Research note

Primary structure and sexual stage-specific expression of a LAMMER protein kinase of *Plasmodium falciparum*[☆]

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Abstract

We have isolated a LAMMER-like gene from *Plasmodium falciparum* by vectorette technique. The gene consists of 3316 bp encoding a protein 881 amino acids with a predicted molecular mass of approximately 106.7 kDa. The encoded protein, termed PfLAMMER, is composed of two distinct domains. The N-terminal domain is not related to any previously described protein kinases and has several interesting features including multiple consensus phosphorylation sites for a range of protein kinases, a number of RS/SR dipeptides, a large proportion of charged amino acids, two putative nuclear localisation signals and 14 copies of a tetramer DKYD repeats. The C-terminal domain is characteristic of a kinase in the LAMMER family with the highest homology to the *Arabidopsis thaliana* AFC3 kinase. Genomic restriction analysis showed that PfLAMMER is encoded by a single copy gene in the parasite genome. A single transcript of approximately 3800 nucleotides is expressed specifically in the sexual stage, indicating that PfLAMMER may be important in regulating the processes of sexual differentiation of the parasite. © 2001 Australian Society for Parasitology Inc. Published by Elsevier Science Ltd. All rights reserved.

Keywords: Malaria; Protein kinase; LAMMER; Gametocyte; *Plasmodium falciparum*

The resistance of *Plasmodium falciparum* to drugs and the resistance of mosquitoes to insecticides have resulted in a resurgence of malaria in the tropics and subtropics. Therefore, development of effective vaccines and new anti-malarial drugs has become a top priority. A better understanding of the various cellular processes at different stages of the parasite will help to identify new targets for vaccine and drug development. Gametocytogenesis within the human erythrocytes and gametogenesis in the mosquito midgut are two key processes in the parasite life cycle. However, the molecular mechanisms involved in control of the sexual development are unclear (Lobo and Kumar, 1998). We are interested in signal transduction pathways involved in the sexual differentiation of *P. falciparum*. The first step toward this goal has been to identify the components, mainly protein kinases and phosphatases, of the signal transduction pathways. Recently, we have reported several sexual stage-specific genes encoding either protein serine/threonine

phosphatases (Li and Baker, 1997, 1998) or protein kinases (Li et al., 1996, 2000). In this paper, we describe the molecular cloning and characterisation of a novel gene encoding a *P. falciparum* LAMMER protein kinase (PfLAMMER). PfLAMMER possesses a large N-terminal extension and is expressed specifically in the sexual stage of the parasite life cycle, indicating that it may be important in regulating the processes of sexual stage development.

Protein kinases possess a catalytic domain of approximately 30 kDa within which some sequences are highly conserved among all members of the family (Hanks et al., 1988). Initially, although our interests lay in the *P. falciparum* protein kinases in general, cyclin-dependent kinases were targeted. Therefore, two primers, cdcI (5′-GAAA-AA/GITIGGIGAA/GGGIAC-3′, 1798–1817) and cdcII (5′-TCCAA/TAACTCCATACATCACT-3′, 2395–2415), were constructed based on the conserved sequences of subdomains I (EKIGEGT) and IX (SDVWSF/YG) and used in PCR. A band of 617 bp was amplified from *P. falciparum* 3D7A genomic DNA (data not shown) and inserted into the pGEM-T vector (Promega). The sequence data revealed that the PCR clone (cdcI-II) encodes 205 amino acids of a kinase-like protein. The A + T content and codon usage are typical of the coding region of *P. falciparum* genes (Weber, 1988). Southern blot analysis

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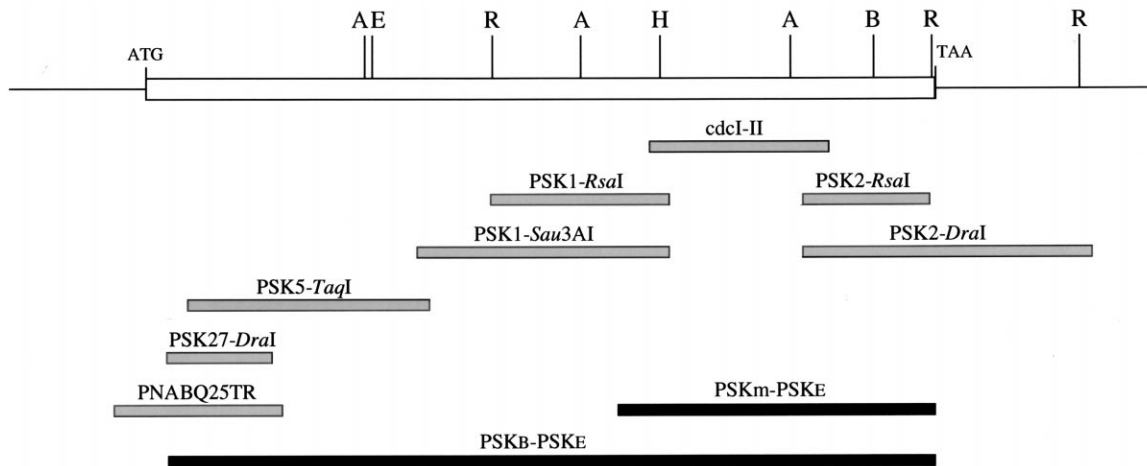*E-mail address:* lij@icrf.icnet.uk (J.-L. Li).

Fig. 1. A schematic representation of a partial restriction map of the *Pflammer* gene and the overlapping fragments used to determine its nucleotide sequence. A, *Acc*I; B, *Bam*HI; E, *Eco*RV; H, *Hinc*II; R, *Rsa*I. The open box represents the coding region of *Pflammer*.

confirmed that the cdcI-II clone is derived from parasite DNA (data not shown). To isolate the full-length gene, two specific primers, PSK1 (5′-GTTTCGAACAACTTT-TACAGCGTAG-3′, 1863–1887) (to obtain further sequence in the 5′ direction) and PSK2 (5′-CTCCA-GAAGTTATATTAAATTTGGG-3′, 2357–2381) (to obtain further sequence in the 3′ direction), were constructed on the basis of known sequences and used in PCR to screen vectorette libraries (Li et al., 1996). Four fragments (PSK1-*Rsa*I, PSK1-*Sau*3AI, PSK2-*Rsa*I and PSK2-*Dra*I) (Fig. 1) were obtained and sequenced. As expected, PSK1-*Sau*3AI contains all sequence of PSK1-*Rsa*I while PSK2-*Dra*I covers a full-length of PSK2-*Rsa*I. The sequence data of PSK1-*Sau*3AI permitted construction of the PSK5 primer (5′-CATACTTGTTGTCA-TATTTGTCCTC-3′, 1048–107-2) and subsequent screening of vectorette libraries. Based on the sequence of PSK5-*Taq*I, the PSK27 primer (5′-CATCTTGTTA-TAACGCTTCTGTTCTC-3′, 518–543) was designed to produce the PSK27-*Dra*I fragment. Database searches revealed that in the *P. falciparum* tag database there was a PNABQ25TR fragment containing all sequence of PSK27-*Dra*I. Nucleotide sequence analysis revealed a putative start codon in PNABQ25TR and a stop codon in PSK2-*Dra*I. In order to confirm the sequence obtained from the overlapping fragments, a pair of primers, PSKв (5′-**AGATCT**CTCGA-GATGGGTTATTCATCGAATTTGTA-3′, 234–257) and PSKᴇ (5′-TTAATAGTACTCATAATT-TTCTTC-3′, 2755–2778), were used to amplify the gene from genomic

DNA and the PCR product was sequenced in both strands (see Fig. 1). The sequence derived from overlapping fragments consists of 3316 bp and contains an open reading frame starting with an ATG codon at nucleotide 133 and terminating with a TAA codon at nucleotide 2776. The sequence and codon usage in the coding region are typical for a *P. falciparum* gene (Weber, 1988). The A + T contents of both flanking regions (91.9 and 84.4%, respectively) are characteristically higher than that of the coding region (75.5%). Two putative polyadenylation signals (AATAAA) are found at nucleotides +242 and +321, downstream of the TAA termination codon. A diffuse G + T-rich sequence occurs at nucleotide +24, downstream of the first polyadenylation signal. The existence of introns in the coding sequence was excluded by reverse transcription-PCRs (Li and Baker, 1997; Li et al., 2000; Li and Cox, 2000) using a number of primer pairs that cover the whole coding region.

The open reading frame encodes a protein of 881 amino acids with a predicted Mr of approximately 106.7 kDa. Database searches revealed that the amino acid sequence of PfLAMMER shares 56–60% similarity and 37–41% identity with kinases in the LAMMER family across the catalytic domain. PfLAMMER has the highest homology to the *Arabidopsis thaliana* AFC3 kinase (41% identity, 60% similarity) (Bender and Fink, 1994). Fig. 2 shows a sequence alignment of LAMMER kinases from diverse species. PfLAMMER is composed of two distinct domains. The N-terminal domain, consisting of 547 amino acid residues, is not related to any previously described protein

Fig. 2. Alignment of the predicted amino acid sequence of PfLAMMER with other members in the LAMMER family. The GenBank™/EMBL/DDJB database accession numbers are as follows: *P. falciparum* PfLAMMER, AF104915; human hCLK1, P49759; hCLK2, P49760; hCLK3, P49761; mouse mCLK1, P22518; mCLK2, O35491; mCLK3, O35492; mCLK4, O35493; rat rCLK3, Q63117; *D. melanogaster* DOA, P49762; *S. cerevisiae* KNS1, P32350; *S. pombe* KATB, Q10156; *A. thaliana* AFC1, P51566; AFC2, P51567; AFC3, P51568; and *N. tabacum* PK12, U73937. Sequences were aligned with the CLUSTAL W (1.60) multiple sequence alignment programme. The amino acid residues are numbered to the left of the sequence. Identical residues are indicated with asterisks and conservative changes indicated with dots. The eleven canonical subdomains of protein kinases (Hanks et al., 1988) are indicated by roman numerals. The residues conserved in the catalytic domain of the protein kinase family are highlighted with bold and the LAMMER kinase signature motifs are shaded with black. The boundary of the N-terminal and catalytic domains is indicated by arrows.

kinases and has several interesting features. Firstly, it is rich in serine/threonine [14% (74/547)] and tyrosine [14% (73/547)], forming a number of potential phosphorylation sites for a range of known protein kinases (Kennelly and Krebs,

1991). Thus, there is a strong possibility for regulation of the PfLAMMER activity by reversible phosphorylation of the N-terminal segment. Secondly, the N-terminal domain contains many serine and arginine residues [15% (92/

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PfLAMMER    1   MTELNIIDNNEKHSKYKLYKYCRNYINDFKNKYMMGYSSNLYKNSPNKYYYDTRNNSNNN

PfLAMMER   61   NNGYYYHNYNNGYQIRNKRNRSFETVSNSKNKHKISKKYKYNNEPHESFRDYDYKSKPSY

PfLAMMER  121   SSVKKIYDREQKRYNKMFVESRYNENHKNKYINNYNLSRKRNYYSYKKKIYTDRRNFDTH

PfLAMMER  181   FGQGNIIYNNEYYLINDGSSLNKKRLHYAKGSFRSRSRSKNYNTYNLEKSKKMKYSSRHN

PfLAMMER  241   IYRYHNGNKNKSISRWEDNYTYKDIYQPSSSLHKYKRRSNFTRDDNSIKSKMYYENKKTD

PfLAMMER  301   HLYSYEDKYDNKYDDKYDDKYDDKYDDKYDDKYDDKYDDKYDDKYDDKYDDNYDDKYDYG

PfLAMMER  361   EEEHRQEYYKKKKISFNNSTNNKSSINYDDVKRELKKRKKKKYSNESKVYDSLKRDETNH

PfLAMMER  421   HTNNNSYLNEINKKNSNLSNNYVAVRNKKRDKEYISDSNKSGFSNKGSYYMQKKKLHVDK
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AFC2       79                                               {78}EDDKDGHY
PK12       77                                               {76}DDDKDGHY
AFC1       96                                               {95}PDDKDGHY
AFC3       52                                               {51}DDDRDGHY
hCLK1     142                                              {141}EDDEEGHL
mCLK1     141                                              {140}EDDEEGHL
mCLK4     140                                              {139}EDDEEGHL
hCLK2     144                                              {143}EDDAEGHL
mCLK2     143                                              {142}EDDAEGHL
mCLK3     137                                              {136}EDDKEGHL
rCLK3     137                                              {136}EDDKEGHL
hCLK3     137                                              {136}EDDKEGHL
DOA       151                                              {150}QDDADGHL
KATB      343                                              {342}FDDDDGHY
KNS1      292                                              {291}KTDKDGHY
PfLAMMER  481   YDNDSHNRTISRTSSDNYSRKKYTHKRNRTKTSDTEDKKERKKKKKKKENNESDDEIVHF
                                                                 *    *
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                N-region ◄─┼─► catalytic domain
AFC2       87   IFELGDDLTP--RYKIYSKMGEGTFGQVLECWDRER-KEMVAVKIVRGVKKYREAAMIEI
PK12       85   MFELGENLTT--RYKILKKIGEGTFGQVLECWDREQ-KGFVAIKIIRSIKKYREAAMVEV
AFC1      104   VFVVGDTLTP--RYQMLSKMGEGTFGQVLECFDNKN-KEVVAIKVIRSINKYREAAMIEI
AFC3       60   VFSLRDNLTP--RYKILSKMGEGTFGRVLECWDRDT-KEYVAIKIIRSIKKYRDAAMIEI
hCLK1     150   ICQSGDVLSA--RYEIVDTLGEGAFGKVVECIDHKAGGRHVAVKIVKNVDRYCEAARSEI
mCLK1     149   ICQSGDVLSA--RYEIVDTLGEGAFGKVVECIDHKVGGRRVAVKIVKNVDRYCEAAQSEI
mCLK4     148   ICQSGDVLRA--RYEIVDTLGEGAFGKVVECIDHGMDGLHVAVKIVKNVGRYREAARSEI
hCLK2     152   IYHVGDWLQE--RYEIVSTLGEGTFGRVVQCVDHRRGGARVALKIIKNVEKYKEAARLEI
mCLK2     151   IYHVGDWLQE--RYEIVSTLGEGTSGRVVQCVDHRRGGTRVALKIIKNVEKYKEAARLEI
mCLK3     145   VCRIGDWLQE--RYEIVGNLGEGTFGKVVECLDHARGKSQVALKIIRNVGKYREAARLEI
rCLK3     145   VCRIGDWLQE--RYEIVGNLGEGTFGKVVECLDHARGKSQVALKIIRNVGKYREAARLEI
hCLK3     145   VCRIGDWLQE--RYEIVGNLGEGTFGKVVECLDHARGKSQVALKIIRNVGKYREAARLEI
DOA       159   IYHTGDILHH--RYKIMATLGEGTFGRVVKVKDMER-DYCMALKIIKNVEKYREAAKLEI
KATB      351   KVVPNSKFAN--RYTVVRLLGHGTFGKVIQCYDQST-GRHCAIKVTRAIPKYREASLIEL
KNS1      300   VYQENDIFGSGGRFVVKDLLGQGTFGKVLKCIDNKYEPNYVAVKVIRAVDRYREAAKTEL
PfLAMMER  541   SWKKGMLLNN--AFLVIRKMGDGTFGRVLLCQHIDN-KKYYAVKVVRRNIKKYTRSAKIEA
                  :     :  : *  *:  *:*: :     :         *:*:  ::  :*  ::  *
                                  I               II            III
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AFC2      144   EMLQQLGKHDKGGN-RCVQIRNWFDYRNHICIVFEKLGSSLYDFLRKNNYRSFPIDLVRE
PK12      142   DVLQLLGRYDRGGT-RCVQLRNWFDYRNHICLVFEKLGPSLFDFLRKNSYRAPPVDLVRE
AFC1      161   DVLQRLTRHDVGGS-RCVQIRNWFDYRNHICIVFEKLGPSLFDFLRKNSYRSFPIDLIRK
AFC3      117   DVLQKLVKSDKGRT-RCVQMKNWFDYRNHICIVFEKLGPSLFDFLRNKYSAFPLALVRD
hCLK1     208   QVLEHLNTTDPNSTFRCVQMLEWFEHRGHICIVFELLGLSTYDFIKENGFLPFRDHIRK
mCLK1     207   QVLEHLNTTDPHSTFRCVQMLEWFEHRGHICIVFELLGLSTYDFIKENSFLPFRMDHIRK
mCLK4     206   QVLEHLNTTDPNSVFRCVQMLEWFDHHGHVCIVFELLGLSTYDFIKENSFLPFQIDHIRQ
hCLK2     210   NVLEKINEKDPDNKNLCVQMFDWFDYHGHMCISFELLGLSTFDFLKDNNYLPYPIHQVRH
mCLK2     209   NVLEKINEKDPDNKNLCVQMFDWFDYHGHMCISFELLGLSTFDFLKDNNYLPYPIHQVRH
mCLK3     203   NVLKKIKEKDKENKFLCVLMSDWFNFHGHMCIAFELLGKNTFEFLKENNFQPYPLPHVRH
rCLK3     203   NVLKKIKEKDKENKFLCVLMSDWFNFHGHMCIAFELLGKNTFEFLKENNFQPYPLPHVRH
hCLK3     203   NVLKKIKEKDKENKFLCVLMSDWFNFHGHMCIAFELLGKNTFEFLKENNFQPYPLPHVRH
DOA       216   NALEKIAQKDPHCDHLCVKMIDWFDYHGHMCIVFELLGLSVFDFLRENNYEPYPLDQVRH
KATB      408   RVLQTIAHSDPTNENKCIQLRDYFDYRNHICIVTDLFGWSVFDFLKNNNYIPFPLKHIQM
KNS1      360   RILQTILNNDPQGQFQCLLLRECFDYKNHICLVTDLYGRSIYDFMCSNGIAGSP-ALISG
PfLAMMER  598   DILKKIQNDDINNN-NIVKYHGKFMYYDHMCLIFEPLGPSLYEIITRNNYNGFHIEDIKL
                  *:  :    *          *        *:*:        *    * :::: *
                              IV              V
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AFC2      203   IGWQLLECVAFMHDLRMI[HIDLKPENI]LVSSDYVKIPEYKGS-RLQ----RDVCY-KR-
PK12      201   IGRQLLECVAFMHDMRLI[HIDLKPENI]FVSADYIKVPDYKGTPWSH----RDRSFSKR-
AFC1      220   LGRQLLESVAYMHDLRLI[HIDLKPENI]LVSSEYIKIPDYKFLSRPT----KDGSYFKN-
AFC3      176   FGCQLLESVAYMHELQLV[HIDLKPENI]LVSSENVKLPDNKRS---A----ANETHFRC-
hCLK1     268   MAYQICKSVNFLHSNKLT[HIDLKPENI]FVQSDYTEAYNPKIK--------RDERT----
mCLK1     267   MAYQICKSVNFLHSNKLT[HIDLKPENI]FVKSDYTEAYNPKMK--------RDERT----
mCLK4     266   MAYQICQSINFLHHNKLT[HIDLKPENI]FVKSDYVVKYNSKMK--------RDERT----
hCLK2     270   MAFQLCQAVKFLHDNKLT[HIDLKPENI]FVNSDYELTYNLEKK--------RDERS----
mCLK2     269   MAFQLCQAVKFLHDNKLT[HIDLKPENI]FVNSDYELTYNLEKK--------RDERS----
mCLK3     263   MAYQLCHALRFLHENQLT[HIDLKPENI]FVNSEFETLYNEHKS--------CEEKS----
rCLK3     263   MAYQLCHALRFLHENQLT[HIDLKPENI]FVNSEFETLYNEHKS--------CEEKS----
hCLK3     263   MAYQLCHALRFLHENQLT[HIDLKPENI]FVNSEFETLYNEHKS--------CEEKS----
DOA       276   MAYQLCYSVKFLHDNRLT[HIDLKPENI]FVDSDYTSHYNHKIN--------REVRR----
KATB      468   LSQQLFKSVAFLHSLGLV[HIDLKPENM]LVSNASRTIRLPYRN--------YSQKV----
KNS1      419   HCRQLIRSVCFLHDLGII[HIDLKPENI]ICDETHIAQKLPLKTVQSLSKRRREASKGKRK
PfLAMMER  657   YCIEILKALNYLRKMSLT[HIDLKPENI]LDDPYFEKSLITVRR-------VTDGKKIQIY
                  ::   : : :::  *:*:*******:*:
                                  VI
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```
AFC2      256   VPKSSAIKVIDFGSTTYERQDQTYIVSTRHYRAPEVILGLGWSYPCDVWSVGCIIVELCT
PK12      256   LPKSSAIKVIDFGSTAYERPDHNYIVSTRHYRAPEVILGLGWSYPCDLWSVGCILIELCS
AFC1      275   LPKSSAIKLIDFGSTTFEHQDHNYIVSTRHYRAPEVILGVGWNYPCDLWSIGCILVELCS
AFC3      228   LPKSSAIKLIDFGSTVCDNRIIHHSIVQTRHYRSPEVILGLGWSYQCDLWSIGCILFELCT
hCLK1     316   -LINPDIKVVDFGSATYDDEHHSTLVSTRHYRAPEVILALGWSQPCDVWSIGCLLIEYYL
mCLK1     315   -IVNPDIKVVDFGSATYDDEHHSTLVSTRHYRAPEVILALGWSQPCDVWSIGCLLIEYYL
mCLK4     314   -LKNTDIKVVDFGSATYDDEHHSTLVSTRHYRAPEVILALGWSQPCDVWSIGCLLIEYYL
hCLK2     318   -VKSTAVRVVDFGSATFDHEHHSTIVSTRHYRAPEVILELGWSQPCDVWSIGCIIFEYYV
mCLK2     317   -VKSTAVRVVDFGSATFDHEHHSTIVSTRHYRAPEVILELGWSQPCDVWSIGCIIFEYYV
mCLK3     311   -VKNTSIRVADFGSATFDHEHHTTIVATRHYRPPEVILELGWAQPCDVWSIGCILFEYYR
rCLK3     311   -VKNTSIRVADFGSATFDHEHHTTIVATRHYRPPEVILELGWAQPCDVWSIGCILFEYYR
hCLK3     311   -VKNTSIRVADFGSATFDHEHHTTIVATRHYRPPEVILELGWAQPCDVWSIGCILFEYYR
DOA       324   -VKNTDVRLIDFGSATFDHEHHSTIVSTRHYRAPEVILELGWSQPCDVWSIGCILFELYL
KATB      516   -LNSCEIRLIDFGSATFEDEYHSSVVSTRHYRAPEIILGLGWSYPCDLWSIGCILVELFT
KNS1      479   ILKNPEIKIIDFGSAIFHYEYHPPVISTRHYRAPEIVLGLGWSFPCDIWSIACVLVELVI
PfLAMMER  710   RTKSTGIKLIDFGCATFKSDYHGSIINTRQYRAPEVILNLGWDVSSDMWSFGCVLAELYT
                  :::  ***  :::   :  :: **:**  **::*  **    *:**  *::  *
                          VII            VIII            IX
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```
AFC2      316   GEALFQTHENL[PFHLAMMER]VG-PFPQQMLKKVDRH--------------SEKYVRRG-
PK12      316   GEALFQTHENL[PFHLAMMER]VG-PLPSQMLKRVDRH--------------AEKYVRRG-
AFC1      335   GEALFQTHENL[PFHLAMMER]VG-PLPPHMVLRADRR--------------SEKYFRRGA
AFC3      288   GEALFQTHDNL[PFHLAMMER]AG-PLPEHMTRKASRG--------------AEKYFRRGC
hCLK1     375   GFTVFPTHDSK[PFHLAMMER]IIG-PLPKHMIQKTR----------------KRKYFHHD-
mCLK1     374   GFTVFPTHDSR[PFHLAMMER]IIG-PLPKHMIQKTR----------------KRRYFHHD-
mCLK4     373   GFTVFQTHDSK[PFHLAMMER]IIG-PIPAHMIQKTR----------------KRKYFHHN-
hCLK2     377   GFTLFQTHDNR[PFHLAMMER]IIG-PIPSRMIRKTR----------------KQKYFYRG-
mCLK2     376   GFTLFQTHDNR[PFHLAMMER]IIG-PVPSRMIRKTR----------------KQKYFYRG-
mCLK3     370   GFTLFQTHENR[PFHLVMMEKII]G-PIPSHMIHRTR---------------KQKYFYKG-
rCLK3     370   GFTLFQTHENR[PFHLVMMEKII]G-PIPSHMIHRTR---------------KQKYFYKG-
hCLK3     370   GFTLFQTHENR[PFHLVMMEKII]G-PIPSHMIHRTR---------------KQKYFYKG-
DOA       383   GITLFQTHDNR[PFHLAMMER]IIG-QIPYRMARKT----------------KTKYFYHG-
KATB      575   GQALFQTHEDS[PFHLGMMEKII]G-PFDRNMISRSSRT--------------SQRFFKSDG
KNS1      539   GESLYPIHENL[PFHLAMHQE]NGTPFPTDIIDKMFYKSKHKLGNSPSDLNSTVIKHFDRKT
PfLAMMER  770   GSLLFRTHEHM[PFHLAMHSL]IQ-PIPKNMLYEATKTN-------------GSKYVNKDE
                *  :: :   *:  **: **:  **:
                                        X
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AFC2      359   -RLDWPDGATSRD------------------SLKAVLKL-----------PRLQNL
PK12      359   -RLDWPEGATSRE------------------SIKSVMKL-----------PRLQNL
AFC1      379   -KLDWPEGATSRD------------------SLKAVWKL-----------PRLPNL
AFC3      332   -RLNWPEGANSRE------------------SIRAVKRL-----------DRLKDM
hCLK1     416   -RLDWDEHSSAGR------------------YVSRACK-----------PLKEFM
mCLK1     415   -RLDWDEHSSAGR------------------YVSRRCK-----------PLKEFM
mCLK4     414   -QLDWDEHSSAGR------------------YVRRRCK-----------PLKEFM
hCLK2     418   -RLDWDENTSAGR------------------YVRENCK-----------PLRRYL
mCLK2     417   -RLDWDENTSAGR------------------YVRENCK-----------PLRRYL
mCLK3     411   -GLVWDENSSDGR------------------YVKENCK-----------PLKSYM
rCLK3     411   -GLVWDENSSDGR------------------YVKENCK-----------PLKSYM
hCLK3     411   -GLVWDENSSDGR------------------YVKENCK-----------PLKSYM
DOA       423   -KLDWDEKSSAGR------------------YVRDHCK-----------PLFLCQ
KATB      619   -KVRYPLSNTPKK------------------SINYLQSL-----------QTLEQI
KNS1      599   LSLQWPEKNKRGDTITTEKSMKRVLQSCDRLDIYISKVLKQDYGDSLSINWNLRPPEKNWS
PfLAMMER  815   LKLAWPENASSIN------------------SIKHVKK--------------CLP--
                : :                            :
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AFC2      385   IM------QHVDHSAGE---------------LINMVQGLLRRFDPSERITAREALRHPF
PK12      385   VM------QHVDHSAGD---------------LIDLLQGLLRFDPSIRMTAHDALRHPF
AFC1      405   IM------QHVDHSAGD---------------LIDLLQGLLRYDPTERFKAREALNHPF
AFC3      358   VS------KHVDNTRSR---------------FADLLYGLLAYDPSERLTANEALDHPF
hCLK1     441   LS------QDVEHER---------------LFDLIQKMLEYDPAKRITLREALKHPF
mCLK1     440   LS------QDAEHEF---------------LFDLVGKILEYDPAKRITLKEALKHPF
mCLK4     439   LC------HDEEHEK---------------LFDLVRRMLEYDPARRITLDEALQHPF
hCLK2     443   TS------EAEEHHQ---------------LFDLIESMLEYEPAKRLTLGEALQHPF
mCLK2     442   TS------EAEDHHQ---------------LFDLIENMLEYEPAKRLTLGEALQHPF
mCLK3     436   LQ------DSLEHVQ---------------LFPDLMRRMLEFDPAQRITLAEALLHPF
rCLK3     436   LQ------DSLEHVQ---------------LFPDLMRRMLEFDPQSQRITLAEALLHPF
hCLK3     436   LQ------DSLEHVQ---------------LFPDLMRRMLEFDPAQRITLAEALLHPF
DOA       448   LS------DSEDHCE---------------LFSLIKKMLEYEPSSRITLGEALHHPF
KATB      645   FA------VSSPEVA-L---------------LLDLLKKVFVYDPKRRITAKEALWHPF
KNS1      659   LINSKLAWKRQTHSSSSSTTDELDKETFLFWYWFIDLLRKMFEFDPTKRITAKDALDHEW
PfLAMMER  838   LY------KIIKHEL----------------FCDFLYSILQIDPTLRPSPAELLKHKF
                :  ::  ::  :*  *   : * * :
                                        XI
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AFC2      423   FARRR---------------------------
PK12      423   FTKDQYRRL-----------------------
AFC1      443   FTRSREQSIPPFNPNPHPFLYNQKN-------
AFC3      396   FKSSS---------------------------
hCLK1     477   FDLLKKSI------------------------
mCLK1     476   FYPLKKHT------------------------
mCLK4     475   FDLLKRK-------------------------
hCLK2     479   FARLRAEPPN-KL--------WDSSRDISR----
mCLK2     478   FACLRTEPPNTKL--------WDSSRDISR----
mCLK3     472   FAGLTPEERS-----------FHSSRNPSR----
rCLK3     472   FAGLTPEERS-----------FHSSRNPSR----
hCLK3     472   FAGLTPEERS-----------FHTSRNPSR----
DOA       484   FDRLPPHHRVGEVSNKQPLSSGSSSRERSHSLSR
KATB      682   FTQPISSNL-----------------------
KNS1      719   FNLGILDDGIATYNNTQG--------------
PfLAMMER  874   LEENYEYY------------------------
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547)], indicating that PfLAMMER is an SR-like protein (Nayler et al., 1997). Close inspection of the sequence revealed 11 RS/SR dipeptides and two RSRS motifs in this region, implying that the N-terminal segment may interact with other SR proteins (Wu and Maniatis, 1993; Kohtz et al., 1994). Thirdly, a large proportion of charged amino acids [37% (203/547) K, R, E and D] exists in the N-terminal segment, suggesting further that this region may be involved in the protein-protein interactions. Fourthly, the N-terminal segment possesses at least two putative nuclear localisation signals (Dingwall and Laskey, 1991), implying that PfLAMMER may be a nuclear protein. Fifthly, there are 14 copies of a tetramer DKYD repeats at residue positions 307 to 482. Finally, in contrast to those of the known kinases in the LAMMER family, the N-terminal segment is the largest extension of LAMMERs yet described, placing PfLAMMER in a distinct category within the LAMMER family. The kinase catalytic domain of PfLAMMER is composed of 334 residues and contains all 11 conserved subdomains of the protein kinase family (Hanks et al., 1988). It has almost all of the characteristic features of a kinase including 15 invariant residues, the ATP-binding motif and the catalytic motif (Hanks et al., 1988; Knighton et al., 1991; DeBondt et al., 1993). The conserved sequences (HTDLKPENIL) in subdomain VI and (EHLAMMESII) in subdomain X are characteristic for LAMMER kinases (Yun et al., 1994). However, a highly conserved R residue in the LAMMER motif is replaced by an S residue in PfLAMMER. A similar change was also found in mammalian CLK3 kinases (Hanes et al., 1994; Becker et al., 1996; Nayler et al., 1997). By analogy with the crystallised structure of two protein kinases (Knighton et al., 1991; DeBondt et al., 1993), the LAMMER motif lies in an α-helix below the substrate-binding cleft, potentially allowing it to contact with substrates. Therefore, the R to S change in the LAMMER motif might affect substrate specificity of PfLAMMER. The sequences (DLKPEN) in subdomain VI indicate that PfLAMMER is a serine/threonine kinase (Hanks et al., 1988); however, the sequences (TRQYRAPE) in subdomain VIII suggest that PfLAMMER could also be a tyrosine kinase (Hanks et al., 1988; Howell et al., 1991). Indeed, many kinases in the LAMMER family have been demonstrated to exhibit a dual-specificity, phosphorylating not only serine and threonine but also tyrosine (Ben-David et al., 1991; Howell et al., 1991; Yun et al., 1994; Duncan et al., 1995; Lee et al., 1996; Sessa et al., 1996). Therefore, PfLAMMER may be a parasite dual-specificity protein kinase. It is worthnoting that, up to date, no tyrosine kinase has been reported in *P. falciparum*.

To investigate the structural organisation of the *Pflammer* gene in the *P. falciparum* genome, clone 3D7A genomic DNA was digested with a number of restriction enzymes and analysed by Southern blotting. Hybridisation of the cdcI-II (see Fig. 1) probe revealed a single band in digests with *Bam*HI, *Bcl*I, *Eco*RI, *Eco*RV, *Hinc*II and *Rsa*I, and two bands in digest with *Acc*I, consistent with the restriction map (Fig. 3a), suggesting strongly that *Pflammer* is encoded by a single copy gene in the parasite genome. However, hybridisation of the same blot with the PSK1-*Sau*3AI fragment detected two bands (one predominant band and the other faint) in digests with *Acc*I, *Bcl*I, *Eco*RI, *Eco*RV and *Hinc*II, and three bands in digest with *Rsa*I, contradictory to
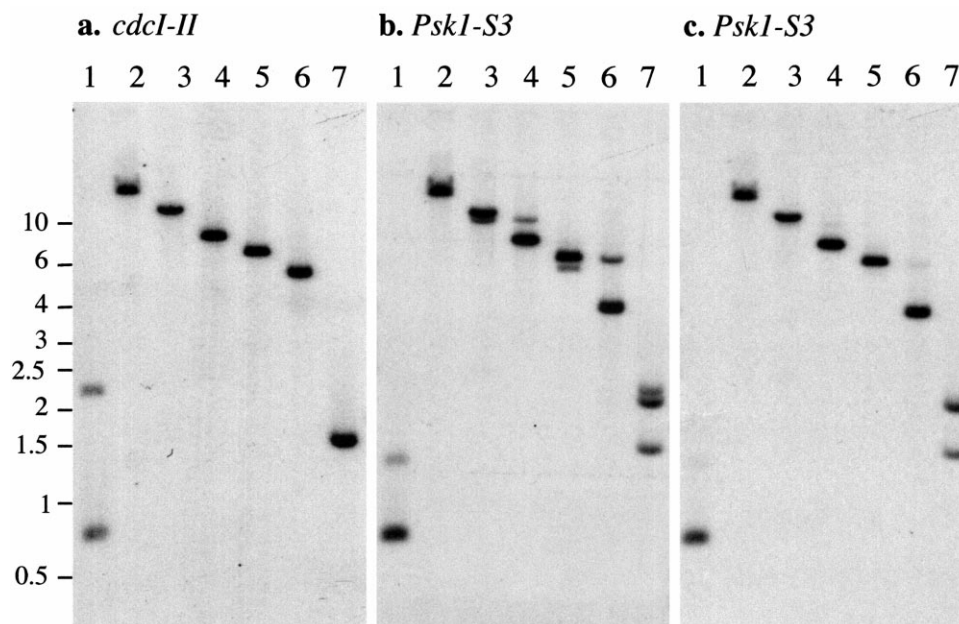
Fig. 3. Southern blot analysis of the *Pflammer* gene. Four micrograms of genomic DNA from *P. falciparum* clone 3D7A were digested with restriction enzymes, electrophoresed on a 1.0% agarose gel, transferred onto a nylon membrane, and probed with the cdcI-II fragment (a) and PSK1-*Sau*3AI fragment (b and c), respectively. The blot was washed either in a normal condition (at 56°C) (a and b) (Li et al., 1996) or in a higher condition (at 65°C) (c). Lanes 1–7 correspond to digests with *Acc*I, *Bam*HI, *Bcl*I, *Eco*RI, *Eco*RV, *Hinc*II and *Rsa*I. The sizes of 1–Kb DNA markers (M) are given in kilobase pairs to the left.
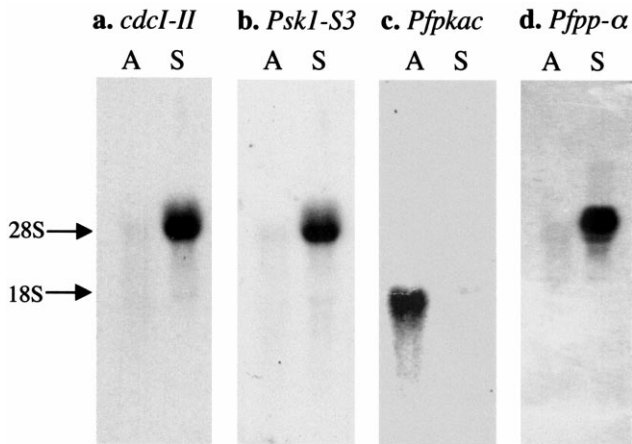
Fig. 4. Northern blot analysis of the *Pflammer* gene. Ten micrograms of total RNA extracted from the asexual erythrocytic stages (A) and the sexual erythrocytic stage (S) of *P. falciparum* (3D7A) were fractionated in a denaturing formaldehyde gel, blotted onto a nylon membrane and hybridised to radiolabeled probes. The positions of *P. falciparum* rRNA subunits (18 and 28S) are indicated by arrows. a, b, c and d are autoradiographs of the membrane probed with the *Pflammer* gene (cdcI-II and PSK1-*Sau*3AI), the *Pfpkac* gene (Li and Cox, 2000) and the *Pfpp*-α gene (Li and Baker, 1998), and exposed for 44, 1 and 24 h, respectively. *Pflammer* detected a transcript of approximately 3800 nucleotides in the sexual stage, *Pfpkac* hybridised with a band of approximately 1800 nucleotides in the asexual stage and *Pfpp*-α hybridised with a band of approximately 3900 nucleotides in the sexual stage.

the restriction map (Fig. 3b). The faint bands disappeared gradually with higher stringency washing conditions (Fig. 3c), indicating the presence of a PSK1-*Sau*3AI-related gene in the *P. falciparum* genome. To date, at least four LAMMER kinases have been identified in mouse (Ben-David et al., 1991; Howell et al., 1991; Nayler et al., 1997), three in human (Johnson and Smith, 1991; Hanes et al., 1994), three in *A. thaliana* (Bender and Fink, 1994), one in rat (Becker et al., 1996), one in *Drosophila melanogaster* (Yun et al., 1994), one in *Saccharomyces cerevisiae* (Padmanabha et al., 1991), one in *Schizosaccharomyces pombe* (the database accession number Q10156) and one in *Nicotiana tabacum* (Sessa et al., 1996). Our southern results suggest that only one LAMMER kinase exists in *P. falciparum*.

In order to obtain some information on how *Pflammer* mRNA levels are regulated during parasite development and differentiation, a northern blot containing equal quantities of total RNA prepared from cultures enriched in stage III to stage V gametocytes and from mixed asexual erythrocytic stages was probed with the cdcI-II clone. A single transcript of approximately 3800 nucleotides in size was detected only in the lane containing the sexual stage RNA, migrating just behind of the 28S rRNA bands (Fig. 4a). This was also confirmed by hybridising the same blot with the PSK1-*Sau*3AI fragment (Fig. 4b). The result is in sharp contrast to the *lammer* genes of humans, mice and insects, for which multiple transcripts were detected due

to the alternative and incomplete splicings (Howell et al., 1991; Johnson and Smith, 1991; Hanes et al., 1994; Yun et al., 1994; Duncan et al., 1995; Nayler et al., 1997). As internal controls for hybridisation to the asexual and sexual stage mRNA, the same blot was hybridised with *Pfpkac*, an asexual stage-specific gene (Li and Cox, 2000), and *Pfpp*-α, a sexual stage-specific gene (Li and Baker, 1998) (Fig. 4c,d). The result suggests that PfLAMMER is involved in sexual stage-specific events. In rodent, it has been shown that mCLK2, mCLK3, mCLK4 and rCLK3 are predominantly expressed in testis, indicating that these kinases regulate a testicular function (Becker et al., 1996; Nayler et al., 1997). In *Arabidopsis*, the AFC1 kinase gene is able to complement yeast signal transduction mutants via activation of the transcription factor STE12 (Bender and Fink, 1994). In *Drosophila*, DOA is critical to the development of the fly embryo and affects eye differentiation and sex determination by phosphorylation of the SR proteins and consequently regulation of alternative pre-mRNA splicing (Yun et al., 1994; Du et al., 1998). In mammalian cells, over expression of the CLK1 kinase in PC12 cells seems to initiate their differentiation into neural derivatives possibly through activation of a mitogen-activated protein kinase cascade (Myers et al., 1994). The mouse CLKs have been demonstrated to bind to and phosphorylate SR-rich mRNA splicing factors (Colwill et al., 1996; Nayler et al., 1997). Overexpression of CLK1 in COS cells leads to the subcellular redistribution of the SR-proteins, and to alternations in mRNA splicing in vivo (Colwill et al., 1996; Duncan et al., 1997). Recently, it has been shown that both mCLK1 and hCLK2 can phosphorylate and activate the tyrosine phosphatases PTP-1B and YPTP1 in vitro (Moeslein et al., 1999). By analogy, therefore, we speculate that PfLAMMER may play an important role in signal transduction pathway(s) involved in the sexual differentiation of the parasite perhaps by phosphorylation of the sexual stage-specific mRNA splicing factors and/or protein phosphatases.

Acknowledgements

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