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Assembly and Function of the *Bacillus anthracis* S-Layer

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Abstract

Bacillus anthracis, the anthrax agent, is a member of the *Bacillus cereus* sensu lato group, which includes invasive pathogens of mammals or insects as well as nonpathogenic environmental strains. The genes for anthrax pathogenesis are located on two large virulence plasmids. Similar virulence plasmids have been acquired by other *B. cereus* strains and enable the pathogenesis of anthrax-like diseases. Among the virulence factors of *B. anthracis* is the S-layer-associated protein BslA, which endows bacilli with invasive attributes for mammalian hosts. BslA surface display and function are dependent on the bacterial S-layer, whose constituents assemble by binding to the secondary cell wall polysaccharide (SCWP) via S-layer homology (SLH) domains. *B. anthracis* and other pathogenic *B. cereus* isolates harbor genes for the secretion of S-layer proteins, for S-layer assembly, and for synthesis of the SCWP. We review here recent insights into the assembly and function of the S-layer and the SCWP.

Keywords

Bacillus cereus sensu lato; virulence plasmid; *Bacillus* S-layer-associated proteins; BSLs; host invasion; S-layer homology domain; SLH domain; protein secretion; crystallization domain; secondary cell wall polysaccharide; SCWP

ORIGINS OF ANTHRAX AND ANTHRAX-LIKE DISEASE-CAUSING ISOLATES

Bacillus anthracis is an aerobic, spore-forming, gram-positive bacterium (139). Spores are the causative agent of anthrax, an infectious disease that is fatal to mammalian species (68). Herbivores are frequent hosts, ingesting spores that contaminate the environment (140). Spores germinate within the host, and their vegetative forms invade and replicate within all organ tissues, eventually with a lethal outcome (140). Vegetative bacilli sporulate in decaying carcasses so that dormant spores can again contaminate the environment, thereby completing the *B. anthracis* life cycle (68). Phylogenetic studies reveal that *B. anthracis* is a member of the *Bacillus cereus* sensu lato (s.l.) group, which includes the insect pathogen *Bacillus thuringiensis* and the soil microbe *Bacillus cereus* (53). Members of the *B. cereus*

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s.l. group are described as common inhabitants and symbionts of the invertebrate gut (56). Only occasionally do specific isolates of *B. cereus* s.l. enter into a pathogenic life cycle, infecting suitable hosts and multiplying without restraint (56). Multiple-locus sequence typing as well as 16S RNA and whole-genome sequencing of different *B. cereus* s.l. isolates demonstrate their relatedness: a core of 1,750 genes is found in every isolate, and an extended core of another 2,150 genes is present in almost every genome (24, 112, 114, 137, 152). Most members of *B. cereus* s.l. are classified into one of two clades yet carry few clade-specific genes (152). A third clade encompasses outliers including environmental isolates with significant gene decay (152). Three additional species, *Bacillus mycoides*, *Bacillus pseudomycooides*, and *Bacillus weihenstephanensis*, have been added to the *B. cereus* s.l. classification (112, 152). The pathogenic attributes of *B. anthracis*, *B. thuringiensis*, and *B. cereus* isolates are encoded on virulence plasmids, which are acquired via horizontal transfer (55, 66, 67, 115). Extensive genome analysis of *B. anthracis* isolates from many different geographic locations demonstrates the clonal relationship of this species (72). Keim & Wagner (60) proposed that members of the *B. cereus* s.l. group represent “hopeful monsters”: following acquisition of toxin-containing plasmids, successful clones rapidly expand and spread around the globe, aided by the mobility of humans.

VIRULENCE PLASMIDS OF *B. ANTHRACIS*

B. anthracis acquired two large plasmids, pXO1 and pXO2 (107). A key feature of the ~182-kb pXO1 plasmid is the presence of three toxin genes (*pag*, *lef*, *cya*) whose secreted products [protective antigen (PA), lethal factor (LF), and edema factor (EF)] assemble into lethal toxin (LT) and edema toxin (ET) (76, 120, 144) (Figure 1). PA binds to the anthrax toxin receptors, TEM8 (ANTRX1) and CMG2 (ANTRX2), on the surface of mammalian host cells (14, 130). The host protease furin cleaves PA (49), generating mature 63-kDa PA, which then oligomerizes into an octameric structure (65, 97). Oligomerization provides for PA association with *B. anthracis* LF and EF (98). The assembled LT (PA+LF) and ET (PA+EF) are endocytosed by clathrin-coated vesicles; acidification of the endosomes promotes membrane insertion of PA and translocation of LF and EF across the endosomal membrane (73, 151). Once translocated into the cytoplasm of target cells, LF exerts zinc protease activity to cleave mitogen-activated protein (MAP) kinase and NLRP1 (31, 77). EF associates with calmodulin, a calcium-regulated host factor, to acquire adenylate cyclase activity, converting ATP into cyclic AMP messenger molecules (76). Both toxins make important contributions to the life cycle of *B. anthracis*, disabling innate and adaptive immune functions of the host and promoting the lethal outcome of anthrax infections (3, 100). The ~95-kb pXO2 plasmid carries the *capBCADE* operon, which provides for the synthesis of the poly-D- γ -glutamic acid (PDGA) capsule (17, 18, 20, 50). The PDGA capsule endows vegetative bacilli with resistance to opsonization by host complement and opsonophagocytosis by granulocytes and macrophages; PDGA capsulation is an essential virulence determinant of *B. anthracis* (30, 111). Temperature shift to 37°C and increased environmental CO₂, as occur following *B. anthracis* entry into mammalian tissues, induce the expression of toxin and capsule genes (45, 119). The transcriptional response to CO₂ requires the *trans*-acting factor AtxA, whose gene (*atxA*) is located on pXO1 (9, 51, 69, 141, 142) (Figure 1). AtxA-mediated regulation of capsulation requires two additional

transcription factors, AcpA and AcpB, whose structural genes are located on pXO2 (28, 29) (Figure 1). CO₂-induced transcriptional activation also encompasses chromosomal genes, suggesting that AtxA is a global regulator of *B. anthracis* (12). Most *B. cereus* s.l. isolates regulate the expression of their secreted products via PlcR, a global transcriptional regulator and quorum sensor that is activated by binding to the PapR autoinducer (2, 11) (see sidebar titled “Virulence Plasmids of *B. cereus* Isolates Causing Anthrax-Like Disease”). Interference between PlcR and AtxA-mediated gene regulation is thought to have selected for the mutation in *plcR* that inactivates quorum sensing and PlcR-mediated gene regulation in *B. anthracis* (96).

S-LAYER AND S-LAYER PROTEINS OF *B. ANTHRACIS*

Two surface (S)-layer proteins, Sap and EA1, were identified in the extracellular medium and in the bacterial envelope of *B. anthracis* cultures (37, 38) (Figure 2). Expression of their structural genes, *sap* and *eag* (EA1), is regulated by CO₂ (95). In response to increased CO₂, the pXO1-encoded virulence regulator AtxA activates transcription of *eag* via PagR (95), the transcriptional regulator of *pag* expression, in addition to activating the expression of capsule genes (51, 94). As a result, encapsulated bacilli produce predominantly EA1 (94), while germinating bacilli or vegetative forms replicating at low CO₂ produce large amounts of Sap and lesser quantities of EA1 (64). When analyzed by electron microscopy, *B. anthracis* variants lacking *sap* or *eag* each form two-dimensional crystalline arrays (23). Replicating at low CO₂, *B. anthracis* forms Sap S-layers mid-cell and along the cylindrical axis of the cell, whereas EA1 S-layers are deposited at cell septa and near future division sites (64). PDGA polymers are anchored to the peptidoglycan of *B. anthracis* (118); electron microscopy studies revealed that PDGA polymers traverse the S-layer of bacilli (91) (see sidebar titled “Bacterial S-Layers”).

Sequence analysis of *sap* and *eag* indicates that both gene products are synthesized as precursors (Sap 814 residues, EA1 862 residues) with a cleavable N-terminal signal peptide followed by three tandem repeats of ~55-residue S-layer homology (SLH) domains and a C-terminal crystallization domain (93). The 29-residue signal peptides differ by a single amino acid residue and are substrates for SecA2 (see below), whereas SLH domains display 66% identity, which is higher than the sequence identity between SLH domains of *B. anthracis* S-layer-associated proteins (BSLs) (63). However, the crystallization domains of Sap and EA1 contain only 22% amino acid identity (93). As reported for SLH domains in other bacteria, three tandem copies of the SLH domains of Sap and EA1 mediate attachment of fusion proteins to the envelope of *B. anthracis* (92). X-ray crystallography experiments show that SLH domains of Sap fold into a three-pronged spindle structure, generating binding sites for the secondary cell wall polysaccharide (SCWP) of *B. anthracis* (63). Binding of the SLH domains of Sap, EA1 and BSLs to the envelope of *B. anthracis* and *B. cereus* G9241 is dependent on ketal-pyruvylation of the terminal ManNAc residue at the nonreducing end of the SCWP (43, 61, 89, 147). Ketal-pyruvylation is thought to be catalyzed by the product of the *csaB* gene, which is located immediately adjacent to *sap* and *eag* on the chromosome of *B. anthracis* and of *B. cereus* isolates causing anthrax-like disease (89, 147) (Figure 2).

SECRETION AND ASSEMBLY OF S-LAYER PROTEINS

B. anthracis grows as chains of incompletely separated, rod-shaped cells, a trait contributing to its escape from phagocytic clearance (122). Mutations that abrogate expression of *sap* cause *B. anthracis* to form elongated chains owing to the mislocalization of the murein hydrolase BslO (64). Screening libraries of mutant bacilli for variants with increased chain length led to the identification of three genes: *secA2*, *slaP*, and *slaQ* (102, 103). These genes are located immediately adjacent to the S-layer gene cluster (*csaA-csaB-sap-eag*) (103) (Figure 2). Mutations in all three genes diminish the abundance of Sap and EA1 in the bacterial S-layer but do not affect the secretion of other products (102). Mutations in *secA2* and *slaP*, but not in *slaQ*, diminish the secretion of mCherry hybrids fused to the N-terminal signal peptide and SLH domains of Sap (103). In contrast, mutations in *slaQ* affect only the secretion of mCherry hybrids fused to full-length Sap including the C-terminal crystallization domain (103). SecA2 and SlaP are thought to recognize the signal peptides and SLH domains of both Sap and EA1, respectively, whereas SlaQ acts on the crystallization domain of Sap (103) (Figure 2).

SecA, the canonical secretion ATPase of eubacteria, has been studied extensively in *Escherichia coli* (109, 138). *E. coli* SecA binds signal peptide-bearing precursors for delivery and translocation at the SecYEG translocon, which leads to precursor movement across the plasma membrane (34, 35, 78). Following signal peptide cleavage by signal peptidase, mature polypeptides are released on the *trans* side of the membrane (25). Auxiliary translocation factors (SecDF, YajC, and YidC) have been shown to increase membrane translocation of specific precursor proteins (32, 123, 129). The genome of *B. anthracis* harbors genes for the canonical SecA secretion ATPase and for components of the translocation machinery (116). Similar to other gram-positive bacterial pathogens, *B. anthracis* encodes an accessory secretion gene, designated *secA2* (15, 126). SecA2, a homolog of the canonical bacterial secretion ATPase SecA, is thought to recognize the signal peptide of Sap and EA1 and facilitate secretion of large amounts of S-layer proteins during vegetative growth (102). SlaP, a peripheral membrane protein of *B. anthracis*, also facilitates secretion of Sap and EA1 (102). Although this has not been tested experimentally, SlaP could play a role in the membrane translocation of SLH domains to prevent their premature association with SCWP precursors in the cytoplasm. Purified SlaQ assembles into high molecular weight structures and promotes Sap S-layer assembly in vitro (103). The *secA2*, *slaP* and *slaQ* genes are conserved in *B. cereus* s.l. isolates expressing S-layer proteins but are absent from isolates lacking S-layer proteins (SLPs) (103) (Figure 2).

VIRULENCE PLASMIDS ENCODE S-LAYER-ASSOCIATED PROTEINS

Vegetative forms of *B. anthracis*, when grown in the presence of increased CO₂, display adherence to mammalian cells (62). This attribute is associated with the expression of *bslA*, a gene that is located on pXO1 (102) (Figure 1). The predicted 652-residue precursor product of *bslA* encompasses an N-terminal cleavable signal peptide, three SLH domains, and a C-terminal domain that, when purified as a recombinant protein, adheres to mammalian cells and functions as a competitive inhibitor for the binding of bacilli (102). BslA is deposited into the *B. anthracis* S-layer and displayed on the bacterial surface (102).

B. anthracis mutants lacking *bslA* display a defect in adherence to mammalian cells. When analyzed in a guinea pig model of subcutaneous spore challenge and compared to wild-type *B. anthracis* Ames, the *bslA* mutants show delayed replication and dissemination into host tissues and an increase in the spore dose required for lethal disease (102). In a mouse model for disseminated anthrax meningitis, *bslA* mutant bacilli were defective in crossing the blood-brain barrier and in establishing anthrax (33). Thus, BslA represents an adhesin and virulence factor for *B. anthracis*. Biochemical experiments suggest that purified BslA binds to laminin, an extracellular matrix protein found in basal membranes (145). *bslA* homologs are also present on the pXO1-like plasmids of *B. cereus* G9241 and *B. cereus* biovar anthracis CI (147) (Figure 1). pXO1 and pXO1-like plasmids carry another gene for a secreted precursor with N-terminal SLH domains, designated *bslB* (147). The gene is absolutely conserved among *B. anthracis* and *B. cereus* isolates. However, a specific function has not yet been assigned to the *bslB* product.

pXO2 carries the *amiA* gene (pXO2–42), whose product is a secreted precursor of 503 amino acid residues with an N-terminal signal peptide, three SLH domains, and a C-terminal murein hydrolase domain (*N*-acetylmuramoyl-L-alanine amidase) (90) (Figure 1). When purified as a recombinant protein without the N-terminal signal peptide and SLH domains, the C-terminal amidase domain of AmiA cleaves *B. anthracis* peptidoglycan (90). The *in vivo* contributions of *amiA* to the life cycle of *B. anthracis* have, however, not yet been studied. *amiA* is conserved in the pXO2-like plasmids of *B. cereus* biovar anthracis CI but is absent from pBC210 of *B. cereus* G9241 (66) (Figure 1). The capsule plasmid (pBC210) of *B. cereus* G9241 encodes three genes for precursors bearing signal peptides and three SLH domains (54, 147). One of these, *bslO2*, is a predicted murein hydrolase with a C-terminal *N*-acetylglucosaminidase domain (147). Thus, genes for S-layer-associated murein hydrolases are a conserved feature of virulence plasmids with capsule genes, suggesting that these enzymes may contribute either to capsule assembly or to controlling the length of chains of capsulating bacterial cells with incompletely separated septal peptidoglycan (see sidebar titled “S-Layer Regulation of Chain Lengths in *B. anthracis*”).

B. ANTHRACIS AND B. CEREUS S-LAYER-ASSOCIATED PROTEINS

In addition to Sap and EA1, the chromosomes of *B. anthracis* and *B. cereus* G9241 encode 19 and 18 genes for precursor proteins with SLH domains, respectively (62, 147). Collectively these gene products have been designated BSLs (*Bacillus* S-layer-associated proteins) (62). Mutations in *bslO* cause mutant bacilli to form chains of vegetative forms with exaggerated lengths, a phenotype that is complemented when purified BslO protein is added *in trans* (5). In wild-type bacilli, the BslO protein is deposited near cell septa (5). BslO localization is dependent on S-layer assembly (64). Further, *patA1* and *patA2* (encoding two acetyltransferases that modify the SCWP) and *wpaA* [whose product belongs to the family of Wzy-/WaaL (also known as O-antigen ligase)-like enzymes] are also required for proper localization of BslO (83, 105) (Figure 2). Mutations in *patA1* and *patA2* affect the deposition of EA1 in the vicinity of cell septa, suggesting that SLP distribution is influenced by chemical modifications to the ligand of SLH domains (83). Computational simulation suggests that BslO effects a nonrandom distribution of *B. anthracis* chain lengths, implying that all septa are not equal candidates for BslO-mediated cell separation (5). BslS

is an *N*-acetylmuramoyl-L-alanine amidase; its structural gene is located downstream of the *wpaA* gene and the S-layer locus (105). Deposition of BslS and BslT, a homolog of BslS, in the S-layer of *B. anthracis* is perturbed in *wpaA* mutant bacilli (105).

The *bslK* gene product carries a NEAT (heme-binding near-iron transporter) domain and binds heme-iron, as reported for other NEAT-domain-containing proteins (88, 136). Hal and IsdX2, secreted hemophores with NEAT domains, scavenge heme-iron from host hemoglobin and transfer the nutrient to BslK (7, 85). Further transport of heme-iron across the bacterial envelope involves other NEAT domain proteins, such as IsdX1 and IsdC, which is anchored by sortase B to the pep-tidoglycan of *B. anthracis* (84, 85). Heme-iron is imported into the bacterial cytoplasm by the IsdE ATP-binding protein and the IsdF membrane transporter (84). A hemoxygenase, IsdG, cleaves the tetrapyrrole of heme, and the released iron is incorporated into iron-containing proteins (117, 132).

STRUCTURE AND FUNCTION OF THE SECONDARY CELL WALL POLYSACCHARIDE

The ~10- to 20-kDa SCWP released by hydrofluoric acid treatment from *B. anthracis* has the repeat structure [\rightarrow 4)- β -ManNAc-(1 \rightarrow 4)- β -GlcNAc(O3- α -Gal)-(1 \rightarrow 6)- α -GlcNAc(O3- α -Gal, O4- β -Gal)-(1 \rightarrow)]₆₋₁₂ (22) (Figure 3*a*). *B. anthracis* CDC684 is an avirulent isolate whose SCWP has the same trisaccharide repeat structure as wild-type *B. anthracis* yet lacks all galactosyl modifications (43, 108). NMR spectroscopy of CDC684 polysaccharide revealed both ketal pyruvyl (blue in Figure 3*a*) and acetyl modifications at the distal (nonreducing) end of the SCWP (43) (red in Figure 3*a*). Of note, only some molecules carry ketal pyruvyl at O4,O6 of ManNAc at the distal end of the SCWP and acetyl at O3 of the penultimate GlcNAc residue (43, 83). *B. cereus* G9241 and other *B. cereus* isolates causing anthrax-like disease synthesize SCWP with similar size and structure as *B. anthracis*, although wall polysaccharide from these isolates carries an additional α -Gal substitution at O3 of ManNAc (44) (Figure 3*b*). The environmental, nonpathogenic isolate *B. cereus* ATCC 10987 synthesizes SCWP with a distinct repeat structure, [\rightarrow 4)- β -ManNAc-(1 \rightarrow 4)- β -GlcNAc-(1 \rightarrow 6)- α -GalNAc-(1 \rightarrow)]_n, and β -Gal substitutions at the O3 of α -GalNAc as well as nonstoichiometrical acetylation at O3 of β -ManNAc (75) (Figure 3*b*). *B. cereus* ATCC 14579 SCWP has the same repeat structure and substitutions of β -GlcNAc at O3 of the β -GlcNAc in addition to β -Glc at O3 and α -ManNAc at O4 of the α -GalNAc residue (19) (Figure 3*b*). The SCWP of *B. cereus* ATCC 14579 is expressed constitutively, while another, structurally dissimilar polysaccharide is synthesized in biofilms but not during planktonic growth (19).

The pyruvylated SCWP of *B. anthracis* and *B. cereus* G9241 serves as the attachment site for the S-layer and its constituents, SLPs and BSLs with three SLH domains (61, 147). However, the SCWP is also a ligand for murein hydrolases of bacteriophages, for example, PlyG and PlyL (81, 127). PlyG, which is encoded by the γ -phage, binds to the SCWP from *B. anthracis* and *B. cereus* G9241, but not to the wall polysaccharides of *B. cereus* ATCC 10987 and ATCC 14597 (46, 81) (Figure 3*b*). Synthetic oligosaccharides variable for the galactosyl modifications of a single SCWP trisaccharide of *B. anthracis* revealed that high-

affinity binding of PlyG and PlyL is dependent on the β -Gal modification at O4 of the α -GlcNAc residue (99). Thus, the SCWP of *B. anthracis* and pathogenic *B. cereus* strains functions as the structural determinant for bacteriophage-mediated lysis of bacteria by serving as ligand for the carbohydrate-binding domain of bacteriophage lysins (42). Furthermore, *B. anthracis* Sap functions as surface receptor for bacteriophage AP50c (110).

ASSEMBLY OF THE *B. ANTHRACIS* SECONDARY CELL WALL POLYSACCHARIDE

Unlike *B. subtilis* and *Staphylococcus aureus*, *B. anthracis* produces SCWP and lipoteichoic acid (LTA) but not wall teichoic acid (WTA) (47, 101). See Figure 4a for a model of SCWP assembly and synthesis reactions and Figure 4b for a summary of operons and genes involved in SCWP synthesis. *B. subtilis* and *S. aureus* synthesize polyribitol-phosphate teichoic acid on murein linkage units (GlcNAc-ManNAc) that are linked to undecaprenyl-pyrophosphate carrier [C₅₅-(PO₄)₂-GlcNAc-ManNAc] (59, 70, 71). Murein linkage unit synthesis requires the products of two genes, *tagO* and *tagA* (74, 134). TagO transfers GlcNAc-1-phosphate from UDP-GlcNAc onto C₅₅-(PO₄)₂ to generate C₅₅-(PO₄)₂-GlcNAc (lipid III) (48, 121) (Figure 4a, Reaction 12); this reaction is inhibited by the antibiotic tunicamycin (52). TagA transfers *N*-acetylmannosamine from UDP-ManNAc onto O4 of GlcNAc within lipid III, releasing C₅₅-(PO₄)₂-GlcNAc-ManNAc (26, 48) (Figure 4a, Reaction 13). *S. aureus tagO* mutants cannot synthesize WTA but also display subtle defects in peptidoglycan synthesis (40, 52, 148). The *B. anthracis* genome contains *tagO* and *tagA* homologs (116). When expressed in *S. aureus tagO* mutants, *B. anthracis tagO* restores WTA synthesis (61). Tunicamycin treatment or genetic repression of *tagO* inhibits vegetative growth, SCWP synthesis, and S-layer assembly of *B. anthracis*, causing expansions in size and spherical shapes of vegetative forms that can no longer divide (61, 82). These data suggest that the SCWP may be synthesized on murein linkage units that are tethered to an undecaprenyl-pyrophosphate carrier (Figure 4a). In agreement with this hypothesis, SCWP synthesis in *B. anthracis* also requires the LytR-CpsA-Psr (LCP) family of enzymes (79, 80) (Figure 4a). In *B. subtilis* and *S. aureus*, LCP enzymes tether teichoic acids to peptidoglycan by forming a phosphodiester bond between O6 of *N*-acetylmuramic acid (MurNAc) in glycan strands and O1 of GlcNAc in murein linkage units (21, 58). As expected, expression of LCPs is essential for *B. anthracis* growth and SCWP synthesis, and expression of some, but not all, *B. anthracis* LCP genes restores WTA synthesis in *S. aureus* mutants lacking endogenous LCPs (79). Genes involved in SCWP synthesis appear to be scattered among four gene clusters: *slc* (S-layer cluster), *sps* (surface polysaccharide), *scwp1*, and *scwp2* (Figures 2 and 4b). Two of these clusters, *sps* (128) and *scwp2*, display variation between *B. anthracis* and various *B. cereus* isolates. We presume that some of these genes are responsible for the galactosylation patterns of *B. anthracis* and *B. cereus* SCWP (Figure 3b). Some of the genes predicted to be involved in SCWP synthesis are present in duplicate; their products often display functional redundancy (105, 106).

Bacteria evolved three biosynthetic pathways for polysaccharide synthesis: synthase, ABC transporter, and Wzy-polymerase-dependent assembly (150). Capsular polysaccharides and cell wall polysaccharides of gram-positive bacteria are frequently synthesized via the Wzy-

polymerase-synthesis pathway. This has been extensively studied for the capsule genes of *Streptococcus pneumoniae* (150), and insights from this work informed our bioinformatic analysis of the *B. anthracis* SCWP synthesis pathway. Spratt and coworkers (10) characterized *S. pneumoniae* genes and putative enzymatic activities of their products involved in Wzy-dependent capsular polysaccharide (CPS) synthesis. *cps* genes are clustered in a single locus of 17–20 genes (10). *S. pneumoniae* evolved more than 90 serological variants with different CPS structures, and 1,999 genes in all of these variants are responsible for the synthesis of various carbohydrates (87). These genes have been classified into homology groups. Bioinformatic schemes have been employed to predict the function of these gene products, identifying proteins within the same homology group, Pfam family, or CAZy glycosyltransferase family. Correlating *cps* gene content with known CPS structures provides tentative assignments of function to the different homology groups: regulatory factors, enzymes for synthesis of polysaccharide constituents, polymerases, flippases, initial sugar transferases, glycosyltransferases, phosphotransferases, acetyltransferases, and pyruvyltransferases (1).

Synthesis of *B. anthracis* SCWP Units

Wzy pathway synthesis begins with the addition of a UDP-linked sugar to undecaprenyl-PO₄ (C₅₅-PO₄) (Figure 4a, Reaction 1). The reaction is catalyzed by the initiating glycosyltransferase WchA (WciI homology group), a member of the polyisoprenyl-phosphate hexose-1-P family (10). In *S. pneumoniae* 9L/9N CPS, the initial sugar is GlcNAc, similar to *B. anthracis* SCWP (10). Bioinformatic searches of *B. anthracis* genome sequences for WchA (WciI, CpsE9N/L) homologs identified the locus tag GBAA_RS27680 upstream of *acrR* in the *scwp2* cluster (Figure 4b). *B. anthracis* C₅₅-(PO₄)₂-GlcNAc-specific glycosyltransferases are expected to extend the product of Reaction 1 first with β-GlcNAc (Reaction 2) and then with β-ManNAc (Reaction 3). Candidate glycosyltransferases for Reactions 2 and 3 are not readily obvious. The identity of the responsible enzymes may be guided from the *S. pneumoniae* CPS database for glycosyltransferases with retaining or inverting stereospecificity (α- or β-linkages) (87) and will need to be confirmed with biochemical tests. Nonetheless, it is likely that these enzymes are encoded within *scwp2* given their genetic association with *wchA*- and *wzx*-like genes in *S. pneumoniae* CPS clusters (87).

UDP-GlcNAc, the substrate for Reactions 1 and 2, is also required for peptidoglycan synthesis and supplied by *N*-acetylglucosamine-1-PO₄ uridylyltransferase (GlmU). SCWP synthesis requires UDP-ManNAc, which is derived from UDP-GlcNAc by UDP-GlcNAc 2-epimerase. GneZ (encoded in *sps*) and GneY (encoded in *scwp1*) have been characterized as the corresponding enzymes in *B. anthracis* (Reaction 7) (128, 146). Galactosyl modifications of the SCWP require *B. anthracis* to synthesize UDP-Gal. The genome of *B. anthracis* encodes two functional UDP-Glc 4-epimerases. One of these genes, *galE1*, is located in the *sps* cluster (Figure 4b). The second gene is located elsewhere on the chromosome and has been shown to be expressed only during sporulation; it functions preferentially as UDP-GlcNAc 4-epimerase to synthesize UDP-GalNAc, a substrate for glycosyltransferases that modify BclA (27). BclA is the exosporium glycoprotein of *B. anthracis* spores (13, 135). *galE1* is expressed during vegetative growth (128) and is the proposed candidate for UDP-

GlcNAc 4-epimerase activity in support of SCWP synthesis. RfaB enzymes (COG0438) function as UDP-Gal-dependent galactosyltransferases (36). The *B. anthracis* genome harbors several genes for RfaB-type enzymes in *sps* and *scwp2* (Figure 4b). These genes are candidate glycosyltransferases for galactosyl transfer to the SCWP trisaccharide repeat (Reactions 4–6).

Translocation and Polymerization of SCWP Units

Translocation of the SCWP precursor across the membrane requires Wzx flippase (RfbX) (Reaction 9); a candidate for the putative SCWP flippase activity is encoded within the *scwp2* cluster (Figure 4b). Wzy polymerase/WaaL (Reaction 10) is responsible for generating SCWP polymers. WpaA (*slc* encoded) and WpaB (*scwp2* encoded) belong to the same clan as members of the Wzy polymerase/WaaL family of proteins. Both genes are involved in the synthesis of the SCWP of *B. anthracis* (105). *B. anthracis* mutants blocking the expression of a single O-antigen ligase-like gene, *wpaA* or *wpaB*, cannot synthesize SCWP and form misshapen cells without S-layers (105). On the other hand, mutants expressing a single gene for Wzy-like/O-antigen ligase-like enzymes display either assembly defects for S-layer-associated murein hydrolases BslS and BslT (*wpaA* mutant) or defects in the length of *B. anthracis* cell chains and in S-layer assembly (*wpaB* mutant) (105). LCP enzymes are thought to transfer SCWP tethered to murein linkage units from C₅₅-(PO₄) to peptidoglycan, thereby generating phosphodiester bonds with the O6 of MurNAc that can be hydrolyzed with hydrofluoric acid and released from the bacterial envelope (61). *B. anthracis* expresses six genes whose products may catalyze this reaction: LcpB1, LcpB2, LcpB3, LcpB4 and LcpC, and LcpD (Reaction 11) (79, 80).

PERSPECTIVE

In addition to the secreted toxins and capsules of *B. anthracis* and pathogenic *B. cereus*, S-layer and S-layer-associated proteins are key virulence factors enabling host invasion and escape from phagocytic clearance during the pathogenesis of anthrax or anthrax-like diseases. However, only some of the S-layer-associated proteins have been studied. Further, the mechanisms whereby S-layer proteins and S-layer-associated proteins are deposited into the bacterial envelope to exert their functions are still poorly understood. The SCWP of *B. anthracis* and pathogenic *B. cereus* assumes strain-specific characteristics that affect the assembly of S-layers and the function of bacteriophage-encoded murein hydrolases. The genes and enzymatic mechanisms of SCWP synthesis and assembly are only beginning to be discovered. Additional insights into this field may explain the unique attributes of pathogenic *B. cereus* isolates, their envelope characteristics, the range of their mammalian hosts, and the tropism of bacteriophages.

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VIRULENCE PLASMIDS OF *B. CEREUS* ISOLATES CAUSING ANTHRAX-LIKE DISEASE

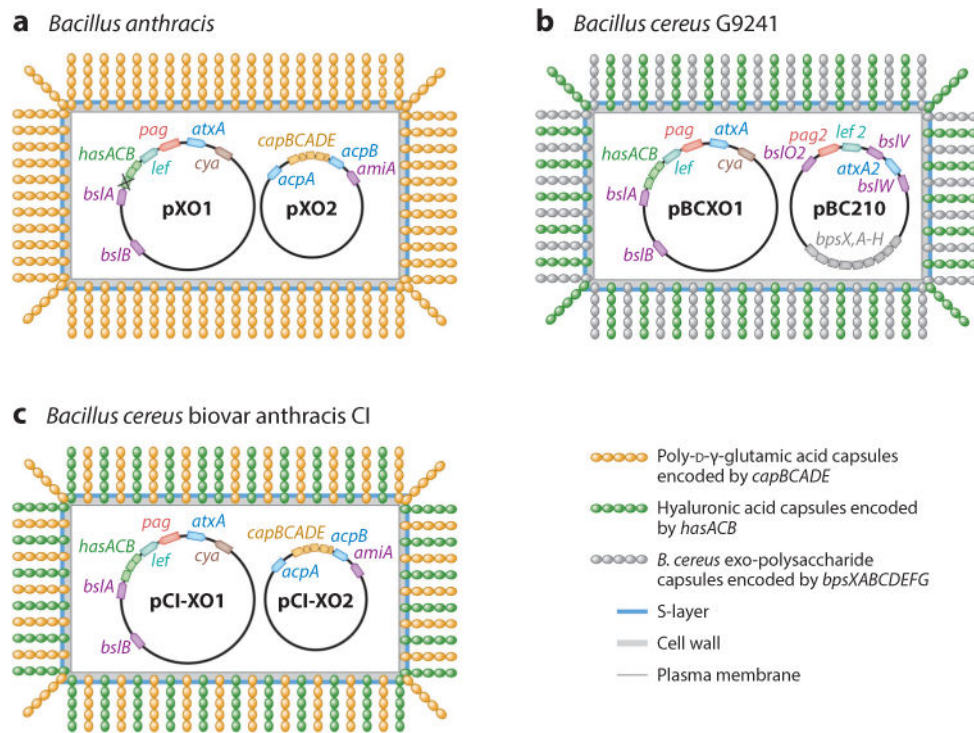
In addition to *B. anthracis*, many other *B. cereus* s.l. isolates, including *B. cereus* biovar anthracis CI and *B. cereus* G9241, have acquired pXO1-like plasmids and with them the ability to secrete anthrax toxins (54, 55, 57, 67, 149) (Figure 1). *B. cereus* G9241 and *B. cereus* biovar anthracis CI, along with *B. anthracis*, belong to clade 1 of *B. cereus* s.l. (152). *B. cereus* biovar anthracis CI was isolated from mammals (gorilla, chimpanzee, elephant, and goat) with anthrax-like disease in West and Central Africa (6, 67). *B. cereus* G9241 and its close relatives were isolated in North America from humans with severe pneumonia or cutaneous anthrax-like disease (55, 86). pXO1-like plasmids of *B. cereus* isolates provide for the expression of hyaluronic acid (HA) capsule via the *hasACB* operon, which is mutated and nonfunctional in *B. anthracis* (16, 104) (Figure 1). *B. cereus* biovar anthracis CI produces both HA and PDGA capsular material, as these strains acquired a pXO2-like plasmid (6, 16, 66). In contrast, *B. cereus* G9241 synthesizes HA in addition to another capsular polysaccharide designated BPS (*B. cereus* exo-polysaccharide), which requires the expression of the *bpsXABCDEFGH* genes located on the pBC210 virulence plasmid (104) (Figure 1). *B. cereus* G9241 and *B. cereus* biovar anthracis CI express *atxA* and carry an inactivating mutation in their chromosomal *plcR* gene that is distinct from that of *B. anthracis* (6, 16, 125). pBC210 carries an *atxA*-like gene (*atxA2*) that contributes to the pathogenesis of anthrax-like disease in *B. cereus* G9241 (125). pBC210 also carries *pag2* (60% identity with *pag*) and *lef2* (36% identity with *lef*) (Figure 1); the *lef2* product shows the presence of a putative PA binding domain; however, the metalloprotease domain of LF is replaced with an ADP-ribosyltransferase domain (41, 143). The *lef2* product, which has been designated Certhrax, is translocated by PA2 into host cells, modifies the host factor vinculin, and disrupts focal adhesion complexes (131, 143). These features of *B. anthracis* and *B. cereus* may account for the observed variation in disease severity and geographic spread associated with spores causing anthrax or anthrax-like disease.

BACTERIAL S-LAYERS

Bacterial S-layers are assembled from proteins that form monomolecular, two-dimensional crystalline arrays on the microbial envelope. S-layer proteins (SLPs) are endowed with a crystallization domain that, when purified and examined in isolation, promotes self-assembly of crystalline arrays (113, 124). A broad range of bacteria and archaea assemble S-layers as their outermost envelope component (4, 39, 133). S-layers form a porous mesh from unit sizes of 3–100 nm and pore sizes of 2–10 nm, thereby functioning as protective barriers with selective permeability (113). The thickness of S-layers varies from 5 to 20 nm in eubacteria but may reach 70 nm in archaea (113). In archaea, SLPs are tethered to the bacterial membrane via a hydrophobic peptide or lipid modification (4). In eubacteria, SLPs are retained on cell surfaces via noncovalent interactions with surface polymers, i.e., lipopolysaccharide (LPS) in gram-negative bacteria and wall polysaccharides in gram-positive bacteria (113). With nanobodies being used to block the formation of two-dimensional crystalline arrays, the structure of the crystallization domain of *Geobacillus stearothermophilus* SbsB was determined via X-ray crystallography (8). The crystallization domain comprises six subdomains with immunoglobulin-like structure (Pfam 02368) folding into a ϕ -shaped, disk-like monomeric unit (8). Interdomain interactions between units are stabilized by calcium ion coordination, which serves as a switch for the formation of a condensed quaternary structure (8). Pores approximately 30 Å in diameter are formed at the interface between three adjacent subunits (8). The crystallization domain of SbsB (residues 202–920) displays 22% amino acid identity with the crystallization domain of *B. anthracis* Sap (residues 211–814), suggesting that Sap may fold into a three-dimensional structure similar to SbsB.

S-LAYER REGULATION OF CHAIN LENGTHS IN *B. ANTHRACIS*

Immediately following germination, *B. anthracis* emerges as a single vegetative cell that begins to elongate and divide, forming chains of 4–16 cells during mid-logarithmic growth (5). As *B. anthracis* vegetative forms approach stationary phase, chain length is decreased (2–4 cells) until eventually short chains and single cells predominate (5). Increased chain length is an impediment to *B. anthracis* host cell invasion and protects bacilli from phagocytosis by macrophages (147). In addition to SLPs, peptidoglycan hydrolases (BslO and BslS) function as determinants of *B. anthracis* chain length. Thus, SLPs and S-layer-associated proteins (BSLs) affect *B. anthracis* invasion of host cells (BslA) and evasion from innate defenses of the mammalian hosts.

**Figure 1.**

B. cereus s.l. isolates causing anthrax and anthrax-like diseases: (a) *B. anthracis* with its two virulence plasmids, pXO1 and pXO2; (b) *B. cereus* G9241, harboring pBCXO1 and pBC210; and (c) *B. cereus* biovar anthracis CI and plasmids pCI-XO1 and pCI-XO2. *B. cereus* G9241 and biovar anthracis CI have a third small plasmid not shown here. In addition to toxin genes (*pag*, *lef*, and *cya*), *B. anthracis* virulence plasmids pXO1 and pXO2 carry genes for poly-D- γ -glutamic acid (PDGA) capsular synthesis (*capBCADE*) and hyaluronic acid (HA) capsular synthesis (*hasACB*); the *hasA* gene, however, carries an inactivating mutation. The plasmids also carry three genes for *B. anthracis* S-layer-associated (BSL) proteins: *bslA*, *bslB*, and *amiA*. Virulence gene expression is regulated by the product of *atxA*, which activates toxin, *bslA*, and *capBCADE* gene expression in response to a host signal (increased CO₂ concentration). The *capBCADE* operon is coregulated by *acpA* and *acpB*. *B. cereus* G9241 synthesizes HA (intact *hasACB* gene cluster). pBC210 encodes genes for three BSLs (*bslO2*, *bslV*, and *bslW*), the Certhrax toxin (*pag2*, *lef2*), an *atxA*-like regulator (*atxA2*), and *B. cereus* exo-polysaccharide (BPS) (*bpsXABCDEFG*). *B. cereus* biovar anthracis capsulation includes both PDGA and HA.

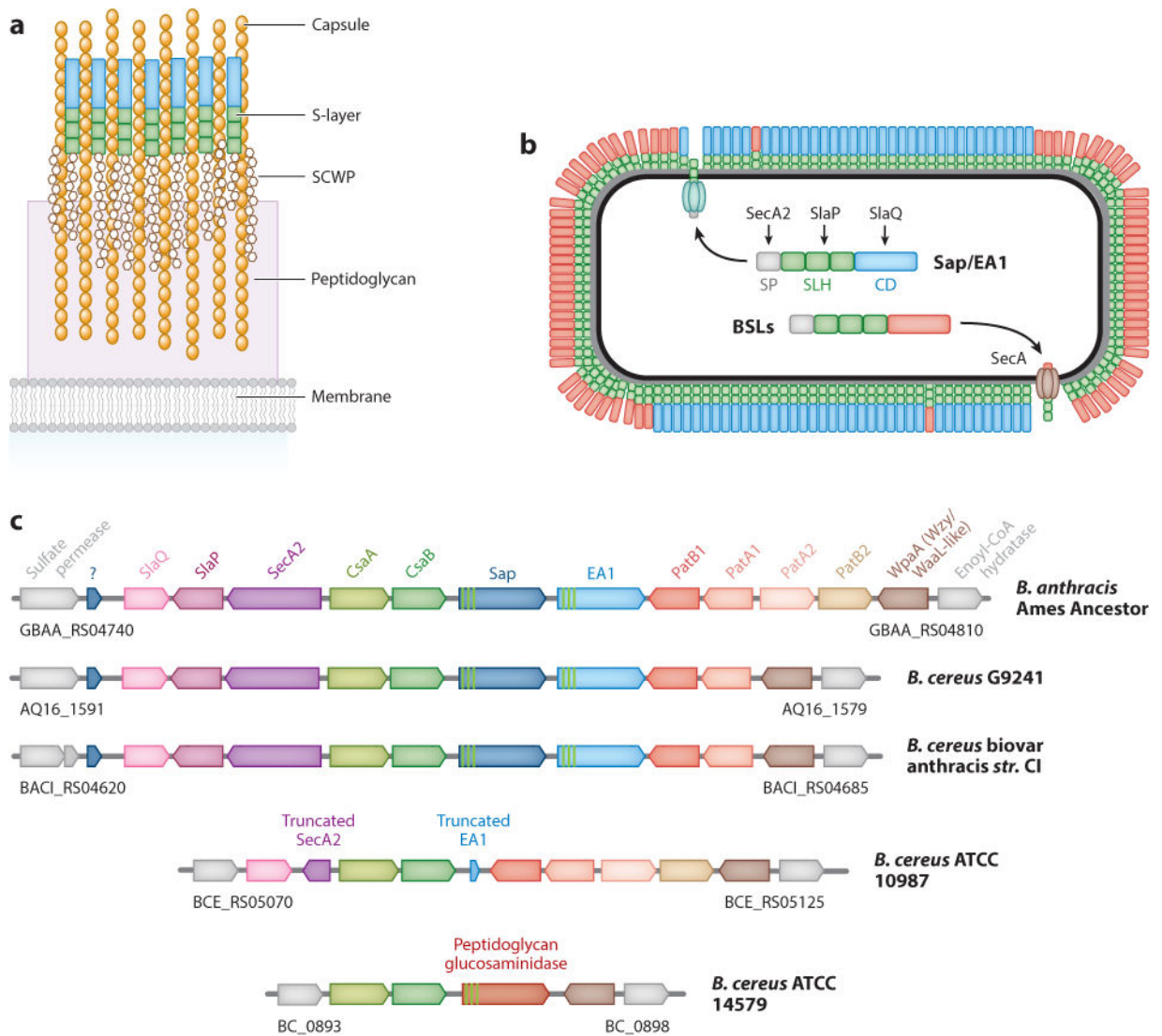


Figure 2.

The S-layer envelope of *B. anthracis*. (a) Model depicting different layers, including the plasma membrane, peptidoglycan, secondary cell wall polysaccharide (SCWP), and S-layer. The SCWP and poly-D- γ -glutamic acid strands (capsule) are tethered to the peptidoglycan of *B. anthracis*. (b) Secretion and assembly of S-layer proteins (SLPs), Sap and EA1, and *B. anthracis* S-layer-associated proteins (BSLs). Each protein is synthesized as a precursor with an N-terminal signal peptide (SP), three S-layer homology (SLH) domains and a C-terminal domain. SecA2 and SlaP contribute to the secretion of Sap and EA1, whereas SlaQ contributes to Sap and EA1 S-layer assembly. In Sap and EA1, the C-terminal domain is a crystallization domain (CD). (c) S-layer gene cluster (*slc*) in *B. anthracis* and other *B. cereus* s.l. isolates. The *slc* is flanked by sulfate permease and enoyl-CoA hydratase genes, shown in gray. The *csaA* and *csaB* (ketal pyruvyltransferase) and *wpaA* [Wzy-/WaaL (O-antigen ligase)-like] genes are absolutely conserved in all species and required for SCWP modifications that enable SLP or BSL assembly. Strains causing anthrax (*B. anthracis*) or anthrax-like disease (*B. cereus* G9241 and *B. cereus* biovar anthracis CI) encode genes for

SLPs (Sap and EA1 with three SLH domains), SLP secretion and assembly (*slaQ*, *slaP*, and *secA2*), and SCWP acetylation (*patA/patB*). The *slc* of *B. cereus* ATCC 10987 harbors truncated *eag* and *secA2* genes and lacks *sap* and *slaP*. The environmental isolate *B. cereus* ATCC 14579 lacks genes for SLPs, SLP secretion and assembly, and SCWP acetylation but encodes a murein hydrolase with three SLH domains in place of Sap and EA1. Gene clusters were obtained from GenBank using accession numbers AE017334.2 for *B. anthracis* str. Ames Ancestor, CP009590.1 for *B. cereus* G9241, CP001746.1 for *B. cereus* biovar anthracis CI, AE016877.1 for *B. cereus* ATCC 14579, and NC_00,3909.8 for *B. cereus* ATCC 10987. Colors of genes were assigned arbitrarily with the exception of flanking genes that are not part of the *slc* and which are shown in gray along with their respective locus number.

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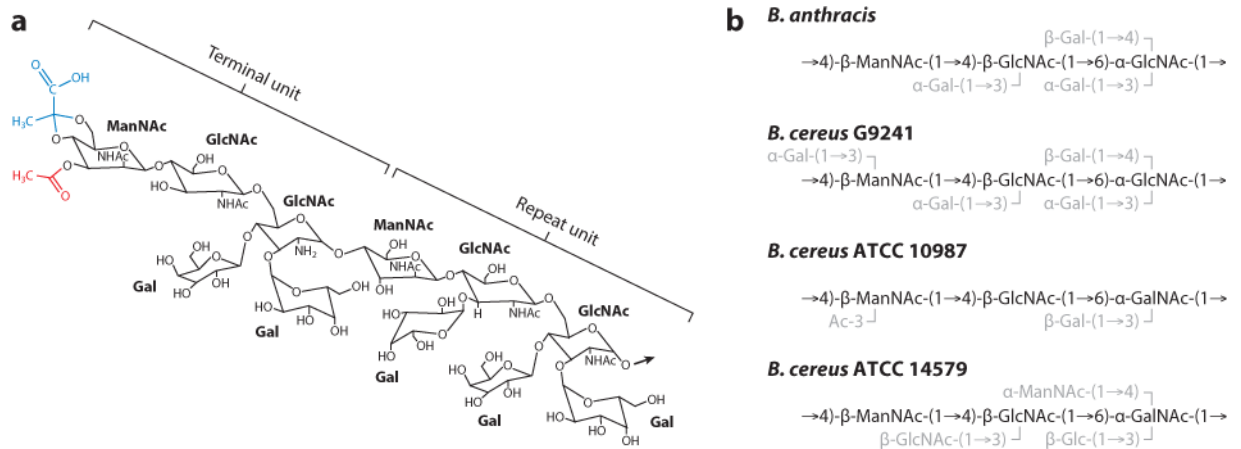


Figure 3.

Structure of the secondary cell wall polysaccharide (SCWP) in *B. cereus* s.l. isolates. (a) Structure of the terminal unit and the repeat unit of *B. anthracis* SCWP. The ketal pyruvyl modification of ManNAc at the nonreducing distal end of the SCWP is in blue. The acetyl group of the penultimate GlcNAc residue in the terminal unit is in red. The arrow identifies the glycosidic linkage to the next repeat unit. (b) Repeat unit structure of the SCWP from *B. anthracis*, *B. cereus* G9241, *B. cereus* ATCC 10987, and *B. cereus* ATCC 14579.

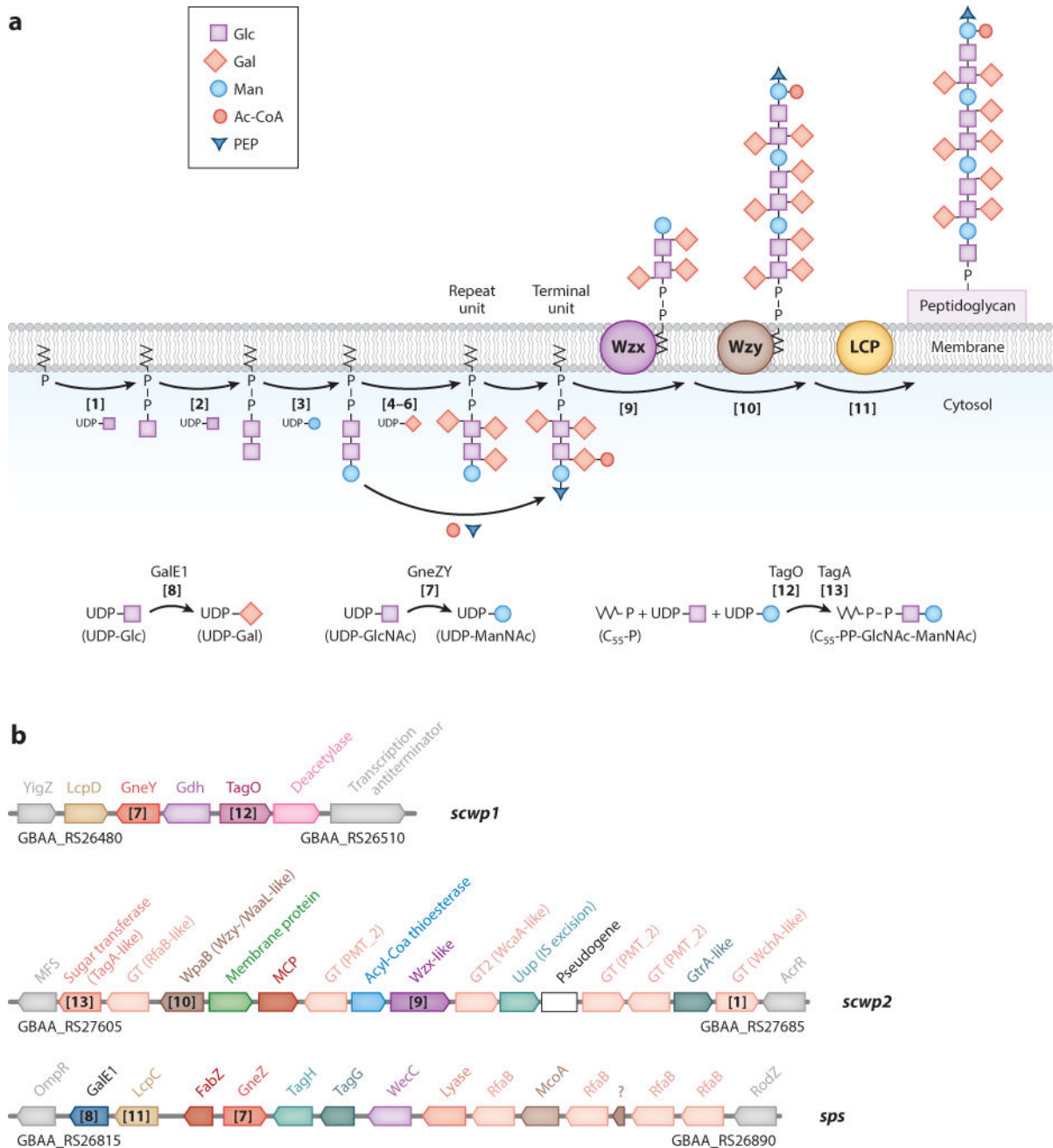


Figure 4.

Model for the biosynthesis of the secondary cell wall polysaccharide (SCWP) in *B. anthracis*. (a) Putative synthesis of undecaprenyl-pyrophosphate-linked SCWP trisaccharide subunits that are modified with galactosyl, ketal pyruvyl, or acetyl groups, translocated across the membrane by Wzx-like and polymerized by Wzy-like catalysts. The SCWP is finally linked to peptidoglycan by LCP enzymes. (b) Gene clusters contributing to SCWP synthesis include *slc* (see Figure 2b) as well as *scwp1*, *scwp2*, and *sps* (surface polysaccharide). Genes in gray flank gene clusters but are not part of them. The GenBank

identifiers of flanking genes are underneath gray gene symbols. Bracketed numbers in both panels indicate putative chemical reactions and enzymes for SCWP synthesis.

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