Genetic Polymorphism of *Plasmodium vivax* msp1p, a Paralog of Merozoite Surface Protein 1, from Worldwide Isolates

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Abstract. *Plasmodium vivax* msp1p, a paralog of the candidate vaccine antigen *P. vivax* merozoite surface protein 1, possesses a signal peptide at its N-terminus and two epidermal growth factor–like domains at its C-terminus with a glycosylphosphatidylinositol attachment site. The msp1p gene locus may have originated by a duplication of the msp1 gene locus in a common ancestor of the analyzed *Plasmodium* species and lost from *P. yoelii, P. berghei*, and *P. falciparum* during their evolutionary history. Full-length sequences of the msp1p gene were generally highly conserved; they had a few amino acid substitutions, one highly polymorphic E/Q-rich region, and a single-to-triple hepta-peptide repeat motif. Twenty-one distinguishable allelic types (A1–A21) of the E/Q-rich region were identified from worldwide isolates. Among them, four types were detected in isolates from South Korea. The length polymorphism of the E/Q-rich region might be useful as a genetic marker for population structure studies in malaria-endemic areas.

INTRODUCTION

Among the species of malarial parasite that infect humans, *Plasmodium vivax* is the most globally prevalent and threatens almost 40% of the world’s population, resulting in approximately 250 million clinical infections each year.† Although *P. vivax* malaria had been considered relatively benign, compared with that of *P. falciparum*, this view is now being challenged. Additionally, resistance to chloroquine is appearing in countries where malaria is endemic.‡ Thus, there are good reasons to pursue an effective *P. vivax* vaccine. In this regard and in contrast to *P. falciparum*, research into *P. vivax* is limited, due in part to difficulties in culturing blood-stage parasites *in vitro*.‡ Nevertheless, several *P. vivax* vaccine candidates from different parasitic stages have been characterized.§ Among them, various merozoite surface proteins (MSPs), apical membrane protein-1, Duffy binding protein, Pvs25, Pvs28, circumsporozoite protein, and thrombospordin-like anonymous proteins have been studied.§ Merozoite surface proteins have been characterized and are highly immunogenic in natural infection.δ–η Among them, several major vaccine candidate antigens (including MSP1, MSP4, MSP5, MSP8, and MSP10) are either known or presumed glycosylphosphatidylinositol–anchored membrane proteins. *Plasmodium vivax* MSP1 is the largest and most abundant protein on the *P. vivax* merozoite surface.†,§,10 The gene that encodes this protein (*Pvmmsp1*) is highly polymorphic and consists of a mosaic of conserved and variable blocks with numerous recombination sites distributed throughout the gene. However, the fragment that encodes the 19-kDa C-terminal epidermal growth factor (EGF)–like domain is relatively conserved.11 This gene has been used as a polymorphic marker for investigations of the genetic structure of *P. vivax* populations and in molecular epidemiology.12

With the completion of *P. vivax* genome sequencing, GPI-anchored proteins of *P. vivax* have been predicted by comparison with validated *P. falciparum* GPI-anchored proteins.13,14 *Plasmodium vivax* msp1p (*Pvmmsp1p*), a novel paralog of the *Pvmmsp1* gene, was found immediately upstream of *Pvmmsp1*.14 This gene is predicted to encode a 1,854-amino-acid protein (predicted molecular mass of 215 kDa) with an N-terminal signal sequence, C-terminal EGF-like domains, and a GPI-attachment motif (Figure 1).† The functions of this molecule remain unknown. Thus, we have analyzed available genomic data from Plasmodb (http://www.plasmodb.org/) to search for distinctive pattern of diversity in *msp1p* and *msp1* genes among *Plasmodium* species. We have also assessed the nature and extent of polymorphisms in *Pvmmsp1p* from worldwide isolates and laboratory lines of *P. vivax*.

MATERIALS AND METHODS

Gene sequences. The following sequences of malarial parasites were used for the analyses: human *P. falciparum* *Pfmmsp1* (CAA27070), *Pfmmsp8* (PFE0120c), and *Pfmmsp10* (PFF0995c), and *P. vivax* *Pvmmsp1* (*PVX_099980*), *Pvmmsp8* (*PVX_097625*), *Pvmmsp10* (*PVX_114145*), and *Pvmmsp1* (*PVX_099975*); rodent malaria *P. berghei* *Pbmmsp1* (AAC28871), *Pbmmsp8* (BPANKA_110220), and *Pbmmsp10* (BPANKA_111960); *P. yoelii* *Pymmsp1* (PY05748), and *P. chabaudi* *Pchmmsp1* (PCAS_083080); primate malaria *P. knowlesi* *Pkmsmp1* (PKH_072850) and *Pkmsmp1p* (PKH_072840), *P. reichenowi* *Prmmsp1* (CAHI0285), and *P. cynomolgi* *Pcymsmp1* (BAI82251); and avian malaria *P. gallinaceum* *Pgmsmp1* (CAHI0838). The *P. gallinaceum* sequence database (http://www.sanger.ac.uk/) was used to search for homologs of *Pvmmsp1p* and *Pvmmsp10*.

Blood samples and DNA preparation. Blood samples were collected, after informed consent had been obtained, from 81

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symptomatic patients diagnosed by microscopic examination with *P. vivax* infection at Korea University Ansan Hospital, local health centers, and clinics in Gyeonggi and Gangwon provinces, Republic of Korea. Genomic DNA was purified from 200 μL of whole blood by using a QIAamp DNA Blood Mini Kit (QIAGEN, Valencia, CA), according to the manufacturer’s protocol. Genomic DNA of *P. vivax* isolates (n = 33) obtained from Thailand (n = 26), Indonesia (n = 3), India (n = 1), Papua New Guinea (n = 1), Western Samoa (n = 1), and Pakistan (n = 1), and nine *P. vivax* laboratory lines (Africa Mauritania, New Guinea, Honduras III, Brazil I, Salvador I, Vietnam IV, Indonesia I, India VII, and Columbia Rio Meta) were used for polymerase chain reaction (PCR).

**Amplification and sequencing of target genes.** Genomic DNA from 20 isolates from the Republic of Korea and 9 isolates from other locations was used for amplification of the *Pvmsp1p* full-length gene. Primers SeqF1 (5′-TGC ATA TTC ATA ATG CAG GAG AAG AAA AAA ATG TAC C-3′) and SeqR8 (5′-GGT GAG CTA ATC GAA CCG G-3′) were designed based on the *Pvmsp1p* sequence of the *P. vivax* Sal I strain. These were used to amplify DNA fragments from 90 basepairs upstream to 80 basepairs downstream of the *Pvmsp1p* coding sequence by using LA Taq DNA polymerase (TaKaRa, Tokyo, Japan). The PCR amplification was performed on a MyCycler Thermal Cycler (Bio-Rad, Hercules, CA) by using the following temperature profile: 94°C for 2 minutes; 35 cycles at 94°C for 30 seconds, 64°C for 30 seconds, 72°C for 6 minutes; and a final extension at 72°C for 10 minutes.

Both strands of the PCR products were directly sequenced by using a series of sequencing forward primers (SeqF1; SeqF2: 5′-ATC AAC CGG AAG AAC TAC CT C-3′; SeqF3: 5′-CAA AGG GAG AAG AAA AAA ATG TAC C-3′; SeqF4: 5′-GGT GAG CTA ATC GAA CCG G-3′; SeqF5: 5′-TGG GGC GCA CAT AAC CT C-3′; SeqF6: 5′-CCC GTG TAC TCT AAC CAG GAT GTG ATG AG C-3′; SeqF7: 5′-TGA AGT GCA ACA CGT GGA AT C-3′; SeqF8: 5′-GTG GAC TACTAC GGG CTA AGG A-3′ and SeqF9: 5′-ATT TTC TCT TAT GCA GAC AAG AAG GAG GTG-3′) and reverse primers (SeqR1: 5′-AAG GCA GGA TTA GAG ACG C-3′; SeqR2: 5′-CGC ACG TTG AGG TGG TAG TC C-3′; SeqR3: 5′-AGC GTC AAG ATG TGG TGG TGG CAG-3′; SeqR4: 5′-TGC GTG ATG ATC TGG CTT GGT ATC AGC-3′; SeqR5: 5′-TCC CCG ATG AAA TAT GC C-3′; SeqR6: 5′-ACT GCA GAT GGA TGG TCA TCT C-3′; SeqR7: 5′-AAC TGC ATC GGC TGG TCA GTA T-3′; and SeqR8) using an ABI Prism 377 DNA sequencer (Genetech, Seoul, South Korea).

Analysis of the full-length gene sequences showed one highly polymorphic region. This region was amplified by PCR from genomic DNA of 61 samples form the Republic of Korea and 24 samples from other locations, as well as nine laboratory strains, and sequenced. To examine variable tandem repeat regions, as found in the *Pvmsp1p* gene sequence, primers (TR-F: 5′-CCT ACA AGG GTT GGG AGA T-3′ and TR-R: 5′-CGG AGA CGG AGT TCG TGA T-3′) were used to amplify a 200-basepair fragment encompassing this region from 33 worldwide isolates and 9 laboratory lines.

**Data analysis.** Sequence data were submitted to GenBank under accession numbers GU556592–GU556620. Amino acid sequence alignments were constructed using the MUSCLE program, with manual corrections. The number of nonsynonymous substitutions per nonsynonymous site (dN) and the number of synonymous substitutions per synonymous site (dS) were computed by using the Nei-Gojobori method with the Jones-Cantor correction, as implemented in the MEGA 4 program. An unrooted tree was constructed by the neighbor-joining method with the Jones-Taylor-Thornton amino acid substitution model, accompanied by bootstrap analysis with 1,000 replicates for the neighbor-joining method and 100 for the maximum parsimony method implemented in PHYLIP version 3.68 after excluding insertions/deletions (indels) and unreliable amino acid sites.

**RESULTS**

An MSP1P homolog can be found in the *P. vivax* and *P. knowlesi* genome databases, but not in the *P. falciparum*, *P. yoelii*, or *P. berghei* genome databases. To investigate the evolutionary relationship of MSP1P with other MSPs, we searched for *Pvmsp1p* homologs in the available *Plasmodium* genome database and analyzed their relationship with MSP1 by using the distantly related MSP8 and MSP10 sequences as outgroups. A TBLASTN search of the *P. gallinaceum* sequence database (http://www.sanger.ac.uk) was conducted by using the *PvMSP1P* amino acid sequence as a query. This search identified a contig (28a.d000066175.Contig1) that contained a putative partial sequence of the *Pgmsp1p* gene (encoding the C-terminal end included EGF-like domains) (Figure 2). Based on the BLOSUM matrix, the amino acid sequence identity/similarity of the EGF-like domains to those of *PvMSP1P* and *PkMSP1P* were 56/71% and 58/75%, respectively (Figure 3A). The identity/similarity of the N-terminal region of this gene sequence of the *P. vivax* merozoite surface protein 1 paralog (PvMSP1P), *P. knowlesi* merozoite surface protein 1 paralog (PkMSP1P), and *P. gallinaceum* merozoite surface protein 1 paralog (PgMSP1P). Dashes indicate deletions. Cys residues with light areas indicate Cys residues conserved among all sequences and those with dark areas and the arrowhead indicate the additional two Cys residues conserved among MSP8, MSP10, MSP1P, and Pf/Msp1P (Figure 3). Asterisks, dots, and colons under the alignment indicate identical, conserved, and semi-conserved substitutions, respectively, based on BLOSUM. The glycosylphosphatidylinositol (GPI) modification site is indicated with arrowhead.
Second, beside the EGF-like domains, N-terminal side of the PgMSP1P and was separated from the MSP1 clade (Figure 3B). cated that PvMSP1P formed one clade with PkMSP1P and sons. First, the dendrogram using EGF-like domains indicated that PvMSP1 might be evolutionarily closer to PfMSP1 than PvMSP1P. However, this appears not to be so for three rea-sons. First, the dendrogram using EGF-like domains indicated that PvMSP1 formed one clade with PkMSP1P and PgMSP1P and was separated from the MSP1 clade (Figure 3B).

Figure 3. Relationship between Plasmodium vivax merozoite surface protein 1 paralog (MSP1P) and other Plasmodium merozoite surface proteins possessing epidermal growth factor (EGF)-like domains. A. Amino acid sequence alignment of the EGF-like domains of Plasmodium MSP1, MSP8, MSP10, and MSP1P. Dashes indicate a deletion. Cys residues with light areas indicate Cys residues conserved among all sequences and those with dark areas masks and the arrowhead indicate the additional two Cys residues conserved among MSP8, MSP10, and Pk/Pf/PgMSP1. Asterisks, dots, and colons under the alignment indicate identical, conserved, and semi-conserved substitutions, respectively, based on BLOSUM. B. Unrooted dendrograms of the EGF-like region of MSP1, MSP8, MSP10, and MSP1P amino acid sequences. Trees were constructed by the neighbor-joining and maximum parsimony methods using amino acid positions 1759, 1782, 1783, 1784, 1785, 1786, 1787, 1812, 1813, 1814, 1815, 1822, 1823, and 1824 (after P. vivax MSP1P amino acid sequence) after excluding indel and unreliable sites. Numbers on branches indicate bootstrap values. C. Schematic diagram of the proposed evolutionary history of the msp1P gene locus in Plasmodium spp. The msp1P gene locus was generated by duplication of the msp1P gene locus in the common ancestor of known Plasmodium species. This locus was then deleted in P. yoelii, P. berghei, and P. falciparum. Sequences of P. falciparum PmMSP1 (CA27070), PmMSP6 (PTF0120c) and PmMSP10 (PFF0995c); P. vivax PmMSP1 (PVX_099980), PmMSP8 (PVX_099975), PmMSP10 (PVX_114145); P. knowlesi KmMSP1 (PKH_072850) and PkMSP1P (PKH_072840); P. berghei PmMSP1 (AAC28871), PmMSP6 (PBANKA_110220) and PmMSP10 (PBANKA_111960); P. reiche-nowi PmMSP1 (CAH10285); P. gallinaceum PgMSP1 (CAH10838), PgMSP1P (encoded in 28a.d000006175.Contig1), PgMSP10 (encoded in 28a.d000005716.Contig1); P. cynomolgi PyMSP1 (BAI82251); P. yoelii PyMSP1 (PY05748); and P. chabaudi PchMSP1 (PCAS_083080) were used.

A product to the corresponding region of PvMSP1P (amino acid positions 1,402–1,675) and PkMSP1P (1,425–1,699) were 37/60% and 39/61%, respectively (Figure 2), where the similarity with PgMSP1 was less than 30%. This gene product to the corresponding region of PvMSP1P (amino acid positions 1759, 1782, 1783, 1784, 1785, 1786, 1787, 1812, 1813, 1814, 1815, 1822, 1823, and 1824 (after P. vivax MSP1P amino acid sequence) after excluding indel and unreliable sites. Numbers on branches indicate bootstrap values. C. Schematic diagram of the proposed evolutionary history of the msp1P gene locus in Plasmodium spp. The msp1P gene locus was generated by duplication of the msp1P gene locus in the common ancestor of known Plasmodium species. This locus was then deleted in P. yoelii, P. berghei, and P. falciparum. Sequences of P. falciparum PmMSP1 (CA27070), PmMSP6 (PTF0120c) and PmMSP10 (PFF0995c); P. vivax PmMSP1 (PVX_099980), PmMSP8 (PVX_099975), PmMSP10 (PVX_114145); P. knowlesi KmMSP1 (PKH_072850) and PkMSP1P (PKH_072840); P. berghei PmMSP1 (AAC28871), PmMSP6 (PBANKA_110220) and PmMSP10 (PBANKA_111960); P. reiche-nowi PmMSP1 (CAH10285); P. gallinaceum PgMSP1 (CAH10838), PgMSP1P (encoded in 28a.d000006175.Contig1), PgMSP10 (encoded in 28a.d000005716.Contig1); P. cynomolgi PyMSP1 (BAI82251); P. yoelii PyMSP1 (PY05748); and P. chabaudi PchMSP1 (PCAS_083080) were used.

In a previous study, Carlton and others reported that the PvMSP1P EGF-like domains contained extra two Cys residues in both P. vivax and P. knowlesi, the lack of a msp1P homolog in the P. yoelii and P. berghei genomes is likely caused by deletion event prior to diversification of these parasite species. Because rodent malaria parasite species form a single clade with P. vivax and P. knowlesi, the lack of a msp1P homolog in the P. yoelii and P. berghei genomes is likely caused by deletion of the msp1P gene locus during their evolution. Deletion of this gene locus may also have occurred in P. falciparum (Figure 2C).

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Of these substitutions, only the S755I or P1686T (or both) mutations were found in 15 of the isolates from the Republic of Korea and Papua New Guinea, whereas more mutants were found in the worldwide isolates (Table 1).

A short and highly diverse region, composed of Glu and Gln as 3–5 Glu residues, followed by one or several basic E/Q residues (n = 1–6) units, was found in the Pvmsp1p gene. Twenty-one distinguishable allelic types (A1–A21) were identified in 127 isolates (clones), based on a comparison with corresponding regions in the P. vivax Sal I strain (Table 2). Type A1 had an identical sequence to that of the Sal I strain, which was found in only two laboratory strains, from Central and South America. Type A2 predominated (33.1%, 42 of 127) in all Pvmsp1p samples, and in the Korean (35.8%, 29 of 81), Thai (26.7%, 8 of 30), Pacific (67%, 2 of 3), and African isolates (100%, 1 of 1), which share 96.7% amino acid identity with type A1. The 81 Korean isolates appeared to have limited diversity because only four genotypes (allelic types A2, A7, A20, A21) were found, whereas isolates from other locations worldwide, and laboratory strains, showed 20 allelic types (the exception being A21). Interestingly, type A21 was detected only in Korean isolates (18.5%, 15 of 81).

Polymerase chain reaction amplification resulted in two or three target bands in each of three Thai isolates, which suggested multi-clone infection. To confirm this finding, PCR products of the polymorphic region amplified from these samples was cloned and sequenced. Three types (A2, A16, A20) were detected from the Thai T21 isolate, two (A7, A15) from isolate T25, and two (A19, A20) from isolate T29.

**DISCUSSION**

We have assessed the evolutionary relationship of the msp1p gene with other msp genes and propose that a duplication event (mspl and msp1p) occurred before the diversification of the clades in *P. vivax* and *P. gallinaceum*. This account requires two independent deletions of msplp, one in the rodent lineage (after its divergence from the primate lineage to *P. knowlesi* and *P. vivax*) and another deletion in the lineage to *P. falciparum* (after its divergence from the primate lineage to *P. knowlesi* and *P. vivax*). We also propose that the common ancestor of *P. vivax*, *P. knowlesi*, *P. yoelii*, and *P. berghei* possessed MSP1 that had 12 Cys residues in the first EGF-like domain, and that two Cys sites were substituted to other amino acids during their evolution. We further investigated the genetic diversity of the Pvmsp1p gene in isolates from locations worldwide, including the Republic of Korea. We found an E/O-rich polymorphic region, a hepta repeat region, and several polymorphic sites. However, no diversifying selection was apparent by comparing dN and dS. Although the molecular data (e.g., size, molecular mass, number, location of Cys residue) were similar to those of PvmSP1, PvmSP1P is not polymorphic and appears to not be under noticeable host immune pressure. However, the repeat-length polymorphism of the E/G-rich region may prove useful as a genetic marker for epidemiologic studies.

High conservation of the double EGF-like domains was also detected in other merozoite surface proteins, such as MSP1 and MSP4. These are involved in putative ligand-receptor interactions during erythrocyte invasion by merozoites. Thus, the lack of variation in the C-terminus sequence of PvmSP1P, especially the high conservation of the double EGF-like domains, suggests that these regions play an important role in this process.

The overall nucleotide diversity of Pvmsp1p is much lower than that of other *P. vivax* antigens, such as MSP1, MSP3β, and apical membrane antigen 1. In the PvmSP1P sequences, the E/O-rich region was shown to be highly polymorphic (21 allelic types in 127 clones/isolates). In the cases of PvmSP1 and Pfs230 (AF269242), the E/Q-rich region was also highly polymorphic and represented the principal source of genetic diversity.

In a low-complexity region analysis of *Plasmodium*, Gln appeared with a somewhat higher frequency in the repetitive than in the non-repetitive motifs. The E/O-rich regions and repeat motif of PvmSP1P and Pfs230 were located in a low-complexity region. These low-complexity regions harbor tandem repeats identified in *Plasmodium* and correspond to species-specific and rapidly diverging regions.

This variation in E/O-rich regions and the number of repeats could be generated by slipped-strand mispairing mechanisms. These result in duplication, deletion, or mutation of certain repeat units. The tandem repeat regions of PvmSP1P may result from rapid diversification, which enables the parasite to evade the immune response of the host by antigenic polymorphism.

Finally, the highly polymorphic E/O-rich region sequence of PvmSP1P might be useful as a genetic marker for studies on the population structure and dynamics of *P. vivax* in malaria-endemic areas.
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| A1           | ·——·——··——·——·——··——··——·——·——·——·——·——·——·——·——··——·——··——·——·——·——·——·——·——·——·——·——·——·——··——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·——·—
REFERENCES


