

Electric Supporting Information (ESI)

Trapping and release of giant unilamellar vesicles in microfluidic wells

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Supplementary Figures

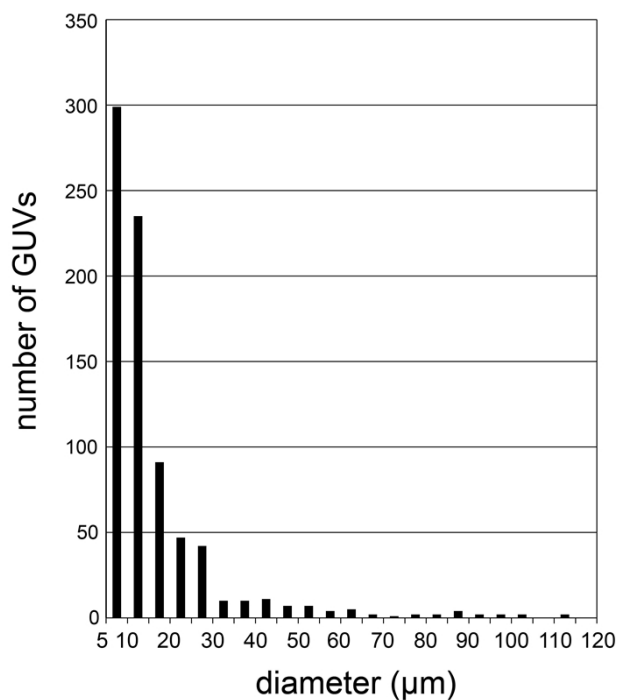


Fig. S1

Size distribution of GUVs prepared by electro-formation method. Immediately after electro-formation, GUVs made of EPC containing 0.5 % Texas Red DHPE (w/w) were transferred to an observation chamber, fluorescence microscopy images were taken, and the diameters measured. GUVs smaller than 5 μm diameter were not counted because of the limitation of image resolution.

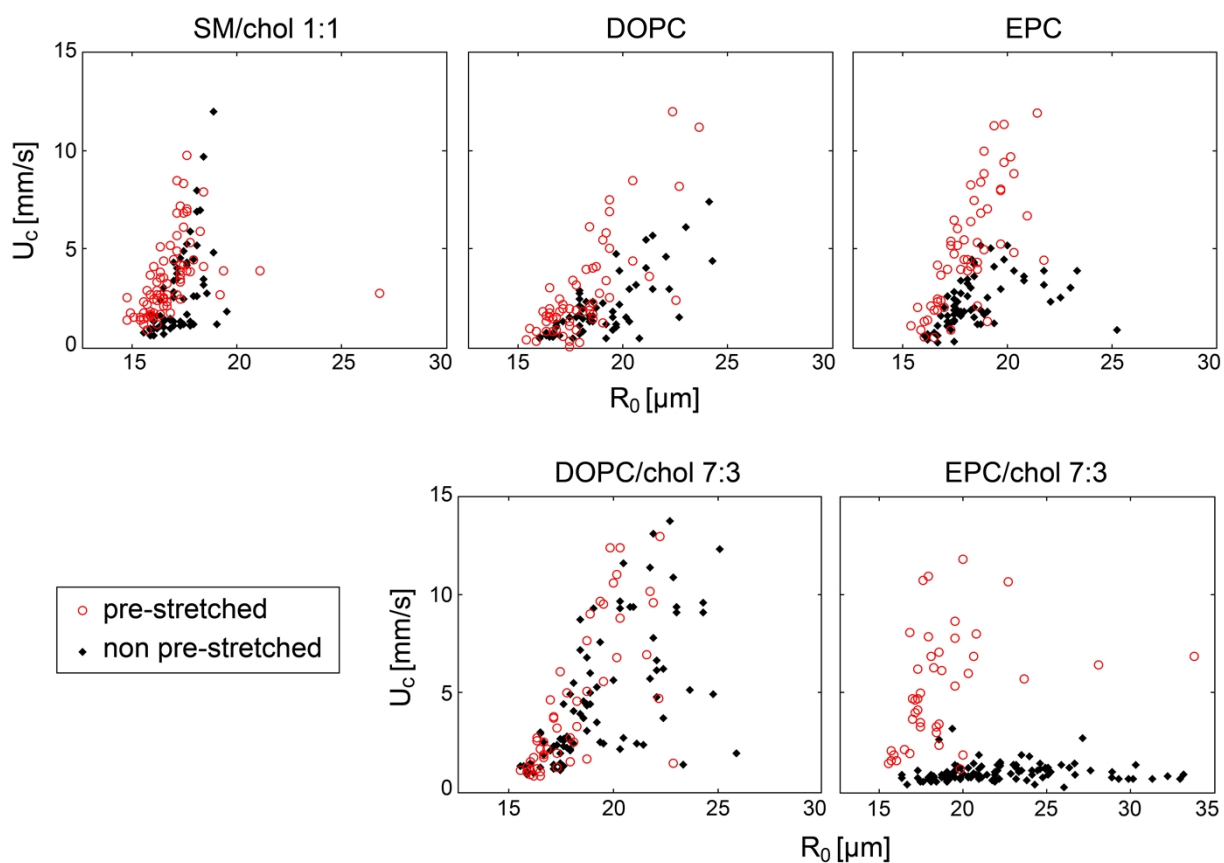


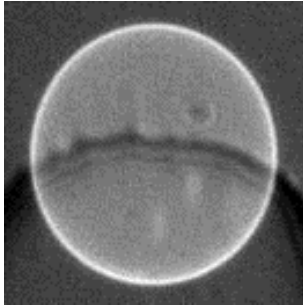
Fig. S2

Difference between pre-stretched and non pre-stretched GUVs. Experimental measurements of the threshold flow velocity U_c for GUV untrapping versus the non-constrained GUV radius R_0 for pre-stretched (red open circles) and non pre-stretched (black filled diamonds) GUVs with different lipid compositions. Data for the pre-stretched GUVs are identical to that in Fig. 5.

Descriptions of Movies

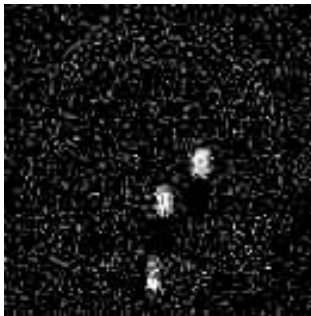
Movie S1

A time-lapse movie showing the flow in the lipid bilayer membrane of a trapped GUV induced by outer medium flow. The fluorescence and transmission images of the GUV in a well were recorded every 100 ms. A still image is shown in Fig. 3 (a). Image size: $84\ \mu\text{m} \times 85\ \mu\text{m}$.



Movie S2

A time-lapse movie showing the extracted fluorescence signals of bright spots in the membrane shown in Movie 1. The images were obtained by sequential subtraction of images from Movie 1. Still images are shown in Fig. 3 (b).



Movie S3

A time-lapse movie showing an escape event of a GUV (fluorescent from a well (dark disks)). The fluorescence and transmission images of GUVs in wells were recorded every 100 ms while ΔP was increased from 5.8 to 29 mbar. The GUV on the right escapes from the well, whereas the left one remains trapped. Note that in this movie, ΔP was increased rapidly for a demonstration, but in normal experiments, ΔP was increased by small increment. Image size: $898\ \mu\text{m} \times 671\ \mu\text{m}$.

