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Expression of Cytosolic Peroxiredoxins in *Plasmodium berghei* Ookinetes Is Regulated by Environmental Factors in the Mosquito Bloodmeal

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**Abstract**

The *Plasmodium* oocinete develops over several hours in the bloodmeal of its mosquito vector where it is exposed to exogenous stresses, including cytotoxic reactive oxygen species (ROS). How the parasite adapts to these challenging conditions is not well understood. We have systematically investigated the expression of three cytosolic antioxidant proteins, thioredoxin-1 (Trx-1), peroxiredoxin-1 (TPx-1), and 1-Cys peroxiredoxin (1-Cys Prx), in developing oocinetes of the rodent parasite *Plasmodium berghei* under various growth conditions. Transcriptional profiling showed that *tpx-1* and 1-cys prx but not *trx-1* are more strongly upregulated in oocinetes developing in the mosquito bloodmeal when compared to oocinetes growing under culture conditions. Confocal immunofluorescence imaging revealed comparable expression patterns on the corresponding proteins. 1-Cys Prx in particular exhibited strong expression in mosquito-derived oocinetes but was not detectable in cultured oocinetes. Furthermore, oocinetes growing in culture upregulated *tpx-1* and 1-cys prx when challenged with exogenous ROS in a dose-dependent fashion. This suggests that environmental factors in the mosquito bloodmeal induce upregulation of cytosolic antioxidant proteins in *Plasmodium* oocinetes. We found that in a parasite line lacking TPx-1 (TPx-1KO), expression of 1-Cys Prx occurred significantly earlier in mosquito-derived TPx-1KO oocinetes when compared to wild type (WT) oocinetes. The protein was also readily detectable in cultured TPx-1KO oocinetes, indicating that 1-Cys Prx at least in part compensates for the loss of TPx-1 *in vivo*. We hypothesize that this dynamic expression of the cytosolic peroxiredoxins reflects the capacity of the developing *Plasmodium* oocinete to rapidly adapt to the changing conditions in the mosquito bloodmeal. This would significantly increase its chances of survival, maturation and subsequent escape. Our results also emphasize that environmental conditions must be taken into account when investigating *Plasmodium*-mosquito interactions.

**Introduction**

Malaria remains a major global threat to human health. One of the reasons for this is the effective transmission and dissemination of the disease-causing protozoan parasite *Plasmodium* by the female *Anopheles* mosquito. For *Plasmodium*, the passage through the mosquito serves two imperative purposes: a) the insect provides the conditions for sexual reproduction and b) it facilitates transport of the new generation of parasites to the next vertebrate host. Following ingestion by the mosquito as part of a blood meal, male and female gametes emerge from their respective host red blood cells (RBCs) to form zygotes. The zygotes convert into highly polarized motile oocinetes, which actively move out of the bloodmeal by invading and crossing the epithelial cell layer to the basal lamina. Here the oocinetes transform into sessile sporozoite-generating oocytes. The sporozoites are released into the hemocoeel and eventually invade the salivary glands.

Although the mosquito bloodmeal environment provides the obligatory conditions for parasite gametogenesis and subsequent fertilization [1], it also represents an increasingly challenging environment, which causes the majority of the parasites to perish within the first 24 hours in the mosquito bloodmeal [2,3]. The reasons for this attrition include mosquito-derived digesting enzymes and the cytotoxic byproducts of their activity, as well as active immune components of the previous vertebrate host [4]. The mosquito also mounts its own innate immune response against the intruding parasite in the form of antimicrobial peptides as well as cytotoxic reactive oxygen (ROS) and reactive nitrogen species (RNS) [5–8]. Moreover, when the gametes egress from their host RBCs *Plasmodium* effectively switches its life style from intra- to extracellular parasite and consequently exposes itself directly to the hostile conditions in the mosquito bloodmeal. While gametogenesis and zygote formation occur within an hour of ingestion, the zygote-to-oocinete transformation requires up to...
20 hours in the increasingly hostile conditions of the bloodmeal before the mature oocyst is capable of moving through the blood bolus [9].

Yet, despite these adverse conditions, more than 200 million annual clinical episodes of malaria are reported worldwide, the majority of which are direct consequences of mosquito bites [10]. Evidently, a sufficient number of oocysts survive and escape the mosquito bloodmeal and facilitate malaria transmission. This suggests that oocysts possess protective mechanisms that help the Plasmodium oocyst to adapt to rapidly changing environmental conditions and thus to increase the probability of survival, maturation and escape from the mosquito midgut.

Figure 1. Cytosolic antioxidant systems of Plasmodium. Shown is a simplified representation of the Trx- and GSH systems highlighting the roles of the cytosolic peroxiredoxins in asexual Plasmodium. The thioredoxin system (black) and the glutathione system (gray) are indicated. Arrows mark the directions of electron flow. Dashed lines indicate that redox reactions have been shown to take place between proteins in vitro and are also hypothesized to occur in vivo. In asexual parasites the food vacuole (FV) represents a major source for endogenous ROS due to hemoglobin digestion. FV = food vacuole; Exo = exogenous; Endo = Endogenous; Hz = hemozoin.

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Peroxiredoxin Expression in Plasmodium Ookinete

The malaria parasite Plasmodium is transmitted by Anopheles mosquitoes. Within the midgut of the insect, it is exposed to multiple environmental stresses, including cytotoxic reactive oxygen species (ROS). To avoid destruction, the parasite develops into a motile oocyst capable of leaving the midgut. Yet, oocyst development lasts over several hours and requires the parasite to adapt to an increasingly challenging environment. Here we show that oocysts of the rodent parasite Plasmodium berghei during development in the mosquito midgut increase the expression of the protective antioxidant proteins peroxiredoxin-1 (TPx-1) and 1-Cys peroxiredoxin (1-Cys Prx). This upregulation was also inducible in cultured oocysts by challenging them with ROS. This suggests that oocysts actively modulate the expression of their antioxidant proteins in response to the changing conditions in the mosquito. We also found that oocysts lacking TPx-1 (TPx-1KO) upregulated 1-Cys Prx expression significantly earlier than wild type oocysts. This indicates that the TPx-1KO parasites compensate for the loss of TPx-1 by altering the expression pattern of the functionally related 1-Cys Prx. The observed dynamic regulation of the cytosolic antioxidant proteins may help the Plasmodium oocyst to adapt to rapidly changing environmental conditions and thus to increase the probability of survival, maturation and escape from the mosquito midgut.

The thioredoxin system and the glutathione system are indicated. Arrows mark the directions of electron flow. Dashed lines indicate that redox reactions have been shown to take place between proteins in vitro and are also hypothesized to occur in vivo. In asexual parasites the food vacuole (FV) represents a major source for endogenous ROS due to hemoglobin digestion. FV = food vacuole; Exo = exogenous; Endo = Endogenous; Hz = hemozoin.
scavenging endogenous oxidative compounds originating from 
Hb-digestion during intraerythrocytic development.

Here we report on the transcription- and protein-expression 
profiles of Trx-1 and the cytosolic thioredoxin-dependent peroxi-
reodoxins TPx-1 and 1-Cys Prx in developing oocinotes of the 
rodent malaria parasite Plasmodium berghei. We compared expres-
sion over time between parasites developing in the mosquito 
bloodmeal to parasites growing under oocinite culture conditions. 
Our results show that P. berghei strongly increases expression of 
the cytosolic peroxiredoxins both at the transcriptional as well as 
at the protein level when developing in the mosquito bloodmeal. 
Peroxiredoxin upregulation was also inducible in oocinite cultures 
that were previously challenged with exogenous ROS. We 
furthermore show evidence that 1-Cys Prx expression is altered 
in TPx-1KO oocinotes possibly to compensate for the loss of TPx-
1. We hypothesize that a flexible antioxidant stress response, 
regulated by factors in the mosquito bloodmeal, contributes to 
the adaptation of the developing oocinotes to the challenging 
environment and increases the probability of parasite survival 
and transmission.

Results

We hypothesize that the Plasmodium oocinite regulates antiox-
idant protein expression in response to environmental changes in 
the mosquito bloodmeal in order to maximize the probability of 
survival, maturation and escape.

tpx-1 and 1-cys prx transcription is upregulated in the 
mosquito blood meal

To assess the effect of the mosquito bloodmeal on antioxidant 
enzyme expression in Plasmodium, we compared the transcription 
profiles of trxr, trx-1, tpx-1 and 1-cys prx (Table 1) in parasites 
that developed either in bloodfed mosquitoes or in oocinite cultures 
over the course of 24 hours (Figure 2). Female Anopheles stephensi 
mosquitoes were allowed to feed on mice previously infected with 
P. berghei ANKA 2.34. Bloodfed mosquitoes were collected at 3, 6, 
12 and 24 hours post infectious bloodmeal (pIBM) and midguts 
were dissected for total RNA isolation. In parallel, oocinite 
cultures were prepared and samples were taken at 3, 6, 12 and 
24 hours post culture setup for total RNA isolation. Candidate 
gene expression relative to 18 s rRNA was assessed using a SYBR 
green based RT-qPCR [34,35]. Transcription profiles were 
generated using the 3 hours sample as a reference point. For 
simplicity, samples from parasites grown in culture will be referred 
to as culture-derived and samples from parasites isolated from 
dissected bloodfed mosquito midguts as mosquito-derived. 

It was observed in the transcriptional profiling of culture-
derived parasites that only tpx-1 and 1-cys prx were significantly 
modulated over time (Figure 2). tpx-1 showed significant 
upregulation at 6 hours (4-fold) and at 12 hours (6-fold) post 
culture setup. A downregulation was seen between the 12 and 24-
hour time points for tpx-1. 1-cys prx transcription was not altered 
during the first 12 hours but showed an upregulation of 2.5 fold 
between the 12 and 24 hour timepoints. Unlike the peroxiredoxin 
genes, trxr and trx-1 transcript abundance did not significantly 
change under culture conditions. In contrast, mosquito-derived 
parasites exhibited a greater upregulation of the peroxiredoxin 
transcripts. tpx-1 transcripts increased between 3 and 6 hours 
pIBM (2.5-fold) and more rapidly between 6 and 12 hours (>10-
fold) pIBM. This was followed by a steady downregulation 
between 12 and 24 pIBM. The resulting profile was similar to that 
observed in culture-derived oocinotes. Conversely, 1-cys prx 
expression showed a large increase (>13-fold) between 3 and 
6 hours pIBM. Significantly increased transcript levels were also 
detected at 12 (11-fold) and 24 hours (4-fold) pIBM. trx-1 was 
initially upregulated, showing a single peak at 6 hours pIBM (3.7-
fold). This was followed by an equal downregulation between 6 
and 12 hours. trx-1 expression levels were not significantly 
modified for the rest of the developmental time frame. Comparable to its transcription profile in culture-derived 
parasites, trx-1 did not exhibit any significant modulation under 
bloodmeal conditions. The transcriptional profiles of 1-cys prx, 
tpx-1, and trxr in parasites developing in mosquitoes demonstrate 
that factors in the bloodmeal environment affect regulation of 
antioxidant gene expression.

1-Cys Prx protein is expressed in mosquito- derived but 
not in culture- derived oocinotes

The parasite populations investigated in oocinite cultures and 
bloodfed mosquitoes consisted of mixed developmental stages, 
including oocinotes. Since it is essential for the oocinite to 
withstand the increasingly hostile conditions in the bloodmeal, we 
examined whether the observed upregulation of the peroxiredoxin 
genes would be directly translated to the functional protein level in 
developing oocinotes. Samples were taken at 3, 12 and 24 hours 
from either oocinite cultures or bloodfed mosquitoes. Following 
fixation, the samples were probed with protein-specific antibodies 
against Trx-1, TPx-1 or 1-Cys Prx and analyzed using laser 
scanning confocal or epifluorescence microscopy. 

Trx-1 was continuously and ubiquitously expressed in all 
developmental stages of culture-derived as well as mosquito-
derived oocinotes (Figure 3A). The uniform distribution of the 
protein in oocinotes indicated cytosolic localization. Quantitative 
analysis of relative fluorescence (RF) in mature oocinotes 
(24 hours) confirmed that there was no significant differential 
expression of Trx-1 protein between culture-derived and mosqui-
to-derived oocinotes (Figure 3A; median RF = 1.7 vs. 2.1). 

TPx-1 was also detected in all oocinite stages in culture-derived 
as well as mosquito-derived parasites (Figure 3B). The protein was 
uniformly distributed in the cytosol of the developing oocinotes 
with the exception of early retort stages from oocinite cultures 
(3 hour pIBM). Here TPx-1 was detected in the main body of the 
parasite but not in the apical protrusion. Subsequent develop-
mental stages showed the protein throughout the oocinite. RF 
analysis showed higher fluorescence intensities in mosquito-
derived oocinotes when compared to culture-derived oocinotes 
(median RF = 1.2 vs. 2.2). However, quantitative analysis revealed 
that this observed difference was not statistically significant 
(p = 0.1). 

Unlike Trx-1 and TPx-1, 1-Cys Prx was not expressed in early 
culture-derived oocinotes (3 and 12 hours) and only at low levels in 
mature oocinotes (24 hours) (Figure 3C). In contrast, in mosquito-

| Table 1. Candidate genes of the cytosolic thioredoxin system of P. berghei. |

<table>
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<th>Putative</th>
<th>Abbr.</th>
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<td>[19,22,23]</td>
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<td>1-Cys Prx</td>
<td>PBANKA_122800</td>
<td>Q4YV50</td>
<td>[23,35]</td>
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doi:10.1371/journal.ppat.1003136.t001
derived ookinetes high cytosolic expression of 1-Cys Prx was observed at 12 and 24 hours pIBM (Figure 3C). No signal was detected in early retort stages (3 hours). RF analysis confirmed that the observed higher expression of 1-Cys Prx in mosquito-derived ookinetes when compared to culture-derived ookinetes at 24 hours pIBM was statistically significant (median RF = 0.2 vs. 2.7; p<0.05). Interestingly, an accumulation of 1-Cys Prx protein at the apical ends of mature mosquito-derived ookinetes was observed (Figure 3C). The reason for this localized buildup or its biological significance is not yet clear.

Upregulation of peroxiredoxins is inducible in ookinete culture

Our data indicate that P. berghei ookinetes actively respond to changes in the bloodmeal of the mosquito midgut by increasing expression of the cytosolic peroxiredoxins. However, the mosquito midgut represents a complex environment that changes throughout the course of ookinete development. This includes an increase of reactive oxygen species (ROS), such as superoxide radicals [36,37] and peroxide compounds [6,7]. We hypothesized that the observed upregulation of cytosolic peroxiredoxins was induced in
We therefore examined the effect of ROS on peroxiredoxin transcription in ookinete cultures by exposing the parasites to increasing concentrations of paraquat (PQ), a superoxide producer [38,39]. 12-hours ookinete cultures were challenged with PQ concentrations between 0.1 and 100 μM [30]. Parasites were harvested after 45 minutes and total RNA was extracted for RT-qPCR analysis (Figure 4). The ratios of \( \text{trxr} \), \( \text{trx-1} \), \( \text{tpx-1} \), and \( \text{1-cys prx} \) transcription levels in treated and untreated samples were determined. Low and moderate PQ concentrations (0.1 to 10 μM) did not alter

<table>
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<td>12 Hours</td>
<td>24 Hours</td>
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**Figure 3. Expression of the Trx-1, TPx-1 and 1-Cys Prx in developing ookinetes from culture or from mosquito bloodmeal.** Samples were isolated from ookinete cultures and blood fed mosquitoes at indicated time points and prepared as described in material and methods. Fixed samples were probed with A) anti-PbTrx-1, B) anti-Pf TPx-1 or C) anti-PbTPx-1. Primary antibodies were probed with secondary antibody coupled to AF 488 (Molecular Probes). Samples were counterstained with the nuclear dye TO-PRO-3 (Life technologies). The source of the ookinetes from either ookinete cultures or bloodfed mosquito midguts is indicated. Images are merged and overlaid onto the respective DIC image. White arrows indicate accumulation of Trx-1, TPx-1 or 1-Cys Prx at the apical ends of developing ookinetes. Scale bar = 5 μM. Quantitative Relative Fluorescence Analysis: Shown is the median boxed by the first and third quartiles with minimum and maximum values displayed as whiskers. Mann-Whitney U-Test was performed with samples of culture-derived ookinetes compared to mosquito-derived ookinetes (p = 0.4551 for Trx-1; p = 0.1061 for TPx-1; p = 0.0413 for 1-Cys Prx). Below each graph are representative images from IFA-epifluorescence experiments. doi:10.1371/journal.ppat.1003136.g003
transcription levels of any of the candidate genes. At 100 μM PQ the transcription levels of tpx-1 and 1-cys prx significantly increased by 2.3-fold and 6.7-fold, respectively. Conversely, trx and trx-1 did not exhibit significant differential expression in treated and non-treated samples. This data supports our hypothesis that ROS present in the bloodmeal of the mosquito contribute to upregulation of the cystolic peroxiredoxins in the parasites.

The expression of 1-Cys Prx protein is altered in TPx-1KO ookinetes

TPx-1 has been hypothesized to play a central role in the antioxidant defense of Plasmodium against exogenous stresses [19,30]. P. berghei TPx-1 knock-out parasites (TPx-1KO) exhibit a reduced number of gametocytes and oocytes, while the number of ookinetes and infective sporozoites remain comparable to the wild-type (WT) parasite. Compensatory mechanisms have been suggested [31,32]. Considering our data we hypothesized that 1-Cys Prx compensates for the loss of TPx-1. We therefore investigated the protein expression profiles of Trx-1 and 1-Cys Prx in culture-derived as well as mosquito-derived TPx-1KO ookinetes.

Trx-1 expression and distribution was comparable in culture-derived as well as in mosquito-derived TPx-1KO ookinetes (Figure 5A). Quantitative RF analysis also indicated comparable Trx-1 protein levels between TPx-1KO and WT ookinetes.

Interestingly, an accumulation of Trx-1 protein at the apical end of the protrusion was observed in early and late retorts in culture-derived as well as in mosquito-derived TPx-1KO ookinetes. These accumulations were not observed in mature ookinetes.

1-Cys Prx was not detected in early or late retort stages of culture-derived TPx-1KO ookinetes (Figure 5B). However, mature TPx-1KO ookinetes showed a strong cytosolic expression of 1-Cys Prx (median RF = 1.2), which was significantly higher than in culture-derived WT ookinetes at 24 hours. In mosquito-derived TPx-1KO ookinetes, 1-Cys Prx was already detected at significant levels in early retort stage (3 hours pIBM). Expression was maintained in the subsequent developmental ookinete stages (median RF = 2.2) (Figure 5B). Accumulation of 1-Cys Prx was also observed in the apical region of the mosquito-derived TPx-1KO ookinetes.

Discussion

Cellular adaptation to new environmental challenges generally occurs on the transcriptional level and leads to a direct increase in functional protein [40]. The developing Plasmodium ookinete in the mosquito bloodmeal has to withstand an increasingly hostile environment for several hours before it reaches its motile stage. The bloodmeal contains a complex combination of cytotoxic compounds, including ROS and RNS ([5–7,36,37]. This would require the ookinete to initiate its stress response in order to maximize its chances of survival. Recent reports have shown that asexual P. falciparum parasites respond to oxidative challenges by upregulating various stress response genes and proteins, including trx, tpx-1 and 1-cys prx [22,41]. This indicates that the parasite has the capacity to respond to sudden environmental changes.

Our data confirm and expand on these observations. We have shown for the first time that antioxidant gene expression is transiently upregulated during parasite development in the mosquito bloodmeal. This upregulation occurred rapidly and peaked between 6 and 12 hours pIBM followed by a steady decline to almost initial levels. In contrast, under ookinete culture conditions we found tpx-1 upregulation to be less strong, while an increase in 1-cys prx transcription was not detected before 24 hours post culture setup. It is likely that the differential expression between mosquito-derived and culture-derived parasites is directly linked to the environmentally more stressful conditions in the mosquito bloodmeal. Complementing this hypothesis are recent observations showing that mosquito-derived ROS and RNS increase in the mosquito bloodmeal over time and reach a maximum at about 12 hours post bloodmeal (pBM), followed by a continuous decrease [7,37,42]. The transcription profiles of trx, tpx-1 as well as 1-cys prx mirror these observations and suggest that oxidants in the bloodmeal stimulate antioxidant gene upregulation. Our results showing induction of tpx-1 and 1-cys prx upregulation in ookinete cultures in response to ROS challenge further support these observations.

We recognize that recent antioxidant studies on P. falciparum were conducted on synchronized asexual parasite cultures [22,41], whereas our transcription data originates from mixed parasite populations under different and, in the case of the mosquito bloodmeal, varying environmental conditions. Consequently, the population dynamics differ significantly between the mosquito bloodmeal and the ookinete culture environment. For example, the majority of the parasites ingested by the mosquito will be dead or dying by 24 hours pIBM [2], while under culture conditions most parasites will survive the first 24 hours. The attrition of parasites in the bloodmeal may also contribute to the decrease in...
the transcription levels of *trx*, *tpx-1* and *1-cys prx*. Yet, the expression profiles reported here were highly reproducible for each gene, indicating that most *Plasmodium* life stages respond to the bloodmeal, including developing ookinetes.

The observed upregulation of the peroxiredoxin genes in culture-derived parasites is most likely due to a slow accumulation of ROS and RNS in the culture medium. Sources of pro-oxidants include actively metabolizing cells in addition to dead as well as dying cells. Remarkably, *trx-1* did not exhibit significant modulation under either experimental condition. Yet, *trx-1* transcripts were detected at high abundance in all investigated samples (Figure S1). This may reflect the central role of *trx-1* in the redox metabolism of the malaria parasite [11,12].

The fact that the observed transcriptional and translational upregulation of each gene occurred simultaneously makes it unlikely that these peroxiredoxins are post-translationally regulated [43,44]. It was furthermore important to show that the observed antioxidant response in the mixed parasite population occurs in the developing ookinete since the main role of this stage is to escape the blood bolus and establish infection of the mosquito vector.

Here we show for the first time expression of *Trx-1*, *TPx-1* and *1-Cys Prx* in developing *Plasmodium* ookinetes [26,29,35]. The increase in *TPx-1* and *1-Cys Prx* protein levels in mosquito-derived ookinetes corresponded very well with the observed transcriptional profile, with the distinction that, following upregulation, protein levels were maintained in all subsequent developmental stages. *Trx-1*, on the other hand, was present at continuous levels throughout ookinete development in mosquito-derived as well as culture-derived parasites, which was also in accordance with the observed transcriptional profile. These data strongly indicate that the expression of the thioredoxin-dependent antioxidant system is not post-transcriptionally regulated in developing ookinetes. Furthermore, ookinetes directly respond to the changing conditions in the bloodmeal by modulating antioxidant protein levels. This supports our hypothesis that the parasite possesses an environmentally regulated oxidative stress response.

Figure 5. Expression of *Trx-1* and *TPx-1* in culture-derived and mosquito-derived TPx-1KO ookinetes. TPx-1KO samples were isolated from ookinete cultures and blood fed mosquitoes at indicated time points and prepared as described in material and methods. Fixed samples were probed with A) anti-PbTrx-1, B) anti-PbTPx-1. Primary antibodies were probed with secondary antibody coupled to AF 488. Samples were counterstained with the nuclear dye TO-PRO-3 (Life technologies). The source of the ookinetes from either ookinete cultures or bloodfed mosquito midguts is indicated. Images are merged and overlaid onto the respective DIC image. White arrows indicate accumulation of *Trx-1* or *1-Cys Prx* at the apical ends of developing ookinetes. Scale bar = 5 μM. Quantitative *Relative Fluorescence Analysis*: Shown is the median boxed by the first and third quartiles with minimum and maximum values displayed as whiskers. Mann-Whitney U-Test was performed with samples compared to WT culture: v TPx-1 KO Culture (*Trx-1*: p = 0.6519; *1-Cys Prx*: p = 0.0124); v TPx-1 KO mosquito (*Trx-1*: p = 0.5777; *1-Cys Prx*: p = 0.0473 ). (*) indicates that *1-Cys Prx* is statistically higher expressed in TPx-1KO ookinetes. Below are representative images from IFA-epifluorescence experiments. doi:10.1371/journal.ppat.1003136.g005
The continuous and ubiquitous expression of Trx-1 observed in gametocytes, gametes and ookinetes expands on previous reports that show ubiquitous Trx-1 expression in asexual stages of *P. falciparum* (PfTrx-1) [26]. Also mass-spectroscopy based proteomics data indicate Trx-1 expression in gametocytes and mature cultured ookinetes of *P. berghei* and *P. gallinaceum*, respectively [45,46]. Our data therefore support the proposed central role of Trx-1 in the metabolism of the malaria parasite [11,12,47,49]. Since catalase and glutathione peroxidase are absent in *Plasmodium* [18] it is likely that peroxiredoxins and thus Trx-1 play a more prominent role in the antioxidant defense of the parasite [15]. Indeed, recent studies strongly indicate that PTrx-1 interacts with multiple target proteins in vitro as well as in vivo, most notably PTP-1 and PfL-Cys Prx [12,14,28].

Our data also show constitutive expression of TPx-1 in *P. berghei* mosquito-derived and culture-derived developing ookinetes as well as in gametocytes and exflagellating gametes. These observations expand on previous studies that show TPx-1 expression in cultured ookinetes of *P. yoelii* [35]. Mass spectroscopy-based proteomics data also indicates TPx-1 expression in cultured mosquito stages of *P. berghei* [45] and *P. gallinaceum* [46]. These data support the hypothesis of *tpx-1* as a housekeeping antioxidant gene that is primarily active against exogenous ROS and RNS [15,23]. As such, TPx-1 expression would be under developmental rather than environmental control and thus relatively independent of exogenous conditions. However, challenging parasites with exogenous ROS leads to the upregulation of *tpx-1* transcript as well as protein [22,41]. We confirmed these findings when challenging culture-derived parasites with ROS. Our studies also show that *tpx-1* transcription and translation is elevated in mosquito-derived parasites when compared to culture-derived parasites, indicating that environmental conditions have an effect on the regulation of this gene. Taken together, these observations suggest that the parasite is continuously exposed to varying concentrations of ROS and RNS throughout its complex life cycle.

Differential regulation at the transcriptional and the translational level under different environmental conditions was most evident for 1-Cys Prx. While the enzyme is hardly detectable in developing cultured ookinetes, we found strong fluorescence signals in ookinetes isolated from mosquito bloodmeals. The timely increase in protein signal coincided with transcriptional upregulation of the 1-cys prx gene, indicating a direct response of the parasite to the bloodmeal conditions. A lack of 1-cys prx upregulation and protein expression was previously reported for gametocytes and cultured ookinetes of several *Plasmodium* species, including *P. berghei* [35,45,46]. Conversely, 1-Cys Prx is strongly expressed during asexual development of the parasite [29,35,49]. It was also shown that the enzyme interacts with heme in vitro [33]. It has therefore been hypothesized that 1-Cys Prx constitutes a bloodstage specific enzyme involved in the detoxification of ROS that originate from the parasites’ food vacuole during hemoglobin (Hb) digestion [15,33]. The absence of 1-Cys Prx in sexual and mosquito stages would then be plausible, as Hb digestion has been completed.

We demonstrate that 1-Cys Prx expression is not limited to the asexual stages and may therefore have a broader function as an antioxidant enzyme throughout the parasite life cycle. Furthermore, it can be upregulated by the parasite in response to changing environmental conditions, which is evident from the diverse transcription and protein expression profiles in mosquito-derived and culture-derived parasites. Our challenge data also suggest that 1-cys prx upregulation may not only be dose-dependent but that a threshold exists that triggers the response in the parasite. Hence, no 1-Cys Prx expression was detected in culture-derived ookinetes as ROS build-up is comparatively slow and initially buffered by TPx-1, which shares substrate specificities with 1-Cys Prx [19,23,49]. In contrast, ROS concentrations rise more quickly and more strongly in the bloodmeal of the mosquito [5,37] and thus trigger an adaptive antioxidant response in the ookinetes involving both peroxiredoxins.

This hypothesis is further supported by our studies on TPx-1KO parasites. The fact that the disruption of the *tpx-1* gene is not lethal in either *P. falciparum* [30] or in *P. berghei* [31,32] suggested that compensatory mechanisms facilitate parasite survival. While *Trx-1* expression did not vary significantly between WT and KO ookinetes, we found that 1-Cys Prx expression was significantly altered in the KO ookinetes. Specifically, enzyme expression was detected significantly earlier in mosquito-derived ookinetes. We also found higher levels of 1-Cys Prx protein in mature culture-derived KO ookinetes, which we attribute to an early upregulation between 12 and 24 hours post culture setup. We suggest that the absence of TPx-1 leads to a quicker build-up of ROS and RNS in the parasite and thus triggers an earlier upregulation of 1-Cys Prx in the KO parasite. Considering our data and the overlapping substrate specificities of the cytosolic peroxiredoxins, we hypothesize that 1-Cys Prx significantly contributes to the compensation for the loss of TPx-1 in the KO parasites. A recent report shows that the mitochondrial homolog of TPx-1, *P. berghei* thioroxygenase peroxidase-2 (TPx-2), is not essential for parasite development in the mammalian host or in the mosquito vector [21,50]. While the lack of a phenotype in the TPx-2KO parasites may be due to compensatory function of other mitochondrial redox proteins [26] it is also possible that ROS leaking from the mitochondria may be detoxified by the cytosolic system. It will therefore be highly informative to investigate the impact of TPx-2KO on the expression of other redox genes and their corresponding proteins.

The present work shows that the *Plasmodium* ookinete utilizes an inducible antioxidant response during adaptation to the challenging and changing conditions in the mosquito bloodmeal. This indicates that TPx-1 and 1-Cys Prx play overlapping roles, and that 1-Cys Prx has the capacity to compensate for the loss of TPx-1. While TPx-1 and 1-Cys Prx have similar antioxidant functions, the respective transcriptional as well as protein expression profiles indicate that these enzymes may be regulated independently. For example, TPx-1 may be inducible by lower ROS concentrations, which would explain its profile in culture-derived parasites. 1-Cys Prx upregulation on the other hand requires higher ROS concentrations; hence the late transcriptional increase in ookinete cultures. Conversely, in the mosquito bloodmeal ROS concentrations rise much more quickly and thus trigger an earlier 1-cys prx upregulation. Alternatively, expression of each peroxiredoxin may be oxidant specific and certain compounds accumulate faster in the mosquito bloodmeal than in the culture medium, such as paraoxonitrile (ONO−) [37]. Upregulation of 1-Cys Prx seems to be ookinete specific, as we detected only low levels of the enzyme in gametocytes and no expression in male gametes in WT as well as in KO parasites.

Considering our data we also propose that a ROS sensing mechanism must be in place to facilitate antioxidant stress response in *Plasmodium*. Oxidant sensor proteins of the glutathione peroxidase family of proteins have recently been implicated in the oxidative stress response of the yeast *Saccharomyces cerevisiae* [51], *Drosophila melanogaster* [52] and human cells [53]. Although the *Plasmodium* genomes do not seem to contain a genuine glutathione peroxidase [11,12,15] a glutathione peroxidase-like thioroxygenase peroxidase (TPxGal) has been characterized in *P. falciparum* [24,26]. TPxGal shares significant sequential as well as structural homologies with the putative redox sensor proteins described in yeast, fruit fly
and human. Work in our lab is under way to investigate the potential signaling function of Plasmodium TPxGl.

It has been stated that the mosquito stages of Plasmodium, specifically the ookinete, represent attractive targets for transmission intervention strategies [54]. It is therefore imperative to understand the mechanisms the ookinete utilizes to adapt to the adverse environment in the mosquito bloodmeal as these may hold the key to close the population bottleneck and to prevent transmission. Our results highlight the general significance of the oxidative stress response for all stages of the malaria parasite. They also emphasize that environmental conditions, especially for the “unprotected” mosquito-stages, must be taken into account when investigating the parasite-vector relationship in search for potential candidate genes.

Materials and Methods

Ethics statement

All experimental protocols were approved by the Institutional Animal Care and Use Committee (IACUC) of Loyola University Chicago (Protocol #148) following the National Institutes of Health guidelines for animal housing and care.

Parasite maintenance and mosquito infections

Plasmodium berghei ANKA 2.34 wild type (WT) and peroxiredoxin-1 knock-out (TPx1KO) [31] parasites were maintained in Harlan ND4 mice for a maximum of four serial passages and passed through Aedes stephensi mosquitoes. Mosquitoes were reared under standard conditions (26°C, 80% RH, 12 hrs light-dark cycle, 5% sucrose solution). Female mosquitoes 5-10 days old were allowed to feed for 15 min on mice infected with WT or TPx-1KO parasites (10%). Bloodfed mosquitoes were maintained at 21°C, 80% RH, 12 hrs light-dark cycle, 5% sucrose solution. Bloodfed mosquitoes were removed at experimental time points and midguts were dissected and prepared for fixation.

Ookinete cultures and challenges

Gametocytes, exflagellating male gametes and ookinetes were obtained following established protocols [55]. Briefly, gametocytes were obtained by pretreating mice with phenylhydrazine (Phz) and subsequently infecting them with WT or TPx-1KO parasites. On day two post-infection, gametocyte-rich blood was harvested and prepared for fixation. To obtain gametes, we incubated gametocyte-rich blood in ookinete medium for 15 min before fixation. Exflagellation rates peaked at two days post infection (>15 exflagellation/20× field). For ookinetes, the blood of three infected mice per experimental culture setup was pooled, diluted 1:5 with ookinete medium (RPMI1640, pH 8.4), and incubated at 19°C for a maximum of 24 hrs. Ookinete development was followed by Giemsa-stained smears (average yield: 1 x 10^6 ookinetes/ml culture) and harvested samples were directly fixed and prepared for confocal analysis. For ookinete cultures, we largely implemented protocols as described by Rodriguez et al. [55] with the following modifications: exflagellation rates of parasites from phenylhydrazine treated mice already peaked on day 2 post infection; blood of three infected mice per experimental culture setup was diluted 1:5 with ookinete medium (RPMI1640, pH 8.4) and incubated at 19°C for a maximum of 24 hours; ookinete development was followed by Giemsa stained smears; culture samples were directly transferred into Tri-Reagent (MRC) for total RNA extraction. For the challenges: Increasing concentrations of paraquat (PQ) were added to ookinete cultures at 12 hours post setup; parasites were harvested following a 45 min incubation time at 19°C and total RNA was prepared for 1st strand synthesis and RT-qPCR analysis.

Quantitative real time RT-PCR and data analysis

Total RNA from bloodfed mosquito midguts and ookinete cultures was extracted using Tri-Reagent (MRC) according to manufacturer’s instructions. Isolated RNA was treated with DNase I (Ambion) and subsequently quantified using the Qubit RNA assay kit and the Qubit fluorometer (Invitrogen). RNA samples were either immediately used for cDNA synthesis or flash frozen and stored at −80°C. cDNA was synthesized from total RNA with the High Capacity RNA-to-cDNA kit (Applied Biosystems) using random hexamer primers. Sequences of target genes for primer design were acquired from Plasmodb (plasmodb.org). For the qRT-PCR, target-specific primers with Tm = 60°C were designed to yield amplicons between 80 and 140 bp (table S1). RT-qPCR was performed on a StepOnePlus machine (Applied Biosystems) using the Fast SYBR Green Master Mix in 10 μl reaction volumes. Each sample was run in triplicates and yielded highly comparable C values (cycle threshold). No primer dimers were detected and amplicons exhibited optimal efficiencies. To test specificity, all primer pairs were tested on uninfected mouse blood, non-fed and uninfected bloodfed mosquitoes. No amplification products could be detected. Expression data was subsequently analyzed with the StepOne Software v2.2 (Applied Biosystems) and normalized against the expression of 18 s rRNA A-Type, which has been established as an internal standard for expression analysis in Plasmodium mosquito stages [29,54]. For analysis of time course expression data, the ΔΔCt method was used with the earliest experimental time point as a reference sample (RQ = 1). Target gene transcription levels were calculated in relation to 18 s rRNA expression by determining the ΔΔCt value for each primer pair and subtracting the measured Ct value of the target from the Ct value of the 18 s rRNA control. ΔCt values were calculated to relative transcription levels using the formula 2−ΔΔCt. The Mann-Whitney U test was used to determine if gene expression at 6, 12, and 24 hours was significantly different from the gene expression at 3 hours (reference point). The Mann-Whitney U test was conducted on each candidate gene from both mosquito-derived and from culture-derived parasites. Significance was assessed at p<0.1 due to the small sample sizes. The non-parametric Mann-Whitney U test was the appropriate statistical analytical approach to use on this dataset due to the violated assumptions of independence in the data and the small and unbalanced sample sizes in gene expression collected at 3, 6, 12, and 24 hours. Statistical analyses were performed in R [56].

Sampling and fixation of parasite cells

Bloodstage parasites were isolated from tail vein blood from mice and were resuspended in RPMI 1640 pH 7.4 media with heparin pre-warmed to 37°C and allowed to adhere to 0.01% poly-L-lysine coated glass slides at 37°C. Gametogenesis was induced by resuspending tail blood in RPMI 1640 pH 8.2 media with heparin pre-warmed to 19°C and allowed to adhere to 0.01% poly-L-lysine coated glass slides at 19°C. Cells were then fixed in 3% para-formaldehyde with 0.1% glutaraldehyde at 4°C. Cultured ookinetes were taken at experimental time points 3, 12 and 24 hours post culture setup. RBCs were lysed with 0.17 M NH₄Cl. Cells were then resuspended in 0.05 M Tris 0.9% NaCl pH 8.2 and allowed to adhere to 0.01% poly-L-lysine coated glass slides. Cells were then fixed in 4% para-formaldehyde at 4°C. Midgit ookinetes were isolated from dissected midguts by crushing midguts with
mortar and pestle in 0.05 M Tris 0.9% NaCl pH 8.2 and allowed to adhere to 0.01% poly-L-lysine coated glass slides. Cells were then fixed in 4% para-formaldehyde at 4°C. Slides were stored at −80°C.

Immunofluorescence staining

Slides were allowed to permeabilize and thaw in 0.05% Triton X-100 in 1× PBS. Slides were then blocked with 4% Bovine Serum Albumin. Epitopes for each protein of interest were retrieved by incubating slides in 10 mM Sodium Citrate, 0.05% Tween 20, pH 6.0 at 95°C. Autofluorescence was quenched by incubating slides in 1% Sodium Borohydride in 1× PBS. Respective antigens were detected with custom polyclonal antisera. (1:300) raised in rabbits (TPx-1, Trx-1, 1-Cys Prx, Open Biosystems) and labeled with anti-rabbit Alexa Fluor 488 (1:1000) (Molecular Probes). Slides were counterstained with TO-PRO 3 (Invitrogen) and sealed with glass cover slips in ProLong Gold AntiFade with DAPI mounting medium (Invitrogen).

Confocal microscopy and quantitative analysis of relative fluorescence

Images were acquired on FluoView FV1000 Confocal Laser Scanning Microscope (Olympus). Images were taken in XY plane and compressed around the Z axis. Images in figures are representative of 50 ookinetes per time point and environment. FV10-ASW 3.0 microscopy software (Olympus) was used to analyze the samples. Quantitative fluorescence images were acquired on Zeiss Axiovert 200 m using a Axiocam MRm. Images were processed using Axiovision Rel. 4.3 software. For quantitative analysis of ookinetes, DAPI was exposed by exposure at 359 nm for 20 milliseconds and AF 488 was excited by exposure at 488 nm for 350 milliseconds. Fluorescent emissions were recorded at 461 and 530 nm respectively. Images were then quantitated using Image J Software (National Institute of Health). Fluorescent Intensities for DAPI and AF488 were assessed by using the histogram function. The ratio of candidate protein fluorescence (AF 488) to DAPI fluorescence was calculated for WT culture (n = 50), TPx-1 KO culture (n = 50), WT mosquito (n = 25), and TPx-1 KO (n = 23).

Accession numbers

PlasmoDB ID numbers for reported genes and proteins are as follows: Thioredoxin Reductase (TrxR;PBANKA_082470), Thioredoxin-1 (PBANKA_132090); Thioredoxin peroxidase-1 (TPx-1; PBANKA_122800), 1-Cysteine Peroxiredoxin (1-Cys Prx; PBANKA_122800).

Supporting Information

Figure S1 Relative transcript abundance of target gene transcripts in culture-derived and in mosquito-derived parasites at the 12 hours time point. RT-qPCR data show relative quantity of target gene transcripts normalized to 18 s rRNA A-type expression [29,34]. The delta Ct values were converted using $2^{-\Delta\Delta Ct}$ x 10^6. Shown are mean values of 3 independent experiments. Error bars indicate STDEV.

(TIF)

Figure S2 Cloning, expression and purification of P. berghei Trx-1 and 1-Cys Prx and specificity of the polyclonal antibodies. A) Translation maps of putative Trx-1 (PBANKA_132090) and putative 1-Cys Prx (PBANKA_122800). The start codons are indicated in bold. Forward and reverse primers are underlined. The characteristic active site motifs including the peroxidatic cysteines are highlighted in yellow. B) Recombinant protein expression and purification. Gene specific primers were designed and PCR was performed to amplify the coding sequences using the following conditions: 40 cycles of 95°C for 30 s, 1 min at 54°C, and 45 s at 63°C. This was followed by a 5 min final extension at 63°C. The verified PCR products were ligated into pQE30 expression vectors (Quagen) and subsequently transformed into E. coli M15 expression cells. Recombinant protein expression was induced by adding isopropyl thio B-galactoside (IPTG) to a final concentration of 1 mM. Bacteria were harvested after a 24-hour incubation time at 37°C. Recombinant proteins were purified via a Ni-NTA column (Life technologies). Protein purity was confirmed via sodium dodecyl sulfate (SDS) polyacrylamide gel electrophoresis. Protein concentrations were assessed via Bradford Assay. Purified recombinant Trx-1 (top) 1-Cys Prx (bottom) from E. coli SDS gel analysis: lane 1) protein ladder, lane 2) 10% SDS gel showing purified HIS-tagged rPb 1-Cys Prx. Lanes 3 to 5 are Western blots testing the primary antibody-containing rabbit antiserum (1/500); lane 3) protein ladder, lane 4) Western blot on purified HIS-tagged rPb 1-Cys Prx, lane 5) Western blot on P. berghei lysate of mixed asexual stages from mouse blood.

(TIF)

Figure S3 Assessment of nuclear stains in ookinetes. The fluorescence intensity of nuclear dyes DAPI and TO-PRO-3 were compared between ookinete nuclei from mixed populations of culture and mosquito (n = 15 each). DAPI (blue, left panel) was selected as the nuclear dye of choice for QF due to its lower standard error when compared to TOPRO-3 (red, right panel) (SE = 3.86 vs. 14.57).

(TIF)

Figure S4 Protein expression of Trx-1, TPx-1 and Peroxiredoxins in gametocytes and male exflagellating gametocytes. P. berghei gametocytes A) and exflagellating microgametes B) were differentiated by morphology. Polyclonal antiseria specific for target protein Thioredoxin-1 (Trx-1), Peroxiredoxin-1 (TPx-1) and 1-Cys Peroxiredoxin (1-Cys Prx) were labeled with donkey anti-rabbit AF 488 (Molecular Probes). Cells are counterstained with TO-PRO-3. Images are merged and overlaid onto the respective DIC image. M designates merozoites also pictured and G designates gametocyte. Scale bar indicates 5 μM.

(TIFF)

Table S1 Orthologous genes in P. falciparum and P. berghei primer sequence information. A) Orthologous genes in P. falciparum. B) P. berghei RT-qPCR primer sequences. C) Protein expression primer sequences.

(DOCX)

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Author Contributions

Conceived and designed the experiments: BAT SMK. Performed the experiments: BAT JJT MAL AZ RK. Analyzed the data: SMK BAT SS ES SIK. Contributed reagents/materials/analysis tools: SMK ES SIK. Wrote the paper: SMK BAT.
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