

RESEARCH ARTICLE

Cofilin regulates actin network homeostasis and microvilli length in mouse oocytes

Anne Bourdais, Benoit Dehapiot* and Guillaume Halet[†]

ABSTRACT

How multiple actin networks coexist in a common cytoplasm while competing for a shared pool of monomers is still an ongoing question. This is exemplified by meiotic maturation in the mouse oocyte, which relies on the dynamic remodeling of distinct cortical and cytoplasmic F-actin networks. Here, we show that the conserved actin-depolymerizing factor cofilin is activated in a switch-like manner upon meiosis resumption from prophase arrest. Interfering with cofilin activation during maturation resulted in widespread elongation of microvilli, while cytoplasmic F-actin was depleted, leading to defects in spindle migration and polar body extrusion. In contrast, cofilin inactivation in metaphase II-arrested oocytes resulted in a shutdown of F-actin dynamics, along with a dramatic overgrowth of the polarized actin cap. However, inhibition of the Arp2/3 complex to promote actin cap disassembly elicited ectopic microvilli outgrowth in the polarized cortex. These data establish cofilin as a key player in actin network homeostasis in oocytes and reveal that microvilli can act as a sink for monomers upon disassembly of a competing network.

KEY WORDS: Cofilin, Oocyte, Actin, Microvilli, LIMK, Mouse

INTRODUCTION

Actin filaments (F-actin) control a vast array of cellular processes requiring membrane deformation or force production, such as cell motility and contractility, cell division, vesicular trafficking and cell polarization (Pollard and Cooper, 2009). Multiple actin networks coexist in the cell, exhibiting elaborate architectures that are defined primarily by their nucleation machinery (Michelot and Drubin, 2011; Blanchoin et al., 2014). Thus, the Arp2/3 complex generates branched actin networks by promoting the nucleation of new filaments from the side of existing ones (Mullins et al., 1998). In contrast, formins nucleate and/or elongate linear actin cables to promote the formation of structures such as stress fibers, filopodia and the cytokinetic ring (Chesarone et al., 2010). An emerging concept in the field is inter-network competition, by which distinct actin structures compete for a finite pool of actin monomers (G-actin), thereby regulating network size and density (Burke et al., 2014; Suarez and Kovar, 2016). How this homeostatic balance is regulated in living cells is not fully understood. So far, emphasis has been placed on the relative amounts and assembly rates

of nucleators (i.e. Arp2/3 versus formins), along with the key role of profilin in allocating G-actin to formins at the expense of Arp2/3 (Rotty et al., 2015; Suarez et al., 2015; Antkowiak et al., 2019; Faust et al., 2019). However, little is known of the contribution of F-actin disassembly mechanisms, particularly in a cellular context.

Actin filaments are structurally polarized, with a fast-growing barbed end and a slow-growing pointed end (Blanchoin et al., 2014). *In vivo*, the barbed end is the preferred site for monomer addition and filament elongation, whereas the pointed end favors monomer disassembly, resulting in a directional shuttling of actin subunits known as treadmilling (Carlier and Shekhar, 2017). Actin-binding proteins of the ADF/cofilin family (herein referred to as cofilin) play an instrumental role in actin network turnover by promoting filament severing and depolymerization from the pointed end, thereby replenishing the cytoplasmic G-actin pool (Lappalainen and Drubin, 1997; Hotulainen et al., 2005; Suarez et al., 2011; Shekhar and Carlier, 2017; Wioland et al., 2017). In addition, cofilin lowers the affinity of Arp2/3 for both the mother and daughter filaments, resulting in debranching and depolymerization of dendritic networks (Blanchoin et al., 2000; Chan et al., 2009). If not rapidly capped, the free barbed ends generated through severing could seed new filament growth (Chan et al., 2000; Ichetovkin et al., 2002). Thus, cofilin plays a dual role in actin homeostasis, promoting either a net increase or decrease in F-actin density. Phosphorylation at Ser3, catalyzed by LIM kinases (LIMKs), prevents cofilin binding to actin, and thereby inhibits its filament-depolymerizing activity (Arber et al., 1998; Yang et al., 1998; Mizuno, 2013). Intriguingly, structural studies have revealed a noncanonical kinase–substrate interaction mode, which may account for the exquisite selectivity of LIMKs toward cofilin (Hamill et al., 2016; Salah et al., 2019).

Oocytes offer an excellent experimental paradigm to address actin network homeostasis at the cell scale. During meiotic maturation, mouse oocytes experience a profound remodeling of their actin cytoskeleton to achieve symmetry breaking and polar body extrusion (reviewed in Uraji et al., 2018; Duan and Sun, 2019). Thus, eccentric positioning of the first meiotic spindle relies on a dynamic network of cytoplasmic F-actin nucleated by Fmn2 (Azoury et al., 2008; Schuh and Ellenberg, 2008; Pfender et al., 2011; Yi et al., 2013). The density of this actin mesh drops steeply at meiosis resumption, and its gradual reassembly drives spindle migration toward the oocyte cortex (Azoury et al., 2011; Holubcová et al., 2013; Wei et al., 2018). Fmn2-dependent actin filaments are also found along spindle microtubules (spindle actin), where they promote kinetochore fiber formation and chromosome alignment (Mogessie and Schuh, 2017). Spindle migration also relies on an Arp2/3-dependent cortical actin thickening, which contributes to decreasing cortical tension (Chaigne et al., 2013). In mature oocytes arrested in metaphase II (MII), a prominent actin cap is established in the cortex overlying the spindle, defining the site

Institut Génétique et Développement de Rennes, CNRS IGDR UMR 6290, Université de Rennes 1, F-35000 Rennes, France.

*Present address: Turing Center for Living Systems, Aix Marseille Université, CNRS IBDM UMR 7288, 13009 Marseille, France.

[†]Author for correspondence (guillaume.halet@univ-rennes1.fr)

 B.D., 0000-0002-7559-5497; G.H., 0000-0002-6204-2105

Handling Editor: Michael Way

Received 9 August 2021; Accepted 18 November 2021

for second polar body protrusion after fertilization (Longo and Chen, 1985). Filaments in the cap are assumed to be arranged as a branched network, as they rely on polarized activation of the Arp2/3 complex (Yi et al., 2011). Remarkably, actin cap formation is invariably associated with the collapse of microvilli in the polarized cortex, but the molecular underpinnings of this cytoskeletal remodeling are still elusive (Maro et al., 1986; Van Blerkom and Bell, 1986; Dehapiot and Halet, 2013).

In this study, we explored the role of cofilin in regulating F-actin dynamics during mouse oocyte maturation. Unexpectedly, we found that cofilin is largely inactivated (i.e. phosphorylated) during prophase arrest and becomes active at meiosis resumption. Inhibition of cofilin during meiosis I via LIMK1 overexpression resulted in a striking elongation of microvilli and a loss of the cytoplasmic F-actin mesh, which was associated with spindle migration and cytokinesis defects. In contrast, cofilin inhibition in MII oocytes resulted in a shutdown of F-actin dynamics together with a considerable expansion of the polarized actin cap. Remarkably, ectopic microvilli outgrowth, restricted to the polarized cortex, was achieved by promoting actin cap disassembly. These data establish cofilin as a critical component in actin network homeostasis in oocytes, in line with cell cycle progression. Furthermore, our findings highlight a role for cofilin in regulating the length of microvillar core bundles and reveal a mechanism for microvilli elongation based on the disassembly of a competing network.

RESULTS

Cofilin activation promotes spindle migration and polar body extrusion

Regulation of cofilin activity is mainly achieved via inhibitory phosphorylation at Ser3, which is catalyzed by LIMKs. Whereas LIMK expression and activation has been detected across meiotic maturation in mouse oocytes (Li et al., 2017; Duan et al., 2018), the time course of cofilin phospho-regulation has not yet been described. We performed western blot analyses using oocyte lysates prepared at four landmark stages of meiotic maturation: prophase arrest (germinal vesicle stage, GV), nuclear envelope breakdown (NEBD), metaphase I (NEBD+6 h; MI) and metaphase II (MII). Phospho-cofilin was readily detected in GV oocytes, and its level decreased by ~50% at meiosis resumption (NEBD) and decreased further to less than 20% of GV level in MI and MII oocytes (Fig. 1A), indicating cofilin activation. The drop in phospho-cofilin level was not due to cofilin downregulation, because total cofilin levels remained unchanged (Fig. 1A).

To explore the functional significance of cofilin activation, we first attempted cofilin depletion via siRNA injection. However, despite successful knockdown of cofilin mRNA, protein levels were little affected (~20% depletion; Fig. S1), suggesting that cofilin is a relatively abundant and stable protein – a recurrent pitfall for RNA interference strategies in oocytes (Pfender et al., 2015). Alternatively, LIMK1 overexpression is a well-established strategy for acute cofilin inactivation in a variety of cell types (Arber et al., 1998; Yang et al., 1998; Sotiropoulos et al., 1999; Bierne et al., 2001; Takahashi et al., 2001; Amano et al., 2002; Higashida et al., 2013). Hence, we injected prophase oocytes with a cRNA encoding mouse LIMK1 and assessed phospho-cofilin levels by western blotting. Surprisingly, phospho-cofilin was not increased further in LIMK1-expressing GV oocytes (Fig. S2, left), indicating that cofilin is already largely inactivated during prophase arrest. In contrast, a substantial increase in phospho-cofilin was observed when LIMK1-injected oocytes were released from prophase arrest and cultured to the MI stage, whereas the catalytically inactive LIMK1^{D460A} (Yang et al., 1998) had no effect

(Fig. S2, middle). Spontaneous meiosis resumption was not overtly affected by LIMK1 or LIMK1^{D460A} overexpression (Fig. 1B). Likewise, the onset and progression of anaphase I was little affected by exogenous LIMK1 or LIMK1^{D460A} (Fig. S3). We noticed, however, that LIMK1-expressing oocytes undergoing anaphase I failed to form a polar body, arguing for a delay or inhibition of cytokinesis (Fig. S3). Accordingly, only 20% of LIMK1-injected oocytes had progressed to the MII stage and emitted a polar body after overnight culture, as compared to 87% of control oocytes and 76% of LIMK1^{D460A}-injected oocytes (Fig. 1C). The majority (72/96 oocytes) of arrested LIMK1-injected oocytes exhibited a bipolar spindle located centrally, pointing to a defect in spindle migration (Fig. 1D). Yet, chromosome spreading revealed that most of these arrested oocytes contained 40 univalents (19/26 oocytes examined; Fig. 1E), demonstrating that they had achieved homologous segregation, but failed to form a polar body.

To further investigate the possible spindle migration defect, oocytes released from prophase arrest were treated with nocodazole to prevent spindle formation, allowing us to monitor chromosome migration powered by cytoplasmic actin dynamics (Van Blerkom and Bell, 1986; Li et al., 2008). As expected, chromosomes migrated to the cortex in the majority of control and LIMK1^{D460A}-injected oocytes, leading to the formation of an actin cap (Fig. 1F). In contrast, chromosomes failed to migrate in oocytes expressing LIMK1 (Fig. 1F), likely reflecting a defect in actin dynamics.

Taken together, these data show that successful oocyte maturation relies on a switch-like activation of cofilin at meiosis resumption in order to execute two major actin-dependent processes: spindle migration and polar body formation. Since a centrally-located spindle does not preclude oocyte cleavage at anaphase I (Pfender et al., 2011; Wei et al., 2018), our data suggest a role for cofilin in cytokinesis, in line with previous findings in cultured somatic cells and *Xenopus* oocytes (Abe et al., 1996; Hotulainen et al., 2005).

Cofilin does not contribute to cytoplasmic F-actin dynamics in prophase oocytes

In line with its canonical role in promoting pointed-end disassembly, cofilin has been suggested to regulate cytoplasmic F-actin dynamics in mouse oocytes (Jang et al., 2014; Jo et al., 2016). However, our finding of cofilin being largely inactivated in GV oocytes is inconsistent with this view. To exclude any contribution from residual active cofilin, we overexpressed LIMK1 in prophase-arrested GV oocytes and examined the integrity of the cytoplasmic F-actin mesh using phalloidin staining in fixed oocytes and EGFP-tagged utrophin calponin homology domain (EGFP-UtrCH) as a probe for F-actin dynamics in live oocytes (Azoury et al., 2008; Schuh and Ellenberg, 2008). Remarkably, GV oocytes overexpressing LIMK1 exhibited a dense cytoplasmic actin network indistinguishable from that of control oocytes (Fig. 2A). In line with previous observations in metaphase oocytes (Azoury et al., 2008; Schuh and Ellenberg, 2008), cytoplasmic F-actin exhibited constant remodeling in a highly dynamic fashion, in both control and LIMK1-expressing oocytes, while at the GV stage (Movie 1). Moreover, and consistent with previous findings (Azoury et al., 2011), both control and LIMK1-expressing oocytes showed a significant drop in cytoplasmic F-actin density shortly before NEBD (Fig. 2B; Movie 1). These observations argue against a significant role of cofilin in promoting cytoplasmic F-actin dynamics during prophase arrest. Surprisingly,

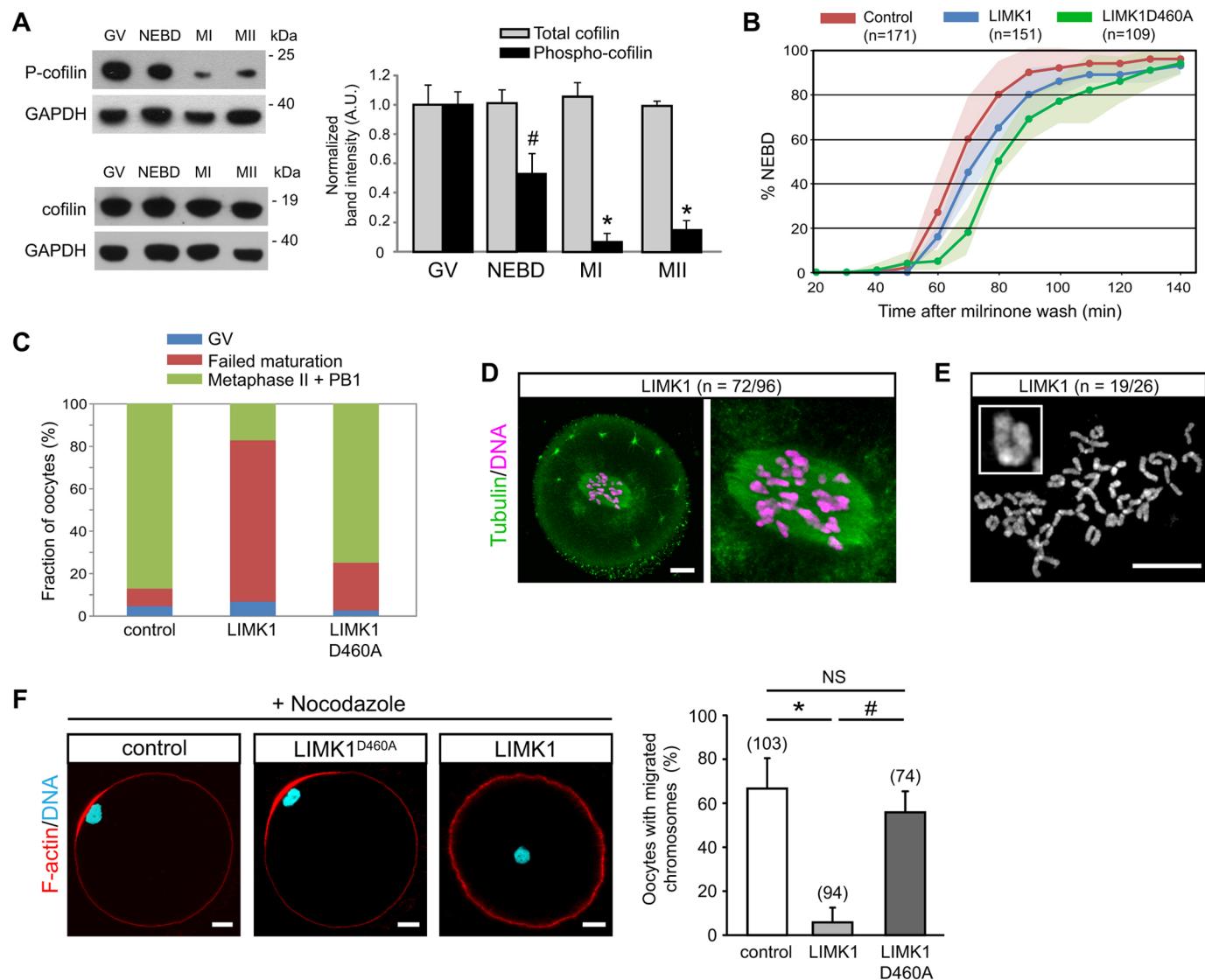


Fig. 1. Cofilin activation promotes oocyte maturation. (A) Detection of phospho-cofilin (P-cofilin; top left) and total cofilin (bottom left) in oocyte lysates at distinct stages of maturation. GAPDH was used as a loading control. The bar graph (right) shows the quantification of band intensities using the GV stage as a reference, and after normalization to the GAPDH signal (A.U., arbitrary units). Data are mean±s.e.m. of three independent experiments. [#] $P=0.026$ compared to GV, ^{*} $P<10^{-4}$ compared to GV (Student's *t*-test). (B) Time course of spontaneous meiosis resumption following milrinone washout in oocytes injected with water (control), LIMK1 cRNA or LIMK1^{D460A} cRNA at the GV stage. Data are mean±s.d. of four independent experiments. The number of oocytes scored is indicated in parentheses. (C) Maturation rate of control, LIMK1- and LIMK1^{D460A}-expressing oocytes. Oocytes were injected at the GV stage. After overnight culture, oocytes were fixed and stained for spindle microtubules and chromatin, for staging. Oocytes that achieved NEBD but failed to emit the first polar body (PB1) are referred to as 'failed maturation'. Over 100 oocytes (from three to five experiments) were scored for each experimental condition. (D) Configuration of the meiotic spindle in a fixed LIMK1-injected oocyte that failed to emit PB1 after overnight culture. Green, anti-tubulin staining; magenta, DNA staining with TO-PRO-3. The right panel shows an expanded view of the spindle. Both images are z-projections of 13 consecutive confocal frames. Scale bar: 10 μm. (E) Chromosome spread of a LIMK1-injected oocyte that failed to emit PB1 after overnight culture. Note the presence of 40 univalents, indicative of homolog separation (19/26 examined oocytes). The inset shows an expanded view of a univalent. Scale bar: 10 μm. (F) Rate of chromosome migration to the cortex in nocodazole (5 μM)-treated oocytes. Representative images of a control oocyte, and oocytes expressing LIMK1^{D460A} or LIMK1, are shown (left). Red, F-actin stained with Alexa Fluor 568-phalloidin; cyan, DNA stained with TO-PRO-3. Scale bars: 10 μm. The bar graph (right) shows the rate of chromosome migration in nocodazole-treated control, LIMK1- and LIMK1^{D460A}-expressing oocytes fixed and stained after overnight culture. Data are mean±s.d. of three to five independent experiments. The number of oocytes scored is indicated in parentheses. * $P<10^{-4}$, # $P<10^{-3}$; NS, not significantly different (Student's *t*-test).

cofilin also appears dispensable for the collapsing of the actin mesh at meiosis resumption. However, we consistently observed bulky F-actin structures in the nucleoplasm of LIMK1-expressing GV oocytes (see Fig. 2A, middle), suggesting that a small pool of active cofilin is engaged in regulating nuclear F-actin turnover during prophase arrest. This is consistent with the reported enrichment of cofilin in the GV nucleoplasm (Jin et al., 2020).

Cofilin inhibition disrupts cytoplasmic and cortical F-actin networks

In view of the spindle migration defect in LIMK1-injected oocytes, we examined the integrity of the cortical and cytoplasmic F-actin networks, which are both required for successful symmetry breaking (Azoury et al., 2008; Schuh and Ellenberg, 2008; Chaigne et al., 2013). Intriguingly, the cortex of MI oocytes

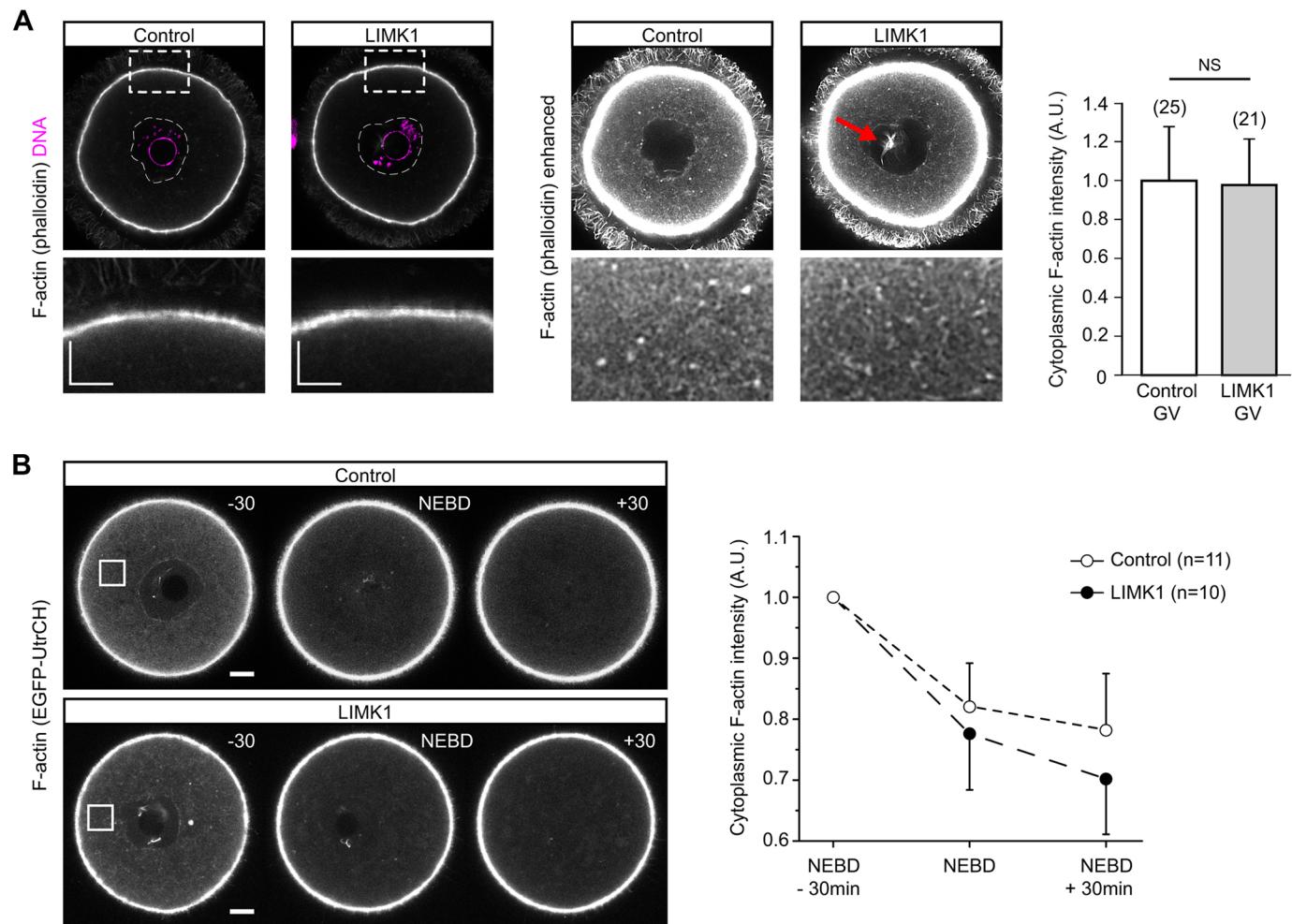


Fig. 2. Cofilin is dispensable for F-actin dynamics in prophase oocytes. (A) F-actin staining with Alexa Fluor 568–phalloidin in prophase-arrested GV oocytes. LIMK1 was overexpressed via cRNA injection followed by culture for 3 h (LIMK1). Left: F-actin imaging with regular settings, avoiding saturation. The GV is depicted by a dashed line. Chromatin is labeled with To-Pro-3 (DNA, magenta). The bottom row shows expanded views of the cortical regions delineated by the dashed boxes. Scale bars: 5 µm. Middle: same images with enhanced brightness for better visualization of the cytoplasmic F-actin mesh. The bottom row shows expanded views of a cytoplasmic region (20×15 µm). Note the thick F-actin bundles in the nucleoplasm of the LIMK1-expressing GV oocyte (arrow). Right: quantification of cytoplasmic F-actin density. The intensity of the phalloidin signal was measured in a cytoplasmic 10×10 µm region. The mean value obtained in control oocytes was normalized to 1. The number of oocytes examined is indicated in parentheses. Data are mean±s.d. of three independent experiments. NS, not significantly different (Student's *t*-test). (B) Live imaging of F-actin networks in control GV (top) and LIMK1-injected GV (bottom) oocytes expressing EGFP–UtrCH and undergoing meiosis resumption. Still images corresponding to 30 min before NEBD (−30), NEBD and 30 min after NEBD (+30) are shown. Fluorescence from the F-actin probe was measured in a 10×10 µm region (white boxes) and plotted against time (right). Data are mean±s.d. The values obtained at 30 min before NEBD were normalized to 1. The number of oocytes examined is indicated in parentheses. Scale bars: 10 µm.

(NEBD+6 h) overexpressing LIMK1 was lined with short (1–3 µm in length) linear actin bundles pointing toward both the extracellular milieu and the cell interior (Fig. 3A, LIMK1), suggestive of an elongation of microvillar core bundles and their cytoplasmic actin rootlets (Wong et al., 1997). Under our experimental conditions, regular microvilli in control oocytes could not be resolved accurately (Fig. 3A, control) due to their fairly short length (<0.5 µm; Benamarra et al., 2017). Interestingly, whereas the cytoplasmic F-actin network was readily detectable in control MI oocytes, it had virtually vanished in LIMK1-expressing MI oocytes (Fig. 3B,C, LIMK1). Live imaging of F-actin dynamics confirmed the absence of the cytoplasmic actin mesh in LIMK1-expressing MI oocytes (Movie 2). In contrast, expression of catalytically inactive LIMK1^{D460A} did not alter microvilli length nor the integrity of the cytoplasmic F-actin mesh (Movie 2), indicating that the observed

F-actin defects were effectively linked to the kinase activity of LIMK1. To confirm the specificity of the phenotype, oocytes were co-injected with LIMK1 and the constitutively active cofilin XAC(A3) (Takahashi et al., 2001) in an attempt to rescue F-actin-severing activity. Expression of the constitutively active non-phosphorylatable mutant cofilin effectively abolished microvilli overgrowth in LIMK1-expressing oocytes (Fig. 3A,B and Fig. 4C,D). Cytoplasmic F-actin staining did not recover however (Fig. 3B,C), consistent with the previous observation that constitutively active cofilin dissolves the F-actin mesh (Jin et al., 2020).

To test whether microvillar actin overgrowth was related to the previously reported Arp2/3-dependent cortical actin thickening (Chaigne et al., 2013), we supplemented the culture medium with the Arp2/3 inhibitor CK-666 (Nolen et al., 2009). Strikingly,

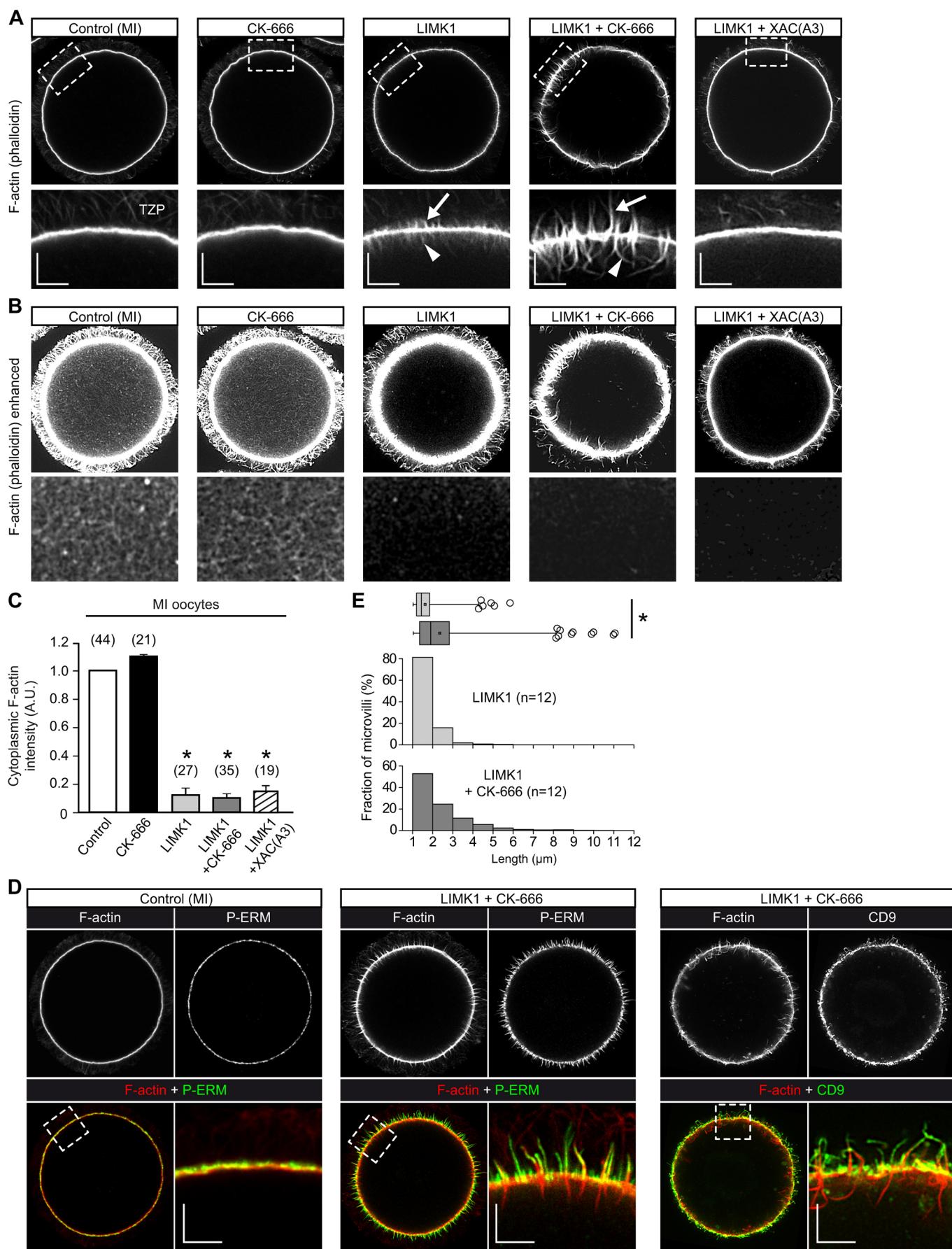


Fig. 3. See next page for legend.

Fig. 3. Cofilin regulates cortical and cytoplasmic F-actin homeostasis in MI oocytes. (A) F-actin staining with Alexa Fluor 568–phalloidin in fixed MI (NEBD+6 h) oocytes, under various experimental conditions. LIMK1 cRNA was injected at the GV stage followed by *in vitro* culture to NEBD+6 h (LIMK1). CK-666 (100 µM) was added at the time of milrinone washout to study the effects of Arp2/3 inhibition alone (CK-666), or in combination with LIMK1 overexpression (LIMK1+CK-666). Co-expression of LIMK1 and XAC(A3) was achieved by co-injecting the respective cRNAs in GV oocytes [LIMK1+XAC(A3)]. The bottom panels are expanded views of the cortical regions delineated by the dashed boxes. The arrows and arrowheads point to elongated microvillar actin bundles and cytoplasmic actin rootlets, respectively. Transzonal projections originating from surrounding follicular cells are also faintly labeled (TZP). Scale bars: 5 µm. (B) Same images as the top row in A, with enhanced brightness for better visualization of the cytoplasmic F-actin mesh (or the absence thereof). The bottom panels are expanded views of a cytoplasmic region (20×15 µm) avoiding the spindle area and actin rootlets. (C) Quantification of cytoplasmic F-actin in fixed MI oocytes, as shown in B. The mean intensity of the phalloidin signal was measured in a cytoplasmic 10×10 µm region. The mean value obtained in control oocytes was normalized to 1. The number of oocytes examined is indicated in parentheses. Data are mean±s.d. of three to five independent experiments. * $P<10^{-4}$ compared to control (Student's *t*-test). (D) Control MI (left) and LIMK1-expressing MI oocytes treated with 100 µM CK-666 (middle and right) were fixed and stained for F-actin (Alexa Fluor 568–phalloidin) together with P-ERM (left and middle) or CD9 (right). Expanded views of the cortical regions (dashed boxes in the merge images) are shown. Scale bars: 5 µm. See also Movie 4. (E) Effect of Arp2/3 inhibition on microvilli length. LIMK1-expressing oocytes were fixed at NEBD+6 h, and microvilli were detected using FiloQuant (with a cut-off at 1 µm) in a single confocal section at the equator of the cell. Microvilli lengths were pooled into 1 µm size bins plotted against frequency. Corresponding box plots (same x axis) are shown, with higher values displayed as outliers. The mean (square), median (line), 25th and 75th percentiles (box), and 5th and 95th percentiles (whiskers) are indicated. Twelve oocytes were analyzed in each condition. * $P<10^{-4}$ (Mann–Whitney test).

Arp2/3 inhibition exacerbated microvillar actin outgrowth, leading to the formation of thick bundles intersecting the cortical actin layer that were tapered at both ends and often heavily bent (Fig. 3A, LIMK1+CK-666; Movie 3). Accordingly, the outward processes were decorated by phosphorylated ezrin/radixin/moesin (P-ERM; Fig. 3D; Movie 4) and the tetraspanin CD9 (Fig. 3D), two classic markers of microvillar identity (Runge et al., 2007; Pelaseyed and Bretscher, 2018). To quantify the effects of CK-666 on microvilli length, we used Filoquant, an ImageJ plugin originally designed to extract filopodia parameters (Jacquemet et al., 2017; Fig. S4). Whereas detected microvilli were mostly in the 1–2 µm length range in LIMK1-overexpressing MI oocytes, CK-666 treatment shifted their length toward the 2–5 µm range, with some microvilli reaching over 10 µm (Fig. 3E). These values are likely an underestimation, because a number of ‘broken’ microvilli could not be resolved along their entire length within the plane of the confocal section (see Fig. S4). Importantly, microvilli overgrowth, and the concurrent loss of cytoplasmic F-actin, were not observed in oocytes treated with CK-666 alone (Fig. 3A–C, CK-666). In contrast, microvilli length occasionally reached 4–6 µm in LIMK1-injected MI oocytes without CK-666 addition (Fig. 3E), indicating that Arp2/3 inhibition was synergistic, but not an absolute requirement.

Collectively, these results show a requirement for cofilin activation at meiosis resumption in order to preserve the integrity of the cortical and cytoplasmic F-actin networks. Interfering with cofilin activation leads to microvilli overgrowth and a reciprocal loss of cytoplasmic F-actin, suggesting that microvilli act as a sink for actin monomers at the expense of F-actin mesh reassembly during maturation.

Microvillar actin elongation relies on G-actin supply, CDK1 and formins

In view of the unaltered F-actin mesh disassembly at meiosis resumption in cofilin-inhibited oocytes (see Fig. 2B), we speculated that the released actin monomers were diverted to microvilli, precluding cytoplasmic mesh reassembly later during maturation. Consistent with this idea, LIMK1-expressing MI oocytes showed significantly lower levels of cytoplasmic G-actin, as visualized by DNase I staining (Fig. 4A). Accordingly, co-expression of XAC(A3) to prevent microvilli overgrowth restored the G-actin pool to near-control levels (Fig. 4A), consistent with microvilli acting as a sink for monomers. To further investigate the possible link between collapsing of the F-actin mesh and microvillar elongation, we injected LIMK1 cRNA after oocytes had reached MI (NEBD+6 h) and recovered a dense cytoplasmic F-actin mesh (as shown in Fig. 3B, control). After additional culture for 3 h to allow for protein expression (NEBD+9.5 h), oocytes were fixed and stained for chromosomes and F-actin. As illustrated in Fig. 4B, a large fraction (35/54 oocytes, 65%) of these oocytes were blocked in a late anaphase I configuration, while the majority of time-matched control oocytes had emitted the first polar body (34/45 oocytes, 76%). This is consistent with the cytokinesis failure described earlier for oocytes injected at the GV stage (Fig. 1C–E). Interestingly, these anaphase I-arrested oocytes showed no sign of microvilli elongation (0/54 oocytes) and exhibited a dense cytoplasmic F-actin network akin to that of control oocytes (Fig. 4B). These findings indicate that in oocytes with an intact cytoplasmic mesh, LIMK1 overexpression is not sufficient to induce microvillar actin overgrowth. This is consistent with the observation that GV oocytes, in which cofilin is largely inhibited, do not show microvillar overgrowth and retain a dense actin mesh.

Consistent with a requirement for actin monomer supply, sequestration of G-actin with latrunculin A (0.3 µM; Coué et al., 1987) prevented microvilli overgrowth in LIMK1-expressing MI oocytes, suggesting that microvillar actin elongation proceeds via monomer addition at filament ends (Fig. 4C,D; +LatA). We next tested the effect of a low dose of cytochalasin D (CD; 100 nM), in order to achieve barbed-end capping (Sampath and Pollard, 1991). However, microvilli elongation was largely resistant to CD treatment (Fig. 4C,D; +CD), arguing that barbed ends may be protected against capping. In this regard, formins exhibit anti-capping activity due to their persistent association with the growing barbed end (Chesarone et al., 2010; Shekhar et al., 2016). Accordingly, the pan-formin inhibitor SMIFH2 (30 µM; Rizvi et al., 2009) abolished microvilli outgrowth, while regular microvilli were preserved (Fig. 4C,D; +SMIFH2). In our search for a candidate formin, we observed that DAAM1, which is constitutively enriched at the oocyte cortex (Lu et al., 2017), accumulated at the distal tips of enlarged microvilli (Fig. S5A,D), possibly reflecting a role in filament elongation. DAAM1 also decorated the entire length of microvillar core bundles and actin rootlets (Fig. S5A), in line with its reported role in filament bundling (Jaiswal et al., 2013). In contrast, Fmn2, which is also enriched at the oocyte cortex (Pfender et al., 2011), did not accumulate along the length nor the tips of elongated microvilli (Fig. S5B).

The fact that prophase oocytes did not show excessive microvilli nor actin rootlet elongation (Fig. 2A) despite a substantial phosphorylation of their cofilin pool (Fig. S2), is suggestive of an additional regulatory mechanism related to cell cycle resumption. To test this idea, LIMK1-injected GV oocytes arrested in prophase by milrinone were treated with the PP1 and PP2A inhibitor okadaic acid (OA, 2.5 µM) in order to accelerate substrate phosphorylation

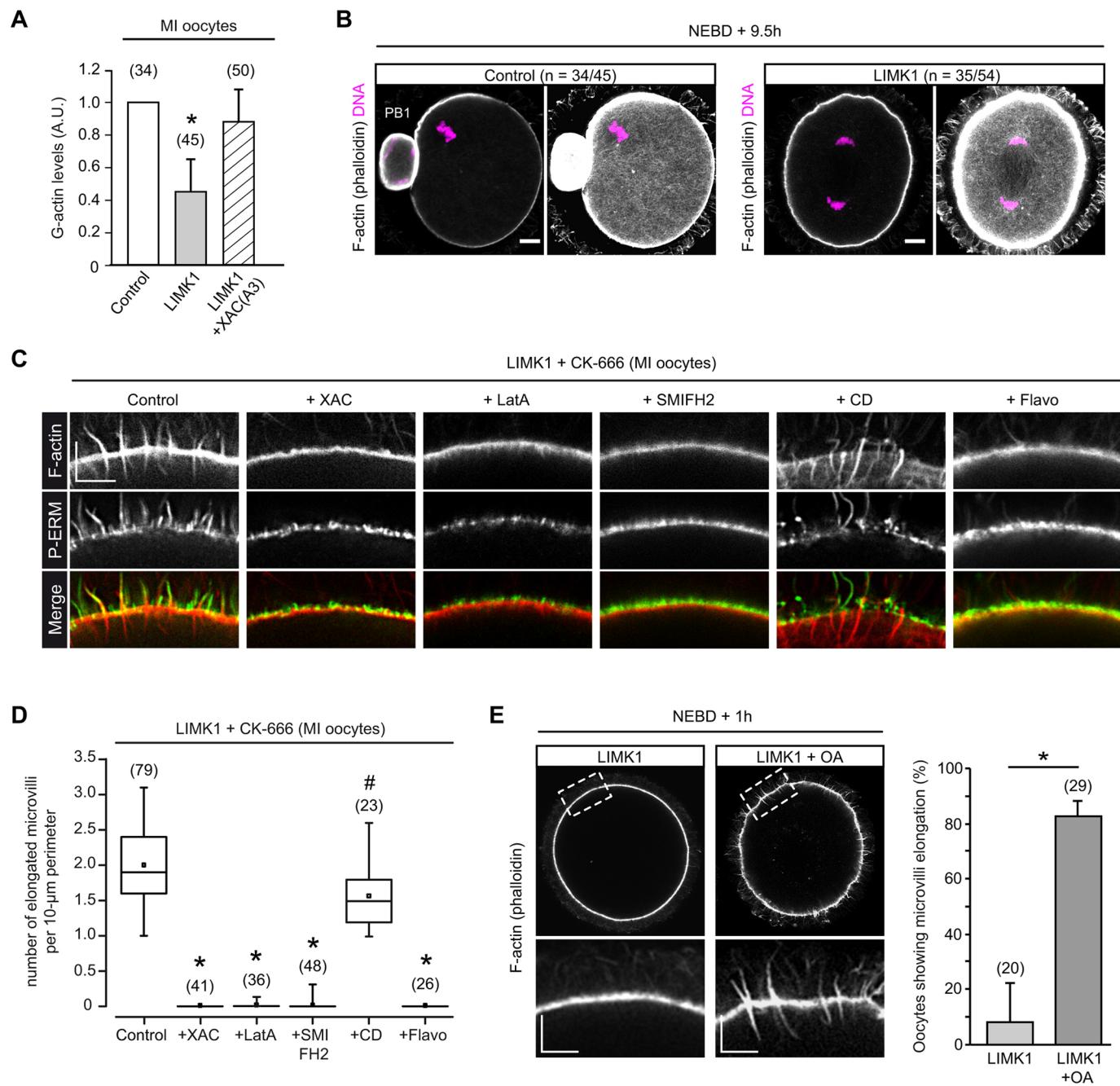


Fig. 4. Microvilli elongation requires G-actin, formin and CDK1 activities. (A) Control oocytes, or oocytes expressing LIMK1 alone or in combination with XAC(A3) were fixed at MI (NEBD+6 h) and processed for DNase I staining to label G-actin. The number of oocytes examined is indicated in parentheses. Data are mean \pm s.d. from four independent experiments. *P<10⁻⁴ compared to control (Student's *t*-test). (B) Chromosome (DNA; To-Pro-3, magenta) and F-actin (Alexa Fluor 568–phalloidin) staining in oocytes fixed at NEBD+9.5 h. Left: control oocyte shortly after polar body (PB1) extrusion. Right: oocyte injected with LIMK1 cRNA at the MI stage (NEBD+6 h). In both panels, F-actin staining is shown with regular settings (left image) or enhanced brightness for better visualization of the cytoplasmic F-actin mesh (right image). The fraction of examined oocytes showing a similar pattern is indicated in parentheses. Scale bars: 10 μ m. (C) Effects of various treatments on microvilli elongation. All images are expanded views of the cortical region of LIMK1-expressing MI oocytes treated with CK-666, stained for F-actin (Alexa Fluor 568–phalloidin, red) and P-ERM (green). XAC, constitutively active XAC(A3); LatA, latrunculin A (0.3 μ M); SMIFH2, SMIFH2 (30 μ M); CD, cytochalasin D (100 nM); Flavo, flavopiridol (5 μ M). The scale bar is 5 μ m and applies to all images. (D) Quantification of the density of elongated microvilli under the conditions shown in C. The number of oocytes examined for each experimental condition is indicated in parentheses. The mean (square), median (line), 25th and 75th percentiles (box), and 5th and 95th percentiles (whiskers) are shown. Statistical significance was calculated against the control condition (LIMK1+CK-666). *P<10⁻⁴; #P=0.001 (Student's *t*-test). (E) F-actin staining (Alexa Fluor 568–phalloidin) in LIMK1-expressing oocytes treated with DMSO (LIMK1) or 2.5 μ M okadaic acid (LIMK1+OA). Oocytes were injected with LIMK1 cRNA at the GV stage and cultured *in vitro* to NEBD+1 h before fixation. Lower panels are expanded views of the cortical regions delineated by the dashed boxes. Scale bars: 5 μ m. The bar graph shows the percentage of oocytes showing microvilli elongation. Data are mean \pm s.d. from three independent experiments. The number of oocytes examined is indicated in parentheses. *P=0.001 (Student's *t*-test).

by CDK1, thereby forcing oocytes into meiosis resumption (Rime and Ozon, 1990). Remarkably, OA treatment accelerated microvilli and actin rootlet overgrowth, which became detectable as early as 1 h after NEBD, at which time vehicle-treated LIMK1-expressing oocytes did not yet show any sign of overgrowth (Fig. 4E). To establish a specific requirement for CDK1, the CDK1 inhibitor flavopiridol (5 μ M; Wei et al., 2018) was added to the culture medium 1 h after NEBD, and oocytes were further cultured until MI (NEBD+6 h). CDK1 inhibition effectively abrogated microvillar actin overgrowth in LIMK1-expressing oocytes (Fig. 4C,D; +Flavo). Regular microvilli were still detectable, however, suggesting that the lack of microvilli elongation was not due to an intrinsic defect in microvillar integrity.

Taken together, these data suggest that microvilli overgrowth relies on a supply of free actin monomers, to which the collapse of the F-actin mesh is likely to contribute. However, the microvillar elongation phenotype is not the mere consequence of cofilin inactivation; it also relies on CDK1 substrate phosphorylation and may involve a formin-dependent elongation machinery. This outgrowth mechanism appears to build upon regular microvilli, which seem otherwise unaffected by CDK1 or formin inhibition.

Cofilin controls F-actin dynamics and the size of the actin cap in MII oocytes

We next explored the role of cofilin in regulating steady-state F-actin dynamics in oocytes that have achieved maturation and arrested at MII. Freshly ovulated MII oocytes were injected with LIMK1 cRNA for acute cofilin inhibition, as reflected in an increase in the phospho-cofilin signal (Fig. S2, right). After 2–3 h of culture, LIMK1-expressing MII oocytes exhibited a dramatic overgrowth of the actin cap, reaching a depth of up to 15 μ m into the inner cytoplasm (Fig. 5A–C). Remarkably, cytoplasmic material was excluded from this actin-filled volume, consistent with a dense network with a small mesh size (Fig. 5B, asterisk). Whereas cortical actin was moderately thickened in the rest of the cortex, there was no sign of microvilli overgrowth (Fig. 5B,D). In contrast, catalytically inactive LIMK1^{D460A} did not alter cortical actin dynamics, in line with the lack of cofilin phosphorylation (Fig. 5C,D; Fig. S2).

As a consequence of actin cap expansion, the MII spindle was distanced from the cortex, yet we never observed spindle detachment (Fig. 5B). Intriguingly, elongated spindle microtubules appeared to infiltrate the oversized actin cap, possibly contributing to a spindle–actin anchoring mechanism (Fig. 5B, arrow). The spindles also appeared highly focused at the poles (Fig. 5B), a feature previously reported in spindles artificially enriched in actin (Mogessie and Schuh, 2017). Accordingly, whereas spindle actin generated a faint signal in control oocytes, it was readily observable in LIMK1-injected MII oocytes, likely reflecting filament stabilization and/or thickening (Fig. 5A,B, lower panels).

We next tested whether the absence of microvilli overgrowth was due to actin monomers being diverted to the expanding actin cap. To this end, MII oocytes were injected with LIMK1 cRNA after prior treatment with CK-666 to inhibit Arp2/3. Consistent with previous observations (Yi et al., 2011), prolonged incubation with CK-666 (100 μ M, 4 h) induced a loss of the actin cap, along with spindle detachment from the cortex (Fig. 5E, left panels). Subsequent LIMK1 expression still failed to induce microvilli elongation (0/24 oocytes), while spindle actin was markedly increased and the cytoplasmic mesh was preserved (Fig. 5E, right panels). Therefore, the absence of microvilli elongation cannot be attributed to monomer sequestration into the oversized actin cap.

In order to appreciate the effects of cofilin inactivation on actin network dynamics, we monitored actin fluxes in live MII oocytes expressing EGFP-UtrCH. Consistent with previous studies (Yi et al., 2011; Mogessie and Schuh, 2017), control oocytes exhibited highly dynamic cytoplasmic and spindle actin networks, as well as a retrograde actin flow originating from the polarized cap (Movie 5). This retrograde actin flow, visualized as a transition zone of milder intensity pervading the subcortical cytoplasm, was readily captured by kymographic analysis, whereas cytoplasmic and spindle actin dynamics were too fast to be resolved (Fig. 5F,G). However, all three actin networks were successfully captured and quantified by spatiotemporal image correlation spectroscopy (STICS) analysis (Yi et al., 2011), revealing highest velocities in the actin cap transition zone and spindle actin (Fig. 5H). In stark contrast, actin networks were virtually static in LIMK1-injected MII oocytes (Movie 5), as reflected in the horizontal lines on the kymographs and a faint STICS signal (Fig. 5G,H). Live imaging also revealed that actin cap overgrowth resulted from the expansion of the transition zone, which remained segregated from the subjacent actin mesh by a sharp interface (Fig. 5F; Movie 5).

Collectively, these data provide three significant insights: (1) cofilin promotes global actin network turnover in mature oocytes and, in particular, restrains actin cap expansion, which is in contrast with the prevailing model by which cofilin amplifies the branching activity of Arp2/3 (Ichetovkin et al., 2002; Kiuchi et al., 2007); (2) in the absence of functional cofilin, the cytoplasmic F-actin network is frozen, in stark contrast with prophase oocytes where it remains highly dynamic, thus reinforcing the idea of distinct regulatory mechanisms in line with the cell cycle stage; and (3) in line with our observations in prophase oocytes and oocytes injected with LIMK1 after reaching the MI stage, cofilin inhibition, per se, is not sufficient to induce microvilli overgrowth in MII oocytes.

Actin cap disassembly promotes microvilli elongation

We reasoned that MII oocytes might experience microvilli elongation if the G-actin pool was raised to a critical threshold. To test this prediction, we treated MII oocytes with CK-666 (100 μ M) in order to promote spontaneous actin cap disassembly and monomer release. Oocytes were fixed after 3 h of CK-666 treatment, at which time the actin cap had not yet fully disassembled. Remarkably, CK-666 triggered the outgrowth of a tuft of elongated microvilli in the polarized cortex overlying the actin cap, a region that was otherwise devoid of microvilli (Fig. 6A,B,I). As shown above for LIMK1-expressing MI oocytes, these ectopic microvilli were positive for P-ERM (Fig. 6B) and CD9 (Movie 6), and were decorated along their length by DAAM1, which also accumulated at the tips (Fig. S5C,D). Accordingly, SMIFH2 prevented the outgrowth of these ectopic microvilli, while leaving regular microvilli unaffected (Fig. 6C,I). Likewise, monomer sequestration by latrunculin A treatment prevented ectopic microvilli elongation (Fig. 6D,I). Moreover, whereas CD (100 nM) effectively collapsed regular microvilli, as evidenced by a substantial loss of P-ERM staining (Fig. 6E, arrow), it did not prevent the outgrowth of ectopic microvilli in response to CK-666 (Fig. 6E,I). In fact, exposure to CD alone resulted in a similar burst of ectopic microvilli, while the regular microvilli were depleted (Fig. 6F,I). As stated above, CD (100 nM) is expected to act as a barbed-end capper, suggesting that the collapse of regular microvilli reflected unbalanced pointed-end disassembly. Accordingly, injection of LIMK1 cRNA shortly (1 h) before CK-666 addition – such as to retain residual cofilin activity for actin cap disassembly – resulted in further elongation of ectopic actin rootlets (Fig. 6G).

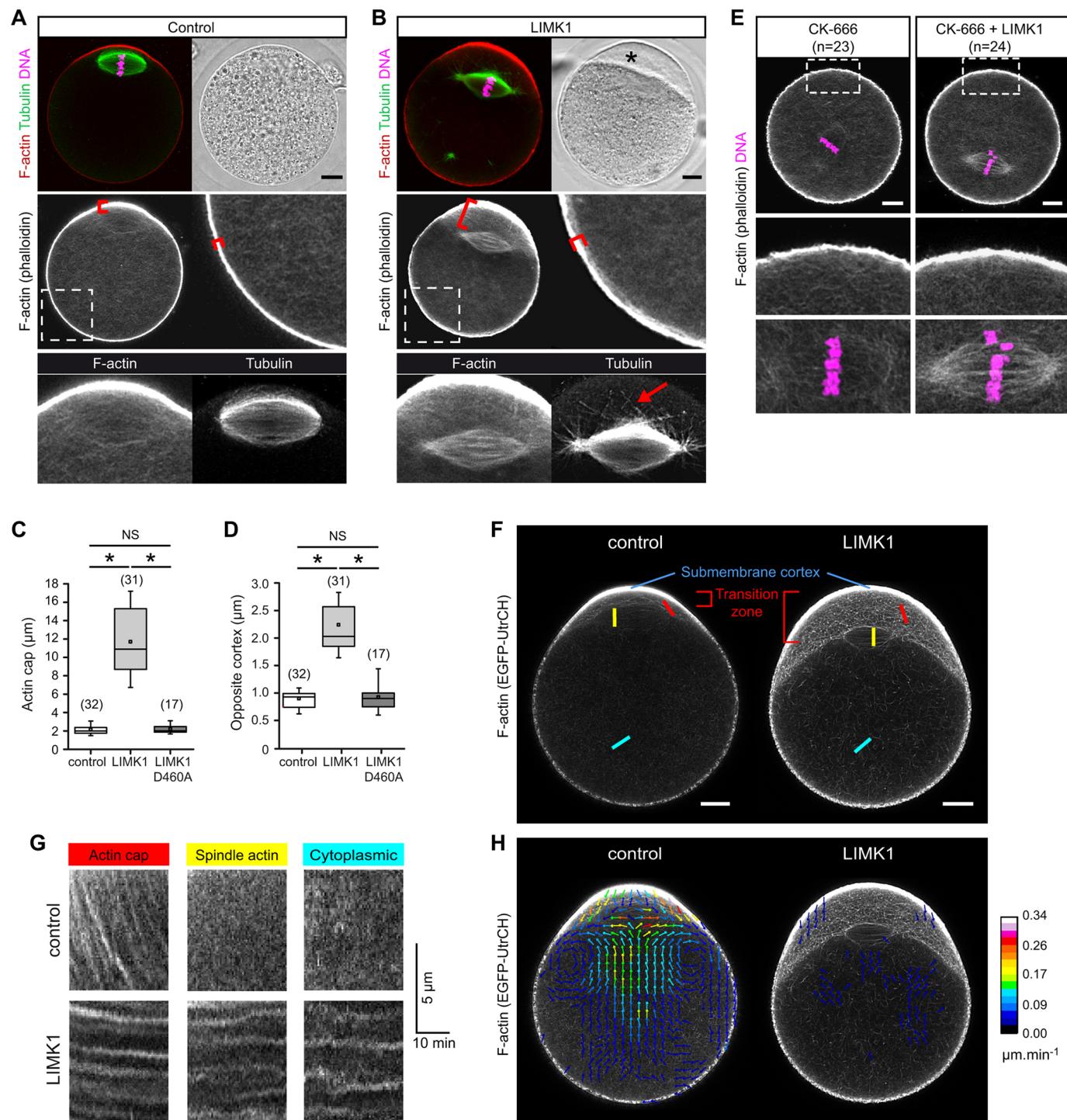


Fig. 5. See next page for legend.

Of note, upon extending the CK-666 treatment to 5 h, ectopic microvilli were no longer detected in oocytes where the actin cap had fully disassembled, and the spindle relocated to the oocyte center (Fig. 6H,I). This observation suggests that ectopic microvilli are transient structures that rely on actin monomers supplied by the disassembling actin cap. Taken together, these results suggest that an increase in G-actin supply can elicit microvilli elongation in MII oocytes, with similar regulatory properties as the elongated microvilli depicted in MI oocytes overexpressing LIMK1. Further

elongation of ectopic actin rootlets upon LIMK1 expression provides compelling evidence that cofilin regulates the length of microvillar actin filaments in oocytes.

DISCUSSION

In this study, we show that meiosis resumption involves a switch-like activation of cofilin, which is instrumental in regulating actin network size and dynamics, for successful oocyte maturation (Fig. 7, left). Considering its universal role in actin homeostasis, our

Fig. 5. Cofilin controls actin network dynamics in MII oocytes.

(A,B) Staining of F-actin and spindle microtubules in a control MII oocyte (A) and a LIMK1-injected MII oocyte (B). Upper panels show the brightfield and merge fluorescence images. Note the exclusion of cytoplasmic material in the region of actin cap expansion (asterisk in B). Red, F-actin (Alexa Fluor 568–phalloidin); green, anti-tubulin; magenta, chromosomes stained with TO-PRO-3 (DNA). Scale bars: 10 µm. Middle panels show the F-actin channel with enhanced brightness for better visualization of cytoplasmic and spindle F-actin. Magnified views of the non-polarized cortex (dashed boxes) are shown (middle right). Red brackets indicate the thickness of the actin cap (middle left) and opposite cortex (middle right). Lower panels are expanded views of the spindle region. Note the increased spindle actin signal and the microtubules projecting toward the expanded actin cap (arrow) in the LIMK1-expressing oocyte. (C,D) Quantification of the thickness of the actin cap (C) and the opposite cortex (D), as illustrated by the red brackets in A and B. The mean (square), median (line), 25th and 75th percentiles (box), and 5th and 95th percentiles (whiskers) are shown. The number of oocytes examined is indicated in parentheses. * $P<10^{-5}$; NS, not significantly different (Student's t-test). (E) F-actin staining (Alexa Fluor 568–phalloidin) in MII oocytes treated with CK-666 (100 µM, 4 h) alone (left), or followed by LIMK1 cRNA injection (right). Middle and bottom panels are expanded views of the cortical area (dashed boxes) and spindle area, respectively. The total number of oocytes examined is indicated in parentheses. Chromosomes are stained with TO-PRO-3 (DNA, magenta). Scale bars: 10 µm. (F) Visualization of F-actin networks in live MII oocytes expressing EGFP–UtrCH alone (left) or together with LIMK1 (right). The polarized actin cap is defined as a submembrane cortical layer of high intensity (submembrane cortex) and a transition zone of milder intensity corresponding to actin filaments flowing away from the cap. Red, yellow and cyan bars indicate sections of the actin cap, spindle actin and cytoplasmic actin, respectively, analyzed in kymographs in G. See also Movie 5. Scale bars: 10 µm. (G) Kymographic analysis of actin flow in the actin cap transition zone, spindle and cytoplasm, as depicted in F. (H) STICS analysis of F-actin dynamics in the oocytes shown in F. The color-coded vector maps were generated from the time-lapse recordings shown in Movie 5. The heat bar indicates filament velocity (µm min⁻¹). Data in F–H are representative of ten (control) and eight (LIMK1) similar observations.

finding that cofilin is largely inactivated in prophase oocytes is intriguing, and it suggests that alternative mechanisms exist to achieve filament turnover. Likewise, *Xenopus* oocytes have been reported to contain mostly phosphorylated cofilin (Abe et al., 1996; Iwase et al., 2013). Cofilin activation at meiosis resumption may involve the upregulation of cofilin phosphatases, such as members of the Slingshot family (Kaji et al., 2003; Iwase et al., 2013).

In contrast to other actin-rich processes, such as lamellipodia and filopodia, for which regulatory mechanisms have been extensively described, microvillar actin nucleation and dynamics are still poorly understood (Crawley et al., 2014; Pelaseyed and Bretscher, 2018). Except for their localized disassembly during cortical polarization (Maro et al., 1986; Van Blerkom and Bell, 1986; Dehapiot and Halet, 2013), mouse oocyte microvilli exhibit little change in size throughout meiotic maturation (Benamar et al., 2017); however, our data demonstrate that they remain dynamic structures amenable to extensive growth in response to actin homeostasis perturbation.

We document a striking elongation of microvilli, and a reciprocal loss of the F-actin mesh, in cofilin-inactivated MI oocytes injected with LIMK1 cRNA at the GV stage. However, our data in GV- and MII-arrested oocytes demonstrate that cofilin inactivation alone is not sufficient to trigger microvilli elongation, nor F-actin mesh breakdown. This is corroborated by the absence of microvillar elongation or mesh disassembly when LIMK1 is injected at the MI stage. The microvillar elongation phenotype seems, therefore, to arise specifically when cofilin is inactivated from meiosis resumption onward. This is precisely the time when the

cytoplasmic F-actin mesh experiences a substantial decrease in density, followed by a slow recovery (Azoury et al., 2011; Holubcová et al., 2013; Wei et al., 2018). Thus, one possible scenario is that the release of actin monomers upon mesh disassembly acts as a cue in driving microvillar actin elongation in cofilin-inactivated oocytes. In line with this idea, microvilli elongation following an increase in G-actin supply has been reported in isolated brush borders (Mooseker et al., 1982; Stidwill and Burgess, 1986). Interestingly, computational modeling of actin-rich protrusion dynamics has predicted that decreased severing at the base of microvilli would eventually increase the steady-state length of both the rootlet and the microvilli protrusion (Orly et al., 2014), which is corroborated by our experimental data. Furthermore, an increase in G-actin has been shown to promote actin filament growth via a direct stimulation of formin activity (Higashida et al., 2013). The synergistic effect of CK-666 may reflect an additional supply of monomers, via the disassembly of an Arp2/3-dependent cortical actin layer (Chaigne et al., 2013). Alternatively, the Arp2/3 complex may restrain microvilli elongation via a direct interaction with a formin machinery, as has previously been shown for mDia2 (DIAPH3)-dependent filopodia formation (Beli et al., 2008). However, it is unknown whether such inhibitory complexes are disassembled by CK-666. While the formin DAAM1 appears as a likely candidate in our system, its role in microvilli elongation remains to be formally established. Furthermore, off-target effects have recently been reported for SMIFH2, and hence data using this compound should be interpreted with caution (Isogai et al., 2015; Nishimura et al., 2021). Other types of elongation factors may contribute to the observed microvilli overgrowth, such as IRTKS (also known as BAIAP2L1) and EPS8, whose roles in microvilli elongation have recently been characterized in epithelial cells (Postema et al., 2018).

Microvilli outgrowth following actin cap disassembly in MII oocytes provides further evidence for microvilli length and/or biogenesis being regulated by G-actin availability. Likewise, a recent study documented epithelial microvilli elongation upon Arp2/3 inhibition in differentiating enterocytes (Faust et al., 2019). It is remarkable that ectopic microvilli in mouse oocytes were restricted to the otherwise amicrovillar polarized cortex. Treatment with CK-666 is expected to release actin monomers locally (Burke et al., 2014; Suarez and Kovar, 2016). Hence, we suggest that, by preventing monomer recycling into new or existing branches (Vitriol et al., 2015), CK-666 and CD both contribute to increasing the G-actin pool locally, leading to microvilli outgrowth in the nearby cortex. In turn, it is tempting to assume that the regulated disassembly of microvilli in the polarized cortex proceeds via a competition for monomers between the Arp2/3 complex and the microvillar machinery, at the expense of the latter. The growth of corresponding ectopic actin rootlets after additional LIMK1 expression provides clear evidence for cofilin regulating microvilli length in oocytes.

We therefore propose that the switch-like activation of cofilin at meiosis resumption provides a means for maintaining the stationary length of microvilli, through pointed-end depolymerization of the core actin filaments, thereby replenishing the cytoplasmic monomer pool. In this model (Fig. 7, right), the failure to activate cofilin leads to sequestration of G-actin into microvilli, acting as a sink for monomers, at the expense of the cytoplasmic F-actin mesh, which fails to reassemble. As a result, the spindle cannot migrate toward the cortex to achieve asymmetric division. Later during maturation, once the F-actin mesh has been re-established, cofilin appears to act mainly as a regulator of actin network dynamics, with no impact on

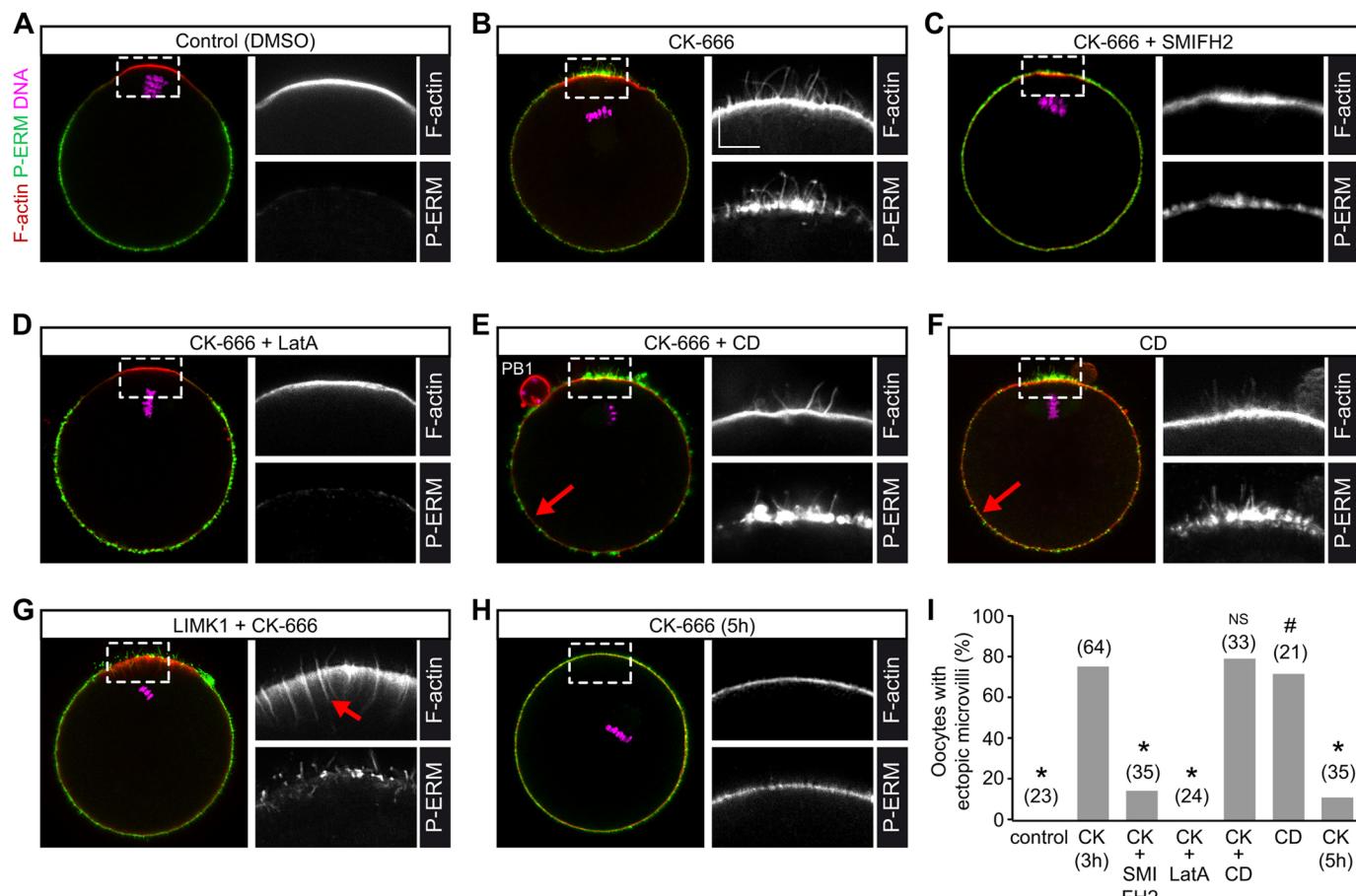


Fig. 6. Actin cap disassembly promotes microvilli outgrowth. MII oocytes were fixed and stained for F-actin (Alexa Fluor 568-phalloidin, red), P-ERM (green) and chromosomes (DNA; TO-PRO-3, magenta). Magnified views of the F-actin and P-ERM signals in the polarized cortex (dashed boxes) are shown. Oocyte treatments were as follows: (A) 0.1% DMSO for 3 h, (B) CK-666 (100 nM) for 3 h, (C) CK-666 (100 nM) and SMIFH2 (30 μM) for 3 h, (D) CK-666 (100 nM) and latrunculin A (LatA; 0.3 μM) for 3 h, (E) CK-666 (100 nM) and cytochalasin D (CD; 100 nM) for 2 h, (F) CD (100 nM) for 1 h, (G) LIMK1 expression and CK-666 (100 nM) for 3 h, and (H) CK-666 (100 nM) for 5 h. All images are z-compressions of three consecutive 1 μm-thick confocal sections. The scale bar in B (F-actin) is 5 μm and applies to all magnified images. In E, treatment was for 2 h only, as CD accelerated the loss of the actin cap. Note the large gaps in P-ERM staining in the non-polarized cortex of CD-treated oocytes (arrows in E,F), reflecting the collapse of regular microvilli. PB1, first polar body. In G, CK-666 was added 1 h after LIMK1 cRNA injection in order to retain residual cofilin activity. Note the cytoplasmic actin rootlets extending from the ectopic microvilli (arrow in G). (I) Rate of ectopic microvilli outgrowth in MII oocytes under the experimental conditions depicted in A–F,H (CK, CK-666). The bars indicate the fraction (%) of oocytes showing ectopic microvilli. The number of oocytes examined is indicated in parentheses. *P<10⁻⁴ compared to CK (3 h); NS, not significantly different compared to CK (3 h); #P=0.01 compared to control (Student's t-test).

microvilli length, unless the G-actin pool is artificially raised via the disassembly of a competing actin network.

Another intriguing finding of this study is that prophase oocytes retain normal cytoplasmic F-actin dynamics despite cofilin inactivation. To explain this discrepancy, we suggest that cytoplasmic F-actin turnover during prophase arrest may rely instead on depolymerization from the barbed end. Indeed, it has been proposed that at one stage of their life cycle, cytoplasmic actin filaments release Fmn2, which may leave the barbed ends unprotected from depolymerization (Montaville et al., 2014). The same scenario could explain the collapse of the cytoplasmic F-actin mesh at NEBD, as this coincides with the regulated destruction of Fmn2 by the proteasome (Azoury et al., 2011). In support of this hypothesis, a truncated Cappuccino (the closest *Drosophila* homolog to Fmn2) has been shown to stabilize the fly oocyte actin mesh against disassembly upon latrunculin A treatment, which implies that depolymerization occurs at the barbed ends (Dahlgaard et al., 2007). Other actin-binding proteins may further accelerate barbed-end depolymerization, such as profilin and the cofilin-

related depolymerizing factor twinfilin (Montaville et al., 2014; Pernier et al., 2016; Hilton et al., 2018; Shekhar et al., 2019). Further investigations are required to verify this hypothesis and establish the molecular basis of the differential regulation of cytoplasmic F-actin during oocyte meiosis.

Notwithstanding subtle changes in morphology and density, microvilli decorate the oocyte surface across all stages of maturation, suggesting that core microvillar components are little affected by the cell cycle (Longo and Chen, 1985; Benamarra et al., 2017). Yet, by interfering with cofilin activation, we uncovered a pathway for microvilli elongation that relies on CDK1 activity, suggesting that microvillar actin dynamics are altered during M phase. Although the relevant CDK1 substrates remain to be characterized, a number of actin regulatory proteins have been identified as putative CDK1 targets in mitotic cells, including various formins and cordon-bleu, a candidate actin nucleator in intestinal microvilli (Dephoure et al., 2008; Grega-Larson et al., 2015). Likewise, CDK1 has been shown to promote actin cable formation by the yeast formin Bni1 and mammalian mDia2

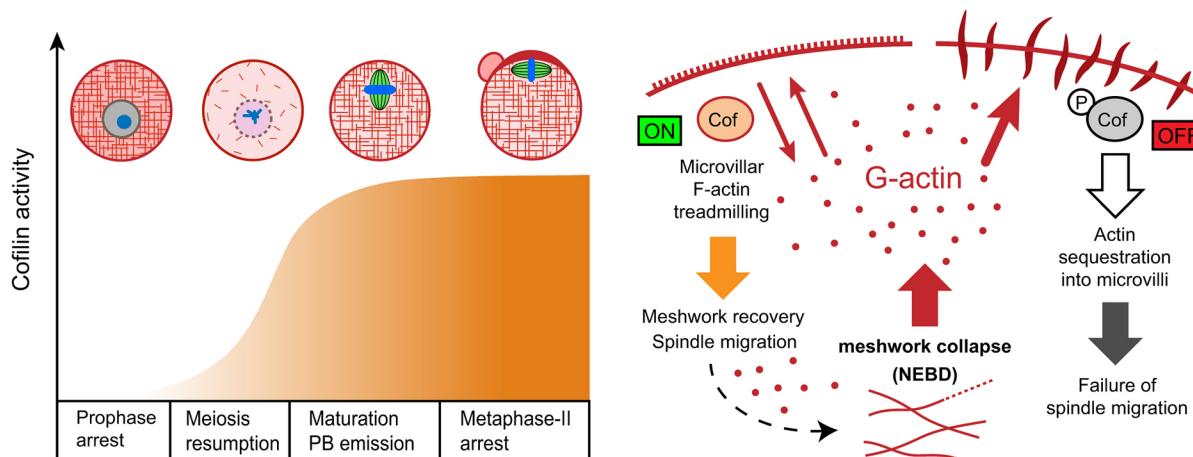


Fig. 7. Model depicting the regulation of actin homeostasis by cofilin. Left: theoretical time course of cofilin activation during oocyte maturation. Actin networks are shown in red. PB, polar body. Right: model depicting how cofilin (Cof) inactivation (phosphorylation, P) leads to microvilli elongation and a loss of the cytoplasmic F-actin mesh. The collapse of the F-actin mesh at meiosis resumption is assumed to release free actin monomers (G-actin, red dots) that are captured by the barbed ends of microvillar actin filaments. Concurrently, cofilin activation (left side; cofilin 'ON') accelerates pointed-end disassembly, thereby maintaining the stationary length of microvilli through treadmilling. In doing so, cofilin also maintains the cytoplasmic G-actin pool for recovery of the F-actin mesh later during maturation, enabling spindle migration. Inactivation of cofilin by phosphorylation (right side; cofilin 'OFF') prevents pointed-end depolymerization, resulting in exaggerated growth of microvillar core bundles, acting as a sink for monomers. As a result, the actin mesh is not rebuilt and spindle migration fails. Microvilli elongation requires CDK1 and formin activities (not depicted).

(DIAPH3) (Miao et al., 2013). Changes in microvilli properties during oocyte maturation (e.g. accelerated treadmilling) may be a prerequisite for achieving microvilli breakdown in the polarized cortex (Dehapiot and Halet, 2013), and/or for productive interaction with the fertilizing sperm (Runge et al., 2007). In this regard, oocyte-derived exosomes, which are assumed to be of microvillar origin, are found in the perivitelline space of MII oocytes but not of GV oocytes, consistent with a change in microvillar dynamics after meiosis resumption (Miyado et al., 2008; Benammar et al., 2017).

Despite a substantial activation of their cofilin pool, MII oocytes do not experience catastrophic disassembly of their actin networks, pointing to additional mechanisms to fine-tune cofilin activity. Two candidates are AIP1 and coronin, which synergize with cofilin to accelerate filament severing and disassembly while inhibiting elongation of the severed fragments (Gressin et al., 2015; Jansen et al., 2015). Although AIP1 (also known as WDR1) has been identified in mouse oocytes, it appears to be dispensable for normal oocyte function (Xiao et al., 2017; Jin et al., 2020). CAP1 (also known as Srv2) is another highly conserved regulator of actin turnover, acting synergistically with cofilin (Kotila et al., 2019; Shekhar et al., 2019). Accordingly, CAP1 knockdown in mouse oocytes is associated with defects in cytoplasmic F-actin, spindle migration and cytokinesis; however, no microvillar outgrowth nor actin cap expansion has been reported (Jin et al., 2018). Thus, the multiple actin networks that populate the oocyte may be differentially regulated by cofilin through the assembly of distinct multi-component machineries. Additional studies are needed to further characterize these accessory proteins, as well as alternative depolymerizing factors (for example, twinfilin), and their regulation across the meiotic cell cycle. While oocytes provide an attractive model for microvilli plasticity and the interplay between actin dynamics and the cell cycle, it will be meaningful to investigate whether the phospho-regulation of cofilin regulates microvilli length in other cells, such as differentiating epithelial cells (Faust et al., 2019) and cells that utilize microvilli as a membrane reservoir (Figard et al., 2016).

MATERIALS AND METHODS

Mice

All animal procedures were conducted in accordance with the European directive for the use and care of laboratory animals (2010/63/EU) and were approved by the local animal ethics committee under the French Ministry of Higher Education, Research and Innovation (Project licence APAFIS#11761). Mice of the MF1 strain were initially purchased from Envigo (Gannat, France) and were maintained as a colony in the local animal facility. To minimize the number of animals, female mice (6–8 weeks old) were primed by intraperitoneal injection of 5–7 units of PMSG (Chronogest, MSD) 48 h prior to GV oocyte collection. To obtain MII oocytes, mice were injected with 5 IU of hCG (Chorulon, MSD) 48 h after the PMSG injection.

Oocyte collection and culture

For GV-stage oocytes, cumulus–oocyte complexes were recovered by puncturing ovarian follicles with a needle and were transferred into M2 medium (M7167, Sigma) supplemented with 2 µM milrinone (Sigma) to maintain prophase arrest. Fully-grown GV-intact oocytes were denuded of surrounding cumulus cells by mouth pipetting. To obtain oocytes at various stages of maturation, GV oocytes were washed twice in milrinone-free M16 medium (M7292, Sigma) to trigger spontaneous meiosis resumption and were then cultured in M16 in an atmosphere of 5% CO₂ in air at 37°C, until the desired stage. MII oocytes were recovered from the oviduct in M2 medium supplemented with 3 mg/ml hyaluronidase (Sigma). The following small molecule inhibitors were used at the indicated concentrations: nocodazole (5 µM; Merck Millipore), CK-666 (100 µM; Sigma), okadaic acid (2.5 µM; Enzo), cytochalasin D (100 nM; Tocris), latrunculin A (0.3 µM; Enzo), SMIFH2 (30 µM; Sigma), flavopiridol (5 µM; Selleckchem). An equivalent amount of DMSO was used for controls. Note that in experiments involving SMIFH2, polyvinyl alcohol (PVA; 0.05%) was substituted for bovine serum albumin (BSA) in the culture medium.

Cloning and mutagenesis of mouse LIMK1

Total RNA was extracted from 112 GV oocytes using a Nucleospin RNA XS Kit (Macherey-Nagel), and first-strand cDNA was generated with SuperScript III Reverse Transcriptase (Thermo Fisher Scientific, 18080-044) and RNaseOUT (Life Technologies), using oligo(dT)15 primers (Promega, C110A). The following primers were used to amplify the full length of LIMK1 cDNA by PCR: forward, 5'-ATGAGGTTGACGCTACTTTGT-3';

reverse, 5'-TCAGTCAGGGACCTCGGGGTG-3'. The LIMK1 fragment was next cloned into pcDNA3.1(+) (Invitrogen), and the construct was validated by sequencing. Catalytically inactive LIMK1^{D460A}, in which Asp460 is replaced by Ala (Yang et al., 1998), was generated by site-directed mutagenesis using the QuickChange II Kit (Stratagene) and then verified by sequencing.

Plasmids, cRNA preparation and microinjection

Constitutively active *Xenopus* ADF/Cofilin XAC(A3) in pSP36T was obtained from Hiroshi Abe (Chiba University, Chiba, Japan) and subcloned into pcDNA3.1(+) for cRNA synthesis. EGFP-UtrCH in pCS2+ was from Addgene (Addgene plasmid #26737, deposited by William Bement; Burkel et al., 2007). Fmn2-mCherry in pCS2 was obtained from Jan Ellenberg (Euroscarf plasmid #P30607; Schuh and Ellenberg, 2008); mCherry was exchanged with EGFP in order to generate pCS2-Fmn2-EGFP. Polyadenylated cRNAs were synthesized *in vitro* from linearized plasmids using the mMessage mMachine T7 or SP6 kit and Poly(A) Tailing kit (Ambion), purified with an RNeasy purification kit (Qiagen), and stored at -80°C. Oocytes were injected with ~5 pl of the cRNA solution (1.5–3.5 µg/µl) using the electrical-assisted microinjection technique, which allows for a high rate of oocyte survival (Fitzharris et al., 2018). For co-expression of LIMK1 with XAC(A3), the cRNA encoding XAC(A3) was co-injected as a 1:1 (volume) mix with 2× concentrated LIMK1 cRNA in order to reach a LIMK1 expression level similar to oocytes injected with LIMK1 only.

RNAi knockdown of cofilin1 transcripts

Transcripts encoding cofilin1 (*Cfl1*), the most abundant cofilin isoform expressed in oocytes (Pfender et al., 2015), were knocked down via cytoplasmic siRNA injection. Denuded prophase oocytes were microinjected with 5–10 pl of ON-TARGETplus mouse cofilin1 SMARTpool siRNA (Dharmacon, L-058638-01-0005; Horizon Discovery) diluted in RNase-free water to a stock concentration of 100 µM. After *in vitro* culture in M16 medium for 22 h, during which oocytes were maintained arrested in prophase with milrinone, total RNA was isolated from 15 oocytes using the NucleoSpin RNA XS kit (Macherey-Nagel) and processed for reverse transcription using oligo(dT)15 primers (Promega, C110A) and SuperScript III reverse transcriptase (Thermo Fisher Scientific, 18080-044). The extent of cofilin1 transcript knockdown was assessed by quantitative real-time PCR performed on a QuantStudio 7 Flex system (Thermo Fisher Scientific, 4485701), using the Power SYBR Green PCR master mix (Thermo Fisher Scientific, 4367659). Detection of β-actin (*Actb*) transcripts was used as an internal control for normalization. Oligo pairs were as follows: cofilin1 forward, 5'-CAGACAAGGACT-GCCGCTAT-3'; cofilin1 reverse, 5'-TTGCTCTTGAGGGGTGCATT-3'; β-actin forward, 5'-AGCGGTTCCGATGCC-3'; β-actin reverse, 5'-CTTACGGATGTCAACGTCACAC-3'.

Western blot analysis

To detect endogenous cofilin and phospho-cofilin, 50–100 oocytes were harvested at the indicated stage of maturation, washed in phosphate-buffered saline (PBS) supplemented with 1% polyvinylpyrrolidone (Sigma) and lysed in Laemmli 2× Concentrate Sample Buffer (Sigma). Lysates were immediately frozen at -80°C until use. After thawing and denaturation at 95°C for 5 min, proteins were separated by SDS-PAGE on NuPAGE Novex 4–12% Bis-Tris gels (Life Technologies) and transferred onto PVDF membranes (Amersham) in Tris-glycine containing 10% ethanol. Membranes were blocked with PBST (PBS containing 0.1% Tween 20) supplemented with 5% BSA or 5% non-fat dry milk for 3 h at room temperature, and then incubated with primary antibodies at 4°C overnight. After washing with PBST three times, membranes were incubated with secondary antibodies conjugated to horseradish peroxidase for 1 h at room temperature, followed by washing with PBST three times. Blots were developed using SuperSignal West DURA Extended Duration Substrate (Thermo Fisher Scientific) following the manufacturer's instructions. Semi-quantitative densitometric analysis of band intensities was performed using the 'Gels' command in FIJI open-source software (<https://fiji.sc/>). The

following primary antibodies were used: rabbit anti-cofilin (Cytoskeleton ACFL02; 1:500), rabbit anti-Ser3-phosphorylated cofilin (Cell Signaling Technology #3311; 1:500) and a mouse anti-GAPDH (Abcam ab8245; 1:5000) used as a control for protein loading. Secondary antibodies were HRP-conjugated goat anti-rabbit IgG (G-21234; 1:25,000; Life Technologies) and goat anti-mouse IgG (G-21040; 1:25,000; Life Technologies). Uncropped images of the western blot data are provided in Fig. S6.

Immunofluorescence

Oocytes were fixed for 16 min at room temperature with 3% paraformaldehyde in PBS, freshly prepared from a 16% methanol-free paraformaldehyde solution (Electron Microscopy Sciences). Fixed oocytes were permeabilized with 0.25% Triton X-100 (Sigma) in PBS for 15 min, blocked with 3% BSA (Sigma) in PBS for 3 h, and incubated overnight at 4°C with primary antibodies diluted in PBS containing 3% BSA. On the next day, oocytes were washed in PBS containing 1% BSA and then were incubated with secondary antibodies diluted in PBS containing 1% BSA for 45 min at 37°C. The following primary antibodies were used at the indicated dilutions: anti-tubulin (rat monoclonal; Abcam, ab6161; 1:200), anti-phospho-ERM (rabbit monoclonal; Cell Signaling Technology, 3726; 1:100), anti-CD9 (rat monoclonal; Santa Cruz, sc-18869; 1:100), anti-DAAM1 (goat polyclonal; Santa Cruz, sc-55929; 1:100). Secondary antibodies were Alexa Fluor 488-conjugated donkey anti-goat, donkey anti-rabbit and goat anti-rat (all 1:1000; Invitrogen). Chromatin was stained with To-Pro-3 (Invitrogen, T3605).

Chromosome staining for monitoring anaphase I

To evaluate the kinetics of anaphase I, GV-stage oocytes (uninjected, or expressing LIMK1 or LIMK1^{D460A}) were washed free from milrinone and allowed to resume meiosis and mature *in vitro* in a small drop of M16 medium layered with mineral oil (Sigma, M8410). At 10 h after milrinone washout, oocytes were stained with Hoechst 33342 (5 µM; Invitrogen, H3570) for 10 min in M2 medium, followed by a wash, and were then transferred to glass-bottom dishes (MatTek, Ashland, MA). Oocytes were examined for chromosome configuration using an inverted Leica DMI4000B microscope equipped with a 365 nm LED module, a digital monochrome DFC345 FX camera (Leica) and a HCX PL S-APO 10× objective. Images were acquired using the LAS AF 2.3.0 software (Leica). Oocytes were then quickly returned to a 37°C hot block and imaged again at 11 h post milrinone washout.

Chromosome spreading

To assess homologous segregation, LIMK1-expressing oocytes that had been cultured overnight in M16 medium were processed for chromosome spreading based on a previously published protocol (Roberts et al., 2005). Briefly, oocytes were treated with a hypotonic solution (fresh 1% sodium citrate) for 5 min, then individually transferred into small (5 µl) drops of fixative (5:1:4 of methanol:acetic acid:water) on a glass-bottom dish (MatTek). After the zona pellucida had gradually dissolved, 10 µl of fixative (3:1 of methanol:acetic acid) was gently dropped onto the oocytes (repeated 2–3 times), followed by air-drying at room temperature. Chromosomes were labeled with Hoechst 33342 (10 µg/ml in water) for 30 min followed by a wash with water. Images of chromosome spreads were acquired using an inverted Leica DMI4000B microscope equipped with a 365 nm LED module, using a HCX PL S-APO 63× oil-immersion objective.

DNase I staining for detection of cytoplasmic G-actin

To assess the levels of G-actin in the cytoplasm, oocytes were stained with DNase I according to a previously published protocol (Yu et al., 2014), with minor modification. Briefly, oocytes were fixed in 4% paraformaldehyde in fixation buffer (130 mM KCl, 3 mM MgCl₂, 25 mM HEPES pH 7, 0.15% glutaraldehyde), for 30 min at room temperature. Fixed oocytes were permeabilized with 0.5% Triton X-100 for 15 min and blocked with 3% BSA in PBS for 2 h. Oocytes were next incubated with Alexa Fluor 488-conjugated DNase I (Thermo Fisher Scientific, D12371), diluted to 10 µg/ml in PBS containing 3% BSA, for 30 min at room

temperature, followed by wash in PBS containing 3% BSA. Confocal images were acquired (see below) and DNase I fluorescence was measured in a 10×10 μm cytoplasmic region, away from the GV or spindle area. To correct for background fluorescence, we performed Triton X-100 extraction on live oocytes before fixation (Kan et al., 2011) in order to remove all soluble G-actin. Thus, live oocytes were incubated for 10 min at room temperature in extraction buffer (100 mM KCl, 20 mM MgCl₂, 3 mM EGTA, 20 mM HEPES pH 6.8, 0.1% Triton X-100), washed in PBS and then processed for fixation as described above. At least five oocytes were extracted in each experimental series in order to calculate an averaged background fluorescence value, which was subtracted from raw values in non-extracted oocytes.

F-actin staining and quantification in fixed oocytes

F-actin was stained with Alexa Fluor 568–phalloidin (Life Technologies). The density of the cytoplasmic F-actin network in MI (NEBD+6 h) oocytes was evaluated by measuring the mean fluorescence intensity in a 10×10 μm region of interest using the Leica LAS AF Lite 2.6.0 software. Care was taken to avoid the spindle region and the elongated cytoplasmic actin rootlets. To calculate the density of elongated microvilli in LIMK1-expressing MI oocytes, a single confocal optical section (thickness 1 μm) obtained at the equator of the oocyte was analyzed. Elongated microvilli actin bundles were counted across the oocyte perimeter, and their density was expressed as per 10-μm perimeter. Only elongated structures showing bidirectional elongation (outward microvilli and inward rootlet) were considered for the analysis. In fixed MII oocytes, the thickness of the actin cap (defined as the distance spanning from the submembrane cortical actin to the surface of the spindle), and of the cortical actin layer in the opposite cortex, were measured offline using Leica LAS AF Lite 2.6.0.

Analysis of microvilli length with FiloQuant

We used the ImageJ plugin FiloQuant (Jacquemet et al., 2017) to detect elongated microvilli and measure their length. A single confocal section (thickness 1 μm) obtained at the equator of the oocyte, was processed using the single-image analysis mode, and outward microvilli were resolved as illustrated in Fig. S4. First, an unsharp filter was applied in FIJI to facilitate microvilli detection. Next, the oocyte contour was manually outlined, and the cytoplasmic region was filled using the ‘Fill’ function in order to allow for thresholding of the cell edge. Transzonal projections contaminating the F-actin signal were manually outlined and removed using the ‘Clear’ function. The resulting image was processed with Filoquant in the single-image mode. Threshold parameters for cell edge and microvilli detection were adjusted iteratively for best results. A lower cut-off at 1 μm was applied to minimize detection of irrelevant bright spots originating mostly from numerous microvilli fragments crossing the plane of the confocal frame (‘broken’ microvilli).

F-actin labeling and quantification in live oocytes

F-actin networks were labeled using EGFP-UtrCH expressed via cRNA microinjection. For GV and MI oocytes, cRNA injection was performed in GV oocytes arrested in prophase with milrinone, followed by culture for 3 h in M16 containing milrinone, to allow for protein expression. For MII oocytes, cRNA injection was followed by a 3 h culture in M16 for protein expression. In LIMK1-expressing oocytes, the cRNA encoding LIMK1 was co-injected as a 1:1 (volume) mix with a 2× concentrated EGFP–UtrCH cRNA in order to reach a similar expression level of the F-actin probe in LIMK1-injected oocytes and control oocytes. In some experiments, chromosomes were labeled with SiR-DNA (Spirochrome), which was added to the culture medium to a final concentration of 0.5 μM. Spatiotemporal image correlation spectroscopy (STICS) analysis was performed in FIJI, using a dedicated plugin written by Jay Unruh (Stowers Institute for Medical Research, Kansas City, MO; <https://research.stowers.org/imagejplugins/index.html>).

Confocal imaging, image processing and analysis

Oocytes were placed on glass-bottom dishes (MatTek, Ashland, MA) and imaged with a Leica SP5 or SP8 confocal microscope, using a 63× oil-

immersion objective. The thickness of optical sections was set at 1 μm. For live imaging, temperature was maintained at 37°C using a stage-top incubator (model INUBG2E-GSI; Tokai Hit, Shizuoka-ken, Japan) fitted on the microscope stage. Confocal settings were kept identical in control and experimental (e.g. LIMK1 overexpression or drug treatment) conditions. Confocal images and time-lapse movies were processed with FIJI. Image sharpness was enhanced by applying an unsharp mask filter in FIJI. Kymographs were generated with the ‘Multi Kymograph’ tool in FIJI. DAAM1 fluorescence profiles were generated using the ‘segmented line’ tool and the ‘Plot Profile’ function in FIJI. Fluorescence intensities (in arbitrary units, A.U.) were expressed as mean±s.d.

Statistical analysis

All experiments were repeated independently at least three times. Mean, standard deviation (s.d.) or standard error of the mean (s.e.m.), and statistical significance ($P<0.05$) were calculated using a two-tailed unpaired Student’s *t*-test or Mann–Whitney test in Origin (OriginLab). The number of oocytes examined and *P* values are indicated in the figures. Line graphs and bar graphs were generated in Excel or Origin. Box plots were generated in Origin and show the mean (square), median (line), 25th and 75th percentiles (box), and 5th and 95th percentiles (whiskers).

Acknowledgements

We thank Professor Hiroshi Abe for the XAC(A3) plasmid and Jay Unruh for advice with STICS analysis. We are grateful to the staff of the ARCHE-Biosit animal facility and MRIC-Biosit microscopy facility for technical assistance and expert advice.

Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: A.B., B.D., G.H.; Methodology: A.B., B.D., G.H.; Validation: G.H.; Formal analysis: A.B., B.D., G.H.; Investigation: A.B., B.D., G.H.; Data curation: G.H.; Writing - original draft: A.B., B.D., G.H.; Visualization: A.B., B.D., G.H.; Supervision: G.H.; Project administration: G.H.; Funding acquisition: G.H.

Funding

This work was supported by an ATIP fellowship to G.H. awarded by the Centre National de la Recherche Scientifique, and by a research grant awarded to G.H. by the Ligue Contre le Cancer Grand Ouest. G.H. wishes to thank the Society for Reproduction and Fertility (SRF) for the award of an Academic Scholarship. B.D. received a PhD scholarship from the Ministère de l’Enseignement Supérieur et de la Recherche and additional funding from the Fondation pour la Recherche Médicale.

Peer review history

The peer review history is available online at <https://journals.biologists.com/jcs/article-lookup/doi/10.1242/jcs.259237>.

References

- Abe, H., Obinata, T., Minamide, L. S. and Bamburg, J. R.** (1996). Xenopus laevis actin-depolymerizing factor/cofilin: a phosphorylation-regulated protein essential for development. *J. Cell Biol.* **132**, 871–885. doi:10.1083/jcb.132.5.871
- Amano, T., Kaji, N., Ohashi, K. and Mizuno, K.** (2002). Mitosis-specific activation of LIM motif-containing protein kinase and roles of cofilin phosphorylation and dephosphorylation in mitosis. *J. Biol. Chem.* **277**, 22093–22102. doi:10.1074/jbc.M201444200
- Antkowiak, A., Guillotin, A., Boiero Sanders, M., Colombo, J., Vincentelli, R. and Michelot, A.** (2019). Sizes of actin networks sharing a common environment are determined by the relative rates of assembly. *PLoS Biol.* **17**: e3000317. doi:10.1371/journal.pbio.3000317
- Arber, S., Barbayannis, F. A., Hanser, H., Schneider, C., Stanyon, C. A., Bernard, O. and Caroni, P.** (1998). Regulation of actin dynamics through phosphorylation of cofilin by LIM-kinase. *Nature* **393**, 805–809. doi:10.1038/31729
- Azoury, J., Lee, K. W., Georget, V., Rassinier, P., Leader, B. and Verlhac, M.-H.** (2008). Spindle positioning in mouse oocytes relies on a dynamic meshwork of actin filaments. *Curr. Biol.* **18**, 1514–1519. doi:10.1016/j.cub.2008.08.044
- Azoury, J., Lee, K. W., Georget, V., Hikal, P. and Verlhac, M.-H.** (2011). Symmetry breaking in mouse oocytes requires transient F-actin meshwork destabilization. *Development* **138**, 2903–2908. doi:10.1242/dev.060269
- Beli, P., Mascheroni, D., Xu, D. and Innocenti, M.** (2008). WAVE and Arp2/3 jointly inhibit filopodium formation by entering into a complex with mDia2. *Nat. Cell Biol.* **10**, 849–857. doi:10.1038/ncb1745

- Benamar, A., Ziyat, A., Lefèvre, B. and Wolf, J. P.** (2017). Tetraspanins and mouse oocyte microvilli related to fertilizing ability. *Reprod. Sci.* **24**, 1062-1069. doi:10.1177/1933719116678688
- Bierne, H., Gouin, E., Roux, P., Caroni, P., Yin, H. L. and Cossart, P.** (2001). A role for cofilin and LIM kinase in Listeria-induced phagocytosis. *J. Cell Biol.* **155**, 101-112. doi:10.1083/jcb.200104037
- Blanchoin, L., Pollard, T. D. and Mullins, R. D.** (2000). Interactions of ADF/cofilin, Arp2/3 complex, capping protein and profilin in remodeling of branched actin filament networks. *Curr. Biol.* **10**, 1273-1282. doi:10.1016/S0960-9822(00)00749-1
- Blanchoin, L., Boujemaa-Paterski, R., Sykes, C. and Plastino, J.** (2014). Actin dynamics, architecture, and mechanics in cell motility. *Physiol. Rev.* **94**, 235-263. doi:10.1152/physrev.00018.2013
- Burke, T. A., Christensen, J. R., Barone, E., Suarez, C., Sirotnik, V. and Kovar, D. R.** (2014). Homeostatic actin cytoskeleton networks are regulated by assembly factor competition for monomers. *Curr. Biol.* **24**, 579-585. doi:10.1016/j.cub.2014.01.072
- Burkel, B. M., von Dassow, G. and Bement, W. M.** (2007). Versatile fluorescent probes for actin filaments based on the actin-binding domain of utrophin. *Cell Motil. Cytoskeleton* **64**, 822-832. doi:10.1002/cm.20226
- Carlier, M.-F. and Shekhar, S.** (2017). Global treadmilling coordinates actin turnover and controls the size of actin networks. *Nat. Rev. Mol. Cell Biol.* **18**, 389-401. doi:10.1038/nrm.2016.172
- Chaigne, A., Campillo, C., Gov, N. S., Voituriez, R., Azoury, J., Umaña-Díaz, C., Almonacid, M., Queguiner, I., Nassoy, P., Sykes, C. et al.** (2013). A soft cortex is essential for asymmetric spindle positioning in mouse oocytes. *Nat. Cell Biol.* **15**, 958-966. doi:10.1038/ncb2799
- Chan, A. Y., Bailly, M., Zebda, N., Segall, J. E. and Condeelis, J. S.** (2000). Role of cofilin in epidermal growth factor-stimulated actin polymerization and lamellipod protrusion. *J. Cell Biol.* **148**, 531-542. doi:10.1083/jcb.148.3.531
- Chan, C., Beltzner, C. C. and Pollard, T. D.** (2009). Cofilin dissociates Arp2/3 complex and branches from actin filaments. *Curr. Biol.* **19**, 537-545. doi:10.1016/j.cub.2009.02.060
- Chesarone, M. A., DuPage, A. G. and Goode, B. L.** (2010). Unleashing formins to remodel the actin and microtubule cytoskeletons. *Nat. Rev. Mol. Cell. Biol.* **11**, 62-74. doi:10.1038/nrm2816
- Coué, M., Brenner, S. L., Spector, I. and Korn, E. D.** (1987). Inhibition of actin polymerization by latrunculin A. *FEBS Lett.* **213**, 316-318. doi:10.1016/0014-5793(87)81513-2
- Crawley, S. W., Mooseker, M. S. and Tyska, M. J.** (2014). Shaping the intestinal brush border. *J. Cell Biol.* **207**, 441-451. doi:10.1083/jcb.201407015
- Dahlgaard, K., Raposo, A. A., Niccoli, T. and St Johnston, D.** (2007). Capu and Spire assemble a cytoplasmic actin mesh that maintains microtubule organization in the Drosophila oocyte. *Dev. Cell* **13**: 539-553. doi:10.1016/j.devcel.2007.09.003
- Dehapiot, B. and Halet, G.** (2013). Ran GTPase promotes oocyte polarization by regulating ERM (Ezrin/Radixin/Moesin) inactivation. *Cell Cycle* **12**, 1672-1678. doi:10.4161/cc.24901
- Dephoure, N., Zhou, C., Villén, J., Beausoleil, S. A., Bakalarski, C. E., Elledge, S. J. and Gygi, S. P.** (2008). A quantitative atlas of mitotic phosphorylation. *Proc. Natl. Acad. Sci. USA* **105**, 10762-10767. doi:10.1073/pnas.0805139105
- Duan, X. and Sun, S.-C.** (2019). Actin cytoskeleton dynamics in mammalian oocyte meiosis. *Biol. Reprod.* **100**, 15-24. doi:10.1093/biolre/roy163
- Duan, X., Zhang, Y., Chen, K. L., Zhang, H. L., Wu, L. L., Liu, H. L., Wang, Z. B. and Sun, S. C.** (2018). The small GTPase RhoA regulates the LIMK1/2-cofilin pathway to modulate cytoskeletal dynamics in oocyte meiosis. *J. Cell. Physiol.* **233**, 6088-6097. doi:10.1002/jcp.26450
- Faust, J. J., Millis, B. A. and Tyska, M. J.** (2019). Profilin-mediated actin allocation regulates the growth of epithelial microvilli. *Curr. Biol.* **29**, 3457-3465. doi:10.1016/j.cub.2019.08.051
- Figard, L., Wang, M., Zheng, L., Golding, I. and Sokac, A. M.** (2016). Membrane supply and demand regulates F-actin in a cell surface reservoir. *Dev. Cell* **37**, 267-278. doi:10.1016/j.devcel.2016.04.010
- Fitzharris, G., Carroll, J. and Swann, K.** (2018). Electrical-assisted microinjection for analysis of fertilization and cell division in mammalian oocytes and early embryos. *Methods Cell Biol.* **144**, 431-440. doi:10.1016/bs.mcb.2018.03.036
- Grega-Larson, N. E., Crawley, S. W., Erwin, A. L. and Tyska, M. J.** (2015). Cordon bleu promotes the assembly of brush border microvilli. *Mol. Biol. Cell* **26**, 3803-3815. doi:10.1091/mbc.E15-06-0443
- Gressin, L., Guillotin, A., Guérin, C., Blanchoin, L. and Michelot, A.** (2015). Architecture dependence of actin filament network disassembly. *Curr. Biol.* **25**, 1437-1447. doi:10.1016/j.cub.2015.04.011
- Hamill, S., Lou, H. J., Turk, B. E. and Boggon, T. J.** (2016). Structural basis for noncanonical substrate recognition of Cofilin/ADF proteins by LIM kinases. *Mol. Cell* **62**, 397-408. doi:10.1016/j.molcel.2016.04.001
- Higashida, C., Kiuchi, T., Akiba, Y., Mizuno, H., Maruoka, M., Narumiya, S., Mizuno, K. and Watanabe, N.** (2013). F- and G-actin homeostasis regulates mechanosensitive actin nucleation by formins. *Nat. Cell Biol.* **15**, 395-405. doi:10.1038/ncb2693
- Hilton, D. M., Aguilar, R. M., Johnston, A. B. and Goode, B. L.** (2018). Species-specific functions of twinfilin in actin filament depolymerization. *J. Mol. Biol.* **430**, 3323-3336. doi:10.1016/j.jmb.2018.06.025
- Holubcová, Z., Howard, G. and Schuh, M.** (2013). Vesicles modulate an actin network for asymmetric spindle positioning. *Nat. Cell Biol.* **15**, 937-947. doi:10.1038/ncb2802
- Hotulainen, P., Paunola, E., Virtainen, M. K. and Lappalainen, P.** (2005). Actin-depolymerizing factor and cofilin-1 play overlapping roles in promoting rapid F-actin depolymerization in mammalian nonmuscle cells. *Mol. Biol. Cell* **16**, 649-664. doi:10.1091/mbc.e04-07-0555
- Ichetovkin, I., Grant, W. and Condeelis, J.** (2002). Cofilin produces newly polymerized actin filaments that are preferred for dendritic nucleation by the Arp2/3 complex. *Curr. Biol.* **12**, 79-84. doi:10.1016/S0960-9822(01)00629-7
- Isogai, T., van der Kammen, R. and Innocenti, M.** (2015). SMIFH2 has effects on formins and p53 that perturb the cell cytoskeleton. *Sci. Rep.* **5**, 9802. doi:10.1038/srep09802
- Iwase, S., Sato, R., De Bock, P.-J., Gevaert, K., Fujiki, S., Tawada, T., Kuchitsu, M., Yamagishi, Y., Ono, S. and Abe, H.** (2013). Activation of ADF/cofilin by phosphorylation-regulated Slingshot phosphatase is required for the meiotic spindle assembly in Xenopus laevis oocytes. *Mol. Biol. Cell* **24**, 1933-1946. doi:10.1091/mbc.e12-12-0851
- Jacquiermet, G., Paatero, I., Carisey, A. F., Padzik, A., Orange, J. S., Hamidi, H. and Ivaska, J.** (2017). FiloQuant reveals increased filopodia density during breast cancer progression. *J. Cell Biol.* **216**, 3387-3403. doi:10.1083/jcb.201704045
- Jaiswal, R., Breitsprecher, D., Collins, A., Corrêa, Jr I. R., Xu, M. Q. and Goode, B. L.** (2013). The formin Daam1 and fascin directly collaborate to promote filopodia formation. *Curr. Biol.* **23**: 1373-1379.
- Jang, W. I., Jo, Y. J., Kim, H. C., Jia, J. L., Namgoong, S. and Kim, N. H.** (2014). Non-muscle tropomyosin (Tpm3) is crucial for asymmetric cell division and maintenance of cortical integrity in mouse oocytes. *Cell Cycle* **13**, 2359-2369. doi:10.4161/cc.29333
- Jansen, S., Collins, A., Chin, S. M., Ydenberg, C. A., Gelles, J. and Goode, B. L.** (2015). Single-molecule imaging of a three-component ordered actin disassembly mechanism. *Nat. Commun.* **6**: 7202.
- Jin, Z.-L., Jo, Y.-J., Namgoong, S. and Kim, N.-H.** (2018). CAP1-mediated actin cycling via ADF/cofilin proteins is essential for asymmetric division in mouse oocytes. *J. Cell Sci.* **131**, jcs222356. doi:10.1242/jcs.222356
- Jin, Z. L., Yao, X. R., Wen, L., Hao, G., Kwon, J. W., Hao, J. and Kim, N. H.** (2020). AIP1 and cofilin control the actin dynamics to modulate the asymmetric division and cytokinesis in mouse oocytes. *FASEB J.* **34**, 11292-11306. doi:10.1093/fj.202000093R
- Jo, Y.-J., Jang, W.-I., Kim, N.-H. and Namgoong, S.** (2016). Tropomodulin-3 is essential in asymmetric division during mouse oocyte maturation. *Sci. Rep.* **6**, 29204. doi:10.1038/srep29204
- Kajii, N., Ohashi, K., Shuin, M., Niwa, R., Uemura, T. and Mizuno, K.** (2003). Cell cycle-associated changes in Slingshot phosphatase activity and roles in cytokinesis in animal cells. *J. Biol. Chem.* **278**, 33450-33455. doi:10.1074/jbc.M305802200
- Kan, R., Yurttas, P., Kim, B., Jin, M., Wo, L., Lee, B., Gosden, R. and Coonrod, S. A.** (2011). Regulation of mouse oocyte microtubule and organelle dynamics by PADI6 and the cytoplasmic lattices. *Dev. Biol.* **350**, 311-322. doi:10.1016/j.ydbio.2010.11.033
- Kiuchi, T., Ohashi, K., Kurita, S. and Mizuno, K.** (2007). Cofilin promotes stimulus-induced lamellipodium formation by generating an abundant supply of actin monomers. *J. Cell Biol.* **177**, 465-476. doi:10.1083/jcb.200610005
- Kotila, T., Wieland, H., Enkavi, G., Kogan, K., Vattulainen, I., Jégou, A., Romet-Lemonne, G. and Lappalainen, P.** (2019). Mechanism of synergistic actin filament pointed end depolymerization by cyclase-associated protein and cofilin. *Nat. Commun.* **10**, 5320. doi:10.1038/s41467-019-13213-2
- Lappalainen, P. and Drubin, D. G.** (1997). Cofilin promotes rapid actin filament turnover in vivo. *Nature* **388**, 78-82. doi:10.1038/40418
- Li, X., Zhu, Y., Cao, Y., Wang, Q., Du, J., Tian, J., Liang, Y. and Ma, W.** (2017). LIM kinase activity is required for microtubule organising center positioning in mouse oocyte meiosis. *Reprod. Fertil. Dev.* **29**, 791-804. doi:10.1071/RF15406
- Li, H., Guo, F., Rubinstein, B. and Li, R.** (2008). Actin-driven chromosomal motility leads to symmetry breaking in mammalian meiotic oocytes. *Nat. Cell Biol.* **10**, 1301-1308. doi:10.1038/ncb1788
- Longo, F. J. and Chen, D.-Y.** (1985). Development of cortical polarity in mouse eggs: involvement of the meiotic apparatus. *Dev. Biol.* **107**, 382-394. doi:10.1016/0012-1606(85)90320-3
- Lu, Y., Zhang, Y., Pan, M.-H., Kim, N.-H., Sun, S.-C. and Cui, X.-S.** (2017). Daam1 regulates fascin for actin assembly in mouse oocyte meiosis. *Cell Cycle* **16**, 1350-1356. doi:10.1080/15384101.2017.1325045
- Maro, B., Johnson, M. H., Webb, M. and Flach, G.** (1986). Mechanism of polar body formation in the mouse oocyte: an interaction between the chromosomes, the cytoskeleton and the plasma membrane. *J. Embryol. Exp. Morphol.* **92**, 11-32.
- Miao, Y., Wong, C. C., Mennella, V., Michelot, A., Agard, D. A., Holt, L. J., Yates, J. R. 3rd and Drubin, D. G.** (2013). Cell-cycle regulation of formin-mediated actin cable assembly. *Proc. Natl. Acad. Sci. USA* **110**: E4446-E4455. doi:10.1073/pnas.1314000110

- Michelot, A. and Drubin, D. G.** (2011). Building distinct actin filament networks in a common cytoplasm. *Curr. Biol.* **21**, R560-R569. doi:10.1016/j.cub.2011.06.019
- Miyado, K., Yoshida, K., Yamagata, K., Sakakibara, K., Okabe, M., Wang, X., Miyamoto, K., Akutsu, H., Kondo, T., Takahashi, Y. et al.** (2008). The fusing ability of sperm is bestowed by CD9-containing vesicles released from eggs in mice. *Proc. Natl. Acad. Sci. USA* **105**: 12921-12926. doi:10.1073/pnas.0710608105
- Mizuno, K.** (2013). Signaling mechanisms and functional roles of cofilin phosphorylation and dephosphorylation. *Cell. Signal.* **25**, 457-469. doi:10.1016/j.cellsig.2012.11.001
- Mogessie, B. and Schuh, M.** (2017). Actin protects mammalian eggs against chromosome segregation errors. *Science* **357**, eaal1647. doi:10.1126/science.aal1647
- Montaville, P., Jégou, A., Pernier, J., Compper, C., Guichard, B., Mogessie, B., Schuh, M., Romet-Lemonne, G. and Carlier, M. F.** (2014). Spire and Formin 2 synergize and antagonize in regulating actin assembly in meiosis by a ping-pong mechanism. *PLoS Biol.* **12**, e1001795. doi:10.1371/journal.pbio.1001795
- Mooseker, M. S., Pollard, T. D. and Wharton, K. A.** (1982). Nucleated polymerization of actin from the membrane-associated ends of microvillar filaments in the intestinal brush border. *J. Cell Biol.* **95**, 223-233. doi:10.1083/jcb.95.1.223
- Mullins, R. D., Heuser, J. A. and Pollard, T. D.** (1998). The interaction of Arp2/3 complex with actin: nucleation, high affinity pointed end capping, and formation of branching networks of filaments. *Proc. Natl. Acad. Sci. USA* **95**, 6181-6186. doi:10.1073/pnas.95.11.6181
- Nishimura, Y., Shi, S., Zhang, F., Liu, R., Takagi, Y., Bershadsky, A. D., Viasnoff, V. and Sellers, J. R.** (2021). The formin inhibitor SMIFH2 inhibits members of the myosin superfamily. *J. Cell Sci.* **134**, jcs253708. doi:10.1242/jcs.253708
- Nolen, B. J., Tomasevic, N., Russell, A., Pierce, D. W., Jia, Z., McCormick, C. D., Hartman, J., Sakowicz, R. and Pollard, T. D.** (2009). Characterization of two classes of small molecule inhibitors of Arp2/3 complex. *Nature* **460**, 1031-1034. doi:10.1038/nature08231
- Orly, G., Naoz, M. and Gov, N. S.** (2014). Physical model for the geometry of actin-based cellular protrusions. *Biophys. J.* **107**, 576-587. doi:10.1016/j.bpj.2014.05.040
- Pelaseyed, T. and Bretscher, A.** (2018). Regulation of actin-based apical structures on epithelial cells. *J. Cell Sci.* **131**, jcs221853. doi:10.1242/jcs.221853
- Pernier, J., Shekhar, S., Jegou, A., Guichard, B. and Carlier, M.-F.** (2016). Profilin interaction with actin filament barbed end controls dynamic instability, capping, branching, and motility. *Dev. Cell* **36**, 201-214. doi:10.1016/j.devcel.2015.12.024
- Pfender, S., Kuznetsov, V., Pleiser, S., Kerkhoff, E. and Schuh, M.** (2011). Spire-type actin nucleators cooperate with Formin-2 to drive asymmetric oocyte division. *Curr. Biol.* **21**, 955-960. doi:10.1016/j.cub.2011.04.029
- Pfender, S., Kuznetsov, V., Pasternak, M., Tischer, T., Santhanam, B. and Schuh, M.** (2015). Live imaging RNAi screen reveals genes essential for meiosis in mammalian oocytes. *Nature* **524**, 239-242. doi:10.1038/nature14568
- Pollard, T. D. and Cooper, J. A.** (2009). Actin, a central player in cell shape and movement. *Science* **326**, 1208-1212. doi:10.1126/science.1175862
- Postema, M. M., Grega-Larson, N. E., Neininger, A. C. and Tyska, M. J.** (2018). IRTKS (BAIAP2L1) elongates epithelial microvilli using EPS8-dependent and independent mechanisms. *Curr. Biol.* **28**, 2876-2888. doi:10.1016/j.cub.2018.07.022
- Rime, H. and Ozon, R.** (1990). Protein phosphatases are involved in the in vivo activation of histone H1 kinase in mouse oocyte. *Dev. Biol.* **141**, 115-122. doi:10.1016/0012-1606(90)90106-S
- Rizvi, S. A., Neidt, E. M., Cui, J., Feiger, Z., Skau, C. T., Gardel, M. L., Kozmin, S. A. and Kovar, D. R.** (2009). Identification and characterization of a small molecule inhibitor of formin-mediated actin assembly. *Chem. Biol.* **16**, 1158-1168. doi:10.1016/j.chembiol.2009.10.006
- Roberts, R., Iatropoulou, A., Ciantar, D., Stark, J., Becker, D. L., Franks, S. and Hardy, K.** (2005). Follicle-stimulating hormone affects metaphase I chromosome alignment and increases aneuploidy in mouse oocytes matured in vitro. *Biol. Reprod.* **72**, 107-118. doi:10.1095/biolreprod.104.03203
- Rotty, J. D., Wu, C., Haynes, E. M., Suarez, C., Winkelmann, J. D., Johnson, H. E., Haugh, J. M., Kovar, D. R. and Bear, J. E.** (2015). Profilin-1 serves as a gatekeeper for actin assembly by Arp2/3-dependent and -independent pathways. *Dev. Cell* **32**, 54-67. doi:10.1016/j.devcel.2014.10.026
- Runge, K. E., Evans, J. E., He, Z.-Y., Gupta, S., McDonald, K. L., Stahlberg, H., Primakoff, P. and Myles, D. G.** (2007). Oocyte CD9 is enriched on the microvillar membrane and required for normal microvillar shape and distribution. *Dev. Biol.* **304**, 317-325. doi:10.1016/j.ydbio.2006.12.041
- Salah, E., Chatterjee, D., Beltrami, A., Tumber, A., Preuss, F., Canning, P., Chaikud, A., Knaus, P., Knapp, S., Bullock, A. N. et al.** (2019). Lessons from LIMK1 enzymology and their impact on inhibitor design. *Biochem. J.* **476**, 3197-3209. doi:10.1042/BCJ20190517
- Sampath, P. and Pollard, T. D.** (1991). Effects of cytochalasin, phalloidin, and pH on the elongation of actin filaments. *Biochemistry* **30**, 1973-1980. doi:10.1021/bi00221a034
- Schuh, M. and Ellenberg, J.** (2008). A new model for asymmetric spindle positioning in mouse oocytes. *Curr. Biol.* **18**, 1986-1992. doi:10.1016/j.cub.2008.11.022
- Shekhar, S. and Carlier, M.-F.** (2017). Enhanced depolymerization of actin filaments by ADF/Cofilin and monomer funneling by capping protein cooperate to accelerate barbed end growth. *Curr. Biol.* **27**, 1990-1998. doi:10.1016/j.cub.2017.05.036
- Shekhar, S., Pernier, J. and Carlier, M.-F.** (2016). Regulators of actin filament barbed ends at a glance. *J. Cell Sci.* **129**, 1085-1091.
- Shekhar, S., Chung, J., Kondev, J., Gelles, J. and Goode, B. L.** (2019). Synergy between Cyclase-associated protein and Cofilin accelerates actin filament depolymerization by two orders of magnitude. *Nat. Commun.* **10**, 5319. doi:10.1038/s41467-019-13268-1
- Sotiropoulos, A., Gineitis, D., Copeland, J. and Treisman, R.** (1999). Signal-regulated activation of serum response factor is mediated by changes in actin dynamics. *Cell* **98**, 159-169. doi:10.1016/S0092-8674(00)81011-9
- Stidwill, R. P. and Burgess, D. R.** (1986). Regulation of intestinal brush border microvillus length during development by the G- to F-actin ratio. *Dev. Biol.* **114**, 381-388. doi:10.1016/0012-1606(86)90202-2
- Suarez, C. and Kovar, D. R.** (2016). Internetwork competition for monomers governs actin cytoskeleton organization. *Nat. Rev. Mol. Cell Biol.* **17**, 799-810. doi:10.1038/nrm.2016.106
- Suarez, C., Roland, J., Boujemaa-Paterski, R., Kang, H., McCullough, B. R., Reymann, A.-C., Guérin, C., Martiel, J.-L., De la Cruz, E. M. and Blanchoin, L.** (2011). Cofilin tunes the nucleotide state of actin filaments and severs at bare and decorated segment boundaries. *Curr. Biol.* **21**, 862-868. doi:10.1016/j.cub.2011.03.064
- Suarez, C., Carroll, R. T., Burke, T. A., Christensen, J. R., Bestul, A. J., Sees, J. A., James, M. L., Sirotnik, V. and Kovar, D. R.** (2015). Profilin regulates F-actin network homeostasis by favoring formin over Arp2/3 complex. *Dev. Cell* **32**, 43-53. doi:10.1016/j.devcel.2014.10.027
- Takahashi, T., Koshimizu, U., Abe, H., Obinata, T. and Nakamura, T.** (2001). Functional involvement of Xenopus LIM kinases in progression of oocyte maturation. *Dev. Biol.* **229**, 554-567. doi:10.1006/dbio.2000.9999
- Uraji, J., Scheffler, K. and Schuh, M.** (2018). Functions of actin in mouse oocytes at a glance. *J. Cell Sci.* **131**, jcs218099. doi:10.1242/jcs.218099
- Van Blerkom, J. and Bell, H.** (1986). Regulation of development in the fully grown mouse oocyte: chromosome-mediated temporal and spatial differentiation of the cytoplasm and plasma membrane. *J. Embryol. Exp. Morphol.* **93**, 213-238.
- Vitriol, E. A., McMillen, L. M., Kapustina, M., Gomez, S. M., Vavylonis, D. and Zheng, J. Q.** (2015). Two functionally distinct sources of actin monomers supply the leading edge of lamellipodia. *Cell Rep.* **11**, 433-445. doi:10.1016/j.celrep.2015.03.033
- Wei, Z., Greaney, J., Zhou, C. and Homer, H. A.** (2018). CDK1 inactivation induces post-anaphase-onset spindle migration and membrane protrusion required for extreme asymmetry in mouse oocytes. *Nat. Commun.* **9**, 4029. doi:10.1038/s41467-018-06510-9
- Wioland, H., Guichard, B., Senju, Y., Myram, S., Lappalainen, P., Jégou, A. and Romet-Lemonne, G.** (2017). ADF/Cofilin accelerates actin dynamics by severing filaments and promoting their depolymerization at both ends. *Curr. Biol.* **27**, 1956-1967. doi:10.1016/j.cub.2017.05.048
- Wong, G. K., Allen, P. G. and Begg, D. A.** (1997). Dynamics of filamentous actin organization in the sea urchin egg cortex during early cleavage divisions: implications for the mechanism of cytokinesis. *Cell Motil. Cytoskeleton* **36**, 30-42. doi:10.1002/(SICI)1097-0169(1997)36:1<30::AID-CM3>3.0.CO;2-L
- Xiao, Y., Ma, H., Wan, P., Qin, D., Wang, X., Zhang, X., Xiang, Y., Liu, W., Chen, J., Yi, Z. et al.** (2017). Trp-Asp (WD) repeat domain 1 is essential for mouse peri-implantation development and regulates cofilin phosphorylation. *J. Biol. Chem.* **292**, 1438-1448. doi:10.1074/jbc.M116.759886
- Yang, N., Higushi, O., Ohashi, K., Nagata, K., Wada, A., Kangawa, K., Nishida, E. and Mizuno, K.** (1998). Cofilin phosphorylation by LIM-kinase 1 and its role in Rac-mediated actin reorganization. *Nature* **393**, 809-812. doi:10.1038/31735
- Yi, K., Unruh, J. R., Deng, M., Slaughter, B. D., Rubinstein, B. and Li, R.** (2011). Dynamic maintenance of asymmetric meiotic spindle position through Arp2/3-complex-driven cytoplasmic streaming in mouse oocytes. *Nat. Cell Biol.* **13**, 1252-1258. doi:10.1038/ncb2320
- Yi, K., Rubinstein, B., Unruh, J. R., Guo, F., Slaughter, B. D. and Li, R.** (2013). Sequential actin-based pushing forces drive meiosis I chromosome migration and symmetry breaking in oocytes. *J. Cell Biol.* **200**, 567-576. doi:10.1083/jcb.201211068
- Yu, X.-J., Yi, Z., Gao, Z., Qin, D., Zhai, Y., Chen, X., Ou-Yang, Y., Wang, Z.-B., Zheng, P., Zhu, M.-S. et al.** (2014). The subcortical maternal complex controls symmetric division of mouse zygotes by regulating F-actin dynamics. *Nat. Commun.* **5**: 4887. doi:10.1038/ncomms5887