

Functional Analysis of Protein Kinase CK2 of the Human Malaria Parasite *Plasmodium falciparum*[†]

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Protein kinase CK2 (casein kinase 2) is a eukaryotic serine/threonine protein kinase with multiple substrates and roles in diverse cellular processes, including differentiation, proliferation, and translation. The mammalian holoenzyme consists of two catalytic alpha or alpha' subunits and two regulatory beta subunits. We report the identification and characterization of a *Plasmodium falciparum* CK2 α orthologue, PfCK2 α , and two PfCK2 β orthologues, PfCK2 β 1 and PfCK2 β 2. Recombinant PfCK2 α possesses protein kinase activity, exhibits similar substrate and cosubstrate preferences to those of CK2 α subunits from other organisms, and interacts with both of the PfCK2 β subunits in vitro. Gene disruption experiments show that the presence of PfCK2 α is crucial to asexual blood stage parasites and thereby validate the enzyme as a possible drug target. PfCK2 α is amenable to inhibitor screening, and we report differential susceptibility between the human and *P. falciparum* CK2 α enzymes to a small molecule inhibitor. Taken together, our data identify PfCK2 α as a potential target for antimalarial chemotherapeutic intervention.

Malaria, caused by infection with intracellular protozoan parasites of the genus *Plasmodium*, is responsible for 300 million to 600 million clinical cases annually (49), resulting in the deaths of up to 3 million people every year (9, 10). The need for novel intervention strategies is rendered more acute by the rise of resistance of the parasites (especially *Plasmodium falciparum*, the causative agent of the most virulent form of the disease) to most available drugs. There is a clear need for further research aimed at identifying novel drug targets (43).

The parasite life cycle is complex, with a succession of proliferation and differentiation events, in the regulation of which protein phosphorylation is likely to play crucial roles. Reversible phosphorylation of proteins is a major regulatory mechanism in most cellular processes, and approximately 30% of all eukaryotic cellular proteins carry a phosphate group. Deregulation of protein phosphorylation underlies various pathologies, including cancers, and protein kinases are considered promising drug targets, comprising as much as 30% of all protein targets under investigation (13). The divergences between human and plasmodial protein kinases suggest that specific inhibition of the latter is an achievable goal (14, 15).

Protein kinase CK2, formerly known as casein kinase II, is a serine/threonine protein kinase ubiquitously expressed in eukaryotes. It has over 300 cellular substrates catalogued to date (34). Consistent with its multiple substrates, the enzyme plays a crucial role in many cellular processes, including differentiation, proliferation, survival, translation, apoptosis, cell cycle

progression, and cellular responses to stress and DNA damage (1, 38); mammalian CK2 has even been found to act as an exokinase, phosphorylating several extracellular proteins (44, 55). CK2 is essential to viability in yeasts and slime molds (24, 37). The human CK2 holoenzyme consists of two catalytic alpha or alpha' subunits and two regulatory beta subunits, and recent evidence indicates that the beta subunits interact with several protein kinases in addition to CK2 α (reviewed in reference 6), pointing to a likely role in the integration of numerous signaling pathways. The human genome encodes three CK2 α subunits and a single version of the beta subunit. Increasing evidence shows that CK2 is an attractive target for antineoplastic and antiviral drugs (46).

The genome of the *P. falciparum* strain 3D7 (18) has been fully sequenced, allowing the discovery of the entire complement of plasmodial protein kinases by analysis of the set of predicted peptides (3, 57). A putative CK2 α orthologue and two predicted CK2 β subunits were identified in these analyses. Here we present the biochemical characterization of the PfCK2 α and both PfCK2 β orthologues and demonstrate by using a reverse genetics approach that the catalytic subunit is essential for completion of the erythrocytic asexual cycle of the parasite.

MATERIALS AND METHODS

Expression and purification of the three PfCK2 subunits. Oligonucleotides were designed to amplify the PfCK2 α open reading frame (ORF) from *P. falciparum* (clone 3D7A) cDNA. The forward (5'-GGGGGGATCCATGTCGGTTAGCTCAATTAATAAAA-3') and reverse (5'-GGGGGTCGACTTATGATTCTCACGGTCTTCTC-3') primers carried BamHI and Sall sites, respectively (underlined). Oligonucleotides were also designed to amplify the PfCK2 β 1 and PfCK2 β 2 ORFs from *P. falciparum* (clone 3D7A) cDNA. The PfCK2 β 1 forward (5'-GGGGAGATCTATGGAAAATAGTGATTCGAATAAAGAC-3') and reverse (5'-GGGGGTCGACTTACGTTTCAGAAATTTGTAGTTCTTCC-3') primers carried BglII and Sall sites, respectively (underlined). The PfCK2 β 2

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sequence has a long N-terminal extension. Oligonucleotides were designed to amplify the PfCK2 β sequence, lacking the N-terminal extension, from *P. falciparum* (clone 3D7A) cDNA. The forward (5'-GGGGGGATCCATGGAAGC AACAGTGTCTTGATTG-3') and reverse (5'-GGGGGTCGACTCATTGA CACTTTCAGAGGATTCCG-3') primers carried BamHI and Sall sites, respectively (underlined). The short version of PfCK2 β , lacking the N-terminal extension, was named shPfCK2 β . Catalytically inactive ("kinase dead") PfCK2 α was obtained by site-directed mutagenesis (K72M) of pGEX-PfCK2 α by overlap extension PCR (23). All cloning primers are provided in Table S1 in the supplemental material. All PCR products were verified by sequencing in the vector pGEM-T Easy (Promega) and then subcloned into the plasmid pGEX-4T3 (GE Healthcare) to generate N-terminal glutathione S-transferase (GST) fusions. The shPfCK2 β sequence was also inserted between the BamHI and NotI sites of the plasmid pQE-30 for expression with an N-terminal His₆ tag. A pET29 vector containing *P. falciparum* CK2 α in frame with a C-terminal His₆ tag sequence was a kind gift from D. Chakrabarti. The pGEX4T3 constructs were expressed in *Escherichia coli* BL21 Gold cells, SG13009 cells were used for expression from the pQE30 plasmid, and the pET29 construct was expressed in *E. coli* BL21(DE3) cells for 20 h at 20°C with 0.1 mM isopropyl- β -D-thiogalactopyranoside (IPTG). GST-tagged proteins were purified on glutathione-agarose beads (Sigma), and His-tagged proteins were purified on Ni²⁺-agarose beads (Qiagen), following the manufacturer's recommendations.

Construction of the knockout (KO) plasmid pCAM-BSD-KOPfCK2 α . A fragment from the PfCK2 α PCR was amplified by PCR and inserted between the BamHI and NotI sites of the pCAM-BSD plasmid (47), which contains the *Aspergillus terreus* blasticidin-S-deaminase gene, whose gene product confers resistance to the drug blasticidin. Cloning primers are listed in Table S1 in the supplemental material.

Construction of the complementation plasmid pCHD-PfCK2 α . A plasmid for in vivo episomal expression of PfCK2 α subunits was constructed as follows: the full-length PfCK2 α coding sequence was first inserted between the BglII and NotI sites of the plasmid pHGB (51) and then transferred into the plasmid pCHD-1/2 (51) by a Gateway LR clonase reaction according to the manufacturer's instructions (Invitrogen). The plasmid pCHD-1/2 includes a cassette encoding human dihydrofolate reductase, conferring resistance to the antifolate drug WR99210. Parasites that were transfected with both a KO plasmid and a complementation plasmid were cultured under double-drug selection (see below).

3'-Tagging plasmid. The 3' end of the PfCK2 α coding sequence (538 bp, omitting the stop codon) was amplified by PCR using primers incorporating PstI and BamHI restriction sites and inserted between the PstI and BamHI sites of the pCAM-BSD-hemagglutinin (HA) plasmid (42).

Kinase assays. Standard kinase reactions (30 μ l) occurred in kinase buffer (20 mM Tris-HCl [pH 7.5], 20 mM MgCl₂, 2 mM MnCl₂, and 10 μ M ATP) containing 0.075 MBq [γ -³²P]ATP (370 MBq/ml; GE Healthcare), 1 μ g of recombinant kinase, and 5 μ g of substrate. Reactions were carried out at 30°C for 30 min and terminated by the addition of Laemmli buffer. Samples were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). The gels were dried and exposed for autoradiography.

Kinase assays were also analyzed by the phosphocellulose method (20). A final assay volume of 18 μ l contained 36 ng of PfCK2 α , 167 mM of peptide substrate, 50 mM Tris-HCl (pH 7.5), 20 mM MgCl₂, 25 μ M ATP, 40 to 150 mM NaCl, and 0.037 MBq [γ -³²P]ATP (220 TBq/mmol; GE Healthcare). Reactions took place for 10 min at room temperature and were terminated by the addition of 60 μ l of 4% trichloroacetic acid. Reaction mixtures were centrifuged at 10,000 \times g for 15 min, and then 60 μ l of supernatant was spotted onto 4- by 4-cm Whatman P81 phosphocellulose paper squares. The squares were washed three times for 15 min in 0.5% orthophosphate, and then the amount of radiolabel incorporated into the peptide substrate was quantified by scintillation counting. Three peptide substrates were used in this study, the NEB peptide p6012 (RRRADDSDDD DD), the custom peptide (RRREDEESDDEE), obtained from NeoMPS, and the eIF2 β -derived peptide (MSGDEMIFDPTMSKKKKKKK) (40, 45), obtained from GeneCust (Evry, France).

The K_m s of PfCK2 α for ATP and GTP were determined by performing kinase assays with ATP concentrations of 100 μ M, 25 μ M, 6.25 μ M, and 1.5625 μ M. Reactions were carried out in triplicate. The [γ -³²P]ATP/GTP was added to the unlabeled ATP/GTP and diluted serially to ensure a constant ratio of labeled-to-unlabeled ATP/GTP. The NEB peptide RRRADDSDDD (100 μ M) was used as the substrate. The K_m s of PfCK2 α for ATP in the presence or absence of PfCK2 β were determined by performing kinase assays in triplicate with ATP concentrations of 100 μ M, 50 μ M, or 25 μ M, with the NeoMPS peptide (100 μ M) as a substrate. Reaction mixtures contained 36 ng of PfCK2 α alone or in combination with 36 ng of PfCK2 β /shPfCK2 β .

Measurement of 50% inhibitory concentrations (IC₅₀s). To test the effect of small molecule inhibitors on PfCK2 α , kinase activity was measured in the presence of increasing concentrations of these molecules. Stocks of the molecules contained dimethyl sulfoxide or ethanol as a solvent, and negative controls for the reactions were provided by reaction mixtures containing ethanol or dimethyl sulfoxide without the small molecule inhibitor. Kinase reactions were performed by the phosphocellulose method as detailed above.

Interaction assay. A mixture of 5 μ g of each recombinant protein was incubated at 4°C for 30 min in 20 mM Tris-HCl (pH 7.5), 0.2 M NaCl, 0.1% Nonidet P40 (IGEPAL), and 10% glycerol. Glutathione-agarose beads were added to each reaction mixture. The tubes were rotated at 4°C for 1 h; the beads were recovered by centrifugation and washed four times in reaction buffer. Laemmli buffer was added to the beads, which were then heated to 100°C. Samples were separated by SDS-PAGE on 12% acrylamide gels and either stained or transferred to the membrane for Western blot analysis.

Western blot analysis. Western blotting was performed according to conventional protocols. Briefly, samples were separated by SDS-PAGE on a 12% acrylamide gel and blotted onto nitrocellulose. The membranes were blocked according to standard methods and incubated with rabbit anti-His antibody (1:1000; Santa Cruz Biotechnology) or subunit-specific antibodies generated in rabbits by BioGenes (Germany) against the PfCK2 α -derived peptide ADVNIH KPKEYDY. A goat anti-rabbit secondary antibody coupled to horseradish peroxidase was used at a ratio of 1:10,000. Antigen was visualized using the enhanced chemiluminescence system (PerkinElmer).

Parasite culture and transfection. Cultures of the *P. falciparum* strain 3D7A (54) were maintained at 37°C in RPMI 1640 medium (Gibco) supplemented with 25 mM sodium bicarbonate, 2 mM L-glutamine, 300 mM hypoxanthine, 10 μ g/ml gentamicin, and AlbuMAX II (Sigma). Cultures were seeded at 5% hematocrit and maintained at a parasitemia of 1 to 10% with daily changes of medium. The incubator was flushed with a gas mixture containing 5% CO₂.

For transfection, asexual blood stage parasites were synchronized by sorbitol treatment (26) to obtain a majority of ring stage parasites. Forty-eight hours later, ring stage parasites were transfected by electroporation with 100 μ g of purified plasmid DNA in Cytomix buffer as described previously (16, 17, 47). Blasticidin (2.5 μ g/ml) was added to the culture medium to select for transfected parasites. Parasites under double selection had WR99210 (5 nM) added to the culture medium in addition to the blasticidin. Parasites were maintained in this supplemented medium from 2 days posttransfection.

Preparation of parasite protein extract. Parasite cultures were lysed in 0.15% saponin. After centrifugation and washing, the parasite pellets were sonicated in RIPA buffer (30 mM Tris, pH 8.0, 150 mM NaCl, 20 mM MgCl₂, 1 mM EDTA, 0.5% Triton X-100, 1% Nonidet P40, 10 mM β -glycerophosphate, 10 mM NaF, 1 mM phenylmethylsulfonyl fluoride, 10 mM benzamidine hydrochlorate hydrate, and Roche complete cocktail protease inhibitors). The lysates were cleared by centrifugation (15,000 rpm for 15 min at 4°C), and the total amount of proteins in the supernatant was measured using the Bradford assay (8).

DNA extraction and Southern blotting. Parasite cultures were lysed in 0.15% saponin. The parasite pellets were resuspended in cold phosphate-buffered saline and treated with proteinase K (150 μ g/ml) and 2% SDS at 55°C for 2 to 3 h. The genomic DNA was extracted in phenol/chloroform/isoamyl alcohol (25:24:1) and precipitated in ethanol and 0.3 M sodium acetate. The DNA was digested with HindIII.

Primers used in genotype characterization. For the detection of integration of the plasmid pCAM-BSD-KOPfCK2 α into genomic DNA, the following primers were used (see Fig. 6A for numbering and see Table S1 in the supplemental material for the sequences): primer 1, CK2a5primeF; primer 2a, CK2aR; primer 2b, CK2a3primeR; primer 3, pCAM-BSDF; and primer 4, pCAM-BSDR.

For the detection of integration of the plasmid pCAM-BSD-HA, the following primers were used (see Fig. 7D for numbering and see Table S1 in the supplemental material for the sequences): primer 1, CK2aF; primer 2, CK2a3primeR; primer 3, pCAM-BSDF; and primer 4, pCAM-BSDR.

RESULTS

Bioinformatics. Phylogenetic analysis of *P. falciparum* protein kinases identified the PlasmoDB (<http://plasmodb.org/plasmo>) sequence PF11_0096 as that of a CK2 α orthologue (3, 57), with 65% amino acid sequence identity to *Homo sapiens* CK2 α . PF11_0096 was therefore named PfCK2 α . An alignment of PfCK2 α with CK2 α subunits from *H. sapiens* and *Zea mays* (see Fig. S1 in the supplemental material) reveals that

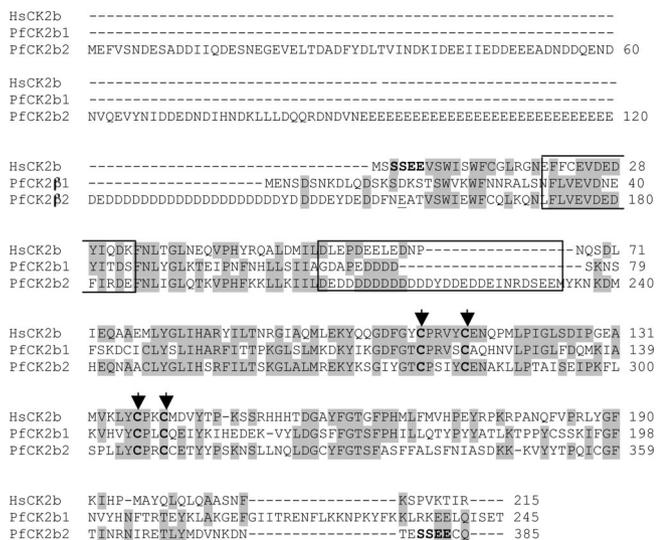


FIG. 1. Alignment of PfCK2β1 and PfCK2β2 with *Homo sapiens* CK2β. ClustalW alignments of the following proteins were performed: *Homo sapiens* CK2β (HsCK2β; AAM50092); PfCK2β1 (AAN35637; PlasmDB PF11_0048; 33% identical to HsCK2β), PfCK2β2 (CAD52554; PlasmDB PF13_0232; 39% identical to HsCK2β). Note the long N-terminal extension and acidic insertion sequence in PfCK2β2. The sequence of shPfCK2β2 begins (after an artificially introduced initiating methionine) with residue E156, underlined. The cysteine residues thought to hold the zinc finger in place are indicated in bold text and with arrowheads. The potential autophosphorylation site on PfCK2β2 (SSEE) is indicated in bold. The export of CK2 as an ectokinase is mediated by CK2β residues 20 to 33 (44) (boxed, along with the equivalent residues in PfCK2β1 and PfCK2β2). The acidic region mentioned in the text is also boxed (D107 to E133 of PfCK2β2).

PfCK2α possesses all 11 of the subdomains conserved across eukaryotic protein kinases (21, 22) and the majority of the conserved features of CK2α subunits (2). Just downstream from subdomain II is a putative nuclear localization signal, Pro-Val-Lys-Lys-Lys-Lys-Ile, conserved across CK2α homologues. PfCK2α also possesses three invariant residues common to CK2 family members; the ATP binding motif present in most other protein kinases is Gly-X-Gly-X-X-Gly, whereas in the CK2 family the motif is Gly-X-Gly-X-X-Ser (PfCK2α, Gly50-Ser55). The most highly conserved amino acid motif specific to members of the CK2 family is Asp179-Trp-Gly181 (notation from PfCK2α; most protein kinases display Asp-Phe-Gly at this position). Likewise, Gly203-Pro-Glu205 (notation from PfCK2α) is a common feature of the family, which diverges from the Ala-Pro-Glu motif present in the vast majority of other protein kinases; thus, all three CK2-specific motifs are present in PfCK2α.

Two putative CK2β subunits were identified in *P. falciparum* (57), hereafter referred to as PfCK2β1 and PfCK2β2 (PlasmDB identifiers PF11_0048 and PF13_0232, respectively). BLASTP searches using the putative PfCK2β1/PfCK2β2 amino acid sequences as queries confirmed their identities as CK2β orthologues. An alignment with the human CK2β sequence (HsCK2β) (Fig. 1) reveals that many of the conserved features of CK2β subunits, including the four cysteine residues responsible for zinc finger formation (12), are present in PfCK2β1 and PfCK2β2 (e.g., Cys117, -122, -145, and -148 for PfCK2β1) (Fig. 1). The human CK2β sequence has a well-

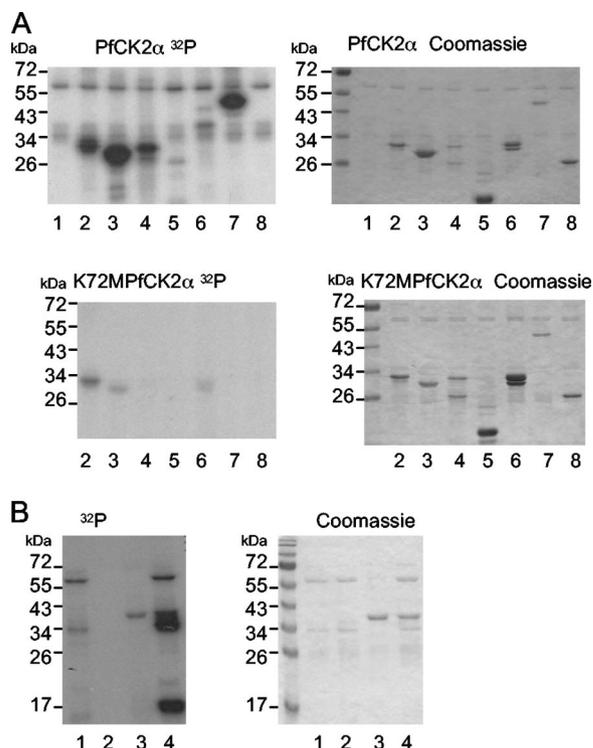


FIG. 2. GST-PfCK2α kinase activity. (A) Autoradiograms (left) and Coomassie blue-stained gels (right) of kinase assays performed with GST-PfCK2α (top) or catalytically inactive GST-K72MPfCK2α (bottom) and the following substrates: lane 1, no substrate; lane 2, α-casein; lane 3, β-casein; lane 4, mixed dephosphorylated caseins; lane 5, myelin basic protein; lane 6, histone H1; lane 7, GST-shPfCK2β2; and lane 8, GST. (B) Autophosphorylation can occur by a transreaction. Left, autoradiogram; right, Coomassie blue-stained gel. The band observed at roughly 66 kDa is the GST-PfCK2α autophosphorylation band, and the band at roughly 40 kDa is the PfCK2α-His autophosphorylation band. Lane 1, GST-PfCK2α; lane 2, GST-K72MPfCK2α; lane 3, PfCK2α-His; and lane 4, GST-K72MPfCK2α plus PfCK2α-His. Radioactive phosphate is incorporated into catalytically inactive PfCK2α (K72MPfCK2α) in the presence of PfCK2α-His, indicating that autophosphorylation can occur by an intermolecular reaction.

documented CK2 phosphorylation site at the N terminus (SSEE). PfCK2β2 possesses several phosphorylatable residues in the N-terminal region that are surrounded by a number of acidic residues, which could therefore be phosphorylated by CK2, and a TESSEE sequence at the C terminus reminiscent of the HsCK2β N-terminal phosphorylation site (MSSEE). The stretch of amino acids found to be necessary for the export of CK2 as an ectokinase (CK2β amino acids E20 to K33) (44) are largely conserved in the PfCK2β sequences, leading to the intriguing possibility that PfCK2 may be exported from the parasite. The acidic stretch responsible for downregulation of CK2 activity and association with the plasma membrane (HsCK2β amino acids D55 to D64) (29, 32) is present in PfCK2β1 (D68 to D75) and extended in PfCK2β2 (D207 to E226). This insertion occurs in a region looping out from the main protein structure (12) and is not unique; for example, *Saccharomyces cerevisiae* CK2β has an insertion sequence of 30 amino acids in this location. Along with the insertion region, PfCK2β2 has a highly acidic and repetitive N-terminal exten-

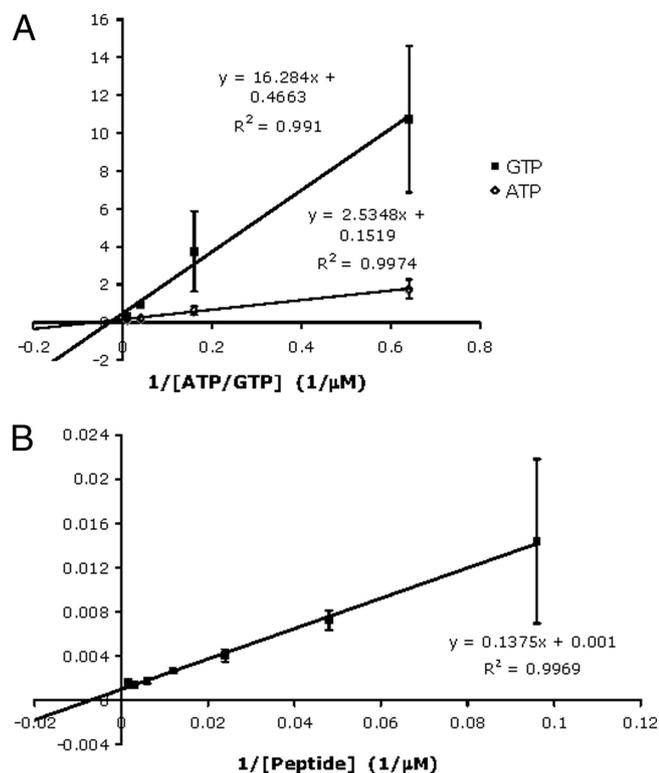


FIG. 3. PfCK2 α kinetics. The enzyme kinetics of PfCK2 α in a Lineweaver-Burke presentation. The experiments were performed in triplicate, the data points represent the means, and the error bars represent three standard deviations. (A) The graph was obtained by linear regression of the enzyme kinetic data for ATP and GTP. The intercepts on the x axis give the negative reciprocal of the K_m , and the intercepts on the y axis give the reciprocal of the V_{max} . (B) The graph was obtained by linear regression of the enzyme kinetic data for the NEB peptide RRRADDSDDDDDD. The intercept on the x axis gives the negative reciprocal of the K_m .

sion that is not found in other CK2 β subunits. The N-terminal region is at the periphery of the 3D structure of the human CK2 β peptide (12, 35) and is not a conserved part of the CK2 β structure and thus may possibly function as a docking site or region that interacts with binding partners or substrates of the PfCK2 holoenzyme. The human CK2 β is phosphorylated at S209 in a cell cycle-dependent manner by p34^{cdc2} (19, 30, 33), although the function of this phosphorylation is unknown. Both PfCK2 β subunits possess serine residues near the C terminus that could be phosphorylated.

Microarray data available on PlasmoDB reveal that the mRNAs encoding all three subunits are detectable throughout the parasite life cycle (7, 28). Expression of the PfCK2 α protein in asexual blood stage parasites was verified by Western blot analysis (see Fig. 7F).

In vitro activity of PfCK2 α . Kinase activity of bacterially expressed recombinant GST-PfCK2 α was detected by an in vitro kinase assay (Fig. 2, top). GST-PfCK2 α autophosphorylates (the 66-kDa band present in each lane in Fig. 2 corresponds to the size of the GST-tagged PfCK2 α subunit) and is capable of phosphorylating a range of exogenous substrates, with strongest activity toward the caseins (Fig. 2, lanes 2 to 4) and recombinant GST-tagged shPfCK2 β 2, a short version of

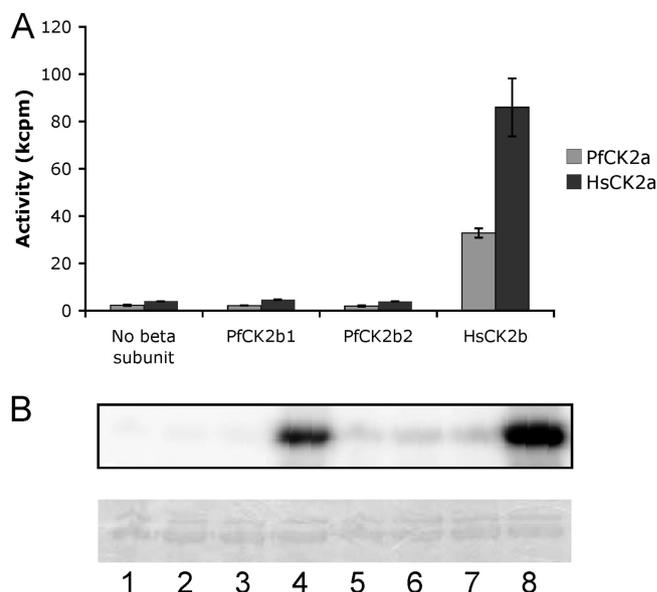


FIG. 4. PfCK2 α and HsCK2 β interact in vitro. To further test the interactions of the alpha and beta subunits, two substrates that are phosphorylated by the CK2 holoenzyme and not by the CK2 α subunit alone were included in kinase assays with mixtures of human and *P. falciparum* alpha and beta subunits. (A) Phosphorylation of the eIF2 β [1-22] peptide (40, 45) by PfCK2 α -His or HsCK2 α in the presence and absence of GST-PfCK2 β 1, GST-shPfCK2 β 2, or HsCK2 β was measured by kinase assays, and the amount of radiolabel incorporated into the peptide was counted by scintillation. Results are shown as the means of two experiments, with the error bars representing the standard deviations. (B) Phosphorylation of the GST-Olig2[1-177] protein (27) by PfCK2 α -His (lanes 1 to 4) or HsCK2 α (lanes 5 to 8) alone (lanes 1 and 5) or in the presence of GST-PfCK2 β 1 (lanes 2 and 6), GST-shPfCK2 β 2 (lanes 3 and 7), or HsCK2 β (lanes 4 and 8). Top, autoradiogram; bottom, corresponding Coomassie blue-stained gel of the kinase assay.

the PfCK2 β 2 subunit lacking the N-terminal extension (see Materials and Methods) (Fig. 2, lane 7). There was no activity against the GST moiety alone (Fig. 2, lane 8), indicating that the activity in lane 7 was against the beta subunit itself and that the autophosphorylation is against the PfCK2 α subunit itself. We did not detect any activity against the PfCK2 β 1 subunit (data not shown). These observations are consistent with general preferences of CK2 homologues for substrates with highly acidic phosphoacceptor sites (11, 25, 39, 52, 56); several potential such sites are present on PfCK2 β 2. Parallel assays performed with an inactive PfCK2 α mutant (Lys72Met; the Lys residue is required for correct orientation of the ATP molecule) were negative for kinase activity, confirming that activity is indeed due to PfCK2 α (Fig. 2, bottom). PfCK2 α autophosphorylates by a transreaction (Fig. 2B); GST-PfCK2 α and PfCK2 α -His autophosphorylate (Fig. 2, lanes 1 and 3), while GST-K72MPfCK2 α does not (Fig. 2, lane 2) but is phosphorylated in the presence of PfCK2 α -His, indicating that at least a proportion of the autophosphorylation of PfCK2 α occurs by an intermolecular reaction.

PfCK2 α shares features in common with CK2 α from other systems. A feature often cited as being characteristic of CK2 enzymes is that they have similar affinities for GTP and ATP. PfCK2 α has a K_m of 16.7 μ M and V_{max} of 6.6 nmol/min for

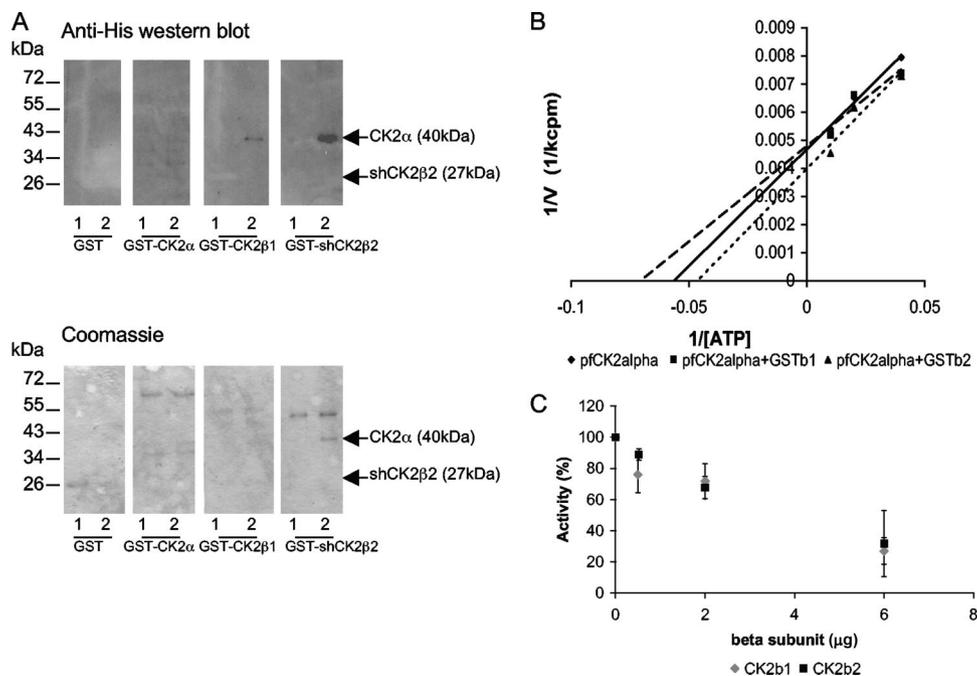


FIG. 5. PfCK2 α and PfCK2 β subunits interact in vitro. (A) Pull-down assays. Each of the GST-tagged subunits (indicated at the bottom of each panel) was incubated with PfCK2 α -His (lane 1) or His-shPfCK2 β 2 (lane 2). Complexes containing the GST-tagged subunits were then purified using glutathione agarose beads, and any bound His-tagged subunits were detected by Western blot analysis. Top, anti-His Western blots. Bottom, Coomassie blue-stained gels. (B) The PfCK2 α K_m for ATP is not greatly altered in the presence of the PfCK2 β subunits. ATP concentrations of 100 μ M, 50 μ M, and 25 μ M were used. Reaction mixtures contained PfCK2 α alone or with equimolar amounts of PfCK2 β 1 or shPfCK2 β 2. (C) Phosphorylation of β -casein by PfCK2 α His in the presence of increasing concentrations of GST-tagged PfCK2 α subunits (and concomitant decreases in the concentration of GST) was quantified by phosphorimaging. Standard kinase assays were performed with 1 μ g of β -casein, 1 μ g of PfCK2 α His, and 6 μ g of GST/GST-PfCK2 β in various proportions. The graph represents the combined results of three experiments (means and standard deviations shown).

ATP and a K_m of 34.9 μ M and V_{max} of 2.1 nmol/min for GTP (Fig. 3A). The enzyme displays a number of other features that confirm it as a true member of the CK2 α family: (i) PfCK2 α is able to phosphorylate the CK2 substrate peptide RRRADDS DDDDD (NEB), with a K_m of 137.5 μ M (Fig. 3B); (ii) CK2 α enzymes are known to have a wide variety of substrates (34), and correspondingly, PfCK2 α (but not the K72M mutant protein used as a negative control) phosphorylates a number of proteins within heat-inactivated parasite protein extract (data not shown); (iii) the activity of PfCK2 α is inhibited by the well-established CK2-specific inhibitor TBB (3,4,5,6-tetrabromobenzotriazole), with a similar IC_{50} curve to that of human CK2 α (IC_{50} for PfCK2 α , 2 μ M; IC_{50} for HsCK2 α , 1.5 μ M) (see Fig. 8C); and (iv) it can be recruited by the human CK2 β subunit to phosphorylate the eIF2 β -derived peptide (40, 45) and the Olig2 protein (27) (Fig. 4). This is in line with the established ability of human CK2, but not CK2 α alone, to phosphorylate the substrates used in this experiment (27, 40).

PfCK2 α and the PfCK2 beta subunits interact in vitro. To assess whether the two regulatory PfCK2 subunits are able to associate with PfCK2 α in vitro, all three subunits were expressed in *E. coli* as His- or GST-tagged proteins and used in pull-down experiments. Mixtures of His- and GST-tagged proteins were prepared, from which proteins were pulled down using glutathione beads. The pulled-down proteins were then subjected to Western blot analysis using an anti-His antibody to detect any bound His-tagged protein that was copurified

with the GST-tagged proteins. PfCK2 α -His was copurified with both GST-tagged beta subunits but not with GST alone (Fig. 5A). The interaction does not significantly alter the K_m for ATP (Fig. 5B) or the phosphorylation of calmodulin or the peptide RRRSDDSDDDDD (data not shown). However, the interaction of the beta subunits with the kinase has functional significance, at least in vitro, since the activity of the kinase toward β -casein is reduced with increasing amounts of GST-PfCK2 β present in the reaction mixtures (Fig. 5C).

PfCK2 α is essential for completion of the erythrocytic asexual cycle. We next wanted to determine whether PfCK2 α plays essential functions in parasite survival. To generate a plasmid able to disrupt the PfCK2 α gene, an internal fragment of the coding sequence, excluding the critical motifs Gly-X-Gly-X-X-Ser (subdomain I, involved in anchoring of the ATP molecule) and Gly-Pro-Glu (subdomain VIII, required for structural stability of the C-terminal lobe) (see Fig. S1 in the supplemental material), was amplified and cloned into the transfection vector pCAM-BSD (47), which confers resistance to blasticidin. Integration of this construct (pCAM-BSD-KOPfCK2a) into the genomic locus by single-crossover homologous recombination is expected to result in a pseudodiploid configuration, where both truncated copies will be unable to express a functional enzyme, since both will lack one of the essential motifs (Fig. 6A).

After two independent transfections of pCAM-BSD-KOPfCK2a into 3D7 parasites, integration was monitored in the blasticidin-resistant populations by PCR (Fig. 6B), using

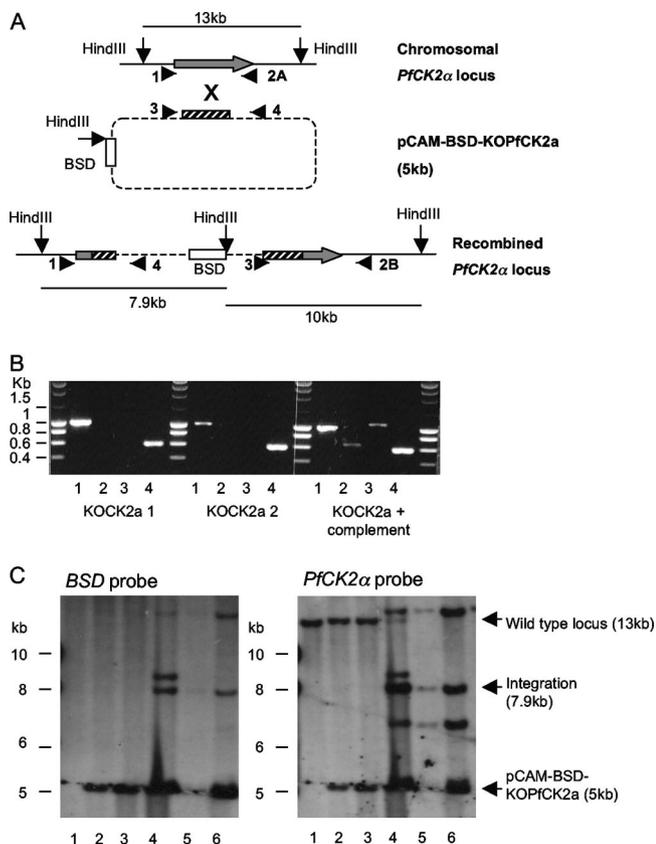


FIG. 6. PfCK2 α is necessary for erythrocytic-stage parasite viability. 3D7 parasites transfected with pCAM-BSD-KOPfCK2 α with or without pCHD-PfCK2 α were analyzed by PCR and Southern blotting. (A) Disruption strategy for PfCK2 α . The locations of the primers used for PCR screening and the HindIII sites used in Southern blotting are shown. (B) PCR screening of genomic DNA (gDNA) from untransfected 3D7 parasites, two separate pCAM-BSD-KOPfCK2 α -transfected lines (KOCK2a 1 and KOCK2a 2), and parasites transfected with both the KO plasmid and the complementation plasmid (KOCK2a + complement). Lane 1, amplification of the wild-type PfCK2 α locus (primers 1 and 2A; expected size, 1,030 bp). Lane 2, amplification over the 5' integration boundary (primers 1 and 4; expected size, 677 bp). Lane 3, amplification over the 3' integration boundary (primers 2B and 3; expected size, 1,060 bp). Lane 4, amplification of the insert in the pCAM-BSD-KOPfCK2 α plasmid (primers 3 and 4; expected size, 610 bp). Evidence of integration is seen only in gDNA from the doubly transfected parasite culture (KOCK2a + complement). (C) This parasite line was cloned by limiting dilution, and the gDNA was analyzed by Southern blotting. The restriction enzyme HindIII was used to digest the gDNA, and the fragments were analyzed by Southern blotting using BSD and PfCK2 α as probes. Lanes: 1, untransfected 3D7; 2, KOCK2a1; 3, KOCK2a2; 4, KOCK2a plus complement; 5, KOCK2a plus complement clone E7; and 6, KOCK2a plus complement clone G9.

primer combinations that allow discrimination between the episome, the wild-type locus, and the disrupted locus. Only the episome and the wild-type locus were detectable, with no sign of integration even after prolonged culturing (16 weeks); in contrast, we regularly observe disruption of nonessential genes 6 to 7 weeks posttransfection (for an example, see reference 42). This might be due either to the fact that the presence of an intact PfCK2 α gene is crucial for parasite asexual multiplication or to the possible nonrecombinogenicity of the locus. To

verify that the PfCK2 α locus is indeed recombinogenic, we proceeded to transfect wild-type parasites with a "3'-tagging" construct whose integration was expected not to cause loss of function of the target protein. We readily observed integration of the tagging construct (Fig. 7A to E) and size increase in the PfCK2 α protein caused by the HA tag (1.1 kDa) (Fig. 7F). This demonstrates that the locus is accessible to recombination if no loss of function is incurred, as is presumably the case with HA tagging, and therefore strengthens the case that PfCK2 α is essential for the parasite's asexual cycle.

We nevertheless wanted to ascertain that PfCK2 α can be disrupted if the enzyme is provided through expression of an extraneous copy of the gene. To this effect, a complementation plasmid was constructed, containing the full-length PfCK2 α coding region under the control of the *Pf*hsp86 promoter and preceding a 3' untranslated region (namely, the *Plasmodium berghei* dihydrofolate reductase terminator sequence). The *P. falciparum* hsp86 gene (PF07_0029) displays a similar mRNA expression profile to the PfCK2 α gene (28); therefore, its promoter is presumably appropriate to drive expression of the complementing protein. In parallel with the transfection of the pCAM-BSD-KOPfCK2 α plasmid alone, further populations of parasites were cotransfected with both pCAM-BSD-KOPfCK2 α and the complementation plasmid. PCR analysis (Fig. 6B, right) showed that disruption of the targeted locus occurred only in the doubly transfected, doubly resistant parasites. Southern blot analysis independently confirmed that integration occurred only in the doubly transfected parasites (Fig. 6C). The 13-kb band that represents the wild-type locus dramatically decreased in the doubly transfected parasites and was undetectable in two clonal lines (E7 and G9) that were derived from this culture by limiting dilution. There are multiple possibilities for the recombination of the KO and complementation plasmids with each other before or after integration, which could account for the additional bands of unexpected size observed (6 kb and 14 kb). The most important observation is that the wild-type band disappears only in the doubly transfected parasites.

Taken together, these data provide strong evidence that PfCK2 α is essential to viability of the asexual erythrocytic stage parasites.

PfCK2 α kinase activity is amenable to inhibition. Using a kinase-directed inhibitor library, we conducted a screen for compounds that inhibit PfCK2 α . This screen identified the compounds Rottlerin and ML-7 as inhibitors of PfCK2 α . The IC₅₀s of these compounds were determined for both PfCK2 α and HsCK2 α (Fig. 8). While ML-7 inhibits both enzymes with an IC₅₀ of roughly 3 to 4 μ M, Rottlerin exhibits differential effects on the orthologues, inhibiting PfCK2 α with an IC₅₀ of 7 μ M and HsCK2 α with an IC₅₀ of \gg 20 μ M. This result indicates that differential inhibition is possible, despite the high percent identity (65%) between the CK2 α amino acid sequences of *P. falciparum* and *Homo sapiens*.

DISCUSSION

We have characterized a *P. falciparum* CK2 α orthologue and confirmed that the recombinant enzyme exhibits kinase activity in vitro and exhibits features in common with other CK2 α enzymes. PfCK2 α contains the major motifs conserved across

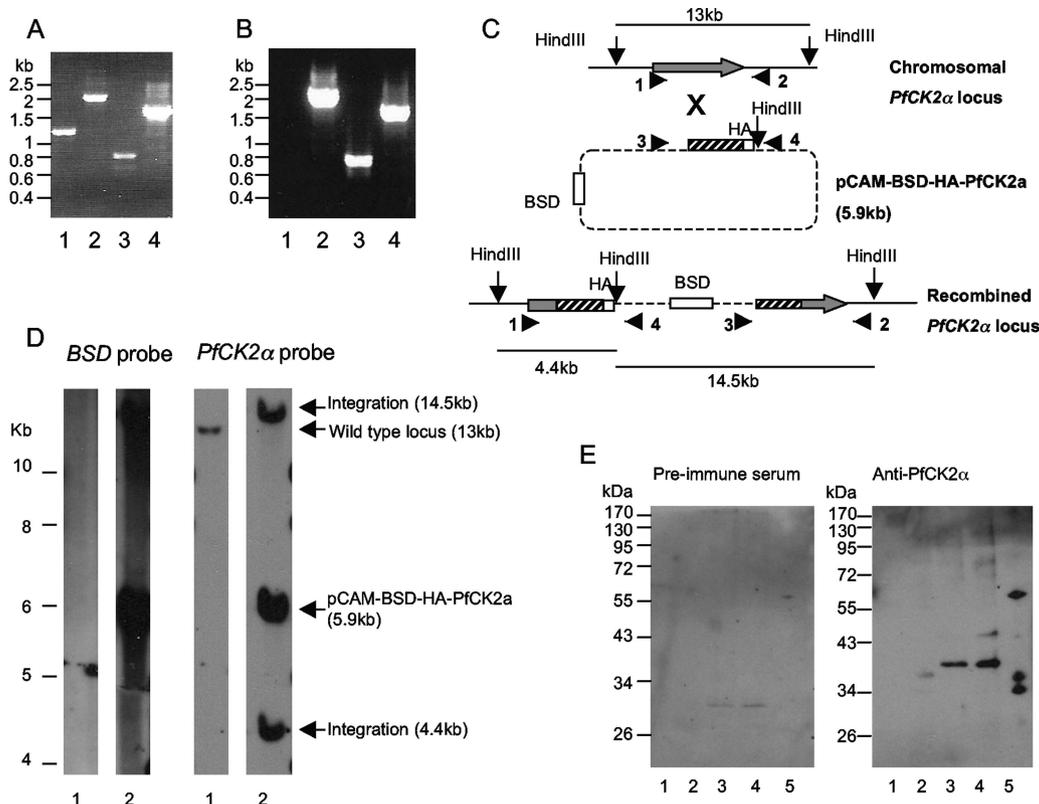


FIG. 7. PfCK2 α can be targeted for recombination. (A) PCR screening for integration in genomic DNA (gDNA) from parasites transfected with pCAM-BSD-HA-PfCK2a revealed the presence of parasites in which integration events had occurred. Lane 1, amplification of the wild-type locus (primers 1 and 2; expected size, 1,176 bp). Lane 2, amplification over the 5' integration boundary (primers 1 and 4; expected size, 1,973 bp). Lane 3, amplification over the 3' integration boundary (primers 2 and 3; expected size, 810 bp). Lane 4, amplification of the insert in the pCAM-BSD-KOPfCK2a plasmid (primers 3 and 4; expected size, 610 bp). This culture still contained the wild-type locus, and therefore clonal lines were derived from the culture by limiting dilution. (B) PCR screening for integration in gDNA from one of the parasite clones revealed that the wild-type band had been lost (lane 1, 1,176 bp) and only the integration bands were seen. (D) This clonal line was further analyzed by Southern blotting. (C) A schematic of the chromosomal gene locus, the pCAM-BSD-HA-PfCK2a plasmid, and the recombined locus, showing the locations of oligonucleotide primers used for the PCR screens (A and B). Oligonucleotide identities are listed in Table S1 in the supplemental material. HindIII sites and the expected sizes of the fragments of gDNA after restriction digestion are shown. (D) Southern blotting. Lanes: 1, untransfected 3D7 parasites; 2, PfCK2 α HA clone B3; and 3, PfCK2 α HA clone E1. (E) Western blot analysis showing PfCK2 α expression in erythrocytic stage parasites. Protein extract from unsynchronized erythrocytic-stage *P. falciparum* parasites was prepared from wild-type 3D7 parasites (lane 2) and from parasites with a sequence encoding a HA tag incorporated at the 3' end of the PfCK2 α gene locus (lane 3, clone B3; lane 4, clone E1). Protein extract from unparasitized red blood cells (lane 1) and recombinant GST-PfCK2 α (lane 5) were included as negative and positive controls. Two identical Western blot analyses were performed with immunopurified rabbit anti-PfCK2 α antibodies (right) or with preimmune serum from the same rabbit as a negative control (left). The expected sizes of the proteins are indicated with arrows.

CK2 catalytic subunits, phosphorylates acidic sequences, interacts with the putative PfCK2 β subunits and the HsCK2 β subunit, is inhibited by the classic CK2 inhibitor TBB (3,4,5,6-tetrabromobenzotriazole), with a similar IC₅₀ to that of HsCK2 α , and is able to utilize GTP or ATP as a cosubstrate. We have also confirmed the identity of two PfCK2 β subunits. The N-terminal extension of PfCK2 β 2 is unusually long for CK2 β proteins, with 160 amino acids before the first conserved residue (Trp161 in PfCK2 β 2). Most CK2 β subunits from vertebrates have only eight amino acids prior to this conserved residue (*Homo sapiens*, *Gallus gallus*, *Mus musculus*, *Xenopus tropicalis*, *Bos taurus*, and *Danio rerio*); this N-terminal extension is expanded in yeast (*Saccharomyces cerevisiae*, 37 residues), trypanosomatids (*Trypanosoma brucei*, 27 residues; *Leishmania major*, 21 residues), plants (*Arabidopsis thaliana*, 100 residues; *Oryza sativa*, 92 residues), and alveolates (*Cryptosporidium parvum*, 27 residues; *Theileria parva*, 34 residues).

Within the alveolates, *Plasmodium yoelii yoelii* (125 residues) and *Plasmodium vivax* (157 residues) also have long extensions, but the extension of *P. falciparum* is the longest known. Homorepeat-containing proteins make up 35.7% of the proteome of *P. falciparum*, although the majority of these homorepeats are asparagines and lysines (48), unlike the polymers of acidic residues present in PfCK2 β 2. One hypothesis for the function of this extension is the downregulation of the alpha subunit. Polyglutamate is a potent CK2 inhibitor (50), and the N-terminal extension of PfCK2 β 2 is rich in polyglutamate and polyaspartate. This beta subunit also possesses an insertion of extra acidic residues (including a stretch of 11 consecutive aspartates) (Fig. 1) in the acidic domain known to downregulate CK2 (32). We have not been able to purify PfCK2 β 2 with the N-terminal extension, and therefore this hypothesis remains to be tested. However, we showed that the presence of either beta subunit reduces the activity of PfCK2 α toward

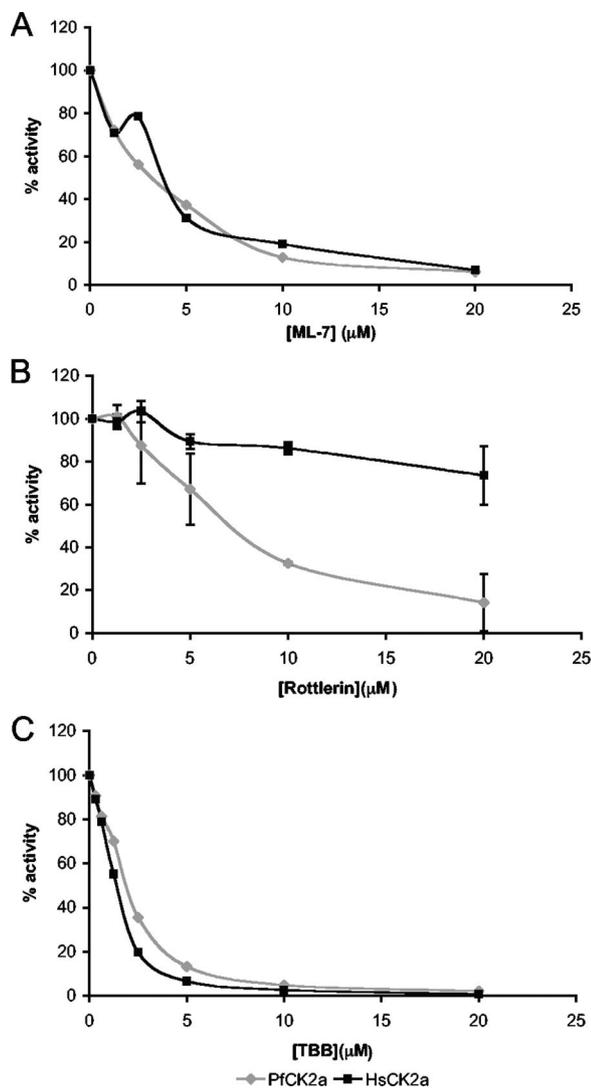


FIG. 8. PfCK2 α can be distinguished from HsCK2 α by small molecule inhibitors. Two small molecules, ML-7 and Rottlerin, were identified in a primary screen as inhibiting the activity of PfCK2 α to below 10% of the uninhibited enzyme. The inhibitors Rottlerin and ML-7 were included in increasing concentrations in kinase assays with 25 μ M ATP, 36 ng of enzyme, and the peptide RRREDEESDDEE as substrate. Activity was measured using the phosphocellulose assay method, and results were scored as a percentage of the control (no inhibitor). (A) ML-7. (B) Rottlerin. Mean values from two experiments are shown, with the error bars representing the standard deviations. (C) The classical CK2 inhibitor, TBB, has a similar inhibitory profile for PfCK2 α and HsCK2 α .

β -casein; such modulation of activity has been seen for other CK2s, often in a substrate-dependent fashion. For example, CK2 β stimulates human CK2 activity toward topoisomerase II and p53 and inhibits activity toward calmodulin (6). Thus, the presence of two beta subunit species in *P. falciparum* (whereas there is only one in human cells) is likely to allow exquisite control of the activity of the catalytic subunit. The interactions we detected in vitro between the recombinant catalytic and regulatory subunits suggest that the *P. falciparum* CK2 subunits may form a similar holoenzyme structure to that seen in other organisms, although the stoichiometry of the complex in vivo

will require detailed analysis of parasite extracts in nondenaturing conditions.

The existence of a number of discrete subpopulations of mammalian CK2 associated with different cellular compartments has been recognized (36), and it has been proposed that this is mediated by assembly of CK2 subunits as well as interaction with many other proteins. Our observation that the alpha and beta subunits possess putative signals for nuclear localization and protein export, respectively, suggests that in *Plasmodium* spp., like in other eukaryotes, CK2 may localize to a variety of compartments. Work is in progress to address this issue.

CK2 α has been shown to be essential for life for a variety of organisms (24, 31, 37). We have demonstrated here that PfCK2 α is required for parasite viability. We show that parasites lacking the enzyme are unable to survive or are impaired in their growth rate to such an extent that they are outcompeted by the parasites which retain wild-type genes. Our approach allows us to conclude that PfCK2 α plays an important role during erythrocytic schizogony but does not provide any information about the molecular basis for essentiality. We are addressing this issue in a number of ways, including conditional expression based on a destabilization domain (4), localization (see above), and identification of interacting partners. Nevertheless, our data validate PfCK2 as a potential drug target. We have also demonstrated that PfCK2 α is amenable to inhibition assays. Active HsCK2 α is present in erythrocytes (53); this raises the question of selectivity of antimalarial inhibitors based on PfCK2 α inhibition. We demonstrated that a small molecule inhibitor, Rottlerin, has a much lower IC₅₀ for PfCK2 α than for HsCK2 α . Although we have identified in Rottlerin a compound that can distinguish between the human and plasmodial CK2 α enzymes, it is unlikely to represent a suitable starting point for antimalarial drug discovery, since Rottlerin has multiple targets (41) and is too weak and non-specific an inhibitor even to be used in cellular assays (5). However, we have established that differential inhibition is possible, despite the 65% identity between the primary sequences of PfCK2 α and HsCK2 α , which suggests that specific inhibition of the plasmodial (versus host) enzyme should be feasible. The level of activity of recombinant PfCK2 α is such that the development of a high-throughput assay should be possible, opening the way for screening of chemical libraries as a first step toward antimalarial drug discovery based on PfCK2 α inhibition.

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