Formin-generated actomyosin arcs propel T cell receptor microcluster movement at the immune synapse

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Actin assembly and inward flow in the plane of the immunological synapse (IS) drives the centralization of T cell receptor microclusters (TCR MCs) and the integrin leukocyte functional antigen 1 (LFA-1). Using structured-illumination microscopy (SIM), we show that actin arcs populating the medial, lamella-like region of the IS arise from linear actin filaments generated by one or more formins present at the IS distal edge. After traversing the outer, Arp2/3-generated, lamellipodia-like region of the IS, these linear filaments are organized by myosin II into antiparallel concentric arcs. Three-dimensional SIM shows that active LFA-1 often aligns with arcs, whereas TCR MCs commonly reside between arcs, and total internal reflection fluorescence SIM shows TCR MCs being swept inward by arcs. Consistently, disrupting actin arc formation via formin inhibition results in less centralized TCR MCs, missegregated integrin clusters, decreased T-B cell adhesion, and diminished TCR signaling. Together, our results define the origin, organization, and functional significance of a major actomyosin contractile structure at the IS that directly propels TCR MC transport.

Introduction

Recognition of antigen on the surface of an antigen-presenting cell (APC) initiates signaling cascades within the T cell that drive large-scale reorganization of its actin cytoskeleton (Bennmiller and Krummel, 2013; Yu et al., 2013; Kumari et al., 2014). This reorganization is essential for the formation of the immunological synapse (IS), the specialized interface between the two cells (Monks et al., 1998; Grakoui et al., 1999). Initially, activation of actin polymerization within the T cell at the periphery of its contact with the APC drives the spreading of the T cell across the surface of the APC. Once spreading is complete, continued actin polymerization begins to drive an inward flow of actin toward the center of the contact site and in the plane of the IS. By coupling this inward flow with depolymerization at the center of the IS, the T cell creates an ongoing centripetal flow of actin that is thought to be a major driving force for the inward movement of T cell receptor microclusters (TCR MCs) and integrin clusters in the T cell’s plasma membrane (Bunnell et al., 2001; Varma et al., 2006; Kaizuka et al., 2007; Babich et al., 2012; Beemiller et al., 2012; Smoligovets et al., 2012; Yi et al., 2012). Over the next 5–10 min, the inward movement of receptor clusters culminates in the formation of a mature IS, in which TCR MCs are concentrated at the center of the IS (the central supramolecular activation cluster [cSMAC]), and leukocyte functional antigen 1 (LFA-1), the T cell’s major integrin, is concentrated in a surrounding ring (the peripheral SMAC [pSMAC]). Importantly, actin assembly and dynamics are intimately linked not just to TCR MC movement, but to virtually every key event during IS formation, including signalosome assembly and tuning (Mattila et al., 2016), integrin activation (Comrie et al., 2015a,b), the mechanical regulation of T cell signaling (Chen and Zhu, 2013), and effector functions such as lytic granule release (Brown et al., 2011; Mace et al., 2012; Basu et al., 2016). Clearly, a full understanding of how actin cytoskeletal forces are created and organized at the IS is required to define the mechanisms by which they drive T cell function.

Numerous laboratories have used diffraction-limited imaging of T cells engaged with planar lipid bilayers containing freely diffusing activators (e.g., anti-CD3 and intercellular adhesion molecule 1 [ICAM-1]) to correlate the dynamics of actin...
flow and receptor cluster movement in an ideal imaging plane (Dustin, 2009). Importantly, these studies revealed robust, polymerization-driven, actin retrograde flow in a ring surrounding the pSMAC now known as the distal SMAC (dSMAC; Kaizuka et al., 2007; Babich et al., 2012; Beemiller et al., 2012; Yi et al., 2012). Moreover, the rate of centripetal TCR MC movement in this radially symmetric dSMAC roughly correlated with the rate of inward actin flow (Kaizuka et al., 2007), and elegant biophysical studies demonstrated frictional coupling between the TCR MCs and actin flow (DeMond et al., 2008; Yu et al., 2010). Less clear, however, is what propels TCR MC movement across the pSMAC, especially as GFP-actin, the reporter typically used to image actin dynamics at the IS, does not reveal obvious actin organization there (Kaizuka et al., 2007). Using F-Actin, an indirect reporter for F-actin, we, in contrast, identified concentric actin arcs in the pSMAC that are decorated with myosin II (Yi et al., 2012). Additionally, we showed that the lamellipodial-like dSMAC and lamella-like pSMAC exhibit distinct rates of inward actin flow and that the rates of centripetal TCR MC movement across these two zones matched their distinct actin flow rates (Yi et al., 2012). Nevertheless, the existence of these arcs has been questioned (Beemiller and Krummel, 2013; Le Floc’h and Huse, 2015), and they have never been observed in primary T cells. Moreover, an alternate mechanism to drive TCR MC movement across the pSMAC has been proposed that involves dynein-dependent, microtubule-based transport (Hashimoto-Tane et al., 2011). Finally, the role of myosin II in TCR MC movement has been controversial (Hammer and Burkhardt, 2013).

In this study, we sought to determine how the actin arcs are created and how they become organized into concentric structures. Beyond that, we sought to define the spatial relationship between the arcs and receptor clusters during IS maturation and to quantify the contributions made by arcs to receptor cluster distribution, T cell–APC adhesion, and proximal TCR signaling. Finally, we sought to directly image the arc-dependent translocation of TCR MCs. Key to accomplishing these goals was the use of total internal reflection fluorescence–structured-illumination microscopy (TIRF-SIM), which, by virtue of its high spatiotemporal resolution (Kner et al., 2009; Beach et al., 2014; Li et al., 2015), proved pivotal in reaching several of our main conclusions.

Results

SIM reveals the architecture and dynamics of F-actin at the IS

To characterize the architecture of actin networks at the IS at higher resolution, we imaged activated, phalloidin-stained Jurkat T cells with 3D-SIM to reveal endogenous F-actin structures at ~120-nm lateral and ~250-nm axial resolution. Consistent with our previous work at lower resolution (Yi et al., 2012), 3D-SIM demonstrated that the IS is comprised of three distinct zones of F-actin: an outer, dense ring corresponding to the branched actin network in the dSMAC, a middle ring comprised of concentric actin arcs corresponding to the pSMAC, and a central, relatively actin-depleted zone corresponding to the cSMAC (Fig. 1 A, first and second panels). Color coding the 3D projection of this cell according to z position (Fig. 1 A, third panel) shows that these actin structures are largely confined to the plane of the IS, indicating that their dynamics will occur in close association with the plasma membrane at the IS. Also consistent with previous work, 3D-SIM images of Jurkats stained for actin and endogenous myosin II A, the dominant myosin II isoform in T cells, showed that the concentric actin arcs in the pSMAC are highly enriched for this contractile motor (Fig. 1 B).

We next sought to visualize the dynamics of IS actin networks using a live-cell compatible SIM technique. Given that these networks are close to the plasma membrane, we used TIRF-SIM, which merges the power of SIM (in this study, ~100-nm xy resolution) with TIRF (enhanced signal-to-noise ratio for structures close to the coverslip) in a format that displays little bleaching or phototoxicity while providing high temporal resolution (Kner et al., 2009; Li et al., 2015). Fig. 1 C (first three panels) shows still images taken from a video of a Jurkat T cell expressing GFP–F-Actin, an indirect, dynamic reporter for F-actin (Johnson and Schell, 2009), 180, 210, and 240 s (Fig. 1 C) after engagement with an activating surface (Video 1). It is immediately apparent that F-Actin faithfully reports the two actin networks and three IS zones identified in fixed images of phalloidin-stained cells. This is most clearly seen in Fig. 1 C (fourth panel), as well as in the corresponding video. Additionally, TIRF-SIM imaging of Jurkats expressing GFP–myosin IIA and tdTomato–F-Actin revealed in dynamic fashion the enrichment of myosin on the actin arcs in the pSMAC (Fig. 1 D, first and second panels). Importantly, close examination of these images, stills from Video 2 (Fig. 1 D, third and fourth panels), and Video 2 itself revealed the presence of individual, 300-nm-long myosin IIA bipolar filaments decorating the actin arcs and moving inward with them. Together, these observations confirm and enhance our previous understanding of actin architecture at the Jurkat T cell IS and provide additional support for a radially symmetric contractile actomyosin network within the pSMAC.

Actomyosin arcs are also prominent structures at the IS of primary mouse T cells

We performed 3D-SIM on activated primary mouse CD8+ T cells stained with phalloidin and an antibody to myosin IIA to reveal endogenous structures. Fig. 2 A shows that the IS of a CD8+ T cell is essentially indistinguishable from that of a Jurkat T cell. Also, like Jurkats, the two major F-actin networks at the CD8+ T cell IS are very close to the IS membrane (Fig. 2 C). Strikingly, enlarged images of typical CD8+ T cell synapses reveal an almost sarcomere-like pattern for myosin IIA embedded in the actin arcs comprising their pSMAC (Fig. 2 B, arrowheads). Perhaps most importantly, rather than being difficult to detect or absent, actomyosin arcs are an even more prominent component of the mouse CD8+ T cell IS than the Jurkat T cell IS (Fig. 2 D). Finally, still images from TIRF-SIM imaging of primary mouse CD8+ T cells expressing GFP–myosin IIA and tdTomato–F-Actin demonstrated the formation of the actomyosin arcs in the pSMAC (Fig. S1, A and B). Together, these results demonstrate in unequivocal fashion the existence of actomyosin arcs at the IS of primary mouse T cells.

3D-SIM and TIRF-SIM reveal linear actin filaments/bundles embedded within the dSMAC that appear to give rise to the actin arcs in the pSMAC

Closer examination of fixed, phalloidin-stained Jurkat synapses using 3D-SIM revealed linear actin filaments/bundles embed-
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Within the branched actin network of the dSMAC. These structures appear to originate at the distal edge of the IS, traverse the dSMAC, and bend upon exiting the dSMAC to give rise to the actin arcs of the pSMAC (Fig. 3A, yellow arrowheads in the bottom panels mark the origin of these structures and red arrowheads mark their exit from the dSMAC). Further support for their existence came from live-cell TIRF-SIM imaging of Jurkat T cells expressing GFP–F-Tractin. Specifically, Fig. 3B shows still images taken from Video 4 of a quadrant of a Jurkat T cell synapse 180, 240, 300, and 360 s after engagement with an activating surface. Close examination reveals the progressive formation of several linear actin filaments/bundles (Fig. 3B, arrowheads) at the distal edge of the IS that are oriented perpendicular to the plasma membrane, traverse the dSMAC, and then bend and splay upon exit from the dSMAC to form arcs (Video 3). Fig. 3C shows four clear examples of such bending upon exit from the inner aspect of the dSMAC (see red arrowheads), and Video 4 shows all of the events described in this section occurring across a broad section of a Jurkat IS.

The linear actin filaments/bundles, and the actin arcs they give rise to, are formin generated.

From a topological standpoint, the linear actin filaments/bundles observed in Fig. 3 are consistent with actin structures built by formins (Kovar, 2006; Breitsprecher and Goode, 2013). Specifically, they originate at the plasma membrane, the site where formins are typically active, and project inwards, one of two fates typically exhibited by formin-generated filaments (the other being projection outward to make structures like filopodia). Additional indirect evidence that these structures are formin generated stems from the observation that the arcs are very poorly labeled by GFP-actin (Kaizuka et al., 2007) because formins are known to incorporate modified actins into filaments very ineffectively (Chen et al., 2012). This is completely consistent with our previous observations (Yi et al., 2012), which we confirmed and extended in this study using colocalization analyses (Fig. S1, C–E).

Recent, paradigm-shifting studies in both yeast (Burke et al., 2014) and mammalian cells (Lomakin et al., 2015; Rotty et al., 2015) have demonstrated that the Arp2/3 complex and formins are competing for a limiting pool of actin monomer, such that when one actin nucleator is inhibited, the network created by the other becomes more robust. We reasoned, therefore, that if the linear actin filaments/bundles embedded within the dSMAC are formin generated, they should be augmented upon Arp2/3 inhibition. Consistently, Video 5 and the still images taken from it (Fig. 4A) show that the addition of the Arp2/3 inhibitor CK666 to activated Jurkat T cells expressing GFP–F-Tractin resulted in the dramatic collapse of the Arp2/3-dependent, branched actin network comprising the dSMAC and the progressive enhancement of the linear actin filaments/bundles, which took on the
appearance of radial actin “spikes” in which actin polymerization and retrograde flow continued (Fig. 4 A, compare second panel [+CK666] to first panel [control]; note that this effect was completely reversible in third panel [washout]; see also Video 5 and Gomez et al., 2007). 3D-SIM imaging of fixed, phalloidin-stained Jurkat T cells treated with CK666 for 1, 2, and 4 min revealed these spikes in greater detail as they emerged from the shrinking lamellipodial network and increased in robustness over time (Fig. 4 B, top). Importantly, actin arcs were still present in these Arp2/3-inhibited cells. Moreover, the corresponding insets in Fig. 4 B (bottom) show that the CK666-enhanced actin filaments/bundles in the dSMAC bend and reorient into actin arcs at the dSMAC/pSMAC boundary (the red arrowheads mark bend points where the spikes marked by yellow arrowheads give rise to arcs). This bending and reorientation process could be observed in real time in Jurkats expressing GFP–F-Tractin and imaged using TIRF-SIM (Video 6 and the still images taken from it in Fig. 4 C). Together, these observations indicate that arc assembly is largely Arp2/3 independent and argue that the arcs are created by one or more formins acting at the distal edge of the IS.

To provide more direct support for the hypothesis that the actin spikes observed upon Arp2/3 inhibition are formin generated, we compared cells treated with CK666 alone to cells treated with both CK666 and the pan-formin inhibitor small-molecule inhibitor of formin homology 2 domain (SMIFH2; Rizvi et al., 2009). As before, 3-min incubation in the presence of CK666 resulted in a considerable diminution in the size of the branched actin array comprising most of the dSMAC and the appearance of actin spikes at the cell edge (Fig. 4 D, compare second panel [+CK666−SMIFH2] to first [DMSO control] panel; the red bracket in the second panel marks a typical actin spike). Importantly, addition of CK666 to cells that had been pretreated with SMIFH2 resulted in a dramatic reduction in the formation of spikes (Fig. 4 D, compare second panel [+CK666−SMIFH2] to first [DMSO control] panel; the red bracket in the second panel marks a typical actin spike). Importantly, addition of CK666 to cells that had been pretreated with SMIFH2 resulted in a considerable diminution in the size of the branched actin array comprising most of the dSMAC and the appearance of actin spikes at the cell edge (Fig. 4 D, compare second panel [+CK666−SMIFH2] to first [DMSO control] panel; the red bracket in the second panel marks a typical actin spike). Importantly, addition of CK666 to cells that had been pretreated with SMIFH2 resulted in a dramatic reduction in the formation of spikes (Fig. 4 D, compare second panel [+CK666−SMIFH2] to first [DMSO control] panel; the red bracket in the second panel marks a typical actin spike).
idea that the linear actin filaments/bundles embedded within the dSMAC and contiguous with the actin arcs in the pSMAC are formin-generated structures.

As a more direct test for the role of formins in actin arc creation, we imaged arc formation with and without SMIFH2. To accomplish this, we allowed Jurkats to activate for 2 min before SMIFH2 addition and then asked if its addition prevented further actin arc formation. We observed that a 5-min incubation in the presence of SMIFH2 (both 10 and 25 µM) resulted in a dramatic decrease in the presence of actin arcs within the pSMAC, as revealed by 3D-SIM imaging of fixed, phalloidin-stained cells (Fig. 5 A, compare second and third panels and the corresponding insets in the fifth and sixth panels to the DMSO control in the first and fourth panels). To quantitate arc loss, we measured total phalloidin fluorescence in the pSMAC area of control cells and cells treated with 10 µM SMIFH2. Fig. 5 A (bottom left) shows that SMIFH2-treated cells exhibited significantly less F-actin in their pSMACs, consistent with the loss of actin arcs after formin inhibition. Importantly, SMIFH2-induced arc loss is likely attributable to formin inhibition rather than off-target effects because treatment with 10 µM KVSM18, a closely related but inactive analogue of SMIFH2 (Rizvi et al., 2009), does not cause arc loss (Fig. 5 A, bottom middle).

Interestingly, formin-inhibited cells also exhibited a dramatic increase in the number of F-actin foci in the dSMAC (Fig. 5 A, arrowheads). We believe these foci correspond to the Arp2/3-dependent F-actin foci described recently (Kumar et al., 2015) and that they become more robust upon formin inhibition as a result of an increase in actin monomer available for formin-dependent nucleation. On the flip side, the inhibition of Arp2/3-dependent nucleation using CK666 increased the content of arcs in the pSMAC (Fig. 5 A, bottom right), presumably because of an increase in actin monomer available for formin-dependent nucleation.

Importantly, the SMIFH2-mediated inhibition of actin arc formation was reversible, as washout of SMIFH2 resulted in the rapid resumption of actin arc formation. This is apparent in still images taken from a TIRF-SIM video (Video 7) 30 (Fig. 5 B, top) and 180 s (Fig. 5 B, bottom) after SMIFH2 washout. Careful examination of the magnified insets in Fig. 5 B (right panels) show that SMIFH2 washout was accompanied not only by the appearance of newly formed arcs in the pSMAC (Fig. 5 B, bottom right), but also by the disappearance of the actin foci (Fig. 5 B, yellow arrowheads in top right), and the reappearance of linear actin filaments/bundles embedded in the dSMAC (yellow arrowheads in bottom right). Within 5 min of washout, the content of F-actin in the pSMAC was restored to control levels (Fig. 5 A, bottom left).

We next tested if one or more formins could be detected at the tips of CK666-enhanced actin spikes by immunofluorescence staining for endogenous proteins. Previous studies (Gomez et al., 2007; Ramabhadran et al., 2011) have provided evidence for the expression of four formins in Jurkat T cells: mDia1, mDia2, formin-like protein 1 (FMNL1), and the non-CAAX version of inverted formin-2 (INF2). Of these, we were able to detect clear signals for mDia1, FMNL1, and non-CAAX INF2 at the tips of CK666-enhanced actin spikes (Fig. S2 A, micrographs). Moreover, the fraction of spike tips that stained for these three formins, which ranged from ~35 to ~55%, was significantly higher than the fraction that stained for mDia2 or with preimmune sera (Fig. S2 A, right panel), arguing that the tip staining seen for mDia1, FMNL1, and
non-CAAX INF2 is specific. Of these three formins, mDia1 appears to be the most highly expressed in Jurkats (Gomez et al., 2007). Importantly, Jurkat T cells in which we achieved near-complete knockdown of mDia1 using shRNA (Fig. S2 B, top left) exhibited a dramatic decrease in the content of arcs in their pSMAC relative to control cells expressing a scrambled shRNA plasmid (Fig. S2 B, micrographs and bottom left). Also like SMIFH2-treated cells, mDia1 knockdown cells exhibited a proliferation of Arp2/3-dependent actin foci (Fig. S2 B, bottom right, arrowheads). Together, these data argue that the formin mDia1 plays a major role in arc formation. Consistent with this conclusion, Jurkats treated with a mem-
brane-permeant pan-activator of Rho GTPases (CN01), which activate most formins including mDia1, exhibited accentuated linear actin structures embedded in their dSMAC, in the amount of arcs in their pSMAC, and in the overall size of their pSMAC (Fig. S3, A–C). Finally, we found that tropomyosin, a universal marker of formin-generated structures (Tojkander et al., 2012), strongly decorates actin arcs. Specifically, Fig. S3 (D–F) show that GFP-tagged nonmuscle tropomyosin TM4 is highly enriched on the arcs in the pSMAC, whereas α-actinin, which typically cross-links Arp2/3-generated branched actin networks, is largely restricted to the dSMAC. Together, the results in this section provide compelling evidence that arcs arise largely, if not entirely, from formin-dependent nucleation at the outer edge of the dSMAC.
Myosin II organizes the linear actin filaments/bundles emanating from the dSMAC into the concentric actin arcs of the pSMAC

Given the enrichment of myosin IIA at the pSMAC and the inherent ability of bipolar myosin II filaments to organize actin networks, we asked if myosin II is required to organize the linear actin filaments/bundles emanating from the dSMAC into the concentric actin arcs of the pSMAC. To selectively and conditionally inhibit myosin IIA, we used para-nitro-blebbistatin (pnBB), a new variant of blebbistatin that lacks the original molecule’s phototoxicity (Képiró et al., 2014). In all experiments, Jurkats were pretreated with pnBB for 30 min followed by activation on antibody-coated coverslips for 10 min. 3D-SIM images of fixed, phalloidin-stained cells show that preincubation with 50 µM pnBB resulted in large-scale disruption of actin arc organization in the pSMAC (Fig. 6 A, compare second and third panels, and the corresponding insets in fifth and sixth panels to the DMSO control in first and fourth panels). This result was borne out by visual scoring of arc organization (Fig. 6 B), as only ~12% of cells treated with 50 µM pnBB and ~5% of cells treated with 100 µM pnBB have highly organized concentric actin arcs, as compared with ~52% of control, DMSO-treated Jurkats (Fig. 6 B, orange bars). In place of organized arcs, the pSMAC region of pnBB-treated cells exhibited either: (1) actin filaments/bundles that projected inward toward the center of the IS (Fig. 6 A, middle panels; and Fig. 6 B, blue bars); or (2) a highly disorganized, meshlike organization (Fig. 6 A, right panels; Fig. 6 B, gray bars; and Video 8).

To further quantify the defects in actin arc organization seen upon myosin II inhibition, we determined the anisotropy of actin structures in the pSMAC using FibrilTool, which measures how well a structure of interest (in this study, actin arcs) in a given region of interest (ROI; in this study, the pSMAC)
is organized in parallel arrays. The values obtained range from 0, when the orientation of the structures in the ROI is random, to 1, when all of the structures are oriented in the same direction. To deal with the fact that the IS is radially symmetric, we divided the pSMAC region in 3D-SIM images of fixed, phalloidin-stained Jurkats into 10–12 trapezoid-shaped ROIs of similar size (Fig. 6 C). Fig. 6 D shows that the anisotropy of pSMAC actin structures is significantly lower in cells treated with 50 µM pnBB (red line) than in control, DMSO-treated cells (black line), corroborating the qualitative assessments described in Fig. 6 (A and B). Together, these data argue strongly that bipolar filamentous of myosin IIA organize formin-generated, linear actin filaments/bundles emanating from the dSMAC into concentric actin arcs populating the pSMAC. Importantly, stochastic variation in the direction the linear filaments bend as they exit the dSMAC should ensure that bipolar myosin II filaments are routinely engaged with actin filaments of opposing orientation, a prerequisite for myosin II–dependent contraction and inward movement of arcs (Fig. 6 E). This self-organizing property should propel the formation of the radially symmetric contractile actomyosin network in the pSMAC.

**Actomyosin arcs facilitate the accumulation of TCR and LFA-1 clusters at the cSMAC and inner aspect of the pSMAC, respectively.** Glass-supported planar lipid bilayers displaying stimulatory molecules to mimic the APC surface have been used extensively to correlate cytoskeletal dynamics in the plane of the IS with TCR MC and integrin cluster position and dynamics in T cells (Kaizuka et al., 2007; Hartman et al., 2009). As a first test for the role of actin arcs in IS formation, we determined the distributions of F-actin, TCR MCs, and the open, active conformation of LFA-1’s β2 subunit in control, DMSO-, and SMIFH2-treated Jurkats after a 10-min engagement with bi-layers containing fluorescent anti-CD3 antibody and unlabeled ICAM (and where antibody M24 against the fixation-sensitive, open, active conformation of the β2 integrin was present for the last 4 min). At 10 min, the cells were fixed, stained with phalloidin, and the distributions of F-actin (Fig. 7, A and B, white), TCR MCs (red), and open, active LFA-1 (green) imaged with 3D-SIM. As expected, control, DMSO-treated Jurkats exhibited a strong accumulation of TCR MCs in their cSMAC (i.e., inside the arcs populating their pSMAC; Fig. 7 A, left), and the accumulation of active LFA-1 in a ring along the inner aspect of their pSMAC/actin arcs (Fig. 7 B, left). Strikingly, SMIFH2 treatment disrupted the centralization of TCR MCs (Fig. 7 A, right) and the formation of the ring of active LFA-1 (Fig. 7 B, right), accompanied by the loss of actin arcs (Fig. 7, A and B, right). These conclusions are supported by radial plot profiles, which report the mean distribution of TCR MCs (Fig. 7 C) and active LFA-1 clusters (Fig. 7 D) between the cell center (0 on the x axis) and the cell edge (1.0 on the x axis). Relative to the DMSO control (black lines), SMIFH2 addition (red lines) reduced the centralization of TCR MCs (Fig. 7 C, as indicated by a flattening and rightward shift in the red line) and broadened the distribution of active LFA-1 clusters (as indicated by a broadening of the red line on both sides of the black peak in Fig. 7 D; P < 0.0001). The effect of SMIFH2 on the distribution of the open, active integrin becomes especially clear when we fit line scans across the IS to Gaussian distributions (Fig. 7 E), in which the normal bimodal distribution of active LFA-1 across the IS that corresponds to the pSMAC ring (black circles and black fitted line) is lost upon SMIFH2 treatment (red circles and red fitted line). Finally, consistent with the fact that arcs require myosin II to assemble into radially symmetric contractile structures, we observed similar defects in the distributions of TCR MCs and active LFA-1 in Jurkats treated with pnBB (Fig. S4 A). Collectively, these results argue strongly that actin arcs, in conjunction with myosin II bipolar filaments, make a major contribution to the centripetal movement and distribution of both TCR MCs and integrin clusters at the IS.

**Actomyosin arcs promote robust T cell–APC adhesion and TCR proximal signaling.** To further gauge the physiological significance of actin arcs at the IS, we measured adhesion and TCR proximal signaling in control and formin-inhibited Jurkats present in conjugates with an APC. For the adhesion assay, Jurkats were mixed with staphylococcal enterotoxin E (SEE)–loaded Raji B cells in the presence or absence of SMIFH2, incubated for 10 min, fixed, immunostained with fluorescent antibodies against OKT3 and human leukocyte antigen (HLA) to mark Raji B cells, and the fraction of total Jurkat cells present in OKT3 and HLA double-positive conjugates was determined by flow cytometry analysis. Fig. 7 F shows that inhibition of arc formation with SMIFH2 results in an approximately fivefold reduction in cell adhesion relative to the DMSO control.

To test for possible defects in TCR proximal signaling after the loss of arcs, Jurkat T cells and Raji B cells expressing plasma membrane markers farnesylated GFP and farnesylated RFP, respectively, were mixed together in the absence or presence of SMIFH2, incubated for 10 min, fixed, and immunostained with fluorescent antibodies specific for the tyrosine phosphorylated versions of Src (Y416), Zap70 (Y319), or Lat (Y191) as proxies for the extent of proximal TCR signaling. To quantitate the signals for pSrc, pZap70, and pLat at the IS, we used an Amnis ImageStream flow cytometer, which combines the sample-size advantages of flow cytometry with the sensitivity of fluorescence microscopy. Specifically, we identified GFP and RFP double-positive T–B cell conjugates and measured the mean fluorescence intensity (MFI) of the three phosphorylated signaling molecules on the T cell side of the IS using a masking strategy to identify the contact area between the two cells (Fig. 7 G and Materials and methods). Fig. 7 H shows that disruption of arc formation using SMIFH2 resulted in marked reductions relative to the DMSO control in the content of pSrc (Y416) and pLAT (Y191) at the IS of Jurkat–Raji cell conjugates and a small but significant reduction in the content of pZAP70 (Y319). Of note, significant reductions in the phosphorylation of these signaling molecules were also seen in Western blots of SMIFH2-treated Jurkats activated using soluble anti-CD3 (Fig. S5, A and B; C shows that KVSM18-treated Jurkats did not show a drop in Lat phosphorylation). Finally, consistent with the data in preceding figures and the work of others (Ilani et al., 2009; Kumari et al., 2012; Yu et al., 2012), Jurkats in which myosin II was inhibited using pnBB phenocopied SMIFH2-treated cells as regards defects in adhesion (Fig. 7 F) and signaling (Fig. 7 H). Together, these results argue that formin-generated actomyosin arcs are required for robust adhesion and TCR proximal signaling in T cell–APC conjugates.
TCR MCs reside between arcs, whereas active LFA-1 coincides with arcs

To further define the physical relationship among actomyosin arcs, TCR MCs, and active LFA-1 in maturing synapses, we engaged Jurkats with bilayers as in Fig. 7. In this case, however, the anti-active LFA-1 antibody M24 was added 2 min after engagement, and cells were fixed at 6 min instead of 10 min so as to capture IS organization during maturation. After staining with phalloidin, the distributions of F-actin (Fig. 8 A, white), TCR MCs (Fig. 8 A, red), and open, active LFA-1 (Fig. 8 A, green) were imaged by 3D-SIM to precisely define the spatial relationships among them. Fig. 8 A (first panel) shows a SIM image of a typical Jurkat 6 min after engagement. Fig. 8 A (second panel) shows an enlargement of a region in this cell’s pSMAC. Fig. 8 A (bottom nine panels) shows the signals in this boxed region as a series of three pairwise comparisons. We made three observations from these comparisons. First, the individual signals for TCR MCs (Fig. 8 A, third panel), F-actin (Fig. 8 A, fourth panel), and their overlay (Fig. 8 A, fifth panel) indicate that TCR MCs (red) tend to reside between actin arcs (white). Second, the individual signals for TCR MCs (Fig. 8 A, sixth panel), active LFA-1 (Fig. 8 A, seventh panel), and their overlay (Fig. 8 A, eighth panel) indicate that TCR MCs (red) also tend to reside between LFA-1-rich structures (green). Finally, the individual signals for F-actin (Fig. 8 A, ninth panel), active LFA-1 (Fig. 8 A, tenth panel), and their overlay (Fig. 8 A, eleventh panel) indicate that active LFA-1 (green) tends to coincide with actin arcs (white). Importantly, these observations were borne out by quantitation. First, Pearson’s coefficient analyses of the entire pSMAC region show that the colocalization of F-actin and active LFA-1 is much more significant than the colocalization of F-actin and TCR MCs, LFA-1 and TCR MCs, and a randomized control in which images were rotated 90° relative to each other (Fig. 8 B). Second, line scans and radial plot profiles across population average projections of individual TCR MCs (Fig. 8, C and D) show overlapping intensity profiles for actin...
Formin-generated actomyosin arcs at the immune synapse

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On average, TCR MCs reside between arcs, whereas active LFA-1 coincides with arcs in maturing synapses. (A) Three-color 3D-SIM imaging of endogenous actin (grayscale), TCR MCs (red), and open, active LFA-1 (green) at a Jurkat T cell IS 6 min after bilayer engagement. (A, bottom nine panels) Pairwise comparisons of the actin, LFA-1, and TCR MC channels and overlays. (B) Pearson’s correlation coefficient; n = 20–40 cells/correlation. Box plots are centered on means and display upper and lower quartile ranges and min to max values. (C) Line scans across population-averaged TCR MCs. n = 110. (D) Radial plot profiles of population-averaged TCR MCs. n = 11 cells; 10–15 TCR MCs/cell. Data are means ± SEM. Note that TCR MCs overlap considerably with F-actin during initial cell spreading (Lam Hui et al., 2014). Bars, 1 µm. ****, P < 0.0001. a.u., arbitrary units; MyoIIA, myosin IIA; Norm. Fluor. Int., normalized fluorescence intensity.

TCR MCs are swept inward by actin arcs

The static SIM images in Fig. 8 demonstrating that TCR MCs commonly reside between actin arcs suggest that TCR MCs may be “swept” inward by these arcs. To look for such sweeping behavior, we used live TIRF-SIM to image Jurkat T cells expressing GFP–F-Actin and engaged with bilayers containing Alexa Fluor 548-labeled anti-CD3 antibody to report the position of TCR MCs. Video 9 and still images taken from it (Fig. 9 B and Fig. S4 B). Together, these highly resolved static SIM images argue that, on average, TCR MCs reside between actin arcs, whereas active LFA-1 (and myosin II) coincide with actin arcs during IS maturation.

by a second arc (Fig. 9 A, fifth through seventh panels, yellow arrowheads), and then slips again to be picked up by a third arc (Fig. 9 A, eighth panel, blue arrowheads). Video 10 and the still images taken from it in Fig. 9 B show a second example of this sweeping behavior, in which a TCR MC (Fig. 9 B, open arrowheads) is swept inward successively by two arcs (Fig. 9 B, first through fourth panels, white closed arrowheads, and fifth and sixth panels, yellow closed arrowheads). Consistent with our previous work (Yi et al., 2012), the rates of inward arc flow and TCR MC movement are not statistically different in these videos (Fig. 9 C), suggesting that the immobile phase arising from slippage is not large enough to significantly uncouple MC movement from arc flow, at least in a bilayer context. Collectively, these findings provide strong evidence that formin-generated, LFA-1– and myosin II–rich actin arcs in the pSMAC sweep TCR MCs inward to facilitate their robust accumulation in the cSMAC.

Discussion

In this study, we used SIM to define the organization of the Jurkat T cell’s actin cytoskeleton in unprecedented clarity and...
demonstrate in unequivocal fashion the existence of actomyosin arcs in the pSMAC region of their IS. By combining SIM imaging with the use of specific inhibitors, we showed how formins and myosin II cooperate to create these actin arcs, and we demonstrated their functional importance. We also used TIRF-SIM to provide the first subdiffraction-limited imaging of actin-driven TCR MC transport. Importantly, we showed that actomyosin arcs are also major components of primary mouse T cell synapses, arguing that they may play significant roles in promoting T cell effector functions in vivo.

Although we presented many lines of evidence that the arcs are formin generated, current models for the formation of anisotropic lamellar actin structures like the arcs described in this study involve not only formin-dependent nucleation, but also the repurposing of the branched actin network in the lamellipodium after debranching and cross-linking (Medeiros et al., 2006; Burnette et al., 2011; Tojkander et al., 2012; Henson et al., 2015; Tee et al., 2015). Although we cannot rule out some contribution to arc formation from this pathway, it appears from our imaging and inhibitor studies that the linear filaments made by formins at the edge of the dSMAC are primarily responsible for arc formation. Regarding the fate of the branched actin network in the dSMAC, the addition of Jasplakinolide, a drug that prevents the depolymerization of actin filaments, causes the rapid formation of a large F-actin ring at the dSMAC/pSMAC boundary (Yi et al., 2012). We think, therefore, that the majority of the dSMAC is disassembled at this boundary, with the ensuing release of monomer supporting continued Arp2/3- and formin-independent actin nucleation at the outer edge of the dSMAC.

With regard to the functional significance of pSMAC arcs, we show in Jurkat T cells that disruption of their formation via formin inhibition results in numerous functional deficiencies, including less centralized TCR MCs, missegregated integrin clusters, decreased T–B cell adhesion frequency, and diminished TCR proximal signaling in conjugates. Moreover, myosin II inhibition phenocopies formin inhibition, consistent with the fact that arcs require myosin II for their organization and contraction. This latter finding is consistent with a subset of previous studies of myosin II function at the IS (Hammer and Burkhardt, 2013). Given that centrosome repositioning is facilitated by formins (Gomez et al., 2007) and helps sustain T cell signaling (Martín-Cófreces et al., 2008), it remains possible that some portion of the defects seen upon formin inhibition are downstream of reduced centrosome repositioning.

Our demonstration that actomyosin arcs are commonly decorated with the open, active conformation of LFA-1 during IS maturation diverges from the recent results of Comrie et al. (2015a), who reported essentially no overlap between active LFA-1 and myosin II at the IS. Although part of this discrepancy could be because of the different T cells studied (Jurkat in this study versus human CD4+), it may have more to do with the time points chosen for imaging (6–10 min in this study versus 30 min). Interestingly, Comrie et al. (2015a) attributed LFA-1 activation primarily to polymerization-driven actin retrograde flow and minimally to myosin II contractility. This conclusion was based in part on their observation that activation was strongly inhibited when retrograde flow was blocked using Jasplakinolide, but not when myosin II contraction was inhibited using blebbistatin. We note, however, that using Jasplakinolide to inhibit actin retrograde flow may also impair actomyosin arc function by rapidly depleting the actin monomer pool, based on our demonstration in this study that this pool is normally limiting for nucleation in T cells. We also note that their use of blebbistatin (no preincubation and addition 5 min after ac-
tivation) may have underestimated myosin II’s role in LFA-1 activation. That said, we also see evidence that actin retrograde flow can promote LFA-1 activation, because open, active LFA-1 was still present at the IS when arc formation had been inhibited (Fig. 7). Nevertheless, our demonstration that the open, active conformation of LFA-1 commonly colocalizes with actomyosin arcs in maturing synapses argues that actomyosin arcs probably play a significant role in integrin activation in T cells, as in mesenchymal cells.

Perhaps our most remarkable observation, made possible by the enhanced spatiotemporal resolution provided by 3D-SIM and TIRF-SIM, was that TCR MCs in the pSMAC appear to be swept inward by the actomyosin arcs. This conclusion has relevance for the longstanding question of how inward actin flow propels the centripetal transport of MCs. Studies by DeMond et al. (2008) and Yu et al. (2010) showed that thwarting the inward movement of MCs using barriers placed in the bilayer causes actin flow to slow transiently at the site of the thwarted MC. This seminal observation argues for a frictional coupling between TCR MCs and a viscoelastic actin cytoskeleton. Our TIRF-SIM videos showing that arcs sweep MCs inward and that slippage events can occur are consistent with this conclusion. Whether the basis for this frictional coupling is simply physical collision between the actin and the MC or involves transient physical tethering between them (or some combination of both) remains to be seen. The imaging described in this study could serve useful in efforts to resolve this latter issue. Finally, these SIM modalities could prove valuable in defining the contributions made by actomyosin force generation to phase-separated cluster formation, ligand discrimination, and mechanotransduction at the IS (Chen and Zhu, 2013; Depoil and Dustin, 2014; Su et al., 2016).

Materials and methods

Cell culture and transfection

E6.1 Jurkat T cells were maintained at 37°C in Iscove’s modified Dulbecco’s media (Thermo Fisher Scientific) supplemented with 10% FBS (Sigma-Aldrich), sodium pyruvate, l-glutamine, penicillin-streptomycin, and MEM nonessential amino acids solution (Thermo Fisher Scientific). Transfections were performed with 1.0 × 10⁶ cells/ml and 50 U/ml recombinant interleukin-2 (R&D Systems) every 48 h. Microbial transduction at the IS

Immunofluorescence

Jurkat T cells were allowed to adhere to the substrate for 7 min at 37°C and then fixed for 15 min in 4% (wt/vol) PFA solution (Electron Microscopy Sciences) in cytoskeletal buffer (300 mM NaCl, 10 mM EGTA, 10 mM glucose, 10 mM MgCl₂, 20 mM Pipes, pH 7.4, and 0.4% sucrose). Samples were then incubated in a blocking solution consisting of 10% FBS (Sigma-Aldrich), 0.01% sodium azide (Sigma-Aldrich), 1× PBS, and 0.2% saponin (Sigma-Aldrich) for 15 min at room temperature (RT). After three washes in 1× PBS, the cells were stained with primary antibody (1:100- to 500-fold dilution in 1× PBS) for 1 h at RT, followed by secondary antibody (1:1,000-fold dilution in 1× PBS) or phalloidin (1:200-fold dilution) for 1 h at RT.

Preparation of supported planar lipid bilayers

Liposomes and glass-supported planar lipid bilayers were prepared as described previously (Varma et al., 2006; Yi et al., 2012; Crites et al., 2015). All lipids were purchased from Avanti Polar Lipids, Inc. Liposomes were created using a mixture of 1,2-dioleoyl-sn-glycero-3-phosphocholine, biotin-CAP-PE (1% molar ratio), and 1,2-dioleoyl-sn-glycero-3-[(N-(5-amino-1-carboxypentyl)iminodiacetic acid)succinyl] (DGS)–NTA (1% molar ratio) lipids. To generate unilamellar liposomes, lipids were lyophilized overnight, reconstituted in nitrogen-purged Tris-saline buffer, sonicated in a water bath, and passed through an extrusion membrane (∼50-nm pore size) using a mini-extruder kit (Avanti Polar Lipids, Inc.). Bilayers were created in flow chambers assembled using glass coverslips that had been cleaned using Piranha solution (70% [vol/vol] sulfuric acid [Sigma-Aldrich]; and 30% [vol/vol] hydrogen peroxide [Thermo Fisher Scientific]). After depositing liposomes onto the flow chambers, the chambers were washed with HBS buffer (4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid buffer saline) and 5% (wt/vol) casein (Sigma-Aldrich). After two washes in HBS buffer, the chambers were stained with primary antibody (1:100- to 500-fold dilution in HBS) and streptavidin (85878; Sigma-Aldrich) were added to the flow chambers along with histidine-tagged ICAM-1, either unlabeled or labeled with Alexa Fluor 647 or rhodamine X–labeled (10 µg/ml) anti-CD3ε antibody and streptavidin (85878; Sigma-Aldrich) were added to the flow chambers along with histidine-tagged ICAM-1, either unlabeled or labeled with Alexa Fluor 647 (0.5 µg/ml). Anti-CD3ε antibody (OKT3; Biovest International Inc.) was monobiotinylated and labeled with fluorescent dyes as described previously (Carrasco et al., 2004). The uniformity and lateral mobility of lipids in the bilayers was assessed by FRAP.

Preparation of stimulatory glass coverslips

Coverslip substrates coated with immobilized antibodies to stimulate Jurkat T cells were prepared as described previously (Bunnell et al., 2003). In brief, eight-well coverglass chamber slides (Lab-Tek; Thermo Fisher Scientific) were washed in HCL and 70% (vol/vol) ethanol. After three washes in 1× PBS, each well was incubated for 30 min at RT with 500 µl of 0.01% (vol/vol) poly-l-lysine (Sigma-Aldrich). Wells were then incubated for 1 h at 37°C with 500 µl of a solution containing 20 µg/ml anti-CD3ε antibody and 20 µg/ml anti-CD28 antibody (BD) diluted in 1× PBS. For primary mouse CD4/CD8 cells, the same procedure was followed except that mouse anti-CD3/CD28 antibodies (BD) were used instead.
Microscopy
3D-SIM was performed on a 3D-SIM Imaging System (GE DeltaVision OMX; Applied Precision Ltd.) equipped with a 60x 1.42 NA objective (Olympus). Raw data were reconstructed using Softworx (Applied Precision Ltd.; GE Healthcare) and 0.003 for the wiener filter constant. TIRF-SIM imaging was performed as described previously (Beach et al., 2014; Li et al., 2015). In brief, images were acquired using an inverted microscope (Axio Observer; ZEISS) fitted with a 100x 1.49 NA objective (Olympus) and a spatial light modulator to provide phase grating for structured illumination (Li et al., 2015). Confocal imaging was performed on an IX81 with a 150× 1.4 NA objective (Olympus) fitted with a spinning disk confocal unit (Yokogawa Electric Corporation). Linear adjustments to images were made using ImageJ software (National Institutes of Health).

Adhesion conjugate assay
The adhesion conjugate assay was performed as previously described (Cannons et al., 2010; Zhao et al., 2012). In brief, Raji B cells were pulsed with 2 µg/ml SEE for 1 h at 37°C and washed three times in medium. Jurkat T cells were mixed with SEE-pulsed Raji B cells at a ratio of 1:4 and incubated for 10 min at 37°C. For myosin II inhibition, Jurkat T cells were preincubated with 50 µM pNbB or DMSO control. For formin inhibition, 25 µM SMI32 treatment was added to the Jurkat T cell and Raji B cell mixture. Samples were fixed using 4% PFA, and stained with FITC-conjugated anti-human CD3 antibody (OKT3; ebioscience) and PE-conjugated anti–human HLA–antigen D related (DR) antibody (LN3; ebioscience). Conjugate frequency was analyzed using flow cytometry (LSR II; BD) after collecting a minimum of 10,000 events, with the OKT3+ HLA-DR+ double-positive population representing Jurkat T–Raji B cell conjugates.

Amnis ImageStream flow cytometry assay
Imaging flow cytometry was performed as previously described (Markey et al., 2015). In brief, Jurkat T and Raji B cells were transfected with GFP- and RFP-farnesyl plasmids, respectively, using a nucleofector kit (Amamax; Lonza). Raji B cells were pulsed with 2 µg/ml SEE for 1 h at 37°C and washed three times in medium. Jurkat T cells were mixed with SEE-pulsed Raji B cells at a ratio of 1:1 and incubated for 10 min at 37°C. Myosin II inhibition was used to preincubate with 50 µM pNbB or DMSO control. For formin inhibition, 25 µM SMI32 was added to the Jurkat T cell and Raji B cell mixture. Samples were fixed using 4% PFA, and stained with FITC-conjugated anti-human CD3 antibody (OKT3; ebioscience) and PE-conjugated anti–human HLA–antigen D related (DR) antibody (LN3; ebioscience). Conjugate frequency was analyzed using flow cytometry (LSR II; BD) after collecting a minimum of 10,000 events, with the OKT3+ HLA-DR+ double-positive population representing Jurkat T–Raji B cell conjugates.

In vitro T cell activation assay
To activate Jurkats using soluble anti-CD3 antibody (Balagopalan et al., 2011), cells were washed with serum-free RPMI 1640, resuspended to a concentration of 3 x 10⁷ cells/ml, and activated with anti-CD3 (1 µg/ml) for the indicated time points. Activation was halted by lysing the cells using 95°C 2x sample buffer and boiling for 3 min. Samples were then passed through a small-gauge syringe needle to shear the DNA and loaded onto a 10% precast polyacrylamide gel (Thermo Fisher Scientific). The separated proteins were transferred to nitrocellulose membrane that was then blocked overnight at 4°C in TBST (10 mM Tris, pH 8.0, 150 mM NaCl, and 0.05% Tween 20) supplemented with 5% milk. All antibodies were diluted in TBST containing 5% milk. Primary antibodies were added to the membrane for 2 h at RT, followed by three 15-min washes in TBST and a 1-h incubation at RT with the appropriate secondary antibody. The proteins were visualized by chemiluminescence. Quantitation of the phosphorylation status of the signaling proteins was performed by measuring the integrated density of the bands on ImageJ (National Institutes of Health) followed by background correction and normalization to an anti–actin antibody (Sigma-Aldrich) loading control.

Image and statistical analyses
All statistical analyses were performed using GraphPad Software. Unless otherwise indicated, all statistical comparisons were performed using unpaired, two-tailed t tests. Box and whiskers plots are centered on the mean and display upper and lower quartile ranges and min to max values. All data represent mean ± SEM except Figs. 4 D and 6 B, which are mean ± SD. To compare the relative pSMAC footprint at the IS of Jurkat T cells versus primary CD8⁺ T cells (Fig. 2 D), the total IS area was determined based on thresholds for F-actin intensity. Outlines for the dSMAC and pSMAC were manually determined using the F-actin channel and superimposed on the thresholded image. The area within the identified regions was measured using ImageJ (National Institutes of Health). To compare the actin spike morphology in Fig. 4 D (bottom panels), the Filotrackr script from MATLAB application CellGeo was used on max projections of phalloidin-stained 3D-SIM images as previously described (Tsyanikov et al., 2014). To qualitatively assess actin arc morphology, maximum projections of 3D-SIM images of phalloidin-stained Jurkat T cells were scored manually into four categories: Organized/concentric, Disorganized/inward pointing, Disorganized/meshwork, and No arcs. Note that Organized/concentric was used to describe cells with robust actin arcs, similar to representative images of control cells. To quantify the content of actin arcs in the pSMAC, maximum projections of phalloidin-stained 3D-SIM images of Jurkat T cells were used to capture the interior of the IS (i.e., everything except the dSMAC). Note that we chose to measure all regions of the IS except the dSMAC because formin inhibited cells do not display enough actin arcs to accurately define a pSMAC/cSMAC boundary. Note also that formin-inhibited cells display an increase in cSMAC phalloidin signal because of an increase in the frequency of F-actin foci. Therefore, our results likely underestimate the loss of pSMAC arc fluorescence upon SMI32 treatment. Imaging parameters across conditions were kept identical. The integrated pixel intensities of the outlined regions of interest for each condition were measured using ImageJ (National Institutes of Health). The FibrilTool plugin for ImageJ was used to measure actin arc morphology as described previously (Boudaoud et al., 2014). In brief, the pSMAC regions in maximum projections of phalloidin-stained 3D-SIM images were divided into 10–12 trapezoid shaped ROIs of similar size to measure the anisotropy of arcs in the radial symmetric pSMAC. To measure the distribution of TCR MCs and LFA-1 at the IS, we used the radial plot profile plugin for ImageJ. The plugin approximated the outline of the
cell based on manual drawing of a quadrangle that best fit the cellular dimensions (identified by F-actin staining). This outline was superimposed on the channel displaying TCR MCs or LFA-1, followed by radial plot profile generation. Radial plot profile intensities were background subtracted and normalized for intensity. To account for cell size variance, cell radii were normalized to 1, and intensities were binned 41 times along the normalized radii. The two-way analysis of variance (ANOVA) was used to test for statistical significance between control and experimental condition datasets. The distribution of TCR MCs and LFA-1 at the IS was also measured by drawing line scans across the IS. After background subtraction, the line scans were normalized as described above for radial plot profiles, but fit to Gaussian distributions and tested for single versus two-Gaussian fit in Prism (GraphPad Software). To compare the spatial associations between F-actin, TCR MCs, LFA-1, and myosin IIA, 3D-SIM images were thresholded using the F-actin channel to find the center of the cell (xy coordinates). The TCR MC channel was then thresholded to identify the MCs and find their centers (xy coordinates). A 40 × 40-pixel box was drawn to crop 10–15 pSMAC TCR MCs per cell. These images were aligned and rotated by measuring the angles between the MC and the center of the cell in Excel using the ArcTan function (Microsoft). The images from each channel were then superimposed to obtain population averaged images for radial plot profiles and line scan analyses as described earlier in this section. To assess colocalizations, Pearson’s coefficient analyses were performed using the JaCop plugin in ImageJ. To measure the speed of centripetal actin and TCR MC movement, kymographs were made from TIRF-SIM videos using the MultiplexKymograph plugin from ImageJ. The dSMAC and pSMAC regions were identified by the relatively abrupt change in slope for F-actin flow, and rates were determined by measuring slope angles.

Online supplemental material

Fig. S1 shows that actomyosin arcs exist in primary mouse CD8+ T cells and that td-Tomato–F-Tractin, but not RFP-actin, labels actin arcs. Fig. S2 shows several forms (including mDia1) are enriched at the tips of actin spikes and that mDia1 knockdown inhibits arc formation. Fig. S3 shows that an activator of Rho GTPases enhances arcs and that actin labels robustly with TM4 but not α-actinin. Fig. S4 shows that myosin II inhibition disrupts receptor cluster distribution and that myosin II coincides with actin and open, active LFA-1 in the pSMAC. Fig. S5 shows that SMIFH2 inhibits signalling in T cells stimulated with soluble antigen and that KVS18 does not inhibit Lat phosphorylation. Video 1 shows the dynamics of F-actin networks at the IS using TIRF-SIM. Video 2 shows the dynamics of actomyosin arcs using two-color TIRF-SIM in a Jurkat. Video 3 shows the formation of linear actin filaments/bundles at the distal edge of the IS. Video 4 shows linear actin filaments/bundles, embedded in the dSMAC, flowing inward and giving rise to pSMAC actin arcs. Video 5 shows the reversible augmentation of linear actin filaments/bundles upon Arp2/3 inhibition. Video 6 shows augmented actin filaments/bundles that flow inward and bend to give rise to actin arcs. Video 7 shows the recovery of actin arc formation upon washout of the formin inhibitor. Video 8 shows that myosin II inhibition disrupts the organization of actin arcs. Videos 9 and 10 show TCR MCs being swept inward by actin arcs. Additional data are available in the JCB DataView@ at http://dx.doi.org/10.1083/jcb.201603080.dv.

Acknowledgments

The authors thank the National Heart, Lung, and Blood Institute FACS Core for help with Amnis ImageStream flow cytometry; Rajat Varma for help with liposome preparations; Henry Higgs, Dan Billadeau, Peter Gunning, David Kovar, and Michael Schell for reagents; and Travis Crites, Alex Ritter, Connie Sommer, Senta Kapnic, Jennifer Cannons, Lakshmi Balagopal, Sarah Hitchcock, Larry Samelson, Pam Schwartzberg, and Clare Waterman for helpful discussions.

This work was supported by Intramural National Heart, Lung, and Blood Institute grant 1ZIAHL006121-04 to J.A. Hammer.

The authors declare no competing financial interests.

Submitted: 23 March 2016
Accepted: 28 September 2016

References


Figure S1. Actomyosin arc formation in a primary mouse CD8+ T cell, and Td Tomato–F-Tractin, but not RFP-actin, reports endogenous, phalloidin-stained actin arcs in the pSMAC. (A) Still images from a two-color TIRF-SIM video of a primary mouse CD8+ T cell expressing GFP–myosin IIA and tdTomato–F-Tractin at the time points indicated after engagement with an activating surface. (B) Radial plot profile showing the distributions of actin and myosin IIA across the IS of a mouse CD8+ T cell 5 min after engagement. SMAC zones are bracketed. Data are represented as mean ± SEM (C) Confocal images of F-actin structures at the IS of a Jurkat T cell expressing RFP-actin and stained with phalloidin. (D) Confocal images of F-actin structures at the IS of a Jurkat T cell expressing tdTomato (tdTom)–F-Tractin and stained with phalloidin. (E) Pearson’s correlation coefficient for tdTomato–F-Tractin versus phalloidin and RFP-Actin versus phalloidin across the pSMAC. n = 15–20 cells/condition. Box and whiskers plot is centered on the mean and display upper and lower quartile ranges and min to max values. The SMAC zones are bracketed in C and D at top. Bars, 5 µm. ****, P < 0.0001. a.u., arbitrary units; Norm., normalized.
Figure S2. Three formins can be detected by immunostaining at the tips of actin spikes in CK666-treated Jurkat T cells, and knockdown of mDia1 greatly attenuates arc formation. (A, first through third panels and magnified insets in fourth through sixth panels) 3D-SIM images of CK666-treated Jurkat T cells immunostained for mDia1, FMNL1, and INF2 non-CAAX and stained with phalloidin. Yellow arrowheads mark formin puncta at spike tips. (A, seventh panel) Percent of spikes enriched with the indicated formin within 1 μm of the tip. n = 10 cells/condition. ****, P < 0.0001 when compared with either mDia2 (black) or preimmune sera (red). Data represent means ± SEM. (B, top left) Western blot of whole-cell extracts of Jurkats 48 h after transfection with either a control shRNA plasmid (lane 1) or an shRNA plasmid for mDia1 (lane 2) and probed for mDia1 and actin as an internal control. (B, right panels) Representative 3D-SIM images of activated shRNA control and mDia1 knockdown Jurkat T cells stained with phalloidin. The yellow arrowheads mark actin foci in knockdown cells. (B, bottom left) Mean phalloidin fluorescence in the pSMAC. n = 10 cells/condition. Box plots are centered on means and display upper and lower quartile ranges and min to max values. Bars, 5 μm. ***, P < 0.001. a.u., arbitrary units; Fluor. Int., fluorescence intensity.
Linear actin filaments in the dSMAC and actin arcs in the pSMAC are accentuated by a pan activator of Rho GTPases, and arcs are enriched for the tropomyosin TM4 but not for α-actinin. (A) Representative 3D-SIM image of a Jurkat cell treated with DMSO (left) or 1 unit/ml CN01 (right), a membrane-permeant pan activator of Rho GTPases. Yellow arrowheads point to the accentuated linear actin filaments/bundles in the dSMAC. (B) Mean phallolidin fluorescence in the pSMAC of control, DMSO-treated, and CN01-treated Jurkats. n = 14–20 cells/condition. Data are means ± SEM (C) The ratio of pSMAC area to total IS area for control, DMSO-treated, and CN01-treated Jurkats. n = 13–24 cells/condition. Data are means ± SEM (D, top) 3D-SIM images of a representative Jurkat T cell immunostained for α-actinin and myosin IIA. (D, bottom) TIRF-SIM images of a representative Jurkat T cell expressing td-Tomato (tdTom)–F-Tractin and GFP-TM4. (E) Radial plot profiles of myosin IIA, TM4, and α-actinin in Jurkat T cells. n = 7 cells; P < 0.0001, two-way ANOVA. (F) Pearson’s correlation coefficient (corr. coeff.). n = 10–16 cells/correlation. Box plots are centered on means and displays upper and lower quartile ranges and min to max values. ***, P < 0.001; ****, P < 0.0001. a.u., arbitrary units; Fluor. Int., fluorescence intensity; Norm., normalized. Bars, 5 μm.
Figure S4. Myosin II inhibition phenocopies formin inhibition with regard to receptor cluster distribution, and myosin II tends to coincide with actin and open, active LFA-1 in the pSMAC. (A, first and second panels) Radial plot profiles of TCR MCs and LFA-1 clusters in Jurkat T cells treated with DMSO or 50 µM pnBB. n = 13 cells; P < 0.0001, two-way ANOVA. (A, third panel) Line scans across cells in A2 fit to Gaussian distributions. (B, top) Four-color 3D-SIM of F-actin (white), TCR MCs (red), open, active LFA-1 clusters (green), and myosin IIA (myoIIA; blue). (B, bottom nine panels) Individual combinations of the actin, myosin IIA, and open, active LFA-1 channels and their overlays. See also Fig. 8 B. Bars, 1 µm. a.u., arbitrary units; Norm. Fluor. Int., normalized fluorescence intensity.
Figure S5. **SMIFH2-treated Jurkats activated with soluble anti-CD3 also exhibit a reduction in TCR proximal signaling, and KVSM18 does not inhibit Lat phosphorylation.** (A) Representative Western blot of whole-cell extracts of DMSO- and SMIFH2-treated Jurkats activated for the indicated times using soluble anti-CD3 and probed for the tyrosine-phosphorylated versions of Src (Y416), Zap70 (Y319), or Lat (Y191). (B) Quantitation of relative phosphoprotein amounts. $n = 3$ independent experiments. Data are means ± SEM. (C) Western blot of whole-cell extracts of DMSO- and KVSM18-treated Jurkats activated and probed with anti-Lat as described in A. *, $P < 0.05$; **, $P < 0.01$. WT, wild-type.

Video 1. **TIRF-SIM reveals the dynamics of F-actin networks at the Jurkat T cell IS.** Jurkat T cell expressing GFP–F-Tractin faithfully reports the two major actin networks at the IS seen with phalloidin staining. Images were collected every 2 s for 1.5 min. See Fig. 1.

Video 2. **Two-color TIRF-SIM reveals the dynamics of actomyosin arcs in the pSMAC.** Jurkat T cell expressing GFP–myosin IIA and tdTomato–F-Tractin showing enrichment of myosin IIA on concentric actin arcs in the pSMAC. Images were collected every 3 s for 2 min. See Fig. 1.

Video 3. **Linear actin filaments/bundles are embedded in the dSMAC.** TIRF-SIM video of a Jurkat T cell expressing GFP–F-Tractin showing the formation of several linear actin filaments/bundles at the distal edge of the IS that span the dSMAC. Images were collected every 1 s for 1 min. See Fig. 3.
Video 4.  **Linear actin filaments/bundles appear to give rise to actin arcs in the pSMAC.** TIRF-SIM video of a Jurkat T cell expressing GFP–F-Tractin showing several linear actin filaments/bundles flowing inward and giving rise to actin arcs in the pSMAC. Images were collected every 3 s for 5 min. See Fig. 3.

Video 5.  **Arp2/3 inhibition reversibly augments linear actin filaments/bundles.** Spinning-disc confocal video of a Jurkat T cell expressing GFP–F-Tractin, before and after treatment with 25 µM CK666, followed by CK666 washout. Images were collected every 2 s for 9 min. See Fig. 4.

Video 6.  **Arp2/3 inhibition augments linear actin filaments/bundles.** TIRF-SIM video of Jurkat T cell expressing GFP–F-Tractin after treatment with 25 µM CK666. Images were collected every 3 s for 2.5 min. See Fig. 4.

Video 7.  **Linear actin filaments/bundles and actin arcs rapidly reappear upon washout from formin inhibition.** TIRF-SIM video of Jurkat T cell expressing GFP–F-Tractin shortly after the washout of 10 µM SMIFH2. Images were collected every 3 s for 5 min. See Fig. 5.

Video 8.  **Myosin II inhibition disrupts actin arc organization.** TIRF-SIM video of a Jurkat T cell expressing tdTomato–F-Tractin pretreated with 50 µM pnBB. Images were collected every 3 s for 2 min. See Fig. 6.

Video 9.  **TCR MCs are swept inward by actin arcs.** TIRF-SIM videos of Jurkat T cells expressing GFP–F-Tractin stimulated on planar lipid bilayers containing Alexa Fluor 568–labeled anti–CD3 antibody (red) to report TCR MC movement. Video shows a single TCR MC (white open arrowhead) being swept inward from right to left by three successive arcs (white closed, yellow closed, and blue closed arrowheads, respectively, outlined momentarily by yellow lines). The cSMAC of this cell is to the left. Images were collected every 3 s for 1 min 39 s, respectively. See Fig. 9.
Video 10. **TCR MCs are swept inward by actin arcs.** TIRF-SIM videos of Jurkat T cells expressing GFP–F-Actin stimulated on planar lipid bilayers containing Alexa Fluor 568–labeled anti-CD3 antibody (red) to report TCR MC movement. Movie shows a single TCR MC (white open arrowhead) being swept inward from top to bottom by two successive arcs (white closed and yellow closed arrowheads, respectively, outlined momentarily by yellow lines). The cSMAC of this cell was toward the bottom. Images were collected every 3 s for 1 min 39 s, respectively. See Fig. 9.