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Confinement induces actin flow in a meiotic cytoplasm

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In vivo, F-actin flows are observed at different cell life stages and participate in various developmental processes during asymmetric divisions in vertebrate oocytes, cell migration, or wound healing. Here, we show that confinement has a dramatic effect on F-actin spatiotemporal organization. We reconstitute in vitro the spontaneous generation of F-actin flow using *Xenopus* meiotic extracts artificially confined within a geometry mimicking the cell boundary. Perturbations of actin polymerization kinetics or F-actin nucleation sites strongly modify the network flow dynamics. A combination of quantitative image analysis and biochemical perturbations shows that both spatial localization of F-actin nucleators and actin turnover play a decisive role in generating flow. Interestingly, our in vitro assay recapitulates several symmetry-breaking processes observed in oocytes and early embryonic cells.

cytoskeleton | self-organization | symmetry breaking

Actin-based dynamics are observed at different stages of embryonic and adult cell life and participate in various essential processes, such as cytokinesis (1), asymmetric divisions in vertebrate oocytes (2–4), cell migration (5, 6), or wound healing (7–9). Recent findings have highlighted key functional roles of cytoplasmic F-actin in oocytes and eggs, such as spindle positioning in mouse oocytes (2–4, 10, 11) or the chromosome congression and intracellular transport in starfish (12, 13). The generation and maintenance of such cellular functions are often correlated with symmetry breaking and spatiotemporal reorganization of F-actin assembly (14, 15). In vitro assays had provided important contributions in the understanding of the spatial organization and dynamics of functional F-actin structures (16–20). In addition, the effect of the geometry of boundary conditions on the cytoskeleton organization have been investigated using in vitro compartmentalization of cytoskeleton proteins (21–26) or using surface patterning techniques (27). *Xenopus* egg extracts had provided a powerful system to dissect mechanisms of many aspects of cell division in vitro and has been extensively used to study bipolar spindle assembly (28–30). F-actin and microtubule interactions were examined in interphase extracts (31) and actomyosin dynamic has been linked to cell cycle regulation using egg extracts (32). However, despite these progresses, the spatiotemporal dynamics of cytoplasmic F-actin network, and in particular of actin flows, are still unclear. In order to gain some insights into the generation and maintenance of cytoplasmic F-actin flow dynamics, we have developed an in vitro system capturing the spatial organization of cytoskeleton filaments within a confined geometry that mimics the cellular environment. This system, based on the cellular reconstitution by confinement of *Xenopus* cytoplasmic extracts, is used to examine functional *Xenopus* egg extracts in a confined environment while keeping some of the advantages of the cell extracts: biochemical manipulations, straight-

forward control over cell cycle progression, and very good optical properties in regard to the difficulty to image large embryonic cells. More precisely, we generate femto-liters of cytoplasmic *Xenopus* egg extracts confined within emulsion droplets stabilized at the boundary with a phospholipid monolayer or an amphiphilic polymer. F-actin spatial organization in confined geometry contrasts drastically from bulk. Whereas bulk F-actin organized in a random network, symmetry in F-actin organization is broken in confined geometry. F-actin self-organized into a contractile ring-like structure surrounded by an actin cloud with the presence of a flow of F-actin. To decipher the mechanisms involved in this process, we quantify the dynamical behavior of F-actin flow. Observations of F-actin dynamics within droplets showed network remodeling with both filaments nucleation at the boundary condition and filament transport directed radially inward. Inhibition or modification of F-actin polymerization dynamics by cytochalasin D or phalloidin alters the flow and symmetry breakings. Addition of a domain of Scar (Scar-WA/pWA), a cellular activator of Arp2/3 that contains both the actin and Arp2/3 binding motifs, stimulates bulk nucleation of F-actin and impedes symmetry breakings and flow dynamics. We finally investigate the influence of the cell cycle state and find that F-actin dynamics in confined interphase extracts show a behavior qualitatively similar to those observed in confined metaphase extracts. Inhibition of myosin-II activity suggests that contractile elements participate in flow dynamics but are not essential to flow maintenance.

Results

Symmetry Breaking in F-actin Organization Within Droplets. To observe F-actin network organization in bulk cell extracts, we have supplemented metaphase II-arrested *Xenopus* oocyte extracts with 50 nM of Alexa488-conjugated phalloidin. After few minutes of incubation at room temperature of such bulk extract, we observed an interconnected network of F-actin filling the entire extract volume (Fig. 1A). F-actin fibers organized into meshworks that were either homogeneous in space or that could exhibit a spatial heterogeneity in F-actin concentration (Fig. 1A, *Left* and *Right*, respectively).

To investigate the dynamics of F-actin network in a confined environment mimicking the cell context, we entrapped functional

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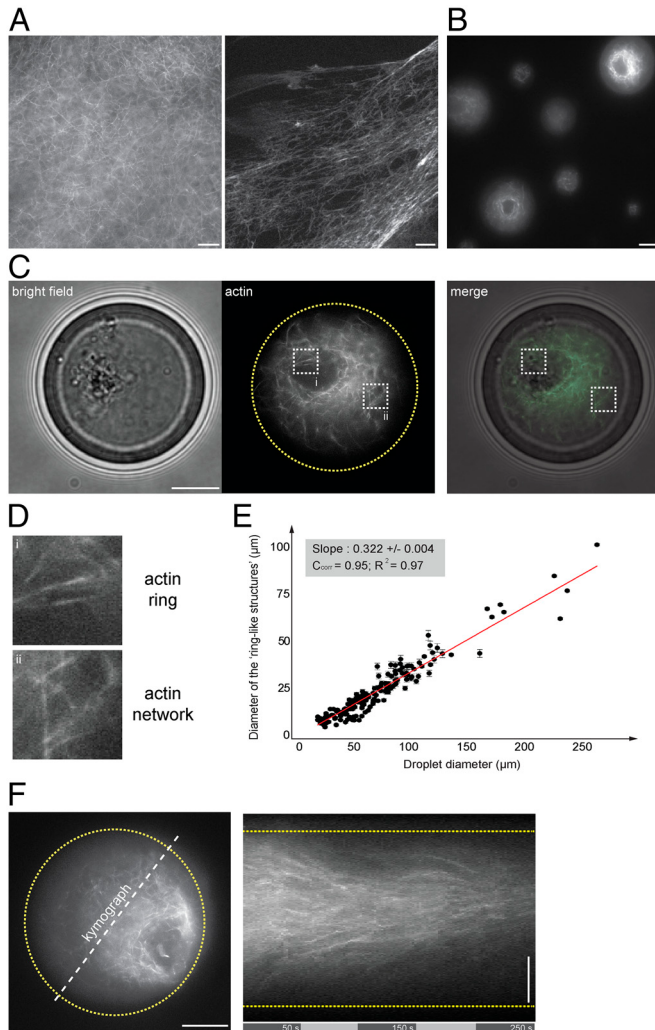


Fig. 1. Symmetry breaking of confined F-actin network. (A) Fluorescent observation of F-actin network (in presence of Alexa488-conjugated phalloidin) generated in bulk extracts (5 min incubation). (B) Fluorescent observation of F-actin network within extract-in-oil droplets. (C, D) Bright field and fluorescent observations of F-actin network confined within a 34 μm -diameter droplet. (D) Actin filaments organized in a ring-like structure asymmetrically positioned within the droplet and surrounded by an actin cloud (actin network). (E) Plot of the ring-like structure diameter as a function of the droplet diameter shows a linear correlation between the ring and the droplet diameter. Quantifications of the ring diameters were performed by extracting a plot profile of the fluorescent structure. The droplet diameter was measured using bright field illumination. We estimated the precision of such measurements to 1 μm . (F) Representative kymograph illustrating the directional flow dynamics of F-actin network within a 40 μm -diameter droplet. Scale bars, 10 μm .

cytoplasm using in vitro compartmentalization methods. These approaches were successfully applied to confine purified proteins (21, 25, 33), and we recently demonstrated that *Xenopus* extracts retain their capability of assembling microtubules when confined within droplets (26, 34). We generated droplets of *Xenopus laevis* egg extracts dispersed in mineral oil (26, 34). This method allows the formation of a large number of spherical droplets with diameters ranging from 10 to 200 μm (Fig. 1B). F-actin spatial organization in confined geometry contrasted drastically from bulk. After few minutes of incubation, the majority of the droplets (more than 90%) contained bright fluorescent actin bundles organized in a very distinct pattern than the ones observed in unconfined conditions (bulk). In droplets, F-actin filaments self-organized in highly stereotyped structures evoking a ring-like

structure that is surrounded by an actin cloud (Fig. 1 *C* and *D* and Fig. S1A). The observation of several droplets in a same field of view showed that the ring-like structures localized randomly in space within droplets (Fig. 1*B* and Fig. S1*B*).

To study the role of the confinement size on actin pattern organization, we measured the apparent diameter of the ring-like structures as a function of the droplet diameter (Fig. 1E, 20 to 250 μm , 169 droplets, nine independent experiments). We found a linear correlation between the ring and the droplet diameter, suggesting a scaling of F-actin self-organization with the droplet size. For smaller droplets, we found that 40% of the droplets exhibit a ring-like structure, whereas 60% failed in organizing a ring-like structure (Fig. S1 C and D, 50 droplets, and seven independent experiments). Altogether, these data suggest that F-actin self-polarized and accumulated asymmetrically within droplets in contrast with unconfined environment (bulk experiments).

Dynamics of F-actin Network: Spontaneous Flow Generation Within Droplets. Interestingly, 1–3 min after the initiation of the F-actin nucleation, we observed the spontaneous generation of F-actin flow within droplets (Fig. 1F and Movies S1 and S2). Closer observations of F-actin dynamics within droplets showed network remodeling with both filament nucleation at the vicinity of the boundary and filament transport directed radially inward (Fig. S2A). In addition, bright field observations permitted to distinguish a local enrichment of cytoplasmic materials entrapped within the dense F-actin meshwork (Fig. 1C). This suggests that F-actin flow may convey cytoplasmic materials through a process that could be linked to F-actin based transport. To provide some insights to this hypothesis, we monitored cytoplasmic material movement during the formation of F-actin ring-like structures using fluorescently labeled beads as passive markers (Fig. S1E). The beads entrapped by the meshwork of filaments eventually accumulated within the ring-structures. These experiments are reminiscent to those recently described in starfish oocytes, where directional transport by F-actin meshwork was proposed to guide passive large objects by steric trapping (13), or where F-actin dynamics were linked to cytoplasmic streaming in mouse oocytes (10). Finally, in some observations, the ring-like structure position moved randomly within the droplet space (Fig. S2B and Movie S2). Altogether these observations suggest that F-actin flow in droplet is mainly directed radially inward, with a centripetal direction from the periphery of the droplet to a convergent area.

In order to get some insights on the mechanisms involved in F-actin flow generation, we first characterized the dynamics behavior of the meshwork in bulk and in droplets. We used optical flow methods to extract the spatiotemporal positions of nodes or bright filaments of the F-actin network (Fig. 2, [SI Materials and Methods](#)). For instance, we reported on Fig. 2A representative examples of the displacement fields of trajectories as a function of time. Computation of the velocity field and vector flow orientation showed variability in velocity distribution and orientation of bulk F-actin networks. We have reported the analysis of approximately 600 trajectories on [Fig. S2C](#) (six independent experiments, mean = $44 \pm 13 \text{ nm s}^{-1}$). More generally, about 60% of experiments exhibited F-actin dynamics associated with a mean velocity of about $10\text{--}20 \text{ nm s}^{-1}$, whereas about 40% were characterized by a velocity reaching approximately $50\text{--}80 \text{ nm s}^{-1}$ (40 independent experiments). In addition, orientation of F-actin displacement fields could be either similar or distinct throughout the same field of observation (examples 1, 2 and 3, 4, respectively). Both heterogeneity due to cell extract preparations and intrinsic variability of the mechanisms involved in F-actin dynamics may account for the variation observed.

Next, we quantified F-actin dynamics in a confined environment. Fig. 2*B* displays the displacement field of F-actin meshwork within a single droplet (approximately 30 trajectories). For this

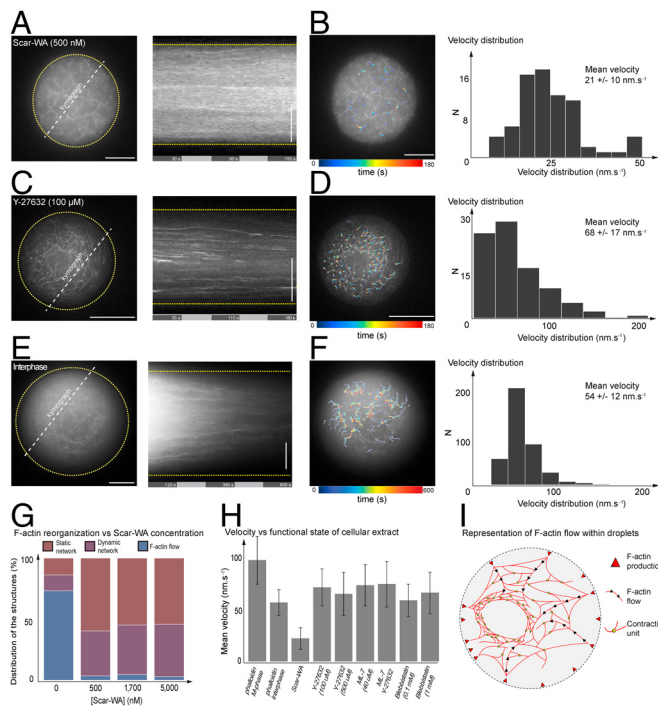


Fig. 4. Testing the role of the spatial localization of F-actin nucleation sites, the requirement of myosin II, and the regulation of the cell cycle in the generation of symmetry breaking and actin flow. (A) Kymograph of F-actin dynamics within droplet in presence of Scar-WA (500 nM). (B) Velocity field in presence of Scar-WA inhibitor (500 nM) and velocity distribution ($N = 3$ droplets, approximately 70 trajectories, mean = 21 ± 10 nm). (C) Kymograph of F-actin network in presence of Rho-Kinase inhibitor Y27632 (100 μ M). (D) Velocity field of F-actin network dynamics in presence of Rho-Kinase inhibitor Y27632 (100 μ M) and velocity distribution ($N = 5$ droplets, approximately 100 trajectories, mean = 68 ± 17 nm). (E) Kymograph of F-actin dynamics within droplet in interphase state. (F) Velocity field in interphase state and velocity distribution ($N = 10$ droplets, approximately 500 trajectories, mean = 54 ± 12 nm). (G) Histogram of events characterizing the network dynamic state for increasing concentration of Scar-WA (0, 0.5, 1.7, and 5 μ M). (H) Histogram of the mean velocity as a function of the functional state of confined cellular extract. (I) Schematic of F-actin-based structure and flow within droplet. Scale bars, 10 μ m.

ing drastically with observations performed with unperturbed extracts (Movie S3). The dynamics of these networks exhibited two main features: We found in average 40% of the droplets that contained a disorganized meshwork fluctuating erratically as a function of time whereas in 57% of the observations, the patterns were rather static (Fig. 4B and G and Fig. S3C). Addition of Scar-WA/pWA promotes the formation of dendritic networks that are known to increase the local actin concentration (47). The modulus of elasticity of semi-flexible polymer networks is expected to scale with approximately $(C_A)^{5/2}$, where C_A is the actin concentration (48). This suggests that F-actin network enhanced by Scar-WA/pWA has a larger elasticity than the one generated in absence of Scar nucleators. This stiffness may counterbalance the compressive forces caused by inward flows until it completely impedes them. The homogeneous nucleation of filaments within the droplet space through ScarWA stimulation may counterbalance the importance of F-actin production at the boundary. These results suggest the role of the spatial localization of F-actin production in the formation of symmetry breakings and flow.

Role of Myosin-II on F-actin Flow Dynamics. In unconfined *Xenopus* extract, it has previously been demonstrated that myosin-II activity powered contraction of F-actin meshwork (32). We therefore investigated the role of myosin-II in the establishment of the F-actin symmetry breaking and flow. Myosin II inhibitors, such

as blebbistatin (0.1 to 1 mM), ML7 (40 μ M), and a Rho-associated kinase (ROCK) inhibitor (Y-27632, 100, or 500 μ M), were added alone or in combination within the cell extracts. Interestingly, F-actin flow was persistent in presence of these myosin II inhibitors (Fig. 4C). A closer look to the quantification of the F-actin network dynamics in presence of the Rho-kinase inhibitors showed a mean velocity reduced by approximately 30% (Fig. 4D). ML7 and blebbistatin also induced slight modifications of the flow dynamics (Fig. S4A). We also found that the evolution of the F-actin mesh size as a function of time was modified when myosin inhibitors were added to cell extracts (Fig. S4B and C). These data support myosin II participation as force generating elements to the contractile behavior of the F-actin network, but is not crucial in generating F-actin flow. However, they can contribute to the cell polarity determination, as observed for instance during retrograde flux in motile keratocytes (6). Other myosin-like motor proteins could be involved in the F-actin network remodeling.

Role of the Cell-Cycle State on F-actin Flow Dynamics. F-actin dynamics and motor proteins are dependent on the cell cycle states of the oocyte cytoplasm (32). For instance, contractility is abolished in interphase extracts contrary to metaphase extracts (M-phase). Since our experiments were performed in M-phase extracts, we have tested the effect of cell cycle regulation on F-actin dynamics using interphase extracts. We observed F-actin flow dynamics directed inward within confined interphase extracts, with a behavior qualitatively similar to those observed in confined M-phase extracts (Fig. 4E). Next, we performed a quantitative analysis of the velocity field of confined interphase extracts and found that the mean velocities were reduced by roughly 40% (Fig. 4F; v approximately 54 nm s^{-1} , 10 droplets, approximately 500 trajectories, and five independent experiments). Therefore, in interphase extracts, e.g., in absence of contraction, vectorial F-actin dynamics are still emerging as observed in M-phase extracts, with a reduced velocity.

Discussion

The combination of biochemical perturbations and quantitative image analyses allowed us to examine how confinement induces the generation of symmetry breaking and F-actin flow. Experiments consisting in perturbing actin polymerization or filament nucleation growth lead to the inhibition of F-actin flow process. On the other hand, inhibiting myosin-II activity or changing the cell cycle state of cell extracts did not significantly perturb F-actin flow dynamics in confined environment (Fig. 4H). Altogether, these data support the primary role of actin dynamics and nucleation in the generation of F-actin flow. In our experiments, the droplet boundary plays the role of structural anisotropy of the environment by localizing filament growth. Indeed, the confinement geometry provided by the droplets participates in the determination of the final actin steady-state organization.

One hypothesis that may explain the formation of ring-like structures is the ability of F-actin flow to concentrate cytoplasmic components within a restricted area of the droplet. The growth of the ring organization may be correlated with the accretion of actin filaments (Fig. 4I). The generation and maintenance of such directional transport of F-actin could involve spatially localized nucleation and actin turnover, such as observed during M-II mouse oocytes and actin retrograde flow of keratocytes and fibroblasts.

Interestingly, our observations evoke F-actin networks involved in physiological functions in oocytes and embryonic systems: maintenance of the spindle asymmetry through cytoplasmic streaming in mouse M-II oocyte (10), chromosome congression and intracellular directional transport in starfish oocyte (12, 13), or pronucleus fusion in *C. elegans* (49).

We anticipate that this assay will be useful to examine cytoplasmic F-actin organization and dynamics involved in oocytes and vertebrate development with a bottom-up approach.

Materials and Methods

Extract-in-oil Droplet Formations. Extract-in-oil droplet formation was prepared as previously described (26, 34). Briefly, egg PC or the block-polymer PHS-PEO-PHS was first dissolved in mineral oil (0.7 mg/mL and 0.25 mg/mL, respectively). The *Xenopus* CSF egg extracts were added to the mixture of mineral oil/polymer [1% (v CSF/v oil)] at 4 °C. The mixture was gently sheared, by pipetting up and down the solution during few seconds, to generate extract-in-oil droplets (*SI Materials and Methods*).

Imaging. Fluorescence imaging of actin droplets was performed using an Ix81 inverted microscope (Olympus) and a x60 (Plan Apo, NA 1.42) objective, equipped with an electron multiplying charge-coupled device (EM-CCD) camera (C9102 or Imagem, Hamamatsu Corp., Sewickley, PA). Microscope settings

and functions were controlled using SimplePCI software (Hamamatsu). Image analysis was performed using ImageJ (Scion Image), SimplePCI, and MATLAB software programs.

Spatiotemporal Image Tracking and Analysis. We used the Lucas–Kanade feature tracking algorithm in order to extract the specific position of F-actin network (*SI Materials and Methods*). The principle of the Kanade–Lucas Tracking (KLT) algorithm is based on optical flow techniques applied to a small number of features detected in the first image of the sequence.

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