Release of Small Molecules during Germination of Spores of Bacillus Species

Barbara Setlow, Paul G. Wahome, and Peter Setlow*

Department of Molecular, Microbial and Structural Biology, University of Connecticut Health Center, Farmington, Connecticut 06030-3305

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Free amino acids, dipicolinic acid, and unidentified small molecules were released early in Bacillus spore germination before hydrolysis of the peptidoglycan cortex, but adenine nucleotides and 3-phosphoglycerate were not. These results indicate that early in germination there is a major selective change in the permeability of the spore’s inner membrane.

Dormant spores of Bacillus species contain a number of small molecules in their central region or core that become the prootoplasm of the growing cell. The small molecule present in spores at the highest level (≈600 μmol/g [dry weight]) is pyridine-2,6-dicarboxylic acid (dipicolinic acid [DPA]), which is largely present as a 1:1 chelate with divalent cations, predominantly Ca$^{2+}$ (Ca-DPA) (17). However, levels of glutamic acid (20 to 25 μmol/g [dry weight]), arginine (≈10 μmol/g [dry weight]), 3-phosphoglyceric acid (3PGA) (5 to 20 μmol/g [dry weight]), AMP (≈1 μmol/g [dry weight]), and ADP (≈0.2 μmol/g [dry weight]) are also significant in spores of Bacillus subtilis and Bacillus megaterium, and there is an even larger amount of sulfolic acid (≈100 μmol/g [dry weight]) in spores of at least one strain of B. subtilis (1, 7, 8, 17, 19).

DPA plays a significant role in dormant spore resistance to a number of treatments (11, 14). However, when spores are triggered to germinate, DPA and its associated cations are excreted rapidly in the initial stage (stage I) of spore germination (15, 18). This DPA release takes place before the hydrolysis of the spore’s peptidoglycan cortex in stage II that completes spore germination and allows progression into outgrowth and ultimately a return to active growth (15, 18). The precise mechanism for DPA release in spore germination is not clear but appears to involve the proteins encoded by the sporulation-specific spoVA operon (23, 25, 26). However, whether these SpoVA proteins comprise all or a part of a gated channel or pore allowing DPA release is not clear.

In addition to DPA, there is a variety of suggestive evidence indicating that other molecules are also released rapidly during spore germination and/or outgrowth. Thus, ≈80% of the spore’s K$^+$ and Na$^+$ are released during germination of B. megaterium spores, and this release appears to precede the release of at least the majority of DPA (21). These monovalent ions may then be reabsorbed, although DPA is not (21). The sulfolic acid in spores of B. subtilis strain SB133 is also released when these spores germinate (1), although the rate of this release has not been studied. Finally, the great majority of the free amino acids generated within spores at some period late in germination and/or early in outgrowth by the hydrolysis of the spore’s large pool of small acid-soluble proteins (SASP) is also released, although these amino acids are also then reabsorbed (20, 22). Since the dormant spore’s pools of small molecules are quite stable in spores incubated in water for years, there clearly must be a radical change in the barriers restricting movement of small molecules into and out of the spore when spores germinate and go into outgrowth. Indeed, there is strong evidence that a molecule as small as methylamine crosses the dormant spore’s inner membrane and enters the spore core extremely slowly, while moving rapidly across this membrane once spores have completed stage II of germination (5, 21). Even water may enter the dormant spore core slowly (29). The dormant spore’s major barrier to movement of small molecules into or out of the spore core appears to be the spore’s inner membrane (21), and fluorescent probes in this membrane are largely immobile in the dormant spore, but become freely mobile when spores complete germination (5). The volume encompassed by the spore’s inner membrane also increases 2-fold upon completion of spore germination, and thus the inner membrane surface area increases −1.5-fold, but without any phospholipid synthesis (5).

While the changes in the inner spore membrane during spore germination noted above require the hydrolysis of the spore’s large peptidoglycan cortex, DPA release takes place in stage I of germination prior to cortex hydrolysis. DPA release can be separated from cortex hydrolysis, since B. subtilis spores that lack both redundant cortex-lytic enzymes (CLEs), CwJ and SleB, or have a cortex that is not recognized by CLEs release DPA relatively normally in response to nutrient germinants (12, 15). As noted above, at least K$^+$ and Na$^+$ release precedes the release of the majority of DPA from B. megaterium spores during germination (21), and a number of other small molecules, including amino acids and sulfolic acid, are also released at some point after the triggering of spore germination (1, 20, 22), although it is not known if the latter releases take place before or after the inner membrane rearrangements that follow completion of spore germination. In this communication, we report studies on the release of small molecules early in the germination of B. megaterium and B. subtilis spores. This work makes the new finding that in addi-
tion to DPA and associated divalent cations, the dormant spore’s free amino acid pools as well as some other unidentified small molecules are also released approximately in parallel with DPA during spore germination. However, several other small molecules present at significant levels in dormant spores were released minimally if at all during this period.

The *Bacillus* strains used in this work are listed in Table 1. All *B. subtilis* strains are derived from strain PS832, which is a prototrophic derivative of strain 168. Strain PS533 (wild type) carries plasmid pUB110 encoding resistance to kanamycin (10 μg/ml) (16). Spores of *B. subtilis* strains were prepared at 37°C on 2× SG medium agar plates, and the spores were harvested, purified, and stored as described previously (9). *B. megaterium* strain PS1029 is a derivative of strain QM B1551, and *B. megaterium* spores were made at 30°C in liquid SNB medium and were purified and stored as described previously (20). Spores of *B. subtilis* strains were germinated at an optical density at 600 nm (OD_{600}) of 2 to 5 in 10 mM KPO_{4} buffer (pH 7.4)–1 mM L-alanine following a heat shock (30 min at 70°C) of spores in water. After incubation of germinating cultures at 37°C, 1-ml aliquots were centrifuged for 1 min in a microcentrifuge or 20-ml aliquots were centrifuged for 5 min at 14,000 × g, the latter for 3PGA analyses. The pellet fractions were isolated, washed twice with 1 ml water, and suspended in ~1 ml water, and the samples were either boiled for 30 min (samples for nuclear magnetic resonance [NMR]) or added to 4 ml boiling 1-propanol and boiled for 5 min (6, 19). In some cases, the supernatant fluid was also boiled in 4 ml of 1-propanol. The samples boiled in water were centrifuged, the supernatant fluid was run through a 1-ml Chelex column to remove divalent cations, most importantly Mn^{2+}, the run-through fractions were pooled and lyophilized, and the dry residue was dissolved in D_{2}O for proton NMR spectroscopy as described previously (6, 24). The samples boiled in 1-propanol were flash evaporated, the residue was dissolved in 1 ml water and centrifuged, and the supernatant fluid was analyzed for 3PGA, ATP, ADP, and AMP as described previously (19), but using a luminometer for assays of the adenine nucleotides. *B. megaterium* spores were germinated following a heat shock for 10 min at 60°C of spores in water. Germination used spores at 1 to 7 mg/ml (dry weight) in 10 mM KPO_{4} buffer (pH 7.5) plus 50 mM KBr and with or without 10 mM NaF to greatly slow 3PGA catabolism as described previously (19). Samples of 1 or 2 ml were harvested (the latter for ultimate analysis of 3PGA) and then processed and analyzed as described for *B. subtilis* spores, except that pellet fractions were not washed, and the supernatant fluid from germinating spores was also boiled directly and passed through Chelex, and the run-through fractions were lyophilized and dissolved in D_{2}O for proton NMR spectroscopy.

As noted above, DPA, glutamic acid, and arginine are present at significant levels in both *B. megaterium* and *B. subtilis* spores, although levels of other amino acids are much lower (7, 8, 17, 19). In addition to the peaks due to these known compounds, NMR spectra of hot water extracts of *B. subtilis* spores contained a number of large peaks not seen in the spectrum of an extract from *B. megaterium* spores (Fig. 1A and B; peaks labeled “X” in Fig. 1A). These additional large peaks in the *B. subtilis* spectrum are due to small molecules, as the compounds giving these peaks eluted after DPA and glutamic acid on a Biogel P2 column (data not shown). At least some of these additional large peaks have provisionally been identified as due to glyceric acid (C. A. Loshon, P. G. Wahome, and P. Setlow, unpublished results). *B. megaterium* and *B. subtilis* dormant spores also contained significant levels of AMP and ADP as well as 3PGA (see below), although these molecules were not seen in NMR spectra of processed extracts. The reason these molecules were not seen in NMR spectra is not known, but may be due to small amounts of residual Mn^{2+} broadening the peaks from these molecules considerably.

Previous work has shown that large amounts of many different amino acids are released during *Bacillus* spore germination, with these amino acids coming primarily from SASP hydrolysis (20). The release of large amounts of amino acids generated by SASP hydrolysis would obscure the release of dormant spores’ pools of free amino acids. Consequently, we examined amino acid release from *B. subtilis* spores in which cortex hydrolysis does not take place, since SASP hydrolysis by the SASP-specific germination protease, GPR, requires at least the initiation of cortex hydrolysis (15, 18). The spores used were from strains FB113 (*cwI/sleB*) and PS2307 (*cwID*) that either lack both CLEs (strain FB113) or have a cortex that does not contain muramic acid–lactam (strain PS2307), a modification essential for CLEs to recognize and act on cortex peptidoglycan (12, 18). While the spores of these two strains do not degrade their cortex in response to nutrient germinants and do not initiate enzyme action in the spore core, they do go through events in stage I of spore germination, most notably DPA release (15). Indeed not only was DPA released upon addition of the nutrient germinant l-alanine to these spores, the spore’s pools of free l-arginine and l-glutamic acid also disappeared from the spores and with approximately the same kinetics as for DPA release (Fig. 2A and B). The unknown small molecule or molecules that give rise to the peaks labeled “X” in NMR spectra of small molecules extracted from *B. subtilis* spores also disappeared from the germinating spores with similar kinetics (Fig. 2A and B).

**Table 1. Bacillus strains used in this study**

<table>
<thead>
<tr>
<th>Strain</th>
<th>Genotype</th>
<th>Source or reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>B. subtilis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>PS533</td>
<td>Wild type</td>
<td>16</td>
</tr>
<tr>
<td>PS2307</td>
<td><em>cwID</em></td>
<td>12</td>
</tr>
<tr>
<td>FB113</td>
<td><em>cwID/sleB</em></td>
<td>10</td>
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<tr>
<td>B. megaterium</td>
<td></td>
<td></td>
</tr>
<tr>
<td>QM B1551</td>
<td>Wild type</td>
<td>H. Levinson, U.S. Army Lab, Natick, MA</td>
</tr>
<tr>
<td>PS1029</td>
<td><em>gpr</em></td>
<td>13</td>
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Mutant strains of *B. megaterium* lacking CLEs or CwID are not available. Consequently, to block release of free amino acids generated by SASP hydrolysis during spore germination and outgrowth, we used an isogenic mutant strain (PS1029) in which the *gpr* gene that encodes the SASP-specific protease has been inactivated. The spores of this strain initiate germination normally, but SASP degradation is greatly slowed (13). These spores were germinated with the salt KBr, which most likely acts by stimulating the spore’s nutrient germinant receptors (3, 4), and use of this germinant allowed analysis of amino acids in
germination exudates by NMR spectroscopy without interference by the germinant itself. Again, these spore’s pools of both glutamic acid and arginine were released with approximately the same kinetics as DPA (Fig. 3) (data not shown). There were also no changes in the level of the total amounts of these compounds present in the germinating spore culture (Fig. 3) (data not shown).

While some small molecules are clearly released approximately in parallel with DPA early in spore germination, it seems extremely unlikely that this should be the fate of all small molecules. Indeed, ATP generation from catabolism of endogenous 3PGA as well as exogenous compounds, including amino acids released after SASP hydrolysis, also begins either late in stage II of spore germination or early in outgrowth (15, 18–20). To examine release of other spore small molecules, levels of ATP in germinating PS533 (wild type) spores were determined in both the spore pellet fraction and in the germination exudate. As expected, <5% of total ATP was found in the germination exudate (Table 2). Similarly, <5% of the pools of AMP and ADP present in dormant cwlJ sleB B. subtilis spores was released into the supernatant fluid upon germination of these spores with L-alanine (Table 2). These cwlJ sleB

FIG. 1. Proton NMR spectra of small molecules in B. megaterium (A) and B. subtilis (B) spores. Small molecules were extracted from 10 mg (dry weight) of dormant spores of B. megaterium QM B1551 (wild type) (A) and B. subtilis strain PS533 (wild type) (B), extracts were processed, and the proton NMR spectra of the small molecules were obtained as described in the text. Peaks due to known compounds are labeled, although some of the peaks in the arginine region are from glutamate and both arginine and glutamate give additional peaks that are not labeled. The peaks labeled “X” that flank the arginine region are due to an unknown compound or compounds in B. subtilis spores. The spectrum in panel A has been magnified ~2.5 times more than the spectrum in panel B.

FIG. 2. Release of small molecules during germination of B. subtilis spores. Spores of B. subtilis strains FB113 (cwlJ sleB) (A) and PS2307 (cwlD) (B) were germinated, and at various times the small molecules remaining in the germinated spores were extracted, extracts were processed, and small molecules were analyzed by NMR spectroscopy to quantitate small molecules as described in the text. ○, DPA; ▲, glutamic acid; △, arginine; and ●, X. Note that whatever gives rise to all of the X peaks was released together. The percentage release of the various molecules was calculated by taking the amount in dormant spores as 100% and assuming that there was no change in this total value throughout the experiment.
spores also does not require cortex hydrolysis. While a number of small molecules were released rapidly during spore germination, clearly this release is at least somewhat selective, as AMP, ADP, ATP, and 3PGA were released minimally if at all during *B. megaterium* and *B. subtilis* spore germination.

It is notable that the release of the additional small molecules during spore germination takes place before hydrolysis of the spore’s cortex and the reorganization of the spore’s inner membrane that takes place in fully germinated spores. Somehow a membrane that has not allowed passage of DPA and several other small molecules, perhaps for years, allows the rapid passage of the same molecules once spore germination is triggered. This triggering can be by nutrients, as shown in the present work, or by elevated pressures or the cationic surfactant dodecylamine, as shown previously (24, 26). The major question then is how this might be accomplished. One formal possibility is that this is via some type of gated mechanosensitive channel (2). However, deletion of all known mechanosensitive channels from *B. subtilis* has no effect on at least DPA release during spore germination (27, 28), and known mechanosensitive channels also allow release of molecules up to the size of ATP and even larger (2). Whatever the mechanisms allowing the rapid release of DPA and other small molecules in the first minute of spore germination, this large increase in the inner membrane’s permeability to small molecules appears to be selective, as a number of small molecules are not released.

![FIG. 3. Release of small molecules during germination of *B. megaterium* spores.](image-url)

**TABLE 2. Levels of adenine nucleotides and 3PGA retained by *Bacillus* spores during spore germination**

<table>
<thead>
<tr>
<th>Strain</th>
<th>Adenine nucleotides or 3PGA</th>
<th>Level of nucleotide or 3PGA in spores (μmol/g spores [dry wt])</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Dormant</td>
</tr>
<tr>
<td><em>B. subtilis</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PS33 (wild type)</td>
<td>ATP</td>
<td>≤0.002</td>
</tr>
<tr>
<td></td>
<td>ADP</td>
<td>0.12</td>
</tr>
<tr>
<td></td>
<td>AMP</td>
<td>0.6</td>
</tr>
<tr>
<td>FB113 (cwIJ sleB)</td>
<td>3PGA</td>
<td>18</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>B. megaterium</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td>QM B1551</td>
<td>3PGA</td>
<td>18</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>B. subtilis</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FB113 (cwIJ sleB)</td>
<td>3PGA</td>
<td>5</td>
</tr>
</tbody>
</table>

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Spores of various strains were germinated and extracted, extracts were processed, and adenine nucleotides and 3PGA were quantitated as described in the text. All values are the result of duplicate determinations.

Values not in parentheses are the levels found in the germinated spore pellet fraction; values in parentheses are the percentage of the total released in the germination exudate.

Values are for spores germinated at an OD_{600} of 2 for 15, 30, and 45 min; DPA release was >85% complete in 30 min.

Values are for spores germinated at an OD_{600} of 2 for 90 min; at this time, DPA release was >90% complete.

Spores were germinated for 15 min at an OD_{600} of 50 with NaF present; at this time, DPA release was 85% complete.

Spores were germinated for 30 min at an OD_{600} of 50 with NaF present; at this time, DPA release was >95% complete.

Spores were germinated for 30 min at an OD_{600} of 50 without NaF; at this time, DPA release was >90% complete.
Some of this selectivity may be based on molecular size, as the molecules released almost completely (DPA, glutamate, and arginine) have a lower molecular weight than those released poorly if at all (3PGA and adenine nucleotides), although this selectivity may depend primarily on the hydrated dimensions of these molecules. In addition, a number of these molecules are chelated to and likely released with divalent ions, which further complicates analysis of selectivity based solely on molecular weight. The large increase in membrane permeability early in spore germination, most likely in stage I, is also almost certainly transient, since a number of these small molecules are subsequently taken back up. Indeed, the continued high permeability of a plasma membrane to hydrophilic small molecules such as amino acids would seem to be incompatible with life. It seems likely that establishing the mechanism for and regulation of this transient change in the permeability properties of the spore’s inner membrane during spore germination will greatly increase our understanding not only of spore germination, but also features of dormant spores as well.

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REFERENCES