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Independent mobility of proteins and lipids in the plasma membrane of *Escherichia coli*

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**Introduction**

The standard structural description for biological membranes for over 30 years has been the fluid mosaic model of Singer and Nicolson (1972) which envisages membrane-integrated proteins immersed in a ‘sea’ of lipid. Unless they are immobilized by other interactions (e.g., with cytoskeletal proteins) membrane proteins are free to diffuse among the surrounding lipid molecules. This model has been modified in recent years as a result of numerous studies showing patchy or compartmentalized distributions of specific lipids and proteins in both eukaryotic and prokaryotic cell membranes (reviewed by Vereb et al., 2003; Engelman, 2005; Matsumoto et al., 2006; Jacobson et al., 2007; Milekovskaya and Dowhan, 2009). Fluorescent protein tagging of various bacterial plasma membrane proteins has revealed their concentration in mobile microdomains (Johnson et al., 2004; Lenn et al., 2008; López and Kolter, 2010; Llorente-Garcia et al., 2014). However, the forces that cause the clustering of specific proteins remain obscure.

Current membrane models suggest that the diffusion of all membrane components must be strongly influenced by the viscosity of the lipid bilayer, which in turn is influenced by temperature and lipid composition (Murata, 1989; Thompson, 1989; Fulbright et al., 1997; Jin et al., 1999; Szalontai et al., 2000; Lindblom et al., 2002; Sarcina et al., 2003). When most biological membranes are cooled to around 10°C below growth temperature, there is a step decrease in the fluidity of the lipid bilayer, attributed to a phase transition from the fluid liquid crystalline state to the more viscous crystalline gel state, observed by measuring the mobility of reporter molecules embedded in the lipid bilayer (Fulbright et al., 1997; Jin et al., 1999; Lindblom et al., 2002; Sarcina et al., 2003). Cooling below phase transition has pronounced effects on the translational and rotational diffusion of lipids, and on the flexing of the fatty acid tails (Szalontai et al., 2000; Lindblom et al., 2002; Denich et al., 2003; Sarcina et al., 2003). Bacteria such as *Escherichia coli* alter their membrane lipid composition in response to environmental and physiological stress to maintain a constant membrane viscosity by ‘homeoviscous adaptation’ (Morein et al., 1996; Rilfors and Lindblom, 2002).

**Summary**

Fluidity is essential for many biological membrane functions. The basis for understanding membrane structure remains the classic Singer-Nicolson model, in which proteins are embedded within a fluid lipid bilayer and able to diffuse laterally within a sea of lipid. Here we report lipid and protein diffusion in the plasma membrane of live cells of the bacterium *Escherichia coli*, using Fluorescence Recovery after Photobleaching (FRAP) and Total Internal Reflection Fluorescence (TIRF) microscopy to measure lateral diffusion coefficients. Lipid and protein mobility within the membrane were probed by visualizing an artificial fluorescent lipid and a simple model membrane protein consisting of a single membrane-spanning alpha-helix with a Green Fluorescent Protein (GFP) tag on the cytoplasmic side. The effective viscosity of the lipid bilayer is strongly temperature-dependent, as indicated by changes in the lipid diffusion coefficient. Surprisingly, the mobility of the model protein was unaffected by changes in the effective viscosity of the bulk lipid, and TIRF microscopy indicates that it clusters in segregated, mobile domains. We suggest that this segregation profoundly influences the physical behaviour of the protein in the membrane, with strong implications for bacterial membrane function and bacterial physiology.
It has been widely assumed that lipid phase transition must also affect the mobility of membrane proteins. However, there is surprisingly little direct evidence that lipid phase transition affects membrane protein mobility. In a classic early experiment (Frye and Edidin, 1970), the mixing of cell surface antigens in fused cultured mammalian cells is retarded at low temperatures, and this is attributed to slower membrane protein diffusion due to greater lipid viscosity. More recent studies using more direct measuring techniques have shown that rotational (Aisenbrey and Bechinger, 2004) and translational (Tominaga et al., 2004) diffusion of proteins in reconstituted model membranes is slower below the lipid phase transition temperature. However real biological membranes have more complicated mixtures of lipids and proteins, and it cannot be assumed that their behaviour is the same.

In a study of the dynamic properties of the cytoplasm of *E. coli*, we used a series of fluorescent protein constructs expressed in the cell to determine the size-dependence of cytoplasmic diffusion. The behaviour of the constructs was simpler and more predictable than that of indigenous proteins; clearly the behaviour of the indigenous proteins was complicated by interactions with partners in the cell (Nenninger et al., 2010). In the present study we have applied a comparable approach to study the 'generic' behaviour of membrane proteins by expressing in *E. coli* cells a simple, non-native membrane protein with a GFP tag. We used confocal Fluorescence Recovery after Photobleaching (FRAP) and video-rate Total Internal Reflection Fluorescence (TIRF) microscopy to examine the distribution and mobility of this protein as a function of temperature. For comparison, we carried out similar measurements on a fluorescent lipid probe and a well-characterized GFP-tagged native membrane protein.

**Results**

**Expression of a model membrane protein in E. coli**

We constructed a gene coding for a model membrane protein using as a starting point a truncated version of open reading frame *sll1021* from the cyanobacterium *Synechocystis* sp. PCC6803, coding for a membrane protein with no identified homologues in *E. coli*. The protein has unknown function but has been identified in the plasma membrane of *Synechocystis* (Huang et al., 2002). The predicted gene product has 673 amino acids with a single predicted transmembrane alpha-helix close to the N-terminus. The ends of the predicted transmembrane helix lack any of the two-residue motifs that appear important for specific interactions with phospholipid head-groups (Hunte, 2005). We took a portion of the *sll1021* sequence coding for 38 amino acids including the predicted transmembrane alpha-helix and fused it in-frame to the gene coding for GFPmut3* (Cormack et al., 1996) with a linker of five asparagine residues. The construct was expressed in *E. coli* cells from the arabinose-inducible pBAD24 vector (Guzman et al., 1995). The predicted topology of the model protein (helix1021-GFP) is illustrated in Fig. 1A. The GFP domain is almost certainly on the cytoplasmic side of the membrane, since there are no Sec or Tat leader sequences present to initiate the translocation of GFP into the periplasm (Thomas et al., 2001; Natale et al., 2008). Cells expressing helix1021-GFP were fractionated and proteins of the periplasmic, cytoplasmic and plasma membrane fractions separated by SDS-PAGE and probed with anti-GFP antibody. GFP was only detected in the plasma membrane fraction, where the antibody recognized a...
protein migrating at 28–29 kDa, consistent with the predicted gene product (Fig. 1B). The band ran at a slightly higher molecular weight than free GFP, confirming that the protein integrates into the plasma membrane as expected, and the GFP tag is not cleaved off the transmembrane helix. Confocal fluorescence micrographs of GFP fluorescence from cells expressing the protein are consistent with membrane localization. Images recorded at high z-axis resolution show green fluorescence confined to a halo at the periphery of the cell (Fig. 1C), a characteristic of confocal images of GFP-tagged membrane proteins in *E. coli* (Mullineaux *et al.*, 2006).

**Localization of the fluorescent lipid probe BODIPY FL-C12 in *E. coli* cells**

For comparison with the behaviour of our model membrane protein we used an artificial fluorescent lipid, BODIPY FL-C12, consisting of a green fluorophore linked to a 12-carbon fatty acyl tail. This molecule was previously used as a probe of membrane fluidity in a cyanobacterium (Sarcina *et al.*, 2003). *E. coli* cells stained with BODIPY FL-C12 again show a fluorescent halo in confocal images (Fig. 2A). We sometimes observed gaps in staining close to sites of cell division (Fig. 2A), as previously observed with lipophilic cytoplasmic membrane stains (Fishov and Woldringh, 1999). The fluorescent halo is consistent with localization in the cytoplasmic membrane; however, at optical resolution it is not possible to directly distinguish the cytoplasmic (inner) and outer membranes in these cells. Using plasmolysis (Fishov and Woldringh, 1999) in cells elongated by growth in the presence of the cell division inhibitor cephalexin (Mullineaux *et al.*, 2006) creates a large enough gap in the plasmolysis bays to permit fluorescence in the inner and outer membranes to be distinguished (Fishov and Woldringh, 1999). As previously observed (Fishov and Woldringh, 1999), plasmolysed cells do not show clear fluorescent halos (Fig. 2B), likely due to cytoplasmic shrinkage and cytoplasmic membrane infolding. No fluorescence could be detected in the outer cell layers in the plasmolysis bays, indicating that BODIPY FL-C12 localizes to the cytoplasmic membrane (Fig. 2B).

**Measuring the lateral mobility of lipids and proteins using confocal FRAP**

We used the techniques described and illustrated in Mullineaux *et al.* (2006) to measure diffusion coefficients for BODIPY FL-C12 and GFP-tagged plasma membrane proteins. Cells were elongated by growth in the presence of cephalexin, without detectable diffusion barriers along the length of the cell (Elowitz *et al.*, 1999; Mullineaux *et al.*, 2006). The use of elongated cells has two advantages. First, it increases the time required for fluorescence to re-equilibrate after bleaching, allowing more accurate determination of diffusion coefficients. In short cells, re-equilibration can be very rapid because of the limited diffusion space available to the fluorophore. Second, provided that the initial bleach is narrow compared to the length of the cell, it allows a simple data analysis in which complications due to cell poles can be ignored (Mullineaux *et al.*, 1997). FRAP measurements were carried out using a wide confocal pinhole to give relatively low z-axis resolution, such that the two-dimensional fluorescence image sums information from the full volume of the cell. Note that this makes the resolution of the membrane halo less clear than in high-resolution images (Figs 1 and 2), which also have the benefit of a better signal-to-noise ratio because of slower image acquisition and signal averaging. The bleach was carried out by scanning a diffraction-limited line at high laser intensity across the short axis of the cell. Cells were then imaged at several time points after the bleach. Note that the initial bleach appears much broader than the diffraction-limited laser line, due to movement of the fluo-
rophore during the bleach and before acquisition of the first post-bleach image. The depth of the bleach that can be obtained is also decreased by movement of the fluorophore during the bleach. Neither effect precludes estimation of the diffusion coefficient, provided that fluorescence distribution is still in disequilibrium when the image sequence commences (Mullineaux et al., 2006). One-dimensional profiles of fluorescence along the long-axis of the cell were extracted, and diffusion coefficients estimated using a previously published protocol that takes into account the initial measured width of the bleach and the time-dependence of fluorescence recovery at the bleach (Mullineaux et al., 1997; 2006). A representative FRAP image sequence for helix1021-GFP is shown in Fig. 3, and full representative measurements with data analysis are shown in Supplementary Fig. S1 (helix1021-GFP) and Supplementary Fig. S2 (BODIPY FL-C12).

**Effect of temperature on the mobility of BODIPY FL-C12**

The green fluorescent fatty acid derivative BODIPY FL-C12 has previously been used as a probe of the fluidity of thylakoid membranes in cyanobacteria, where there is a pronounced dependence of its diffusion coefficient on temperature and the extent of fatty acid desaturation in the membrane (Sarcina et al., 2003). We used confocal FRAP to measure the diffusion of BODIPY FL-C12 in the cytoplasmic membrane of *E. coli* as a function of temperature, for cells grown at 37°C. Supplementary Fig. S2 shows a representative measurement with data analysis. When cells were cooled we observed a step decrease in the lateral diffusion coefficient at about 28°C (Fig. 4). This resembles a classic phase transition from the liquid crystal to the crystalline gel states of the lipid bilayer, which in biological membranes usually occurs around 5–10°C below growth temperature (Hazel and Williams, 1990). However, as the *E. coli* cytoplasmic membrane is a very complex mixture of lipids and proteins, we cannot exclude the possibility that the change in BODIPY FL-C12 mobility is due to effects other than a simple lipid phase transition.

Cells generally acclimate to sustained growth at different temperatures by changing their membrane lipid composition, resulting in changes in fluidity and phase transition temperature (Thompson, 1989). Growth of *E. coli* at lower temperatures led to differences in the temperature dependence of BODIPY FL-C12 mobility (see Supplementary Fig. S3). Some effects of growth at different temperatures on the BODIPY FL-C12 diffusion coefficient are shown in Fig. 5A: note the higher BODIPY FL-C12 diffusion coefficient in cells grown at lower temperatures. At the extremes of the conditions tested, the diffusion coefficient of BODIPY FL-C12 differed by a factor of more than 4, ranging from 0.6 ± 0.2 μm² s⁻¹ (mean ± SD) in cells grown at 37°C and measured at 17°C to 2.7 ± 0.5 μm² s⁻¹ in cells grown and measured at 32°C (Figs 4 and 5).

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Effects of membrane protein expression and lipid biosynthesis mutations on the mobility of BODIPY FL-C_12

To further explore the factors controlling the mobility of BODIPY FL-C_12, we measured its diffusion coefficient in cells with different membrane protein and lipid compositions (summarized in Table 1). To perturb membrane protein composition, we used a mutant lacking the entire twin-arginine translocation (tat) operon (Wexler et al., 2000), with and without a pBad plasmid overexpressing tatABC (Bolhuis et al., 2001). After arabinose induction, the TatABC proteins pack the membrane (Bolhuis et al., 2001). However, neither condition measurably perturbed the BODIPY FL-C_12 diffusion coefficient (Table 1). We then also tried a series of mutants with deletions of genes coding for lipid biosynthesis enzymes (Dowhan, 1997): cardiolipin synthetase (cls); pssA (required for phosphatidylserine synthesis); pgsA (phosphatidylglycerophosphate synthetase) and phosphatidylserine decarboxylase (psd). Some of these mutants showed significant alteration of the BODIPY FL-C_12 diffusion coefficient, by a factor of up to \( \sim 2 \) in either direction (Table 1).

**Mobility of membrane proteins probed by FRAP**

The same confocal FRAP techniques were used to measure lateral diffusion coefficients for two GFP-tagged proteins probed by FRAP.

**Table 1.** Effects of mutations affecting membrane composition on protein and lipid mobility.

<table>
<thead>
<tr>
<th>E. coli strain</th>
<th>Effect on membrane composition</th>
<th>BODIPY FL-C_12 diffusion coefficient (( \mu m^2 s^{-1} )) ± SD</th>
<th>Helix1021-GFP diffusion coefficient (( \mu m^2 s^{-1} )) ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild-type (DH5(\alpha))</td>
<td></td>
<td>1.2 ± 0.30</td>
<td>0.43 ± 1.3</td>
</tr>
<tr>
<td>(\Delta \text{tat})</td>
<td>Loss of twin-arginine translocon (Tat) subunits (Wexler et al., 2000)</td>
<td>1.3 ± 0.58</td>
<td>0.32 ± 0.08</td>
</tr>
<tr>
<td>(\Delta \text{tat}) overexpressing tatABC</td>
<td>Dense packing of the membrane with Tat subunits (Bolhuis et al., 2001)</td>
<td>1.3 ± 0.25</td>
<td></td>
</tr>
<tr>
<td>(\Delta \text{pp})</td>
<td>Loss of lipoprotein Lpp</td>
<td>1.06 ± 0.71</td>
<td>0.38 ± 0.15</td>
</tr>
<tr>
<td>(\Delta \text{pp} \Delta \text{pgsA})</td>
<td>Loss of lipoprotein Lpp and reduced levels of phosphatidyglycerol and cardiolipin (Dowhan, 1997)</td>
<td>0.66 ± 0.22</td>
<td>0.43 ± 0.22</td>
</tr>
<tr>
<td>(\Delta \text{cls})</td>
<td>Loss of the major pathway for cardiolipin synthesis (Dowhan, 1997)</td>
<td>1.04 ± 0.41</td>
<td>0.50 ± 0.19</td>
</tr>
<tr>
<td>(\Delta \text{psdA})</td>
<td>Loss of phosphatidyserine and phosphatidylethanolamine (Dowhan, 1997)</td>
<td>2.4 ± 0.84</td>
<td>0.32 ± 0.07</td>
</tr>
<tr>
<td>(\Delta \text{psd})</td>
<td>Loss of phosphatidylethanolamine (Dowhan, 1997)</td>
<td>1.6 ± 0.53</td>
<td>0.52 ± 0.10</td>
</tr>
</tbody>
</table>

All cells were grown and measured at 37°C. Mean diffusion coefficients (± SD, \( n = 10 \)) are shown.
plasma membrane proteins; the simple model membrane protein helix1021-GFP and TatA-GFP, a tagged version of one of the three essential membrane-spanning proteins of the twin-arginine translocon (Tat) in *E. coli* (Ray et al., 2005). The mobility of TatA-GFP in the membrane has previously been investigated (Mullineaux et al., 2006; Leake et al., 2008). In both cases, the GFP tag is expected to be located on the cytoplasmic side of the membrane (Ray et al., 2005). Lateral diffusion coefficients were measured for helix1021-GFP and TatA-GFP at a range of measurement temperatures, for cells grown at different temperatures (Fig. 5). Both proteins diffused slower under all conditions than BODIPY FL-C12, and under most conditions diffusion of TatA-GFP was significantly slower than helix1021-GFP (Fig. 5). More surprisingly, the mobility of the two proteins did not show the same temperature dependence as BODIPY FL-C12, with no measurable change in mobility at the phase transition temperature (Fig. 5B). The standard deviations largely reflect real differences between individual cells, as the estimated errors in individual measurements are small. The mean diffusion coefficient of helix1021-GFP showed no significant temperature dependence over the range tested (one-way ANOVA, *P* = 0.179, *n* = 10). Pooling the entire set of data for different growth and measurement temperatures (Fig. 5), helix1021-GFP showed a mean diffusion coefficient (± SD) of 0.4 ± 0.1 μm² s⁻¹ (n = 60) with no significant differences under any of the conditions tested (one-way ANOVA, *P* = 0.043, *n* = 10).

We measured helix1021-GFP diffusion in the set of lipid biosynthesis mutants also used to measure effects of membrane lipid composition on BODIPY FL-C12 diffusion (Table 1). Effects on helix1021-GFP diffusion were generally much smaller than with BODIPY FL-C12 diffusion, and there was no obvious correlation between the effects on helix1021-GFP and BODIPY FL-C12 diffusion (Table 1). For example, the BODIPY FL-C12 diffusion coefficient was respectively increased and decreased by (Table 1). For example, the BODIPY FL-C12 diffusion coefficient was respectively increased and decreased by factors of about 2 in the ΔpssA and ΔppΔpgsA mutants, with no significant effects on helix1021-GFP diffusion (paired *t*-test, *P* = 0.129, *n* = 10 (Table 1).

The diffusion coefficient of TatA-GFP at different measuring temperatures (Fig. 5B) showed significant variation over the temperature range from 17°C to 37°C (one-way ANOVA, *P* < 0.0005, *n* = 10). However, there was no significant trend over the range 17–29°C (one-way ANOVA, *P* = 0.088, *n* = 10). Thus there is no significant change in mobility over the range where the lipid phase transition occurs (Fig. 5B). However, diffusion becomes significantly faster at 37°C (Tukey’s test with 95% simultaneous confidence intervals). The mean diffusion coefficient for TatA-GFP over all the other conditions tested was 0.13 ± 0.03 μm² s⁻¹ (n = 50), which is consistent to within experimental error with values previously obtained from FRAP (Mullineaux et al., 2006) and single-particle tracking (Leake et al., 2008).

**Video-rate TIRF microscopy of the helix1021-GFP model membrane protein**

We further investigated the distribution and mobility of helix1021-GFP using video-rate TIRF microscopy, a technique previously used to quantify the behaviour of TatA in the *E. coli* cytoplasmic membrane (Leake et al., 2008). When expression of helix1021-GFP is strongly induced, it appears at optical resolution to be evenly distributed in the membrane (as seen by confocal microscopy in Fig. 1C). However, we found that when expression is weakly induced, GFP fluorescence was concentrated in distinct mobile foci in the membrane (typically one to two observed per cell), with no measurable localization bias to specific regions of the membrane around the cell. The foci could be observed most clearly by high-speed TIRF microscopy (Fig. 6A and Supplementary Movie). At higher protein concentrations these distinct spots of fluorescence were no longer detectable and the cellular fluorescence intensity increased. We applied automated single particle tracking to the motions of the spots to generate particle displacement trajectories, to quantify the intensity of the foci as a function of time after the start of the bleach, and to estimate the full width at half maximum (FWMH) of the intensity profile of each particle. Each displacement trajectory was converted into a mean-squared displacement (MSD) versus time interval trace (Leake et al., 2008), and an average MSD trace (Fig. 6B) was compiled from all automatically detected trajectories having a duration of at least 1 s (Robson et al., 2013). Individual MSD traces indicated a broad range of lateral diffusion coefficients from 0.05 μm² s⁻¹ to 0.40 μm² s⁻¹ with a mean value of 0.2 ± 0.1 μm² s⁻¹. Intensity analysis on the individual distinct foci was performed as described previously (Leake et al., 2006; 2008; Reyes-Lamothe et al., 2010; Badarinraranayan et al., 2012). In brief, this consisted of automated foci detection followed by iterative two-dimensional Gaussian fitting to determine the foci intensity centroid to a precision of a few tens of nm (Llorente-Garcia et al., 2014) followed by local background intensity subtraction in the immediate vicinity of each focus based upon an unbiased average pixel intensity estimate pixels in a square region of interest of area ~ 800 × 800 nm centred on each focus, excluding pixels from the circular two-dimensional point spread function image region of each focus within that square. The intensity analysis indicated a range of initial unbleached intensity values, with an average of 16 200 ± 6100 counts (± SD) on our camera detector. The estimated signal-to-noise ratio, with regard to the integrated foci pixel intensity signal and the local background pixel noise, was typically lower by a factor of ~ 2 (an
example of a pixel intensity profile of a focus in a live cell is given in Supplementary Fig. S4) than had been measured from previous studies involving a similar imaging set-up but investigating CytdB-GFP (Lenn et al., 2008) or TatA-YFP (Leake et al., 2008). However, this level of signal-to-noise ratio was consistent with other previous in vivo single-molecule fluorescence imaging studies using E. coli investigating lower stoichiometry molecular complexes (Leake, 2014) used in DNA replication (Reyes-Lamothe et al., 2010), DNA remodelling (Badrinarayanan et al., 2012), oxidative phosphorylation (Llorente-Garcia et al., 2014) and purified fluorescent protein (Plank et al., 2009). Prior calibration experiments using immobilized GFP on the surface of the glass coverslip indicated a unitary brightness of single GFP molecule of 670 ± 150 counts (± SD). We measured the penetration depth of the TIRF evanescent field as 110 ± 10 nm (± SD) and estimated previously that protein complexes embedded in the cytoplasmic membrane of E. coli were typically ~ 90 nm in height from the coverslip surface using the same cell immobilization protocol (Leake et al., 2006). This indicates that the number of GFP molecules associated with the distinct foci of helix1021-GFP is in the range of 30–70 molecules. At the lowest levels of expression used for the single-particle tracking TIRF microscopy data, we estimate ~ 180–840 helix1021-GFP molecules per cell, based on one to two foci observed per cell from TIRF images that are likely to encompass ~ 1/6 of the total cytoplasmic membrane area of the cell (Leake et al., 2006), whose stoichiometry is in the range ~ 30–70 molecules per focus. Measuring the intensity profile of the distinct foci indicated FWHM in the range 300–400 nm, 50–150 nm larger than measured from the surface-immobilized GFP molecules measured using the same imaging set-up (Plank et al., 2009; Reyes-Lamothe et al., 2010; Badrinarayanan et al., 2012; Llorente-Garcia et al., 2014). Therefore we can estimate that the real diameter of the foci is in the range 50–150 nm. Comparison of E. coli cell strains with a variety of different OXPHOS proteins (Llorente-Garcia et al., 2014) or components of the MukBEF DNA remodelling system (Badrinarayanan et al., 2012) labelled either with the standard Clontech enhanced GFP (as we use here), or with the ‘definitively’ monomeric fluorescent protein mCherry, yielded no statistically significant differences between stoichiometry distributions, with evidence of a unimodal population from surface immobilization assays of GFP (Plank et al., 2009). This indicates that putative GFP dimerization is unlikely to be an issue in our study.

Similar video-rate TIRF experiments using E. coli cells treated with BODIPY FL-C12 indicated a reasonably uniform fluorescence in the cell membrane with some putative low-level heterogeneity in the intensity but no distinct foci even at low concentrations of the dye.

Discussion

Rather than examining the diffusion of proteins in artificial membranes divorced from the native biological context, our approach here was to examine the diffusion of a simplified model protein in a real biological membrane, the cytoplasmic membrane of living E. coli cells, with the full native biological context intact. We created a fluorescent probe by expressing within the cell a fusion construct of a single membrane-spanning α-helix with a GFP reporter domain predicted to be exposed on the cytoplasmic side of the membrane. The protein localizes to the cytoplasmic membrane, and the GFP tag allows the use of FRAP to measure the lateral diffusion of the protein in the membrane, and to visualize the probe using video-rate, high-contrast TIRF microscopy. For comparison we also monitored the diffusion of BODIPY FL-C12, a fluorescent lipid analogue with a 12-carbon fatty acid tail. In live E. coli cells this localizes preferentially to the plasma

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membrane, so we are able to monitor diffusion of a model protein and a model lipid in the same membrane. The model lipid shows the expected temperature behaviour; for cells grown at 37°C its diffusion coefficient changes by a factor of about 2 over a narrow temperature range between 27°C and 30°C. This suggests a phase transition in the lipid (Hazel and Williams, 1990; Sarcina et al., 2003) although the change in translational diffusion coefficient is small in comparison to that observed in simple lipid bilayers in vitro, where translational diffusion coefficients can change by orders of magnitude due to phase transition (Hazel and Williams, 1990).

Although the model membrane protein carries a relatively large GFP tag on the cytoplasmic side its diffusion is expected to be mainly controlled by the environment of the transmembrane domain, since the effective viscosity of the cell membrane is significantly greater than that of the cytoplasm. The diffusion coefficient of free GFP in the E. coli cytoplasm is typically 5–10 μm² s⁻¹ (Elowitz et al., 1999; Mullineaux et al., 2006; Nenninger et al., 2010), ~ 30 times faster than the diffusion coefficient we measured for the model membrane protein. In cells grown and measured at 37°C, the tagged protein diffuses roughly three times slower than the BODIPY label. The diffusion coefficient of a native membrane protein, GFP-tagged TatA, proved to be slower than the model protein under all the conditions that we tested. This could be explained by supramolecular interactions, including a tendency to homo-oligomerize (Leake et al., 2008).

We examined the effect on protein diffusion of treatments that alter the effective viscosity of the membrane, as judged from the mobility of the BODIPY lipid probe. As compared to the standard situation (E. coli cells grown and measured at 37°C) the BODIPY diffusion coefficient can be decreased by a factor of about 2 by cooling below 27°C, or increased by a factor of about 2 by growing cells at lower temperatures, presumably due to changes in the lipid composition of the membrane (Thompson, 1989). However, none of these treatments had any significant effect on the diffusion coefficient of the model membrane protein. The BODIPY diffusion coefficient could be perturbed to a similar extent in mutants deficient in specific lipid biosynthesis enzymes, but again without concomitant effects on protein diffusion (Table 1).

The use of high-speed TIRF microscopy allowed us to observe a very inhomogeneous distribution of the protein in the membrane plane. When the model protein is weakly induced, and therefore present only at low concentrations in the membrane, GFP fluorescence was concentrated in mobile spots in the membrane. At higher protein concentrations these distinct spots of fluorescence were no longer detectable. This behaviour indicates that the formation of the fluorescent spots is not due to the concentration-dependent aggregation frequently observed when membrane proteins are overexpressed (Ray et al., 2005). Rather, the protein has a tendency to segregate into domains of effective diameter of about 100 nm even at low concentrations. Each domain contains in the order of 50 helix1021-GFP proteins (Fig. 7). These domains are probably not observable at higher protein concentrations simply because they are too numerous to allow the spaces between them to be detected at optical resolution. We can estimate that the concentrated protein typically occupies less than 1% of a domain area (Fig. 7). Therefore it may well be undergoing faster micro-diffusion within each domain at timescales below our sampling time. The forces that hold these loose domains together are unclear and a topic for future study.

It is established that membrane proteins are surrounded by a shell of lipid molecules which is analogous to the solvent layer surrounding water-soluble proteins; several membrane proteins show greater affinity towards lipid in the liquid crystalline phase over that in the gel phase, consistent with packing considerations. For example, liquid crystalline phase lipid binding to Ca²⁺-ATPase is ~ 25 times more likely than for gel phase lipid binding (Lee, 2003). As a working model for the behaviour of our model protein, we suggest that the protein has the ability to reorganize the surrounding lipid to such an extent that its diffusion is unaffected by changes in the
effective viscosity of the bulk lipid in the membrane. A preference of the protein for a particular lipid phase could also account for its tendency to segregate out into what appear to be loosely organized mobile domains in the membrane. An open question is whether our model protein is influencing the segregation of the membrane, or whether it is simply partitioning into microdomains that are already defined by specific interactions of the native lipids and proteins.

Several recent studies have examined the behaviour of native fluorescently tagged proteins in bacterial membranes (Johnson et al., 2004; Ray et al., 2005; Leake et al., 2006; 2008; Lenn et al., 2008; López and Kolter, 2010), and have observed a tendency for the protein to concentrate in mobile domains. These domains were characterized in some detail in the case of the cytochrome bd complex in E. coli, which is found in mobile domains on average approximately 100 nm in diameter, containing around 75 cytochrome bd complexes, and differing with a mean lateral diffusion coefficient of about 0.05 μm² s⁻¹ (Lenn et al., 2008). Other OXPHOS complexes partition into separate domains which share similar characteristics (Llorente-Garcia et al., 2014). These results already suggested that the plasma membrane of E. coli is a highly compartmentalized fluid. Our results here indicate that lateral segregation of the membrane can occur even with a very simple model membrane protein, and suggest that this may have a significant effect on the physical behaviour of the protein in the membrane. The precise factors that cause lateral segregation in the membrane, and the physiological consequences, are subjects for future study. It is likely that the observed robustness in membrane protein diffusion confers functional advantages in cells that are exposed to fluctuations in temperature.

Experimental procedures

Cell strains and molecular biology

All experiments were performed using the E. coli strain DH5x [fhuA Δ(argF-lacZ)U169 phoA glnV44 φ80 Δ( lacZ)M15 gyrA96 recA1 relA1 endA1 thi-1 hsdR17]. The GFPmut3* (Cormack et al., 1996) labelled membrane probes were expressed from the arabinose-inducible pBAD24 vector (Guzman et al., 1995). TatA-GFP was expressed from pBAD24-TatA-GFP-BCs (Ray et al., 2005). For the helix1021-GFP construct the truncated version of sll1021 was amplified via PCR using genomic DNA from Synechocystis sp. PCC 6803 creating an EcoRI site at the 5′ end and a gfp overlap on the 3′ end. The fluorescent tag was amplified from pJD1 (Thomas et al., 2001) creating a sll1021 overlap at the 5′ end and a HindIII site at the 3′ end. To fuse the two PCR products an overlap extension PCR (Shevchuk et al., 2004) was performed using external primer and the end-product cloned into pBAD24 via EcoRI and HindIII and transformed into DH5α. Restriction enzymes were FastDigest® (Fermentas). PCR was with PfuUltra II Fusion HS DNA Polymerase (Stratagene). Ligations used the Quick Ligase Kit from NEB. For primer sequences see Supplementary Table S1.

The lipid biosynthesis deletion mutants were created using the λ Red and FLP recombinase system (Datsenko and Wanner, 2000). Kanamycin (ΔpsaA, Δcis, ΔpsaA, Δpsd) and chloramphenicol (Δppp) resistant cassettes were amplified via PCR from pKD13 or pKD3 respectively. They were inserted into the E. coli genome to replace corresponding genes. The correct insertion was confirmed by PCR and the stability of the deletions was verified several times during the period of the experiments.

Preparation of cells for confocal FRAP measurements

Bacterial cultures were grown aerobically overnight in Luria–Bertani (LB) medium (Sambrook and Russell, 2001) at 37°C under constant shaking (180 r.p.m.) and supplemented with ampicillin (50 μg ml⁻¹) where needed. For FRAP measurements, overnight cultures were diluted 1:100 and long non-septated cells were produced by adding cephalexin to a final concentration of 30 μg ml⁻¹ (Ishihara et al., 1983) and grown for up to 120 min at the different temperatures under constant shaking to mid-exponential phase (OD₆₀₀ = 0.3–0.4). Wild-type cultures were grown to mid-exponential phase with 1 μM BODIPY FL-C₁₂ (Invitrogen) and washed several times in fresh LB. For induction of the GFP-tagged membrane proteins, L-arabinose was added to the diluted culture [500 μM for TatA-GFP and 133 mM (2%) for helix1021-GFP] and cells were grown at the appropriate temperature to mid-exponential phase. For transfer to the microscope drops of cell culture were spotted onto LB-agar plates and allowed to adsorb onto the agar surface. Small blocks of agar holding adsorbed cells were cut from the plate and placed in a laboratory-built sample holder connected to a temperature-controlled circulating water-bath, covered with a coverslip and placed under the microscope (Mullineaux et al., 2006). Sample preparation and temperature equilibration took about 20 min in total.

Confocal imaging and FRAP data analysis

FRAP measurements and data analysis were carried out according to Mullineaux et al. (2006) using a Nikon PCM2000 laser-scanning confocal microscope equipped with an Argon laser run at 100 mW. A wide (50 μm) confocal pinhole was used to reduce the z-axis resolution and thus collect information from the full depth of the cell. Excitation at 488 nm was reduced by a factor of 32 via neutral-density filters for imaging, ensuring that repeat imaging did not measurably bleach the sample. The filters were manually lifted (for about 1 s) for the high laser power bleach. For the bleach across the short axis of the elongated cell the x-scanning mode was selected. Pre- and post-bleach scans were recorded separately at 1.64 s intervals over a 512 by 512 pixel area using the xy mode for scanning. The measurement area was either 29 μm or 58 μm square. Fluorescence emission was detected between 500 and 527 nm. Image analysis used Image-Pro 6.2 (Media Cybernetics). Pre- and post-bleach image sequences were combined and a one-dimensional fluores-
cience profile along the long axis of the cell was extracted as a line-profile summing data widthways across the cell for the entire sequence. In SigmaPlot 10.0 (Jandel Scientific) the diffusion coefficient D was obtained by fitting to a one-dimensional diffusion equation (Mullineaux et al., 1997; 2006). SigmaPlot was also used to perform Student’s t-tests and for fitting of sigmoidal curves to plots of D versus temperature. Further statistical analyses were made using MINITAB.14 (Minitab). We conducted one-way ANOVAs followed by Tukey’s tests to assess significant differences, considering a P value of < 0.005 as significant. High-resolution fluorescence and DIC images were recorded using a Leica TCS SP5 laser-scanning confocal microscope with excitation at 488 nm and emission at 500–520 nm. A 6× line-average was used, and the confocal pinhole was set to give a z-axis resolution of about 0.8 μm.

Preparation of cells for TIRF microscopy

Cells were sub-cultured into M63-glucose medium (Atlas, 1993) with 1 mM arabinose for induction of helix1021-GFP expression, then grown with shaking at 30°C for 3 h. The cell suspension was injected through a poly-L-lysine-coated tunnel slide (Leake et al., 2006) and left to incubate for 20 min. Excess medium was flushed through the tunnel slide to remove unbound cells.

TIRF microscopy and single particle tracking

A home-built inverted TIRF microscope with 473 nm excitation wavelength was used with specifications as described (Leake et al., 2006; 2008; Lenn et al., 2008). Fluorescence emission was imaged at 50 nm/pixel in frame-transfer mode at 25 Hz by a 512 × 512-pixel, cooled, back-thinned electron-multiplying charge-coupled device camera (iXon+ DV885-BI, Andor Technology). Images were sampled for ~10 s at room temperature (20°C). Fluorescent particles were automatically detected and tracked on each consecutive image frame using automated custom-written image-analysis software and a mean-squared displacement (MSD) calculated for each particle trajectory, with an estimate for lateral diffusion coefficient generated using a linear fit to the MSD versus time interval relation (Leake et al., 2008).

Cell fractionation and SDS-PAGE with Western blotting

Spheroplasts were produced according to Randall and Hardy (1986) by lysing the cell wall with lysozyme and stabilizing spheroplasts with MgSO4. They were harvested by gentle centrifugation and the supernatant was centrifuged a second time to ensure removal of all spheroplasts. After an additional wash step, spheroplasts were fractioned into a membrane and soluble cytoplasmic fractions according to Thomas et al. (2001). Spheroplasts were burst by sonication and the membranes were collected by ultracentrifugation (250 000 g for 30 min). The supernatant corresponded to the cytoplasmic fraction. The membrane fraction was washed once in buffer and, after a second centrifugation step, resuspended in buffer. The proteins of the membrane fraction were solubilized overnight by adding a small droplet of Triton X-100 to the buffer. Proteins were separated by 15% SDS-PAGE, semi-dry electrophoresis onto Hybond-R polyvinylidene difluoride (PVDF) membrane (GE Healthcare) and probed with antibodies against GFP (Invitrogen). A horseradish peroxidase-conjugated anti-mouse IgG secondary antibody and ECL enhanced chemiluminescence detection kit (GE Healthcare) permitted visualization of GFP.

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References


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**Supporting information**

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