Sequential Membrane Rupture and Vesiculation during *Plasmodium berghei* Gametocyte Egress from the Red Blood Cell

Maria Andreadaki\(^1\),\(^2\), Eric Hanssen\(^3\),\(^4\), Elena Deligianni\(^2\), Cyrille Claudet\(^5\), Kai Wengelnik\(^6\), Vanessa Mollard\(^7\), Geoffrey I. McFadden\(^7\), Manouk Abkarian\(^5\), Catherine Braun-Breton\(^6\) & Inga Siden-Kiamos\(^1\)

Malaria parasites alternate between intracellular and extracellular stages and successful egress from the host cell is crucial for continuation of the life cycle. We investigated egress of *Plasmodium berghei* gametocytes, an essential process taking place within a few minutes after uptake of a blood meal by the mosquito. Egress entails the rupture of two membranes surrounding the parasite: the parasitophorous vacuole membrane (PVM), and the red blood cell membrane (RBCM). High-speed video microscopy of 56 events revealed that egress in both genders comprises four well-defined phases, although each event is slightly different. The first phase is swelling of the host cell, followed by rupture and immediate vesiculation of the PVM. These vesicles are extruded through a single stabilized pore of the RBCM, and the latter is subsequently vesiculated releasing the free gametes. The time from PVM vesiculation to completion of egress varies between events. These observations were supported by immunofluorescence microscopy using antibodies against proteins of the RBCM and PVM. The combined results reveal dynamic re-organization of the membranes and the cortical cytoskeleton of the erythrocyte during egress.

Egress from the host cell is essential for the completion of the life cycle of intracellular pathogens. *Plasmodium* parasites, alternating between intracellular stages and extracellular invasive stages, must exit host cells during key transition stages. One of these is gametogenesis, which takes place during transmission to the mosquito. During a brief interval, male and female gametocytes, having developed inside the parasitophorous vacuole (PV) in the red blood cell (RBC), are released. Microscopy studies suggest that gametocyte egress is similar to merozoite egress in that the PV membrane (PVM) ruptures first and the RBC membrane (RBCM) opens later, which is consistent with the so called inside-out model of egress\(^1\).\(^2\).

The PVM is formed during invasion of the red blood cell and is believed to contain lipids originating both from the host cell and secreted from the invading parasite (for a review see\(^3\)). A limited number of proteins of parasite origin are residents in the PVM, while proteins of the host cell are mostly excluded. Some PVM proteins are shared between gametocytes and the asexual blood stages, such as subunits of the translocon\(^4\), but there are also important differences. One example is the different members of the ETRAMP protein family\(^3\). In *P. falciparum* the ETRAMP family protein *Pfs16* is specifically present in the gametocyte PVM\(^3\)\(^6\) and gene disruption studies suggest that it is important for successful development of gametocytes\(^7\).

Egress has been studied by reverse-genetic approaches, and several proteins are confirmed to have roles in egress. Most studies have been performed in *P. berghei*. For instance, MDV/PEG3\(^8\) and PbGEST\(^9\) function in the rupture of the PVM in both genders, and are localized to specialized secretory organelles called osmiophilic

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\(^1\)Institute of Molecular Biology and Biotechnology, Foundation for Research and Technology - Hellas, Heraklion, Greece. \(^2\)Department of Biology, University of Crete, Heraklion, Greece. \(^3\)Bio21 Molecular Science and Biotechnology Institute, Electron Microscopy Unit, University of Melbourne, Melbourne, Victoria, Australia. \(^4\)Department of Biochemistry and Molecular Biology, University of Melbourne, Melbourne, Victoria, Australia. \(^5\)Université de Montpellier, CNRS UMR 5048, INSERM UMR, 1054, Montpellier, France. \(^6\)Université de Montpellier, CNRS UMR 5235, Montpellier Cedex, France. \(^7\)School of BioSciences, University of Melbourne, Melbourne, Victoria, Australia.

Correspondence and requests for materials should be addressed to I.S.-K. (email: inga@imbb.forth.gr)
bodies. Mutant male gametocytes lacking the stage-specific actin isoform, actin 2, are also unable to rupture the PVM, but egress of the female gametocytes is not affected in this mutant\(^1\). In these three mutants both surrounding membranes remain intact. Another mutant lacking PPLP2, one of the five perforin-like proteins, in \textit{P. berghei}, also has a male-specific phenotype. However, in this disruption mutant of \textit{pplp2} the PVM ruptures normally while the RBCM remains intact\(^1\). In many mutant cells lacking PPLP2 the cytoplasm of the red blood cell is also visible in EM pictures (unpublished\(^1\)). A female-specific protein called G377 also has a minor role in egress\(^1\). Inhibitor studies suggest that proteases have important roles during egress, but no protease has so far been identified with a role in this process\(^1\). Recently a proteomics approach identified a trap-like protein, called MTRAP, residing in osmiophilic bodies. In its absence were trapped in the host cell\(^1\). Another study confirmed this role of MTRAP in both \textit{P. berghei} and \textit{P. falciparum} gametogenesis and also showed that in disruption mutants neither the PVM nor the RBCM were ruptured. Furthermore, the protein was important for egress of both genders\(^4\)

Gametocytegenesis is initiated when merozoites already committed to sexual development through the activity of the transcription factor ApiAP2-G, invade RBCs\(^1,2\). The early stages of the developing gametocytes are molecularly and morphologically similar to the asexual stages; they develop within the PVM, digest hemoglobin and modify the host red blood cell, but in later stages gametocytes follow a distinct developmental program\(^7\). After uptake of the mature gametocytes in a blood meal by the mosquito, gametogenesis is activated by the decrease in temperature and by xanthurenic acid (XA)\(^18\). XA triggers activation of cGMP dependent protein kinase (PKG)\(^29\) which mediates an increase in intracellular Ca\(^{++}\) levels\(^20\) and hydrolysis of PIP\(_2\) and the production of IP\(_3\),\(^21\) thus initiating egress of both genders. Effector proteins important for membrane rupture reside in osmiophilic bodies and their release is dependent upon intracellular Ca\(^{++}\) levels\(^1\). The male gametocyte also undergoes three mitotic divisions, and assembly and activation of eight axonemes leads to the formation of eight motile flagellar gametes from each gametocyte. The flagella of gametocytes are believed to be required for fertilization without any re-organization of the cell. Gametogenesis is completed 10–15 min after uptake of the blood by a mosquito.

In the asexual blood cycle invasive merozoites are released after initial rupture of the PVM and then subsequent opening of the RBCM\(^22\). Merozoite egress is also dependent on PKG activity\(^22,23\), elevated Ca\(^{++}\) levels\(^24–26\) and PIP\(_2\) hydrolysis\(^20\). Parasite-derived proteases have important roles\(^27\) and host factors are necessary for disrupting the host cytoskeleton, a crucial step for RBCM rupture\(^24,28,29\). Detailed morphological studies by live imaging of \textit{P. falciparum} merozoite egress revealed that a single stabilized pore in the RBCM is followed by outward curling and buckling of the membrane\(^30\). A similar outward curling has also been detected in egressing \textit{P. falciparum} gametocytes\(^31\). An ultrastructural study revealed that egress is preceded by the loss of mechanical integrity of the RBC cytoskeleton, and at the same time the PVM ruptures into multilamellar vesicles. Next the RBCM becomes permeable followed by the explosive egress of the merozoites\(^22\). In the study presented here we use the term vesicles of the ruptured membranes, in line with that publication\(^22\).

Here, we investigated \textit{P. berghei} gametocyte egress combining high speed video observations with immunofluorescence analysis using antibodies against the RBCM and PVM. Although each event differed, four main phases were identified: swelling; PVM rupture and vesiculation; a single pore opening in the RBCM; and, finally, rupture and vesiculation of the RBCM. The sequence of phases was similar between the two genders.

**Results**

High speed video recordings of gametocyte egress reveal rupture and vesiculation of the two surrounding membranes. We used a high-speed video camera to follow egress in real time. Blood samples from infected mice were diluted in medium, transferred to microscope slides and videos were recorded, beginning a few minutes after activation. Due to the short interval between activation and the first signs of egress the beginning was not always recorded. 56 events, of which 34 were females and 22 males, were captured. Four representative movies of each gender are presented (Snapshots Figs 1 and 2 and Supplementary Information Video 1–8). Egress does not follow a stereotyped pattern, but despite this, the series of events have common features. The first discernible feature is swelling of the erythrocyte (Figs 1b,m and 2k) with the parasite asymmetrically positioned in the host cell. Then the PVM ruptures and forms vesicles internal to the erythrocyte membrane (Figs 1f,i,j,n and 2a,b,e,h,i). The vesicles are freely moving within the erythrocyte, suggesting that the RBC cytoplasm is largely digested or has leaked out of the cell. Next, the erythrocyte membrane opens at a single pore and in the females the internal vesicles are discharged (Fig. 1c,o) or the gamete is released with the vesicles positioned on the opposite side of the pore (Fig. 1g,k). In the males the vesicles are accompanied by the exit of a single motile male flagellum (Fig. 2c,i), but in two cases we detected vesicle extrusion slightly earlier than activation of the flagella (Fig. 2f,m, Supplementary Information Video S6, S8). The pore is then widened with the expulsion of the gamete followed by destabilization of the pore's rim, buckling of the RBCM and its shredding into small vesicles. The vesicles remain in the vicinity of the male residual gametocyte or female gamete for several minutes (Figs 1d,h,l,p and 2d,g,j,n).

We recorded the approximate timing of the different phases. In total we were able to time at least some of them for 10 females and 9 males. In only two events did we detect the beginning of the swelling and the PVM in these cases ruptured 66 sec and 375 sec after swelling. The interval between PVM rupture and pore opening varies from a few seconds to five minutes, whilst the RBCM vesiculation in eight cases happens within seconds but in two cases ~2 minutes after the opening (Supplementary Information Fig. S1a). Thus the intervals between the phases are highly variable, and this is the case for both genders.

To quantify the swelling that we observed we measured the area of the RBC and gametocyte in Giemsa stained smears of non-activated and activated gametocytes (Supplementary Information Fig. S1b). The ratio of the two values was plotted, and the difference comparing non-activated and activated cells was significant. The results are consistent with the live imaging although we cannot exclude that weakening of the host cell membrane after activation may make the cells more sensitive to the smearing on the slide, thus overestimating swelling.
Live immunolabeling confirms single pore opening in the erythrocyte membrane and vesiculation of the membrane. We used the TER-119 monoclonal antibody recognizing a glycophorin-A associated protein on the surface of the red blood cell to label the erythrocyte membrane. Blood samples were diluted in medium containing the antibody, and the cells were imaged in an epifluorescence microscope with an attached CCD camera. We used the \textit{P. berghei} strain 820 in which female gametocytes express red fluorescent protein (RFP), while the males express a weak green fluorescent protein (GFP) not detectable in these experiments. The flagellar male gametes move rapidly, precluding good quality live imaging. Therefore, we focused on female cells. Female gametocytes were observed that were enclosed in the RBCM being adjacent to the parasite (Fig. 3a). In other cells swelling of the host cell was seen as the RBCM was now separate from the parasite (Fig. 3b). The actual opening of the pore was not captured, possibly due to the very brief time before the wider opening of the pore. However, we did detect a partially opened membrane (Fig. 3c). Typically, the final opening of the membrane is accompanied by vesiculation of the membrane (Fig. 3d, upper panel). However, we also detected a gamete having almost exited the

Figure 1. Snapshots from four movies of female gamete egress. Representative snapshots chosen to illustrate the stages of egress. Upper broken lines represent time in sec between each frame of the movies in real time. The time of each frame from the start of the movies is indicated above the snapshots (min:sec). For technical reasons this does not correspond with the time of the start of the experiment (activation). (a–d) Female 1. (a) An infected RBC in the initial phase of swelling. The parasite occupies most of the volume of the RBC. (b) The RBCM is more extended. (c) The PVM vesicles are extruded through a single pore. (d) Egress is complete. Vesiculated PVM and RBCM are localized in proximity to the gamete. (e–h) Female 2. (e,f) The PVM is already vesiculated inside the RBCM. (g) The RBCM is extended and the gamete slides out of the host membrane. (h) Egress is complete with vesicles next to the gamete. (i–l) Female 3. (i) The RBCM is largely extended. (j) PVM is ruptured and vesiculated. (k) The gamete slides out of the extended RBCM. (l) Egress is complete, the RBCM is vesiculated. (m–p) Female 4. (m) RBC is swollen. (n) PVM vesicles are seen inside the RBCM towards the viewer. (o) Expulsion of PVM vesicles through a single pore. (p) Egress is complete. Black arrows indicate the area of interest in each snapshot, and white double arrows direction of gamete expulsion in (g) and (k). Scale bar, 5 μm. See also Supplementary Information Videos S1–S4.
intact membrane (Fig. 3d, lower panel, compare to Fig. 1g,k and Supplementary Information Video S2,S3). We did not unequivocally recognize the characteristic curling of the membrane, previously described for egress of P. falciparum merozoites30 and gametocyte31, although in Fig. 3c and d there is a slight thickening of the membrane at the opening. However, the observed buckling of the RBC membrane following female gamete expulsion is indicative of curling. This is most clearly visible in Supplementary Information Video S2.

After completion of gametogenesis we detected labeled membrane vesicles in the proximity of the female gamete (Fig. 3e) and the residual gametocyte of exflagellating cells despite the vigorous beating of the flagellar gametes (Fig. 3f). These observations confirm that the RBCM is vesiculated after egress and that the vesicles remain close to the cells in both genders.

**Outside-out vesiculation of the red cell membrane.** We next performed immunofluorescence assays using the TER-119 antibody to label specifically the outer leaflet of the RBCM. Strain 820 gametocytes were fixed...
Figure 3. Live imaging reveals RBCM rupture. Samples were imaged without prior fixation. (a–e) Female cells expressing RFP (red) and labeled with TER-119 (green). (a) A female gametocyte still enclosed in the intact RBCM. The membrane is adjacent to the parasites. (b) Swelling of the RBCM membrane is evident. Insets show the adjacent (a) and displaced RBCM (b) compared to the parasite. (c) RBCM opened at a single point. Arrow points to a slight thickening of the membrane at the opening. (d) Two cells in the process of egress. The arrows point to a slight thickening of the membrane at the opening. Top panel: The RBCM is opening and simultaneously forming vesicles (asterisks, out of focus), the most common RBCM rupture detected. Lower panel: The RBCM opens as an intact membrane. Compare Fig. 1, female 2 and 3 and Supplementary Information Videos S2, S3. (e) Rupture of RBCM completed, and RBCM vesicles remain next to the gamete (white arrow). White arrowheads in (c–e) point to opened or vesiculated RBCM. (f) Egressed male which is exflagellating. RBCM vesicles remain in proximity to the residual gametocyte and two flagellar gametes are visible (BF image, white arrows). The cells are motile and thus slightly differently located between panels. Scale bar, 5 µm.
in formaldehyde without permeabilization at different time points after activation. We were not able to visualize the membrane just after opening of the pore, which is not unexpected given that our live imaging shows this to be a very rapid event. However, one example is shown where the membrane has opened but not yet vesiculated (Fig. 4a). We readily detected cells associated with free TER-119 labeled vesicles (Fig. 4b,c), confirming the live imaging results and indicating that the vesicles are of the outside-out type, that is, the outer leaflet of the membrane remains on the outside of the vesicles. However, we cannot exclude that inside-out vesicles are also formed but we were unable to test this possibility due to the lack of a marker for the inner leaflet of the RBCM.

**PVM visualized by SEP1 immunolabeling.** We used an antibody directed against the SEP1 protein, a member of the ETRAMP (early transcribed membrane proteins) protein family, previously shown to be located in the PVM, to visualize directly the PVM32. The immunolabelling of SEP1 revealed the closed PVM in non-activated samples (Fig. 5a) and SEP1 labeled vesicles inside intact RBCM in activated gametocytes (Fig. 5b). We also captured labeled vesicles in the process of exiting the RBCM (Fig. 5c,e). This confirms the results from the live imaging that PVM vesicles are extruded through a single pore opening of the RBCM. In addition a thickening of the RBCM around the pore was seen suggesting curling of the membrane. In another cell the PVM was seen close to the vesiculated RBCM (Fig. 5d).

To visualize the gametocyte during egress we used PSOP12 as a marker; this protein has been shown to be located on the gametocyte surface33. Gametocytes of a strain expressing PbSOP12 fused to GFP33 were labeled with the SEP1 and TER-119 antibodies as above. In non-activated cells the signals of the three proteins were largely overlapping (Fig. 6a). In a cell where swelling had taken place SEP1 and TER-119 were detected adjacent to each other, while the SOP12 labeled the gametocyte inside the extended PVM (Fig. 6b). However, in exflagellating gametocytes we failed to detect PSOP12 circumferential to the gametocyte/gamete. Instead staining was seen in the vicinity of the cell suggesting that the protein is secreted after activation and trapped between the ruptured PVM and RBC membranes (Fig. 6c). Hence it was not possible to use this marker for labeling the egressing gamete.

![Figure 4.](image-url) Outside-out vesiculation revealed with TER-119 label. (a–c) Females (expressing RFP, red) were fixed at different time points after activation and the RBCM labeled with TER-119 (green). (a) 7.5 min after activation the RBCM has opened at a single site. (b) Three females having completed gametogenesis 9 min after activation. Labeled outside-out vesicles remain in proximity to the gametes. (c) Completed gametogenesis at 11.5 min. Labeled vesicles remain close to the gamete. An intact RBC is indicated with an asterisk. White arrows point to opened or vesiculated RBCM. Scale bar, 5 μm.
The PVM is vesicularized in parasites lacking Perforin-like protein 2. We reported previously that in mutant male gametocytes lacking *P. berghei* perforin-like protein 2 (PPLP2) the PVM ruptured during activation of gametogenesis while the RBCM remained intact. However, in that study we were not able to image the PVM. To confirm the earlier study and verify that the rupture of PVM is independent of that of RBCM we imaged activated

**Figure 5.** Labeling with SEP1 antibodies reveal PVM vesicles. Cells were labeled with the SEP1 antibody (red) and TER-119 (green) highlighting the PVM and RBCM, respectively. DNA is stained with Hoechst 33342 (blue). (a–d) Are projections of sections from confocal imaging. (a) Non-activated cell. (b) An activated cell. PVM vesicles have formed inside the intact RBCM. (c) The RBCM is opened at a single pore and SEP1 labeled vesicles are in the process of exiting. A montage of the same cell is shown in (e). (d) A cell where egress is completed. Vesicles of RBCM and PVM are intermingled. (e) Montage of single sections of the cell in (c). Scale bar, 5µm.
pplp2(−) mutant gametocytes. As expected, PVM vesicles were formed inside the RBC membrane (Fig. 7a). In these experiments equina toxin was used to permeabilize the RBC for efficient antibody labeling and this led in certain cases to collapse of the RBCM and release of PVM vesicles (Fig. 7b). In this case we did not detect the typical RBCM vesicles. We also detected free membrane whorls and spirals (Fig. 8c); it is not however possible to determine whether they are derived from the PVM or the RBCM. A scanning EM

Electron microscopy analysis of egressing parasites. Our light microscopy analysis was supported by electron microscopy of activated gametocytes. An example of a gametocyte still enclosed in the intact RBCM but with no visible PVM is presented (Fig. 8a); three membrane vesicles, one of which is flattened, are seen in the space outside the parasite. We interpret this as vesicles derived from the PVM consistent with the live imaging experiments and the SEP1 localization. In other cells the RBC cytoplasm was no longer visible, showing the gametocytes surrounded by the RBCM (Fig. 8b). We also detected free membrane whorls and spirals (Fig. 8c);
(SEM) analysis of samples from a time point when exflagellation took place revealed ruptured membranes (Fig. 8d,e), possibly derived from the RBCM. In one case (Fig. 8d) the membrane has a rough surface which can be interpreted to be the inner leaflet of the RBCM. Another membrane collapsed over a cell has a smoother surface suggesting the outer leaflet in this case facing outwards (Fig. 8e). This membrane is visible as one sheet further strengthening the view that the membrane is opened at a single pore which is then extended before vesiculation takes place.

Discussion
Malaria parasites use unique processes to escape from the red blood cell. In this study we applied complementary imaging methods for a detailed analysis of P. berghei gametocyte egress from the host cell. The process does not follow a stereotyped pattern as each event that we observed was different and the timing of events also varies. However, egress can be divided into discreet steps which are similar in both genders. Based on our results we suggest a model for gametocyte egress (Fig. 9). The first sign of egress is extension of the membranes and swelling of the erythrocyte. Then the PVM ruptures and vesiculates. This is followed by the opening of a single stabilized pore in the RBCM. The next step had two distinct morphologies in females. In some cases vesicles of the PVM are extruded through the pore followed by vesiculation of the red blood cell membrane. In other cases, the membrane remained in one piece while the gamete gently slid out. After the gamete exit was complete the extrusion of the vesicles was accompanied by a single flagellum, although in two movies a vesicle was extruded before the flagellum (Fig. 2f,m, Supplementary Information Videos S6,S8). Again, vesiculation of the red blood cell membrane happened rapidly. Interestingly, in one unusual
also showed that although motile flagella were formed they remained trapped within the RBCM. Gamete egress normally although motile flagella are not formed. Our results from a mutant lacking PPLP2 also showed that although motile flagella were formed they remained trapped within the RBCM. A recent study34 has been suggested to be the result of water and ion re-distribution between the two compartments36. In schizont rupture the swelling takes place several minutes before egress35. In our analysis swelling had usually already taken place when recording started. However, in a few events we could time the interval to PVM vesiculation which lasted between approximately 1–6 minutes. This is roughly similar to what has been reported for PVM rupture in P. falciparum gametocyte egress37. The coincidence of this time interval in schizont and gametocyte egress may point to common processes necessary for the continuation of membrane rupture. The swelling of the cell is the first stage of egress. This has also been observed during merozoite egress30,35,36, and similar to that we found the PV compartment was extended while the RBC cytoplasm shrunk (Fig. 6). This has been suggested to be the result of water and ion re-distribution between the two compartments36. In schizont rupture the swelling takes place several minutes before egress35. In our analysis swelling had usually already taken place when recording started. However, in a few events we could time the interval to PVM vesiculation which lasted between approximately 1–6 minutes. This is roughly similar to what has been reported for PVM rupture in P. falciparum gametocyte egress37. The coincidence of this time interval in schizont and gametocyte egress may point to common processes necessary for the continuation of membrane rupture.

The next step, the rupture and consequent vesiculation of the PVM, was detected in all of the events, suggesting that this is a required step in egress. In schizont rupture the opening of a single stabilized pore in the RBCM, and the outward curling of the membrane results in expulsion of the non-motile merozoites thus assisting in their dispersal35. Gamete egress is different; male gametes are highly motile due to the beating of the flagellum and after collision with the non-motile female gametes, fusion of the gametes takes place. Therefore, there is no need for assisted dispersal of these stages. In all cases observed here a single pore was opened in the RBCM, but the subsequent events had two distinct outcomes. On most occasions the opening of the membrane was rapid, and vesiculation occurred quickly following gamete release (Supplementary Information Videos S1,S4-S8). In two cases female gametes gently slid out of the RBCM, which for a few seconds remained intact, buckled and then vesiculated (Supplementary Information Videos S2.S3). Although buckling strongly suggests curling, we were not able to clearly detect the characteristic curling of the pore’s rim, which has been described during merozoite and gametocyte egress of P. falciparum30,31, despite many attempts of visualization of the RBCM using the TER-119 antibody. In some images there is a thickening of the membrane at the pore opening (Figs 3b,c and 5c) but we hesitate to unequivocally state that this is outward curling. The lack of detection of unequivocal curling, however, might be due to the fact that the RBCM destabilizes into vesicles faster than it curls.

After the membrane opening, the RBCM is vesiculated. The time interval between pore opening and RBCM vesiculation varies between a few seconds and several minutes (Fig. S1a). In all cases membrane vesicles were formed. To determine if these vesicles were of the outside-out type, that is the outer leaflet of the RBCM remains outside the vesicle, we used the TER-119 antibody to label the vesicles on fixed but not permeabilized samples at a time point when egress was complete. We detected many labeled vesicles, suggesting that the outer RBCM leaflet is facing outwards (Fig. 4). This observation is not contradictory to curling as the destabilized membrane can re-seal in an outside-out fashion. The lack of a marker for the inner leaflet prevented us from identifying whether any vesicles of inside-out type occur.

In the movies we especially noticed the immediate vesiculation of the RBCM after curling and buckling. In P. falciparum merozoite egress vesiculation of the RBCM takes place later30. Possibly this difference can be explained by differences in the organization of infected cells and/or the molecular events taking place. The stabilization of the pore and the vesiculation is governed by the state of the cortical cytoskeleton, which is altered in the parasitized red blood cell30. RBCM ruptured at one point with the characteristic curling at the pore opening of a P. falciparum gametocyte has been described31. However, the RBCM seems to remain intact in one piece attached to the emerging gametocyte2,21,40, which is different to the vesiculation that we detected here. Less is known about PVM rupture although three studies on P. falciparum have revealed similar vesicles to those described here.
Immunofluorescence assays of fixed infected blood samples. The TER-119 antibody was used to label the outer leaflet of the RBCM. Gametocytes were activated and samples were fixed in 4% formaldehyde at different time points after the activation. The samples were not permeabilized. All steps were carried out at room temperature. The parasites were added to poly-L-lysine coated cover slips placed in 24-well plates, centrifuged at 500 x g for 15 min, and the fixative removed, followed by washing twice with 1 × PBS. The TER-119 antibody was added diluted 1:200 in PBS with 2% normal goat serum for 1 h incubation. Samples are washed twice with 1 × PBS before mounting in Vectashield (vector laboratories). Image analysis was performed as above.

For immunolabeling of WT and PbSOP12-EGFP parasites with anti-SEP1 antibody non-activated and activated for 8 min samples were permeabilized with 0.1% saponin for 30 sec; at later time points no permeabilization was carried out. Fixation was as described above. Hoechst 33342 was used to stain DNA. For PbSOP12-EGFP parasites anti-GFP antibody was also added.

The ppp2(−) mutants were immunolabeled with anti- SEP1 as above. Permeabilization was done using equinatoxin.

Electron microscopy. Samples were fixed in 2.5% glutaraldehyde and processed using the ROTO method and observed on a FEI Tecnai F30 at 300 kV (Advanced Microscopy Facility, Bio21 Institute, the university of...
Melbourne, Australia). Micrographs were acquired with a Gatan Ultrascan 1000 2 × 2k. Alternatively the samples were visualized by block face imaging in a Zeiss Sigma equipped with a Gatan 3View system (Cenier for Microscopy and Microanalysis, University of Queensland, Australia) with a 50 nm section thickness.

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**Acknowledgements**

We thank Simon Conway for essential help with the scanning EM and Theodossia Bitzou for artwork. We are grateful to Robyn Webb and Richard Webb, of the Australian Microscopy & Microanalysis Research Facility at the Center for Microscopy and Microanalysis at the University of Queensland, for access to the 3View. Support was provided by a grant from the Program Bilateral Research and Technology Cooperation Greece-France, The General Secretariat for Research and Technology, Ministry of Education, Greece (grant number 2013SE01380004). The post-doctoral research of MA was supported by a fellowship from IKY (State Scholarship Foundation) funded by the action “Support of Post-Doctoral Researchers” from the Operational Program “Development of Human Resources, Education and Life-Long Learning” with priority areas 6, 8, 9 and co-funded by the European Social Fund – ECB and Greek public funds. ISK acknowledges the receipt of an OzMalNet fellowship from the EviMalar network of excellence. GIM is supported by a Discovery Project and Laureate Fellowship from the ARC and a Program Grant from the NHMRC. This work was performed in the framework of the BIOSYS research project, Action KRIPIIS, project No MIS-448301 (2013SE01380036) funded by the General Secretariat for Research and Technology, Ministry of Education, Greece and the European Regional Development Fund (Sectoral Operational Programme: Competitiveness and Entrepreneurship, NSRF 2007–2013)/ European Commission.

**Author Contributions**

I.S.K., M.Abk., C.B.B. conceived the study, M.A., E.H., E.D., C.C., K.W., V.M. performed the experiments, I.S.K. and M.A. analyzed the data, I.S.K., M.A., E.D., C.B.B. and G.I.M. wrote the paper. All authors reviewed the manuscript.

**Additional Information**

**Supplementary information** accompanies this paper at https://doi.org/10.1038/s41598-018-21801-3.

**Competing Interests:** The authors declare no competing interests.

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Author/s:
Andreadaki, M; Hanssen, E; Deligianni, E; Claudet, C; Wengelnik, K; Mollard, V; McFadden, GI; Abkarian, M; Braun-Breton, C; Siden-Kiamos, I

Title:
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Date:
2018-02-23

Citation:
Andreadaki, M., Hanssen, E., Deligianni, E., Claudet, C., Wengelnik, K., Mollard, V., McFadden, G. I., Abkarian, M., Braun-Breton, C. & Siden-Kiamos, I. (2018). Sequential Membrane Rupture and Vesiculation during Plasmodium berghei Gametocyte Egress from the Red Blood Cell. SCIENTIFIC REPORTS, 8 (1), https://doi.org/10.1038/s41598-018-21801-3.

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