# Journal Name

### ARTICLE

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#### **Supporting Information**

## Driving a Planar Model System into the 3<sup>rd</sup> Dimension: Generation and Control of Curved Pore-Spanning Membrane Arrays

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**Fig. S1** depicts a representative scanning electron microscope (SEM) image (top view) of a porous substrate used for the experiments. SEM measurements were performed with a LEO Gemini 1530 SEM with acceleration voltages between 1 and 6 kV.





**Fig. S2** displays fluorescence images of Texas Red DHPE (A) labelled pore-spanning membranes (DPhPC/PIP<sub>2</sub> 99:1) after the addition of Alexa488-labelled ENTH domain (B). The membranes were imaged 40 min after ENTH domain addition. The overlay of both fluorescence images (C) show the co-localization of both fluorophores indicating a homogeneous binding of ENTH domain to the membranes.



**Fig. S2.** Confocal laser scanning fluorescence microscopy image of **A**. Texas Red DHPE labelled protruded pore-spanning DPhPC/PIP<sub>2</sub> (99:1) membranes after applying an osmolarity gradient of 19 mOsmol/L, and **B**. Alexa488-labelled ENTH domain ( $c = 3 \mu$ M) bound to the membranes. **C.** Overlay of the fluorescence images of **A**. and **B**. Scale bars: 5  $\mu$ m.

**Fig. S3** shows individual time traces of the relative radius  $r/r_0$  of pore-spanning DPhPC/PIP<sub>2</sub> (99:1,  $\mu$ m,  $n_0 = 73$ ) to show the deviation within an experiment.



**Fig. S3.** Change of the radius of individual protruded pore spanning membranes composed of DPhPC/PIP<sub>2</sub> (99:1) as a function of time (gray solid lines) after the addition of 3  $\mu$ M ENTH domain (t = 0 min). The osmolarity gradient was set to 19 mOsmol/L. The thick black line is the average curve calculated from the  $n_0 = 73$  individual curves.

**Fig. S4** illustrates the calculated change in height *h* of the protrusions as a function of the lateral membrane tension  $\sigma$  as obtained from eq. (5) and (6) for an osmolarity gradient of  $\Delta O = 19$  mOsmol/L.



**Movie S1** shows a time series of three dimensional confocal fluorescence *z*-stack images of pore-spanning membranes (DPhPC/PIP<sub>2</sub> 99:1, labelled with Texas Red DHPE after applying an osmolarity gradient of 19 mOsmol/L). Data were acquired with a spinning disk confocal fluorescence microscope and the images were rendered using the program IMARIS. ENTH domain (3  $\mu$ M) was added at *t* = 0 s. Shrinking as well as growing of pore-spanning membranes is visible.