Membrane curvature underlies actin reorganization in response to nanoscale surface topography

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Edited by Charles M. Lieber, Harvard University, Cambridge, MA, and approved September 17, 2019 (received for review June 14, 2019)

Surface topography profoundly influences cell adhesion, differentiation, and stem cell fate control. Numerous studies using a variety of materials demonstrate that nanoscale topographies change the intracellular organization of actin cytoskeleton and therefore a broad range of cellular dynamics in live cells. However, the underlying molecular mechanism is not well understood, leaving why actin cytoskeleton responds to topographical features unexplained and therefore preventing researchers from predicting optimal topographic features for desired cell behavior. Here we demonstrate that topography-induced membrane curvature plays a crucial role in modulating intracellular actin organization. By inducing precisely controlled membrane curvatures using engineered vertical nanostructures as topographies, we find that actin fibers form at the sites of nanostructures in a curvature-dependent manner with an upper limit for the diameter of curvature at ~400 nm. Nanotopography-induced actin fibers are branched actin nucleated by the Arp2/3 complex and are mediated by a curvature-sensing protein FBP17. Our study reveals that the formation of nanotopography-induced actin fibers drastically reduces the amount of stress fibers and mature focal adhesions to result in the reorganization of actin cytoskeleton in the entire cell. These findings establish the membrane curvature as a key linkage between surface topography and topography-induced cell signaling and behavior.

Results and Discussion

Nanotopography Induced F-Actin Accumulation Is Curvature Dependent. Arrays of SiO2 nanopillars with well-defined geometry (diameter of 100 to 1000 nm, height of 1 to 2 μm, and interpillar distance of 2 to 5 μm) are fabricated using top-down e-beam lithography.

Significance

Despite a large body of work, little is known about the origin or underlying mechanisms of how cells interact with surface topographic patterns, which has resulted in a primarily phenomenological approach in studying cell–nanotopography interactions. This study illustrates that membrane curvatures induced by nanoscale surface topography can serve as a direct biochemical signal to activate curvature-sensing protein and regulate actin polymerization and mechanotransduction in the intracellular space.
(EBL) and reactive ion etching on SiO2 substrates (Fig. 1A). Here, the surface chemistry is identical across both the nanostructures and flat areas as they are made of the same material and processed through the same chemical procedure. When U2OS cells are cultured on nanopillar substrates, their plasma membranes wrap around nanopillars as shown by both scanning electron microscopy (SEM) (Fig. 1B) and focused ion beam (FIB)/SEM in our previous study (25). Therefore, vertical nanopillars induce local membrane curvatures with a curvature value determined by the nanopillar diameter (29). The largest contribution to membrane curvature is from tight wrapping around the sidewall, which has a 10 times larger area than the top. Confocal fluorescence imaging of U2OS cells transfected with Lifeact-RFP (an F-actin marker) shows that F-actin distributes along the sidewall and the top of nanopillars (SI Appendix, Fig. S1A). F-actin visualized by phallolidin staining shows significant accumulation at nanopillars of 3 different diameters (arrowheads in Fig. 1C) compared to the surrounding flat area.

To study the effect of different membrane curvatures on F-actin accumulation within the same cell, we engineered a gradient array with nanopillar diameters ranging from 100 nm to 1,000 nm (Fig. 1D and SI Appendix, Fig. S1C). For a single U2OS cell covering a range of nanopillar diameters, F-actin accumulation is often stronger on small nanopillars than on larger ones in the same cell (Fig. 1E). We note that a vertical nanopillar presents an increased membrane area in a projected 2D fluorescence image, and thus the enhanced F-actin signal at nanopillar locations could be due to the increased actin cortex associated with the projected plasma membrane area. To quantify the differences on nanopillars of different diameters, we separately averaged the fluorescence signals of F-actin and GFP-CAAX, a plasma membrane marker, over hundreds of nanopillars for each diameter (Fig. 1F and SI Appendix, Fig. S1 D–F). Both F-actin and GFP-CAAX appear as rings on large-diameter nanopillars and dots on small-diameter ones due to the diffraction limit of the light. Above 400 nm, the normalized F-actin signal by the GFP-CAAX signal stays relatively constant as the nanopillar diameter increases further (Fig. 1G), suggesting that the F-actin rings around large nanopillars are likely due to the increased membrane area at nanopillar locations. In contrast, as the nanopillar diameter decreases from 400 nm to 100 nm, the normalized F-actin signal increases significantly. This trend is preserved when using the supported lipid bilayer as the reference for surface area (SI Appendix, Fig. S1 G and H). The increased F-actin accumulation...
Membrane Curvature Induces the Formation of Branched F-Actin Nucleated by the Arp2/3 Complex. Actin polymerization is mainly initiated by 2 types of actin nucleators: the Arp2/3 complex and formins (Fig. 2A). The Arp2/3 complex induces the formation of branched F-actin while formins promote the growth of linear F-actin (30). When U2OS cells were transfected with Arp3-GFP, a subunit of the Arp2/3 complex, we observed distinct local accumulation of Arp3-GFP at the ends of nanobars with very little signal along the sidewalls (Fig. 2B). A similar distribution of endogenous Arp2/3 complex with a strong bias toward the ends of nanobars is also observed by anti-Arp2/3 immunostaining in untransfected cells (SI Appendix, Fig. S2A). In sharp contrast to Arp3, mDia1-Emerald and mDia2-Emerald, 2 members of the formin family proteins, show no preferential accumulation at the ends of nanobars (Fig. 2C and the corresponding full images in SI Appendix, Fig. S2B).

Moreover, Arp2/3 nucleation promoting factors, N-WASP and Cortactin (31) also exhibit curvature-dependent accumulation at the ends of the nanobars (Fig. 2D and the corresponding full images in SI Appendix, Fig. S2C), which show much higher end-to-side ratio than the membrane control (mCherry-CAAX) does (Fig. 2E). We note that the visibly higher end-to-side ratio of Arp3-GFP, N-WASP-GFP, and GFP-Cortactin in comparison to mCherry-CAAX were always confirmed in the same batch of cotransfection experiments, but the quantified values for the same protein vary from experiment to experiment due to large variations in protein expression levels. Cotransfection of Arp3-GFP, N-WASP-GFP, or GFP-Cortactin with Lifeact-RFP shows that all 3 proteins strongly colocalize with F-actin on nanobars (Fig. 2F–H). The spatial preference of Arp2/3, N-WASP, and Cortactin on the ends of nanobars and their colocalization with F-actin suggests that Arp2/3-nucleated branched actin, rather...
than formin-nucleated linear actin, accumulates on nanostructures in a curvature-dependent manner.

Nanostructure-Induced F-Actin Accumulation on Nanostructures Is Highly Dynamic. In cells, F-actin usually undergoes rapid assembly or disassembly in order to afford cell adhesion, migration, morphogenesis, or force generation. Here, we show that nanostructure-induced F-actin accumulation also exhibits fast dynamics. As shown in Fig. 3A, the kymograph of Lifeact-RFP exhibits characteristic “blinking events” on nanopillars, indicating that F-actin on nanopillars is not stationary but constantly depolymerizes and repolymerizes in a time-dependent manner. The dynamic on-off behavior of F-actin accumulation is apparent on small nanopillars (Movie S1). On the other hand, the F-actin rings around large nanopillars repolymerizes in a time-dependent manner. The dynamic on nanopillars is not stationary but constantly depolymerizes and repolymerizes in a time-dependent manner. The dynamic on-off behavior of F-actin accumulation is apparent on small nanopillars (Movie S1). On the other hand, the F-actin rings around large nanopillars are of a different process and likely due to the increased membrane area around large nanopillars. Quantitative analysis of Lifeact-RFP dynamics indicates that F-actin assembles and disassembles with a frequency of ∼0.4/min (Fig. 3C). Single cell measurements show that drug inhibition of Arp2/3 using inhibitor CK666 significantly reduces the dynamic fluctuations of F-actin on nanopillars (Fig. 3C). In contrast, a formin FH2 inhibitor SMIIFH2 did not perturb F-actin dynamics on nanopillars (Fig. 3D).

Furthermore, Arp2/3, N-WASP, and Cortactin also exhibit highly synchronized dynamic with F-actin on nanopillars (Fig. 3E and F and Movies S2–S4). On the other hand, neither mDia1-Emerald nor mDia2-Emerald shows observable accumulations or dynamic fluctuations on nanopillars. The dynamic correlation between F-actin and actin-related proteins on nanopillars was further evaluated by using Pearson correlation coefficients (Fig. 3G). The correlation coefficients between Lifeact-RFP and Arp3-GFP (0.68 ± 0.02, SEM), GFP-Cortactin (0.69 ± 0.02), and N-WASP-GFP (0.54 ± 0.02) indicate high-level correlation. On the other hand, mDia1-Emerald (0.04 ± 0.02) and mDia2-Emerald (0.09 ± 0.01) show very little correlation. The temporal correlation of F-actin with Arp3, N-WASP, and Cortactin further confirms the involvement of these proteins in nanotopography-induced F-actin assembly.

Curvature Sensing Protein, FBPI7, Mediates the Branched F-Actin Formation at Nanotopography-Induced Membrane Curvatures. Neither Arp2/3 complex nor its nucleation promoting factors are known to recognize membrane curvatures. Their connecting factor is the key to understand the underlying molecular mechanism of the topography-triggered F-actin formation. Here, we hypothesize that curvature-sensitive BAR proteins (32) can serve the role of bridging actin polymerization with membrane curvature. In mammalian cells, the Toca F-BAR family, including FBPI7, Toca-1, and CIP4, are known to contain an N-terminal F-BAR domain that senses membrane curvature, and a C-terminal SH3 domain that interacts with N-WASP and cdc42.

Fig. 3. Curvature-dependent F-actin accumulation on nanostructures is highly dynamic. (A, Left) A time-lapse image of a cell transfected with Lifeact-RFP (Movie S1). (A, Right) A kymograph plot shows that Lifeact-RFP signals on nanopillars repeatedly appear and disappear. (B) Representative trajectories of F-actin dynamics before and after CK666 treatment. (C) Kinetic analysis shows that F-actin dynamics on nanopillars are significantly reduced by Arp2/3 inhibitor CK666. (D) A formin inhibitor SMIFH2 does not alter F-actin dynamics on nanopillars. (E and F) Time-lapse images of Lifeact-RFP with Arp3-GFP, Lifeact-RFP with GFP-Cortactin, and Lifeact-RFP with N-WASP-GFP on individual nanopillars with 15-s time interval show correlated dynamic fluctuations. (G) The quantified Pearson correlation coefficient between the dynamics of Lifeact-RFP and 5 F-actin related proteins, n = 89 to 220 trajectories for each protein (see SI Appendix, Table S3 for detailed statistics). Welch’s t test, P values indicated in C, D, and G, ****P < 0.0001; ns = 0.08. Error bars represent SEM. (Scale bar, 10 μm [A, full image], 2 μm [A, kymograph, horizontal], 2 min [A, kymograph, vertical].)
(33) to modulate actin cytoskeleton (34, 35) (Fig. 4A). A previous in vitro study indicates that they are able to induce curvature-dependent F-actin assembly on vesicle membranes (36). Therefore, we hypothesized that these proteins recognize the topology-induced membrane curvatures to induce F-actin accumulation on nanostructures.

Three Toca family proteins GFP-FBP17, Toca1-GFP, or CIP4-GFP were all shown to accumulate at the ends of nanobars similar to Arp2/3 complex, Cortactin, and N-WASP (Fig. 4B and SI Appendix, Fig. S3 A and B). The accumulation is often more pronounced under the nucleus due to the lower fluorescence background inside the nucleus, and high protein expression induces tubule formation (37). For the following studies, we focus on widely expressed FBP17 that also shows the strongest response among the 3 members. We further confirmed the curvature response of FBP17 using wedged nanobars which are narrow (high curvature) at one end and wide (low curvature) at the other end (Fig. 4C). GFP-FBP17 has a much higher fluorescence intensity at the narrow end than the wide end on the same nanobar (Fig. 4 D and E).

Taking advantage of the gradient geometry generation by EBL fabrication, we quantitatively probed the range of curvature values that can induce FBP17 accumulation using gradient nanobar arrays with the bar width varying from 100 nm to 1,000 nm (Fig. 4F). The width of each nanobar determines the curvature value at its ends. To corroborate the transient transfection studies used in Fig. 4 B and D, we used anti-FBP17 immunostaining to probe the spatial distribution of endogenous FBP17. The averaged nanobar images of different widths show that endogenous FBP17 accumulates strongly to the 2 ends of nanobars with a width smaller than 400 nm, but do not show an obvious response to a width of 500 nm or above (Fig. 4G). The curvature threshold for FBP17 accumulation is similar to that for F-actin. The trend is more clearly shown in the quantitative analysis in Fig. 4H.

The role of FBP17 is validated by constructing a truncated FBP mutant (FBP17ΔSH3), which contains the curvature-sensing F-BAR domain but not the C-terminal SH3 domain for inducing actin polymerization (Fig. 5A). Spatial distribution of both GFP-FBP17 and mCherry-FBP17ΔSH3 show preferential accumulation on the ends of nanobars (Fig. 5B), indicating that FBP17ΔSH3 maintains the ability to recognize nanostructure-induced plasma membrane curvature (Fig. 5C).

Interestingly, overexpression of mCherry-FBP17ΔSH3 abolishes the nanostructure-induced F-actin accumulation on nanopillars and nanobar ends. As shown in Fig. 5D, U2OS cells cotransfected with GFP-FBP17 and Lifeact-RFP show clear accumulation of both

![Fig. 4. FBP17 preferentially accumulates at high curvature locations induced by nanostructures. (A) A schematic drawing of Toca family proteins activating Arp2/3-mediated actin polymerization. (B) GFP-FBP17 preferentially accumulates to the ends of nanobars. (C) An SEM image of wedged nanobars. (D) On wedged nanobars, GFP-FBP17 shows a much higher signal at the narrow end (high curvature) than the wide end (low curvature). (E) An intensity plot along a line of wedged nanobars illustrates that GFP-FBP17 prefers the narrow ends of wedged nanobars. (F) A gradient nanobar array with 2-μm length, 800-nm height, 5-μm center-to-center distance between nanobars, and a variable width from 100 to 1,000 nm with 100-nm increment. Each width is represented by 2 bar columns in the array. (G) The averaged nanobar images of anti-FBP17 immunostaining for 10 different nanobar widths, from the largest 1,000 nm to the smallest 100 nm. n = 1,278 to 1,728 nanobars (see SI Appendix, Table S4 for detailed statistics). (H) The measured nanobar end-to-side ratio of anti-FBP17 decreases with the bar width but levels off when the bar width is 500 nm or larger. Error bars represent SEM.]
GFP-FBP17 and Lifeact-RFP at the ends of nanobars. On the other hand, in cells transfected with mCherry-FBP17ΔSH3, the preferential accumulation of Lifeact-GFP at the same locations are largely eliminated (Fig. 5E). Quantitative analysis shows that the nanobar end-to-side ratio of Lifeact in mCherry-FBP17ΔSH3 transfected cells is significantly lower than that of GFP-FBP17 transfected cells (Fig. 5F). The decrease of F-actin intensity on nanopillars positively correlates with the expression level of FBP17ΔSH3 (SI Appendix, Fig. S3 C and D). Similarly, the reduction of Arp2/3 at nanobar ends was also observed in cells overexpressing mCherry-FBP17ΔSH3 (Fig. 5G and H) and the corresponding full images in SI Appendix, Fig. S4). These results indicate that the accumulation of FBP17 on membrane curvatures interferes with nanostructure-induced activation of Arp2/3 complex and prevents F-actin polymerization.

**Nanostructure-Induced F-Actin Polymerization Reduces the Formation of Actin Stress Fibers and Mature Focal Adhesions.** Actin polymerization is involved in many cellular processes. Our previous study shows that nanostructure-induced membrane curvature enhances clathrin-mediated endocytosis (28), so we examine whether F-actin assembly at nanostructures is related to the endocytosis process. Inhibiting clathrin-mediated endocytosis by Pitstop 2 does not significantly affect F-actin accumulation or its dynamic fluctuations on nanopillars (SI Appendix, Fig. S5 A–C). Similarly, the spatially biased accumulation toward the ends of nanobars persists after Pitstop 2 treatment (SI Appendix, Fig. S5 D and F). When the nanostructure-induced F-actin accumulation is blocked by overexpression of mCherry-FBP17ΔSH3, the preference of the clathrin-adaptor protein AP2-GFP toward the ends of nanobars is reduced but does not fully disappear (SI Appendix, Fig. S5 E and G). This result suggests that nanostructure-induced F-actin is not exclusively involved in the endocytosis process.

In addition to endocytosis, we also probed how nanostructure-induced formation of branched F-actin affects the formation of linear F-actin stress fibers and their associated focal adhesion complexes, which are essential for mechanotransduction, cell adhesion, contraction, and migration. For this purpose, we examined the cellular distribution of F-actin and 3 core protein components of mature focal adhesions, vinculin, paxillin, and Myosin II. The results show that, unlike F-actin that preferentially accumulates at the ends of nanobars, vinculin, paxillin, and Myosin IIb show very little signal either at the ends or along the sidewalls of nanobars (Fig. 6A and B). Occasionally, vinculin and paxillin show weak fluorescence signal on nanobars, but there is no preference to the ends. Therefore, nanostructure-induced F-actin fibers do not appear to participate in mature focal adhesions.

When stained by phalloidin and anti-paxillin, cells cultured on flat areas show a network of long and straight stress fibers whose ends colocalize with large paxillin patches of similarly oriented and elongated focal adhesions (Fig. 6C). On the other hand, cells cultured on nanopillar areas show very different F-actin patterns with clear accumulation around nanopillars, illustrating the regularly

![Fig. 5](https://www.pnas.org/cgi/doi/10.1073/pnas.1910166116)
spaced arrays but much fewer stress fibers. Paxillin staining also shows a highly diffusive signal in cell and the paxillin patches are much smaller in size. Statistical analysis shows that the ratio of stress fiber area to cell area is reduced by 45% on nanopillars (Fig. 6E) and the number of large focal adhesion (>0.5 μm²) per cell decreased by 90% (Fig. 6F). When treated with blebbistatin, a specific Myosin II inhibitor that is known to cause disassembly of actin stress fibers (38), the long F-actin stress fibers and the large paxillin patches on the flat surface have mostly disappeared (Fig. 6D). Interestingly, blebbistatin treatment did not block nanostructure-induced F-actin formation on nanopillars (Fig. 6D). Furthermore, quantitative analysis shows that the total intensity of F-actin on nanopillars per cell has almost doubled after blebbistatin treatment (Fig. 6G). Quantitative analysis shows that nanostructure-induced F-actin formation and stress fiber F-actin negatively correlated with each other to induce whole-cell F-actin reorganization on nanopillars (SI Appendix, Fig. S6). This anticorrelation could be due to their competition for the same pool of actin monomer or due to other nanostructure-induced changes of mechanotransduction (39). These results also agree with the previous report that impairment of stress fibers prohibits the stabilization of nascent focal adhesions and therefore the formation of mature focal adhesions (40).

Conclusions

In this work, we show that local membrane curvature induced by surface topography serves as a biochemical signal to induce actin reorganization inside cells (Fig. 7). Curvature-sensing proteins, FBP17 in particular, are able to recognize topography-induced curvature of a limited range (diameter of the curvature <400 nm) to activate downstream signaling components including N-WASP, Cortactin, and the Arp2/3 complex to nucleate the formation of branched F-actin. This topography-induced F-actin actively impacts multiple cellular processes, such as endocytosis, stress fiber organization, and focal adhesion maturation. While this work does not preclude other potential mechanisms, it supports the

Fig. 6. Nanostructure-induced F-actin accumulation reduces the formation of stress fibers and mature focal adhesions. (A) Nanostructure-induced F-actin formed at the ends of nanobars does not colocalize with Myosin IIb, paxillin, or vinculin, core protein components of mature focal adhesion. (B) Quantified nanobar end-to-side ratios for F-actin, Myosin-IIb, paxillin, and vinculin. (C) Anti-paxillin staining and phalloidin staining of F-actin show a clear reduction of F-actin stress fibers and the size of focal adhesion patches on nanopillars as compared with flat areas. (D) Blebbistatin treatment (25 μM, 30 min) has largely removed the F-actin stress fibers and paxillin patches in U2OS cells on both flat areas and nanopillars. On the other hand, the F-actin accumulation on nanopillars still persists. (E) The percentage of stress fiber area for U2OS cells cultured on flat surfaces is much larger than that on nanopillar arrays. (F) The number of large paxillin focal adhesion patches (>0.5 μm²) per cell is significantly reduced on nanopillar arrays as compared with that on flat areas. (G) The F-actin intensity per nanopillar per cell increased dramatically upon blebbistatin treatment. Welch’s t test with P values are indicated. Error bars represent SEM. (Scale bars, 10 μm [A, C, and D].)
hypothesis that topography-induced membrane curvature is a key mechanism underlying how cells recognize and respond to surface topography. The knowledge gained here will assist the effort to design material surface topography to achieve desired cell behavior.

Materials and Methods

Nanostructure Fabrication. The SiO2 nanostructures in this work were fabricated on quartz coverslips by electron-beam lithography and reactive ion etching (28, 41). The patterns of nanostructures were written by electron-beam lithography (Raith 150) and developed in xylene. Metal masks were formed by sputtering with 100 nm of Cr and liftoff in acetone. The quartz nanostructures were generated by anisotropic reactive ion etching with CHF3 and O2 mixture (AMT 8100 etcher, Applied Materials).

Cell Culture, Transfection, and Live Cell Imaging. U2OS cells were used in this study. Before cell plating, nanostructure substrates were sterilized by O2 plasma and coated in 0.1% gelatin. U2OS cells were cultured for at least 18 h before fixation or imaging. Electroporation was used for plasmid transfection. Live-cell imaging was carried out using an epifluorescence microscope (Leica DMI 6000B). During the imaging, cells were maintained at 37 °C with 5% CO2 in an onstage incubator (INUBSF-ZILCS, Tokai Hit).

Immunostaining and Imaging. Cells were fixed in 4% paraformaldehyde, washed by PBS, permeabilized in 0.1% Triton X-100, blocked in 1% BSA, and followed by staining with 2 μg/mL secondary antibodies. The prepared samples were imaged by epifluorescence microscope (Leica DMI 6000B) or confocal microscopy (Olympus FV1000).

Quantitative Analysis. For the quantitative analysis, the images were processed by a rolling ball algorithm with a radius of 12 pixels. The circle masks were created to cover the pillar area. The pixel intensities within the masks were integrated and the intensity with the same nanopillar diameter was averaged. The ratios of F-actin were calculated by dividing the mean intensities of F-actin with GFP-CAAX.

Statistics. Sample sizes were validated using power analysis similar to previous work (28). Welch’s t tests (unpaired, tailed, not assuming equal SD) and the Kolmogorov-Smirnov test (unpaired, not assuming Gaussian distribution) were used to evaluate the significance. All tests were performed using Prism (GraphPad Software). Data are presented as mean ± SEM or mean ± SD as stated in the figure captions.

ACKNOWLEDGMENTS. We thank C. Kaplan in D.G.D. group, Q. Ong in B.C. group, and E. C. Norby in J. Theriot group for helpful advice. Nanofabrication and characterization were performed in Stanford Nano Shared Facilities (SNF) and Stanford Nanofabrication Facility (SNF). This work was funded by NIH (RO1GM117263 to B.C. and R35GM118114 to D.G.D.), the Packard Foundation (to B.C.), Heart Rhythm Society (to F.S.), and Arnold and Mabel Beckman Foundation (to M.A.).
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