



SUPPLEMENTAL FIGURE 1. Amplification curves. The linear regression for the amplification curves of (A) Pf18S DNA, (B) Pfs25, (C) SBP1, (D) RESA, and (E) Hyp8 to allow an assessment of polymerase chain reaction efficiency. Presented is the CT values (y axis) and the corresponding Log<sub>10</sub> transformed plasmid copy numbers per μL (x axis). Amplification efficiency was determined from the slope of the linear curve (indicated on the top right corners).

SUPPLEMENTAL TABLE 1  
 Selection of candidate marker

GeneID	Frequency	Intron	Beta-ring stages	Beta-mature gametocytes	Product description	Rings/MG
PF11735c	6	1	11.04258	2.152472	Ring-exported protein 1 (REX1)	5.130184
PF10_0016	5	0	5.538458	1.0868	Acyl-CoA binding protein, isoform 2, ACBP2	5.096117
PFD1170c	24	1	8.970933	2.077469	<i>Plasmodium</i> -exported protein (PHISTb), unknown function	4.318204
PFL2560c	2	1	8.053292	1.893178	<i>Plasmodium</i> -exported protein, unknown function	4.253848
MAL13P1.273	2	0	3.206335	0.784526		4.086971
PF08_0003	8	1	5.818946	1.461252	Tryptophan/threonine-rich antigen	3.982164
PFD0095c	0	1	6.810232	1.906783	<i>Plasmodium</i> -exported protein (PHISTb), unknown function	3.571582
MAL7P1.6	1	1	9.048624	2.575236	<i>Plasmodium</i> -exported protein (hyp12), unknown function	3.513707
MAL13P1.61	17	1	11.47698	3.29632	<i>Plasmodium</i> -exported protein (hyp8), unknown function	3.481757
MAL7P1.171	16	1	8.620577	2.562053	<i>Plasmodium</i> -exported protein, unknown function	3.364715
PFB0095c	18	1	9.215172	2.805933	Erythrocyte membrane protein 3	3.284175
PF10_0022	18	1	5.936012	1.838252	<i>Plasmodium</i> -exported protein (PHISTc), unknown function	3.229161
PFB0256w	12	0	3.34374	1.042435		3.207625
PFC0055w	11	1	2.23945	0.700442	<i>Plasmodium</i> -exported protein (hyp13), unknown function	3.197195
PF11_0503	13	1	9.200204	2.914172	<i>Plasmodium</i> -exported protein (PHISTc), unknown function	3.157056
MAL7P1.170	1	1	11.19367	3.556579	<i>Plasmodium</i> -exported protein, unknown function	3.147315

SUPPLEMENTAL TABLE 2  
Primer efficiency and limit of detection

Target	Slope	Efficiency (%)	LOD (copies/ $\mu$ L)	$R^2$	Cutoff CT values for positivity
Pf18S DNA	-3.3790	98	1	0.9984	36.0
Pfs25	-3.2845	102	10	0.9997	33.0
SBP1	-3.3295	100	10	0.9932	34.5
RESA	-3.4865	94	1	0.9924	ND
Hyp8	-3.3592	91	1	0.9995	ND
PHISTb	ND	ND	ND	ND	35.0
REX1	ND	ND	ND	ND	35.0

LOD = limit of detection; ND = not determined; RESA = ring-infected erythrocyte surface antigen; REX1 = ring-exported protein 1; SBP1 = skeleton-binding protein 1. Primer efficiency and limit of detection were assessed by cloning polymerase chain reaction products of the respective primers into p-TOPO-TA vectors. Constructs were used as quantitative controls/1standards as well.

SUPPLEMENTAL TABLE 3  
Primer sequence

Primer	GeneID	Direction	Sequence	Annealing temperature ( $^{\circ}$ C)	Reference
18S DNA	PF3D7_1148600	Forward	GTAATTGGAATGATAGGAATTTACAAGGT	60	21
		Reverse	TCAACTACGAACGTTTTAACTGCAAC		
18S RNA	PF3D7_1148600	Forward	TCCGATAACGAACGAGATCTTAAC	60	12
		Reverse	ATGTATAGTTACCTATGTTCAATTTCA		
Pfs25	PF3D7_1031000	Forward	GAAATCCCGTTTCATACGCTTG	60	12
		Reverse	AGTTTTAACAGGATTGCTTGATCTAA		
SBP1	PF3D7_0501300	Forward	GCAAAACAAGCCGTACATGTTG	60	14
		Reverse	TTGCTAGGTAATATCCTTTTCTTTTCC		
RESA	PF3D7_0102200	Forward	CAGCATATGGGTTTACTGGC	60	
		Reverse	TGGAAGTTCATCTTCTGGCGTACA		
REX1	PF3D7_0935900	Forward	GCTCCCTTTATTCTTGTG	58	
		Reverse	CATTATCATTTCCCTTCC		
PHISTb	PF3D7_0424600	Forward	GGATAGCAAGAATCTTAAGAAC	56	
		Reverse	AACATACGTTCAATAAGACAAC		
Hyp8	Pf3D7_1301700	Forward	CTTAATAATGGAACCTTTAAGAAC	60	
		Reverse	GTGTAGATTGTTCTGTTTCGAC		

RESA = ring-infected erythrocyte surface antigen; REX1 = ring-exported protein 1; SBP1 = skeleton-binding protein 1.