Monitoring PfMDR1 transport in *Plasmodium falciparum*

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**Abstract**

**Background:** The *Plasmodium falciparum* multidrug resistance 1 transporter, PfMDR1, contains five amino acid polymorphisms that are suggested to be involved in altered drug transport from the parasite’s cytosol into the digestive vacuole (DV). Transport of a substrate into another intracellular compartment influences drug availability at its site of action, therefore making the parasite more susceptible or resistant to a drug. Fluo-4 is a known fluorescent substrate that can be used as a molecular tool to investigate transport dynamics of PfMDR1 in many parasite strains.

**Methods:** Six *P. falciparum* strains with varying PfMDR1 mutations were loaded with Fluo-4 AM. Accumulation of the fluorophore in the DV was measured using confocal microscopy. The role of a key amino acid mutation was verified using selected parasite clones with point mutations at PfMDR1 amino acid position 1042. Equal expression of PfMDR1 was confirmed by Western blot.

**Results:** Fluo-4 was transported by PfMDR1 into the DV of most drug-sensitive and -resistant parasites. Asparagine at PfMDR1 amino acid position 1042 was crucial for Fluo-4 transport, while the N1042D substitution abolished Fluo-4 transport. Competition studies of Fluo-4 with chloroquine, quinine and mefloquine were performed on parasites harbouring asparagine at position 1042. A distinct Fluo-4 transport inhibition pattern for each tested anti-malarial drug was observed in parasite strains of different genetic background.

**Conclusion:** This study demonstrates that Fluo-4 can be used to investigate PfMDR1 transport dynamics in both drug-sensitive and -resistant parasites. Furthermore, direct evidence of altered Fluo-4 transport in PfMDR1 is linked to a single amino acid mutation in the substrate binding pocket. This system offers a great tool to investigate the role of substrate transport by PfMDR1 and the mutations necessary to support transport, which would lead to new insights for the development of novel anti-malarial drugs.

**Keywords:** *Plasmodium falciparum*, Malaria, Fluo-4, Anti-malarial drugs, Chloroquine, Mefloquine, Quinine, Digestive vacuole, Transport

**Background**

Emerging resistance to commonly used anti-malarial drugs is a major setback in the fight against malaria worldwide [1]. Understanding the molecular mechanisms behind drug resistance is of high importance in ongoing efforts to control this disease.

Early on, researchers found a correlation between antimalarial resistance and the *Plasmodium falciparum* multidrug resistance 1 transporter (PfMDR1) [2, 3]. PfMDR1 is a P-glycoprotein homologue (Pgh1) and belongs to the ATP binding cassette (ABC) transporter superfamily. It is a 162 kDa protein with two nucleotide binding domains (NBD) and twelve transmembrane domains (TMDs), with a putative substrate binding pocket in TMD11 [4]. The transporter is located in the membrane of the digestive vacuole (DV) [5], transporting substrates from the parasite’s cytoplasm into the DV [6].

Two factors have been suggested to play a role in altered drug susceptibility to specific anti-malarial drugs: PfMDR1 polymorphisms and *pfmdr1* gene duplications. Resistance has been associated with one or more variations at five amino acid positions in the *PfMDR1* transporter, with wild-type PfMDR1 containing the amino
acids N86, Y184, S1034, N1042, D1246. Increased pfmdr1 copy numbers have been linked to mefloquine (MQ), lumefantrine (LF), halofantrine (HF), quinine (QN) and artemisinin (AS) resistance [7–9], while PfMDR1 amino acid mutations S1034C, N1042D, D1246Y were found to enhance parasite susceptibility to MQ, HF and AS, independent of gene copy number [2, 10].

Additional evidence for altered drug transport in wild-type versus mutant PfMDR1 was shown through expression of different pfmdr1 variants in Xenopus laevis oocytes. Wild-type PfMDR1 transported QN and chloroquine (CQ) but not HF, while mutant PfMDR1 transported HF but not QN or CQ [11]. Furthermore, residue 184 altered transport kinetics independent of drug binding specificity [12].

Computational models of PfMDR1 describe a substrate-binding pocket that includes the amino acid residues 1034 and 1042 [4, 13]. The binding of several anti-malarial drugs was investigated using docking simulations within the PfMDR1 substrate-binding pocket. Among the tested drugs, MQ was the only candidate whose ability to form an H-bond with residue 1042 was completely abolished through the N1042D substitution [13].

Apart from anti-malarial drug transport, it was shown that several PfMDR1 drug-resistant variants could transport the fluorescent substrate Fluo-4 into the DV of the parasite [6]. In contrast, Fluo-4 transport was abolished in drug-sensitive HB3 parasites harbouring the PfMDR1 variant N86, F184, S1034, D1042, D1246. In a follow-up paper, the pump rate of PfMDR1 (F/Y86, Y184, S1034, N1042, D1246) was determined for Dd2 parasites using live cell imaging of intact infected erythrocytes [14].

In this study, drug-sensitive and -resistant P. falciparum strains of different genetic background and varying PfMDR1 polymorphisms were used to investigate the crucial mutations required for Fluo-4 transport. In addition, P. falciparum clones harbouring a single mutation at position 1042 were analysed. Aspartic acid at position 1042 was found to abolish Fluo-4 transport into the DV. This could be restored by replacing aspartic acid at position 1042 with asparagine. Furthermore, the N1042D substitution resulted in increased sensitivity to MQ. Using Fluo-4 as a competitive substrate offers a powerful tool to investigate the role of PfMDR1 in transport of currently used anti-malarial drugs for both drug-sensitive and -resistant parasites.

Methods
Parasite strains and culture conditions
Three CQ-sensitive (CQS) (3D7, D10, HB3) and three CQ-resistant (CQR) (Dd2, FCR3, FCB) P. falciparum strains, as well as stably transfected P. falciparum clones, derived from the parental lines GC03 and 3BA6, were used in this study. The two strains GC03 and 3BA6 are progeny of the HB3 × Dd2 genetic cross [15] and harbour the PfMDR1 variant from HB3 but differ in their PfCRT phenotype and genotype (Table 1). PfMDR1 mutants derived from GC03 and 3BA6 were produced by Sidhu and colleagues through partial pfmdr1 gene replacement that substituted aspartic acid at position 1042 with asparagine while leaving amino acids at residues 1034 and 1246 unchanged. The resulting clones were: SNDGC03 (S1034, N1042, D1246 in a GC03 genetic background), SND3BA6 (S1034, N1042, D1246 in a 3BA6 genetic background).

Table 1 Main PfCRT and PfMDR1 mutations of Plasmodium falciparum parasites used in this study

| Strains | PICRT | PfMDR1 |
|---------|-------|--------|
| HB3     | K     | N      | F     | S     | D     | D     |
| 3D7     | K     | N      | Y     | S     | N     | D     |
| D10     | K     | N      | Y     | S     | N     | D     |
| Dd2     | T     | Y/F    | Y     | S     | N     | D     |
| FCR3    | T     | Y      | Y     | S     | N     | D     |
| FCB     | T     | Y      | Y     | S     | N     | D     |
| Clones  |       |        |       |       |       |       |
| GC03    | K     | N      | F     | S     | D     | D     |
| SDDGC03 | K     | N      | F     | S     | D     | D     |
| SNDGC03 | K     | N      | F     | S     | N     | D     |
| 3BA6    | T     | N      | F     | S     | D     | D     |
| SDD3BA6 | T     | N      | F     | S     | D     | D     |
| SND3BA6 | T     | N      | F     | S     | N     | D     |
genetic background), as well as the recombinant controls SDDGC05 and SDD3BRA6 [10]. All strains were cultured continuously, as described by Trager and Jensen [16], with modifications. Briefly, parasites were propagated at 5% haematocrit in culture medium containing RPMI 1640 (Life Technologies, Burlington, ON, Canada) supplemented with 25 mM HEPES, 2 mM L-glutamine, gentamicin (20 µg/ml) (Life Technologies, Burlington, ON, Canada), 100 µM hypoxanthine (Sigma-Aldrich, Oakville, ON, Canada), 0.5% AlbuMAX I (Life Technologies, Burlington, ON, Canada). Parasites were maintained at 37°C with an atmosphere of 5% CO2, 3% O2 and 92% N2. A+ red blood cells were obtained from the Interstate Blood Bank (Memphis, TN, USA). Giemsa-stained blood smears were prepared daily to monitor parasite growth. For synchronization, parasites were treated with 5% d-sorbitol (BioShop Canada, Burlington, ON, Canada) for 10 min at 37°C; sorbitol was removed and parasites were washed once before putting them back into culture. To obtain highly synchronous parasite cultures, sorbitol treatment was repeated after 6–8 h.

**DNA isolation and sequence analysis**

The full-length sequence of pfmdr1 and partial sequence of pfcrt was verified for all strains. Parasite strains were grown to ≥5% parasitaemia and DNA was extracted using the QIAamp DNA Blood Mini Kit (Qiagen, Toronto, ON, Canada) according to manufacturer’s instructions. DNA was amplified in overlapping PCR fractions using HotStarTaq DNA polymerase (Qiagen, Toronto, ON, Canada) according to manufacturer’s instructions. To account for the AT-rich nucleotide content in the P. falciparum genome, dNTPs (Invitrogen Canada, Burlington, ON, Canada) were mixed at 75% AT and 25% GC. For PCR optimization, 2 mM MgCl2, 300 µM dNTPs and 300 nM primers were used for the reaction. For each reaction mix, 20 ng genomic DNA was used. PCR reactions consisted of an initial activation step of 94°C for 3 min, followed by 35 cycles of 94°C for 60 s, 52°C for 30 s, and 72°C for 1 min. Primers used for pfmdr1 gene sequencing are described in [14]. Primers used for sequencing the pfcrt gene region containing the main mutation sites were: pfcrt F: 5'-GGAGGTTCTTGCTTGGTAAATG, pfcrt R: 5'-TGAGTTGGAAATAGTTCTTTTAAA. Samples were sent for sequencing to Genome Quebec, Canada and analysed using the BioEdit software [17].

**Protein expression**

To determine PfMDR1 protein levels, 20 µg whole cell protein lysates were loaded in each lane of an 8% acrylamide gel containing SDS. The proteins were transferred onto a PVDF membrane, which was then blocked O/N at 4°C with 5% milk (w/v) and 0.05% Tween-20 (ACP Chemicals, St-Leonard, QC, Canada) in phosphate buffered saline (PBS). The membrane was further incubated with the appropriate dilution of primary anti-PfMDR1 (kindly provided by Prof Cowman, Walter and Eliza Hall Institute, VIC, Australia) or anti-PfHSP70 (GenWay Biotech, San Diego, CA, USA) antibody (1:2,000) in PBS-T + 5% milk for 1 h (RT), then washed and incubated with HRP-conjugated anti-rabbit IgG secondary antibody (Abcam, Toronto, ON, Canada) (1:20,000) for 1 h (RT). Immunoreactive bands were detected with ImmunStar WesternC Chemiluminescent Kit (Bio-Rad Laboratories, Mississauga, ON, Canada) using a myECL Imager (Thermo Scientific, Burlington, ON, Canada). To confirm equal protein loading, chemiluminescence intensities of PfMDR1 were calculated for each parasite clone relative to the respective PHSP70 chemiluminescence using ImageJ 1.47q (National Institutes of Health, USA).

**Growth inhibition assay**

Growth inhibition assays were performed as described previously [18], with modifications. Briefly, synchronized ring stage parasites were diluted to a final parasitaemia of 0.5% and a haematocrit of 2%. A total of 100 µl culture medium per well was prepared in a 96-well plate assay, with a drug dilution series of 1:3, ranging from 1 µM to 0.15 nM. Plates were incubated at 37°C, 5% CO2, 3% O2 and 92% N2 for 72 h, then frozen and stored at −80°C. Plates were thawed at room temperature and 100 µl 2× lysis buffer (20 mM Tris pH 7.5, 5 mM EDTA, 0.008% saponin, 0.08% Triton X-100, 0.2 µl SYBR Green 1/µl) was added to each well. Plates were incubated in the dark for at least 1 h. Fluorescence intensity was determined using a Synergy H4 plate reader (Fisher Scientific, Nepean, ON, Canada) with 485 nm excitation and 520 nm emission wavelengths. IC50 values were determined by fitting concentration response curves with a custom-made procedure for IGOR Pro 6.2 based on an R script kindly provided by Le Nagard [19, 20].

**Live cell imaging**

Synchronized trophozoite stage parasites were loaded with 5 µM Fluo-4 AM (Life Technologies, Burlington, ON, Canada) in Ringer’s solution (122.5 mM NaCl, 5.4 mM KCl, 1.2 mM CaCl2, 0.8 mM MgCl2, 11 mM d-glucose, 10 mM HEPES, 1 mM NaH2PO4, pH 7.4) for 50 min at 37°C. Cells were then washed twice with Ring er’s solution and transferred to a microscope chamber. Parasites were kept at 37°C during microscopy using a stage-top incubator (Tokai Hit, Shizuoaka-ken, Japan). A series of four images per parasite was taken using a Zeiss LSM710 confocal microscope (Carl Zeiss, Oberkochen, Germany) equipped with a water-corrected objective (C-apolochromat 63×/1.20 W Korr M27) and a 488 nm
laser (12.5 mW, 2% intensity). The range of emitted fluorescence was measured from 493 to 622 nm. Images were analysed using ImageJ 1.47q (National Institutes of Health, USA).

Results and discussion

**PfMDR1 transports Fluo-4 into the digestive vacuole**

A genetic linkage between Fluo-4 accumulation in the DV and the PfMDR1 transporter was previously described [6]. Although variations in Fluo-4 accumulation were observed between parasite strains harbouring different PfMDR1 mutations, the amino acid mutation(s) responsible for Fluo-4 transport remained to be determined. To identify PfMDR1 mutation(s) crucial for Fluo-4 transport, several drug-sensitive and -resistant *P. falciparum* strains of different genetic backgrounds were tested for accumulation of Fluo-4 in the DV. Three CQS and three CQR strains harbouring different PfMDR1 mutations were selected for these experiments (Table 1). While PfMDR1 has been suggested to play a role in CQ resistance, the key genetic indicator for CQS versus CQR parasites is attributed to the amino acid mutation K76T in the *P. falciparum* chloroquine resistance transporter (PfCRT) [21]. This mutation was taken into consideration in this study and the relevance of PfMDR1 polymorphisms is discussed for strains harbouring either lysine (K76) or threonine (K76T) in PfCRT.

Of the five mutation sites in PfMDR1 known to play a role in drug resistance, the parasite strains used for these experiments contained mutations at amino acid positions 86, 184 and 1042, which are mainly found in African and Asian isolates [7, 22–24]. To determine if mutations at these residues influence transport of Fluo-4 via PfMDR1, accumulation of Fluo-4 in the DV was measured in live parasites using confocal microscopy. All parasite strains, except HB3, showed high accumulation of Fluo-4 in the DV (Figure 1a, b). Fluo-4 accumulation in the DV was impeded by pre-incubation of the parasites with tariquidar (TQ) (Figure 1b), a specific inhibitor of Pgp transporters [25]. Importantly, Fluo-4 accumulation in the DV was shown to be independent of the PfCRT K76T mutation, suggesting that the PfCRT mutation alone has no effect Fluo-4 accumulation.

**Fluo-4 transport is associated with PfMDR1 N1042**

The parasite strain HB3, which does not accumulate Fluo-4 in the DV, harbours two PfMDR1 polymorphisms—Y184F and N1042D—not found in the other tested *P. falciparum* strains (Table 1), suggesting that one of these two residues is responsible for the altered Fluo-4 transport seen in HB3 parasites. It has previously been demonstrated for the human homolog of PfMDR1 that the Y184F mutation does not influence drug specificity [12]. Furthermore, this residue only mildly increased drug resistance in comparison to other PfMDR1 mutation sites [4]. In contrast, amino acid position 1042 is thought to be part of the transporter binding pocket and forms electrostatic interactions with amodiaquine, CQ, MQ (only in the presence of asparagine), HF, vinblastine and vincristine [4, 10, 13]. Therefore, the role of PfMDR1 mutations at position 1042 was investigated in more detail.

To verify the influence of PfCRT on Fluo-4 transport, existing parasite clones containing either lysine or threonine at residue 76 in PfCRT were selected and compared. For this, stably transfected *P. falciparum* clones derived from the parental lines GC03 and 3BA6 were used, where GC03 harbours the K76 PfCRT genotype and is CQ sensitive, and 3BA6 contains the PfCRT K76T mutation that confers CQ resistance (Table 1). Both GC03 and 3BA6 contain the PfMDR1 sequence N86, F184, S1034, D1042, D1246, identical to the HB3 strain. Experiments verified that neither GC03 nor 3BA6 were able to accumulate Fluo-4 in the DV, as expected (Figure 2a, b).

As a control, parasite clones SDDGC03 and SSD3BA6, harbouring the same mutation as GC03 and 3BA6, revealed that the PfMDR1 N1042D mutation did not transport Fluo-4 into the DV (Figure 2a, b). Fluo-4 transport was only established with the introduction of asparagine at residue 1042 for both clones (SNDGC03 and SND3BA6). Transport was independent of the PfCRT K76T mutation, since Fluo-4 transport was equally impaired in GC03 and 3BA6 parental lines as well as the SDDGC03 and SSD3BA6 controls, and only seen in the clones SNDGC03 and SND3BA6 (Figure 2a, b). Transport specificity for PfMDR1 in clones SNDGC03 and SND3BA6 was confirmed through pre-incubation of the samples with the Pgp inhibitor TQ (Figure 2b; Additional file 1: Figure S1). These results indicate a pivotal role for asparagine at PfMDR1 residue 1042 in Fluo-4 transport, independent of PfCRT.

In addition to PfMDR1 polymorphisms, increased *pfmdr1* gene copy number [8, 9, 26] and protein expression levels have also been suggested to play a role in drug resistance or susceptibility. To verify that Fluo-4 accumulation in the DV of clones SNDGC03 and SND3BA6 was not linked to increased PfMDR1 expression, PfMDR1 protein levels were analysed for these parasite clones (Figure 2c). No significant increase in PfMDR1 expression was detected for the GC03 and 3BA6 parental lines or their clones harbouring the D1042N substitution (p > 0.05). Therefore, Fluo-4 accumulation in the DV of these clones was only associated with the PfMDR1 mutation at residue 1042.
Since transport of Fluo-4 was dependent on specific PfMDR1 polymorphisms, it follows that the N1042D substitution in the PfMDR1 substrate-binding pocket may not only alter the binding and transport of Fluo-4 but also influence other substrates, such as anti-malarial drugs. To determine if the amino acid substitution at residue 1042 resulted in altered drug sensitivity or resistance of the parasite clones, growth inhibition assays were performed. A change in a given drug’s IC$_{50}$ would indicate a role for PfMDR1 amino acid position 1042 in its transport. CQ IC$_{50}$ values for CQS clones were found to
be 47 ± 2.6 nM for GC03, 46 ± 2.2 nM for SDDGC03 and 49 ± 2.8 nM for SNDGC03, while CQ IC50 values for CQR clones were 262 ± 5.7 nM for 3BA6, 204 ± 9.4 nM for SDD3BA6 and 274 ± 5.6 nM for SND3BA6 (Table 2). This suggests that an amino acid mutation at PfMDR1 residue 1042 does not affect susceptibility or resistance to CQ. CQ resistance could be reversed in the CQR strains through the addition of 1 µM verapamil, while no significant difference was observed in CQS strains (p > 0.05), as expected. Similarly, CQS clones were more susceptible to quinacrine (QC) with IC50 values of 11 ± 0.3 nM for GC03, 13 ± 0.3 nM for SDDGC03 and 14 ± 0.4 nM for SNDGC03 compared to the higher QC IC50 values of 43 ± 3.4 nM for 3BA6, 41 ± 1.0 nM for SDD3BA6 and 42 ± 2.3 nM for SND3BA6. Here again, as for CQ, an amino acid mutation at PfMDR1 residue 1042 did not affect susceptibility or resistance to QC. The same was found for dihydroartemisinin (DHA). CQS clones were fourfold more resistant than CQR clones (4 ± 0.1 vs 1 ± 0.1 nM) and DHA resistance was not reversible by
the addition of 1µM verapamil. Therefore, the PfMDR1 N1042D mutation alone does not appear to be a major determinant for CQ, QC or DHA resistance in these parasites (Table 2).

Interestingly, the N1042D mutation did provide changes in drug susceptibility for the anti-malarial drugs QN and MQ. A significant decrease in QN IC50 values was observed in the GC03 and 3BA6-derived clones SNDGC03 and SND3BA6 compared to the parental lines GC03 and 3BA6 or their recombinant controls SDDGC03 and SDD3BA6 (p < 0.005) (Table 2). The opposite effect was observed for MQ. For the clones SNDGC03 and SND3BA6, the D1042N substitution led to an approximate threefold increase in resistance to MQ. MQ IC50 values increased from 6±0.9 nM in GC03 to 19±0.6 nM in SNDGC03, and from 5±0.3 nM in 3BA6 to 14±0.5 nM in SND3BA6. This was a significant increase in MQ IC50 values for the clones SNDGC03 and SND3BA6 compared to the parental lines (p = 0.0003 for GC03 vs SNDGC03; p = 0.0001 for 3BA6 vs SND3BA6). Similar increases in MQ resistance (approximately threefold) have been found in field isolates from Africa and Asia [27, 28]. A threefold increase in MQ resistance was also confirmed in vitro through allelic exchange of pfmdr1 [2]. Therefore, the PfMDR1 mutation crucial for MQ resistance in vitro is likely to play a role in increased MQ resistance in the field. Nevertheless, growth inhibition experiments do not fully elucidate altered transport of MQ by PfMDR1. For this reason, co-incubation of anti-malarial drugs with Fluo-4 can provide additional insight into the role of PfMDR1 transport in drug resistance.

Table 2 IC50 values of Plasmodium falciparum parasites used in this study

|          | CQ   | CQ + VP | DHA | DHA + VP | QC   | QC + VP | MN  | MQ  |
|----------|------|---------|-----|----------|------|---------|-----|-----|
| 3D7      | 24±5.6 | 17±1.2  | n.d.| n.d.     | n.d. | n.d.    | 23±1.8 | 8±1.2 |
| Dd2      | 169±3.9| 53±6.5  | n.d.| n.d.     | n.d. | n.d.    | 283±39.8 | 27±40.0 |
| GC03     | 47±2.6 | 46±2.1  | 4±0.1 | 4±0.1 | 11±0.3 | 18±0.4 | 15±0.7 | 76±4.9 | 7±0.6 |
| SDDGC03  | 46±2.2 | 47±0.6  | 4±0.1 | 3±0.2 | 13±0.3 | 15±0.7 | 52±3.5 | 19±0.6 |
| SNDGC03  | 49±2.8 | 47±2.2  | 4±0.1 | 3±0.1 | 14±0.4 | 17±0.3 | 32±1.8 | 5±0.3 |
| 3BA6     | 262±5.7| 41±1.8  | 1±0.1 | 1±0.1 | 43±3.4 | 47±2.2 | 83±4.8 | 5±0.3 |
| SDD3BA6  | 204±9.4| 63±1.5  | 1±0.0 | 1±0.1 | 41±1.0 | 39±0.5 | 78±5.3 | 6±0.7 |
| SND3BA6  | 274±5.6| 69±5.6  | 1±0.1 | 1±0.0 | 42±2.3 | 39±0.4 | 48±2.4 | 14±0.5 |

All values are given in nM. Experiments were done in triplicate on three separate days. CQ chloroquine, VP verapamil, DHA dihydroartemisinin, QC quinacrine, QN quinine, MQ mefloquine, n.d. not determined.

Further insight into the importance of PfMDR1 in drug transport can be achieved through competition studies using two potentially competitive substrates, e.g., fixed amounts of Fluo-4 and increasing drug concentrations. For this purpose, two parasite strains were chosen to evaluate potential differences in anti-malarial drug transport in sensitive (3D7) and resistant (Dd2) parasites. 3D7 is sensitive to CQ, QC and MQ (Table 2), while Dd2 is resistant to these drugs. HB3 was not used, since it is not able to transport Flu-4 into the DV. The measured changes of Flu-4 accumulation in the DV in the presence of anti-malarial drugs were compared for 3D7 and Dd2 parasites.

For 3D7 parasites, increasing CQ concentration led to the strong decrease in Flu-4 accumulation in the DV (Figure 3a). While TQ is a non-competitive inhibitor of MDR1 through inhibition of the ATPase activity [25], CQ is thought to bind to the transporter’s substrate-binding pocket and is likely a competitive inhibitor of Flu-4. Pre-incubation of 3D7 parasites with 250 nM CQ resulted in decreased Flu-4 accumulation in the DV, where only 5±1.0% Flu-4 fluorescence was measured when compared to non drug-treated parasites. In Dd2 parasites, pre-incubation with 250 nM CQ reduced Flu-4...
Figure 3  Competition of Fluo-4 transport with anti-malarial drugs. a Parasites were pre-incubated with different concentrations of chloroquine (CQ), mefloquine (MQ) or quinine (QN), or left untreated before adding Fluo-4 AM. b Parasites were pre-incubated as described with 250 nM of CQ, MQ or QN. Experiments were done in triplicate 3 independent days. Error bars represent SEM. *P < 0.05; ***P < 0.0001.
fluorescence in the DV by half (48 ± 3.0%). MQ reduced Fluo-4 transport into the DV to 45 ± 2.0% in 3D7 parasites and only 82 ± 6.0% in Dd2 parasites (Figure 3a). For MQ, a decrease in Fluo-4 transport in 3D7 compared to Dd2 parasites cannot be attributed to a PfMDR1 amino acid mutation at position 1042 alone since both strains harbour wild-type N1042. However, Dd2 parasites harbour an amino acid mutation at PfMDR1 residue 86, which was described to influence MQ sensitivity in African field isolates [24, 30, 31]. This N86Y mutation may alter MQ transport by PfMDR1, as shown in the experiments presented here. QN was less effective in reducing Fluo-4 accumulation than CQ or MQ and only decreased Fluo-4 fluorescence in the DV to 59 ± 2.9% in 3D7 parasites and not significantly in Dd2 parasites (92 ± 3.4%) (Figure 3a). The differences between 3D7 and Dd2 parasites at the level of Fluo-4 reduction may be linked to PfMDR1 residue 86, which is suggested to influence CQ susceptibility [29]. It is conceivable that N86Y can allosterically influence the drug binding site at TMD11, as suggested in [4]. Residue 86 was not examined in this study.

Fluo-4 competition with anti-malarial drugs was also tested on the parasite clones SNDGC03 and SNDJRA6. Using a single drug concentration of 250 nM, Fluo-4 accumulation was decreased in the DV of SNDGC03 and SNDJRA6 parasites to 33 ± 1.9% and 39 ± 3.0% for CQ, 61 ± 3.1% and 65 ± 4.8% for MQ, and 58 ± 3.9% and 91 ± 4.4% for QN, respectively. Almost all tested anti-malarial drugs decreased Fluo-4 accumulation in the DV in both SNDGC03 and SNDJRA6 clones, except for QN (Figure 3b). This again suggests that the influence of anti-malarial drugs on substrate transport by PfMDR1 is not only dependent on PfMDR1 polymorphisms but also on the varying genetic background of the parasite strain.

The influence of the N1042D substitution on Fluo-4 transport is explained in a proposed model (Figure 4). Fluo-4 is negatively charged and replacement of a neutral asparagine at residue 1042 with a negatively charged aspartic acid alters the local environment, making it less favourable to Fluo-4 transport. TQ does not influence Fluo-4 affinity for the substrate-binding pocket but prevents adenosine triphosphate (ATP) hydrolysis of PfMDR1. Since both NBDs act in concert [32], complete inhibition of ATPase activity by TQ can be achieved when one nucleotide-binding site is blocked.

Detailed transport kinetics for parasite strains of different origin will provide additional information on substrate transport via PfMDR1. Using Fluo-4 as a competitive substrate circumvents the issue of direct labeling of anti-malarial drugs with a fluorescent tag that may alter their transport properties. Therefore, Fluo-4 is a powerful tool to selectively study transport kinetics of PfMDR1 in intact P. falciparum-infected erythrocytes.

**Conclusion**

This study provides direct evidence for reduced PfMDR1-driven substrate transport through the single amino acid mutation N1042D, located in the substrate-binding pocket. The relevance of this amino acid...
mutation may have been underestimated and needs further investigation. Furthermore, it is now possible to test additional mutations within the binding pocket of PfMDR1 to verify their potential role in drug transport. Accordingly, using Fluo-4 in drug competition assays provides a powerful tool to better examine drug transport kinetics via PfMDR1. This newly acquired information may help elucidate the role of PfMDR1 in drug transport, which has remained controversial for decades. For example, while earlier investigations have suggested a link between MQ resistance and asparaginase at PfMDR1 amino acid position 86 in isolates from Thailand and West Africa [30, 33], others have associated wild-type PfMDR1 and increased gene copy numbers with MQ resistance [3, 34]. More recent publications have found strong evidence for a role for N1042 in MQ resistance in Thai isolates [35, 36], suggesting that the importance of this mutation has been underestimated in previous field studies. The importance of N1042 for MQ resistance was similarly demonstrated in vitro through combined amino acid substitutions that generated a parasite strain harbouring the mutations S1042C, N1042D, D1246Y [2]. This newly generated strain was approximately threefold more susceptible to MQ exposure compared to the parental strain [2]. The results presented here support a potential role of PfMDR1 N1042 in parasite resistance to MQ. Further investigations will help elucidate the significance of this polymorphism on substrate transport.

Additional file

**Additional file 1: Figure S1.** Inhibition of Fluo-4 transport through tariquidar. *Plasmodium falciparum* clones were pre-incubated with 100 nM tariquidar for 10 min at 37°C, then 5 µM Fluo-4 AM was added for 50 min. Fluo-4 accumulation in the digestive vacuole was diminished. Experiments were done in triplicate on different days. These images are supplementary to the data obtained for Figure 2. Scale bar, 5 µm.

Authors’ contributions

SJR and PR designed the study; SJR performed experiments and quantified the data; SJR and PR wrote the manuscript. Both authors read and approved the final manuscript.

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Compliance with ethical guidelines

Competing interests

The authors declare that they have no competing interests.

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