Premature targeting of a cell division protein to midcell allows dissection of divisome assembly in *Escherichia coli*

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Cell division in *Escherichia coli* requires the recruitment of at least 10 essential proteins to the bacterial midcell. Recruitment of these proteins follows a largely linear dependency pathway in which depletion of one cell division protein leads to the absence from the division site of “downstream” proteins in the pathway. Analysis of events that underlie this pathway is complicated by the fact that a protein’s ability to recruit “downstream” proteins is dependent on its own recruitment by “upstream” proteins. Hence, one cannot separate the individual contributions of various upstream proteins to any specific recruitment step. Here we present a method—premature targeting—for bypassing the normal localization requirements of a cell division protein and apply it to FtsQ, a protein recruited midway through the pathway. We fused FtsQ to the FtsZ-binding protein ZapA such that FtsQ was targeted to FtsZ rings independently of proteins FtsA and FtsK, which are normally required for FtsQ localization. Analysis of the resulting ZapA–FtsQ fusion suggests that FtsQ associates with a large complex of cell division proteins and that premature targeting of FtsQ can restore localization of this complex under conditions in which neither FtsQ nor the associated proteins would normally be localized.

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In *Escherichia coli*, cell division requires the coordinated constriction of the three layers of the Gram-negative cell envelope. Invagination is mediated by the septal ring, an assembly of proteins that localizes to the site of constriction and appears to remain at the site until separation of the two daughter cells. The septal ring is composed of multiple proteins, both essential and nonessential, which localize to the septum and/or are required for optimal cell division. The 10 known essential proteins are strictly required for cell division and localize to the constriction site in a defined hierarchical order (Fig. 1A; Buddeleijer and Beckwith 2002). This hierarchy defines the septal recruitment “pathway,” in which a given protein requires all proteins upstream of it to localize to the division site, and is itself required for the localization of those that are farther downstream. The mode of localization to the septum within the pathway, however, is understood for only a few of the proteins, leaving open the questions of how the cell assembles the “divisome” into its final state and ultimately whether the order of assembly is critical to the cell division process.

The assembly of the most upstream components of the septal ring is best understood. FtsZ is a widely conserved bacterial homolog of tubulin [Mukherjee et al. 1993; Lowe and Amos 1998] and was the first cell division protein shown to be localized to the bacterial midcell [Bi and Lutkenhaus 1991]. FtsZ polymers form a ring or tight spiral at the bacterial midcell, which is thought to provide both a scaffold for assembly of subsequent septal ring components and possibly the force required for constriction [Romberg and Levin 2003]. FtsA, a conserved member of the actin superfamily [Bork et al. 1992; van den Ent and Lowe 2000], and ZipA have both been shown to bind directly to the C terminus of FtsZ [Hale and de Boer 1997; Wang et al. 1997; Ma and Margolin 1999; Mosyak et al. 2000; Moy et al. 2000; Yan et al. 2000] and are thought to be required for stabilizing polymerized Z rings at the midcell [Pichoff and Lutkenhaus 2002]. The molecular events that underlie recruitment among later proteins remain speculative.

Interestingly, the almost strict linearity of recruitment observed in *E. coli* does not appear to be conserved in the Gram-positive bacteria *Bacillus subtilis* and *Streptococcus pneumoniae* [Errington et al. 2003; Morlot et al. 3Present address: Departamento de Bioquímica, Instituto de Química, Universidade de São Paulo, Av. Prof. Lineu Prestes 748, São Paulo, SP 05508-000, Brazil.
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In B. subtilis, localization of late cell division proteins is interdependent, suggesting they are assembled and recruited in a cooperative fashion (Errington et al. 2003; Morlot et al. 2004). Recent information derived from bacterial two-hybrid approaches suggests that the pathway in E. coli may in fact be more complex than originally thought. Indeed, it is suggested that many of the cell division proteins interact with at least one other cell division protein, and that several proteins interact with multiple proteins within the septal ring assembly (Fig. 1B; Di Lallo et al. 2003; G. Karimova and D. Ladant, in prep.). Interpretation of these two-hybrid results and understanding the ultimate role of these apparent interactions in divisome assembly are complicated as these interactions are not necessarily direct and should be confirmed by in vitro pull-down assays; however, if these two-hybrid interactions actually reflect the protein-binding events that direct assembly of these proteins at the septum, the significance of the linear assembly pathway and the molecular events responsible for the linearity seem even more mysterious.

The simplest model to explain recruitment in the E. coli system envisions a largely linear assembly via protein–protein interactions, such as cars in a train. In this model, each protein binds only to immediately adjacent proteins in the pathway, and interactions between adjacent proteins are independent of proteins localizing either farther upstream or downstream (Fig. 1C). Under this scenario, many of the positive signals seen by two-hybrid analysis could simply be a result of weak interactions or the close proximity of these cell division proteins when assembled into a complex rather than strong direct interactions. Alternatively, the two-hybrid data may be interpreted as indicating that recruitment involves cooperative binding of a recruited protein by a preassembled complex of upstream proteins. Under this model, only this complex of early proteins is able to provide the multiple required interactions that allow recruitment of the subsequent protein (Fig. 1D). Finally, one could envision that the linear pathway actually reflects a series of sequentially ordered septum modification events. In such a model, upstream proteins modify the septum (e.g., by modifying the cell wall murein), to create a localized signal at the septum that can be recognized by the next protein in the pathway (Fig. 1E). The assembly of high-molecular-weight penicillin-binding proteins (HMW PBPs) at the division site in S. pneumoniæ (Morlot et al. 2004) seems to follow such a model.

All three models make specific predictions about the nature of recruitment events that occur between adjacent proteins in the pathway. In particular, the latter two models would predict that a protein must necessarily localize before the next protein in the pathway, as it must associate with other early proteins or perform some function at the septum before the next protein can be recruited. In the first model (linear assembly), however, the order of localization would not necessarily be important because recruitment would be simply a consequence of binding between two proteins. In this scenario, localization of either of a pair of interacting proteins would allow the recruitment of the other, and this interaction would be independent of upstream proteins in the pathway.

The linearity of the pathway, however, makes distinguishing among these models difficult. Since a protein within the pathway requires all proteins localizing up-

Figure 1. (A) The septal recruitment pathway for cell division proteins in E. coli. (B) Bacterial two-hybrid results summarized from several groups (Di Lallo et al. 2003; G. Karimova and D. Ladant, in prep.; M. Gonzalez and J. Beckwith, unpubl.). Cell division proteins are listed clockwise corresponding to their respective order in A. Lines indicate proposed interactions; circular arrows indicate proposed homodimerization. (C–E) Potential models for divisome assembly. (C) Linear assembly. (D) Cooperative binding by an upstream complex. (E) Sequential activities plus substrate recognition. (F) ZapA–FtsQ constructs used in this work. (Cyto) Cytoplasmic domain of FtsQ; (TM) transmembrane segment of FtsQ; (Peri) periplasmic domain of FtsQ. YFP and non-YFP constructs are identical except for the addition of YFP fused in frame to the N terminus of ZapA. (G) Proposed premature targeting of ZapA–FtsQ construct.
stream of it for its proper localization and a delocalized protein presumably cannot recruit the next protein in the pathway to the septum, it is difficult to identify which proteins are in fact strictly required for a given recruitment event. Hence, testing these predictions requires a method to divorce a given protein from its normal localization requirements, allowing recruitment by that protein to be analyzed in cells lacking upstream proteins. In this work we devised such a method and apply it to analyze the role of FtsQ in the assembly pathway.

FtsQ is a bitopic membrane protein with a short cytoplasmic sequence at its N terminus and a relatively large periplasmic sequence (Carson et al. 1991). It is in the middle of the localization pathway (Buddelmeijer and Beckwith 2002) and according to two-hybrid analysis [Di Lalio et al. 2003; G. Karimova and D. Ladant, in prep.] makes multiple contacts with other proteins in the pathway. These data raise the possibility that FtsQ could coordinate recruitment of many of the downstream proteins. Coimmune precipitations also indicate that FtsQ is likely part of a complex with downstream proteins FtsL and FtsB, although FtsQ localizes efficiently in their absence [Buddelmeijer and Beckwith 2004]. Normally FtsQ requires FtsZ, FtsA, ZipA, and FtsK in order to localize to the septum, and its localization in turn is required for recruitment of downstream proteins such as FtsL and FtsI [Chen et al. 1999; Ghigo et al. 1999; Weiss et al. 1999; Chen and Beckwith 2001; Hale and de Boer 2002; Pichoff and Lutkenhaus 2002]. If an early protein such as FtsA or FtsK is selectively depleted from cells, FtsQ is no longer localized to the septum and its ability to recruit proteins cannot be assayed. However, if FtsQ could somehow be localized “prematurely” to the septum in the absence of upstream proteins, one could analyze whether these upstream proteins contribute to the recruitment of downstream proteins by FtsQ or are simply required for the localization of FtsQ itself.

In this paper, we describe a technique for premature targeting of FtsQ to the nascent Z ring by fusing it to the protein ZapA. ZapA is a small cytoplasmic protein that appears to be a nonessential component of the septal ring. It localizes to FtsZ rings in both *E. coli* and *B. subtilis*, and purified ZapA protein can bind to FtsZ in vitro and promote bundling of FtsZ filaments [Gueiros-Filho and Losick 2002; Anderson et al. 2004]. These data suggest that ZapA binds directly to FtsZ in vivo independently of other cell division proteins. Therefore, we predict that FtsQ, if fused to ZapA [diagrammed in Fig. 1F], should localize in the absence of FtsA, ZipA, and FtsK [Fig. 1G]. The fact that ZapA is nonessential in both *B. subtilis* and *E. coli* [Gueiros-Filho and Losick 2002; Johnson et al. 2004] indicates that it is not required for localization of other cell division proteins and hence is unlikely to be responsible for any observed recruitment by a ZapA fusion protein.

The results presented here demonstrate that FtsQ when fused to ZapA is prematurely targeted to midcell. Our studies on the functioning of the prematurely targeted FtsQ in the assembly pathway provide new insights into this pathway and help to distinguish between various models for how proteins are assembled at the nascent division site.

### Results

Based on the known properties of the ZapA protein, we reasoned that fusion of full-length *E. coli* ZapA to the N terminus of FtsQ would result in the localization of FtsQ to the sites of FtsZ rings independently of FtsA, ZipA, and FtsK, three proteins normally required for FtsQ localization. Critical to this experiment, however, is the confirmation that, in *E. coli*, ZapA is part of the septal ring and localizes to FtsZ rings independently of other cell division proteins.

#### Positioning ZapA in the *E. coli* septal recruitment pathway

In order to determine whether localization of ZapA is dependent on FtsZ and not other downstream proteins, we introduced the plasmid pNG53, which expresses a YFP–ZapA fusion under IPTG control, into strains carrying either the *ftsA12*(Ts) or *ftsZ84*(Ts) temperature-sensitive mutations to create NWG128 and NWG129. Strains carrying either mutation grow and divide normally at 30°C, but fail to divide when grown at 42°C, resulting in the formation of filamentous, undivided cells. A GFP fusion to a cell division protein such as FtsQ that depends on both FtsZ and FtsA for localization fails to localize in cells carrying these mutations at 42°C, but localizes normally at 30°C [Fig. 2; Chen et al. 1999]. The YFP–ZapA fusion protein, although failing to localize in *ftsZ84* cells grown at the restrictive temperature, localizes efficiently in *ftsA12* cells [Fig. 2]. These results indicate that, in *E. coli*, ZapA localizes to the septal ring and its localization is dependent on FtsZ, independent of FtsA, and presumably independent of proteins further downstream such as FtsK and FtsQ, all of which require FtsA for their normal localization.

#### Construction and localization of a ZapA–FtsQ fusion

To test the ability of ZapA to localize FtsQ prematurely, we constructed the plasmid pNG107, which expresses a

![Figure 2. Localization of YFP–ZapA requires FtsZ but not FtsA. YFP–ZapA and GFP–FtsQ localize in *ftsZ84(Ts)* or *ftsA12(Ts)* cells grown at 30°C (permissive temperature). When grown at 42°C (restrictive temperature), YFP–ZapA localizes in cells carrying the *ftsA12(Ts)* allele but not in those containing *ftsZ84(Ts).* GFP–FtsQ does not localize in either background at 42°C. The strains used are JOE95, JOE97, NWG128, and NWG129.](image-url)
YFP–ZapA–FtsQ fusion [Fig. 1F] at low levels from an IPTG-inducible \( P_{\text{Trc}} \) promoter carrying −10 and −35 down mutations [Weiss et al. 1999]. The YFP–ZapA–FtsQ fusion expressed from pNG107 localized to the midcell in wild-type cells and was able to complement a strain depleted for FtsQ [data not shown]. To confirm that YFP–ZapA–FtsQ localization was independent of FtsA and FtsK, pNG107 was introduced into a strain carrying the \( ftsA12[Ts] \) mutation or into the FtsK depletion strain NWG326. The latter strain carries a chloramphenicol replacement of the \( ftsK \) gene complemented by a low-copy plasmid expressing FtsK under an arabinose-inducible \( P_{\text{BAD}} \) promoter. When grown in arabinose, FtsK is expressed and the cells grow and divide normally. Upon switching to glucose, \( ftsK \) expression is repressed, FtsK protein is depleted, and cells become unable to divide, resulting in filamentous growth [Chen and Beckwith 2001]. Both strains also carry an FtsZ–CFP fusion construct under IPTG control integrated into the chromosome [Chen and Beckwith 2001]. This allowed us to confirm that the observed localization of YFP signal overlapped the sites of FtsZ rings as visualized by CFP. To confirm that our cells were properly depleted, isogenic strains expressing YFP–FtsQ from pJC64 were always grown in parallel for comparison.

As expected, both YFP–FtsQ and YFP–ZapA–FtsQ appeared as rings overlapping the CFP–FtsZ signal at the bacterial midcell [Fig. 3] when expressed in \( ftsA12[Ts] \) cells grown at 30°C or cells complemented for FtsK by growth in arabinose. Upon depletion for FtsA at 42°C or for FtsK by growth in glucose, YFP–ZapA–FtsQ remained colocalized with CFP–FtsZ, forming regularly spaced rings along the filamentous cell, whereas the control, YFP–FtsQ, was no longer localized. Western blots indicated that the lack of YFP–FtsQ localization was not due to lower amounts of protein [data not shown]. Since the YFP is fused to the N terminus of the ZapA–FtsQ fusion, it is possible that the septal signal we see is due to a breakdown product consisting of YFP–ZapA rather than the full-length fusion; however, Western blot analysis [data not shown] in combination with the results described below indicate that the majority of full-length fusion protein is recruited to the septum under these conditions. These results suggest that the ZapA moiety is effectively recruiting the ZapA–FtsQ fusion to the septum independently of FtsA and FtsK, both of which are normally required for FtsQ localization.

Recruitment of FtsL

Previous results indicated that FtsQ, FtsL, and FtsB exist in a stable complex in the cell and that the FtsL–FtsB complex, presumably including FtsQ, can form in the absence of FtsK, a condition in which none of the three proteins are normally localized to the septum [Buddelmeijer and Beckwith 2004]. In the previous section we showed that fusion to ZapA results in the localization of FtsQ to the septum independently of FtsK. Therefore, if FtsQ, FtsL, and FtsB are part of an FtsK-independent complex, FtsL and FtsB should be recruited to the septum by the ZapA–FtsQ fusion protein in cells depleted for FtsK. To test this prediction, an IPTG-inducible construct expressing the ZapA–FtsQ fusion [construct diagrammed in Fig. 1F] was integrated into the chromosome at the \( \lambda \) \( att \) site using \( \lambda \) InCh and expression confirmed by Western blot analysis [data not shown]. This construct was then introduced into an FtsK depletion strain expressing a GFP–FtsL fusion integrated into the chromosome at the \( att \) 800 site as described previously [Chen and Beckwith 2001]. In order to ensure that the recruitment was specific to the ZapA–FtsQ fusion protein, we also introduced identical chromosomal constructs expressing either ZapA or FtsQ alone into isogenic FtsK depletion strains for comparison.

As predicted, when the ZapA–FtsQ protein is expressed in the FtsK depletion strain described above, GFP–FtsL localizes efficiently to regularly spaced rings when compared to control cells expressing either ZapA or FtsQ alone [Fig. 4A–C, Table 1, rows 1–3]. The ring spacing (an inverse measure of ring frequency) for GFP–FtsL in the ZapA–FtsQ-expressing cells of 9.5 \( \mu \)m/ring [Table 1, row 1] is very similar to the published spacing of 8.8 \( \mu \)m/ring for FtsZ localization in FtsK-depleted cells and 13 \( \mu \)m/ring for FtsL localization in FtsN-depleted cells [Chen and Beckwith 2001]. This indicates that ZapA–FtsQ is recruiting GFP–FtsL at a near maximal expected level if it were to recruit GFP–FtsL to every available Z ring. In contrast, expression of FtsQ or ZapA alone from an identical chromosomal construct in isogenic FtsK-depleted cells resulted in significantly reduced levels of localization of GFP–FtsL [Fig. 4B,C, Table 1, rows 2–3]. Thus, ZapA–FtsQ appears able to direct the recruitment of FtsL to FtsZ rings in the absence of FtsK, and this recruitment is not due to the individual effects of expressing either ZapA or FtsQ alone.
It remained possible that this ability of ZapA–FtsQ to recruit FtsL was specific in its ability to bypass FtsK but still depended on proteins farther upstream. We considered FtsA as the most likely candidate for such a contributor as two-hybrid data suggest it may contact downstream proteins FtsI and FtsN (Di Lallo et al. 2003; G. Karimova and D. Ladant, in prep.) and is widely conserved among bacteria [Margolin 2000]. To test the ability of ZapA–FtsQ to recruit FtsL in the absence of FtsA, we combined the ZapA–FtsQ fusion and GFP–FtsL constructs with the ftsA12(Ts) mutation and analyzed localization of GFP–FtsL at the restrictive temperature. Under these conditions, expression of ZapA–FtsQ is able to restore localization of GFP–FtsL to regularly spaced rings (Fig. 4D; Table 1, row 4). As is the case in FtsK-depleted cells, expression of either FtsQ or ZapA alone failed to restore localization of GFP–FtsL, as indicated by reduced frequency and fluorescence intensity of observed rings (Fig. 4E,F; Table 1, rows 5–6). Thus, neither FtsA nor FtsK is required for recruitment of FtsL in cells expressing the ZapA–FtsQ fusion.

Finally, it remained formally possible that FtsL requires FtsQ and either FtsA or FtsK for its localization. To rule out such redundant roles for FtsA and FtsK, we performed the same experiment in cells depleted for both FtsK and FtsA. To do this, we introduced the ftsA12(Ts) mutation into the FtsK depletion strains expressing GFP–FtsL and either the ZapA–FtsQ fusion or FtsQ alone as described above. By growing the resulting cells at 42°C to deplete FtsA and in glucose to repress expression of FtsK from the complementing plasmid, we were able to analyze recruitment in cells missing both proteins. GFP–FtsL localized in doubly depleted cells expressing ZapA–FtsQ (Fig. 4G; Table 1, row 5), indicating that ZapA–FtsQ is in fact capable of recruiting FtsL in the absence of both FtsA and FtsK. In light of previous work indicating that FtsB and FtsL are dependent on each other for localization and can be coimmunoprecipitated [Buddelmeijer et al. 2002; Buddelmeijer and Beckwith 2004], these data are consistent with a model in which FtsL, FtsB, and FtsQ exist in a stable complex that can form independently of proteins upstream of FtsQ. Hence, under normal circumstances, the simple presence of FtsQ at the septum is sufficient to allow localization of FtsL and, presumably, FtsB.

Recruitment of FtsI and FtsN

Having established the ability of ZapA–FtsQ to recruit FtsL in the absence of upstream proteins, we proceeded to examine the extent to which simply localizing FtsQ to the septum is able to restore localization of the remainder of downstream proteins in the cell division pathway, including FtsI and FtsN. To this end, we repeated the above experiments with GFP–FtsI and GFP–FtsN in ZapA–FtsQ-expressing cells that were depleted for FtsK, FtsA, or both.

FtsI was efficiently localized in cells expressing ZapA–FtsQ but depleted for either FtsK or FtsA (Fig. 5A,B). In cells depleted for both FtsA and FtsK, localization also seems to occur at regular intervals in filaments [Fig. 5C]. Such localization was not seen in control cells expressing FtsQ (Table 1). This suggests that ZapA–FtsQ not only restores FtsL and FtsB to the septum in FtsA/FtsK-depleted cells, but also restores the localization pathway from FtsQ through FtsI as shown in Figure 8A (below). Since FtsI strictly requires FtsL, FtsB, and FtsW to localize to the septum under normal conditions, we assume these proteins are also required for recruitment of FtsI when FtsQ is localized via fusion to ZapA. If such is the case, FtsQ, FtsL, FtsB, FtsW, and FtsI are likely capable of assembling into a stable complex in the cell independently of upstream proteins and hence independently of the proteins’ localization to midcell.

FtsN presents quite a different story. ftsN was originally identified as a multicopy suppressor of the ftsA12(Ts) mutation at 37°C. Subsequently, overexpression of FtsN was shown to suppress Ts mutations in ftsA, ftsK, ftsQ, and ftsL as well as partially suppressing FtsK depletion strains [Dai et al. 1993; Draper et al. 1998]. This property of FtsN complicates the analysis of FtsN recruitment by ZapA–FtsQ in FtsK depletion strains. Notably, in cells depleted for FtsK, GFP–FtsN localizes well regardless of whether FtsQ or ZapA–FtsQ is expressed [Fig. 5; J. Chen and J. Beckwith, unpubl.]. In fact, average cell length is much reduced compared to FtsK depletion strains expressing GFP fusions to other cell division proteins [Fig. 5D; Table 1, rows 15–16 vs. 1–3, 9–10], and these cells readily form colonies on glucose plates, suggesting gfp–ftsN is acting as an overexpression suppressor in these strains. We have some indications that this apparent suppression may only be possible because of residual FtsK expression under non-inducing conditions [data not shown].

To obviate the problem of ftsN overexpression suppression, we analyzed recruitment of FtsN in ZapA–FtsQ-expressing strains depleted for FtsA or both FtsK and FtsA, since, at 42°C, the GFP–FtsN fusion is unable to suppress the cell division defects of these strains. In both strains, GFP–FtsN did not localize to regularly spaced rings [Fig. 5E,F; Table 1, rows 17, 19]. In contrast,
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Table 1. Quantitative summary of localization data

| GFP fusion fusion | FtsA | FtsK | ZapA–FtsQ or FtsQ(wt) | Strain     | Cells | Avg cell length (µm) | Fraction of cells w/rings | Rings/cell | Ring spacingb | Avg cell length (µm) | Avg cell length (µm) |
|-------------------|------|------|------------------------|------------|-------|----------------------|--------------------------|------------|----------------|----------------------|----------------------|
|                   |      |      | FtsQ                  | NWG252     | 65    | 22.1                 | 0.95                     | 2.3        | 9.5           |                      |                      |
| 1                 |      |      | FtsQ                  | NWG390     | 44    | 28.6                 | 0.39                     | 0.5        | 52.4          |                      |                      |
| 2                 |      |      | ZapA                  | NWG448     | 46    | 27.3                 | 0.61                     | 0.9        | 29.9          |                      |                      |
| 3                 |      |      | ZapA–FtsQ             | NWG341     | 38    | 40.5                 | 1.00                     | 3.1        | 13.3          |                      |                      |
| 4                 |      |      | FtsQ                  | NWG418     | 36    | 41.6                 | 0.56                     | 0.9        | 48.3          |                      |                      |
| 5                 |      |      | ZapA                  | NWG447     | 20    | 33.0                 | 0.8                      | 1.2        | 28.7          |                      |                      |
| 6                 |      |      | ZapA–FtsQ             | NWG259     | 77    | 49.3                 | 1.00                     | 4.2        | 11.8          |                      |                      |
| 7                 |      |      | FtsQ                  | NWG411     | 47    | 47.9                 | 0.36                     | 0.4        | 112.6         |                      |                      |
| 8                 |      |      | ZapA–FtsQ             | NWG277     | 53    | 25.1                 | 0.94                     | 2.3        | 10.8          |                      |                      |
| 9                 |      |      | ZapA–FtsQ             | NWG394     | 54    | 24.4                 | 0.65                     | 0.9        | 28.0          |                      |                      |
| 10                |      |      | ZapA–FtsQ             | NWG340     | 18    | 39.1                 | 1.00                     | 2.9        | 13.5          |                      |                      |
| 11                |      |      | ZapA–FtsQ             | NWG417     | 32    | 45.0                 | 0.69                     | 0.8        | 55.4          |                      |                      |
| 12                |      |      | ZapA–FtsQ             | NWG299     | 30    | 51.2                 | 1.00                     | 3.5        | 14.5          |                      |                      |
| 13                |      |      | FtsQ                  | NWG413     | 44    | 49.4                 | 0.36                     | 0.5        | 103.6         |                      |                      |
| 14                |      |      | ZapA–FtsQ             | NWG257     | 207   | 7.0                  | 0.85                     | 1.0        | 7.1           |                      |                      |
| 15                |      |      | FtsQ                  | NWG391     | 107   | 10.5                 | 0.7                      | 1.1        | 9.1           |                      |                      |
| 16                |      |      | ZapA–FtsQ             | NWG342     | 46    | 38.9                 | 0.70                     | 1.1        | 35.8          |                      |                      |
| 17                |      |      | FtsQ                  | NWG419     | 27    | 46.3                 | 0.85                     | 1.4        | 32.9          |                      |                      |
| 18                |      |      | ZapA–FtsQ             | NWG269     | 41    | 38.8                 | 0.8                      | 1.2        | 33.1          |                      |                      |
| 19                |      |      | FtsQ                  | NWG428     | 32    | 42.0                 | 0.75                     | 1.2        | 36.3          |                      |                      |
| 20                |      |      | ZapA–FtsQ             | NWG352     | 28    | 40.0                 | 1.00                     | 4.7        | 8.5           |                      |                      |
| 21                |      |      | FtsQ                  | NWG421     | 30    | 42.6                 | 0.83                     | 1.6        | 26.6          |                      |                      |
| 22                |      |      | ZapA                  | NWG446     | 24    | 31.9                 | 0.89                     | 2.2        | 15.3          |                      |                      |
| 23                |      |      | ZapA–FtsQ             | NWG255     | 99    | 21.4                 | 0.44                     | 0.4        | 38.4          |                      |                      |
| 24                |      |      | FtsQ                  | NWG343     | 41    | 43.5                 | 0.85                     | 1.3        | 32.4          |                      |                      |
| 25                |      |      | ZapA–FtsQ             | NWG261     | 26    | 55.2                 | 0.50                     | 0.7        | 79.8          |                      |                      |
| 26                |      |      | FtsQ                  | NWG414     | 31    | 22.8                 | 0.45                     | 0.5        | 41.6          |                      |                      |
| 27                |      |      | ZapA–FtsQ             | NWG415     | 50    | 21.1                 | 0.9                      | 1.9        | 11.0          |                      |                      |
| 28                |      |      | FtsQ                  | NWG423     | 23    | 41.9                 | 0.87                     | 1.6        | 26.8          |                      |                      |
| 29                |      |      | FtsQ                  | NWG258     | 107   | 15.8                 | 0.70                     | 1.1        | 35.8          |                      |                      |
| 30                |      |      | ZapA–FtsQ             | NWG394     | 54    | 24.4                 | 0.65                     | 0.9        | 28.0          |                      |                      |
| 31                |      |      | ZapA–FtsQ             | NWG340     | 18    | 39.1                 | 1.00                     | 2.9        | 13.5          |                      |                      |
| 32                |      |      | ZapA–FtsQ             | NWG417     | 32    | 45.0                 | 0.69                     | 0.8        | 55.4          |                      |                      |
| 33                |      |      | ZapA–FtsQ             | NWG299     | 30    | 51.2                 | 1.00                     | 3.5        | 14.5          |                      |                      |
| 34                |      |      | ZapA–FtsQ             | NWG413     | 44    | 49.4                 | 0.36                     | 0.5        | 103.6         |                      |                      |

*a+ indicates endogenous FtsK expression; +++ indicates endogenous FtsK plus expression from pBAD44.

*bRing spacing = total length [microns]/number rings; bold indicates positive localization score based on comparison with control cells (FtsQ or ZapA) and with published values of 8.1–14.5 µm/ring for localization of fts proteins in FtsN-depleted cells [Chen and Beckwith 2001].

under identical conditions, GFP–FtsN localizes normally in wild-type cells and can complement a ΔftsN-null mutation [data not shown], indicating the GFP–FtsN fusion protein is stable and functional under our experimental conditions.

The fact that FtsN fails to localize under conditions in which FtsQ, FtsL, FtsB, and FtsI are recruited suggests that whether or not FtsN makes direct contacts with FtsI or FtsQ, it must rely on additional signals at the septum to localize. Since FtsN requires the presence of FtsI and does not localize in cells depleted for FtsI, it is likely that its localization must integrate signals from both FtsI and early proteins. While previous studies were consistent with interactions between FtsI and FtsN being sufficient for recruitment of FtsN, our results point to a more complex process [Di Lallo et al. 2003; Wissel and Weiss 2004; G. Karimova and D. Ladant, in prep.].

Back-recruitment of FtsK

FtsK immediately precedes FtsQ in the cell division pathway and is ordinarily required for the localization of FtsQ [Chen and Beckwith 2001], suggesting that it likely plays a direct role in recruiting FtsQ. As proposed above, one can envision several models to explain how FtsQ is recruited by FtsK. In the linear association model, a simple interaction between FtsK and FtsQ would allow recruitment. In this case, upstream proteins such as FtsA would only be required in order to localize FtsK. Moreover, if a simple physical interaction links FtsK and FtsQ, localization of either protein should restore the localization of the other in cells lacking FtsA. Under alternative models, FtsK would necessarily need to localize first, either to associate with other upstream proteins or to modify the septum to produce an FtsQ recruitment signal.

To distinguish among these models, we determined whether localization of FtsQ via the premature targeting method could restore localization of FtsK. To do this, we examined localization of YFP–FtsK in ftsA12[Ts] cells expressing either ZapA–FtsQ or wild-type FtsQ constructs from the λ att site. Compared to expression of FtsQ or ZapA alone, expression of ZapA–FtsQ promotes efficient localization of YFP–FtsK, leading to both in-
crease ring frequency (decrease in average ring spacing) and increased localization signal intensity (Fig. 6; Table 1, rows 21–23). These results indicate that, for the recruitment of FtsQ, there is no underlying fundamental temporal order or integration of upstream signals. Rather, this evidence suggests that FtsQ can physically associate with FtsK within the cell. Although further experiments will be required to address whether this association requires FtsZ, it is possible that FtsK may be part of a cell division complex along with FtsQ, FtsL, FtsB, FtsW, and FtsI that can form independently of lo-
sociation requires FtsZ, it is possible that FtsK may be

tact the recruitment of multiple FtsQ molecules.

Prior to this work, it was unclear whether FtsA and
FtsK played a role in cell division other than facilitating
the recruitment of downstream proteins. Interestingly,
although premature recruitment of FtsQ to the septum
via fusion to ZapA was able to restore recruitment of
most of the cell division proteins downstream of FtsA
and FtsK, expression of ZapA–FtsQ failed to restore the
ability of ftsA12[Ts] or ∆ftsK strains to divide under re-
strictive conditions (Table 1, average cell length ZapA–
FtsQ vs. FtsQ-expressing cells, data not shown).

Discussion

The study of an individual step within a sequentially
ordered pathway is complicated by the fact that such a
step requires the prior activity of upstream proteins. In a
synthetic reaction, for example, activity of early en-

Initially, we concluded that either the two-hybrid data
were not reflecting direct interactions in vivo or that for
some reason our ZapA–FtsQ and GFP–FtsQ fusions were
incompatible. One possible explanation for the latter hy-
pothesis lies in the recent observation that ZapA, itself,
is present as a dimer in the cell [Low et al. 2004]. This
could result in strongly favoring a ZapA–FtsQ ho-
modimer rather than the ZapA–FtsQ:GFP–FtsQ heter-
odimer, making any potential dimerization impossible
to detect.

In the course of related experiments, however, we oc-
casionally saw localization of GFP–FtsQ in ZapA–FtsQ-
expressing cells when FtsK was expressed from a multi-
copy plasmid, suggesting that FtsK may somehow facili-
tate the recruitment of multiple FtsQ molecules.

To test this idea, we introduced a multicopy plasmid
expressing excess FtsK into a strain carrying the
ftsA12[Ts] allele and expressing both ZapA–FtsQ and
GFP–FtsQ constructs. Under conditions of FtsA deple-
tion, GFP–FtsQ was recruited in these cells [Fig. 7B;
Table 1, row 28]. If we either replaced the FtsK plasmid
with empty vector, or expressed wild-type FtsQ instead
of ZapA–FtsQ, GFP–FtsQ localization was reduced (Fig.
7A,C; Table 1, rows 27, 29). This result suggests that
FtsK may play some role in facilitating the association of
multiple FtsQ molecules, perhaps through stabilizing
the assembled proteins within a single large complex.

Recent work using bacterial two-hybrid systems has
raised the possibility that FtsQ interacts with itself [Di
Lallo et al. 2003; G. Karimova and D. Ladant, in prep.; M.
Gonzalez and J. Beckwith, unpubl.]. It remains unclear
from these results whether FtsQ is acting as a ho-
modimer or higher-order oligomer, or if the two-hybrid
data reflect the fact that FtsQ is simply present in mul-
tiple copies within a single divisome complex. Given
these results, it seemed reasonable to ask whether
ZapA–FtsQ could recruit GFP–FtsQ to the septum in
cells depleted for FtsK, FtsA, or both; however, our ex-
periments indicated that GFP–FtsQ fails to localize in
such cells expressing ZapA–FtsQ, conditions under
which FtsL and FtsI were efficiently localized (Table 1,
rows 24–26, data not shown).

"Divisome" may contain multiple copies of FtsQ

Recent work using bacterial two-hybrid systems has
raised the possibility that FtsQ interacts with itself [Di
Lallo et al. 2003; G. Karimova and D. Ladant, in prep.; M.
Gonzalez and J. Beckwith, unpubl.]. It remains unclear
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periments indicated that GFP–FtsQ fails to localize in
such cells expressing ZapA–FtsQ, conditions under
which FtsL and FtsI were efficiently localized (Table 1,
rows 24–26, data not shown).
enzymes is required to produce intermediate substrates for
the enzyme of interest. In such a case, the addition of
exogenous substrate diverts the reaction of interest
from its normal place in the pathway, allowing the
reaction to be analyzed in isolation. For a sequential as-
sembly pathway, such as the septal recruitment pathway
in *E. coli*, the ability of a protein to recruit downstream
proteins to the septum requires a priori the localization
of that protein by proteins farther upstream. Hence,
within the normal in vivo pathway, it is difficult to ana-
lyze recruitment by a particular protein separately from
the upstream events that are required for its localization.

In this work we have developed a method, which we
term premature targeting, of divorcing a particular pro-
tein in the cell division pathway of *E. coli* from its nor-
mal localization requirements. Here, fusion of the FtsZ-
binding protein ZapA to FtsQ allowed the premature tar-
getting of FtsQ directly to FtsZ rings. FtsQ, when
recruited prematurely to the septum by ZapA, can effi-
ciently recruit two downstream proteins, FtsL and FtsI,
in the absence of FtsK and/or FtsA, two upstream pro-
teins that are normally strictly required for localization
of FtsQ and all proteins downstream of it (Fig. 8A). We
also show that prematurely targeted FtsQ is even capable
of "back-recruitment" of FtsK in the absence of FtsA
(Fig. 8B).

These results are consistent with prior studies using
two-hybrid analysis and coimmune precipitations,
which established that FtsQ, FtsL, and FtsB can form a
complex within the cell and that this complex can form
even in the absence of FtsK, a condition under which
these proteins are not localized to the septum. Those
findings suggest that these proteins are always associ-
ated within the cell regardless of localization (Di Lallo et
al. 2003; Buddelmeijer and Beckwith 2004; G. Karimova
and D. Ladant, in prep.). Our results indicate further that
neither FtsK nor FtsA is required for the formation of
this complex or necessarily for its localization to the
septum if FtsQ is localized artificially (i.e., by fusion to
ZapA). Rather, since the simple localization of FtsQ by
premature targeting is sufficient to direct recruitment of
this FtsQ–FtsL–FtsB complex to the septum, FtsK and
FtsA presumably are only required for the localization of
the members of this complex insofar as they allow lo-
calization of FtsQ. We also show that this complex is not
limited to FtsQ, FtsL, and FtsB, but likely includes FtsI
and by inference FtsW, which is "upstream" of FtsI and
is normally required for FtsI localization. While the mo-
lecular mechanism by which FtsW and FtsI are recruited
to this complex is unclear, the association does not ap-
pear to involve FtsK or FtsA.

Although the results presented on FtsN localization
are not unambiguous, they suggest that FtsN, the last
known essential protein in the pathway, may follow a
more complex path to the septum. The finding that FtsN
is not simply recruited by contact with FtsI raises the
intriguing possibility that it plays a role in integrating
signals from both early and late proteins. Prior work has
speculated on such a role for FtsN in the pathway, as
overexpression of FtsN is capable of suppressing tem-
perature-sensitive mutations in various other genes
within the cell division pathway including *ftsA*, *ftsK*,
*ftsQ*, and *ftsI* (Dai et al. 1993; Draper et al. 1998).
The apparent FtsA–FtsN interaction (Fig. 1B) raises the
question as to whether FtsA directly contributes to FtsN lo-
calization, providing this "early signal." However, the
cytoplasmic and membrane-spanning segments of FtsN
are not required for FtsN localization and function [Dai
et al. 1996], making it difficult to imagine how a direct
interaction between cytoplasmic FtsA and FtsN would
be required for cell division. Instead, it seems reasonable
to speculate that FtsN somehow recognizes either pro-
tein conformations or a modification of the septum that
is generated only by properly assembled ring. By recog-
nizing such a signal, FtsN could prevent activation of the
division machinery until assembly of the "divisome" is
complete.

The "back-recruitment" of FtsK by ZapA–FtsQ sug-
gests a possible interaction between these two proteins,
which under normal circumstances allows forward re-
cruitment of FtsQ by a localized FtsK. The fact that lo-

Figure 7. Overexpression of FtsK promotes association of mul-
tiple FtsQ molecules. (β) When FtsK was overexpressed from
multicopy plasmid pBAD44 (pNG137) in cells expressing both
GFP–FtsQ and ZapA–FtsQ under conditions of FtsA depletion,
localization of GFP–FtsQ was seen. If either wild-type FtsQ was
expressed instead of ZapA–FtsQ (C) or empty vector was sub-
stituted for the FtsK plasmid (A), localization was significantly
reduced. Representative images from paired samples are shown.
A quantitation of results is presented in Table 1, rows 24–29.

Figure 8. Summary of observed recruitment by prematurely
targeted FtsQ in cells depleted for FtsA and FtsK (A) or FtsA
alone (B). (A) ZapA–FtsQ is able to recruit FtsL and likely FtsB,
which restores the pathway through FtsI. (B) ZapA–FtsQ is also
able to interact with FtsK, and this interaction with FtsK in turn
supports association of multiple FtsQ molecules. FtsB and FtsW
are shown in outline as their presence is not explicitly tested in
this work, but implied from previous data, which indicate FtsI
requires both FtsB and FtsW to localize to the septum.
calization of either protein results in the localization of the other, argues against a model in which activity of a localized FtsK must precede FtsQ localization. Additionally, since this interaction is independent of FtsA and since FtsQ localization under normal circumstances is independent of FtsL, FtsB, FtsW, FtsI, and FtsN (Chen et al. 1999; Chen and Beckwith 2001; Buddelmeijer et al. 2002, Di Lallo et al. 2003), this FtsK–FtsQ interaction is likely not dependent on any other identified proteins in the cell division pathway. Given these results in combination with those discussed above, it is reasonable to speculate that FtsQ, FtsL, FtsB, FtsW, and FtsI could assemble into a complex, which is then recruited to the septum via interaction of FtsQ with FtsK or that all of these proteins could possibly exist as a preassembled complex within the cell, even when the cell is not dividing.

Finally, we have presented preliminary evidence that multiple copies of FtsQ protein are localized together within the cell division complex and that this association of multiple FtsQ proteins is dependent on the amount of FtsK in the cell. One must interpret these results with some degree of caution as we have been unable to determine whether our data reflect the stabilization of FtsQ dimers or oligomers by FtsK. Models in which FtsK is capable of multimerization or in which a single FtsK molecule can recruit multiple FtsQ molecules are equally plausible interpretations of these data. Rather, we include these data only to raise the idea that the “divisome” complex could include multiple copies of the FtsQ protein, suggesting the divisome may, in fact, represent an assembly of subcomplexes.

Recent bacterial two-hybrid results suggested an unexpectedly high number of interactions among cell division proteins within the E. coli cell (Di Lallo et al. 2003). The interpretation of these results suggested that despite the linear assembly pathway in E. coli, the cell division proteins may actually be assembled in a cooperative manner into a large divisome complex and hence assembly may in fact be more similar to that in B. subtilis than originally thought. Our results suggest these two-hybrid interactions should be interpreted with caution as they do not necessarily reflect direct interactions that are required for recruitment or assembly of the ring complex. For example, although FtsI appears to interact with both FtsK and FtsA, neither protein appears to be required for FtsI localization provided FtsQ is localized efficiently. Hence, although our data support a view based on two-hybrid data that these proteins ultimately are assembled within a large divisome complex, assembly of the divisome in E. coli would seem to be driven ultimately by a specific subset of these observed interactions.

It is interesting that ZapA–FtsQ, although restoring localization of downstream cell division proteins [with the exception of FtsN] in cells depleted for FtsA and/or FtsK, does not restore cell division in these cells. This result implies that FtsK and FtsA perform critical functions at the division site, including but not necessarily limited to facilitating the localization of FtsN. If FtsA and FtsK were only required to facilitate localization of FtsK and FtsQ, respectively, restoring the localization to the septum of FtsQ and the proteins it interacts with would seem to be sufficient to rescue cells deficient for either protein. Whatever this function is, it must be lacking in cells in which cell division proteins are recruited by ZapA–FtsQ rather than by the endogenous pathway.

This method of recruiting proteins prematurely to the septum using ZapA should be applicable to the study of other cell division proteins. Presumably, any given cytoplasmic or transmembrane cell division protein could be targeted to FtsZ independently of normally required proteins. This would allow for analysis of recruitment by prematurely targeted cell division proteins as performed here for FtsQ. We think this approach will be particularly useful in analyzing the recruitment steps among the late proteins in the cell division pathway.

In trying to dissect the role of a particular protein in the cell division pathway, our lab and others have turned to genetics to isolate mutations in cell division proteins that affect localization, recruitment, and potential enzymatic activity. The analysis of such mutants in vivo is often complicated by the fact that these proteins are essential and it is often difficult to determine which of a protein’s functions are affected by a given mutation. The premature targeting method provides a potential tool for characterizing mutants affected in localization and recruitment. For example, one could analyze the behavior of mutant alleles of cell division genes when incorporated into a ZapA fusion protein [which would be predicted to restore function of mutant proteins specifically affected in localization] or when introduced into strains expressing ZapA fusions to downstream cell division proteins [which would be predicted to restore function of mutant proteins specifically defective in recruitment].

Materials and methods

Bacterial strains, plasmids, and media

The bacterial strains and plasmids used in this work are listed in Supplementary Table S1. All experiments were performed in NZY medium (Guzman et al. 1992). Antibiotics were added when appropriate at the indicated concentrations: ampicillin, 200 µg/ml [plasmid] or 25 µg/ml [chromosome], chloramphenicol, 10 µg/ml, kanamycin, 40 µg/ml, spectinomycin, 100 µg/ml [plasmid] or 50 µg/ml [chromosome], and tetracycline, 15 µg/ml. D-Glucose and L-arabinose were added at 0.2% to repress or induce the expression of genes under control of the PBAD promoter. IPTG [isopropyl-β-D-thiogalactoside] was added at the appropriate concentrations as indicated.

Standard laboratory techniques were used for cloning and analysis of DNA, PCR, electroporation, transformation, and P1 transduction (Miller 1992). Chromosomals constructs were integrated using λ InCh [Boyd et al. 2000]. Individual bacterial strains and plasmids were constructed as described or as indicated in Supplementary Table S1. To transduce GFP and CFP fusions integrated at the attB80 site into ftsK depletion strains, JOE563 was made trp [by transduction of a linked Tn5 (tetR)] and screening for tryptophan auxotrophy. The resulting strain was transduced with P1 lysates from trp strains carrying the appropriate attB80 integrated fusion constructs and transductants selected on minimal media. Resulting transductants were
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screened for expression of GFP/CFP fusions by growth in liquid NZY containing 10 μM IPTG. In all other strains, *att* integrated constructs were transduced by P1 using resistance to spectinomycin. *katt* integrated constructs were transduced by P1 using resistance to ampicillin (25 μg/mL). The *ftsA12* allele was transduced from the strain MM61 by linkage to *leu*::Tn10 (tet*), and resulting transductants were screened for growth sensitivity.

To construct pNG52 and pNG53, *zapA* (pNG52) or *yfp-zapa* (pNG53) was PCR-amplified from pFSG5 using primers NWG43 (YFP_ygfE_fwd, 5'-CGGATCTCCATCTAATATAGTCTGCACAACCCGTCGATA-3') or NWG44 (YgfE_Eco_fwd, 5'-CGGA TCGAATTCTGCTCAACACGGTTCTGGATTA-3') and NWG45 (YgfE_Xba_rev, 5'-CGGGGCTCTAGATTTTGTTAGTT-3'). The products were digested with EcoRI and XbaI and ligated into the same sites of pTrc99a- *his*, resulting in the construction of pNG52 and pNG53. For localization of GFP fusions under conditions of normal cell division, bacteria were grown to midlog phase at 37°C, washed twice in PBS before mounting. Cells were examined for fluorescence using an Axioskop 2 microscope (Zeiss) equipped with 63× or 100× plan-Apochromat oil immersion objective and a 100-W mercury lamp. Filter sets to visualize GFP [HQ:FITC/ Bodipy/Fluo3], EYFP, ECFP, and DAPI (UV) were from Chroma Technology Corp. Images were captured using an Orca-100 CCD camera (Hamamatsu Photonics) and Openlab (Improvison) and subsequently processed and analyzed in Openlab. All measurements were limited to intact cells exhibiting normal segregation of nucleoids as evidenced by DAPI staining. Final processing for presentation was performed in Adobe Photoshop.

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Premature targeting of a cell division protein to midcell allows dissection of divisome assembly in *Escherichia coli*

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