A multidrug resistance transporter serine protease gene is required for prestalk specialization in *Dictyostelium*

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The prestalk-specific gene, *tagB*, was disrupted by restriction enzyme-mediated integration (REMI) mutagenesis. Mutant aggregates exhibit a cell-autonomous defect in specialization of PST-A cells, a prestalk subpopulation that forms the tip and eventually forms the stalk of the fruiting body. Cooperative (non-cell-autonomous) defects were found in sporulation and in specialization of prestalk cells that eventually form the upper cup of the fruiting body (PST-O). The pattern of *ecmA::*lacZ expression in mutant *tagB*− cells defines a primary prestalk population, PST-I, from which other prestalk cells differentiate. After PST-A cells differentiate, they induce remaining PST-I cells to become PST-O cells. Subsequently, prestalk cells induce encapsulation of prespore cells during culmination. *tagB* is homologous to serine protease and to multidrug resistance (MDR) transporter genes, implying a mechanism of action that includes proteolysis and export of peptide signals. Intercellular communication via TagB may mediate integration of cellular differentiation with morphogenesis.

[Key Words: Dictyostelium, differentiation, MDR transporter genes, serine proteases, prestalk specialization]

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The initial events in cell-type divergence in *Dictyostelium discoideum* are spatially independent and can be recognized as early as 8 hr after the onset of development (for review, see Loomis 1993). Prespore cells that are marked with the *Escherichia coli* β-galactosidase (lacZ) fused to a *cotB* regulatory element (*cotB::*lacZ) are initially dispersed throughout the loose aggregate at 8 hr, but 4 hr later they sort out to the lower part of the mound. Prestalk cells that are labeled with *ecmA::*lacZ also differentiate in a spatially random manner shortly after the initiation of prespore development. At the tight aggregate stage, 12 hr after the initiation of prespore development, the proportions of the two cell types are well defined as 70% prespore cells and 30% prestalk cells. Spatial sorting is also complete at this stage and morphogenesis starts to take place as prestalk cells sort to the top of the aggregate and form a tip. The tip extends to form a finger-like structure that can fall over and migrate on the substratum (for review, see Loomis 1993). The anterior 20% of the migrating slug consists of prestalk cells that can be divided into two subpopulations on the basis of the pattern of expression of lacZ gene fusions with subdomains of the *ecmA* regulatory element. The most anterior portion of the slug is occupied by PST-A cells that will eventually give rise to the cellular stalk. The rest of the prestalk region consists of PST-O cells that will eventually give rise to the upper cup of the sorus and may participate in formation of the cellular stalk. The posterior 80% contains mainly prespore cells. The few prestalk cells found in the posterior region are called anterior-like cells and they ultimately participate in formation of the lower cup and the basal disk of the fruiting body (Jermyn et al. 1989; Williams et al. 1989).

Cell-type proportioning in *Dictyostelium* is size invariant and is dynamically maintained through intercellular communication (Loomis 1993). Slugs ranging in size from 10^3 to 10^5 cells consist of 70% prespore and 30% prestalk cells, but when prespore cells are surgically removed from a slug (Raper 1940), or when they are poisoned by expression of *cotB::*ricinA (Shaulsky and Loomis 1993), prestalk cells regulate to regenerate the proper proportion of prespore cells. A limited number of prespore cells have been shown to regulate and take on a prestalk cell fate following prolonged slug migration (Sternfeld and David 1982, Harwood et al. 1991, H. MacWilliams, pers. comm.), but poisoning prestalk cells with an *ecmA::*ricinA gene fusion does not result in conversion of prespore cells to replace the dying prestalk cells (Shaulsky and Loomis 1993). Strains carrying *ecmA::*ricinA, however, suffered from a 50% reduction in sporulation efficiency, indicating that prestalk cells have a cooperative role in sporulation (Shaulsky and Loomis 1993). We proposed that all the cells in a developing aggregate have an initial tendency to become pre-
spore cells but once a critical proportion of prespore cells is produced, the rest of the cells become prestalk. The tendency to become prespore cells is retained in the prestalk cells, and prespore cells suppress this tendency by producing and secreting an inhibitor of prespore differentiation to which they themselves are resistant [Loomis 1993; Shaulsky and Loomis 1993]. Another mechanism of intercellular communication was inferred from the expression of a *spiA::lacZ* gene fusion in sporulating cells during culmination. As the stalk tube elongates down through the prespore cell mass, a wave of *spiA::lacZ* expression propagates from the top of the sorus down [Richardson et al. 1994]. Expression of *spiA* can also be induced by addition of 8-Br-cAMP to cells dissociated at the culmination stage and shaken in buffer [Richardson et al. 1991]. It was therefore proposed that prestalk cells signal the initiation of sporulation, probably by activating the cAMP-dependent protein kinase A (PKA).

Genes that affect differentiation in *Dictyostelium* have been cloned on the basis of sequence similarity to homologous genes in other organisms, but cloning of novel genes in the cell-type divergence pathway has required a different approach (Loomis et al. 1994). Random mutagenesis by plasmid insertion allows us to screen for morphologically aberrant strains and to subsequently clone the disrupted genes. This approach, referred to as restriction enzyme-mediated integration (REMI), involves electroporation of cells with linear vector DNA carrying a selectable marker, together with a restriction endonuclease that produces ends compatible to those of the linear vector [Kuspa and Loomis 1992]. The enzyme directs integration of the plasmid into cognate restriction sites. Transformants are then visually screened for developmental defects after growth on bacteria as a source of nutrients. Mutants that are blocked at a certain stage in development can be mixed with wild-type cells to find ones that will then progress past the developmental block. These mutants are by definition cooperative (non-cell autonomous) and are suspected to have defects in intercellular communication.

Here we describe a mutant in cell-type divergence that suffered an insertion in the *tagB* locus. The gene has a unique structure, combining features of both serine proteases and of ATP-binding transporters of the multidrug resistance (MDR) family. *tagB* null cells are blocked at the tight aggregate stage of development and fail to differentiate as PST-A cells. Progression through the morphogenetic stages can be restored by cooperation with wild-type cells, but the block to PST-A development cannot be rescued. The results led us to propose a model in which a primary prestalk population (PST-I) is formed from which PST-A cells can then develop. Established PST-A cells induce remaining PST-I cells to differentiate as PST-O cells. This specialization is essential for further development of the fruiting body. The structure of the *tagB* gene implies that one of the signals emanating from PST-A cells could be a peptide that is processed by the protease part of the predicted TagB protein and secreted via the MDR homology transporter. This notion may provide a clue as to the processing and secretion of peptide hormones in other systems.

**Results**

Mutants in *tagB* were found in a screen of *BamHI–REMI* transformants as cells that develop normally until the tight aggregate stage but fail to develop further. Figure 1 compares the wild-type strain, which forms fruiting bodies after 24 hr of development, with *tagB* mutant mutants that do not develop past the tight aggregate stage.

**Insertions in the *tagB* locus result in lack of expression**

Genomic *tagB* DNA was cloned from the mutant strain AK228, and probes were made from regions flanking the insertion site. The probes were used to map the gene to chromosome 4 (position 5.2) [Kuspa and Loomis 1994] and to clone the rest of the gene from genomic DNA sublibraries as well as from a cDNA library and by inverse PCR. A map of the *tagB* region in Figure 2a shows the *tagB* gene and two additional loci that cross-hybridize with several of the *tagB* probes. These loci, *tagC* and *tagD*, are located within 20 kb of *tagB*. Probes that hybridize with *tagB* but not with the other two genes were also found (Fig. 2a). Homologous recombination with the *BgIII* fragment around IS228 was used to recapitulate the mutation in a fresh host strain and an additional allele was generated in a different host by homologous recombination with the insertion placed 1.2 kb downstream, at IS517 [Fig. 2a]. Figure 2b shows a Southern blot analysis of DNA from wild-type cells and from a *tagB* mutant [S17] with a probe that preferentially recognizes *tagB* (probe B) and a probe that recognizes *tagC* and *tagD* as well (probe A). The data indicate that the insert into IS517 disrupted the *tagB* gene but not the other two. Insertions at both IS228 and IS517 resulted in similar morphological phenotypes and were identical in most of the physiological parameters. The IS228 insertion involved a complex genetic event that consisted of a small deletion in the *tagB*-coding region and an insertion of uncharacterized DNA; therefore, all the experiments re-

![Figure 1. Morphological arrest of *tagB* mutant at the tight aggregate stage of development. Wild-type (a) and *tagB* mutant (b) cells were washed from the nutrient source and developed on filters for 36 hr.](image-url)
Figure 2. Insertional mutation in tagB results in lack of expression. (a) Map of tagB and the two closely related genes tagC and tagD. The original REMI disruption of tagB occurred at IS228. The insert into IS517 (shown to scale) was generated by homologous recombination of pGEM3 plasmid DNA that contained 2.1 kb of tagB genomic DNA around IS517 and the pyr5-6 gene as selectable marker. The predicted coding region of tagB is shown in thick boxes and the filled pattern indicates cross-hybridization between the three genes. Boxes at the bottom represent a unique probe [probe B, empty box] and a redundant probe [probe A, full box]. Restriction sites: (B) BamHI; (B) BglII; (B) BglII; (C) ClaI; (E) EcoRI. (b) Genomic DNA from wild-type and from tagBmutant cells was digested with ClaI and resolved by electrophoresis through a 0.8% agarose gel. Southern blots were hybridized with the redundant probe A (see Fig. 2a) to visualize the three genes. The blots were later stripped and reprobed with probe B (see Fig. 2a) to visualize tagB. Locus names are indicated on the left; size standards are indicated on the right (kilobases). (c) Polyadenylated RNA was prepared from wild-type and from tagBmutant cells at different stages of development as indicated. RNA was resolved by electrophoresis through a 1.0% agarose gel containing formaldehyde. Northern blots were hybridized with probe B (see Fig. 2a). Size standards are indicated on the right (kilobases). (d) Total protein was prepared from vegetative cells (V) or from wild-type migrating slugs (wt) and from tagBmutant aggregates after 48 hr of development on agar plates (D). Protein (10 μg/lane) was resolved by electrophoresis through a 6% polyacrylamide gel containing SDS and electrotransferred to a nitrocellulose membrane. Western blots were reacted with an anti-peptide antibody for TagB and with a secondary, alkaline-phosphatase-conjugated antibody and developed with NBT and BCIP. The size marker is indicated on the left (kilodaltons).

Probes specific to tagB were used to determine the size and the developmental regulation of the mRNA. Figure 2c shows Northern blots of RNA prepared from wild-type and tagB mutant cells at various stages of development. mRNA of 6.5–7.0 kb was first observed after 8 hr of starvation and continued to accumulate throughout development in the wild-type strain, but was not detected in the tagB− mutant [Fig. 2c]. An antibody was raised against a synthetic peptide from the predicted amino-acid sequence of one of the regions in tagB that did not cross-hybridize with tagC and tagD. The anti-peptide antibody revealed a protein of about 210 kD in Western blot analysis of wild-type cells but not in tagB− mutants [Fig. 2d]. These results show that the insertion into IS517 resulted in lack of tagB expression at the mRNA and at the protein level.

Mutants in tagB respond to intercellular signals but fail to produce DIF-1

The onset of tagB expression occurs concomitantly with the beginning of cell-type divergence at the loose aggregate stage. The morphological arrest in the mutant is at the tight aggregate stage, just before tip formation by prestalk cells. To understand the role of tagB in cell-type divergence and tip formation, we examined the effects of the mutation on expression of the prestalk-specific gene ecmA and the prespore-specific gene cotB. The data in
Figure 3a show that whereas the $tagB$ mutation did not alter $cotB$ expression, accumulation of $ecmA$ mRNA was markedly reduced in the mutant. An apparent decrease in $cotB$ and $ecmA$ mRNA levels can be observed in the 24-hr samples from wild-type cells but not in the mutants [Fig. 3a]. This is because of reduced efficiency of RNA extraction from encapsulated spores and stalk cells in wild-type cells [Richardson and Loomis 1992; Shaulsky and Loomis 1993] and reflects the observation that $tagB^-$ mutants do not form spores and stalks. The continued accumulation of $cotB$ and $ecmA$ RNA in the mutant cells shows that, in spite of the morphological arrest at the tight aggregate stage, $tagB^-$ mutants continue to progress along some developmental pathways.

The reduced accumulation of $ecmA$ RNA in the mutants suggested that prestalk development was impaired. Because $ecmA$ expression can be induced by the natural morphogen DIF-1, we tested the possibility that $tagB$ was involved in the ability of developing cells to respond to the morphogen. Addition of DIF-1 to wild-type cells in suspension, in the presence of $Ca^{2+}$ ions and cAMP, results in increased accumulation of $ecmA$ and $ecmB$ RNA and decreased levels of the prespore marker D19 [Berks and Kay 1990]. Fosnaugh and Loomis [1991] showed that accumulation of $cotB$ mRNA was increased in the presence of cAMP and $Ca^{2+}$ ions and decreased following addition of DIF-1. The data in Figure 3b show that $tagB^-$ mutants were able to respond normally to the addition of DIF-1 in the presence of cAMP and $Ca^{2+}$ ions by accumulating higher levels of $ecmA$ mRNA and reduced levels of $cotB$ mRNA. These experiments also revealed that the mutant cells respond normally to cAMP and to $Ca^{2+}$ ions by the accumulation of $cotB$ mRNA. DIF-1 breakdown in the $tagB^-$ mutants was also found to be unaltered relative to the wild-type strain (R.R. Kay, pers. comm.).

To quantify the effects of the mutation on development, we generated $tagB^-$ mutant strains that carry the $lacZ$ gene under control of various cell-type-specific regulatory elements. Total enzyme activity was used to es-

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**Figure 3.** Response of $tagB^-$ mutants to extracellular morphogenic signals. (a) Northern blot analysis of total RNA prepared from wild-type and from $tagB^-$ mutant cells at different stages of development as indicated. Blots were hybridized with a probe for $ecmA$, stripped, and reprobed for $cotB$ as indicated. (b) $tagB^-$ mutant cells were developed on filters for 12 hr, collected, disassembled, and resuspended in buffer. Cells were shaken in suspension with supplements as indicated and samples were collected 2 and 4 hr later [14 hr, 16 hr, respectively]. Total RNA was prepared from the samples and from cells that were developed on filters without supplements. Northern blots were prepared and hybridized with probes for $ecmA$ and $cotB$. Supplements: (cAMP) 20 μM/hr; (Ca$^{2+}$) 0.2 mM CaCl$_2$; (DIF) 50 nM DIF-1/hr. (c) Cells were mixed at 1:1 ratios and developed on filters. Samples were collected at the indicated time points and resuspended in buffer containing 1% Triton X-100. Cell-free extracts were incubated with ONPG, and β-galactosidase activity was measured by absorbance at 420 nm. Results were normalized to protein concentration and to reaction time. (d) $tagB^-$ $ecmA::lacZ$ mutant cells mixed with $tagB^-$ mutant cells without the reporter gene, (●) wild-type $ecmA::lacZ$ cells mixed with wild-type cells without the reporter gene, (□) $tagB^-$ $ecmA::lacZ$ mutant cells mixed with wild-type cells without the reporter gene. (d) $tagB^-$ $ecmA::lacZ$ cells were developed on filters or in suspension with supplements as in b. β-Galactosidase activity was measured as in c. (Solid bars) 12-hr cells; (stippled bars) 14-hr cells; (hatched bars) 16-hr cells.
timate the effects of extracellular signals on specific gene expression. The results in Figure 3c show that the enzyme activity detected in the tagB− ecmA::lacZ reporter strain [open boxes] was consistent with the expression pattern of the endogenous ecmA mRNA, and was about fourfold reduced compared to the parental wild-type strain [circles]. Mixing tagB− ecmA::lacZ cells with unlabeled wild-type cells resulted in normal overall morphology during filter development [see below] and increased the enzyme activity in the mutant strain back to the wild-type level of expression [Fig. 3c, solid boxes]. Addition of DIF-1 in the presence of cAMP and Ca2++ to shaking suspensions of the tagB− ecmA::lacZ cells induced the enzyme activity about fivefold as well [Fig. 3d], indicating that the increased level of ecmA expression in the mixed aggregate may be fully attributed to DIF-1 production by the wild-type cells. The effect of DIF-1 on the increase in ecmA::lacZ expression was due in part to a twofold increase in the number of cells stained by X-gal [data not shown]. It is impossible to determine whether the increased number of positively stained cells was attributable to recruitment of undifferentiated cells to the prestalk pathway, or to increase of enzyme activity in already existing prestalk cells above the threshold of detection. On the basis of the results presented in Figure 3 we conclude that tagB− mutants have developed the ability to respond to cAMP, Ca2++ and DIF-1, indicating that the response to extracellular signals is not likely to be the primary defect induced by the mutation.

The possibility that tagB− mutants were defective in production of DIF-1 was also tested. In wild-type cells, DIF-1 starts to accumulate at the tight aggregate stage and maximal levels are observed at the finger stage, ~15–20 hr after the initiation of development, whereas vegetative cells contain no measurable amounts of DIF-1 [Brookman et al. 1987]. Hexane extracts were therefore prepared from wild-type cells and several tagB− cell lines at the vegetative stage and after 17 hr of development. The extracts were tested for their ability to induce vacuolization of V12M2 cells in submerged cultures [Brookman et al. 1987]. The results were normalized to the number of cells extracted and compared to the effect of adding 200 nM pure synthetic DIF-1 to the test plates. Table 1 shows that extracts from 1.4×10⁷ developing wild-type cells contained the equivalent of 100 nm DIF-1, but extracts prepared from the same number of tagB− mutant cells contained no measurable amounts of DIF-1.

The results in Figure 3 and Table 1 indicate that tagB− cells have the ability to respond to DIF-1 but fail to produce it. Wild-type cells were able to rescue the effects of the tagB− mutation on general morphology and on ecmA expression, presumably by secreting an exogenous factor that tagB− mutants fail to produce or secrete themselves. If DIF-1 was that factor, it would be expected to rescue the morphological defect in tagB− mutants. To test that possibility, tagB− mutant cells were developed on filters and DIF-1 was added at a range of concentrations from 50 to 200 nm at various time points after starvation or in 50 nm doses at 2-hr intervals starting from 8 hr of development. Added DIF-1 caused an increase in ecmA::lacZ expression, but none of the conditions was sufficient to induce sporulation or any significant changes in the overall morphology [data not shown]. We conclude that the defect in DIF-1 production in the mutant does not account for the developmental arrest and is probably due to a dependent sequence [epistasis] effect rather than a direct consequence of the tagB gene disruption.

**Table 1. DIF-1 synthesis in wild-type and tagB− mutants**

| Hexane extract (1.4×10⁷ cells) | Percent vacuolization×a |
|-------------------------------|------------------------|
| Wild type, vegetative         | 0                      |
| Wild type, 17-hr development  | 11 ± 2                 |
| tagB− mutant, vegetative      | 0                      |
| tagB− mutant, 17-hr development | 0.2 ± 0.3b           |
| 200 nM purified DIF-1         | 26 ± 4                 |

*aAverage ± standard deviation of six independent tagB− mutant strains.

Failure of tagB− mutants to develop as PST-A cells reveals a new type of prestalk cells

The same lacZ reporter strains were used to follow the distribution of all prestalk cells by X-gal staining of pure populations of mutant cells or in mixed populations with equal numbers of wild-type cells. Figure 4a shows that prestalk cells were sorted to the upper part of the tight aggregates in pure tagB− mutant populations, a normal pattern for the tight aggregate stage at which development was arrested. When mixed with wild-type cells, tagB− mutants were able to cooperate and participate in formation of a mature fruiting body [Fig. 4b], but a significant difference from the wild-type prestalk pattern was observed in such mixed populations. The mutant prestalk cells were not found in the tip of the anterior region of the finger stage [Fig. 4b] and were also absent from the stalk of the mature fruiting body [Fig. 4c], indicating a defect in differentiation of PST-A cells. Two possible explanations could account for the lack of staining in the tip and in the stalk of the mixed structure: Either the tagB− mutant cells were physically excluded from the tip and from the stalk, or they were present in the tip but did not express the ecmA::lacZ gene. To distinguish between the two possibilities we constructed a tagB− mutant strain expressing actin 15 promoter. Figure 4d shows that all of the cells were stained in a pure population of tagB− mutants carrying act15::lacZ. In the mixed population, X-gal-stained cells were absent from the tip [Fig. 4e] and from the stalk [Fig. 4f], indicating that the mutant cells were excluded from those regions.

Mutant tagB− cells appear to have a cell-autonomous defect in PST-A differentiation. It was interesting to know whether the other prestalk subpopulation, namely
Figure 4. Prestalk specialization of tagB− mutants. lacZ-expressing cells were developed on filters for 16 and 24 hr in pure populations or mixed at 1:1 ratios with wild-type cells without the reporter gene. tagB− ecmA::lacZ cells were developed as a pure population for 24 hr [a] or mixed with wild-type cells for 16 [b] and 24 hr [c]. tagB− act15::lacZ cells were developed as a pure population for 24 hr [d] or mixed with wild-type cells for 16 hr [e] and 24 hr [f]. tagB− pstO::lacZ cells were developed as a pure population for 24 hr [g] or mixed with wild-type cells for 16 hr [h] and 24 hr [i]. Wild-type pstO::lacZ cells were developed as a pure population for 16 hr [j] and 24 hr [k]. Samples were fixed, permeabilized, and stained with X-gal. lacZ-Expressing cells are blue.

PST-O, was also affected. In wild-type aggregates pstO::lacZ staining is apparent in scattered cells throughout the mound (Early et al. 1993), but in tagB− mutants we observed no staining until 20–24 hr after the initiation of development. Figure 4g shows that in pure populations of tagB− mutant cells carrying pstO::lacZ, most aggregates contained no stained cells whereas a few had one or two. When mixed with wild-type cells, the staining pattern of the mutant tagB− pstO::lacZ after 16 and 24 hr of development (Fig. 4h and i, respectively) was indistinguishable from that of the wild-type pstO::lacZ strain (Fig. 4) and k, respectively). Because PST-O cells were normally distributed in the mixed aggregates but were absent in mutant aggregates, we conclude that PST-O development is dependent on tagB in a cooperative (non-cell-autonomous) manner and is probably dependent on the presence of PST-A cells.

By use of the ecmA::lacZ marker we found that tagB− mutants are able to differentiate and sort as pre-stalk cells (Fig. 4a) but fail to develop as PST-A cells (Fig. 4b,c) or as PST-O cells (Fig. 4g). This observation indicates the existence of a previously unrecognized type of pre-stalk cells from which the two other types may develop. We propose the name PST-I to describe these cells.

The primary defect in pre-stalk differentiation of tagB− mutants is cell autonomous and is consistent with a lack of the PST-A subpopulation. We tested RNA preparations enriched for prespore and pre-stalk cells and found that the cell-type specificity of tagB RNA expression is supportive of that notion. Cell-type-specific RNA was prepared by centrifugation of dissociated wild-type cells prepared from slugs and extraction of RNA from the pre-stalk and prespore fractions. Figure 5a shows a Northern blot of RNA from the fractions, hybridized to probes for ecmA and for cotB as cell-type-specific markers to estimate the degree of purification. The tagB-specific probe in Figure 5b shows that the mRNA is enriched in the pre-stalk fraction as much as the ecmA marker.

tagB contains sequences that are homologous to serine protease and to MDR transporter genes

Additional insight into the possible mechanism of action of the tagB gene in development was gained from the primary sequence of the predicted protein shown in Figure 6a. The tagB gene contains an open reading frame (ORF) of 5715 bp without any introns and is separated from the tagC gene by about 1 kb (see map in Fig. 2). The
upstream region and ORF were cloned into a plasmid carrying a G418-resistance marker. Upon transfection of the plasmid into tagB− mutants, G418− colonies developed that regained the ability to form normal fruiting bodies [data not shown]. This result indicates that the sequenced region is sufficient to complement the tagB− mutation and can therefore account for the complete sequence of the gene. The calculated molecular weight of the TagB protein is 210 kD, consistent with the measured size found by Western blot analysis [Fig. 2d]. Comparison of the TagB sequence against several protein data bases using the BLAST algorithm [Altschul et al. 1990] revealed two major domains of homology to distinct gene families. In the amino-terminal half of TagB, we found sequence similarity to the essential components of the active site in a family of serine proteases [Fig. 6b]. The similarities are in the predicted charge relay system that includes D387, H432, and the active serine S695, as well as N538 in the predicted orientation determination pocket [Kraut 1977; Vos et al. 1989]. In the carboxy-terminal half of TagB we found extensive similarity to proteins from the MDR family of ATP-dependent transporters [for review, see Ames 1986; Pastan and Gottesman 1991]. Sequence similarity was found in the bipartite ATP-binding site [Fig. 6b] and in the six transmembrane domain [data not shown] as well as in the overall hydrophobicity pattern of TagB that indicated a plausible six transmembrane domain between amino acids 1000−1400 [Fig. 6c]. A hydrophobic region that could serve as a signal sequence for membrane insertion was found in the amino terminus of TagB. Our search of the data base did not reveal other examples of proteins carrying a fusion of these two regions. On the basis of the homology to serine proteases and to ATP-binding transporters, we suggest that TagB could be a membrane protein involved in proteolytic processing and transport of a peptide.

tagB allows cooperation of prestalk and prespore cells during culmination

Microscopic examination of dissociated cells from tagB− aggregates revealed no spores or stalk cells [data not shown]. The lack of spores is demonstrated in Table 2 where detergent treatment of developed tagB− cells resulted in a complete loss of viability, showing that <1 in 5×107 mutant cells had sporulated. Interestingly, tagB− cells can develop into spores in mixed aggregates with wild-type cells [Table 2] indicating that, like the PST-O defect, the sporulation deficiency is cooperative [non-cell autonomous]. This situation is similar to another prestalk specific defect, namely that which results from over expression of ecmA::Rm. In the latter, a mutated regulatory subunit [Rm] of the cAMP-dependent protein kinase A is expressed under the ecmA promoter such that prestalk cells are expected to have reduced levels of PKA activity, whereas prespore cells are not affected directly [Harwood et al. 1992]. ecmA::Rm cells develop normally until the finger stage, at which they arrest. Spores and stalks are not produced when these cells are developed as pure populations, but sporulation of the mutant cells can be induced by cooperation with wild-type cells [Table 2] as in the case of tagB− mutant cells. In the mixed aggregates, ecmA::Rm cells fail to enter the PST-A region and do not participate in stalk formation. We mixed tagB− mutants with ecmA::Rm cells and examined their ability to form spores. The cells in such experiments formed mixed aggregates and developed to the finger stage, but neither of them sporulated, despite their ability to sporulate in the presence of wild-type cells [Table 2]. This experiment indicates that both tagB− and ecmA::Rm cells are deficient in the same pathway that leads to differentiation of PST-A cells, which are subsequently required for development of PST-O cells as well as for sporulation.

Discussion

The primary defect in tagB− mutants appears to be a complete lack of differentiation of PST-A cells that cannot be overcome even by the presence of wild-type cells in mixed aggregates. This cell-autonomous phenotype may result from a failure to remove an endogenous inhibitor produced in early prestalk differentiation. The serine protease homology domain of TagB suggests that it may be responsible for proteolytic inactivation of an inhibitor protein, whereas the MDR transporter homology suggests that it may remove an inhibitor by secreting it from the cell. In fact, TagB may carry out both functions by generating peptides and then transporting them out of the cells. This raises the possibility that the se-

Figure 5. Cell-type specificity of tagB gene expression. Wild-type cells were developed on agar plates for 48 hr and slugs were collected and dissaggregated. Cells were harvested through Percoll gradients, and two fractions were collected separately. The lower fraction [heavy (H)] is enriched with prespore cells and the upper fraction [light (L)] is enriched with prestalk cells. Northern blots were made with total RNA from the respective fractions, ecmA and cotB probes were used to determine the level of enrichment of prestalk and prespore cells, respectively [a], and probe B [Fig. 2a] was used to detect tagB RNA [b] as indicated.

Prestalk specialization in Dictyostelium
Figure 6. Sequence analysis of the tagB gene. (a) The complete amino acid sequence was deduced from the nucleotide sequence of cDNA and genomic DNA clones (GenBank accession no. U20432). Underlined sequences indicate regions of homology to serine protease and to ATP-dependent transporters of the MDR family. (See details below.) (b) Comparison of TagB predicted amino acid sequence to the serine protease proteinase K (endopeptidase K precursor from Trypanosoma cruzi) and to ATP-dependent transporters of the MDR family, from Chinese hamster, Cricetulus griseus, accession no. A27126 in PIR data base) and to the ATP-dependent transporter MDR 1 protein (MDR1) from Chinese hamster, Cricetulus griseus, accession no. A27126 in PIR data base). Underlined sequences indicate regions of homology to serine protease and to ATP-dependent transporters of the MDR family. (c) Comparison of TagB nucleotide sequence of cDNA and genomic DNA clones (GenBank accession no. U20432). Underlined sequences are essential for activity of the respective proteins. Numbers above the box indicate position in the complete amino acid sequence was deduced from the nucleotide sequence. Numbers on the left indicate relative hydrophobicity. Note the hydrophobic region at the amino-terminus indicating a predicted six transmembrane domain.

creted peptides could function as signaling molecules, controlling the differentiation of other cell types in the aggregate.

Allopathic systems that combine proteolytic processing and membrane transport of the peptide products have been described in other organisms. In Saccharomyces cerevisiae, the mating pheromone α-factor is synthesized as a 36- or 38-amino-acid precursor form the MFA1 and MFA2 genes, respectively (for review, see Michaelis 1993). Following proteolytic processing by yet-undefined proteases, isoprenylation by the gene products of RAM1 and RAM2, and methylation by the STE14 gene product, the mature pheromone is transported across the plasma membrane and out of the cell via the STE6 gene product that is a member of the MDR gene family (Kuchler et al. 1989; McGarth and Varshavsky 1989). In mammalian cells, foreign proteins are fragmented by proteolytic activities in the cytoplasm, and the products are transported into a pre-Golgi compartment via a heterodimeric

Table 2. Cooperative sporulation

| Strains (%) | Mutant spores [% of total] |
|-------------|---------------------------|
| Wild type   | tagB          | mutant |
| 0           | 100           | 0      |
| 10          | 90            | 10     |
| 50          | 50            | 0      |

Note: Wild type refers to the reference strain, while ecmA:Rm indicates a mutation in the ecmA gene.
membrane transporter of the MDR family that is composed of the TAP1 and TAP2 gene products (Townsend et al. 1985; Deverson et al. 1990; Monaco et al. 1990; Spies et al. 1990; Trowsdale et al. 1990). The peptides are then associated with major histocompatibility complex (MHC) class I proteins and presented on the cell surface. In Dictyostelium, the two functions are linked because in TagB the protease and the transporter homology domains are encoded by the same gene.

MDR genes are internally duplicated such that each gene product consists of two sets of a highly conserved bipartite ATP-binding domain and two sets of a less-conserved putative six transmembrane domain [Chen et al. 1986]. The MHC transporters TAP1 and TAP2 are not internally duplicated, but the proteins associate in the membrane into a functional dimer (Spies et al. 1992). Likewise, tagB encodes a single putative six transmembrane domain and a single bipartite ATP-binding domain, indicating that it may also function as a dimer. TagB may form homodimers, or heterodimers with the proteins encoded by the cohybridizing genes tagC and tagD. We have partially sequenced the tagC gene and find that the putative nucleotide binding domains of tagB and tagC are ~90% identical and that the putative transmembrane domains are ~60% identical [G. Shaulsky and W.F. Loomis, unpubl.]. Thus, tagC may function as a transporter protein as well. Indeed, insertion inactivation of the tagC gene results in a phenotype that is almost identical to that of tagB− mutants [G. Shaulsky and W.F. Loomis, unpubl.]. Although both tagB− and tagC− cells can differentiate into spores when they develop in mixed aggregates with wild-type cells, they do not form spores when mixed with each other in the absence of wild-type cells. Thus, both TagB and TagC appear to be required to generate the signal necessary for sporulation, as would be expected if TagB and TagC function in a heterodimer.

We found that in addition to the cell-autonomous lack of PST-A cell specialization, tagB− mutants exhibit defects that can be overcome by the presence of wild-type cells. Development of mixed populations showed that both sporulation and PST-O specialization of the tagB− mutant cells were restored in aggregates containing wild-type cells. Therefore, wild-type cells produce signals that tagB− mutants can receive and respond to. These signals could be peptides that are produced and secreted by different dimers of TagB, TagC, and TagD in the wild-type PST-A cells, each combination producing a different signal. The cooperative failure of ecmA::lacZ cells to sporulate and their inability to cooperate with tagB− mutants indicates that the signaling pathway involves the function of PKA in PST-A cells.

Prestalk differentiation of tagB− mutant cells was apparently normal during early development up to the time of developmental arrest. Induction of ecmA expression occurred at the normal time and mutant prestalk cells sorted to the top of the aggregate as would wild-type cells at that stage. In wild-type cells, there are several subpopulations of prestalk cells, including PST-A cells that are defined by localization to the most anterior part of the prestalk zone [Jermyn et al. 1989, Williams et al. 1989] and PST-O cells that are defined by activation of specific regions in the ecmA promoter [Early et al. 1993]. In pure populations of tagB− mutant cells, PST-O cell differentiation failed to occur. Therefore, PST-O cells could not account for the presence of ecmA::lacZ expressing cells in the developing tagB− mutants. Because tagB− mutants do not form PST-A cells either, we conclude that tagB defines an initial population of prestalk cells, namely PST-I, from which both PST-A and PST-O cells specialize. Because tagB− mutant cells do not accumulate DIF-I, PST-I cells can be defined as ecmA-expressing cells that develop independently of the prestalk morphogen but can respond to it by accumulating increased levels of ecmA RNA. PST-I cells are also sensitive to the effects of cAMP and calcium ions. Because PST-O specialization was cooperatively induced by wild-type cells, we propose that in normal prestalk differentiation, PST-O specialization is dependent on the presence of PST-A cells. We propose that following accumulation of the tagB gene product, some of the PST-I cells specialize and become PST-A cells that can produce a signal that induces yet unspecialized PST-I cells to become PST-O cells (Fig. 7). This model could account for proportioning of these prestalk cell types by a mechanism of lateral inhibition.

Following aggregate formation, prestalk cells sort to the top where a tip is formed. From the phenotype of tagB− mutants we learn that tip formation is dependent on the presence of PST-A cells, PST-O cells, or both. On the basis of the spatial localization of the cell types in slugs, PST-A cells are probably responsible for the tip, whereas PST-O cells define the base of the tip. PST-A cells subsequently form the stalk in culminants [Jermyn et al. 1989, Williams et al. 1989], so it is not surprising that tagB− mutants do not differentiate into stalk cells even when developed in mixed aggregates. However, it is not obvious why tagB− mutants developing in pure populations fail to sporulate because prespore cells appear to differentiate normally and they are fully capable of forming spores when developed in the presence of wild-type cells. It appears that spore maturation is gated by specialization of prestalk cells and resulting morphogenesis.

Figure 7. The involvement of tagB in prestalk specialization and in cell–cell communication. [Plain arrows] Cell lineage; [stippled arrows] intercellular communication [see text for details].
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Because TagB is essential for triggering encapsulation, peptides generated by its serine protease domain or those of the closely related proteases encoded by the linked genes, tagC and tagD, may be used to signal prespore cells to encapsulate. Such peptide signals could be released from prespore cells by the MDR transporter domains. A signal emanating from prespore cells at the tip of the stalk cells and surrounding tissue has been previously recognized from the pattern of expression of a late prespore gene, spiA (Richardson et al. 1994). This gene is initially expressed at the boundary of prestalk and prespore cells and subsequently in more distal cells over a period of several hours just prior to encapsulation. Our data identify the source of the signal as PST-A cells and suggest that the signal is a peptide that is processed and secreted by the TagB protein. Proteolytic processing and secretion of peptides could be a mechanism that integrates morphogenesis and differentiation via communication between cells and tissues during development in Dictostelium and in other organisms.

**Materials and methods**

**Cells, growth, transformation and development**

*Dictostelium discoideum* V12M2 cells were grown on SM agar plates (Sussman 1987) in association with *Klebsiella aerogenes*. AX4 and its pyrS-6 derivative HL328 were grown in HL5 medium (Sussman 1987) supplemented with 20 μg/ml of uracil. G418-resistant strains TL43 (pyrS-6 actS-15::lacZ), TL51 (pyrS-6 ecmA::lacZ), TL52 (ecmA::Rm), TL68 (AX2 pstO::lacZ), a kind gift from Dr. J. Williams, University College, London, UK), TL88 (AX2 pyrS-6 pstO::lacZ), and the respective tagB pyrS-6 derivatives were grown in HL5 medium supplemented with 5 μg/ml of geneticin. Restriction enzyme-mediated integration, disruption of tagB by homologous recombination, and selection for uracil auxotrophs in FM medium were carried out as described by Kuspa and Loomis (1992). Transformation and G418 selection, development on filters, migration on agar plates, mixed development, detergent treatment of spores and sporulation assays were carried out as described by Shaulsky and Loomis (1993).

**Staining, enzyme activity, and bioassays**

X-Gal staining was performed as before (Insall et al. 1994). β-Galactosidase was assayed as described by Fosnaugh and Loomis (1993). In mixing experiments, 50% lacZ transformants were mixed with 50% of isogenic cells that did not carry the lacZ gene as a control for the dilution effect. Comparison of tagB to wild-type cells was always to the parental tagB+ cell line. Response of tagB+ mutants to DIF-1, CAMP, and Ca2+ ions was determined as described by Fosnaugh and Loomis (1993), with the exception that cells were dissociated after 12 hr of filter development and that DIF-1 was added at 50 nm/hr, CAMP was added at 20 μM/hr and samples were collected every 2 hr. Note that Ca2+ ions were added to selected samples rather than to the buffer of all samples as in Berks and Kay (1990). Hexane extraction of DIF-1 from AX4 cells and from tagB+ mutants, as well as the DIF-1 bioassay on V12M2 cells was performed according to Brookman et al. (1987 and references therein).

**Molecular cloning of tagB**

Cloning of flanking DNA from the REMI strain AK228 cut with EcoRI was performed as described (Kuspa and Loomis 1992), and the entire insert was sequenced. The plasmid (p228Ec) was digested with BamHI and BgIII and a 1.2-kb fragment was gel purified and used as a probe to clone DNA from the rest of the gene. For inverse PCR, AX4 genomic DNA was digested with BgIII and size selected to 1.4–2.8 kb by agarose gel electrophoresis and reelectrophoresion. DNA was diluted, ligated to promote circularization, and amplified with oligonucleotides from the sequenced portion of the gene. The PCR product (1 kb) was cloned into pGEM3 (Promega) and sequenced. A cDNA library was prepared (Stratagene) from polycloned RNA extracted from AX4 cells that have developed on filters for 10 hr mixed with poly[A]-selected RNA extracted from AX4 slugs that had migrated for 45 hr. The library was screened with probes from the genomic DNA clones, and positive clones were sequenced. AX4 DNA was digested with Clal, size selected by gel electrophoresis to 5–7 kb, digested with BamHI, and size selected again to 2.5–3.5 kb. DNA was ligated with AccI–BamHI-digested pGEM3 and transformed intoSURE cells (Stratagene). Colonies were screened with probes from the 5’ end cDNA clones and positive clones were sequenced. A similar approach was used to clone an EcoRI–Clal fragment between tagB and tagC and genomic DNA and cDNA from tagC (see map in Fig. 2). Both DNA strands from all of the clones were sequenced by use of the Sequenase system (U.S. Biochemical). The expression vector for tagB was made by combining overlapping genomic clones from the EcoRI site in tagC to the BgIII site at the 3’ end of the inverse PCR product and mapping tagB into an XhoI–Xbal-digested pDdGal17 (Harwood and Drury 1990), deleting the lacZ sequences but retaining the Neo-resistance cassette, such that the tagB sequence was followed by a transcriptional termination signal from actin 8.

**Nucleic acid analysis**

Genomic DNA was prepared and analyzed on Southern blots and RNA was prepared and analyzed on Northern blots as described (Shaulsky and Loomis 1993). Separation of cell types on Percoll (Pharmacia) gradients was performed according to Ratner and Borth (1983). RNA was extracted from the respective fractions and analyzed as above. Selection of polyadenylated RNA from total RNA was performed on 100 μg of total RNA aliquots with oligo(dT)-cellulose (batch affinity chromatography) according to Sambrook et al. (1989). Probes for ecmA and cotB were as described (Shaulsky and Loomis 1993). Probe A for tagB (Fig. 2a) was prepared by cloning an EcoRI–BglII DNA fragment 3’ of the BglII site in tagB into pGEM3 (Promega), digestion with Sau3AI and transcription in vitro with T7 RNA polymerase [BRL] in the presence of [32P]CTP to generate a radioactive 150-bp long transcript. Probe B (Fig. 2a) was prepared from an EcoRI–BamHI DNA fragment 5’ of IS228.

**Protein analysis**

The synthetic peptide KARSNGENSTDQAQDGSLP represents amino acid 666–683 of TagB, with an added lysine (K) at the amino terminus and a substitution of cysteine (C) with an alanine (A) at amino acid 13 of the peptide. The peptide was synthesized at the peptide synthesis facility, Center of Molecular Genetics at University of California, San Diego, an aliquot was conjugated to PPD [Statens Seruminstitute, Copenhagen, Denmark] according to Lachmann et al. (1986) and antibodies were...
raised in rabbits by Scantibodies Lab. Inc. (Ramona, CA). IgG was purified from crude antisera by affinity chromatography on protein A–cellulose (Sigma). Cells were collected at different stages of development, resuspended in 50 mM Tris-HCl (pH 8.0), 5 mM EDTA, 150 mM NaCl, 0.5% Nonidet-40, 1 mM PMSF, and protein concentration was determined using Bio-Rad protein assay reagent. Equal amounts of protein were resolved on 6% polyacrylamide–SDS gels (Laemmli 1970), electrotransferred to nitrocellulose membranes and detected with 2 μg/ml of the purified IgG antibody against TagB followed by incubation with alkaline-phosphatase-conjugated goat anti-rabbit antibody and color development according to the ProteoBlot [Promega] protocol.

Computer sequence tools

Predicted amino acid sequence and hydropathy plot (Kyte and Doolittle 1982) were performed with DNA Strider I.2 and with MacVector 3.5 [IBI]. Protein data base searches were carried out with BLAST [Altschul et al. 1990].

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