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RESEARCH

Genomic and Bioinformatic Analysis of NADPH-Cytochrome P450 Reductase in *Anopheles stephensi* (Diptera: Culicidae)

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ABSTRACT. The cytochrome P450 monooxygenase (P450) enzyme system is a major mechanism of xenobiotic biotransformation. The nicotinamide adenine dinucleotide phosphate (NADPH)-cytochrome P450 reductase (CPR) is required for transfer of electrons from NADPH to P450. One CPR gene was identified in the genome of the malaria-transmitting mosquito *Anopheles stephensi* Liston (Diptera: Culicidae). The gene encodes a polypeptide containing highly conserved flavin mononucleotide-, flavin adenine dinucleotide-, and NADPH-binding domains, a unique characteristic of the reductase. Phylogenetic analysis revealed that the *A. stephensi* and other known mosquito CPRs belong to a monophyletic group distinctly separated from other insects in the same order, Diptera. Amino acid residues of CPRs involved in binding of P450 and cytochrome *c* are conserved between *A. stephensi* and the Norway rat *Rattus norvegicus* Berkenhout (Rodentia: Muridae). However, gene structure particularly within the coding region is evidently different between the two organisms. Such difference might arise during the evolution process as also seen in the difference of P450 families and isoforms found in these organisms. CPR in the mosquito *A. stephensi* is expected to be active and serve as an essential component of the P450 system.

Key Words: binding domain, gene structure, phylogenetic tree, sequence analysis

The mosquito *Anopheles stephensi* Liston (Diptera: Culicidae) is one of the major vectors that transmit the human malaria parasite *Plasmodium falciparum* Welch. Geographical distribution of the mosquito ranges from the Middle East to the Indian subcontinent (Kiszewski et al. 2004). Various types of insecticides have been used in controlling populations of mosquito vectors. Insecticide resistance or changes in insecticide susceptibility have been documented in *A. stephensi* (Ganesh et al. 2003, Enayati and Ladonni 2006, Tiwari et al. 2010, Tikar et al. 2011, Shetty et al. 2012).

The cytochrome P450 monooxygenase (P450) system is well known as a major mechanism of xenobiotic biotransformation. Involvement of specific P450 isoforms in insecticide resistance has been characterized in several insects, including mosquitoes (Nikou et al. 2003, Shen et al. 2003, Djouaka et al. 2008, Yang and Liu 2011, Bariami et al. 2012). As part of the redox reaction, a nicotinamide adenine dinucleotide phosphate (NADPH)-cytochrome P450 reductase (CPR) is needed in transferring electrons to P450s. CPR is a flavoprotein containing binding domains for flavin mononucleotide (FMN), flavin adenine dinucleotide (FAD), and NADPH, as determined by X-ray crystallography of rat CPR (Wang et al. 1997). NADPH serves as electron donor, whereas FAD and FMN serve as cofactors in the pathway of electron transfer. In eukaryotic microsomes, both P450s and CPRs are integral membrane proteins. Although there is such an extensive diversity of P450 isoforms identified in animals (Nelson 2011), each animal's genome generally carries only one CPR gene (Porter et al. 1990). Thus, interaction between the two enzymes in the microsomal environment is probably in the ratio of multiple P450s to one CPR. Feyereisen (2005, 2012) suggests a ratio of 6–18 P450s to 1 reductase in insects. Lycett et al. (2006) showed that silencing the CPR gene could impact insecticide susceptibility in the mosquito *Anopheles gambiae*.

To date, only a few mosquito CPRs have been identified and deposited in the NCBI GenBank database: *Aedes sollicitans* (D. Sun and L.B.B., unpublished data), *Aedes aegypti* (Nene et al. 2007), *Anopheles funestus* (Matambo et al. 2010), *A. gambiae* (Nikou et al. 2003), *Anopheles minimus* (Kaewpa et al. 2007), and *Culex*

quinquefasciatus (Arensburger et al. 2010). By using *A. gambiae* microarray platforms, a few possible P450 transcripts were identified in *A. stephensi* (Vontas et al. 2007). A small nucleotide sequence, possibly representing a fragment of a CPR gene, was obtained from *A. stephensi* by multilocus DNA sequencing as part of a phylogenetic study among different anopheline mosquitoes (Dixit et al. 2010). However, a complete sequence of the *A. stephensi* CPR gene has not, to date, been determined.

In this study, a complete coding sequence of a CPR gene was identified by comparative genomic analysis of the *A. stephensi* genome; the whole genome shotgun sequence of *A. stephensi* was originally deposited in the NCBI GenBank database by Hall and Jiang, Virginia Tech University, in 2012. The deduced amino acid sequence was analyzed by using bioinformatic and phylogenetic analyses. Conserved residues were further characterized in relation to rat and known mosquito CPRs. This study was undertaken because of our long-standing interest in the role of the P450 enzyme system in the responses of insects, especially mosquitoes, to insecticides applied in an attempt to control their populations and also in their responses to allelochemical substances in their ecological habitats (Suwanchaichinda and Brattsten 2002). (Note: the nucleotide sequence of the *A. stephensi* CPR transcript is available in the Third Party Annotation Section of the DDBJ/EMBL/GenBank databases under the accession number TPA:BK008720.)

Materials and Methods

Genomic Analysis. To screen the NCBI nonredundant nucleotide database for any possible CPR gene identified in *A. stephensi*, the nucleotide sequence of the *A. gambiae* CPR gene was used as a query in a BlastN search. The *A. gambiae* CPR nucleotide and amino acid sequences were then used in BlastN and TblastN searches, respectively, against the genome sequences in the NCBI database. An identified genome contig was compared with the *A. funestus* CPR gene to reconfirm its identity. Additionally, the *Ae. sollicitans* CPR cDNA, which was cloned and sequenced in our laboratory, was used as an inferential evidence for the annotation of the *A. stephensi* CPR gene.

Sequences obtained as results of both nucleotide and genome sequence database searches were subsequently used for further analysis.

Sequence Characterization. An identified genome contig containing a possible *A. stephensi* CPR gene was manually analyzed by comparing it with previously identified mosquito CPR genes to obtain a complete coding region. The intron–exon organization (from initiation codon to stop codon) and the sizes of exons of the *A. stephensi* CPR gene were determined according to CPR genes in the genome releases of *A. gambiae*, *Ae. aegypti*, and *C. quinquefasciatus* available in VectorBase (<http://www.vectorbase.org>). The CPR gene of the Norway rat (*Rattus norvegicus*) was used as a representative of vertebrates for comparison. Information on the intron–exon organization of the rat CPR gene was obtained from Ensemble (<http://www.ensembl.org>).

A hypothetical translation of the complete open reading frame of the *A. stephensi* CPR transcript was performed to obtain a deduced amino acid sequence. The translated protein was then used in a BlastP search against the NCBI nonredundant protein database to confirm the *A. stephensi* CPR identity compared with other known CPRs in invertebrates and vertebrates. The isoelectric point (pI) and molecular mass of the *A. stephensi* CPR were calculated by a proteomics tool (Gasteiger et al. 2005). A transmembrane spanning region was predicted by TMpred (http://embnet.vital-it.ch/software/TMPRED_form.html) and TMHMM Server 2.0 (<http://www.cbs.dtu.dk/services/TMHMM>).

In addition to putative conserved domains identified as part of the BlastP search, the identification of binding domains (for FMN, FAD, and NADPH) and catalytic residues were predicted by Pfam 26.0 (<http://pfam.sanger.ac.uk>). A multiple sequence alignment between *A. stephensi* CPR and other mosquito CPRs was performed by ClustalW (Larkin et al. 2007). Amino acid residues involved in binding to cytochrome P450s and cytochrome *c* were predicted based on the rat CPR previously identified by Shen et al. (1989) and Shen and Kasper (1995). ClustalW was also used to create a multiple sequence alignment between mosquito and rat CPRs.

Phylogenetic Analysis. The deduced amino acid sequence of *A. stephensi* CPR was used in a BlastP search for known homologous CPRs deposited in the GenBank. Subsequently, CPRs representing vertebrates and arthropods with complete open reading frames were selected for phylogenetic analysis. The CPR amino acid sequences were first analyzed using T-Coffee multiple sequence alignment with default parameters (Notredame et al. 2000) to determine variable and conserved regions. A region of low homology at the N-termini was manually removed, and the remaining portion of the sequences was realigned using ClustalW embedded in the MEGA 4.0 software (Tamura et al. 2007). The created alignment file was then used for construction of a neighbor-joining phylogenetic tree. Gaps were eliminated from the algorithmic analysis, and 1,000 replications were performed as a bootstrap test of phylogeny. A cut-off value of 50% was set for the bootstrap consensus tree.

Results

Genomic and Phylogenetic Analysis. The BlastN search, using the nucleotide sequence of the *A. gambiae* CPR gene against the genome sequences in the NCBI GenBank database, identified a specific contig (GenBank ALPR01027092) of the *A. stephensi* genome. This contig was confirmed to contain one CPR gene by comparative genomic analysis. A small fragment of nucleotide sequence (GenBank HM171646) was acquired as a result of a BlastN search against the nonredundant nucleotide sequence. This partial sequence was incorporated in this study to correct a genomic sequencing error. The corrected genomic sequence was then compared with CPR genes previously identified in other mosquito species, which yielded a complete coding region of the *A. stephensi* CPR gene from the initiation codon (ATG) to the stop codon (TAA). The existence and expression of the gene were confirmed by the presence of an expressed sequence tag (EST) (Patil et al. 2009).

The identified *A. stephensi* CPR gene contained an open reading frame of 2,040 bp, coding for a polypeptide of 679 amino acids (Fig. 1).

```

atggacgcccagacagacagcgaaatgccacggcgcaactgagcgacgagcccttctc 60
M D A Q T E T E M P T G N V S D E P F L 20
ggcccggtggacatctctgctcgtcagcctgctggccgacggcgacgctgctg 120
G P L D I I L L L V S L L A G T A W Y L L 40
aaggcgcaagaaagaaatcaagctagctcagttcaatctctactcgatcccaacgacg 180
K K K K E N Q A S Q F K S Y S I Q P T 60
accgtcaaacagcagtgacaatgggtggagaactcgttcaatcaagaagctgcaatctccggt 240
T V N T M T M V E N S F I K K L Q S S G 80
cgccgtttggtggtgttttacggttcccaaacaggcagcgagaggaattgcccgtgct 300
R R L V V F Y G S Q T G T A E E F A G R 100
ctggcgcaaggaagaaatccgctatcagatgaaggcgatggcgccgagcgaggtgctg 360
L A K E G I R Y Q M K G M V A D P E E C 120
aacatggaagaactgctgctggttgaggacatcgacaaatcggtgctggtgtttgctta 420
N M E E L L M L K D I D K S L A V F C L 140
gcccgctacggcaggggtgaccgacggacaattgtatggagttttacgactggtacag 480
A T Y G E G D P T D N C M E F Y D W I Q 160
aacaacgatctgacatgacgggtttgaattacggcgttatgtggcctcggaataagacg 540
N N D L D M T G L N Y A T T G L G N K T 180
tacgagcatctacaacaggtcgccatctcagtggaataagcggtcgagagctggtgctg 600
Y E H Y N K V G I Y V D K R L E L G A 200
aaccgagttgttgagctagagctcgttgacagatgacgacattgagactatctatc 660
N R V F E L G L G D D A A N I E D Y F I 220
acgtggaagaaagttttgccaacgggtgctgcatctactcgccatcgaaagtagcggtc 720
T W K E K F W P T V C D Y F G I E S T G 240
gagatgctgctgacgcccagtcacgtcgtcgtggaagcagcgacgagcgagcgacgcg 780
E D V L M R Q Y R L L E Q P D V S A D R 260
gtgtataccggcaggggtggcagcgctccactcgtcccaaacgacgagcgctccacgttctgat 840
V Y T G E V A R L H S L Q T Q R P P F D 280
gcaaaagaaacccgttctgccaacgatcaaggttaaacgggagctgcacaaagccggcgcc 900
A K N P F L A P I K V N R E L H K A G G 300
cggtctgcatcgacgctgagttcgatatacgaaggtcggaagtcggttagcgagcggtc 960
R S C M H V E F D I E G S K M R Y E A G 320
gaccatctcgcatctgacccggttaacagatcggtgctgagcggtcggtcggaagcgtg 1020
D H L A M Y P V N D R D L V E R L G K L 340
tgcaaacgacgctggaagcggttctctcgtctcaacacggcagcgacgacgacgcaag 1080
C K A D L E T V F S L I N T D T D S S K 360
aagcatccgttccgtgcccacacgctacggacgcccctgacgactacgttgaaata 1140
K H P F P C P T T Y R T A L T H Y L E I 380
acggccctcgccgctaccacatcctgaaggtgtggcggaatactgctcgagagaaag 1200
T A L P R T H I L K E L A E Y C S E E K 400
gacaagagttctcgtggtttatctcgtcgtcagcagcgacgagcggtcgaagctaccag 1260
D K E F L R F I S S T A P E G K V K Y Q 420
gaatgggtcgagcagctgcccgaacatgtgtgacgtgctggagacatccgtctctgt 1320
E W V Q D S C R N I V H V L E D I P S C 440
cagccacgcatcgatcagctgtgagctgtgcccgtttgcaaccccgctactactcc 1380
Q P P I D H V C E L P R L Q P R Y Y S 460
atctctctctcgtccaaagatccatccgacgagcggtgacgtgacggtgctggtacgg 1440
I S S S S K I H P T T V H V T A V L V R 480
tacgaaacgaaacggcgccgctgaacagggcggtggcgacgttctctcgtcggaag 1500
Y E T K T G R L N K G V A T T F L A E K 500
caccggaacgatggggaacgggttacgcgcgtgcccactcttcatcgaagagcgagttc 1560
H P N D G E P L P R V P I F I R K S Q F 520
cgccctgccacggaacgggaaacggcggtgataatggtagggcccggtgacgggttgca 1620
R L P P K P E T P V I M V G P G T G L A 540
ccgttccgtggtctatccagagcggtgactctcgtcgaagcaggggtgaaagacatcgga 1680
P F R G F I Q E R D F C K Q E G K D I G 560
cagacgactctactttggtgcccgaacgggtccgaggtacatcatagcaggaacgag 1740
Q T T L Y F G C R K R S E D Y I Y E D E 580
ctggaagactactccaaggggacatcatcaatctgcccgtggtgctctcgcgcgacag 1800
L E D Y S K R G I N L R V A F S R D Q 600
gacaagaggtgtacgtgacgcatctcgtagagcaagcagcctgacgtatggacgtt 1860
D K K V Y V T H L L E Q D S D L I W N 620
atcgccgaaacagggacacttttcatctcgtggtgatgacaaataatggccacggt 1920
I G E N K G H F Y I C G D A N M A T D 640
gtgcaaacattctgctcaggtcatagcgtcgaagggtgggttaagcgaacacggaagcc 1980
V R N I L L K V I R S K G G L S E T E A 660
cagcagtcacataaaagatggaagcacaagcgatctactcggcgagcgtttggagctaa 2040
Q Q Y I K K M E A Q K R Y S A D V W S * 679

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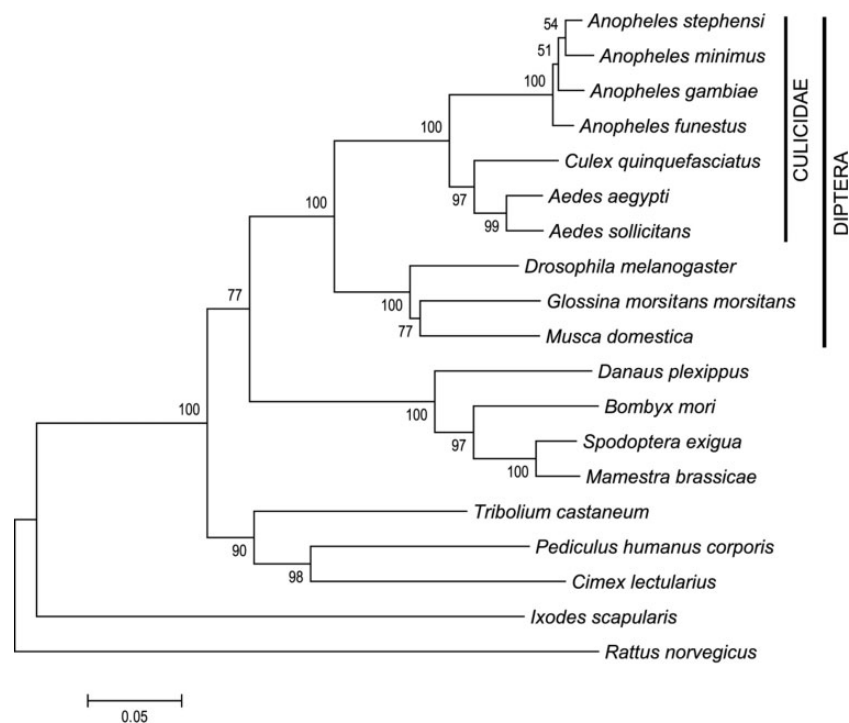
Fig. 1. The predicted sequence of *A. stephensi* CPR transcript. The sequence extends from the initiation codon to the stop codon. The upper and lower lines represent nucleotide and deduced amino acid sequences, respectively. The one-letter code for each amino acid is aligned with the second nucleotide of each codon. The predicted transmembrane region is underlined.

The calculated molecular mass of the deduced amino acid sequence was 77.51 kDa, with a pI of 5.62. The CPR apparently did not contain an N-terminal signal peptide. Instead, a transmembrane helix was found at the N-terminal region, indicating that this was a possible anchored site to the endoplasmic reticulum. The transmembrane hydrophobic region was predicted to be from Phe¹⁹ to Leu⁴⁰ (Fig. 1). The *A. stephensi* CPR showed the highest identity to CPRs in *A. funestus*, *A. gambiae*, and *A. minimus* and a lower identity to CPRs in *Culex* and *Aedes* mosquitoes (Table 1). Evolutionarily, the *A. stephensi* CPR showed higher percentages of identity with dipteran insects than with those in other arthropod groups and rat (Table 1).

Amino acid sequences were used for phylogenetic analysis. A neighbor-joining phylogenetic tree was constructed representing *A. stephensi* CPR along with previously characterized CPRs of

Table 1. Amino acid sequence identity of CPRs between *A. stephensi* and other selected organisms

GenBank accession no.	Species	Common name	Total score	Max identity (%) ^a
ABO77954	<i>A. funestus</i>	African malaria mosquito	1,392	97
AAO24765	<i>A. gambiae</i>	African malaria mosquito	1,386	97
ABL75156	<i>A. minimus</i>	Southeast Asian malaria mosquito	1,378	97
XP_001656715	<i>Ae. aegypti</i>	Yellow fever mosquito	1,273	88
ACL01092	<i>Ae. sollicitans</i>	Eastern saltmarsh mosquito	1,264	87
XP_001865801	<i>C. quinquefasciatus</i>	Southern house mosquito	1,243	87
NP_477158	<i>Drosophila melanogaster</i>	Fruit fly	1,120	78
ADD19306	<i>Glossina morsitans morsitans</i>	Tsetse fly	1,102	77
Q07994	<i>Musca domestica</i>	House fly	1,094	77
XP_971174	<i>Tribolium castaneum</i>	Red flour beetle	998	70
ADX95746	<i>Spodoptera exigua</i>	Beet armyworm	991	69
AAR26515	<i>Mamestra brassicae</i>	Cabbage moth	985	68
NP_001104834	<i>Bombyx mori</i>	Silkworm	978	68
EHJ63867	<i>Danaus plexippus</i>	Monarch butterfly	976	69
XP_002423980	<i>Pediculus humanus corporis</i>	Human body louse	972	67
AFD50507	<i>Cimex lectularius</i>	Bed bug	970	66
XP_002400171	<i>Ixodes scapularis</i>	Black-legged tick	816	59
AAA41683	<i>R. norvegicus</i>	Norway rat	792	55

^aE value = 0.0.**Fig. 2.** A neighbor-joining phylogenetic tree of CPRs in *A. stephensi* and other selected organisms. Gaps were excluded ($n = 1,000$). The branch length represents the accumulation of amino acid changes. The scale indicates 0.05 amino acid change per site. The vertical bars indicate phylogenetic clusters.

mosquitoes and other selected arthropods. A rat CPR was used as an outgroup. As expected, all mosquito CPRs formed a cluster distinctly separated from other organisms (Fig. 2). In particular, CPRs of *A. stephensi* and other *Anopheles* mosquitoes were grouped together within a monophyletic group. The branch representing mosquito CPRs bifurcated from the same node as did the branch that represented other dipteran CPRs. This indicated that they possibly descended from a common ancestor.

The genomic DNA sequences of mosquito CPRs were analyzed for intron–exon organizations. Only species with publicly released genomes were included in this analysis. Based on the region of nucleotides from the initiation codon to the stop codon, seven exons were identified in the mosquito CPR genes (Fig. 3). The sizes of the exons of

the *A. stephensi* CPR gene are generally similar to those of *A. gambiae*, *Ae. aegypti*, and *C. quinquefasciatus*. On the other hand, 15 exons were identified in the rat CPR within the coding region. The sizes of exons 3, 4, and 7 of the *A. stephensi* CPR were identical to exons 4, 5, and 15 of the rat CPR, respectively. There was only a slight difference in exon 1 between the two groups. The rat CPR exons 2 and 3 were comparable with the mosquito CPR exon 2, whereas the rat CPR exons 6–14 corresponded to the mosquito CPR exons 5 and 6. The last exon of the *A. stephensi* CPR appeared to be conserved, when compared with the last exon of the rat CPR (Fig. 4). CPRs in insects representing Diptera, Coleoptera, and Lepidoptera also contained some amino acid residues identical to those in *A. stephensi* and rat within this particular region.

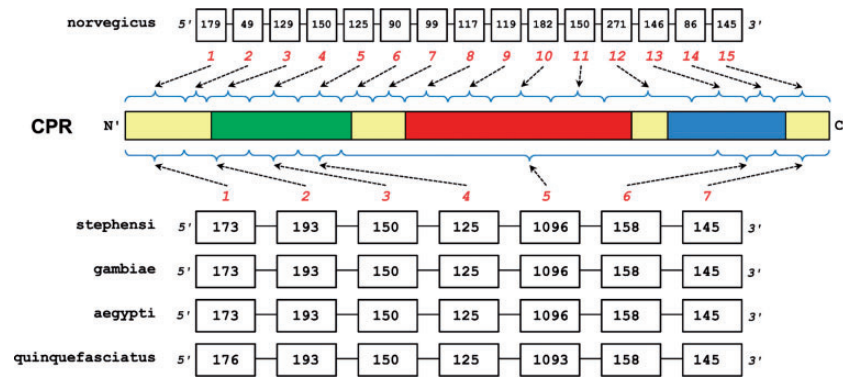


Fig. 3. Intron–exon organization of representative CPR genes. Only mosquitoes with their genomes completed and released to the public were analyzed for gene structure. Rat CPR was included for genetic comparison. The coding regions shown here start from the initiation codon to the stop codon. The square boxes represent exons, and the lines connecting the boxes represent introns. The sizes of both exons and introns are not drawn to scale. The nucleotide length of each exon is presented in each box. Sequential exon numbers (in red) are shown either under (in the case of rat CPR gene) or above (in the case of mosquito CPR genes) the corresponding exons. Binding domains for FMN, FAD, and NADPH are shown in green, red, and blue, respectively.



Fig. 4. Nucleotide and amino acid sequence comparisons of the last exons of the CPR coding regions between *A. stephensi* and other organisms. The one-letter code for each amino acid is aligned with the second nucleotide of each codon. The predicted secondary structures of the *A. stephensi* CPR are shown under the amino acid sequences. The green bar represents the alpha helix, and the blue arrow represents the beta sheet. Identical residues in amino acid sequences are highlighted. Organisms included in the analysis are *R. norvegicus* (representing a vertebrate); *Drosophila melanogaster* and *Musca domestica* (representing Diptera along with *A. stephensi*), *Tribolium castaneum* (representing Coleoptera), and *Spodoptera exigua* (representing Lepidoptera).

Amino Acid Sequence Analysis. A multiple sequence alignment among mosquito CPRs was carried out to determine conserved regions involved in ligand bindings. The *A. stephensi* CPR clearly contained all conserved FMN-, FAD-, and NADPH-binding domains (Fig. 5). Catalytic residues (Ser⁴⁶⁰, Cys⁶³¹, Asp⁶⁷⁶, and Trp⁶⁷⁸) were identified in the mosquito CPR. These residues may be essential in the hydride transfer reaction as previously characterized in the rat CPR (Shen et al. 1999, Hubbard et al. 2001). The FMN-binding domain of the mosquito CPR contained two conserved tyrosine residues (Tyr¹⁴³ and Tyr¹⁸¹) that might be critical in FMN binding (Shen et al. 1989; Fig. 5). In addition, Phe residues at positions 86 and 219 were conserved in all mosquitoes except *A. minimus*, which instead contained Leu at both positions. Phe⁸⁶ and Phe²¹⁹ are involved in FMN binding (Saraputit et al. 2008, 2010).

The FAD- and NADPH-binding domains carry conserved sequence motifs recognized within the ferredoxin reductase structural family. One of the most conserved motifs is Arg-x-Tyr-Ser(Thr), which is found in all members of the family (Dym and Eisenberg 2001). Indeed, ⁴⁵⁷Arg-Tyr-Tyr-Ser⁴⁶⁰ was detected within the FAD-binding domain of the *A. stephensi* CPR. This motif was also conserved in *A. fumestus*, *Ae. aegypti*, *Ae. sollicitans*, and *C. quinquefasciatus*. However, the *A. minimus* and *A. gambiae* CPRs contained Ser and His, respectively, instead of Tyr at the position 459. Another conserved sequence motif within this particular protein family is Met-x-x-x-Gly-Thr(Ser)-Gly(Ala)-Ile-x-Pro, identified in the NADPH-binding domain of all mosquito CPRs (Fig. 5). With one exception, the mosquito CPRs carried Leu instead of Ile in the motif.

One major function of microsomal CPRs is the electron transfer from electron donors to P450s. The interaction between the two

stephensi	MDAQTEEMPTGNVS-DEPFLGPLDIIILLVSLLAGTAWYLLKGKKKENQASQFKSYSIQP	59
minimus	MDAQTEEMPTGVS-DEPFLGPLDIIILLVCLLAGTAWYLLKGKKKENQASQFKSYSIQP	59
funestus	MDAQTEEMPTGNVS-DEPFLGPLDIIILLVSLLAGTAWYLLKGKKKENQASQFKSYSIQP	59
gambiae	MDAQTEVPAGSVS-DEPFLGPLDIVLLVSLLAGTAWYLLKGKKKESQASQFKSYSIQP	59
aegypti	MDAQTEPEVPPAPIS-DEPFLGPLDIIILLAVLIGGAAYFLKSKKKTQTSQFKSYSIQP	59
sollicitans	MDAQTEPEVPPAPVS-DEPFLGPLDIIILLTVLIGGAAYFLKNNKKTQASQFKSYSIQP	59
quinquefasciatus	MDAQTEPEVPPAAATEEPFLGPLDIVLLVLVAGAAWYLLKNNKKEAQTQFKSYSIQP	60
*****:*.*:.*: :*****:* *:*.*:.*:.*: :*****		
stephensi	TTVNTMTMVENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGIRYQMKGMVADPEE	119
minimus	TTVNTMTMVENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGIRYQMKGMVADPEE	119
funestus	TTVNTMTMVENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGIRYQMKGMVADPEE	119
gambiae	TTVNTMTMVENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGIRYQMKGMVADPEE	119
aegypti	TTVNTMTMAENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGLRYQMKGMVADPEE	119
sollicitans	TTVNTMTMAENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGLRYQMKGMVADPEE	119
quinquefasciatus	TAVNTMTMAENSFIKKLQSSGRRLLVYGSQTGTAEAFAGRLAKEGLRYQMKGMVADPEE	120
*:*****.*****:*****:*****:*****:*****:*****		
stephensi	CNMEELMLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNDLDMTGLNYAVFGLGNK	179
minimus	CNMEELMLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNDLDMTGLNYAVFGLGNK	179
funestus	CNMEELMLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNDLDMTGLNYAVFGLGNK	179
gambiae	CNMEELMLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNDLDMTGLNYAVFGLGNK	179
aegypti	CDMEELSLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNVDVDFSLNYAVFGLGNK	179
sollicitans	CDMEELSLKIDKSLAVFCLATGEGDPTDNCMEFYDWIQNNVDVDFSLNYAVFGLGNK	179
quinquefasciatus	CDMEELTLKIDKSLAVFCLATGEGDPTDNCMEFYEWIQNNVDVDFGLNYAVFGLGNK	180
*:*****.*****:*****:*****:*****:*****:*****		
stephensi	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPTVCDYFGIEST	239
minimus	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPTVCDYFGIEST	239
funestus	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPTVCDYFGIEST	239
gambiae	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPTVCDYFGIEST	239
aegypti	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPACDHFIESS	239
sollicitans	TEHYNKVGIYVDKRLEELGANRVFELGLGDDANI EDDYITWKEKFWPACDHFIESS	239
quinquefasciatus	TEHYNKVGIYVDKRLEELGASRVFELGLGDDANI EDDYITWKEKFWPACDHFIEST	240
*****:*****.*****:*****:*****:*****:*****:*****		
stephensi	GEDVLMRQYRLLEQPDVSADRYTGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
minimus	GEDVLMRQYRLLEQPEVGADRIYTGGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
funestus	GEDVLMRQYRLLEQPDVGADRIYTGGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
gambiae	GEDVLMRQYRLLEQPDVSADRIYTGGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
aegypti	GEEVLMRQYRLLEQPETPTERLYTGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
sollicitans	GEDVLMRQYRLLEQPETPAERLYTGEVARLHSLQTRPPFDKPNFLAPIKVNRELHKAG	299
quinquefasciatus	GDEVLRQYRLLEQPETSPERLYTGEVARLHSLQTRPPFDKPNFMAPIKINRELHKG	300
*:** *****:..*:*****:*****:*****:*****:*****		
stephensi	GRSCMHVEFDIEGSKMRYEAGDHLAMPVNDRLVERLGLKCNADLETVFSINTDTS	359
minimus	GRSCMHVEFDIEGSKMRYEAGDHLAMPVNDRLVERLGLKCNADLETVFSINTDTS	359
funestus	DRSCMHVEFDIEGSKMRYEAGDHLAMPVNDRLVERLGLKCNADLETIFSLINTDTS	359
gambiae	GRSCMHVEFDIEGSKMRYEAGDHLAMPVNDRLVERLGLKCNADLETVFSINTDTS	359
aegypti	GRSCMHIEFDIEGSKMRYEAGDHLAMPVNDQDLVRLGLKCNADLDTIFSLINTDTS	359
sollicitans	DRSCMHIEFDIEGSKMRYEAGDHLAMPVNDQDLVRLGLKCNADLDTIFSLINTDTS	359
quinquefasciatus	GRSCMHIEFDIEGSKMRYEAGDHLAMPVNDSDLVRLGLKCNADLDTIFSLINTDTS	360
*****:*****:*****:*****:*****:*****:*****		
stephensi	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFISSTAPEGKVKY	419
minimus	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFISSTAPEGKAKY	419
funestus	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFISSTAPEGKAKY	419
gambiae	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFISSTAPEGKAKY	419
aegypti	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFMCSTNPEGKAKY	419
sollicitans	KKHPFPCTTYRTALTHYLEITALPRTHILKELAEYCSEKDKFLRFMCSTNPEGKAKY	419
quinquefasciatus	KKHPFPCTTYKTALTHYLEITALPRTHILKELAEYCSEKDKFLRFMCSTAPEGKAKY	420
*****:*****:*****:*****:*****:*****:*****		
stephensi	QEWVQDSRNIVHVLEDIPSCPPIDHVCELLPRLQPRYYSISSSSKIHPTTVHVTAVLV	479
minimus	QEWVQDSRNIVHVLEDIPSCPPIDHVCELLPRLQPRYYSISSSSKIHPTTVHVTAVLV	479
funestus	QEWVQDSRNIVHVLEDIPSCPPIDHVCELLPRLQPRYYSISSSSKIHPTTVHVTAVLV	479
gambiae	QEWVQDSRNIVHVLEDIPSCPPIDHVCELLPRLQPRYYSISSSSKIHPTTVHVTAVLV	479
aegypti	QEWVQDSRNIVHVLEDLPSCRPPIDHICELLPRLQPRYYSISSSSKLYPTTVHVTAVLV	479
sollicitans	QEWVQDSRNIVHVLEDLPSCRPPVDHICELLPRLQPRYYSISSSSKLYPTTVHVTAVLV	479
quinquefasciatus	QEWVQDSRNIVHVLEDVPSCHPPIDHICELLPRLQPRYYSISSSSKLYPTTVHVTAVLV	480
:*.*.*:***:*****:*****:*****:*****:*****		
stephensi	RYETKTGRNLKGVATTFLAEKHPNDGEPLPRVPIFIRKSQFRLPPKPETPVI MVGPGTGL	539
minimus	KYETKTGRNLKGVATTFLAEKHPNDGEPLPRVPIFIRKSQFRLPPKPETPVI MVGPGTGL	539
funestus	KYETKTGRNLKGVATTFLAEKHPNDGEPLPRVPIFIRKSQFRLPPKPETPVI MVGPGTGL	539
gambiae	KYETKTGRNLKGVATTFLAEKHPNDGEPLPRVPIFIRKSQFRLPPKPETPVI MVGPGTGL	539
aegypti	KYETKTGRNVNGVATTFLSQKHPLDGEPLPRVPIFIRKSQFRLPAKTETPVI MVGPGTGL	539
sollicitans	KYQTKTGRVNGVATTFLSMKHPLNGEPLPRVPIFIRKSQFRLPAKTETPVI MVGPGTGL	539
quinquefasciatus	KYVTKTGRVNGVATTFLAQKKVN-GESLPRVPIFIRKSQFRLPAKTETPVI MVGPGTGL	539
*:*****.*****:*****:*****:*****:*****:*****		
stephensi	AFFRGFIQERDFCKQEGKDIGQTTLYFGCRKRSEDIYEDELEDYSKRGINLRFVAFSRD	599
minimus	AFFRGFIQERDFSKQEGKDIGQTTLYFGCRKRSEDIYEDELEDYSKRGINLRFVAFSRD	599
funestus	AFFRGFIQERDFSKQEGKDIGQTTLYFGCRKRSEDIYEDELEDYSKRGINLRFVAFSRD	599
gambiae	AFFRGFIQERDHCQEGKDIGQTTLYFGCRKRSEDIYEDELEDYSKRGINLRFVAFSRD	599
aegypti	AFFRGFIQERDFNKKDGKEVGQTTLYFGCRKRSEDIYEDELEDYVQGRIMKLRFAFSRD	599
sollicitans	AFFRGFIQERDFHKKDGKDIGQTTLYFGCRKRSEDIYEDELEDYVQSGTIKLRTAFSRD	599
quinquefasciatus	AFFRGFIQERDFNKKQEGKDIGQTTVMYFGCRKRSEDIYEDELEDYVVKRGVLSLRTAFSRD	599
*****:*****:*****:*****:*****:*****:*****		

Fig. 5. Conserved regions for ligand bindings in the *A. stephensi* CPR. The multiple sequence alignment illustrates FMN-, FAD-, and NADPH-binding domains of mosquito CPRs according to Pfam. The binding domains are boxed (green, FMN; blue, FAD; purple, NADPH). Triangles above amino acids indicate catalytic residues involved in hydride transfer. Essential residues involved in ligand bindings are highlighted.

<i>stephensi</i>	201	NRVFELGLG	DDD	ANI	EDY	FITWKEKFWPTV	230
<i>minimus</i>	201	NRVFELGLG	DDD	ANI	EDY	LITWKEKFWPTV	230
<i>funestus</i>	201	NRVFELGLG	DDD	ANI	EDY	FITWKEKFWPTV	230
<i>gambiae</i>	201	NRVFELGLG	DDD	ANI	EDY	FITWKEKFWPTV	230
<i>aegypti</i>	201	NRVFELGLG	DDD	ANI	EDD	FITWKDKFWPAV	230
<i>sollicitans</i>	201	NRVFELGLG	DDD	ANI	EDD	FITWKDKFWPAV	230
<i>quinquefasciatus</i>	202	SRVFELGLG	DDD	ANI	EDD	FITWKDKFWPAV	231
<i>norvegicus</i>	198	QRIFELGLG	DDD	GNL	EED	FITWREQFWPAV	227

Fig. 6. Conserved residues involved in cytochrome P450 or cytochrome *c* interactions. The multiple sequence alignment shows two clusters of acidic residues of mosquito CPRs in comparison with rat CPR. The clusters are boxed, and identical residues are highlighted.

(²⁰⁷Asp-Asp-Asp²⁰⁹ and ²¹³Glu-Glu-Asp²¹⁵) that are important in the interaction with cytochrome *c*. In comparison with the rat enzyme, two acidic clusters (²¹⁰Asp-Asp-Asp²¹² and ²¹⁶Glu-Asp-Tyr²¹⁸) are found in the *A. stephensi* CPR. The first cluster is entirely identical in the two species, but there are two differences in the second cluster. Specifically, one of the two (Asp²¹⁷) is a conserved substitution that is found in the bacterium *Desulfovibrio vulgaris* Hildenborough flavodoxin (Dubourdieu and Fox 1977). A study based on site-directed mutagenesis in a rat CPR suggests that the first acidic cluster interacts primarily with P450s, whereas the second cluster interacts primarily with cytochrome *c* (Shen and Kasper 1995). It is likely that the *A. stephensi* CPR can perform electron transfers to P450s in the mosquito. Whether the *A. stephensi* enzyme can efficiently transfer electrons to cytochrome *c* will require further investigation. The assumption is supported by activity studies of recombinant CPRs in *A. minimus* (Kaewpa et al. 2007), *A. gambiae* (Lian et al. 2011), and *Ae. sollicitans* (C.S., unpublished data). Although there are similarities between mosquito and vertebrate CPRs, variations in terms of ligand-binding capacities and enzyme kinetics between the two groups are likely (Lian et al. 2011).

Similar to P450s, microsomal CPRs are typically present as integral proteins anchored on the membrane of the endoplasmic reticulum. The anchoring segment of CPRs is a type I signal-anchor sequence with N'-lumen and C'-cytoplasm topology (Kida et al. 1998). The membrane anchoring is also critical for the reductase function. Solubilized reductase without the N'-hydrophobic segment seems to lose its ability to interact with P450s (Black and Coon 1982). As expected, in the case of the *A. stephensi* CPR, the protein does not have a secretion signal peptide but does carry an N'-hydrophobic portion predicted to be a transmembrane region. This suggests that the mosquito CPR can function as part of the electron transfer system to terminal electron acceptors like P450s, which also reside in the microsomal membrane.

In conclusion, analysis of the *A. stephensi* genome reveals a gene encoding for a CPR. The enzyme is highly conserved among mosquitoes and also comparatively similar to CPR enzymes in other organisms. These findings facilitate the understanding of CPR evolution and the role the enzyme plays in electron transfer to its electron acceptor including the P450 family of enzymes, which are often of critical importance in the metabolism of xenobiotic compounds. The results of this study facilitate further elucidation of the mosquito *A. stephensi* CPR functions, including their expected involvement in xenobiotic metabolism.

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