Introduction of F446I mutation in the K13 propeller gene leads to increased ring survival rates in *Plasmodium falciparum* isolates

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**Abstract**

**Background:** Mutations in the *Plasmodium falciparum* k13 gene are associated with artemisinin (ART) resistance. However, it is unclear whether the F446I mutation, the most prevalent allele at the China–Myanmar border and north of Myanmar, is associated with ART resistance. Therefore, the aim of this study was to investigate the role of this mutation in ART resistance by generating transgenic parasites expressing the F446I mutant allele.

**Methods:** The transgenic parasites carrying the F446I or C580Y mutation in both 3D7 and FCC1/HN isolates were generated by single crossing-over recombination and verified using PCR and gene sequencing. The ring-stage survival assay of 0–3 h (RSA0–3 h) was used to evaluate ART susceptibility of the transgenic parasites in vitro.

**Results:** Four transgenic parasite lines named 3D7F446I mut, 3D7C580Y mut, FCC1/HNF446I mut and FCC1/HNC580Y mut were successfully generated. These parasite lines showed no changes in the expression level of k13 when compared with their parent parasite isolates. However, introduction of the F446I mutation in k13 of the 3D7 and FCC1/HN isolates led to elevated ring survival rates detected using RSA0–3 h when subjected to both 700 and 20 nM concentrations of dihydroartemisinin. The survival rates were similar to those detected in the parasite lines with the C580Y mutation.

**Conclusions:** Insertion of the F446I mutation in k13 led to increased ring survival, suggesting that this mutation may be associated with ART resistance and could be used as a molecular marker for monitoring ART-resistant parasites. The results also highlights the importance of surveillance of F446I mutants for containing the resistant parasite.

**Keywords:** *Plasmodium falciparum*, K13 propeller gene, F446I mutation, Artemisinin resistance

**Background**

Artemisinin (ART) is a quick, effective anti-malarial drug that was discovered in China [1, 2]. Since ART-based combination therapies were recommended by the World Health Organization (WHO) as a first-line treatment of malaria in 2005, the global malaria burden has significantly declined [3]. However, in 2008, ART resistance was reported in *Plasmodium falciparum* parasites in western Cambodia; a delay in clearance time was observed in infected patients who had received the standard 3-day treatment [4, 5]. The resistance has spread across the mainland of Southeast Asia, including Cambodia, Thailand, Vietnam, Myanmar, and China–Myanmar border areas [6, 7]. The spread of ART resistance, particularly in Africa, will certainly threaten global malaria control programmes. Therefore, efficient tools, such as molecular markers, for the detection and surveillance of ART-resistant parasites are urgently required to contain the spread of the resistance.

Currently, tools available for the detection and surveillance of ART-resistant malaria parasites include measurement of in vivo parasite clearance rates and ring-stage survival assay of 0–3 h (RSA0–3 h) that is well correlated with in vivo parasite clearance rates [8]. The K13 propeller gene of *P. falciparum* (k13; PF3D7_1343700) has been identified as a molecular marker for monitoring ART-resistant parasites [9]. A single mutation after position

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440 in \( k13 \) is associated with slow parasite clearance rates in vivo and high parasite survival rates in vitro \([7, 10]\). The prevalent mutant alleles of \( k13 \) in the Thai–Cambodian/Thai–Myanmar border areas are C580Y, Y493H, R539T, and I543T, while that in the northern Myanmar and China–Myanmar border areas is F446I \([7, 9, 11]\). Mutant alleles such as C580Y and R539T have been confirmed to be correlated to ART resistance characterized by long in vivo parasite clearance half-lives after ART treatment and high RSA\(_{0.3\, h}\) survival rates \([9, 12]\). In addition, the association of the C580Y mutation with ART resistance was also confirmed through introduction of the mutation into \( k13 \) \([13]\). However, F446I is the most prevalent mutant allele of \( k13 \) at the China–Myanmar border and in northern Myanmar. Previous study reported that of 392 isolates from the China–Myanmar border, 128 had the F446I mutation with a prevalence of 32.7% \([11]\). Two other studies showed the F446I mutation with a prevalence of 36.5 and 27.2% in this area \([14, 15]\). It is unclear whether this mutation is associated with ART resistance. Therefore, homology recombinant technology was used to introduce the F446I mutation in \( k13 \) gene of 2 distinct laboratory strains, 3D7 and FCC1/HN, of \( P. falciparum \) and investigate the impact of the mutation on ART susceptibility.

**Methods**

**Parasite culture**

The parasite culture was maintained in vitro in RPMI 1640 (Sigma-Aldrich, USA) with 10 mM HEPES (Invitrogen, California), 20 \( \mu \)g/ml hypoxanthine (Sigma-Aldrich, USA), 50 \( \mu \)g/ml gentamicin (Sigma-Aldrich, USA), 0.17% NaHCO\(_3\) (Sigma-Aldrich, USA), and 5 mg/ml AlbumaxII (Gibco, USA). *Plasmodium falciparum* lines were propagated in human O\(^+\) erythrocytes (obtained from Shanghai Hospital) and maintained at 37 °C in 5% CO\(_2\) and 90% N\(_2\). The parasites were synchronized using 5\% d-sorbitol, as described previously \([16]\). Briefly, the cultured parasites were pelleted and treated with sorbitol at room temperature for 15 min and washed with non-serum RPMI1640 medium and maintained in standard culture medium.

**Construction of the pARL-PfK13\(^{F446I}\) and pARL-PfK13\(^{C580Y}\) recombinant plasmids**

The pARL-GFP plasmid containing the drug screening gene hDHFR was used to construct the pARL-PfK13\(^{F446I}\) and pARL-PfK13\(^{C580Y}\) recombinant plasmid. The full-length \( k13 \) gene with F446I mutation and 1563 bp of the C-terminal of \( k13 \) gene was amplified using overlapping PCR. The primers for amplifying fragments with F446I were as follows: 1350 bp 5′-terminal fragment was amplified by primer pair of PARL-F (GCagatctATGGAA GGAGAAAAAGTAAAAACAAAAAG) and K13-446-R (TCCACCTTACAAATTACTAATGGG); while 831 bp 3′-terminal fragment was amplified by primer pair of K13-446-F (CCCCCCATTGTTTGTAGTGGGA) and PARL-R (GGcgttacCTATAATTGCTATTTAAAC GGAATG). The two fragments were combined by PCR through overlapping sequences to generate the full length \( k13 \) gene with F446I mutation by primer pair of PARL-F and PARL-R. Similar strategy was used to generate the 1563 fragment of \( k13 \) gene with C580Y mutation: the 1128 bp upstream fragment was amplified by primer pair of PARL-A-F (GCGagtctAACGAAATTAAGTAGTGC TAGTG) and K13-580-R (AGCCACATACTAGCTGA TGA) while the 435 bp downstream fragment by K13- 580-F (GCTATGATGTGCTTTTTGTAG) and PARL-R. The resulting fragments were cloned into the pARL-GFP plasmid by using the restriction sites BglII and KpnI (Fermentas, USA) to generate recombinant plasmids of pARL-PfK13\(^{F446I}\) and pARL-PfK13\(^{C580Y}\). The recombinant plasmids were purified using the Endo Free Plasmid Maxi Kit (Qiagen, Germany).

**Parasite transfection**

Synchronous *P. falciparum* 3D7 and FCC1/HN (isolated from Hainan Province and established for in vitro continuous cultivation in 1978) parasites were transfected with the recombinant plasmids as described previously \([17]\). Briefly, *P. falciparum* parasite culture with a high synchronous ring stage was prepared at 5% parasitaemia. 270 \( \mu \)l Cytomix buffer (120 mM KCl, 10 mM KH\(_2\)PO\(_4\), 25 mM HEPES, 2 mM EGTA, 0.15 mM CaCl\(_2\), and 5 mM MgCl\(_2\)) was mixed with 100 \( \mu \)g of the plasmid, and then with 50% haematocrit of the infected erythrocytes. The mixture was transferred to a 0.2-cm cuvette (Bio-Rad) and electroporated using the Bio-Rad MicroPulser with voltage 310 V and 950 \( \mu \)F. The transfected culture was selected using 2.5 nM WR99210 (Jacobus Pharmaceuticals) after 48 h until the WR99210-resistant parasites were obtained. DNA was extracted using QIAmp Mini kits (Qiagen, Germany) and detected by PCR using primers hDHFR, P3/P4, P4/P5, and P3/P6. The primers for identification of transfected parasites were as follows: hDHFR-F: TTAATGTATATTCCAATGTGCA TAT, hDHFR-R: GTACCTTATGCCCTTCTCTCTCTG; P3: ACCATGATTAGGCCAACG; P4: AAAACGGG ATTAACGTGCC; P5: AACAAAAAGGATATAT TCG TG; P6: ATAAAAGACATTTAAAAATTCTTCTCATT. The sequences of the modified \( k13 \) gene were confirmed by DNA sequencing. RNA extracted from the F446I mutant parasite was digested by DNaseI (Sigma-Aldrich, USA) for 15 min in room temperature before transcribed to cDNA. The full-length of \( k13 \) gene was amplified from the cDNA of F446I mutant parasite and subcloned to
pEASY-T1 cloning Vector (purchased from Transgen Biotech, China). The primers for amplification were as follows: K13-F: ATGGAGGCAGAAAAATGAAAAACA AAAG; K13-R: TTATATTTGTCTATTTAAAAGCCA GTG. The RNA of F446I mutant parasite after digestion was used as template to amplified k13 gene as a no-RT control. The k13 gene in the pEASY-T1-K13 recombinant plasmid was sequenced using the primers of M13 as follows: M13F-GTAAAAACGACGGCCAGT; M13R-CAG GAAACAGCTATGAC.

Antibody and western blot
The antibody against k13 protein was produced by GenScript Corporation (USA). Briefly, a synthetic peptide (MEGEKVKTKANSISC) was conjugated with keyhole limpet haemocyanin and formulated with adjuvant. Rabbits were immunized for 4 times at 2-week intervals. The polyclonal antibodies were affinity-purified with peptides.

Western blot was performed as described previously [18]. Briefly, the parasite culture was pelleted and resuspended with 10 volumes of phosphate-buffered saline (PBS). Next, 3% saponin was added to lyse the erythrocytes at 0.03% final concentration on ice for 15 min, and the mixture was centrifuged. The pellets were washed 2 times with PBS and resuspended with a suitable volume of buffer A (20 mM HEPES, 10 mM KCl, 1.5 mM MgCl₂, 1 mM EDTA, 1 mM EGTA, 50 mM NaCl, 0.65% NP-40, 1 mM DTT, and 1× protease inhibitor cocktail). The mixture was incubated for 1 h at 4 °C and centrifuged at 14,000 rpm to obtain the supernatant. The supernatant was separated by 10% SDS-PAGE gel and detected using Pk13 polyclonal antibody (1:200). The Actin (Abcam ab40854) was used as the loading control (1:1000). The secondary antibody was IRDye800CW goat anti-rabbit (LI-COR) diluted at 1:5000.

RSA₀₃₉ assay
RSA₀₃₉ was performed as described previously [8]. Briefly, P. falciparum parasites were synchronized 2 times with 5% sorbitol to obtain tightly synchronous early-ring stage. The initial culture was prepared at 1% parasitaemia and 2% haematocrit, and then was exposed to 700 or 20 nM DHA (Sigma-Aldrich, USA) for 6 h. Then, the culture mixture was washed 3 times with non-serum 1640 culture medium, and cultured again for 66 h. The viable parasites were counted using a microscope in Giemsa-stained thin smears. The survival rate was calculated using Ps/Pc × 100%, where Ps is the parasitaemia in the DHA treatment group, while Pc is the parasitaemia in the control group. The counting was performed by two different experienced laboratories. Each assay was repeated for 3 times independently, with each group setting three technical repetitions. Each technical repetition was counted for 40,000 erythrocytes.

Results
Generation of constructs for transfection of the parasites
To generate the recombinant plasmids for the transfection, the fragments of k13 gene were amplified with either F446I or C580Y mutations by using PCR. For construction of the replacement F446I, 2 DNA fragments covering the entire k13 gene (2181 bp) were amplified from the genomic DNA of P. falciparum. The two fragments were combined using PCR with an overlapping sequence of 15 bp in which a point mutation of T to A was introduced, resulting in changes to the corresponding amino acid residue from phenylalanine (F) to isoleucine (I) (designated as Pfk13F446I mut gene). The full-length Pfk13F446I mut gene contains the 5′ cleavage site of KpnI and 3′ cleavage site of BglII, which allowed cloning of the entire gene into the vector pARL-GFP through corresponding sites in the vector and generation of the recombinant plasmid designated as pARL-K13F446I. The substitution of T to A in the Pfk13F446I mut gene was verified by DNA sequencing. The overall strategy for the preparation of Pfk13F446I mut is outlined in Fig. 1a.

A similar strategy was used for the generation of the Pfk13C580Y mut gene. However, in this case, only a truncated k13 gene (1563 bp) was amplified instead of the full gene. This is because of the position of the C580Y mutation near the C-termini and that only a partial sequence of the gene is needed for single crossing-over recombination. Similarly, two fragments covering the truncated k13 gene were amplified and combined using PCR. A point mutation of G to A at nucleotide position 1739 was introduced through the overlapped neighbouring sequences of the 2 fragments, resulting in a change from cysteine (C) to tyrosine (Y) (designated as Pfk13C580Y gene).

Generation of transgenic parasites
Two distinct laboratory strains of P. falciparum, 3D7 (designated as 3D7mut) and FCC1/HN (FCC1/HNmut), were used for the transfection. Parasites in the high synchronous ring stage were transfected with individual recombinant plasmids by electroporation, as described above (Fig. 1b). The transfected culture was selected using 2.5 nM WR99210 at 48 h post-electroporation until the WR99210-resistant parasites were visible. Cloning of the resistant parasites was performed by limiting dilution. The resulting clones with the replacement F446I were designated as 3D7F446I mut and FCC1/HN F446I mut lines (Fig. 1c), and the clones with the replacement C580Y, as 3D7C580Y mut and FCC1/HN C580Y mut lines (Fig. 1d).

The genomic DNA of each parasite clone was prepared for PCR verification of the integrations and mutant
Fig. 1 Strategy for generation of constructs for parasite transfection. 

**a** Generation of PfK13 fragment with the mutations

**b** Recombinant plasmid

**c** Replacement F446I

**d** Replacement C580Y

Fig. 1. Strategy for generation of constructs for parasite transfection. 

**a** PCR amplification of k13 gene with F446I mutation.  
**b** Insertion of PfK13F446I mut into the vector to generate the pARL-K13F446I recombinant plasmid.  
**c** Integration of PfK13F446I mut into the target locus of the parasite chromosome.  
**d** Integration of PfK13C580Y mut into the target locus of the parasite chromosome.
allele replacements. The P5 primer is located in 5' UTR of k13 in the chromosome while the P4 primer in 3'UTR of Pb DHFR-TS gene that is located in the plasmid. As shown in Fig. 2a, a 2371-bp fragment was amplified from the transgenic parasite lines with either F446I or C580Y mutation by using the P4/P5 primers, indicating that both PfK13\textsuperscript{F446I mut} and PfK13\textsuperscript{C580Y mut} were inserted into the target locus by single crossing-over recombination. No fragment was detected from their parent parasites, 3D7\textsuperscript{wt} and FCC1/HN\textsuperscript{wt} strains.

Similarly, the P6 primer is located in 3' UTR of the gene in the chromosome, and the P3 primer, in the plasmid. Successful amplification of 2195- and 1614-bp fragments further demonstrated the correct integration into the loci of PfK13 with F446I and C580Y mutations, respectively (Fig. 2b). In addition, the fragments amplified by the primer pair P4/P5 were sequenced, and the correct substitution of A for T for the F446I mutation and A for G for the C580Y mutation was confirmed (Fig. 2c).

As shown in Fig. 1c, the transgenic parasite with the F446I mutation has two full-length k13 coding sequences, the mutant and wild-type k13 alleles. To exclude the possibility that they are both expressed, we generated the full-length k13 cDNA from the 3D7\textsuperscript{F446I mut} parasite and

![Diagram of PCR amplification and sequencing](image_url)

**Fig. 2** Verification of the integration of the F446I mutation into the target locus. a PCR amplification using the indicated primers produced the expected fragments in lanes 4, 5, 7, and 8 for the transgenic parasite lines, whereas no fragments were detected in the other lanes for the parent parasites. Lanes 1–9: negative control, pARL-K13\textsuperscript{F446I} plasmid, pARL-K13\textsuperscript{C580Y} plasmid, FCC1/HN\textsuperscript{F446I mut}, FCC1/HN\textsuperscript{C580Y mut}, 3D7\textsuperscript{C580Y mut}, 3D7\textsuperscript{F446I mut}, and 3D7\textsuperscript{wt}. b PCR analysis showed that the expected fragments were amplified from the transgenic parasites. Lanes 1–9, as indicated above. c Sequencing verification of the mutations from T to A for F446I and G to A for C580Y.
subcloned the fragments into pEASY-T1 cloning vector for sequencing. As shown in result no k13 gene was amplified by PCR in the no-RT controls (Fig. 3a, lane 1), indicating no contaminating of genomic DNA in the sample. 12 clones of the recombinant plasmids containing the k13 gene were sequenced across the F446I mutation. Sequencing data showed that all the 12 clones showed the F446I mutant alleles (none of them are wild-type), indicating that only the mutant allele is expressed.

To investigate whether the integration of the mutant alleles affected the expression of k13, the k13 expression of each transgenic parasite clone were determined with western blot. As shown in Fig. 3b, the expression levels of k13 in both 3D7 C580Y mut and 3D7 F446I mut clones were

Fig. 3  Verification of only k13 with the F446I mutation expressed. a Analysis of expression of both the wild-type and mutant k13 alleles from the 3D7 C580Ymut parasite. The gel blot shows the K13 gene amplified from the RNA extracted from the transgenic parasite (lane 1, no-RT control) and the cDNA (lane 2); sequence analysis shows all the 12 clones have the F446I mutant alleles. b Western blot analysis of PfK13 expression in the transgenic parasite lines and their parent strains.
similar to those in the 3D7 wt line. Similar expression levels of k13 were detected in FCC1/HN C580Y mut and FCC1/HN F446I mut.

**Ring-stage survival of the transgenic parasites with the F446I mutation**

RSA0–3 h was performed to evaluate the transgenic parasite lines expressing F446I and C580Y mutants of k13 for their susceptibility to DHA. First, ring survival rates of both transgenic parasites and their parent lines were measured when parasites were exposed to elevated concentrations of DHA. As shown in Fig. 4a, at the lowest concentration of DHA (i.e., 5.5 nM), no significant differences in the ring survival rates of the 3D7 wt and 3D7 F446I mut parasite lines were observed (p > 0.05), while at 11, 22, and 44 nM DHA, the ring survival rates of 3D7 F446I mut were significantly higher than that of 3D7 wt (p < 0.05, 0.01, and 0.001, respectively). At the concentrations beyond 44 nM, i.e., from 88 to 700 nM, no parasite was observed in the 3D7 wt parasite line, and different ring survival rates were detected in the 3D7 F446I mut line. A similar result was obtained for the FCC1/HN F446I mut line when compared with its parent parasite strain (Fig. 4b).

Then 2 DHA concentrations (i.e., 20 and 700 nM) from the above-mentioned experiment were selected to further measure the ring survival rates of the F446I and C580Y mutant parasite lines. No parasite of the wild type strain of 3D7 could survive (survival rate: 0) when 700 nM DHA was used, while the ring survival rates for the transgenic parasite lines of both 3D7 F446I mut and 3D7 C580Y mut were similar at 0.73 ± 0.10% and 0.91 ± 0.11%, respectively (Table 1). However, when 20 nM DHA was used, both 3D7 F446I mut and 3D7 C580Y mut transgenic parasite lines showed high ring-survival rates at 26.0 ± 1.48% and 33.08 ± 1.20%, respectively, whereas the 3D7 wt parasites showed a significantly lower survival rate at 3.15 ± 0.13% (p < 0.05). Similar results were obtained after measuring the ring survival rates of FCC1/HN F446I mut and FCC1/HN C580Y mut lines when compared with the FCC1/HN wt strain (Table 1). F08-27 isolate of *P. falciparum* carrying the C580Y mutation collected from China to Myanmar border showed high RSA survival rates [11] and is used as a positive control in this study. The survival rate of the F08-27 isolate was 8.27 ± 0.63% and 65.8 ± 4.06% when the isolate was treated with 700 and 20 nM DHA, respectively.

For generating transgenic 3D7 parasites with the F446I mutation, 7 clones were obtained by sub-cloning the WR99210-resistant parasite by limiting dilution. Then these clones were sequenced to confirm the presence of F446I. All the clones showed increased levels of ring survival from 17.6 to 29.5% at 20 nM DHA when compared with 3.15% survival of 3D7 wt (Fig. 5). Interestingly, the survival rate of one clone, 3D7 F446I mut-F1, expressing the F446I mutant allele was 0 at 700 nM DHA, which is the same as the wild type of the strain. The survival rate was also lower than that of the other clones (17.6%) at 20 nM DHA. The sequencing of the Pf K13 gene of the 3D7 F446I mut-F1 clone showed an additional mutation at nucleotide position 1156 from G to T, resulting in a change in amino acid residue from valine to leucine at position 386.
It is rare for multisite mutation to occur in \( k13 \) in field isolates. Further investigation is required to understand whether more than 1 mutation in \( k13 \) affects the ART susceptibility of the parasite.

**Discussion**

ART resistance is evaluated in vivo by delayed parasite clearance and in vitro by high ring survival rates with RSA\(_{0-3\ h}\) [8]. The resistant phenotypes are strongly associated with a single mutation after amino acid position 440 in \( k13 \) [7, 10]. Some \( k13 \) mutations, such as C580Y, R539T, Y493H, and I543T, have been reported to be associated with the resistance of parasites to ART [9], and they are highly prevalent in the border areas of Thai/Cambodia and Thai/Myanmar but rare north of Southeast Asia, including north of Myanmar and China–Myanmar border region [7, 19, 20]. Interestingly, north of Southeast Asia, the most prevalent mutant allele of \( k13 \) is F446I. In previous study, the result showed that the prevalence of this mutation is 32.7% in the China–Myanmar border area and this mutation was not detected in Thai/Cambodia and Thai/Myanmar border regions [11]. From 2009 to 2012, Huang et al. [15] conducted a therapeutic efficacy study of artesunate in the China–Myanmar border region and showed that the single mutation of F446I with a prevalence of 36.5% was significantly associated with prolonged parasite half-life and parasitaemia on day 3 after treatment. However, several other studies have found no correlation between this mutation and prolonged parasite clearance time [7, 21]; however, the reason may be the small sample sizes. A previous study found that 2 of 3 parasites with this mutation showed a parasite clearance time < 5 h [22]. A recent study on the therapeutic efficacy of artemether–lumefantrine conducted in the northeast region of India also found no correlation of this mutation with delayed parasite clearance [23].

In previous study, the ring survival rate of 27 isolates with F446I mutation were measured by using RSA\(_{0-3\ h}\). Of them, only 6 had a high ring survival rate (> P\(_{95}\) value) that was defined as the association with delayed parasite clearance in vivo, and the ring survival rates of the other isolates were below the P\(_{95}\) value. The statistical analysis indicated that the F446I mutant was not associated with high ring-stage survival. Thus, there is much debate on whether the F446I mutation is associated with ART resistance. In this study, F446I mutation in \( k13 \) was introduced to the ART-sensitive 3D7 and FCC1/HN strains, and the resulting transgenic parasite lines FCC1/HN F446I mut and 3D7 F446I mut showed increased ring survival rates at both 20 and 700 nM DHA concentrations. Moreover, the survival rates of the transgenic parasites with the F446I mutation were similar to those of the parasites with the C580Y mutation demonstrated to be correlated with ART resistance. Thus, the result suggested the F446I mutation may be associated with ART resistance.

The C580Y mutation in \( k13 \) is a primary marker for ART resistance and the most prevalent mutation in western Cambodia where the resistant parasite was first reported. Several transgenic parasite clones have been generated in which the C580Y mutation was introduced into the \( k13 \) gene of various parasite isolates by using the CRISPR–Cas9 technology or zinc-finger nucleases [10, 13]. These transgenic clones showed elevated ring-stage survival when compared with their parent parasites; however, the ring survival rates of these transgenic parasite clones were variable. The 2 transgenic clones expressing \( k13 \) C580Y showed survival rates of 11–15%, while another transgenic FCB parasite clone expressing the C580Y allele showed a survival rate of 1.9% when compared with the survival rate of 0.3% in the parent FCB isolate. We generated both FCC1/HNC580Y mut and 3D7C580Y mut transgenic lines that showed 0.63 ± 0.07% and 0.91 ± 0.11% RSA\(_{0-3\ h}\) survival at 700 nM DHA concentration, whereas the survival rates of their parent parasites were 0%. Interestingly, when the DHA concentration reduced to 20 nM, both transgenic lines showed 17.5 ± 0.80% and 33.08 ± 1.20% survival rates, whereas their parent parasites showed only 2.86 ± 0.44% and 3.15 ± 0.13% survival rates, respectively. It could be concluded that the introduction of both C580Y and F446I mutations in \( k13 \) of 2 distinct laboratory parasite strains resulted in increased ring survival indicating that, similar to the C580Y substitution, F446I may play role in parasite response to ART.
Conclusions

In summary, the F446I mutation in k13 is the most prevalent allele at the China–Myanmar border and north of Myanmar. In this study, this mutation is associated with the increased ring survival rates by genetically introducing the mutation into the k13 gene of distinct ART-sensitive laboratory parasite strains, suggesting that the F446I mutation be used as a molecular marker for monitoring ART-resistant parasites and highlighting the importance of surveillance of F446I mutants for resistant parasites.

Abbreviations

k13: P. falciparum K13 propeller gene; ART: artemisinin; RSA0–3h: ring-stage survival assay of 0–3 h; DHA: dihydroartemisinin.

Authors’ contributions

JW, DZ and WP conceived and designed the experiments. JW, YH, YZ and RY performed the experiments. JW and WP analysed the data and drafted the manuscript. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

Consent for publication

Not applicable.

Ethics approval and consent to participate

Not applicable.

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