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A signal sequence suppressor mutant that stabilizes an assembled state of the twin arginine translocase

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28 **Abstract**

29 The twin-arginine protein translocation (Tat) system mediates transport of folded proteins

30 across the cytoplasmic membrane of bacteria and the thylakoid membrane of chloroplasts.

31 The Tat system of *Escherichia coli* is made up of TatA, TatB and TatC components. TatBC

32 comprise the substrate receptor complex, and active Tat translocases are formed by the

33 substrate-induced association of TatA oligomers with this receptor. Proteins are targeted to

34 TatBC by signal peptides containing an essential pair of arginine residues. We isolated

35 substitutions, locating to the transmembrane helix of TatB that restored transport activity to

36 Tat signal peptides with inactivating twin arginine substitutions. A subset of these variants also

37 suppressed inactivating substitutions in the signal peptide binding site on TatC. The

38 suppressors did not function by restoring detectable signal peptide binding to the TatBC

39 complex. Instead, site specific crosslinking experiments indicate that the suppressor

40 substitutions induce conformational change in the complex and movement of the TatB subunit.

41 The TatB F13Y substitution was associated with the strongest suppressing activity, even

42 allowing transport of a Tat substrate lacking a signal peptide. *In vivo* analysis using a TatA-

43 YFP fusion showed that the TatB F13Y substitution resulted in signal peptide independent

44 assembly of the Tat translocase. We conclude that Tat signal peptides play roles in substrate

45 targeting and in triggering assembly of the active translocase.

46

47 **Significance statement**

48 The twin-arginine translocation (Tat) system transports folded proteins across the prokaryotic

49 inner membrane and the thylakoid membrane of plant chloroplasts. Proteins are targeted to

50 the Tat system by signal peptides containing a highly conserved twin arginine motif. We

51 isolated suppressors in the TatB component that allowed a Tat substrate with a defective twin

52 arginine motif to be transported. The strongest of these suppressors, TatBF13Y, resulted in

53 the constitutive assembly of the Tat translocase in the absence of signal peptide binding.

54 These results show that Tat signal peptides have two separable roles – they target their

55   passenger proteins to the Tat machinery but they also trigger the assembly of the active Tat

56   transporter.

57

58   \body

59   **Introduction**.

60   A large proportion of prokaryotic proteins are trafficked into or across the cytoplasmic

61   membrane. Extracytoplasmic proteins are synthesized with cleavable N-terminal signal

62   peptides to address them to export machineries located in the cytoplasmic membrane. Signal

63   peptides are generally between 20-30 amino acids in length and have a recognizable tripartite

64   structure comprising a basic n-region, a hydrophobic h-region and a polar c-region with a

65   signal peptidase cleavage site.

66   The Sec pathway is the major route of protein export in most prokaryotes, transporting

67   unfolded polypeptides across the cytoplasmic membrane. In bacteria it is comprised of a

68   SecYEG channel complex and a peripheral membrane ATPase, SecA (see (1, 2) for recent

69   reviews). The initial discovery of Sec components was driven by genetic approaches using

70   *Escherichia coli* to isolate suppressors of defective Sec substrate proteins and inactive signal

71   peptides (e.g. 3, 4). These genetic suppressors also contributed significantly to mechanistic

72   understanding of Sec-dependent protein translocation (5-7). More recently it has been shown

73   that Sec signal peptides have dual roles – they serve to target their passenger proteins to the

74   Sec machinery (3, 8) but also to allosterically activate the SecY channel (9).

75   The twin-arginine protein translocation (Tat) pathway operates in parallel to the Sec pathway

76   to transport folded proteins across the prokaryotic cytoplasmic membrane and the thylakoid

77   membrane of plant chloroplasts (10, 11). Proteins are targeted to the Tat pathway by N-

78   terminal signal peptides that contain a conserved twin arginine motif at the n-region/h-region

79   boundary (12, 13; Fig 1A). The Tat system in the model bacterium *E. coli* requires three

80   membrane-bound subunits, TatA, TatB and TatC (14-17). The TatB and TatC proteins form a

81   multivalent complex that binds Tat substrates through their twin arginine signal peptides (e.g.

82   (18-20). Numerous experiments have shown that the TatC component recognizes the twin

83   arginine motif (21-26) whereas TatB is close to the signal peptide h-region (27, 28). Signal

84   peptides have been shown to penetrate deeply into the TatBC complex (29) and in thylakoids

4

85　at least this deep-binding mode may be modulated by the transmembrane proton

86　electrochemical gradient (PMF) (30).

87　It is generally accepted that the TatA protein forms the protein-conducting element of the Tat

88　pathway. TatA oligomers assemble at the substrate-bound TatBC complex, dependent on the

89　PMF (27, 31-35). Current models for Tat transport propose that TatA oligomers either provide

90　form-fitting channels of varying diameter that adapt to the size of the folded passenger domain,

91　or that oligomeric assemblies of TatA cause a localized weakening of the membrane and

92　transient bilayer disruption accompanied by substrate transport (reviewed in 10, 11). An

93　implicit prediction of the latter model is that transient membrane rupture would be expected to

94　be accompanied by ion leakage.

95　In this study we have addressed the function of the twin-arginine signal peptide in the Tat

96　transport process by isolating genetic suppressors that either restore transport to signal

97　peptides harbouring transport inactivating twin arginine substitutions, or that restore Tat

98　activity to a TatC variant that has an inactive signal sequence binding site. Our results

99　identified a common set of substitutions, primarily located in the transmembrane helix of TatB

100　that can suppress both types of transport defect. Biochemical analysis of Tat translocases

101　harboring these substitutions indicates that at least one of them, TatB F13Y, promotes signal-

102　peptide independent TatA assembly. Our findings show that, like Sec signal peptides, Tat

103　targeting sequences also play two roles in the transport process.

104

Results.

Single amino-acid substitutions in TatB permit export of twin-arginine substituted signal peptides.

Previous genetic screens using maltose binding protein (21, 22) or GFP (23) fused to the Tat signal sequence of *E. coli* TorA identified mutations that were able to restore some level of Tat transport to fusions with export defective twin arginine substitutions. These substitutions were located towards the N-terminal end of the TatB transmembrane helix or within the cytoplasmic loops of TatC. To shed light on potential functions of signal peptides during Tat transport, we initiated an independent genetic screen using a native *E. coli* Tat substrate, AmiA, as our reporter. AmiA is an *N*-acetylmuramoyl-L-alanine amidase that remodels the cell wall during growth. In the absence of a functional Tat system, the cell envelope is impaired due to the inability to correctly localize AmiA and the related Tat substrate AmiC, rendering *E. coli* sensitive to killing by SDS (36). Utilizing a strain lacking chromosomal *tat* genes and *amiA/amiC* (37), co-production of plasmid-encoded AmiA (from the medium copy construct pSUAmiA; 36) alongside a separate plasmid, pTAT1d (producing TatABC from a compatible medium copy plasmid; 38), permits the strain to grow on LB medium containing 2% SDS (SI Appendix, Fig S1). We then mutated the consecutive arginines in the Tat signal peptide of plasmid-encoded AmiA to each of the amino acid pairs RD, RE, RH, RN, RQ, KH, KQ or HH. As expected, each of these substitutions abolished growth on SDS (SI Appendix, Fig S1).

Next we screened a *tatB* mutant library, generated in pTAT1d by error-prone PCR, for clones that supported growth of strains carrying AmiA with variant signal peptides (38). Each AmiA twin arginine variant was challenged with this library, screening approximately 10,000 clones for each construct. In total across the screening campaign we isolated thirty individual clones. Upon re-screening twenty of these retained the ability to suppress the inactive signal peptide variant of AmiA against which they were originally isolated. These clones are listed in SI Appendix, Table S1. Substitutions appeared to cluster within the transmembrane region of TatB, including L9Q that appeared in seven of the clones and F13Y that occurred in five clones, whilst F6Y, E8K, L9P and L10P were each found once. It should be noted that E8K,

6

133 L9P and L9Q substitutions have previously been identified as suppressors of inactive signal

134 peptides (21, 22). We also found clones, each isolated twice, that contained no substitutions

135 in the transmembrane domain, where the first substitution was in the amphipathic helix of TatB

136 (clones BRQ1, BRQ2, BRQ3 and BRQ5).

137 We introduced the individual amino acid substitutions F6Y, E8K, L9P, L9Q, L10P, F13Y, K30I

138 and I36N into TatB encoded within the *tatABC* operon on the very low copy number plasmid

139 pTAT101 (39) and tested their ability to support export of wild type AmiA and to suppress each

140 of the RD, RE, RH, RN, RQ, KH, KQ or HH AmiA signal sequence variants (Fig 1B, 1C, SI

141 Appendix, Fig S2, Table 1). Each of the substitutions was able to support transport of wild type

142 AmiA, but they varied in their ability to permit export of the AmiA signal sequence variants. For

143 example the F13Y variant of TatB supported growth on SDS for all of the signal peptide

144 variants tested, whereas the L9Q, L10P, K30I and I36N TatB substitutions could suppress

145 some of the signal sequence variants but not KH, KQ and HH substitutions. To determine

146 whether the suppression observed was specific for the AmiA signal peptide, we generated a

147 further construct where the signal peptide of the Tat substrate SufI was fused to the mature

148 portion of AmiA and the same twin arginine substitutions were introduced (SI Appendix, Fig

149 S3, Table 1). The TatB variants generally showed the same pattern of suppression of SufI

150 signal peptide substitutions, although the TatBL10P, K30I and I36N variants could not

151 suppress RD or RE substitutions in the SufI signal peptide.

152

153 **A subset of TatB signal peptide suppressors also suppress TatC signal peptide binding**

154 **site mutants.**

155 In a complementary approach we asked whether it was possible to select suppressors of

156 defects in the signal peptide binding site on TatC. Residue F94 in *E. coli* TatC is highly

157 conserved and lies within the signal peptide binding site (25, 40; Fig 2A), and substitution to

158 other amino acids is poorly tolerated (41). We constructed substitutions of F94 to each of the

159 small neutral amino acids Ala and Gly, helix breaking Pro, polar residues Ser and Gln,

160 positively charged Arg and Lys and negatively charged Asp. These substitutions were

7

161   introduced into TatC encoded on both the medium copy plasmid pTAT1d and the very low

162   copy plasmid pTAT101 that also carry wild type *tatA* and *tatB*. SI Appendix, Fig S4 shows that

163   substitutions to Asp, Gln or Pro resulted in a complete inability of strain DADE (Δ*tatABCD*,

164   Δ*tatE*; 42) to grow in the presence of SDS at both medium and very low copy number. We

165   selected the F94Q substitution of TatC and constructed three mutant libraries in the pTAT1d

166   vector by error prone PCR in an attempt to identify suppressors of this inactivating *tatC*

167   mutation. LibC1 carried mutations in the first 93 codons of *tatC*, LibC2 carried mutations in

168   *tatC* from residue 95 onwards and LibAB contained mutations in the *tatA* and *tatB* genes.

169   A number of suppressors were identified from screening the LibC1 and LibC2 libraries for

170   growth on SDS plates. However sequence analysis indicated that for each of these Tat active

171   mutants there was substitution at the *tatC* F94Q codon to *tatC* F94Y, W or L codons. By

172   contrast, after screening more than 180 000 clones from the LibAB library, eleven mutants

173   were isolated which were able to rescue the growth defect of TatC F94Q on SDS plates, of

174   which five were still able to support growth on SDS following fresh transformation of strain

175   DADE with the isolated plasmid. These clones are listed in SI Appendix, Table S2.

176   Interestingly, each of these suppressors encoded either TatB L10P, F13Y or I36N that had

177   been identified in our prior screen for signal sequence suppressors. Introduction of each of

178   these substitutions, individually, into the very low copy number pTAT101CF94Q plasmid

179   supported growth of strain DADE on SDS-containing media (Fig 2B), indicating that these

180   TatB variants are each able to rescue the Tat-inactivating F94Q TatC substitution.

181   Since these three TatB substitutions that suppress the TatC F94Q defect were previously

182   isolated as suppressors of inactive Tat signal sequences, we asked whether any of the other

183   *tatB* signal sequence suppressors we had found could also rescue the TatC F94Q substitution.

184   Fig 2C shows that in addition to L10P, F13Y and I36N, L9Q could also restore Tat activity to

185   cells producing TatC F94Q. The location of each these residues on a model of TatB is shown

186   in Fig 2D. We next asked the question whether any of these TatC F94Q suppressors could

187   restore Tat activity to other inactivating substitutions in TatC. Fig 2E shows that two different

188 inactivating substitutions of E103 in the signal peptide binding site, either E103A or E103K

189 (25, 40, 41; Fig 2A), could also be complemented by three of the four suppressors of *tatCF94Q*

190 (I36N did not suppress these substitutions), but they could not restore Tat activity caused by

191 inactivating TatC substitutions located outside the signal peptide binding site (SI Appendix,

192 Fig S5). Finally we tested whether F94Q suppressing substitutions were additive, i.e. whether

193 when combined they resulted in a stronger suppressing activity. However, SI Appendix, Fig

194 S6A shows that none of the pairwise combinations we tested gave any suppression of

195 *tatCF94Q* and, with the exception of the F13Y, I36N substitution which showed some

196 suppression of the RN signal peptide variant, the combined suppressors lost suppressive

197 function of RN or KK substitutions of the SufI signal peptide (SI Appendix, Fig S6B). We

198 therefore conclude that the suppressor mutations, when combined, have detrimental rather

199 than additive effects on suppression activity.

200

201 **The TatB F13Y and L9Q substitutions support export of AmiA lacking a signal peptide.**

202 The results above indicate that the TatBF13Y substitution is the strongest suppressor of

203 inactive Tat signal peptides, allowing the *E. coli* Tat system to recognize all eight of the

204 different twin arginine motif substitutions tested as well as suppressing the TatC F94Q

205 mutation. We therefore tested whether more severe signal peptide defects could be

206 suppressed by this TatB variant. Fig 2F shows that, remarkably, even after truncation of the

207 signal peptide by removal of the h-region, or indeed complete removal of the entire signal

208 peptide-coding sequence of AmiA, we could still detect some Tat-dependent translocation of

209 the AmiA passenger domain in the presence of TatBF13Y. The TatB L9Q substitution, which

210 was the strongest suppressor of signal peptide defects after F13Y, also supported some

211 translocation of mature AmiA, however we were not able to detect export of mature AmiC in

212 the presence of either of these two suppressor substitutions (SI Appendix, Fig S6C). We

213 conclude that TatB L9Q and F13Y allow at least one Tat substrate to be transported

214 independent of any signal peptide.

215

Variant TatB proteins support good transport activity of a native Tat substrate but much poorer transport when the signal peptide is altered.

We next addressed whether the TatB substitutions were detrimental to the activity of the Tat system. Using overproduced his-tagged, but otherwise native SufI as a substrate it was seen that mature SufI was clearly detected in the periplasmic fractions of all of the strains tested, although TatBI36N seemed to support only low levels of transport (SI Appendix, Fig S7A). However, we were unable to detect transport of the RD, RN or KQ signal peptide variants of SufI in the presence of TatBF13Y (SI Appendix, Fig S7B) or of a twin-lysine substituted his-tagged CueO in the presence of TatBE8K or F13Y (Fig 3). It therefore appears that there is very low export efficiency of substrates with variant signal peptides in the suppressor mutant strains.

The TatB suppressors do not restore biochemically detectable signal peptide binding to TatBC.

We subsequently sought to understand the biochemical basis for the action of the TatB suppressors. Our initial hypothesis was that they acted to increase the affinity of the TatBC complex for the variant signal peptides, or to restore binding to complexes containing the TatC F94Q or E103A/E103K substitutions. First we produced his-tagged GFP with variants of the SufI signal peptide at its N-terminus and assessed how much TatBC could be co-purified with this from detergent-solubilized membrane fractions. Fig 4A shows that when the wild type signal peptide was fused to GFP, wild type TatBC or variants harbouring the TatB E8K, F13Y or I36N substitutions were co-eluted with his-tagged SufI-GFP. However no TatBC was detected when the RR motif in the signal peptide was mutated to RD, RN, or KK, even in the presence of the TatB suppressor substitutions (despite the fact TatBC and GFP were clearly present in all of the input samples; SI Appendix, Fig S8A and B). Very similar behavior was also seen when his-tagged AmiA variants were used as substrate for co-purification experiments (SI Appendix, Fig S9). Thus TatBC and TatBF13YTatC co-purified with the his-tagged wild type AmiA precursor, but no TatBC was detected when RD, RN, KK or KQ

10

244 substitutions were introduced into the signal peptide, or when the AmiA signal peptide was

245 lacking. We conclude that the TatB suppressors do not detectably restore binding of variant

246 signal peptides to the TatBC complex. Since several TatB variants can transport substrates

247 with defective signal peptides, but not without signal peptides (Fig 3), we infer that the

248 defective signal peptides must still weakly interact with the TatBC complex at a level that is

249 not detected by our co-purification assay.

250 We then tested whether any of the four suppressors, TatB L9Q, L10P, F13Y and I36N, that

251 allow Tat transport in the presence of the TatC F94Q and E103K mutations, acted to restore

252 substrate binding to TatBC complexes containing these signal sequence binding site

253 substitutions. Although high GFP fluorescence and strong TatBC signals were detected in

254 whole cells (SI Appendix, Fig S8C and D), only wild type TatBC was found to co-purify with

255 his-tagged SufIss-GFP (Fig 4B). Thus, as expected, TatC substitutions F94Q or E103K

256 prevented co-purification of TatBC-substrate complex, consistent with loss of signal peptide

257 binding detected for substitutions at these amino acid positions (25, 40). However, detectable

258 signal peptide binding was not restored by introduction of the individual TatB suppressor

259 substitutions. We conclude that the TatB suppressors do not act by rescuing signal peptide

260 binding.

261

262 **The TatBC complexes harboring TatB suppressor substitutions are conformationally**

263 **altered.**

264 We next investigated whether TatBC complexes could still be detected when any of the

265 TatBL9Q, L10P, F13Y or I36N substitutions were present in TatB. Membranes harboring wild

266 type TatA and TatC along with each of these TatB variants were solubilized with digitonin and

267 analysed by blue-native gel electrophoresis (BN-PAGE). As shown in Fig 5A, the wild type

268 TatBC complex solubilized with digitonin migrated close to the 440 kD marker, as reported

269 previously (e.g. 43). The TatB L9Q, F13Y and I36N variants were also associated with a

270 complex of apparently identical size to wild type TatBC, whereas for membranes producing

271 TatBL10P, very little TatBC complex could be detected, even though both proteins were

272 solubilized from the membrane (SI Appendix, Fig S10). Interestingly, the L9Q and F13Y TatB

273 substitutions also resulted in the appearance of a second band of apparently higher mass that

274 was absent from the sample containing wild type TatBC (Fig 5A, B).

275 We wondered whether this additional band might arise due to the presence of excess TatA

276 bound to the variant complexes. However, blotting the BN gels for TatA showed the distinct

277 TatA-laddering pattern reported previously (44, 45) was detectable for all of the samples, but

278 there was no obvious TatA cross-reactive material migrating at the same position as the higher

279 mass TatBC-containing complex (Fig 5B). To examine whether the presence of TatA was

280 required for these higher molecular weight variant TatBC complexes to form, we repeated the

281 BN-PAGE analysis in the absence of TatA (or its paralog TatE; 16). Surprisingly, this resulted

282 in the apparent aggregation of the variant TatBC complexes, yielding a series of bands of

283 apparent masses well above 440kDa that were not seen for the wild type (Fig 5C). We infer

284 from this that there is a conformational alteration in the TatBC complex induced by the

285 presence of the L9Q or F13Y TatB substitutions that in the absence of TatA causes further

286 oligomerisation.

287 Conformational alterations in the TatBC complex have been previously detected by disulfide

288 crosslinking (46, 47). Cléon *et al*. (47) reported that when a Tat substrate was overproduced,

289 a disulfide crosslink between M205C in transmembrane helix 5 of neighboring TatC proteins

290 could be detected *in vivo*, suggesting the formation of a transient TatC dimer in response to

291 substrate binding. Fig 5D confirms that dimerization through TatC M205C is not observed

292 unless cells also harbor an overproduced Tat substrate, in this case CueO. The TatC M205C

293 dimer induced by CueO is almost completely absent when the F94Q substitution is introduced

294 into TatC, again supporting the conclusion that substrate binding promotes TatC dimerization

295 (Fig 5D). Interestingly, however, when either the TatB L10P or F13Y substitutions were

296 present, a TatC M205C crosslink was detected in the absence of overexpressed substrate.

297 We wondered whether these TatB substitutions rendered the TatBC complex more responsive

298 to the presence of endogenous substrates. To test this, we also introduced the signal peptide

299 binding defective F94Q substitution into TatC M205C. However, as Fig 5D shows, the TatC

300   M205C dimer can still be detected in the presence of TatB L10P or F13Y substitutions, even

301   when the F94Q inactivating substitution is present, and is therefore independent of signal

302   peptide binding. We conclude that at least a subset of the TatB suppressors induce

303   conformational changes in the TatBC complex, and that the TatBL10P and F13Y substitutions

304   potentially mimic the substrate-bound form of the complex.

305

306   **The TatBF13Y substitution promotes signal peptide-independent oligomerisation of**

307   **TatA *in vivo*.**

308   Substrate binding to the TatBC complex is a pre-requisite for the assembly of a TatA oligomer.

309   TatA oligomer assembly *in vivo* can be followed by fluorescence microscopy in cells producing

310   a chromosomally-encoded TatA-YFP fusion protein (34). When Tat substrates are present at

311   native level, TatA oligomers are found with low frequency, but this frequency can be

312   significantly increased by overproduction of a Tat substrate protein with a functional signal

313   peptide (34, 35). This finding is confirmed in Fig 6A, where clusters of TatA-YFP can be seen

314   in cells overproducing AmiA from a plasmid. As expected, introduction of the F94Q codon

315   substitution into chromosomally-encoded *tatC* prevented the AmiA-induced clustering of TatA-

316   YFP resulting in a halo of delocalized TatA around the cell periphery (Fig 6A), consistent with

317   the inability of the TatC variant to bind substrates. We next assessed whether any of the TatC

318   F94Q suppressors, TatB L9Q, L10P, F13Y or I36N (introduced into chromosomal *tatB*)

319   affected the oligomerisation of TatA-YFP (Fig 6B, SI Appendix, Fig S11). Remarkably we

320   found that the presence of the TatB F13Y substitution promoted constitutive assembly of TatA-

321   YFP in the absence of overproduced Tat substrates (Fig 6B), and the TatA-YFP assemblies

322   persisted even in the presence of the TatC F94Q substitution for this variant (but not for L9Q,

323   L10P or I36N; SI Appendix, Fig S11). Taken together, these results indicate that the TatB

324   F13Y substitution triggers signal peptide-independent assembly of TatA oligomers.

325

326   **No leak across the cytoplasmic membrane when cells produce the Tat system**

327   **containing TatB F13Y.**

328   One of the current models for Tat transport posits that TatA oligomers facilitate transport of

329   substrates by causing a localized weakening of the bilayer and transient disruption (discussed

330   in 10, 11). Such a mechanism might be expected to be accompanied by increased

331   permeability of small molecules associated with assembled TatA. The availability of a TatB

332   variant (F13Y) that causes TatA to accumulate in the assembled state provides an

333   experimental tool to investigate this issue.

334   First we asked whether overexpression of Tat systems containing the TatB suppressors L9Q,

335   L10P, F13Y or I36N from an arabinose-inducible promoter had any effect on the growth rate

336   of *E. coli*. Fig 7A shows that when production of each of these variant Tat systems was induced

337   by the addition of arabinose, cells grew more slowly than when the wild-type Tat system was

338   overexpressed, with the TatBL9Q substitution having a particularly detrimental effect on

339   growth rate. This indicates that some level of toxicity is associated with overproduction of these

340   variants. We next assessed whether the TatB variants facilitated membrane permeability

341   using an osmotic lysis method previously used to monitor solute movement through the Sec

342   protein transport channel (48). Here spheroplasts containing wild type or variant Tat

343   translocases were diluted into an iso-osmotic solution of the uncharged sugar xylitol and

344   permeation of xylitol into the cells was assessed by monitoring turbidity associated with

345   osmotically-induced spheroplast lysis. Spheroplasts expressing a SecY variant that is known

346   to increase permeability (48, 49) rapidly lysed following dilution into xylitol solution (Fig 7B).

347   However, no lysis was observed for spheroplasts producing any of the variant Tat

348   translocases, even those harboring the TatA-oligomerizing TatB F13Y variant (Fig 7B).

349   Western blotting confirmed that the Tat proteins were present in these membranes (Fig 7C).

350   These results show the TatA assemblies induced by the TatB F13Y substitution do not result

351   in a small molecule leak across the cytoplasmic membrane.

352

353

14

Discussion

In this work a genetic approach has been taken to shed light on functions of twin arginine signal peptides during Tat transport. Two complementary screens, the first to identify substitutions in TatB permitting export of substrates with inactivating substitutions at the signal peptide arginine pair, and the second to identify rescue mutations of the TatC signal peptide binding site converged on a similar group of *tatB* suppressors. Four TatB substitutions were identified – three in the transmembrane domain, L9Q, L10P and F13Y, and one in the amphipathic helix (I36N) that restored Tat transport in the presence of the inactivating TatC F94Q substitution. The same three substitutions in the transmembrane helix could suppress inactivating substitutions at E103, also in the signal peptide binding site. Of these three, the F13Y substitution displayed the strongest suppressing activity, allowing export all of the twin arginine substitutions tested, and even allowing some translocation of AmiA completely devoid of a signal sequence.

A combined bioinformatics and mutagenesis approach has shown the TatB transmembrane helix to bind along transmembrane helix 5 of TatC (50), and this is consistent with *in vitro* disulfide crosslinking studies (25, 39). Signal peptide binding to the TatBC complex is suggested to cause movement of TatB from its resting state binding site on TatC to a site elsewhere on the protein. This is proposed to prime TatA to occupy the same binding site, which in turn triggers assembly of further TatA molecules to form the active translocase (50; Fig 8). After our biochemical experiments revealed that the suppressors did not function by restoring detectable binding of signal peptides to the TatBC complex (Fig 4, S9), we considered whether the TatB substitutions were mimicking the substrate-driven conformational changes which prime the translocase for TatA recruitment, but in the absence of substrate binding. Our analysis using BN-PAGE showed that the TatB L9Q, L10P and F13Y substitutions caused conformational alterations in the resting TatBC complex (Figure 5A,B). The complex containing the L10P substitution appeared more labile as very little full-sized TatBC complex could be detected, whereas the L9Q and F13Y substitutions yielded a subset

381   of TatBC complexes with apparently increased mass which may be indicative of altered

382   subunit composition or significant conformational change. Substrate-induced conformational

383   changes in the wild-type complex can also be monitored by appearance of a crosslink between

384   cysteine residues at position 205 at the periplasmic end of TatC transmembrane helix 5. This

385   residue forms part of the TatB resting-state binding site (50), so is occluded from dimerization

386   with a neighboring TatC molecule in the resting state, but has been shown to dimerize in

387   response to overproduction of a Tat substrate (39). Similarly, assembly of TatA-YFP oligomers

388   (indicating assembled translocation sites) which can be monitored by fluorescence

389   microscopy is only seen for the wild-type translocase upon overproduction of Tat substrates

390   (34). For one of the TatB substitutions, F13Y, both TatC M205C dimerization and TatA-YFP

391   assembly were observed not only in the absence exogenous substrates, but also in the

392   presence of a TatC F94Q substitution which disrupts interaction with signal peptides. We

393   therefore conclude that TatB F13Y has decreased affinity for the 'resting' TatC binding site,

394   and an increased affinity for the 'activated' binding site, such that it is able to trigger recruitment

395   of TatA and transport of precursors in the absence of signal peptide binding. For the other

396   TatB variants we propose that each differs in affinity for the resting and activated binding sites,

397   leading to slight differences in conformation for these translocases, and we assume that in

398   these cases, weak residual binding of a signal peptide lacking its twin arginine motif is

399   sufficient to trigger TatA recruitment, whereas in the wild-type system the higher energy of

400   binding of the twin arginine residues are strictly required for this. Hence the signal peptide

401   plays two distinct roles- in precursor targeting and translocase activation.

402   A favored model for Tat-mediated protein translocation is that protein passage across the

403   membrane is facilitated by bilayer disruption arising from TatA oligomerization. Interestingly,

404   it was noted that there was a reduced growth rate associated with overproduction of Tat

405   systems containing TatB suppressors, including F13Y, suggesting apparent toxicity. However,

406   no leak of the small uncharged sugar, xylitol, could be detected in membranes harboring Tat

407   complexes containing TatB F13Y. This may suggest the presence of a substrate precursor is

16

408   necessary to provide the force required to disrupt the bilayer. Alternatively, it remains possible

409   that the foci of TatA-YFP observed in cells producing TatBF13Y do not correspond to fully

410   assembled translocases and that further recruitment of TatA(-YFP) (for example mediated by

411   the folded mature domain of a Tat substrate (51)) is required. Indeed the export of mature

412   AmiA in the presence of the TatB F13Y substitution might suggest that some Tat substrates

413   have internal targeting information.

414   In summary, our findings support the notion that Tat signal peptides have two distinct roles.

415   They serve to target their passenger domains to the export machinery, but also to trigger

416   assembly of the active translocase. The isolation of substitutions in the Tat machinery that

417   bypass these steps should prove very useful to dissect the mechanism by which folded protein

418   translocation is achieved.

419

Materials and Methods

Strain construction. The *E. coli* strains used in this work are listed in SI Appendix, Table S3.

Strain JM109 was used for regular cloning and transformation of Quickchange products, and

ultracompetent cells of XL10-Gold® (Agilent) were used for construction of the random

mutagenesis libraries.

Strain DADE (as MC4100, Δ*tatABCD*, Δ*tatE* (42)) was used as the background strain for Tat

transport activity tests and production of Tat proteins for membrane protein extraction, *in vivo*

disulfide crosslinking and Blue-Native PAGE, with the exception of Fig 3 where strain MΔBC

(MC4100 Δ*tatBC*; 33) was used. Strain DADE-P (as DADE, *pcnB1 zad*-981*::*Tn*10*d (Kan^r);

(52)) was used to co-produce TatB and TatC along with AmiA for co-purification experiments.

Strain MCDSSAC Δ*tatABC* (37), in which the 2-33 codon of *amiA* and 2-32 codons for *amiC*

are deleted and the *tatABC* operon was replaced with an apramycin resistance cassette, was

used as the background strain for AmiA signal sequence library screening and to analyse

transport of AmiA mediated by AmiA or SufI signal peptide variants.

Transport of AmiA mediated by signal sequence truncations was assessed in strain MC4100

Δ*amiA* Δ*amiC* Δ*tatABC*, which was constructed as follows. The Δ*amiA:*kan^r allele from the

Keio collection (53) was moved into MC4100 by phage P1 transduction, after which the

kanamycin resistance cassette was eliminated according to (54). Subsequently the *amiC*

deletion was introduced and the kanamycin cassette subsequently eliminated using the same

approach. Finally, the Δ*tatABC::*Apra allele was introduced from strain BW25113

Δ*tatABC::*Apra (54) by P1 transduction. Strain BW25113 Δ*glpF* Δ*tatABC* was used in osmotic

lysis experiments and was constructed by P1 transduction of the Δ*glpF:*kan^r allele from the

Keio collection (53), elimination of the kanamycin resistance and P1 transduction of the

Δ*tatABC::*Apra allele as described above.

Strain AyBCE (34), which lacks *tatA* at the native locus and has a *tatA-YFP* fusion integrated

into the chromosomal *att* site, was used in fluorescence imaging. Chromosomal point

447   substitutions in *tatB* and *tatC* were introduced into this strain via plasmid pMAK-AupBC and

448   its variants using the approach of Hamilton *et al.* (55).

449   Strain BL21(DE3) Δ*tatABC* was used to co-produce TatB and TatC along with SufIss-GFPhis

450   for the co-purification experiments. This strain is a derivative of BL21(DE3) where the *tatABC*

451   genes have been replaced with the apramycin resistance cassette, and was constructed by

452   recombination as described previously (56).

453

454   **Plasmid construction**. The plasmids used and constructed in this work are listed in SI

455   Appendix, Table S4. All point mutations in plasmids, as well as insertion of the flag sequence

456   to create p101C*BCflag, were introduced by Quickchange site-directed mutagenesis

457   (Stratagene) using the primers listed in SI Appendix, Table S5.

458   Plasmids pTAT101 (39) and pTAT1d (38) were used to express *tatABC* under the control of

459   the native *tatA* promoter at very low and medium copy number, respectively. pTAT101 cys

460   less (47) was used as the backbone to introduce single cys substitutions for *in vivo* disulfide

461   crosslinking experiments. Plasmid pTATBC1d encodes TatBC and was constructed following

462   amplification of *tatBC* pTAT1d using primers STIPE-ISH and pT7.5R (SI Appendix, Table S5)

463   was digested using *Bam*HI and *Pst*I and cloned into similarly digested pUNIPROM (57).

464   Plasmid pBADTatABChis codes for *tatABC* with a hexahistag coding sequence at the 3' end

465   of *tatC* in pBAD24 (58). It was constructed following amplification of *tatABChis* from pUNITAT2

466   (59), digestion with *Nco*I and *Xba*I and cloning into similarly digested pBAD24.

467   pSUAmiA (36) was used to produce full-length AmiA from a vector specifying chloramphenicol

468   resistance. pSUSufIss-mAmiA was used to produce SufIss-mAmiA and was constructed

469   following separate amplification of DNA encoding the SufI signal sequence including the *sufI*

470   ribosome binding site using primers SufIssFE and SufIssR, and the mature region of AmiA

471   (mAmiA) using primers AmiA-mF and AmiA-mRX from the chromosome, and fusing the two

472   fragments by overlap extension PCR according to reference (60). The resultant DNA fragment

473   was then cloned into the pSU18 vector (61) using *Eco*RI and *Hind*III sites to generate

474   pSUSufIss-mAmiA plasmid. Plasmids pSUSufIssnoH-mAmiA were used for production of

19

475   truncated SufIss-mAmiA lacking the signal peptide h-region. It was constructed by removal of

476   codons 11-21 of the SufI signal sequence via Quickchange using pSUSufIss-mAmiA as

477   template with primers SufI-noHF and SufI-noHR. pSUmAmiA was used to produce signal-less

478   AmiA and was constructed as follows. A DNA fragment containing the ribosome binding site

479   of *amiA* and the coding sequence for mature AmiA was amplified using pSUAmiA as template

480   with primers AmiA-nossFE and AmiA-mRX. The DNA fragment was subsequently cloned into

481   pSU18 using *Eco*RI and *Hind*III sites to generate pSUmAmiA. To express the mature domains

482   of AmiA or AmiC from pQE70, DNA covering these regions were amplified with primer pairs

483   mAmiA-SphI-F/AmiAnostopBamHI-R or mAmiC-SphI-F/AmiCnostopBamHI-R, respectively,

484   using chromosomal DNA as template. The DNA fragments were digested with *Sph*I and

485   *Bam*HI and cloned into *Sph*I/*Bgl*II digested pQE70 vector (Qiagen, Manchester, UK). For

486   fractionation experiments, SufI was produced with a C-terminal histag from pQE80 (Qiagen,

487   Manchester, UK). It was cloned by excision of DNA covering a C-terminally his-tagged SufI

488   from pQE60-SufI (62) as an *Nhe*I-*Xho*I fragment and ligation into similarly digested pQE80.

489   Plasmid pMAK-AupBC was used to introduce the mutations into the AyBCE chromosome and

490   was constructed by amplification of 500 bp of *tatA* upstream DNA from the chromosome of

491   strain AyBCE using primers TatAup1-*Xba*I and TatAup2-*Cla*I, which was cloned into

492   pBluescript KS(+) using *Xba*I and *Cla*I sites to give pKS-Aup. Next a DNA fragment covering

493   the whole of *tatBC* was amplified from the chromosome of AyBCE strain with primers

494   TatA6B7-*Cla*I and TatCrev-*Kpn*I, and cloned into pKS-Aup using *Cla*I and *Kpn*I sites to

495   generate pKS-AupBC. Subsequently the DNA covering the *tatA* upstream sequence along

496   with *tatBC* was excised using *Xba*I and *Kpn*I and cloned into similarly digested pMAK705 (55)

497   to give pMAK-AupBC.

498   Plasmid pFAT75ΔA-BC, which is a pQE-based plasmid expressing TatB and TatC without a

499   histag (19), and pSufIss-GFPhis, which is a pCFDuet-based plasmid expressing synthetic SufI

500   signal sequence-fused GFP with a histag at its C-terminus under the control of T7 promoter,

501   were used in co-purification experiments. Plasmid pFAT75ΔA-BC-AmiAhis coproduces

502   untagged TatBC along with his-tagged AmiA and was constructed as follows. A DNA fragment

503 covering *amiA* was amplified from MC4100 genomic DNA using primers AmiAFATApaI-F and

504 AmiAnostopBamHI-R, digested with *Apa*I and *Bam*HI and cloned into *Apa*I - *Bgl*II digested

505 pFAT75-SufIhis (47). pFAT75ΔA-BC-mAmiAhis was constructed similarly, using primer

506 mAmiAFATApaI-F and AmiAnostopBamHI-R to amply DNA covering the mature region of

507 AmiA.

508

509 **Mutant library construction and screening**.

510 To screen for Tat signal sequence suppressors, substitutions of the twin arginine sequence of

511 the AmiA signal peptide (to -RD, -RE, -RN, -RQ, -RH, -HH, -KH and –KQ) were constructed

512 in the pSUAmiA plasmid. The resulting plasmids were individually introduced into strain

513 MCDSSAC Δ*tatABC* and each resulting strain served as host to screen an existing *tatB* mutant

514 library (in plasmid pTAT1d, 600,000 individual clones, 0.25% error rate, (38)).

515 To screen for suppressors of the TatC F94Q substitution, three separate mutagenesis

516 libraries, each of which carried the *tatC* F94Q codon substitution, were constructed that

517 contained random mutations in either *tatAB*, codons 1-93 of *tatC* (*tatC*1) or codons 95-258 of

518 *tatC* (*tatC*2), respectively using a modified MEGAWHOP (megaprimer PCR of whole plasmid)

519 method as described (63). DNA fragments of *tatAB*, *tatC1* and *tatC2* containing random

520 mutations were generated using error-prone PCR. Error-prone PCR was carried out in 1x

521 GoTaq buffer, 7mM MgCl$_2$, 0.2 mM dATP, 0.2mM dGTP, 1mM dCTP, 1mM dTTP, 0.4 µM

522 each primer, 0 to 0.1 mM MnCl$_2$, 50 ng pTAT1dCF94Q plasmid as template and 5 U GoTaq®

523 DNA Polymerase (Promega) in a total volume of 50 µl using a PCR program: 94 °C for 2min

524 followed by 20 cycles of incubation at 94 °C for 30s, 50 °C for 30s, and 72 °C for 3min, and a

525 final incubation at 72 °C for 5min. Primer pairs TatA-FB and TatB-RS were used to amplify

526 *tatAB*, TatCm6 and TatC93R to amplify *tatC*1 and TatC95F and TatCR1d were used to amplify

527 *tatC*2. The DNA fragments were then used as megaprimers to amplify the whole plasmid,

528 which was carried out in a 50 µl mixture containing 1x Herculase II reaction buffer, 0.5 mM

529 each dNTP, 100ng pTAT1dCF94Q plasmid as template, 500 ng DNA fragment obtained

530 above as megaprimers and 5 U Herculase II Fusion DNA Polymerase (Agilent) using the PCR

21

531 program : incubation at 68 °C for 5min, 95 °C for 2min, followed by 20 cycles of incubation at

532 95 °C for 30s, 55 °C for 30s, and 68 °C for 6min. The resultant whole plasmid PCR products

533 were digested with *Dpn* I to remove the template DNA and incubated with T4 polynucleotide

534 kinase and T4 DNA ligase to repair the nicks. Finally, the whole plasmids were separately

535 transformed into XL10-Gold® Ultracompetent Cells (Agilent) resulting in three mutagenesis

536 libraries. Subsequent sequencing of 10 randomly selected colonies from each library revealed

537 an average error rate of approximately 2 nucleotides per 1000 base pair. Screening of these

538 three libraries was carried out in strain DADE ($\Delta tatABCD$, $\Delta tatE$).

539 For screening experiments the libraries were transformed into the respective host strains and

540 subsequently plated onto solid LB medium containing appropriate antibiotics and 2% SDS for

541 selection. Colonies able to grow under these conditions were isolated, the mutations in the *tat*

542 gene(s) identified by sequencing and retested for growth in the presence and absence of 2%

543 SDS. To further verify isolated candidates, individual mutations were introduced into low copy

544 number plasmid pTAT101 by site-directed mutagenesis and the activity was again assessed

545 on 2% SDS-containing plates.

546

547 **Protein methods.** For co-purification of TatBC-substrate complexes, cultures of strain

548 BL21(DE3) $\Delta tatABC$ harboring pFAT75ΔA (or a point-substituted variant) and pSufI-GFPhis

549 were incubated at 37°C for 7 hours with shaking, after which they were supplemented with

550 0.2mM isopropyl β-D-1-thiogalactopyranoside (IPTG) and incubated for a further 17 hours at

551 37°C. The cells were subsequently harvested and resuspended in 1 x PBS, the fluorescence

552 intensity of the suspension was recorded, after which the cells were re-pelleted, resuspended

553 in 2 x lysis buffer (100 mM NaH_2PO_4 pH 8.0, 600 mM NaCl, 40 mM imidazole, 50 mg lysozyme,

554 80 U benzonase, and protease inhibitor) and mixed gently at room temperature for one hour.

555 Cells were snap frozen at -80°C, thawed at room temperature and an equivalent amount of

556 2.5% digitonin was added and the sample gently mixed at room temperature for one hour. Cell

557 debris was pelleted by centrifugation and the supernatant was transferred to a 96-well plate

558 and mixed with 20 μl Ni-NTA Magnetic Agarose Beads (Qiagen) for one hour. After the beads

22

559   were washed three times with wash buffer (50 mM NaH_2PO_4, pH 8.0, 300 mM NaCl , 40 mM

560   imidazole, 0.03 % digitonin), bound proteins were eluted with 50 µl elution buffer (50 mM

561   NaH_2PO_4, pH 8.0, 300 mM NaCl , 250 mM imidazole, 0.03 % digitonin).

562   *In vivo* disulfide crosslinking was carried out in strain DADE harboring pTAT101 cys less

563   CM205C, as described in (47). Blue-Native PAGE was undertaken according to (47, 64).

564   Subcellular fractionation was according to (65). Preparation of membrane fractions was as

565   described previously (39). For analysis of SufI export, *E. coli* strain DADE harboring wild-type

566   or signal peptide variants of pQE80SufIhis alone or alongside wild-type or TatB variants of

567   pTAT101 was cultured in the presence of 1mM IPTG until OD_{600} of 1 was reached. Samples

568   (equivalent to 150 µl of whole cells from an OD_{600} = 1, or periplasm fractions from the

569   equivalent of 300 µl of cells from an OD_{600} = 1) were separated by SDS PAGE and analysed

570   by Western blot with anti-6X His tag® or anti-RNA polymerase β subunit antibodies

571   (cytoplasmic control protein). For analysis of CueO export, strain MΔBC harbouring wild-type

572   and KK variants of pQE80-CueO alongside wild-type, *tatB*^{E8K} or *tatB*^{F13Y} variants of

573   p101C*BCflag were cultured and fractionated as previously described (34). Immunoblotting

574   was according to the methodology of (66), and antibodies to TatA, TatB and TatC have been

575   described previously (47, 67). Anti-6X His tag® antibody (HRP-conjugated) was purchased

576   from Abcam (Cambridge, UK, catalog number ab184607), anti-DnaK mouse monoclonal

577   8E2/2 antibody was also from Abcam (catalog number ab69617) and a mouse monoclonal

578   anti RNA polymerase β-subunit antibody was purchased from NeoClone Biotechnology

579   (Madison, USA; catalog number W0023). Secondary antibodies were goat anti-Rabbit IgG

580   (HRP Conjugate, catalog number 170-6515) or Goat Anti-Mouse IgG (HRP conjugate, catalog

581   number170-6516), both from Biorad (Hemel Hempstead, UK). Immunoreactive bands were

582   visualized with the Clarity Western ECL Substrate Kit (BioRad) and captured either on light-

583   sensitive film or using the GeneGNOME camera (Syngene).

584

585 **Cell permeability experiments**. Cell permeability experiments were performed according to

586 (48). Briefly, cells were grown aerobically at 37°C with 1:100 inoculation of an overnight culture

587 for 2 hours. Production of TatABC harboring TatB variants and of SecY(Δplug)EG was induced

588 by addition of 0.2% arabinose at 37°C for 3 hours. Subsequently, the OD_{600nm} of each sample

589 was normalized using LB, a small volume was withdrawn for Western blotting and equal

590 volumes of each culture were then harvested and resuspended in fractionation buffer (50 mM

591 Tris-HCl buffer, 20% sucrose, pH 7.5, 5 mM EDTA, 0.6 mg/ml lysozyme) and incubated at

592 room temperature for 20min to obtain spheroplasts. The spheroplast samples were then

593 adjusted to the same OD_{600} and a 19-fold excess of 0.616 M xylitol solution was added. The

594 samples were rapidly transferred to a 96-well plate and OD_{600} was measured every 30

595 seconds for 300 seconds.

596

597 **Fluorescence microscopy.** Cells were prepared for fluorescence microscopy and imaged as

598 previously described (47), with the exception that a 550nm LP emission filter was used.

599
600

601

612

Figure Legends

615 **Figure 1. Isolation of signal sequence suppressors in *tatB*.** A. Schematic representation

616 of a twin arginine signal peptide. The signal peptide sequences of *E. coli* Tat substrates SufI

617 and AmiA are given underneath, with residues matching the Tat consensus motif in red, the

618 consecutive arginines in red underline and the signal peptidase cleavage site in black

619 underline. B and C. An example of screening results. Growth of MCDSSAC Δ*tatABC*

620 coproducing the indicated TatB variants (with wild type *tatA* and *tatC*) from pTAT101,

621 alongside B. the HH or C. RE-substituted signal peptide variants of AmiA, on LB agar

622 supplemented with chloramphenicol and kanamycin, with or without the addition of 2% SDS

623 as indicated. An 8μl aliquot of each strain/plasmid combination following aerobic growth to an

624 OD$_{600}$ of 1.0 was spotted and incubated for 16 hr at 37°C.

625 **Figure 2. Isolation of suppressors of the TatC F94Q inactivating substitution.** A. Model

626 of *E. coli* TatC (from (47)) showing the location of the F94 and E103 residues (in red) that form

627 part of the signal peptide binding. B., C. and E. Growth of DADE (Δ*tatABCD*, Δ*tatE*)

628 coproducing wild type TatA alongside; B. and C. F94Q-substituted TatC or E. E103A-

629 substituted TatC, and the indicated substitution in TatB from plasmid pTAT101 on LB agar or

630 LB agar containing 2% SDS. D. Structure of *E. coli* TatB (from (68)) with the locations of the

631 TatCF94Q suppressor substitutions shown in red. F. Strain MC4100 Δ*ami*A Δ*amiC* Δ*tatABC*

632 coproducing the indicated TatB variants (with wild type *tatA* and *tatC*) from pTAT101,

633 alongside either a signal peptide variant of SufI lacking the h-region fused to the AmiA mature

634 domain, or the mature AmiA domain alone on LB agar or LB agar containing 1% SDS. For all

635 growth tests, a single colony of each strain/plasmid combination was resuspended in 30μl of

636 PBS and an 8μl aliquot was spotted onto LB agar supplemented with appropriate antibiotics,

637 along with SDS as indicated and incubated for 16 hr at 37°C.

639 **Figure 3. TatB suppressors support export of a Tat substrate with its native signal**

640 **peptide.** *E. coli* strains producing native levels of TatA, TatC and the indicated TatB variants,

641 and overproducing his-tagged CueO with a wild-type (top panel) or KK-substituted (bottom

642 panel) signal peptide were fractionated into whole cell (W), spheroplast (S) and periplasm (P)

643 fractions. Equivalent amounts of each fraction were separated by SDS PAGE and analysed

644 by Western blot with antibodies against CueO and the cytosolic marker DnaK.

645 **Figure 4. The TatB suppressors do not restore signal peptide binding to the TatBC**

646 **complex.** A. C-terminally his-tagged GFP with the wild type (RR) or twin-arginine substituted

647 SufI signal peptide at its N-terminus, as indicated, was purified by magnetic nickel beads from

648 digitonin–treated cell extracts co-expressing TatC along with either wild type TatB or the E8K,

649 F13Y or I36N substituted variants. B. C-terminally his-tagged GFP with the wild type SufI

650 signal peptide at its N-terminus was purified by magnetic nickel beads from digitonin–treated

651 cell extracts co-expressing TatB and TatC with the indicated amino acid substitutions. For A.

652 and B. the elution fractions from each sample were normalized for GFP fluorescence and an

653 equivalent amount of purified SufIss-GFPhis was loaded onto SDS-PAGE (4-15% Mini-

654 PROTEAN® TGX™ precast gradient gel) followed by western blot using TatB and TatC mixed

655 antibodies.

656 **Figure 5. A constitutive disulfide crosslink and aberrant Blue-native PAGE migration**

657 **induced by a subset of TatB suppressors.** A-B. Membranes from *E. coli* strain DADE

658 (Δ*tatABCD* Δ*tatE*) producing the indicated TatB variants alongside wild-type TatA and TatC

659 from plasmid pTAT1d were solubilized by addition of 2% digitonin and analysed by BN-PAGE

660 (4-16% Bis-Tris NativePAGE gels) followed by Western blot with anti-TatA, anti-tatB or anti-

661 TatC antibodies as indicated. 20 μg solubilized membrane was loaded in each lane. C.

662 Membranes from strain DADE producing the indicated TatB variant alongside wild-type TatC

663 from plasmid pTATBC1d were solubilized and analysed as in A-B. D. Whole cells of DADE

664 harboring pTAT101 co-producing either wild type TatA, the M205C single cysteine variant of

665 TatC and the indicated substitution in TatB, or wild type TatA, the TatC F94Q M205C variant

27

666   of TatC and the indicated substitution in TatB were subjected to oxidizing (O) or reducing (R)

667   conditions. Where indicated the additional plasmid pQE80-CueO, (producing His-tagged

668   CueO) was also present. Membranes were prepared from equal quantities of cells following

669   treatment and equivalent amounts of material from each sample were resolved by non-

670   reducing SDS-PAGE (12% acrylamide). TatC was visualized by western blotting using an anti-

671   TatC antibody and CueO-His with an anti-His antibody.

672   **Figure 6. The TatB F13Y substitution promotes constitutive oligomerisation of TatA *in***

673   ***vivo.*** Fluorescence images of TatA-YFP in representative cells of A. strains AyBCE or

674   AyBC$_{F94Q}$E (encoding the TatC F94Q substitution in chromosomal *tatC*) in the presence

675   (pAmiA) or absence of plasmid-encoded wild type AmiA, as indicated. Production of AmiA was

676   induced by addition of 1mM IPTG to mid-log cell cultures 45 min before harvesting. B. strains

677   AyB$_{F13Y}$CE (encoding the TatB F13Y substitution in chromosomal *tatB*) and AyBC$_{F94Q}$E

678   (encoding the TatB F13Y substitution in chromosomal *tatB*, alongside the TatC F94Q

679   substitution in chromosomal *tatC*). Representative micrographs are shown for each sample;

680   Scale bar: 1 μm.

681   **Figure 7. No detectable leak of xylitol across the cytoplasmic membrane when cells**

682   **produce Tat translocases harboring TatB suppressors.** A. Overnight cultures of *E. coli*

683   strain BW25113 Δ*glpF* Δ*tatABC* harboring pBAD24 encoding TatA and TatC-his along with

684   wild type TatB or each of the L9Q, L10P, F13Y or I36N point substitutions, were subcultured

685   at 1:100 dilution into fresh LB medium containing ampicillin, which was supplemented after

686   120 min with 0.2% of glucose or arabinose, as indicated. Growth of the strains was followed

687   for a further 6.5h. Error bars represent standard deviation, *n* = 3 (biological replicates). B. The

688   same strain and plasmid combinations as in part A, alongside BW25113 Δ*glpF* harboring

689   pBAD22SecY(Δplug)EG. were subcultured and supplemented with 0.2% of arabinose as

690   described in part A and grown for a further 3 h after which spheroplasts were prepared and

691   incubated in the presence of xylitol. C. An aliquot of each sample producing plasmid-encoded

692   Tat proteins was analysed by SDS PAGE and western blotting to confirm expression of TatA,

693   TatB and TatC-his.

694   **Figure 8. Tat translocases containing TatB suppressor variants may more readily**

695   **transition to the signal peptide-activated state.** Top panel: Model for Tat transport. A signal

696   peptide bound through its n-region to the cytoplasmic surface of TatC (step 1) transitions to a

697   deep binding mode (step 2). The deep insertion of the signal peptide displaces TatB from its

698   resting state binding site on TatC (grey arrow). TatB movement allows polymerisation of TatA

699   to be nucleated (step 3). The substrate passes across the membrane facilitated by the TatA

700   oligomer (step 4). Bottom panel: TatB variants that suppress signal sequence defects

701   (represented as *B**) may be more easily displaced from the resting state binding site. The TatB

702   variants appear to be on a continuum with TatB F13Y pushing the Tat system into an

703   assembled state (step 4), whereas Tat systems harboring the weaker suppressing variants

704   are more likely to correspond to step 3.

705

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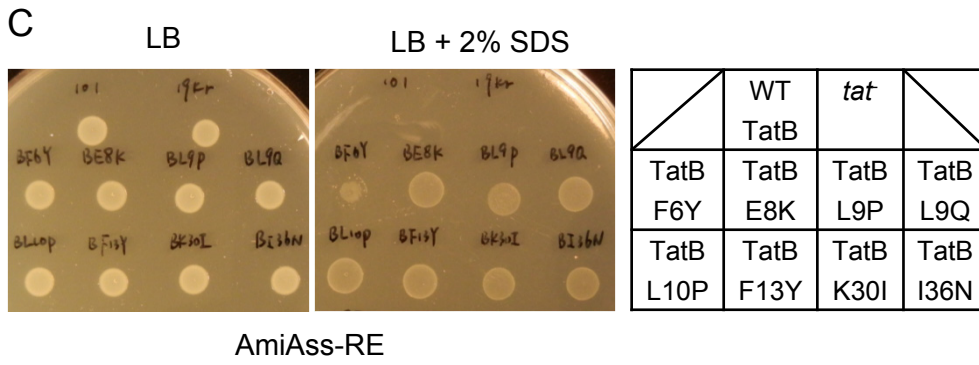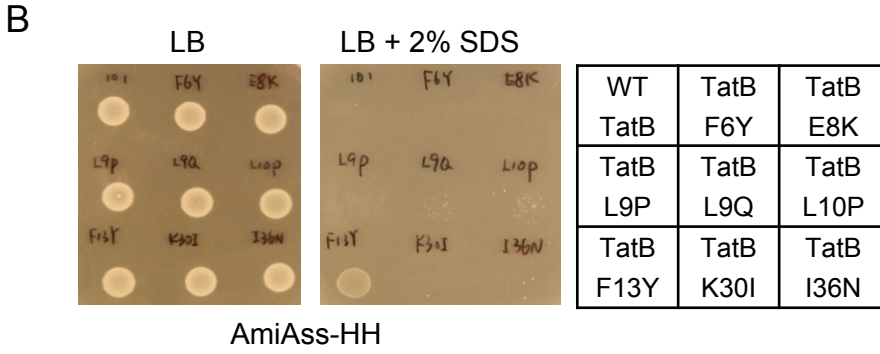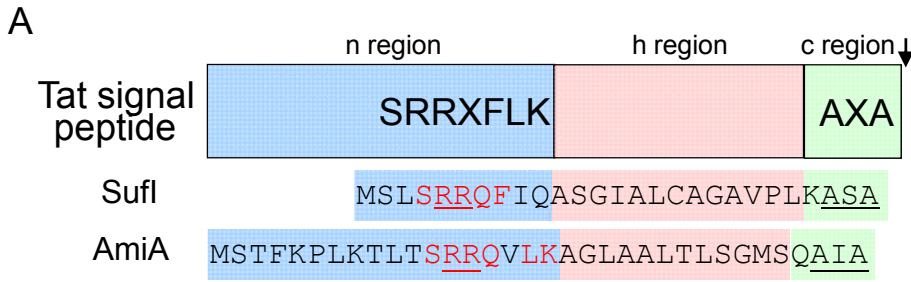
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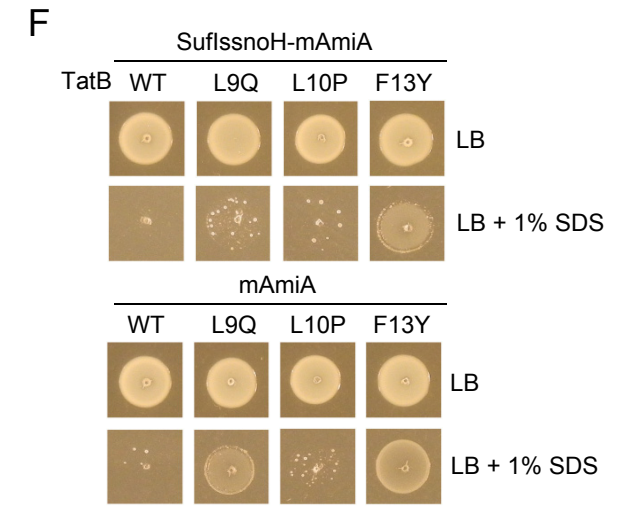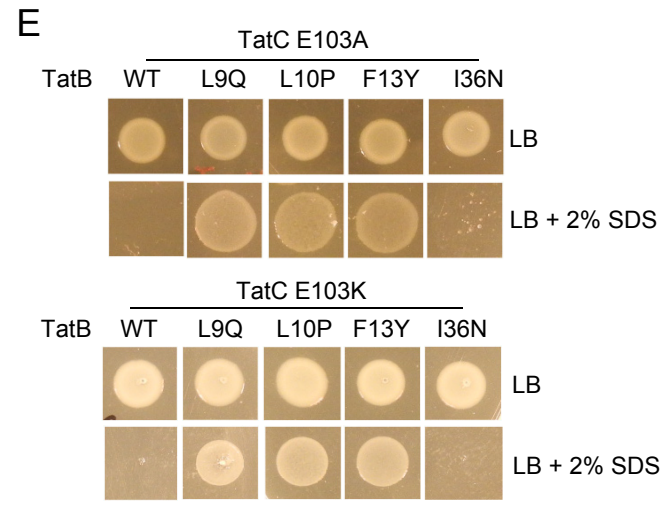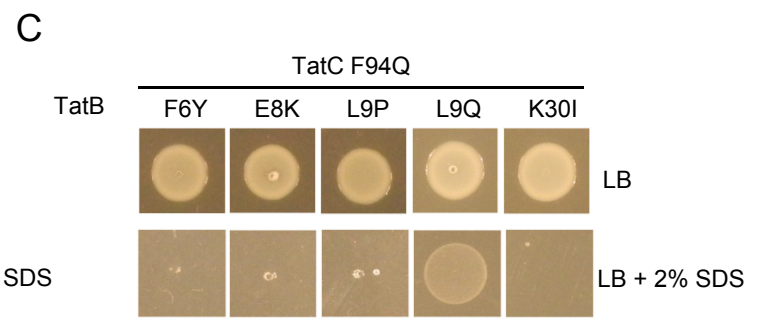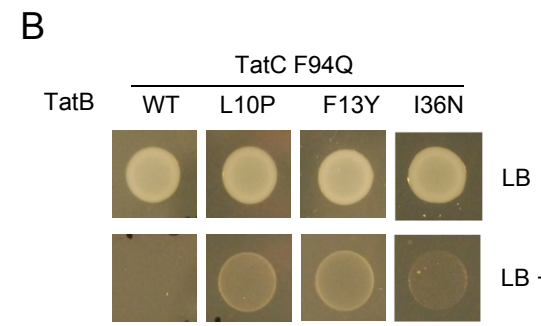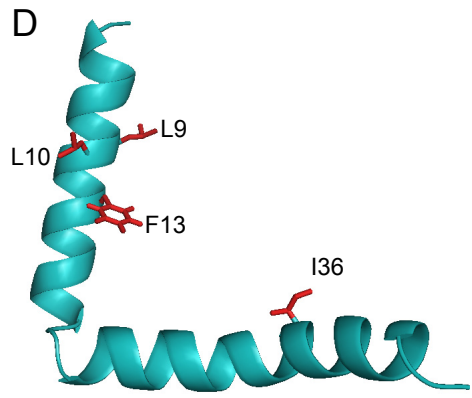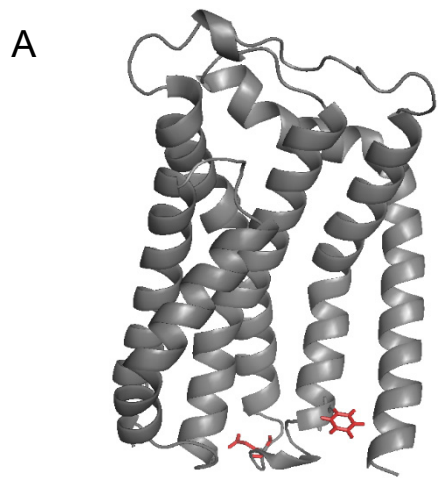
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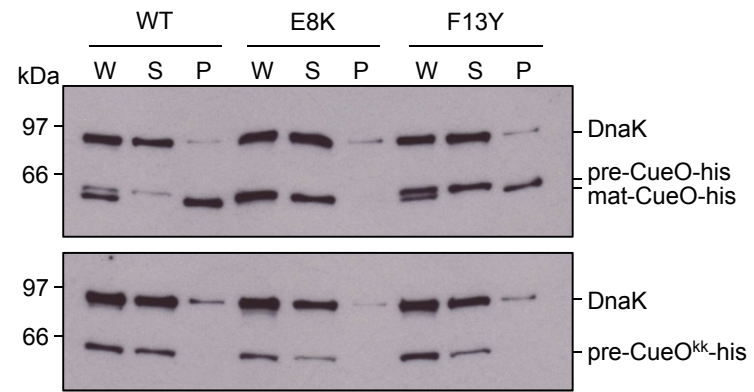812   210.

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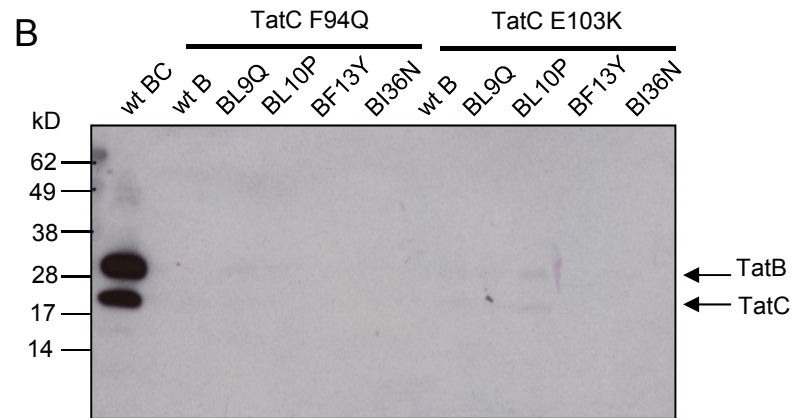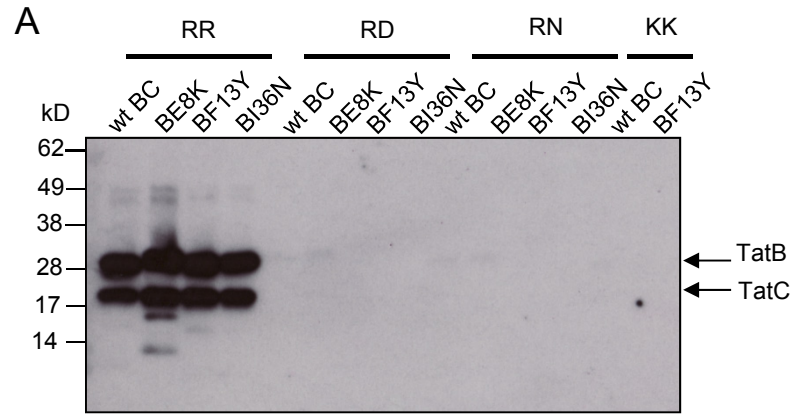866   functions as an obligate oligomer. *Mol Microbiol* 98(1):111-129.
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A

Tat signal peptide

| | n region | h region | c region ↓ |

SufI   MSL**SRRQ**FIQASGIALCAGAVPL**KASA**

AmiA   MSTFKPLKTLT**SRRQVLK**AGLAALTLSGMS**QAIA**

B

LB   LB + 2% SDS

WT TatB	TatB F6Y	TatB E8K
TatB L9P	TatB L9Q	TatB L10P
TatB F13Y	TatB K30I	TatB I36N

AmiAss-HH

C

LB   LB + 2% SDS

	WT TatB	*tat*	
TatB F6Y	TatB E8K	TatB L9P	TatB L9Q
TatB L10P	TatB F13Y	TatB K30I	TatB I36N

AmiAss-RE

A

B

TatC F94Q

TatB	WT	L10P	F13Y	I36N	
	○	○	○	○	LB
		○	○	○	LB + 2% SDS

C

TatC F94Q

TatB	F6Y	E8K	L9P	L9Q	K30I	
	○	○	○	○	○	LB
				○		LB + 2% SDS

D

L9
L10
F13
I36

E

TatC E103A

TatB	WT	L9Q	L10P	F13Y	I36N	
	○	○	○	○	○	LB
		○	○	○		LB + 2% SDS

TatC E103K

TatB	WT	L9Q	L10P	F13Y	I36N	
	○	○	○	○	○	LB
		○	○	○		LB + 2% SDS

F

SufIssnoH-mAmiA

TatB	WT	L9Q	L10P	F13Y	
	○	○	○	○	LB
		○	○	○	LB + 1% SDS

mAmiA

TatB	WT	L9Q	L10P	F13Y	
	○	○	○	○	LB
		○	○	○	LB + 1% SDS

A

	RR				RD				RN				KK	
kD	wt BC	BE8K	BF13Y	BI36N	wt BC	BE8K	BF13Y	BI36N	wt BC	BE8K	BF13Y	BI36N	wt BC	BF13Y

62 —
49 —
38 —
28 —   ← TatB
17 —   ← TatC
14 —

B

		TatC F94Q				TatC E103K				
kD	wt BC	wt B	BL9Q	BL10P	BF13Y	BI36N	wt B	BL9Q	BL10P	BF13Y	BI36N

62 —
49 —
38 —
28 —   ← TatB
17 —   ← TatC
14 —

A AyBCE

AyBCE pSUAmiA

AyBC$_{F94Q}$E

AyBC$_{F94Q}$E pSUAmiA

B AyB$_{F13Y}$CE

AyB$_{F13Y}$C$_{F94Q}$E

A

WT Tat — TatB-L9Q — TatB-L10P — TatB-F13Y — TatB-I36N

0.2% glucose

0.2% arabinose

B

WT Tat — TatB-L9Q — TatB-L10P — TatB-F13Y — TatB-I36N — SecYΔplug

C

kD

15 — TatA

25 — TatB

25 — TatC-his

WT L9Q L10P F13Y I36N

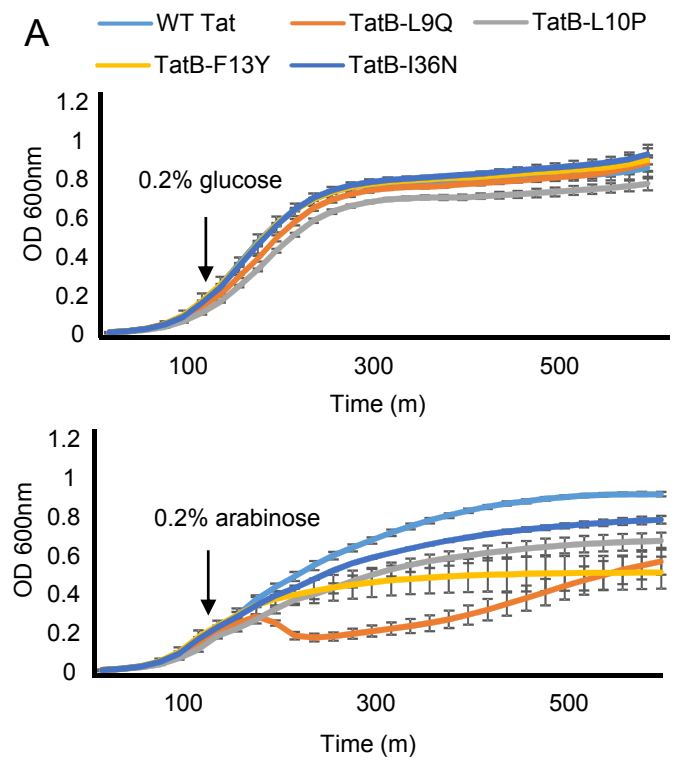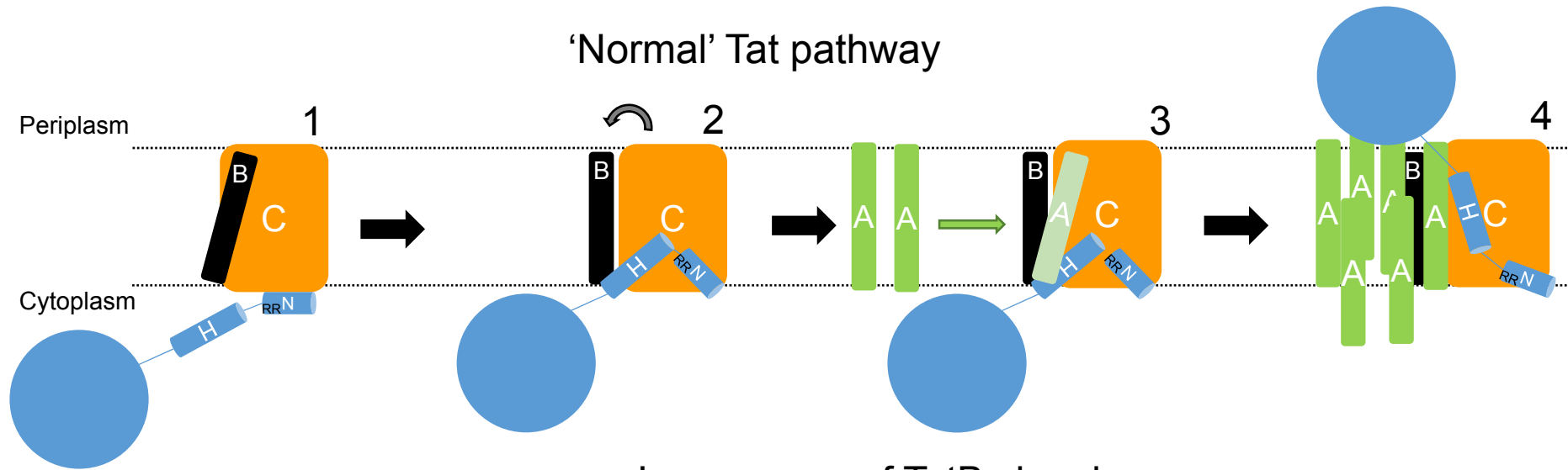'Normal' Tat pathway
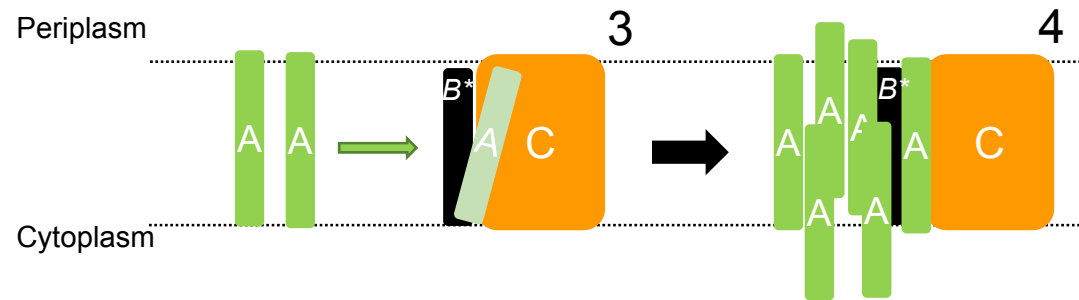
Periplasm

Cytoplasm

In presence of TatB signal sequence suppressors

Periplasm

Cytoplasm

A signal sequence suppressor mutant that stabilizes an assembled state of the twin arginine translocase

Qi Huang, Felicity Alcock, Holger Kneuper, Justin C. Deme[3], Sarah Rollauer, Susan M. Lea,

Ben C. Berks and Tracy Palmer

SUPPORTING INFORMATION

SI APPENDIX

Clone	TatB substitution/s
BRE1	L9Q K103R
BRE2*	L9Q
BRN2	L9Q Q134V
BRN3*	L9Q
BRN4	L10P V12M
BRN5	L9P
BRQ1*	K30I K65R N99D
BRQ2*	K30I K65R N99D
BRQ3*	I36N S41T
BRQ4	L9Q T72A
BRQ5*	I36N S41T
BRH1	L9Q N73K
BRH2	E8K L71H
BRH4	F13Y P138L K159R
BRH5*	F13Y
BRH6*	F13Y
BRH7*	L9Q
BRH8	F6Y V32A A69V
BRH9*	F13Y
BKQ1*	F13Y

Table S1. Clones isolated from a *tatB* mutant library following screening for suppression of transport defects of inactive signal peptides. The BRE, BRN, BRQ, BRH, BHH, BKH or BKQ clone nomenclature signify substitutions isolated following screening against RE, RN or KQ variants of the AmiA signal peptide RR motif, respectively.

*identical clones

Clone	TatAB substitution/s
AB-1	TatA K23N,TatB I36N, TatB N119I, TatB S164C
AB-5	TatB L10P
AB-16	TatB L10P, TatB N73Y
AB-157	TatB F13Y
AB-172	TatA A60E, TatA A76R, TatB F13Y

Table S2. Clones isolated from a *tatAB* mutant library following screening for suppression of the transport defect arising from the TatC F94Q substitution.

Strain	Relevant genotype	Source
JM109	$endA1$, $recA1$, $gyrA96$, thi, $hsdR17$ (r_k^-, m_k^+), $relA1$, $supE44$, $\Delta(lac\text{-}proAB)$, [F′ $traD36$, $proAB$, $laqI^qZ\Delta M15$]	Promega
XL10-gold	$Tet^r\Delta(mcrA)183$ $\Delta(mcrCB\text{-}hsdSMR\text{-}mrr)173$ $endA1$ $supE44$ $thi\text{-}1$ $recA1$ $gyrA96$ $relA1$ lac Hte [F′ $proAB$ $lacI^qZDM15$ Tn10 (Tet^r) Amy Cam^r]	Agilent
MC4100	F-$\Delta lacU169$ $araD139$ $rpsL150$ $relA1$ $ptsF$ rbs $flbB5301$	(1)
DADE	As MC4100, $\Delta tatABC$, $\Delta tatE$	(2)
DADE-P	as DADE, $pcnB1$ zad-$981::$Tn10d (Kan^r)	(3)
MΔBC	MC4100 $\Delta tatBC$	(4)
MCDSSAC $\Delta tatABC$	MC4100, $amiA\Delta 2$-33 $amiC\Delta 2$-32, $\Delta tatABC::$Apra	(5)
MC4100 $\Delta amiA$ $\Delta amiC$ $\Delta tatABC$	MC4100, $\Delta amiA$, $\Delta amiC$, $\Delta tatABC::$Apra	This work
AyBCE	MC4100 $\Delta tatA$, $attB::P_{tatA}tatA\text{-}EAK\text{-}eyfp_{A206K}$	(4)
AyBCE ($tatB_{L9Q}$)	As AyBCE, $tatB_{L9Q}$	This work
AyBCE ($tatB_{L10P}$)	As AyBCE, $tatB_{L10P}$	This work
AyBCE ($tatB_{F13Y}$)	As AyBCE, $tatB_{F13Y}$	This work
AyBCE ($tatB_{I36N}$)	As AyBCE, $tatB_{I36N}$	This work
AyBCE ($tatC_{F94Q}$)	As AyBCE, $tatC_{F94Q}$	This work
AyBCE ($tatB_{L9Q}$ $tatC_{F94Q}$)	As AyBCE ($tatB_{L9Q}$, $tatC_{F94Q}$)	This work
AyBCE ($tatB_{L10P}$ $tatC_{F94Q}$)	As AyBCE ($tatB_{L10P}$, $tatC_{F94Q}$)	This work
AyBCE ($tatB_{F13Y}$ $tatC_{F94Q}$)	As AyBCE ($tatB_{F13Y}$, $tatC_{F94Q}$)	This work
AyBCE ($tatB_{I36N}$ $tatC_{F94Q}$)	As AyBCE ($tatB_{I36N}$, $tatC_{F94Q}$)	This work
BL21(DE3) $\Delta tatABC$	BL21(DE3), $\Delta tatABC::$Apra	This work
BW25113	$lacI^q$ $rrnB_{T14}$ $\Delta lacZ_{WJ16}$ $hsdR514$ $\Delta araBAD_{AH33}$ $\Delta rhaBAD_{LD78}$	(6)
BW25113 $\Delta glpF$ $\Delta tatABC$	BW25113, $\Delta glpF$, $\Delta tatABC::$Apra	This work

Table S3. Strains used and constructed in this study.

Plasmid	Relevant genotype	Source
pTAT101	Low copy number vector expressing TatABC under the control of *tat* promoter. Kan^r.	(7)
pTH19kr	Low copy-number cloning vector. Backbone of pTAT101.	(8)
pTAT101-BF6Y	As pTAT101, TatB F6Y exchange	This work
pTAT101-BE8K	As pTAT101, TatB E8K exchange	This work
pTAT101-BL9P	As pTAT101, TatB L9P exchange	This work
pTAT101-BL9Q	As pTAT101, TatB L9Q exchange	This work
pTAT101-BL10P	As pTAT101, TatB L10P exchange	This work
pTAT101-BF13Y	As pTAT101, TatB F13Y exchange	This work
pTAT101-BK30I	As pTAT101, TatB K30I exchange	This work
pTAT101-BI36N	As pTAT101, TatB I36N exchange	This work
pTAT101-CF94Q	As pTAT101, TatC F94Q exchange	This work
pTAT101-CF94A	As pTAT101, TatC F94A exchange	This work
pTAT101-CF94D	As pTAT101, TatC F94D exchange	This work
pTAT101-CF94G	As pTAT101, TatC F94G exchange	This work
pTAT101-CF94K	As pTAT101, TatC F94K exchange	This work
pTAT101-CF94P	As pTAT101, TatC F94P exchange	This work
pTAT101-CF94R	As pTAT101, TatC F94R exchange	This work
pTAT101-CF94S	As pTAT101, TatC F94S exchange	This work
pTAT101-CF94Q-BF6Y	As pTAT101-CF94Q, TatB F6Y exchange	This work
pTAT101-CF94Q-BE8K	As pTAT101-CF94Q, TatB E8K exchange	This work
pTAT101-CF94Q-BL9P	As pTAT101-CF94Q, TatB L9P exchange	This work
pTAT101-CF94Q-BL9Q	As pTAT101-CF94Q, TatB L9Q exchange	This work
pTAT101-CF94Q-BL10Q	As pTAT101-CF94Q, TatB L10P exchange	This work
pTAT101-CF94Q-BF13Y	As pTAT101-CF94Q, TatB F13Y exchange	This work
pTAT101-CF94Q-BK30I	As pTAT101-CF94Q, TatB K30I exchange	This work
pTAT101-CF94Q-BI36N	As pTAT101-CF94Q, TatB I36N exchange	This work
pTAT101-BL9Q F13Y	As pTAT101, TatB L9Q, F13Y exchange	This work
pTAT101-BL10P F13Y	As pTAT101, TatB L10P, F13Y exchange	This work
pTAT101-BL9Q I36N	As pTAT101, TatB L9Q, I36N exchange	This work
pTAT101-BL10P I36N	As pTAT101, TatB L10P, I36N exchange	This work
pTAT101-CE103A	As pTAT101, TatC E103A exchange	This work
pTAT101-CE103A-BL9Q	As pTAT101-CE103A, TatB L9Q exchange	This work
pTAT101-CE103A-BL10P	As pTAT101-CE103A, TatB L10P exchange	This work
pTAT101-CE103A-BF13Y	As pTAT101-CE103A, TatB F13Y exchange	This work
pTAT101-CE103A-BI36N	As pTAT101-CE103A, TatB I36N exchange	This work
pTAT101-CE103K	As pTAT101, TatC E103K exchange	(7)
pTAT101-CE103K-BL9Q	As pTAT101-CE103K, TatB L9Q exchange	This work
pTAT101-CE103K-BL10P	As pTAT101-CE103K, TatB L10P exchange	This work

pTAT101-CE103K-BF13Y	As pTAT101-CE103A, TatB F13Y exchange	This work
pTAT101-CE103K-BI36N	As pTAT101-CE103A, TatB I36N exchange	This work
pTAT101-CP48L	As pTAT101, TatC P48L exchange	(9)
pTAT101-CP48L- BL9Q	As pTAT101-CP48L, TatB L9Q exchange	This work
pTAT101-CP48L -BL10P	As pTAT101-CP48L, TatB L10P exchange	This work
pTAT101-CP48L -BF13Y	As pTAT101-CP48L, TatB F13Y exchange	This work
pTAT101-CM59K	As pTAT101, TatC M59K exchange	(9)
pTAT101-CM59K-BL9Q	As pTAT101-CM59K, TatB L9Q exchange	This work
pTAT101-CM59K -BL10P	As pTAT101-CM59K, TatB L10P exchange	This work
pTAT101-CM59K -BF13Y	As pTAT101-CM59K, TatB F13Y exchange	This work
pTAT101-CV145E	As pTAT101, TatC V145E exchange	(9)
pTAT101-CV145E-BL9Q	As pTAT101-CV145E, TatB L9Q exchange	This work
pTAT101-CV145E -BL10P	As pTAT101-CV145E, TatB L10P exchange	This work
pTAT101-CV145E -BF13Y	As pTAT101-CV145E, TatB F13Y exchange	This work
pTAT101-CD211K	As pTAT101, TatC D211K exchange	This work
pTAT101-CD211K-BL9Q	As pTAT101-CD211K, TatB L9Q exchange	This work
pTAT101-CD211K -BL10P	As pTAT101-CD211K, TatB L10P exchange	This work
pTAT101-CD211K -BF13Y	As pTAT101-CD211K, TatB F13Y exchange	This work
pTAT101-CQ215K	As pTAT101, TatC Q215K exchange	This work
pTAT101-CQ215K-BL9Q	As pTAT101-CQ215K, TatB L9Q exchange	This work
pTAT101-CQ215K -BL10P	As pTAT101-CQ215K, TatB L10P exchange	This work
pTAT101-CQ215K -BF13Y	As pTAT101-CQ215K, TatB F13Y exchange	This work
pTAT101 cys less	As pTAT101, All 4 cys codons in *tatC* substituted with ala	(9)
pTAT101 cys less CM205C	As pTAT101 cys less, TatC M205C exchange	(9)
pTAT101 cys less BL9Q CM205C	As pTAT101 cys less CM205C, TatB L9Q exchange	This work
pTAT101 cys less BL10P CM205C	As pTAT101 cys less CM205C, TatB L10P exchange	This work
pTAT101 cys less BF13Y CM205C	As pTAT101 cys less CM205C, TatB F13Y exchange	This work
pTAT101 cys less BI36N CM205C	As pTAT101 cys less CM205C, TatB I36N exchange	This work
pTAT101 cys less CF94Q M205C	As pTAT101 cys less CM205C, TatC F94Q	This work
pTAT101 cys less BL10P CF94Q M205C	As pTAT101 cys less CF94Q M205C, TatB L10P	This work

7

pTAT101 cys less BF13Y CF94Q M205C	As pTAT101 cys less CF94Q M205C, TatB F13Y	This work
pQE80-CueO	As pQE80, carrying $cueO_{his}$	(4)
pQE80-CueO^{KK}h	As pQE80-CueO, CueO R3K, R4K exchange	(4)
pTAT1d	Medium copy number vector expressing TatABC under the control of *tat* promoter. Amp^r.	(10)
pUNIPROM	pT7.5 vector carrying a *tat* promoter. Backbone of pTAT1d	(11)
pTAT1d-CF94Q	As pTAT1d, TatC F94Q exchange	This work
pTAT1d-CF94A	As pTAT1d, TatC F94A exchange	This work
pTAT1d-CF94D	As pTAT1d, TatC F94D exchange	This work
pTAT1d-CF94G	As pTAT1d, TatC F94G exchange	This work
pTAT1d-CF94K	As pTAT1d, TatC F94K exchange	This work
pTAT1d-CF94P	As pTAT1d, TatC F94P exchange	This work
pTAT1d-CF94R	As pTAT1d, TatC F94R exchange	This work
pTAT1d-CF94S	As pTAT1d, TatC F94S exchange	This work
pTAT1d-CF94Q-BL9Q	As pTAT1d-CF94Q, TatB L9Q exchange	This work
pTAT1d-CF94Q-BL10P	As pTAT1d-CF94Q, TatB L10P exchange	This work
pTAT1d-CF94Q-BF13Y	As pTAT1d-CF94Q, TatB F13Y exchange	This work
pTAT1d-CF94Q-BI36N	As pTAT1d-CF94Q, TatB I36N exchange	This work
pTATBC1d	pUNIPROM carrying *tatBC*	This work
pSUAmiA	pSU18 carrying *amiA*	(12)
pSUAmiA-RD	As pSUAmiA, R14D exchange	This work
pSUAmiA-RE	As pSUAmiA, R14E exchange	This work
pSUAmiA-RH	As pSUAmiA, R14H exchange	This work
pSUAmiA-RN	As pSUAmiA, R14N exchange	This work
pSUAmiA-RQ	As pSUAmiA, R14Q exchange	This work
pSUAmiA-KH	As pSUAmiA, R13K, R14H exchange	This work
pSUAmiA-KQ	As pSUAmiA, R13K, R14Q exchange	This work
pSUAmiA-HH	As pSUAmiA, R13H, R14H exchange	This work
pSUSufIss-mAmiA	pSU18, carrying SufIss-mAmiA	This work
pSUSufIss-mAmiA-RD	As pSUSufIss-mAmiA, SufI R6D exchange	This work
pSUSufIss-mAmiA-RE	As pSUSufIss-mAmiA, SufI R6E exchange	This work
pSUSufIss-mAmiA-RH	As pSUSufIss-mAmiA, SufI R6H exchange	This work
pSUSufIss-mAmiA-RN	As pSUSufIss-mAmiA, SufI R6N exchange	This work
pSUSufIss-mAmiA-RQ	As pSUSufIss-mAmiA, SufI R6Q exchange	This work
pSUSufIss-mAmiA-KH	As pSUSufIss-mAmiA, SufI R5K, R6H exchange	This work
pSUSufIss-mAmiA-KQ	As pSUSufIss-mAmiA, SufI R5K, R6Q exchange	This work
pSUSufIss-mAmiA-KK	As pSUSufIss-mAmiA, SufI R5K, R6K exchange	This work
pSUSufIss-mAmiA-HH	As pSUSufIss-mAmiA, SufI R5H, R6H exchange	This work
pSUSufIssnoH-mAmiA	As pSUSufIss-mAmiA, *sufI* Δ11-21	This work
pSUmAmiA	As pSUAmiA, *amiA* Δ2-34	This work
pFAT75ΔA-BC	As pQEABC, but with *tatA* gene in frame deleted	(13)
pFAT75ΔA-BC BE8K	As pFAT75ΔA-BC, TatB E8K exchange	This work
pFAT75ΔA-BC BF13Y	As pFAT75ΔA-BC, TatB F13Y exchange	This work
pFAT75ΔA-BC BI36N	As pFAT75ΔA-BC, TatB I36N exchange	This work
pFAT75ΔA-BC	As pFAT75ΔA-BC, TatC F94Q exchange	This work

8

CF94Q		
pFAT75ΔA-BC BL9Q CF94Q	As pFAT75ΔA-BC CF94Q, TatB L9Q exchange	This work
pFAT75ΔA-BC BL10P CF94Q	As pFAT75ΔA-BC CF94Q, TatB L10P exchange	This work
pFAT75ΔA-BC BF13Y CF94Q	As pFAT75ΔA-BC CF94Q, TatB F13Y exchange	This work
pFAT75ΔA-BC BI36NC CF94Q	As pFAT75ΔA-BC CF94Q, TatB I36N exchange	This work
pFAT75ΔA-BC-AmiAhis	As pFAT75ΔA-BC also producing C-terminally his-tagged AmiA	This work
pFAT75ΔA-BC-AmiARDhis	As pFAT75ΔA-BC-AmiAhis, AmiA R14D exchange	This work
pFAT75ΔA-BC-AmiARNhis	As pFAT75ΔA-BC-AmiAhis, AmiA R14N exchange	This work
pFAT75ΔA-BC-AmiAKKhis	As pFAT75ΔA-BC-AmiAhis, AmiA R13K, R14K exchange	This work
pFAT75ΔA-BC-AmiAKQhis	As pFAT75ΔA-BC-AmiAhis, AmiA R13K, R14Q exchange	This work
pFAT75ΔA-BF13YC-AmiAhis	As pFAT75ΔA-BC, TatBF13Y exchange	This work
pFAT75ΔA-BF13YC-AmiARDhis	As pFAT75ΔA-BF13YC-AmiAhis, AmiA R14D exchange	This work
pFAT75ΔA-BF13YC-AmiARNhis	As pFAT75ΔA-BF13YC-AmiAhis, AmiA R14N exchange	This work
pFAT75ΔA-BF13YC-AmiAKKhis	As pFAT75ΔA-BF13YC-AmiAhis, AmiA R13K, R14K exchange	This work
pFAT75ΔA-BF13YC-AmiAKQhis	As pFAT75ΔA-BF13YC-AmiAhis, AmiA R13K, R14Q exchange	This work
pFAT75ΔA-BC-mAmiAhis	As pFAT75ΔA-BC also producing C-terminally his-tagged mature AmiA	This work
pFAT75ΔA-BF13YC-mAmiAhis	As pFAT75ΔA-BC-mAmiAhis, TatBF13Y exchange	This work
pQE70-mAmiA	pQE70 producing C-terminally his-tagged mature AmiA	This work
pQE70-mAmiC	pQE70 producing C-terminally his-tagged mature AmiC	This work
pSufIss-GFPhis	As pCDFDuet-1, carrying synthetic SufI signal sequence-fused GFPhis	This work
pSufIssRD-GFPhis	As pSufIss-GFPhis, SufI R6D exchange	This work
pSufIssRN-GFPhis	As pSufIss-GFPhis, SufI R6N exchange	This work
pSufIssKK-GFPhis	As pSufIss-GFPhis, SufI R5K, R6K exchange	This work
pQE80 sufIhis	pQE80 carrying $sufI_{his}$	This work
pQE80 RDsufIhis	pQE80 sufIhis SufI R6D exchange	This work
pQE80 RNsufIhis	pQE80 sufIhis SufI R6N exchange	This work
pQE80 KQsufIhis	pQE80 sufIhis SufI R5K, R6Q exchange	This work
pMAK705	Cloning vector with a temperature-sensitive replicon	(14)
pMAK-AupBC	As pMAK705, carrying 500 bp upstream sequence of *tatA* and *tatBC* sequence	This work
pMAK-AupBC-BL9Q	As pMAK-AupBC, TatB L9Q exchange	This work
pMAK-AupBC-BL10P	As pMAK-AupBC, TatB L10P exchange	This work
pMAK-AupBC-BF13Y	As pMAK-AupBC, TatB F13Y exchange	This work

pMAK-AupBC-BI36N	As pMAK-AupBC, TatB I36N exchange	This work
pMAK-AupBC-CF94Q	As pMAK-AupBC, TatC F94Q exchange	This work
pMAK-AupBC- BL9Q CF94Q	As pMAK-AupBC, TatB L9Q, TatC F94Q exchange	This work
pMAK-AupBC- BL10P CF94Q	As pMAK-AupBC, TatB L10P, TatC F94Q exchange	This work
pMAK-AupBC- BF13Y CF94Q	As pMAK-AupBC, TatB F13Y, TatC F94Q exchange	This work
pMAK-AupBC- BI36N CF94Q	As pMAK-AupBC, TatB I36N, TatC F94Q exchange	This work
pBAD24	Arabinose-inducible protein expression vector	(15)
pBADTatABChis	As pBAD24, carrying *tatABC*his	This work
pBADTatABChis-BL9Q	As pBADTatABChis, TatB L9Q exchange	This work
pBADTatABChis-BL10P	As pBADTatABChis, TatB L10P exchange	This work
pBADTatABChis-BF13Y	As pBADTatABChis, TatB F13Y exchange	This work
pBADTatABChis-BI36N	As pBADTatABChis, TatB I36N exchange	This work
pBAD22SecY(Δplug)EG	pBAD22, producing SecY(Δcodons62-72)EG	Ian Collinson
p101C*TatBC	Low copy vector for expression of *tatBC* from the *tatA* promoter with a modified RBS	(4)
p101C*BCflag	p101C*BC derivative producing TatB and C-terminally flag-tagged TatC	This work
p101C*BCflag E8K	As p101C*BCflag, TatB E8K exchange	This work
p101C*BCflag F13Y	As p101C*BCflag, TatB F13Y exchange	This work

Table S4. Plasmids used and constructed in this study

Primer name	Sequence (5'-3')
AmiARDf	CTCACTTCGCGCGACCAGGTGCTG
AmiARDr	CAGCACCTGGTCGCGCGAAGTGAG
AmiAREf	CTCACTTCGCGCGAACAGGTGCTG
AmiAREr	CAGCACCTGTTCGCGCGAAGTGAG
AmiARNf	CTCACTTCGCGCAACCAGGTGCTG
AmiARNr	CAGCACCTGGTTGCGCGAAGTGAG
AmiARQf	CTCACTTCGCGCCAACAGGTGCTG
AmiARQr	CAGCACCTGTTGGCGCGAAGTGAG
AmiARHf	CTCACTTCGCGCCACCAGGTGCTG
AmiARHr	CAGCACCTGGTGGCGCGAAGTGAG
AmiAHHf	CTCACTTCGCACCACCAGGTGCTG
AmiAHHr	CAGCACCTGGTGGTGCGAAGTGAG
AmiAKHf	CTCACTTCGAAACACCAGGTGCTG
AmiAKHr	CAGCACCTGGTGTTTCGAAGTGAG
AmiAKQf	CTCACTTCGAAACAACAGGTGCTG
suflssFE	CCGGAATTCGTTTTACATGGAGCAAATATG
suflssR	GTTCGTCTTTTGCGCTGGCCTTCAGGG
amiA-mF	GGCCAGCGCAAAAGACGAACTTTTAAAAACC
amiA-mRX	GACTCTAGATTATCGCTT TTTC
AmiAKQr	CAGCACCTGTTGTTTCGAAGTGAG
Suflss-RDf	GTCACTCAGTCGGGATCAGTTCATTCAGGC
Suflss-RDr	GCCTGAATGAACTGATCCCGACTGAGTGAC
Suflss-RHf	GTCACTCAGTCGGCATCAGTTCATTCAGGC
Suflss-RHr	GCCTGAATGAACTGATGCCGACTGAGTGAC
Suflss-RNf	GTCACTCAGTCGGAACCAGTTCATTCAGGC
Suflss-RNr	GCCTGAATGAACTGGTTCCGACTGAGTGAC
Suflss-RQf	GTCACTCAGTCGGCAGCAGTTCATTCAGGC
Suflss-RQr	GCCTGAATGAACTGCTGCCGACTGAGTGAC
Suflss-RKf	GTCACTCAGTCGGAACAGTTCATTCAGGC
Suflss-RKr	GCCTGAATGAACTGTTTCCGACTGAGTGAC
Suflss-KHf	GTCACTCAGTAAACATCAGTTCATTCAGGC
Suflss-KHr	GCCTGAATGAACTGATGTTTACTGAGTGAC
Suflss-KQf	GTCACTCAGTAAACAGCAGTTCATTCAGGC
Suflss-KQr	GCCTGAATGAACTGCTGTTTACTGAGTGAC
Suflss-KKf	GTCACTCAGTAAAAAACAGTTCATTCAGGC
Suflss-KKr	GCCTGAATGAACTGTTTTTTACTGAGTGAC
Suflss-HHf	GTCACTCAGTCATCATCAGTTCATTCAGGC
Suflss-HHr	GCCTGAATGAACTGATGATGACTGAGTGAC
Sufl-noHF	CGGCGTCAGTTCATTCAGCCCCTGAAGGCCAGCGCA
Sufl-noHR	TGCGCTGGCCTTCAGGGGCTGAATGAACTGACGCCG
AmiA-nossFE	CCGGAATTCTATTACAACTCAGGCCGTATGAAAGACGAACTTTT AAAAACCAG
AmiAFATApaI-F	GCGCGGGCCCATTAAAGAGGAGAAATTAACCATGAGCACTTTT AAACCACTAAAAAC
mAmiAFATApaI-F	GCGCGGGCCCATTAAAGAGGAGAAATTAACCATGAAAGACGAA CTTTTAAAAACCAG

mAmiA-SphI-F	GCGCGCATGCGAAAAGACGAACTTTTAAAAACC
AmiAnostopBamHI-R	CGCGGATCCTCGCTTTTTCGAATGTGCTTTC
mAmiC-SphI-F	GCGCGCATGCGAGCGGTCAGCCAGGTCGTG
AmiCnostopBamHI-R	CGCGGATCCTCCCCTTCTCGCCAGCGTC
C-F94S1	GGTGTGGGCATCTATCGCCCCAG
C-F94S2	CTGGGGCGATAGATGCCCACACC
C-F94A1	GGTGTGGGCAGCGATCGCCCCAG
C-F94A2	CTGGGGCGATCGCTGCCCACACC
C-F94K1	GGTGTGGGCAAAAATCGCCCCAG
C-F94K2	CTGGGGCGATTTTTGCCCACACC
C-F94Q1	GGTGTGGGCACAGATCGCCCCAG
C-F94Q2	CTGGGGCGATCTGTGCCCACACC
C-F94R1	GGTGTGGGCACGCATCGCCCCAG
C-F94R2	CTGGGGCGATGCGTGCCCACACC
C-F94P1	GGTGTGGGCACCGATCGCCCCAG
C-F94P2	CTGGGGCGATCGGTGCCCACACC
C-F94D1	GGTGTGGGCAGATATCGCCCCAG
C-F94D2	CTGGGGCGATATCTGCCCACACC
C-F94G1	GGTGTGGGCAGGCATCGCCCCAG
C-F94G2	CTGGGGCGATGCCTGCCCACACC
C-F94C1	GGTGTGGGCATGCATCGCCCCAG
C-F94C2	CTGGGGCGATGCATGCCCACACC
B-F6Yf	GTTTGATATCGGTTATAGCGAACTGC
B-F6Yr	GCAGTTCGCTATAACCGATATCAAAC
B-L9Qf	GGTTTTAGCGAACAGCTATTGGTG
B-L9Qr	CACCAATAGCTGTTCGCTAAAACC
B-L9Pf	GGTTTTAGCGAACCGCTATTGGTG
B-L9Pr	CACCAATAGCGGTTCGCTAAAACC
B-L10Pf	GCGAACTGCCATTGGTGTTCATC
B-L10Pr	GATGAACACCAATGGCAGTTCGC
B-F13Yf	GCTATTGGTGTACATCATCGGCC
B-F13Yr	GGCCGATGATGTACACCAATAGC
B-K30If	GTGGCGGTAATTACGGTAGCGG
B-K30Ir	CCGCTACCGTAATTACCGCCAC
B-I36Nf	GTAGCGGGCTGGAATCGCGCGTTGC
B-I36Nr	GCAACGCGCGATTCCAGCCCGCTAC
tatB E8K 1	GATATCGGTTTTAGCAAACTGCTATTGG
tatB E8K 2	CCAATAGCAGTTTGCTAAAACCGATATC
C-E103A1	CGCTGTATAAGCATGCGCGTCGCCTGGTGGTGC
C-E103A2	GCACCACCAGGCGACGCGCATGCTTATACAGCG
C-P48L1	GGTATCCGCGCTGTTGATCAAGC
C-P48L2	GCTTGATCAACAGCGCGGATACC
C-M59K1	GGTTCAACGAAGATCGCCACCG
C-M59K2	CGGTGGCGATCTTCGTTGAACC

C-V145E1	CGGAAGGGGAACAGGTATCCAC
C-V145E2	GTGGATACCTGTTCCCCTTCCG
C-D211K1	CTGACGCCGCCGAAAGTCTTCTCGCAAAC
C-D211K2	GTTTGCGAGAAGACTTTCGGCGGCGTCAG
C-Q215K1	GTCTTCTCGAAAACGCTGTTG
C-Q215K2	CAACAGCGTTTTCGAGAAGAC
TatB-L9QL10P-F	GGTTTTAGCGAACAGCCATTGGTGTTCATC
TatB-L9QL10P-R	GATGAACACCAATGGCTGTTCGCTAAAACC
TatB-L10PF13Y-F	GCGAACTGCCATTGGTGTACATCATCGG
TatB-L10PF13Y-R	CCGATGATGTACACCAATGGCAGTTCGC
TatA-FB	CAGAGGAGGATCCATGGG
TatB-RS	GTATCGTCGACAGACATGC
TatCR1d	CTTGGGCTGCAGCCTTATTCTTC
TatC93R	TGCCCACACCTGATAGAG
TatCm6	CTTCCTCGAGTGATAAACCTTAAGCATG
TatC95F	ATCGCCCCAGCGCTGTAT
TatAup1-Xba I	CGCTCTAGAGAAAACCCTGCTCTACGTC
TatAup2-ClaI	GCGCATCGATAAGCTTGATATCGAAT
TatA6B7-ClaI	GCGCATCGATGATAAAGAGCAGGTGTAATCCGTGTTTGATATCGGTTTTAGC
TatCrev-KpnI	CGGGGTACCTTATTCTTCAGTTTTTTCGCTTTC
TatANcoI	GCGCCCATGGGTGGTATCAGTATTTGG
HisXba	GCGCTCTAGATTAGTGATGGTGATGGTG
STIPE-ISH	GCGGATACGAATCAGGAACAG
pT7.5R	CGCTGAGATAGGTGCC.
p101C*BCflag_F	GAAAGCGAAAAAACTGAAGAAGACTACAAGGACGATGACAAGTAAGGCTGCAGGCATGCAAG
p101C*BCflag_R	CTTGCATGCCTGCAGCCTTACTTGTCATCGTCCTTGTAGTCTTCTTCAGTTTTTTCGCTTTC

Table S5. Oligonucleotides used in this study

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Figure S1. Substitutions of the twin arginines in the AmiA signal peptide prevent growth in the presence of SDS. Strain MCDSSAC Δ*tatABC* producing wild type *tatABC* from plasmid pTAT1d and either wild type ('RR') or signal peptide point-substituted AmiA, as indicated, from pSUAmiA. The strain and plasmid combinations were cultured overnight in LB medium supplemented with chloramphenicol and ampicillin (for plasmid selection), after which they were streaked onto LB agar containing the same antibiotics, with and without the addition of 2% SDS and incubated for 16 hr at 37°C.

Figure S2. TatB variants are able to restore Tat transport to a range of defective twin arginine substitutions in the AmiA signal sequence. Growth of MCDSSAC Δ*tatABC* coproducing the indicated TatB variants (with wild type *tatA* and *tatC*) from pTAT101, or the empty plasmid pTH19kr (indicated by 'Δ*tat*') alongside signal peptide variants of AmiA, on LB agar supplemented with chloramphenicol and kanamycin, with and without the addition of 2% SDS as indicated. An 8μl aliquot of each strain/plasmid combination following aerobic growth to an OD$_{600}$ of 1.0 was spotted and incubated for 16 hr at 37°C. A. Individual signal peptide substitutions of AmiA (indicated to the left of each panel) were tested against the TatB suppressors F6Y, L9P, L9Q, L10P, F13Y, K30I and I36N. B. The TatB E8K suppressor was tested for the ability to suppress the indicated AmiA signal peptide substitutions.

Figure S3. TatB variants are able to restore Tat transport to a range of defective twin arginine substitutions in the SufI signal sequence. Growth of MCDSSAC Δ*tatABC* coproducing the indicated TatB variants (with wild type *tatA* and *tatC*) from pTAT101, or the empty plasmid pTH19kr (indicated by 'Δ*tat*') alongside signal peptide variants of SufI fused to the AmiA mature domain, on LB agar supplemented with chloramphenicol and kanamycin, with and without the addition of 2% SDS as indicated. An 8μl aliquot of each strain/plasmid combination following aerobic growth to an OD$_{600}$ of 1.0 was spotted and incubated for 16 hr at 37°C. A. Individual signal peptide substitutions of AmiA (indicated to the left of each panel) were tested against the TatB suppressors F6Y, L9P, L9Q, L10P, F13Y, K30I and I36N. B. The

TatB E8K suppressor was tested for the ability to suppress the indicated AmiA signal peptide substitutions

Figure S4. A subset of amino acid substitutions at TatCF94 abolish Tat activity when produced at medium and low copy number. A and C. Growth of DADE coproducing either wild type TatABC (Tat⁺), wild type TatAB alongside F94-substituted TatC or the cognate empty plasmid (Tat⁻) on LB agar containing 2% SDS. A single colony of each strain/plasmid combination was resuspended in 30µl of PBS and an 8µl aliquot was spotted onto LB agar supplemented with appropriate antibiotics, along with 2% SDS as indicated. Plates were incubated for 16 hr at 37°C. B and D. Detection of TatC protein present in membrane fractions of the same strain and plasmid combinations as in A. and C., respectively, by Western immunoblot with anti-TatC antiserum. A total of 5µg membranes was loaded per lane for TatC produced from pTAT1d (B) and 20µg per lane for membranes produced from strains harboring pTAT101 derivatives (D).

Figure S5. TatB variants cannot supress TatC inactivating substitutions outside of the signal peptide binding site. Growth of DADE (Δ*tatABCD*, Δ*tatE*) coproducing wild type TatA alongside and the indicated substitution in TatB alongside either of TatC P48L, TatC M59K, TatC V145E, TatC D211K or TatC Q215K as indicated, from plasmid pTAT101 on LB agar or LB agar containing 2% SDS. A single colony of each strain/plasmid combination was resuspended in 30µl of PBS and an 8µl aliquot was spotted onto LB agar supplemented with appropriate antibiotics, along with 2% SDS as indicated, and incubated for 16 hr at 37°C.

Figure S6. The suppressive effect of the TatB variants is not additive and mature AmiC is not exported in the presence of the TatB F13Y suppressor. A. Growth of DADE coproducing either wild type TatABC (Tat⁺), wild type TatAB alongside F94-substituted TatC or the cognate empty plasmid (Tat⁻) on LB agar or LB agar containing 2% SDS. B. Growth of MCDSSAC Δ*tatABC* coproducing the indicated TatB variants (with wild type *tatA* and *tatC*) from pTAT101, or the empty plasmid pTH19kr (indicated by 'Δ*tat*') alongside the RN or KK

17

signal peptide variants of SufI fused to the AmiA mature domain, as indicated, on LB agar with or without the addition of 2% SDS. C. Strain MC4100 Δ*ami*A Δ*amiC* Δ*tatABC* coproducing either wild-type TatB or TatB F13Y (with wild type *tatA* and *tatC*) from pTAT101 and the AmiA or AmiC mature domains (from pQE70-mAmiA or pQE70-mAmiC, respectively) on LB agar or LB agar containing 2% SDS. In each case a single colony of each strain/plasmid combination was resuspended in 30µl of PBS and an 8µl aliquot was spotted onto LB agar supplemented with appropriate antibiotics, along with 2% SDS where indicated. Plates were incubated for 16 hr at 37°C.

Figure S7. The TatB suppressors support export of his-tagged SufI with its native signal peptide. A. and B. *E. coli* strain DADE producing wild type TatA and TatC and the indicated TatB variants alongside wild-type SufI-his or the indicated signal-peptide variants were fractionated into whole cell (upper panels) and periplasm (lower panels) fractions, then analysed by Western blot with anti-6X His tag® or anti-RNA polymerase β subunit antibodies (cytoplasmic control protein). wc – whole cell.

Figure S8. TatBC and SufIss-GFP-His twin-arginine variants are detectable in whole cell samples. A. and B. Cells producing SufIss-GFP-His with the wild type (RR) or twin-arginine substituted SufI signal peptide, as indicated, alongside TatC and either wild type TatB or the E8K, F13Y or I36N substituted variants, or C. and D. Cells producing SufIss-GFP-His with the wild type SufI signal peptide along with either wild type TatBC, the TatC F94Q allele along with either wild type TatB or the L9Q, L10P, F13Y or I36N substituted variants, or the TatC E103K allele along with either wild type TatB or the L9Q, L10P, F13Y or I36N substituted variants, as indicated were harvested and resuspended in PBS. A. and C. The fluorescence intensity and OD_{600} of the samples were measured using a plate reader and the Fluorescence/OD_{600} plotted for each sample. B. and D. 20 µl of each cell suspension was taken, all samples were normalized to the same OD_{600} and then analysed by SDS-PAGE followed by western blot using a TatB-TatC mixed antibody.

18

Figure S9. TatBC complexes containing the TatB F13Y suppressor do not co-purify with signal peptide variants of AmiA. C-terminally his-tagged wild type AmiA, twin-arginine substituted AmiA or signal sequence-less AmiA, as indicated was co-produced alongside wild type TatBC or TatBF13Y/TatC and purified using nickel beads from digitonin–treated cell extracts. Aliquots of the load and elution fractions were subject to SDS-PAGE followed by Western blot using either anti-His, anti-TatB and TatC antibodies.

Figure S10. TatB variants are extracted from the membrane with digitonin. Membrane suspensions (containing equivalent amounts of total protein) from strain DADE coproducing either wild type TatABC or wild type TatA and TatC alongside the indicated amino acid variant of TatB were solubilized by addition of 2% digitonin and incubation on ice for 30 min. Samples total membranes and digitonin solubilized material (each containing 10μg protein) were analysed by SDS-PAGE followed by western blotting with anti-TatA, anti-TatB or anti-TatC antibodies as indicated.

Figure S11. Constitutive oligomerisation of TatA is not promoted by the TatB L9Q, L10P or I36N substitutions. Fluorescence images of TatA-YFP in representative cells of A. strains AyBCE or AyBC$_{F94Q}$E (encoding chromosomal TatC F94Q) in the presence (pAmiA) or absence of plasmid-encoded wild type AmiA, as indicated (reproduced from Fig 5A). B. strains AyB$_{L9Q}$CE (encoding chromosomal TatB L9Q), AyB$_{L10P}$CE (encoding chromosomal TatB L10P) and AyB$_{I36N}$CE (encoding chromosomal TatB I36N) or the same strains additionally harboring the chromosomally-encoded TatC F94Q substitution. Scale bar: 1 μm. Note that the pictures in panel A are identical to those in Fig 5A and were included here to provide a direct comparison with panel B.

LB   LB + 2% SDS

Fig S1

A

WT TatB	Δtat	TatB F6Y
TatB L9P	TatB L9Q	TatB L10P
TatB F13Y	TatB K30I	TatB I36N

WT TatB	Δtat	TatB F6Y
TatB L9P	TatB L9Q	TatB L10P
TatB F13Y	TatB K30I	TatB I36N

RN

RR

RQ

RD

KH

RE

KQ

RH

HH

LB   LB + 2% SDS   LB   LB + 2% SDS

B   TatB E8K

LB   LB + 2% SDS

AmiA RR	AmiA RD	AmiA RE
AmiA RN	AmiA RQ	AmiA RH
AmiA HH	AmiA KH	AmiA KQ

Fig S2

A

WT TatB	Δ*tat*	TatB F6Y
TatB L9P	TatB L9Q	TatB L10P
TatB F13Y	TatB K30I	TatB I36N

WT TatB	Δ*tat*	TatB F6Y
TatB L9P	TatB L9Q	TatB L10P
TatB F13Y	TatB K30I	TatB I36N

RN

RR

RQ

RD

KH

RE

KQ

RH

HH

LB   LB + 2% SDS   LB   LB + 2% SDS

B

TatB E8K

Sufl RR	Sufl RD	Sufl RE
Sufl RN	Sufl RQ	Sufl RH
Sufl HH	Sufl KH	Sufl KQ

LB   LB + 2% SDS

Fig S3

Fig S4

Fig S5

Fig S6

Fig S7

Fig S8

Fig S9

Fig S10

A AyBCE   AyBCE pSUAmiA   AyBC$_{F94Q}$E   AyBC$_{F94Q}$E pSUAmiA

B AyB$_{L9Q}$CE   AyB$_{L9Q}$C$_{F94Q}$E   AyB$_{L10P}$CE   AyB$_{L10P}$C$_{F94Q}$E

AyB$_{I36N}$CE   AyB$_{I36N}$C$_{F94Q}$E

Fig S11