

# Dynamic molecular confinement in the plasma membrane by microdomains and the cytoskeleton meshwork

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It is by now widely recognized that cell membranes show complex patterns of lateral organization. Two mechanisms involving either a lipid-dependent (microdomain model) or cytoskeleton-based (meshwork model) process are thought to be responsible for these plasma membrane organizations. In the present study, fluorescence correlation spectroscopy measurements on various spatial scales were performed in order to directly identify and characterize these two processes in live cells with a high temporal resolution, without any loss of spatial information. Putative raft markers were found to be dynamically compartmented within tens of milliseconds into small microdomains ( $\varnothing < 120$  nm) that are sensitive to the cholesterol and sphingomyelin levels, whereas actin-based cytoskeleton barriers are responsible for the confinement of the transferrin receptor protein. A free-like diffusion was observed when both the lipid-dependent and cytoskeleton-based organizations were disrupted, which suggests that these are two main compartmentalizing forces at work in the plasma membrane.

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## Introduction

Although it is generally assumed that lateral heterogeneities exist in cell membranes (Edidin, 2003; Simons and Vaz, 2004), the factors responsible for the local organization of

the plasma membrane are poorly understood. Various mechanisms have been suggested to explain the significant discrepancies observed in the diffusion rates of lipids and proteins between artificial membranes and plasma membranes. For instance, membrane protein diffusion differs from pure Brownian diffusion because of the transient interactions occurring with large multimeric complexes (Marguet *et al.*, 1999), the obstacles resulting from direct and indirect molecular immobilization (Sako and Kusumi, 1994; Fujiwara *et al.*, 2002), and the corrals formed on various submicrometric scales (Yechiel and Edidin, 1987). There is still some controversy about the ability of lipids to form domains, which are also known as lipid rafts, in plasma membranes (Simons and Ikonen, 1997; Brown and London, 1998; Edidin, 2003; Munro, 2003; Simons and Vaz, 2004; van Meer, 2005; Marguet *et al.*, 2006). In these lipid domains, cholesterol is thought to be tightly packed with long saturated fatty acid chains of specific glycosphingolipids, thus favoring lipid phase separation processes. It has been suggested that these domains may be insoluble in nonionic detergents (Brown and London, 1998; Drevot *et al.*, 2002). In addition to studies characterizing biochemical rafts, morphological approaches have been used to prove the existence of these rafts in cell membranes, but the conclusions drawn have been rather controversial, since the results depended strongly on the methods used. When the lateral motion of lipid probes was analyzed using single dye tracing methods, the domain size was found to range from 0.2 to 2  $\mu\text{m}$  (Schutz *et al.*, 2000). These data are in the same range as those obtained on the barrier-free paths of membrane proteins, but the latter depend mainly on how the cytoskeleton is organized (Sako and Kusumi, 1994). Other studies using biophysical approaches on live cells have strongly supported the idea that rafts are small diffusive entities (Pralle *et al.*, 2000; Sharma *et al.*, 2004). Fluorescence resonance energy transfer methods (FRET) have been used to probe the molecular aggregates present in lipid rafts. Experiments of this kind using different lipid-modified proteins expressed in the apical membrane of MDCK cells have led to divergent conclusions (Kenworthy and Edidin, 1998; Zacharias *et al.*, 2002; Hess *et al.*, 2005). FRAP measurements on live cells have provided evidence that the anchoring mode determines the protein-raft interaction dynamics (Shvartsman *et al.*, 2003). A direct visualization of liquid-ordered domains was recently obtained using fluorescent cell staining procedures, although information is still lacking about the dynamic aspects of these structures (Gaus *et al.*, 2003). In fact, these methods span different spatiotemporal scales, and the data obtained about the features of rafts, and even their existence in live cells, may therefore not always seem to be in agreement.

Improving our knowledge of membrane organization requires developing new methods in which the temporal resolution is improved without any loss of spatial information. For instance, FRAP measurements performed on different

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spatial scales have made it possible to identify micrometer-scale domains present in the plasma membrane of fibroblasts (Yechiel and Edidin, 1987). By applying a similar strategy using fluorescence correlation spectroscopy (FCS) methods, we recently managed to distinguish between different submicron confinement models. In the latter study, we described a novel method of performing FCS on various observation scales (Wawrezynieck *et al*, 2004) and developed a simple analytical method of investigating the complexity of the cell membrane organization and the dynamic processes involved (Wawrezynieck *et al*, 2005). This novel method could be used to identify and discriminate between variously organized submicroscopic domains preventing the free lateral diffusion of membrane components (Saxton, 2005). The *FCS Diffusion Law* was defined in the latter study as the plot of the diffusion time (i.e. the average time a fluorescent molecule stays within the illuminated area) versus the observation area. Based on extensive confined diffusion modeling studies, it was established that the  $y$ -intercept  $t_0$  in the graph of the FCS diffusion laws is positive in the case of confining in microdomains (dynamic partitioning into microdomains) but negative in that of trapping in a meshwork (actin-based cytoskeleton corrals).

Here, it was proposed to investigate the dynamics of membrane organization in live cells. The diffusion behavior of various membrane components was described and it was established whether molecular confinement depends on the specific cholesterol/sphingomyelin membrane content or on the presence of actin-based cytoskeleton barriers.

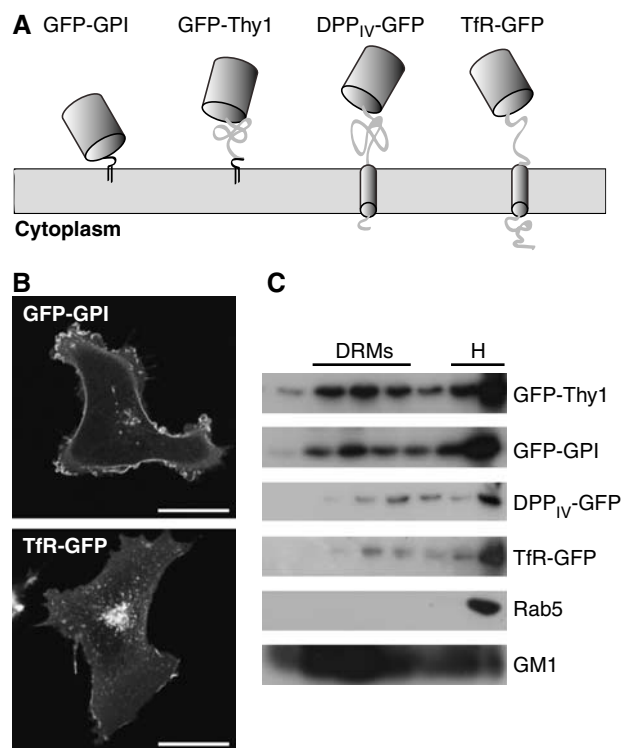
## Results

### Molecules used to establish the presence of domains in plasma membranes

To establish the presence of domains in the plasma membrane, the diffusion behavior of four different groups of membrane components was investigated: (1) a glycerophospholipid group with two fluorescent lipid analogs, BODIPY- $C_5$ -phosphatidylcholine (FL-PC) and BODIPY-dihexadecanoyl-phospho-ethanolamine (FL-PE), (2) a sphingolipid group including the BODIPY- $C_5$ -sphingomyelin (FL-SM) and BODIPY- $C_5$ -ganglioside- $G_{M1}$  (FL- $G_{M1}$ ) analogs, (3) a glycosylphosphatidylinositol (GPI)-anchored protein group in which the GFP was either tagged onto the GPI-anchoring signal originating from the decay accelerating factor (GFP-GPI) or inserted just after the signal peptide of Thy1 (GFP-Thy1), and (4) a GFP-tagged transmembrane protein group including the dipeptidyl peptidase IV (DPP<sub>IV</sub>-GFP) and the transferrin receptor (TfR-GFP) (Figure 1A).

To examine the mobility of lipid analogs, COS-7 cells were labeled using the lipid exchange method (Martin and Pagano, 1994). Laser scanning confocal microscopy showed that the probes were uniformly distributed in the plasma membrane. A vesicular staining pattern consisting of small intracellular dots was also observed (Figure 2A–C). All the GFP-tagged proteins were efficiently addressed to the plasma membrane when transiently expressed in COS-7 cells (Figure 1B and data not shown). Vesicular staining was also detected in the cytosol, especially in the perinuclear area.

Based on previous studies, these molecules are known to be either present in high levels or depleted from detergent resistant membranes (DRMs). This was confirmed by Brij 98

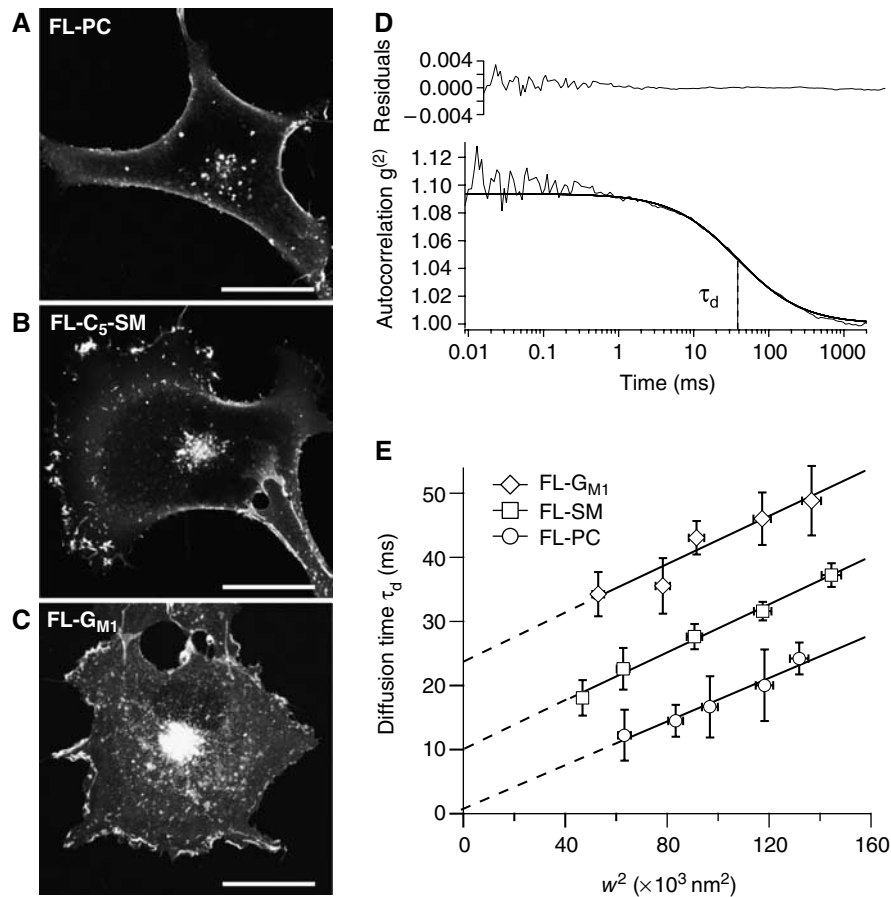


**Figure 1** GFP-tagged proteins used in this study. (A) Membrane topology of the GFP-tagged proteins. (B) Confocal images of COS-7 cells expressing the GFP-GPI and TfR-GFP proteins. Scale bar is 20  $\mu$ m. (C) Brij 98 solubilized PNS from transfected COS-7 cells was fractionated on the sucrose gradient and blotted with anti-GFP, anti-Rab 5 antibodies or cholera toxin B.

detergent-insolubility experiments performed at physiological temperatures (Drevot *et al*, 2002). Although the BODIPY fluorophore can affect the properties of lipids, yielding analogs with quite different characteristics from those of their natural counterparts (Kuerschner *et al*, 2005), both FL- $G_{M1}$  and FL-SM were readily detected in DRMs, where they accounted for 11 and 17% of the total fluorescence, respectively. This association is not negligible, since under the same experimental conditions, about 20% of the GFP-Thy1 or GFP-GPI molecules were detected in DRMs, whereas those anchored by a classical transmembrane peptide were barely detectable (Figure 1C).

### The different groups of membrane components follow distinct FCS diffusion laws

As described previously, the *FCS diffusion law* giving the diffusion time  $\tau_d$  (i.e. the average time a fluorescent molecule stays within the illuminated area) versus the beam area, where  $w$  denotes the beam radius, should provide a suitable analytical method for determining the diffusion behavior of a molecule in the membrane (Saxton, 2005; Wawrezynieck *et al*, 2005). This approach is based on the fact that if molecules diffuse freely, one can expect to obtain a linear function between  $\tau_d$  and  $w^2$ : the curve  $\tau_d = f(w^2)$  will intercept the time origin and its slope will correspond to the diffusion coefficient  $D$  in this environment. In most situations where diffusion is hindered, the FCS diffusion law will no longer fit this scheme. By sizing the laser beam filling the microscope objective, we varied  $w$  experimentally from 200



**Figure 2** Diffusion behavior of lipid analogs. Confocal images of COS-7 cells after labeling with lipid analogs: (A) FL-PC, (B) FL-SM and (C) FL-G<sub>M1</sub>. Scale bar is 20  $\mu$ m. (D) ACF obtained with FL-G<sub>M1</sub> lipids. The curve was satisfactorily fitted with a 1-species free diffusion model (see Supplementary data). The diffusion time  $\tau_d$  was given by the lag time at half maximum. (E) Diffusion behavior of lipid analogs. Confinement times were determined from the position at which the diffusion curves and the time-axis intersected. Error bars in  $x$  and  $y$  give standard deviations (s.d.) of the means.

to 400 nm; the lower limit was imposed by the diffraction law, whereas the upper one was based on a compromise between the spatial resolution and FCS constraints.

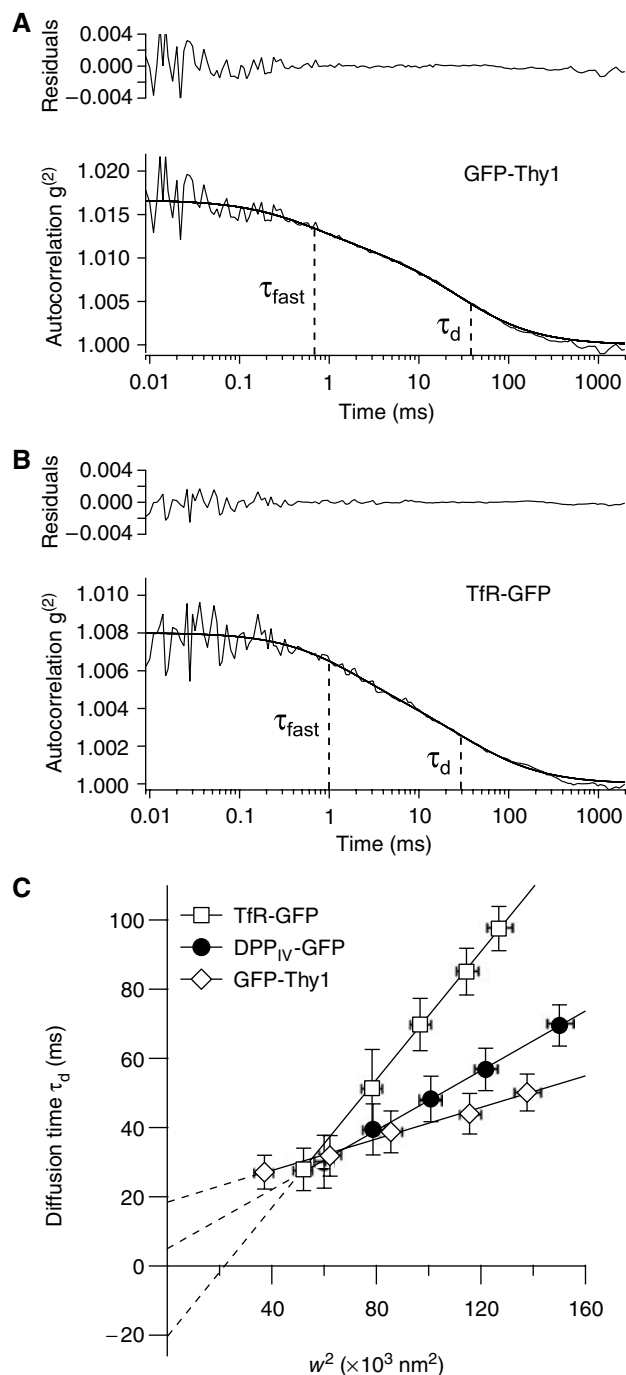
We first investigated the lipid dynamics using trace mole fractions of probes relative to the total lipids in order to minimize any membrane perturbations. Based on experimental autocorrelation functions (ACF), the data were efficiently fitted using a single component, giving a single decay time (see Supplementary data and Figure 2D). In this case,  $\tau_d$  was obtained by measuring the lag time at half the maximum ACF value. With all the lipid probes used,  $\tau_d$  increased linearly with  $w^2$  but the intersections with the time axis differed (Figure 2E). In the case of the glycerophospholipid analogs FL-PC and FL-PE, the FCS diffusion laws intercepted the time axis at the origin at almost null  $t_0$  values ( $-0.4 \pm 0.8$  and  $0.2 \pm 0.7$  ms, respectively). By contrast,  $t_0$  was strictly positive in the case of the sphingolipid analogs FL-G<sub>M1</sub>, FL-C<sub>5</sub>-SM and FL-C<sub>12</sub>-SM, where values of  $24.3 \pm 1.9$ ,  $10.2 \pm 0.6$  and  $9.7 \pm 0.5$  ms were obtained, respectively.

With all the chimeric proteins used, two decay times were clearly identified from the ACFs (Figure 3A–B and data not shown): a short time  $\tau_{\text{fast}}$  and a longer one  $\tau_d$ . We checked that the slow decay time corresponded to the diffusion time of plasma membrane-bound molecules, whereas the fast one corresponded to that of the intracellular pool. When detected

from outside by an Alexa488-conjugated anti-Thy1 Fab fragment, the lateral mobility of Thy1 showed a similar single long relaxation time to the longest one observed in the case of GFP-Thy1 (with a spot radius of  $\sim 240$  nm, these times were  $31.8 \pm 5.8$  and  $27.4 \pm 2.9$  ms, respectively).

The FCS diffusion laws were then analyzed (Figure 3C). In the case of the two GPI-anchored proteins, GFP-Thy1 and GFP-GPI,  $t_0$  was found to be significantly positive ( $18.4 \pm 1.7$  and  $12.1 \pm 2.5$  ms, respectively), whereas the reverse was observed with the TfR-GFP, which gave a  $t_0$  value of  $-20.4 \pm 1.9$  ms. Interestingly, the DPP<sub>IV</sub>-GFP diffusion behavior was in between that observed with the GPI-anchored proteins and the TfR-GFP. The reason for this small positive intercept ( $4.8 \pm 1.7$  ms) will be discussed below.

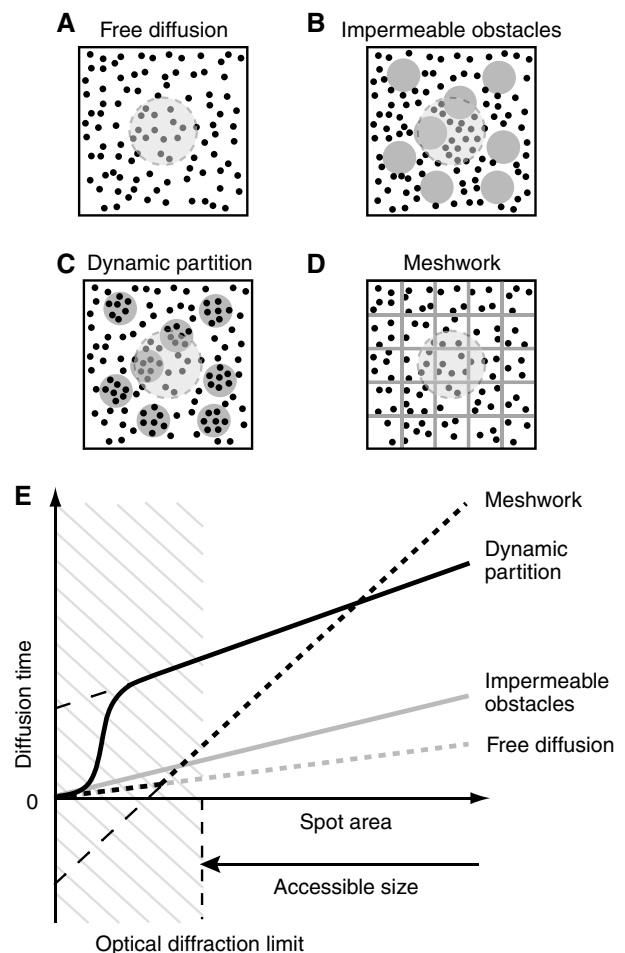
In order to check whether the diffusion behavior reflected in the FCS corresponded to a significant fraction of the molecules tested, confocal FRAP measurements (Kenworthy *et al*, 2004) were performed. The mobile fraction  $M_f$  was assessed in terms of the percentage of the fluorescent molecules recovered in the bleached area during the FRAP experiment. All the lipid analogs were nearly 100% mobile, and  $M_f > 95\%$  with FL-PC, FL-SM and FL-G<sub>M1</sub>. In the case of the GFP-tagged proteins, the  $M_f$  value ranged between 77 and 92% (77, 85, 90 and 92% with TfR-GFP, DPP<sub>IV</sub>-GFP, GFP-GPI and GFP-Thy1, respectively). The FCS diffusion laws given



**Figure 3** Diffusion behavior of GFP-tagged proteins. ACFs obtained with GFP-Thy1 (A) and TfR-GFP (B). In both cases, two decay times were clearly identified with the various probes used: a short time  $\tau_{fast}$  and a longer one  $\tau_d$ . (C) FCS diffusion laws in the case of TfR-GFP, DPP<sub>IV</sub>-GFP and GFP-Thy1. Error bars on the x and y axes give the s.d.s of the means.

above therefore accurately describe the behavior of a very large fraction of the lipid analogs and GFP-tagged proteins.

A broad range of  $t_0$  values was obtained with the various molecules used to explore the occurrence of domains in the plasma membrane. Based on this parameter, three subsets of molecules were identified: the glycerophospholipid analogs (null  $t_0$ ), the sphingolipid and GPI-anchored protein groups (positive  $t_0$ ) and the transmembrane TfR-GFP (negative  $t_0$ ).



**Figure 4** Simulated FCS diffusion laws corresponding to different membrane models. (A–D) Diffusion models for membrane organization. (A) In the free diffusion model, fluorescent molecules (black dots) show pure Brownian motion and fluoresce under the laser excitation spot (large gray circle). (B) In the presence of impermeable obstacles (gray spots), the diffusion is restricted to the free space. (C) When the domains are permeable (as the result of dynamic partition processes), the molecules can diffuse into and out of the microdomains and be transiently trapped (gray spots). (D) In a meshwork model, multiple adjacent domains are separated by barriers (gray lines) preventing the diffusion of the molecules. (E) Characteristic diffusion laws obtained for the different models. The diffusion time  $\tau_d$  was analyzed as a function of the squared radius  $w^2$ . Diffusion laws applying to the free diffusion model (dotted gray curve), the diffusion process in the presence of impermeable obstacles (gray curve), the dynamic partition model (dotted black curve) and the meshwork model (black curve). Diffusion laws cannot be determined experimentally below the diffraction limit. For detailed analyses, see the Supplementary data and Wawrezinieck *et al* (2004, 2005).

In view of the above data and the results of our previous numerical simulations (Wawrezinieck *et al*, 2005), where the change in the sign of the  $y$ -intercept  $t_0$  was correlated with different modes of membrane compartmentalization, we hypothesized that simple confinement models would explain the deviations from free diffusion (Figure 4; see also the Supplementary data). A positive  $t_0$  was linked, for example, to the dynamic partition model, which may provide a basis for explaining the diffusion behavior of the sphingolipid and GPI-anchored protein groups, whereas with a negative  $t_0$ , TfR molecules may be subjected to membrane confinement due

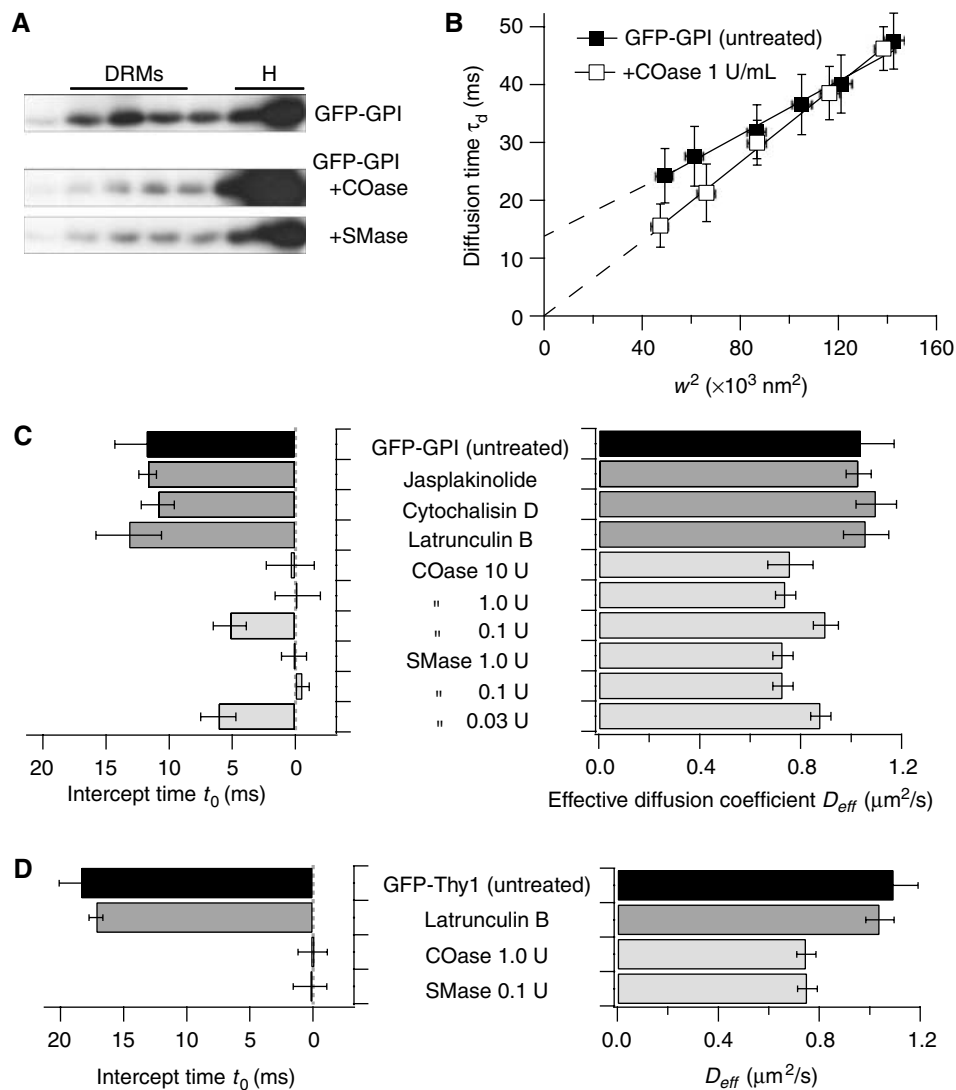
to meshwork effects. Lastly, because of the null  $t_0$ , the glycerophospholipid analogs may neither be confined within discrete domains nor be hindered by a meshwork.

**A lipid-dependent microdomain organization confines sphingolipid analogs and GPI-anchored proteins within the plasma membrane of resting cells**

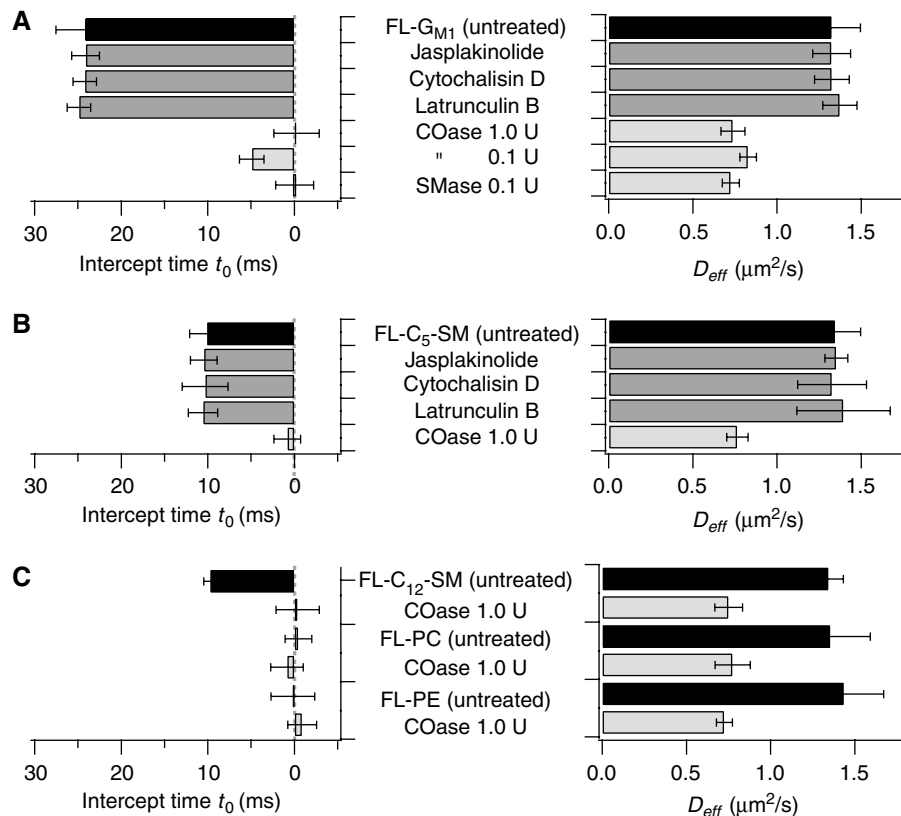
Among the mechanisms possibly responsible for confinement in discrete microdomains, several lines of evidence combine to support the idea that cholesterol and sphingomyelin are involved in the control and the organization of local heterogeneities: the differential packing characteristics of glycerophospholipids and sphingolipids with cholesterol promote the formation of a liquid-ordered phase or a similar state, which coexists in the membrane plane with a liquid-disordered phase (Brown and London, 1998; Rietveld and Simons, 1998). To test whether this hypothesis holds in the case of

sphingolipid analogs and GPI-anchored proteins, cells were incubated in the presence of exogenous cholesterol oxidase (COase) or sphingomyelinase (SMase) to decrease the cholesterol and sphingomyelin membrane contents, respectively.

With 1 U/ml of COase,  $23.3 \pm 2.2\%$  of the total cell cholesterol was converted into cholestenone. The ability of GPI-anchored proteins and sphingolipid analogs to float to the buoyant fractions of a sucrose gradient was subsequently found to be severely impaired (Figure 5A), and the FL-SM and FL-G<sub>M1</sub> present in DRMs, as measured by spectrofluorimetry, decreased by 58 and 63%, respectively. This decrease in the cholesterol levels also induced significant changes in the behavior of GFP-GPI, since the curve of the corresponding FCS diffusion law then intercepted the time origin ( $0.2 \pm 0.8$  ms) (Figure 5B). Similar results were obtained with GFP-Thy1 and the sphingolipid analogs (Figures 5C, D



**Figure 5** The confinement of GFP-tagged GPI-anchored proteins in microdomains is lipid-dependent. (A) Biochemical determination of GFP-GPI localization in DRMs after COase or SMase treatment. (B) FCS diffusion laws governing GFP-GPI proteins after cell treatment with COase. Error bars on the x and y axes are the s.d.s of the means. (C) Time intercept  $t_0$  and effective diffusion coefficient  $D_{eff}$  obtained with GFP-GPI in untreated COS-7 cells (black bars), after cytoskeleton drug treatments (dark gray bars) or lipid modifications (light gray bars). (D) Idem in the case of GFP-Thy1 protein.



**Figure 6** The confinement of lipid probes in the plasma membrane of COS-7 cells is lipid-dependent. Time intercept  $t_0$  and effective diffusion coefficient  $D_{eff}$  obtained with FL-G<sub>M1</sub> (A), FL-C<sub>5</sub>-SM (B) and the other lipid analogs used in this study (C) in untreated COS-7 cells (black bars), after applying cytoskeleton drug treatment (dark gray bars) or lipid modifications (light gray bars). Error bars give the s.d.s of the means.

and 6, respectively). Importantly, a 10-fold lower enzyme concentration, which converted  $<10\%$  of the total cholesterol into cholestenone ( $8.2 \pm 4.4\%$ ), led to an intermediate value between those obtained with cells treated and not with 1 U/ml of COase, as illustrated here in the case of GFP-GPI and FL-G<sub>M1</sub>, (Figures 5C and 6A, respectively). The effects of the larger changes in the cholesterol content ( $56.9 \pm 4.7\%$ ) induced by 10 U/ml of COase did not differ from those observed with 1 U/ml. All in all, these results suggest that the microdomain organization generated by dynamic partitioning processes no longer existed in these cell membranes.

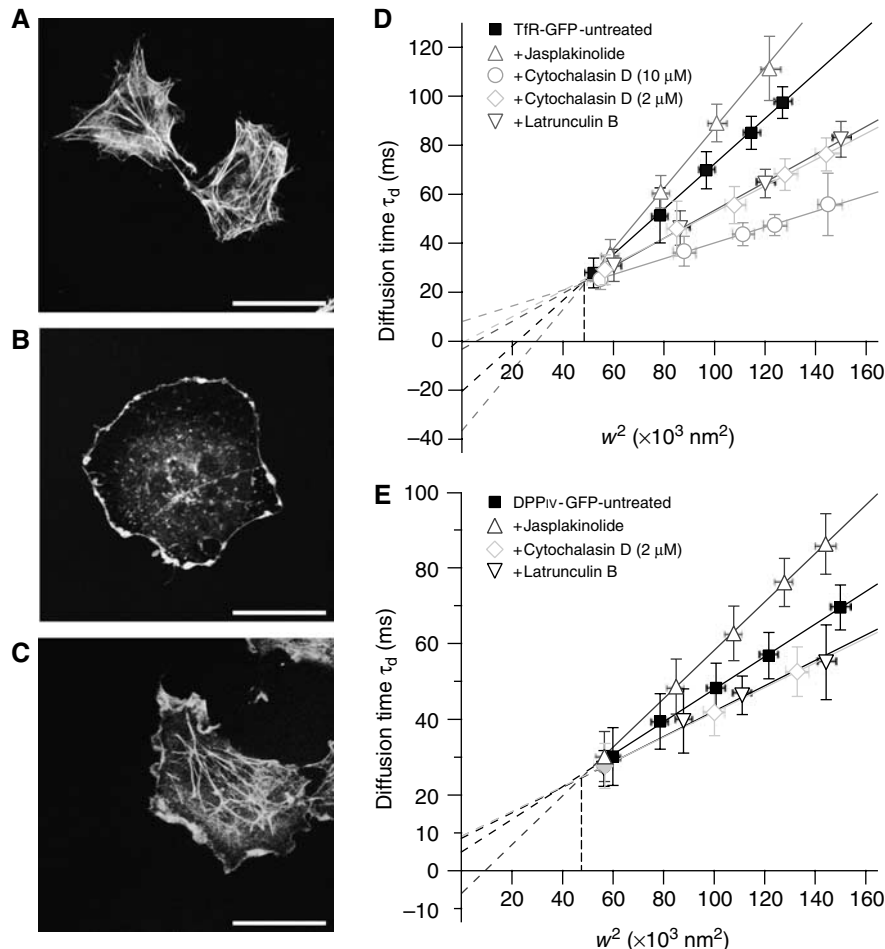
We then hydrolyzed SM into ceramide by adding exogenous SMase to COS-7 cells. With 0.1 U/ml of SMase,  $68 \pm 4\%$  of the total SM was hydrolyzed. At the same time, we noted that the presence of GFP-GPI within DRMs was also severely impaired after enzymatic treatment (Figure 5A). More importantly, the curves of all the FCS diffusion laws governing GFP-GPI, GFP-Thy1 and FL-G<sub>M1</sub> intercepted the time origin (Figures 5C–D and 6A). Once again, we observed that the SM membrane concentration was critical, since titration experiments showed that the confinement of GFP-GPI disappeared in a dose-dependent manner (Figure 5C). Lower concentrations ( $0.03$  U/ml) converting  $54 \pm 4\%$  of the total SM led to an intermediate value between those obtained with cells treated or not with 0.1 U/ml of SMase, as illustrated in the case of GFP-GPI and FL-G<sub>M1</sub>. The effects of higher SMase concentrations (1 U/ml) did not differ from those observed with 0.1 U/ml.

It was remarkable that the lipid-mediated microdomain organization was entirely abolished when only  $\sim 20\%$  of the total cholesterol was converted into cholestenone by COase. This could be due to a combined effect of cholesterol hydrolysis and cholestenone production, since an antagonizing action of the cholestenone to ordered domains has been reported (Xu and London, 2000). A similar scenario might occur upon SMase treatment as well. Ceramide has recently been shown to displace cholesterol from sphingomyelin-cholesterol domains as well as to induce strong alterations of the lateral organization in model membranes (Megha and London, 2004).

After COase or SMase treatment, a similar decrease in  $D_{eff}$  occurred in both the GPI-anchored proteins and sphingolipid analogs ( $\sim 0.75 \mu\text{m}^2/\text{s}$ ) (Figures 5 and 6). The fact that this was also the case with the glycerophospholipid analogs although the  $t_0$  value was still null (Figure 6C) suggested that an overall change in the plasma membrane viscosity had also occurred due to the enzymatic change in the lipid membrane composition. All in all, these results strongly support the idea that in the absence of molecular crosslinking processes, sphingolipid analogs and GPI-anchored proteins are confined by a cholesterol and sphingomyelin-dependent process of microdomain formation.

#### **A cytoskeleton-mediated meshwork hinders the lateral mobility of TfR-GFP**

The TfR-GFP diffusion law curve, which showed a negative  $t_0$ , suggested that TfR molecules diffused in a meshwork



**Figure 7** The GFP-tagged transmembrane protein diffusion laws are actin-dependent. Confocal images of rhodamine phalloidin labeling of F-actin in COS-7: (A) untreated; (B) 1 μM latrunculin B; (C) 0.4 μM jasplakinolide. Diffusion laws governing TfR-GFP (D) and DPP<sub>IV</sub>-GFP (E) after applying cytoskeleton drug treatment. Error bars on the x and y axes are the s.d.s of the means.

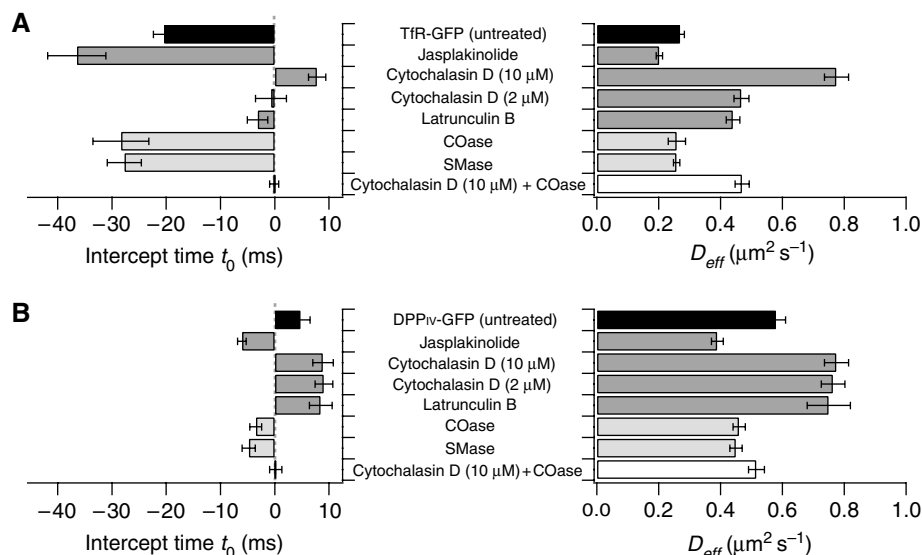
context (Figure 4D and E). To investigate the possible involvement of the cytoskeleton, COS-7 cells were treated with drugs known to alter the actin-based cytoskeleton, such as latrunculin-B (an actin polymerization inhibitor), cytochalasin D (an actin depolymerization inducer) or jasplakinolide (an actin filament stabilizer). Suitable experimental conditions were first setup to ensure that these drugs changed the pattern of cytoskeleton organization without significantly affecting the cell morphology (Figure 7A–C).

The lateral mobility of TfR-GFP was then assessed (Figure 7D). Under latrunculin B (1 μM) or cytochalasin D (2 μM) treatment, the FCS diffusion law curve intercepted the y-axis near the time origin ( $-3.2 \pm 1.8$  and  $0.7 \pm 1.8$  ms, respectively). These findings indicate that the physical barriers confining TfR-GFP were lowered. At the same time, the  $D_{\text{eff}}$  value increased from  $0.27 \pm 0.01$  to  $0.44 \pm 0.02$  μm $^2$ /s, which suggests that the long-range diffusion was less strongly impeded by the barriers between adjacent domains. Conversely, treatment with jasplakinolide (0.4 μM) decreased both the  $D_{\text{eff}}$  and  $t_0$  values to  $0.20 \pm 0.01$  μm $^2$ /s and  $-36.5 \pm 5.4$  ms, respectively. This decrease suggests that the barriers confining TfR-GFP were reinforced by jasplakinolide treatment.

By contrast, it is noteworthy that neither the actin-based skeleton alterations induced by latrunculin B nor its stabilization by jasplakinolide had any effect on the molecular confinement observed with the GPI-anchored proteins and the sphingolipid analogs (Figures 5 and 6). Neither the  $t_0$  nor  $D_{\text{eff}}$  values were affected under the experimental conditions that drastically altered the diffusion of TfR-GFP.

**Meshwork and discrete domains contribute concomitantly to the dynamic compartmentalization of transmembrane proteins in the cell membrane**

The shape of the diffusion law curve obtained for DPP<sub>IV</sub>-GFP suggested that confinement by discrete domains may have occurred since, as observed in the case of GPI-anchored proteins,  $t_0$  was positive. However, a similar study to that carried out on TfR-GFP showed that the FCS diffusion law was actin-dependent (Figure 7E). Treatment with latrunculin B or cytochalasin D led to increases in both  $t_0$  and  $D_{\text{eff}}$ , whereas jasplakinolide treatment decreased both  $t_0$  and  $D_{\text{eff}}$ , and even led to a negative  $t_0$  value. More intriguingly, although TfR-GFP and DPP<sub>IV</sub>-GFP were obviously sensitive to cytoskeleton-based compartmentalization, the possibility that discrete domain confinement may have occurred was



**Figure 8** Both actin meshwork and lipid domains contribute to transmembrane protein compartmentalization. Time intercept  $t_0$  and effective diffusion coefficient  $D_{eff}$  obtained with TfR-GFP (A) and DPP<sub>IV</sub>-GFP (B) in untreated COS-7 cells (black bars), after applying cytoskeleton drug treatment (dark gray bars), after lipid modifications (light gray bars) or combined treatment (open bars). COase and SMase concentrations were 1 and 0.1 U/ml, respectively. Error bars are the s.d.s of the means.

suggested by the fact that high drug concentrations (cytochalasin D 10  $\mu$ M) disrupted the actin network organization and gave a  $t_0$  of  $7.8 \pm 1.6$  and  $8.9 \pm 1.9$  ms, respectively. In addition, the lipid modifications induced by COase or SMase systematically decreased the respective  $t_0$  values (Figure 8). These two findings suggest that both meshwork and isolated domains contributed simultaneously to the DPP<sub>IV</sub>-GFP and TfR-GFP diffusion process. To test this hypothesis, we then established the corresponding FCS diffusion laws under experimental conditions where actin-based cytoskeleton barriers and cholesterol were simultaneously modified (Figure 8). Significantly,  $\tau_d$  for DPP<sub>IV</sub>-GFP and TfR-GFP was found to be linear in  $w^2$ . Thus, when both confinement forces were relaxed, these proteins recovered free-like diffusion behavior.

Our study also indicated that the microdomain and the meshwork confinement may mask each other in the FCS law analysis. Indeed, we could interpret the differences in the diffusion modes between TfR-GFP and DPP<sub>IV</sub>-GFP in terms of the differential strength of the actin-based and lipid-dependent confinement processes. COase and SMase treatments both abolished the lipid-dependent confinement process but unraveled the cytoskeleton-dependent one, which was in turn abolished by cytochalasin D or latrunculin B (Figure 8). With these TfR-GFP and DPP<sub>IV</sub>-GFP proteins, COase treatment decreased the  $t_0$  values to  $-28.4 \pm 5.2$  and  $-3.5 \pm 1.1$  ms, respectively. It was therefore deduced that the confinement of the cytoskeleton was about eight times stronger with TfR-GFP than DPP<sub>IV</sub>-GFP. On the opposite, upon impairing the actin network, it was observed that both proteins underwent an almost identical weak lipid-dependent microdomain confinement process ( $7.8 \pm 1.6$  and  $8.9 \pm 1.9$  ms for TfR-GFP and DPP<sub>IV</sub>-GFP, respectively). It seems likely that the effects of cytoskeleton and lipid confinement may have been additive, but this hypothesis remains to be confirmed in further studies. The actin-based confinement process was stronger than the lipid-dependent one in the case of TfR-GFP, whereas the reverse was true in that of DPP<sub>IV</sub>-GFP.

## Discussion

The present study describing the diffusion behavior of diverse membrane components provides insights on the role of lipids in compartmentalizing the cell membrane. First, based on FCS data obtained on various spatial scales, we established that the sphingolipid analogs and GPI-anchored proteins undergo transient confinements in isolated microdomains. Secondly, the molecular origin of these confinement processes was found to be cholesterol- and sphingomyelin-dependent. Thirdly, their compartmentalization in isolated domains is independent of the actin-based cytoskeleton organization. Lastly, when the actin-based cytoskeleton confinement processes are relaxed, the contribution of lipid-dependent microdomains to confining transmembrane proteins was also readily observed. Overall, our data provide strong evidence supporting the idea that in live cells, a lipid-dependent microdomain organization is responsible for the dynamic molecular confinement of membrane components.

### **FCS diffusion laws provide a robust experimental frame of analysis for investigating the presence of submicron domains in live cells**

Among the methods used to characterize the cell membrane, our novel experimental approach constitutes a significant advance, since it can be used to quantify relevant parameters to explore domain structure in the plasma membrane. Although FRAP and FCS methods involve similar experimental setups, the FCS method provides unique information, mainly because it combines high temporal resolution with single molecule sensitivity and robust statistical analysis (Schwille *et al*, 1999; Bacia *et al*, 2004). By comparison, the FRET methods frequently used to study membrane domains probe direct molecular interactions (Kenworthy and Edidin, 1998; Sharma *et al*, 2004), but these methods are not appropriate for studying long-range interactions and diffusive processes. SPT certainly competes with FCS, since it can be used to determine particle positions to within tens of nm and



in some cases, with a high temporal resolution (Schutz *et al*, 2000; Fujiwara *et al*, 2002). However, accurate interpretation of the SPT results generally requires analyzing large numbers of individual trajectories. Besides, in our FCS experiments, the direct conjugation of the fluorophore on the tracked molecules circumvents unwanted crosslinking and aggregation. Overall, the FCS approach described here for recording measurements on different spatial scales provides a highly appropriate means of exploring domain structure and dynamics in the plasma membranes of live cells.

The diffusion behavior of different subclasses of membrane components was determined here experimentally. Although the sphingolipid analogs show a lower affinity for DRMs than their natural counterparts (our data and Kuerschner *et al*, 2005), they are still useful to probe lipid raft organization, since FCS is sufficiently sensitive to be able to detect even small amounts of the fluorescent molecules in the coexisting phase (Korlach *et al*, 1999). The wide range of time intercepts  $t_0$  measured (from  $-20$  to  $+20$  ms) suggests that different mechanisms may hinder the diffusion of membrane components. Based on our previous simulations (Wawrezynieck *et al*, 2005), these findings indicated that a strictly positive time intercept  $t_0$  resulted from confinement into isolated domains, whereas a strictly negative value reflected the presence of a meshwork. It is significant that the components thought to be associated with lipid rafts, such as GPI-anchored proteins or sphingolipid analogs, showed strictly positive time intercepts, which suggests that a dynamic process of molecular partition into discrete domains mainly hinders their diffusion. By contrast, barriers predominantly impede the diffusion of the TfR-GFP since  $t_0$  was strictly negative. Owing to the null  $t_0$ , the glycerophospholipid analogs apparently underwent unhindered diffusion.

#### **Evidence supporting a lipid-dependent microdomain organization in live cells**

A clearly visible signature of lipid-dependent compartmentalization is provided by COase and SMase enzymatic treatments. Our finding is consistent with previous data showing that both cholesterol and SM make a pivotal structural contribution to order domain formation (Brown and London, 1998; Rietveld and Simons, 1998). These results indicate that the lipid-dependent microdomain organization dictates the overall mobility of sphingolipid analogs and GPI-anchored proteins. It is also worth pointing out that the microdomains detected in the present study do actually exist in the plasma membrane of cells under steady-state conditions. This observation is in line with recently published data showing that phase separation and the formation of lipid domains do in fact occur in live cells (Meder *et al*, 2006).

Although an actin-based system of cytoskeleton compartmentalization has been described in the case of molecules located in the outer leaflet of the plasma membrane (Sako and Kusumi, 1994; Fujiwara *et al*, 2002; Kusumi *et al*, 2005), no such system was observed here in the mode of confinement of GPI-anchored proteins or sphingolipid analogs. Indeed, although cholesterol and SM modifications completely abolished the membrane confinement for these molecules, neither the actin-based cytoskeleton disorganization nor its stabilization had any such effects. Further studies are now required to determine whether the above discrepancies

may be attributable to differences in the cell types, experimental conditions or techniques used. Our results are nevertheless in agreement with those of SPT analyses performed on GPI-anchored MHC class I and DiI molecules (Vrljic *et al*, 2005) as well as with the finding that cholesterol and/or SM modifications reduced the molecular mobility (Kenworthy *et al*, 2004; Nishimura *et al*, 2005; Vrljic *et al*, 2005).

The free diffusion behavior observed in the case of the glycerophospholipid analogs is similar to that previously observed in the case of Cy3-DOPE in the plasma membrane of various cell types at a recording rate of 100 ms/frame (Fujiwara *et al*, 2002; Kusumi *et al*, 2005). However, we were unable to detect any submicron-sized compartments such as those previously observed by authors performing SPT at an enhanced time resolution (25  $\mu$ s/frame) using gold-tagged Cy3-DOPE molecules to label COS-7 cells (Kusumi *et al*, 2005).

#### **A lipid-dependent microdomain organization also impedes the diffusion of transmembrane proteins**

Our data clearly confirm the existence of fences confining transmembrane proteins, even if these molecules are not interacting directly with the cytoskeleton. For instance, the motion of TfR-GFP is mainly controlled by the actin-based membrane skeleton fences, and  $t_0$  decreased when this meshwork was strengthened. It is also worth noting that all these experimental FCS diffusion laws form a bundle of lines with a common crossover point (Figure 7D). Previous simulations have shown that (i) these lines intersect at a single point in the meshwork at different barrier strengths but with the same mesh sizes; (ii) the crossover occurs when the spot area is about one to twice that of the mesh. Since the experimental crossover was obtained with  $w^2 = 5 \times 10^4 \pm 1 \times 10^4$  nm<sup>2</sup>, the area of the mesh was around  $8 \times 10^4 \pm 2 \times 10^4$  nm<sup>2</sup>. Assuming a square mesh, its sides would measure  $240 \pm 60$  nm: this value is in good agreement with the 260 nm mesh size measured in rat kidney fibroblasts, which was determined by SPT experiments performed at 25  $\mu$ s/image (Fujiwara *et al*, 2002). However, in our case, we did not detect any enlarged meshes after a partial disruption of the cytoskeleton. In addition, we observed that a treatment by cytochalasin D at 10  $\mu$ M released TfR-GFP and DPP<sub>IV</sub>-GFP proteins from the meshwork-based state of confinement. Nevertheless, the latter proteins still underwent the lipid domain-based constrained diffusion on a much smaller compartmentalization scale.

Some extremely interesting features have been brought to light by monitoring the motion of these proteins after subjecting them to treatments that disrupt the actin-based cytoskeleton and/or decrease the membrane cholesterol level. This combined treatment applied here induced the recovery of an FCS diffusion law curve, which intercepted the time origin. A single treatment abolishing one of the two major hindrance mechanisms served to show up the other one.

#### **Main features of lipid-dependent microdomains**

Our observations of the sphingolipid and cholesterol dependency for the lipid-dependent microdomains revealed by the FCS diffusion study suggest that they correspond to lipid rafts, a type of microdomains proposed to be involved in numerous cellular functions, but whose existence has been matter of controversy. The current view is that 'membrane

rafts are small (10–200 nm), heterogeneous, highly dynamic, sterol- and sphingolipid-enriched domains that compartmentalize cellular processes' (Pike, 2006). The analysis of our experimental data in the light of the simulation studies allow us to further conclude that the lipid-mediated microdomains described here fit the above description very nicely in many respects.

As mentioned above, our FCS method is sensitive to nanoscale confining structures. Based on dynamic molecular partition simulations, we were able to determine the upper limit of the microdomain radius  $r$  (Supplementary Figure S1). We previously observed that the graph line between  $\tau_d$  and  $w^2$  is solely observed if the observation spot is significantly larger than the domains ( $w^2/r^2 > 10$  in the case of isolated microdomains) (Wawrezynieck *et al*, 2005). In our experiments, the smallest observation spot was close to the transverse resolution limit of the microscope ( $\sim 200$  nm); the upper limit of the microdomain radius  $r$  is therefore around 60 nm: this value is consistent with data published in the literature (Pralle *et al*, 2000; Zacharias *et al*, 2002; Prior *et al*, 2003; Sharma *et al*, 2004).

The  $t_0$  intercept could be of special interest, since it could be used to delineate the temporal characteristics of these microdomains. In the simulation models, this value depends on both the partition factor  $\alpha$  of a given molecule in the microdomains and the confinement time within one microdomain  $\tau_{\text{conf}}$  (see equation (2) in the Supplementary data). The value of  $\alpha$  is not yet available, but based on DRM experiments, we might tentatively take this value to range between 10 and 30%. For instance, with a  $t_0$  value of 18.4 ms in the case of GFP-Thy1, we have 31 ms  $< (\tau_{\text{conf}} - \tau_d^{\text{domain}}) < 92$  ms. Since we measured that  $\tau_d > 30$  ms when  $(w/r)^2 > 10$ , the diffusion time in a domain should be at most 30/10  $\sim 3$  ms. It can therefore be concluded that  $\tau_{\text{conf}}$  ranges between 34 and 95 ms. Confinement in microdomains is therefore an extremely transient process taking probably only some tens to hundreds of milliseconds. Reaching a deeper understanding of  $t_0$  obviously constitutes one of most important and challenging issues awaiting future theoretical analysis. Our experiments showed (Figures 5 and 6) that before reaching the maximum effect (i.e.  $t_0 \sim \text{null}$ ), a parallel existed between the change in the sphingomyelin or cholesterol levels and the decrease in  $t_0$ . It is therefore possible to consider this parameter as a *microdomain confinement index*, since the partition factor  $\alpha$  and the confinement time  $\tau_{\text{conf}}$  are essential parameters characterizing the confinement into microdomains.

We would like to stress the fact that our results do not tell us whether membrane molecules partition dynamically into pre-existing raft domains or whether they directly participate in assembly/disassembly of the domains forming self-promoting membrane complexes. This is because the confinement time  $\tau_{\text{conf}}$  is always to be either shorter than or equal to the lifetime of the microdomain in our models. In the first case, the molecule is partitioning dynamically into a pre-existing domain, whereas in the second case, the molecule accompanies/participates in the assembly and disassembly of the domain. In the same context, we observed in our FCS measurements that the diffusion time  $\tau_d$  remained constant within the limits of the experimental error bars although the amount of fluorescent probe used varied by a factor of 20 (i.e. from  $\sim 100$  to  $\sim 2000 \mu\text{m}^{-2}$  in the case of GFP-GPI)

(Supplementary Figure S2). Consequently, the probe concentration does not greatly affect the diffusion behavior of the probe. However, this is not relevant to the question as to whether we are dealing with pre-existing or self-assembling raft models, because our FCS analysis did not include the probe density, contrary to recent FRET and immuno-EM studies on nanoscale membrane organization (Sharma *et al*, 2004; Hess *et al*, 2005; Plowman *et al*, 2005).

Overall, our data strongly support a dynamic picture where nanometer-scale microdomains confine constituents for up to a few tens to hundreds of milliseconds and undergo continuous changes as molecular diffusion processes occur. These findings also show that studies on diffusion behavior throw particularly useful light on the modes of lateral membrane organization, which are likely to be largely governed by weak molecular interactions.

## Materials and methods

### DNA constructs and cell transfection procedures

GFP-GPI was provided by A LeBivic (IBDM, Marseille, France). DPP<sub>IV</sub>-GFP was obtained from G Trugnan (INSERM, Paris, France). The human TfR cDNA was provided by S Mèresse (CIML, Marseille, France). The Thy1 gene expression vector and the Alexa488-conjugated anti-Thy1 Fab fragment were a kind gift from R Morris (King's College London, UK). To construct GFP-Thy1, three overlapping fragments were PCR amplified and cloned into the pEGFP-C1 plasmid. This yielded a recombinant GFP-Thy1 cDNA encoding the 19th aa of Thy1 (signal peptide), the EGFP and the remaining sequence of Thy1 (aa 20–162). To construct TfR-GFP, the TfR cDNA (aa 1–112) was cloned inframe with the GFP cDNA into the pEGFP-N3 plasmid.

All experiments were carried out on COS-7 cells (ATCC, CRL-1657). Cells were grown in DME supplemented with 10% FCS, glutamine and sodium pyruvate. Transient transfections were performed with ExGen 500 as per the manufacturer's instructions (Euromedex) and 24 h before FCS measurements.

### Fluorescent staining with lipid analogs

Complexes of BODIPY-lipids (Invitrogen) and BSA were prepared in line with the manufacturer's instructions and as described in Martin and Pagano (1994). Briefly, 500 nmol of FL-SM or FL-GM1 stock solution (CHCl<sub>3</sub>:MeOH 19:1) were dried under a stream of nitrogen, and then under a vacuum for at least 1 h. The dried lipid was dissolved in 200  $\mu\text{l}$  of absolute ethanol and injected into 10 ml of defatted BSA solution (5  $\mu\text{M}$  defatted BSA in Hanks buffered salt solution containing 10 mM HEPES pH 7.4, HBSS/HEPES) while vortexing vigorously. FL-PC and FL-PE were prepared in a similar way, except that the lipid to BSA ratio was 2:1 (mol:mol). Cell cultures were washed in HBSS/HEPES, incubated with 0.05  $\mu\text{M}$  lipid/BSA complex in HBSS/HEPES for 30 min at 4°C, washed and further incubated in HBSS/HEPES at 37°C.

### Treatment of cells with enzymes and drugs

To modify the cholesterol or sphingomyelin contents of plasma membrane, cells were treated with *Streptomyces* sp. COase or *Staphylococcus aureus* SMase (Calbiochem) in serum-free HBSS/HEPES buffer at 1 U/ml for 30 min or at 0.1 U/ml for 5 min, respectively. Cells were washed in HBSS/HEPES prior to the diffusion measurements. Prior to pharmacological treatments, cells were washed in HBSS/HEPES and incubated at 37°C with 1  $\mu\text{M}$  latrunculin B (5 min), 2–10  $\mu\text{M}$  cytochalasin D (30 min) or 0.4  $\mu\text{M}$  jaspilkinolide (5 min). All FCS measurements were completed within 30 min of the cell treatment, before any significant morphological changes in the cell could take place.

### Biochemical analyses

To determine the cholesterol levels, cells were scraped off and lysed at 4°C for 30 min in a buffer containing 1% Nonidet P-40, 10 mM Tris pH 7.4, 150 mM NaCl, 1 mM EDTA and protease inhibitors. Samples were cleared by centrifugation at a rate of 12 000 g before

undergoing fluorometric measurements using the Amplex<sup>®</sup>Red Cholesterol kit (Invitrogen).

To determine the sphingomyelin levels, a similar procedure was used, except that the cells were lysed at 4°C by sonication and the PNS harvested by centrifugation. Lipid extraction was performed using the Bligh-Dyer method (Bligh and Dyer, 1959) and the sphingomyelin levels were determined using a fluorometric method with the Amplex<sup>®</sup>Red sphingomyelinase assay kit (Invitrogen) as described in He *et al* (2002). With each test sample, a negative control assay was run on the sample and the reaction mixture devoid of SMase. Sphingomyelin amounts were calculated from the difference in fluorescence between the test and control samples, and compared with a standard.

To isolate DRMs, Brij98 (1%) solubilized postnuclear supernatant was fractionated on the sucrose gradient by centrifugation at 38 000 r.p.m. for 16 h in a SW41 rotor (Beckman Instruments Inc.) (Drevot *et al*, 2002). Fractions were resolved on SDS-PAGE and blotted with anti-GFP mAbs (Roche Molecular Biochemicals). The presence of DRMs was monitored, based on the distribution of Rab5 and ganglioside G<sub>M1</sub> over the sucrose gradient. Spectrofluorimetry measurements were performed on the different fractions of the lipid analogs.

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## Fluorescence microscopy, FCS measurements and numerical analysis

Details of the experimental fluorescence microscopy, FRAP, FCS and numerical analysis procedures used are given in the Supplementary data, as well as in previous studies (Kenworthy *et al*, 2004; Wawrezinieck *et al*, 2004, 2005).

## Supplementary data

Supplementary data are available at *The EMBO Journal* Online.

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