Plasmodium falciparum Rab5B Is an N-Terminally Myristoylated Rab GTPase That Is Targeted to the Parasite’s Plasma and Food Vacuole Membranes

Carinne Ndjembo Ezougou1,2, Fathia Ben-Rached1,2, David K. Moss4, Jing-wen Lin3,4, Sally Black4, Ellen Knuepfer4, Judith L. Green4, Shahid M. Khan3, Amitabha Mukhopadhyay5, Chris J. Janse3, Isabelle Coppens4, Hélène Yera1,2, Anthony A. Holder4,*, Gordon Langsley1,2,*

1 Laboratoire de Biologie Cellulaire Comparée des Apicomplexes, Département d’Immunologie, Inflammation et Infection, Faculté de Médecine, Université Paris Descartes - Sorbonne Paris Cité, Paris, France, 2 Institut National de Recherche Médecale U1016, Centre National Recherche Scientifique UMR8104, Cochin Institute, Paris, France, 3 Parasitology, Center of Infectious Diseases, Leiden University Medical Center, Leiden, The Netherlands, 4 Division of Parasitology, MRC National Institute for Medical Research, Mill Hill, London, United Kingdom, 5 National Institute of Immunology, Aruna Asaf Ali Marg, New Delhi, India, 6 Department of Molecular Microbiology and Immunology, Johns Hopkins University Bloomberg School of Public Health, Baltimore, Maryland, United States of America

Abstract

Plasmodium falciparum (Pf) has a family of 11 Rab GTPases to regulate its vesicular transport. However, PfRab5B is unique in lacking a C-terminal geranyl-geranylation motif, while having N-terminal palmitoylation and myristoylation motifs. We show that the N-terminal glycine is required for PfRab5B myristoylation in vitro and when an N-terminal PfRab5B fragment possessing both acylation motifs is fused to GFP and expressed in transgenic P. falciparum parasites, the chimeric PfRab5B protein localizes to the plasma membrane. Upon substitution of the modified glycine by alanine the staining becomes diffuse and GFP is found in soluble subcellular fractions. Immuno-electron microscopy shows endogenous PfRab5B decorating the parasite’s plasma and food vacuole membranes. Using reverse genetics rabs5 couldn’t be deleted from the haploid genome of axenial blood stage P. berghei parasites. The failure of PbRab5A or PbRab5C to complement for loss of PbRab5B function indicates non-overlapping roles for the three Plasmodium Rabs5, with PfRab5B involved in trafficking MSP1 to the food vacuole membrane and CK1 to the plasma membrane. We discuss similarities between Plasmodium Rab5B and Arabidopsis thaliana ARA6, a similarly unusual Rab5-like GTPase of plants.

Introduction

Plasmodium falciparum is a parasite that infects red blood cells (RBC) and causes severe human malaria, a disease that kills approximately 750,000 people per year, mostly children living in tropical Africa [1]. The malaria parasite spends much of its life cycle inside RBC, cells that provide it with an abundant food supply in the form of haemoglobin and a degree of protection from the host’s immune system. For survival inside the RBC the parasite has to both import nutrients and to export metabolic waste products and like other eukaryotes it uses Rab GTPases to regulate vesicular trafficking [2] [3] [4].

Rabs are molecular switches belonging to the Ras-superfamily that are conserved from yeast to humans and regulate in space and time the budding and fusion of intracellular vesicles from donor to acceptor membranes [5]. Rabs vary in size from 20 to 29 kDa and were first identified in yeast and named Ypt proteins. Eleven Ypts were characterized in Saccharomyces cerevisiae and seven in Schizosaccharomyces pombe [6], whereas higher eukaryotes possess more than 60 different Rabs [7]. Rab proteins exist in GTP-bound and GDP-bound states with switching from one state to another regulated by GTP Exchange Factors and GTPase Activating Proteins [8] [9]. GDP-bound Rabs are considered to be inactive and are associated with Rab GDP Dissociation Inhibitors (rabGDIs), whereas GTP-bound Rabs are considered to be active and able to recruit and interact with their effectors, which is a key specific Rab-mediated function [10]. Rabs from different species display a significant degree of sequence conservation in the GTP-binding domain, but they possess a variable N-terminal sequence responsible for recruitment of effectors and a hyper-variable C-terminal domain. The C-terminal domain confers the specific intracellular location for each Rab and has a motif containing one or two cysteines (CAAX, CC, CXC, CCX, CCXX) necessary for
The PfRab-interactome predicted that Casein Kinase 1 (PICK1; PF3D7_1136000.1) is a specific PfRab5B-interacting protein and indeed, PfRab5B physically interacts with PICK1 in vitro [20]. The subcellular distribution of PICK1 has not yet been described [26], but in T. gondii CK1 is found both in the cytosol and at the plasma membrane [27]. We decided to test therefore, whether in vivo PfRab5B is competent to traffic PICK1 to different subcellular locations in P. falciparum. Moreover, as PfRab5B lacks the necessary motif for isoprenylation, yet possesses recognisable myristoylation and palmitoylation motifs [19], we examined directly whether the N-terminal glycine residue of PfRab5B is a bona fide substrate for P. falciparum N-myristoyltransferase (NMT) [28] in vitro and whether in vivo its myristoylation is involved in targeting PfRab5B to specific subcellular membranes. Based on our results we propose that myristoylation (and palmitoylation) of PfRab5B allows it to traffic specific cargo to the parasite’s plasma and food vacuole membranes.

Results

The N-terminal glycine residue in PfRab5B is required for N-myristoylation

To test the validity of the myristoylation motif prediction we produced recombinant PfRab5B protein and variants in which either the glycine or the glycine and cysteine residues at the N-terminus of PfRab5B (MGCSS) were changed to alanine. These His-tagged full-length recombinant wild type and variant PfRab5B proteins were purified and used as substrates in an N-myristoylation assay using recombinant P. falciparum N-myristoyltransferase (PfNMT) [28] and tritiated myristoyl-CoA [29] (see Figure 1). Wild type and G2A-variant recombinant PfARF protein (PF3D7_1020900) were used as controls [30] [29]. The results showed that PfRab5B is a substrate for PfNMT (although the efficiency of myristoyl transfer was less than that for PfARF) and the presence of a glycine at position two is required for PfRab5B myristoylation in vitro, which occurs via peptide bond formation with the α-amino group of the N-terminal glycine. As expected, modification of both the myristoylation and the palmitoylation site (G2AC3A) had no additional effect.

Figure 1. Recombinant PfRab5B is N-myristoylated by PfNMT in vitro. PfRab5B and the G2A and G2AC3A variants were incubated with [3H]-myristoyl CoA in the presence of N-myristoyl transferase. Recombinant ARF and ARFG2A were treated in the same way, acting as controls. N-myristoylation was detected by the incorporation of radiolabel into the substrate following SDS-PAGE and fluorography [30] [29]. The results showed that PfRab5B is a substrate for PfNMT (although the efficiency of myristoyl transfer was less than that for PfARF) and the presence of a glycine at position two is required for PfRab5B myristoylation in vitro, which occurs via peptide bond formation with the α-amino group of the N-terminal glycine. As expected, modification of both the myristoylation and the palmitoylation site (G2AC3A) had no additional effect.
PfRab5B myristoylation in the parasite is essential for its targeting to membranes

We generated a transgenic *P. falciparum* line that expresses GFP fused to the N-terminal 28 amino acids of PfRab5B (see Table S1 in File S1). An episomal maintained plasmid was used for transfection, which expresses the fusion protein under the control of the *msp3* promoter [31] [32]. Asexual blood stages of this transgenic line were analysed by fluorescence microscopy. GFP-fluorescence was consistent with a location of the fusion protein in association with the parasite’s plasma membrane (Figure 2, top left hand panel). As a positive control, we used a second transgenic *P. falciparum* line that expresses GFP fused to 29 amino acids from the N-terminus of glideosome associated protein (GAP45 [Table S1 in File S1], GAP45 is known to be associated with the plasma membrane at its N-terminus [31]. In asexual blood stages of this transgenic parasite GFP was observed at the plasma membrane (Figure 2, top right hand panel). In transgenic parasites that expressed GFP fused to G2A variants of both PfRab5B and GAP45 the association of GFP with the plasma membrane was lost and we observed a diffuse GFP-staining throughout the cytoplasm of the parasites (Figure 2, second row panels). This cytoplasmic GFP-staining was also observed in blood stages expressing GFP-fusion proteins in which additional substitutions had been introduced (Table S1 in File S1): the C3A change in PfRab5B or C5A in GAP45 to give the double G+C variants (Figure 2, third row panels). The C to A variants alone allowed some membrane association (Figure 2, bottom row). Clearly, the N-terminal 28 or 29 amino acids in combination with myristoylation of G2 of both PfRab5B and GAP45, respectively, are sufficient to target the GFP-fusion proteins to the parasite’s plasma membrane.

PfRab5B-GFP and GAP45-GFP fusion proteins are embedded in membranes of *P. falciparum* parasites

Schizonts of transgenic parasites expressing GFP fused to N-terminal sequences of PfRab5B and GAP45 (28 or 29 amino acids, respectively) were purified and submitted to subcellular fractionation using hypotonic and high salt buffers to release cytosolic proteins, followed by high pH carbonate extraction to fractionate peripheral membrane (carbonate soluble) and integral membrane (carbonate insoluble) proteins. The proteins in the four different fractions were resolved by SDS-PAGE and analysed by western blotting with an anti-GFP antibody (Figure 3). When the wild type PfRab5B N-terminal sequence was fused to GFP (PfRab5B28-GFP), the protein was found in all subcellular fractions (to differing degrees) indicating that not all of this fusion protein was associated with membranes despite the presence of both G2 and C3 residues. In contrast, both the GFP-fusion proteins containing the G2A and C3A variants were located solely in the hypotonic lysis soluble fraction, as was the variant with both substitutions (Figure 3, track 1). The GFP protein fused to the wild type GAP45 N-terminal sequence (GAP4529-GFP) was located exclusively in the carbonate-insoluble fraction (track 4) typical for a membrane-associated protein. Similar to the PfRab5B28-G2A variant of the GFP fusion protein, GAP4529-G2A was found exclusively in the soluble fraction consistent with myristoylation alone being necessary and sufficient for membrane association.

Endogenous PfRab5B is associated with both the food vacuole and plasma membranes

We generated antibodies to PfRab5B and verified their specificity by western blot (see Figure S1). These specific PfRab5B-specific antibodies were then used to determine the subcellular location of PfRab5B by immuno-electron microscopy. In Figure 4 micrographs are shown of two different stages of intraerythrocytic development, mature schizonts (Panel A) and a trophozoite (Panel B). Gold-labelled anti-PfRab5B antibodies decorated the parasite plasma membrane (PPM, white arrows) and were also observed associated with the food vacuole (FV) and neighbouring vesicles, as shown for the trophozoite in panel B. The distribution of endogenous PfRab5B was therefore similar to that of PfRab5B-GFP suggesting that the plasma- and food vacuole membrane associations are the result of myristoylation and/or palmitoylation at the N-terminus of PfRab5B.

Endogenous PfRab5B colocalises with merozoite surface protein (MSP)1 and PfCK1, but not with haemoglobin

GFP-tagged PfRab5A has been described as associated with vesicles harbouring haemoglobin imported by the parasite from the erythrocyte cytosol [21]. The association of PfRab5B with the food vacuole membrane and adjacent vesicles led us, therefore, to look at an association with haemoglobin-containing structures using indirect immunofluorescence. Specific anti-PfRab5A antibodies (see Figure S1) were used as a positive control. In contrast to PfRab5A (*r* = 0.724; *n* = 3) we could find no evidence for co-localisation between PfRab5B and haemoglobin (HBA1, *r* = 0.081; *n* = 3) in trophozoites, the developmental stage that imports haemoglobin (Figure 5, panel A).

Following merozoite invasion, a C-terminal fragment of MSP1 is associated with the developing food vacuole membrane [33], a localisation that is similar to that observed for PfRab5B, as shown in Figure 4. We confirmed therefore that antibodies to MSP1 and PfRab5B significantly (*r* = 0.849; *n* = 3) decorate the same structure (Figure 5, panel B). As PfRab5B was found to physically interact with PfCK1 *in vitro* [20] we looked for evidence that PfRab5B-positive vesicles might traffic the kinase to different subcellular locations. Depending on the developmental stage of the parasite (indicated by the number and size of DAPI-stained nuclei; Figure 5, panel C) the degree of association between PfRab5B and PfCK1 varied (*r* = 0.615; *n* = 3 in late schizonts and *r* = 0.475; *n* = 3 in young schizonts). Clearly, PfRab5B-positive vesicles can traffic more than one type of cargo, however the failure to observe any association of PfRab5B with haemoglobin suggests that PfRab5B and PfRab5A perform separate trafficking functions.

*Plasmodium* Rab5B is essential for asexual blood stage parasite development

Over-expression of individual Rab5s and/or Rab5 GDP-ON variants in transgenic cells has the potential to generate phenotypes due to sequestering effector proteins that are shared by the different Rab isoforms, and therefore the use of such ‘over-expressing’ transgenic cells may hamper discrimination between the different functions of the Rab isoforms. Therefore, we decided to analyse the function of *Plasmodium rab5b* by attempting to switch off its expression using targeted gene-deletion by reverse genetics. We targeted the orthologous rab5b gene of the rodent malaria parasite *P. berghei* for gene deletion using standardized transfection methods [34]. In multiple experiments (Figure S2 in File S1) we were unable to select for mutants lacking the *Pbrab5b* gene indicating that *Pbrab5b* has an essential function during asexual

Myristoylated Rab5B of *Plasmodium falciparum* parasites

Plasmodium rab5b gene of the rodent malaria parasite *P. berghei* for gene deletion using standardized transfection methods [34]. In multiple experiments (Figure S2 in File S1) we were unable to select for mutants lacking the *Pbrab5b* gene indicating that *Pbrab5b* has an essential function during asexual
blood stage growth and multiplication. Moreover, these results also indicate that PbRab5A and PbRab5C cannot compensate for the function of PbRab5B.

**Discussion**

*P. falciparum* has a small family of 11 PfRabs and all but PfRab5B possess classical geranyl-geranylation motifs at the C-terminus. This isoprenyl modification is present in 10 of the PfRabs presumably anchoring these GTPases to membrane. In contrast, we have shown here that PfRab5B is myristoylated allowing membrane attachment at the N-terminal glycine. PfRab5B peptides have recently been detected in a global *in vivo* myristome of *P. falciparum* [35]. Our inability to knock out the *rab5b* gene of *P. berghei* indicates that *Plasmodium* Rab5B performs functions essential for parasite asexual blood stage development. Potential functions may include trafficking MSP1-tagged vesicles from the plasma membrane to form the food vacuole membrane following RBC invasion [33] and PfCK1 to the plasma membrane. However, in contrast to PfRab5A [21] we never observed PfRab5B associated with haemoglobin-positive structures indicating that these two *Plasmodium*-specific Rab5 isoforms perform separate functions.

The presence of the wild type PfRab5B-GFP fusion protein in different subcellular fractions suggests that endogenous PfRab5B associates with varying affinity to membranes. However, as the wild type PfRab5B-GFP fusion protein was also found in the soluble fraction the results suggest that not all of the endogenous PfRab5B is myristoylated and/or palmitoylated at a given time. Substitution of the N-terminal glycine with alanine abolished the association of the PfRab5B-GFP fusion protein with membrane, clearly demonstrating that myristoylation of the PfRab5B N-terminal amino acid sequence is necessary and sufficient to attach it to membranes. Although we did not directly examine the palmitoylation status of PfRab5B substituting a cysteine with an alanine ablated membrane-targeting of the GFP-fusion protein as judged by solubility analysis, although some apparent association with the plasma membrane was detected by live cell imaging. Palmitoylation of endogenous PfRab5B therefore, seems to contribute to its subcellular location and indeed, a recent analysis of the *P. falciparum* palmitoylation proteome identified PfRab5B, as a component [36,37]. Protein palmitoylation is reversible and

![Figure 2. Localisation of PfRab5B-GFP fusion proteins in *P. falciparum*. Parasites were transfected with constructs expressing the N-terminal 28 amino acids of PfRab5B or 29 amino acids of PfGAP45 fused to GFP under a schizont stage-specific promoter (*msp3* 3′UTR region). Localisation of PfRab5B-GFP (A) and GAP45-GFP fusions (B) as well as myristoylation (G2A) and palmitoylation-deficient fusions (C3A in PfRab5B-GFP and CSA in GAP45-GFP) were investigated. The first image in a series corresponds to GFP fluorescence, the second a merge of GFP fluorescence with nuclear DAPI stain, and the third a merge of GFP fluorescence, DAPI and bright field images. Size bars are 2 μm. doi:10.1371/journal.pone.0087695.g002](http://www.plosone.org/figure2.html)
Substitution of glycine at position two with alanine (PfRab5B28 G2A) GFP is found to varying degrees in the four fractions studied. endogenous PfRab5B (Fig. 4), and as only the first N-terminal 28 amino acids of PfRab5B are present in the GFP-chimera this implies that the C-terminus of PfRab5B might contribute to its food vacuole location. However, unlike PfRab5B, palmitylation of ARA6 is not required for membrane association per se, only for endosome-specific targeting. It is possible that additional targeting information is encoded in the N-terminal amino acid sequence of myristoylated and palmitylated proteins, since the subcellular fractionation of PfRab5B-GFP and GAP45-GFP fusion proteins was not identical following replacement of the palmitylated cysteine residue. For PfRab5B with the C3A substitution the fusion protein was found only in the soluble fraction, in contrast to GAP45 with the C5A substitution, for which the fusion protein was distributed across all subcellular fractions. The N-terminal sequence of GAP45 fused to GFP has ten basic and six acidic residues, whereas the N-terminal sequence of PfRab5B fused to GFP has only four basic and one acidic residue, so the GAP45 N-terminal sequence is much more highly charged and this greater charge might contribute to greater affinity for membrane phospholipid head groups.

ARA6 acts in an endosomal trafficking pathway by modulating the assembly of a distinct SNARE complex that includes VAMP727 [16]. A family of P. falciparum SNAREs has been described [39] [40] and PfVAMP8 (PF3D7_1303200.1) displays the highest identity (5.6e-35) to VAMP727 suggesting that perhaps PRab5B regulates trafficking via assembly of PfVAMP8 complex. Whether a P. falciparum PRab5B/PfVAMP8 complex is involved in regulating a stress response, as is the ARA6/VAM727 complex in Arabidopsis [17] is an interesting and open question. Unlike PRab5A we never observed PfRab5B associated with haemoglobin containing vesicles, implying that haemoglobin uptake would likely not involve the PfVAMP8 complex. Clearly, in malaria causing parasites in spite of having a small Rab family of only 11 GTPases, PfRab5A, PfRab5B and PfRab5C isoforms regulate independent and crucial functions.

Materials and Methods

Attempts to generate P. berghei rab5b gene deletion mutants

Female Swiss OF1 mice (6-8 weeks old; Charles River/Janvier) were used. All mouse experiments received approval from the Animal Experiments Committee at the Medical Centre of Leiden University (DEC 0717; DEC 10099). The Dutch Experiments on Animal Act is established under European guidelines (EU directive no. 86/609/EEC regarding the Protection of Animals used for Experimental and Other Scientific Purposes). In order to create P. berghei rab5b (PBANKA_140910) gene deletion mutants, a DNA construct pL1709 was generated that would target Pb rab5b gene deletion (Figure S1), as double crossover homologous recombination (Figure S1), as described [34] and PbVAMP8 (PF3D7_1303200.1) displays the highest identity (5.6e-35) to VAMP727 suggesting that perhaps PRab5B regulates trafficking via assembly of PfVAMP8 complex. Whether a P. falciparum PRab5B/PfVAMP8 complex is involved in regulating a stress response, as is the ARA6/VAM727 complex in Arabidopsis [17] is an interesting and open question. Unlike PRab5A we never observed PfRab5B associated with haemoglobin containing vesicles, implying that haemoglobin uptake would likely not involve the PfVAMP8 complex. Clearly, in malaria causing parasites in spite of having a small Rab family of only 11 GTPases, PfRab5A, PfRab5B and PfRab5C isoforms regulate independent and crucial functions.

Materials and Methods

Attempts to generate P. berghei rab5b gene deletion mutants

Female Swiss OF1 mice (6-8 weeks old; Charles River/Janvier) were used. All mouse experiments received approval from the Animal Experiments Committee at the Medical Centre of Leiden University (DEC 0717; DEC 10099). The Dutch Experiments on Animal Act is established under European guidelines (EU directive no. 86/609/EEC regarding the Protection of Animals used for Experimental and Other Scientific Purposes). In order to create P. berghei rab5b (PBANKA_140910) gene deletion mutants, a DNA construct pL1709 was generated that would target Pb rab5b gene deletion (Figure S1), as previously described [34]. The targeting plasmid was made to double crossover homologous recombination (Figure S1), as previously described [34]. The targeting plasmid was made to target both the 5’ and 3’ regions of Pb rab5b gene and contains the drug selectable marker human dhfr (dhfr) (See Table S2 in File S1 for details of the primers). Prior to transfection, the plasmid was linearized by digestion with enzymes HindIII and EcoRI. The P. berghei ANKA reference reporter parasite line Pb/GFP-Luc+ of the ANKA strain of P. berghei was used for two independent transfection experiments (See RMgm-29 in www.pberghei.eu for details of the Pb/GFP-Luc+ line). All experimental procedures including animal work and transfection and selection protocols to generate gene knock-out parasites were performed as described [34].

Myristoylated Rab5B of Plasmodium falciparum

Figure 3. Subcellular fractionation of PfRab5B and GAP45 GFP chimeras. When fused to GFP, the first 28 amino acids of PfRab5B confer a partial membrane association to the fusion protein. PfRab5B28-GFP is found to varying degrees in the four fractions studied. Substitution of glycine at position two with alanine (PfRab5B28 G2A) results in loss of all membrane association, as does substitution of cysteine at position three (PfRab5B28 C3A). A known myristoylated and palmitylated protein, GAP4529-GFP is found exclusively in membrane fractions and a G2A substitution results in an entirely soluble protein (GAP4529 C5A). Double substitutions of glycine and cysteine result in a reduction in the proportion of protein associated with membranes (GAP4529 G2A). A substitution of cysteine at position five causes a reduction in the proportion of protein associated with membranes (GAP4529 C5A). Double substitutions of glycine and cysteine result in a cytosolic protein for both PfRab5B28 and GAP4529 GFP fusion proteins. HL: hypotonic lysis buffer supernatant, HS: high salt buffer supernatant, CS: cytosolic buffer supernatant, Cl: cytosolic buffer insoluble.

doi:10.1371/journal.pone.0087695.g003

dynamic and can result in proteins targeted to specific membrane domains [38].

PfRab5B resembles ARA6 a Rab5-like GTPase considered unique to plants such as A. thaliana [16,17,18]. When ARA6 was fused to GFP and expressed in protoplasts prepared from cultured Arabidopsis cells, both punctate and plasma membrane localisation was observed [18]. Interestingly, when the N+3 cysteine was changed to serine leaving the N+2 glycine intact ARA6 became uniquely myristoylated and the GFP-fusion no longer localised to endosomes, indicating that additional palmitoylation is necessary for endosome targeting [18]. When the N-terminal glycine was changed to an alanine ARA6-GFP was found in the cytosol and 3 glycine intact ARA6 became uniquely myristoylated and the GFP-fusion no longer localised to endosomes, indicating that additional palmitoylation is necessary for endosome targeting [18].
Expression constructs and purification of recombinant proteins

To make C-terminal-tagged PfRab5A-hexaHis and PfRab5B-hexaHis constructs for expression in *Escherichia coli*, the CDS of *Pfrab5a* (PF3D7_0211200) and *Pfrab5b* (PF3D7_1310600) were PCR amplified from *P. falciparum* 3D7 cDNA and cloned into *BamHI*/*XhoI* sites of the *E. coli* expression vector pET-21d (Novagen, EMD Biosciences). The sequences of all expression constructs were verified.

*Pfrab5a* and *Pfrab5b* constructs were transformed into BL21-CodonPlus (DE3)–RIL strain (Stratagene). BL21 cells were grown at 37°C in LB medium containing 34 mg/ml chloramphenicol and 100 mg/ml ampicillin to an absorbance of approximately 0.6 at 600 nm. Protein expression was induced by adding 0.2 mM IPTG, and the cultures were incubated overnight at 20°C. Cells were harvested by centrifugation at 5000 g for 20 min and re-suspended in 6 M urea buffer supplemented with protease inhibitor cocktail (Roche), then stored at -80°C. The His-tagged proteins were purified on Ni-NTA agarose (Qiagen).

To generate *Pfrab5B<sup>G2A</sup>*, *Pfrab5B<sup>C3A</sup>* and *Pfrab5B<sup>G2A,C3A</sup>* mutations, PCR site-directed mutagenesis was performed using *Pfrab5b* cloned into pET-21d vector as a template. Thereafter, PCR products were incubated at 37°C with DpnI enzyme for an hour. The sequences of all expression constructs were verified.

Specific antibodies

Specific PfRab5A antibodies were raised in rabbits and specific anti-PfRab5B antibodies were raised in rats (Eurogentec) to full-length His-tagged recombinant PfRab5A and PfRab5B (see above). Specific anti-PICK1 antibodies were a gift of Dominique Dorin-Semblat and Christian Doerig and anti-MSP1<sub>19</sub> antibodies have been described [41].

In vitro N-myristoylation assay

The *in vitro* myristoylation assay was performed essentially as described previously [29] using bacterially expressed recombinant PNMT, recombinant substrate protein and [<sup>3</sup>H]myristoyl-CoA. Recombinant PIARF and G2A PIARF were used as controls. 0.5 μg PNMT was incubated with either 5 μg PfRab5B or 2.5 μg ARF proteins and myristoyl-CoA (0.007 MBq) for 2 h at room temperature in reaction buffer (30 mM Tris-Cl, pH 6.8, 0.5 mM EGTA, 0.45 mM EDTA, 4.5 mM 2-Mercaptoethanol, 1% Triton X-100). The amount of myristoylated protein was determined by SDS-PAGE and autoradiography.

Figure 4. Ultrastructural detection of PfRab5b in *P. falciparum*-infected RBC. Immuno-electron microscopy of schizonts (A) and a large trophozoite (B) of *P. falciparum* labelled with specific anti-PfRab5B antibodies, revealing the presence of gold particles both on the food vacuole (FV) and the parasite plasma membrane (PPM, white arrows) in trophozoites and exclusively on parasite plasma membrane for late stages. P, parasite; r, rhoptry. Scale bars, 150 nm.

doi:10.1371/journal.pone.0087695.g004
X-100). Fractions from each reaction were resolved by SDS-PAGE under reducing conditions on precast 12% NuPAGE acrylamide gels. The gels were treated for fluorography, dried and exposed to X-ray film to detect the tritium label.

Parasite culture

*Plasmodium falciparum* 3D7 was maintained in human O+ erythrocytes using RPMI 1640 medium supplemented with 1% Albumax at 3% hematocrit in gassed (90% nitrogen, 5% oxygen, 5% carbon dioxide) in flasks at 37°C. Parasites were synchronized using Percoll gradient centrifugation.

**Figure 5.** PfRab5B colocalises with PfMSP1 and PfCK1, but not with haemoglobin. (A) PfRab5A colocalises with haemoglobin (HBA1) containing vesicles \( r = 0.724 \), unlike PfRab5B \( r = 0.081; n = 3 \). (B) PfRab5B colocalises to differing degrees \( r = 0.849; n = 3 \) with the C-terminal 19 kDa fragment of PfMSP1 on structures close to the food vacuole and the parasite nucleus shown in blue by DAPI staining. (C) PfRab5B colocalises with PICK1 on intracellular structures \( r = 0.615; n = 3 \) and at the parasite plasma membrane \( r = 0.475; n = 3 \). Areas of colocalisation are shown in white and used to calculate Pearson’s \( r \) coefficients. Scale bars, 2 \( \mu \)m.

Plasmid constructs and transfection of *P. falciparum*

To generate GFP-fusion protein expressing parasites we used primers to amplify the sequence encoding the N-terminal fragments of PfRab5B and GAP45, and inserting nucleotide changes to introduce alanine instead of glycine or cysteine as indicated in Table S1 in File S1. The PCR reactions were purified using a QIAGEN quick purification kit and digested with Sall and AvrII. The digested inserts were gel extracted then ligated with digested pHH3 vector distal to the msp3 promoter [31]. Resulting plasmids were transformed into DH5α and the sequence of each construct was established using DNA sequencing (Geneservice). 100 µg of sterile plasmid DNA in 100 mM Tris-HCl, 10 mM EDTA, pH 8.0 buffer was transfected by electroporation into
parasites at 10% ring stage parasitaemia and culture was continued in the presence of blasticidin (2.5 μg/ml).

Subcellular fractionation of schizonts and western blotting
The method described previously [32] was used with minor modifications; all incubations were carried out at 4°C and the buffers contained complete protease inhibitor cocktail (Roche). Parasite pellets were solubilised in 10 times the volume of hypotonic lysis buffer (10 mM Tris, 5 mM EDTA, pH 8.0) followed by centrifugation at 100,000 g for 30 min at 4°C and recovery of the supernatant. The pellet was washed once by repetition of the process and subsequently extracted with high-salt buffer (10 mM Tris, 5 mM EDTA, 500 mM NaCl, pH 7.5), followed by centrifugation. The supernatant was retained and the pellet was extracted again in carbonate buffer (100 mM sodium carbonate, pH 11.0). Following a final centrifugation the three supernatants and the remaining carbonate-insoluble material were each mixed with SDS sample loading buffer and subjected to SDS-PAGE and western blotting with anti-GFP antibodies. Samples were heated at 95°C for 5 min prior to being separated by SDS-PAGE on a pre-cast 12% Bis-Tris NuPAGE polyacrylamide gel (Invitrogen) and transferred to nitrocellulose membrane according to standard protocols. Blots were blocked in a solution of 5% w/v of non-fat milk powder and 0.2% Tween20 in PBS for 1 h then probed with a mouse anti-GFP antibody (Roche) diluted in blocking solution for 1 h. Bound antibody was detected by incubation with a horseradish peroxidase conjugated anti-mouse secondary antibody (Biorad) and visualised using enhanced chemiluminescence western blotting detection reagents (GE Healthcare).

Fluorescence Microscopy
Live synchronized parasite populations expressing GFP-tagged proteins were examined by epifluorescence microscopy on an Axioplan 2 microscope (Zeiss), equipped with a Plan Apochromat 100×/1.4 oil immersion objective and an AxioCam HRc camera [31]. Parasites were labelled with DAPI DNA stain, then a cell suspension was placed on a slide and overlaid with a Vaseline rimmed cover slip. Parasitized erythrocytes were viewed live, and the dual colour fluorescence images were captured using Axiovision 4.6.3 software and edited using Adobe Photoshop CS4.

For indirect immunofluorescence smears of red blood cells infected with P. falciparum 3D7 strain were fixed for 5 min using 100% cold methanol. In the case of haemoglobin labelling, cells were pre-treated with saponin prior to fixation. Cells were washed with PBS, and then permeabilized with 0.1% Triton X100 in PBS for 5 min. After washing with PBS, slides were blocked in PBS containing 3%(w/v) BSA for 1 h at room temperature. The slides were incubated successively for 1 h with primary antibodies, containing 3%(w/v) BSA for 1 h at room temperature. The slides for 5 min. After washing with PBS, slides were blocked in PBS and then permeabilized with 0.1% Triton X100 in PBS for 5 min. After washing with PBS, slides were blocked in PBS containing 3%(w/v) BSA for 1 h at room temperature. The slides were incubated successively for 1 h with primary antibodies, containing 3%(w/v) BSA for 1 h at room temperature. The slides were incubated successively for 1 h with primary antibodies, containing 3%(w/v) BSA for 1 h at room temperature. The slides for 5 min. After washing with PBS, slides were blocked in PBS

Immu-no-electron microscopy
P. falciparum-infected red blood cells (mixed development stages) were fixed and sectioned as described [43]. The sections were immunolabelled with anti-PfRab5B antibody at 1/10 dilution in PBS/1% fish skin gelatin, then incubated with anti-IgG antibodies, followed directly by 10 nm protein A-gold particles before examination with a Philips CM120 Electron Microscope (Eindhoven, the Netherlands) at 80 kV.

Supporting Information
File S1 Supporting Information contains Supplementary Tables S1 and S2 and legends for Supplementary Figures S1 and S2. Table S1: N-terminal sequences of PfRab5B, GAP45, and their variants used in the study. Under “Construct name” are listed all the different wild type and mutant GFP chimera expression plasmids used in the study. The subscript number corresponds to the number of amino acids fused to GFP. Under “Sequence fused to GFP” are the different wild type and mutant amino acids (underlined) sequences fused to the N-terminus of GFP. Table S2: Primers used for generation of the gene-deletion construct and for genotyping parasites. Listed in the number of each primer, its sequence with the restriction site underline, its description and the name of the restriction enzyme. Figure S1: Specificity of anti-PfRab5A and PfRab5B antibodies. Different amounts of purified His-tagged PfRabs recombinant proteins were separated using a 15% polyacrylamide gel and transferred to a nitrocellulose membrane, (A) The blot was first incubated with the rabbit anti-PfRab5A (1:5000) and then incubated with an anti-rabbit peroxidase-conjugated secondary antibody (1:10000, Sigma Aldrich). The lower panel shows protein loading by Ponceau S staining. (B) The blot was first incubated with the rat anti-PfRab5B antibody (1:1000) and then with an anti-rat peroxidase-conjugated secondary antibody (1:4000, Sigma Aldrich). The lower panel shows protein loading by Ponceau S staining. Each anti-PfRab5 antibody specifically reacted only with its corresponding recombinant protein. Figure S2: Unsuccessful attempts to generate P. berghei rab3b (PBANKA_140910) gene-deletion mutants. (A) Schematic representation of the gene-deletion construct used for targeting the rab3b gene for deletion and the expected gene locus before and after disruption. The construct that has bldh as a drug selectable marker (SM, black) is designed to disrupt the open reading frame (ORF) of the Phob3b genes by double crossover homologous recombination. The expected genomic integration of the construct into the P. berghei genome is indicated, and the size and location of the Phob3b targeting regions (hatched boxes) are shown in relation to the Phob3b gene ORF. The targeting regions are indicated as +/- bp distance from the putative start codon. The location and name of the primers used for diagnostic PCR are shown. (B) Diagnostic PCR of genomic DNA of parasites selected after transfection with gene-deletion construct pL1709 (see A) showing that rab3b ORF was not disrupted in the selected parasites. Two independent transfection experiments were performed and the parasites that survived drug selection with pyrimethamine contained both the selectable marker and the intact ORF. The following primers were used: 5' integration (5'): 6909/3189; 3' integration (3') 4592/ 6910; amplification of the bldh cassette (M): 307C/3187; ORF (O): 6911/6912. Genomic DNA of wild type P. berghei parasites

Myristoylated Rab5B of Plasmodium falciparum
Myristoylated Rab5B of Plasmodium falciparum

Author Contributions
Conceived and designed the experiments: GL AAH AM CJJ. Performed the experiments: CNE FBR DKM JWLB EK JLG SMK IC HY. Analyzed the data: GL AAH AM CJJ SMK IC HY. Wrote the paper: GL AAH. Rab5B his-tagged expression construct, CNE, FBR, HY. Rab5B Specific antibodies and imaging, CNE, FBR HY. Rab5B-GFP constructs and imaging, EK JLG. NMT assays: DKM SB. Immuno-ecs IC. Mouse experiments: JWLB SMK CJJ.

References
1. White NJ, Pukrittayakamee S, Hen TT, Faiz MA, Mokashy OA, et al. (2013) Malaria. Lancet 381: 27–38.
2. de Castro FA, Ward GE, Jambou R, Attal G, Mayou A, et al. (1996) Identification of a family of Rab G-proteins in Plasmodium falciparum and a detailed characterisation of prf06. Mol Biochem Parasitol 80: 77–88.
3. Ward GE, Tinley LG, Langley G (1997) Rab GTAPases and the unusual secretory pathway of plasmodium. Parasitol Today 13: 57–62.
4. Langley G, van Noort V, Carret C, Meissner M, de Villiers EP, et al. (2008) Comparative genomics of the Rab protein family in Apicomplexan parasites. Microbes Infect 10: 462–476.
5. Zerial M, McBride H (2000) Rab proteins as membrane organizers. Nat Rev Mol Cell Biol 2: 107–117.
6. Novick PJ, Goud B, Salminen A, Walshworth NC, Nair J, et al. (1998) Regulation of vesicular traffic by a GTP-binding protein on the cytoplasmic surface of secretory vesicles in yeast. Cold Spring Harb Symp Quant Biol 63 Pt 2: 637–647.
7. Grosshans BL, Ortiz D, Novick P (2006) Rab5 and their effectors: achieving complexity in membrane traffic. Proc Natl Acad Sci U S A 103: 11021–11027.
8. Pfeiffer SR (2001) Rab GTAPases: specifying and deciphering organelle identity and function. Trends Cell Biol 11: 487–491.
9. Pfeiffer SR (2001) Vesicle tethering factors united. Mol Cell 8: 729–730.
10. Stenmark H, Olkkonen VM (2001) The Rab GTPase family. Genome Biol 2: 2001109.
11. Chavrier P, Gorvel JP, Stelzer E, Simons K, Gruenberg J, et al. (1991) Comparative genomics of the Rab protein family in Apicomplexan parasites. Mol Biochem Parasitol 47: 57–70.
12. Hirota Y, Kuronita T, Fujita H, Tanaka Y (2007) A role for Rab5 activity in the biogenesis of endosomal and lysosomal compartments. Biochem Biophys Res Commun 356: 40–47.
13. Novick PJ, Chavrier P, Zerial M, Grunenberg J (1991) rab5 controls early endosome fusion in vitro. Cell 66: 915–925.
14. Chen PI, Kong C, Su X, Stahl PD (2009) Rab5 isoforms differentially regulate endosome fusion in vitro. Cell 64: 915–925.
15. Zeigerer A, Gilleron J, Bogorad RL, Marsico G, Nonaka H, et al. (2012) Rab5 is necessary for the biogenesis of the endosomoidal system in vivo. Nature 485: 465–470.
16. Elune K, Fujimoto M, Okatani Y, Nishiyama T, Goh T, et al. (2011) A membrane trafficking pathway regulated by the plant-specific Rab GTAPase ARA56. Nat Cell Biol 13: 833–839.
17. Elune K, Miyakawa N, Fujimoto M, Uemura T, Nakano A, et al. (2012) Endosomal trafficking pathway regulated by ARA56, a RAB5 GTAPase unique to plants. Small GTAPases 3: 23–27.
18. Zeigerer A, Gilleron J, Bogorad RL, Marsico G, Nonaka H, et al. (2012) Rab5 is necessary for the biogenesis of the endosomoidal system in vivo. Nature 485: 465–470.
19. Elune K, Miyakawa N, Fujimoto M, Uemura T, Nakano A, et al. (2012) A membrane trafficking pathway regulated by the plant-specific Rab GTAPase ARA56. Nat Cell Biol 13: 833–839.
20. Elune K, Miyakawa N, Fujimoto M, Uemura T, Nakano A, et al. (2012) Endosomal trafficking pathway regulated by ARA56, a RAB5 GTAPase unique to plants. Small GTAPases 3: 23–27.
21. Ueda T, Yamaguchi M, Uchiyama H, Nakano A (2001) Araf, a plant-unique novel type Rab GTAPase, functions in the endocytic pathway of Arabidopsis thaliana. EMBO J 20: 4730–4741.
22. Chevallier F, Spielmann T, Brabandt D, Yeronimidis I, et al. (2003) The Plasmodium falciparum family of SNAREs. Mol Biochem Parasitol 152: 113–128.
23. Chavrier P, Gorvel JP, Stelzer E, Simons K, Gruenberg J, et al. (1991) Comparative genomics of the Rab protein family in Apicomplexan parasites. Mol Biochem Parasitol 47: 57–70.
24. Agop-Nersesian C, Egarter S, Langley G, Foth BJ, Ferguson DJ, et al. (2010) Biogenesis of the inner membrane complex is dependent on vescular transport by the alveolat specific GTAPase Rab11B. PLoS Pathog 6: e1000109.
25. Kremer K, Kamin D, Wittweger E, Wilkes J, Jammes H, et al. (2013) An overexpression screen of Toxoplasma gondii Rab-GTPases reveals distinct transport routes to the micronemes. PLoS Pathog 9: e1003213.
26. Barik N, Taylor RE, Chakrabarti D (1997) Identification, cloning, and mutational analysis of the casein kinase 1 cDNA of the malaria parasite, Plasmodium falciparum. Stage-specific expression of the gene. J Biol Chem 272: 26132–26138.
27. Donald RG, Zhong T, Meijer L, Liberatore PA (2005) Characterization of two T. gondii CK1 isoforms. Mol Biochem Parasitol 141: 15–27.
28. Gunaratne RS, Sajid M, Ling IT, Tripathi R, Pachelbat JA, et al. (2000) Characterization of N-myristoyltransferase from Plasmodium falciparum. Biochem J 348 Pt 2: 459–463.
29. Rees-Channer RR, Martin SR, Green JL, Bowyer PW, Grainger M, et al. (2006) Dual acylation of the 45 kDa gliding-associated protein (GAP5) in Plasmodium falciparum merozoites. Mol Biochem Parasitol 149: 113–116.
30. Stafford WH, Stockley RW, Landbrook SB, Holder AA (1996) Isolation, expression and characterization of the gene for an ADP-ribosylation factor from the human malaria parasite, Plasmodium falciparum. Eur J Biochem 242: 104–113.
31. Ridzuan MA, Moore RW, Knupfer E, Black S, Holder AA, et al. (2012) Subcellular location, phosphorylation and assembly into the motor complex of GAP55 during Plasmodium falciparum schizont development. PLoS One 7: e38845.
32. Knupfer E, Suleyman O, Dluzsowski AR, Straschil U, O’Keeffe AH, et al. (2013) RON12, a novel Plasmodium-specific rhoptry neck protein important for parasite proliferation. Cell Microbiol.
33. Dluzsowski AR, Ling IT, Hopkins JM, Grainger M, Marges G, et al. (2008) Formation of the food vacuole in Plasmodium falciparum: a potential role for the 19 kDa fragment of merozoite surface protein 1 (MSPI[19]). PLoS One 3: e3085.
34. Jones CJ, Ramesar J, Waters AP (2006) High-efficiency transfection and drug selection of genetically transformed blood stages of the rodent malaria parasite Plasmodium berghei. Nat Protoc 1: 346–356.
35. Wright MH, Clough B, Ruckham MD, Raugachari K, Brannigan JA, et al. (2013) Validation of N-myristoyltransferase as a potential target for drug discovery. Nature Chemistry.
36. Jones ML, Collins MO, Goulding D, Cloughary JS, Rayner JC (2012) Analysis of protein palmitoylation reveals a pervasive role in Plasmodium development and pathogenesis. Cell Host Microbe 12: 246–258.
37. Jones ML, Tay CL, Rayner JC (2012) Getting stuck in: protein palmitoylation in Plasmodium. Trends Parasitol 28: 496–503.
38. Blasink S, Blane M, van der Goes FG (2013) What does S-palmitoylation do to membrane proteins? FERS J 289: 2766–2774.
39. Ayong L, Pagnotti G, Tobon AB, Chakrabarti D (2007) Identification of a family of Rab G-proteins in Plasmodium falciparum. Mol Biochem Parasitol 152: 113–122.
40. Palmis LA, Rayner JC (2009) Plasmodium falciparum secretory pathway: characterization of Pls51, a plasma membrane Qu-SNARE. Mol Biochem Parasitol 164: 153–156.
41. Holder AA (2009) The carbohydrate-terminus of merozoite surface protein 1: structure, specific antibodies and immunity to malaria. Parasitology 136: 1445–1456.
42. Bolte S, Cordelieres FP (2006) A guided tour into subcellular colocalization analysis in light microscopy. J Microsc 224: 213–232.
43. Tomlins AM, Ben-Rached F, Williams RA, Proto WR, Coppens I, et al. (2013) Plasmodium falciparum ATG8 implicated in both autophagy and apicoplast formation. Autophagy 9.