GDV1 C-terminal truncation of 39 amino acids disrupts sexual commitment in *Plasmodium falciparum*.

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Abstract

Malaria is a mosquito-borne disease caused by apicomplexan parasites of the genus *Plasmodium*. Completion of the parasite's life cycle depends on the transmission of sexual stages, the gametocytes, from an infected human host to the mosquito vector. Sexual commitment occurs in only a small fraction of asexual blood stage parasites and is initiated by external cues. The gametocyte development protein 1 (GDV1) has been described as a key facilitator to trigger sexual commitment. GDV1 interacts with the silencing factor heterochromatin protein 1 (HP1), leading to its dissociation from heterochromatic DNA at the genomic locus encoding AP2-G, the master transcription factor of gametocytogenesis. How this process is regulated is not known. In this study we have addressed the role of protein kinases implicated in gametocyte development. From a pool of available protein kinase KO lines, we identified two kinase knockout lines which fail to produce gametocytes. However, independent genetic verification revealed that both kinases are not required for gametocytogenesis but both lines harbour the same mutation that leads to a truncation in the extreme C-terminus of GDV1. Introduction of the identified nonsense mutation into the genome of wild type parasite lines replicates the observed phenotype. Using a GDV1 overexpression line we show that the truncation in the GDV1 C-terminus does neither interfere with the nuclear import of GDV1 nor its interaction with HP1 *in vitro*, but appears important to sustain GDV1 protein levels and thereby sexual commitment.

Importance

Transmission of malaria causing *Plasmodium* species by mosquitoes requires the parasite to change from a continuously growing asexual parasite form growing in the blood, to a sexually differentiated form, the gametocyte. Only a small subset of asexual parasites differentiates into gametocytes that are taken up by the mosquito. Transmission represents a bottleneck in the lifecycle of the parasite, so a molecular

understanding of the events that lead to stage conversion may identify novel intervention points. Here we screened a subset of kinases we hypothesized to play a role in this process. While we did not identify kinases required for sexual conversion, we identified a mutation in the C-terminus of the Gametocyte Development 1 protein (GDV1), which abrogates sexual development. The mutation destabilises the protein but not its interaction with its cognate binding partner HP1. This suggest an important role for the GDV1 C-terminus beyond trafficking and protein stability.

Introduction

Malaria is a devastating disease caused by parasites of the genus *Plasmodium*, leading to ~ 405,000 deaths per year ¹. *Plasmodium falciparum* causes the most severe and life-threatening form of human malaria. The complex life cycle involves interactions with multiple tissues in two different organisms, the human host and the mosquito vector. Inside the human host *P. falciparum* predominantly infects red blood cells (RBC) where it asexually replicates or, a small fraction (0-20%), commits to sexual development (gametocytogenesis) ². Gametocytogenesis occurs preferentially in the extravascular compartment in the bone marrow and spleen ³⁻⁹. After 10-12 days mature stage V gametocytes are released into the peripheral circulation to allow transmission to mosquitoes.

Sexual commitment can be initiated by metabolic cues in the human host. Specifically, it has been described that depletion of lysophosphatidylcholine (LysoPC), a common component of human serum, leads to increased rates of gametocyte production and therefore represents the first molecularly defined factor known to inhibit or trigger sexual conversion ¹⁰. Sexual commitment depends on upregulation of the *ap2-g* gene ^{2,11}, which requires removal of heterochromatin protein 1 (HP1) from chromatin. HP1 interacts directly with the gametocyte development 1 protein (GDV1), which causes HP1 to dissociate ¹². HP1 is responsible for repression of a range of genes ¹³, while GDV1 specifically acts on the *ap2-g* locus. How this

specificity is achieved is not known. Furthermore, how a drop in LysoPC levels is sensed and transduced into GDV1-mediated HP1 removal is not understood. Kinases are key transducers of signals in cellular processes in various stages of the *Plasmodium* life cycle^{14,15} and are likely candidates to play important roles in gametocyte commitment and development. A study by Solyakov et al¹⁴ has identified a panel of likely and confirmed non-essential protein kinases, some of which are transcribed during sexual development (PlasmoDB) or in gametocytes¹⁶⁻¹⁹. Aiming to identify protein kinases involved in sexual development we screened eight KO lines for phenotypes in gametocyte induction and/or maturation. Two lines made no gametocytes, but subsequent validation showed that their gametocytogenesis defect was not due to the absence of these kinases. Instead, we found that both lines shared the same truncation in the C-terminal end of GDV1, which caused the loss of gametocyte development. Here, we address the importance and role of the GDV1 C-terminal for sexual commitment and interaction with HP1. We show that the loss of the C-terminal 39 amino acids of GDV1 does not interfere with nuclear import and interaction with HP1 in vitro, but prevents GDV1 from triggering efficient sexual commitment.

Results

Identification and characterization of *Plasmodium falciparum* kinase KO lines with a gametocytogenesis phenotype

It has been shown previously in *P. berghei* that protein kinases non-essential during the asexual blood stages are essential in other lifecycle stages, for example during parasite transmission in the mosquito¹⁵. To identify kinases important for gametocytogenesis we investigated the role of a group of likely non-essential kinases¹⁴ during asexual blood stages development. Using the lines described by Solyakov et al¹⁴, which have been generated by single crossover gene disruption, we induced sexual development using conditioned medium²⁰ and followed progression through

the stages I to V of gametocytogenesis (Figure 1a). Six of the eight KO lines displayed normal gametocyte development, while two, TKL2 (PF3D7_1121300) and eIK2 (PF3D7_0107600) kinase KO lines, produced very few ($\leq 0.1\%$) gametocytes (Figure 1b). Of these, one has a disrupted tyrosine kinase like 2 (*tkl2*) locus, which has been characterized as a protein kinase secreted outside of the red blood cell¹⁷. Gene loss and accumulation of mutations is frequently observed in parasite lines kept in continuous *in vitro* culture over time and the loss of the ability to form gametocytes is not uncommon²¹. To exclude mutations in the *ap2-g* gene, which was identified through a loss of function mutant previously², we sequenced the *ap2-g* locus in the 3D7/TKL2 KO parasite line. The sequencing results confirmed that the phenotype observed was not associated with mutations in *ap2-g*, leading us to conclude that the deletion of TKL2 was possibly the cause for the observed phenotype. In order to verify the role of TKL2 in gametocyte induction, we generated a DiCre-mediated TKL2 conditional KO line in NF54 parasites (NF54/TKL2:loxPint). We used CRISPR/Cas9 to simultaneously introduce a DiCre cassette into the *pfs47* locus, as previously described^{22,23}, and to flank the kinases domain of *tkl2* with two loxPints (Figure 1c, Supplementary Figure 1a and b). To address the role of TKL2 in gametocyte development we treated the NF54/TKL2:loxPint line with DMSO (control) or rapamycin (KO) (Supplementary Figure 1b). We then induced sexual commitment using conditioned medium²⁰, and monitored gametocyte development. No difference in commitment or development between the control and the rapamycin-induced NF54/TKL2:loxPint parasites (Figure 1d) were observed. These results show that TKL2 is not involved in sexual commitment or gametocyte development/maturation and another mutation is likely the cause for the observed phenotype.

A common GDV1 truncation is found in both kinase KO lines deficient in gametocyte formation.

The second kinase KO where a gametocytogenesis defect was identified was the eukaryotic initiation factor serine/threonine kinase 2 (eIK2) KO line (3D7/eIK2 KO)

(Supplementary Figure 2a). eIK2 has previously been characterized as non-essential during sexual development in *P. falciparum* and *P. berghei* and eIK2 KO lines appeared to undergo normal gametocyte development in rodent *Plasmodium* species²⁴. This indicated that, as 3D7/TKL2 KO, the 3D7/eIK2 KO line also harbours a mutation preventing efficient gametocyte development. Sequencing of the *ap2-g* locus in this parasite line as described above showed no mutations in the coding region of *ap2-g*. Therefore, a potentially unknown mutation underlies the loss of gametocytes in these parasite lines.

To understand the nature of the block in sexual development we analysed the transcriptome of induced wildtype 3D7 parasites and in two eIK2 KO clones using RNAseq. Samples were collected for RNA extraction between 28-32 hours post-invasion (hpi) after induction with conditioned medium (Figure 2a). The RNAseq analysis revealed a significant downregulation in 3D7/eIK2 KO parasites of genes known to be upregulated during gametocytogenesis, including genes that have been shown to be AP2-G-dependent^{2,10,12,25-27} (Figure 2b and Supplementary Table 1). We found *ap2-g* itself to be downregulated in 3D7/eIK2 KO parasites, but this reached significance only in one of the clones. Together with the lack of mutations in *ap2-g* itself, these results suggested that the block in gametocytogenesis was upstream of AP2-G function during sexual commitment. At that time, GDV1 was shown to be an upstream activator of AP2-G expression¹² so we sequenced the *gdv1* locus in the eIK KO clones and identified a nonsense mutation in *gdv1* that results in a premature stop codon leading to a C-terminal truncation of 39 amino acids (GDV1Δ39) (Figure 2c and Supplementary Figure 2b). Sequencing of the 3D7/TKL2 KO parasite clones showed the same mutation (Supplementary Figure 2b), suggesting that the deletion of the last 39 amino acids of GDV1 in both mutant lines is responsible for the gametocytogenesis phenotype observed in both kinase KO lines.

The carboxy-terminal 39 amino acids of GDV1 are important for its function

170 To verify genetically the identified mutation in *gdv1* we generated an 3xHA-tagged
 171 version of GDV1Δ39 and introduced it in the endogenous *gdv1* locus in the NF54
 172 parasite line (NF54/GDV1Δ39:HA) (Figure 3a and Supplementary Figure 3a and b).
 173 GDV1Δ39:HA parasites lost the ability to form gametocytes (Figure 3b), suggesting
 174 the GDV1 C-terminus plays an essential role during sexual commitment or
 175 development. Determination of the localisation or expression levels of GDV1Δ39:HA
 176 was not possible, as we could not confidently distinguish true signal from background
 177 fluorescence. We repeatedly failed to obtain parasites expressing 3xHAtagged full
 178 length GDV1 from the endogenous locus to compare expression levels and the
 179 localisation of the truncated GDV1 version. Notably, direct C-terminal tagging of
 180 GDV1 at the endogenous locus was also not successful in other studies, unless when
 181 in combination with a destabilisation domain ^{12,28}.
 182 Therefore, we resorted to a system that allows robust testing of GDV1-dependent
 183 gametocyte induction. We introduced an ectopic *gdv1-gfp* fusion gene under control
 184 of the calmodulin promoter and a *glmS* ribozyme in the 3' untranslated region has
 185 been introduced into the *cg6* (*glp3*, PF3D7_0709200) locus, allowing conditional
 186 overexpression of GDV1-GFP to trigger high gametocyte conversion rates
 187 (NF54/iGP2 line) ²⁹. In the presence of glucosamine, the *glmS* ribozyme destabilizes
 188 the mRNA preventing GDV1:GFP expression, while in the absence of glucosamine
 189 GDV1:GFP is overexpressed, leading to gametocyte induction ³⁰. For simplicity, the
 190 NF54/iGP2 line described by Boltryk and colleagues ²⁹ has been renamed
 191 NF54/GDV1:GFP_cOE in this study (cOE stands for conditional over expression). To
 192 test GDV1Δ39 function in this assay, we introduced a *gdv1Δ39-gfp-glmS* cassette
 193 into the *cg6* locus, generating a conditional GDV1Δ39:GFP overexpression parasite
 194 line (NF54/GDV1Δ39:GFP_cOE) (Figure 3c and Supplementary Figure 3c and d). We
 195 then compared the sexual conversion rates in the NF54/GDV1:GFP_cOE and
 196 NF54/GDV1Δ39:GFP_cOE parasite lines in the presence and absence of

glucosamine. Unlike the NF54/GDV1:GFP_cOE control line, the induced overexpression of GDV1 Δ 39:GFP in the NF54/GDV1 Δ 39:GFP_cOE parasite line failed to trigger a significant increase in sexual commitment (Figure 3d). These results suggest that the full integrity of the GDV1 C-terminus is important for sexual commitment.

GDV1 Δ 39 is imported into the nucleus and retains the ability to interact with HP1

GDV1 is a nuclear protein and we hypothesized that the deletion of a predicted C-terminal nuclear bipartite localisation sequence (cNLS mapper, http://nls-mapper.iab.keio.ac.jp/cgi-bin/NLS_Mapper_form.cgi) may interfere with GDV1 nuclear localisation and hence its ability to interact with HP1 at heterchromatic loci. Therefore we localised GDV1 Δ 39:GFP in the NF54/GDV1 Δ 39:GFP_cOE parasites by immunofluorescence at 28-32 hpi (Figure 4a and Supplementary Figure 3e). The results show a clear punctate and nuclear signal in the induced NF54/GDV1:GFP_cOE parasite line, as previously reported (Supplementary Figure 3e)¹². NF54/GDV1 Δ 39:GFP_cOE parasites also show a localized GFP signal in the nucleus, but the signal is weaker and more diffuse when compared with NF54/GDV1:GFP_cOE (Supplementary Figure 3e). In order to quantify and compare GFP levels in NF54/GDV1:GFP_cOE and NF54/GDV1 Δ 39:GFP_cOE parasites, we performed a whole cell protein extraction for Western blot (WB) analysis using anti-GFP using anti-HSP70 antibodies as a loading control (Figure 4b and c). The WB showed a clear reduction of GDV1 Δ 39:GFP levels compared to GDV1:GFP (Figure 4c). To quantify the localization of GDV1 Δ 39:GFP in the cytoplasm compared to the nucleus we prepared cytosolic and nuclear protein extracts using subcellular fractionation (Figure 4b and d). We determined the cytoplasmic fraction using anti-aldolase antibodies³¹ and anti-histone 3 antibodies were used to determine the nuclear fraction³². GDV1 Δ 39:GFP was only detected in the nuclear fraction further

supporting that its nuclear localisation is not affected by the C-terminal truncation (Figure 4d). Thus, GDV1 Δ 39:GFP protein level is much reduced compared to GDV1:GFP, despite being expressed from the same locus and driven by the same promoter. To test if the GDV1 Δ 39 deletion affects its interaction with HP1, we performed an in vitro assay where 6xHIS-tagged GDV1 WT and Δ 39 versions were co-expressed with Strep-tagged HP1 in *Escherichia coli* bacteria. Interaction between GDV1 and HP1 is detected by affinity purification of HIS:GDV1 and analysis of co-eluted proteins by Coomassie staining¹². HIS-tagged SIP2 does not interact with HP1 and was used as a negative control (Figure 4e). As previously shown, HIS-GDV1 pulled down HP1, which was not observed when SIP2 was used as a bait¹². Interestingly, the Δ 39:GFP of GDV1 also pulled down HP1, indicating GDV1 C-terminus was not essential for the interaction in *E. coli* (Figure 4e). This observation indicates that the interaction of GDV1 Δ 39:GFP and HP1 can still occur in the parasite, but that it is insufficient to trigger gametocytogenesis. An explanation for this could be that GDV1 Δ 39:GFP levels do not reach the threshold required for efficient gametocyte induction. To examine expression of the GDV1 Δ 39 mutant, we analysed the mean fluorescence intensity in uninduced and induced NF54/GDV1:GFP_cOE and NF54/GDV1 Δ 39:GFP_cOE parasites at the single cell level using flow cytometry (Figure 4f and g). As expected, NF54/GDV1:GFP_cOE parasites show a robust increase of GFP fluorescence upon induction of GDV1:GFP expression through glucosamine removal. A measurable increase of the mean fluorescence was also observed upon induction in most NF54/GDV1 Δ 39:GFP_cOE parasites, but well below the levels observed for NF54/GDV1:GFP_cOE parasites. However, a small proportion of NF54/GDV1 Δ 39:GFP_cOE parasites displayed GFP fluorescence at the level observed in the NF54/GDV1:GFP_cOE control line. In line with its nuclear localisation,

GDV1 Δ 39:GFP may contribute to form gametocytes in these parasites. This is further discussed below.

Discussion

The aim of this study was to identify non-essential kinases as regulators of gametocyte commitment/development in *P. falciparum*. While several parasite lines of the kinase knock-out collection¹⁴ were able to form gametocytes, two kinase KO lines showed a gametocytogenesis phenotype that led to the identification of a nonsense mutation in *gdv1* that results in a 39 aa truncation of the GDV1 C-terminus. This mutation may have been acquired by the common parental line prior to generation of the original transgenic lines, although several other clones from the Solyakov study we tested here are able to form gametocytes, possibly reflecting that only a proportion of the parasite population in the parental line carried the mutation. Alternatively, it cannot be excluded that the mutation arose independently in these two lines. This might be clarified by carrying out WGS, but this lies outside the scope of the present study. Our results show that the premature stop codon mutation in *gdv1* resulting in a 39 amino acid C-terminal truncation in the *tlk2* and *elk2* KO lines is sufficient to abolish sexual commitment. Based on our analysis of inducible GDV1 overexpression lines we observed that the truncated GDV1 Δ 39-GFP protein was present at substantially reduced protein levels compared to full-length GDV1-GFP. We propose that it is the loss of GDV1 stability that is the underlying cause for the lack of gametocytes in the GDV1 mutants. Although we have not further tested this here, the reduced amount of GDV1 protein shown in Western blots is likely caused by a destabilisation of the GDV1 protein due to the truncation. Alternatively, although unlikely, it could be caused by a reduction in *gdv1* transcripts. Regardless of the observed decrease of GDV1 protein levels, the truncation does neither result in a strong nuclear localisation defect when overexpressed as a GFP fusion protein, nor in a failure to interact with HP1 expressed in bacteria. It will be important to show in the future whether the few NF54/GDV1 Δ 39:GFP_cOE parasites, which show similar

levels of GDV1 Δ 39:GFP compared to GDV1:GFP in NF54/GDV1:GFP_cOE parasites, are able to induce gametocytogenesis. If they fail to do so, it would indicate additional functions of the GDV1 C-terminus, potentially contributing to bringing GDV1 to the *ap2-g* locus. In this respect, it would be of great interest to identify possible interactors of the GDV1 c-terminus.

Material and Methods

Plasmid construction and transfection.

The construction of each of the ePK knockout plasmids here characterized has been described in ¹⁴. The *pMK-RQ-tkl2-loxPint* donor plasmid (synthesized by Geneart) contains a recodonized version of sequence containing the glycine-rich loop in the kinase domain of *tkl2* (rc. Gly Loop) flanked by two loxPints and homology regions for homology-directed repair. The *pDC2-Cas9-hDHFRyFCU* guideRNA plasmid targeting *tkl2* locus (pDC2_TKL2_gRNA) was generated using the primer pairs pDC2_TKL2_gRNA1_FOR/pDC2_TKL2_gRNA1_REV. Because we didn't have a 3D7::DiCre line, we generated the 3D7/TKL2:loxPint conditional KO line by doing, for the first time, a double transfection with the *pMK-RQ-tkl2-loxPint* and pDC2_TKL2_gRNA, together with the pBSPfs47DiCre (containing the DiCre cassette) and the CRISPR/Cas9 plasmid pDC287 containing the guide RNA targeting the Pfs47 locus, as previously described ²³. The plasmids were suspended in 100uL of P3 primary cell solution, 40ug of each rescue plasmid and 20ug of *pDC2-Cas9-hDHFRyFCU guide RNA* for each respective rescue plasmid, and transfected into the 3D7 parasites. Briefly, purified *P. falciparum* 3D7 schizont stages were electroporated using using Amaxa 4D-Nucleofector™ (Lonza) - program FP158 ³³. Selection of parasites transfected was done using 5nM WR99210 (Jacobus Pharmaceutical) and after a first round of selection, cloned.

To generate the *pMK-RQ-gdv1 Δ 39-HA* plasmid, which upon integration into the endogenous *gdv1* locus mimics the mutation found in the kinase KO lines, the *gdv1*

(PF3D7_0935400) 3' homology region was PCR amplified from NF54 genomic DNA with primers #268/#269 (Supplementary Table 2). The amplified PCR fragment was Gibson-cloned into an AflIII-digested plasmid synthesized by Geneart that contains a *gdv1* 5' homology sequence followed by a recodonized truncated *gdv1Δ39* version and the sequence encoding the 3xHA tag (Supplementary Table 2). To generate the pD_cg6_cam-gdv1Δ39-gfp-glmS plasmid we amplified the *gdv1Δ39* sequence from the pMK-RQ-gdv1Δ39-HA plasmid using primers #383/#384 (Supplementary Table 2) and introduced the PCR fragment using Gibson assembly into the donor plasmid pD_cg6_cam-gdv1-gfp-glmS²⁹ digested with EagI and BsaBI. The guideRNA cassette to mutate endogenous *gdv1* was generated using the primer pairs pDC2_GDV1Δ39_gRNA1_FOR/pDC2_GDV1Δ39_gRNA1_REV and cloned into the pDC2-Cas9-hDHFRyFCU plasmid as previously described²². The rescue plasmid *pMK-RQ-gdv1Δ39-HA* and the CRISPR/Cas9 plasmid *pDC2-Cas9-hDHFRyFCU* were suspended in 100uL of P3 primary cell solution, 40ug and 20ug DNA respectively, and transfected using Amaxa 4D-Nucleofector™ (Lonza). Briefly, purified *P. falciparum* NF54::DiCre schizont stages were electroporated using program FP158³³. Selection of parasites transfected was done using 5nM WR99210 (Jacobus Pharmaceutical) and after a first round of selection, cloned. Transfection of NF54 parasites using the CRISPR/Cas9 pHF_gC-cg6 suicide plasmid²⁹ and the *pD_cg6_cam-gdv1Δ39-gfp-glmS* donor construct was performed as described previously¹². 50 μg each of the suicide plasmid and donor plasmid were transfected and parasites cultured in the presence of glucosamine to block NF54/GDV1Δ39:GFP_cOE protein overexpression. 24 hours after transfection and for six subsequent days in total, the transfected populations were treated with 4 nM WR99210 and then cultured in absence of drug selection until a stably propagating transgenic population was obtained. All primers, guide RNAs and fragments used in the construction and integration of the constructs as well as confirmation of rapamycin mediated excision are described in Supplementary Table 2.

335

336 *Plasmodium falciparum* in vitro culture of asexual and sexual blood stages.

337 *Plasmodium falciparum* parasite lines used in this study were all derived from the
 338 NF54 strain (originally isolated from an imported malaria case in the Netherlands in
 339 the 1980s; BEI Resources, cat. no. MRA-1000)³⁴. Asexual parasites were cultured in
 340 human blood (UK National Blood Transfusion Service) and RPMI 1640 medium
 341 containing 0.5% w/v AlbumaxII (Invitrogen) at 37°C, as previously described³⁵.
 342 Asexual parasites were used to produce gametocytes by seeding asexual rings at
 343 1% or 3% parasitaemia and 4% haematocrit on day 0 and feeding the parasites once
 344 a day during 15 days (day 0 to day 14) in 3% O₂-5% CO₂-92% N₂ gas, in RPMI
 345 complemented with 25mM HEPES, 50mg/liter hypoxanthine, 2g/L sodium
 346 bicarbonate, 10% human serum^{35,36}.

347

348 *Plasmodium falciparum* sexual induction

349 Sexual induction of parasite lines was done by following Trager protocol³⁵. More
 350 specifically, gametocyte induction was started with a 3% asexual ring culture where
 351 sexual commitment was induced by using 50% spent medium, expecting the sexually
 352 committed merozoites to invade and develop during the next cycle^{20,35}. The
 353 overexpressing NF54/GDV1:GFP_cOE and NF54/GDV1Δ39:GFP_cOE parasite lines
 354 were kept in the constant presence of 2.5 mM glucosamine to block ectopic GDV1
 355 expression and therefore sexual induction, while sexual induction was achieved
 356 culturing the parasites in the absence of glucosamine, as previously described²⁹.

357

358 Time course of gametocyte induction, RNA extraction and RNA-seq library 359 preparation.

360 The samples were collected during the asexual cycle at 28-32 hpi and in the
 361 matching cycle at 28-32 hpi after induction of sexual commitment. The infected RBCs
 362 pellets were collected at the respective time point, centrifuged and solubilized in ten
 363 volumes of TRIzol (Ambion) prewarmed to 37°C, lysed for 5 minutes by mixing

vigorously at 37°C and immediately frozen at -80°C until extraction. Complete RNA was isolated from the samples using Trizol/chloroform extraction followed by isopropanol precipitation²² and its concentration and integrity was verified using Agilent Bioanalyzer (RNA 6000 Nano kit) and NanoDrop 1000 spectrophotometer. 1-2 µg of total RNA from each sample (or complete sample if the yield was lower) was used for mRNA isolation (Magnetic mRNA Isolation Kit, NEB). First strand cDNA synthesis was performed using the SuperScript III First-Strand Synthesis System and a 1:1 mix of Oligo(dT) and random primers (Invitrogen). The DNA-RNA hybrids were purified using Agencourt RNACleanXP beads (Beckman Coulter) and the second cDNA strand was synthesized using a 10 mM dUTP nucleotide mix, DNA Polymerase I (Invitrogen) and RNaseH (NEB) for 2.5 h at 16°C. The long cDNA fragments were purified and fragmented using a Covaris S220 system (duty cycles = 20, intensity = 5, cycles/burst = 200, time = 30s). The ~200 bp long fragments were end-repaired, dA-tailed and ligated to “PCR-free” adapters³⁷ with index tags using NEBNext according to the manufacturer’s instructions. Excess adapters were removed by two rounds of clean-up with 1 volume of Agencourt AMPure XP beads. Final libraries were eluted in 30 µl water, quality-controlled using Agilent Bioanalyzer (High Sensitivity DNA chip) digested with USER enzyme (NEB) and quantified by qPCR. For some libraries additional 5 cycles of PCR amplification were performed, using KAPA HiFi HotStart PCR mix and Illumina tag-specific primers to obtain enough material for sequencing. Pools of indexed libraries were sequenced using an Illumina HiSeq2500 system (100 bp paired-end reads) according to manufacturer’s manual. All samples were generated in duplicates or triplicates and uninduced controls were always generated and processed in parallel. Raw data is available through GEO database repository (study GSE158689).

RNAseq data analysis

The generation of raw data in the form of *.cram files quality control and adapter trimming was performed using the default analysis pipelines of the Sanger Institute.

The raw data was transformed into paired *. fastq files using Samtools software (ver. 1.3.1). The generated reads were re-aligned to *Plasmodium falciparum* genome (PlasmoDB-30 release) in a splice aware manner with HISAT2³⁸ using --known-splicesite-infile option within the splicing sites file generated based on the current genome annotation. Resulting *.bam files were sorted and indexed using Samtools and inspected visually using Integrated Genome Viewer (ver. 2.3.91). HT-seq python library³⁸ was used to generate reads counts for all genes for further processing. Raw counts were normalised to median-ratio and then tested against linear models of time nested in line and line nested within time using a negative binomial model for the normalised counts using DESeq2, differential genes being selected for a false discovery rate of < 0.1³⁹.

Saponin lysis and whole cell, cytoplasmic and nuclear protein extraction

10mL parasite culture (2-5% parasitemia, 4% haematocrit) was transferred to a 15 mL tube and centrifuged at 600 g for 5 min. The supernatant was aspirated and the RBC pellet resuspended in 5 volumes 0.15% saponin solution (2.5 mL for 500 µL RBC). After an incubation on ice of max. 10 min, the parasites were centrifuged at 1503 g for 5 min at 4°C. Subsequent steps were performed on ice in order to prevent protein degradation. The supernatant was aspirated and the parasite pellet resuspended in 1 mL cold phosphate buffered saline (PBS) and transferred to an Eppendorf tube. The parasite pellet was centrifuged at 1503 g for 30 sec at 4°C and washed with cold PBS until the supernatant was clear.

For whole cell protein extraction, one pellet volume (30-50 µL) of whole cell protein lysis buffer (8 M Urea, 5% SDS, 50 mM Bis-Tris, 2 mM EDTA, 25 mM HCl, pH 6.5) complemented with 1x protease inhibitor cocktail (Merck) and 1 mM DTT was added to the pellet at RT in order to lyse the parasites. The tube was vortexed, heated to 94°C for 5 min, sonicated for 2 min (5 cycles of 30 sec ON/ 30sec OFF), vortexed and heated again. Subsequently, the protein sample was centrifuged at

20238 g for 5 min at RT and the supernatant was transferred into a new tube, which was frozen at -20°C and stored until use.

For cytoplasmic and nuclear protein extraction, the parasite pellet was lysed in 300µL cytoplasmic lysis buffer (20 mM Hepes (pH 7.9), 10 mM KCl, 1 mM EDTA, 0.65% Igepal) complemented with 1x protease inhibitor cocktail (Merck) and 1 mM DTT (leaving the nucleus intact) and incubated on ice for 5 min (Voss et al., JBC, 2002). The lysed parasites were centrifuged at 845 g for 3 min, the supernatant representing the cytoplasmic protein fraction was transferred into a new tube and placed on ice. The remaining nuclear pellet was washed in 500 µL cytoplasmic lysis buffer and centrifuged at 845 g for 3 min. The washing was repeated until the supernatant was clear. The nuclear pellet was resuspended in 60 µL whole cell lysis buffer and vortexed at high speed at RT for 10-20 min. The insoluble material was centrifuged at 20238 g for 3 min, the supernatant representing the nuclear protein fraction was transferred to a new tube and placed on ice. Both protein fractions were frozen at -20°C and stored until use.

Western Blot.

Parasite extracts were solubilized in protein loading buffer, denatured at 95°C for 10min, subjected to SDS-PAGE and transferred onto a nitrocellulose membrane. Membranes were immunostained with mouse anti-GFP (1:250 dilution; Roche, 11814460001), rabbit anti-Aldolase-HRP conjugated (1:5000 dilution, abcam ab38905) and rabbit anti-Histone 3 (1:2000 dilution, abcam ab1791) primary antibodies. Antibody detection was done using chemiluminescent western blot detection using goat anti-mouse secondary antibody conjugated with HRP and the ECL western blotting detection reagents (Amersham RPN2106) or by direct infrared fluorescence detection on the Odyssey Infrared Imaging System (Odyssey CLx, LI-COR) using IRDye 680LT goat anti-rat IgG (1:10000 dilution; LI-COR) and IRDye 800CW goat anti-rabbit IgG (1:10000 dilution; LI-COR).

Immunofluorescence assay at different parasite stages. Air-dried thin blood films of asexual parasites were fixed with 4% paraformaldehyde containing 0.0075% glutaraldehyde for 15 min and permeabilized in 0.1% (v/v) Triton X-100 (Sigma) for 10 min⁴⁰. Blocking was performed in 3% BSA for 1 h. Slides were incubated with rat anti-HA high-affinity (1:1000 dilution; Roche, clone 3F10) at room temperature for 30 min, followed by Alexa fluor conjugated goat anti-rat IgG (1:1000 dilution; Thermo Fisher Scientific) at room temperature for 30 min. Parasite nuclei were stained with 4', 6-diamidino-2-phenylindole (DAPI; Invitrogen). Slides were mounted in ProLong® Gold antifade reagent (Invitrogen) and images were obtained with the inverted fluorescent microscope (Ti-E; Nikon, Japan) and processed using NIS-Elements software (Nikon, Japan).

Flow Cytometry

NF54/GDV1:GFP_cOE and NF54/GDV1Δ39:GFP_cOE parasites were grown in the presence or absence of glucosamine in order to block or allow sexual commitment, respectively. Schizonts from the 4 conditions were purified by percoll gradient and allowed to invade fresh red blood cells for 4h, before uninvaded schizonts were removed. Flow cytometry analysis was performed at approximately 44h post invasion, in 4 biological replicates. For one replicate, parasites were fixed for 1h in 4% paraformaldehyde in PBS, stained with Hoechst 33342 (1:1000 in PBS) for 10 minutes and analysed on an LSRFortessa flow cytometer (Becton Dickinson) using FACSDiva software. For the other three replicates, live parasites were stained with Hoeschst 33342 and analysed on a BD FACSAria II flow cytometer (Becton Dickinson) using FACSDiva software. Hoechst fluorescence was detected using a 355nm (UV) excitation laser with a 450/50nm bandpass filter, while GFP fluorescence was detected with a 488nm (blue) excitation laser, a 505nm longpass filter and a 530/30nm bandpass filter. At least 30,000 cells were counted for each sample. Data were analysed using FCS Express 7 (Research Edition) software. The population was first gated on single cells based on the side and forward scatter, then on highly

Hoechst-positive infected schizonts, before the median fluorescence intensity (MFI) of the GFP-fluorescence was calculated for each line. An example of the gating strategy for infected cells is shown in Supplementary Figure 4. Due to the variation in fluorescence intensity between different experiments, MFI values were normalised by dividing the MFI of each infected sample by the average MFI of the uninfected samples within the same experiment (n=4). Statistical analysis was performed using Holm-Sidak corrected multiple comparison analysis of variance (ANOVA) on samples paired within each experiment using Graphpad Prism version 8.

In vitro protein-protein interaction experiments

In order to co-express Strep(II)-tagged HP1 with a His-SUMO-tagged truncated version of GDV1, we deleted the 39 C-terminal amino acids of the coding sequence of GDV1 in the vector pStrep-HP1_HS-GDV1¹². For this purpose, we circularized a PCR product amplified from this vector with the primers D39F and D39R using Gibson assembly. The proteins were expressed and the in vitro interaction assay was performed as previously described (14) using full-length GDV1 as positive and SIP2 as negative control.

FUNDING

This work was supported by the Marie Skłodowska-Curie Individual Fellowship to Marta Tibúrcio (grant agreement 661167 — PFSEXOME — H2020-MSCA-IF-2014) and core funding to MT by the Francis Crick Institute (<https://www.crick.ac.uk/>), which receives its core funding from Cancer Research UK (FC001189; <https://www.cancerresearchuk.org>), the UK Medical Research Council (FC001189; <https://www.mrc.ac.uk/>), and the Wellcome Trust (FC001189; <https://wellcome.ac.uk/>). The Bioinformatics and Flow Cytometry STPs are supported through Crick Core funding (FC001999). This work was further supported by a research grant to T.V from the Swiss National Science Foundation (BSCGI0_157729).

OB acknowledges funding by Wellcome core grant 206194/Z/17/Z to the Sanger Institute.

ACKNOWLEDGEMENTS

We would like to thank Christian Doerig for the original *Plasmodium falciparum* kinase KO lines. We would like to thank Frank Schwach and Mandy Sanders for preparing, running and initial quality control of RNAseq samples. We would like to thank Ellen Knuepfer and Christiaan van Ooij for the pBSPfs47DiCre and pDC287 plasmids, as well as scientific advice. We would like to thank Kostas Kousis for his help with the Flow Cytometry data acquisition.

AUTHOR CONTRIBUTIONS

M. Tibúrcio and M. Treeck conceived the study. M. Tibúrcio performed most of the parasite genetic manipulations and all the parasite line phenotyping experiments, as well as RNAseq material collection. E. Hitz generated the NF54/GDV1Δ39:GFP_cOE parasite line, I. Niederwieser performed the in vitro protein-protein interaction experiments and T. S. Voss supervised these experiments and provided conceptual advice and resources. Gavin Kelly performed RNAseq analysis. RNAseq samples were run in Oliver Billkers group at the Sanger Institute. H. Davies performed the Flow Cytometry Data Analysis. Christian Doerig provided the original *P. falciparum* kinase KO cell lines. All authors contributed to experimental design and interpretation of the results. M. Tibúrcio and M. Treeck wrote the article with contributions from all authors.

Competing interests

We declare that we have no competing interests.

References

1. World Malaria Report 2019. World Health Organization, 2019.
2. Kafsack BF, Rovira-Graells N, Clark TG, et al. A transcriptional switch underlies commitment to sexual development in malaria parasites. *Nature* 2014; 507(7491): 248-52.
3. Aguilar R, Magallon-Tejada A, Achtman AH, et al. Molecular evidence for the localization of *Plasmodium falciparum* immature gametocytes in bone marrow. *Blood* 2014; 123(7): 959-66.
4. De Niz M, Meibalan E, Mejia P, et al. *Plasmodium* gametocytes display homing and vascular transmigration in the host bone marrow. *Sci Adv* 2018; 4(5): eaat3775.
5. Farfour E, Charlotte F, Settegrana C, Miyara M, Buffet P. The extravascular compartment of the bone marrow: a niche for *Plasmodium falciparum* gametocyte maturation? *Malar J* 2012; 11: 285.
6. Joice R, Nilsson SK, Montgomery J, et al. *Plasmodium falciparum* transmission stages accumulate in the human bone marrow. *Sci Transl Med* 2014; 6(244): 244re5.
7. Smalley MEA, S.; Brown, J. The distribution of *Plasmodium falciparum* in the peripheral blood and bone marrow of Gambian children. *Trans R Soc Trop Med Hyg* 1981; 75: 103-5.
8. Thomson JGR, A. The structure and development of *Plasmodium falciparum* gametocytes in the internal organs and peripheral circulation. *Trans R Soc Trop Med Hyg* 1935; 29 31–40.
9. Marchiafava EAB, A. Sulle Febbri Malariche Estivo Autunnali Innocenzo Artero; 1892.
10. Brancucci NMB, Gerdt JP, Wang C, et al. Lysophosphatidylcholine Regulates Sexual Stage Differentiation in the Human Malaria Parasite *Plasmodium falciparum*. *Cell* 2017; 171(7): 1532-44 e15.
11. Sinha A, Hughes KR, Modrzynska KK, et al. A cascade of DNA-binding proteins for sexual commitment and development in *Plasmodium*. *Nature* 2014; 507(7491): 253-7.

- 561 12. Filarsky M, Frschka SA, Niederwieser I, et al. GDV1 induces sexual
562 commitment of malaria parasites by antagonizing HP1-dependent gene silencing.
563 Science 2018; 359(6381): 1259-63.
- 564 13. Brancucci NMB, Bertschi NL, Zhu L, et al. Heterochromatin protein 1 secures
565 survival and transmission of malaria parasites. Cell Host Microbe 2014; 16(2): 165-
566 76.
- 567 14. Solyakov L, Halbert J, Alam MM, et al. Global kinomic and phospho-proteomic
568 analyses of the human malaria parasite Plasmodium falciparum. Nat Commun 2011;
569 2: 565.
- 570 15. Tewari R, Straschil U, Bateman A, et al. The systematic functional analysis of
571 Plasmodium protein kinases identifies essential regulators of mosquito transmission.
572 Cell Host Microbe 2010; 8(4): 377-87.
- 573 16. Lopez-Barragan MJ, Lemieux J, Quinones M, et al. Directional gene
574 expression and antisense transcripts in sexual and asexual stages of Plasmodium
575 falciparum. BMC Genomics 2011; 12: 587.
- 576 17. Abdi AI, Carvalho TG, Wilkes JM, Doerig C. A secreted Plasmodium falciparum
577 kinase reveals a signature motif for classification of tyrosine kinase-like kinases.
578 Microbiology 2013; 159(Pt 12): 2533-47.
- 579 18. Pelle KG, Oh K, Buchholz K, et al. Transcriptional profiling defines dynamics
580 of parasite tissue sequestration during malaria infection. Genome Med 2015; 7(1):
581 19.
- 582 19. Lasonder E, Rijpma SR, van Schaijk BC, et al. Integrated transcriptomic and
583 proteomic analyses of P. falciparum gametocytes: molecular insight into sex-specific
584 processes and translational repression. Nucleic Acids Res 2016; 44(13): 6087-101.
- 585 20. Fivelman QL, McRobert L, Sharp S, et al. Improved synchronous production
586 of Plasmodium falciparum gametocytes in vitro. Mol Biochem Parasitol 2007; 154(1):
587 119-23.

- 588 21. Claessens A, Affara M, Assefa SA, Kwiatkowski DP, Conway DJ. Culture
589 adaptation of malaria parasites selects for convergent loss-of-function mutants. Sci
590 Rep 2017; 7: 41303.
- 591 22. Knuepfer E, Napiorkowska M, van Ooij C, Holder AA. Generating conditional
592 gene knockouts in Plasmodium - a toolkit to produce stable DiCre recombinase-
593 expressing parasite lines using CRISPR/Cas9. Sci Rep 2017; 7(1): 3881.
- 594 23. Tiburcio M, Yang ASP, Yahata K, et al. A Novel Tool for the Generation of
595 Conditional Knockouts To Study Gene Function across the Plasmodium falciparum
596 Life Cycle. mBio 2019; 10(5).
- 597 24. Zhang M, Fennell C, Ranford-Cartwright L, et al. The Plasmodium eukaryotic
598 initiation factor-2alpha kinase IK2 controls the latency of sporozoites in the mosquito
599 salivary glands. J Exp Med 2010; 207(7): 1465-74.
- 600 25. Poran A, Notzel C, Aly O, et al. Single-cell RNA sequencing reveals a signature
601 of sexual commitment in malaria parasites. Nature 2017; 551(7678): 95-9.
- 602 26. Bancells C, Llorca-Batlle O, Poran A, et al. Revisiting the initial steps of sexual
603 development in the malaria parasite Plasmodium falciparum. Nat Microbiol 2019;
604 4(1): 144-54.
- 605 27. Josling GA, Russell TJ, Venezia J, et al. Dissecting the role of PfAP2-G in
606 malaria gametocytogenesis. Nat Commun 2020; 11(1): 1503.
- 607 28. Usui M, Prajapati SK, Ayanful-Torgby R, et al. Plasmodium falciparum sexual
608 differentiation in malaria patients is associated with host factors and GDV1-
609 dependent genes. Nat Commun 2019; 10(1): 2140.
- 610 29. Boltryk SD, Passecker A, Alder A, et al. CRISPR/Cas9-engineered inducible
611 gametocyte producer lines as a novel tool for basic and applied research on
612 Plasmodium falciparum malaria transmission stages. bioRxiv 2020.
- 613 30. Prommana P, Uthapibull C, Wongsombat C, et al. Inducible knockdown of
614 Plasmodium gene expression using the glmS ribozyme. PLoS One 2013; 8(8):
615 e73783.

31. Knapp B, Hundt E, Kupper HA. Plasmodium falciparum aldolase: gene structure and localization. Mol Biochem Parasitol 1990; 40(1): 1-12.
32. Salcedo-Amaya AM, van Driel MA, Alako BT, et al. Dynamic histone H3 epigenome marking during the intraerythrocytic cycle of Plasmodium falciparum. Proc Natl Acad Sci U S A 2009; 106(24): 9655-60.
33. Moon RW, Hall J, Rangkuti F, et al. Adaptation of the genetically tractable malaria pathogen Plasmodium knowlesi to continuous culture in human erythrocytes. Proc Natl Acad Sci U S A 2013; 110(2): 531-6.
34. Delves MJ, Straschil U, Ruecker A, et al. Routine in vitro culture of P. falciparum gametocytes to evaluate novel transmission-blocking interventions. Nat Protoc 2016; 11(9): 1668-80.
35. Trager W, Jensen JB. Human malaria parasites in continuous culture. Science 1976; 193(4254): 673-5.
36. Delves MJ, Ruecker A, Straschil U, et al. Male and female Plasmodium falciparum mature gametocytes show different responses to antimalarial drugs. Antimicrob Agents Chemother 2013; 57(7): 3268-74.
37. Kozarewa I, Ning Z, Quail MA, Sanders MJ, Berriman M, Turner DJ. Amplification-free Illumina sequencing-library preparation facilitates improved mapping and assembly of (G+C)-biased genomes. Nat Methods 2009; 6(4): 291-5.
38. Anders S, Pyl PT, Huber W. HTSeq--a Python framework to work with high-throughput sequencing data. Bioinformatics 2015; 31(2): 166-9.
39. Love MI, Huber W, Anders S. Moderated estimation of fold change and dispersion for RNA-seq data with DESeq2. Genome Biol 2014; 15(12): 550.
40. Tonkin CJ, van Dooren GG, Spurck TP, et al. Localization of organellar proteins in Plasmodium falciparum using a novel set of transfection vectors and a new immunofluorescence fixation method. Mol Biochem Parasitol 2004; 137(1): 13-21.

Figure Legends

Figure 1. Screening of Plasmodium falciparum non-essential kinases during sexual commitment and development. A) List of non-essential kinases characterized during sexual development in this study (PF3D7_0107600 – eIK2; PF3D7_0525900 - NEK2; PF3D7_0615500 – CRK5; PF3D7_0623800 – TKL4; PF3D7_0715300 - calcium/calmodulin-dependent protein kinase, putative; PF3D7_1121300 – TKL2; PF3D7_1201600 - NEK3; PF3D7_1423600 - calcium-dependent protein kinase, putative; PF3D7_1450000 - serine/threonine protein kinase, putative) ¹⁴. B) Comparison of the percentage of gametocytaemia between the 3D7 WT line and the PftKL2 kinase KO clones of the same transfection (clones B10 and B12) generated by single crossover integration ¹⁴. Each column represents the mean of triplicate microscope counts, each of at least 500 cells, analysed using paired t test, $\pm$ SD, (*; $p < 0.05$; 3D7 versus TKL2 KO clones, $P = 0.0377$). C) Schematic of the CRISPR/Cas9 strategy used to generate a TKL2 conditional knockout (KO) line (3D7/TKL2:loxPint) as well as the primers used to confirm successful gene editing (Supplementary Table 2); The *pMK-RQ-tkl2-loxPint* donor plasmid contains a recodonized version of the

glycine loop in the kinase domains of *tkl2*(rc. Gly Loop) flanked by two loxPints and
homology regions for homology-directed repair. D) Sexual conversion rates in
3D7/TKL2:loxPint parasites treated with DMSO (control) or rapamycin (KO). Each
column represents the mean of triplicate microscope counts, each of at least 500
cells, analysed using paired t test $\pm$ SD, (ns, $p \geq 0.05$; 3D7/TKL2:loxPint treated with
DMSO versus Rapamycin, $P=0.4017$).

Figure 2. RNA sequencing analysis comparing WT and PfeiK2 gametocyte-less kinase KO line. A) Representation of the interactive cycles of asexual and sexual differentiation upon sexual induction; dotted box illustrates the time point and asexual and sexual stages of the parasite collected for RNAseq. B) Heatmap showing genes previously described as being associated with AP2-G expression and significantly

downregulated in the PfkK2 kinase KO clones (log2 fold change >1). C) DNA sequence trace showing the stop mutation identified in the PfkK2 and PfkKL2 KO clones which is absent in the 3D7 reference parasite line.

Figure 3. Quantification of gametocyte production in GDV1Δ39 mutant parasite lines.

A) Illustration of the strategy used to generate the 3xHA-tagged GDV1Δ39 mutant line (NF54/GDV1Δ39:HA) as well as the primers used to confirm integration (Supplementary Table 2); The *pMK-RQ-gdv1Δ39-HA* donor plasmid contains a recodonized version of the *gdv1Δ39* mutant and a 3xHA tag, flanked by homology regions. B) Comparison of sexual conversion rates between NF54 and NF54/GDV1Δ39:HA parasite lines. Each column represents the mean of duplicate (NF54) and triplicate (NF54/GDV1Δ39:HA) microscope counts, each of at least 500

689 cells, analysed using paired t test, $\pm$ SD, (*, $p < 0.05$; NF54 versus NF54/GDV1 Δ 39:HA,
690 $P = 0.0489$). C) Schematic of the strategy used to make the NF54/GDV1 Δ 39:GFP_cOE
691 overexpressing line as well as the primers used to verify integration of the transgene
692 cassette into the *cg6* (*gfp3*) locus (Supplementary Table 2). The *pD_cg6_cam-*
693 *gdv1 Δ 39-gfp-glmS* donor plasmid contains a recodonized version of the *gdv1 Δ 39*
694 mutant followed by the in-frame *gfp* sequence and the *glmS* ribozyme element,
695 flanked by homology regions. D) Comparison of sexual conversion rates between
696 NF54/GDV1:GFP_cOE and NF54/GDV1 Δ 39:GFP_cOE parasite lines in the presence
697 (prevents sexual conversion) or absence of glucosamine (induces sexual
698 conversion). Each column represents the mean of triplicate counts of at least 500
699 cells, analysed using paired t test, $\pm$ SD, (*, $p < 0.05$; ns, not significant, $p \geq 0.05$;
700 NF54/GDV1:GFP_cOE non-induced versus induced, $P = 0.0094$; non-induced
701 NF54/GDV1:GFP_cOE versus NF54/GDV1 Δ 39:GFP_cOE, $P = 0.2276$;
702 NF54/GDV1 Δ 39:GFP_cOE non-induced versus induced, $P = 0.4038$; induced
703 NF54/GDV1:GFP_cOE versus NF54/GDV1 Δ 39:GFP_cOE, $P = 0.0281$).
704

Figure 4. GDV1Δ39 expression, localization and interaction with HP1. A) Representation of the protocol used to collect the samples used to characterize expression and localization of GDV1Δ39:GFP. B) Illustration of the subcellular fractionation workflow. C) Western blot showing the levels of GDV1:GFP expression

in NF54/GDV1:GFP_cOE and NF54/GDV1Δ39:GFP_cOE parasites grown in the absence of glucosamine (induces expression); GDV1:GFP/ GDV1Δ39:GFP expression is detected using an anti-GFP antibody while anti-HSP70 antibodies have been used as controls. D) Western blot showing GDV1:GFP expression levels in the cytoplasmic and nuclear fraction in NF54/GDV1:GFP_cOE and NF54/GDV1Δ39:GFP_cOE parasites cultured in the absence of glucosamine (induces expression). E) Strep-HP1 co-purifies with both HIS-GDV1 and HIS-GDV1Δ39 but not with the HIS-SIP2 control. Coomassie-stained SDS-polyacrylamide gel from pull down experiment with HIS-GDV1/Strep-HP1 and HIS-SIP2/Strep-HP1. Lane 4: protein size standard. F) Representative normalised flow cytometry histograms quantifying GDV1:GFP fluorescence for each parasite line. The experiment was repeated 4 times with similar results. Dotted lines indicate the position of the peaks for the wild-type NF54/GDV1:GFP_cOE line. G) Quantification of the median fluorescence intensity of GDV1:GFP in induced or uninduced NF54/GDV1:GFP_cOE and NF54/GDV1Δ39:GFP_cOE parasite lines, normalised to uninfected parasites from each experiment, $\pm$ SD, $n = 4$. (*, $p < 0.05$; ns, not significant, $p \geq 0.05$; NF54/GDV1:GFP_cOE uninduced vs induced, $p = 0.0227$; NF54/GDV1Δ39:GFP_cOE uninduced vs induced, $p = 0.0318$; NF54/GDV1:GFP_cOE induced vs NF54/GDV1Δ39:GFP_cOE induced $p = 0.0227$; NF54/GDV1:GFP_cOE uninduced vs NF54/GDV1Δ39:GFP_cOE uninduced, $p = 0.1235$); Statistical analysis was performed using Holm-Sidak corrected multiple comparison analysis of variance (ANOVA).

Supplementary Figure 1. Confirmation of *tkI2-loxPint* cassette integration into the *Plasmodium falciparum* 3D7 line and the efficiency of DiCre mediated excision. A) Representation of the primer pairs used to test correct integration of *tkI2-loxPint* cassette and efficient rapamycin mediated excision. B) PCR analysis shows correct integration of *tkI2-loxPint* cassette and near complete excision of TKL2:LoxPint in two different clones from the same transfection (clone 13 and 33) after rapamycin treatment. The sequences of the primers used are in Supplementary Table 2.

Supplementary Figure 2. PfkI2 kinase KO parasites fail to produce gametocytes. A) Comparison of gametocytemia upon sexual induction between the 3D7 WT line (n=4) and two PfkI2 kinase KO clones (n=2 for each clone, both from the same transfection). Parasite line and clones measured in at least two biological experiments with single replicates. Each column represents the mean number of gametocytes in at least 500 cells. B) Representation of GDV1 sequences of the NF54, 3D7, PfkI2 and TKL2 parasite lines and the identified point mutation in the PfkI2 and PfkTKL2 KO clones (red arrow) which is absent in the 3D7 and NF54 reference parasite lines.

Supplementary Figure 3. Generation of the NF54/GDV1Δ39:HA and NF54/GDV1Δ39:GFP_cOE *Plasmodium falciparum* lines. A) Illustration of the strategy used to generate the 3xHA-tagged GDV1Δ39 mutant line (NF54/GDV1Δ39:HA) as well as the primers used to confirm donor sequence integration. B) PCR analysis of integration of the *gdv1*Δ39:ha construct into the *gdv1* locus in the NF54 *P. falciparum* parasite line. C) Schematic of the strategy used to generate the NF54/GDV1Δ39:GFP_cOE overexpressing line as well as the primers

used to verify integration of the transgene cassette into the *cg6 (gfp3)* locus. The pGDV1Δ39:GFP_cOE donor plasmid contains a 5' Cam sequence followed by a recodonized version of the *gdv1Δ39* mutant in-frame with *gfp* sequence, the *glmS* ribozyme element and *Plasmodium berghei* dihydrofolate reductase (PfDHFR), flanked by two homology regions. D) PCR analysis of *gdv1Δ39-gfp-glmS* cassette integration in the *cg6* locus in the NF54 *P. falciparum* parasite line and the presence of the CRISPR/Cas9/Suicide plasmid. E) Quantification of GFP expression in the GDV1:GFP and GDV1Δ39:GFP lines cultured in the presence (prevents GDV1 expression) or absence of glucosamine (induces GDV1 overexpression).

Supplementary Figure 4. A) Gating strategy for flow cytometry experiments. B) Flow cytometry histograms quantifying GFP fluorescence in uninfected RBC and RBC infected with ring stage parasites, trophozoites and schizonts. Stacked plots are shown for both GDV1:GFP uninduced and induced parasites. C) Normalised flow cytometry histograms quantifying GFP fluorescence for each line. The experiment was repeated 4 times with similar results. Dotted lines indicate the position of the peaks for the wild-type NF54/GDV1:GFP_cOE line.

Supplementary Table 1. Comparison of gene expression, based on RNAseq data, of samples from 3D7 WT and PflK2 C3 and F12 kinase KO clones.

Log2 fold changes of induced vs. non-induced parental (3D7) and the 2 *gdv1* mutant parasite clones from the PflK2 KO clones C3 and F12.

Supplementary Table 2. Primers, fragment and guide RNA sequences used to generate and confirm integration and rapamycin-induced excision of *pMK-RQ-tkl2-loxPint*, *pMK-RQ-gdv1Δ39-HA* and *pD_cg6_cam-gdv1Δ39-gfp-glmS*.

Primer	Sequence
#145	AGCAAAAGTCTTAAGCTCATGGAGG
#146	ATGAATAGCAGGAGATTTAGATACC
pDC2_TKL2_gRNA_FOR	attgATAATAGAATTGCAAAAGGA
pDC2_TKL2_gRNA_REV	aaacTCCTTTTGCAATTCTATTAT
<i>tkl2-loxPint</i>	CTGGAAAGCGGGCAGTGAAAGGAAGGCCCATGAGGCC CAGATAATAGGGGTCTGCATATCCTGGGGTTCCTCCTGT CAAGTTAAACACATTATTGTTGTTTCATGTAAACAAAGGAA AGCCCAAATCACCAAGCTTAGCATTAAATTGATCATCTA TTAAATGTTTGCAGATTTTAAATCTCTATGATATACAATA GGAGAAGATGTATGTAAATAACATAAGACATTAATTATTT GTACTAATATATTTATTCTTATATTAAATGATAAAAATAGA GGTGTGTTATTATCAAATGAATTTTGTTGAAATGGAGAAT ATTTTGAATTATTGAAATATGGGAAAATGTTTTGTTTATAA AACGTTGAAGTATAATAATTTGTAGAAGAACAAGAAGAA GAACTTTTTTTTTTACGTAGATAATTTTCATAACAACCTTATA TTATAATAACTTAATTCTGGATTTTCTTTGTTTTTTGAATTAT AATAATAATAATGATTAAATAATAATGTTCTTAAATCACCT AGATTAACATATTCATATATTAAATAAAAATTATTTTTATTC GTAGCATAACCTAATAAAGATCTACAAAAAAGAATAAAAA TAAAAAATAAAAAATAAAAAAATAAACATATATATATATA

	TACATATATATACATATATATATCCATATATACATATTAATT CATTTCATAACTTCGTATAATGTATGCTATACGAAGTTAT AACTTGCATATATTTTATACATATTCTGTATTACAAGATGTT GTTGTGTCTGTATCTGGACATGATAATGATTTTCGTTTTCG AAACCGTTGTTTTCGTTCTTTTTCAaGACCTTAATGGCAAC GTTGATGCAGTTCTTCAAAACaCCTTTGTAAACAGTACCG TTACCACCCTTGGCGATTCTAAAAGAATATAAAATATAT AAATATATATATATATACATATATAATAACTTCGTATAATGT ATGCTATACGAAGTTATTGTATATTATTTTTTTTATTTACTAT TATATTCTGAAAAATTATTTGTTGCTTCTACTAAATCATTAA AATCATATTTATTAGATAAAATCTTTAAAGGAATAAATATA TTCTCTTCACGTTTTCTATTCTGTAAATTCCATACCTTCA TTTGGTGTCTTGTATTCGACATTTTCTCTATTTATTTTCATCT TCATAAATCTCATTTTCTTCATCATACATATTTAAAATGTAT TTATCATAAGTATCATATGTTTCTATATATACTACCTTTCTA TTCTTATGTCTATTTCGAATTCTTTCTATTAATATCGTTGTTA TTTTTGATATGATTATTATTTATATGGTTATTGCTCACATGA ATATTATTAATATGGTTATTGCTCACATGGATATTATTAATA TGGTCATtGKTCACATGGATATTTTAAATATGtCTATTATTC ATATAATTGTTTATAtATTTATTATTAATATGTTTATCATTCA CATGATGATTATTAATATTGAAATTATGGTTATTATTTATAT CAATCTCTTTCTTCTTTTCATCCCCCTTCC
#268 (GDV1_TAA_GIB FOR1)	TAG GCC CAG GAA GGA AAA CAA AAA CGT CAG CAT TCA TCA TC
#269 (GDV1_TAA_GIB REV1)	AGT GAA AGG AAG GCC CAT GAG GCC CAG TGA ACA TGC TAT ATT CTT ATG TAT GTA CC
#383 (GDV1_TAA_GIB FOR2)	TGTTCCCTAATATAAACTCAGATATAAATGTTGATACC
#384 (GDV1_TAA_GIB REV2)	GTTCTTCTCCTTTACTCATGCGGCCATAGATAATATCGTC CAGCAGCAGAAC
pDC2_GDV1Δ39_gRNA1_FOR	ATTGCTACAAATTTTGTAATTAAT
pDC2_GDV1Δ39_gRNA1_REV	AAACATTAATTACAAAATTTGTAG
<i>gdv1Δ39-HA</i>	GATTCCGATGTAATATCTTATAGTTATCATTTTCGACCATAC AATCATTTATTGTATTAAGATAAATATAAAAAATGTTACAA ATGTTCCCTAATATAAACTCAGATATAAATGTTGATACCATA ACACATCCTGAAACAACAACAACATATCCAATCATTATA AAAATCATAAACACTATTATGTTTAAAAAACCTGTGATTCTT ATCTGTTCTTTGAATAATTTACCCTTAAAAAATAATAAAAT ACATGATGATATAATTCTTGACGATTTCTGTTATATGACCT ACATTATTAAACTTTGTATTTTCGTTTTCAATCAAGAAGT ATACGCAACATGCTCAAAAGCAATCTGAATAACACCACC AACTTTGTGATTAACCGTATCACCCAGATGAATAATACCA ATCGCCTGCTGAAAAACAAGATTAATGTTCTGCTGCTGG ACGATATTATCTATTACCCATACGATGTTCCAGATTACGC TTATCCGTATGATGTGCCGGATTATGCGTACCCATACGA TGTTCCAGATTACGCTTAAATTAGGCCCAGGAAGGAAAA CAAAAACGTCAGCATTTCATCATCTTCTACAAAAGGCTTAT TCTCCATTCTATTATCCAATTCATTATTATATAAAAGAAAA ATGTACATATAAATAAATAAATGAAAAAAAAAAAAAAAAAGAA AAATAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAGAAAAA GAAAAAAGAAAAAAGTAAAAAGAAAAAAGAAAAAATTTA TATGTAATATAATTAATAAATTTGTATGCTAATATATGTGTA TAATATGTATCTCATCGTGTACATGTATATAATATATATAT ATTTTGTAGGTAAATATTATATCTACTTTTAAACATACATATA TATGTAATATATATATATATATATTACTAAAAAATATAAAGG TTTCAAAACATATATGTTTGCTTTTATAATTTTAAAAAAAAA

	TTCTTATCTATGTTGGTACATACATAAGAATATAGCATGTT CA
#123	AAAGGTGTTTTGAAGAACTGCATCAACG
#124	TATCTGGACATGATAATGATTTTCG
#145	AGCAAAAGTCTTAAGCTCATGGAGG
#146	ATGAATAGCAGGAGATTTAGATACC
#147	GGAGGGAATGGAACAGTATATAAA
#148	TTTATATACTGTTCCATTCCCTCC
#332	ACAATACTACAAATTTTGTAATTAATCGG
#333	ACAAAATTTGTAGTATTGTTGAGGTTAC
#334	AAGGATATTAATAATCATAGAAAACG
#336	TCAATTAATAATATACAGAACAAGTATCC
#402	AAGAGGTAGAGTTCAATTCATCAAACC
#403	ATCTTTAATTTTATTTTGGTCATGC
#404	AAGGCTTTTTCCATTTTCAAGTGTTTCAGG
#406	ACATTGAAGATGGAAGCGTTCAACTAGC
#600 (CG6 integration rv)	ATTATGGGAAAATAATCCTTAC
#601 (Cam rv)	AGAAGCTCAGAGGCATGC
#602 (CG6 integration fw)	CTTTAATTTTATTTTGGTCATG
#603 (PbDT 3' For)	GGGAAGGTGTTGCTCAAATAGTG
#604 (CG6 WT fw)	GTTTCATGCTCCTCAACAAAG
#606 (CG6 WT rv)	GAACAAATACATAAGAGCGC
#607 (Armin 73)	GCTCAATTCTTTATGTCCACAAC
#608 (Armin 124)	CATGTTTTGTAATTTATGGGATAGCG
#609 (Amp ORI seq fw)	GCGAGGAAGCGGAAGAGC

790

791