Characterization of *Bacillus subtilis* \(\gamma\)-glutamyltransferase and its involvement in the degradation of capsule poly-\(\gamma\)-glutamate

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INTRODUCTION

\(\gamma\)-Glutamyltransferase (EC 2.3.2.2; GGT) is widely distributed in nature, from bacteria to animals (Tate & Meister, 1981, 1985). Animal GGT is located on the external surface of epithelial cells, where it catalyses transfer of the \(\gamma\)-glutamyl moiety from glutathione to amino acid or peptide acceptors (transferase), or to \(H_2O\) (hydrolysis). The resultant \(\gamma\)-glutamyl product can be recruited for glutathione synthesis to maintain appropriate cellular pools of glutamate and the dipeptide produced by the hydrolytic reaction, are metabolized as amino acid sources well as glutamate and the dipeptide produced by the reaction. Some strains of *B. subtilis* and *Bacillus anthracis* produce capsular poly\((\gamma\)-glutamic acid\) (\(\gamma\)PGA; Thorne, 1993). In both *B. subtilis* and *B. anthracis*, the membrane \(\gamma\)PGA synthetic proteins encoded by the *capBCA* (also referred to *ywsC–ywtAB* or *pgsBCA* in *B. subtilis*) operon catalyse synthesis of the capsule polypeptide (Ashiuchi et al., 1999; Makino et al., 1989; Urushibata et al., 2002). However, the stereochemistry of these bacterial \(\gamma\)PGAs depends upon the structure of the \(\gamma\)-glutamyl linkage, and the synthesis of each \(\gamma\)PGA is regulated differently. *B. anthracis* \(\gamma\)PGA consists of D-glutamate only, and it is produced in the presence of serum or under high atmospheric CO\(_2\) concentrations, circumstances that mimic host environments where the capsule functions as a protective barrier against phagocytosis by macrophages (Makino et al., 1988, 1989, 2002). On the other hand, *B. subtilis* produces capsule \(\gamma\)PGA consisting of both D- and L-glutamate specifically during the early stationary phase. This growth-phase-dependent synthesis of the capsule is mediated through the ComQXPA quorum-sensing mechanism, which also controls the expression of other stationary-phase-specific traits (Dubnau, 1999; Lazazzera et al., 1999; Tran et al., 2000).

During early stationary phase, *Bacillus subtilis* NAFM5 produces capsular poly\((\gamma\)-glutamic acid\) (\(\gamma\)PGA, \(2 \times 10^6\) Da), which contains D- and L-glutamate, and then degrades it during late stationary phase. The \(\gamma\)-glutamyltransferase (EC 2.3.2.2; GGT) of this strain successively hydrolysed \(\gamma\)PGA from the amino-terminal end, to yield both D- and L-glutamate. This enzyme was specifically synthesized during the stationary phase through transcriptional activation of the corresponding *ggt* gene by the ComQXPA quorum-sensing system. A *ggt* knockout mutant degraded \(\gamma\)PGA into \(1 \times 10^5\) Da fragments, but not any further, indicating that the capsule \(\gamma\)PGA is first internally degraded by an endo-type of \(\gamma\)PGA hydrolase into \(1 \times 10^5\) Da intermediates, then externally into glutamates via GGT. Due to its inability to generate the glutamates from the capsule, the *ggt* mutant sporulated more frequently than the wild-type strain. The results show that *B. subtilis* GGT has a powerful exo-\(\gamma\)-glutamyl hydrolase activity that participates in capsule \(\gamma\)PGA degradation to supply stationary-phase cells with constituent glutamates.

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Abbreviations: GGT, \(\gamma\)-glutamyltransferase; GNA, \(\gamma\)-glutamyl-p-nitroanilide; \(\gamma\)PGA, poly\((\gamma\)-glutamic acid\).
Both *B. anthracis* and *B. subtilis* degrade their capsule \( \gamma \)PGAs, but probably via different enzymes. *B. anthracis* degrades the capsule polypeptide into 2–14 kDa fragments via a depolymerase encoded by *capD*, which lies immediately downstream of the *cap* operon (Makino et al., 2002). The resultant polypeptide fragments appear to be required for the pathogen to flourish in hosts (Makino et al., 2002). The *B. subtilis* capsule is degraded during late stationary phase. This bacterium has the *ywtD* gene encoding \( \gamma \)-D-glutamyl hydrolase at a locus corresponding to *capD* (Suzuki & Tahara, 2003). The *ywtD* product, however, has no amino acid sequence similarity to the CapD depolymerase, and it cleaves \( \gamma \)PGA in vitro into fragments of 490 and 11 kDa (Suzuki & Tahara, 2003). *B. subtilis* GGT appears to be capable of generating D- and L-glutamate in vitro from \( \gamma \)PGA (Abe et al., 1997). However, the precise hydrolytic mechanism of this enzyme has not been defined, and whether Ywtd and GGT participate in the in vivo degradation process remains unknown.

*B. subtilis* can utilize both D- and L-glutamate as nitrogen sources (Kimura et al., 2004). D-Glutamate catabolism by this bacterium proceeds after conversion to the L-form by glutamate racemases (the *rac*E and *yrpC* products). Mutants of *racE* or *yrpC* accumulate D-glutamate in late-stationary-phase cultures (Kimura et al., 2004), indicating that *B. subtilis* cells degrade capsule \( \gamma \)PGA into its constituent glutamates outside the cells, and utilize them as nitrogen sources during late stationary phase.

We report here that *B. subtilis* GGT has powerful exo-\( \gamma \)-glutamyl hydrolase activity towards \( \gamma \)PGA, and generates both the amino-terminal D- and L-glutamate of the polypeptide. Experiments with a mutant lacking GGT activity demonstrated that this enzyme is involved in \( \gamma \)PGA degradation in vivo to yield the constituent amino acids, and that *B. subtilis* has, in addition to Ywtd, a second endo-\( \gamma \)PGA hydrolase that degrades the capsule polypeptide into 1 × 10^5 Da fragments. Furthermore, we showed that when the nitrogen supply is limited, mutant cells lacking GGT sporulate more frequently than the wild-type strain, suggesting that capsule glutamates serve *B. subtilis* as nitrogen sources during the stationary phase.

### METHODS

**Bacterial strains and media.** *B. subtilis* NAFM5 (Rif^\*^) is a derivative of the commercial starter strain Miyagino, which is used in the fermentation of soybeans to produce natto (a Japanese foodstuff) by introducing rifampicin resistance (Rif^\*^) and by curing the plasmids pUH1 (=pTA1015) and pNGL1 (=pLS20) (Kimura & Itoh, 2003; Meijer et al., 1995, 1998; Nagai et al., 1997). *B. subtilis* NAFM65 (comP::Spc), NAFM90 (ggt::Spc), and NAFM96 (ggt::Spye::ggt) were constructed from strain NAFM5 as described below. *B. subtilis* strains were cultured in Luria–Bertani (LB) medium, or in E9 minimal medium (Birrer et al., 1994) with the appropriate antibiotics and supplements (Tran et al., 2000).

**Construction of mutants.** A DNA region corresponding to the mature part of GGT was amplified using KOD DNA polymerase (Toyobo Biochemicals), primers [5’-GATGAGTCAAAAAGAGTA-GATGTGGGA-3’], nt 106–132 relative to the translation initiation codon (1) of *ggt* (DDBJ/EMBL/GenBank accession number AB095984) and 5’-TATTATTGTAAAAAATTATGCGGATCGC-3’, complementary to nt 1734–1762 of *ggt*, and the chromosomal DNA of *B. subtilis* NAFM5 (Kimura & Itoh, 2003) as the template. The amplified DNA region was then cloned into the *HincII* site on plasmid pUC118 (Veira & Messing, 1987) to verify the nucleotides by sequencing. A spectinomycin (Spc)-resistance cassette, isolated from plasmid pDG1726 (Guérout-Fleury et al., 1995) as an EcoRV–*HincII* fragment, was then inserted into the *Stu*I site of *ggt* on the resultant plasmid. After linearization at the unique *SacI* site on the vector sequence, the plasmid DNA was used to knock out the *ggt* of strain NAFM5 by double-crossover recombination, generating strain NAFM90 (ggt::Spc). The Sop–SpbI fragment carrying the entire *ggt* gene was integrated into the *amyE* locus of strain NAFM90 via plasmid pDG1661 (Guérout-Fleury et al., 1996) by homologous recombination to create strain NAFM96 (ggt::Spc *amyE::ggt*). Replacing *comP* in strain NAFM5 with *comP::Spc* using a pUC118 derivative carrying a 4·4 kb *HindIII* fragment containing *comP* (Tran et al., 2000), which had been inactivated by insertion of the EcoRV–*HincII* Spc-resistance cassette (see above) at the *ClaI* site, resulted in strain NAFM65 (comP::Spc). Southern blotting (Nakada & Itoh, 2002) confirmed that the Spc-resistance cassette and *ggt* at the target loci were correctly inserted.

**Preparation of \( \gamma \)PGA and \( \gamma \)-glutamyltetrapeptides.** We purified \( \gamma \)PGA from *B. subtilis* NAFM90 (ggt::Spc) cultures incubated for either 2 or 7 days on E9 agar (without glutamate) containing 0·5 µg biotin ml^-1^, as described by Nagai et al. (1997). We determined the molecular masses of the polypeptides by gel-permeation HPLC using an Asahipak GFA-7M column (Asahi Chemical Industry) (Nagai et al., 1997). The content of D- and L-glutamate in the polypeptides was determined after hydrolysis with 1 M HCl for 3 h, and by using CrownPack CR (+) and CrownPack CR (−) chiral columns (Daicel Chemical Industry) (Nagai et al., 1997). The molecular masses of the polypeptides isolated from the 2 and 7 day cultures were 2 × 10^6 and 1 × 10^5 Da, respectively, and they both comprised 54% D-glutamate. \( \gamma \)PGA of 1 × 10^5 Da, with a high D-glutamate content (76%), was also prepared from strain NAFM90 cultures incubated on GSP agar containing 0·1 mM MnCl_2_ (Nagai et al., 1997) for 7 days. The synthetic glutamyltetrapeptides \( \gamma \)-D-Glu (\( \gamma \)-L-Glu)_2, \( \gamma \)-D-Glu (\( \gamma \)-L-Glu)_3, \( \gamma \)-D-Glu (\( \gamma \)-L-Glu)_4, and \( \gamma \)-D-Glu, \( \gamma \)-L-Glu, \( \gamma \)-D-Glu) were obtained from Hokkaido System Science (Sapporo, Japan). The oligopeptides were constructed using a Pioneer Dual Column Peptide Synthesizer (Applied Biosystems). We confirmed the molecular mass and purity (99%) of the synthetic peptides by mass spectrometry (Apex II 70e, Bruker Daltonics) and gel-permeation HPLC, respectively (Nagai et al., 1997).

**Enzyme assays.** We measured GGT activity using \( \gamma \)-glutamylp-nitroanilide (\( \gamma \)GNA) as the substrate in the presence of the acceptor glycglycine, according to Suzuki et al. (1986). One unit was defined as the amount of enzyme that was required to produce 1 µmol p-nitroaniline \((a_{\text{1000}} 8800 \text{~M}^{-1} \text{~cm}^{-1})\) per min. We assayed the hydrolytic activity of GGT towards \( \gamma \)PGA and \( \gamma \)-glutamyltetrapeptide in a reaction mixture (400 µl) containing 2 mg ml^-1^ 1 × 10^5 Da \( \gamma \)PGA or 0·5 mM synthetic \( \gamma \)-glutamyltetrapeptide, 20 mM sodium phosphate buffer (pH 6·9), 150 mM NaCl (omitted from the reaction with the tetrapeptide) and GGT (0·4 µg), at 37 °C. Portions (60 µl) of the reaction mixtures were withdrawn at 0, 5, 10, 20, 30 and 45 min, and then boiled for 10 min to terminate the reaction. The D- and L-glutamate reaction products were separated using HPLC chiral columns (see above), and quantified using a Shimadzu RF-10AXL fluorescent detector (excitation at 345 nm, emission at 455 nm) after coupling with o-phthalaldehyde.

**Purification of GGT.** *B. subtilis* GGTs have been purified from...
strains NR-1 and 168, and amino acid sequencing of the large and small subunits has confirmed that they are the products of ggt (Minami et al., 2003; Ogawa et al., 1991, 1997; Kunst et al., 1997).

We purified GGT from stationary-phase cultures (2 l) of B. subtilis NAFM5 in E9 medium. The culture supernatant was dialysed against 25 mM Tris/HCl buffer (pH 7.5) containing 0.5 mM DTT, and eluted through a Hiprep 16/10 DEAE column (Amersham Biosciences) using a linear gradient of NaCl (0–4 M). After dialysis against 10 mM sodium phosphate buffer (pH 6.8) containing 0.5 mM DTT, fractions containing the enzyme were applied to a hydroxyapatite column (CHT5-I; Bio-Rad), and the enzyme was eluted with a gradient (0–0.2 M) of sodium phosphate, pH 6.8. Active fractions were dialysed against 10 mM sodium phosphate (pH 6.8) containing 0.5 mM DTT, and then eluted through a MonoQ column (HR 5/5; Amersham Biosciences) using a linear NaCl gradient (0–0.35 M) in the same buffer. Combined active fractions were concentrated using Centricon-10 (Millipore), and finally gel filtered through a Superose12 column (Amersham Biosciences) using 10 mM sodium phosphate buffer (pH 6.8) containing 0.15 M NaCl as the running buffer. These procedures resulted in a 2 M yield of GGT that was purified 122-fold. The purified GGT (56 kDa) was apparently homogeneous, and consisted of 44 and 23 kDa subunits when determined using a Protein Assay kit (Bio-Rad) with bovine serum albumin as the standard. We performed SDS-PAGE using 20 % (w/v) protoplasts were quickly sedimented by centrifugation, and sus-

RESULTS

Hydrolitic activities of GGTs towards γPGA

We initially examined the hydrolitic activity of B. subtilis GGT towards γPGA in vitro by measuring the amounts of L-glutamate generated by NAD-dependent glutamate dehydrogenase. The enzyme yielded L-glutamate from 1 × 10^5 Da γPGA, containing 54 % D-glutamate, at a rate of 4.4 μmol min^{-1} (mg protein)^{-1}, and with a K_m value of 9.0 μM. To determine whether B. subtilis GGT could also generate D-glutamate from the γPGA, we separated the D- and L-isomers using HPLC chiral columns. D- and L-Glutamate were generated in amounts corresponding to their proportions in the substrate (Fig. 1a). When 1 × 10^5 Da γPGA containing 76 % D-glutamate was the substrate, the enzyme yielded approximately three times more D-than L-glutamate (Fig. 1b). B. subtilis GGT thus appeared to have no apparent specificity in terms of the D- or L-configuration of the γ-glutamyl linkage. As determined by

Two-dimensional immunoelectrophoresis. γPGA was extracted from portions (1 ml) of B. subtilis strains NAFM5 (wild-type) and NAFM90 (ggt::Spc) incubated in medium E9 (100 ml) as described by Nagai et al. (1997). After dissolution in 10 μl 20 mM sodium phosphate buffer (pH 6.9), 8 μl portions of the samples were resolved by electrophoresis through 1.2–2 % (w/v) agarose gels containing 0.1 M Tris/HCl (pH 8.5) at 2 mA cm^(-1) for 6 h. Second-

Primer extension and Northern blotting. Cells cultivated in the media specified in Results (100 ml) were incubated in 15 ml of 20 % (w/v) sucrose containing 6 mg egg-white lysozyme per ml, 50 mM Tris/HCl (pH 7.5) and 50 mM EDTA, at 37 °C for 3 min. The resul-
tant protoplasts were quickly sedimented by centrifugation, and sus-
pended in 10 ml acetate/EDTA buffer (pH 4.8) containing 30 mM sodium acetate, 1 mM EDTA and 10 mM Tris. Thereafter, total RNA was extracted with hot phenol (Nakada & Itoh, 2002). For primer extension analysis, RNA samples (20 μg) were annealed with an oligonucleotide (5'-AGCGACTAAACAGAACTAAGCAGAGC-3', complementary to nt 31–58 of ggt) labelled with β-32P at the 5' end by using [α-32P]dCTP (220 TBq mmol^(-1); Amersham Biosciences) and T4 polymerase (Toyobo Biochemicals). Complementary strands were synthesized using AMV reverse transcriptase XL (Toyobo Biochemicals), and resolved on a denatured 6 % (w/v) polyacrylamide gel. Sequence ladders were generated using a BcaBest sequencing kit (Takara Shuzo; http://www.takara-bio.co.jp) with the oligonucleotide as the primer, and plasmid pNAG201 carrying an SpI–BglII ggt fragment as the template. Total RNA (10 μg) was resolved for Northern blotting on 1.2–2 % (w/v) agarose gels, and blotted onto nylon membranes (Hybond-N+; Amersham Biosciences). A ggt DNA fragment amplified by PCR, as described above, was labelled using a random-prime labelling kit (Nippon Gene) and [α-32P]dCTP (220 TBq mmol^(-1); Amersham Biosciences), and hybridized with membrane ggt mRNA. Hybridized probes were visualized on X-ray films.

Fig. 1. Hydrolysis of γPGA by B. subtilis GGT. B. subtilis GGT was incubated with γPGA (1 × 10^5 Da) consisting of 54 % (a) and 76 % (b) D-glutamate. Portions of reaction mixtures were withdrawn after 0, 15, 30, 45 and 60 min, and the amounts of D-glutamate (●) and L-glutamate (○) were determined using CrownPack CR(+), and CrownPack CR(−) HPLC chiral columns, as well as standard curves for D- and L-glutamate. Values are means of two measurements; SD values are below 5 % of the corresponding means.
the total amounts of D- and L-glutamate generated from 1 × 10^5 Da γPGA (54 % D-glutamate), the specific activity of *B. subtilis* GGT towards this polypeptide was 8.6 μmol min⁻¹ (mg protein)⁻¹. Similar activity was determined with 2 × 10^6 Da γPGA containing 54 % D-glutamate. Agarose gel electrophoresis and gel-permeation HPLC detected only a marginal reduction in the molecular sizes of the substrate polypeptides at the end of the incubation (60 min), suggesting that *B. subtilis* GGT externally cleaves γPGA. In contrast, the amounts of L-glutamate generated were negligible with either *E. coli* or bovine GGT, even when the reactions included 0.06 γ-GNA-hydrolase units of the enzymes, which were equivalent to 2.4 μg *B. subtilis* GGT.

### Hydrolysis of the N-terminal γ-glutamyl bond

We investigated the direction of hydrolysis, as well as the preferred configuration of the terminal residues and γ-glutamyl linkages, using a set of γ-L-glutamyltripeptides labelled with γ-D-glutamate or α-L-glutamate at either the amino or carboxyl terminal, γ-D-Glu-(γ-L-Glu)₃, (γ-L-Glu)₃-γ-D-Glu, α-L-Glu-(γ-L-Glu)₃, and (γ-L-Glu)₃-α-L-Glu, as the substrates. When γ-D-Glu-(γ-L-Glu)₃ was incubated with the enzyme, D-glutamate was generated from the start of the incubation (Fig. 2a). In contrast, the D-isomer of (γ-L-Glu)₃-γ-D-Glu appeared at a later stage of incubation (Fig. 2a), showing that hydrolysis proceeds at the amino terminal. The hydrolytic rates of the tetrapeptides determined as the total amounts of D- and L-glutamate were almost identical (data not shown), supporting the notion that *B. subtilis* GGT has no significant stereospecificity for the terminal residue or the γ-peptide bond. The enzyme was active towards (γ-L-Glu)₃-α-L-Glu, but inert to α-L-Glu-(γ-L-Glu)₃ (Fig. 2b). GGT was also active towards the γ-glutamyltetrapeptides, showing 2.8-fold greater activity than it did towards γPGA [the specific activities, as determined by the total amounts of D- and L-glutamate generated from 0.5 μM synthetic γ-glutamyltetrapeptide, were 25.6 μmol min⁻¹ (mg protein)⁻¹ for γ-D-Glu-(γ-L-Glu)₃ and 24.5 μmol min⁻¹ (mg protein)⁻¹ for (γ-L-Glu)₃-γ-D-Glu], although the *Kₘ* values were similar (9.0 μM for γPGA, and 8.0 μM for each γ-glutamyltetrapeptide).

### Accumulation of degradation intermediates in a ggt mutant culture

We constructed a ggt knockout mutant of *B. subtilis* NAFM5 by inserting a Spc-resistance cassette, and examined whether this mutant can degrade the capsule. The mutant NAFM90 (ggt::Spc) produced no detectable GGT (<0.01 × 10⁻³ units ml⁻¹), even after 6 days incubation when the wild-type had accumulated as much as 28 × 10⁻³ units ml⁻¹ (Fig. 3), but it thrived normally, like the wild-type, in minimal medium. Two-dimensional immunoelectrophoresis showed that γPGA synthesis by both the wild-type and NAFM90 (ggt::Spc) strains began after about 2 days of incubation, and continued similarly.

![Fig. 2. Direction (a) and specificity (b) of γ-peptide hydrolysis by *B. subtilis* GGT. (a) Synthetic γ-tetrapeptides, γ-D-Glu-(γ-L-Glu)₃ (●) and (γ-L-Glu)₃-γ-D-Glu (○), were incubated with *B. subtilis* GGT, and the amounts of D- and L-glutamate generated were quantified as in Fig. 1(b). (b) Tetrapeptides, (γ-L-Glu)₃-α-L-Glu ( ●) and α-L-Glu-(γ-L-Glu)₃ (○), were incubated with the GGT, and the L-glutamate liberated was quantified by HPLC. Values are means of two measurements; SD values are below 5 % of the corresponding means.](image1)

![Fig. 3. Regulation of GGT synthesis by the quorum-sensing system and exogenous L-glutamate. *B. subtilis* NAFM5 (wild-type) was cultured in E9 medium with (●) or without (○) 2 %, w/v, L-glutamate, and strain NAFM65 (comP::Spc) (△) was incubated in E9 medium without L-glutamate. Activities of GGT in culture supernatants were determined every day for 6 days. Transition from exponential to stationary phase. Values are means of two independent measurements; SD values are below 5 % of the corresponding means.](image2)
Regulation of GGT synthesis

B. subtilis produces GGT like other exoenzymes, during the stationary phase (Xu & Strauch, 1996). Growth-phase-specific exoenzyme synthesis is controlled by a quorum-sensing system that consists of four elements: ComX pheromone (a cell-density signal peptide), ComQ (responsible for processing, modification and secretion of the pheromone), ComP (a pheromone-sensor and histidine kinase) and ComA (a cognate response regulator of ComP) (Lazazzera et al., 1999; Tran et al., 2000). The accumulation of $1 \times 10^5$ Da intermediates in L-glutamate cultures suggests that this amino acid inhibits either the activity or the synthesis of

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**Fig. 4.** Formation and degradation of $\gamma$PGA in wild-type strain NAFM5 (a) and ggt::Spc mutant NAFM90 (b) cultures. $\gamma$PGA extracted from NAFM5 (wild-type) and NAFM90 (ggt::Spc) cultures after incubation for 2, 4 and 7 days was analysed by two-dimensional immunoelectrophoresis (Uchida et al., 1993). Molecular masses of the polypeptides were determined by gel-permeation HPLC (see Fig. 5). Sizes of $\gamma$PAs ($2 \times 10^6$ Da) in 4 day cultures were slightly smaller than those ($5 \times 10^6$ Da) in 2 day cultures, perhaps due to spontaneous fragmentation.

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**Fig. 5.** HPLC analysis of $\gamma$PGA in wild-type and ggt::Spc mutant cultures. $\gamma$PGA extracted from strains NAFM5 (wild-type) and NAFM90 (ggt::Spc) cultured with or without 2% (w/v) L-glutamate for 4 (a) and 7 (b) days was quantified using Asahipak GFA-7M HPLC gel-permeation columns. Amounts and molecular masses of polypeptides were determined using standard curves of purified $\gamma$PGA and retention times relative to those of pullulan molecular standards (Nagai et al., 1997), respectively. Trace 1, wild-type culture with L-glutamate (L-Glu); trace 2, mutant culture without L-glutamate; trace 3, wild-type culture without L-glutamate.

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up to 4 days (Fig. 4a, b); the cultures entered the stationary phase after 1-5 days of incubation. During an additional 3 days of incubation, the size and amount of $\gamma$PGA in the wild-type culture became minimal (Fig. 4a). In the mutant culture, the polypeptides also became shorter, but they appeared to be significantly larger than those in the wild-type culture (Fig. 4b). The $\gamma$PGA in the 7 days cultures of the mutant formed a cross-reactive area larger than that in the 4 day cultures. Since smaller polypeptides form larger cross-reactive areas on agarose gels due to faster migration rates, the cross-reactive areas are not proportional to the amounts of polypeptide, unless the polypeptide is the same size (Uchida et al., 1993). HPLC analysis of the polypeptides from wild-type and mutant cultures revealed that 0.6 mg ml$^{-1}$ of $2 \times 10^6$ Da $\gamma$PGA accumulated in the wild-type culture during the first 4 days (Fig. 5a), and that the amounts and size of the polypeptides in the 7 day cultures decreased to below 0.1 mg ml$^{-1}$ and to $5 \times 10^5$ Da, respectively (Fig. 5b). The mutant culture produced similar amounts (0.7 mg ml$^{-1}$) of $2 \times 10^6$ Da $\gamma$PGA during the first 4 days (Fig. 5a). This amount remained unchanged after 7 days, but the size of the polypeptide decreased to $1 \times 10^5$ Da (Fig. 5b). These results suggest that $\gamma$PGA is first degraded into $1 \times 10^5$ Da intermediates by an end-type hydrolytic enzyme, and then to glutamates by GGT. Strain NAFM63 (ggt::Spc amyE::ggt), harbouring an intact ggt allele at the amyE locus, degraded $\gamma$PGA in a similar way to the wild-type strain (data not shown), confirming that GGT is responsible for degradation of the $1 \times 10^5$ Da intermediates. Adding 2% (w/v) L-glutamate, the precursor of $\gamma$PGA (Urushibata et al., 2002), to the medium caused a 2.5-fold increase in $\gamma$PGA production (Fig. 5a). After 7 days of incubation, the quantities of polypeptides in the glutamate medium remained unchanged, even in the wild-type culture. However, the polypeptides became similarly smaller to those of the degradation intermediates that accumulated in the ggt mutant culture (Fig. 5b).
GGT. L-Glutamate (5 mM) inhibited the hydrolytic activity of GGT towards γ-PGA, but only by 6 %, suggesting that L-glutamate hampers the degradation of 1 × 10^5 Da intermediates mainly by preventing GGT synthesis.

To test whether GGT synthesis is under the control of the quorum-sensing system and L-glutamate, we measured GGT activities in NAFM5 (wild-type) cells cultured with or without L-glutamate, and in comP mutant NAFM65 (comP::Spc) cells incubated without L-glutamate. In the absence of L-glutamate, the wild-type strain initiated GGT synthesis immediately after the culture entered the stationary phase, and this continued for another 3 days, producing maximal levels of the enzyme after 5 days (Fig. 3). Exogenous L-glutamate reduced GGT synthesis to marginal levels (2.1 × 10^{-3} units ml^{-1}). The comP mutant produced negligible amounts (0.08 × 10^{-3} units ml^{-1}) of the enzyme during the entire stationary phase (up to 6 days; Fig. 3), showing that the quorum-sensing system positively controls GGT synthesis and that L-glutamate antagonizes this positive control.

**Regulation of ggt transcription**

The ggt gene shares a 184 bp intergenic promoter region with the upstream divergent yoeD gene (Kunst et al., 1997). Primer extension experiments with total RNA from the stationary-phase wild-type cells incubated without L-glutamate identified the 5' end of the ggt transcript at position −34 relative to the translation initiation codon (Fig. 6, lane 1). The −35 (5'-TTGTCA-3') and −10 (5'-TTTTAC-3') sequences proceed at the corresponding sites relative to the inferred transcription initiation point. In contrast, ggt cDNA was not detected using total RNA from comP mutant cells cultured under the same conditions (Fig. 6, lane 2). Northern blots (not shown) showed that ggt was scarcely transcribed during the exponential phase (i.e. 1 day culture), but became actively transcribed after the culture entered the stationary phase (2 days incubation). The amounts of the ggt transcripts reached maximal levels after 3 days, and these were maintained for at least 1 day. Very small amounts of ggt mRNA were detected in stationary-phase cells incubated with L-glutamate, indicating that exogenous L-glutamate inhibits GGT synthesis at the level of transcription.

**Knockout of ggt promotes sporulation under nitrogen limitation**

Since *B. subtilis* NAFM5 can use both D- and L-glutamate as nitrogen sources (Kimura et al., 2004), this strain will utilize the amino acids generated from capsule degradation. In fact we detected negligible amounts of glutamate in late-stationary-phase cultures in which over 80 % γ-PGA had been degraded (Fig. 5b). These findings confirmed that the cells had internalized the resultant amino acids for metabolism (Kimura et al., 2004).

* B. subtilis cells develop spores during the nutrient-poor stationary phase (Phillips & Strauch, 2002). We assumed that in the absence of any other nitrogen source, ggt mutant cells unable to utilize capsule glutamate as a nitrogen source would sporulate more frequently than wild-type cells. To test this hypothesis, we counted spores in wild-type (strain NAFM5) and ggt mutant (NAFM90) cultures during the stationary phase. In the presence of excess NH_4Cl (e.g. 100 mM), both wild-type and mutant culture spores constituted less than 2 % of the total cells, even after 7 days of incubation (data not shown). At 10 mM NH_4Cl, the wild-type cells initiated sporulation after 5 days, and spores accounted for 25 % of the total cells after 7 days (Fig. 7). In contrast, the mutant culture began to develop spores after 2 days, and 40 % cells sporulated after 4 days, when most wild-type cells remained in a vegetative stage (Fig. 7). Limiting the carbon source did not significantly change sporulation frequencies between these strains.

**DISCUSSION**

*B. subtilis* GGT has a powerful exo-γPGA hydrolase activity. Enzymes that can hydrolyse a γ-glutamyl linkage include CapD depolymerase, γPGA hydrolase, γ-glutamyl hydrolase (EC 3.4.19.9) and glutamate carboxypeptidase II (EC 3.4.17.21). *B. anthracis* CapD depolymerase (Makino et al., 2002) and γPGA hydrolases of a fungus and *B. subtilis*...
and the results described herein (Figs 4 and 5) favour the view that γPGA is a natural substrate of *B. subtilis* GGT. In minimal medium, *B. subtilis* strains produce about 1 mg γPGA ml⁻¹. However, because of the high molecular mass of γPGA (2 × 10⁶ Da), molar concentrations of the polypeptide in the culture are around 0.5 μM. This concentration is far below the *Kₘ* value (90 μM). To degrade γPGA by GGT, *B. subtilis* must fragment the polypeptide to increase the molar concentrations of substrate to nearer the *Kₘ* value. The endo-type γPGA hydrolase inferred by this study appears to perform such fragmentation (Figs 3 and 4). Degradation of γPGA by the endo-type enzyme yields 1 × 10⁴ Da fragments at a concentration around 10 μM, which is appropriate for hydrolysis by GGT. When γPGA is completely hydrolysed by the combined action of the two hydrolytic enzymes, the total concentrations of D- and L-glutamate in the growth medium would reach about 8 mM. This concentration would represent a significant nitrogen source for *B. subtilis* in the stationary phase (Fig. 7). During *in vivo* capsule degradation, GGT levels would be properly modulated through the negative control of ggt transcription by the product L-glutamate (Fig. 3). This feedback regulation should prevent overdegradation of γPGA, and steadily supply *B. subtilis* cells with the required amounts of glutamate.

The ComQXPA quorum-sensing system plays a pivotal role in the mechanism through which *B. subtilis* adapts to nutrient starvation during the stationary phase (Lazazzera et al., 1999). This system monitors increasing cell population, and expresses an array of cellular processes, including exoenzyme production and flagellation, through which the cells cope with the nutrient shortage imposed by a dense cell population (Lazazzera et al., 1999; Phillips & Strauch, 2002). Exoenzymes enable the cells to utilize energetically less favourable polysaccharides, proteins or lipids, whereas flagella allow the cells to translocate to nutritionally favourable sites. Integration of capsule γPGA and GGT synthesis by *B. subtilis* into the regulatory circuit of the quorum-sensing system (Figs 3 and 6; Tran et al., 2000) enables them to fulfil their respective roles as an extracellular glutamate reserve and as a cognate degradation enzyme. Thus, *B. subtilis* can adapt to starvation during the stationary phase, not only by utilizing polymer nutrients in the environment or moving to other sites, but also by preserving nutrients as capsule γPGA.

The response regulator ComA of the quorum-sensing system either directly or indirectly expresses a set of genes that determine the stationary-phase-specific phenotypes (Lazazzera et al., 1999). This regulatory protein stimulates expression of the relevant genes through binding to specific sites having the consensus sequence 5'-TTGCGGNNNN-CGGCAA-3' in the promoters (Lazazzera et al., 1999). Neither the capBCD operon nor ggt has a ComA-binding site in its promoter, implying that the quorum-sensing system indirectly regulates the γPGA synthetic and degradation systems. Identification of the cascade pathways that

![Graph](image-url)

**Fig. 7.** Increase of sporulation frequency in *ggt* mutant cultures. *B. subtilis* NAFM5 (wild-type; ○) and NAFM90 (*ggt*:Spac; ●) were cultured in E9 medium containing 8% (w/v) glycerol and 10 mM NH₄Cl as carbon and nitrogen sources, respectively. Aliquots of cultures taken at days 1, 2, 3, 4, 5, 6 and 7 were divided into two portions. One aliquot was heated at 80 °C for 30 min to kill vegetative cells, and numbers of spores and total viable cells were determined on LB agar. Values are means of three determinations; SE values are below 5% of the corresponding means.
transduce the quorum-sensing signal to the regulatory machineries of the γPGA synthetic and degradation enzyme genes would provide further insight into the regulatory mechanisms of capsule γPGA, a unique extracellular reserve of glutamate.

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REFERENCES


