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1 **Subtilisin A production is influenced by surfactin levels in *Bacillus subtilis***

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9 **Abstract**

10 Although not essential for their growth, the production of secondary metabolites increases
11 the fitness of the producing microorganisms in their natural habitat by enhancing
12 establishment, competition and nutrient acquisition. The Gram-positive soil-dwelling
13 bacterium, *Bacillus subtilis* produces a variety of secondary metabolites. Here, we
14 investigated the regulatory relationship between the non-ribosomal peptide surfactin and
15 the lactopeptide bacteriocin subtilisin A. We discovered that *B. subtilis* mutants lacking
16 surfactin production exhibited higher production of subtilisin A compared to their parental
17 wild-type strain. Additionally, spatial visualization of *B. subtilis* production of metabolites
18 demonstrated that surfactin secreted by a wild-type colony could suppress subtilisin A
19 production in an adjacent mutant colony lacking surfactin production. Reporter assays were
20 performed using mutants in specific transcriptional regulators that confirmed the role of
21 ResD as an activator of the subtilisin A encoding BGC, while removal of Rok and AbrB
22 repressors increased expression of the BGC that was further enhanced by additional
23 deletion of surfactin, suggesting that a so far unidentified regulator might mediate the
24 influence of surfactin on production of subtilisin A. Our study reveals a regulatory influence
25 of one secondary metabolite on another, highlighting that the function of secondary
26 metabolites could be more complex than its influence on other organisms and interactions
27 among secondary metabolites could also contribute to their ecological significance.

28 Introduction

29 Biosynthetic gene clusters (BGCs) that are involved in secondary metabolite (SM) production
30 are prevalent across bacterial genera [1,2]. While the production of SMs may not be
31 essential in laboratory settings [3], they likely play a crucial role in the establishment of
32 bacteria within natural niches [4,5]. In the past, the role of SMs in nature has predominately
33 been classified as microbial weapons, likely due to the industrial use of SMs to combat
34 microbial infections [6–8]. However, in recent years, this notion has been adjusted. While
35 the antimicrobial properties of SMs are still acknowledged, more research is being directed
36 toward understanding their ecological function rather than being a direct inhibitor of
37 cellular processes [9–12]. The soil-dwelling, plant growth-promoting bacterium, *Bacillus*
38 *subtilis* harbors a diverse array of BGCs, with surfactin and plipastatin being the most
39 studied non-ribosomal lipopeptides [13–15]. Particularly, surfactin has a strong
40 biosurfactant activity in addition to its antimicrobial properties [16–18]. Surfactin facilitates
41 *B. subtilis* motility through swarming and sliding, thereby playing an important role in *B.*
42 *subtilis* root colonization in soil [19–21]. In addition to non-ribosomally synthesized peptides
43 (NRPs), *B. subtilis* also produces ribosomally synthesized and post-translationally modified
44 peptides (RiPPs). One of the *B. subtilis* specific RiPPs, the bacteriocin subtilosin A was first
45 isolated in 1985 [22] and it displays antibacterial activity towards both Gram-positive and
46 Gram-negative bacteria [22–24]. Other functions of subtilosin A have been reported such as
47 suppression of biofilm formation in *Listeria monocytogenes*, *Gardnerella vaginalis*, and
48 *Escherichia coli* [25]. Furthermore, Schoenborn et al. found delayed sporulation in a mutant
49 lacking subtilosin A compared to its parental wild type strain [14].

50 Whereas surfactin production has been extensively studied across a plethora of *B. subtilis*
51 isolates, research on the production of subtilosin A has predominantly concentrated on the
52 domesticated *B. subtilis* 168 strain or its derivative JH642 [22,23,26]. Domesticated *B.*
53 *subtilis* strains lack surfactin production due to mutation in *sfp* gene [27]. Importantly,
54 natural isolates of *B. subtilis* encode the intact BGC for subtilosin A production (BGC^{Sbo}) [13]
55 and the presence of this BGC is fully conserved among all isolates of *B. subtilis* [28],
56 nevertheless, the production of subtilosin A has not been reported in undomesticated
57 strains.

58 The BGC^{Sbo}, the *sbo-alb* operon encodes the proteins SboA, SboX, and AlbA-AlbG proteins
59 involved in post-translational modifications, processing, and export of the peptide,
60 respectively [23]. The BGC^{Sbo} is regulated by several transcription factors, including Rok,
61 AbrB, and ResD, in addition to the sigma factor SigA. Rok and AbrB repress, while ResD
62 activates the expression of the BGC^{Sbo} [29]. Production of subtilisin A is linked to later
63 growth stages, characterized by nutrient starvation and oxygen limitation [14,30]. Nakano
64 and colleagues demonstrated that the two-component response regulator, ResDE is
65 essential for activating the subtilisin A BGC in response to oxygen limitation [31].
66 Several starvation or stationary phase-specific genes are repressed during exponential
67 growth by AbrB, which directly binds to the respective promoters of those genes, as
68 demonstrated for the *sboA* gene. AbrB-mediated repression is alleviated by Spo0A during
69 starvation [32]. Additionally, AbrB also represses the transcription of *rok* [33,34]. Similarly,
70 Rok binds directly to the promoter of *sboA* and represses its expression [34]. While no
71 specific signal or environmental condition has been correlated with the activity of Rok, it is
72 noteworthy that sRok, an interaction partner of Rok, exhibits altered binding affinity during
73 salt stress [35]. sRok and DnaA, another interaction partner of Rok, affect the binding
74 affinity of Rok, which may affect Rok's regulatory role [36]. Moreover, Rok regulates several
75 genes [34,37], including *sboA*, as well as the biofilm gene *bslA* (*yuaB*) in *B. subtilis* [38].
76 While these studies provide detailed molecular insights into the transcriptional regulation of
77 BGC^{Sbo}, the regulation and production of subtilisin A have only been explored in *B. subtilis*
78 168 and its derivatives that lack surfactin production, excluding any insights into potential
79 co-dependencies or conflicting expression related to subtilisin A and surfactin production.
80 In this study, we demonstrate that while the two SMs, surfactin and subtilisin A, are not
81 produced simultaneously, the presence of surfactin regulates the production of subtilisin A
82 in *B. subtilis*. Additionally, we investigate the regulatory mechanism by which surfactin
83 suppresses the expression of the BGC^{Sbo} using knockout mutants in gene encoding
84 transcriptional regulators. Employing GFP reporter assays, analytical chemistry and spatial
85 detection of SMs, we demonstrate that extracellular surfactin inhibits the production of
86 subtilisin A in mutants that otherwise lack surfactin production.

Results

Production of subtilisin A is increased in mutant strains lacking surfactin

We previously investigated SM production in 12 natural isolates of *B. subtilis* and tested mutants unable to produce specific NRP [13]. Analysis of SM production in one of these isolates, P8_B1 and its derivative NRP-related BGC mutants using liquid chromatography–mass spectrometry (LC-MS) revealed a varying presence of subtilisin A between P8_B1 and mutants (Fig. 1A). The chemical extractions originating from the mutant derivatives lacking surfactin production (Δ *srfAC* and Δ *sfp*) displayed an additional LC-MS peak corresponding to a *m/z* of 1134.1963 ($[M+3H]^{3+}$) that was identified as subtilisin A [39]. The same peak is observed in the LC-MS profiles of other isolates that corresponds to mutants lacking surfactin production [13]. We previously validated that the corresponding LC-MS peaks corresponds to surfactin [13] and highlighted with the green box in Fig. 1A. Indeed, LC-MS peaks corresponding to surfactin are absent in the Δ *srfAC* and Δ *sfp* strains. To confirm this observation in the most frequently used undomesticated *B. subtilis* strain (DK1042 the naturally competent derivative of NCIB3610), samples were extracted from strains DK1042 and Δ *srfAC* to quantify the level of subtilisin A using the peak area. This approach showed an 8.7-fold increase between the peak area in the mutant strain compared to the wild-type DK1042 (*P* = 0.0193, *t*-student) (Fig. 1B).

Surfactin attenuates the expression of BGC^{Sbo}

To determine whether the lack of subtilisin A in LC-MS samples from surfactin producers, was due to differentiated production or degradation of subtilisin A, we tested the expression of BGC^{Sbo} in the wild type and mutant derivatives using the GFP signal normalized by OD_{600nm} as proxy. For this, the promoter region of *sboA* gene of the BGC^{Sbo} was inserted before the promoter-less *gfp* gene and the construct was introduced into the *amyE* locus of the wild type and mutant strains (see Materials and methods). Green fluorescence was followed in plate reader assays (see Materials and methods. Here, the expression of the BGC^{Sbo} was increased in both the Δ *srfAC* and the Δ *sfp* strains compared to the wild type (*p* < 0.0001, *p* < 0.0001, ANOVA and Tukey HSD) (Fig. 2A). To evaluate whether the influence of the lack of surfactin production can be extracellularly complemented,

commercially available purified surfactin was supplemented to the $\Delta srfAC$ strain in varying concentration, showing a reduction in expression of the BGC^{Sbo} with increasing concentration of surfactin (Fig. 2B). Externally added surfactin of $400 \mu g \cdot ml^{-1}$ almost reduced the BGC^{Sbo} expression level in the $\Delta srfAC$ to the levels observed in the wild type ($p = 0.8196$, ANOVA and Tukey HSD).

Complementation of diminished surfactin production in the $\Delta srfAC$ mutant colony by a neighboring wild-type colony

As external complementation with surfactin can reduce the BGC^{Sbo} expression in the $\Delta srfAC$ strain similar to the levels seen in the wild type, we investigated whether surfactin production by a wild-type colony could downregulate the expression of BGC^{Sbo} in a neighboring $\Delta srfAC$ colony. Wild type and $\Delta srfAC$ strains were spotted next to each other on potato dextrose agar (PDA) medium and sampled for visual detection of SMs. Spatial mapping of metabolites allowed visualization of surfactin production and secretion into the agar by the wild-type strain reaching the proximal edge of the $\Delta srfAC$ colony. Subtilosin A was detected in a reverse distribution, with high abundance on the distal part of the $\Delta srfAC$ colony (zone I) with a gradual decrease towards the wild type neighboring edge of the colony (zone II) (Fig. 3AB). Samples were harvested in a line crossing the middle of both colonies (I – IV) and subjected to semi-quantitative LC-MS analysis that verified the diffusion of surfactin from the wild-type strain in its environment, in addition to gradually decreasing subtilosin A levels in the $\Delta srfAC$ colony at increasing surfactin concentrations (Fig. 3C). Additionally, analysis of the spatial metabolite distribution also revealed that production of the sporulation killing factor (SKF) was absent in the $\Delta srfAC$ colony while abundant in the wild-type strain (Fig S1).

Influence of lack of surfactin production on regulation of BGC^{Sbo} by known global regulators

To evaluate whether surfactin downregulates BGC^{Sbo} expression through one of the known global regulators of BGC^{Sbo} , *resD*, *rok*, and *abrB* genes (Fig. 4) were disrupted in the wild type and $\Delta srfAC$ carrying the $P_{sboA}-gfp$. Deletion of *resD* prevented the expression of *sboA* (Fig. 5A), whereas introduction of Δrok and $\Delta abrB$ increased expression of BGC^{Sbo} ($p = <0.0001$

and $p = <0.0001$, ANOVA and Tukey HSD) (Fig. 5BC). The combination of $\Delta srfAC$ with $\Delta resD$ did not influence the already diminished expression of BGC^{Sbo} (Fig. 5A). In contrast, deletion of the BGC for surfactin production in the Δrok further increased the expression level of BGC^{Sbo} ($p = <0.0001$, ANOVA and Tukey HSD) (Fig. 5BC). While in the absence of *rok* gene, this increase was maintained throughout the experiment, BGC^{Sbo} expression in the $\Delta abrB$ $\Delta srfAC$ mutant was only enhanced in the first 20 h, whereas afterward was comparable to the single $\Delta abrB$ strain with no statistical difference ($p = 0.5463$, ANOVA and Tukey HSD) (Fig. 5C).

Discussion

SMs have been extensively investigated and harnessed, playing a pivotal role in combating microbial infections and improving human health [4,5], with a growing interest in the application of SMs beyond medicine [6,7]. In particular, understanding the ecological functions of SMs can enhance the utilization of SM producing bacteria in agricultural applications, where production of SM is important for the efficiency of biocontrol bacteria. Identification of the underlying regulatory mechanisms influencing SM production may facilitate elucidating their role in nature.

Here, we dissected the influence of the lipopeptide surfactin on expression and production of the bacteriocin subtilisin A in *B. subtilis*. Surfactin decreased the production of subtilisin A in *B. subtilis*, while strains lacking surfactin production had increased level of subtilisin A. The lack of surfactin production and therefore enhanced subtilisin A level could be reverted by pure surfactin or inoculating a neighboring wild-type colony next to the $\Delta srfAC$ mutant strain. Testing the expression of BGC^{Sbo} demonstrated a transcriptionally regulatory mechanism behind surfactin-mediated repression of subtilisin A production. Previous studies have demonstrated both overlapping and dissimilar production of SMs in *B. subtilis* [40]. For example, Yannarell et al. reported little overlap of cells expressing both BGCs for surfactin and subtilisin A production in biofilm colonies [40]. Spatial detection of key SMs in *B. subtilis* biofilm colonies has been previously reported using MALDI-MSI [41]. Although not specifically reported, the MALDI-MSI images suggest increased subtilisin A production in

the $\Delta srfAA$ mutant colony confirming our results. Similarly, reduced SKF level was noticeable in the $\Delta srfAA$ mutant used by Si and colleagues [41], which again confirms our data.

The lack of simultaneous production of surfactin and subtilisin A might suggest that their roles in *B. subtilis* are distinctive and these SMs might contribute to different developmental stages or specific environmental conditions. Or rather, that in the absence of surfactin, subtilisin A antibacterial properties are replacing that of surfactins. While experimental validation is required to demonstrate such possibility, various roles of RIPPs have previously been reported, such as growth inhibition, nutrient competition and quorum sensing [42]. Notably, surfactin plays a pivotal role in the early stages of root colonization in soil and during initiation of biofilm formation [19–21], congruent with the early exponential phase expression of the *srfA* operon, around 7 hours after inoculation [43]. On the contrary, production of subtilisin A is correlated with the end of the exponential phase/starting stationary phase [30]. The quantities in which *B. subtilis* produce these SMs is also different. The level of surfactin has been quantified in different *B. subtilis* strains, ranging from 1.25 – 6.45 g·l⁻¹ [44–47], while subtilisin A concentration in different strains and conditions has been reported to be between 0.5 – 7.8 mg·l⁻¹ [22,30]. The production of surfactin and subtilisin A was not measured quantitatively in our study; however, the LC-MS data suggest that surfactin was produced in higher quantities than subtilisin A. The difference in production quantity might be related to their role in the environment, since the function of surfactin as a bio-surfactant may require higher quantities compared to the primarily antibiotic role of subtilisin A.

The gene cluster encoding subtilisin A synthesis is known to be regulated by the global regulators ResD, Rok, and ArbB [29]. Our analysis with *sboA* promoter coupled *gfp* reporter strains confirmed current knowledge on the role of ResD, Rok, and AbrB in the transcriptional regulation of BGC^{Sbo} [29]. Disruption of surfactin production further increased the expression of BGC^{Sbo} on a Δrok background, suggesting that Rok is not involved in perceiving the presence of surfactin. Since ResD works as an activator of BGC^{Sbo} expression, deletion of both *resD* and *srfAC* does not permit the demonstration whether surfactin influences ResD. AbrB functions as a repressor of BGC^{Sbo} transcription, with its repression being relieved during starvation. While disruption of surfactin production in a $\Delta abrB$ background hastened the expression of BGC^{Sbo} in the first 20h compared to the single

212 deletion of *abrB*, the expression levels of BGC^{Sbo} were comparable in the two strains from
213 20h onwards. The enhanced expression of BGC^{Sbo} observed in the earlier phase of the
214 population growth in a Δ *abrB* background, when expression of surfactin related BGC is
215 prominent, suggests that surfactin is not regulating subtilisin A production through *AbrB*.
216 Interestingly, while deletion of either *srfAC*, *sfp*, or *rok* increases BGC^{Sbo} expression from the
217 first few hours of the population growth, disruption of *abrB* only influences BGC^{Sbo}
218 expression 20 h after inoculation of the culture. These experiments suggest an additional
219 regulatory system might be involved in perceiving the presence of surfactin in *B. subtilis*. It is
220 possible that surfactin influence membrane fluidity that results in downregulation of
221 expression BGC^{Sbo}. This is also suggested by slightly increased BGC^{Sbo} expression in the strain
222 lacking plipastatin, the other lipopeptide produced by *B. subtilis* in addition to surfactin.
223 Indeed, surfactin influences gene expression related to biofilm development of *B. subtilis*
224 [48–50] and influence membranous structures in other microorganisms [51]. Examination of
225 the wild-type and Δ *srfAC* *B. subtilis* transcriptome could potentially reveal which genes and
226 regulatory pathways are primarily influenced by surfactin. This could additionally reveal if
227 the transcription of BGCs other than BGC^{Sbo}, are differentially regulated in the absence of
228 surfactin, in accordance with the decreased SKF level detected in the Δ *srfAC* mutant colony.
229 Identifying possible correlations and differences in the production of SMs in *B. subtilis*, such
230 as that described here between subtilisin A and surfactin, could further increase our
231 understanding of the ecological roles of SMs.

Materials and methods

Bacterial strains and culture media

All strains used in this study, including genomic DNA (gDNA) donors, are listed in Table 1. Overnight starter cultures were grown in lysogeny broth (LB, Carl Roth, Germany; 10 g·l⁻¹ tryptone, 5 g·l⁻¹ yeast extract, and 5 g·l⁻¹ NaCl) medium. If not stated otherwise, experiments were performed in potato dextrose broth (PDB; BD, USA; potato infusion at 4 g·l⁻¹, glucose at 20 g·l⁻¹), supplemented with 1.5 % agar when required.

Generation of mutant *B. subtilis* strains

DK1042 *P_{sboA}-gfp* was obtained with gDNA from the gDNA donor 168 *amyE::P_{sboA}-gfp*. Mutant strains in DK1042 *P_{sboA}-gfp* were obtained by natural competence [57], by transforming gDNA and selecting for antibiotic (AB) resistance on AB containing LB agar medium. gDNA was extracted from the donors using the EURx Bacterial & Yeast Genomic DNA Purification Kit (EURx, Gdansk, Poland), following the manufacturer's instructions. To verify transformation and lack of SM production, overnight grown cultures were directly extracted with acetonitrile using a 1:1 acetonitrile:culture dilution, where after the solution was centrifuged and supernatant transferred to HPLC vials and analyzed by ultrahigh performance liquid chromatography coupled to high-resolution mass spectrometry (UHPLC-HRMS).

Expression assay in *B. subtilis* BGC mutants

The effect of SM production on the expression BGC^{Sbo} was evaluated in plate reader assays. Fluorescence and optical density were detected in cultures grown in 96-well microtiter plates with 200 µl PDB including the reporter strains with a final optical density of 0.01 at 600nm (OD₆₀₀). To test the influence of surfactin on the expression of *sboA*, a similar setup was used, except the *P_{sboA}-gfp ΔsrfAC* strain was supplemented with surfactin at a final concentration of 50, 100, 200 and 400 µg·ml⁻¹. PDB medium without surfactin served as a control. Cultivation was performed in Synergy XHT multi-mode reader (Biotek Instruments, Winooski, VT, US), at 30°C with orbital continuous shaking (3 mm), monitoring the OD₆₀₀ as well as GFP (Ex: 482/20; Em:528/20; Gain: 60) fluorescence every 5 min.

Detection of subtilisin A and surfactin in neighboring colonies of wild-type and Δ srfAC strains

Complementation of surfactin production by the wild-type colony towards the neighboring Δ srfAC mutant was tested on PDA medium. 2 μ l overnight grown bacterial cultures were inoculated on PDA medium using a 2.5 cm distance between the inoculation points of the two strains. The plates were incubated at 37°C for 3 days. To assess the level of surfactin and subtilisin A, four plugs were transferred from the plates distributed from the distal region of the Δ srfAC colony to the distal edge of the wild-type colony (see Fig. 3). 1.5 ml isopropanyl:ethyl acetate (1.3 v/v) with 1% formic acid was added to each plug and sonicated for 60 min before centrifugation (3 min, 13400 rpm). The supernatant was extracted and transferred under N₂ with no heat before resuspension in 250 μ l methanol and centrifugation (3 min, 13400 rpm). Supernatant was transferred to HPLC vials and tested by UHPLC-HRMS.

UHPLC-HRMS was performed on an Agilent Infinity 1290 UHPLC system with a diode array detector. UV-visible spectra were recorded from 190 to 640 nm. Liquid chromatography of 1 μ L extract (or standard solution) was performed using an Agilent Poroshell 120 phenyl-hexyl column (2.1 $\times$ 150 mm, 1.9 μ m) at 40 °C using acetonitrile (ACN) and H₂O, both containing 20 mM formic acid, as mobile phases. Initially, a gradient of 10% ACN/H₂O to 100% acetonitrile over 10 min was employed, followed by isocratic elution of 100% ACN for 2 min. The gradient was returned to 10% ACN/H₂O in 0.1 min, and finally isocratic condition of 10% ACN/H₂O for 2.9 min, at a flow rate of 0.35 ml·min⁻¹. HRMS spectra were acquired in positive ionization mode on an Agilent 6545 QTOF MS equipped with an Agilent Dual Jet Stream electrospray ion source with a drying gas temperature of 250 °C, drying gas flow of 8 l·min⁻¹, sheath gas temperature of 300 °C, and sheath gas flow of 12 l·min⁻¹. Capillary voltage was set to 4000 V and nozzle voltage to 500 V. MS data analysis and processing were performed using Agilent MassHunter Qualitative Analysis B.07.00.

Mass spectrometry imaging of pairwise interactions between Δ srfAC and wild-type colonies

Samples were prepared as described above for quantification of SMs from PDA grown colonies. Interaction zone of the two colonies were excised from agar plates and adhered to

MALDI IntelliSlides (Bruker, Billerica, Massachusetts, USA) using a 2-Way Glue Pen (Kuretake Co., Ltd, Nara-Shi, Japan). Slides were covered by spraying 1.5 ml of 2,5-dihydrobenzoic acid (40 mg·ml⁻¹ in MeOH/H₂O (80:20, v/v, 0.1% trifluoroacetic acid)) and dried prior to MSI acquisition. MALDI-MSI data was acquired using a timsTOF flex (Bruker Daltonik GmbH) mass spectrometer operating in a positive mode with 30 µm raster width and a m/z range of 500–4000. Calibration was performed by using red phosphorus. The settings in the timsControl were as follow: Laser: imaging 30 µm, Power Boost 3.0%, scan range 26 µm in the XY interval, and laser power 70%; Tune: Funnel 1 RF 300 Vpp, Funnel 2 RF 300 Vpp, Multipole RF 300 Vpp, isCID 0 eV, Deflection Delta 70 V, MALDI plate offset 100 V, quadrupole ion energy 5 eV, quadrupole loss mass 100 m/z, collision energy 10 eV, focus pre TOF transfer time 75 µs, pre-pulse storage 8 µs. Data was root mean square normalized and visualized in SCiLS software (Bruker Daltonik GmbH, Bremen, Germany).

Statistics

Data was analyzed and graphically represented using R 4.3.2 and the package ggplot2 [58]. Student's t-test was used to test for statistical differences in experiments with two groups. Statistical significance (α) was set at 0.05. For multiple comparisons (more than two treatments), one-way analysis of variance (ANOVA) and Tukey's honestly significant difference (HSD) were performed. In all the cases, normality and equal variance were assessed using the Shapiro - Wilks and Levene test, respectively.

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Author contributions

Designed research: CD, ÁTK; performed the experiments: CD; analysis of strains: CNLA; performed the chemical detection and analysis: MV, SJ, AJCA; analyzed data: CD; wrote the manuscript: CNLA, ÁTK with corrections by co-authors.

Competing interests

The authors declare no competing interests.

Data availability

All raw data used to generate figures in the study are available from the corresponding author upon request.

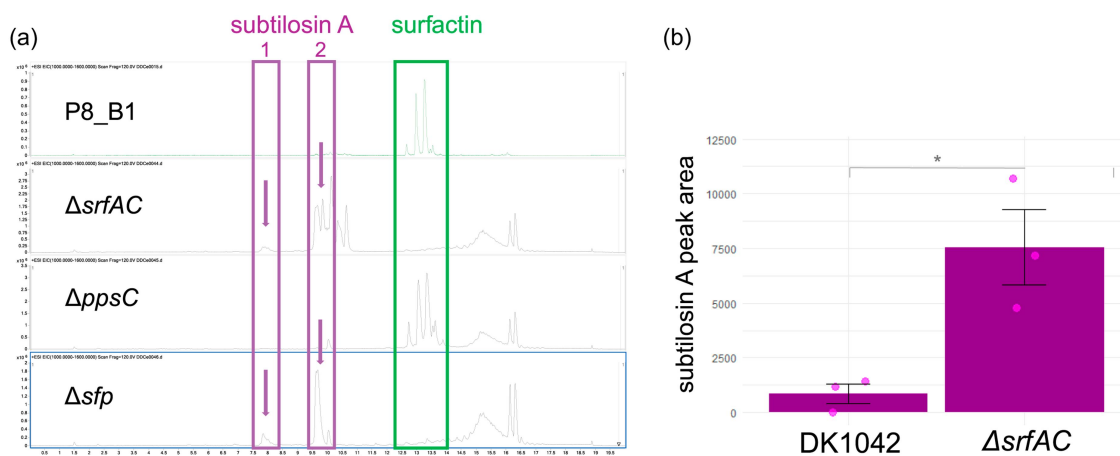
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467 **Fig. 1** (A) LC-MS chromatogram (EIC: m/z 1000-1600) for *B. subtilis* P8_B1 and its derivative
 468 mutants $\Delta srfAC$ (lacking surfactin), $\Delta ppsC$ (lacking plipastatin) and Δsfp (lacking all NRPs).
 469 Subtilisin A's peak (1134.1963 [M+3H]³⁺) is highlighted in the purple boxes. The box
 470 labeled with 1 highlights the hydrolyzed form of subtilisin A, while box 2 refers to the
 471 cyclized subtilisin A product. The isomers of surfactin are highlighted with the green box.
 472 (B) The production of subtilisin A in DK1042 and $\Delta srfAC$ estimated by peak area from EIC
 473 data, statistical difference was tested using students t-test ($p = 0.0193$, t-student, $n=3$).

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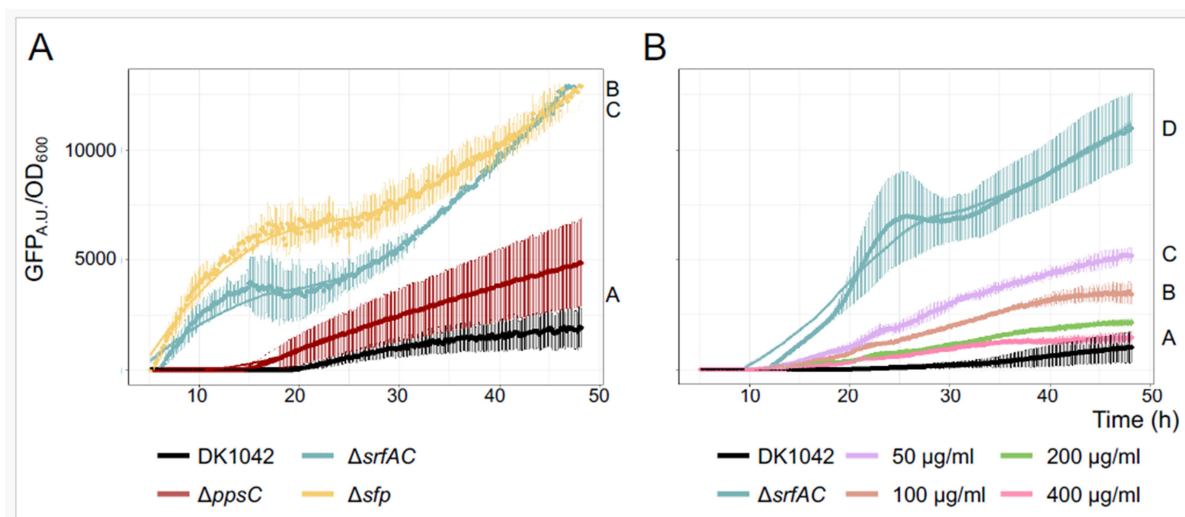


Fig. 2 (A) Expression of BGC^{Sbo} in wild type and derived BGC mutants compared using corresponding strains carrying P_{sboA} -*gfp* reporter fusion. The fluorescence was normalized by growth (optical density at 600nm, OD₆₀₀). (B) Expression of BGC^{Sbo} in $\Delta srfAC$ strain carrying P_{sboA} -*gfp* reporter fusion supplemented with varying concentrations of surfactin (50 to 400 $\mu g \cdot ml^{-1}$). Normalized GFP expression between different strains and treatments was compared using the area under the curve (AUC) using one-way ANOVA and Tukey honest test, letters present significant difference between strains (Table S1).

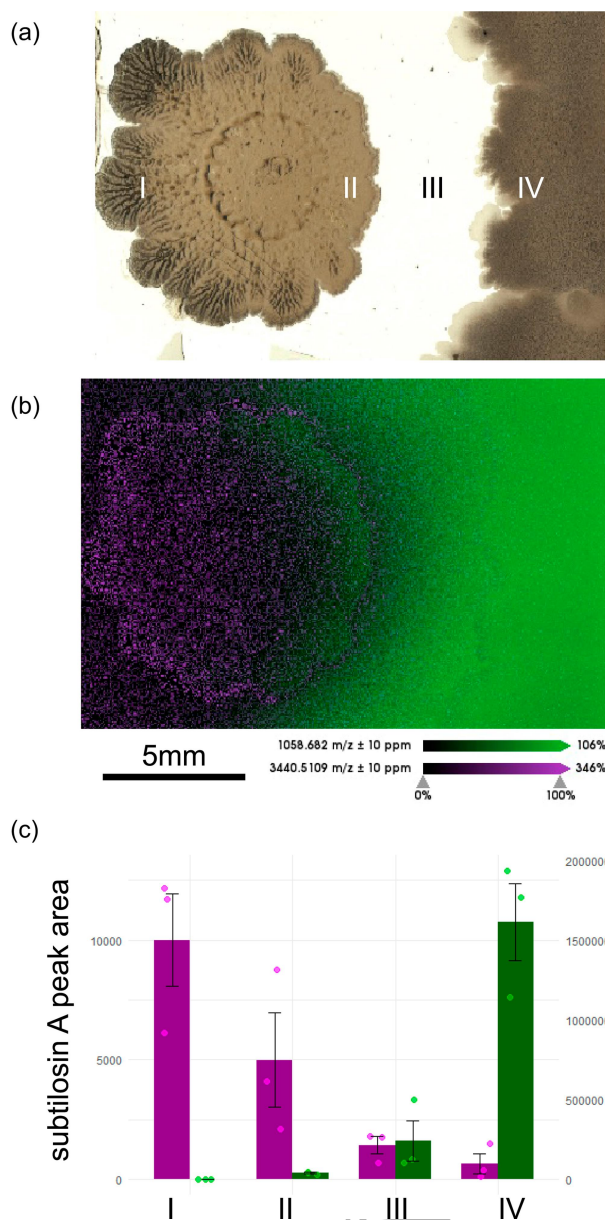
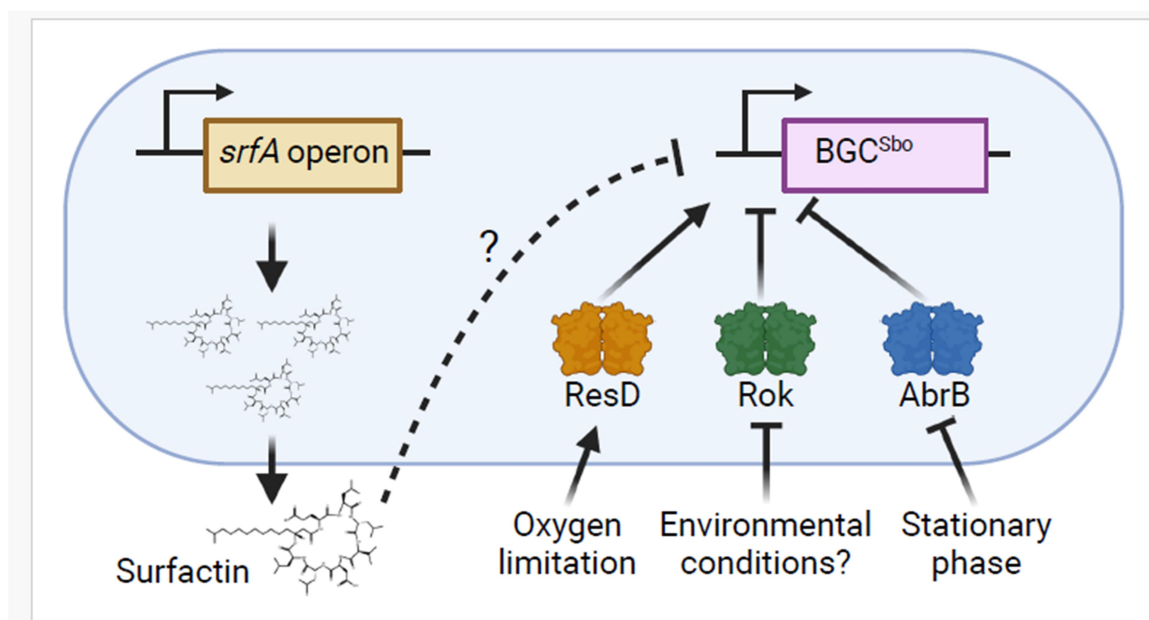


Fig. 3 Spatial mapping of subtilisin A and surfactin distribution in neighboring colonies. (A) Light image of $\Delta srfAC$ (left) and wild type (right) colonies, including the approximate positions of samples taken for LC-MS analysis (I-IV) on a replicate. Scalebar indicates 5 mm. (B) MALDI mass spectrometry imaging-based localization of subtilisin A (magenta) and surfactin (green) in the neighboring colonies of $\Delta srfAC$ and wild-type strains. (C) Relative amount of subtilisin A (magenta) and surfactin (green) estimated by peak area from the LC-MS EIC data in the samples taken at the different positions depicted in panel A.



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497 Fig. 4 Schematic representation of BGC^{Sbo} expression in *B. subtilis*. BGC^{Sbo} expression is
 498 transcriptionally regulated by three known regulators: ResD, Rok, and AbrB that respond to
 499 various signals. The figure also depicts the influence of surfactin on BGC^{Sbo} expression.
 500 Arrows indicate production and activation, while T lines denote repression. Dashed line
 501 refers to either direct or indirect influence.

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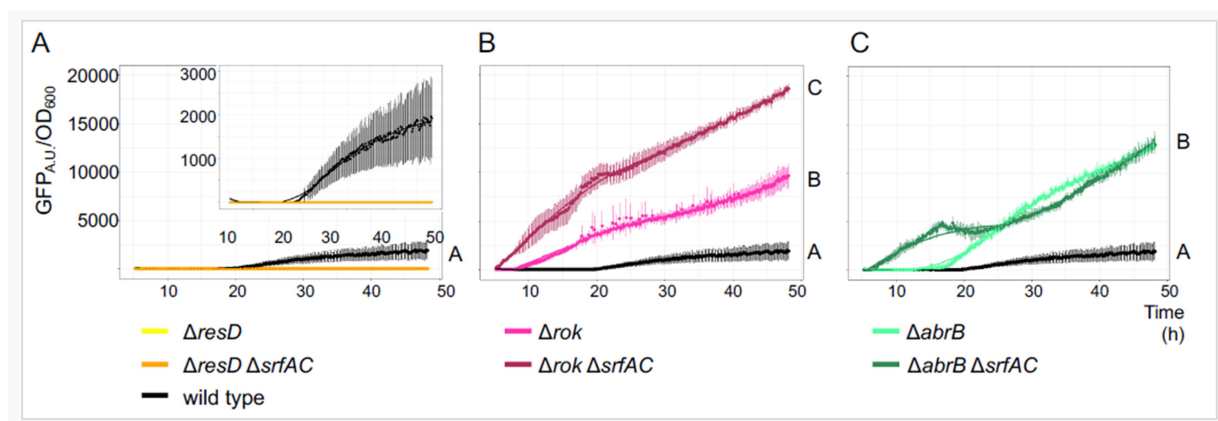


Fig. 5 Expression of BGC^{Sbo} in wild type (black line) and derived regulator mutants compared using corresponding strains carrying *P_{sboA}-gfp* reporter fusion. The fluorescence was normalized by growth (optical density at 600 nm, OD₆₀₀). Expression was assayed in *ΔresD* (A), *Δrok* (B), and *ΔabrB* (C) single mutants (light line colors) or in combination with *ΔsrfAC* (dark line colors). Normalized GFP expression between different strains and treatments was compared using the area under the curve (AUC) using one-way ANOVA and Tukey honest test, letters represent significant difference between strains (Table S2).

513 **Table 1. Detailed information about strains used in this study.**

Strain	Description	Reference
168	<i>amyE::P_{sboA}-gfp</i> (Cm ^R)	[52]
DK1042	NCIB 3610 <i>comI</i> ^{Q12}	[53]
DS1122	3610 Δ <i>srfAC</i> (Mls ^R)	[54]
DS4114	3610 Δ <i>ppsC</i> (Tet ^R)	[55]
DS3337	3610 Δ <i>sfp</i> (Mls ^R)	[56]
P8_B1	WT	[13]
P8_B1	Δ <i>srfAC</i> (Mls ^R)	[13]
P8_B1	Δ <i>ppsC</i> (Tet ^R)	[13]
P8_B1	Δ <i>sfp</i> (Mls ^R)	[13]
DTUB366	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R)	This study
DTUB367	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>srfAC</i> (Mls ^R)	
DTUB368	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>ppsC</i> (Tet ^R)	
DTUB369	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>sfp</i> (Mls ^R)	
DTUB370	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>rok</i> (Km ^R)	
DTUB371	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>resD</i> (Km ^R)	
DTUB372	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>abrB</i> (Km ^R)	
DTUB373	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>rok</i> (Km ^R), Δ <i>srfAC</i> (Mls ^R)	
DTUB374	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>resD</i> (Km ^R), Δ <i>srfAC</i> (Mls ^R)	
DTUB375	DK1042 <i>amyE::P_{sboA}-gfp</i> (Chl ^R); Δ <i>abrB</i> (Km ^R), Δ <i>srfAC</i> (Mls ^R)	

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