Compositional and expression analyses of the glideosome during the *Plasmodium* life cycle reveal an additional myosin light chain required for maximum motility

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Running title: The *Plasmodium* glideosome needs two light chains for maximum motility

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ABSTRACT
Myosin A (MyoA) is a Class XIV myosin implicated in gliding motility and host cell and tissue invasion by malaria parasites. MyoA is part of a membrane-associated protein complex called the glideosome, which is essential for parasite motility and includes the MyoA light chain MTIP, and several glideosome-associated proteins (GAPs). However, most studies of MyoA have focused on single stages of the parasite life cycle. We examined MyoA expression throughout the Plasmodium berghei life cycle in both mammalian and insect hosts. In extracellular ookinetes, sporozoites and merozoites, MyoA was located at the parasite periphery. In the sexual stages, zygote formation and initial ookinete differentiation precede MyoA synthesis and deposition, which occurred only in the developing protuberance. In developing intracellular asexual blood stages, MyoA was synthesized in mature schizonts and was located at the periphery of segmenting merozoites, where it remained throughout maturation, merozoite egress and host cell invasion. Besides the known GAPs in the malaria parasite, the complex included GAP40, an additional myosin light chain designated essential light chain (ELC) and several other candidate components. This ELC bound the MyoA neck region adjacent to the MTIP binding site, and both myosin light chains co-located to the glideosome. Co-expression of MyoA with its two light chains revealed that the presence of both light chains enhances MyoA-dependent actin motility. In conclusion, we have established a system to study the interplay and function of the three glideosome components, enabling the assessment of inhibitors that target this motor complex to block host cell invasion.

The abbreviations used are:
ABC, ammonium bicarbonate; ACN, acetonitrile; CTD, C-terminal domain; ELC, essential light chain; GAP, glideosome associated protein; IFA, indirect immunofluorescence assay; IMC, inner membrane complex; IPTG, Isopropyl β-D-1-thiogalactopyranoside; MLC-B, myosin B light chain; Myo, myosin; MTIP, myosin tail domain interacting protein; PV, parasitophorous vacuole; PVM, parasitophorous vacuole membrane; TCEP, tris(2-carboxyethyl)phosphine;

Three stages of the malaria parasite, sporozoites, ookinetes and merozoites, are invasive to host cells or tissue and two stages, sporozoites and ookinetes, are motile. Motility and invasion are active processes and in Plasmodium as in other apicomplexan parasites such as Toxoplasma gondii (Tg), an actomyosin motor is central to these processes (reviewed in (1, 2)). Myosin proteins are typically composed of a conserved globular head domain, a neck domain comprising variable numbers of myosin light chain binding sites, and a diverse tail domain that is often responsible for dimerization and/or cargo binding. The myosin head binds to actin and also contains the motor activity, where ATP hydrolysis is translated into molecular movement (3-6). Myosin A (MyoA), an Apicomplexa-specific class XIV myosin comprised of a head and neck region, but lacking a tail domain, is the motor component of the glideosome protein complex is attached to the inner membrane complex (IMC), a network of large, flattened vesicles located just below the surface plasma membrane of the parasite (7-9).

Some components of the glideosome have been defined in T. gondii and also in Plasmodium species. In addition to MyoA, there is a myosin light chain (MLC) that binds to the last few amino acid residues of MyoA, corresponding to part of the myosin neck region and called T. gondii MLC1 (10), or myosin tail domain interacting protein (MTIP) in Plasmodium spp. (11). This MLC is involved in anchoring MyoA to the IMC via a palmitoyl modification (12) and thus appears to take on a role usually fulfilled by a myosin tail, that is anchoring myosin at a specific subcellular location. Recently, two more T. gondii MyoA MLCs have been described and called essential light chains (ELC) 1 and 2. The terminology follows that used for mammalian muscle myosins with two light chains in which the most proximal is an ELC and the distal is a regulatory light chain (RLC). ELC1 and 2 have been shown to bind to the MyoA neck adjacent to MLC1 (which binds at the RLC site) in a mutually exclusive manner and are important for motor activity (13, 14). Bookwalter and colleagues have demonstrated that a heterologously expressed
MyoA and its associated light chains, a number of glideosome-associated proteins (GAPs) have been described, including GAP45 (16-18) (and its Coccidia-specific homolog GAP70 (19)), GAP50 and a family of three proteins with multiple membrane spans known as GAPMs (20). Another glideosome component, GAP40, has been described in T. gondii (19) and has a homologue in Plasmodium based on genomic (21) and proteomic studies (22-24), but it has not been shown experimentally to be part of the glideosome in this genus. In Plasmodium, most studies on MyoA have been limited to the asexual blood stage parasites in the mammalian host, although its essential role in gliding motility in P. berghei ookinetes has been demonstrated in promoter swap experiments (25).

In this study, we have examined the expression and location of MyoA throughout the malaria parasite life cycle, including all invasive stages, and its temporal profile during merozoite and ookinete development. We have examined the composition of the glideosome in asexual blood stages by immunoprecipitation and mass spectrometry of proteins associated with GFP-tagged MyoA, confirming the presence of GAP40 and identifying a new myosin light chain. This essential-type light chain binds to the MyoA neck adjacent to, and in addition to, MTIP. Recombinant PfMyoA binds actin and with both light chains bound demonstrates increased sliding velocity of actin compared to MyoA with just MTIP bound. This is a first important step to reconstituting the Plasmodium glideosome in vitro, which will be an invaluable system with which to dissect the function of, and interplay between, different components of the complex.

RESULTS
Myosin A is a predominantly membrane-associated protein at the periphery of all invasive stages of the P. berghei life cycle
Expression of the myoA gene was detected by qRT-PCR throughout the parasite life cycle, particularly in schizonts and sporozoites, but also in non-activated gametocytes (Fig. 1A). To examine the synthesis and location of the protein throughout the parasite life cycle we produced a C-terminal green fluorescent protein (GFP)-tagged MyoA expressed from the endogenous genomic locus using a strategy we described recently for P. falciparum MyoA and P. berghei MyoB (26) (Fig. 1B). Integration of sequence coding for GFP into the endogenous MyoA locus and expression of the tagged protein was confirmed by PCR amplification from genomic DNA using primers that only produce a product following specific integration (Fig. 1C), by western blotting (Fig. 1D), and by fluorescence microscopy (Fig. 1F-H). Tagging MyoA with GFP resulted in no detectable phenotype: there was no effect on the rate of asexual parasite growth, on male gametogenesis (the number of exflagellation centres), the number of oocysts per mosquito gut or the efficiency of infection via sporozoite inoculation (as judged by the number of days following infected mosquito bite that parasites were observed by microscopy in blood smears). In a subcellular fractionation of schizonts, the protein was largely associated with the peripheral membrane (carbonate soluble) and membrane (carbonate insoluble) fractions rather than in the soluble cell lysate (Fig. 1E), consistent with its association with the glideosome and the IMC membrane compartments. Using live parasite microscopy, the protein was detected uniformly at the periphery of segmenting schizonts and extracellular merozoites in asexual blood stages and at a similar location in extracellular ookinetes and salivary gland sporozoites (Fig. 1F), consistent with location at the IMC. Interestingly, in sporozoites there was also a clear perinuclear localization. In early cytomere stages of liver-stage schizonts the signal appeared to be cytosolic, rather than membrane associated. In late cytomere stages, 55 h after invasion by sporozoites, a clear peripheral location associated with the hepatic merozoites was observed (Fig. 1G). The presence of myoA mRNA in ookinete stages, as well as in non-activated gametocytes as a translationally repressed transcript, prompted us to examine the temporal profile of protein expression during the 24 hours of ookinete development divided into six stages (27) (Fig. 1H). MyoA-GFP was not visible in either activated or non-activated gametocytes and barely detectable in the zygote and stage I and II ookinetes. Following the establishment of morphological polarity at about 10 hours after fertilization (Stage III) the protein was detectable in the growing protuberance rather
than the spherical body in the intermediate retort forms, so-called because of their shape (stages III–V). This location was retained in later stages, with the MyoA-GFP associated with the parasite periphery in fully formed motile ookinetes. There was a concentration of fluorescence detected at the apical tip of the mature oocinete. Immunofluorescence using an antibody against the surface marker P28 was used to visualize the oocinete at all stages of development.

Expression of MyoA during P. falciparum asexual blood stage development and formation of the glideosome complex

The use of P. falciparum allows a more detailed analysis of asexual blood stages than can be achieved with P. berghei. MyoA-GFP was largely undetectable by live fluorescence microscopy until approximately 38 hours after merozoite invasion of an erythrocyte, but was present in later stages at the periphery of developing segmented schizonts and free merozoites (but not associated with the food vacuole/residual body), consistent with its proposed location at the IMC (Fig. 2A). No GFP signal was detected in ring or trophozoite stages indicating that the protein is not synthesised at this time and any MyoA carried through with the invading merozoite is degraded rapidly following erythrocyte invasion. Interestingly, the protein was also not detected in early schizonts when the first formation of the IMC can already be detected with, for example GFP-tagged GAP50 (28) or GAP45 (29) and seen as small ring-shaped structures at the tips of developing merozoites. Western blot analysis of parasite extracts at the corresponding time points (Fig. 2B) confirmed that GAP50 is present in early schizonts (detectable from 24 h and strongly expressed 30 h post-invasion). Whilst a small amount of MyoA-GFP can be detected 27–36 h after merozoite invasion, the strongest expression is detected from 38 h post-invasion. A western blot analysis of the MyoA-GFP immunoprecipitate obtained from these extracts (Fig. 2Bi) indicates that known components of the glideosome complex, MTIP, GAP45 and GAP50 are associated with MyoA from 38 h after erythrocyte invasion by a merozoite. Tagging MyoA with GFP had no effect on the rate of asexual blood stage parasite growth.

Location of MyoA in P. falciparum during erythrocyte invasion

The peripheral location of MyoA-GFP in mature schizonts and free merozoites was confirmed by indirect immunofluorescence assay (Fig. 3A). Antibodies against GFP co-localised with antibodies against MTIP, GAP45 and GAP50 consistent with the presence of these proteins in a complex. During erythrocyte invasion, the peripheral location of MyoA did not change during the transition from initial attachment of the merozoite to the intracellular ring stage at the completion of invasion (Fig. 3B). However, thereafter the signal disappeared until resynthesis during schizogony.

Identification of GAP40 and discovery of a second MyoA light chain in the P. falciparum glideosome complex

A number of proteins associated with the glideosome have been identified in both Plasmodium and Toxoplasma species and the presence of GAP45, GAP50, and MTIP, some of the known components in P. falciparum asexual blood stages, has been confirmed above. To examine what other proteins might be associated we examined further the protein complex immunoprecipitated with GFP-specific antibodies. To do this we first used parasites in which GAP45 was tagged internally with GFP (29) and then those in which MyoA was tagged at its C-terminus with GFP. In the first instance we were interested to see whether or not GAP40, which is a highly phosphorylated protein (23, 30-32) present in the T. gondii glideosome (19), is present in the Plasmodium complex. In order to visualise phosphorylated proteins associated with GFP-GAP45 we metabolically labelled P. falciparum schizont phosphoproteins using $^{32}$P-phosphate (Fig. 4). As a control, 3D7 schizonts in which GAP45 was unmodified were used (Fig. 4, lane 1 in panels A and B). The glideosome was precipitated from schizont lysates using antibodies to GFP, and $^{32}$P-labelled phosphoproteins were visualised by autoradiography (Fig. 4A). In addition to GFP-GAP45 itself, there were five labelled proteins that were absent from the control samples. The identities of some were confirmed by immunoprecipitation from schizont lysates with GFP antibodies followed by western blotting with antibodies against known glideosome components MyoA,
GAP45, GAP50, and MTIP (Fig. 4A). The most prominent of the phosphoproteins, with an apparent molecular weight of just under 50 kDa, was distinct from these four proteins and so its identity was examined further by mass spectroscopy following fractionation of the protein precipitate by SDS-PAGE and tryptic digestion of specific bands (Fig. 4B and C). This analysis clearly identified the protein as GAP40 and confirmed the presence of GAP45 and GAP50 in the immunoprecipitate. The high molecular weight phosphoprotein, marked with an asterisk in Fig. 4A, was not detected in the SYPRO Ruby stained immunoprecipitate. High level phosphorylation of a low abundance protein is one possible explanation for this discrepancy.

An immunoprecipitate from parasites expressing GFP-tagged MyoA was also analysed by tryptic digestion, with LC-MS/MS analysis of resultant peptides in order to identify associated proteins. Again, 3D7 schizonts in which MyoA was unmodified were used as a negative control. Table 1 shows the proteins identified only in the MyoA-GFP precipitate, by the presence of two or more specific peptides. This approach confirmed the presence of MTIP, GAP50, GAP45 and GAP40. It also identified GAPM2 and GAPM3, which are two of the three GAPM proteins previously shown to associate with the glideosome and IMC (20). The absence of GAPM1 in the precipitate could be explained by the extremely detergent-resistant property of this protein (20). In addition to these known or expected components of the glideosome, 13 additional proteins were identified; whilst it is likely that some of them represent co-purified contaminants, it cannot be excluded that some represent real components of the complex. For example it is unlikely that signal peptide peptidase, MESA and the hexose transporter are part of the glideosome because of their known or predicted subcellular location: the endoplasmic reticulum (33), the parasite-infected erythrocyte surface (34), and the parasite plasma membrane (35), respectively. On the other hand, the protein encoded by PF3D7_1420200 is a tetratricopeptide repeat protein related to TgUNC, demonstrated to be a member of the UCS family of myosin-specific chaperones (15). In addition, of particular interest was the protein encoded by PF3D7_1017500, which is a 134 residue polypeptide with sequence homology to members of the EF-hand superfamily (36) and reminiscent of myosin light chains and other calmodulin-like proteins. This protein is a candidate second light chain for MyoA, particularly because it has been proposed that there is ‘space’ for a second light chain located adjacent to the MTIP binding site (37) and additional light chains have been characterised for MyoA in T. gondii (13, 14). Because of this, and data that will be shown below, we subsequently refer to PF3D7_1017500 as PfELC.

Bioinformatic and functional analysis of PfELC, a novel P. falciparum MyoA light chain

The ELC protein is highly conserved across the Plasmodium genus, with identities ranging from 59 % with P. berghei to 98 % with P. reichenowi (Fig. 5A), however there is only moderate sequence similarity with the T. gondii MyoA ELCs that have been described recently (13, 14). The Toxoplasma gene database, ToxoDB, has recently split the gene encoding TgELC1 into 2 genes—TGME49_269438 and TGME49_269442 in the ME49 strain, erroneously in our opinion. Previous releases showed a single gene—TGME49_069440—and it was using this release that Nebl and colleagues were able to assign peptides to TGME49_069440 from immunoprecipitates of the glideosome (13). Alignment of PfELC and TgELC1 (TGME49_069440) proteins shows 20 % identity. Similarly, a second putative essential light chain, TgELC2 (TGME49_305050) that can bind to TgMyoA in place of TgELC1, shows 21 % identity with PfELC (Fig. 5B). Homology modelling using the Phyre2 structure prediction server (38) suggested considerable structural similarity with calmodulin that is consistent with PfELC being a functional myosin light chain. The entire sequence was modelled on the structure of the calmodulin moiety of a genetically encoded calcium indicator protein (pdb ID: 3U0K) with 76 % confidence (Fig. 5C).

Interactions of PfELC with the glideosome and MyoA

Several approaches were taken to obtain further information on the properties of PfELC. Antibodies were generated to the recombinant protein and used to test for the presence of the protein in immunoprecipitated...
protein complexes from *P. falciparum* schizont lysates. The immunoprecipitate with GFP-specific antibodies from cells expressing either MyoA-GFP or MyoB-GFP was probed with antibodies raised against recombinant PfELC. This analysis revealed a 14 kDa protein associated with MyoA in the glideosome complex, with very little free unbound protein, and none was detected associated with the MyoB complex (Fig. 5D). When the immunoprecipitation was performed with anti-MTIP antibodies and a lysate of schizonts or merozoites from 3D7 parasites, western blotting with the PfELC-specific antibodies detected a 14 kDa protein in the complex, not only confirming the presence of PfELC but also demonstrating that it is present with MTIP in the same complex (Fig. 5E).

PfELC was expressed in recombinant form as a His-tagged fusion protein (Fig. 6A). Far UV-CD spectra indicated that PfELC is mainly helical with 32% alpha helix, 17% beta sheet, and 20% turn (Fig. 6Bi). The addition of 1 mM calcium ions had no effect on either the near- or far-UV spectrum suggesting that calcium does not bind to the protein, or that there is no change in secondary or tertiary structure of the protein upon binding. This is supported by the fact that there are no appropriately placed acidic residues that would normally be involved in the coordination of calcium ions in calmodulin-related proteins with canonical EF-hands (39). To examine whether or not PfELC interacts directly with MyoA, the binding of the recombinant protein to peptides derived from the neck region of MyoA was examined. Amino acid residues 760 to 818 of MyoA constitute its neck region and MTIP has previously been shown to interact strongly with peptides from the C-terminus of the neck (40, 41). Using biolayer interferometry we measured the affinity of PfELC to overlapping biotinylated peptides corresponding to residues 770–787, 786–803, and 801–818 of MyoA. There was no observable binding to peptide 770–787, but both peptides 786–803 and 801–818 bound to PfELC, with calculated $K_d$ of 0.48 [± 0.15] and 0.32 [± 0.07] µM respectively (Fig. 6C). There was no effect of 1 mM calcium on the affinity to any of the three peptides. As MTIP has been shown to bind to the more distal MLC binding site, but not to a peptide representing residues 781–801 of PfMyoA (41), we propose that PfELC binds to a region within 786–803, while MTIP binds to a region within 801–818 (Fig. 6D).

### The subcellular location of PfELC

A parasite line expressing an HA-tagged PfELC was generated by single homologous recombination into the endogenous gene locus (Fig. 7A) and integration into the PfELC locus confirmed by PCR in two independent clones (Fig. 7B). The 17-kDa tagged protein could be detected by western blot analysis of parasite extracts and the time of synthesis corresponded to that of MTIP and GAP45 in mature schizonts, peaking at 44–46 h post-invasion (Fig. 7C). By 48 h after invasion, many of the schizonts had ruptured and the merozoites gone on to invade fresh erythrocytes, explaining the apparent drop in expression of all glideosome components at this time point. Immunofluorescence analysis of fixed blood-stage *P. falciparum* with HA-specific antibodies showed co-localisation of the protein with GAP45 and MTIP in late, segmented schizonts (Fig. 7B).

### Recombinant PfMyoA binds to actin

As a first step to reconstitute the functional complex in vitro we expressed PfMyoA using an *in vitro* transcription/translation system and examined its ability to bind to actin. Soluble GST-tagged PfMyoA was purified (Fig. 8A) and its ability to bind actin was examined using an optical tweezer-based single molecule system (Fig. 8B and C). A total of 244 acto-MyoA binding events were recorded in 2 µM ATP, a concentration chosen based on the affinity of TgMyoA for 2'-3'-O-(N'methylanthraniloyl)-ADP (mant ADP (10). The MyoA working-stroke was 3 nm, as measured from the shift in mean position of the observed event distribution (Fig. 8D).

Specificity of the acto-MyoA interaction was established using two different ATP concentrations of 0.3 µM and 8 µM and measuring the duration of acto-MyoA binding events fitted to a single exponential decay. The data were consistent with a single rate-limiting step, due to ATP binding to the acto-MyoA rigor complex. At 0.3 µM and 8 µM ATP we observed interactions lasting on average 0.1 s ($N_{obs}=79$) and 0.013 s ($N_{obs}=165$) respectively and consistent with Mg,ATP binding and dissociating the acto-MyoA rigor complex with second order kinetics of $\sim 15 \times 10^6 \text{ M}^{-1}.\text{s}^{-1}$ (Fig. 8E).
**MTIP and PfELC are required for maximum PfMyoA-dependent motility in vitro**

To examine the effects of light chain composition on MyoA motility, we expressed and purified recombinant MyoA together with MTIP and MTIP/PfELC. For these studies, we required more material than was possible to generate using the in vitro transcription/translation system that we used in the single molecule PfMyoA studies described in the previous section. We opted for a heterologous expression system that would allow co-expression of PfMyoA and its light chains. Bookwalter and colleagues showed recently that TgMyoA can be expressed well in insect cells, but requires Unc45 to fold correctly (15). Therefore, we followed the same strategy and co-expressed the P. falciparum Unc45 homologue and PfHsp90 together with PfMyoA, and were able to purify functional MyoA in complex with MTIP or with both MTIP and PfELC. Figure 9A shows size exclusion chromatography traces of the two purifications, indicating a pure, homogeneous complex. Fig. 9B shows the complexes separated by SDS-PAGE, indicating that the complexes contain the full complement of light chains included in the co-expression system. A peculiarity is that MTIP migrates as two bands on the SDS-PAGE. We have performed mass spectrometry on each of the two bands to confirm they are both MTIP. We are unsure what is the cause of this aberrant migration. We studied MyoA complexes using an *in vitro* motility assay. MyoA was captured on a cover slip either by coating the surface with nitrocellulose (Fig. 9C) or with an anti His-tag antibody (Fig. 9D), which orients the head domain away from the surface. Without addition of ATP, actin filaments bound to the surface and were non-motile. Following ATP addition the filaments started moving, showing ATP-driven motility (supplemental movie file). We also observed significant actin filament fragmentation even at the lowest surface densities used that would still support smooth filament sliding motility (50 µg/ml myosin protein added to the flow-cell surface). In our assay more than 95% of actin filaments showed smooth gliding movement with only a small number of non-moving filaments. This observation is firm evidence that our MyoA preparation contained very few so-called “dead-heads” (myosin “dead-heads” bind actin in an irreversible, tightly bound state). Nitrocellulose and antibody captured MyoA/MTIP gliding velocities were $0.13 \pm 0.04 \mu m s^{-1}$ ($N_{obs}=414$), and $0.25 \pm 0.06 \mu m s^{-1}$ ($N_{obs}=761$), respectively; showing antibody immobilization increased the gliding velocity approximately two-fold. Furthermore, the MyoA/PfELC/MTIP complex moved filaments approximately two-fold faster than the MyoA/MTIP complex and produced actin gliding velocities of $0.27 \pm 0.06 \mu m s^{-1}$ ($N_{obs}=404$) on nitrocellulose and $0.46 \pm 0.07 \mu m s^{-1}$ ($N_{obs}=329$) when immobilized via anti-His antibody. The maximum actin-activated ATPase activity ($V_{max}$) and actin affinity ($K_m$) were $12.9 \mu M$ and $42 \mu M$ for MyoA/MTIP and 11.2 s$^{-1}$ and 37 µM for MyoA/MTIP/PfELC, respectively (Fig. 9E).

**DISCUSSION**

We have taken advantage of the ability to GFP-tag MyoA to study in detail its time of synthesis and location in the life cycle of the malaria parasite, using fluorescence microscopy with live parasites expressing the tagged gene from the endogenous genomic locus. No growth phenotype was associated with the tagging of MyoA with GFP. The protein is most abundant in the invasive and motile stages. MyoA is an integral component of the glideosome complex, which is associated with the IMC located below the parasite plasma membrane. In the *P. falciparum* asexual blood stage there is no IMC in the ring and trophozoite stages, and genesis of the IMC commences soon after the onset of nuclear division, as part of the establishment of polarity for each developing merozoite and detected as intracellular ring-shaped structures associated with the individual nuclei (28, 29). At this stage MyoA-GFP cannot be detected in the cell. However, as the IMC continues to grow, leading eventually to cell segmentation and individual merozoite formation, MyoA accumulates on the IMC, which can be distinguished from the parasite plasma membrane in mature schizonts because it does not surround the food vacuole. In released extracellular merozoites, MyoA-GFP has the characteristic peripheral location but then rapidly disappears following erythrocyte invasion. The mechanism by which the disappearance of the glideosome...
occurs is undetermined. It is likely that proteasome-mediated degradation of the protein components occurs, though the signalling mechanisms that underlie this are unknown. Throughout the *P. berghei* life cycle MyoA is most abundant in the extracellular motile and invasive forms (merozoites, ookinetes and sporozoites). In the sexual stages although *myoa* mRNA is detected in gametocytes the protein is barely detectable in gametocytes or the zygote, and protein is only evident once ookinete differentiation is underway. Translational repression is a mechanism of post-transcriptional gene regulation in *Plasmodium*, and *myoa* has previously been shown to be regulated in this way (42, 43), with protein synthesis occurring after fertilisation. Following formation of the protuberance that heralds development of the polarised ookinete, MyoA-GFP begins to be deposited at the periphery of the growing protuberance through the retort forms until finally the entire mature motile ookinete is labelled with GFP fluorescence. The deposition of MyoA in this way provides insight into the formation of the IMC in the developing ookinete. It is possible that MyoA activity is important in providing the force required for the developing protuberance to extend away from the spherical body and acquire the characteristic shape of the ookinete. Similarly, the periphery of the salivary gland sporozoite is uniformly labelled with MyoA-GFP. Interestingly, the sporozoite was the only parasite stage in which we observed an additional perinuclear MyoA-GFP localization. It may be that the IMC plays a role in maintaining the shape of the sporozoite. A gene knockout in *P. berghei* of *PbIMC1a*, which encodes a sporozoite-specific protein with homology to articulins and localizes to the sporozoite IMC, resulted in salivary gland sporozoites with an abnormal morphology. The sporozoites had a characteristic bulge at the position of the nucleus, and the nucleus appeared more rounded than the usual slender nucleus of sporozoites (44). It is possible that MyoA plays a role in maintaining the cellular architecture and shape of the sporozoite in addition to its role in cell motility. The early cytomere stage of liver-stage parasites exhibits cytosolic MyoA-GFP fluorescence, with peripheral localization only detected in the late, segmented cytomere stage. It may be that in the early cytomere stage there is a pool of MyoA that is present prior to localization to the IMC in the late cytomere stage.

The GFP-tag has also proved very useful to purify MyoA and associated proteins in *P. falciparum*. This approach confirmed the presence of MTIP, GAP45 and GAP50, identified previously by western blotting (26), and GAPM2 and 3—previously identified as components of the Apicomplexan glideosome (20). A third member of this family (GAPM1) was not detected here, likely because of its poor solubility in the detergent used. We also identified GAP40 as a component of the glideosome and confirmed that it is one of the major phosphorylated proteins in the complex. Although GAP40 has been described as a component of the *T. gondii* glideosome it has not previously been shown to be a component of this structure in *Plasmodium*. The fact that we only detected two peptides attributable to GAP40 (3.3 % sequence coverage) in our immunoprecipitation/mass spectrometry studies is likely due to the hydrophobic nature of this protein. GAP40 has ten predicted membrane-spanning regions, with very small loops between them. The two peptides we detected map to the short loop between predicted transmembrane domains 2 and 3. Hydrophobic peptides, such as those that would make up the majority of a GAP40 tryptic digest, are often very poorly ionised in the mass spectrometry workflow, making detection very difficult. This could also explain why GAP40 has, until now, not been detected as a component of the *Plasmodium* glideosome.

In addition to these known or expected components of the *Plasmodium* glideosome, several additional proteins were also identified. Whilst some such as the signal peptide peptidase or MESA are likely to be simple contaminants due to their known subcellular location, this cannot be said for some of the other proteins. Their presence or absence in complex with MyoA or other components of the glideosome complex needs to be examined in reciprocal immunoprecipitation experiments using, for example a variety of GFP-tagged proteins. The tetratricopeptide repeat-containing protein encoded by gene PF3D7_1420200 is similar in sequence to TgUNC in *T. gondii* (TGME49_249480), a member of the UCS family of myosin chaperones (15). Its presence in our immunoprecipitations supports a
conserved mechanism of myosin folding in a diverse range of organisms (45). Another of the proteins identified by mass spectrometry, coded by the gene PF3D7_1017500 and annotated as a conserved protein of unknown function, has a sequence suggesting that it might be a myosin light chain. It shows relatively weak homology with T. gondii ELCs, indicating that in this case it would not have been possible to identify the Plasmodium ELC by this criterion alone. Interestingly, the abundance of this protein—like that of other glideosome components including MyoA and MTIP—was significantly decreased in parasites blocked during schizont development (46). Unlike the situation in T. gondii tachyzoites, where two ELCs have been identified that bind to TgMyoA in a mutually exclusive manner (14), we found no evidence of light chains other than MTIP and ELC in our immunoprecipitations. This may indicate different modes of regulation and control of the glideosome in these species. Interestingly, there is redundancy in the ELCs of TgMyoA. Either of the genes encoding TgELC1 or 2 can be deleted with no effect on parasite viability, however deletion of both genes causes a marked reduction in the parasite’s ability to invade host cells (14). We have been unable to disrupt the elc gene in either P. berghei or P. falciparum, suggesting that the ELC is essential for parasite viability and that this redundancy does not exist in Plasmodium species. This is supported by the inability to disrupt Pfelec in the PlasmoGEM project (Plasmodb.org, RMgm-1936) (47). Utilizing approaches that are now available in P. falciparum, such as the expression of a Cre recombinase that dimerizes in response to the presence of a small molecule to rapidly and inducibly delete genes (48), will be essential in order to further analyze the specific deficits that occur in parasite growth and development in the absence of PfELC.

We tested the binding of PfELC to several peptides derived from the neck region of MyoA, which encompass the MTIP binding site at the C-terminus of the molecule and the adjacent putative light chain binding site previously identified by homology modelling (37). PfELC bound to both the known MTIP binding site and the adjacent proximal site. It has been shown previously that MTIP cannot bind to a peptide comprised of residues 781–801 of the MyoA neck (41). This covers all of the sequence of peptide 786–801 used in these studies, which we have shown binds strongly to PfELC. These data, along with the co-precipitation of PfELC with MTIP and MyoA from parasite lysates, confirm that MyoA is decorated with two light chains and that PfELC binds to sequences of MyoA adjacent to the MTIP binding site. A 30 amino acid region of the T. gondii MyoA neck (residues 770–799) has been demonstrated to be the binding site for TgELC1 and 2 (14). The region of PfMyoA that we have demonstrated binds to PfELC is at the C-terminus of the equivalent region of PfMyoA, suggesting a similar arrangement of the two light chains on both TgMyoA and PfMyoA. We found no evidence of an effect of Ca2+ on the binding of PfELC to MyoA neck peptides or on the structure of PfELC itself, as judged by biolayer interferometry and circular dichroism, respectively. Studies in T. gondii have demonstrated Ca2+-binding by ELC1 and, by sequence homology it is assumed that TgELC2 will also bind Ca2+. Specific amino acid residues have been suggested (D15, D17 and D19) or demonstrated (D15) to be required for Ca2+ coordination in TgELC1 (14). In the equivalent region of PfELC these residues are S17, C19, and D21 respectively. The sequence differences here support our observations that Ca2+ has no effect on the overall structure of PfELC or on its binding to PfMyoA neck peptides, in that the acidic residues required to coordinate Ca2+ in canonical EF-hand motifs (39) are absent. This suggests that there may be subtle differences in the regulation of ELC binding in the Plasmodium versus Toxoplasma glideosome.

Additionally, we expressed and purified recombinant MyoA that bound to actin, and recombinant MyoA decorated with MTIP or MTIP/ELC that we used to measure sliding velocities by in vitro motility assays. The motility was induced by the addition of ATP, and only a small proportion of actin filaments were static, suggesting only a small fraction of non-functional or incorrectly folded myosin heads after a two-step purification. Importantly, MyoA with both MTIP and ELC bound had about a two times higher velocity in comparison to MyoA with MTIP alone. This suggests that ELC plays a significant role in Plasmodium motility. Our results are in line with results using TgMyoA, which moves actin with approximately two times higher
velocity when decorated by two light chains (15). It would be interesting to know whether the step size of the MyoA/MTIP/ELC complex is increased compared to that of the MyoA/MTIP complex, as this could be a possible explanation for the increased velocity that we observe in our motility studies for MyoA with its full complement of light chains. Since the actin-activated ATPase activity of MyoA/MTIP and MyoA/MTIP/PfELC was virtually identical, our conclusion is that the increased in vitro motility sliding velocity seen with MyoA/MTIP/PfELC compared to MyoA/MTIP alone cannot be explained by a change in catalytic turnover rate (ATPase activity). It is most easily explained by the MTIP/ELC light-chain combination having a longer effective lever arm and "working-stroke" distance, meaning that actin is moved approximately twice as far for each molecule of ATP hydrolyzed.

The glideosome is a molecular motor that has a key role throughout the parasite life cycle. The key components of the glideosome for moving actin filaments are MyoA and its two associated light chains, and we have now established a system where the interplay and function of these three glideosome components can be studied. This will also allow the development and assessment of inhibitors that target the glideosome function and block invasion of host cells.

**EXPERIMENTAL PROCEDURES**

**Ethics statement for animal work**
Animal work at the University of Nottingham was approved by the local Animal Welfare and Ethical Review Body and by the United Kingdom Home Office under project licence numbers 40/3344 and 30/3248 in accordance with the United Kingdom Animals (Scientific Procedures) Act 1986 and in compliance with European Directive 86/609/EEC for the protection of animals used for experimental purposes. Six to eight-week-old female Tuck-Ordinary (TO) outbred mice (Harlan) were used for these experiments. At the University of Bern experiments were conducted in strict accordance with the guidelines of the Swiss Tierschutzgesetz (TSchG; Animal Rights Laws) and approved by local authorities. The BALB/c mice used in these experiments were between 6 and 10 weeks of age and bred in the central animal facility of the University of Bern.

**Analysis of PbMyoA gene expression by quantitative (q) RT-PCR.**
Blood stage *P. berghei* parasites from infected mice (day 4 post-infection) were cultured *in vitro* (49) and the following day schizont-infected cells were purified on a 60 % v/v Nycodenz/PBS gradient. Gametocytes were purified on a 48 % (v/v) Nycodenz gradient, washed twice in RPMI 1640 medium and gamete formation activated by transfer to ookinete medium (RPMI 1640 containing 25 mM HEPES, 25 % fetal bovine serum, 10 mM sodium bicarbonate, 50 mM xanthurenic acid at pH 7.6) for 30 min at 20 °C. For ookinete preparation, parasites from mice day 5 post-infection were placed in ookinete medium for 24 h at 20 °C, other parasite stages were lysed in red blood cell lysis buffer for 30 min and then ookinetes were purified on a 63 % Nycodenz gradient (v/v in PBS). On day 14 post-feeding, mosquito guts were dissected and crushed in a loosely fitting homogeniser to release sporozoites, which were then quantified using a haemocytometer (50).

Total RNA isolation, cDNA synthesis and PCR analysis were as described previously (49). Expression was determined using the Pfaffl method (51), using *hsp70* (PBANKA_0818900) and arginyl-tRNA synthetase (PBANKA_1434200) as reference genes. All primers sequences are available on request.

**Parasite genetic modification**

**GFP-tagging of P. berghei genes**
*P. berghei myoa* (PBANKA_1355700) was modified by single homologous recombination to insert a sequence coding for a C-terminal GFP tag (Fig. 1B), using the p277 vector that contains a human dhfr selection cassette (52). An 833 bp region of *Pbmyoa* starting 1.86 kb downstream of the ATG start codon and omitting the stop codon was amplified using primers 7 and 8 and inserted into the vector using KpnI and ApaI restriction sites. The construct was linearized by digestion with PacI and used to transfect the *P. berghei* ANKA 2.34 line by electroporation (53). The electroporated parasites were injected into naïve mice and transgenic parasites were selected with pyrimethamine (49) and analyzed by diagnostic PCR using the primers
9 and 10 to determine correct integration of the gfp sequence at the targeted locus (Fig. 1C). Western blotting with GFP-specific antibodies was used to confirm expression of full-length tagged protein of the predicted size (Fig. 1D).

**Tagging of *P. falciparum* genes**

*P. falciparum* (3D7) was cultured in RPMI 1640 medium containing Albumax II according to previously described methods (54). GFP-tagging of *P. falciparum* MyoB, MyoA, and GAP45 has been described previously (26, 29). For HA-tagging of *P. falciparum* ELC at its C-terminus, a 632 bp DNA fragment of the gene was amplified using primers 11 and 12 and cloned via XmaI and AvrII sites into the pHH4-HA vector (E. Knuepfer, unpublished), a derivative of pHH1 (55). 100 μg DNA was used to transfect ring-stage *P. falciparum* as described previously (56). Transgenic lines were selected with 10 nM WR99210 (a gift from Jacobus Pharmaceuticals). Plasmid integration was monitored by diagnostic PCR screening with primers 13 and 14 to detect the unmodified locus and primer pair 13 and 15 to detect integration (Fig. 7B).

**Antibodies**

Polyclonal rat anti-PfMyosin A antibodies were raised to a peptide of the sequence MVNKINELNNYFRINSTFINKSENE by Peptides Ltd. according to their standard protocol. Mouse anti-PfELC antibodies were raised by immunisation of female BALB/c mice with recombinant His-tagged protein (production of which is described elsewhere in this manuscript). Other primary polyclonal antibodies and monoclonal antibodies (mAbs) used were anti-PfMTIP (40), anti-PfGAP45 (18), anti-PfGAP50 (26) anti-PfRON4 mAb (57), and anti-PbP28 mAb 13.1 (58), which are described in the referenced manuscripts. Rabbit anti-GFP (Knuepfer and Holder, unpublished) and rat anti-HA (monoclonal 3F10; Roche) antibodies were used to detect tagged proteins. Species-specific Alexafluor-488 and 594 conjugated secondary antibodies (Life Technologies) were used to visualize primary antibody binding. All antibody dilutions were carried out with 3 % BSA (w/v) in PBS at 4 °C overnight. Primary antibodies were diluted as required; species-specific Alexafluor-488 and 594 conjugated secondary antibodies (Life Technologies) were used to visualize primary antibody binding. All antibody dilutions were carried out with 3 % BSA (w/v) in PBS. Slides were mounted for microscopic examination using Prolong Gold anti-fade reagent with DAPI (4',6-diamidino-2-phenylindole, Life Technologies). Slides were viewed using a Zeiss Axioplan 2 microscope, images captured using a Zeiss AxioCam MRC digital camera and Axiovision 4.8.2 software, and prepared for publication using Adobe Photoshop.

**Microscopy**

**Live imaging of GFP-tagged parasite lines**

*P. berghei* asexual blood stage infections in mice were initiated by intraperitoneal (i.p.) injection of parasite-infected blood. The asexual, gametocyte and ookinete stages were observed using Hoechst 33342 dye (Molecular Probes) in ookinete medium and Cy3-conjugated mouse mAb 13.1 to recognise the p28 protein on the surface of activated female gametes, zygotes and ookinete. To examine mid-gut infection, guts were dissected 14 days after mosquito feeding on infected blood and mounted under Vaseline-rimmed cover slips after staining with Hoechst 33342 for 10–15 min. On day 21 post-feeding, guts and salivary glands were dissected and crushed separately in a loosely fitting homogeniser to release sporozoites, which were then quantified using a haemocytometer and used for imaging. Microscopy was performed using a Zeiss AxioImager M2 microscope (Carl Zeiss, Inc.) fitted with an AxioCam MRm digital camera. *P. berghei* liver stage parasites were prepared and imaged as described previously (26), with the exception that infections were performed in HeLa cells.

*P. falciparum*-infected red blood cells were prepared for live imaging as described previously (26, 29) and viewed using an Axio Imager M1 microscope (Zeiss).

**Indirect immunofluorescence assay**

Thin smears of parasites were fixed with 4 % paraformaldehyde for 30 min at room temperature. The fixed cells were permeabilized with PBS containing 0.1 % Triton X-100 for 5 min followed by blocking in 3 % BSA (w/v) in PBS at 4 °C overnight. Primary antibodies were diluted as required; species-specific Alexafluor-488 and 594 conjugated secondary antibodies (Life Technologies) were used to visualize primary antibody binding. All antibody dilutions were carried out with 3 % BSA (w/v) in PBS. Slides were mounted for microscopic examination using Prolong Gold anti-fade reagent with DAPI (4',6-diamidino-2-phenylindole, Life Technologies). Slides were viewed using a Zeiss Axioplan 2 microscope, images captured using a Zeiss AxioCam MRc digital camera and Axiovision 4.8.2 software, and prepared for publication using Adobe Photoshop.

To image stages of invasion of erythrocytes, tightly synchronized PfMyoA-GFP-expressing late stage schizonts were mixed with erythrocytes and samples taken after 2, 5, 8, 10 and 30 min and fixed in solution with 4 % paraformaldehyde with
0.01 % glutaraldehyde for 60 min at room temperature. Further processing was carried out as described previously (59).

**Preparation of parasite lysates, immunoprecipitation of GFP-fusion proteins and analysis of the precipitates by western blotting and LC-MS/MS**

**Parasite lysates**

Schizonts from MyoA-GFP, GAP45-GFP and 3D7 *P. falciparum* parasite lines were first lysed in 0.15 % (w/v) saponin in PBS, harvested by centrifugation and then proteins were extracted using ten cell pellet volumes of ice-cold lysis buffer (0.5 % NP40, 150 mM NaCl, 10 mM Tris HCl pH 7.5, 1x Complete protease inhibitors [Roche]). Samples were cleared by centrifugation at 100,000 x g for 10 min. The detergent concentration in the samples was reduced to 0.2 % by dilution with 150 mM NaCl, 10 mM Tris pH 7.5, 1x Complete protease inhibitors. For *P. berghei*, subcellular fractionation was performed as described previously (60). Briefly, cell pellets of MyoA-GFP or WT-GFP schizonts were resuspended in hypotonic lysis buffer (10 mM Tris-HCl pH 8.4, containing 5 mM EDTA and protease inhibitors), freeze/thawed twice and then incubated for 1 hr at 4°C. Following centrifugation at 100,000 g for 30 min, supernatant was collected as the soluble protein fraction (S). The remaining pellet was re-suspended in carbonate solution (0.1M Na2CO3, pH 11.0, containing protease inhibitors), incubated for 30 min at 4°C and then centrifuged at 100,000 g for 30 min. The resulting supernatant was classified as the peripheral membrane fraction (PM), and the pellet was washed and solubilised in 4% SDS and 0.5% Triton X-100 in PBS and classified as the integral membrane fraction (IM). Equal amounts of these three fractions were analyzed by western blot using anti-GFP antibody.

**Immunoprecipitation**

Parasite lysates were pre-cleared by incubation with blocked agarose beads for 1 h, then the supernatant was incubated with GFP-Trap agarose beads for 2 h at 4 °C with end-over-end rotation. The beads were washed extensively with 150 mM NaCl, 10 mM Tris pH 7.5, then subjected to a further wash in 300 mM NaCl, 10 mM Tris pH 7.5. Finally, the beads were resuspended in 5 volumes of 2x SDS-PAGE sample buffer and boiled for 5 min prior to fractionation of the proteins by SDS-PAGE.

**Western blotting**

Cell lysates were separated under reducing conditions on NuPAGE gels (Life Technologies). Proteins were transferred to nitrocellulose and blocked in 5 % w/v non-fat milk in PBS containing 0.2 % Tween-20. Primary antibodies were diluted in 5 % w/v non-fat milk in PBS containing 0.2 % Tween-20. Species-specific horseradish peroxidase-conjugated secondary antibodies (Biorad) were used and the signal developed using enhanced chemiluminescence (GE Healthcare). Blots were exposed to Biomax MR film (Kodak) and images prepared for publication using Adobe Photoshop.

**Protein identification by LC-MS/MS**

Proteins immunoprecipitated from GFP-GAP45 parasite lysates were run on a 10 % NuPAGE BisTris gel (Life Technologies) and stained with SYPRO Ruby (Thermo Scientific). The protein bands were visualised using a Pharors FX Plus Molecular Imager and excised using EXQuest Spot Cutter (both Biorad). Proteins immunoprecipitated from MyoA-GFP or 3D7 parasite lysates were run 4 mm into a 10 % NuPAGE BisTris gel (Life Technologies), then excised using a clean scalpel blade. In both cases proteins were reduced for 1 h in a solution of 20 mM DTT, 200 mM ammonium bicarbonate (ABC), 50 % acetonitrile (ACN), and alkylated for 20 min in a solution of 5 mM iodoacetamide, 200 mM ABC, 50 % ACN, prior to overnight trypsin
digestion. The resulting digests were analyzed by LC-MS/MS using an Ultimate 3000 nanoRSLC HPLC, equipped with a 50 cm x 75 μm Acclaim Pepmap C18 column, coupled to an LTQ Orbitrap Velos Pro equipped with a Nanoflex electrospray source (all Thermo Scientific). A gradient of 6–32 % ACN/0.1 % formic acid over 48 min was used at a flow rate of 0.3 μl/min. The Orbitrap was operated in Data Dependent Acquisition mode with a survey scan at 60,000 resolution and up to the 10 most intense ions selected for MS/MS. Raw files were processed using Proteome Discoverer (PD) 1.3 (Thermo Scientific) with Mascot 2.4 (Matrix Science, UK) as the search engine against the appropriate protein database. A decoy database of reversed sequences was used to filter the results at a false detection rate of 1 %.

Bioinformatic analyses
Alignment of ELC sequences was performed using CLC Sequence Viewer 6 (Qiagen) and graphics prepared using BoxShade. A structural model of PfELC was constructed using the Phyre 2 protein fold recognition server (38). Images were prepared using CCP4 Molecular Graphics software (61).

Cloning and expression of recombinant PfELC
RNA was prepared from late P. falciparum parasites (approximately 36 h post-invasion) using TRizol (Life Technologies) and cDNA made from this using the Reverse Transcription system with oligo dT primers (Promega). Primers 16 and 17 were used to amplify elc from cDNA, and the product was cloned into the pET46-Ek-LIC expression vector (Novagen) using ligation-independent cloning. After verification by sequencing, the plasmid was transformed into BL21 (DE3) cells (Stratagene), and protein expression induced by the addition of isopropyl β-D-1-thiogalactopyranoside (IPTG) to 1 mM for 3 h at 37 °C. The cell pellet was lysed using Bugbuster (Novagen) with 25 U Benzonase/ml and 1x Complete Protease Inhibitors without EDTA (Roche). The lysate was cleared by centrifugation at 20,000 x g and was applied to a column of Ni-NTA agarose (Qiagen), followed by extensive washing of the column with 20 mM imidazole in phosphate buffer (GE Healthcare). The protein was eluted in three column-volumes phosphate buffer with 0.5 M imidazole. The eluate containing the His-ELC protein was collected and concentrated using a Vivaspin 20 concentrator with a 5 kDa molecular weight cut-off (Sartorius). Buffer exchange into 50 mM NaCl, 10 mM Tris HCl, pH 8.0 was carried out using a PD10 column (GE Healthcare).

Biophysical analysis of PfELC and its interaction with MyoA neck peptides
Peptides
Overlapping peptides representing amino acids Q770–A787, A786–S803, and 1801–Q818 of the PfMyoA neck were synthesised by Mimotopes with biotin at the N-terminus. Each peptide had an SGSG linker between the biotin and 18 amino acids of the MyoA neck. The C-termini of the peptides were capped with an amide to remove unnatural charge, apart from peptide 801–818, which was left as a free acid.

Far-UV CD
Far-UV CD spectra were recorded on a Jasco J-815 spectropolarimeter fitted with a cell holder thermostatted by a CDF-426S Peltier unit. CD measurements were made at a protein concentration of 10 µM PfELC in 25 mM Tris HCl pH 8.0, 50 mM NaCl using fused silica cuvettes with 1 mm path length (Hellma). All spectra were recorded with 0.1 nm resolution and baseline corrected by subtraction of the buffer spectrum. Secondary structure content was estimated using methods described by Sreerama and Woody (62).

Biolayer interferometry
P. falciparum ELC binding to peptides was analyzed by biolayer interferometry using the Octet Red system (ForteBio) at 25 °C in 96-well microplates, as described previously (63). All assays were performed in 100 mM Tris HCl pH 8.0 containing 0.005 % v/v Tween-20, according to the manufacturer’s instructions. A baseline was established over 4 min, and biotinylated peptides (0.4 µg/ml) were loaded onto streptavidin sensors for 10 min. The sensors were washed for 4 min and then incubated with PfELC at different concentrations (0.04–5 μM) for 800 s. The data were analyzed to calculate a dissociation constant (K_d) using non-linear regression with a one-site binding model and a least-squares fit (GraphPad Prism 6).
In vitro expression and purification of recombinant MyoA

MyoA was expressed in a wheat germ cell free protein synthesis system (ENDEX® Technology, CellFree Sciences) (64). Recombinant plasmid DNA using recodomer template DNA was amplified using specific primers and cloned into the vector PEU-E01G-TEV-N1 to express N-terminal GST-tagged protein. For the in vitro transcription reaction, 2 µg DNA was added to a transcription premix (11.5 µl nuclease free water, 1x transcription buffer, 2.5 mM NTP mix, 1 U/µl RNase inhibitor, 1 U/µl SP6 RNA Polymerase) in a total volume of 20 µl, and incubated at 37 °C for 6 h, according to the manufacturer’s instructions. The quality of mRNA produced was monitored by gel electrophoresis. The Bilayer Reaction System (CellFree Sciences) was used to translate mRNA into protein. mRNA (10 µl) was added to 10 µl of wheat germ extract (WEPRO® 1240) supplemented with 40 ng/µl creatine kinase, and carefully transferred directly below the SUB-AMIX solution (0.3 mM of each L-amino acid, 100 mM potassium acetate, 2.7 mM magnesium acetate, 16 mM disodium creatine phosphate, 0.4 mM spermidine, 1.2 mM ATP, 0.25 mM GTP, 4 mM dithiothreitol, and 30 mM HEPES-KOH [pH 7.8]), and incubated for 20 h at 15-16 °C. Soluble GST-tagged MyoA was purified using glutathione sepharose beads (GE Healthcare); supernatant was incubated with the resin for 2 h at 4 °C, followed by extensive washing with 20 mM Tris HCl pH 8.0, 250 mM NaCl, 0.5 mM TCEP. The beads were resuspended in 3 volumes 20 mM Tris HCl pH 8.0, 100 mM NaCl, 10 mM reduced glutathione in order to elute the GST-tagged protein. This was followed by buffer exchange into PBS on a PD10 column (GE Healthcare).

Single molecule optical tweezer studies to examine the MyoA interaction with actin

The optical tweezer (OT) assay, developed to measure the force and movement produced by individual myosin molecules interacting with an actin filament, was used in the well-established three bead format with the experimental conditions as described (65-67). The experimental apparatus is based around a Zeiss inverted microscope and experiments were performed in a flow-cell chamber created using a microscope slide and cover-slip. The solution buffers and other conditions were identical to those used in earlier studies (67). A single, rhodamine-phalloidin labelled, actin filament, comprising 1:5 biotin-labelled actin to plain actin, was captured between two neutravidin-coated beads (~1 µm diameter) using the dual-beam optical tweezers. A third, surface-immobilised bead that was sparsely coated with MyoA was then positioned in close proximity to the actin filament. Interactions between the surface-bound MyoA and the actin filament were recorded by monitoring movement of the beads held in the optical tweezers using two 4-quadrant, photodiode detectors. To facilitate visualization, the beads were moved backwards and forwards using a 100 nm amplitude, 10 Hz triangular wave. Individual events were then automatically detected by measuring changes in amplitude of the imposed triangular oscillation and event amplitudes were histogrammed. Data were acquired and saved to computer memory using custom software, with a sampling rate of 10 kHz, and later analyzed using IgorPro software (WaveMetrics Inc., Lake Oswego, Or, USA). Individual acto-MyoA force-producing events were identified as described previously (65) and the MyoA power stroke amplitude and kinetics were analyzed by histogram analysis of the data.

Expression in insect cells and purification of recombinant MyoA with light chains

Following the example of Bookwalter et al. (15), we co-expressed PfHsp90 and Unc45 chaperones together with full-length MyoA, PfELC and MTIP in Sf21 cells. By sequence homology, PF3D7_1420200 is the P. falciparum orthologue of Toxoplasma Unc45 (PlasmoDB.org). Codon-optimised genes were designed and synthesised (Life Technologies) for insect cell expression. A codon-optimised Hsp90 (PF3D7_0708400) was a kind gift from Dr. Will Stanley. PfUnc45 was cloned under the polyhedrin promter in the pFL vector and Hsp90 under the control of p10 promoter in the same vector. MyoA (PF3D7_1342600) was cloned into the pFL vector and fused with N-terminal StrepTagII and gfp under the polyhedrin promoter. MTIP (PF3D7_1246400) and ELC (PF3D7_1017500) genes were...
cloned into the pFL vectors under the control of the polyhedrin promoter with N-terminal hexa-histidine tags. Each construct was transformed in MAX Efficiency DH10Bac cells (Invitrogen), bacmids were isolated, and the gene integrations were confirmed using standard protocols (Bac-to-Bac, Invitrogen). Sf21 cells (Gibco) were transfected using FuGene 6 (Promega) as described (69). After two rounds of virus amplification, Sf21 cells were co-infected with untitered recombinant viruses in the ratios of 0.75:1:1:1 for MyoA:Unc45:MTIP:ELC or 0.75:1:1 for MyoA:Unc45:MTIP, and harvested cells 3 days post-infection. We used 1.5 ml of MyoA-expressing virus per 1 x 10^8 cells, the others 2 ml per 1 x 10^8 cells. Cells were collected, washed with PBS, resuspended in the lysis buffer (20 mM Hepes pH 7.5, 5 % trehalose-dihydrate (w/v), 1 mM ATP, 0.5 mM TCEP, 2.5 mM MgCl2, 0.5 mg avidin, 1X SigmaFast EDTA free protease inhibitor), sonicated, and clarified at 38,000 g. Remaining supernatant was applied into two inter-connected 1 ml Strept-Tactin superfow columns (IBA GmbH), washed with lysis buffer, washing buffer 1 (20 mM Hpes pH 7.5, 5 % trehalose-dihydrate(w/v), 150 mM KCl, 0.5 mM TCEP), and washing buffer 2 (20 mM Hpes pH 7.5, 5 % trehalose-dihydrate(w/v), 0.5 mM TCEP). MyoA complexes were eluted with 20 mM Hepes pH 7.5, 5 % trehalose-dihydrate(w/v), 0.5 mM TCEP). MyoA complexes were applied for 2 min, before washing the flow cell twice with AB/BSA, and applying rhodamine-phalloidin labelled F-actin in AB/COG for 2 min. We used MyoA/MTIP and MyoA/MTIP/PfELC at concentrations of 0.25 and 0.27 mg/ml for nitrocellulose capture, and 0.11 and 0.05 mg/ml for anti His-tag capture, respectively. The flow-cell was washed twice with AB/COG and then viewed by fluorescence microscopy. Assays were performed using an Axioskop 40 fluorescence microscope with a Zeiss PlanNeofluar 100x 1.3 NA objective lens. Fluorescence was excited by a mercury arc lamp using a rhodamine filter set (Excitation filter HQ535/50, dichroic mirror Q565LP and emission filter HQ605/75; Chroma technology) and light emitted from the rhodamine-phalloidin labelled actin filaments specimen was imaged onto an image-intensified charge-coupled device camera (IC-310 Photon Technology International) where, at the magnification used, each pixel corresponded to 100 nm at the object plane. Sequences of video frames were captured every 40 ms using a frame grabber card (Multipix Imaging Ltd). The laboratory was air-conditioned and the experimental temperature kept at 23 °C. Filament specimens were tracked automatically with GMimPro (71) using non-default settings Q-threshold 3, and track length 5, after saving each image sequence as a median of five consecutive image blocks. Non-motile or short tracks were a hypodermic syringe to prevent oxygen contamination. An oxygen scavenger, COG, (20 mM DTT, 0.2 mg/ml glucose oxidase, 0.5 mg/ml catalase, 3 mg/ml glucose, 0.5 mg/ml BSA) was added to buffers used during fluorescence imaging, and stored at room temperature during experiments. Microscope flow-cell chambers were constructed using double-sided tape (Tesa), and flow-cell volume was approximately 8–12 µl. To capture MyoA/light chain complexes by anti-His antibody, nitrocellulose-coated coverslips were incubated with neutravidin (Thermo Fisher Scientific) for 4 min. The flow-chamber was then washed four times with buffer AB and then blocked (2x) with AB/BSA (AB with 0.5 mg/ml BSA) for 2 min. Biotinylated anti-His antibody (Qiagen) was diluted 1:5 in AB/BSA, and applied into the flow cell for 4 min. The non-bound antibodies were removed by washing twice with AB/BSA, and MyoA complexes were applied for 2 min, before washing the flow cell twice with AB/BSA, and applying rhodamine-phalloidin labelled F-actin in AB/COG for 2 min. Motility assays were essentially performed as in Butt et al. (70). Our assay buffer, AB (25 mM imidazole-HCl, 25 mM KCl, 1 mM EGTA, 4 mM MgCl2; pH 7.4) was repeatedly degassed, and flushed with argon and stored in

In vitro motility assays

Motility assays were essentially performed as in Butt et al. (70). Our assay buffer, AB (25 mM imidazole-HCl, 25 mM KCl, 1 mM EGTA, 4 mM MgCl2; pH 7.4) was repeatedly degassed, and flushed with argon and stored in

a hypodermic syringe to prevent oxygen contamination. An oxygen scavenger, COG, (20 mM DTT, 0.2 mg/ml glucose oxidase, 0.5 mg/ml catalase, 3 mg/ml glucose, 0.5 mg/ml BSA) was added to buffers used during fluorescence imaging, and stored at room temperature during experiments. Microscope flow-cell chambers were constructed using double-sided tape (Tesa), and flow-cell volume was approximately 8–12 µl. To capture MyoA/light chain complexes by anti-His antibody, nitrocellulose-coated coverslips were incubated with neutravidin (Thermo Fisher Scientific) for 4 min. The flow-chamber was then washed four times with buffer AB and then blocked (2x) with AB/BSA (AB with 0.5 mg/ml BSA) for 2 min. Biotinylated anti-His antibody (Qiagen) was diluted 1:5 in AB/BSA, and applied into the flow cell for 4 min. The non-bound antibodies were removed by washing twice with AB/BSA, and MyoA complexes were applied for 2 min, before washing the flow cell twice with AB/BSA, and applying rhodamine-phalloidin labelled F-actin in AB/COG for 2 min. We used MyoA/MTIP and MyoA/MTIP/PfELC at concentrations of 0.25 and 0.27 mg/ml for nitrocellulose capture, and 0.11 and 0.05 mg/ml for anti His-tag capture, respectively. The flow-cell was washed twice with AB/COG and then viewed by fluorescence microscopy. Assays were performed using an Axioskop 40 fluorescence microscope with a Zeiss PlanNeofluar 100x 1.3 NA objective lens. Fluorescence was excited by a mercury arc lamp using a rhodamine filter set (Excitation filter HQ535/50, dichroic mirror Q565LP and emission filter HQ605/75; Chroma technology) and light emitted from the rhodamine-phalloidin labelled actin filaments specimen was imaged onto an image-intensified charge-coupled device camera (IC-310 Photon Technology International) where, at the magnification used, each pixel corresponded to 100 nm at the object plane. Sequences of video frames were captured every 40 ms using a frame grabber card (Multipix Imaging Ltd). The laboratory was air-conditioned and the experimental temperature kept at 23 °C. Filament specimens were tracked automatically with GMimPro (71) using non-default settings Q-threshold 3, and track length 5, after saving each image sequence as a median of five consecutive image blocks. Non-motile or short tracks were
removed manually. From resulting average velocities, frequency distribution histograms were calculated (a bin size of 0.02) and a Gaussian curve was fitted with GraphPad Prism 7.

**Actin-activated ATPase assay**
The actin-activated ATPase of MyoA/MTIP and MyoA/MTIP/PfELC was estimated using an NADH-coupled assay to quantify the rate of ATP hydrolysis. The assay buffer contained 160 U/ml lactate dehydrogenase, 400 U/ml pyruvate kinase, 1 mM phospho(enol) pyruvate, 0.5 mM ATP and 0.3 mM NADH in the motility assay buffer, AB. The myosins (MyoA/MTIP or MyoA/MTIP/PfELC) were added to a final concentration of 0.13 µM and phalloidin-stabilized filamentous actin was added over the range of 7 to 130 µM. The rate of ATP hydrolysis was calculated from the rate of change in NADH absorbance measured in a temperature-controlled spectrophotometer at 340nm. Experiments were conducted at 23 °C. The data were fit to the Michaelis-Menten equation using non-linear regression with a least squares fit in GraphPad Prism.

**Primers**
Sequences of primers used in these studies are available on request.

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**Competing interests**
The authors declare that they have no conflicts of interest with the contents of this article.

**Authors' contributions**
JLG assisted with conception of the project, experimental design, interpretation of results, preparation of figures and writing of the manuscript, as well as the design, execution and analysis of experiments on the novel ELC. RJW constructed the MyoA-GFP *P. berghei* parasite and analyzed MyoA expression throughout the asexual blood and insect stages of the life cycle. JV performed experiments with insect cell expressed recombinant MyoA, and assisted with the writing of the manuscript. NAY constructed the MyoA-GFP *P. falciparum* parasite and analyzed MyoA-GFP expression and interaction with other glideosome partners. MAMR designed and executed the experiments to identify the expression of GAP40 in the glideosome. RRS analyzed the expression of MyoA-GFP in the hepatic stage of the parasite life cycle. JS analyzed the expression of MyoA at all stages of the *P. berghei* life cycle and was responsible for preparation of these figures. EK designed plasmids to genetically modify *P. falciparum*, developed the methods to examine erythrocyte invasion and contributed to the analysis of these data. DB assisted with all the work to produce genetically modified *P. berghei* and their maintenance and analysis. SRM was responsible for the design and analysis of the data from all the biophysical experiments with ELC. SAH performed and interpreted all the mass spectrometry analysis of proteins and their complexes. IP prepared the MTIP and ELC baculovirus constructs and viruses. RWM assisted with the experiments to generate tagged MyoA in *P. falciparum* and the analysis and interpretation of the expression by microscopy. JEM and IK
analyzed and interpreted the work involving recombinant MyoA and assisted with the writing of the manuscript. RT organised, coordinated, analyzed and interpreted all the work involving the use of *P. berghei* and assisted with the writing of the manuscript. AAH conceived and coordinated the project, analyzed and interpreted the data, and organised and wrote large sections of the manuscript. All authors read and approved the final manuscript.

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Table 1. Identification of proteins immunoprecipitated by GFP-TRAP affinity resin together with *P. falciparum* MyoA-GFP using mass spectrometry.
All proteins were absent from a control *P. falciparum* immunoprecipitation where MyoA was unmodified. Proteins are ranked by the percentage sequence coverage and previously known components of the glideosome/IMC are shaded grey.

| Accession number | Protein description | MW [kDa] | Coverage [%] | Peptides |
|------------------|---------------------|---------|-------------|---------|
| PF3D7_1342600    | myosin A (MyoA)     | 92.2    | 72.5        | 66      |
| PF3D7_1246400    | myosin light chain 1, myosin A tail domain interacting protein (MTIP) | 23.5 | 31.4 | 7 |
| PF3D7_0918000    | glideosome-associated protein 50 (GAP50) | 44.6 | 29.8 | 9 |
| PF3D7_1017500    | conserved Plasmodium protein, unknown function | 15.7 | 24.6 | 2 |
| PF3D7_1033200    | early transcribed membrane protein 10.2 (ETRAMP 10.2) | 38.9 | 22.8 | 6 |
| PF3D7_1222700    | glideosome-associated protein 45 (GAP45) | 23.6 | 19.6 | 4 |
| PF3D7_0822900    | conserved Plasmodium protein, unknown function | 138.2 | 16.2 | 15 |
| PF3D7_1444300    | putative 1-acyl-sn-glycerol-3-phosphate acyltransferase | 33.7 | 12.7 | 2 |
| PF3D7_0423500    | glideosome-associated protein with multiple membrane spans 2 (GAPM2) | 42.6 | 12.4 | 2 |
| PF3D7_1416100    | putative GTP-binding protein, root hair defective 3 (RHD3) homologue | 110.5 | 12.2 | 10 |
| PF3D7_1409400    | conserved Plasmodium protein, unknown function | 30.5 | 10.7 | 5 |
| PF3D7_1032500    | putative DER1-like protein, (Der1-2) | 31.0 | 10.3 | 3 |
| PF3D7_1457000    | signal peptide peptidase (SPP) | 47.5 | 8.0 | 3 |
| PF3D7_0515700    | putative glideosome-associated protein 40 (GAP40) | 51.8 | 7.5 | 4 |
| PF3D7_0500800    | mature parasite-infected erythrocyte surface antigen, protein 2 (MESA) | 168.2 | 6.9 | 6 |
| PF3D7_1406800    | glideosome-associated protein with multiple membrane spans 3 (GAPM3) | 32.9 | 6.7 | 2 |
| PF3D7_0912400    | alkaline phosphatase | 52.7 | 5.8 | 2 |
| PF3D7_0204700    | hexose transporter (HT) | 56.4 | 5.4 | 2 |
| PF3D7_0824400    | nucleoside transporter 2 (NT2) | 67.6 | 3.8 | 2 |
| PF3D7_1420200    | tetratricopeptide repeat family protein, putative | 147.9 | 1.9 | 2 |
Figure 1. Expression of GFP-tagged MyoA through the *P. berghei* life cycle. (A) Transcription of *myoa* analyzed by qRT-PCR normalised against two control genes, *hsp70* and *arginine-tRNA synthetase* at different stages of the life cycle. The mean normalized *myoa* expression is shown as a horizontal bar and standard deviations are indicated by error bars. AS: asynchronous asexual blood stages, Sch: schizonts, NAG: non-activated gametocytes, AG: activated gametocytes, Ook: ookinetes, and Spor: sporozoites. (B) A plasmid containing the sequence for the 3’ end of the MyoA coding region fused in frame to the *gfp* sequence, together with a human *dhfr* selectable marker, was used to insert the *gfp* sequence into the endogenous *myoa* gene. (C) Successful integration into the *P. berghei* genome was confirmed by diagnostic PCR on genomic DNA prepared from wild type parasites and those with integrated plasmid sequence using primers 9 and 10, which amplify a 1.1 kb DNA fragment from the modified *myoa* locus following correct integration. (D) Expression of the GFP-tagged MyoA was confirmed by western blotting of lysates from parasites expressing either the unfused GFP or the MyoA-GFP proteins; as indicated GFP-specific antibodies reacted with either a 29 kDa or a 120 kDa protein, respectively. (E) Extracts of schizonts following hypotonic lysis in the soluble fraction (S) and further fractionation of the insoluble material by carbonate buffer into carbonate-soluble (PM, peripheral membrane) and insoluble (IM, integral membrane) fractions. Schizonts from parasites expressing either GFP (upper row) or MyoA-GFP (lower row) were used. (F) Expression of MyoA-GFP at the invasive and motile stages of the parasite life cycle, detected by live fluorescence microscopy. The parasite nuclei are stained with Hoechst 33342 and the ookinete surface with Cy3-conjugated anti-P28 antibody 13.1. In the merged colour image these are blue and red respectively, and the MyoA-GFP is green. The differential interference contrast (DIC) bright field images are also shown. Scale bar: 5 µm. (G) Expression of MyoA-GFP in liver stage parasites at the early and late cytomere stages; GFP fluorescence (green) and Hoechst 33342 staining (blue). Scale bar: 10 µm. (H) Temporal profile of MyoA-GFP expression during the six stages of *Plasmodium berghei* ookinete development. MyoA-GFP was detected by live fluorescence microscopy associated with the protuberance that grows out of the spherical body only at stage III and eventually forming the motile ookinete. The parasites were co-stained with Hoechst 33342 and with Cy 3-conjugated anti-P28 antibody 13.1 as a marker for the zygote and ookinete surface. The merged (MyoA-GFP:green; P28: red; Hoechst: blue) and differential interference contrast (DIC) bright field images are also shown. Scale bar: 5 µm.

Figure 2. MyoA-GFP expression and glideosome complex formation during the late stages of intracellular development of *P. falciparum* in the red blood cell. (A) MyoA-GFP expression was detected by live fluorescence microscopy and the nuclei were detected by staining with DAPI. The merged colour image with MyoA-GFP (green) and DAPI (blue) and differential interference contrast (DIC) bright field images are also shown. Schizogony starts at around 30 hours post-invasion and MyoA-GFP is detected from 38–40 hours in multinucleated forms. Scale bar: 2 µm. (B) Parasites expressing MyoA-GFP were collected at indicated time points post-invasion and lysates prepared; samples of these lysates (upper panel) and of proteins precipitated from them with a GFP-specific antibody (lower panel) were fractionated by SDS-PAGE and probed on western blots with antibodies to GFP, MTIP, GAP45 and GAP50. Molecular weight markers are indicated in kDa.

Figure 3. The location of MyoA during *P. falciparum* merozoite invasion. (A) MyoA-GFP is located at the periphery of developing intracellular (top row of each pair of images) and free extracellular merozoites (bottom row), and is colocalized with antibodies specific for IMC proteins MTIP, GAP45 and GAP50. In the merged colour image the MyoA-GFP signal is green and antibodies specific for the IMC proteins are red; nuclei are stained blue with DAPI. The DIC image is also shown. (B) Individual merozoites are captured at different stages of invasion from initial attachment, through early and late invasion to the intracellular ring stage. MyoA-GFP remains peripheral whereas RON4, initially in the apical rhoptry neck, relocates during invasion. Merged colour images with MyoA-GFP (green), RON4 (red), and nuclei (blue) and DIC images are also shown, together with a schematic of each cell-pair. Scale bar: 2 µm.

Figure 4. Identification of GAP40 in the *P. falciparum* glideosome complex. (A) Wild-type (lane 1) or GAP45-GFP expressing (lane 2) *P. falciparum* 3D7 parasites were incubated with radiolabelled
phosphate and proteins binding to a GFP-specific antibody were immunoprecipitated, resolved on a SDS-PAGE gel and detected by autoradiography. Bands corresponding to two unidentified proteins are labelled with one or two asterisks. In parallel the same samples were probed with antibodies to known components of the glideosome complex: MyoA, GAP45, GAP50 and MTIP. (B) The immunoprecipitation with anti-GFP antibody resin was repeated with unlabelled schizont lysate from wild-type (lane 1) or GAP45-GFP expressing (lane 2) *P. falciparum* 3D7 parasites. Immunoprecipitated proteins were fractionated by SDS-PAGE, detected with SYPRO Ruby and the indicated bands were excised. (C) Proteins identified in the three excised bands by tryptic digestion and mass spectrometry. The number of unique peptides identified for each protein is indicated, along with the percentage of the protein sequence that these peptides cover.

**Figure 5.** The putative ELC binds to MyoA but not MyoB and is in the *P. falciparum* glideosome complex together with MTIP. Alignment of (A) *Plasmodium* ELC homologues, and (B) PfELC with the two *T. gondii* MyoA ELCs. Identical residues are shaded black, biochemically similar residues are shaded grey. Percentage identities to PfELC are displayed. Gene identifiers are as follows: PfELC: PF3D7_1017500; PrELC: *P. reichenowi* PRCDC_1016900; PvELC: *P. vivax* PVX_001745; PcyELC: *P. cynomolgi* PCYB_061180; PeELC: *P. knowlesi* PKNH_0601700; PchELC: *P. chabaudi* PCHAS_0501900; PyELC: *P. yoelii* PY02639; PbgELC: *P. berghei* PBANKA_0501800; TgELC1: *T. gondii* TGME49_069440; TgELC2: *T. gondii* TGME49_305050. (C) Structural homology model of ELC generated using Phyre 2. (D) Proteins in extracts from wild type (3D7) schizonts and those expressing either MyoA-GFP or MyoB-GFP were immunoprecipitated with resin-bound antibodies to GFP, and then probed by western blotting with antibodies to GFP and ELC. I: input lysate, U: unbound protein, E: protein eluted from the resin. (E) The protein complex containing MTIP also contains ELC. The glideosome complex from lysates of schizonts (S) and merozoites (M) of 3D7 was precipitated with anti-MTIP antibodies, resolved by SDS-PAGE and the presence of ELC was determined by western blot with specific antibodies. A pre-immune antibody sample was used to immunoprecipitate from the lysates as a control.

**Figure 6.** The putative MyoA ELC binds to the neck region of MyoA. (A) Purification of recombinant His-PfELC protein. (Bi) Far- and (ii) near-UV circular dichroism spectra of recombinant protein indicate that it is highly structured, with 32% alpha helix, 17% beta sheet, 20% turn. (C) PfELC binding to immobilised peptides corresponding to residues 770–787, 786–803 and 801–818 of MyoA was measured by biolayer interferometry, using increasing concentrations of PfELC. The dissociation constant (Kd) with standard error was calculated from each of the curves. (D) The amino acid sequence of the neck region of MyoA, showing the location of the peptides used for analysis, the MTIP binding site, and the proposed ELC binding site.

**Figure 7.** PfELC expression matches that of known glideosome components in *P. falciparum* schizonts. (A) Scheme for modifying the PF3D7_1017500 gene locus by single crossover homologous recombination in order to C-terminally modify the ELC protein with a triple-HA tag. (B) PCR screening of parental 3D7 parasites and two independently generated clones. PCR product from the wild-type locus was amplified using primers 13 and 14 (WT), while a product (I) from the modified locus was amplified using primers 13 and 15. (C) Western blot analysis of HA-tagged ELC, MTIP and GAP45 proteins present during a time course of schizont development. (D) Immunofluorescent detection of HA-tagged ELC and either MTIP or GAP45 in mature schizonts and merozoites. Merged colour images are also shown, with ELC-HA (green) and MTIP or GAP45 (red). Nuclei are stained with DAPI (blue). Scale bar: 2 µm.

**Figure 8.** Recombinant MyoA binds actin. (A) Coomassie-stained SDS-PAGE showing affinity purified GST-MyoA, marked with an asterisk. (B) An actomyosin event as determined using optical tweezers, with an actin filament held between 2 beads: a left-hand bead (blue) and a right-hand bead (red); with the MyoA molecule on a third bead. Because of the high-sensitivity required in these single molecule measurements, Brownian motion causes a significant amount of displacement noise and when MyoA binds to actin that event is evident by a sudden decrease in the standard deviation of the displacement noise (shown in black). Events are scored by thresholding data falling below a fixed
value (dotted red/blue line). Note: movement of the left-hand bead and its standard deviation are offset by -200 nm and -30 nm, respectively, for clarity. (C) Improved visualisation of binding events is achieved by moving the beads back-and-forth using a 10 Hz triangular wave function of 100 nm amplitude. (D) Displacement histogram generated from 244 acto-MyoA binding events with a fitted Gaussian distribution. The magnitude of the displacement (nm) is displayed on the x-axis and the number of observations at that value (N_{obs}) on the y-axis. The average movement caused by the myosin “power-stroke” was measured as 3 nm. (E) Acto-MyoA event lifetime distributions measured at two Mg.ATP concentrations, 8 µM (blue symbols) and 0.3 µM (red symbols), confirm that the event lifetimes are strongly dependent on [Mg.ATP]. The x-axis shows the duration of the binding event (in seconds); the y-axis, is the number of observations (N_{obs}) at that value. The lines are single exponential least-squares fits with rate constants 53 s^{-1} (blue) and 6.9 s^{-1} (red).

**Figure 9.** MTIP/ELC decorated MyoA shows increased in vitro sliding motility. (A) Size exclusion chromatography of co-expressed MyoA and MTIP (solid line) and MyoA, MTIP and ELC (dotted line) (B) Coomassie-stained SDS-PAGE of the purified MyoA-light chain complexes. (C) Motility of nitrocellulose-captured MyoA complexes, and (D) the corresponding capture of MyoA complexes via an anti His-tag antibody measuring the velocity of actin filament movement in the presence of ATP. A solid line represents a Gaussian fit to each corresponding histogram. (E) The actin-activated ATPase of MyoA/MTIP and MyoA/MTIP/PfELC was estimated using an NADH-coupled assay to quantify the rate of ATP hydrolysis.

**Supplemental movie.** The four panels show the movement of fluorescent actin filaments by MyoA/MTIP bound by nitrocellulose (top left panel) or by anti-His antibody (lower left panel), and MyoA/MTIP/ELC bound by nitrocellulose (top right panel) or by anti-His antibody (lower right panel).
Figure 1

A

B

C

D

E

F

G

H

MyoA-GFP
P28
Hoechst
MyoAP28
DIC

S
PM
IM

GFP
MyoA-GFP

MyoA-GFP
Hoechst
MyoA/Hoechst

Early cytoplasmic stage
Late cytoplasmic stage
Figure 2
Figure 3
### Figure 4

#### A

[Image of a gel with bands labeled as MycA, GFP-GAP45, GAP50, and MTIP.]

#### B

[Image of a gel with bands labeled as GFP-GAP45 and GAP50.]

#### C

| Protein identification | Number of unique peptides | Coverage (%) |
|------------------------|---------------------------|--------------|
| PF3D7_0515700 (GAP40)  | 2                         | 3.3          |
| PF3D7_1222700 (GAP45)  | 13                        | 38.2         |
| P42212 (GFP)           | 6                         | 23.5         |
| PF3D7_0918000 (GAP50)  | 3                         | 8.3          |
Figure 5
Figure 6

A  

B i)

ii)

Wavelength, nm

C i)

ii)

Response, nm

MeyA 770–787, µM

0 1 2 3 4 5

MeyA 786–803, µM

0 1 2 3 4 5

MeyA 801–818, µM

0 1 2 3 4 5

D  

770–787 QREKLVWENCVSVDAA  
786–803 AAILKHKYRQQVKNIPS  
801–818 ISSLLRVQAHIRKHMKVAQ

MyoA neck QEGKILTKI QREKLVWENCVSVDAA ILEHKKYRQQVKNIPS LLLRVQAHIRKHMKVAQ

Lever Arm  

ELC binding  

MTIP binding
Figure 7
Figure 8
Figure 9

A

B

C

D

E

Retention (min)

100

0

0.2

0.4

0.6

0.8

Velocity / \mu \text{m s}^{-1}

0

20

40

60

80

100

120

140

160

ATPase activity, n

[Actin], \mu \text{M}

MyoAMTP

MyoAMTP/ELC