Fast lysis of *Escherichia coli* filament cells requires differentiation of potential division sites

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**INTRODUCTION**

The peptidoglycan (murein) layer of the bacterial cell envelope forms a giant, bag-shaped macromolecule, the sacculus, which surrounds the cell. The sacculus is the principal stress-bearing element of the cell wall and plays an essential role in bacterial morphogenesis (Ayala et al., 1994; Höltje, 1998; Nanninga, 1998). Metabolism of the sacculus requires a large number of enzymes and metabolites and is subjected to complex regulatory pathways (Glauner & Höltje, 1990; Höltje, 1998; Höltje & Glauner, 1990; Nanninga, 1991). The physical integrity of the sacculus is of vital importance to the cell because it is the only structure able to withstand the turgor pressure of the cell. Therefore, growth and septation of the sacculus must proceed avoiding the generation of discontinuities to prevent cell lysis (Koch, 2000).

In *Escherichia coli* the mode of insertion of new precursors into the sacculus apparently shifts from diffuse to zonal at the initiation of septation (de Pedro et al., 1997). Lateral wall elongation is accomplished by intercalation of new and pre-existing murein all over the cylindrical surface of the sacculus. At the time of septation an area of highly localized murein synthesis develops into the potential division site and remains active until septum completion (SMS, septal murein synthesis). Once the dividing sacculus is split, SMS stops and the new cell wall poles become metabolically inert. Activation of SMS requires the activity of FtsZ, but not the essential division proteins FtsA, FtsQ and PBP3 (penicillin-binding protein 3). Inactivation of FtsZ leads to the generation of filamentous cells in which murein precursors are inserted in a diffuse way for as long as the filament elongates. In contrast, cells from *ftsA*, *ftsQ* or *ftsI* mutants growing at the restrictive temperature are able to trigger SMS in due time. However, SMS remains active only for a defined period of time. Then it is switched off and murein synthesis reverts to a diffuse insertion mode. When the time for the next round of...

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**Periodic activation of zonal peptidoglycan (murein) synthesis at division sites in *Escherichia coli* has been reported recently. Zonal synthesis is responsible for septum formation, whereas elongation of the cell sacculus is performed by diffuse insertion of precursors. Zonal synthesis can be triggered in *ftsA*, *ftsQ* and *ftsI* (*pbpB*) division mutants growing as filaments at the restrictive temperature, but not in *ftsZ* mutant strains. The lytic response to β-lactams of cells able or unable to periodically trigger a zonal mode of murein synthesis could be substantially different. Therefore, we investigated the response to the bacteriolytic β-lactam cefsulodin of *ftsZ* and *ftsI* mutants growing at the restrictive (42°C) temperature. The *ftsI* cells lysed early and quickly after addition of the antibiotic. Sacculi of lysed cells were transversely cut in a very sharp way. In contrast the *ftsZ* strain lysed late and slowly after addition of the antibiotic and sacculi showed a generalized weakening of the murein network and extended breaks with a frayed appearance. No transversely cut sacculi comparable to those seen in the *ftsI* samples were found. Our results strongly support that β-lactam-induced lysis occurs preferentially at division sites because of the activation of zonal murein synthesis at the initiation of septation.**

**Keywords:** *ftsZ*, *pbpB*, penicillin-binding protein, beta-lactam, murein
Growth of the bacterial cell sacculus is thought to require the concerted action of biosynthetic and hydrolytic enzymes (Höltje, 1996, 1998; Höltje & Heidrich, 2001; Weidel & Pelzer, 1964). The former are essential enzymes which mediate elongation of nascent peptidoglycan strands as well as their insertion into the sacculus. An essential role for murein hydrolases has been repeatedly postulated on theoretical grounds. As septation itself needs the chemical energy present in pre-existing peptidoglycan bridges. Such enzymes would cleave bonds in the sacculus concomitantly with the insertion of new murein.

Induction of bacterial lysis by β-lactams and other murein synthesis inhibitors normally requires active growth of the cells and the activity of murein hydrolases (autolysins) (Tomasz, 1974; Tuomanen & Tomasz, 1990; Leduc & van Heijenoort, 1980; Leduc et al., 1982). These observations have led to models proposing that lysis occurs because the inhibitors break the balance between biosynthetic and hydrolytic enzymes, favouring uncontrolled murein degradation by the latter (Koch, 2000; Rogers et al., 1980; Weidel & Pelzer, 1964). However, the lack of hard evidence in support of the implication of hydrolases in elongation of the sacculus and the demonstration of β-lactam-induced lysis in non-growing cells under specific conditions (Ishiguro & Kusser, 1988; Kusser & Ishiguro, 1986, 1987) has led to alternative models in which lysis is the consequence of the activation of lytic enzymes which under normal growth conditions would be partially or totally inhibited (Ishiguro & Kusser, 1988; Tomasz, 1983, 1984).

Early observations of β-lactam-induced bacteriolysis showed the preferential occurrence of lysis at potential division sites (Staugaard et al., 1976; Tomasz, 1974), which, in the case of E. coli, often resulted in sacculi with a sharp cut at the corresponding location (Schwarz et al., 1969). These observations suggest that SMS might be particularly prone to provoke lysis when disturbed. The zonal nature of SMS suggests that the local concentration of biosynthetic complexes might be higher at division sites than anywhere else in the cell. If biosynthetic complexes require the concomitant activity of hydrolases, damage inflicted by bacteriolytic agents should be more pronounced at the places with higher local densities. Furthermore, as septation itself needs the activity of murein hydrolases to split nascent septa (Heidrich et al. 2001; Höltje & Heidrich, 2001), bacteriolytic agents could trigger lysis more effectively at those areas where hydrolases are actually performing their function and loss of control would have immediate consequences. According to such ideas, the mode of lysis of filament cells should be strongly dependent on whether or not SMS is activated during growth under restrictive conditions. In the former case lysis should occur as sharp cuts at the position of the rings of ‘all new’ murein generated by the activation of SMS. In the latter, as no regions of localized synthesis develop, a

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**Fig. 1.** Effect of cefsulodin on filament cells of E. coli MC6RP41 (ftsI) and MC6RP62 (ftsZ) growing at 42 °C. Cultures of both strains growing exponentially at 30 °C for three mass doublings were diluted 1:4 in the same medium prewarmed at 42 °C (time 0) and allowed to grow for one more doubling in mass. At that moment (arrow) cultures were divided into two equal parts. Cefsulodin (50 µg ml⁻¹) was added to one subculture of each strain and the OD₅₅₀ of the cultures was measured periodically. At the times indicated by the arrowheads samples were removed from each subculture to purify sacculi for electron microscopy. ●, MC6 RP41 control; ○, MC6RP41 plus cefsulodin; ■, MC6RP62 control; □, MC6RP62 plus cefsulodin.
general weakening of the sacculus network, with essentially random break points, is more likely to be observed. Examination of cefsulodin-induced lysis in sacculi from filament cells of E. coli ftsZ and ftsI mutants confirmed these predictions.

METHODS

Strains and growth conditions. E. coli strains used in this work were MC6RP1 (K-12, F’ dra drm leuA lysA proA thi tbr) (Prats & de Pedro, 1989), and derivatives MC6RP41 (ftsZ+) and MC6RP62 (ftsZ-) (Garcia del Portillo & de Pedro, 1990). Cultures were routinely grown in LB medium (Lennox, 1955) at 30 or 42 °C, as indicated, in gyratory water baths. Growth was monitored by measuring OD_{595} in a Spectronic 20 (Bausch & Lomb) spectrophotometer.

Purification of sacculi. Culture samples (ca 10^{10} cells) were harvested by centrifugation (10000 g, 3 min) at the growth temperature. The pellets were quickly resuspended into 3 ml 0.9 g NaCl l^{-1}, transferred into test tubes containing 6 ml 8% (w/v) SDS and kept closed on a boiling water bath for 6 h with magnetic stirring. Sacculi were further processed for electron microscopy as described by de Pedro et al. (1997).

Electron microscopy of sacculi. Carbon-pioloform-coated copper grids (200# mesh) were glow-discharged (10 min) and floated for 15 min on drops of sacculi suspensions appropriately diluted in double-distilled water. Grids were then removed, excess liquid was absorbed on filter paper and grids were allowed to air dry for 10 min. Grids were washed (five times) by flotation on distilled water drops, stained by floating for 1 min on 1% (w/v) uranyl acetate in water, briefly washed in water and air-dried. To better visualize the flatness of sacculi, grids were in some instances subjected to carbon/platinum shadowing at an angle of 15°. Microscopic observations were performed on a Philips CM10 transmission electron microscope at an acceleration voltage of 60 kV.

Muramidase digestion of murein sacculi. ‘On the grid’ digestion of sacculi was performed as described by de Pedro et al. (1997). In short, purified sacculi were deposited on electron microscope copper grids and washed as described above. Grids were floated for 1 min on drops (50 µl) of a solution of Cellosyl muramidase (Aventis) at 5 µg ml^{-1} in 20 mM sodium phosphate buffer, pH 4.9, at room temperature. To stop the reaction the enzyme solution was removed from the grid with filter paper, grids were rapidly washed three times in drops of precooled (4–6 °C) double-distilled water and stained for 1 min by flotation on drops of 1% (w/v) uranyl acetate in water.

RESULTS

Cefsulodin-induced lysis in filament cells of E. coli MC6RP41 (ftsI) and MC6RP62 (ftsZ)

Lysis was induced by addition of cefsulodin, a highly bacteriolytic β-lactam that binds preferentially to PBPs 1a and 1b in E. coli and that has been used to study the mechanism of bacterial lysis (Garcia del Portillo et al., 1989; Jacoby & Young, 1991; Powell & Young, 1991). Cultures of both strains were grown in LB at 30 °C for three mass doublings and were then diluted 1:4 into new medium pre-warmed at 42 °C. Cultures were allowed to grow for one doubling in mass, divided into two halves and cefsulodin (50 µg ml^{-1}) was added to one of the subcultures. Lysis started earlier and was clearly faster in the ftsI strain than in the ftsZ strain (Fig. 1).

Analysis of sacculi from lysing cultures of E. coli MC6RP41 (ftsI) and MC6RP62 (ftsZ)

At the times indicated by the arrowheads in Fig. 1, samples of filamenting cells were withdrawn from the cultures and processed for sacculi purification and electron microscopy. Sacculi from cefsulodin-treated ftsI cells showed evident differences compared to those from intact cells (Fig. 2a, b). Sacculi from treated cells were often transversely cut in an extremely sharp way (Fig. 2e, f). Rings of murein with a partially degraded appearance were often observed at positions apparently corresponding to division sites (Fig. 2b, e, f). The rest of the sacculus surface showed signs of general degradation in the form of a murein network mesh opening (Fig. 2c, d). In samples from similarly treated ftsZ cultures only a few sacculi showed clear signs of degradation. Not a single one was found with cuts comparable to those in ftsI sacculi. Instead extended areas of partially degraded murein with a frayed appearance were found in sacculi from the ftsZ mutant (Fig. 3a). However, the murein of most sacculi also showed clear signs of general degradation, in the form of small openings and a fibrillar appearance (Fig. 3b, c), similar to that commented on above for ftsI strain filaments (Fig. 2c, d). The appearance of murein in antibiotic-treated ftsZ sacculi was very similar to that of control sacculi subjected to mild digestion (1 min, 5 µg ml^{-1} at room temperature) with the murein hydrolytic enzyme Cellosyl, an endo-N-acetylmuramidase able to fully degrade murein into soluble muropeptides (Fig. 3c, d).

Size distribution of sacculi from cefsulodin-lysed filamentous cells of E. coli MC6RP41

An immediate consequence of the apparently preferential lysis at division sites in the ftsI strain is that the length distribution of lysed sacculi and the distribution of lengths from the cell poles to division sites should correspond. Therefore, the lengths of lysed sacculi from a culture of MC6RP41 grown at 42 °C for one doubling time and then treated with cefsulodin for 60 min were compared to the polar distances of constrictions in sacculi from filaments of a parallel antibiotic-free culture. For cefsulodin-treated sacculi, length was calculated from the rectangular coordinates of the sacculi ends using the microscope built-in system for grid position determination. To define the polar distance distribution of constrictions, the distance of each constriction to each pole was measured in the same way. Only straight sacculi were considered suitable. Repeated (10-fold) measurements of selected sacculi indicated a σ of ±0.1 µm. The results (Fig. 4) showed a clear correlation between the two distributions, at least for sacculi up to 6 µm, and therefore support the idea that
Fig. 2. For legend see facing page.
cells do lyse at potential septation sites under our experimental conditions.

DISCUSSION

Analysis of sacculi from ftsZ and ftsI cell division mutants induced to lyse under restrictive conditions further confirmed that cell lysis is preferentially triggered at active growth sites (Hölting, 1995; Hölting & Heidrich, 2001; Rogers et al., 1980; Schwarz et al., 1969; Tuomanen & Tomasz, 1990; Tomasz, 1983; Weidel & Pelzer, 1964) and suggests a co-localization of biosynthetic complexes and hydrolytic activities. Induction of cell lysis in filament cells of the ftsZ mutant by cefsulodin treatment resulted in rather delayed onset of lysis, as asserted by optical density measurements and in the generation of areas of extended damage apparently at random in the murein sacculus. On the assumption that hydrolases are associated to biosynthetic complexes (Hölting, 1996), such a pattern fits well with a diffuse mode of growth where many biosynthetic complexes are active at dispersed and random locations of the sacculus. Inhibition of each one would be insufficient to start lysis at that particular site, at least in the short term. However, accumulation of relatively small lesions would progressively reduce the mechanical resistance of the sacculus below the limit necessary to withstand the turgor pressure of the cell and lysis would ensue. In contrast, similarly treated ftsI cells lysed faster and sacculi exhibited extremely sharp transverse cuts at positions which correspond with SMS activation sites in the ftsI cells. These characteristics suggest that the simultaneous inhibition of the synthetic enzymes in a relatively large number of densely packed biosynthetic complexes leads to rapid murein degradation at the precise place that breaks apart under the turgor pressure of the cell. In the ftsI filaments, β-lactams also inhibit complexes involved in lateral wall synthesis. However, development of critical damage in these sites would be much slower than in regions of active SMS and cells would preferentially lyse at the latter sites as observed. An alternative possibility would be the activation of specific hydrolases at potential division sites in an FtsZ-dependent fashion. However, this alternative would not explain damage to murein outside the division site and so far there is no indication for direct regulation of murein hydrolases by FtsZ.

The periodic alternating use of diffuse and zonal modes of murein growth helps to explain the old observation that β-lactams apparently induce lysis in accordance with cell division in normally dividing cells. Induction of lysis by β-lactams proceeds at a rate related to the growth rate and very often cells blow up at positions corresponding to division sites (Leduc & van Heijenoort, 1980; Leduc et al., 1982; Schwarz et al., 1969; Staunagard et al., 1976). According to our interpretation in any cell able to trigger SMS, lysis should occur preferentially at the division site. As generation of division sites is intimately related to growth rate, the rate of lysis should be automatically linked to the rate of growth.

The link between induction of fast, localized cell lysis and activation of SMS at division sites could explain previous, and apparently contradictory, results about the lytic response to cefsulodin of synchronized cultures of E. coli. In experiments performed with cells synchronized by thymine/amino acid starvation, addition of cefsulodin at any time prior to termination of DNA replication resulted in cell lysis at the time of division, but not earlier (Garcia del Portillo et al., 1989). In contrast, in similar experiments performed with cells synchronized either by the ‘Baby Cell Machine’ (membrane elution) (Jacoby & Young, 1991) or centrifugal elutriation (Wientjes & Nanninga 1991 methods, lysis started a short time (10 and 20 min, respectively) after addition of the β-lactam with no apparent connection to the division event. The key fact might be that the thymine/amino acid starvation synchronization method blocks division via FtsZ, whereas both the Baby Cell Machine and centrifugal elutriation methods do not. Therefore, in the former case no cell in the culture was able to trigger SMS before the end of the ongoing round of DNA replication. Addition of cefsulodin during the replication period surely affected diffuse lateral wall synthesis. However, as shown above, cells take a long time to lyse when only lateral wall synthesis is affected. Therefore, cells could terminate replication, trigger SMS and, as cefsulodin was present, a fast lysis would then follow. In Baby Cell Machine and centrifugal elutriation synchronized cultures alignment of division initiation is not quite as strict as it is for thymine/amino acid starvation where the cells have to go through a complete round of replication (around 40 min under the conditions used) before SMS can actually start. Furthermore, the doubling time in the experiments was relatively short (30–60 min) and cefsulodin, as for any other β-lactam, requires a few minutes before the onset of lysis. Therefore, it is quite possible that SMS was triggered in some cells before or shortly after addition of cefsulodin to the synchronized cultures even at the earlier times of drug addition. Then lysis could start soon after cefsulodin addition in all instances, as was indeed observed. The longer lysis delay observed in centrifugal elutriation synchronized cultures could reflect the better synchrony apparently obtained by this method.

Fig. 2. Electron microscopy of sacculi from cefsulodin-treated filament cells of E. coli MC6RP41 (ftsI) growing at 42 °C. Samples from control and cefsulodin-treated cultures of MC6RP41 growing at 42 °C were collected as indicated in Fig. 1 and processed for electron microscopy. (a) Control sacculi; (b) cefsulodin-treated sacculi (arrowheads indicate break points); (c) detail of murein from control sacculi; (d) detail of murein from cefsulodin-treated sacculi; (e) ftsI lysed sacculi from cefsulodin-treated samples (arrowheads indicate partially degraded constriction sites). (a–e) Uranyl-acetate-stained sacculi were shadow-cast with carbon-platinum at an angle of 15°.
Fig. 3. Electron microscopy of sacculi from cefsulodin-treated filament cells of *E. coli* MC6RP62 (ftsZ) growing at 42 °C. Samples from control and cefsulodin-treated cultures of MC6RP62 growing at 42 °C were collected as indicated in Fig. 1 and processed for electron microscopy. (a) Cefsulodin-treated sacculi displaying extended damaged areas; (b) control sacculi; (c) cefsulodin-treated sacculi; (d) muramidase-digested control sacculi.
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REFERENCES


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