A new suite of tnaA mutants suggests that Escherichia coli tryptophanase is regulated by intracellular sequestration and by occlusion of its active site

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**Abstract**

**Background:** The *Escherichia coli* enzyme tryptophanase (TnaA) converts tryptophan to indole, which triggers physiological changes and regulates interactions between bacteria and their mammalian hosts. Tryptophanase production is induced by external tryptophan, but the activity of TnaA is also regulated by other, more poorly understood mechanisms. For example, the enzyme accumulates as a spherical inclusion (focus) at midcell or at one pole, but how or why this localization occurs is unknown.

**Results:** TnaA activity is low when the protein forms foci during mid-logarithmic growth but its activity increases as the protein becomes more diffuse, suggesting that foci may represent clusters of inactive (or less active) enzyme. To determine what protein characteristics might mediate these localization effects, we constructed 42 TnaA variants: 6 truncated forms and 36 missense mutants in which different combinations of 83 surface-exposed residues were converted to alanine. A truncated TnaA protein containing only domains D1 and D3 (D1D3) localized to the pole. Mutations affecting the D1D3-to-D1D3 interface did not affect polar localization of D1D3 but did delay assembly of wild type TnaA foci. In contrast, alterations to the D1D3-to-D2 domain interface produced diffuse localization of the D1D3 variant but did not affect the wild type protein. Altering several surface-exposed residues decreased TnaA activity, implying that tetramer assembly may depend on interactions involving these sites. Interestingly, changing any of three amino acids at the base of a loop near the catalytic pocket decreased TnaA activity and caused it to form elongated ovoid foci in vivo, indicating that the alterations affect focus formation and may regulate how frequently tryptophan reaches the active site.

**Conclusions:** The results suggest that TnaA activity is regulated by subcellular localization and by a loop-associated occlusion of its active site. Equally important, these new TnaA variants are immediately available to the research community and should be useful for investigating how tryptophanase is localized and assembled, how substrate accesses its active site, the functional role of acetylation, and other structural and functional questions.

**Keywords:** TnaA, tryptophanase, Cell pole, Protein localization, Enzyme regulation

**Background**

The enzyme tryptophanase hydrolyzes tryptophan to form indole, pyruvate and ammonia, and is produced by over 85 gram-positive and gram-negative bacteria [1,2]. The indole thus produced diffuses across cell membranes independent of specific transporters and affects the physiology of nearby members of the microbial community or their animal hosts [2-6]. For example, indole regulates such diverse microbial processes as motility, biofilm formation, antibiotic resistance, persister formation and host cell invasion [2]. Recent observations continue to underscore the biological significance of this compound, including the fact that some bacteria which do not themselves produce indole nevertheless detect its presence and become more antibiotic resistant [7,8], and...
in others indole inhibits quorum sensing and growth [9]. Finally, in situ bacterial production of indole helps create and maintain proper epithelial cell function in the mammalian intestine [10,11]. This broad range of intra- and inter-kingdom effects makes it important to understand more fully how indole synthesis is regulated.

In Escherichia coli, tryptophanase is encoded by the tnaA gene in the tnaCAB operon, which also includes the tryptophan-specific transporter TnaB [12-14]. Transcription of tnaAB is activated by the cAMP-CRP complex [15], and is induced by tryptophan via a Rho-dependent terminator and a TnaC leader peptide [16,17]. Induction of the tnaA gene and production of indole requires high concentrations of tryptophan, achieved mainly by the import of exogenous tryptophan via the TnaB transporter [18,19]. Functional TnaA is tetrameric, with a pyridoxal phosphate (PLP) coenzyme covalently linked to the Lys270 residue at each of the four active sites [20,21]. Incubation of the enzyme at low temperature results in reversible loss of PLP, with the tetramer disassociating into inactive dimers [22]. Two crystal structures of apo TnaA have been solved; both are tetramers in which each of the four TnaA proteins has a similar quaternary structure comprised of three subdomains: D1, D2 and D3 [23,24]. In turn, subdomains D1 and D3 form a small domain, with D2 forming a single large domain. Interactions between these two domains and with the other three TnaA subunits create the tightly assembled tetramer. The two extant TnaA crystal structures differ in that the catalytic pocket of one is closed [23] while that of the other is open [24]. The existence of these two forms suggests that the structure of TnaA is flexible, though the causes and consequences of this difference are unknown.

Recently, we found that wild type TnaA co-purifies with membrane vesicles derived from the cell poles of E. coli and that TnaA-sfGFP forms a single focus at midcell or at one of the poles during mid-log growth but becomes diffusely localized as the cells approach stationary phase [25]. Untagged TnaA competes with TnaA-sfGFP for polar localization, the foci do not have the characteristics of inclusion bodies, and TnaA-sfGFP is as active as the untagged protein with regard to indole production, all of which suggest that TnaA naturally localizes to the pole [25]. However, the mechanism that drives the formation of these foci is unknown, as is the question of whether they have a biological function.

Here, we show that indole production is associated with the disappearance of discrete TnaA foci and the concomitant rise of diffusely localized cytoplasmic TnaA. In tandem, we created six truncated variants of TnaA and 36 missense mutants in which different combinations of 83 surface-exposed polar amino acid residues were converted to alanine. Several variants displayed different localization characteristics and/or reduced enzymatic activities, suggesting that the sequestration and release of TnaA from intracellular foci may represent a novel post-translational mechanism to regulate tryptophanase activity. Finally, three mutations at the base of an external loop may alter TnaA activity by regulating the frequency with which this loop occludes the enzyme’s active site.

Results

Diffuse TnaA protein is functional

TnaA forms a single polar or midcell focus in all cells during mid-logarithmic growth, but the protein gradually diffuses into the cytoplasm as the culture nears stationary phase so that the majority of cells do not contain TnaA foci [25]. Interestingly, indole production is low during early exponential growth but increases dramatically as TnaA becomes more diffuse [25,26]. Under these same conditions, TnaA forms polar foci in 100% of the cells during exponential growth, but when these cells enter stationary phase TnaA diffuses so that only 31-58% of cells contain individual foci [25]. Thus, indole production may increase because more enzyme is made or because TnaA is moved from intracellular foci to become soluble and active in the cytoplasm. In these previous experiments, E. coli was grown in LB medium and transited rapidly from mid-logarithmic growth into stationary phase, so it was difficult to determine if the diffuse TnaA fraction was responsible for indole production. Therefore, to address this relationship, we incubated E. coli in M9 minimal medium to slow cell growth. We also examined TnaA function in E. coli GL619, a strain in which the chromosomal tnaA-sfGFP gene is controlled by a tufA promoter. In this strain, the TnaA-sfGFP fusion protein is produced steadily and constitutively [19], allowing us to observe the regulation of TnaA activity in the absence of any effects that might be caused by alterations in tnaA gene expression. As expected, TnaA-sfGFP formed single polar or mid-cell foci during early mid-logarithmic growth (Figure 1A and B, 2 h) but became completely diffuse as the cells entered stationary phase and thereafter (Figure 1B, 4 h and afterwards). Most interestingly, indole was not produced during mid-logarithmic growth (Figure 1C, 2 h) but was produced continuously after TnaA had become diffuse in stationary phase (Figure 1C, 4 h and afterwards). These results suggested that indole was produced by the diffuse TnaA fraction but was not produced by TnaA sequestered in a focus.

TnaA probably dimerizes before assembling into the functional tetramer. Thus, the diffuse and focus fractions might be composed of different multimeric forms of TnaA, and, if so, the formation of TnaA foci might regulate TnaA function. Unfortunately, repeated attempts to isolate and analyze intact foci were unsuccessful because the foci dissolved and disappeared when the cells were disrupted. This prevented us from
analyzing the multimerization status of TnaA in foci and also prevented us from determining the precise amount of soluble TnaA in cells as they progressed through the growth cycle. However, because these lysates did convert tryptophan to indole (not shown), the results indicated that foci were easily disrupted to give fully active TnaA-sfGFP tetramers.

The TnaA D1D3 two domain dimer localizes to cell poles

Because the TnaA in discrete foci could not be tested separately from the diffuse protein, we used a truncation and mutagenesis approach to address the question of how TnaA multimerization might affect enzyme localization and activity. We rearranged the tnaA gene to create a set of derivatives that encoded all six possible combinations of the D1, D2 and D3 protein subdomains (Figure 2). The sfGFP gene was appended to the 3′ end of each rearranged gene, and the fusions were inserted into the chromosome to replace the wild type tnaA gene. The resulting gene constructs were expressed under control of the native tnaA promoter and Shine-Dalgarno sequence. We modeled the structures of these TnaA variants by subtracting individual subdomains from the known crystal structure (Figure 3A). The truncations predicted the existence of three potential dimers (D1-D1, D3-D3 and D1D3-D1D3) (Figure 3A, upper right three structures) and three tetramers (created by self-association of the D2, D1D2 or D2D3 domains) (Figure 3A, lower right three structures). Note that the tetramer form of TnaA required the presence of the D2 domain, but that different dimers could assemble in its absence.

Cells expressing the D1, D1D2 and D1D3 proteins produced fluorescent products (Figure 3C), but little or no fluorescent signal was observed in cells carrying the D2, D3 and D2D3 fusion products (not shown), suggesting that these latter proteins were not expressed or were degraded. SDS-PAGE analysis of the fusion proteins was
consistent with these results (Figure 3B). Among the constructs, D1D3 was the most stable (e.g., exhibited the least degradation), D1D2 was of intermediate stability, while most of the D1 domain construct was degraded into smaller fragments (Figure 3B). The D2, D3 and D2D3 subdomain variants were not visible on SDS-PAGE, strongly suggesting that the D1 domain was essential for the folding or stability of TnaA. None of the truncation variants could produce indole (not shown), confirming that the complete TnaA protein was required for enzyme activity. The D1D3-sfGFP fusion protein formed a single focus at midcell or at one of the poles, as was observed for full length TnaA-sfGFP (Figure 3C). In contrast, the D1 and the D1D2 variants were diffuse throughout the cytoplasm (Figure 3C). Thus, the D1D3 dimer was sufficient for directing polar localization. Because the D1-only and D3-only variants were not stable, we could not determine directly if one or the other subdomain contained the major localization signal.

The polar localizations of TnaA and D1D3 are not artifactual. TnaA-sfGFP appears to localize specifically to cell poles because the protein does not co-localize with IbpA, which binds to inclusion bodies, and because TnaA-sfGFP foci are chased from the poles by untagged TnaA [25]. In addition, the dissipation of TnaA foci when the cells were broken open (see above) is unlike the behavior of inclusion bodies, which cohere and can be pelleted by

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**Figure 3** The D1D3 domain is sufficient for polar localization. A. Models of the oligomerization status of TnaA and the subdomain constructs. The four subunits in the TnaA tetramer are designated as TnaA$^1$, TnaA$^2$, TnaA$^3$, and TnaA$^4$. The deleted subdomains were removed from the TnaA tetrameric crystal structure (PDB 2OQX). The positions of the D1 (magenta), D2 (grey) and D3 (pink) subdomains in TnaA are illustrated in the upper right molecule of the tetramer. The other three TnaA molecules that make up the tetramer are shown in blue, cyan and orange. The four catalytic pockets are located at the left and right sides of the tetramer, and are not visible in this view. B. Protein stability of the subdomain constructs. E. coli strains expressing each TnaA truncation mutant were grown in LB to an OD$_{600}$ of 1.0 and analyzed by SDS-PAGE. The sfGFP fusion proteins were visualized by in-gel GFP fluorescence imaging. The top-most band in each lane represents the expected size of the respective full-length fusion protein. C. Cellular localization of the subdomain constructs. GFP and phase contrast images are overlaid.
low speed centrifugation. Nonetheless, to determine if the D1D3-sfGFP foci were inclusion bodies, the fusion protein was expressed in the presence of IbpA-mCherry, which binds to misfolded proteins and marks such artifacts [27,28]. Very little IbpA-mCherry signal was observed (not shown), indicating that few misfolded proteins were present to induce ibpA. In addition, when the fluorescence images were artificially enhanced, any IbpA-mCherry foci that were present did not overlap with the D1D3-sfGFP foci, confirming that the latter were not non-specific inclusion bodies (Figure 4A). In addition, we overproduced untagged wild type TnaA (from plasmid pTnaA) and found that this protein displaced polar D1D3-sfGFP foci so that the fusion protein was distributed diffusely throughout the cytoplasm (Figure 4B). These results strongly suggest that, like wild type TnaA, the localization of D1D3-sfGFP at the poles was real and specific.

Recent observations indicate that some fluorescent reporter proteins, including sfGFP, may skew the in vivo localization behavior of proteins on a case-by-case basis [29]. For example, sfGFP-tagged ClpP forms a single focus but untagged ClpP and a ClpP-mGFPmut3 fusion protein are instead spread diffusely throughout the cytoplasm [29]. Thus, to be doubly certain that polar localization of TnaA was not an artifact caused by attaching sfGFP, we appended monomeric mGFPmut3 to the carboxyl termini of TnaA and D1D3. As before, the fusion proteins localized to the cell poles (Figure 4C), providing further evidence that the polar localization of TnaA was not an artifact caused by adding the sfGFP reporter. In subsequent localization experiments, we observed the behavior of sfGFP-fused proteins because mGFPmut3 produced a much weaker fluorescent signal.

**Residues that affect assembly and localization of TnaA foci**
To identify the functional roles of the regions and residues of TnaA, we mutated 79 charged and polar residues on the surface of the D1D3 construct, reasoning that one or more of these residues might form ionic or hydrogen bonds with neighboring TnaA subunits or with other molecules. These residues covered almost the entire surface of the D1D3 protein (Figure 5A, residues colored yellow, blue, and cyan). Non-polar residues were not mutated, although they might contribute additional interactions (Figure 5A, residues in magenta). In addition, six lysine residues (Lys5, Lys450, Lys459 on the D1D3 domains; and Lys115, Lys156, Lys239 on the D2 domain) are known to be acetylated, though the physiological functions of this modification are not known [30,31]. Therefore, in addition to the lysine residues already changed on D1D3, we also mutated the three lysine residues on the D2 domain. Finally, we mutated the active site Lys270 on the D2 domain to determine if PLP or substrate binding might affect protein localization. In sum, 83 different residues were replaced with alanine. As a preliminary screen, we constructed TnaA variants containing one to four mutations near enough to one another that they could be altered by using a single oligonucleotide during site-directed mutagenesis (Tables 1, 2 and 3, mutants A1-TnaA to A32-TnaA). A corresponding series of mutagenized variants was engineered into the D1D3 fusion protein (Tables 1, 2 and 3, A1-D1D3 to A32-D1D3). sfGFP was fused to the carboxyl terminus of each TnaA or D1D3 variant, and the gene for each mutant was inserted into the original chromosomal position so that gene expression was controlled by the native tnaA promoter.

The localization and enzymatic phenotypes of the mutant proteins fell into three categories. In the first category, the mutant proteins behaved like wild type TnaA (Table 1). Each of these variants formed a single polar or
midcell focus during the mid-to-late growth phase, after which the proteins became diffuse as the cells entered stationary phase. In addition, these proteins produced indole at wild type levels, indicating that they had normal enzymatic activity. Of particular note is that neither localization nor enzyme activity was affected by mutating individual lysine residues that would normally be acetylated (Table 1, mutants A1, A10, A11, A12, A28 and A30). In fact, a mutant in which all six acetylated lysine residues were mutated continued to exhibit wild type localization and its enzymatic activity was affected very little (Table 1, mutant A33). Thus, many surface residues did not contribute significantly to TnaA function, and the purpose of lysine acetylation remains unknown.

The second mutant category consisted of TnaA variants that exhibited altered localization and/or enzymatic properties in cells as they reached the mid-to-late growth phase (Table 2). These mutants could be classified into three subgroups. The first subgroup consisted of full length proteins that exhibited delayed focus formation (Table 2, variants A2, A24 and A25). These variants were distributed diffusely in the cytoplasm during mid-log growth and formed a single polar or midcell focus about 30 min later than did wild type TnaA (e.g., Figure 6A). The mutated residues in these variants were located along the interface of neighboring D1D3-to-D1D3 domains (Figure 5A, deep blue residues), suggesting that interactions along the non-catalytic axis of the TnaA tetramer [32] might be affecting in vivo protein sequestration. The second subgroup, represented by only one mutant, localized normally when incorporated into the full length TnaA protein (Table 2, mutant A9, and Figure 6C). However, the corresponding A9-D1D3 truncation variant produced no foci but was, instead, distributed throughout the cytoplasm at all times (Table 2, and Figure 6D). The mutations in this variant were clustered tightly at the D1D3-D2 interface (Figure 5A, residues in cyan). However, the D2 domain was not present in this construct so that these residues were exposed on the surface of the A9-D1D3 truncation protein. Thus, these residues might be required for formation of dimers and foci. Finally, members of the third subgroup consisted of TnaA variants that produced highly elongated ovoid foci in the context of the full length mutant protein (Table 2, variants A21, A21a, A23, A23a, A31 and A31a). These ovoid foci (e.g., Figures 6E and 7A) were much different from the spherical foci associated with wild type TnaA and other variants (e.g., Figure 6C and F). These variants formed a single focus at midcell or at
Table 1 Mutations that do not affect TnaA localization or enzyme activity (Category 1)

| Variant | Residue(s) changed to alanine | Residue location | TnaA-sfGFP localization | D1D3-sfGFP Localization | Indole (\%) |
|---------|-----------------------------|-----------------|------------------------|-------------------------|-----------|
| A1      | K5                          | D1 surface      | Wild type              | N/A                     | 96.5      |
| A3      | E17, K20, R21               | D1 surface      | Wild type              | Wild type               | 101.3     |
| A4      | T23, R24, Y26               | D1 surface      | Wild type              | Wild type               | 99.5      |
| A6      | K33, S34                    | D1 surface      | Wild type              | Wild type               | 102.7     |
| A10     | K115                        | D2 surface      | Wild type              | N/A                     | 93.0      |
| A11     | K156                        | D2 surface      | Wild type              | N/A                     | 98.7      |
| A12     | K239                        | D2 surface      | Wild type              | N/A                     | 100.1     |
| A14     | N327, D329                  | D3 surface      | Wild type              | Wild type               | 96.8      |
| A15     | Q339, Y340, D343            | D3 surface      | Wild type              | Wild type               | 99.6      |
| A16     | E346, E347                  | D3 surface      | Wild type              | Wild type               | 93.1      |
| A18     | D363, K366                  | D3 surface      | Wild type              | Wild type               | 97.7      |
| A19     | H370, D374, Q375            | D3 surface      | Wild type              | Wild type               | 99.2      |
| A21b    | R403                        | D3 loop         | Wild type              | N/A                     | 103.3     |
| A21c    | D404                        | D3 loop         | Wild type              | N/A                     | 105.4     |
| A22     | K405, K409, Q410            | D3 loop         | Wild type              | Wild type               | 100.9     |
| A26     | E437, K440, H441            | D3 surface      | Wild type              | Wild type               | 92.7      |
| A27     | K443, E444, N445, N448      | D3 surface      | Wild type              | Wild type               | 103.4     |
| A28     | K450                        | D3 surface      | Wild type              | N/A                     | 97.9      |
| A30     | K459                        | D3 surface      | Wild type              | N/A                     | 102.9     |
| A31c    | T465                        | D3 surface      | Wild type              | N/A                     | 72.7      |
| A32     | K467, K469, E470            | D3 surface      | Wild type              | Wild type               | 92.4      |
| A33     | K5, K115, K156, K239, K450, K459 | Surface | Wild type              | Wild type               | 82.5      |

*Residue location is based on the crystal structure of TnaA (PDB ID 2OQX). Pocket, the active site cavity; loop, the loop comprised of aa 398 to 416 at the edge of the catalytic pocket.

*Wild type: spherical focus at midcell or one of the poles at mid-to-late exponential phase, becoming more diffuse at stationary phase as wild type TnaA.

*Wild type: spherical focus at midcell or one of the poles at mid-to-late exponential phase. D1D3 constructs were mostly degraded at stationary phase.

*The amount of indole produced by each TnaA-sfGFP variant compared to that produced by wild type TnaA in LB at an OD$_{600}$ of 1.5. Values are reported as the percentage of wild type, and are the average of two independent cultures. Growth rates of all cells carrying wild type TnaA or any of its variants were nearly identical.

*N/A indicates that the variant was not constructed or that data was not available.

Table 2 Mutations that alter TnaA focus formation during exponential growth (Category 2)

| Variant | Residue(s) changed to alanine | Residue location | TnaA-sfGFP localization | D1D3-sfGFP Localization | Indole (\%) |
|---------|-----------------------------|-----------------|------------------------|-------------------------|-----------|
| A2      | E9, R12, R14                | D1D3$^{-1}$-D1D3$^{2}$ interface | Delayed               | Wild type              | 7.2       |
| A24     | T426, Y427, T428            | D1D3$^{1}$-D1D3$^{2}$ interface | Delayed               | Wild type              | 8.5       |
| A25     | Q429, T430, H431, D433      | D1D3$^{1}$-D1D3$^{2}$ interface | Delayed               | Wild type              | 13.3      |
| A9      | T60, Q61, S62, Q64          | D1D3$^{1}$-D2$^{3}$ interface | Wild type             | Diffuse                | 5.3       |
| A21     | S398, R403, D404            | Edge of the pocket | Ovoid                  | Wild type              | 5.8       |
| A21a    | S398                        | Bottom of the loop | Ovoid                  | Wild type              | 43.4      |
| A23     | E416, R419                  | Edge of the pocket | Ovoid                  | Wild type              | 5.1       |
| A23a    | E416                        | Bottom of the loop | Ovoid                  | N/A$^a$                | 12.1      |
| A31     | R462, H463, T465            | Edge of the pocket | Ovoid                  | Wild type              | 2.1       |
| A31a    | R462                        | Bottom of the loop | Ovoid                  | N/A                    | 57.1      |

Legend is the same as for Table 1.
one pole, but the assemblies were elongated and oriented parallel to the long axis of the cell or, in some cases, were slightly tilted relative to the poles (Figures 6E and 7A). The mutations in these variant proteins clustered near the edge of the catalytic pocket, though they were not themselves part of the active site. A surprising observation was that some full-length TnaA variants localized differently than did their corresponding D1D3 truncation proteins (Table 2). For example, unlike their full length counterparts, focus formation was not

Table 3 Mutations that delay TnaA focus disassembly in stationary phase (Category 3)

| Variant | Residue(s) changed to alanine | Residue location | TnaA-sfGFP localization | D1D3-sfGFP localization | Indole<sup>d</sup> (%) |
|---------|------------------------------|-----------------|-------------------------|-------------------------|-----------------------|
| A2      | E9, R12, R14                 | D1D3<sup>i</sup>-D1D3<sup>j</sup> interface | Delayed                | Wild type               | 7.2                   |
| A5      | R27, E28, E29                | D1 surface      | Wild type               | Wild type               | 14.9                  |
| A7      | D42, S43, E44, D45           | D1 surface      | Wild type               | Wild type               | 36.8                  |
| A8      | D49, T52, D53, S54           | D1D3<sup>i</sup>-D2<sup>j</sup> | Wild type               | Wild type               | 6.6                   |
| A9      | T60, Q61, S62, Q64           | D1D3<sup>i</sup>-D2<sup>j</sup> | Wild type               | Diffuse                 | 5.3                   |
| A13     | K270                         | D2 active site  | Wild type               | N/A                     | 10.9                  |
| A17     | C352, Q353, Q354             | D3 pocket       | Wild type               | Wild type               | 6.0                   |
| A20     | E384, K387, R392             | D3 surface (EK) | Wild type               | Wild type               | 4.8                   |
| A21     | S398, R403, D404             | Edge of the pocket | Ovoid                 | Wild type               | 5.8                   |
| A21a    | S398                         | Bottom of the loop | Ovoid                | Wild type               | 43.4                  |
| A29     | T453, T455, Y456, E457       | D3 surface      | Wild type               | Wild type               | 11.7                  |

Legend is the same as for Table 1.

Figure 6 Altered localization of TnaA and D1D3 variants. Images of altered localization, one representative variant from each subgroup in Table 2. The TnaA variants are shown in A, C and E, and their corresponding D1D3 variants are shown in B, D and F. The cells were grown in LB for 2.5 or 3 h (OD<sub>600</sub> ~ 0.6 and 1.0, respectively).
delayed for the A2-D1D3, A24-D1D3 and A25-D1D3 truncation variants (Table 2 and Figure 6B). Also, as stated above, mutations at the D1D3-D2 interface produced a diffuse localization of A9-D1D3 (Figure 6D) but the A9-TnaA full-length variant continued to localize normally at the poles (Figure 6C). Finally, the mutations that caused full length TnaA-sfGFP to form elongated ovoid foci had no effect on the shape of foci produced by the truncation proteins A21-D1D3, A23-D1D3 or A31-D1D3 (Table 2, Figure 6F). These results suggest that polar localization is a complicated process with multiple interactions involving both the D1D3 and D2 domains.

The third major mutant category was composed of TnaA variants for which the disassembly of foci was delayed during stationary phase (Table 3). Previously, we showed that when wild type TnaA is expressed from its native promoter, the protein becomes more diffuse when
entering stationary phase [25], an observation that was confirmed here because TnaA foci begin to disappear in early stationary phase cells (e.g., after 4–6 h, Table 4). In contrast, this third set of mutant proteins continued to form polar foci in almost 100% of cells at 6 h and began to become diffuse only at 12 h (Tables 3 and 4). Three of these variants also displayed altered focus formation (Table 2), suggesting that in some cases common factors may affect both assembly and disassembly of foci. In addition, A13-TnaA, which carries a Lys270Ala substitution at the active site, localized at poles even though it had no enzymatic activity (i.e., indole was produced at background levels) (Table 3, mutant A13). Other mutations in the catalytic pocket also eliminated indole production but had no effect on polar localization (e.g., A17-TnaA, Table 3). These results indicated that localization did not depend on the presence of the PLP cofactor or on the ability of TnaA to hydrolyze tryptophan. However, altering some active site residues did affect the disassembly of foci during stationary phase.

We particularly wished to know if delayed disassembly of TnaA foci was associated with a delay in the development of enzymatic activity, as measured by the production of indole. Therefore, we examined four of these variants (A2, A9, A17 and A21a) for enzymatic activity and for the presence of TnaA-sfGFP foci (Table 4). Each of the variants described in Tables 3 and 4 produced less indole at all points in the growth cycle, even though the amount of each variant protein was 2- to 3-fold higher per cell after 12 h incubation (not shown). In terms of TnaA localization, the selected proteins exhibited three types of behavior. First, the A21a variant most closely replicated the cycle of wild type TnaA-sfGFP. Although A21a foci disappeared later than did wild type TnaA foci, after 12 h the number of foci and enzymatic activity paralleled the results observed for wild type TnaA (Table 4, variant A21a). The A21a variant carries the S398A mutation, which is unable to close the peptide loop adjacent to the active site (Figure 7C). This implies that although the loop may inhibit substrate access to the active site (see Discussion), it may not be required for enzymatic activity per se. The A2 and A9 variants displayed a second type of behavior. For each of these proteins the number of foci decreased at 12 h, similar to wild type TnaA, but the enzymatic activity of the mutants remained low (Table 4). Therefore, these mutations may affect both localization and enzymatic activity. Finally, the A17 variant exhibited a third class of behavior – the number of cells with intact foci remained at almost 100%, even at 12 h, while enzymatic activity also remained low. This phenotype was consistent with the possibility that sequestered TnaA is inactive (see Discussion), though more work would have to be done to prove that this protein was active outside of foci.

### Three loop-associated substitutions produce elongated ovoid foci

We were very intrigued by the formation of elongated ovoid foci in the A21-TnaA, A23-TnaA and A31-TnaA variants. Each of these proteins contained two to four amino acid substitutions, so we constructed single substitution variants to determine the residues responsible for this curious phenomenon. Each of three alanine substitutions replacing Ser398 (A21a-TnaA), Glu416 (A23a-TnaA) or Arg462 (A31a-TnaA) resulted in protein variants that formed elongated ovoid foci and decreased indole production by 43-78% (Table 2, Figure 7A). These altered localizations and enzyme activities were not due to decreased protein stability (Figure 8).

The three relevant residues mapped to the bottom of a loop, encompassing residues 398–416, located just to the side of the TnaA catalytic pocket (Figure 7B). This loop obscures the catalytic pocket in the closed conformation of TnaA (Figure 7B, right), but is cantled away from the active site cavity in the open conformation of the protein (Figure 7B, left), making it likely that movement of this loop shifts the enzyme between its open-and-closed forms. The three residues are in close contact, possibly forming hydrogen bonds to maintain the position of this loop (Figure 7C). Surprisingly, the local arrangement of these residues and the loop itself in the closed form of TnaA are almost identical to the arrangements in the open form, suggesting that the movement of the loop is mediated mainly by moving the D1D3 domain towards the D2 domain. Mutations of the three residues may alter the position or orientation of the loop or may disrupt how the loop is tethered to the body of TnaA, thereby altering the protein’s enzymatic activity and localization. Interestingly, mutations that changed the composition of the loop itself (e.g., in variants A21b-TnaA, A21c-TnaA and A22-TnaA) did not affect polar localization or enzymatic activity (Table 1), suggesting that protein function

### Table 4 Indole activity of selected disassembly-delay mutants of TnaA

| Variant | Cells with foci (%) | Indole production (mM) |
|---------|---------------------|------------------------|
|         | 2 h  | 3 h  | 4 h  | 6 h  | 12 h | 2 h  | 3 h  | 4 h  | 6 h  | 12 h |
| Wild type | 100  | 100  | 100  | 75   | 67   | 0.02 | 0.25 | 0.55 | 0.59 | 0.50 |
| A2      | 41   | 100  | 100  | 100  | 64   | 0.00 | 0.04 | 0.04 | 0.13 | 0.05 |
| A9      | 100  | 100  | 100  | 100  | 72   | 0.01 | 0.05 | 0.03 | 0.12 | 0.07 |
| A17     | 100  | 100  | 100  | 100  | 100  | 0.02 | 0.06 | 0.03 | 0.16 | 0.15 |
| A21a    | 100  | 100  | 100  | 100  | 87   | 0.01 | 0.06 | 0.24 | 0.43 | 0.45 |

*The number of foci and enzyme activity of specified mutants of TnaA-sfGFP fusion proteins.
depends more on loop placement than on loop composition. The results imply that the mutated residues at the base of the loop may alter the equilibrium between the open and closed conformations of TnaA by affecting loop dynamics.

**Discussion**

A noteworthy number of proteins are distributed asymmetrically in bacterial cells, and these different locations may affect protein activity or function [33,34]. Such relationships are most obvious for proteins within macromolecular complexes, such as those that comprise the divisome, flagella, secretion systems and the chromosomal segregation apparatus [35-42]. It is less clear how or why other proteins are directed to specific locations, or whether differential localization affects protein function. However, a few examples lend credence to the idea that cellular localization causes or is at least associated with alterations in enzymatic activity. For example, the *E. coli* MurG protein participates in synthesizing peptidoglycan precursors for insertion into the cell wall, and excess amounts of this protein are localized to the cell poles, where the enzyme is apparently inactive [43]. Similarly, the activity of the *Bacillus subtilis* LicT antiterminator is accompanied by an overt change in its cellular localization: inactive enzyme (in the absence of the substrate salicin) is distributed diffusely in the cytoplasm, whereas active enzyme (in the presence of salicin) moves to a subpolar localization [44]. Finally, the *Caulobacter crescentus* CtpS protein, a CTP synthase, performs its enzymatic role when diffuse in the cytoplasm but also helps determine cell shape by polymerizing into a cytoskeletal filament along the cell membrane, this latter function being independent of its enzymatic activity [45]. We now report that the cellular location and activity of the TnaA tryptophanase are correlated, suggesting that the sequestration of TnaA in discrete foci may regulate its enzyme activity.

A strong indication that TnaA localization influences its enzymatic activity is that *E. coli* produces very little indole when TnaA is located in foci but that it produces large amounts of indole when the foci disappear and TnaA becomes diffuse throughout the cytoplasm. This suggests that the fraction of TnaA in foci is inactive and that the diffuse population is functional. A second argument supporting a relationship between protein function and focus formation is the existence of three mutations that reduce enzyme activity and change the geometry of TnaA foci from spherical to elongated ovals. The mechanism that creates these effects is unknown but the three altered residues are located at the base of a loop near the catalytic cavity, indicating that the disposition of this loop affects both characteristics. A third argument in support of the focus-function premise is that several mutations (i.e., in the mutants A2, A24 and A25) alter the kinetics or timing of focus formation and also reduce enzyme activity. The location of these mutated residues at the D1D3-to-D1D3 dimer interface, well away from the catalytic pocket, puts them in a position to affect the equilibrium among the different multimeric forms of TnaA, which may affect both sequestration and activity of the enzyme. Finally, several mutants delay the time when TnaA moves from foci to become diffuse while also delaying the time at which the enzyme becomes active. This strong genetic correlation suggests the two characteristics are causally related and corroborates the relationships observed for the wild type protein.

Overall, the results are consistent with a model in which individual foci contain inactive TnaA monomers or dimers while the diffuse fraction contains active tetramers (Figure 9). If true, then *E. coli* may regulate TnaA activity by altering the equilibrium between these protein states. This model is supported by previous work in which the TnaA C298S mutant, which is near the active site, disassociates more readily into inactive dimers, thus reinforcing the relationship between activity and tetramerization [32]. The fact that focus assembly is reversible, both in *vivo* during growth and in *vitro* when cells are disrupted, suggests that one or more cellular factors might help orchestrate the focus-to-diffuse shift. The physiological purpose of this regulation might be to restrict indole signaling to a particular time late in the growth cycle.

One of the most intriguing findings was that three individual missense mutations decrease the tryptophanase...
activity of TnaA and are associated with the formation of curiously elongated ovoid foci. The affected amino acids (Ser398, Glu416 and Arg462) are located at the base of a loop (from residues 398-416) that sits to one side of the catalytic pocket. In the two crystal structures of TnaA, this catalytic site is closed in one conformation and open in the other. The major cause of the closed pocket is a movement of the small domain (D1D3) towards the large domain (D2), re-orientating the loop so that it controls access to the active site by physically occluding the catalytic cavity. The three mutations that do change the behavior of TnaA foci are perfectly positioned to affect the orientation of the loop and may regulate the equilibrium between the open and closed versions of TnaA. Such alterations may also favor intermolecular interactions that lead to the formation of oddly shaped foci. Alternately, the assembly of TnaA into foci may drive the loop to adopt one or the other orientation, which could determine whether the active site is more or less available to its substrate. Mutations affecting residues within the loop itself did not alter polar localization or enzymatic activity, implying that the loop’s orientation is the more important regulatory feature. However, combining any one of the three mutations at the base of the loop with mutations within the loop produced enzymes with even lower activity, suggesting that once the base is altered additional considerations come into play (Table 2, compare variants A21a, A23a and A31a to variants A21, A23 and A31). The tryptophanase of Proteus vulgaris and the tyrosine phenol-lyases of Citrobacter freundii and Erwinia herbicola are 51.7%, 41.1% and 43% identical to E. coli TnaA, respectively (EMBOSS Needle software analysis, via http://www.ebi.ac.uk/Tools/psa/emboss_needle/index.html), and all three have crystal structures with quaternary structures similar to that of E. coli TnaA (PDB entries 1AX4, 2EZ1 and 1C7G). Significantly, in all three enzymes the residues homologous to Ser398, Glu416 and Arg462 in E. coli TnaA are conserved and map to the base of a protruding loop near the active site, suggesting that the loop might play a similar structural and regulatory role in these enzymes, as well.

Lastly, we again address the question of whether polar assembly is an artifact caused by tagging TnaA with the fluorescent protein sfGFP. Mis-localization of analogous constructs made with the Clp protease [29] and MreB [46] does not mean that all protein fusion localizations are artifactual. The lesson is that such results need to be interpreted with caution when this kind of tagging is the sole approach for studying protein localization. In the case of TnaA, additional evidence suggests that polar assembly of TnaA-sfGFP is specific and likely represents its native localization [25]. Such evidence includes the fact that untagged TnaA associates with membrane vesicles derived from poles, that it competes with TnaA-sfGFP for polar localization, and that TnaA-sfGFP foci do not overlap with inclusion bodies [25]. Here, we find that the TnaA-sfGFP foci disappear when cells are lysed, again indicating that these are not inclusion bodies. Finally, when we fused TnaA to monomeric GFP (mGFPmut3) as recommended by Landgraf et al. [29], the composite protein formed the same type of polar foci, providing additional evidence that localization is not due to artifacts caused by sfGFP.

**Conclusions**

The subcellular location and enzymatic activity of the TnaA tryptophanase are correlated, strongly suggesting...
| Strain | Description* | Source or Reference |
|--------|--------------|---------------------|
| MG1655 | Wild type F’ λ ilvG rfb-50 rph-1 | Laboratory collection |
| GL38   | MG1655 aer-sfgfp::kan | [25] |
| GL40   | MG1655 tnaA-sfgfp::kan | [25] |
| GL60   | MG1655 ibpA-mCherry::kan | [25] |
| GL504  | MG1655 ΔtnaA::frt | [25] |
| GL619  | GL607 ΔmaA::frt PtnA-sfgfp tna8::frt | [19] |
| GL676  | MG1655 tnaA-mgfpmut3::kan | This study |
| GL678  | MG1655 D1-sfgfp::kan | This study |
| GL680  | MG1655 D1D2-sfgfp::kan | This study |
| GL682  | GL504 D3-sfgfp::kan | This study |
| GL684  | GL504 D2D3-sfgfp::kan | This study |
| GL686  | GL504 D1D3-sfgfp::kan | This study |
| GL688  | GL504 D2-sfgfp::kan | This study |
| GL726  | GL726 D1D3-sfgfp::frt ibpA-mCherry::kan | This study |
| GL730  | GL726 D1D3-sfgfp::frt | This study |
| GL734  | GL504 D1D3-mgfpmut3::kan | This study |
| GL801  | GL504 A1-tnaA-sfgfp::kan (K5A) | This study |
| GL802  | GL504 A2-tnaA-sfgfp::kan (E9A, R12A, R14A) | This study |
| GL803  | GL504 A3-tnaA-sfgfp::kan (E17A, K20A, R21A) | This study |
| GL804  | GL504 A4-tnaA-sfgfp::kan (T23A, R24A, Y26A) | This study |
| GL805  | GL504 A5-tnaA-sfgfp::kan (R27A, E28A, E29A) | This study |
| GL806  | GL504 A6-tnaA-sfgfp::kan (K33A, S34A) | This study |
| GL807  | GL504 A7-tnaA-sfgfp::kan (D42A, S43A, E44A, D45A) | This study |
| GL808  | GL504 A8-tnaA-sfgfp::kan (D49A, T52A, D53A, S54A) | This study |
| GL809  | GL504 A9-tnaA-sfgfp::kan (T60A, Q61A, S62A, Q64A) | This study |
| GL810  | GL504 A10-tnaA-sfgfp::kan (K115A) | This study |
| GL811  | GL504 A11-tnaA-sfgfp::kan (K156A) | This study |
| GL812  | GL504 A12-tnaA-sfgfp::kan (K239A) | This study |
| GL813  | GL504 A13-tnaA-sfgfp::kan (K270A) | This study |
| GL814  | GL504 A14-tnaA-sfgfp::kan (N327A, D329A) | This study |
| GL815  | GL504 A15-tnaA-sfgfp::kan (Q339A, Y340A, D343A) | This study |
| GL816  | GL504 A16-tnaA-sfgfp::kan (E346A, E347A) | This study |
| GL817  | GL504 A17-tnaA-sfgfp::kan (C352A, Q353A, Q354A) | This study |
| GL818  | GL504 A18-tnaA-sfgfp::kan (E363A, K366A) | This study |
| GL819  | GL504 A19-tnaA-sfgfp::kan (H370A, D374A, Q375A) | This study |
| GL820  | GL504 A20-tnaA-sfgfp::kan (E384A, K387A, R392A) | This study |
| GL821  | GL504 A21-tnaA-sfgfp::kan (S398A, R403A, D404A) | This study |
| GL822  | GL504 A22-tnaA-sfgfp::kan (K406A, K409A, Q410A) | This study |
| GL823  | GL504 A23-tnaA-sfgfp::kan (E416A, R419A) | This study |
| GL824  | GL504 A24-tnaA-sfgfp::kan (T426A, Y427A, T428A) | This study |
| GL825  | GL504 A25-tnaA-sfgfp::kan (Q429A, T430A, H431A, D433A) | This study |
| GL826  | GL504 A26-tnaA-sfgfp::kan (E437A, K440A, H441A) | This study |
| GL827  | GL504 A27-tnaA-sfgfp::kan (K443A, E444A, N445A, N448A) | This study |
| GL828  | GL504 A28-tnaA-sfgfp::kan (K450A) | This study |
that the sequestration of TnaA in discrete foci may regulate its enzyme activity. In addition, altering any one of three residues at the base of a loop near the catalytic pocket produces aberrant TnaA foci that exhibit greatly decreased tryptophanase activity. These latter results strongly imply that the disposition of this loop controls the formation of TnaA foci and may regulate the rate at which tryptophan can access the active site.

| GL829 | GL504 A29-tnaA-sfgfp::kan (T453A, T455A, Y456A, E457A) | This study |
| GL830 | GL504 A30-tnaA-sfgfp::kan (K459A) | This study |
| GL831 | GL504 A31-tnaA-sfgfp::kan (R462A, H463A, T465A) | This study |
| GL832 | MG1655 A32-tnaA-sfgfp::kan (K467A, K469A, E470A) | This study |
| GL833 | GL504 A21a-tnaA-sfgfp::kan (S398A) | This study |
| GL834 | GL504 A21b-tnaA-sfgfp::kan (R403A) | This study |
| GL835 | GL504 A21c-tnaA-sfgfp::kan (D404A) | This study |
| GL836 | GL504 A23a-tnaA-sfgfp::kan (E416A) | This study |
| GL837 | GL504 A23b-tnaA-sfgfp::kan (R419A) | This study |
| GL838 | GL504 A31a-tnaA-sfgfp::kan (R462A) | This study |
| GL839 | GL504 A31b-tnaA-sfgfp::kan (H463A) | This study |
| GL840 | GL504 A31c-tnaA-sfgfp::kan (T465A) | This study |
| GL844 | GL504 A33-tnaA-sfgfp::kan (KSA, K115A, K156A, K239A, K450A, K459A) | This study |

*frt* indicates the presence of the following oligonucleotide scar that was left after removing the kan cassette: GAAGTTCCTATACTTTCTAGAGAATAGGAACTTC. The "A##" numbers correspond to the site-directed mutations described in Tables 1 and 2.
Methods

Strains, plasmids and growth conditions

All strains were derivatives of *E. coli* MG1655, and are listed in Table 5. Strain GL60 was the source for the *ibpa-mCherry-kan* allele [25]. The plasmid pTnaA (KanR) carries the wild type *tnaA* gene under control of the *lac* promoter [25]. The plasmid pJD1 (AmpR) contains *torA-gfpmut3* and was used as the template to amplify *gfpmut3* [47]. Strains were selected by adding kanamycin (Kan, 50 μg/ml) or ampicillin (Amp, 100 μg/ml). Cells were grown overnight in LB medium supplemented with 0.4% glucose. Overnight cultures were diluted 1:100 into 2 ml LB or M9 minimal medium supplemented with 2 mM MgSO₄, 0.1 mM CaCl₂, 33 μM thiamine, 1% Bacto Casamino acids and 0.5 mM tryptophan. Cells were incubated with shaking at 37°C for TnaA localization and indole production experiments. When applicable, 50 μM IPTG was added to induce expression of the plasmid-encoded *tnaA* gene.

Strain construction

Chromosomal gene deletions or insertions were performed by using λ-Red recombination [48]. The DNA fragments were amplified by one-step or sequential PCR using designated primers (see Additional file 1). To construct the D1-sfGFP and D1D2-sfGFP strains, the *sfgfpl::kan* cassettes were amplified from the chromosome of *E. coli* GL38 (*aer-sfGFPl::kan*) and inserted into the chromosome of MG1655 to delete the appropriate subdomain fragments and to fuse *sfgfpl* to D1 and D1D2. To construct the D2-sfGFP, D3-sfGFP, D2D3-sfGFP and D1D3-sfGFP strains, the respective *sfgfpl*-containing fragments were amplified from the chromosome of GL680 (D1D2-*sfgfpl::kan*) or GL40 (*tnaA-sfGFPl::kan*) and inserted into the chromosome of GL504 (Δ*tnaA::*frt) under control of the original *tna* promoter. The A1-TnaA-sfGFP to A31-TnaA-sfGFP strains were constructed by amplifying the *tnaA-sfGFPl::kan* fragments from the chromosome of GL40 and inserting them into the chromosome of GL504 at the original *tnaA* gene location. To construct the A32-TnaA-sfGFP strain, the *sfgfpl* fragment was amplified from the chromosome of *E. coli* GL38 and inserted into the chromosome of MG1655 to introduce the mutation and to fuse *sfgfpl* to *tnaA*. To construct the A33-TnaA-sfGFP strain, the DNA fragment was generated by six rounds of PCR reactions to introduce the mutations for the six codons one by one. The A1-D1D3-sfGFP to A32-D1D3-sfGFP fusions were constructed in the same way as described for the D1D3-sfGFP fusions, except that the chromosomes of corresponding A1-TnaA-sfGFP to A32-TnaA-sfGFP strains were used as the templates for PCR. To construct the TnaA-mGFPmut3 fusion, the fragment encoding mGFPmut3 was amplified from *gfpmut3* in pJD1 by using two-step PCR to introduce the A206K mutation [29]. The *mGFPmut3* fragment was joined to the *kan* cassette by an additional PCR and then inserted into the chromosome of MG1655 to replace the stop codon of *tnaA*. The D1D3-mGFPmut3 fusion was constructed as described for the D1D3-sfGFP fusion, except that the relevant gene fragment was amplified from the chromosome of GL676 (*tnaA-mGFPmut3::kan*). The entire *tnaA* region in all strains was PCR amplified and sequenced to confirm the presence of the correct gene constructions.

TnaA structure modeling

The crystal structures of TnaA (open conformation, PDB 2OQX; closed conformation, PDB 2C44) were assembled into tetramers by using the PISA software (http://www.ebi.ac.uk/pdbe/prot_int/pistart.html) [49]. In silico analysis and visualization of TnaA structures were performed by using the PyMOL Molecular Graphics System, Version 1.3. Schrödinger, LLC (http://www.pymol.org).

Microscopy

Phase contrast and fluorescence images of cells were obtained by using an Olympus microscope BX60 with a 100× oil objective (1.3 NA PH3), a 1.4 MP MONO CCD camera XM10, and illuminated by the fluorescence source X-Cite120. Live cells were loaded to 1% agarose-coated slides for imaging. GFP was visualized by using the OSF-0008Z filter set (471 nm excitation/520 nm emission).

TnaA enzyme activity and protein stability

TnaA activity was assayed by measuring indole production [19]. Standard curves with known concentrations of indole confirmed that the assay was linear for indole amounts down to 0.01 mM. Samples were collected at different points during cell growth and the amounts of indole were determined as described previously [19]. Cellular proteins were separated by SDS-PAGE, and TnaA and its various protein fragments that had been fused to sfGFP were visualized by in-gel GFP fluorescence imaging [19].

Additional file

Additional file 1: Table S1. Title of dataset: Oligonucleotides used to construct mutant genes. Description of dataset: list of oligonucleotides used in making mutant genes.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

GL and KDY conceived the project, devised the experiments, analyzed the data, interpreted the results and wrote the paper. GL performed the experiments. Both authors read and approved the final manuscript.
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