

A *Plasmodium falciparum* C-mannosyltransferase is dispensable for parasite asexual blood stage development

Research Article

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Abstract

C-mannosylation was recently identified in the thrombospondin-related anonymous protein (TRAP) from *Plasmodium falciparum* salivary gland sporozoites. A candidate *P. falciparum* C-mannosyltransferase (*PfDPY-19*) was demonstrated to modify thrombospondin type 1 repeat (TSR) domains *in vitro*, exhibiting a different acceptor specificity than their mammalian counterparts. According to the described minimal acceptor of *PfDPY19*, several TSR domain-containing proteins of *P. falciparum* could be C-mannosylated *in vivo*. However, the relevance of this protein modification for the parasite viability remains unknown. In the present study, we used CRISPR/Cas9 technology to generate a *PfDPY19* null mutant, demonstrating that this glycosyltransferase is not essential for the asexual blood development of the parasite. *PfDPY19* gene disruption was not associated with a growth phenotype, not even under endoplasmic reticulum-stressing conditions that could impair protein folding. The data presented in this work strongly suggest that *PfDPY19* is unlikely to play a critical role in the asexual blood stages of the parasite, at least under *in vitro* conditions.

Introduction

Malaria remains one of the most serious global infectious diseases. Only in 2017, it was responsible for 219 million cases and 435 000 deaths worldwide (World Health Organization, 2018). It is generally accepted that malaria elimination from the remaining endemic countries will not be possible with currently available strategies (Alonso *et al.*, 2011). In this regard, a better understanding of the numerous gaps of knowledge in the complex biology of the parasite would facilitate the rationale design of new interventions.

The thrombospondin type 1 repeat (TSR) superfamily is a diverse group of proteins often involved in interactions with the extracellular matrix (Tucker, 2004; Morahan *et al.*, 2009). Particularly, the functional characterization of *P. falciparum* TSR domain-containing proteins revealed the essential roles that these effectors play in processes involved in cell invasion, egression and transmission of the parasite through the mosquito stages. Thus, circumsporozoite- and TRAP-related protein (CTRP) is essential for ookinete invasion of the mosquito midgut epithelium and oocyst development (Dessens *et al.*, 1999), whereas thrombospondin-related protein 1 is required for the subsequent sporozoite egress from oocysts (Klug and Frischknecht, 2017). TRAP, circumsporozoite protein (CSP) and TRAP-related protein are all involved in the sporozoite invasion of the mosquito salivary glands; in the vertebrate host, CSP and TRAP also bind to the hepatocyte surface during sporozoite invasion (Ménard *et al.*, 1997; Sultan *et al.*, 1997; Combe *et al.*, 2009). Finally, TRAP-like protein and thrombospondin-related sporozoite protein have been described to play an important role in hepatocyte cell traversal (Labaied *et al.*, 2007; Moreira *et al.*, 2008). Other TSR domain-containing proteins are expressed in the blood stages of the parasite. Previously described to be essential for the merozoite invasion of erythrocytes (Baum *et al.*, 2006), more recent studies revealed that the merozoite TRAP family protein (MTRAP) is completely dispensable for the viability of asexual parasites. Instead, MTRAP was demonstrated to be essential for gametocyte egress and, thus, for transmission to the mosquito (Bargieri *et al.*, 2016). On the other hand, the thrombospondin-related apical merozoite protein (TRAMP) seems to be involved in the invasion of erythrocytes, since antibodies raised against this protein block merozoite invasion (Siddiqui *et al.*, 2013). Interestingly, a secreted protein with altered thrombospondin repeat domain (SPATR) is expressed at multiple stages of the parasite such as sporozoites, asexual blood stages and gametocytes. Antibodies raised against SPATR block hepatocyte invasion by sporozoites; however, the biological relevance of SPATR during other stages of the parasite has not been addressed (Chattopadhyay *et al.*, 2003).

The elucidation of the crystal structure of TSR domains revealed an unusual three-stranded fold known as the tryptophan ladder, also found in type 1 cytokine receptors and characterized by a motif of stacked aromatic and basic amino acids stabilized by disulfide bonds (Tan *et al.*, 2002; Olsen and Kragelund, 2014). In other organisms, TSR domains can be modified by two types of glycosylation, namely O-fucosylation and C-mannosylation. The protein O-fucosyltransferase 2 (PoFUT2) catalyses the O-fucose modification of TSR domains, which in turn may be elongated by a β 3-glycosyltransferase (Kozma *et al.*, 2006; Luo *et al.*,

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Table 1. Conservation of the apicomplexan DPY19 C-mannosyltransferase consensus sequence among TSR domain-containing proteins expressed in *P. falciparum*

	Protein	Gene ID	W	X	X	W	X	X	C	*
Sporozoite	TRAP	PF3D7_1335900	W	D	E	W	S	P	C	Y
	TLP	PF3D7_0616500	W	S	E	W	S	P	C	Y
	TRSP	PF3D7_0104000	W	S	E	W	S	A	C	Y
	TRP1	PF3D7_0822700	W	S	E	W	S	T	C	Y
	CSP	PF3D7_0304600	S	T	E	W	S	P	C	N
	TREP	PF3D7_1442600	W	S	E	W	S	G	P	N
Ookinete	CTRP	PF3D7_0315200	W	E	E	W	S	P	C	Y
			F	G	E	W	S	E	C	N
			W	S	D	W	S	S	C	Y
			W	E	E	W	N	E	W	N
			W	S	D	W	S	E	C	Y
			W	E	T	W	V	E	C	Y
Blood	MTRAP	PF3D7_1028700	W	S	E	W	S	A	C	Y
			W	G	E	W	S	E	C	Y
	SPATR	PF3D7_0212600	W	S	D	W	S	A	C	Y
	PTRAMP	PF3D7_1218000	W	G	E	W	S	N	C	Y

CSP, circumsporozoite protein; CTRP, circumsporozoite and TRAP-related protein; MTRAP, merozoite TRAP-like protein; PTRAMP, *Plasmodium* thrombospondin-related apical merozoite protein; SPATR, secreted protein with altered thrombospondin repeat; TLP, TRAP-like protein; TRAP, thrombospondin-related anonymous protein; TREP, TRAP-related protein; TRP1, thrombospondin-related protein 1; TRSP, thrombospondin-related sporozoite protein. Residues of the apicomplexan C-mannosylation consensus sequence are marked in bold, when conserved. *TSR domains with or without a conserved C-mannosylation consensus sequence are marked as Y and N, respectively.

2006). On the other hand, DPY19 enzymes have been identified as the glycosyltransferases responsible for TSR C-mannosylation (Buettner *et al.*, 2013; Shcherbakova *et al.*, 2017). Recent works revealed that TRAP and CSP are both modified *in vivo* by a deoxyhexose and a hexose (presumably forming a glucose- β 1,3-fucose O-glycan) in *P. falciparum* and *P. vivax* salivary gland sporozoites. Additionally, TRAP was also shown to be modified by an additional hexose (presumably C-linked mannose) in *P. falciparum* (Swearingen *et al.*, 2016) but, interestingly, not in *P. vivax* (Swearingen *et al.*, 2017). These, and other works (Sanz *et al.*, 2013; Cova *et al.*, 2015; Bandini *et al.*, 2019), are renewing the interest of the *Plasmodium* research community on the extent and relevance of the parasite glycosylation.

*Pf*PoFUT2 was demonstrated to be dispensable for the asexual and sexual blood stage development of the parasite, in agreement with the absence of predicted acceptors with the TSR O-fucosylation consensus sequence (Lopaticki *et al.*, 2017; Sanz *et al.*, 2019). On the contrary, the relevance of *Pf*PoFUT2 for the establishment of the infection in the mosquito host remains a controversial issue. While Lopaticki *et al.* reported that *Pf*PoFUT2 gene disruption resulted in a deficient ookinete invasion and a reduced sporozoite gliding motility and cell traversal activity, Sanz *et al.* did not observe any evident phenotype secondary to PoFUT2 ablation, neither in *P. falciparum* nor in the murine model *P. berghei*. Interestingly, the phenotype described by Lopaticki *et al.* was attributed to a deficient secretion of certain PoFUT2 acceptors, in accordance with the proposed role of O-fucosylation in a protein folding quality control system (Vasudevan and Haltiwanger, 2014). However, to our knowledge, the relevance of the C-mannosylation of *P. falciparum* TSR-containing proteins has not been addressed before.

*Pf*DPY19 was recently demonstrated to harbour a C-mannosyltransferase activity *in vitro* (Hoppe *et al.*, 2018). Interestingly, *Pf*DPY19 exhibited a different acceptor specificity than the mammalian enzymes (see Table 1 for a list of putative

*Pf*DPY19 acceptors). In other organisms, C-mannosylation has been shown to be required for the efficient secretion of certain acceptors, in accordance with the role played by this modification in the stabilization of the tryptophan ladder (Buettner *et al.*, 2013; Shcherbakova *et al.*, 2017). Taking into consideration the proposed roles for some of the putative acceptors of *Pf*DPY19 in the asexual blood stages, we aimed to describe the relevance of the gene for the viability of *P. falciparum* by disrupting it and analysing the growth phenotype of Δ *Pf*DPY19 mutant parasites.

Material and methods

Construction of plasmids for CRISPR/Cas9-mediated *Pf*DPY19 knockout

A single guide RNA (sgRNA) targeting the *Pf*DPY19 genomic locus (PF3D7_0806200) was designed with the Eukaryotic Pathogen gRNA Design Tool (Peng and Tarleton, 2015). To generate the plasmid expressing the *Streptococcus pyogenes* Cas9 and the sgRNA, the primers 5'-TAAGTATATAATATTTAAATTTAGGCCTTCCATAGTTTATAGAGCTAGAA-3' and 5'-TTC TAGCTCTAAAACATATGGAAAGGCCTAAATTTAAATATTAT-ATACTTA-3' were annealed and ligated into a BbsI-digested pDC2-Cas9-hDHFryFCU plasmid (a generous gift from Ellen Knuepfer) (Knuepfer *et al.*, 2017). To generate the rescue plasmid, two homology regions for *Pf*DPY19 of ~530 bp were amplified from *Pf*3D7 1.2B genomic DNA with the primer pairs 5'-ATTCGAGCTCGGTACCCGGGAAATATGTTGAGCAGA-AACTTTCC-3'/5'-CAGGTCGACTCTAGAGGATCCACGTCG-GAGCCCCAAGAATAATTTCTAAAAACAT-3' and 5'-TTCTT GGGCTCGAGACGTGCGGCCGCATTATATGTTTTTCAGCT-TGTCTTCC-3'/5'-CAGGTCGACTCTAGAGGATCCGCAATA-TGGAATATTACTTCCACAT-3' and were cloned into a pUC19 backbone.

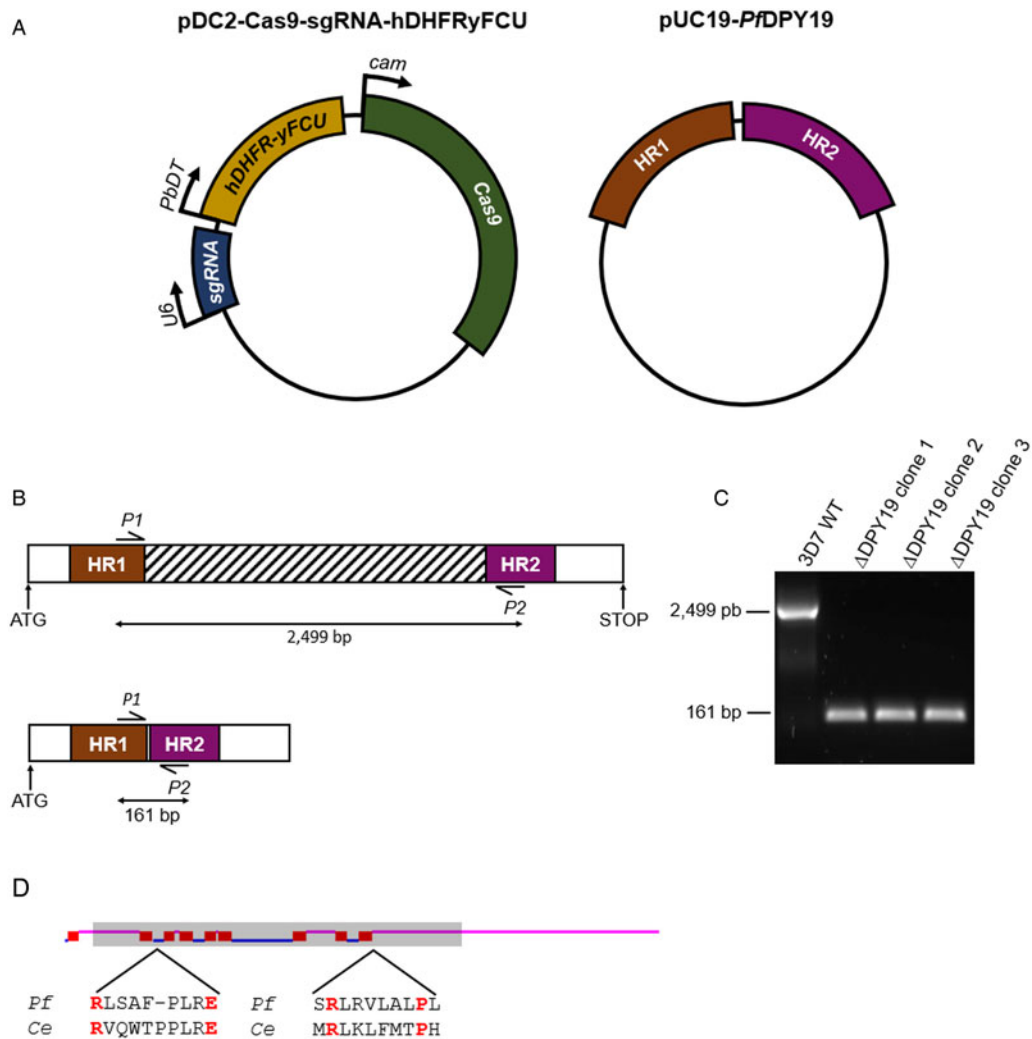


Fig. 1. *PfDPY19* gene disruption. (A) The pDC2-Cas9-sgRNA-hDHFRyFCU expresses the *S. pyogenes* Cas9 endonuclease, the sgRNA targeting the *PfDPY19* genomic locus and a fusion protein of the positive selectable marker human dihydrofolate reductase (hDHFR) and the negative selectable marker yeast cytosine deaminase/uridyl phosphoribosyl transferase (yFCU). The rescue plasmid (pUC19-*PfDPY19*) contains two homology regions of approximately 530 bp corresponding to the 5' and 3' ends of the *PfDPY19* gene (PF3D7_0806200). (B) Schematic representation of *PfDPY19* wild-type (up) and modified loci (down). In total, 2365 bp of the *PfDPY19* coding region (which contain the locus targeted by the designed sgRNA) are excised after double homologous recombination with homology region 1 (HR1) and 2 (HR2) from the rescue plasmid. (C) Polymerase chain reaction (PCR) screening of *PfDPY19* disruption of wild-type parasites (3D7 WT) and three different Δ *PfDPY19* clones obtained by limited dilution. A negative control reaction (NC) was performed in the absence of template DNA to discard unspecific amplifications. (D) Predicted transmembrane helices (red boxes) of *PfDPY19* protein with the TMHMM Server v. 2.0 software (Krogh *et al.*, 2001). Loops predicted to localize in the lumen of the ER and the cytoplasm are marked in pink and blue, respectively. The shaded area indicates the deleted gene fragment in Δ *PfDPY19* mutants. Amino acids predicted to bind to dolichol-phosphate mannose in *C. elegans* (*Ce*) DPY19 are marked in red and are conserved in the *P. falciparum* (*Pf*) homologue (Buettner *et al.*, 2013).

Parasite culture and transfection

Plasmodium falciparum 3D7 1.2B line (kindly provided by Cortés) (Cortés, 2005) was used for transfection and parasite maintenance. *Pf*3D7 1.2B was cultured with B+ human erythrocytes in Roswell Park Memorial Institute (RPMI) medium supplemented with Albumax-II at 37 °C under an atmosphere of 92% N₂, 3% O₂ and 5% CO₂, following standard methods (Trager and Jensen, 1976). For *PfDPY19* gene disruption, 60 µg of pDC2-Cas9-sgRNA-hDHFRyFCU plasmids and 15 µg of ScaI-linearized pUC19-*PfDPY19* plasmids (Fig. 1A) were transfected into Percoll-purified segmented schizonts by electroporation with the Amaxa 4D system, as previously described (Moon *et al.*, 2013). Drug selection (*i.e.* 10 nM WR99210 from Jacobus Pharmaceuticals) was first applied ~20 h post-transfection and maintained for 4 days with daily media changes. The emergence of resistant parasites was monitored by visualizing Giemsa-stained blood smears by light microscopy. Viable parasites were screened by PCR for

PfDPY19 gene disruption and treated for 1 week with 1 µM 5-fluorocytosine for negative selection of parasites containing the pDC2-Cas9-sgRNA-hDHFRyFCU plasmid, prior to sub-cloning by limiting dilution.

In vitro growth assay of parasite asexual development

To assess the biological relevance of *PfDPY19* for the asexual blood stage development of the parasite, the growth of two independent clones of *PfDPY19* null mutants was compared with that of the wild-type parental line in three biological replicates. Cultures were synchronized at ring stages by sorbitol synchronization and parasitaemias were determined by flow cytometry (FACSCalibur, BD Biosciences, Franklin Lakes, NJ, USA) using the Syto 11 dye to discriminate between infected and uninfected erythrocytes, as described elsewhere (Urbán *et al.*, 2011). Cultures were adjusted to ~1.5% parasitaemia and 1% haematocrit. After incubation for one complete intraerythrocytic cycle

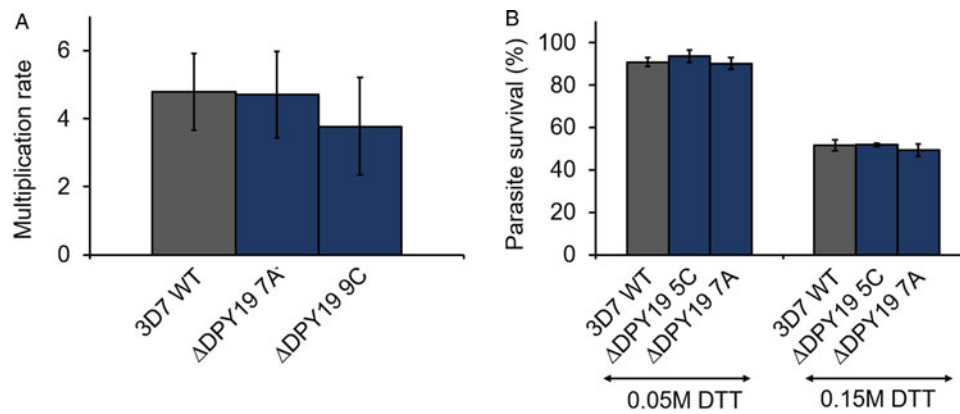


Fig. 2. *PfDPY19* disruption does not alter parasite growth. (A) The growth rate of 3D7 1.2B wild-type (3D7 WT) parasites and *PfDPY19* null mutants (Δ DPY19 7A and 9C clones) was monitored over a complete intraerythrocytic cycle. Values are the mean of three biological replicates and error bars represent the standard variation. (B) Growth inhibition of 3D7 1.2B wild-type (3D7 WT) parasites and *PfDPY19* null mutants (Δ DPY19 5C and 7A clones) at two DTT concentrations (0.05 and 0.15 M). Values are the mean of three technical replicates and error bars represent the standard deviation.

(~48 h), parasitaemias were determined again by flow cytometry. Multiplication rates for each strain were calculated as the ratio between the final and the initial parasitaemia.

Oxidative stress induction and parasite survival assay

To assess the ability of *PfDPY19* null mutants to cope with the endoplasmic reticulum (ER)-stressing conditions, parasite cultures were subjected to different concentrations of dithiothreitol (DTT) as an oxidative stress proxy. Cultures were treated with DTT at a final concentration of 0.05 and 0.15 M or, alternatively, with phosphate buffered saline (PBS) as a negative control. After ~48 h of incubation, parasitaemia was determined by flow cytometry as described previously. The survival rate for each strain was expressed as the ratio between the parasitaemia of the cultures treated with DTT and the negative controls incubated with PBS.

Results

PfDPY19 is not essential for the asexual blood development of *P. falciparum*

To investigate the relevance of the *PfDPY19* C-mannosyltransferase, we disrupted the gene coding for this glycosyltransferase (Fig. 1). As demonstrated by PCR, *PfDPY19* gene was truncated in edited parasites. Remarkably, the deleted gene fragment (see striped box in Fig. 1B) codes for eight out of the nine predicted transmembrane domains of the native protein. Moreover, the deleted fragment also codes for the conserved amino acids that have been predicted to bind to the dolichol-phosphate mannose precursor (Buettner *et al.*, 2013). Given the magnitude of the gene deletion, it is highly unlikely that the generated *PfDPY19* null mutants could yield a functional C-mannosyltransferase activity. Thus, the viability of these mutants strongly suggests that *PfDPY19* is not essential for the asexual blood stage development of the parasite.

PfDPY19 gene disruption does not affect parasite growth

Plasmodium falciparum expresses three TSR domain-containing proteins during the blood stage development of the parasite, all of which contain at least one TSR domain that conserves the apicomplexan C-mannosylation consensus sequence W-X-X-W-X-X-C (see Table 1). Specifically, antibodies raised against TRAMP block the erythrocyte invasion of *P. falciparum*

merozoites (Siddiqui *et al.*, 2013). To assess if *PfDPY19* gene disruption could have an impact on the secretion of TRAMP (or other acceptors) and, consequently, on parasite growth, we compared the multiplication rate of *PfDPY19* null mutants with that of the parental wild-type line (Fig. 2A). However, we did not observe any difference across three biological replicates, suggesting that *PfDPY19* disruption does not affect the parasite's asexual blood stage *in vitro* growth rate.

During the intraerythrocytic development, the parasite is exposed to oxidative stress due to the host immune responses and the products generated during haemoglobin degradation. Thus, the parasite is especially vulnerable to redox imbalances that affect protein folding in the ER (Chaubey *et al.*, 2014). Given the alleged role of C-mannosylation in the stabilization of the tryptophan ladder of TSR domains (Buettner *et al.*, 2013; Shcherbakova *et al.*, 2017), we compared the survival rate of *PfDPY19* null mutants with that of the parental wild-type line when subjected to different concentrations of DTT (Fig. 2B) to assess the impact of the absence of a *PfDPY19* C-mannosyltransferase activity when coping with such conditions. However, no significant differences were observed at any of the DTT concentrations tested.

Discussion

The recent functional characterization of *P. falciparum* PoFUT2 suggests a critical role of O-fucosylation during the infection of the mosquito host (Lopaticki *et al.*, 2017), although these results could not be replicated in an independent study (Sanz *et al.*, 2019). According to the work conducted by Lopaticki *et al.*, PoFUT2 disruption results in a deficient secretion of TRAP, which probably affects the parasite ability to invade hepatocytes. Moreover, it is likely that other acceptors are also affected by the absence of O-fucosyltransferase activity, as evidenced by the deficient invasion of the mosquito midgut epithelium. Although TRAP has also been demonstrated to be C-mannosylated in salivary gland sporozoites (Swearingen *et al.*, 2016), the relevance of this modification remains unknown. The essential role of C-mannosylation for protein secretion observed in other organisms (Buettner *et al.*, 2013; Shcherbakova *et al.*, 2017) prompted us to investigate the contribution of *PfDPY19* to parasite survival and infectivity.

Contrary to O-fucosylation (Lopaticki *et al.*, 2017; Sanz *et al.*, 2019), C-mannosylation is also expected to occur during the blood stages of the parasite, as supported by the conservation of the apicomplexan consensus sequence (Hoppe *et al.*, 2018) in the TSR domain-containing proteins PTRAMP, MTRAP and

SPATR (Table 1). However, the successful generation of a parasite line with a severely truncated *PfDPY19* gene strongly suggests that this modification is not essential for the asexual development of the parasite. Furthermore, parasite growth was not affected in Δ *PfDPY19* mutants, suggesting that the gene is not relevant during the asexual multiplication of the parasite, at least under the tested conditions. This finding is in partial agreement with a previous high-throughput screening study that described the gene coding for *PfDPY19* as dispensable, but conferring significant fitness to *P. falciparum* in the asexual blood stages (Zhang *et al.*, 2018). The discrepancy might arise from slight deviations associated with the comparison of sequencing reads as a proxy for competitive growth fitness in this particular gene or genomic region. The contribution of *PfDPY19* to the secretion of the putative acceptors expressed in the blood stages has not been directly assessed. Nevertheless, if there is an effect on protein secretion secondary to *PfDPY19* gene disruption, it does not seem to affect neither parasite growth nor the parasite's ability to cope with DTT-induced oxidative stress. Contrary to this, a recent work reports that *Toxoplasma gondii* DPY19 may be important for the growth fitness of *T. gondii* tachyzoites, a related apicomplexan (Gas-Pascual *et al.*, 2019).

The O-fucosylation dependence for protein secretion seems to be variable among the different acceptors that are expressed in the mosquito stages of *P. falciparum* (Lopaticki *et al.*, 2017). These findings are in agreement with similar observations made with human TSR domain-containing proteins after the loss of the enzyme responsible for O-fucose elongation with glucose (Vasudevan *et al.*, 2015). Similarly, C-mannosylation seems to be required for the secretion of only a subset of its acceptors in *C. elegans* (Buettner *et al.*, 2013; Shcherbakova *et al.*, 2017). Thus, this modification may be more relevant in other stages of the parasite, where other putative effectors have also been described to play essential roles for parasite viability and infectivity. For instance, given MTRAP essentiality for gametocyte egress (Bargieri *et al.*, 2016), a reduced secretion of this protein may result in a deficient transmission to the mosquito. Likewise, a reduced secretion of CTRP or TRAP as it has been reported after *PfPoFUT2* gene disruption (Lopaticki *et al.*, 2017) may cause a deficient mosquito midgut colonization and hepatocyte invasion, respectively. Furthermore, specific *in vivo* environmental conditions, such as temperature variations or other ER stress inducers, might affect the secretion of different acceptors along the parasite life cycle, including the asexual blood stages. Hence, further studies are required to define the function of *PfDPY19* C-mannosyltransferase in *Plasmodium* parasites.

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Conflict of interest. None.

Ethical standards. Erythrocytes were obtained from the Banc de Sang i Teixits (Catalonia, Spain), after approval from the Comitè Ètic Investigació Clínica Hospital Clínic de Barcelona (Reg. HCB/2017/0413).

References

- Alonso PL, Brown G, Arevalo-Herrera M, Binka F, Chitnis C, Collins F, Doumbo OK, Greenwood B, Hall BF, Levine MM, Mendis K, Newman RD, Plowe CV, Rodríguez MH, Sinden R, Slutsker I and Tanner M (2011) A research agenda to underpin malaria eradication. *PLoS Medicine* **8**, e1000406.
- Bandini G, Albuquerque-Wendt A, Hegermann J, Samuelson J and Routier FH (2019) Protein O- and C-glycosylation pathways in *Toxoplasma gondii* and *Plasmodium falciparum*. *Parasitology*, 1–12. doi: 10.1017/S0031182019000040.
- Bargieri DY, Thiberge S, Tay CL, Carey AF, Rantz A, Hischen F, Lorthois A, Straschil U, Singh P, Singh S, Triglia T, Tsuboi T, Cowman A, Chitnis C, Alano P, Baum J, Pradel G, Lavazec C and Ménard R (2016) Plasmodium merozoite TRAP family protein is essential for vacuole membrane disruption and gamete egress from erythrocytes. *Cell Host and Microbe* **20**, 618–630.
- Baum J, Richard D, Healer J, Rug M, Krnajsiki Z, Gilberger T-W, Green JL, Holder AA and Cowman AF (2006) A conserved molecular motor drives cell invasion and gliding motility across malaria life cycle stages and other apicomplexan parasites. *The Journal of Biological Chemistry* **281**, 5197–5208.
- Buettner FFR, Ashikov A, Tiemann B, Lehle L and Bakker H (2013) *C. elegans* DPY-19 Is a C-mannosyltransferase glycosylating thrombospondin repeats. *Molecular Cell* **50**, 295–302.
- Chattopadhyay R, Rathore D, Fujioka H, Kumar S, de la Vega P, Haynes D, Moch K, Fryauff D, Wang R, Carucci DJ and Hoffman SL (2003) PISPATR, a *Plasmodium falciparum* protein containing an altered thrombospondin type I repeat domain is expressed at several stages of the parasite life cycle and is the target of inhibitory antibodies. *The Journal of Biological Chemistry* **278**, 25977–25981.
- Chaubey S, Grover M and Tatu U (2014) Endoplasmic reticulum stress triggers gametocytogenesis in the malaria parasite. *The Journal of Biological Chemistry* **289**, 16662–16674.
- Combe A, Moreira C, Ackerman S, Thiberge S, Templeton TJ and Menard R (2009) TREP, a novel protein necessary for gliding motility of the malaria sporozoite. *International Journal for Parasitology* **39**, 489–496.
- Cortés A (2005) A chimeric *Plasmodium falciparum* Pfnbp2b/Pfnbp2a gene originated during asexual growth. *International Journal for Parasitology* **35**, 125–130.
- Cova M, Rodrigues JA, Smith TK and Izquierdo L (2015) Sugar activation and glycosylation in Plasmodium. *Malaria Journal* **14**, 427.
- Dessens JT, Beetsma AL, Dimopoulos G, Wengelnik K, Crisanti A, Kafatos FC and Sinden RE (1999) CTRP is essential for mosquito infection by malaria ookinetes. *The EMBO Journal* **18**, 6221–6227.
- Gas-Pascual E, Ichikawa HT, Sheikh MO, Serji MI, Deng B, Mandalasi M, Bandini G, Samuelson J, Wells L and West CM (2019) CRISPR/cas9 and glycomics tools for *Toxoplasma* glycobiology. *The Journal of Biological Chemistry* **294**, 1104–1125.
- Hoppe CM, Albuquerque-Wendt A, Bandini G, Leon DR, Shcherbakova A, Buettner FFR, Izquierdo L, Costello CE, Bakker H and Routier FH (2018) Apicomplexan C-mannosyltransferases modify thrombospondin type I-containing adhesins of the TRAP family. *Glycobiology* **28**, 333–343.
- Klug D and Frischknecht F (2017) Motility precedes egress of malaria parasites from oocysts. *eLife* **6**, 1–32.
- Knuepfer E, Napierkowska M, van Ooij C and Holder AA (2017) Generating conditional gene knockouts in Plasmodium – a toolkit to produce stable DiCre recombinase-expressing parasite lines using CRISPR/Cas9. *Scientific Reports* **7**, 3881.
- Kozma K, Keusch JJ, Hegemann B, Luther KB, Klein D, Hess D, Haltiwanger RS and Hofsteenge J (2006) Identification and characterization of abeta1,3-glycosyltransferase that synthesizes the Glc-beta1,3-Fuc disaccharide on thrombospondin type 1 repeats. *The Journal of Biological Chemistry* **281**, 36742–36751.
- Krogh A, Larsson B, von Heijne G and Sonnhammer EL (2001) Predicting transmembrane protein topology with a hidden Markov model: application to complete genomes. *Journal of Molecular Biology* **305**, 567–580.
- Labaied M, Camargo N and Kappe SHI (2007) Depletion of the *Plasmodium berghei* thrombospondin-related sporozoite protein reveals a role in host cell entry by sporozoites. *Molecular and Biochemical Parasitology* **153**, 158–166.
- Lopaticki S, Yang ASP, John A, Scott NE, Lingford JP, O'Neill MT, Erickson SM, McKenzie NC, Jennison C, Whitehead LW, Douglas DN, Kneteman NM, Goddard-Borger ED and Boddey JA (2017) Protein O-fucosylation in *Plasmodium falciparum* ensures efficient infection of mosquito and vertebrate hosts. *Nature Communications* **8**, 561. doi: 10.1038/s41467-017-00571-y.

- Luo Y, Koles K, Vorndam W, Haltiwanger RS and Panin VM (2006) Protein O-fucosyltransferase 2 adds O-fucose to thrombospondin type 1 repeats. *Journal of Biological Chemistry* **281**, 9393–9399.
- Ménard R, Sultan AA, Cortes C, Altszuler R, van Dijk MR, Janse CJ, Waters AP, Nussenzweig RS and Nussenzweig V (1997) Circumsporozoite protein is required for development of malaria sporozoites in mosquitoes. *Nature* **385**, 336–340.
- Moon RW, Hall J, Rangkuti F, Ho YS, Almond N, Mitchell GH, Pain A, Holder AA and Blackman MJ (2013) Adaptation of the genetically tractable malaria pathogen *Plasmodium knowlesi* to continuous culture in human erythrocytes. *Proceedings of the National Academy of Sciences* **110**, 531–536.
- Morahan BJ, Wang L and Coppel RL (2009) No TRAP, no invasion. *Trends in Parasitology* **25**, 77–84.
- Moreira CK, Templeton TJ, Lavazec C, Hayward RE, Hobbs CV, Kroeze H, Janse CJ, Waters AP, Sinnis P and Coppi A (2008) The Plasmodium TRAP/MIC2 family member, TRAP-Like Protein (TLP), is involved in tissue traversal by sporozoites. *Cellular Microbiology* **10**, 1505–1516.
- Olsen JG and Kragelund BB (2014) Who climbs the tryptophan ladder? On the structure and function of the WSXWS motif in cytokine receptors and thrombospondin repeats. *Cytokine & Growth Factor Reviews* **25**, 337–341.
- Peng D and Tarleton R (2015) EuPaGDT: a web tool tailored to design CRISPR guide RNAs for eukaryotic pathogens. *Microbial Genomics* **1**, e000033.
- Sanz S, Bandini G, Ospina D, Bernabeu M, Marino K, Fernandez-Becerra C and Izquierdo L (2013) Biosynthesis of GDP-fucose and other sugar nucleotides in the blood stages of *Plasmodium falciparum*. *Journal of Biological Chemistry* **288**, 16506–16517.
- Sanz S, Aquilini E, Tweedell RE, Verma G, Hamerly T, Hritzo B, Tripathi A, Machado M, Churcher TS, Rodrigues JA, Izquierdo L and Dinglasan RR (2019) Protein O-fucosyltransferase 2 is not essential for *Plasmodium berghei* development. *Frontiers in Cellular and Infection Microbiology* **9**, 238.
- Shcherbakova A, Tiemann B, Buettner FFR and Bakker H (2017) Distinct C-mannosylation of netrin receptor thrombospondin type 1 repeats by mammalian DPY19L1 and DPY19L3. *Proceedings of the National Academy of Sciences of the USA* **114**, 2574–2579.
- Siddiqui FA, Dhawan S, Singh S, Singh B, Gupta P, Pandey A, Mohammed A, Gaur D and Chitnis CE (2013) A thrombospondin structural repeat containing rhoptry protein from *Plasmodium falciparum* mediates erythrocyte invasion. *Cellular Microbiology* **15**, 1341–1356.
- Sultan AA, Thathy V, Frevert U, Robson KJ, Crisanti A, Nussenzweig V, Nussenzweig RS and Menard R (1997) TRAP is necessary for gliding motility and infectivity of plasmodium sporozoites. *Cell* **90**, 511–522.
- Swearingen KE, Lindner SE, Shi L, Shears MJ, Harupa A, Hopp CS, Vaughan AM, Springer TA, Moritz RL, Kappe SHI and Sinnis P (2016) Interrogating the plasmodium sporozoite surface: identification of surface-exposed proteins and demonstration of glycosylation on CSP and TRAP by mass spectrometry-based proteomics. *PLoS Pathogens* **12**, e1005606.
- Swearingen KE, Lindner SE, Flannery EL, Vaughan AM, Morrison RD, Patrapuvich R, Koepfli C, Muller I, Jex A, Moritz RL, Kappe SHI, Sattabongkot J and Mikolajczak SA (2017) Proteogenomic analysis of the total and surface-exposed proteomes of Plasmodium vivax salivary gland sporozoites. *PLoS Neglected Tropical Diseases* **11**, e0005791.
- Tan K, Duquette M, Liu J, Dong Y, Zhang R, Joachimiak A, Lawler J and Wang J (2002) Crystal structure of the TSP-1 type 1 repeats: a novel layered fold and its biological implication. *The Journal of Cell Biology* **159**, 373–382.
- Trager W and Jensen JB (1976) Human malaria parasites in continuous culture. *Science (New York, NY)* **193**, 673–675.
- Tucker RP (2004) The thrombospondin type 1 repeat superfamily. *The International Journal of Biochemistry & Cell Biology* **36**, 969–974.
- Urbán P, Estelrich J, Cortés A and Fernández-Busquets X (2011) A nanovector with complete discrimination for targeted delivery to *Plasmodium falciparum*-infected versus non-infected red blood cells in vitro. *Journal of Controlled Release* **151**, 202–211.
- Vasudevan D and Haltiwanger RS (2014) Novel roles for O-linked glycans in protein folding. *Glycoconjugate Journal* **31**, 417–426.
- Vasudevan D, Takeuchi H, Johar SS, Majerus E and Haltiwanger RS (2015) Peters plus syndrome mutations disrupt a noncanonical ER quality-control mechanism. *Current Biology* **25**, 286–295.
- World Health Organization (2018) *World Malaria Report 2018*. Geneva.
- Zhang M, Wang C, Otto TD, Oberstaller J, Liao X, Adapa SR, Udenze K, Bronner IF, Casandra D, Mayho M, Brown J, Li S, Swanson J, Rayner JC, Jiang RHY and Adams JH (2018) Uncovering the essential genes of the human malaria parasite *Plasmodium falciparum* by saturation mutagenesis. *Science* **360**, eaap7847. doi: 10.1126/science.aap7847.