A bacterial chaperone is required for plastid function in malaria parasites

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Abstract

Apicomplexan parasites such as *Plasmodium falciparum*, the causative agent of malaria, contain a non-photosynthetic plastid known as the apicoplast that functions to produce essential metabolic compounds. It was previously reported that several members of the Clp family of chaperones and proteases localize to the apicoplast. In bacteria and in chloroplasts these proteins form complexes that degrade proteins in a proteasome-like manner to regulate key cellular processes, but their function in the apicoplast is completely unknown. In this study, we generated a conditional mutant of the *P. falciparum* apicoplast-targeted pfclpc gene and found that under normal conditions it localizes to the apicoplast. Knockdown of PfClpC results in growth inhibition and morphological defects, indicating that PfClpC is essential for parasite viability. Upon inhibition, PfClpC loses its apicoplast localization and appears in vesicle-like structures. Other apicoplast-targeted proteins also localize to these structures, suggesting that organelle integrity is compromised. Addition of isopentynyl pyrophosphate completely rescued the growth inhibition, indicating that the only essential function of PfClpC is related to the apicoplast. Moreover, cellular assays suggest that PfClpC inhibition interferes with the ability of the schizont-stage parasites to properly sort functional apicoplast organelles into daughter-merozoites. These data show that PfClpC is an essential gene that functions to maintain apicoplast integrity.
Author Summary

The deadly human malaria parasite, *Plasmodium falciparum*, contains a unique organelle called the apicoplast, a non-photosynthetic plastid that produces vital metabolites. Members of the prokaryotic-derived Clp family were previously reported to localize to the apicoplast. In bacteria and plant chloroplasts, Clp homologs form a proteasome-like complex that degrades proteins but their function in parasite biology is unknown. Here we took a conditional knockdown approach to study an apicoplast localized Clp protein, PfClpC, which we found to be essential for parasite viability. Inhibition of PfClpC results in a growth arrest phenotype that correlates with a reduced replication rate. We observed that PfClpC localizes to the apicoplast, however upon inhibition it is found dispersed in vesicle-like structures suggesting a complete breakdown of organelle integrity. Our ability to rescue the phenotype by adding an essential apicoplast-derived metabolite proved that the only essential function of PfClpC is linked to apicoplast function. Furthermore, we have found evidence supporting a role for PfClpC in apicoplast sorting into daughter cells. Therefore, we propose PfClpC as a potential drug target due to its essentiality, prokaryotic origin and absence from the human host.
Introduction

Malaria is a devastating human disease caused by obligate intracellular parasites of the genus *Plasmodium*. This disease results in nearly 450,000 deaths each year, which are mostly caused by one species, *Plasmodium falciparum* [1]. The life cycle of the parasite is remarkably complex, moving between different cellular niches in a mosquito vector and in the human host. Upon infection, the parasite initially invades liver cells followed by invasion into red blood cells (RBCs), where the parasite numbers expand exponentially via asexual replication. This part of the cycle initiates with the invasion of a diminutive merozoite into the RBC, developing first into the early ring, then a metabolically active trophozoite, and finally a multinucleated schizont that will give rise to multiple merozoite progeny which egress and reinvade fresh RBCs.

This blood stage form of the parasite is responsible for the entirety of malaria-associated morbidity and mortality. Currently, the parasite has gained resistance to all clinically available antimalarial drugs, generating an urgent need to identify new drugs and potential new drug targets [2,3]. The minute eukaryotic cell of *P. falciparum* is remarkably complex with two organelles that carry their own genetic material, the mitochondrion and a unique algal endosymbiont known as the apicoplast [4]. The apicoplast harbors vital metabolic pathways that are required for parasite growth and survival [5]. Importantly, drugs that target cellular processes in the apicoplast are clinically effective [6-8]. Therefore, understanding the function, structure and biogenesis of the apicoplast are essential areas of research that provide a rich vein of antimalarial drug targets.
The parasite specific and essential nature of the apicoplast makes the development of anti-malarial drugs targeting this organelle extremely attractive. Very little is known about the biogenesis and maintenance of the apicoplast organelle, although factors that are required for or regulate these processes may be an important source of anti-malarial drug targets. One potential class of such targets are the caseinolytic protease (Clp) family of proteins that act as key regulators of the biology of bacterial cells, the evolutionary ancestors of the apicoplast. In bacteria and plant chloroplasts, Clp proteins play vital roles in cell/organelle division, segregation, protein homeostasis and protein transport [9]. Typically, they form a regulated proteolytic complex in which a Clp protease is paired with a Clp chaperone that has a AAA+ ATPase domain (also known as the Hsp100 family of chaperones) such as ClpC or ClpA [10]. There are several putative clp genes encoded in the *P. falciparum* genome and it was recently shown that one of them (PfHSP101) is located at the host-parasite interface and is required to transport parasite virulence factors into the infected host cell [11-13]. Two other Clp proteins have been localized to the mitochondria [14] and six putative Clp’s have been localized to the apicoplast [15], yet very little is known about their roles in parasite biology. Although some of their enzymatic activities have been studied *in vitro* [16], their roles *in vivo* remain poorly understood due to the challenging genetics of *P. falciparum* and the difficulty in targeting organelle localized genes for molecular study. Here we studied the role of the previously uncharacterized PfClpC (PF3D7_1406600), an Hsp100 chaperone with a triple AAA+ ATPase domain.
The *pfclpc* gene is conserved in all *Plasmodium* species, as well as in other apicoplast containing *Apicomplexa*. It encodes a large protein (156 kDa) with a predicted apicoplast transit peptide. Using a conditional knockdown approach, we show that PfClpC activity is essential for parasite survival and growth. Conditional inhibition of PfClpC interferes with apicoplast integrity and the localization of apicoplast proteins. We observe a delayed growth inhibition phenotype, which results from a reduced replication rate that compounds with each subsequent replication cycle. Importantly, we can chemically rescue PfClpC-associated phenotypes using isopentenyl pyrophosphate (IPP) indicating that the only essential role of PfClpC is linked to apicoplast function. Finally, we demonstrate that even after prolonged inhibition and a general apicoplast loss, parasites are able to recover from undetectable levels and grow when PfClpC inhibition is reversed. Overall, we show that PfClpC is a vital apicoplast-targeted protein that is essential for parasite viability through its role in apicoplast function.
Results

Generating conditional mutants of PfClpC

In order to examine the biological role of the putative chaperone PfClpC in apicoplast function and parasite biology, we took a genetic approach based on conditional chaperone auto-inhibition. We inserted a dihydrofolate reductase (DHFR)-based destabilization domain (DDD) into the pfclpc genomic locus, a technique that was previously used successfully to conditionally inhibit chaperone function \[13,17\], as well as to knockdown other Plasmodium proteins \[18,19\] (Fig 1A). In the chaperone-DDD fusion protein, the unfolded DDD binds the chaperone intra-molecularly, thereby excluding client proteins and inhibiting normal chaperone function (Fig 1A). A small molecule ligand, trimethoprim (TMP) is used to stabilize and refold the DDD, releasing the chaperone to resume its normal function (Fig 1A). Using a single crossover homologous recombination strategy, we tagged the C-terminus of the pfclpc gene with a triple-hemagglutinin (HA) epitope tag and the DDD (Fig 1B). We isolated two clones from two independent transfections and analyzed the pfclpc locus using a southern blot analysis (Fig 1B). Both isolated clones were correctly tagged, and subsequent experiments were done using the two clones, 1G8 and 2E10 (Fig 1B). Henceforth, these parasite lines will also be referred to as PfClpC-DDD.

Using an anti-HA antibody, we confirmed expression of the PfClpC-DDD fusion protein at the expected molecular size in tagged clones but not in the parental line (Fig 1C). The subcellular localization of PfClpC in PfClpC-DDD parasites was observed using immunofluorescence microscopy (Fig 1D). We detected
expression of tagged PfClpC in distinctive structures throughout the asexual blood stages; punctate in early parasites, elongated in mid-stage trophozoites and multiple foci in individual merozoites in the late schizonts stages (Fig 1D).

PfClpC is essential for intraerythrocytic growth of malaria parasites
To test the requirement of PfClpC for parasite growth during the asexual blood stages, we removed the stabilizing ligand (TMP) from culturing media and monitored the growth of unsynchronized PfClpC-DDD parasites. A severe and dramatic growth arrest was seen in parasites after TMP withdrawal (Fig 2A). We found that inhibition of PfClpC relies on TMP in a dose-dependent manner with an EC50 of 80nM (Fig 2B). As expected, based on previously described chaperone-DDD auto-inhibition model, TMP removal did not lead to PfClpC degradation, as protein levels remained constant over time (Fig 2C). Overall, these results demonstrate that PfClpC activity is essential for parasite survival and growth within human RBCs.

PfClpC is required for apicoplast integrity
In order to verify the subcellular localization of PfClpC, we stained cells with anti-HA and anti-acyl carrier protein (ACP), a known apicoplast marker [20], and found that the two co-localize, indicating that under normal conditions PfClpC localizes to the apicoplast (Fig 3A). After TMP removal, we observed that the integrity of the apicoplast was compromised, as observed by staining of PfClpC and ACP (Fig 3B). Inhibition of PfClpC function resulted in loss of the canonical apicoplast
morphology and in the appearance of PfClpC in a punctate, vesicle-like, pattern throughout the cell (Fig 3B). Moreover, this abnormal localization was also observed for ACP, which was detected in similar vesicle-like structures upon PfClpC inhibition, suggesting damage to apicoplast integrity (Fig 3B). Apicoplast targeting of nuclear encoded proteins is mediated through an N-terminal transit peptide that is cleaved in the apicoplast to produce the mature protein [21,22]. In correlation with the mislocalization of PfClpC, a second higher band for PfClpC appeared upon TMP removal, suggesting that the N-terminal transit peptide was not cleaved because it did not reach the apicoplast (S1 Fig).

**Intraerythrocytic development requires PfClpC**

In agreement with apicoplast dysfunction, PfClpC mutants developed normally during the early stages of rings and trophozoite (S2 Fig), but late schizont stages (≤6 nuclei) developed aberrant morphology, including irregular cellular shape, empty vacuoles and fewer nuclei suggesting that these parasites are nonviable (Fig 4A). These morphologically abnormal parasites appeared on the 3rd replication cycle and their fraction increased over time (Fig 4B). Analysis of the entire population using flow cytometry revealed that instead of a single peak that usually characterizes a synchronized culture, these late-stage parasites had a wider distribution, further suggesting variation in DNA content (Fig 4C). To test the viability and replication efficiency of the mixed parasites population, we used a synchronized culture and monitored the rate of schizonts to ring conversion. In agreement with our previous results showing the presence of morphologically
abnormal schizonts, TMP removal resulted in a significant decrease in the numbers of parasites that were formed in each successive generation (Fig 4D). This reduced replication rate accounts for the observed growth inhibition as well as the increase in the numbers of morphologically abnormal parasites with each generation.

Chemical rescue of PfClpC-DDD parasites using IPP

The only essential function of the apicoplast during the blood stages is the biosynthesis of isopentenyl pyrophosphate (IPP), the precursor for all isoprenoids, through the non-mevalonate pathway [23]. To test the effect of IPP on PfClpC inhibition, we removed TMP and added IPP to the growth media of PfClpC-DDD parasites and observed normal growth as well as typical cellular morphology (Figs 5A and B). Immunofluorescence microscopy revealed that IPP treated PfClpC-DDD parasites survived in the absence of a functional apicoplast and still retained the multiple vesicle-like structures containing PfClpC and ACP (Fig 5C). Moreover, quantitative Real Time PCR (qRT-PCR) analysis supported the visual observation that the apicoplast disappears from these parasites (Fig 5D). Importantly, the mitochondrial genome was not similarly affected by PfClpC inhibition indicating that, in addition to a functional damage to the apicoplast, there was an actual loss of the plastid genome (Fig 5D). Overall, we concluded that the only essential activity of PfClpC is linked to apicoplast function.

The apicoplast is lost from most but not all PfClpC-DDD parasites
In order to link the cellular observation of a morphologically mixed PfClpC-DDD population to the phenotypic evidence of apicoplast dysfunction, we aimed to visualize apicoplast presence or absence in early stage parasites (≤5 hours post invasion). We used Image Stream analysis, which combines flow cytometry with imaging, to analyze these parasites and found that all parasites were positive for PfClpC (S3 Fig and S1 Table). Higher resolution microscopy confirmed that these cells contained PfClpC and that its localization appeared, even in these early stages, in a vesicle-like pattern rather than as a typical discrete apicoplast structure (S4 Fig). This suggests that either newly formed rings inherit these vesicles from their mother cell or that these vesicles represent very early *de novo* synthesis of apicoplast proteins.

Due to the fact that apicoplast proteins are synthesized even in absence of the organelle, they are not indicative of apicoplast loss. Therefore, we employed a functional assay to investigate whether PfClpC inhibition results in a mixed population of parasites with and without an apicoplast. We wanted to know, whether or not a small, yet viable, population of parasites containing an intact and functional apicoplast remained in PfClpC-DDD parasites cultured in the absence of TMP. To test this, we first removed TMP and allowed PfClpC-DDD parasites to grow in the presence or absence of IPP for two weeks (Fig 6A). As expected, PfClpC-DDD parasites grown without TMP and supplemented with IPP grew normally, whereas parasites incubated without TMP and without IPP were unable to grow and were undetectable for several days. On day 14 we removed IPP and relieved PfClpC inhibition by adding back TMP, and monitored the growth of the
parasites (Fig 6B). Upon addition of TMP to the media, parasites grown in the absence of IPP (Fig 6B) recovered and resumed normal growth, indicating that a small fraction of these parasites indeed possessed a functional apicoplast. Conversely, parasites that were grown with IPP started dying 48 hours after removing IPP and adding back TMP (Fig 6B), suggesting that these apicoplast-less parasites through continued growth had outcompeted the few remaining parasites that contained a functional apicoplast. Despite restoration of PfClpC activity, once lost, the parasite is not able to generate, de novo, the four-membrane apicoplast. Overall, these data indicate that PfClpC-DDD mutants grown in the absence of TMP lose the apicoplast from most parasite progeny with a small, yet observable population still retaining the plastid.
Discussion

The deadly malaria parasite, *Plasmodium falciparum*, is a eukaryotic pathogen and as such, it shares conserved basic biology with its human host. It is therefore both challenging and essential, in the search for potential drug targets, to identify key components that are absent or significantly different from the human host.

One such potential candidate is the apicoplast-associated prokaryotic Clp family of chaperones and proteases. In the bacterial ancestors, as well as in other organelar descendants such as the mitochondria and chloroplast, these proteins serve a variety of basic molecular functions ranging from protein degradation, transport across membranes, protein folding, cell division, stress response and pathogenicity [24]. These have placed bacterial Clp proteins at the center of several drug discovery programs, and have led to the identification of potent and specific inhibitors [25,26]. Very little is known, however, about the functional roles of the apicoplast-resident Clp proteins in the biology of *Plasmodium falciparum*.

In this study we identified an essential Clp family member, PfClpC, a nuclear encoded gene that is transported to the apicoplast where it is required for the integrity of the organelle. Using a conditional knockdown approach, we demonstrated that PfClpC activity is essential for parasite growth and viability (Figs 1 and 2).

The growth defect of PfClpC-DDD parasites is linked to a failure in apicoplast integrity in several ways. We showed that PfClpC localizes to the apicoplast under normal conditions, but appears, along with other apicoplast proteins, in vesicle like structures upon PfClpC inhibition (Fig 3). Several studies reported the appearance
of such structures when the apicoplast integrity is compromised, for example with
the use of certain antibiotics [5,23,27]. This was interpreted as stalled vesicular
transport that has left the ER but cannot dock to the apicoplast surface due to loss
of the organelle.
Moreover, chemical rescue using IPP restored PfClpC-DDD growth as well as
recovered normal cellular morphology (Fig 5). It has been shown that isoprenoid
biosynthesis is the only essential metabolic function of the apicoplast, and
supplementing IPP can replace a non-functional apicoplast in living parasites [23].
Indeed, the IPP supplemented parasites grew normally but microscopic imaging
revealed the presence of punctate structures, which are indicative of apicoplast
loss (Fig 5C). This was also supported by qRT-PCR analysis that demonstrated a
decrease in the ratio of nuclear to apicoplast genome, highlighting an actual loss
of the organelle (Fig 5D). We therefore concluded that the only essential function
of PfClpC is linked to apicoplast biology.
Inhibition of essential apicoplast metabolic pathways with drugs like
Fosmidomycin, kills parasites immediately and does not lead to the loss of the
organelle [28]. Conversely, inhibition of apicoplast translation or replication with
drugs like Doxycycline, allows the parasites to complete one asexual cycle,
proceed through the second replication cycle, and die only at the schizont stage of
the second cycle [7,23]. Similar to the effect of drugs that inhibit apicoplast
replication, PfClpC mutants develop normally during the early stages of rings and
trophozoite (S2 Fig), but late schizont stages (≤6 nuclei) exhibited aberrant
morphology (Figs 4A and C). These non-viable parasites did not manifest
uniformly at the end of the second cycle but appeared on the 3rd replication cycle, and their fraction increased over time (Fig 4B). One possible explanation for the delayed growth arrest, as well as the gradual increase in abnormal parasites, is that PfClpC inhibition interferes with the segregation or division of functional apicoplasts into daughter merozoites. As a consequence, a mixed population of viable and non-viable daughter cells is forming after each cycle, diluting over time the viable parasites in the total culture. Indeed, we observed a significant decrease in the rate of schizont to ring conversion in each successive generation (Fig 4D), clarifying the delayed growth inhibition, and suggesting a possible defect in apicoplast sorting.

Since most apicoplast resident proteins are encoded by the nucleus [20], they are expressed and sorted into vesicles despite the absence of an apicoplast and therefore could not serve as a proper indicator of apicoplast presence, even in the earliest stages (S3 and S4 Figs and S1 table). We hypothesized that inhibition of PfClpC resulted in an apicoplast segregation or division defect that leads to improper sorting of the organelle into daughter merozoites. Therefore, during every schizogony event only a small fraction of progeny receives an apicoplast from the mother cell. To test the possibility of an apicoplast-related sorting defect, we removed TMP for several replication cycles, inducing persistent growth arrest, and then adding it back and monitoring parasite growth. In the event of a uniform functional damage to all parasites in the culture, PfClpC re-activation would not lead to viable parasites, as de novo synthesis of the apicoplast is impossible. Nonetheless, we observed that re-addition of TMP could restore parasites growth,
indicating the presence of a small but undetectable sub-population of parasites
that contains a functional apicoplast (Fig 6). This observation further supports a
sorting defect rather than a general apicoplast dysfunction in the entire parasite
population. Interestingly, TMP addition had the opposite effect on parasites that
were rescued with IPP. These parasites started dying 48 hours after removing IPP
despite addition of TMP (Fig 6), indicating that re-activation of PfClpC was not
effective enough to sustain viability in a population of parasites that permanently lost the
apicoplast. Future work will be needed in order to reveal how exactly PfClpC
inhibition affects organelle sorting.

The question remains as to what is the molecular mechanism of PfClpC activity
and how does it affect apicoplast function. There are two ClpC homologs in \textit{P. falciparum}, PfClpC that is a nuclear encoded protein targeted to the apicoplast
and PfClpM (PFC10_API0060) that is encoded by the apicoplast genome. Our
data show that their functions are not redundant and that PfClpC is essential for
parasite survival and apicoplast function (Figs 2, 3, 4, and 5). In cyanobacteria
and plant chloroplasts, ClpC orthologs, together with a ClpP protease, typically
form an ATP-dependent proteolytic complex that degrades proteins to maintain
protein homeostasis in the cell/ organelle [10,29]. PfClpC, but not PfClpM,
possesses the entire conserved motif that is required to interact with ClpP
proteases [30]. The \textit{Plasmodium} ClpP homolog, PfClpP, was shown to have
protease activity \textit{in vitro} and localize to the apicoplast \textit{in vivo} [16]. Several other
studies provided structural data and binding analysis for two other apicoplast Clp
proteins, the inactive protease PfClpR [31] and the adaptor protein PfClpS [32].
As expected, these two proteins have bacterial and chloroplast homologs, which are essential components of the Clp proteolytic complex [31,32]. Interestingly, it was shown that interfering with ClpP activity inhibits cell division in Gram-positive bacteria due to changes in rates of substrate degradation [33]. Further studies are required in order to understand whether a similar complex is indeed formed in the apicoplast of *P. falciparum*, and whether it plays an essential role in maintaining a functional apicoplast. Such investigation may reveal the link between the apicoplast sorting defect and the proteolytic ClpP/C complex in the form of unknown substrate/s that need to be degraded in order to facilitate proper organelle division and segregation.
Fig 1: Generating PfClpC-DDD Conditional Mutants.

A. Mechanism of PfClpC conditional inhibition. The pfclpc locus was modified to contain a triple hemagglutinin (HA) tag and a DHFR-based destabilization domain (DDD). In the presence of trimethoprim (TMP) the DDD is stable and the chaperone is active. Upon TMP removal the chaperone binds the DDD intramolecularly and cannot interact with client proteins, inhibiting normal activity.

B. Single crossover homologous recombination enables the integration of the plasmid into the 3' end of the pfclpc gene (upper panel). Southern blot analysis of genomic DNA (bottom panel) isolated from parasite lines indicated above the lanes. The genomic DNA was digested with NcoI and XmnI. Bands expected from integration of the plasmid into the 3' end of the pfclpc gene were observed in two clones (1G8 and 2E10), isolated from two independent transfections (Red). A plasmid band was observed in the clones (blue), suggesting that a plasmid concatamer integrated into the gene. A single band indicative of the parental allele was observed for the parental strain (black) and it was absent in the integrant clones.

C. Western blot of parasite lysates from parental line and two independent clones (1G8 and 2E10) probed with antibodies against HA (green) and EF1α (loading control, red). The protein marker sizes that co-migrated with the probed protein are shown on the left.

D. Immunofluorescence imaging of fixed unsynchronized PfClpC-DDD parasites stained with antibodies against HA (green) and DAPI (blue). Z-stack images were
deconvolved and projected as a combined single image. Scale bar, 5μm. One representative experiment out of four (two for each clone) is shown.

Fig 2: PfClpC Activity is Essential for Parasite Growth.

A. Asynchronous PfClpC-DDD clones, 1G8 and 2E10, were grown with or without 10μM TMP and parasitemia was monitored every 24 hours over 12 days via flow cytometry. During the course of the experiment cultures were cut back and data were calculated using the actual parasitemia multiplied by the dilution factors of each individual culture. 100% of relative parasitemia represents the highest value of calculated parasitemia. Growth inhibition is observed after 7-8 days post TMP removal, corresponding to roughly 3-4 asexual cycles. Data are fit to an exponential growth equation and are represented as mean ± S.E.M. (n=3). One representative experiment out of three is shown.

B. Asynchronous PfClpC-DDD parasites were incubated for 11 days without TMP, and on day 12 were seeded in a 96 well plate with varying concentrations of TMP. Parasitemia was measured after 5 days using flow cytometry showing an EC50 of 80nM. Data are fit to a dose-response equation and are represented as mean ± S.E.M. (n=3). One representative experiment out of four is shown.

C. TMP was removed from PfClpC-DDD parasites and parasite lysates were isolated every 24 or 48 hours over 11 days. PfClpC and EF1α were visualized on Western blots using antibodies against HA (PfClpC-DDD, green) and EF1α (loading control, red). The protein marker sizes that co-migrated with the probed
protein are shown on the left. One representative experiment out of four is shown
two for each clone).

**Fig 3: PfClpC Inhibition Disrupts the Localization of Apicoplast Proteins**

A. PfClpC-DDD parasites were fixed and stained with antibodies against HA (red) and ACP (green). Both clones of PfClpC-DDD parasites are shown, indicated next to the images. Images from left to right are anti-HA, anti-ACP, DAPI, fluorescence merge and phase. Z-stack images were deconvolved and projected as a combined single image. Scale bar, 5μM.

B. PfClpC-DDD parasites were incubated for 10 days without TMP and then fixed and stained with antibodies against HA (red), ACP (green) and DAPI (blue). Both clones of PfClpC-DDD parasites are shown, indicated next to the images. Images from left to right are anti-HA, anti-ACP, DAPI, fluorescence merge and phase. Z-stack images were deconvolved and projected as a combined single image. Scale bar, 5μM.

**Fig 4: PfClpC Inhibition is Associated with Reduced Replication Efficiency**

A. Hema 3 stained thin blood smears of PfClpC-DDD parasites that were grown for 10 days without TMP. Both clones of PfClpC-DDD parasites are shown, as indicated above the images.

B. TMP was removed from synchronized PfClpC-DDD parasites and thin blood smears of late stage schizonts were stained and analyzed using light microscopy. Parasites were counted, and the fraction of defective cells (as seen
in 4A) was calculated out of the total late-stage population (≤6 nuclei). Data are shown from one representative experiment with clone 2E10. Two experiments (with technical triplicates) were performed with each clone.

4C. TMP was removed from synchronized PfClpC-DDD parasites and the DNA of the parasites was stained using Acridine Orange and was analyzed by flow cytometry during the 6th replication cycle. One representative image is shown (out of four experiments, two with each clone).

8D. TMP was removed from synchronized PfClpC-DDD parasites and the numbers of rings and late schizont stages was determined by flow cytometry. The ratio of rings to schizonts was calculated using the number of rings arising from schizonts in the previous generation. Cultures were re-synchronized at the beginning of each replication cycle using sorbitol. Data were normalized to the ring: schizont ratio in the presence of TMP. Data are shown for 1G8 clone. The experiment was performed twice for each clone (with technical triplicates in each experiment).

**Fig 5: IPP Rescues PfClpC Auto-inhibition**

18A. PfClpC-DDD parasites were grown for 10 days without TMP and supplemented with isopentenyl pyrophosphate (IPP). Parasitemia was measured using flow cytometry. During the course of the experiment cultures were subcultured and data were calculated using the actual parasitemia multiplied by the dilution factors of each individual culture. 100% of relative parasitemia represents the highest value of calculated parasitemia at the end of the experiment. Data are fit
to an exponential growth equation and are represented as mean ± S.E.M. (n=3).

One of two (one for each clone) representative experiments is shown. Graph
denotes data collected for 1G8 clone.

4B. Hema 3 stained thin blood smears of PfClpC-DDD parasites (1G8 clone) grown
for 10 days with TMP (upper), without TMP (middle) or without TMP and
supplemented with IPP (bottom). Two representative images for each condition
are shown.

8C. PfClpC-DDD parasites (1G8 clone) grown for 10 days without TMP and
supplemented with IPP. These PfClpC-DDD parasites were fixed and stained
with antibodies against HA (red) and ACP (green). Images from left to right are
anti-HA, anti-ACP, DAPI, fluorescence merge and phase. Z-stack images were
deconvolved and projected as a combined single image. Scale bar, 5μM.

13D. Synchronized PfClpC-DDD parasites were grown in the absence of TMP and
presence of IPP and DNA samples were taken at the beginning of each
replication cycle for quantitative Real Time PCR analysis. Apicoplast: nuclear
genome ratio was calculated for each replication cycle. Mitochondria: nuclear
genome ratio served as a control. Genome ratios were normalized to parasites
grown in the presence of TMP. Data are represented as mean ± S.E.M (n=3)
from one representative experiment (out of four, two for each clone).

20E.

Fig 6: PfClpC Inhibition is Reversible

22A. PfClpC tagged parasites were grown for 14 days without TMP (red line), or
without TMP and supplemented with IPP (green line). During the course of the
experiment cultures were subcultured and data were calculated using the actual parasitemia multiplied by the dilution factors of each individual culture. 100% of relative parasitemia represents the highest value of calculated parasitemia at the end of the experiment. Data are fit to an exponential growth equation and are represented as mean ± S.E.M. (n=3). One representative experiment out of two for each clone is shown.

7B. On day 14 IPP was removed from the media and TMP was added back to all parasite cultures and parasitemia was measured using flow cytometry. 100% of relative parasitemia represents the highest parasitemia value at the beginning of the experiment. Data are represented as mean ± S.E.M. (n=3). One representative experiment out of two for each clone is shown.

S1 Fig: PfClpC Auto-inhibition Interferes with Transit Peptide Processing
TMP was removed from PfClpC-DDD parasites and parasite lysates were isolated every 24 to 48 hours over 13 days. PfClpC and Plasmepsin V (PMV) were visualized on Western blots using antibodies against HA (PfClpC-DDD) and PMV (loading control).

S2 Fig: PfClpC inhibition does not affect early stage development
Hema 3 stained thin blood smears of early stage PfClpC-DDD parasites that were grown for 10 days without TMP. Both clones of PfClpC-DDD parasites are shown, indicated above the images.
S3 Fig: Early Stage PfClpC-DDD parasites are positive for HA

Synchronized PfClpC-DDD parasites incubated for 10 days without TMP. Schizont stage parasites were isolated on a percoll gradient following by a sorbitol treatment 5 hours later to obtain early rings (0-5 hours post invasion). Parasites were fixed and stained with DAPI to visualize DNA and antibodies against HA to observe PfClpC-DDD. Parasites were detected using Imaging flow cytometry (ImageStream). Images from left to right are anti-HA, DAPI, fluorescence merge and phase. Representative images of fixed parasitized RBCs as observes via the imaging flow cytometry.

<table>
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<tr>
<th>Clone/ condition</th>
<th>Infected cells positive for HA (%)</th>
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S1 Table: Early Stage PfClpC-DDD Parasites are Positive for HA

Synchronized PfClpC-DDD parasites incubated for 10 days without TMP. Schizont stage PfClpC-DDD parasites were isolated on a percoll gradient following by a sorbitol treatment 5 hours later to obtain early rings (0-5 hours post invasion). Parasites were fixed and stained with DAPI to visualize DNA and antibodies against HA to observe PfClpC-DDD. Parasites were detected using Imaging flow cytometry (ImageStream). The able shows a summary of the results as percentage of parasitized RBCs positive for HA staining.
S4 Fig: Vesicle-like Structures are Present in Early Stage PfClpC-DDD Parasites

Synchronized PfClpC-DDD parasites incubated for 10 days without TMP. Schizont stage parasites were isolated on a percoll gradient following by a sorbitol treatment 5 hours later to obtain early rings (0-5 hours post invasion). Parasites were fixed and stained with DAPI to visualize DNA and antibodies against HA to observe PfClpC-DDD. Images from left to right are phase, anti-HA, anti-ACP, DAPI, and fluorescence merge. Z-stack images were deconvolved and projected as a combined single image. Scale bar, 5μM.
Materials and Methods

Plasmid construction

Genomic DNAs were isolated from *P. falciparum* using the QIAamp DNA blood kit (QIAGEN). Constructs utilized in this study were confirmed by sequencing. PCR products were inserted into the respective plasmids using the In-Fusion cloning system (Clonetech). For generation of PfClpC-DDD a 1-kb homologous sequence from the 3’-end of the *pfclpc* gene (not including the stop codon) was amplified by PCR using primers 5’-cactataagaactcagCCAAATAAACCTATTGGTACTCTTCTATTATGTGGTTTCATCAGG-3’ and 5’-cgtatgggtacctaggAGATGAAAATTGTTGAACTGGTGCTTTTATTAATTGTACTTTAA-3’ and was inserted into pHADB [13] using restriction sites XhoI and AvrII (New England Biolabs).

Parasite culture and transfections

Commercially purchased de-identified and anonymized units of human blood were utilized for growing *P. falciparum* (Interstate Blood Bank, Memphis, TN). Parasites were grown in human red blood cells at 2% hematocrit in RPMI medium supplemented with Albumax I (Gibco) and transfected as described earlier [34,35]. pPfClpC-HADB was transfected in duplicates into 3D7-derived parental strain PM1KO which contains a hDHFR expression cassette conferring resistance to TMP [36]. Selection, drug cycling and cloning were performed as described [17] in the presence of 10 μM of TMP (Sigma). Integration was detected after one round of drug cycling with blasticidin (Sigma). Two clones
from 2 independent transfections, 1G8 and 2E10, were isolated via limiting
dilutions and used for subsequent experiments.
For IPP rescue, media was supplemented with 200 μM of IPP (Isoprenoids LC) in
PBS.

**Growth assays**
For asynchronous growth assays parasites were washed twice and incubated in
the required medium. Throughout the course of the experiment parasites were
sub-cultured to avoid high parasite density and parasitemia was monitored every
24 hours via flow cytometry. Relative parasitemia at each time point was back
calculated based on actual parasitemia multiplied by the relevant dilution factors.
Parasitemia in the presence of TMP at the end of each experiment was set as
the highest relative parasitemia and was used to normalize parasites growth.
Data were fit to exponential growth equations using Prism (GraphPad Software,
Inc.)

To generate an EC50 curve for TMP, asynchronous PfClpC-DDD parasites were
incubated for 11 days without TMP, and on day 12 were seeded in a 96 well
plate with varying concentrations of TMP. Parasitemia was measured after 5
days using flow cytometry. Data were fit to a dose-response equation using
Prism (GraphPad Software, Inc.).

To determine replication rate (rings: schizonts ratio), TMP was removed from
percoll-isolated schizonts-stage parasites and parasites were allowed to egress
and rein invade fresh RBCs. Parasitemia was monitored by flow cytometry and
microscopy. The ratio of rings to schizonts was calculated using number of rings
arising from schizonts in the previous generation. At the beginning of each
replication cycle parasites were re-synchronized using Sorbitol, and sub-cultured
when required. For each replication cycle data were normalized to rings:
schizonts ratio in the presence of TMP.

To determine the fraction of morphologically aberrant schizonts thin blood
smears of synchronized PfClpC-DDD parasites were performed at the final
stages of each replication cycle and the fraction of defective cells was calculated
based on the total late schizont stage parasite counts.

**Southern blot**

Southern blots were performed with genomic DNA isolated using the Qiagen
Blood and Cell Culture kit. 10 μg of DNA was digested overnight with Ncol and
XmnI (New England Biolabs) and integrants were screened using biotin-labeled
probes against the 3'-end of the *pfclpc* ORF. Southern blot was performed as
described earlier [37]. The probe was labeled using biotinylated Biotin-16-dUTP
(Sigma). The biotinylated probe was detected on blots using IRDye 800CW
Streptavidin conjugated dye (LICOR Biosciences) and was imaged, processed
and analyzed using the Odyssey infrared imaging system software (LICOR
Biosciences).

**Western blot**

Western blots were performed as described previously [18]. Briefly, parasites
were collected and host red blood cells were permeabilized selectively by
treatment with ice-cold 0.04% saponin in PBS for 10 min, followed by a wash in
ice-cold PBS. Cells were lysed using RIPA buffer, sonicated, and cleared by
centrifugation at 4°C. The antibodies used in this study were rat monoclonal anti-HA, 3F10 (1:3000) (Roche), mouse monoclonal anti-PMV (from D. Goldberg, 1:400) and rabbit polyclonal anti-EF1α (from D. Goldberg, 1:2000). The secondary antibodies that were used are IRDye 680CW goat anti-rabbit IgG and IRDye 800CW goat anti-mouse IgG (LICOR Biosciences, 1:20,000). The Western blot images were processed and analyzed using the Odyssey infrared imaging system software (LICOR Biosciences).

Microscopy and image processing

For IFA cells were fixed using a mix of 4% Paraformaldehyde and 0.015% glutaraldehyde and permeabilized using 0.1% Triton-X100. Primary antibodies used are rat anti-HA clone 3F10 (Roche, 1:100) and rabbit anti-ACP (from G. Mcfadden, 1:10,000). Secondary antibodies used are Alexa Fluor 488 and Alexa Fluor 546 (Life Technologies, 1:100). Cells were mounted on ProLong Diamond with DAPI (Invitrogen) and were imaged using DeltaVision II microscope system with an Olympus IX-71 inverted microscope using a 100X objective. All images were collected as Z-stack, were deconvolved using the DVII acquisition software SoftWorx and displayed as maximum intensity projection. Image processing, analysis and display were preformed using SoftWorx and Adobe Photoshop. Adjustments to brightness and contrast were made for display purposes. Thin blood smears were stained using Hema 3 stain set (PROTOCOL/ Fisher Diagnostics) and were imaged on a Nikon Eclipse E400 microscope.

Flow cytometry
Aliquots of parasite cultures (5 μl) were stained with 1.5 mg/ml Acridine Orange (Molecular Probes) in PBS. The fluorescence profiles of infected erythrocytes were measured by flow cytometry on a CyAn ADP (Beckman Coulter, Hialeah, Florida) and analyzed by FlowJo software (Treestar, Inc., Ashland, Oregon). The parasitemia data were fit to standard growth curve or dose–response using Prism (GraphPad Software, Inc.).

Quantitative Real Time PCR

Synchronized ring stage parasites samples were collected at the beginning of each replication cycle and genomic DNA was isolated by saponin lysis to remove extracellular DNA. Genomic DNA was purified using QIAamp blood kits (Qiagen). Primers that amplify segments from genes encoded by nuclear or organelles genomes were designed using RealTime qPCR Assay Entry (IDT). cht1 (nuclear): 5'-TCCATTGGGTGATTTTGTAAAGACTG-3' and 5'-CTAATGTTCATTATGTGCAAGCATTATC-3'. tufA (apicoplast): 5'-AATTAACACAAGCACAATCCGG-3' and 5'-GGTTTATGACGACCACCTTCT-3'. cytB3 (mitochondria): 5'-CTGCTTTCGTTGGTTATGTCTTAC-3' and 5'-CTCACAGTATATCTCCACATATCC-3'. Reactions contained template DNA, 0.5 μM of gene specific primers, and IQ™ SYBR Green Supermix (BIORAD). Quantitative real-time PCR was carried out in triplicates and was performed at a 2-step reaction with 95°C denaturation and 56°C annealing and extension for 35 cycles on a CFX96 Real-Time System (BIORAD). Relative quantification of target genes was determined using Bio-Rad CFX manager 3.1 software. Standard curves for each primers set were obtained by using different dilutions of control
gDNA isolated from parasites grown in the presence of TMP (20 to 0.2 ng) as template, and these standard curves were used to determine primers efficiency. For each replication cycle number, the organelle: nuclear genome ratio of the – TMP+IPP treated parasites was calculated relative to that of the +TMP control.

Imaging flow cytometry

Synchronized PfClpC-DDD parasites incubated for 10 days without TMP and then were isolated on a percoll gradient following by a sorbitol treatment 5 hours later to obtain early rings (0-5 hours post invasion). Cells were fixed and stained with anti HA antibody as described above and nuclei were stained using DAPI from Amnis Intracellular staining kit (EMD MILIPORE). Data were collected on ImageStream X Mark II (EMD MILIPORE) and an automated collection of a statistically large number of cells (10,000) was performed. Data were analyzed using IDEAS software version 6.2.
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A) A graph showing normalized parasitemia (%) over time (days) for different conditions: +TMP, +TMP+IPP, -TMP, and -TMP+IPP.

B) Images showing the effect of TMP and IPP on parasite morphology.

C) Images of stained cells showing HA, ACP, and DAPI staining under different conditions: +TMP and -TMP+IPP.

D) A bar chart showing the ratio of organelles to nuclear genomes under different conditions: +TMP and -TMP (Asexual Replication Cycle). The conditions include apicoplast:nuclear and mitochondria:nuclear.