MINI REVIEW

Genetic Diversity of *Plasmodium vivax* Causing Epidemic Malaria in the Republic of Korea

Young Yil Bahk¹, Jeonga Kim², Seong Kyu Ahn³, Byoung-Kuk Na⁴, Jong-Yil Chai⁵*, Tong-Soo Kim⁵*

¹Department of Biotechnology, College of Biomedical and Health Science, Konkuk University, Chungju 27478, Korea; ²Department of Medicine, Division of Endocrinology, Diabetes, and Metabolism, UAB Comprehensive Diabetes Center, University of Alabama at Birmingham, Birmingham, AL 35294, USA; ³Department of Parasitology and Tropical Medicine, Inha University School of Medicine, Incheon 22212, Korea; ⁴Department of Parasitology and Tropical Medicine and Institute of Health Sciences, Gyeongsang National University College of Medicine, Jinju 52727, Korea; ⁵Korea Association of Health Promotion, Seoul 07853, Korea

Abstract: *Plasmodium vivax* is more challenging to control and eliminate than *P. falciparum* due to its more asymptomatic infections with low parasite densities making diagnosis more difficult, in addition to its unique biological characteristics. The potential re-introduction of incidence cases, either through borders or via human migrations, is another major hurdle to sustained control and elimination. The Republic of Korea has experienced re-emergence of vivax malaria in 1993 but is one of the 32 malaria-eliminating countries to-date. Despite achieving successful nationwide control and elimination of vivax malaria, the evolutionary characteristics of vivax malaria isolates in the Republic of Korea have not been fully understood. In this review, we present an overview of the genetic variability of such isolates to increase understanding of the epidemiology, diversity, and dynamics of vivax populations in the Republic of Korea.

Key words: Genetic diversity, vivax population, Korea

INTRODUCTION

The Republic of Korea has recently made significant progress in eliminating the burden of vivax malaria since the re-emergence of infectious vivax malaria in 1993, and has been categorized as one of 32 malaria-eliminating countries [1]. Malaria is an infectious disease caused by the *Plasmodium* parasite which is transmitted via the bite of infected female *anopheline* mosquitoes. Of the 4 *Plasmodium* species comprising *P. falciparum*, *P. vivax*, *P. malariae*, and *P. ovale* which routinely cause malaria in humans, *P. vivax*, a relatively neglected human malaria parasite, is the most widely distributed species in humans across the world causing an estimated 80-90 million cases each year [2]. Among the protozoa that cause malaria, *P. falciparum* is the greatest menace because of its high rate of mortality, mostly among children under the age of 5 in Africa. However, malaria caused by *P. vivax* is less lethal but has accounted for an enormous share of global malaria burden in recent years along with *P. falciparum*, greatly affecting the quality of life of several human populations in tropical, subtropical, and temperate countries [3,4]. *P. vivax* has evolved into a strain group with highly unique biological features, including a hypnozoite stage as a dormant and relapsing stage in the liver and a preference for invading reticulocytes (Duffy red blood cells) in as little as 0.5-1.5% normal blood; the lack of a convenient and continuous in vitro culture system for blood stages is an additional challenge [5]. In general, vivax malaria was thought to be benign and self-limiting, was not considered a priority in comparison with the significant burden of morbidity and mortality presented by *P. falciparum*, and was found to be harder to eliminate than *P. falciparum* [1,6,7]. However, the most geographically widespread distribution of *P. vivax* and recent evidence of severe and complicated vivax malaria across several endemic regions of the world suggest that this disease may have been overlooked as benign [8].

In the Republic of Korea, *P. vivax* is the most prevalent malaria species [9]. Vivax malaria has been endemic in the Republic of Korea for more than 2 decades after re-emergence. Since its re-emergence in 1993 [10], it has accounted for a total of 33,972 cases reported from 1993 to 2017 [9]. Ascertaining the *P. vivax* population structure is essential prior to establishing goals for control, elimination, and the rollout of interven-
tions, and detailed knowledge of the spatial distribution, transmission, and clinical burden of *P. vivax* is required to act as a benchmark against which control targets can be set and measured [11-13]. The majority of studies on the genetic diversity of *Plasmodium* spp. have been based on *P. falciparum*, which is responsible for the most severe disease form, and on genes coding for antigenic determinants such as the circumsporozoite surface protein (CSP) and the merozoite surface protein (MSP). These antigenic genes are non-synonymous nucleotide polymorphisms and the multiple allelic forms differ in their ability to abrogate recognition by the host’s immune response [14]. At the time of re-emergence, no heterogeneity was observed in malaria incidence or in the distribution of *P. vivax* cases across the Republic of Korea. In July 1993, a Korean Army soldier stationed near the DMZ in northern Gyeonggi Province was diagnosed with vivax malaria [15]. After this first re-emergence case, there was an exponential increase in vivax malaria cases with epidemic outbreaks from 1995 to 2000. Since then, vivax malaria has been recognized as a significant public health problem [16-18]. In this short review article, we will discuss the characteristics of the well-known polymorphisms in the genes from South Korean isolates of *P. vivax*. The genetic diversity of the *Plasmodium* spp. population can be a valuable resource to analyze parasite populations. The degree of genetic diversity and the distribution of that diversity, or its population structure, offer insight into drifts in parasite transmission and parasite population history, information that is pivotal to malaria control and elimination programs and assessing the impact of such interventions [19].

**GENETIC DIVERSITY OF WELL-CHARACTERIZED POLYMORPHIC REGIONS IN *P. VIVAX* ISOLATES FROM THE REPUBLIC OF KOREA**

Because the high levels of polymorphism in *Plasmodium* parasites indicate the sophisticated immune evasion strategies of malaria parasites [20], understanding parasite population diversity is important for the strategic development and deployment of control measures [21,24]. Moreover, genetic diversity based on multiple gene polymorphisms in *vivax* parasites has been shown to greatly contribute to drug resistance [25]. Due to polyclonal infection of the same host by multiple parasite clones, to elucidate local population structure and genetic diversity, it is essential to be able to distinguish between distinct clones as well as between infections. In addition, *P. vivax* in the Republic of Korea displays relatively low parasitemia and low male-to-female gametocyte ratio, contributing to low transmission efficacy [26,27]. Further, the frequency of the polyclonal infection rate in the Republic of Korea is relatively very low (~3% vs 30-65% in Papua New Guinea, Thailand, and India) [28]. These unique biological characteristics are not well-understood. In fact, it is well-known that *P. vivax* displays much greater genetic diversity in terms of SNP diversity and microsatellite and gene family variability than *P. falciparum* [29], and this means that the global population of *P. vivax* may have a capacity for greater functional variation [29]. Understanding the genetic variation patterns of the parasite may help in analyzing trends and assessing the extent of malaria endemicity in the Republic of Korea [30]. This knowledge of the extent of correlation between the parasite population diversity and transmission intensity could also provide early warnings to identify populations living in endemic areas, facilitate the identification of new drug resistance genes, and eventually provide information essential for malaria control and elimination [31]. The well-characterized polymorphic regions in genes such as *gam1* coding for the gametocyte antigen 1, *csp* coding for the circumsporozoite protein (CSP), *msp1* and *msp3α* coding for the merozoite surface proteins (MSP)-1 and 3α, respectively, *ama1* coding for the apical membrane antigen (AMA) 1, and *dbp* coding for the Duffy binding protein (DBP) have been extensively used to analyze genetic diversity patterns of natural populations of *P. vivax* (Table 1) [32,33].

The genetic diversity of South Korean *P. vivax* isolates has been extensively studied and characterized by several research groups. The population structure of *P. vivax* parasites in the Republic of Korea has been largely based on genetic analyses focusing on the mutation of antigen-coding genes. Initially, a combinational analysis of *PvMSP-1* and *PvMSP-3α* clearly demonstrated the drastic genetic changes occurring in the South Korean *P. vivax* population. The MSP-1 and 3α are blood-stage vaccine-candidate antigens with repetitive arrays. In particular, *PvMSP-3α* is highly polymorphic and has been confirmed to be a molecular marker in several geographic malaria parasite populations [32]. From early re-emergence to 2000, *PvDBP, PvMSP-1*, and *PvMSP-3α* genotypes were found to be highly homogenous. However, since 2001, they have rapidly diversified. Based on the investigation of 30 samples from *P. vivax* patients in 1998, it was found that *PvDBP* had little polymorphism and only 2 genotypes of *P. vivax* coexisted
in the Republic of Korea [34]. *PvDBP* is involved in the formation of an irreversible junction between *P. vivax* merozoites and its receptor, the Duffy antigen/receptor for chemokines at the surface of immature reticulocytes, and is a prime blood stage vaccine candidate antigen for *P. vivax*. *PvMSP-1* in South Korean isolates analyzed from blood samples collected in 1996 and 1997 was reported to be highly homogenous [35]. The *PvMSP-3* locus of re-emerging Korean *P. vivax* is dimorphic with 2 allelic types coexisting in the endemic area [36]. In addition, 2 genotypes of *PvAMA-1* genes coexist in the re-emerging South Korean *P. vivax* [37], suggesting that 2 genotypes of *P. vivax* coexist in the endemic area of the Republic of Korea. *PvAMA-1* is an immunogenic, type I integral membrane protein, which is expressed at the apical surface of merozoites and sporozoites. Because of the geographical linkage of the genetic variation of these genes in *Plasmodium* spp, *P. vivax* found in the Republic of Korea was unique and, based on molecular structure analysis, could be divided into 2 strains [38]. Of course, the molecular basis for the geographic linkage is poorly understood to-date. Based on the analysis of nucleotide diversity across the entire *PvAMA-1* gene, South Korean *PvAMA-1* was found to have limited diversity compared to global isolates and differed from other global isolates in its pattern of genetic polymorphism [39]. In case of polymorphism in GAM-1, a *P. vivax* transmission-blocking candidate antigen [40] no longer used as a reliable genetic marker [32], Korean isolates from 30 patients in 1998 showed only one genotype containing 4 33-bp motifs identical to the Belem strain [41]. However, an analysis of South Korean isolates from 1996-2007 based on the sequence analysis of *PvMSP-1* and *PvCSP* genes revealed that

### Table 1. Genes encoding antigens used as molecular markers to study field populations of *Plasmodium vivax* in the Republic of Korea

| Antigen    | Total sample No. | Protein fragments                   | No. of subtypes (Years of blood collection) | References |
|------------|------------------|------------------------------------|---------------------------------------------|------------|
| *PvAMA-1*  | 22               | Domain I                           | 2 (1999-2000)                               | Han et al. (2002) [37] |
|            | 30               | Domain I                           | 2 (1998)                                    | Chung et al. (2003) [73] |
|            | 66               | Domain I, III                      | 30 (2009-2011)                              | Kang et al. (2015) [39] |
| *PvCSP*    | 21               | Tandem- and post repeat region      | 2 (1995-2000)                               | Kim et al. (2002) [73] |
|            | 30               | Post-repeat region                  | 2 (1997)                                    | Kho et al. (1999) [74] |
|            | 632              | Central repeating domain           | 7 (1996-2007)                               | Choi et al. (2010) [42] |
|            | 121              | Central repeating domain           | 2 (1998-2001)                               | Nam et al. (2014) [44] |
| *PvDBP*    | 8                | Domain II, III                     | 1 (1997)                                    | Suh et al. (2001) [38] |
|            | 30               | Domain II, IV                      | 2 (1998)                                    | Kho et al. (2001) [34] |
|            | 70               | N-terminal Cys-rich region         | 13 (2005-2010)                              | Ju et al. (2013) [46] |
| *PvMSP-1*  | 25               | Interspecies conserved blocks 5, 6 | 1 (1966-1997)                               | Lim et al. (2000) [35] |
|            | 30               | Interspecies conserved blocks 4, 5 | 2 (1998)                                    | Kim et al. (2009) [75] |
|            | 632              | Interspecies conserved blocks 5, 6 | 6 (1996-2007)                               | Choi et al. (2010) [42] |
|            | 45               | Whole                              | 2 (1996-2000)                               | Han et al. (2011) [76] |
|            | 255              | Interspecies conserved blocks 5, 6 | 2 (1998-2001)                               | Kang et al. (2016) [45] |
|            | 149              | C-terminal 42 kDa region           | 11 (1999-2010)                              | Kang et al. (2012) [77] |
| *PvMSP-3*  | 24               | No applied RFLP-PCR                | 2 (1996)                                    | Han et al. (2004) [36] |
|            | 255              | Blocks I & II                      | 3 (1998-2001)                               | Kang et al. (2016) [45] |
| *PvWtr*    | 85               | Wtr27, Wtr21, Wtr12, Wtr4          | 10.25 (2011-2013)*                          | Son et al. (2017) [78] |
| *PvGAM-1*  | 30               | Nt3792-4029 of *PvGAM-1*           | 1 (1998)                                    | Kho et al. (2001) [41] |
| *PvMdr-1*  | 55               | SNPs of *PvMdr-1*                  | 4 (2011-2012)                               | Chung et al. (2015) [61] |
| *PvTARAg55*| 42               | Exon 2 of *PvTARAg55*              | 4 (2007-2012)                               | Jang et al. (2015) [79] |
| *PvS25 & Pv38* | 87          | Whole for *PvS25*                  | 3 in *PvS25* (2006-2011)                    | Jang et al. (2014) [80] |
|            |                  | Whole for *Pv38*                   | 2 in *Pv38* (2006-2011)                     |             |
| *PvS48/45 & PvS47* | 40 | Whole for *PvS48/45* | 7 in *PvS48/45* (2006-2011) | Woo et al. (2013) [81] |
| *PvS48/45 & PvS47* | 40 | Whole for *PvS48/45* | 10 in *PvS47* (2006-2010) |             |
| *PvMt-1 & PvMt-2* | 11 | Mitochondrial DNA                | 2 (1999)                                    | Iwagami et al. (2010) [82] |
| *PvLDH*    | 20               | Whole                              | 2 (2010-2011)                               | Shin et al. (2013) [47] |

*The average numbers of subtypes based on the antigens studied.*
most isolates after 2000 were different from the isolates during the initial re-emergence period [42]. The PvCSP is an abundant antigen on the surface of sporozoites, which is extensively used as a vaccine development target [43]. In the above study, 6 PvMSP-1 subtypes were identified in genomic parasite DNA from 632 patients and a new PvCSP subtype was found based on polymorphism of repeat regions. These authors also found that malaria parasites in the Republic of Korea were becoming progressively more genetically diverse. Interestingly, when PvCSP was loaded onto an RDT kit and applied with 121 patient sera, CSP antigen-antibody reactions were seen in up to 57% of cases [44]. Similarly, Kang et al. [45] suggested that the \( P. \) vivax population in the Republic of Korea has been diversifying rapidly and that new genotypes are appearing based on combinational genetic analysis of polymorphic patterns of 2 major antigenic proteins, \( \text{PvMSP-1} \) and \( \text{PvMSP-3α} \). They also found that the \( \text{PvDBP II} \) in Korean isolates had a polymorphic nature and that 13 different \( \text{PvDBP II} \) haplotypes were present in Korean \( P. \) vivax isolates. This population is under natural selection, and \( \text{PvDBP II} \) showed limited genetic polymorphism at restricted positions with a lower genetic diversity [46]. However, \( \text{PvLDH} \) in Korean isolates, a useful antigen for a rapid diagnostic test (RDT), did not show any genetic variation except a single-nucleotide polymorphism with no effect on the amino acid sequence [47].

In summary, genetic analyses using antigen-coding genes such as \( \text{PvDBP}, \text{PvCSP}, \text{PvMSP (MSP3α and MSP3β)}, \text{PvAMA}, \) and \( \text{PvGAM-I} \) suggest that only 1 or 2 alleles existed in South Korean isolates before 2000. However, these alleles have shown their patterns by year genetic diversity since then. It is not certain why the South Korean \( P. \) vivax population structure has changed so drastically since the early 2000s despite rapid reductions in transmission and low multiple infection rates. The rarity of polyclonal \( P. \) vivax infections in the Republic of Korea [48] suggests that local outcrossing alone is unlikely to explain the increased genetic divergence observed in the antigen-coding genes. Thus, cases introduced from an external reservoir likely contributed to the increased diversity [28]. Because changing antigenicity is a major strategy for parasites to be protected from host immunity, antigenic diversity/polymorphism, which links the expression of different alleles of a gene in different parasite populations with outcomes of selection under immune pressure, would selectively expand [49].

**A Spectrum of Population Structures Identified via Microsatellite Marker Analysis**

\( P. \) vivax populations are more genetically diverse and more frequently comprise polyclonal infections than do sympatric \( P. \) falciparum, suggesting that frequent recombination between the clones in the populations may contribute to genetic diversity [50]. Although \( P. \) vivax surface antigens show high genetic diversity, antigen-encoding genes can be subject to strong natural selection due to the host immune system. In contrast, microsatellites (MSs) are putatively neutral markers that can be used to monitor parasite transmission patterns without the potentially confounding impact of host immunity or other selective pressures. Simple MSs (sequence tandem repeats) are a valuable resource for the study of genetic diversity. They are useful for fingerprinting an organism because they are genetically hypervariable, co-dominant and locus-specific, and are not directly subject to host immunity, making them compatible with an important assumption of population genetic theory which states that molecular markers should be selectively neutral [51]. In case of \( P. \) falciparum, microsatellite markers have been known to be extensively widespread occurring every 2-3 kb throughout the genome [52]. In spite of a limited number of polymorphic MS loci in \( P. \) vivax (a few dozens) rather than that in \( P. \) falciparum (~1,000), MS-based studies of \( P. \) vivax have provided pivotal information regarding population structure and diversity [32,53,54]. The recent markers of choice for large-scale population studies are highly polymorphic and short (1-6 bp-long) MSs [55]. \( P. \) vivax contains 14 chromosomes. There are now 5 reference genomes of \( P. \) vivax (Salvador I, North Korean, India VII, Mauritania I, and Brazil I) [29] and \( P. \) vivax chromosomes appear to significantly larger and have more stable GC-rich genomes (approximately 40%) than those of \( P. \) falciparum (19%) [56,57]. Using 13 MS DNA markers across 12 chromosomes in \( P. \) vivax from the Republic of Korea, only 2 major multilocus MS genotypes were found during 1997-2000 while numerous genotypes appeared in 2007, indicating a marked expansion in the genome-wide genetic diversity of the \( P. \) vivax population structure in the Republic of Korea [26,51]. In addition, Iwagami et al. [58,59] investigated \( P. \) vivax in South Korean soldiers using 14 highly polymorphic MS DNA loci and dividing the samples from soldiers and veterans into 3 different time periods spanning 1993-1998, 1999-2003, and 2004-2008 in a 15-year-long longitudinal study on...
P. vivax population genetics. Further, another population genetic approach using P. vivax positive blood samples from 98 autochthonous cases and nine tandem repeat markers showed a trend of increasing parasite divergence since re-emergence [48]. It is also reported that polyclonal infections were rare (~3%) suggesting that local out-crossing alone was unlikely to explain the high diversity. The South Korean Army introduced a strategy of mass chemoprophylaxis with chloroquine during the malaria transmission season and primaquine after the transmission season to reduce the number of vivax malaria cases in both the military and civilian populations. Chemoprophylaxis with chloroquine (300 mg/wk) and primaquine (30 mg/wk) began with approximately 16,000 soldiers in 1997 and was expanded to more than 200,000 military personnel in zones at high risk for vivax malaria [60]. In a process of monitoring for chloroquine-resistant P. vivax using single nucleotide polymorphisms in Pmdr1 and microsatellite genotypes, P. vivax with newly identified SNPs in Pmdr1 was introduced and the microsatellite-based study revealed higher genetic diversity in samples collected in 2012 than in those collected in 2011 [61]. The study determined microsatellite genotypes of the 55 isolates based on a combination of allelic data of 10 microsatellite loci. The findings revealed that 2 dominant MS haplotypes had been transmitted for several years, providing decisive evidence for a low recombination rate. This may be due to recent migrations of different genotypes from other geographic regions to the Republic of Korea, and suggests an increase in the genetic diversity of the P. vivax population in the Republic of Korea [45,48]. While the level of genetic diversity in the P. vivax population in the Republic of Korea is significantly lower than that observed in tropical and subtropical areas [62-65], the population diversity is consistently increasing and is associated with a decline in multi-locus linkage disequilibrium (LD) over time [28,48,51,59]. Declining LD may also result from inter-allelic recombination. For example, PvMSP-1 displays heterogeneity in its frequency of inter-allelic recombination events, leading to rapid decline in multilocus LD between pairs of synonymous sites [66]. This is to be possible that the appearance of a P. vivax parasite population with new genotypes from other external reservoirs has accelerated genetic diversity in the South Korean P. vivax population through recombination events. These facts might be consistent with the low polyclonal infections and the small numbers of imported malaria cases in the Republic of Korea (<20 cases per year except 52 in 2000) [9].

**CONCLUSION**

Based on the above-mentioned studies on genetic diversity and incidence, the evidence obtained supports a link between vivax malaria in South and North Korea [67]. Annual WHO Malaria Reports show that North Korea had a restricted capacity to eliminate malaria until 1998 and experienced its highest reported incidence of malaria in 2001, peaking at 296,540 microscopically confirmed and indigenous cases (43.4 cases per 1,000 population) [68]. The vivax malaria cases in North Korea coincided with peak incidence in South Korea (3,932 in 1998, 3,621 in 1999, 4,142 in 2000, and 2,556 in 2001) [9]. Several independent genetic studies have reached the conclusion that the extant P. vivax population is shaped by continuous introductions from an external source, possibly North Korea, although the exact genetic data on North Korean P. vivax populations is not available to-date. This scenario does not exclude the consideration that local transmission events may also occur in South Korea and contribute to the population structure. Importantly, this interpretation allows for local propagation to occur largely via inbreeding, which is concordant with the low incidence, low prevalence of infections, and limited seasonality (July-August) of the opportunity for recombination in the An. sinensis vector. This scenario also supports the longitudinal trends in the genetic diversity and structure of P. vivax in the Republic of Korea. Indeed, there is an example elsewhere similar to such genetic diversity in the Republic of Korea. Sri Lanka has remarkably reduced its vivax malaria incidence case by 99.99% with high levels of heterozygosity of P. vivax and large effective population sizes [69]. Moreover, local vivax populations are characterized by high levels of allelic polymorphism and significant LD indicative of reduced recombination. However, approximately 70% isolates in that study had polyclonal infections. The authors suggested that the imported malaria could by the most likely source of vivax malaria transmission. Overall, the significant diversity of the P. vivax population observed hampers the development of an effective vaccine against South Korean isolates of vivax malaria. In comparison to the very low endemic, pre-elimination settings of Sabah, Malaysia [70] and Southern Iran [71], there is lesser evidence of population bottlenecks in the Republic of Korea.

Therefore, further continuous monitoring of malaria parasites using various molecular markers including polymorphic DNA markers in the P. vivax genome is needed for the control
and elimination of vivax malaria in the Republic of Korea. Such monitoring can also assess the population structure and dynamics of the parasite transmission.

ACKNOWLEDGMENTS

This research was supported by the grants of a Research Grant from Inha University (56866-01, TSK).

CONFLICTS OF INTERESTS

The authors declare they have no competing interests.

REFERENCES

1. Feachem RG, Phillips AA, Hwang J, Cotter C, Wielgosz B, Greenwood BM, Sabot O, Rodriguez MH, Abeyasinghe RR, Gheneypeus TA, Snow RW. Malaria elimination 1: Shrinking the malaria map: progress and prospects. Lancet 2010; 376: 1566-1578.

2. WHO. World malaria report. Geneva: World Health Organization 2017.

3. Mendis K, Sina BJ, Marchesini P, Carter R. The neglected burden of Plasmodium vivax malaria. Am J Trop Med Hyg 2001; 64: 97-106.

4. Carlton JM, Sina BJ, Adams JH. Why is Plasmodium vivax a neglected tropical disease? PLoS Negl Trop Dis 2011; 5: e1160.

5. Luo Z, Sullivan SA, Carlton JM. The biology of Plasmodium vivax explored through genomics. Ann NY Acad Sci 2015; 1342: 53-61.

6. Price RN, Tjitra E, Guerra CA, Yeung S, White NJ, Anstey NM. Vivax malaria: neglected and not benign. Am J Trop Med Hyg 2007; 77: 79-87.

7. Mueller I, Galinski MR, Baird JK, Carlton JM, Kochar DK, Alonso PL, del Portillo HA. Key gaps in the knowledge of Plasmodium vivax, a neglected human malaria parasite. Lancet Infect Dis 2009; 9: 555-566.

8. Ashley EA, Pyae PA, Woodrow CJ. Malaria. Lancet 2018; 391: 1608-1621.

9. Bahk YY, Lee HW, Na BK, Kim J, Jin K, Hong YS, Kim TS. Epidemiological characteristics of re-emerging vivax malaria in the Republic of Korea. Korean J Parasitol 2001; 39: 293-299.

10. Chai JY. Re-emerging Plasmodium vivax malaria in the Republic of Korea. Korean J Parasitol 1999; 37: 129-143.

11. Guerra CA, Howes RE, Pattil AP, Geting PW, Van Boeckel TP, Temperley WH, Kabaria CW, Tatem AJ, Manh BH, Elyazar IR, Baird JK, Snow RW, Hay SI. The international limits and population at risk of Plasmodium vivax transmission in 2009. PLoS Negl Trop Dis 2010; 4: e774.

12. Schultz L, Wapling J, Mueller I, Nsuke PO, Senn N, Nale J, Kimboro B, Buckee CO, Tavul L, Siba PA, Reeder JC, Barry AE. Multilocus haplotypes reveal levels of diversity and population structure of Plasmodium falciparum in Papua New Guinea, a region of intense perennial transmission. Malar J 2010; 9: 336.

13. Cui L, Escalante AA, Imwong M, Snounou G. The genetic diversity of Plasmodium vivax populations. Trends Parasitol 2003; 19: 220-226.

14. Rich SM, Ayala FJ. Population structure and recent evolution of Plasmodium falciparum. Proc Natl Acad Sci USA 2000; 97: 6994-7001.

15. Chai IH, Lim GI, Yoon SN, Oh WJ, Kim SJ, Chai JY. Occurrence of tertian malaria in a male patient who has never been abroad. Korean J Parasitol 1994; 32: 195-200 (in Korean).

16. Lee KJ, Kim CB, Choi BJ, Park KH, Park JK. Analysis of vivax malaria cases in Gangwon-do (province), Korea in the year 2000. Korean J Parasitol 2001; 39: 293-299.

17. Moon JJ, Cho SY. Incidence patterns of vivax malaria in civilians residing in a high-risk county of Kyonggi-do (province), Republic of Korea. Korean J Parasitol 2001; 39: 232-239.

18. Lee JS, Lee WJ, Cho SH, Ree HL. Outbreak of vivax malaria in areas adjacent to the demilitarized zone, South Korea. Am J Trop Med Hyg 2002; 66: 13-17.

19. Nkhoma SC, Nair S, Al-Saai S, Ashley E, McGready R, Phyo AP, Nosten F, Anderson TJ. Population genetic correlates of declining transmission in a human pathogen. Mol Ecol 2013; 22: 273-285.

20. Roy SW, Ferreira MJ, Hartl DL. Evolution of allelic dimorphism in malarial surface antigens. Heredity 2008; 100: 103-110.

21. Barry AE, Arnott A. Strategies for designing and monitoring malaria vaccines targeting diverse antigens. Front Immunol 2014; 5: 359.

22. Barry AE, Wältmann A, Koepfli C, Barnadas C, Mueller I. Uncovering the transmission dynamics of Plasmodium vivax using population genetics. Pathog Glob Health 2015; 109: 142-152.

23. Auburn S, Barry AE. Dissecting malaria biology and epidemiology using population genetics and genomics. Int J Parasitol 2017; 47: 77-85.

24. Casares S, Richie TL. Immune evasion by malaria parasites: a challenge for vaccine development. Curr Opin Immunol 2009; 21: 321-330.

25. Suwanarusk R, Russell B, Chavchich M, Chalfein F, Kenangalem E, Kosaisavee V, Prasetyorini B, Piera KA, Barends M, Brockman A, Lek-Uthai U, Anstey NM, Tjitra E, Nosten F, Cheng Q, Price RN. Chloroquine resistant Plasmodium vivax: in vitro characterization and association with molecular polymorphisms. PLoS One 2007; 2: e1089.

26. Huh AJ, Kwak YG, Kim ES, Lee KS, Yeon JS, Cho YK, Kim CS, Park JW. Parasitaemia characteristics of Plasmodium vivax malaria patients in the Republic of Korea. J Korean Med Sci 2011; 26: 42-46.

27. Oh MD, Shin H, Shin D, Kim U, Lee S, Kim N, Choi MH, Chai JY, Choe K. Clinical features of vivax malaria. Am J Trop Med Hyg 2001; 65: 143-146.

28. Kim JY, Goo YK, Zo YG, Ji SY, Trirmarsanto H, To S, Clark TG, Price RN, Auburn S. Further evidence of increasing diversity of Plasmodium vivax in the Republic of Korea in recent years. PLoS
Bahk et al.: Genetic diversity of Plasmodium vivax from Korean isolates

One 2016; 11: e0151514.

29. Neafsey DE, Galinsky K, Jiang RH, Young L, Sykes SM, Saif S, Gujia S, Goldberg JM, Young S, Zeng Q, Chapman SB, Dash AP, Anvikar AR, Sutton PL, Birren BW, Escalante AA, Barnwell JW, Carlton JM. The malaria parasite Plasmodium vivax exhibits greater genetic diversity than Plasmodium falciparum. Nat Genet 2012; 44: 1046-1050.

30. Choi YK, Choi KM, Park MH, Lee EG, Kim YJ, Lee BC, Cho SH, Rhie HG, Lee HS, Yu JR, Lee JS, Kim TS, Kim JY. Rapid dissemination of newly introduced Plasmodium vivax genotypes in South Korea. Am J Trop Med Hyg 2010; 82: 426-432.

31. Arnott A, Barry AE, Reeder JC. Understanding the population genetics of Plasmodium vivax is essential for malaria control and elimination. Malar J 2012; 11: 14.

32. Brito CF, Ferreira MU. Molecular markers and genetic diversity of Plasmodium vivax. Mem Inst Oswaldo Cruz 2011; 106: 12-26.

33. Hwang SY, Kim SH, Kho WG. Genetic characteristics of polymorphic antigenic markers among Korean isolates of Plasmodium vivax. Korean J Parasitol 2009; 47 (suppl): 51-58.

34. Kho W, Chung J, Sim E, Kim D, Chung W. Analysis of polymorphic regions of Plasmodium vivax Duffy binding protein of Korean isolates. Korean J Parasitol 2001; 39: 143-150.

35. Lim CS, Kim SH, Kwon SI, Song JW, Song KJ, Lee KN. Analysis of Plasmodium vivax merozoite surface protein-1 gene sequences from resurgent Korean isolates. Am J Trop Med Hyg 2000; 62: 261-265.

36. Han ET, Song TE, Park JH, Shin EH, Guk SM, Kim TY, Chai JY. Allelic dimorphism in the merozoite surface protein-3 gene of Plasmodium vivax. Korean J Parasitol 2001; 39: 157-162.

37. Suh IB, Hoffman KJ, Kim SH, Song KJ, Song JW, Lee JS, Lim CS. The analysis of Plasmodium vivax Duffy receptor binding domain gene sequence from resurgent Korean isolates. Parasitol Res 2001; 87: 1007-1010.

38. Kang JM, Lee J, Cho PY, Moon SU, Bahk YY, Cho PY, Sohn WM, Park JW, Park JW, Kim TS, Na BK. Genetic diversity and natural selection of Duffy binding protein of Plasmodium vivax Korean isolates. Acta Trop 2013; 125: 67-74.

39. Shin HI, Kim JY, Lee WJ, Sonh Y, Lee SW, Kang YJ, Lee HW. Polymorphism of the parasite lactate dehydrogenase gene from Plasmodium vivax Korean isolates. Malar J 2013; 12: 166.

40. Kim JY, Goo YK, Zo YG, Ji SY, Timansanto H, To S, Clark TG, Price RN, Auburn S. Further evidence of increasing diversity of Plasmodium vivax in the Republic of Korea in recent years. PLoS One 2016; 11: e0151514.

41. Hisaeda H, Yasutomo K, Himeno K. Malaria: immune evasion by parasites. Int J Biochem Cell Biol 2005; 37: 700-706.

42. Ferreira MU, Karunaweera ND, da Silva-Nunes M, da Silva NS, Wirth DF, Hartl DL. Population structure and transmission dynamics of Plasmodium vivax in rural Amazonia. J Infect Dis 2007; 195: 1218-1226.

43. Russell B, Suwanarusk R, Lek-Uthai U. Plasmodium vivax genetic diversity: microsatellite length matters. Trends Parasitol 2006; 22: 399-401.

44. Su X, Ferdig MT, Huang Y, Huynh CQ, Liu A, Wootton JC, Wellens TE. A genetic map and recombination parameters of the human malaria parasite Plasmodium vivax. Science 1999; 286: 1351-1353.

45. Karunaweera ND, Ferreira MU, Hartl DL, Wirth DF. Fourteen polymorphic microsatellite DNA markers for the human malaria parasite Plasmodium vivax. Mol Ecol Notes 2007; 7: 172-175.

46. Rezende AM, Tarazona-Santos E, Fontes CJ, Souza JM, Couto AD, Carvalho LH, Brito CF. Microsatellite loci: determining the genetic variability of Plasmodium vivax. Trop Med Int Health 2010; 15: 718-726.

47. Schlötterer C. Genome evolution: are microsatellites really simple sequences? Curr Biol 1998; 8: 132-134.

48. Gomez JC, Bockarie MJ, Baird JK, Carlton JM, Zimmerman PA. Identification of a polymorphic Plasmodium vivax microsatellite marker. Am J Trop Med Hyg 2003; 69: 377-379.

49. Carlton JM, Galinski MR, Barnwell JW, Dame JB. Karyotype and syntenies on the chromosomes of all four species of human malaria parasites. Mol Biochem Parasitol 1999; 101: 23-32.

50. Iwagami M, Fukumoto M, Hwang SY, Kim SH, Kho WG, Kano S. Population structure and transmission dynamics of Plasmodium vivax.
vivax in the Republic of Korea based on microsatellite DNA analysis. PLoS Negl Trop Dis 2012; 6: e1592.

59. Iwagami M, Hwang SY, Kim SH, Park SJ, Lee GY, Louise E, Matsumoto-Takahashi A, Kho WG, Kano S. Microsatellite DNA analysis revealed a drastic genetic change of Plasmodium vivax population in the Republic of Korea during 2002 and 2003. PLoS Negl Trop Dis 2013; 10: e2522.

60. Yeom JS, Ryu SH, Oh S, Choi DH, Song KJ, Oh YH, Lee JH, Kim YA, Ahn SY, Yang HY, Cha JE, Park JW. Evaluation of anti-malarial effects of mass chemoprophylaxis in the Republic of Korea. J Korean Med Sci 2005; 20: 707-712.

61. Chung DI, Jeong S, Dinzouna-Boutamba SD, Yang HW, Yeo SG, Hong Y, Goo YK. Evaluation of single nucleotide polymorphisms of pmmdr1 and microsatellite genotype in Plasmodium vivax isolates from Republic of Korea military personnel. Malar J 2015; 14: 336.

62. Imwong M, Nair S, Pukrittayakamee S, Sudimack D, Williams JT, Mayxay M, Newton PN, Kim JR, Nandy A, Orsorio L, Carlton JM, White NJ, Day NP, Anderson TJ. Contrasting genetic structure in Plasmodium vivax populations from Asia and South America. Int J Parasitol 2007; 37: 1013-1022.

63. Karunaweera ND, Ferreira MJ, Haril DL, Wirth DE. Fourteen polymorphic microsatellite DNA markers for the human malaria parasite Plasmodium vivax. Mol Ecol Notes 2007; 7: 172-175.

64. Orjuela-Sánchez P, da Silva NS, da Silva-Nunes M, Ferreira MJ. Recurrent parasitaemias and population dynamics of Plasmodium vivax polymorphisms in rural Amazonia. Am J Trop Med Hyg 2009; 81: 961-968.

65. Auburn S, Barry AE. Dissecting malaria biology and epidemiology using population genetics and genomics. Int J Parasitol 2017; 47: 77-85.

66. Putaporntip C, Jongvutivises S, Sakihama N, Ferreira MJ, Kho WG, Kaneko Akira, Kanbara H, Hattori T, Tanabe K. Mosaic organization and heterogeneity in frequency of allelic recombination of the Plasmodium vivax merozoite surface protein-1 locus. Proc Natl Acad Sci USA 2002; 99: 16348-16353.

67. Ree HI. Unstable vivax malaria in Korea. Korean J Parasitol 2000; 38: 119-138.

68. World Health Organization. World Malaria Report 2010. Geneva, Switzerland. World Health Organization. 2010.

69. Gunawardena S, Ferreira MJ, Kapilandana GMG, Wirth DE, Karunaweera ND. The Sri Lankan paradox: high genetic diversity in Plasmodium vivax populations despite decreasing levels of malaria transmission. Parasitology 2014; 141: 880-890.

70. Abdullah NR, Barber BE, William T, Norahmad NA, Satsu UR, Munirandy PK, Ismail Z, Grigg MJ, Jelip J, Piera K, von Seidlein L, Yeo TW, Anstey NM, Price RN, Auburn S. Plasmodium vivax population structure and transmission dynamics in Sabah Malaysia. PLoS One 2013; 8: e82553.

71. Hamedi Y, Sharifi-Sarasiabli K, Dehghan E, Safari R, To S, Han-dayuni I, Trimmersanto H, Price RN, Auburn S. Molecular epidemiology of P. vivax in Iran: High diversity and complex substructure using neutral markers, but no evidence of Y976F mutation at PvMDR1. PLoS One 2016; 11: e0166124.

72. Chung Ji, Chun EH, Chun JH, Kho WG. Analysis of the Plasmodium vivax apical membrane antigen-1 gene from re-emerging Korean isolates. Parasitol Res 2003; 90: 325-329.

73. Kim T, Kim YJ, Song KJ, Song JW, Cha SH, Kim YK, Shin YK, Sih IB, Lim CS. The molecular characteristics of circumsporozoite protein gene subtypes from Plasmodium vivax isolates in the Republic of Korea. Parasitol Res 2002; 88: 1051-1054.

74. Kho WK, Park YH, Chung JY, Kim JP, Hong YL, Lee WJ, Kim TS, Lee JS. Two new genotypes of Plasmodium vivax circumsporozoite protein found in the Republic of Korea. Korean J Parasitol 1999; 37: 265-270.

75. Kim SH, Hwang SY, Shin JW, Moon CS, Kim DW, Kho WK. Molecular genetic characterization of the merozoite surface protein 1 gene of Plasmodium vivax from reemerging Korean isolates. Clin Vaccine Immunol 2009; 16: 733-738.

76. Han ET, Wang Y, Lim CS, Cho JH, Chai JY. Genetic diversity of the malaria vaccine candidate merozoite surface protein 1 gene of Plasmodium vivax field isolates in the Republic of Korea. Parasitol Res 2011; 109: 1571-1576.

77. Kang JM, Ji HJ, Kang YM, Lee DH, Moon SU, Sohn WM, Park JW, Kim TS, Na BK. Genetic polymorphism and natural selection in the C-terminal 42 kDa region of merozoite surface protein-1 among Plasmodium vivax Korean isolates. Malaria J 2012; 11: 206.

78. Son UH, Dinzozona-Boutamba SD, Lee S, Yun HS, Kim JY, Joo SY, Jeong S, Rhee MH, Hong Y, Chung DI, Kwak D, Goo YK. Diversity of Vir genes in Plasmodium vivax from endemic regions in the Republic of Korea: an initial evaluation. Korean J Parasitol 2017; 55: 149-158.

79. Jang JW, Yun SG, Woo MK, Han ET, Lu E, Gao Q, Yoon SY, An SSA, Lim CS. Sequence polymorphisms of Plasmodium vivax tryptophan and alanine rich antigen (PvTARAg55). Acta Trop 2015; 142: 122-126.

80. Jang JW, Cho CH, Kim JY, Koh YE, Woo MK, Kim KA, Yoon SY, Lim MS, Han ET, An SSA, Lim CS. Sequence polymorphisms in Pvs48/45 and Pvs47 gametocyte and gamete surface proteins in Plasmodium vivax isolated in Korea. Mem Inst Oswaldo Cruz 2013; 108: 359-367.

81. Iwagami M, Hwang SY, Fukumoto M, Hayakawa T, Tanabe K, Kim SH, Kho WG, Kano S. Geographic origin of Plasmodium vivax in the Republic of Korea: haplotype network analysis based on the parasite’s mitochondrial genome. Malaria J 2010; 9: 184.