The Role of SwrA, DegU and $P_{D3}$ in fla/che Expression in B. subtilis

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Abstract

In B. subtilis swarming and robust swimming motility require the positive trigger of SwrA on fla/che operon expression. Despite having an essential and specific activity, how SwrA executes this task has remained elusive thus far. We demonstrate here that SwrA acts at the main $\sigma^A$-dependent fla/che promoter $P_{A(\text{fla/che})}$, through DegU. Electrophoretic mobility shift assays (EMSA) reveal that SwrA forms a complex with the phosphorylated form of DegU (DegU~P) at $P_{A(\text{fla/che})}$ while it is unable to do so with either unphosphorylated DegU or the DegU32(Hy) mutant protein. Motility assays show that a highly phosphorylated DegU is not detrimental for flagellar motility provided that SwrA is present; however, DegU~P represses $P_{A(\text{fla/che})}$ in the absence of SwrA. Overall, our data support a model in which DegU~P is a dual regulator, acting either as a repressor when alone or as a positive regulator of $P_{A(\text{fla/che})}$ when combined with SwrA. Finally, we demonstrate that the $\sigma^D$-dependent $P_{D3(\text{fla/che})}$ promoter plays an important role in motility, representing a contingent feedback loop necessary to maintain basal motility when swrA is switched to the non-functional swrA* status.

Introduction

Regulation of flagellar motility in Bacillus subtilis appears to be exerted at the level of the fla/che operon in which most flagellum components are encoded (reviewed in 1). fla/che contains the gene for the alternative sigma factor $\sigma^D$, which is needed for the transcription of the flagellin gene $hag$, as well as of a number of additional genes. Two promoter sequences drive fla/che transcription: $P_{D3(\text{fla/che})}$ and $P_{A(\text{fla/che})}$ [2]. The $\sigma^D$-dependent $P_{A(\text{fla/che})}$ is necessary and sufficient for fla/che expression and motility whereas $P_{D3(\text{fla/che})}$, which is dependent on $\sigma^D$ for activation, is not sufficient to promote motility and its involvement in a positive feedback effect on fla/che expression could not be demonstrated [2,3].

Flagella are necessary for both swimming and swimming motility. Swarming is the typical motility in liquid media, while swarming occurs on semi-solid surfaces. The latter form, described in B. subtilis by Keams and Losick in 2003 [4], requires SwrA to ensure the optimal activation of $P_{A(\text{fla/che})}$ transcription which occurs through a still unclear mechanism [5,6]. Additionally, swimming requires surfactin production, which does not occur in laboratory strains because of a mutation in the surfactin biosynthetic pathway [7]. Thus swarming can be analyzed in laboratory strains if two conditions are met: i) the swrA allele is in the functional swrA+ form; ii) surfactin is added in the medium during the assay [5].

Swimming is boosted by SwrA but it also takes place in its absence, albeit at a reduced rate [8]. The wild-type swrA allele is typically found in undomesticated strains; in most laboratory strains, e.g. 168, the swrA coding sequence contains a nucleotide insertion that prematurely interrupts its reading frame [5,9]. The inactivating mutation occurs in a mononucleotide repeat sequence and can easily shift back and forth with very high frequency ($10^4$). Thus, the alternation between the functional and non-functional swrA alleles is more typical of phase variation mechanisms than point mutations [1,5] and B. subtilis cultures are likely to include both swrA- and swrA* cells.
Transcription of swrA is mainly σ^5-dependent and is positively autoregulated through a circuit that also sustains fla/che expression: SwrA promotes fla/che and thus sigD - transcription and σ^5 transcribes swrA (Figure 1A) [8].

Besides the regulatory effect of SwrA, several studies have established that fla/che operon expression is also regulated in a complex way by DegS/DegU, a two-component system (TCS) that controls important stationary-phase behaviours in *B. subtilis*. DegU undergoes phosphorylation and dephosphorylation by DegS, which is both a kinase and a phosphatase [10]. It was observed that a non-phosphorylatable DegU mutant causes a non-swarming phenotype in swrA^- strains, leading to the conclusion that DegU phosphorylation is required for swarming motility [11,12]. However, employing the degU32(Hy) mutant allele (described below), the repressive effect of DegU^-P on motility has been repeatedly observed both in swrA^- and swrA^+ strains [11-16]. It has been proposed that the response regulator DegU can be phosphorylated at different levels allowing it to tune gene expression as a kind of rheostat according to the extent of its phosphorylation. Specifically, at a low level of phosphorylation DegU^-P is necessary for swarming; conversely, a highly phosphorylated DegU, as DegU32(Hy), inhibits fla/che transcription [10].

degU32(Hy) belongs to a group of several mutants that share the pleiotropic “Hy” phenotype which is characterized by the ability to secrete high levels of degradative enzymes, such as proteases and levansucrase, low or no competence, as well as by the lack of flagella [13,17,18]. The “Hy” phenotype is caused by single nucleotide changes in the coding sequence of degU or degS that perturb the DegS/DegU signal transduction pathway, increasing the stability of phosphorylated DegU in the cell. In degU32(Hy) a H12L amino acid substitution in DegU [18] extends the half-life of the phosphorylated form from 20 to 140 min leading to a higher steady-state level of DegU^-P [19]. A very similar phenotype can also be obtained by the over-expression of DegQ or DegR which are small proteins that interfere with the trans-phosphorylation/dephosphorylation cycle between DegS and DegU [13,17,18,20-22]: DegR stabilizes DegU^-P [23] and DegQ facilitates phosphate transfer from DegS to DegU [11]. Currently degU32(Hy) is the “Hy” mutant most often used in studies on the DegS/DegU TCS and it is considered as an otherwise wild-type more stable DegU^-P.

We have previously hypothesized that DegU can regulate motility cooperatively with SwrA based on evidence that SwrA cannot induce swarming in the absence of a functional degSU operon and that deletion of degS/degU has no motility phenotype in swrA^- strains [8,13]. Furthermore, it was recently shown that SwrA is able to modulate the ability of DegU to activate transcription of ycdA [24]. In this work we demonstrate that SwrA forms a complex with DegU^-P at P_{A(fla/che)} under conditions that correspond to those leading to activation of fla/che transcription. We show that DegU^-P obtained through the use of DegS200(Hy), a different “Hy” mutant, is not inhibitory on motility when SwrA is functional; conversely, without SwrA, DegU^-P produced by degS200(Hy) or degU32(Hy) is equally repressing P_{A(fla/che)}. We
also show that DegU32(Hy)-P behaves as a constitutive repressor since it does not interact with SwrA on P_{fla/che}. Overall, our data suggest that DegU phosphorylation, independently of its level, is required for both activation and repression of P_{fla/che}. Upon phosphorylation, the positive or negative outcome of DegU→P on fla/che expression depends on the presence or absence respectively of SwrA.

Finally, we demonstrate that when the SwrA-based autoregulatory loop is non-functional P_{D3} contributes to maintaining fla/che expression through an ancillary positive feedback loop (Figure 1B). Thus, P_{D3} represents a sort of contingency plan for ensuring a basal level of motility that constitutes a primary survival resource for B. subtilis.

Materials and Methods

Bacterial strains and growth conditions

*Bacillus subtilis* strains used in this study with their relevant genotype are listed in Table S1 in Supporting Information. Strains were routinely grown on LB (Luria-Bertani broth: 10 g of tryptone, 5 g of yeast extract, 10 g of NaCl per liter) supplemented with 1.5% agar unless otherwise stated. When required, selective agents were added at the following concentrations: erythromycin 1 or 50 μg ml\(^{-1}\), kanamycin 2 μg ml\(^{-1}\), spectinomycin 60 μg ml\(^{-1}\). The *E. coli* DH5α or BL21 and BL21(DE3) strains were grown at 37°C in LB with 100 μg ml\(^{-1}\) ampicillin and 20 μg ml\(^{-1}\) chloramphenicol when required.

Strain construction

Plasmids and oligonucleotides used in this work are listed in Tables S2 and S3 respectively, in Supporting Information. The markerless P_{D3} deletion, that removes 118 bp from the fla/che upstream region (between positions -302 and -184 from the flgB translation start site) (Figure 2A), was generated with pMADΔPD3. For its construction two separate PCR fragments containing the two flanking regions of P_{D3} were amplified from genomic DNA using the primer pairs CodYF and 8465 and 8601 and nRflgBN (downstream region). The two fragments were blunt-end ligated and re-amplified with the external primers CodYF and 8465 (upstream region) and 8601 and nRflgBN (downstream region).

Recombinant protein production and purification

For DegU and DegU32(Hy) production, competent *E. coli* BL21(DE3) cells were transformed with pET16Uwt or pET16Uhy plasmids (Table S2), grown at 37°C and induced with 1 mM of IPTG for 3 hours at 28°C. Cells were collected by centrifugation, resuspended in 0.02 volumes of buffer A (20 mM Tris-HCl pH 8, 300 mM NaCl, 1% Triton X-100, 0.5 mM PMSF, 5 μg ml\(^{-1}\) DNaseI, 20 mM imidazole) and sonicated. Cell lysate was centrifuged at 10,000 g at 4°C for 15 minutes to remove cellular debris; the supernatant was incubated with Co-IDA agarose beads (Biontex), previously equilibrated in buffer B (20 mM Tris-HCl pH 8, 300 mM NaCl, 20 mM imidazole), at 4°C for 1 hour. After extensive washing with buffer B, proteins were eluted with 500 mM imidazole in buffer B; protein-containing fractions were pooled and dialyzed against buffer C (20 mM Tris-HCl pH 8, 1 mM EDTA). The dialyzed proteins were loaded on a 5 ml DEAE column (GE Healthcare) to remove any contaminant DNA and eluted with a discontinuous NaCl gradient; after SDS-PAGE analysis, selected fractions were dialyzed against buffer D (20 mM Tris-HCl pH 8, 200 mM NaCl, 1 mM EDTA, 50% [v/v] glycerol) and stored at -80°C in aliquots.

degS and degS200(Hy) genes were amplified from PB5249 or PB5390 (Table S1), respectively, with primers DegSFcacc and DegSRblunt. Products were cloned into the directional pET101/D-TOPO plasmid (Invitrogen) and plasmids were sequenced. The resulting pETOPO-DegS or pETOPO-DegSHy together with pGro7 (Takara Bio) were used to transform *E. coli* BL21(DE3) according to the manufacturer's protocol. DegS and DegS200Hy were purified according to the same protocol used for DegU and DegU32Hy, omitting the DEAE step. GST-SwrA and SwrA purification has already been described [27]. All proteins were dialyzed against buffer D, which was also used to normalize protein volumes in EMSA. Protein purity was assessed by SDS-PAGE and concentration was calculated...
Figure 2. SwrA is bound to P_{fla/che} through DegU-P. (A) The region of the fla/che promoter, including the σ^A and σ^D-dependent promoters, is drawn to scale. The coding sequences of the preceding gene (codY) and of the first gene of fla/che (flgB) are in black. The span of the P_{fla/che} probe, of the K probe used in Figure S1 and of the markerless P_{D3} deletion are indicated. The bar corresponds to 50 nt. (B) EMSA with fluorescently labeled P_{fla/che} probe. DegU, DegS and SwrA (listed on the left above the gel) were added to reactions labeled with a + above the lane. In each 10 μL-reaction proteins had the following concentrations: 0.35 μM for DegU; 0.4 μM for DegS; 1 μM for SwrA. ATP was omitted from the reaction in lane 9. On the right of the gel, the arrowhead labeled with S points to the probe shifted by DegU-P (lane 3); the SS arrow marks the super-shifted complex formed upon SwrA addition (lane 8). (C) Competition reactions with an excess (20 ng) of unlabeled P_{fla/che} DNA or with the same amount of unlabeled non-specific (N-S) DNA are indicated above the gel. The EMSA probe was radioactively labeled. In each reaction, the proteins had following concentration: 0.2 μM DegU, 0.1 μM DegS and 0.2 μM SwrA. ATP was omitted from the reaction in lane 9. On the right of the gel, the arrowhead labeled with S points to the probe shifted by DegU-P (lane 3); the SS arrow marks the super-shifted complex formed upon SwrA addition (lane 8). (D) A GST-tag fused to SwrA reduces the migration of the SS complex. Radioactively labeled P_{fla/che} was incubated with DegU (0.1 μM), DegS (0.1 μM), and either SwrA (lane 6), GST-SwrA (lane 7) or GST (lane 8), each at 1.5 μM. The grey arrow labeled with SS^* points at the retarded band obtained with GTS-SwrA (lane 7).

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through absorbance at 280 nm and with a Bradford Protein Assay (Bio-Rad).

EMSA

EMSA were performed using two different labeling and detection procedures. The P_{A(fla/che)} probe (Figure 2A) was amplified from PB5249 DNA either with primers cs8610 and nRflgBN (285 bp) and end-labeled with [γ-32P]-ATP using T4 polynucleotide kinase, or with 6-FAM-fluorescent-labeled primers 8601 and nRflgB (299 bp). A shorter K probe (156 bp, Figure 2A), amplified with 6-FAM-fluorescent-labeled primer 8601 and a non-labeled primer 8756, was also used in EMSA (Figure S1). Probes were purified by silica microcolumns. Binding reactions (10 μl) contained: 12 fmol of the labeled dsDNA, 0.1 μg of sonicated salmon sperm DNA, 0.1 mM ATP, binding buffer BB (50 mM Tris-HCl pH 7.6, 10 mM MgCl₂, 1 mM DTT, 25 mM KCl, 0.1 mg ml⁻¹ BSA) and a total volume of 3 μl of the required proteins in different final concentrations, as specified in each figure legend; volumes were normalized to 3 μl using buffer D. Reaction mixtures were incubated for 10 min at room temperature (26°C), in the dark for FAM-labeled probes, and were then resolved on a native 5% polyacrylamide gel (6.7 mM Tris-HCl pH 7.6, 3.3 mM Na-acetate, 1 mM EDTA, 5% acrylamide, 2.5% glycerol). The gel was pre-run at 120 V at 4°C, in a low ionic strength TAE buffer (6.7 mM Tris-HCl pH 7.6, 3.3 mM Na-acetate, 1 mM EDTA) for 1 hour; samples were separated at 190 V at 4°C for 3 hours. Gels were either dried for radioactive signal acquisition with a Cyclone Plus Phosphor Imager system (Perkin Elmer), or directly analyzed with a Typhoon scanner (GE Healthcare) for FAM-labeled substrates.

Results

SwrA binds to the fla/che promoter together with DegU-P

Previous work has demonstrated a positive effect of the functional SwrA protein on fla/che transcription [6] but the method of operation of SwrA has remained unclear. SwrA does not contain any known DNA binding domain, therefore its effect on fla/che is unlikely to be due to direct binding to the P_{A(fla/che)} promoter. However, an indirect interaction with P_{A(fla/che)} through a bridging factor has never been explored. The discovery that DegU binds to P_{A(fla/che)} [11,16] as well as the evidences suggesting a cooperative effect between DegU and SwrA [8,24], prompted us to investigate the possible interaction of the two proteins at the P_{A(fla/che)} locus. Initially, pull-down assays were attempted using SwrA, DegU and DegS proteins in several different conditions and prey/bait combinations but the observed strength of interaction was not satisfactory. Indeed, difficulties to assess an interaction between SwrA and DegU have been previously noted [24]. Two possible reasons account for this result: i) the interaction between these two proteins, which is mediated by a transient phosphorylation event (described below), is intrinsically unstable in vitro; ii) DegU, DegS and SwrA bind to Glutathione and Nickel-charged resins aspecifically, in all the conditions tested, causing residual signals in the negative control reactions. As an alternative, EMSA are reported to be more sensitive than other types of assay for the detection of unstable interactions as the gel matrix intervenes to stabilize labile complexes through a "caging" effect [28]. Therefore, SwrA, DegU and DegS were challenged with a PCR probe (ca 300 bp) spanning the P_{A(fla/che)} region (Figure 2A) in EMSA.

We observed that DegU showed higher affinity for P_{A(fla/che)} when phosphorylated by DegS kinase (Figure 2B) as previously reported [11], and formed a complex with DNA whose mobility is marked with an "S" in Figure 2. ATP was required for the interaction (data not shown). However, when SwrA was added to the reaction containing DegS and DegU the probe was super-shifted in a complex (SS complex in Figure 2) with a slower mobility than the S complex (DegU-P:DNA). The SS complex was not assembled when ATP was omitted from the reaction, thus indicating active DegU phosphorylation was required. SwrA did not bind the DNA probe, either alone or in combination with DegS or with unphosphorylated DegU. To verify the specificity of both S and SS complexes an excess of either the unlabelled P_{A(fla/che)} probe or an unrelated DNA fragment was added to the reactions. Whereas the unlabeled P_{A(fla/che)} was able to outcompete the labeled probe, the non-specific DNA fragment could not (Figure 2C) thus confirming that both complexes were specifically occurring on P_{A(fla/che)}. Both the S and SS complexes were also obtained with a shorter probe, K, spanning a narrower region (156 bp) around P_{A(fla/che)} (Figure 2A and Figure S1). The robustness of S and SS complexes was corroborated by their reproducibility over a wide range of concentrations of the three proteins used in the reactions, as indicated in the legends of Figures 2, 4, S1 and S2.

Next we sought to determine whether the slower migration of the SS complex was dependent on an SwrA-induced conformational change of DegU-P:DNA or rather on the association of SwrA with the S complex. To discriminate between these two possibilities SwrA (14 kDa) was substituted in EMSA by a higher molecular mass GST-SwrA fusion protein (41 kDa). No differences with the untagged SwrA were expected in the case that SwrA had an indirect effect on mobility of the complexes, whereas a further decrease in migration of the SS complex was anticipated in the case that SwrA was integral to the DNA-bound complex. As shown in Figure 2D, the sole GST moiety used as a control did not interfere with the assay. GST-SwrA behaved as the untagged SwrA: it did not interact with DNA alone (not shown) or in combination with non-phosphorylated DegU, whereas it was able to super-shift the probe in the presence of DegU-P (SS* in Figure 2C, lane 7). In this case, however, the mobility of the complex was indeed slower than the SS complex obtained with untagged SwrA, demonstrating that SwrA was physically associated with the DNA-bound complex. Moreover, in reactions in which both DegU and DegS concentrations were limiting we observed a substantial increase in the efficiency of DNA binding in the presence of SwrA (Figure S2).
Thus, SwrA modifies the mobility of DegU~P:P<sub>fla/che</sub> by taking part in the complex and increasing its affinity for DNA. DegU phosphorylation is required for such an interaction.

The role of phosphorylation on DegU and the nature of DegU32(Hy)

Our results indicate that DegU phosphorylation is necessary for the interaction with the swarming factor SwrA, in line with previous work that highlighted the necessity of a phosphorylatable DegU for swarming [11,12]. However, it has also been repeatedly observed that high levels of DegU~P, as obtained with the degU32(Hy) allele, cause a non-motile phenotype independent of the status of the swrA gene [11-16]. In order to investigate the relevance of the extent of phosphorylation of the transcription factor DegU we analyzed motility under conditions in which a wild-type DegU is highly phosphorylated. For this purpose we used the degS200(Hy) mutant allele which causes a pleiotropic phenotype whose characteristics and severity are indistinguishable from that of degU32(Hy) [13,17]. degS200(Hy) leads to a G218E substitution in DegS that impairs its dephosphorylation activity leading to accumulation of a wild-type DegU~P [29]. When strains were transformed with the degS200(Hy) allele swimming and swarming motility (on surfactin-supplemented plates) greatly differed between the isogenic swrA<sup>+</sup> and swrA<sup>−</sup> backgrounds (Figure 3). In the swrA<sup>−</sup> degS200(Hy) strain swimming was repressed compared to the wild-type degS/degU sibling, in line with previous reports and with what occurs with degU32(Hy). Conversely, in the swrA<sup>+</sup> background the accumulation of the DegU~P form mediated by degS200(Hy) did not preclude robust swimming and swarming motility. These findings indicate that the higher extent of DegU phosphorylation caused by impairment of the phosphatase activity of DegS does not hinder motility when SwrA is present.

**Figure 4.** DegU32(Hy) and DegS200(Hy) behave differently in EMSA. Fluorescently labeled P<sub>fla/che</sub> was challenged with the mutant DegU32(Hy) or DegS200(Hy) proteins in EMSA. DegS200(Hy) was used to phosphorylated wild-type DegU in lanes 10 and 11. DegU32(Hy) was phosphorylated by wild-type DegS in lanes 5 and 6. Protein concentrations were the following: 0.2 μM for DegS; 0.4 μM for DegS200(Hy); 0.36 μM for DegU and 1 μM for SwrA. Different concentrations of DegU32(Hy) were used: 0.3 μM in lane 2; 0.6 μM in lanes 3, 5 and 6; 1.8 μM in lane 4. Control reactions with wild-type proteins are shown in lanes 7, 8, 9. An asterisk marks the particular pattern of bands, bracketed on the left-hand side of the gel, produced by DegU32(Hy) and DegU32(Hy)~P. Other symbols are the same as in Figure 2B.

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but is indeed repressive when SwrA is absent. Therefore, the level of phosphorylation attained by DegU is not the cause of motility inhibition per se. Indeed, wild B. subtilis strains carry the degQ36(Hy) allele that increases the phosphorylation rate of DegU but are nonetheless swarming proficient [11].

To elucidate the reason for the different behavior of degU32(Hy) and degS200(Hy) alleles, the corresponding mutant proteins were purified and challenged with the P_{A(fla/che)} probe in EMSA. As shown in Figure 4, DegS200(Hy) behaved exactly as its wild-type counterpart: it was able to phosphorylate wild-type DegU and induced the formation of the SS complexes in the presence of SwrA. Conversely, DegU32(Hy) bound DNA efficiently, even at low concentration and in the absence of DegS, causing a distinctive band-shift pattern (indicated by an asterisk in Figure 4). This pattern was not substantially modified by the addition of DegS to the reaction and was not efficiently converted into the SS complex upon addition of both DegS and SwrA. Since the mutant DegU32(Hy) protein can be trans-phosphorylated and dephosphorylated by DegS in vivo [13] and in vitro [19], the absence of the SS complex was not due to a lack of phosphorylation. Thus, the most likely explanation is that the H12L amino acid substitution in DegU32(Hy) not only causes a slow dephosphorylation rate, as already assessed [19], but also impairs its interaction with SwrA at P_{A(fla/che)} in vitro, preventing the assembly of the SS complex. In agreement with this hypothesis, the H12L mutation is localized in the N-terminal domain of DegU, which is the domain that interacts with SwrA in Yeast-2-Hybrid assays [24].

The results obtained in vivo and in vitro with DegU32(Hy) and DegS200(Hy) demonstrate that DegU phosphorylation, even at high levels, is not necessarily linked to motility repression; rather, that the ability to interact with SwrA is crucial for swarming and swimming. Indeed, in the absence of SwrA repression of fla/che transcription occurs with both DegU32(Hy) and DegS200(Hy), indicating that DegU~P by itself acts as a P_{A(fla/che)} repressor. The full motility of the degS200(Hy) swrA+ strain in vivo coupled with the ability of wild-type DegU~P [but not DegU32(Hy)] to interact with SwrA at P_{A(fla/che)} in vitro suggests that the SS complex is the positive stimulator of fla/che transcription.

To test whether such a trigger is sufficient to relieve DegU32(Hy)-mediated swimming inhibition we sought to analyze motility in a degU32(Hy) swrA- background in which an optimized P_{A(fla/che)} promoter, with an increased similarity to the σ^A consensus sequence, was introduced. To this end we used the dhsA6 (degU_Hy suppressor) single-nucleotide mutation in the -35 box of P_{A(fla/che)} that has been shown to suppress the motility-null phenotype caused by degU32(Hy) [16]. Swarming was not assessed in the original dhsA6 mutant since SwrA had not been identified yet. As shown in Figure 5, both swimming and swarming were fully recovered in the strain carrying the dhsA6-optimized P_{A(fla/che)} promoter, although the constitutive degU32(Hy) mutation was still active and the functional swrA allele was absent. This result confirms that the enhancement of fla/che transcription is sufficient for swarming.
The role of PD3(fla/che)

After having investigated the mechanisms which promote fla/che transcription from PA(flac/che), the role of the secondary σD-dependent fla/che promoter was evaluated. Although a feedback effect of PD3(flac/che) in maintaining fla/che expression is postulated (Figure 1B), previous studies have failed to detect it [3]. We have already shown the existence of a powerful SwrA-dependent autoregulatory circuitry (Figure 1A) [8]. We reasoned that the existence of such an efficient loop might mask the effect of PD3(flac/che) and would therefore account for the lack of evidence reported in the past.

To understand the role of PD3(flac/che), we used strains in which the SwrA-based loop was impaired, either through a mutation in the σD-dependent SwrA promoter PswrA in front of the swrA+ allele to interrupt the circuitry at the level of swrA transcription (Figure 1A), or through the frameshifting mutation in swrA that is found in laboratory strains [5]. The effect on swimming motility of the PswrA+ mutation is mild because it is partially relieved by the presence of the PswrA promoter; in fact, this strain is swimming proficient because the PswrA promoter becomes fully active on solid surfaces [8]. Conversely, the frameshifting mutation causes a more severe phenotype in swimming and prevents swarming [5]. A markerless 118-nucleotide deletion of PD3(flac/che) (ΔPD3 in Figure 2A) was created in the two strains as well as in a swrA+ control. We decided to analyzed swimming in order to measure motility performances even in the swrA- backgrounds in which swarming is precluded. As shown in Figure 6A, swimming was not affected by the ΔPD3 mutation in the control swrA+ strain. However, deletion of PD3 causes a progressively more severe phenotype in the swrA+ backgrounds, exacerbating the effect of mutations in swrA. A time course of swimming expansion allowed a better appreciation of the increasingly greater differences between each ΔPD3 strain and its PD3+ wild-type counterpart (Figure 6B). A parallel decrease in the amount of flagellin produced by these strains was also observed during growth in shaking cultures (Figure 6C).

These data demonstrate that PD3(flac/che) is indeed involved in positive feedback regulation of fla/che transcription, thereby increasing flagellin production and swimming speed, but its...
The effect can be appreciated only in the absence of SwrA. The severe motility phenotype of our swrA-ΔP₃D₃ mutant strongly corroborates the importance of P₃D₃ in motility autoregulation. Thus, P₃D₃(fla/che) is redundant in swrA+ strains but it ensures basal motility in case the swrA allele switches to the non-functional status.

We surmise that the discrepancy between our results and those of previous investigations [3] lies in the fact that those experiments were performed in swrA- laboratory strains containing spontaneous swrA+ revertants; thus, the heterogeneous swrA background would have masked the contribution of P₃D₃. Indeed, several swrA+ revertants arose during motility evaluation of swrA strains due to the very high frequency of forward and reverse mutations that occur in the “slippery” polyadenine tract of swrA [5], and scrupulous analysis of the status of the swrA allele was performed immediately before and after each experiment (see also 30).

Discussion

Results contributing to a clearer picture of the regulatory signals mediating fla/che transcription have been presented. We have demonstrated that SwrA interacts with phosphorylated DegU and forms a complex on the fla/che promoter. This complex presumably represents the SwrA-mediated fla/che positive trigger that is mandatory for activating swarming motility. Thus, although flagella are assembled and support basal swimming even the absence of SwrA and DegU [8, 13], phosphorylation of DegU is necessary for swarming motility as earlier demonstrated [11, 12, 14], its phosphorylation being necessary for the interaction with SwrA, although it is repressive in the absence of SwrA as previously observed [13, 15, 16]. The molecular function of SwrA is to act as a modifier of DegU-P, converting it from a repressor to a booster of fla/che expression. Interestingly, the availability of SwrA is modulated through a phase variation ON/OFF mechanism, thereby ensuring diversification of motility performances upon expansion of a clonal population.

Our findings define a situation in which DegU is a dual regulator: upon phosphorylation, which possibly occurs through inhibition of flagella rotation [31], it can act as a repressor or a stimulator of fla/che transcription; the decision is made through an association with the regulator SwrA. Therefore SwrA, together with DegQ and DegR which modulate the phosphorylation/dephosphorylation rate of DegU, represent distinct auxiliary factors that intervene in the fine tuning of the DegS/DegU response, possibly merging different signals directed to this crucial TCS. The action of auxiliary factors is not unusual in TCS signaling pathways, as recently reviewed [32].
Presumably, SwrA also has a more general effect in addition to its specific effect on P_{A(fla/che)} illustrated here, and might contribute to modulating the expression of other genes belonging to the DegU–P regulon. Indeed, besides ycdA [24], preliminary data indicate that in a wild-type degS/degU background the production of extracellular proteases, that is normally triggered by “Hy” mutations [13,17,19], can be modulated by SwrA (Figure S3). Unfortunately, most of the studies on the DegU regulon have been carried out using the degU32(Hy) allele which we have demonstrated to be a mutant protein impaired in its interaction with SwrA; thus, the effects of SwrA on DegU-regulated genes have probably been missed. It would be worthwhile analyzing DegU–P-dependent processes in swrA backgrounds by using the degS200(Hy) mutant allele.

We have also shown that P_{D3(fla/che)} allows the establishment of a positive autoregulatory loop that is used in maintaining fla/che transcription when SwrA is absent (Figure 1B). The importance of this feedback loop is that it could contribute to explain the bistable switch that governs motility in B. subtilis. In fact, motility is virtually null in the swrAΔP_{D3} (Figure 6); the few residual motile cells could be due to the limited amount of functional SwrA produced through transcriptional slippage, as recently proposed by Gordon and collaborators [33]. These authors have elegantly shown that in heritable ON/OFF epigenetic switches whose regulator is inactivated by a slippery A_R tract in its coding sequence (such as swrA) transcription errors can generate wild-type mRNAs, and thereby enough active regulator, to promote epigenetic ON-switching. Further work is required to gather data to verify this hypothesis.

Finally, it is remarkable that although a single nucleotide change might have turned P_{D3(fla/che)} into a self-sufficient promoter, this promoter has been maintained as intrinsically suboptimal: the possibility of regulating energetically-demanding flagellar motility over a wider dynamic range probably has an essential adaptive value.

Supporting Information

Figure S1. The S and SS complexes assemble also on a shorter probe K. EMSA were performed as described in Material and Methods with FAM-labeled probes. The final concentration (μM) of the proteins present in each reaction is indicated on top of the figure. In the first seven lanes reactions were performed with the standard P_{A(fla/che)} probe used in Figures 2 and 4 of the manuscript; in the following seven lanes the shorter 156bp-probe K was used (depicted in Figure 2A). Probe K was amplified with FAM-labeled primer pair 8601 and 8756 (Table S3).

Table S1. Strains used in this study.

Table S2. Plasmids used in this study.

Table S3. Primers used in this study.

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Author Contributions

Conceived and designed the experiments: CC AG. Performed the experiments: CC S. Mordini CO S. Marini FS. Analyzed the data: CC AG. Contributed reagents/materials/analysis tools: RB. Wrote the manuscript: CC.

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