. (D) $\sigma_4$ in SigW/RsiW structure (PDB ID: 5WUQ) is superposed onto that of the $\sigma_4$-35W structure. SigW in SigW/RsiW structure is colored green. E157 in $\sigma_4$ interacts with T71 in $\sigma_2$ and does not participate in DNA binding.
(TIF)
Author Contributions

Conceptualization: Eunju Kwon, Dong Young Kim.

Data curation: Eunju Kwon.

Formal analysis: Eunju Kwon, Shankar Raj Devkota, Deepak Pathak, Pawan Dahal, Dong Young Kim.

Funding acquisition: Dong Young Kim.

Investigation: Eunju Kwon, Shankar Raj Devkota, Deepak Pathak, Pawan Dahal, Dong Young Kim.

Methodology: Eunju Kwon, Shankar Raj Devkota, Deepak Pathak, Pawan Dahal.

Project administration: Eunju Kwon, Dong Young Kim.

Resources: Eunju Kwon.

Software: Eunju Kwon.

Supervision: Dong Young Kim.

Validation: Dong Young Kim.

Visualization: Eunju Kwon.

Writing – original draft: Eunju Kwon, Dong Young Kim.

Writing – review & editing: Eunju Kwon, Dong Young Kim.

References

1. Burgess RR, Travers AA, Dunn JJ, Bautz EK. Factor stimulating transcription by RNA polymerase. Nature. 1969; 221(5175):43–6. Epub 1969/01/04. https://doi.org/10.1038/221043a0 PMID: 4882047.

2. Saecker RM, Record MT Jr., Dehaseth PL. Mechanism of bacterial transcription initiation: RNA polymerase—promoter binding, isomerization to initiation-competent open complexes, and initiation of RNA synthesis. J Mol Biol. 2011; 412(5):754–71. Epub 2011/03/05. https://doi.org/10.1016/j.jmb.2011.01.018 PMID: 21371479; PubMed Central PMCID: PMC3440003.

3. Feklistov A, Sharon BD, Darst SA, Gross CA. Bacterial sigma factors: a historical, structural, and genomic perspective. Annu Rev Microbiol. 2014; 68:357–76. https://doi.org/10.1146/annurev-micro-092412-155737 PMID: 25002089.

4. Paget MS. Bacterial Sigma Factors and Anti-Sigma Factors: Structure, Function and Distribution. Biomolecules. 2015; 5(3):1245–65. Epub 2015/07/02. https://doi.org/10.3390/biom5031245 PMID: 26131973; PubMed Central PMCID: PMC4598750.

5. Feklistov A, Darst SA. Structural basis for promoter-10 element recognition by the bacterial RNA polymerase sigma subunit. Cell. 2011; 147(6):1257–69. https://doi.org/10.1016/j.cell.2011.10.041 PMID: 22136875; PubMed Central PMCID: PMC3245737.

6. Campagne S, Marsh ME, Capitani G, Vorholt JA, Allain FH. Structural basis for -10 promoter element melting by environmentally induced sigma factors. Nat Struct Mol Biol. 2014; 21(3):269–76. https://doi.org/10.1038/nsmb.2777 PMID: 24531660.

7. Campbell EA, Muzzin O, Chlenov M, Sun JL, Olson CA, Weinman O, et al. Structure of the bacterial RNA polymerase promoter specificity sigma subunit. Mol Cell. 2002; 9(3):527–39. PMID: 11931761.

8. Lane WJ, Darst SA. The structural basis for promoter -35 element recognition by the group IV sigma factors. PLoS Biol. 2006; 4(9):e269. https://doi.org/10.1371/journal.pbio.0040269 PMID: 16903784; PubMed Central PMCID: PMC1540707.

9. Wigneshweraraj S, Bose D, Burrows PC, Joly N, Schumacher J, Rappas M, et al. Module operandi of the bacterial RNA polymerase containing the sigma54 promoter-specificity factor. Mol Microbiol. 2008; 68(3):538–46. Epub 2008/03/12. https://doi.org/10.1111/j.1365-2958.2008.06181.x PMID: 18331472.

10. Helmann JD. The extracytoplasmic function (ECF) sigma factors. Adv Microb Physiol. 2002; 46:47–110. Epub 2002/06/21. PMID: 12073657.
11. Osterberg S, del Peso-Santos T, Shinger V. Regulation of alternative sigma factor use. Annu Rev Microbiol. 2011; 65:37–55. Epub 2011/06/07. https://doi.org/10.1146/annurev.micro.112408.134219 PMID: 21639785.

12. Staron A, Sofia HJ, Dietrich S, Ulrich LE, Liesegang H, Mascher T. The third pillar of bacterial signal transduction: classification of the extracytoplasmic function (ECF) sigma factor protein family. Mol Microbiol. 2009; 74(3):557–81. Epub 2009/09/10. https://doi.org/10.1111/j.1365-3958.2009.06870.x PMID: 19737356.

13. Lonetto M, Gribskov M, Gross CA. The sigma 70 family: sequence conservation and evolutionary relationships. J Bacteriol. 1992; 174(12):3843–9. Epub 1992/06/01. https://doi.org/10.1128/jb.174.12.3843-3849.1992 PMID: 1597408; PubMed Central PMCID: PMC206090.

14. Zellmeier S, Schumann W, Wiegert T. Involvement of Clp protease activity in modulating the Bacillus subtilis sigma(W) regulon. Mol Microbiol. 2009; 74(3):1569–82. Epub 2009/09/05. https://doi.org/10.1111/j.1365-3958.2009.06509.x PMID: 19752122.

15. Steil L, Hoffman T, Budde I, Volker U, Bremer E. Genome-wide transcriptional profiling analysis of adaptation of Bacillus subtilis to high salinity. J Bacteriol. 2003; 185(21):6358–70. Epub 2003/10/18. https://doi.org/10.1128/JB.185.21.6358-6370.2003 PMID: 14563871; PubMed Central PMCID: PMC2612467.

16. Haugen SP, Berkmen MB, Ross W, Gaal T, Ward C, Gourse RL. rRNA promoter regulation by nonoptimal binding of sigma region 1.2: an additional recognition element for RNA polymerase. Cell. 2006; 125(6):1069–82. Epub 2006/06/17. https://doi.org/10.1016/j.cell.2006.04.034 PMID: 16777598.

17. Pietiainen M, Gardemeister M, Mecklin M, Leskelä S, Sarvas M, Kontinen VP. Cationic antimicrobial peptides elicit a complex stress response in Bacillus subtilis that involves ECF-type sigma factors and two-component signal transduction systems. Microbiology. 2005; 151(8):1577–92. https://doi.org/10.1099/mic.0.27761-0 PMID: 15870467.

18. Asai K. Anti-sigma factor-mediated cell surface stress responses in Bacillus subtilis sigmaW stress response. Mol Microbiol. 2004; 52(3):591–9. https://doi.org/10.1046/j.1365-2958.2004.03050.x PMID: 12207695.

19. Kunst F, Ogasawara N, Moszer I, Albertini AM, Alloni G, Azevedo V, et al. The complete genome sequence of the gram-positive bacterium Bacillus subtilis. Nature. 1997; 390(6657):249–56. https://doi.org/10.1038/36786 PMID: 9384377.

20. Liu L, Fang C, Zhuang N, Wang T, Zhang Y. Structural basis for transcription initiation by bacterial ECF sigma factors. Nat Commun. 2015; 10(1):1153. Epub 2015/09/13. https://doi.org/10.1038/ncomms11553 PMID: 26393243; PubMed Central PMCID: PMC4411747.
30. Hubin EA, Fay A, Xu C, Bean JM, Saecker RM, Glickman MS, et al. Structure and function of the mycobacterial transcription initiation complex with the essential regulator RbpA. Elife. 2017; 6. Epub 2017/01/10. https://doi.org/10.7554/eLife.22520 PMID: 28067618; PubMed Central PMCID: PMC5302886.

31. Park SY, Ha SC, Kim YG. The protein crystallography beamlines at the pohang light source II. Biodesign. 2017; 5:30–4.

32. Battye TG, Kontogiannis L, Johnson O, Powell HR, Leslie AG. iMOSFLM: a new graphical interface for diffraction-image processing with MOSFLM. Acta Crystalogr D Biol Crystalogr. 2011; 67(Pt 4):271–81. Epub 2011/04/05. https://doi.org/10.1107/S0907444910048675 PMID: 21460445; PubMed Central PMCID: PMC3069742.

33. McCoy AJ, Grosse-Kunstleve RW, Adams PD, Winn MD, Storoni LC, Read RJ. Phaser crystallographic software. J Appl Crystallogr. 2007; 40(Pt 4):658–74. Epub 2007/08/01. https://doi.org/10.1107/S0021889807021206 PMID: 19461845; PubMed Central PMCID: PMC2483472.

34. Afonine PV, Grosse-Kunstleve RW, Echols N, Headd JJ, Moriarty NW, Mustyakov M, et al. Towards automated crystallographic structure refinement with phenix.refine. Acta Crystallogr D Biol Crystallogr. 2012; 68(Pt 4):352–67. https://doi.org/10.1107/S0907444912001308 PMID: 22505256; PubMed Central PMCID: PMC3322595.

35. Emsley P, Lohkamp B, Scott WG, Cowtan K. Features and development of Coot. Acta Crystallogr D Biol Crystallogr. 2010; 66(Pt 4):486–501. https://doi.org/10.1107/S0907444910007493 PMID: 20383002; PubMed Central PMCID: PMC2852313.

36. Holm L, Laakso LM. Dali server update. Nucleic Acids Res. 2016; 44(W1):W351–5. Epub 2016/05/01. https://doi.org/10.1093/nar/gkw357 PMID: 27131377; PubMed Central PMCID: PMC4987910.

37. Laskowski RA, Swindells MB. LigPlot+: multiple ligand-protein interaction diagrams for drug discovery. J Chem Inf Model. 2011; 51(10):2778–86. https://doi.org/10.1021/ci200227u PMID: 21919503.

38. Krissinel E, Henrick K. Inference of macromolecular assemblies from crystalline state. J Mol Biol. 2007; 372(3):774–97. https://doi.org/10.1016/j.jmb.2007.05.022 PMID: 17681537.

39. Lu XJ, Olson WK. 3DNA: a versatile, integrated software system for the analysis, rebuilding and visualization of three-dimensional nucleic-acid structures. Nat Protoc. 2008; 3(7):1213–27. Epub 2008/07/05. https://doi.org/10.1038/nprot.2008.104 PMID: 18600227; PubMed Central PMCID: PMC3065354.

40. Jurrus E, Engel D, Star K, Monson K, Brandi J, Felberg LE, et al. Improvements to the APBS biomolecular solvation software suite. Protein Sci. 2018; 27(1):112–28. Epub 2017/08/25. https://doi.org/10.1002/pro.3280 PMID: 28836357; PubMed Central PMCID: PMC5734301.

41. Schrodinger, LLC. The PyMOL Molecular Graphics System, Version 1.8. 2015.

42. Barton GJ. ALSCRIPT: a tool to format multiple sequence alignments. "Protein Engineering, Design and Selection". 1993; 6(1):37–40. https://doi.org/10.1093/protein/6.1.37

43. Huang X, Fredrick KL, Helmann JD. Promoter recognition by Bacillus subtilis sigmaw: autoregulation and partial overlap with the sigmaw regulon. J Bacteriol. 1998; 180(15):3765–70. PMID: 9695469; PubMed Central PMCID: PMC107356.

44. Crooks GE, Hon G, Chandonia JM, Brenner SE. WebLogo: a sequence logo generator. Genome Res. 2004; 14(6):1188–90. Epub 2004/06/03. https://doi.org/10.1101/gr.849004 PMID: 15173120; PubMed Central PMCID: PMC419797.

45. Erickson JW, Gross CA. Identification of the sigma E subunit of Escherichia coli RNA polymerase: a second alternate sigma factor involved in high-temperature gene expression. Genes Dev. 1989; 3 (9):1462–71. Epub 1989/09/01. https://doi.org/10.1101/gad.3.9.1462 PMID: 2691330.

46. Rhodius VA, Suh WC, Nonaka G, West J, Gross CA. Conserved and variable functions of the sigmaE stress response in related genomes. PLoS Biol. 2006; 4(1):e2. https://doi.org/10.1371/journal.pbio.0040002 PMID: 16336047; PubMed Central PMCID: PMC1312014.

47. Miticka H, Rezuchova B, Homorova D, Roberts M, Kormanec J. Identification of nucleotides critical for activity of the sigmaE-dependent rpoEp3 promoter in Salmonella enterica serovar Typhimurium. FEMS Microbiol Lett. 2004; 238(1):227–33. Epub 2004/09/01. https://doi.org/10.1016/j.femsle.2004.07.039 PMID: 15336426.

48. Haran TE, Mohanty U. The unique structure of A-tracts and intrinsic DNA bending. Q Rev Biophys. 2009; 42(1):41–81. Epub 2009/06/11. https://doi.org/10.1017/S0033583509004752 PMID: 19508739.