

Review

Inside view of cell locomotion through single-molecule: fast F-/G-actin cycle and G-actin regulation of polymer restoration

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Abstract: The actin cytoskeleton drives cell locomotion and tissue remodeling. The invention of live-cell fluorescence single-molecule imaging opened a window for direct viewing of the actin remodeling processes in the cell. Since then, a number of unanticipated molecular functions have been revealed. One is the mechanism of F-actin network breakdown. In lamellipodia, one third of newly polymerized F-actin disassembles within 10 seconds. This fast F-actin turnover is facilitated by the filament severing/disrupting activity involving cofilin and AIP1. Astoundingly fast dissociation kinetics of the barbed end interactors including capping protein suggests that F-actin turnover might proceed through repetitive disruption/reassembly of the filament near the barbed end. The picture of actin polymerization is also being revealed. At the leading edge of the cell, Arp2/3 complex is highly activated in a narrow edge region. In contrast, mDia1 and its related Formin homology proteins display a long-distance directional molecular movement using their processive actin capping ability. Recently, these two independently-developed projects converged into a discovery of the spatiotemporal coupling between mDia1-mediated filament nucleation and actin disassembly. Presumably, the local concentration fluctuation of G-actin regulates the actin nucleation efficiency of specific actin nucleators including mDia1. Pharmacological perturbation and quantitative molecular behavior analysis synergize to reveal hidden molecular linkages in the actin turnover cycle and cell signaling.

Keywords: single-molecule imaging, actin turnover, mDia1, pharmacokinetic simulation, G-actin

1. Introduction

The actin cytoskeleton is the major component beneath the surface of eukaryotic cells. Remodeling of the actin cytoskeleton is directly coupled to cell shape changes. To accomplish body functions such as morphogenesis, migration of immune cells, formation of the neural network, endocytosis, exocytosis

and cytokinesis, cells must reorganize cell surface structures through mobilization of the actin remodeling machinery. The actin cytoskeleton not only facilitates transformation of cell structures but also generates forces that drive translocation of the whole cell body and even tissue remodeling.

Recent development of live-cell fluorescence single-molecule imaging has given us a window for direct viewing of cellular biochemical reactions.¹⁾ Currently, the prevailing method to examine molecular functions is to knock out or knock down gene expression and to observe resultant phenotypes. For example, my research group has recently carried out comprehensive screening to elucidate the role of Rho family GTPases in the PDGF-induced chemotaxis of fibroblast cells. This study identified Cdc42, Rac1 and RhoG as crucial regulators of fibroblast chemotaxis. Knock down of three GTPases had severe and distinct morphological effects on chemotactic cells.

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Abbreviations: CCD: charge-coupled device; Formins: formin homology proteins; PAF: photoactivation of fluorescence; FRAP: fluorescence recovery after photobleaching; CP: capping protein; qFSM: quantitative fluorescence speckle microscopy; EGFP: enhanced green fluorescent protein; Pi: γ -phosphate; FH1: formin homology 1; FH2: formin homology 2; Arp2: actin related protein 2; LatB: latrunculin B.

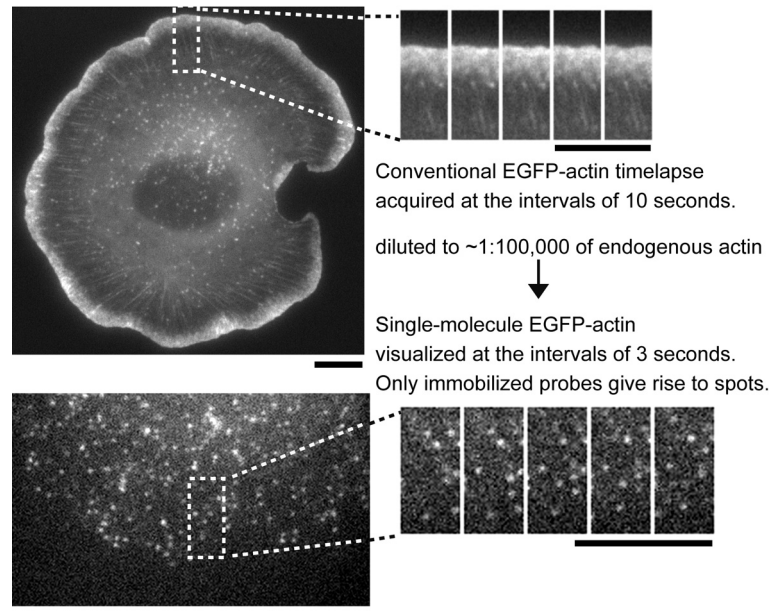


Fig. 1. Overview of fluorescence single-molecule imaging. The upper panels show conventional live-cell imaging of EGFP-actin in a *Xenopus* fibroblast cell. A part of the timelapse images are paneled on the right. The retrograde flow of bulk F-actin structures can be visualized although movement of individual F-actin subunits and local F-actin assembly and disassembly can not be assessed by the conventional imaging of GFP probes. The lower panels show fluorescence single-molecule imaging employing the same EGFP-actin probe. The density of labeled probes is much less than that in the upper example. Low noise CCD cameras can capture the images of individual cell structure-anchored fluorochromes with long exposure times, typically 0.2 to 2 seconds. Signals from soluble probes are blurred. By this way, the association-dissociation kinetics of cytoskeletal proteins can be monitored with high resolution. Scale bars, 5 μm .

Interestingly, impaired chemotaxis in all situations arises from the reduced cell migration speed but not loss in the directionality.²⁾ As seen in this example, genetic manipulation provides a powerful means to elucidate molecular functions. However, due to its slowness, genetic manipulation does not always clarify the primary site of action of the molecule. Success in elucidating mechanistic aspects of complex biological systems such as actin-based motility relies on the methods enabling real time monitoring of the process.

In 2002, my study with Tim Mitchison convincingly demonstrated that intracellular molecules can be visualized at the single-molecule level.³⁾ It was fortunate that my research subject was actin filament dynamics. The assembly to disassembly step of cytoskeletal proteins is a suitable target for fluorescence single-molecule observation. With the aid of the low-noise charge-coupled device (CCD) cameras, it is possible to visualize a single fluorochrome attached to filamentous actin (F-actin). Signals from a non-diffusing fluorochrome add up in a small spot on the CCD chip with the exposure time of ~ 2 seconds. In

contrast, signals from a freely diffusing fluorochrome are blurred on the image. Thus, cytoskeletal proteins which cease random walk for a certain period can be visualized as single-molecules in a state-specific manner (Fig. 1).

Using this approach, my research has elucidated detailed F-actin turnover kinetics at the leading edge of fibroblasts³⁾ as well as F-actin association kinetics of major actin regulators.^{4),5)} It is now possible to compare kinetics data obtained in living cells directly with data from *in vitro* actin biochemistry. In a number of cases, single-molecule observation has revealed unexpectedly fast actin remodeling processes. In the former part of this review, I overview the molecular kinetics involved in breakdown of the actin network.

In the latter, I describe the feedback F-actin restoration mechanism regulated by monomeric actin (G-actin). In the cell, actin exists in two forms, monomeric G-actin and filamentous F-actin (Fig. 2). F-actin turnover is very dynamic. In cultured cells, the conversion between G- and F-actin occurs in every a half to a few minutes.^{3),6)-8)} G-actin and F-actin

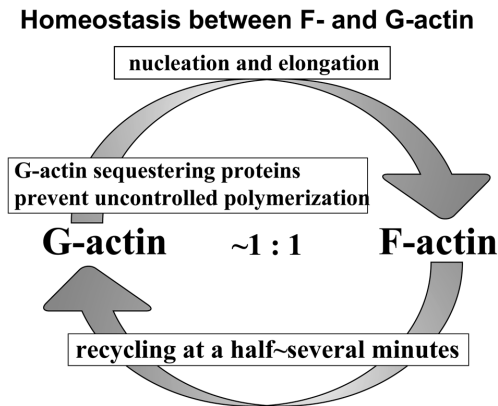


Fig. 2. Homeostasis between G-actin and F-actin. In typical cultured cells, conversion between monomeric ‘G-actin’ and filamentous ‘F-actin’ occurs in every a half to a few minutes. G-actin and F-actin exist roughly at the 1:1 ratio. G-actin sequestering proteins bind G-actin and prevent uncontrolled actin nucleation by lowering the concentration of ‘free’ G-actin. The concentration of total G-actin may exceed 100 μM while the concentration of free G-actin is kept low at a range of 0.1–1 μM .^{9),170)} Because of the large size of the G-actin pool, little has been investigated about the role of the G-actin pool in actin remodeling. However, my two independently-developed researches have converged into an intriguing finding that the actin disassembly and F-actin nucleation activities are tightly coupled in times and places within the cell.⁹⁾

exist roughly at the 1:1 ratio. The cellular concentration of G-actin may exceed a hundred micromolar. Similarly abundant G-actin sequestering proteins prevent G-actin from uncontrolled polymerization. The large mass of the G-actin pool and its sequestering proteins has kept researchers’ eyes off their regulatory roles. However, two independently-developed projects in my research recently converged into a discovery of the dynamic association of filament nucleation to actin disassembly mechanisms in the cell.⁹⁾ I discuss the spatiotemporal coupling between actin disassembly and reassembly through the concentration fluctuation and heterogeneity of free G-actin.

2. Force generation by actin system in non-muscle cells

To generate the force to drive cell locomotion, the actin cytoskeleton operates two major force-generating systems.¹⁰⁾ One uses the actin polymerization force and the other is the actomyosin contraction mechanism. The leading edge of motile cells

forms two morphologically distinct pseudopods called filopodia and lamellipodia. In both, actin forms polarized arrays of filaments directing fast-growing barbed ends towards the cell edge.^{11),12)} In lamellipodia, F-actin free from barbed end inhibitors such as capping protein exists in abundance.^{4),13)–15)} Many free barbed ends are presumably in contact with the leading edge plasma membrane. Due to the flexible nature of either the plasma membrane¹⁶⁾ or the lateral bending of actin filaments,¹⁷⁾ G-actin is allowed to occasionally assemble onto the barbed end in contact with the plasma membrane. Such insertion events lead to generation of force pushing the cell edge forward. This mechano-chemical coupling has been demonstrated using pharmacological intervention of actin elongation¹⁸⁾ and theorized in the context of the Brownian ratchet mechanism.¹⁹⁾

In the more central region of the cell at the back of lamellipodia, actin forms an array of long, often anti-parallel filaments. This type of F-actin is connected to the extracellular matrix and the neighboring cells, through actin stress fibers to focal adhesions and circumference bundles to adherence junctions, respectively. The conventional type-II myosin associating with such actin arrays generates contractile forces. Cells exhibiting the amoeboid mode of migration use the actomyosin mechanism.²⁰⁾ Actomyosin also plays a crucial role in migration of epithelial sheets during embryo morphogenesis.²¹⁾ In addition, blebbing, the third form of cell edge protrusion, is driven by actomyosin.^{22),23)} The local cortical rupture as a consequence of increased actomyosin tension²⁴⁾ or uneven transient pressure formed by a poroelastic cytoplasm²⁵⁾ may lead to the formation of blebs. Blebbing is frequently observed in cells exhibiting amoeboid migration and cytokinesis accompanied by an increase in cortical tension.

Here I propose another potent regulatory step conducted by actin polymerization that is the formation of structure-bridging actin filaments. Thus far, much of our attention to the role of actin polymerization has been paid in cell edge protrusion. However, in the central region, F-actin turnover is ubiquitously found.^{7),26)} Anti-parallel F-actin structures such as actin stress fibers and the cytokinetic contractile ring provide the scaffold for conventional myosin. The amount of myosin-generated force and its transmission should be modulated by the density and the polarity of surrounding actin filaments. Therefore, assembly of cell structure-bridging F-actin structures

would be a vital step to control the force generation for cell locomotion and tissue remodeling.

Under many conditions, rapid remodeling of actin stress fibers is observed in cultured cells. Extracellular ligands such as lysophosphatidic acid stimulate the rapid formation of actin stress fibers. Upon wounding the chick embryonic skin, thick actin cables are formed at the front of the marginal epithelial cells within 5 min.²⁷⁾ Rho, a small GTPase, is a molecular switch in the formation of actin stress fibers²⁸⁾ and the contractile ring during cytokinesis.^{29),30)} In mid 90's, my work in graduate school identified mDia1, a member of formin homology proteins (Formins) as an effector of Rho.³¹⁾ Further analysis revealed that its FH1-FH2 unit structure, which is conserved among Formins, is capable of inducing the formation of long thin actin fibers.³²⁾ These actin fibers often bridge across the diameter of the cell. This activity cooperates with ROCK, a Rho effector kinase that upregulates actomyosin contractility,^{33)–37)} to induce morphologically-varied actin fibers. It is noteworthy that overexpression of active mDia1 induces the bleb formation in a fraction of cells,³²⁾ suggesting a close link between mDia1 and actomyosin-derived cortical tension. Thus, cell morphology-based analysis provided a number of insights into the function of mDia1. At the same time, I realized the difficulty in capturing the precise site of action of this potent actin fiber inducer because of the complex nature of cellular F-actin remodeling. Direct viewing of the molecular behavior of mDia1 and its related molecules has now turned out to be a powerful tool to clarify how structure-bridging actin arrays are generated.

3. F-actin turnover measurement: different interpretations

Before the invention of live-cell fluorescence single-molecule observation,³⁾ cellular F-actin turnover had been characterized mainly by two methods. One observes primary incorporation sites of newly-administrated actin probes. Researchers fixed cells 20–30 seconds after microinjection of labeled actin and observed distribution of the probe incorporation either by electron microscopy¹⁴⁾ or by fluorescence microscopy.¹³⁾ These studies observed that the extreme leading edge of the cell assembles F-actin at the fastest rate.

The other approach uses photoactivation of fluorescence (PAF) of caged fluorescent actin^{6),8)} or

fluorescence recovery after photobleaching (FRAP) of fluorescently-labeled actin.^{7),38),39)} These methods simultaneously visualize movement and disintegration of the cytoskeletal network. In an early study, Yu-li Wang³⁸⁾ discovered the continuous centripetal movement of the lamellipodium actin network which is called the retrograde actin flow. This finding illuminated the main body of the long known centripetal movement of the cell peripheral architecture.^{40),41)} This centripetal actin flow is thought to serve for adhesion-induced cell edge protrusion, as proposed in a hypothetical model called the 'clutch model'.⁴²⁾ In this model, strengthening linkage between cell adhesion molecules and the retrograde actin flow reduces the flow speed. If the filament growth rate is unaltered at the leading edge, the reduction in the actin flow rate may lead to cell edge extension. Recently, the interaction either between Shootin-1 and L1-CAM⁴³⁾ or between N-cadherin and F-actin⁴⁴⁾ has been proposed to constitute the molecular 'clutch' to promote neurite outgrowth. Since this massive actin flow is found in most of cultured cells, there must be more adhesion molecules that function in a similar manner. Notably, a similar flow of E-cadherin complexes between two adjacent cells is also found at the cell-to-cell junction.⁴⁵⁾ The relationship between this 'physical' clutch mechanism and the adhesion-induced 'chemical' cell signaling remains to be clarified.

With regard to F-actin turnover, an important notion was brought by PAF experiments using caged fluorescent actin. Based on the F-actin disintegration faster than the filament density decay expected from the retrograde flow rate, Theriot and Mitchison proposed that actin frequently polymerizes away from the leading edge.⁶⁾ This proposal contradicted the treadmilling model which had prevailed in the research field. Actin forms a polarized filament with the barbed end growing and the pointed end shrinking at steady state,⁴⁶⁾ which is referred to as treadmilling. At the leading edge, the majority of actin filaments direct their fast-growing barbed ends toward the cell edge.¹¹⁾ F-actin assembles the fastest at the leading edge^{13),14)} and then flows inward.^{18),38)} From these observations, the treadmilling model, in which actin polymerizes exclusively at the leading edge while disassembling at the back of lamellipodia, was widely accepted. The proposal by Theriot and Mitchison provoked debate over the treadmilling and their nucleation release model.^{6),47)} I reinvestigated this issue using fluorescence single-molecule

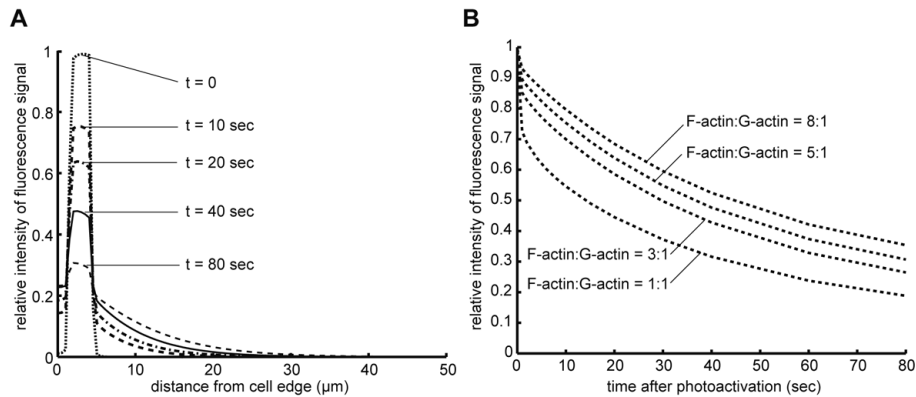


Fig. 3. Simulation of PAF (photoactivation of fluorescence) experiments using the Tardy model: a delay in the recovery kinetics due to local reincorporation. Tardy *et al.*⁵²⁾ mathematically analyzed actin-based PAF and FRAP experiments incorporating the exchange between F-actin and G-actin. Although the model does not consider uneven distribution of F-actin and the retrograde actin flow, the early decay or recovery kinetics in the local cell environment can be estimated by this model. (A) The time-resolved distribution of PAF signals in a 50 μm wide cell. The photoactivation label is placed between 1.5 μm and 4.5 μm away from the cell edge. Diffusion constant of G-actin, the turnover rate from F-actin to G-actin and the ratio between F- and G-actin are $4.0 \times 10^{-8} \text{ cm}^2/\text{s}$,⁷⁾ 0.03 s^{-1} and 5:1, respectively. Although the overall ratio between F- and G-actin is about 1:1, F-actin exists at $\sim 1000 \mu\text{M}$ in lamellipodia,⁵³⁾ which should lead to a local high F-/G-actin concentration ratio. (B) Dependency of PAF signal decay on the ratio between F- and G-actin. The conditions are the same as above except for the F-/G-actin ratios. Note that the simulated decay rate of PAF labels becomes substantially slower (apparent $T_{1/2}$ is approximately 30~50 sec) than the true F-actin disassembly rate ($T_{1/2} = 23 \text{ sec}$).

observation with Mitchison, and found that about one third of newly assembled F-actin subunits had a short lifetime less than 10 seconds in lamellipodia of *Xenopus* XTC fibroblasts.³⁾ Since it takes 2–5 minutes for the actin network to travel through the entire width of a lamellipodium, the results support Theriot and Mitchison's conclusion. The fast disassembling population of F-actin was also observed by a computer assisted fluorescence speckle analysis of X-rhodamine-labeled actin.²⁶⁾

FRAP experiments often show slow recovery kinetics of F-actin in lamellipodia. A recent study³⁹⁾ strongly argues that FRAP data showing slow recovery of the photobleached actin are inconsistent with single-molecule observation results showing fast disassembling population.³⁾ This discrepancy might arise in part from the intrinsic problems in FRAP and PAF experiments.

First, single-molecule analysis detects fast F-actin species repeatedly within the observed time window. By contrast, FRAP and PAF analyses measure the recovery kinetics of the label existing at a given moment. Hence direct comparison between single-molecule lifetime distribution and the FRAP/PAF decay rate may not be valid. Reinterpreted to the FRAP/PAF situation by weighing lifetime

of each population, the single-molecule lifetime data yield that at a given moment, nearly a half of F-actin subunits have lifetime of >100 seconds. F-actin forms high order structures and its stability is enhanced by association with tropomyosin,^{48),49)} fascin^{8),50),51)} and so on.

Second, as shown by mathematical simulation,⁵²⁾ reincorporation of dissociated labels into the marked zone in FRAP and PAF experiments may retard the decay rate substantially. In lamellipodia, F-actin exists at $\approx 1000 \mu\text{M}$ ⁵³⁾ and therefore the ratio of F- and G-actin could become as high as 5~10:1. In such situations, the reincorporation mechanism retards the apparent FRAP decay ~ 2 times slower than true actin disassembly as duration in the G-actin state becomes very short in the actin turnover cycle (Fig. 3). These two problems need special attention when FRAP/PAF analysis is applied for molecules which rapidly switch between fast and slow diffusion states.

In addition, as I discuss in the following sections, actin disassembly may proceed in part through release of actin oligomers.⁴⁾ This mechanism, which is now under investigation in my research group, may further slow actin diffusing out of the FRAP and PAF areas.

4. Single-molecule kinetics of F-actin turnover and actin regulators in lamellipodia

I summarize single-molecule kinetics of actin and its regulators in a single defined cell system, XTC *Xenopus* fibroblast cells. These cells are suitable for live cell imaging as they grow at room temperature in ambient atmosphere. A condition where cells form a wide flat lamellipodia was developed.³⁾ Since thickness of lamellipodia is less than 200 nm,⁵³⁾ it is possible to observe molecules in the entire lamellipodium using conventional epifluorescence microscopy. This method can also be applied to other mammalian cell types with modification.⁴³⁾ I have found that several types of mammalian cells including mouse NIH3T3 cells and rat 3Y1 cells tend to lose activity rapidly under strong xenon illumination for unknown reasons (unpublished observations).

The dendritic nucleation model is believed to constitute the major mechanism controlling lamellipod actin dynamics.^{54),55)} F-actin is nucleated by Arp2/3 complex off the side of preexisting filaments.^{56)–59)} Those nucleated filaments grow outward as actin filaments in lamellipodia direct their fast growing barbed end toward the cell periphery.¹¹⁾ Growth of barbed ends is terminated by capping protein.⁶⁰⁾ The filament network moves toward the cell center and disassembles.

Single-molecule observation of EGFP-actin revealed F-actin turnover kinetics in lamellipodia of XTC fibroblasts in detail. The observation of single-molecule fluorescent actin revealed that (i) 34% of newly assembled filaments disassemble within 10 seconds, (ii) F-actin lifetime distributes over a wide range of 4–148 seconds and (iii) F-actin migrates parallel to each other except for subtle local deformation of the actin network (See Supplementary Movie 3 in ref. 3). The observation (iii) is expected from the dense, crosslinked filament network in EM studies.^{47),61)} The observation (iii), however, does not agree with the conclusion drawn in a study led by Waterman-Storer and Danuser which claimed a large population of long-lived F-actin migrating at a slow speed in lamellipodia.²⁶⁾ Their method called qFSM (quantitative fluorescence speckle microscopy) analyzes behavior of fluorescence speckles, a cluster of fluorochromes, by computer-assisted algorithms. Recently, it was disputed that several manual or automatic analyses, designed to eliminate tracking errors, fail to detect such long-lived, slowly-migrating F-actin species in the actin speckle images.⁶²⁾ In general,

errors in automatic particle tracking may dramatically increase at high particle densities and for molecules that appear and disappear quickly. Moreover, for particles localized in narrow cell structures such as focal adhesions, correlation-based tracking may detect displacement of structures but not of the particles if the label density is too high. Therefore, care also needs to be taken in the interpretation of other qFSM-based flow rate analyses for focal adhesion proteins,⁶³⁾ actin regulators⁶⁴⁾ and so on. I suggest that reanalysis using image samples labeled at various particle densities may help understand the mechanics of error generation by computer-assisted algorithms. I also suggest that reanalysis by single-molecule observation may be required for validating the conclusions drawn by qFSM approaches. For such a purpose, it is important to carefully develop conditions that eliminate phototoxicity problems in each cell system.

Apart from these possible tracking problems, the original actin speckle images provided by Ponti *et al.*²⁶⁾ and other studies^{65),66)} show coexistence of quickly appearing and disappearing actin speckles in the body of lamellipodia, which is consistent with single-molecule observation of EGFP-actin.³⁾ The average dissociation rate of F-actin is $\approx 0.03 \text{ s}^{-1}$. Slightly faster is the dissociation of Arp2/3 complex from the actin network which occurs at 0.048 s^{-1} (Fig. 4). Thus the lag between uncapping at the pointed end and disassembly of the entire filament is very limited. On the other hand, the dissociation of capping protein (CP) occurs astoundingly fast at 0.58 s^{-1} . Importantly, this fast CP dissociation is specific to *in vivo* as the EGFP-tagged capping protein probe tightly caps the barbed end with the dissociation rate of 0.005 s^{-1} *in vitro*, which is comparable to native capping protein.⁶⁷⁾ The recent study, which strongly argued that lamellipod actin turnover is slow,³⁹⁾ also detected fast dissociation of CP with $T_{1/2}$ of 7.2 seconds using FRAP. Thus the previously prevailing view that CP blocks barbed end growth until the end of lifetime of its bound filament needs to be revised. None of the other barbed end interacting proteins, Eps8, VASP, gelsolin,⁴⁾ wild-type mDia1⁹⁾ and AIP1,⁵⁾ shows persistent association with the lamellipod actin network. Consistently, the concentration of free barbed ends is high at $\sim 1 \mu\text{M}$ in lamellipodia of permeabilized XTC cells.⁴⁾ Furthermore, the actin elongation rate determined from the speed of processive actin polymeriza-

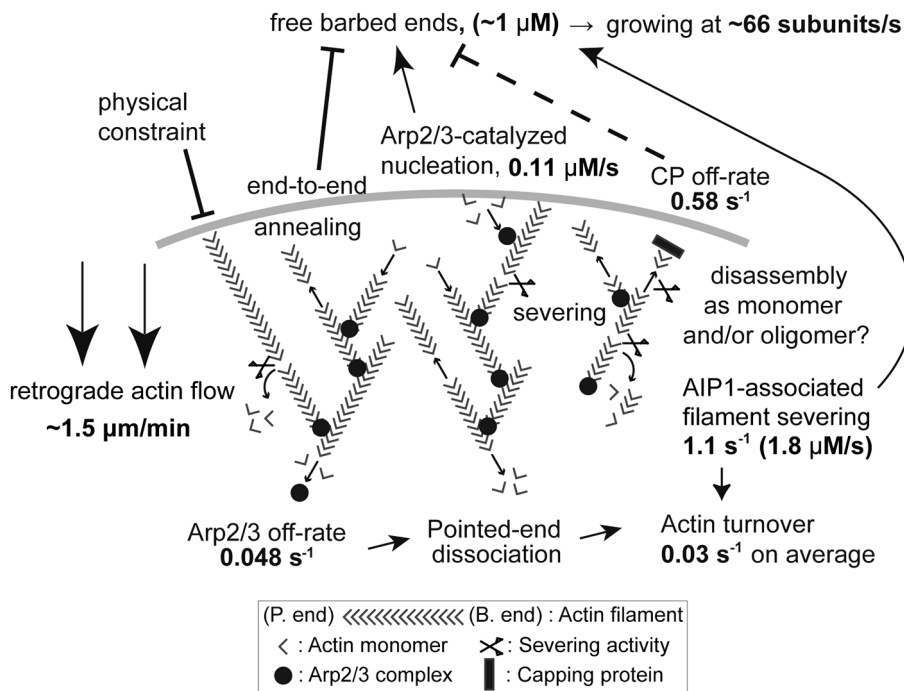


Fig. 4. Summary of single-molecule kinetics in the dendritic actin network. The numbers in bold letters show kinetic parameters obtained in fluorescence single-molecule studies.³⁾⁻⁵⁾ Overall, dynamics at the barbed end has turned out to be faster than previously thought. The dissociation of capping protein occurs at 0.58 s^{-1} . Fast elongation of the free barbed end is derived from the speed of mDia1-catalyzed processive actin elongation in lamellipodia. In contrast, the pointed end side of F-actin, where Arp2/3 complex associates, disassembles much slower at the rate of 0.048 s^{-1} . Certain mechanisms other than simple treadmilling must compensate the imbalance between fast growing barbed ends with slowly disassembling pointed ends. AIP1-mediated actin severing/disruption might be one of the mechanisms. It turned out to be ≈ 15 times more frequent than F-actin nucleation by Arp2/3 complex.

tion by the mDia1 FH2 domain^{4),68)} and its biochemical property⁶⁹⁾ is very fast (66 s^{-1}). Combined, growth of barbed ends in lamellipodia is not restricted as thought before. Mathematical simulation of the treadmilling model using the above parameters points to the requirement of >90 fold acceleration of pointed end disassembly (unpublished results). *In vitro*, pointed end disassembly can be accelerated by a maximum of 22–30 fold with ADF and less efficiently with cofilin.^{70),71)} Thus the single-molecule kinetics data suggest a mechanism other than the treadmilling model in which cofilin/ADF disassembles F-actin solely from the pointed end.

5. How might filament disassembly proceed in the cell?

1) Recent advance in actin depolymerization biochemistry. F-actin disassembly to G-actin recycling remains one of the most obscurely understood steps in the actin turnover cycle. The

cofilin/ADF family^{72),73)} is essential in cellular F-actin turnover.^{4),74),75)} However, how actin disassembly proceeds (i.e. end depolymerization versus filament severing, the barbed end versus the pointed end) remains an open issue. Whether F-actin is frequently severed or not has attracted researchers' interest because severing is a measure for the cell to generate newly-growing filaments. Currently it remains elusive whether cofilin/ADF functions exclusively by accelerating pointed end disassembly or its weak 'non-catalyzing' filament severing activity⁷⁶⁾ substantially contributes to actin disassembly processes. In addition, cofilin also promotes filament nucleation at a high concentration.⁷⁷⁾

Cofilin/ADF by itself may not accomplish the fast actin disassembly observed in cells. Recent work, however, succeeded to purify a highly active tri-component actin disassembling system consisting of cofilin, AIP1 and coronin from calf thymus.^{78),79)} The high actin disassembling activity by these three

proteins appears to be promising towards the goal to elucidate the fast actin recycling mechanism.

AIP1 (actin interacting protein 1) was originally identified as a binding partner of both actin and cofilin (reviewed in ref. 80). AIP1 shows genetic interaction with cofilin in *yeast*,^{81),82)} *C. elegans*⁸³⁾ and *Drosophila*,⁸⁴⁾ where they collaborate to disassemble F-actin.^{85),86)} In *mouse*, two amino acid deletion in AIP1 causes neutrophil dysfunction and macrothrombocytopenia.⁸⁷⁾ AIP1 caps the barbed end in a manner dependent on cofilin and inhibits elongation.⁸⁸⁾ AIP1 does not prevent reannealing of mechanically fragmented filaments.⁸⁸⁾ Hence AIP1 specifically recognizes and caps the barbed end generated by cofilin-catalyzed filament severing.

Recently, bipolar effects of coronin on cofilin-catalyzed actin disassembly have been demonstrated.⁸⁹⁾ Coronin prevents cofilin-induced severing of F-actin assembled from ATP-actin but promotes severing of F-actin assembled from ADP-actin. The authors interpreted that coronin protects ATP/ADP-Pi-F-actin from cofilin while helping cofilin sever the filament after ATP hydrolysis and γ -phosphate (Pi) release. Although this work provided important notions, their data (Fig. 2B in ref. 89) might not simply agree with the author's interpretation. Hydrolysis of Mg-ATP occurs quickly upon polymerization and Pi release proceeds over minutes.⁹⁰⁾ Pi release is prerequisite for filament disassembly because Pi stabilizes F-actin.⁹¹⁾ Pi release is greatly accelerated by cofilin.⁹²⁾ At the start of the above filament disassembly experiments, a large fraction of F-actin should already be ADP-F-actin. Therefore coronin's bipolar effects may not simply depend on the nucleotide bound to F-actin. Their microscopic observation (Fig. 2G and Movies S3 and S4 in ref. 89) shows weak F-actin bundling activity of coronin. I therefore speculate that the inhibitory effects of coronin on ATP-F-actin disassembly might be caused by other mechanisms such as coronin-induced filament bundling.⁹³⁾

It is also known that coronin interferes with Arp2/3 complex-induced actin nucleation^{94),95)} and promote debranching.⁹⁶⁾ These effects on Arp2/3 complex are weak (typically, ≈ 2 fold changes) even with a saturating concentration of coronin (50–500 nM).^{94),96)} Coronin may not exist free at these concentrations in cells because the dissociation constant of coronin with F-actin is $\approx 6 \times 10^{-9}$ M.⁹⁷⁾ Thus these activities toward the Arp2/3 complex may not

be the primary function of coronin under intracellular environment. Rather, the following actin disassembly activity might be the more important function of coronin because coronin, cofilin and AIP1 synergize effectively.

When combined together, cofilin, AIP1 and coronin achieve fast actin disassembly in a manner insensitive to polymerizable monomer.⁷⁸⁾ This system facilitates actin disassembly from both barbed and pointed ends by abruptly removing filaments of a mean size of 260 subunits.⁷⁹⁾ The authors of this study hypothesized that instantaneous subunit loss proceeds through cooperative strand separation. Further biochemical analysis of this tri-component actin disassembling system should provide a clue to solve how fast actin disassembly might be achieved in the cell.

2) Frequent severing triggers fast capping protein dissociation in living cells. In parallel with the above biochemical analyses, single-molecule analysis by my group has revealed frequent F-actin severing/disruption in lamellipodia.^{4),5)} As mentioned, capping protein (CP) observed at the single-molecule level dissociates at 0.58 s^{-1} , approximately 20 times faster than actin disassembly. Notably, F-actin stabilizing treatments such as jasplakinolide,⁹⁸⁾ overexpression of hLIMK1 which inactivates cofilin,^{99),100)} and association with actin stress fibers all block the fast dissociation of CP. Jasplakinolide inhibits the interaction between cofilin and F-actin in live XTC cells.⁵⁾ These data strongly indicate that filament severing by cofilin triggers dissociation of CP from the actin network.

The dissociation of CP from the barbed has been reported to be induced by phosphatidylinositol-4,5-bisphosphate (PIP₂)⁶⁷⁾ or CARMIL.^{101),102)} However, recent studies employing direct microscopic observation of the uncapping reaction reported that PIP₂ induced-dissociation of CP is undetectable¹⁰³⁾ or occurs at a very slow rate ($\sim 10\%$ uncapped by 50 μM PIP₂ after 60 min).¹⁰⁴⁾ This markedly contrasts with the instantaneous uncapping by PIP₂ observed in the previous reports,⁶⁷⁾ one of which also described the fast uncapping effect of CARMIL.¹⁰¹⁾ The previous observations^{67),101)} might be caused by filament shearing upon mixing and simultaneous blockade of free CP by PIP₂. It would be necessary to reevaluate the effects of phospholipids and CARMIL using several other methods including direct microscopic observation.^{58),91),103)} The binding site for phospholipids¹⁰⁴⁾

overlaps with the actin binding surface¹⁰⁵⁾ on the 3D structure of CP.¹⁰⁶⁾

The finding of actin turnover-dependent fast CP dissociation led my research group to propose the frequent filament severing/end-to-end annealing hypothesis.⁴⁾ In this model, CP dissociates accompanied by a barbed end actin oligomer. The other part of the filament is rescued by similarly frequent end-to-end annealing, escaping rapid collapse. The size of actin oligomers capable of diffusing off the actin network is unknown. An EM study⁴⁷⁾ observed the filament intercross every 14–16 nm, ≈ 5 actin subunits. The detergent-extracted actin comet tail induced by *Listeria monocytogenes* in macrophages largely consists of short filaments, typically $\approx 0.1 \mu\text{m}$ (Fig. 8 in ref. 107). The size of diffusible actin oligomers could thus be within the range between 5 and 30 subunits. To achieve fragmentation of F-actin to these sizes, a severing rate of $0.1\text{--}0.02 \text{ s}^{-1}$ might be required. This problem was challenged in the following study.

3) AIP1-associated filament disruption in lamellipodia: 15 times more frequent than actin nucleation by Arp2/3 complex. A number of attempts have been made to quantitatively model actin-based cell edge protrusion. Regrettably, modeling studies^{108),109)} omit filament severing despite the accumulating evidence.^{4),110)} Such omission might have arisen from the lack of the knowledge regarding the *in vivo* frequency of filament severing. A recent study introducing an open source editable simulation database¹¹¹⁾ incorporated filament severing and annealing. This study built an extensive simulation with a large set of parameters. Yet the parameters used for several reactions including filament severing and uncapping are distinct from the data obtained in live cells (Fig. 4). The difficulty in predicting kinetic parameters only from *in vitro* reconstitution is evident.

To investigate filament severing *in vivo*, my research group turned to AIP1. AIP1 specifically recognizes the barbed end generated after cofilin-catalyzed filament severing/disruption. Several lines of evidence from single-molecule observation of AIP1 suggest the same property operating *in vivo*. First, LIMK1 overexpression decreased the F-actin association of AIP1. Second, the F-actin association of AIP1 is attenuated by jasplakinolide which blocks the cofilin-F-actin interaction simultaneously. Third, the dissociation of AIP1 slows upon jasplakinolide

treatment, which is analogous to the reaction of CP to this drug.⁴⁾ These notions support the use of AIP1 probes for monitoring cofilin/ADF-mediated filament severing/disruption.

Single-molecule observation of two AIP1 probes yielded $1.8 \mu\text{M s}^{-1}$ for the filament disruption rate in lamellipodia. There F-actin exists at $\approx 1000 \mu\text{M}$.⁵³⁾ The frequency of AIP1-associated filament severing/disruption corresponds to 1 s^{-1} per ≈ 560 subunits long filament on average. Surprisingly, this frequency is ≈ 15 times more than that of actin nucleation by Arp2/3 complex, $0.11 \mu\text{M s}^{-1}$.⁴⁾ Does AIP1 dissociate rapidly enough to allow barbed end growth?

In vitro, AIP1 dissociates from barbed end at $\tau < 10 \text{ s}$.⁷⁹⁾ AIP1 dissociates from barbed end at $0.1\text{--}0.2 \text{ s}^{-1}$ in jasplakinolide-treated cells where severing-based release of CP is severely attenuated.⁴⁾ If these observations reflect spontaneous dissociation of AIP1, $10\text{--}20\%$ of dissociation of AIP1 may lead to the formation of free barbed ends. This mode of barbed end generation may account for the ubiquitous actin polymerization^{3),6),26)} and the abundant free barbed ends^{4),15)} throughout lamellipodia. At the leading edge, Arp2/3 complex adds new filament nuclei receiving signals downstream of Rac-WAVE/Scar and Cdc42-WASP.¹¹²⁾ At the back of the leading edge, the AIP1-associated mechanism operates repeatedly along the retrograde flow. It is interesting to ask how the two opposite actions of AIP1, generation of the barbed end and breakdown of the polymer mass, might propagate in the actin array behind the leading edge.

4) Possible dynamic instability-like behavior of actin barbed end. To my regret, however, the above frequency of AIP1-associated filament disruption falls short of the rate predicted in the frequent filament severing-annealing theory by one order. Instead, this discrepancy opened up another possibility in the actin turnover regulation.⁵⁾ The filament disassembly mechanism involving AIP1 might preferentially attack the filament near the barbed end. *In vitro*, both barbed and pointed ends are disassembled by cofilin, AIP1 and coronin faster than the other portion of the filament. In lamellipodia, the pointed-end side disassembles much more slowly than the barbed end as the dissociation of Arp2/3 complex occurs at 0.048 s^{-1} (Fig. 4).⁴⁾ Thus the pointed end is more stabilized against the cellular actin disassembly activity than the barbed end. It is worthy mentioning that the dissociation rate of Arp2/3 complex

is almost equal to the decay curve of lifetime distribution of fast disassembling F-actin species. Hence the whole actin network other than the barbed end filament disassembles with similar ‘slow’ kinetics ($T_{1/2}$, ≈ 15 seconds) while the barbed end portion may undergo rapid filament disruption/severing cycles. This raises an intriguing possibility that the barbed end of cellular actin filaments may display growth and shrinkage behavior similar to dynamic instability (Fig. 5).

Dynamic instability is characteristic of microtubules dynamics.^{113),114)} At any point of time, a subset of microtubules are growing while others are rapidly shrinking. GTP-cap is thought to protect the plus end from disassembly. Once the plus end microtubules hydrolyze GTP, GDP-tubulin tends to be in a bent conformation, which triggers rapid disassembly from the plus end. While several molecules such as MCAK and XMAP215 enhance this growth and shrinkage dynamics,¹¹⁵⁾ dynamic instability is an intrinsic property of microtubules.

In contrast, the actin filament is believed to undergo treadmilling. The dissociation of actin is the fastest at the barbed end of ADP-F-actin (7.2 s^{-1} ; ref. 46). However, in the presence of ATP-G-actin, this fast barbed end dissociation may not occur. ATP hydrolysis is fast but Pi release is slow after ATP-G-actin assembles at the barbed end.^{46),92)} Pi abrogates the dissociation of ADP-F-actin at the barbed end.⁹¹⁾ ATP, which is abundant in the cytoplasm, also prevents the barbed end depolymerization of ADP-F-actin.¹¹⁶⁾ In the cell, profilin:actin complex exists at $22\text{--}40 \mu\text{M}$.^{9),117),118)} This complex can add to the barbed end as fast as free G-actin. Indeed, from the speed of processive movement of mDia1 FH2, the barbed end growth rate can be estimated to be 66 s^{-1} in lamellipodia.⁴⁾ All together, depolymerization from the barbed end seems unlikely to contribute to filament disassembly. However, the identification of cofilin/AIP1/coronin system and its ultra-fast filament disruption from both ends of F-actin^{78),79)} has now opened up the possibility that the barbed end may be the major site for filament disassembly *in vivo*. The fast barbed end and slow pointed end single-molecule kinetics of many filament end factors^{4),5)} suggest the dynamic instability-like behavior of the barbed end (Fig. 5). Further extensive studies are required to prove this intriguing possibility.

6. How does actin reassemble? : actin nucleating factors

To form filaments *de novo*, actin undergoes two steps, nucleation and elongation.⁴⁶⁾ Filament nucleation occurs spontaneously in the physiological salt condition, but this step requires long time in typical *in vitro* experiments. The formation of actin dimers and trimers occurs frequently in solution. However, dissociation of subunits is also fast and the nucleation step barely proceeds forward. Once filament nuclei consisting of 4 actin subunits are formed, they act as a filament and actin elongation proceeds steadily. The rate of actin nucleation depends on the cubic of the free G-actin concentration. In the cell, actin nucleating factors are required for efficient filament nucleation.

1) Arp2/3 complex. The major actin nucleator at the leading edge is Arp2/3 complex. Arp2/3 complex nucleates the filaments off the side of pre-existing filaments.^{56),58),59)} This branched actin network is abundantly found in lamellipodia.⁶¹⁾ As revealed by single-molecule imaging, its activation zone is highly restricted in the narrow ($< 0.65 \mu\text{m}$ wide) area proximal to the leading edge.⁴⁾ For activation, Arp2/3 complex requires three factors, the mother filament, actin monomer and activators such as WASP/WAVE family of proteins.¹¹²⁾ Especially the WAVE complex¹¹⁹⁾ is highly concentrated to the lamellipodium tip.¹²⁰⁾ Single-molecule observation revealed that distribution of the duration time of Arp2/3 association to the actin network fits with a single exponential curve. Thus a single rate-limiting step probably controls Arp2/3 complex dissociation. A cofilin/ADF protein actophorin¹²¹⁾ and coronin⁹⁶⁾ accelerate debranching whereas cortactin inhibits debranching.¹²²⁾ Le Clainche *et al.*¹²³⁾ showed that ATP hydrolysis by Arp2/3 complex coincides with debranching, whereas Dayel and Mullins¹²⁴⁾ argued that Arp2/3 complex hydrolyzes ATP immediately upon nucleation. It remains to be solved how the timing of the dissociation of Arp2/3 complex is determined in the cell.

2) Formin family. On the other hand, the Formin family (Formins) is responsible for the formation of a different set of actin filaments such as actin stress fibers and cytokinetic contractile rings.^{125),126)} Formins share conserved FH1 and FH2 domains in their C-terminal halves.¹²⁷⁾ With the FH2

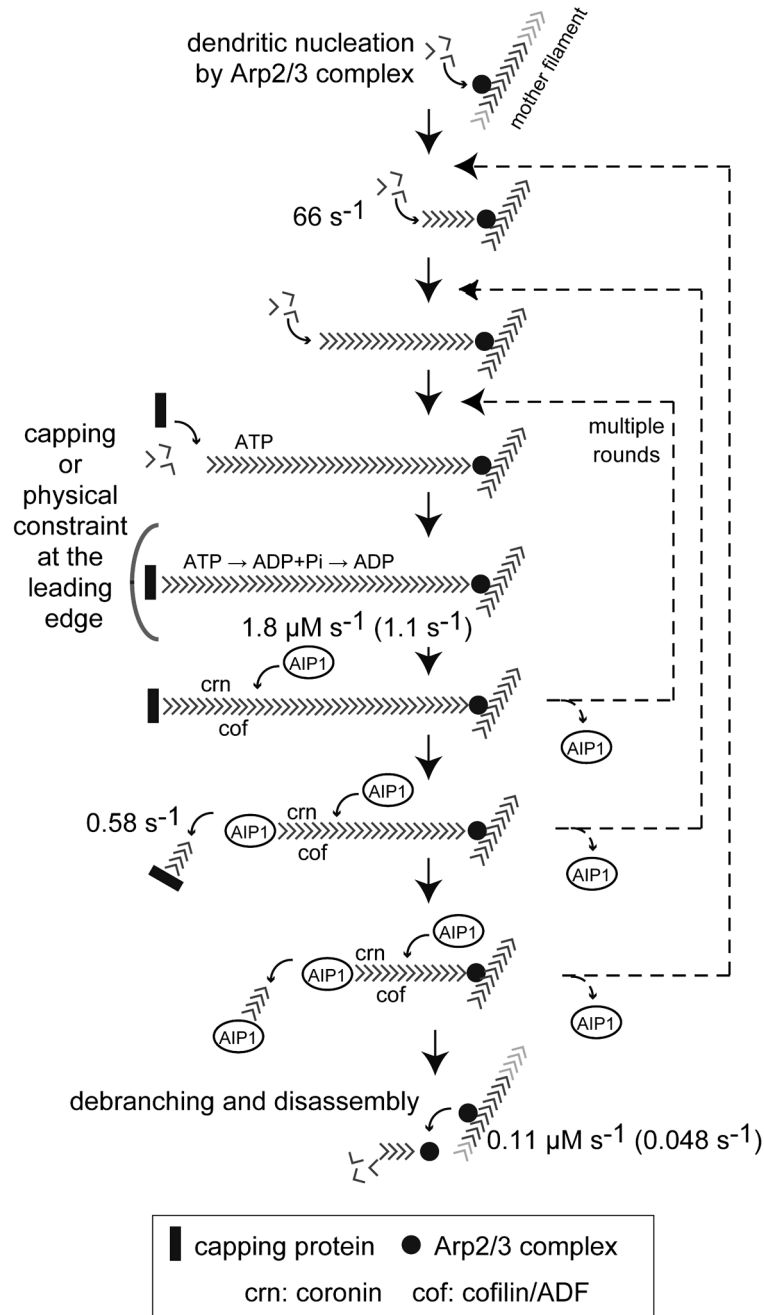


Fig. 5. Possible dynamic instability-like behavior of F-actin plus end. Combined with recent progress in biochemical analysis of the actin disassembling machinery consisting of cofilin, AIP1 and coronin,⁷⁹⁾ the fast single-molecule kinetics of capping protein⁴⁾ and AIP1⁵⁾ implies one-end growth and shrinkage behavior of F-actin, reminiscent of dynamic instability of the microtubules plus end.

domain, Formins nucleate actin filaments.^{128),129)} Remarkably FH2 domains stay processively associated with the growing barbed end while the filament elongates. This property was initially recognized as

'leaky cap'. At low concentrations, FH2 domains decrease the rate of actin subunit exchange at the barbed end to a certain extent while allowing subunit addition at higher concentrations.^{130),131)} This leaky

cap property led researchers to hypothesize the processive capping of the growing barbed end by Formins.¹³¹⁾

A proof of the processive actin capping by Formins was brought by the experiments demonstrating antagonism between Formins and CP.¹³²⁾ However, a possibility still remained for non-processive mechanisms such as a cluster of molecules associating with the side of the barbed end filament (See Fig. 1 in ref. 125). Recently, such a mechanism has been demonstrated for VASP.¹³³⁾ The VASP family¹³⁴⁾ has an actin nucleation activity which becomes apparent under non-physiological low salt conditions. VASP prevents capping protein from the barbed end. Certainly VASP and Formins share common properties. However, processive motion of VASP has not been observed (See for example supplementary video 6 in ref. 4; a cluster of VASP move forward in this movie, but this does not prove processive movement of individual VASP molecules).

A recent work provided compelling evidence for non-processive barbed end elongation by VASP.¹³³⁾ When VASP was clustered on the surface of beads, long range barbed end growth occurred in a manner insensitive to CP. Probably a cluster of VASP interacts one after the other with the barbed end and protects it from CP without processively moving along the growing barbed end. Formins can protect the barbed end at lower concentrations than VASP. This difference made it less likely that Formins use the same clustering mechanism as VASP. However in 2003, a direct proof was awaited for the processive mechanism of Formins.

My research brought a conclusion to this problem by direct visualization. Single-molecule observation of the FH1-FH2 domain of mDia1 directly demonstrated its directional movement in living cells.⁶⁸⁾ The FH1-FH2 unit structure moves over several tens of microns at 2 $\mu\text{m/s}$, which is comparable to the speed of kinesin and myosin in motility assays. The FH2 domain alone can move processively albeit at a slower speed of 0.13 $\mu\text{m/s}$. Moreover, processive actin assembly at the contact of recombinant mDia1 FH1-FH2 was reconstituted *in vitro*. Later, other groups employing similar *in vitro* single-filament elongation assays elucidated the acceleration mechanism of the barbed end growth by Formins. Profilin accelerates elongation of the FH1-FH2 bound barbed end remarkably fast, 5–15 times faster than the native filament^{69),135)–137)} as it was predicted from the

fast speed of mDia1 FH1-FH2.⁶⁸⁾ Formins can thus produce a long actin filament over the diameter of the cell in a short duration. Formins probably have evolved to enable rapid assembly of actin stress fibers and other structure-bridging F-actin.

Genetic studies have identified the roles of Formins *in vivo*. mDia1 is a mammalian homolog of *Drosophila* diaphanous, which is essential in cytokinesis. Among 3 mammalian isoforms, mDia2 is specifically essential in cytokinesis of certain cell types.¹³⁸⁾ mDia1^{-/-} mice are viable, but exhibit abnormal proliferation and migration of T lymphocytes.^{139),140)} mDia1^{-/-} mice develop age-dependent myeloproliferative disorder.¹⁴¹⁾ This phenotype is reminiscent of myelodysplastic syndrome (MDS) which is often associated with deletion in chromosome 5q. Human Dia1 gene is located in 5q31.3 and adds the list of candidates for a tumor suppressor such as *EGR1*, *RPS14* and α -catenin in 5q⁻MDS.¹⁴²⁾ Mouse Formin-2 is essential in positioning of the meiotic spindle,¹⁴³⁾ an actin-dependent process. Over wide eukaryotic species including yeast, *C. elegans* and *Drosophila*, Formins play important roles in the cell polarity formation in early development and cytokinesis. Please refer to other review articles to follow the progress of the entire Formin field.^{125)–127),144),145)}

3) Nucleators with tandem WH2 domains.

This recently emerged group includes Spire,¹⁴⁶⁾ Cobl,¹⁴⁷⁾ VopF,¹⁴⁸⁾ VopL¹⁴⁹⁾ and Lmod¹⁵⁰⁾ which contain three or four tandem WH2 domains. The WH2 domain^{151),152)} is also found in other types of actin regulators such as WASP, VASP, MIM,^{153)–155)} IRSp53,¹⁵⁶⁾ thymosin β 4 family and its related molecule ciboulot.^{157),158)} Through WH2 domains, these molecules interact with either G-actin or F-actin and exert various effects such as monomer sequestration, nucleation, barbed end elongation and scaffolding. The WH2 domain consists of a short amino acid stretch (17–43 aa) with a conserved N-terminal helix and a four amino acid motif corresponding to the LKKT in thymosin β 4.¹⁵²⁾ The N-terminal helix interacts with the cleft between actin subdomains 1 and 3 near the barbed end side of actin. The LKKT motif binds to a hydrophobic pocket on the actin surface.^{158)–160)} The two to three WH2 domains in tandem stabilize an actin dimer or trimer along the long pitch helix of F-actin,¹⁶¹⁾ illuminating the mode of filament nucleation. In N-WASP, the WH2 domain is followed by its closely related C motif. The

C motif binds Arp2 (actin related protein 2) and stabilizes the association of the WH2-bound actin subunit with Arp2. Dominguez and his colleagues proposed that the C motif is a variant of the WH2 domain which has specifically evolved for Arp2. They further proposed that WASP/WAVE is conceptually viewed as actin filament nucleators that specifically use the actin related protein (Arp2) instead of actin to build filament nuclei.¹⁶²⁾

On the other hand, the WCA region of WASP/WAVE dramatically increases the efficiency for Arp2/3 complex activation upon homo- or heterodimerization.¹⁶³⁾ The 2:1 complex formation between GST-WCA (VCA) and Arp2/3 complex implies an additional interface on the surface of Arp2/3 complex for the interaction with the second WCA region. Furthermore, the WH2 motif in N-WASP interacts with the barbed end filament, supporting attachment of F-actin to moving membranes.¹⁶⁴⁾ Thus diverse functions are brought by distinct combinations of WH2 domains. Together with the non-processive actin elongation by a cluster of VASP,¹³³⁾ a variety of activities by the cooperative actions of multiple WH2 domains within a single or combinations of WH2 proteins operate in the actin system, which are now being revealed.

7. A clue for activation mechanism of mDia1 and Formins: G-actin inhibitors

I describe a novel role of homeostasis between G- and F-actin which bridges the gap between actin disassembly and polymer reassembly.

The GTP-bound form of Rho binds N-terminal region of mDia1. mDia1 is an autoinhibited molecule, and the binding of Rho opens mDia1 to release the activity of the FH1-FH2 unit.^{32),165)} On a mission to seek small molecule compounds that evoke processive actin assembly by mDia1, latrunculin B (LatB) was unexpectedly identified.⁹⁾ LatB is a G-actin inhibitor and is widely used to probe actin-dependent cellular processes. LatB binds G-actin near its ATP binding site and inhibits nucleotide exchange and filament assembly.¹⁶⁶⁾ When administrated at a high concentration ($>1 \mu\text{M}$), LatB stops the movement of mDia1 because of a loss in the polymerization-competent G-actin.⁶⁸⁾ But at a lower dose, LatB dramatically increases the frequency of processively-moving mDia1. This induction is observed as early as 10 s after the perfusion of 100 nM LatB, accompanied by a reduction in the speed of mDia1.⁹⁾

The induction of processive mDia1 speckles is reproduced by analogous treatments such as swinholide A and unpolymerizable G13R and R62D actins. Swinholide A-bound G-actin forms an anti-parallel dimer.¹⁶⁷⁾ This dimer species is distinct from a readily polymerizing dimer obtained by cross-linking adjacent subunits in the F-actin helix.¹⁶⁸⁾ The anti-parallel dimer neither fits with the structure of actin in complex with the FH2 domain of Bni1p¹⁶⁹⁾ nor promotes actin nucleation.¹⁶⁸⁾ I therefore predict that swinholide A promotes actin nucleation by mDia1 not through dimer stabilization.

The FH1-FH2 or FH2 region of mDia1 alone responds to low-dose LatB and display increased processive movement. Rho is required for full-length mDia1 being in an open conformation, but is not a direct mediator of LatB-increased actin nucleation by mDia1. LatB thus increases the rate of actin nucleation by acting on the FH2 catalytic core domain of mDia1 in cells. In marked contrast, LatB has only negative effects on mDia1 FH2-catalyzed actin filament assembly *in vitro*.⁹⁾

8. Latrunculin paradox: inhibitor-induced increase in drug-free targets

The opposite effects of LatB on the mDia1-catalyzed actin nucleation between in cells and *in vitro* puzzled me. While profilin binds G-actin noncompetitively with latrunculin, the binding of latrunculin reduces the affinity between T β 4 and G-actin.¹⁶⁶⁾ In one study, latrunculin-induced concentration changes of G-actin either free or bound to partners were extensively analyzed.¹⁷⁰⁾ The authors of this study concluded that latrunculin has little effect on the concentration of free G-actin. Even so, I reevaluated this issue by kinetic modeling. My research group came to notice the paradoxical increase in the free G-actin concentration induced by LatB.⁹⁾ The simulation analysis predicted that LatB induces a several-fold increase in the concentration of free G-actin (Fig. 6). This increase in free G-actin is accompanied by the accumulation of several micromolar LatB in the cell compartment. The underlying key mechanism for this paradoxical drug effect is the accumulation of total G-actin. In addition to the primary effect of LatB to inactivate G-actin, LatB causes a 1.2–1.3 fold increase in the total G-actin concentration as a result of decreased actin assembly, which in turn raises the free G-actin concentration several fold. This latter effect is due to the saturation

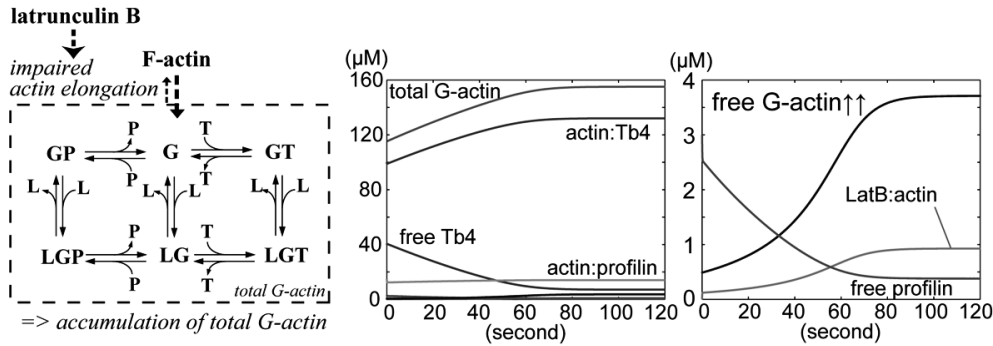


Fig. 6. Latrunculin paradox.⁹⁾ The left panel depicts the scheme of kinetic modeling used for estimating time-dependent concentration changes of each G-actin species either free or bound to LatB, Tβ4 and/or profilin. G, L, T, and P represent G-actin, LatB, thymosin-β4 (Tβ4) and profilin, respectively. The simulation results (middle and right graphs) revealed the paradoxical effect of latrunculin B. The primary action of latrunculin B is to bind G-actin and inhibit actin polymerization. In experiments, slowdown of actin elongation by mDia1 and a ~30% F-actin decrease were observed in cells treated with low-dose LatB. The change in the balance between F- and G-actin leads to the paradoxical, several fold increase in free G-actin accompanied by saturation of the G-actin sequestering activities in the simulation. Adopted with permission from J. Cell Sci. **121**, 3403-3412, Fig. 4B (2008).

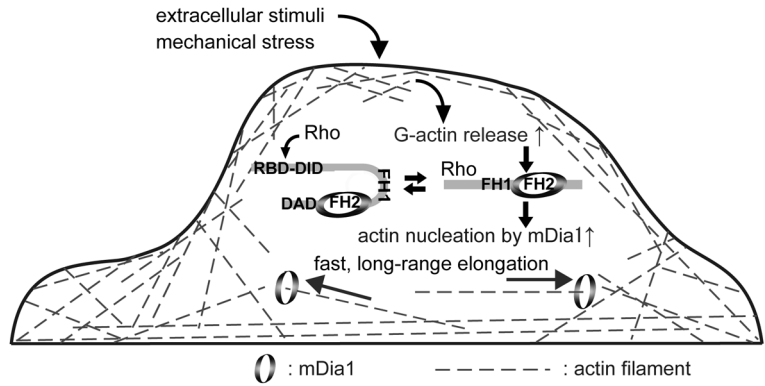


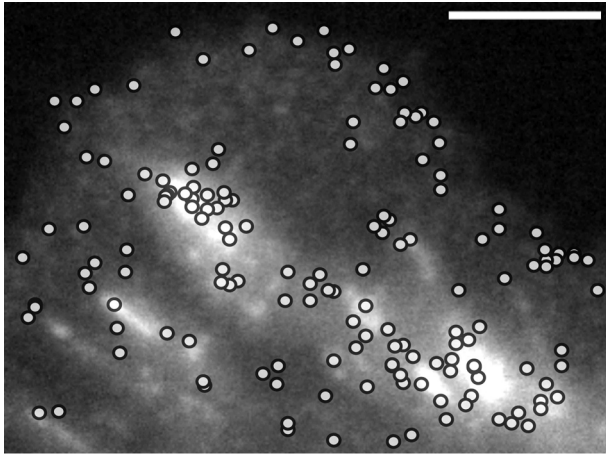
Fig. 7. Feedback actin polymer restoration mechanism involving mDia1. Rho signaling determines the ratio between the closed (left) and the opened (right) form of mDia1. An increase in free G-actin efficiently converts “opened” mDia1 to the fully-active actin nucleating state, leading to fast long-range actin filament assembly in cells. With this property, mDia1 responds to acute disruption of F-actin structures evoked by physical stress or by extracellular stimuli-induced active actin remodeling, and rapidly restores cellular actin polymers by their fast actin elongation ability.

of the buffering capacity of G-actin sequestering proteins. This study highlight an unprecedented example of pharmacological effects in which an antagonist increases the concentration of drug-free targets in a complex biological system.⁹⁾

9. G-actin regulation of F-actin nucleation and reassembly

Figure 7 summarizes the model for dual regulation of mDia1-catalyzed actin assembly. Rho signaling determines the ratio of the opened and the closed

forms of mDia1, whereas an increase in free G-actin effectively converts the opened mDia1 to the actin nucleating state, leading to fast long-range actin filament assembly. Since the efficiency of actin nucleation by Formins depends on the cubic of the free G-actin concentration *in vitro*, FH2 domains may serve as sensitive sensors for the concentration fluctuation of free G-actin in the cell. In accordance with this idea, a distantly-related member of Formins, FRL1, also exhibits frequent processive movement in response to low-dose LatB. In contrast, Arp2/3 com-



fluorescence image: mPlum-AIP1

○ : site of emergence of single-molecule mDia1 FH2

Fig. 8. Enhanced F-actin nucleation by mDia1 around the sites of vigorous actin disassembly in live XTC cells (adopted from ref. 9). Colocalization between the emergence site of the single-molecule mDia1 FH2 probe (circles), the catalytic core for its actin nucleation activity, and the distribution of AIP1 (mPlum-tagged; fluorescence image) is shown. AIP1 is a cofactor of cofilin and was used to monitor the distribution of cofilin-catalyzed actin disassembly. Scale bar, 5 μm . Adopted with permission from *J. Cell Sci.* **121**, 3403-3412, Fig. 6 (2008).

plex does not respond to LatB.⁹⁾ Activities of actin nucleators may thus be regulated differentially by the G-actin concentration in the cell.

Another important notion is that the emergence frequency of processive mDia1 correlates with the local concentration of AIP1. During the course of the initial work that proved the processive actin elongation by mDia1,⁶⁸⁾ it was noticed that the FH2 domain construct of mDia1 frequently appears in discrete foci found at the base of lamellipodia. From these spots, mDia1 FH2 mutants appear and move away as if such foci were a fountain of processive mDia1. This finding had not been explored for a while. Later, the similarity in the spatiotemporal dynamics between mDia1 FH2 and AIP1 was noticed. It was then confirmed that the local concentration of AIP1 correlates with the emergence frequency of both full-length and the FH2 mutant of mDia1 (Fig. 8).⁹⁾ This coincidence is also observed between AIP1 and other processive Formins including FRL1 (unpublished results). The association of AIP1 with F-actin depends on the activity of cofilin. *In vitro*, the association of Formins to free barbed ends was described by direct microscopic observation, but this

event appears to be much less frequent than filament nucleation on the image.⁶⁹⁾ Therefore appearance of processive mDia1 in the cell may mostly represent *de novo* filament nucleation. The above findings are best interpreted as showing that an increase in the local G-actin release upregulates the efficiency of filament nucleation by mDia1.

mDia1 demonstrates the greatest rates of accelerated actin elongation among Formins.^{69),135)} mDia1 ΔN3 elongates actin at the rate of 2 $\mu\text{m}/\text{s}$ (≈ 720 subunits/s) in cells.⁶⁸⁾ Eukaryotic cells, whose diameter reaches several tens of microns, have probably evolved to obtain these fast actin elongating systems. For example, leukocytes in the blood flow need to adhere the vessel wall within a few seconds after stimulation by leukotrienes. Even with mDia1, it takes five seconds to form *de novo* actin filaments which bridge over the diameter of leukocytes. The property of mDia1 may also provide an efficient way to rapidly reassemble actin polymers when cells are subjected to physical stress. The alignment of actin stress fibers remarkably alters when cells are subjected to cyclic stretch.^{171),172)} Rho is activated upon stretch of smooth muscle cells¹⁷³⁾ and plays a role in stretch-induced reorientation of stress fibers in endothelial cells.¹⁷⁴⁾ Mechanical stress-induced disassembly of actin stress fibers may possibly upregulate actin nucleation by mDia1 through an increase in G-actin. The role of the G-actin regulated actin polymer restoration mechanism is currently under investigation by the use of fluorescence single-molecule observation.

10. Perspective and future directions

In this article, I have focused on the accomplishment of live-cell fluorescence single-molecule imaging developed in my research, and described its implications in relation to findings from other studies. As the readers may notice, most of comparison is made against *in vitro* actin biochemistry rather than *in vivo* phenotype analysis. Direct comparison between the single-molecule data and the biochemical knowledge has led us to find the important links such as filament severing-mediated barbed end disruption (Fig. 5) and G-actin regulation of actin nucleation by Formins (Figs. 7 and 8). Single-molecule imaging has an enormous advantage in elucidating the coupling between the probe and its surrounding molecules under the complex intracellular environment. With its speeds and quantitative characteristics,

single-molecule approach can be a powerful tool to elucidate elementary process in many dynamic biological systems.

On the other hand, there still remains a gap between molecular kinetics and determination of the cell behavior. To bridge over the gap, modeling and computational analysis are becoming more and more important. The example of the paradoxical latrunculin effect (chapter 7–9) highlights the importance of quantitative modeling. At the same time, this example warns us to be cautious before moving on to modeling of the body processes on the computer chip. We still do not fully understand biological systems as simple as the ‘latrunculin paradox’ case (Figs. 6 and 7).

Finally, I introduce another use of single-molecule imaging for bridging the gap over elementary and whole body processes. Pharmacological perturbation and real time *in vivo* molecular behavior analysis often synergize to reveal hidden molecular linkages. Recently, my research group has found the first example of inhibitor-induced conformational regulation of kinases.¹⁷⁵⁾ Imatinib, a very successful kinase inhibitor in the treatment of chronic myelogenous leukemia, induces rapid translocation of its target, Abelson kinase (*c-abl*) to the cell edge. This translocation is triggered by the drug-induced complex formation with unknown cofactor(s). Despite extensive studies employing biochemistry, genetics, cell-based assays and structural analysis, our findings had not been recognized for many years. Two characteristics of live-cell single-molecule imaging, (i) a whole set of molecules reconstituted in the assay and (ii) its capability of observing a specific molecular event, enabled us to discover this unanticipated effect of kinase inhibitors. Real time observation of the effect of small compounds on the target behavior may thus help elucidate the mechanism of action of the drugs. Conversely, small compounds provide tools to perturb molecular functions rapidly. This combination should deepen our understanding of molecular functions *in vivo*. Small compounds may be the key to bridge the gap of our knowledge between elementary and whole body processes in the post genomic era.

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Profile

Naoki Watanabe was born in 1964. After graduating from Kyoto University Faculty of Medicine, he worked as a clinical doctor for three years. He then started his research career in 1993 with studies on cell signaling under the supervision of Prof. Shuh Narumiya. He received Ph.D. from Kyoto University in 1998 for the studies on the cell signaling cascades downstream of the small GTPase Rho. He became a JSPS research fellow, and then he was promoted to assistant professor in 1998. From 1999 to 2002, he joined Prof. Timothy J. Mitchison's laboratory at Harvard Medical School as a postdoctoral research fellow and developed fluorescence single-molecule speckle microscopy. He moved back to Kyoto University and was promoted to associate professor in 2002. He also worked as a researcher at Precursory Research for Embryonic Science and Technology (PRESTO) for three years from 2002. His research focuses on the complexity and dynamicity of actin cytoskeleton remodeling in living cells. The invention of live-cell fluorescence single-molecule imaging in 2002 and the discovery of the processive actin elongation mechanism by formin homology proteins in 2004 are the highlights of his study. He was awarded the first JSPS prize (FY2004).

