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Myo2p is the major motor involved in actomyosin ring contraction in fission yeast

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Cytokinesis in many eukaryotes requires an actomyosin-based contractile ring [1]. In fission yeast, cytokinesis involves the type II myosins Myo2p and Myp2p and the type V myosin Myo51p [2]. A recent study by Laplante et al. [3], using deletion mutants of myp2 and myo51 and the mis-sense mutant myo2-E1 [4], concluded that each myosin has distinct functions and proposed that Myp2p plays the dominant role in actomyosin ring contraction. Here we present evidence that Myo2p, not Myp2p, is likely to be the major motor driving actomyosin ring contractility. Since the previous work [3] was performed at 25°C, the permissive temperature for myo2-E1, we compared cytokinesis timings in myo2-E1 and myo2Δ at 25°C and found that myo2-E1 is only partially compromised at 25°C. Furthermore, we find that myp2Δ and myp2Δ myo51Δ double mutants contract actomyosin rings at ~90% of the rate of wild-type cells at 30°C and 36°C, suggesting that Myp2p plays a minimal role in ring contraction at these temperatures. Finally, ring contraction in our myo2-E1 strain took longer at 25°C than previously reported [3]. Although faster-acting alleles of myo2 will be required to evaluate its contribution at 25°C, our work establishes that Myo2p is the major motor involved in ring contraction, under most, if not all, conditions.

From several laboratories has shown that myo2-E1 forms healthy colonies at 25°C but fails to do so at increased temperatures [4,5]. Since previous work has shown that Myo2-E1p (the product of myo2-E1) neither binds actin filaments nor has ATPase activity in vitro at 25°C [6], Laplante et al. [3] inferred that myo2-E1 should also be severely compromised in vivo at 25°C. It is known that in vitro activity and in vivo function of products of mutant alleles are not necessarily correlated [7]. Furthermore, Laplante et al. [3] extrapolate that, since myo2-E1 is a severely compromised allele of myo2, the mGFP-tagged version (mGFP-myo2-E1) used in their work would also be severely compromised. The data in Figure 1D of the accompanying response of Laplante and Pollard and in Figure S1A in Laplante et al. [3] suggest, however, that the mGFP tag partially rescues myo2-E1.

We first investigated whether myo2-E1 is an appropriate allele to understand Myo2p function at 25°C. We imaged the dynamics of the actomyosin ring component Bt1p, which localizes to precursor nodes and actomyosin rings [8], in germinating myo2Δ spores, with the rationale that for its use in comparative studies it should have a phenotype comparable to myo2Δ. Actomyosin rings never assembled in myo2Δ cells, even after 4 hours of imaging, whereas germinated wild-type spores assembled normal actomyosin rings (Figure S1A and Movie S1 in Supplemental Information). However, contractile actomyosin rings assembled in myo2-E1 cells and importantly even in myo2-E1 myo51Δ cells [3] (Figure S1B, see 36˚–54˚ time points, and Movie S2, see 35˚–55˚ time points), suggesting that Myo2-E1p should retain some activity for ring assembly through the search, capture, pull and release (SCPR) mechanism [9]. Thus, myo2-E1 is not a severely compromised allele at 25°C and is inappropriate for investigating the contribution of Myo2p at 25°C.

We then compared the time taken for various aspects of cytokinesis in wild-type cells, the partially active myo2-E1 mutant, the myp2Δ myo51Δ double mutant, and double mutants combining the partially compromised myo2-E1 and myp2Δ or myo51Δ at 25°C (as a control), and at 30°C and 36°C. Through DNA sequencing and

**Figure 1. Actomyosin ring kinetics in S. pombe myosin mutants.**

(A,C,E) Time taken for various steps in cytokinesis (coalescence of nodes into a ring, dwell time before contraction, and contraction). (B,D,F) Ring contraction rates. In all cases, Rlc1p-3GFP was used as a marker of cytokinetic nodes and the actomyosin ring. Note that improper rings underwent aberrant contraction were detected in myo2-E1 mutants (at 30°C and 36°C), myo2-E1 myp2Δ mutants (at 25°C, 30°C and 36°C) and myo2-E1 myo51Δ mutants (at 25°C, 30°C and 36°C).

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At 25°C, we made similar observations to Laplante et al. [3] when ring assembly, maturation, and contraction times were scored in wild-type cells, and in myo2Δ, and myo51Δ mutants, but not in myo2-E1 and myo2-E1 myo51Δ mutants (Figures 1A and S1B, and Movie S2). Ring assembly time doubled in myo51Δ compared with wild-type cells, consistent with previous work reporting an ancillary role for Myo51p in ring assembly [10]. Ring contraction was slower in myo2Δ mutants (53.9 ± 7.5 minutes). In our analysis, ring contraction was not as severely affected in myo2-E1 myo51Δ as reported in Laplante et al. [3]. Importantly, even the partially compromised myo2-E1 (containing Myp2p and Myo51p) had a stronger defect than myo2Δ in ring contraction and took ~86 mins for ring contraction, suggesting that Myo2p rather than Myp2p played a major role in ring contraction even at 25°C, with Myp2p playing an ancillary role in ring contraction at 25°C (Figures 1A,B and S1B and Movie S2). The strong additive effect in myo2-E1 myo51Δ and myo2-E1 myo2Δ mutants made it difficult to demarcate different steps in cytokinesis, but the entire process of improper cytokinesis took a comparable amount of time in myo2-E1, myo2-E1 myo2Δ, and myo2-E1 myo51Δ mutants (125–150 minutes).

Ring assembly was defective in myo2-E1, myo2-E1 myo51Δ, and myo2-E1 myo2Δ mutants grown at 30°C or 36°C, confirming the essential role of Myo2p in actomyosin ring assembly (Figures 1C,E and S1C,D and Movie S2). Ring contraction times in myo2Δ, myo51Δ, and myo2Δ myo51Δ mutants grown at 30°C or 36°C were comparable to those in wild-type cells, with ring contraction rates at least 90% of that seen in wild-type cells (Figures 1D,F and S1C,D and Movie S2). These data further reinforced the view that Myp2p is not the major motor involved in ring contraction at 30°C and 36°C.

Laplante and Pollard (accompanying response) have shown that Rlc1p-3GFP shows adverse interactions specifically with myo2-E1 and suggest this may have increased the ring contraction time in myo2-E1 rlc1-3GFP at 25°C in our work. However, the fact that rlc1-3GFP exacerbates the cytokinesis defect of the myo2-E1 mutant (but not wild-type cells) is itself further evidence that myo2-E1 is a weak allele that is further weakened by its interaction with Rlc1p-3GFP in vivo. This further weakened Myo2p-E1p-Rlc1p-3GFP complex causes a significant reduction in ring contraction rate, strengthening our conclusion that Myo2p is the main motor involved in ring contraction.

What then are the specialized functions of Myo2p, Myp2p, and Myo51p? We and others have established that Myo2p is central to ring assembly at all temperatures (current work and [2]), with Myo51p playing an ancillary role [10]. Our analysis has established that ring contraction is independent of Myo2p at 30°C or 36°C. Myp2p may perform an ancillary role in ring contraction at these temperatures. As reported [3], ring contraction is slower in cells lacking Myp2p at 25°C, consistent with previous work that Myp2p is specialized for cytokinesis at lower temperatures [2]. In our work, actomyosin ring contraction in the myo2-E1 mutant took longer than in the myo2Δ mutant at 25°C, suggesting that Myo2p is likely the major motor involved in ring contraction even at 25°C. However, analysis of myo2-E1 mutants to inform the role of Myo2p in ring contraction should be considered to underestimate Myp2p’s function, since myo2Δ cells and myo2-E1 rlc1-3GFP cells have a stronger phenotype than myo2-E1 at 25°C. Fast-acting stronger conditional mutant alleles of Myo2p will be essential to investigate the precise role of Myo2p in ring contraction at 25°C.

SUPPLEMENTAL INFORMATION

Supplemental Information includes experimental procedures, one figure and two movies and can be found with this article online at http://dx.doi.org/10.1016/j.cub.2016.12.024.

AUTHOR CONTRIBUTIONS

P.Z. and S.P., conception and design, acquisition of data, analysis and interpretation of data, drafting or revising the article; A.K., analysis tool development and interpretation of data; M.K.B., conceived the project, conception and design, analysis and interpretation of data; S.P. and M.K.B., wrote the manuscript; all authors reviewed the manuscript.

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