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Abstract

Obligate intracellular parasites of the phylum Apicomplexa exhibit gliding motility, a unique form of substrate-dependent locomotion essential for host cell invasion and shown to involve the parasite actin cytoskeleton and myosin motor(s). Toxoplasma gondii has been shown to express three class XIV myosins, TgM-A, -B, and -C. We identified an additional such myosin, TgM-D, and completed the sequences of a related Plasmodium falciparum myosin, PfM-A. Despite divergent structural features, TgM-A purified from parasites bound actin in an ATP-dependent manner. Isoform-specific antibodies revealed that TgM-A and recombinant mycTgM-A were localized right beneath the plasma membrane, and subcellular fractionation indicated a tight membrane association. Recombinant TgM-D also had a peripheral although not as sharply defined localization. Truncation of their respective tail domains abolished peripheral localization and tight membrane association. Conversely, fusion of the tails to green fluorescent protein (GFP) was sufficient to confer plasma membrane localization and sedimentability. The peripheral localization of TgM-A and of [...]
A Dibasic Motif in the Tail of a Class XIV
Apicomplexan Myosin Is an Essential Determinant of
Plasma Membrane Localization

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Isoform-specific antibodies revealed that TgM-A and recombinant mycTgM-A were localized
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sufficient to confer plasma membrane localization and sedimentability. The peripheral localization
of TgM-A and of the GFP-tail fusion did not depend on an intact F-actin cytoskeleton, and the GFP
chimera did not localize to the plasma membrane of HeLa cells. Finally, we showed that the
specific localization determinants were in the very C terminus of the TgM-A tail, and site-directed
mutagenesis revealed two essential arginine residues. We discuss the evidence for a proteinaceous
plasma membrane receptor and the implications for the invasion process.

INTRODUCTION

The phylum of Apicomplexa comprises numerous patho-
gens of medical and veterinary significance. Among them, Plasmodium falciparum is the most virulent of the Plasmodium
species responsible for malaria in human. The opportunistic
pathogen Toxoplasma gondii causes diseases in immunocom-
promised patients and in congenitally infected infants. Al-
though members of the Apicomplexa differ significantly in
their host range and cell type specificity, they do share a
similar mechanism for host cell entry. Host cell penetration
is a prerequisite for the survival of these obligate intracellu-
lar parasites and does not occur by induced phagocytosis
but via an active process from the parasites (Sibley, 1995). In
the absence of locomotive organelles, apicomplexan para-
sites have developed an unusual mode of substrate-depen-
dent gliding locomotion that is necessary for invasion (Sib-
aley et al., 1998). The inhibitory functions of cytochalasins on
invasion by Plasmodium (Miller et al., 1979) and Eimeria (Rus-
sell, 1983) have been reported and argue for a participation
of actin filaments. More recently, a direct involvement of T. gondii actin has been demonstrated, establishing that motil-
ity is essential for invasion and depends on the parasite’s
own cytoskeleton (Dobrowolski and Sibley, 1996). An actin-
based motor is predicted to generate the force during inva-
sion by Apicomplexa, and this hypothesis is corroborated by
the inability of parasites to glide and to invade in the pres-
ence of the inhibitor of myosin heavy chain ATPase butane-
dione monoxime (Dobrowolski et al., 1997a; Pinder et al.,
1998). Similarly, tachyzoites treated with jasplakinolide, a
membrane-permeable actin-polymerizing and filament-sta-
bilizing drug, reversibly inhibit host cell invasion (Shaw and
Tilney, 1999). Members of the myosin superfamily are
mechanoenzymes that convert chemical energy stored in
ATP into a directed force along actin filaments (Spudich,
1994). In the past few years evidence has emerged for roles
played by actin-based molecular motors in a wide range of

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membrane movements (Hasson and Mooseker, 1995; Mermall et al., 1998). In the case of host cell invasion by apicomplexan parasites, a capping model proposes that gliding motility shall be driven by the redistribution of transmembrane proteins from the apical to the posterior pole of the parasite along the subcortical actomyosin system. Correspondingly, the molecular motor powering gliding locomotion and capping is expected to localize right beneath the plasma membrane. Recently, three myosins were identified in T. gondii (Heintzelman and Schwartzman, 1997) and shown to build a 14th phylogenetic and structural class of myosins (Mermall et al., 1998). Antibodies cross-reacting with heterologous myosins were used to immunolocalize a myosin at the anterior pole of T. gondii tachyzoites (Schwartzman and Pfefferkorn, 1983), and more recent work from the same group potentially identified this myosin as TgM-A (Heintzelman and Schwartzman, 1999). In addition, antibodies raised against the conserved peptide LEAF revealed one or more myosins localized beneath the plasma membrane and also scattered throughout the cytosol of T. gondii tachyzoites (Dobrowolski et al., 1997a). Similarly, antibodies generated against a small fragment of the motor domain of Pf-myol showed labeling concentrated under the plasma membrane of P. falciparum merozoite but absent from the apical prominence itself (Webb et al., 1996). The P. falciparum myosin recognized by these antibodies migrates slightly above 100 kDa in SDS-PAGE, is expressed in mature schizonts and merozoites, and localizes predominantly around the periphery of the cell (Pinder et al., 1998).

Here, we report of the identification of a 91-kDa T. gondii myosin of class XIV, TgM-D, and of the completion of the PfM-A sequence, the likely P. falciparum homologue of TgM-A. The exquisite amenability of T. gondii to molecular genetics allowed us to investigate the determinants of subcellular localization of two class XIV myosins. A pair of basic residues is essential to target TgM-A to the periphery, defining its cargo-binding site down to the amino acid level. Such precise mapping has only been achieved for two other myosins, NINAC, a Drosophila myosin III interacting with INAD (Wes et al., 1999), and Myo2, a Saccharomyces cerevisiae myosin V binding to the vacuole (Cafleit and Weisman, 1998). In addition, the first purification of such a myosin is presented, allowing us to show that, despite its structural divergences, TgM-A had the ability to bind F-actin in an ATP-dependent manner. Evidence is presented that the peripheral localization is largely independent of the actin cytoskeleton and is likely due to specific and saturable interaction with a proteinaceous component of the plasma membrane.

**MATERIALS AND METHODS**

**Strains and Reagents**

The bacterial strains for recombinant DNA techniques were Escherichia coli XL1-Blue and XLOR. The helper phage ExAssist, from Stratagene (La Jolla, CA), was used for the in vivo excision of the phagemid vectors from the AZAPII clones. Restriction enzymes were purchased from New England Biolabs (Beverly, MA). The secondary antibodies for Western blotting were from Bio-Rad (Hercules, CA), and those for immunofluorescence were from BioTrend (Cologne, Germany).

**Growth of Parasites and Isolation of DNA and RNA**

T. gondii tachyzoites (RH strain wild-type and RHinxgfrt) were grown in human foreskin fibroblasts (HFFs) or in vero cells (African green monkey kidney cells) maintained in Dulbecco’s modified Eagle’s medium supplemented with 10% FCS, 2 mM glutamine, and 25 μg/ml gentamicin. Parasites were harvested after complete lysis of the host cells and purified by passage through 3.0-μm filters and centrifugation in PBS. Genomic DNA was isolated from purified parasites by SDS-proteinase K lysis followed by phenol-chloroform, chloroform extractions and ethanol precipitation (Sibley and Boothroyd, 1992). Total RNAs were prepared using RNA Clean from AGS (Heidelberg, Germany) according to the manufacturer’s instructions. P. falciparum total RNA was purified from the isolate FCBR, Columbia grown in vitro (Knapp et al., 1989) and kindly provided by Dr. H. del Portillo (University of Sao Paulo, Sao Paulo, Brazil).

**PCR Screening**

PCR was performed on T. gondii genomic DNA and yielded two distinct myosin fragments. PCR reactions were carried out in a GeneAmp PCR System 2400 instrument (Perkin-Elmer, Norwalk, CT). Primers myol and myo5 (corresponding to the conserved motifs GESGAGKT and LEAFNAKT, respectively) and the parameters used for the PCR reaction are described elsewhere (Schwarz et al., 1999). The PCR products were purified over a spin column (QIAquick PCR purification kit; Qiagen, Hilden, Germany) and the whole sample was cloned into PCR Script (Stratagene) as described by the manufacturer. Double-stranded DNA sequencing of the products was performed by the dideoxy-termination method using Sequenase version 2.0 (United States Biochemical, Cleveland, OH) with T3 and 17 primers.

**Toxoplasma cDNA and Genomic Library Screening**

The digoxigenin system (Boehringer Mannheim, Mannheim, Germany) for nonradioactive labeling and detection of nucleic acids was used to screen a T. gondii cDNA library. The 300-bp myosin fragment was digoxigenin-dUTP labeled by PCR and used as a probe to screen the RH (Elmer Pfefferkorn) cDNA library in AZAPII from National Institutes of Health AIDS reagents (kindly provided by D.S. Roos, University of Pennsylvania, Philadelphia, PA). The detection of positive clones was achieved by chemiluminescence with CSPD (Boehringer Mannheim) according to the manufacturer. Plaques from 15 X 10⁴ phage were screened, and positive clones were identified on both duplicates. After two additional cycles of hybridization, positive clones were excised in vivo, and the insert sizes were determined by restriction digests. One clone contained a full-length cDNA corresponding to the previously described TgM-A and predicting a protein of 93 kDa (Heintzelman and Schwartzman, 1997). The second clone encoded another myosin lacking a start codon (TgM-D). Further screening of cDNA libraries failed to identify full-length TgM-D cDNA clones. The missing 5’ sequences of TgM-D were finally obtained by sequencing genomic cosmid clones encompassing the TgM-D locus. The cosmid library used the SuperCos vector modified with an SAG1/ble Toxoplasma selection cassette inserted into its HindIII site. The library was prepared from a SacAI partial digestion of RH genomic DNA ligated into the BamHI cloning site and was kindly provided by D. Sibley and D. Howe (Washington University, St. Louis, MO). The gene coding for TgM-D was interrupted by 11 introns of various sizes (our unpublished results). Southern blotting analysis of T. gondii genomic DNA digested with several restriction enzymes yielded patterns of bands matching those predicted by the restriction map of the cosmid clone, indicating that TgM-D is a single-copy gene.

We also obtained and sequenced a cosmid clone encompassing the TgM-B/C locus (GenBank accession number AF202535). This confirmed that the two proteins, which have a common N terminus
and differ only in their C terminus, are really translated from two transcripts with alternatively spliced 3' ends (as suggested by Heinzelman and Schwartzman, 1997). This also reveals that PCR using the reported common 5' termini were cDNA library cloning artifacts, resulting from an in frame fusion at an EcoRI site with the 3' end of GRA7. The correct cDNA could then be amplified by reverse transcription (RT-PCR), cloned, and sequenced.

Identification of a Full-Length P. falciparum Myosin

The sequences of the short and conserved, positively charged tail domains of TgM-A and TgM-D showed a significant homology with an expressed sequence tag derived from the closely related organism *P. falciparum* (BLAST software search). An RT-PCR strategy using an antisense primer corresponding to the tail and a sense primer corresponding to a conserved motif in the head domain allowed us to amplify, clone, and sequence a large fragment of the myosin. The sequence corresponded to the partial sequence of a *P. falciparum* myosin present in the database (Ptmpro-1). The progress on the genome sequencing project allowed us to complete and to confirm the sequence of this myosin (PJM-A, accession number AF105118). The PJM-A gene contains only two introns clustered immediately downstream of the AUG, suggesting a possible regulatory function of 5'-splice-site splicing. This also revealed that the sequences of the short and conserved, positively charged tail transcripts with alternatively spliced 3' ends (as suggested by Donald et al., 1997) and differ only in their C terminus, are really translated from two transcripts with alternatively spliced 3' ends (as suggested by Heinzelman and Schwartzman, 1997). This also revealed that PCR using the reported common 5' termini were cDNA library cloning artifacts, resulting from an in frame fusion at an EcoRI site with the 3' end of GRA7. The correct cDNA could then be amplified by reverse transcription (RT-PCR), cloned, and sequenced.

Construction of *T. gondii* Expression Vectors

TgM-A and TgM-D were amplified by PCR to introduce a SacI site before the stop codon, and a BamHI site before the stop codon, using the sense oligonucleotide 5'-gacctgcggacaaagggcatcctg-3' combined with reverse oligonucleotide primer 5'-ggatcggagacgactcctg-3' for TgM-A and 5'-gacgacggacacgactcctg-3' combined with 5'-ggatcggacaaagttgctgctc-3', for TgM-D. The PCR products were cloned into bacterial expression vectors (pET-TgM-A and pET-TgM-D) and analyzed for expression of the recombinant protein in BL21 (DE3, expressing T7 polymerase) after isopropyl-1-thio-β-D-galactopyranoside induction. The deduced amino acid sequence of TgM-A predicts the presence of 9 Da for TgM-A and 91 kDa for TgM-D were produced (our unpublished results). Vectors used to express TgM-A and TgM-D in *T. gondii* were derived from the pS214SCAT vector containing 5' and 3' flanking sequences of the SAG1 gene and pTSRCAT containing 5' flanking sequence of the TUB1 gene described previously (Soldati and Boothroyd, 1995). pS'TgM-A was generated by subcloning TgM-A from pET-TgM-A into pS'/4R between NsiI and PacI sites (Soldati and Boothroyd, 1995). The vector p'TgM-D was obtained by cloning TgM-D coding sequence into NsiI and PacI sites of pT/R230 (Soldati and Boothroyd, 1995). In these constructions, both myosins are fused at the N terminus to a c-myc epitope tag and seven His residues (these sequences were imported from a modified myosin vector). The mycTgM-A and mycTgM-D proteins. The construct p'TgM-A contains the mycTgM-A except that sequences corresponding to the last 53 amino acids of TgM-A are lacking. This deletion is fluorescent as nonfusion protein in tachyzoites (M. Soete, C. Heitmam, and D. Soldati, unpublished data) and has been extensively modified in the codon usage (Haseloff et al., 1997). The constructs p'TGFP-TgM-Atail, p'TGFP-TgM-Dtail, and p'TGFP-PM-Atail were generated by cloning the PCR products corresponding to the last 82 amino acids of TgM-A (as in Figures 7 and 9, or 62, as in Figures 1A, 5, and 8) and the last 57 amino acids of TgM-D. This also revealed that the sequences of the short and conserved, positively charged tail are really translated from two transcripts with alternatively spliced 3' ends (as suggested by Donald et al., 1997) and differ only in their C terminus, are really translated from two transcripts with alternatively spliced 3' ends (as suggested by Heinzelman and Schwartzman, 1997). This also revealed that PCR using the reported common 5' termini were cDNA library cloning artifacts, resulting from an in frame fusion at an EcoRI site with the 3' end of GRA7. The correct cDNA could then be amplified by reverse transcription (RT-PCR), cloned, and sequenced.

Generation of a Serum Specific for the Tail of TgM-A

Two peptides covering the tail of TgM-A ([C]VLEAYYAGRRH) and ([C]VLEAYYAGRRH) were synthesized and coupled to keyhole limpet hemocyanin according to the instructions of the manufacturer (Pierce, Rockford, IL). Two rabbits were first immunized with 400 μg of both peptides with the complete Freund's adjuvant. Four successive subcutaneous injections with 200 μg of both peptides in incomplete adjuvant LQ (Gerbus, Heidelberg, Germany) were performed at 24-day intervals. The serum was immunoaffinity purified against the peptides previously coupled to Affi-Gel 10 according to the manufacturer (Bio-Rad). Four milligrams of each peptide were used to affinity purify 2 ml of sera.

Parasite and HeLa Cell Transfection and Selection of Transformants

*T. gondii* tachyzoites (RHhxgprt1) were transfected by electroporation as previously described. Selection of stable transformants expressing pS'TgM-A and p'TGFP-TgM-Atail were achieved by cotransfection of the expression vector with the selectable plasmid (pminHXGPRT). Restriction enzyme-mediated integration was applied as previously described (Black et al., 1995). The hypoxanthine-xanthine-guanine-phosphoribosyltransferase (HXGPRT) was used as a positive selectable marker gene in the presence of mycophenolic acid and xanthine as described (Donald et al., 1996). Freshly released parasites (5 x 10^7) of the RHhxgprt1 strain were resuspended in cytomix buffer in the presence of 100 U of BanHI and were cotransfected with 10–20 μg of the plasmid pminHXGPRT, which carries the HXGPRT gene, and 80–100 μg of the plasmids expressing GFP fusions or myosins. Parasites expressing p'TgM-D, p'TGFP-PM-A, and p'TGFP-PG-Atail could not be cotransfected; therefore, the selectable marker gene dhfrtsXHGPRTdhfrts was introduced into the expression vector in a unique SacI site before selection. HeLa cells were transfected with pUHD15-1GFP-TgM-M.
Actin and a calcium phosphate method as described previously (Graham and Eb, 1973).

**Cytochalasin D Treatment**

Cytochalasin D (Sigma, St. Louis, MO) was stored as DMSO stock at −20°C. Intracellular parasites grown for 24 h in HFFs on glass coverslips were incubated with 10 µg/ml cytochalasin D in medium at 37°C for up to 3 h. At the indicated time, the coverslips were taken out and rinsed twice with PBS, and cells were fixed as described below.

**Indirect Immunofluorescence Microscopy and Detection of GFP in T. gondii**

All manipulations were carried out at room temperature. Intracellular parasites grown for 24 h in HFFs on glass slides were fixed with 4% paraformaldehyde and 0.05% glutaraldehyde for 20 min. After fixation, slides were rinsed in PBS and 0.1 M glycine. Cells were then permeabilized in PBS and 0.2% Triton X-100 for 20 min and blocked in the same buffer with 2% FCS. Slides were incubated for 60 min with primary antibodies diluted in PBS and 1% FCS, washed, and incubated for 60 min with Alexa488- or FITC-labeled goat anti-mouse immunoglobulin Gs diluted in PBS and 1% FCS, washed, and incubated for 60 min with Alexa488- or FITC-labeled goat anti-mouse immunoglobulin Gs diluted in PBS and 1% FCS. After cytochalasin D treatment, stainings were carried out with the following modifications. Fixation was with 4% paraformaldehyde only; all subsequent buffers contained 5 mM EGTA to chelate Ca2+. Staining of F-actin was performed with Oregon Green-phalloidin (Molecular Probes, Eugene, OR) for 60 min. The rapid freezing combined with fixation and permeabilization in ultracold methanol was performed as described (Neuhaus et al., 1998). Slides were mounted in Vectashield (Vector Laboratories, Burlingame, CA) and kept at 4°C in the dark. The mAb anti-myc was an ascites preparation of 9E10 used at the dilution 1:1000. The mAb DG52, recognizing SAG1, coupled to biotin was generously provided by Dr. J.F. Dubremetz (Institut Pasteur de Lille). Intracellular parasites expressing GFP were fixed according to the above protocol and mounted immediately for microscopic analysis. Confocal images were collected with a Leica (Nussloch, Germany) laser scanning confocal microscope (TCS-NT DM/IRB) using a 100× Plan-Apo objective, numerical aperture 1.30. Single optical sections were recorded with an optimal pinhole of 1.0 (according to Leica’s instructions) and 16 times averaging. All other micrographs were obtained with a Zeiss (Thornwood, NY) Axioptih microscope with a camera (CH-250; Photometrics, Tucson, AZ). Adobe Photoshop (Adobe Systems, Mountain View, CA) was used for image processing.

**Western Analysis of Parasite Lysates**

SDS-PAGE was performed using standard methods (Leahmili, 1970). Crude extracts from T. gondii tachyzoites were separated by SDS-PAGE and transferred to nitrocellulose. Western blot analysis was carried out essentially as described previously (Soldati et al., 1998), using 8–12% polyacrylamide gels run under reducing condition with 144 mM β-mercaptoethanol or 100 mM DTT in the loading samples. After electrophoresis, proteins were transferred to Hybond ECL nitrocellulose (Amersham, Arlington Heights, IL). For detection, the membranes were incubated with the mAb 9E10 (mouse ascites fluid diluted 1:1000 in PBS and 0.5% Tween 20) and then with the affinity-purified horseradish peroxidase-conjugated goat anti-mouse immunoglobulin G (1:2000), and bound antibodies were visualized using the ECL system (Amersham). Direct recording of chemoluminescent signals and densitometry by an LAS-1000 luminescent image analyser (Fuji, Tokyo, Japan) allowed quantification of signal intensities within a broad linear range.

**Cell Fractionation**

Freshly released parasites (108) from infected cells were resuspended into 300 µl of buffer and lysed by freezing and thawing three times followed by sonication (four times for 30 s on ice). Pellet and soluble fractions were separated by ultracentrifugation 1 h at 55,000 rpm at 4°C. The buffers were PBS, PBS and 1 M NaCl, PBS and 0.1 M Na2CO3, pH 11.5, PBS and 2% Triton X-100, and 3 M urea. In addition, all buffers contained 5 mM ATP to minimize the interaction of myosins with F-actin, and 5 mM EDTA.

**Purification of Recombinant Myosin from T. gondii**

Freshly lysed transgenic parasites expressing TgM-Δtail (1010 parasites) were lysed under native conditions by freezing and thawing followed by sonication. The soluble fraction was separated by ultracentrifugation at 65,000 rpm, and the supernatant was applied to an Ni-nitrioltriacetic acid resin column and eluted according to the manufacturer (The QIAexpressionist; Janknecht et al., 1991); determination of protein concentration was performed by a Bradford assay (Bio-Rad).

**Actin Binding Assays**

The binding of TgM-Δtail to F-actin was measured using an F-actin sedimentation assay as described previously (Jung and Hammer, 1994). F-actin from rabbit skeletal muscle was purified according to the method of Pardee and Spudich (1982). Binding assays were performed using a 4 μM final concentration of F-actin and 0.5 μg of TgM-Δtail (corresponding to ~0.06 μM). The buffer (plus ATP) contained 10 mM Tris-HCl, pH 7.5, 130 mM KCl, 2 mM MgCl2, 2 mM ATP, and 0.1 mM DTT. The washing steps were performed by resuspending the pellet in washing buffer and were followed by centrifugation at 50,000 rpm. Pellets and supernatants were analyzed by SDS-PAGE and Coomassie blue staining.

**RESULTS**

**Identification and Cloning of Two Members of the Class XIV Myosins**

To identify myosins in T. gondii, we designed a PCR screening strategy taking advantage of the highly conserved motifs present in the head domain (Schwarz et al., 1999). The approach identified TgM-A, a myosin already described by Heintzelman and Schwartzman (1997), and TgM-D, an additional member of the apicomplexan-specific class XIV of
myosins (Figure 1B). The entire TgM-D gene was sequenced. The protein predicted from the open reading frame has 823 amino acid residues (Figure 1A) with a mass of 91 kDa (accession number AF105117).

The sequences of the short and conserved, positively charged tail domains of TgM-A and TgM-D showed a significant homology with an expressed sequence tag derived from the closely related organism P. falciparum. This allowed us to identify and subsequently clone a complete myosin corresponding to a partial P. falciparum sequence present in the database (PfMyo-1; see MATERIALS AND METHODS). We named this myosin PfM-A (Figure 1A) because its predicted sequence of 90 kDa has highest homology to TgM-A (Figure 1B). The degree of conservation at the amino acid level between the myosins is high and extends throughout the entire coding region. There is 55.7% identity between TgM-A and TgM-D, 63.1% between TgM-A and PfM-A, and 52.0% between TgM-D and PfM-A. Pairwise comparisons between any of these myosins with the other two, alternatively spliced, class XIV myosins from T. gondii, TgM-B and -C, give lower identity scores of ~47–49%.

Like the other apicomplexan myosins TgM-D and PfM-A, have a very short tail, do not contain the strictly conserved glycine residue at the proposed fulcrum point of the lever arm, and appear to lack conserved “IQ” motifs that bind calmodulin and calmodulin-related proteins. Among their particularities, and contrary to the P. falciparum molecule, all the T. gondii myosins do not follow the TEDS rule (Heintzelman and Schwartzman, 1997; Figure 1A), which describes the presence of an acidic or phosphorylatable residue at a precise position close to the actin-binding region (originally mapped by Brzeska et al., 1989, 1990; for review, see Bement and Mooseker, 1995). In lower eukaryotes, phosphorylation of this conserved serine or threonine was shown to be crucial for the stimulation of the ATPase activity of class I myosins (Bement and Mooseker, 1995; Carragher et al., 1998; Novak and Titus, 1998). It is unclear at the moment whether and how these class XIV motors are activated and how conformational changes in the molecules occur.

The different recombinant constructs of TgM-A and TgM-D used in this study are presented schematically in Figure 1C.

**Antibodies to the Tail of TgM-A Are Isoform Specific**

To investigate the expression and distribution of TgM-A, we raised polyclonal sera against two peptides covering its entire tail. The specificity of the antibodies was assessed by Western blotting of parasites expressing GFP-TgM-Atail and GFP-TgM-Dtail (Figure 2A). The anti-myc antibodies detected both GFP chimeras, but despite high overall sequence homologies, the anti-tail antibodies recognized only the TgM-A tail (Figure 2A). In wild-type parasites, the affinity-purified antibody recognized a single band migrating above 90 kDa, the predicted size of TgM-A (Figure 2B). Recombinant mycTgM-A was detected by the anti-tail antibodies and was slightly bigger than endogenous TgM-A. Confirming the antibody specificity, only the endogenous protein was detectable in parasites expressing mycTgM-A
tail (Figure 2B).

**The Expression of Endogenous TgM-A Is Down-regulated in the Transgenic Parasites Expressing Full mycTgM-A**

Surprisingly, in recombinant parasites expressing full-length mycTgM-A, the endogenous TgM-A was barely detectable (Figure 2B). To obtain a quantitative impression of the phenomenon, equal numbers of wild-type and different recombinant parasites were analyzed (Figure 2B). MycTgM-A appeared to be expressed at approximately two- to threefold the level of endogenous protein found in wild-type parasites and parasites expressing mycTgM-A
tail. We conclude that expression of full-length mycTgM-A, but not of the head domain alone, led to down-regulation of the endogenous TgM-A protein and expression. In parasites expressing GFP-TgM-Atail, the anti TgM-A tail antibodies simultaneously detected the 40-kDa GFP chimera and the endogenous TgM-A. Expression of the endogenous protein appeared slightly reduced compared with wild-type cells, indicating that the tail alone mimicked the down-regulation effect observed with the full-length myosin. Because the signal for the GFP chimera was approximately half as intense as for mycTgM-A (Figure 2B, first lane), we suggest that the effect is partial because of a lower expression level.

**The Presence of Its Tail Influences the Expression of TgM-D**

TgM-D was well expressed transiently, but our initial efforts to obtain stable transformants were unsuccessful, potentially because mycTgM-D overexpression was not well tolerated by the parasites. To circumvent this problem, we introduced the selectable marker directly in the expression vector and obtained a few stable cell lines expressing mycTgM-D. Ex-
subcellular localization of TgM-A, TgM-D and mutants thereof, two approaches were used. First we generated isoform-specific antibodies. Second, we used an epitope-tagging strategy, taking advantage of the easy accessibility of T. gondii to genetic manipulation. The anti-myc tag antibody revealed that, aside from a relatively diffuse cytoplasmic distribution, a proportion of mycTgM-A localized precisely beneath the plasma membrane (Figure 4A, a), as visualized by the almost perfect colocalization with SAG1, the major surface antigen of T. gondii tachyzoites (Figure 4A, b). Because SAG1 is anchored at the plasma membrane via a glycosylphosphatidylinositol, the staining suggests that TgM-A associates closely with this membrane, although the resolution of light microscopy cannot exclude that the myosin interacts with the inner membrane complex. This latter structure is composed of flattened membrane cisternae found in close apposition with the plasma membrane of all Apicomplexa. It covers the elaborate basket of microtubules and contributes to the maintenance of cell shape and polarity (Morrisette et al., 1997).

Surprisingly, despite strong and specific signals obtained on Western blots (Figure 2), immunofluorescence using the anti-TgM-A tail antibodies failed to detect endogenous TgM-A in wild-type parasites (our unpublished results) and parasites expressing mycTgM-AΔtail (Figure 4B, e) and also failed to corroborate the peripheral localization of mycTgM-A (Figure 4B, compare a and b). Nevertheless, in the latter case a signal was clearly perceived and was similar in intensity (but not in staining pattern) to the cytoplasmic signal obtained for mycTgM-AΔtail (Figure 4B, compare d and b). It is reasonable to assume that the tail of TgM-A, being very rich in arginine and lysine residues, aldehyde cross-linking to plasma membrane proteins, may lead to complete epitope masking. Therefore, we used an alternative fixation method involving rapid freezing fixation and permeabilization in ultra-cold methanol, an ideal method to preserve both structure and antigenicity (Neuhaus et al., 1998). Figure 4C illustrates that both the endogenous TgM-A (Figure 4C, b) and the myc-tagged protein (Figure 4C, a) were now detected by the anti-tail antibody. As expected, the signal for TgM-A was slightly lower in wild-type parasites than in recombinants expressing the myc-tagged protein. The cytoplasmic staining for mycTgM-A (Figure 4B, b) was stronger than for endogenous TgM-A (Figure 4B, e). Finally, irrespective of the expression level, both proteins localized similarly all around the parasite periphery, confirming the epitope-tagging data.

**The Short Tail Domains Are Necessary for Plasma Membrane Localization of the T. gondii Myosins**

Myosin molecules are modular motors made up of three domains. The N-terminal domain is the actin binding motor unit per se. The middle neck domain bears the light chains and acts as a lever arm. The tail domain is exceptionally divergent and reflects the diversity in myosin functions. The tail is thought to target a given myosin to its cargo or site of action and thereby to determine the specific task of the motor. To assess the role of the short tail domains in the subcellular distribution of the two proteins, we constructed vectors expressing myosins lacking their tail. Recombinant parasites expressing mycTgM-AΔtail and mycTgM-DΔtail were analyzed by confocal microscopy (Figure 4, B and D).

**TgM-A Localization Is Confined to the Parasite Periphery**

TgM-A and TgM-D share a high degree of homology, and it is not yet known whether T. gondii expresses additional similar myosins (in addition to the related TgM-B and -C, which harbor longer and different tail domains). To determine unambiguously, without risk of cross-reaction, the
Figure 4. Immunofluorescence localization of TgM-A and TgM-D and their respective tail-less constructs. Classical (A and D) and confocal (B and C) immunofluorescence microscopic analysis of wild-type parasites (C, b) or parasites expressing mycTgM-A (A, B, a–c, and C, a), mycTgM-AΔtail (B, d–f), mycTgM-D (D, a), or mycTgM-DΔtail (D, b) is shown. Fixation was performed either by paraformaldehyde-glutaraldehyde (A, B, and D) or ultracold methanol (C). (A) MycTgM-A, visualized by anti-myc antibodies (a), colocalized at the parasite periphery (c) with SAG1, the major glycosylphosphatidylinositol-anchored surface antigen of *T. gondii* (b). (B and C) The antibody against the tail of TgM-A did not detect TgM-A (B, e) or mycTgM-A (B, b) at the plasma membrane when cells were fixed by paraformaldehyde-glutaraldehyde, even though mycTgM-A was detected at the periphery by anti-myc staining (B, a). The rapid freezing and fixation in ultracold methanol allowed peripheral localization of endogenous TgM-A (C, b) and mycTgM-A (C, a). Myc-TgM-AΔtail was mainly found in the cytoplasm (B, d). The anti-tail antibodies detected a cytosolic pool of mycTgM-A (B, b). (D) As detected by anti-myc antibodies, mycTgM-D was enriched at the parasite periphery (a) but in a way distinct from TgM-A (compare with A–C). MycTgM-DΔtail was mainly cytoplasmic (b).
As already observed above, absence of tail in mycTgM-AΔtail caused the disappearance of the membrane-associated pool (Figure 4B, d).

In a way similar to mycTgM-A, mycTgM-D appeared to distribute predominantly at the parasite periphery (Figure 4D, a), even though not as sharply as mycTgM-A. The subtle difference in distribution may indicate a different mode or mechanism of localization. As in the case of mycTgM-A, the absence of tail caused an apparent redistribution of mycTgM-D to the cytoplasm (Figure 4D, b), and, as mentioned above (Figure 3B), it also led to a significant increase in the level of the recombinant product compared with the full-length protein.

The Tail of TgM-A Is Necessary for Distribution in the Particulate Fraction

To corroborate the localization data with biochemical fractionation, cells were first lysed in PBS, separated into soluble and sedimentable fractions by high-speed centrifugation, and analyzed by Western blotting. A comprehensive analysis was undertaken on cells expressing GFP-TgM-Atail, allowing for simultaneous investigation of endogenous TgM-A and of the GFP tail chimera. In agreement with the immunofluorescence, TgM-A partitioned with the particulate fraction (Figure 5A). The nature of this interaction was further investigated. TgM-A behaved as a strongly associated peripheral membrane protein, because it was resistant to solubilization by high salt. Full solubilization was achieved only by carbonate treatment (pH 11.5) and extraction by detergent (1% Triton X-100) or 3 M urea (our unpublished results).

Even though mycTgM-AΔtail appeared cytoplasmic (Figure 4B, d), it is formally possible that TgM-A localized at the periphery and, despite inclusion of ATP during fractionation, interacted with the particulate fraction through binding of its head domain to the actin cytoskeleton. Therefore, the role of the tail domain was investigated for recombinant TgM-A (Figure 5C). When parasites were lysed in PBS, about one-third to one-half of mycTgM-A sedimented with the particulate, membrane fraction whereas mycTgM-AΔtail partitioned essentially with the soluble cytosolic fraction, in perfect agreement with the immunofluorescence data. This indicated that the sedimentation of TgM-A was not due to interactions through the motor domain but was likely mediated by the tail domain (also see Figure 7 and accompanying text). Similar results were obtained for mycTgM-D (Figure 5C). Moreover, whereas endogenous TgM-A is quantitatively recovered with the particulate fraction (Figure 5A and B), approximately half of mycTgM-A, which is overexpressed two- to threefold, is found in the soluble fraction (Figure 5C), possibly because a plasma membrane receptor is saturated.

TgM-A Binds to F-actin in an ATP-dependent Manner

The peripheral targeting of TgM-A appears not to depend on the interaction of the head domain with the actin cytoskeleton. This could potentially be caused by the fact that the motor domains of class XIV myosins are predicted to exhibit important structural divergences. Therefore, it was crucial to assess experimentally whether these proteins truly act as ATP-driven, actin-dependent motors. In addition, according to the capping model of invasion, the myosin involved would have to be able to grab onto F-actin and exert ATP-dependent traction. The complete solubility of recombinant mycTgM-AΔtail in tachyzoites allowed us to purify biochemical quantities of the protein under native conditions. The degree of purity of the recombinant mycTgM-AΔtail was examined by SDS-PAGE and Coomassie blue staining (Figure 6A). The identity of two low-molecular-mass bands (~30 kDa) has not been fully determined yet (our unpublished results), but they were distinct from degradation products of TgM-A and could potentially represent copurifying light chain(s). To
gain first insights into the biochemical characteristics of this mycTgM-A\_Dtail, a cosedimentation assay with F-actin was performed (Figure 6B). The results showed that mycTgM-A\_Dtail coprecipitated with actin only in the absence of ATP and that the sedimented myosin was subsequently, even though only incompletely, released by the addition of 10 mM ATP, demonstrating that this purified myosin was able to reversibly bind actin in an ATP-dependent manner. Similar results have been obtained with full-length mycTgM-A (our unpublished results). Together, our results also demonstrated the functionality of both recombinant TgM-A constructs.

**The Short TgM-A and TgM-D Tail Domains Are Sufficient to Target a Reporter Protein to the Cell Periphery**

The short basic tails of TgM-A, TgM-D and PfM-A are very similar in length and in amino acid composition. A comparison of the myosin tails is depicted in Figure 7A. The conserved spots of basic residues are highlighted. Analysis of the predicted secondary structure of these domains also revealed a conserved overall organization in three helices interrupted by short loops. To determine whether the short tail domains are sufficient to confer specific cellular distri-

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**Figure 6.** MycTgM-A\_Dtail sediments with F-actin in an ATP-dependent manner. (A) SDS-PAGE and Coomassie blue staining of mycTgM-A\_Dtail purified from an extract of recombinant parasites. (B) SDS-PAGE analysis and Coomassie blue staining of the supernatant (S) and pellets (P) obtained after cosedimentation of mycTgM-A\_Dtail with F-actin in the presence (\(+\)) or absence (\(\text{-}\)) of ATP. MycTgM-A\_Dtail was completely soluble in the absence of F-actin (Control), MycTgM-A\_Dtail could be released from F-actin by subsequent incubation with ATP. Rabbit muscle F-actin was used at 4 \(\mu\)M and ATP at 10 mM in the presence of 130 mM KCl.

**Figure 7.** The tails of TgM-A and TgM-D are sufficient to confer to GFP both peripheral localization and association with the particulate fraction. (A) Alignment of three highly related apicomplexan myosin tails. The basic residues are boxed. (B) Localization of GFP (a), GFP\_TgM-Atail (b), GFP\_TgM-Dtail (c), and GFP\_PfM-Atail (d). Fluorescence was observed directly (a and b) or after indirect immunofluorescence with anti-myc antibodies (c and d). (C) Recombinant parasites expressing GFP, GFP\_TgM-Atail, GFP\_TgM-Dtail, and GFP\_PfM-Atail were lysed in PBS and separated by high-speed centrifugations in soluble (S) and particulate (P) fraction. The partitioning of GFP and the tail chimeras reflected their respective intracellular localization observed by immunofluorescence microscopy.
bution to the myosins, the tails of TgM-A (last 82 amino acids) TgM-D (last 57 amino acids), and PfM-A (last 85 amino acids) were fused to GFP. The plasmids pTGF and pTGF-TgM-Atail were stably integrated into tachyzoites by cotransfection with HXGPR expression vector. We had to introduce the selectable marker gene on pTGF-TgM-Dtail and pTGF-PfM-Atail to obtain stable transformants. The transgenic parasites expressing GFP and GFP-TgM-Atail were analyzed by direct GFP fluorescence, whereas GFP-TgM-Dtail and GFP-PfM-Atail required indirect immunofluorescence analysis using anti-myc antibodies to definitely visualize the recombinant protein (Figure 7B). The nonfusion GFP was abundantly expressed, homogeneously distributed in the cytosol with a significant accumulation in the nucleus (Figure 7B, a). In contrast, GFP-TgM-Atail was found almost exclusively closely associated with the plasma membrane, with barely detectable cytosolic staining (Figure 7B, b). In a way reminiscent of the respective localization of the full-length proteins, GFP-TgM-Dtail showed a slightly more diffuse and speckled peripheral signal than the GFP-TgM-Atail chimera (Figure 7B, c). The difficulty reported above to express high amounts of mycTgM-D appeared indeed to be dependent of the tail domain, because parasites expressing GFP-TgM-Dtail were also difficult to obtain, and the fusion was expressed at a very reduced level compared with both GFP and GFP-TgM-Atail. This was visible on the immunofluorescence stainings and was confirmed by Western blotting (our unpublished results). Similarly, the GFP-PfM-Atail fusion was expressed at a low level and localized essentially to the parasite cytosol (Figure 7B, d).

The GFP-TgM-Atail and GFP-TgM-Dtail Chimeras Partition with the Membrane Fraction

The transgenic parasites expressing GFP and chimeras were lysed in PBS, separated into soluble and particulate fractions, and subsequently analyzed by Western blot. As illustrated in Figure 7C, the migration on SDS-PAGE of the GFP chimera was in agreement with their predicted sizes. GFP-TgM-Atail and GFP-TgM-Dtail completely partitioned with the particulate fraction, whereas GFP and GFP-PfM-Atail were entirely recovered in the supernatant. The quantitative association of GFP with the particulate fraction was comparable with the behavior of endogenous TgM-A. In contrast, GFP-TgM-Dtail sediments despite the fact that the main part of mycTgM-D was soluble. This could potentially indicate that both GFP chimeras were expressed at levels that did not saturate their respective peripheral binding sites. In addition, it is important to note that both chimeras were fully solubilized by carbonate treatment, demonstrating that they were not simply aggregated in the cell (Figure 7C). In fact, the solubilization characteristics of GFP-TgM-Atail were extremely similar to the ones of endogenous TgM-A (see Figure 5A).

The Localization of TgM-A Is Not Due to Interactions with the Actin Cytoskeleton

Altogether, the data indicate a strong interaction of the tail of TgM-A with components of the parasite cortex, likely with proteins associated peripherally with the plasma membrane, even though an interaction with the closely apposed inner membrane complex cannot be excluded. To definitively eliminate the potential targeting role of the cortical actin cytoskeleton, F-actin was severely compromised by cytochalasin D incubation. It has been previously reported that phalloidin fails to stain the actin filaments in T. gondii (Dobrowolski et al., 1997b). Therefore, as indication of the treatment efficacy, the peripheral localizations of mycTgM-A (Figure 8A, a and b) and GFP-TgM-Atail (Figure 8A, c and d) are shown simultaneously with a phalloidin staining of the host actin cytoskeleton. No noticeable change of localization of both mycTgM-A and GFP-TgM-Atail was observed despite severe disturbance of the actin cytoskeleton (Figure 8A, compare b with a and d with c).

The Tail of TgM-A Does Not Localize to the Plasma Membrane of HeLa Cells

Stretches of basic residues in the tail of class I myosins have been shown to contain high-affinity phospholipid binding sites (Dobberstein and Pollard, 1992). If TgM-A were to interact with the plasma membrane solely through unspecific interaction with lipids, the peripheral localization should be observed in unrelated cellular systems. When expressed in HeLa cells (Figure 8B), GFP-TgM-Atail is homogeneously distributed in the cytoplasm, confirming that the association with the particulate fraction in T. gondii was not due to improper folding. More importantly, the chimera showed no sign of membrane association. Indeed, its cytoplasmic localization (Figure 8B, d) was indistinguishable from the one of GFP (Figure 8B, b).

Identification of the Molecular Determinants of Peripheral Membrane Localization

To define precisely, at the amino acid level, the determinants of membrane localization, deletion analysis and site-specific mutagenesis of the basic residues in the tail of TgM-A fused to GFP were undertaken (Figure 9A). Faithful expression and stability of the respective constructs were confirmed by immunoblotting (Figure 9C). The GFP fusions with mutants of TgM-A tail were also examined by indirect immunofluorescence (Figure 9B). Deletion of the C-terminal extension composed of the last 14 amino acids (TgM-AtailA14) did not alter the association of mycTgM-A with the cell membrane. In contrast, a further deletion encompassing the last 22 amino acids (TgM-AtailA22) caused a complete loss of membrane localization. The mutagenesis of two arginine residues within the last 22 amino acids into alanine residues (TgM-Atail mut III) completely abolished localization. However, the conversion of the three other sets of basic residues into neutral amino acids (TgM-Atail mut I and TgM-Atail mut II) did not alter significantly the plasma membrane distribution. These data indicate that the localization of TgM-A is directed by two precise residues rather than by the overall positive charge of the tail domain.

DISCUSSION

A Family of Apicomplexan Myosins Presenting Divergent Structural Features

T. gondii and P. falciparum are both obligate intracellular parasites sharing a similar mechanism of host cell penetra-
tion. According to the capping model for host cell invasion, a myosin is necessary to mediate the ATP-dependent actin-based motility (Sibley et al., 1998). The molecular motor mediating this process has not been identified so far. In the present study, we have cloned and characterized two apicomplexan myosins that share a high degree of homology with the previously identified class XIV of myosins (Heintzelman and Schwartzman, 1997). With a molecular mass ranging between 90 and 93 kDa, these myosins are the smallest molecular motors identified so far. Each of the T. gondii myosins exhibit either a glutamine or an asparagine at the TEDS site, which is neither acidic nor phosphorylatable, potentially implying either different structural requirements for activation or a different post-translational modification involved in the regulation of these myosins. We are exploring the speculative hypothesis that a deamidase might convert asparagine or glutamine into acidic residues at the TEDS site. Among the >100 known myosins, there are few others making exception to the TEDS rule, such as two Acetabularia class XIII myosins, myosin IB of Entamoeba histolytica (Vargas et al., 1997), and a Drosophila class III myosin (Montell and Rubin, 1988). Like some T. gondii myosins, the Drosophila myosin IA has an Asn (Strom Morgan et al., 1994). Another striking feature of all apicomplexan myosins is the presence of a serine instead of the highly conserved glycine corresponding to Gly699 in the chicken myosin II (Kinose et al., 1996). This structural feature has only been reported for a plant myosin (HaMyok3; D. Menzel, unpublished data) and a Caenorhabditis elegans class XII myosin (Baker and Titus, 1997). The conserved glycine residue likely plays an essential role in the conformational changes around the actin- and nucleotide-binding sites, maybe functioning as a pivot point for the lever arm. Mutation of this glycine into an alanine residue dramatically alters the motor activity of the skeletal muscle myosin and the Dictyostelium discoideum myosin II, inhibiting the velocity of actin filament movement by 100-fold (Kinose et al., 1996; Patterson and Spudich, 1996; Patterson et al., 1997). Finally, class XIV myosins do not carry classical IQ motifs for the binding of light chains but harbor a conserved stretch of amino acids that might represent a very divergent form of this motif.

Because of these important structural divergences, it was crucial to assess experimentally whether the apicomplexan myosins truly act as ATP-driven, actin-dependent motors. Also, a requirement of the capping model of invasion is that the motor involved has to grab onto F-actin and exert ATP-dependent traction. The ability to express and purify recombinant myosins from T. gondii offers a unique opportunity to undertake combined in vivo and in vitro structure–function analyses of this novel class of myosins. Taking advantage of the solubility of mycTgM-A tail, we purified the recombinant protein from parasites under native conditions. This material was used in pilot experiments aiming at the biochemical characterization of TgM-A. Indeed, F-actin sedimentation assay showed that TgM-A binds actin in an ATP-dependent manner. The ability to express and purify recombinant myosins from T. gondii offers a unique opportunity to undertake combined in vivo and in vitro structure–function analyses of this novel class of myosins. Taking advantage of the solubility of mycTgM-A tail, we purified the recombinant protein from parasites under native conditions. This material was used in pilot experiments aiming at the biochemical characterization of TgM-A. Indeed, F-actin sedimentation assay showed that TgM-A binds actin in an ATP-dependent manner. Preliminary transient ATPase kinetics and motility experiments indicate further that TgM-A purified from parasites is fully functional (our unpublished results).

**TgM-A Localizes at the Plasma Membrane via Its Tail Domain**

By using a reverse genetic approach and by generating a tail-specific antibody, we have determined the localization of the T. gondii myosin TgM-A. It distributed beneath the
Figure 9. A dibasic motif in the tail of TgM-A is essential for the cell membrane localization of GFP-tail chimera. (A) The wild-type sequence of the tail of TgM-A (wt) is shown aligned with the site-specific mutants (I–III) and the two deletion mutants (Δ14 and Δ22). The mutated residues are indicated in red. (B) Confocal microscopic analysis of parasites stably transformed with the respective mutant tail constructs fused to GFP. Indirect immunofluorescence was performed using the anti-myc antibodies. (C) Anti-myc immunoblot analysis of recombinant parasites expressing the different constructs presented in A.
plasma membrane of tachyzoites, an ideal position to transmit mechanical energy in forward motion and thus to propel cell invasion. At this point, we have to mention that Heinzelman and Schwartzman (1999) recently reported similar data concerning the association of TgM-A with the particulate fraction of *T. gondii*, but contrary to our results, they documented a punctate localization of TgM-A near the apical pole of tachyzoites, possibly associated with some intracellular structure or organelle such as the rhoptries. The discrepancy is probably related to our problems in detecting peripheral localization after aldehyde fixation associated with the spurious detection of a cross-reactive rhoptry antigen.

TgM-D appears enriched in the peripheral region of the parasite but not as sharply defined as for TgM-A. The loss of membrane targeting of TgM-A and TgM-D resulting from deletion of their tail supports the idea that this domain mediates interaction with cargo and brings the myosin to its site of action. Conversely, fusion of the TgM-A tail to GFP confers membrane association to that otherwise soluble cytosolic protein. These data strongly support that the tail domain is necessary and sufficient determinant mediating specific interactions of the myosin with integral or peripheral membrane constituents.

Expression of mycTgM-A Down-regulates the Level of Endogenous TgM-A

MycTgM-A and GFP-TgM-Atail transgenes were well expressed in the parasites. Whereas the GFP chimera was essentially associated with the periphery and the particulate fraction, only approximately half of mycTgM-A behaved that way, suggesting that the specific membrane “receptor” is saturable. Intriguingly, drastic down-regulation of endogenous TgM-A was observed upon expression of recombinant mycTgM-A. This phenomenon appeared to be associated with overexpression of the full TgM-A and at least partially also of GFP-TgM-Atail but was not seen with mycTgM-Atail. We hypothesize that the parasites autoregulate the level of TgM-Â in accordance with the abundance of a receptor present in limiting amounts at the cell periphery.

The difficulties encountered in the generation of recombinant parasites expressing TgM-D and the very low level of protein produced suggest a similar phenomenon. The high expression levels of TgM-DÂtai contrast with the low expression levels of GFP-TgM-Dtail and argue for a regulatory effect of the tail domain. Analogously, the tail of PIM-A seems to induce a comparable effect. It is conceivable that the limited expression of TgM-D, GFP-TgM-Dtail, and GFP-PIM-Atail are due to their inappropriate presence in cells lacking either a stage- or species-specific partner, respectively. The potential role of stage-specific regulation is not well understood yet. Preliminary studies on the developmental pattern of *T. gondii* myosin gene expression suggest that TgM-A is constitutively expressed in tachyzoites and bradyzoites, whereas TgM-D transcripts accumulate predominantly in bradyzoites (F. Debac, A. Sänger, C. Toursel, S. Tomavo, and D. Soldati, unpublished results).

Note that in all cases expression is limited, but we did not notice deleterious effects on the morphology and proliferation of parasites. This contrasts with observations reported for the expression of other myosins. Overexpression of entire unconventional myosin myoIB from *D. discoideum* is an example of deleterious effects leading to a phenotype as severe as observed in null cells (Novak and Titus, 1997). Similarly, the overexpression of MYO4 or MYO2 in *S. cerevisiae* led to morphological abnormalities (Haarer et al., 1994).

The exact nature of the interaction between TgM-A tail and the cell membrane is not elucidated yet; however, in addition to the down-regulation phenomenon discussed above, the following evidence further points toward a parasite-specific proteinaceous receptor. First, the fractionation experiments with mycTgM-A as well as GFP-TgM-Atail argue for a very strong interaction of the tail with the constituents of the plasma membrane. Second, the tails of TgM-A and PPM-A have similar overall charges but markedly differ in their association with the periphery. Third, the mutagenesis experiments indicate that two specific residues and not simply the overall charge of the tail domain are important for localization, rendering it unlikely that it interacts nonspecifically with phospholipids. This is corroborated by the fact that the GFP-TgM-Atail had an exclusively cytoplasmic distribution in HeLa cells. Finally, after aldehyde fixation, the failure to detect the tail antigen specifically when TgM-A is at the plasma membrane suggests that it is engaged in strong interactions with a protein component.

Is TgM-A or TgM-D Powering Gliding Locomotion?

To accommodate the capping model for invasion, the myosin shall lie beneath the plasma membrane and interact with transmembrane proteins exposed at the cell surface at the time of invasion. The motility shall then be driven by the concerted redistribution of these transmembrane proteins toward the posterior pole of the parasite. The TRAP protein (trombospondin-related adhesive protein) expressed in the sporozoites of *Plasmodium* species and its homologue MIC2 in *T. gondii* (Wan et al., 1997) are exported at the surface of the parasite and relocate to the posterior pole during invasion (Carruthers et al., 1999). This protein is an excellent candidate to interact directly or indirectly with the actomyosin system. Disruption of the TRAP gene in *Plasmodium berghei* demonstrates that this protein is necessary not only for sporozoite infection of the mosquito salivary glands and the rat liver but also for gliding motility of the sporozoites in vitro (Sultan et al., 1997). The TgMIC2 C-terminal domain has recently been shown to functionally complement TRAP mutants lacking their C-terminal domain in *P. berghei* (Kappe et al., 1999). Although MIC2 constitutes an attractive partner for TgM-D during invasion, it is only transiently present at the surface of the parasite during host cell penetration and therefore is unlikely to serve as receptor for TgM-A at the plasma membrane during the intracellular replicative phase of the life cycle (Carruthers and Sibley, 1997). It is conceivable that an abundant protein serves as a docking partner, ideally positioning the myosin close to its later site of action. Once at the plasma membrane, the myosin could respond immediately to an activation signal and trigger invasion by establishing a connection with proteins such as MIC2.

A recent study provides evidence for a myosin II responsible for the capping of surface receptors in *E. histolytica* (Arhets et al., 1998). In light of such a finding, it is rather surprising that no conventional myosin of type II has been
identified in Apicomplexa so far. In comparison, despite its simplicity, even \textit{S. cerevisiae} relies on the function of five myosins from three distinct classes (Brown, 1997). Our most recent studies revealed the presence of a fifth myosin of class XIV in \textit{T. gondii} (our unpublished results). Nevertheless, because our screening is not exhaustive, we cannot yet exclude the presence of other classes of myosins in \textit{T. gondii}. The genome sequencing project for \textit{P. falciparum} is still in progress, and the current status reveals that at least two additional myosin genes are present in this parasite, both of which appear to be closest to class XIV. Although not yet confirmed by functional data, TgM-A is definitely guilty by localization. Besides unraveling its potentially crucial role in invasion, the investigation of the biochemical details of enzymatic and motile properties of this very divergent class of unconventional myosins holds the promise of shedding light on the fundamental requirements of myosin function.

\textbf{Note added in proof.} A fragment corresponding to the complete PfM-A cDNA reported here was previously described as Pfmyo-1 (Pinder et al., 1998). We named all the class XIV myosins according to the nomenclature introduced by Heintzelman and Schwartzman (1997). To minimize the confusion, Pinder and colleagues now kindly agree to rename Pfmyo-1 Pfmyo-A.

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