



Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:

Barry, AE;Leliwa-Sytek, A;Tavul, L;Imrie, H;Migot-Nabias, F;Brown, SM;McVean, GAV;Day, KP

Title:

Population genomics of the immune evasion (var) genes of Plasmodium falciparum

Date:

2007-03-01

Citation:

Barry, A. E., Leliwa-Sytek, A., Tavul, L., Imrie, H., Migot-Nabias, F., Brown, S. M., McVean, G. A. V. & Day, K. P. (2007). Population genomics of the immune evasion (var) genes of Plasmodium falciparum. Plos Pathogens, 3 (3), pp.690-690. <https://doi.org/10.1371/journal.ppat.0030034>.

Persistent Link:

<https://hdl.handle.net/11343/265004>

License:

CC BY

Population Genomics of the Immune Evasion (*var*) Genes of *Plasmodium falciparum*

Alyssa E. Barry^{1,2^{‡a}*}, Aleksandra Leliwa-Sytek^{1,2^{‡b}}, Livingston Tavul³, Heather Imrie^{1,2}, Florence Migot-Nabias^{4^{‡c}}, Stuart M. Brown⁵, Gilean A. V. McVean⁶, Karen P. Day^{1,2^{‡b}}

1 Peter Medawar Building for Pathogen Research, University of Oxford, Oxford, United Kingdom, **2** Department of Zoology, University of Oxford, Oxford, United Kingdom, **3** Papua New Guinea Institute for Medical Research, Madang, Papua New Guinea, **4** Centre International de Recherches Médicales de Franceville (CIRMF), Franceville, Gabon, **5** Research Computing Resource, New York University School of Medicine, New York, New York, United States of America, **6** Department of Statistics, University of Oxford, Oxford, United Kingdom

Var genes encode the major surface antigen (PfEMP1) of the blood stages of the human malaria parasite *Plasmodium falciparum*. Differential expression of up to 60 diverse *var* genes in each parasite genome underlies immune evasion. We compared the diversity of the DBL α domain of *var* genes sampled from 30 parasite isolates from a malaria endemic area of Papua New Guinea (PNG) and 59 from widespread geographic origins (global). Overall, we obtained over 8,000 quality-controlled DBL α sequences. Within our sampling frame, the global population had a total of 895 distinct DBL α “types” and negligible overlap among repertoires. This indicated that *var* gene diversity on a global scale is so immense that many genomes would need to be sequenced to capture its true extent. In contrast, we found a much lower diversity in PNG of 185 DBL α types, with an average of approximately 7% overlap among repertoires. While we identify marked geographic structuring, nearly 40% of types identified in PNG were also found in samples from different countries showing a cosmopolitan distribution for much of the diversity. We also present evidence to suggest that recombination plays a key role in maintaining the unprecedented levels of polymorphism found in these immune evasion genes. This population genomic framework provides a cost effective molecular epidemiological tool to rapidly explore the geographic diversity of *var* genes.

Citation: Barry AE, Leliwa-Sytek A, Tavul L, Imrie H, Migot-Nabias F, et al. (2007) Population genomics of the immune evasion (*var*) genes of *Plasmodium falciparum*. PLoS Pathog 3(3): e34. doi:10.1371/journal.ppat.0030034

Introduction

Quantifying the diversity of major surface antigens underlying immune evasion of HIV 1 and 2 and Influenza A has been central to characterizing the transmission dynamics of these important human pathogens. In addition, documentation of variation data has provided a basis for the development of candidate vaccine targets [1,2]. Surprisingly, in-depth molecular epidemiological sampling and population genomic analyses of the *var* genes encoding the major blood stage surface antigen of the malaria parasite, *Plasmodium falciparum* erythrocyte membrane protein 1 (PfEMP1), has not been done. This is largely due to the inherent difficulties in the population genomic analysis of highly diverse multigene families. We set out to develop and evaluate a rapid, molecular epidemiological population genomic framework to investigate *var* gene diversity in natural parasite populations, due to the importance of these genes to the biology of *P. falciparum*.

To achieve chronic infection, malaria parasites evade the host immune response by switching PfEMP1 isoforms through differential expression of members of the *var* multigene family [3–5]. PfEMP1 is expressed on the surface of blood stage parasites known as trophozoites [6] and the transmission (early gametocyte) stages [7]. Parasite adhesion occurs in the deep vasculature of host tissues by binding of PfEMP1 to host endothelial cell receptors. Some PfEMP1-adhesion interactions are proposed to lead to severe disease manifestations such as cerebral and placental malaria (reviewed in [8]). Variant specific anti-PfEMP1 antibodies are believed to contribute to the regulation of parasite

density in a manner that decreases the incidence of clinical disease [9–13]. This immunity may reduce the duration of infection in a variant-specific manner to drive the dynamics of multiple infections in semi-immune children [14] and induced infections in humans [15]. Immunity to PfEMP1 can thereby influence transmission by reducing persistence. It may also reduce transmission by regulating the density of asexual blood stages with potential to become transmission stages and by directly targeting early gametocytes to prevent the maturation of transmission stages [16]. Consequently, diversity of PfEMP1 or *var* genes is able to promote transmission success by immune evasion.

Describing the diversity of *var* genes presents a more

Editor: Margaret J. Mackinnon, University of Cambridge, United Kingdom

Received: June 7, 2006; **Accepted:** January 24, 2007; **Published:** March 16, 2007

Copyright: © 2007 Barry et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Abbreviations: CIDR, cysteine interdomain rich region; DBL, Duffy binding like; PfEMP1, *Plasmodium falciparum* erythrocyte membrane protein 1; PNG, Papua New Guinea; PTS, pairwise type sharing

* To whom correspondence should be addressed. E-mail: alyssa.barry@burnet.edu.au

^{‡a} Current address: MacFarlane Burnet Institute for Medical Research and Public Health, Melbourne, Victoria, Australia

^{‡b} Current address: Department of Medical Parasitology, New York University School of Medicine, New York, New York, United States of America

^{‡c} Current address: Institut de Recherche pour le Développement, Cotonou, République du Bénin

Author Summary

Malaria parasites live in red blood cells of the human host for part of the life cycle, during which a family of diverse antigens known as PfEMP1 are placed on the surface. PfEMP1 variants switch by sequential expression of up to 60 *var* genes. This allows the parasite to evade immune detection within an individual host, enhancing its chances to be transmitted to the mosquito vector in situations where mosquitoes are seasonally available. Methods to rapidly assess *var* gene diversity in parasite populations are needed to measure antigenic diversity and define relationships with malaria transmission. Using a specialized framework, we completed the first systematic sampling of *var* genes from parasite genomes obtained from the same (Papua New Guinea [PNG]) and different (global) populations. Globally, there was no limit to the number of *var* genes because parasites rarely shared *var* genes. In PNG, *var* gene numbers were restricted due to high levels of sharing, and most were only found in that population. Recombination was important to the evolution of *var* genes in PNG. The data suggest there are distinct *var* genes in different populations, which may have consequences for the spread of malarial disease from one geographic area to another.

complex problem than assessing the diversity of the single copy major surface antigen genes of HIV and Influenza A. Individual *P. falciparum* genomes have repertoires of *var* genes that can recombine with other repertoires during the obligatory sexual phase of the life cycle in the mosquito [17–19]. There is also circumstantial evidence for ectopic recombination among *var* genes within the same genome, possibly during both meiosis and mitosis [20–22]. Therefore, there is enormous potential to generate diversity, even among closely related genomes. *Var* genes are large (5–12 kb) and complex [23–26], encoding variable numbers and classes of the adhesion domains, Duffy binding like (DBL: α , β , δ , ϵ , and γ classes) and cysteine interdomain rich (CIDR: α , β , and γ classes) [26]. Both size and complexity preclude population genomic analysis of the full *var* gene sequences. The DBL α encoding domain was used previously as a marker of *var* genes in investigations of diversity [21,22,27–30] and expression [31–33]. The small size (~1 kb) and ubiquitous presence of DBL α among *var* genes [24–26,34] make this domain a suitable population genomic marker.

Previous analyses of *var* gene diversity have examined DBL α domain sequences from the *var* gene repertoires of allopatric (distantly related) [22,27,30] or just a few sympatric (closely related) [28,29] isolates. These studies have established that any pair of DBL α sequences from the same genome were on average as diverse as any pair from two different genomes, with a range of 45%–100% amino acid identity [3,22,27,28]. This has made it impossible to identify *var* gene orthologs among genomes. Limited overlap (shared DBL α sequences) among *var* repertoires from sympatric isolates has also been reported [27–29], suggesting that many genomes must be sampled to see the extent of diversity of these genes in natural populations. Given the importance of *var* genes to transmission, a systematic sequencing effort and population genomic analysis is needed to examine *var* gene diversity in sufficient depth to estimate levels of antigenic diversity within natural populations.

A high-throughput population genomic framework was developed to address sampling issues specific to the molec-

ular epidemiological analysis of diverse multigene families. This allowed the random sampling of the *var* gene repertoires of culture-adapted and field isolates of *P. falciparum* by sequencing DBL α domains as population genomic markers (Figure 1). *Var* genes were sampled from a “global” collection of isolates, including clones 3D7 and HB3 used in genome sequencing projects ([34]; D. Wirth, personal communication). DBL α sequences from these isolates were used to validate the framework. The “global” DBL α sequences were combined with available data from previous studies and compared to that obtained from a local population of Papua New Guinea (PNG). The results show immense levels of diversity among the *var* genes with strong evidence of geographic structuring of variation. We demonstrate patterns of similarity among sequences that suggest the widespread action of recombination in creating and maintaining diversity.

Results

Population Sampling

We tested a population genomic framework (Figures S1–S4; Text S1) to sample and analyze the *var* gene sequences of 25 cloned *P. falciparum* lines/isolates from widespread geographic origins (global; see Materials and Methods). This gave 4,754 reads (median = 161.5; range = 43–311 per isolate) of high quality DBL α sequences, which after removing redundancy, resulted in 608 consensus sequences (median = 20; range = 10–53 per isolate), with an average of seven times coverage. The GenBank accession numbers for these sequences can be found in Table S1. Artifacts were limited due to the multiple reads contributing to each DBL α sequence and by the use of a 96% DNA sequence identity cut-off (grouping reads from a single isolate with less than 4% sequence differences into a single consensus). To calculate the number of distinct DBL α sequences among isolates, individual repertoires were compared with each other, again using an arbitrary 96% DNA sequence identity cut-off. This identified shared DBL α sequences (i.e., overlap) among *var* repertoires. It also defined all of the distinct DBL α sequences in the global population sample. We named these distinct sequences “DBL α types” rather than alleles because the possibility of ectopic recombination means that the ancestry of a particular *var* gene sequence will be unlikely to be linked to physical location. It should be noted that by defining types in this way, synonymous and nonsynonymous mutations were treated equally. We identified 534 distinct DBL α types from the 608 sequences obtained from the global isolates. This large number of types confirmed the wide-ranging amplification capacity of the degenerate primers (see Materials and Methods, Text S2). Some *var* gene types were shared among isolates as shown by the fact that the number of DBL α sequences was greater than the number of DBL α types in the population. When 480 DBL α sequences (404 types) obtained from the GenBank database (see Materials and Methods) were combined with these data, 1,088 DBL α sequences and 895 DBL α types were defined. We did not have information for the number of reads contributing to this additional data.

Local sampling of 30 *P. falciparum* isolates from the population of Amele, PNG (see Materials and Methods) gave 3,080 reads (median = 85.5; range = 8–238 per isolate). Analysis of the sequence data resulted in 460 DBL α sequences

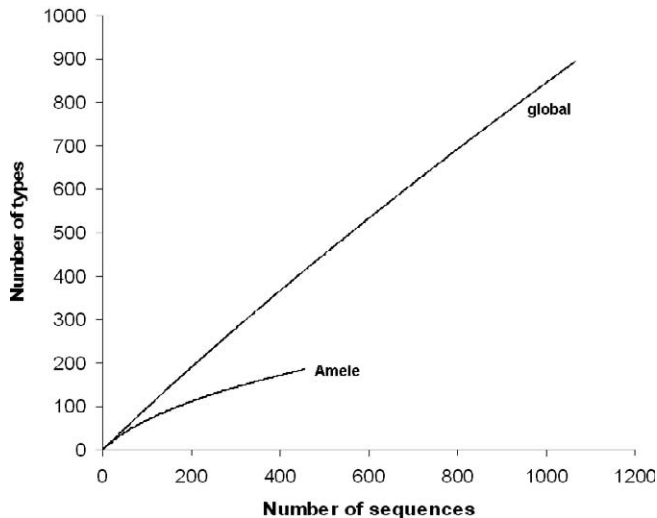


Figure 1. Cumulative Diversity Curves for DBL α from the Amele and Global Populations

The cumulative diversity of DBL α was determined by comparing each new DBL α repertoire to previous repertoire(s) using a 96% DNA sequence identity cut-off. We plotted the number of accumulating “types” (distinct sequences) against the number of “sequences” compared. For example, the first point on the plot will be the number of types found in both isolates A and B on the y-axis against the total number of sequences compared on the x-axis. The next point will be the number of types found in isolates A, B, and C against the total number of sequences compared. The plot shown here is the average curve of 1,000 permutations of the order that isolates were compared (i.e., isolates A, B, then C; or A, C, then B; or B, C, then A).
doi:10.1371/journal.ppat.0030034.g001

(median = 15; range = 2–30 per isolate) with each sequence having an average of 6.7 times coverage. The GenBank accession numbers for these sequences can be found in Table S1. We defined 185 DBL α types from the sequences obtained. The ratio of sequences to types for the global sampling (ratio = 1.21) was much less than that of Amele local sampling (ratio = 2.5), demonstrating greater overlap amongst *var* gene repertoires from the Amele isolates than those from the global collection. This overlap is defined more precisely below.

Cumulative Diversity

An important question in malaria population genomics is “how many isolates need to be sampled to find the majority of *var* gene diversity in a population?” To answer this question, we applied an ecological method to the data. We measured how the rate of DBL α type discovery changed with sample size in a manner similar to a species richness curve [35]. This method was previously used to measure heterozygosity in the human blood group antigens [36]. The number of DBL α types was plotted against the number of DBL α sequences as the *var* gene repertoire from each isolate was sampled. For sampling of global isolates, the curve did not plateau even though we sampled more than 1,000 DBL α sequences from 59 isolates (the 25 global isolates we sampled plus 34 *var* repertoires available from GenBank including the entire 3D7 *var* repertoire), showing that the majority of the *var* gene diversity in the global *P. falciparum* population was not defined (Figure 1). When we plotted the cumulative diversity for the Amele local population, the curve began to deviate

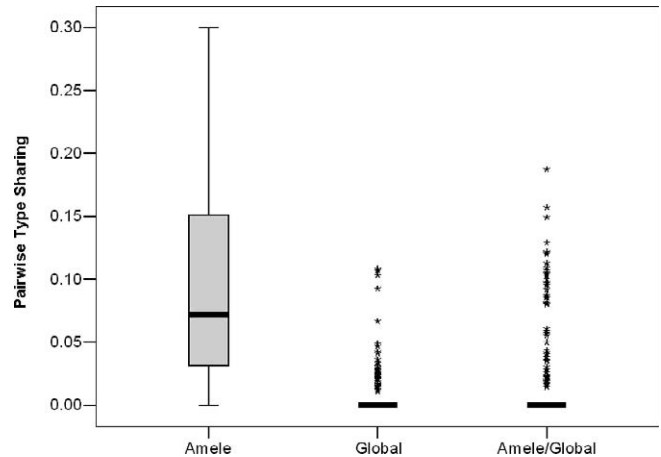


Figure 2. PTS among DBL α Repertoires from Amele and Global Populations

PTS was plotted for pairs of Amele DBL α repertoires, global DBL α repertoires, and Amele DBL α repertoires paired with global DBL α repertoires. The median (thick line), interquartile range (boxes), approximate 95% sampling interval (whiskers = 1.5 times the interquartile range), and suspected outliers (asterisks) are shown. The raw data for these plots can be found in Figure S5.
doi:10.1371/journal.ppat.0030034.g002

from that of the global sample after sampling just 50 DBL α sequences (approximately three isolates) and tended toward a plateau after sampling 460 DBL α sequences (30 isolates). Thus, while *var* gene diversity was immense on a global scale it was restricted in this local population of PNG. The shape of the curve suggested that a large proportion (around 50%–60%) of common types were sampled in the local population of Amele, although this may not represent all of the diversity in the Madang region or in PNG.

Pairwise Type Sharing

The higher ratio of DBL α sequences to DBL α types and the flattening of the cumulative curve for Amele in comparison to that for global indicated a greater degree of *var* repertoire overlap among Amele isolates than among global isolates. To quantify this overlap in the two populations, we considered the proportion of those types that were found in both of a pair of isolates; a statistic designated the pairwise type sharing (PTS) (see Materials and Methods). We found that average PTS among Amele isolates (or within the population) was significantly greater than for global isolates (Figures 2 and S5). The PTS statistic estimated that 7% of *var* genes were shared among Amele isolates, but that a negligible number of *var* genes were shared among global isolates. When the two populations were compared, again the PTS was negligible (Figure 2). Note that we removed other PNG isolates from the global population (see Materials and Methods) for this analysis as these showed significantly higher PTS values with Amele isolates (unpublished data). The higher degree of overlap in Amele in relation to that among global isolates demonstrated the presence of geographic population structure. In particular, there was higher overlap among the contemporary Amele isolates and among those isolates and PNG isolates, also from Amele, but collected at different time periods (see Materials and Methods). Some of this will be due to the presence of a few high-frequency types in Amele (Table S2).

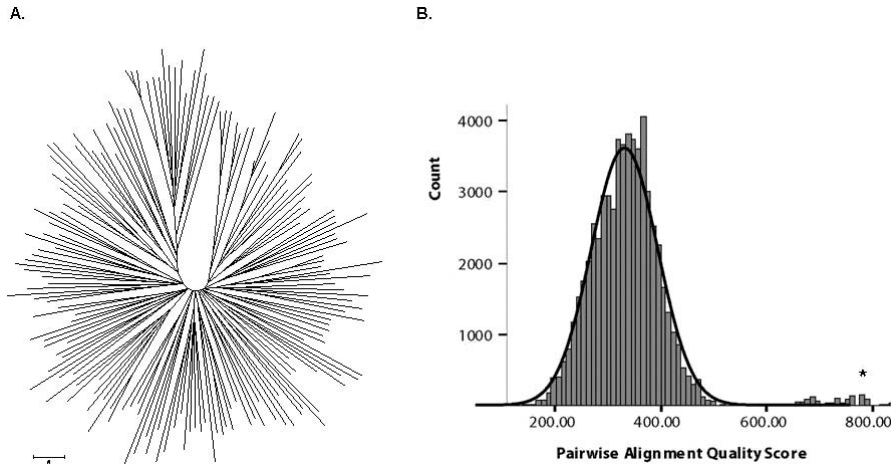


Figure 3. Diversity of Amele DBL α Sequences

Relationships among the *var* gene sequences from Amele were defined by (A) phylogenetic analysis of Amele DBL α types. A multiple alignment of amino acid sequences was constructed in Clustal W [70] with parameters set at gap opening = 5 and gap extension = 0.05, for all translated DBL α types ($n = 177$ after removal of those with frame shifts) in the Amele population. A neighbor-joining tree was constructed in MEGA version 3.0 [71] for Amele types using the p-distance algorithm, pairwise alignment, and 1,000 bootstrapping repetitions. The consensus tree is shown.

(B) Analysis of the pairwise alignment quality scores for Amele DBL α amino acid sequences. Here, the frequency distribution of scores is shown. Note the approximately Normal distribution of scores for the cluster of scores to the left, i.e., the dark line indicates maximum likelihood estimate for the Normal distribution. The asterisk indicates a cluster of pairs with high quality scores, which correspond to the shared sequences among isolates (types). doi:10.1371/journal.ppat.0030034.g003

Geographic Distribution of Amele Types

To test whether there was any geographic population structure, we examined the distribution of Amele types in the global population sample. The majority of the defined Amele types (61.6%) was unique to PNG. The remaining 71 types were found in Africa (24.8%), Asia (3.2%), and Latin America (3.8%); two were found in all malaria endemic regions and ten types were found in either Africa and Asia (2.2%); Africa and Latin America (2.7%); or Asia and Latin America (0.5%) (Table S2). The geographic restriction of Amele types to PNG demonstrates the existence of geographic population structure. Nevertheless, a substantial fraction of types has a cosmopolitan distribution indicating that different geographic regions do not have completely different repertoires of *var* genes.

Sequence Diversity among Amele Types

Having demonstrated extremely high levels of diversity among the *var* gene sequences, we wanted to explore potential structuring within the data. For example, do sequences broadly fall into a few groups? Or is every sequence equally different from all others? A natural approach to analyzing such structure within the data is to construct a phylogenetic tree for the sequences from a multiple-sequence alignment of the translated DNA sequences (Figure S6). The phylogeny estimated from the aligned DBL α types from Amele showed mostly distant relationships with long external branches and short, unresolved internal branches (Figure 3A). This “star-like” phylogeny suggests that the sequences are so diverse that little structuring to the variation is apparent. One evolutionary force known to result in “star-like” trees is recombination [37], suggesting that frequent exchange between different *var* gene copies may be occurring. This tree shape may also result from a rapidly expanding population or other forms of natural selection [37]. However, it is also possible that the “star-like” quality may simply

reflect a low quality multiple sequence alignment (Figure S6). This showed it was difficult to assess the relatedness of *var* gene sequences using multiple sequence alignments. Alternatively, pairwise analyses will maximize the quality of an alignment and highlight the most highly related fragments of a pair. Quality scores from these alignments are better estimates of the relationship among potentially recombined sequences because they are based on the percent similarity as well as the length of the alignment (i.e., the partial alignment of a pair of recombined sequences will have a higher score than a pair of sequences with an equal number of polymorphisms across their entire length; see Materials and Methods). The distribution of the quality scores can then be used to identify potential clustering of even weakly related sequences. Therefore, we also considered the distribution of the quality scores from pairwise sequence alignments.

Pairwise alignments of the translated DBL α sequences (i.e., prior to type definition) showed an average amino acid similarity of 66% (range = 22%–100%). The distribution of quality scores among Amele DBL α sequences showed a bimodal distribution with a small cluster of high scoring alignments to the right (Figure 3B). These clusters corresponded to the sequences that were shared among isolates (i.e., types) and those that had a high degree of similarity. The majority of sequences, however, show more distant relationships, resulting in a distribution of quality scores that appears approximately Normal (Figure 3B). One consequence of recombination is that different regions of pairs of sequences will be related to each other to different degrees, and the more recombination, the stronger the homogenizing force. High levels of recombination, therefore, might “average” out the quality score across the sequence and result in the approximately Normal distribution, while low levels of recombination are expected to result in more “jagged” distributions. Overall, these analyses suggested that recomb-

nation is a major force in the generation and maintenance of variation in the *var* genes. Given the antigenic nature of the *var* genes, we suggest that recombination is an important source of novelty for the *P. falciparum* *var* gene repertoire. However, the evidence presented here is only circumstantial, and the extreme diversity of the sequences makes application of existing population genetic approaches to detecting recombination problematic. We are currently developing novel population genetic approaches to learn about genetic exchange among the *var* gene sequences.

Var Gene Groups

To determine whether previously observed *var* gene structural groups found in the 3D7 genome [24,25] were represented within the Amele sample, we used a phylogenetic approach to identify *var* gene groupings among translated DBL α types obtained from the Amele population. This was done by observing the Amele types that clustered with 3D7 DBL α sequences upon which the groupings were partly based [24,25]. We observed that Amele DBL α types clustered significantly with all six groups identified for the DBL α CIDR α region of *var* genes [25] (Figure S7). Therefore, sequences with high levels of similarity to the DBL α domains of *var* gene structural groups previously defined in the 3D7 genome [25] were present in the Amele population. It should be noted that the structural groupings were based upon the DBL α plus CIDR α head-structure sequences [25], whereas we compared only DBL α sequences; therefore, the only group that formed a significant cluster was group A (Figures 3A and S7). Recently, an alternative grouping of *var* genes has been published based on the number of cysteine residues and short amino acid motifs within the DBL α domain [31]. We found all six of these alternative groups within the Amele population (Table S2). Therefore, the degenerate PCR primers amplified a broad range of DBL α sequences from this natural population. These results demonstrate that the functional specialization and restricted evolution of *var* gene groups suggested from analysis of a single genome [24] and African clinical isolates [31] were also observed in a random sample of a natural population of *P. falciparum* from PNG.

Serological analyses have predicted that the *var* genes expressed during severe disease are conserved among clinical parasite isolates [38]. Recent investigations have shown group A *var* genes to be associated with the rosetting phenotype and severe disease [31,39–41]. Although the most conserved (i.e., high prevalence) *var* genes in the Amele population were not restricted to a particular *var* gene group, at least four group A *var* genes were found in more than 20% of isolates (Table S2). These are clearly of interest as candidate vaccine targets.

The *var1csa* gene, previously found to have a global distribution [42–44], was detected in our dataset. Our analyses confirmed that *var1csa* had a high prevalence globally, but this gene was rare among Amele isolates (Table S2). This further demonstrates geographic differences among the two populations.

Discussion

While *var* gene sequences have been extensively explored in terms of conserved features associated with adhesion [24–26,31,43,45–47], we have examined the primary sequence diversity because of its role in evasion of host immunity and

consequent *P. falciparum* transmission. We present a population genomic framework to randomly sample the diversity of *var* genes by sequencing the ubiquitous marker, DBL α , in natural *P. falciparum* populations. The framework can be considered a molecular epidemiological tool for malaria with the specific application to diverse multigene families. It defines the diversity and distribution of *var* genes in natural populations. The framework could also be applicable to other diverse systems such as additional *var* gene domains (e.g., DBL α , CIDR); other multigene families encoding putative malaria surface antigens (e.g., *vir* genes in *Plasmodium vivax*); and the immune evasion genes of other pathogens (e.g., variant surface glycoprotein [*vsug*] genes of *Trypanosoma brucei*). Importantly, the cumulative diversity curve addressed the issue of incomplete sampling of individual repertoires by defining the number of isolates that were needed to see the majority of diversity in a local population (i.e., 30 isolates for Amele). If multiple infections are prevalent, which is true for many malaria populations [48–50], the extent of *var* gene diversity for the population could still be defined by plotting the cumulative curve.

Analysis of *var* gene diversity on a global and local scale using the above-described framework showed very different patterns of diversity. We were unable to sample all of the diversity in the global population even though we compared over 1,000 DBL α sequences. The global range of *var* gene diversity is likely to be so large that it would take an enormous number of isolates to observe its full extent. When *var* genes were sampled from the local population of Amele, a region less than 10 km in diameter in Madang Province, PNG, the majority of *var* gene diversity was defined. Whether this is a representation of the *var* gene diversity of Madang, or PNG in general, remains to be determined by further spatial sampling within PNG. The variation between populations indicates important differences in their evolutionary history. For example, the presence of high frequency types in Amele might indicate the effect of selective sweeps in parts of the genome that led to the fixation of specific *var* types. In addition, reduced population size, population bottlenecks, and lower levels of multiple infection may all potentially have played a role in reducing *var* gene repertoire diversity within PNG. Unfortunately, it is only by obtaining full genomic sequences for multiple isolates for each locality that the different hypotheses can be compared. Nevertheless, our findings have important implications for the understanding of local immunity in different geographic localities.

Analysis of relationships among the DBL α types found in Amele showed that *P. falciparum* has unprecedented levels of genetic diversity of a major surface antigen on a small geographic scale. This finding is particularly striking when compared to data on other prevalent human pathogens sampled locally and globally. The linear phylogenies of *ha* and *env* genes of Influenza A and HIV, respectively, are highly structured such that ancestral sequences can be traced even among isolates from distant locations [1,2], whereas the DBL α types from Amele were so diverse that the phylogeny showed only distant relationships. We have sequenced only a 500-bp fragment (DBL α) as a marker of *var* genes (>5 kb) in a limited number of global isolates and in one local population. Thus, we have described only the “tip of the iceberg” of total *var* gene diversity. This gives an indication of the vast archive of

antigenic diversity available to *P. falciparum* and may explain why immunity to malaria is nonsterilizing and develops slowly.

To determine whether all *var* genes sampled are under similar selective pressures, it is important to know whether or not they are expressed during natural infection. In our dataset, the presence of pseudogenes could only be defined by frame shifts or stop codons in the DBL α domain. *T. brucei* has many *vsg* pseudogenes that have been proposed to be an archive of further antigenic diversity [51]. It is conceivable that *P. falciparum* also recombines fragments of *var* pseudogenes into functional *var* genes to increase its antigenic repertoire. For this reason, we included putative pseudogenes in the population genomic analysis of *var* genes where possible. In addition, only one or a few members of the *var* multigene family is expressed at a time [6]; therefore, sampling all possible transcripts from the population will require many times more sampling than for the present study. We therefore compared the Amele types to those found to be expressed in clinical isolates from the nearby population of Maiwara [32]. Interestingly, we found that 4.3% of the Amele DBL α types were expressed in the Maiwara isolates (Table S2). When a neighbor-joining tree was constructed, 22.4% of Amele DBL α types clustered with significant bootstrap support with the expressed sequences from Maiwara (unpublished data). This indicates that many of the *var* genes sampled in the Amele population were likely to be functional genes.

Our investigation of the sequence diversity among *var* genes in the Amele local population suggested that obtaining further types from Amele will only identify additional diverse *var* genes with presumably novel immunological properties. Previous studies have suggested that recombination occurs among *var* genes by analyzing distantly related genomes [21,22] and the progeny of a genetic cross [20]. Our results, namely the phylogenetic and pairwise alignment quality score analyses, suggested that recombination is an important factor in generating polymorphism among the *var* genes within a natural population. Ultimately, mutation must be the source of amino acid changes. However, the generation of novel combinations to which the host immune system has not previously been exposed is likely to be equally important to immune evasion. The complex nature of the sequence data requires new statistical tools to further define recombination among *var* genes in natural populations. As mentioned earlier, we are currently addressing this issue.

The restriction of *var* gene diversity in PNG, *var* genes in PNG which were predominantly unique to that location and the extremely limited overlap among global repertoires points to the existence of geographic population structure. Such structure has been observed for microsatellite [52] and single nucleotide polymorphism [53] markers. The majority of the broadly distributed *var* genes were found in Africa, but this may be an artifact because almost half of the global population (29 isolates) was from Africa giving a higher probability of finding matching *var* genes in that region. As *P. falciparum* originated in Africa [54], it is likely that the PNG population is a subpopulation of the ancestral population. The possibility of such geographic population structure warrants further investigation because genomes with different *var* repertoires may be introduced to other transmission systems with consequences for immune evasion and malarial disease. An alternative explanation is temporal variation

among the Amele isolates collected in 1999 and the global isolates collected over a period of 20 y. However, our data also showed that certain *var* genes were maintained in the PNG population over long periods of time. Given the relative importance of *var* genes to *P. falciparum* survival [10–11], host immunity [12–19], and the potential utility of certain PfEMP1 subgroups [39,45,55] and domains [56–58] for malaria vaccination, it will now be a priority to observe spatial and temporal variation on small and large geographic scales using our validated molecular epidemiological framework.

Materials and Methods

Parasite isolates and cloned lines. For global sampling, 25 *P. falciparum* isolates or cloned lines were chosen for their single distinct genotypes and diverse origins. From Africa there were three field isolates from Zimbabwe (collected by S. Mharakurwa from Sahumani Province, Zimbabwe in 1999. Ethical approval was obtained from the Medical Research Council of Zimbabwe); four field isolates from Dienga in Southeast Gabon, 50 km from Bakoumba (collected by FMN, ethical approval was obtained by the Ethics Committee for Research in Human Medicine, International Center for Medical Research, Gabon); D6 cloned from isolate Sierra Leone I/CDC from Sierra Leone, West Africa [59]; isolates K39, M24, and 3118 from Kenya (donated by W. Watkins, Wellcome Trust Research Laboratories, Kenyatta National Hospital, Nairobi, Kenya). PNG isolates were KF1776 (B. Tiwari, unpublished data), KF1916 [13], and Muz12, Muz37, Muz51, and Muz106 [60] collected from Amele, Madang Province, PNG, in 1986 and 1990, respectively. From Asia there were isolates SK44, PO18, PO19, and AMB47 from Thailand (donated by Wellcome Trust Research Laboratories, Faculty of Tropical Medicine, Mahidol University, Bangkok, Thailand); W2 cloned from Indochina III/CDC, isolated from a Lao refugee in 1980 [59]; isolate MRC20 from India (donated by H. Joshi, S. K. Subbarao, and C. R. Pillai, Malaria Research Centre, Delhi). From Latin America there was HB3 cloned from the isolate Honduras-I/CDC and isolated in Honduras, Central America, in 1980 [61]; and 7G8 cloned from IMTM22 isolated in Manaus, Brazil, in 1980 [62]. Parasites were cultured as described previously [13]. W2, 7G8, and D6 clones were propagated in the laboratory of D. F. Wirth, Harvard School of Public Health.

DBL α sequences downloaded from the GenBank database consisted of a total of 480 sequences from 34 isolates. These included 3D7 (number of DBL α sequences (n) = 57) from the Malaria Genome Project [34]; three isolates from India: R35 (n = 9), R15 (n = 9), and R1 (n = 7) [63]; four isolates from Sudan: SD101 (n = 2), SD105 (n = 14), SD106 (n = 2), and SD126 (n = 4) [22]; ten isolates from Kenya: KEN1 (n = 2), KEN2 (n = 8), KEN3 (n = 9), KEN4 (n = 6), KEN5 (n = 1), KEN13 (n = 5), KEN15 (n = 5), KEN16 (n = 6), KEN19 (n = 11), and KEN20 (n = 2) [33]; one isolate from Cape Verde: CV (n = 8) [33]; four isolates from the Solomon Islands: S2 (n = 28), S55 (n = 22), S99 (n = 25), and N72 (n = 15) [27]; two isolates from the Philippines: PH1 (n = 26) and PH4 (n = 29) [27]; two isolates from Vanuatu: VAN232 (n = 4) and VAN230 (n = 3) [34]; one isolate from PNG: AN143 (n = 22); one isolate from Thailand: Dd2 (n = 22) [27]; one isolate from Africa: NF36 (n = 23) [27]; three isolates from Venezuela: VZ1 (n = 32), VZ2 (n = 15), and VZ3 (n = 14) [28]; and one isolate from Brazil: A4 (n = 33) [33].

Epidemiological sampling of a local population involved the selection of isolates from 1,082 field isolates collected for other studies in our laboratory [64]. This local population was called "Amele." Field isolates were randomly collected in November/December of 1999 from residents of a cluster of six Amele villages in Madang on the north coast of PNG, where intense transmission of *P. falciparum* malaria occurs [65]. Daily inoculation rates range from 0.19–1.44 infective bites/person/night (68–526 per y) [66]. Amele isolates were collected as blood samples from children aged between 6 mo and 11 y, with asymptomatic infection. Ethical approval for the study was obtained from the Medical Research Advisory Committee of PNG.

DNA extractions. Whole genomic DNA was obtained from laboratory isolates as previously described [67]. Field isolate genomic DNA was obtained by one of two methods. Isolates from Gabon and Zimbabwe (from the global collection) were stored as whole blood spots on 3MM filter paper (Whatman, <http://www.whatman.com>), and DNA was extracted using the Chelex method. Field isolates from Amele were stored in Guanidine and extracted using Glassmilk (Qbiogene, <http://www.qbiogene.com>). Isolates or cloned lines were

genotyped to confirm that they contained single, distinct genomes. This was done by analysis with 12 genome-wide microsatellite markers [68] and/or MSP2 fingerprinting [69].

High-throughput DBL α sequencing. The DBL α sequencing protocol is outlined in Figure S1A. DBL α repertoires were amplified with two sets of degenerate primers to homology blocks D (forward primer) and H (reverse primer) [26]. One set of primers known as “AF” and “BR” were described previously [33]. PCR conditions were 10 ng (cultured isolate) or 1 μ l (field sample) genomic DNA, 2 mM dNTPs, 2.4 mM MgCl₂, 50 pmol each primer, 2.5 Units Taq Polymerase (Promega, <http://www.promega.com>). Reactions were performed in a Perkin Elmer 9600 thermal cycler at 95 °C for 4 min, 30 cycles of 95 °C for 1 min, 42 °C for 1 min, 60 °C for 1 min, and a final cycle of 60 °C for 7 min. A second set of primers, known as “AB” (DBL α AB-F: AGRAGYTTYGCNGAYATHGG and DBL α AB-R: AACCACTYAARTAYTGNGG) were used to correct the amplification of nonspecific human sequences from field samples, which we found to be a problem with the AF/BR primers. PCR conditions were 10 ng (cultured isolate) or 1 μ l (field sample) genomic DNA, 2 mM dNTPs, 3.5 mM MgCl₂, 20 pmol each primer, 2 units AmpliTaq Gold DNA Polymerase (Applied Biosystems, <http://www.appliedbiosystems.com>). Reactions were performed in a Perkin Elmer 9600 thermal cycler at 95 °C for 10 min; 35 cycles of 95 °C for 30 s; 48 °C for 30 s; 60 °C for 45 s; and a final cycle at 60 °C for 7 min. PCR products of approximately 350–450 bp were cloned using the TopoTA PCR cloning kit (Invitrogen, <http://www.invitrogen.com>) according to the manufacturer’s instructions. At least 96 colonies were picked from each cloning experiment and cultured in 1 ml of LB broth supplemented with 25 μ g/ml of Kanamycin in deep-well culture plates. Plasmid DNA was extracted using the Millipore 96-well plasmid extraction kit (Millipore, <http://www.millipore.com>) according to the manufacturer’s instructions, except for the vacuum steps that were replaced with a single centrifugation step encompassing both lysate removal and DNA binding. The bacterial lysate was added to the lysate clearing plate, which was then stacked on a plasmid plate with a waste collection plate on the bottom. Centrifuge adapters for stable stacking of plates are available from Millipore. Centrifugation for 40 min resulted in the plasmid DNA binding to the plasmid plate. This was followed by two wash cycles as for the vacuum protocol. Up to eight plates (768 clones) could be extracted in a few hours using this adaptation. Cycle sequencing was performed using the Big Dye Terminator Sequence Reaction mix (Applied Biosystems) according to the manufacturer’s instructions, for both strands using M13 Universal forward and reverse primers. Sequencing reactions were run on an ABI3700 (Applied Biosystems) at the core sequencing center at the Department of Zoology, University of Oxford.

Sequence analysis. The primary sequence analysis protocol is outlined in Figure S1B and describes how raw sequence data was processed. Raw sequence data for each isolate was screened for vector contamination and end-trimmed where necessary using the automated functions in the DNA sequence analysis program Sequencher (Gene Codes, <http://www.genecodes.com>). Alignment was then performed at 96% DNA sequence identity to allow DBL α sequences from the same gene to align yet separate sequences from different genes. This limit was based on an analysis of data from two isolates: 3D7 [33] and 1776 (B. Tiwari, unpublished data) available from GenBank at the time of project design. Available sequences from these two isolates were aligned at incremented percentage similarity until they no longer aligned to each other, giving a value of 96%. If duplicate genes are present and diverged only within the 96%–100% identity range, these will be sampled as one *var* gene. Hence, the number of DBL α sequences within an isolate may also be a measure of within-isolate diversity if each isolate is sampled equally. Note that these numbers may also be normalized for sequence input by dividing by the number of reads. To determine whether this was a possibility, the 59 3D7 gene sequences were aligned at 96% identity. A total of 54 types remained after alignment. This showed that there was redundancy within this genome, and thus there may be redundancy within other genomes. After alignment as described, the resulting consensus DBL α sequences were considered a sample of the *var* gene repertoire. Multiple sequence reads for each consensus sequence reduced the possibility of PCR, cloning, and sequencing artifacts.

For definition of types, comparison among different *var* gene repertoires was performed. Again a 96% identity cut-off was used to align matching consensus DBL α sequences among repertoires. The number of unique DBL α sequences (types) was then defined for the population (i.e., a group of isolates). *Var* genotypes were then defined based on the range of DBL α types (Figure S1B).

To summarize the relatedness between the *var* gene repertoires from two isolates, we used a simple statistic PTS. If isolate A has a repertoire

of n_A distinct types and isolate B has a repertoire of n_B distinct types, and a total of n_{AB} types are shared by the two isolates, we define:

$$PTS_{AB} = \frac{2n_{AB}}{n_A + n_B}$$

Because the isolate repertoires are not fully sampled, estimates of PTS from empirical data are likely to be downward biased. However, as long as there are not systematic differences between the number of reads obtained for isolates from different geographic locations the sample estimates of PTS are still useful statistics on which to base a comparative analysis of diversity. Note that if each isolate had a repertoire of exactly one *var* gene, sample PTS is equivalent to expected homozygosity, and the apportionment of PTS within and between populations would be equivalent to measuring Wright’s fixation indices. PTS can therefore be thought of as a simple extension of F_{ST} to the case where multiple gene family members of unknown allelic status have been collected.

DBL α types were translated to amino acid sequences using the program Transeq from the EMBOSS toolkit at the European Bioinformatics Institute (<http://www.ebi.ac.uk>). Translations that contained two or more stop codons were not analyzed further as this may indicate a frame shift and may artificially result in a lack of homology to other DBL α protein sequences.

Phylogenetic analysis of DBL α types was done by constructing multiple alignments of translated DBL α types using stand-alone Clustal W downloaded from the European Bioinformatics Institute Website (<http://www.ebi.ac.uk>) [70]. Multiple alignments were then imported into MEGA version 3.0 [71] where neighbor-joining trees were constructed using the p-distance model to define pairwise distances between sequences. Primer sequences were removed for the phylogenetic analysis.

Pairwise alignments of “good” protein sequences (described above) were constructed using the GCG (Genetics Computer Group, <http://www.accelrys.com/products/gcg>) program “BestFit.” BestFit makes an optimal alignment of the best segment of similarity between two sequences using a Smith and Waterman local homology algorithm. Percent identity and percent similarity are indicators of the quality of the alignment. For scoring similarities among protein sequences, BestFit uses the blosum62 comparison matrix [72]. We also examined quality scores because percent identity or similarity scores ignore the length of the alignment. Quality scores for pairwise alignments are related to the degree of similarity but also account for length of the alignment and gaps. These will more accurately quantify the relationship among potentially recombined sequences. For example, a mosaic sequence may have 50% similarity to each of its donors across the entire length of the alignment. This would be considered equivalent to the similarity among a pair of sequences with 50% differences spread across the length of the alignment. However, the pairing of a mosaic sequence with its donor will have a higher quality score because it will score 100% similarity for half of the alignment. The calculation of pairwise quality scores was done using the following formula:

$$\begin{aligned} \text{Quality} = & [\text{CmpVal}(\text{AA}) \times \text{Total}(\text{AA}) + \text{CmpVal}(\text{AB}) \times \text{Total}(\text{AB}) + \\ & \text{CmpVal}(\text{AC}) \times \text{Total}(\text{AC}) \dots + \text{CmpVal}(\text{ZZ}) \times \text{Total}(\text{ZZ})] \\ & - (\text{GapCreationPenalty} \times \text{GapNumber}) - (\text{GapExtensionPenalty} \times \text{TotalLengthOfGaps}) \end{aligned}$$

Where: “CmpVal” is the comparison value for an amino acid pair based on an amino acid comparison table (PAM matrix) and “Total” is the total number of matches for that pair in the alignment. Therefore, each match in the alignment adds points to the quality score and mismatches and gaps subtract from it.

Supporting Information

Figure S1. A Sampling Framework for the *var* Multigene Family

(A) High-throughput sequencing and (B) population genomic analysis of *var* gene sequences.

Found at doi:10.1371/journal.ppat.0030034.sg001 (47 KB PPT).

Figure S2. Sampling Results for Clone 3D7 and HB3

Distribution of DBL α sequences among reads using (A) 3D7 with AFBR primers [2]; (B) 3D7 with AB primers; (C) HB3 with AFBR primers [33]; or (D) HB3 with AB primers (see Materials and Methods). Bars are colored to represent distinct DBL α CIDR α groups as per Lavsten et al. [25]. Asterisks show sequences that were degenerate or recombined in our data compared to the genome sequence.

Found at doi:10.1371/journal.ppat.0030034.sg002 (328 KB PPT).

Figure S3. Sampling Efficiency

The number of sequences obtained for each repertoire was plotted against the number of sequence reads (both forward and reverse) for global and Amele populations. Each point on the plot represents the data obtained for one PCR reaction after cloning. Up to two microtitre plates (284 sequence reads) of clones were sequenced for some isolates and lines, therefore a few points on the plot have more than 192 sequence reads. The variable number of sequence reads shown was partially due to failed sequencing reactions or erroneous amplification of non-DBL α sequences, which were discarded. Each *var* repertoire size (number of DBL α sequences obtained) varied as a result of this and other variables, such as actual repertoire size [27], PCR amplification, and DNA quantity in the field isolates.

Found at doi:10.1371/journal.ppat.0030034.sg003 (52 KB PPT).

Figure S4. Analysis of Primer Bias

Scatterplot of the relationship between the average number of reads contributing to a type and the frequency of that type in the population. Extreme outliers can be seen above the dashed line. Arrows denote high frequency outliers.

Found at doi:10.1371/journal.ppat.0030034.sg004 (30 KB JPG).

Figure S5. Raw Data for PTS

Matrix of the results for pairwise comparisons of *var* gene (DBL α) repertoires showing the absolute number of DBL α sequences found to be shared among a pair of isolates in the upper right hand side of the matrix; and the PTS statistic for the pair in the lower left hand side of the matrix. Self-self values are shaded in yellow.

Found at doi:10.1371/journal.ppat.0030034.sg005 (87 KB XLS).

Figure S6. Multiple Alignment of Amele DBL α Types

Amino acid sequences for a total of 177 “good” DBL α types, defined by the absence of frame shifts and aligned using Clustal X version 1.83 [70] using the parameters: gap opening penalty = 5; gap extension penalty = 0.05.

Found at doi:10.1371/journal.ppat.0030034.sg006 (55 KB PDF).

Figure S7. Prediction of *var* Gene Structural Groups within the Amele Population

A multiple alignment of the translated DBL α types was constructed in Clustal W [70] with parameters set at gap opening = 5 and gap extension = 0.05, for all “good” DBL α types in the Amele population ($n = 177$) with DBL α amino acid sequences from the 3D7 *var* genes

References

- McCutchan FE, Artenstein AW, Sanders-Buell E, Salminen MO, Carr JK, et al. (1996) Diversity of the envelope glycoprotein among human immunodeficiency virus type 1 isolates of clade E from Asia and Africa. *J Virol* 70: 3331–3338.
- Zambon MC (1999) Epidemiology and pathogenesis of influenza. *J Antimicrob Chemother* 44 (Suppl B): 3–9.
- Baruch DI, Pasloske BL, Singh HB, Bi X, Ma XC, et al. (1995) Cloning the *P. falciparum* gene encoding PfEMP1, a malarial variant antigen and adherence receptor on the surface of parasitized human erythrocytes. *Cell* 82: 77–87.
- Smith JD, Chitnis CE, Craig AG, Roberts DJ, Hudson-Taylor DE, et al. (1995) Switches in expression of *Plasmodium falciparum var* genes correlate with changes in antigenic and cytoadherent phenotypes of infected erythrocytes. *Cell* 82: 101–110.
- Su XZ, Heatwole VM, Wertheimer SP, Guinet F, Herrfeldt J A, et al. (1995) The large diverse gene family *var* encodes proteins involved in cytoadherence and antigenic variation of *Plasmodium falciparum*-infected erythrocytes. *Cell* 82: 89–100.
- Chen Q, Fernandez V, Sundstrom A, Schlichtherle M, Datta S, et al. (1998) Developmental selection of *var* gene expression in *Plasmodium falciparum*. *Nature* 394: 392–395.
- Hayward RE, Tiwari B, Piper KP, Baruch DI, Day KP (1999) Virulence and transmission success of the malarial parasite *Plasmodium falciparum*. *Proc Natl Acad Sci U S A* 96: 4563–4568.
- Chen Q, Schlichtherle M, Wahlgren M (2000) Molecular aspects of severe malaria. *Clin Microbiol Rev* 13: 439–450.
- Day KP, Marsh K (1991) Naturally acquired immunity to *Plasmodium falciparum*. *Immunol Today* 12: A68–A71.
- Gupta S, Trenholme K, Anderson RM, Day KP (1994) Antigenic diversity and the transmission dynamics of *Plasmodium falciparum*. *Science* 263: 961–963.
- Marsh K, Howard RJ (1986) Antigens induced on erythrocytes by *P.*

[34]. The neighbor-joining tree was constructed from this alignment in MEGA version 3.0 [71] using the p-distance algorithm, pairwise alignment, and 1,000 bootstrapping repetitions. The consensus tree was condensed to show only clusters with >50% bootstrap support. Primer sequences were removed prior to drawing the tree. A key is shown to indicate the specific DBL α CIDR α sequence groupings (A–E and X) [25].

Found at doi:10.1371/journal.ppat.0030034.sg007 (942 KB PPT).

Table S1. GenBank Accession Numbers for New Sequences

Found at doi:10.1371/journal.ppat.0030034.st001 (95 KB XLS).

Table S2. Properties of Amele DBL α Types

Found at doi:10.1371/journal.ppat.0030034.st002 (343 KB DOC).

Text S1. Validation of a Sampling Framework for the *var* Multigene Family

Found at doi:10.1371/journal.ppat.0030034.sd001 (55 KB DOC).

Text S2. Primer Bias

Found at doi:10.1371/journal.ppat.0030034.sd002 (33 KB DOC).

Acknowledgments

The authors wish to acknowledge the excellent technical support of L. Richardson, M. Golding, L. Pilling, C. Williams, and T. Dickinson. We are grateful to the Broad Institute for making HB3 genome sequence data publicly available before publication (*Plasmodium falciparum* HB3 Sequencing Project, Broad Institute of Harvard and MIT, <http://www.broad.mit.edu>). We would also like to acknowledge the anonymous reviewers whose comments resulted in a much-improved manuscript.

Author contributions. AEB and KPD conceived and designed the experiments. AEB and ALS performed the experiments. AEB, GAVM, and KPD analyzed the data. AEB, LT, HI, FMN, SMB, and GAVM contributed reagents/materials/analysis tools. AEB, GAVM, and KPD wrote the paper.

Funding. This work was supported by a Wellcome Trust program grant and US National Institutes of Health/National Institute of General Medical Sciences grant GM061351-07. AEB was supported by a Howard Florey Postdoctoral Fellowship from the Royal Society (UK) and National Health and Medical Research Council (Australia), 2001–2003.

Competing interests. The authors have declared that no competing interests exist.

- falciparum*: Expression of diverse and conserved determinants. *Science* 231: 150–153.
- Bull PC, Lowe BS, Kortok M, Molyneux CS, Newbold CI, et al. (1998) Parasite antigens on the infected red cell surface are targets for naturally acquired immunity to malaria. *Nat Med* 4: 358–360.
- Forsyth KP, Philip G, Smith T, Kum E, Southwell B, et al. (1989) Diversity of antigens expressed on the surface of erythrocytes infected with mature *Plasmodium falciparum* parasites in Papua New Guinea. *Am J Trop Med Hyg* 41: 259–265.
- Bruce MC, Day KP (2003) Cross-species regulation of *Plasmodium* parasitemia in semi-immune children from Papua New Guinea. *Trends Parasitol* 19: 271–277.
- Molineaux L, Trauble M, Collins WE, Jeffery GM, Dietz K (2002) Malaria therapy reinoculation data suggest individual variation of an innate immune response and independent acquisition of antiparasitic and antitoxic immunities. *Trans R Soc Trop Med Hyg* 96: 205–209.
- Piper KP, Roberts DJ, Day KP (1999) *Plasmodium falciparum*: Analysis of the antibody specificity to the surface of the trophozoite-infected erythrocyte. *Exp Parasitol* 91: 161–169.
- Babiker HA, Ranford-Cartwright LC, Currie D, Charlwood JD, Billingsley P, et al. (1994) Random mating in a natural population of the malaria parasite *Plasmodium falciparum*. *Parasitology* 109 (Pt 4): 413–421.
- Paul RE, Packer MJ, Walmsley M, Lagog M, Ranford-Cartwright LC, et al. (1995) Mating patterns in malaria parasite populations of Papua New Guinea. *Science* 269: 1709–1711.
- Su X, Ferdig MT, Huang Y, Huynh CQ, Liu A, et al. (1999) A genetic map and recombination parameters of the human malaria parasite *Plasmodium falciparum*. *Science* 286: 1351–1353.
- Freitas-Junior LH, Bottius E, Pirrit LA, Deitsch KW, Scheidig C, et al. (2000) Frequent ectopic recombination of virulence factor genes in telomeric chromosome clusters of *P. falciparum*. *Nature* 407: 1018–1022.
- Taylor HM, Kyes SA, Newbold CI (2000) *Var* gene diversity in *Plasmodium*

- falciparum* is generated by frequent recombination events. *Mol Biochem Parasitol* 110: 391–397.
22. Ward CP, Clotey GT, Dorris M, Ji DD, Arnot DE (1999) Analysis of *Plasmodium falciparum* PfEMP-1/var genes suggests that recombination rearranges constrained sequences. *Mol Biochem Parasitol* 102: 167–177.
 23. WHO-UNAIDS Vaccine Advisory Committee (2001) Approaches to the development of broadly protective HIV vaccines: Challenges posed by the genetic, biological, and antigenic variability of HIV-1. Report from a meeting of the WHO-UNAIDS Vaccine Advisory Committee. Geneva, Switzerland. 21–23 February 2000. *AIDS* 15: W1–W25.
 24. Kraemer SM, Smith JD (2003) Evidence for the importance of genetic structuring to the structural and functional specialization of the *Plasmodium falciparum* var gene family. *Mol Microbiol* 50: 1527–1538.
 25. Lavstsen T, Salanti A, Jensen AT, Arnot DE, Theander TG (2003) Subgrouping of *Plasmodium falciparum* 3D7 var genes based on sequence analysis of coding and noncoding regions. *Malar J* 2: 27.
 26. Smith JD, Subramanian G, Gamain B, Baruch DI, Miller LH (2000) Classification of adhesive domains in the *Plasmodium falciparum* erythrocyte membrane protein 1 family. *Mol Biochem Parasitol* 110: 293–310.
 27. Fowler EV, Peters JM, Gattton ML, Chen N, Cheng Q (2002) Genetic diversity of the DBLalpha region in *Plasmodium falciparum* var genes among Asia-Pacific isolates. *Mol Biochem Parasitol* 120: 117–126.
 28. Tami A, Ord R, Targett GA, Sutherland CJ (2003) Sympatric *Plasmodium falciparum* isolates from Venezuela have structured var gene repertoires. *Malar J* 2: 7.
 29. Kirchgatter K, Mosbach R, del Portillo HA (2000) *Plasmodium falciparum*: DBL-1 var sequence analysis in field isolates from central Brazil. *Exp Parasitol* 95: 154–157.
 30. Kyes S, Taylor H, Craig A, Marsh K, Newbold C (1997) Genomic representation of var gene sequences in *Plasmodium falciparum* field isolates from different geographic regions. *Mol Biochem Parasitol* 87: 235–238.
 31. Bull PC, Berriman M, Kyes S, Quail MA, Hall N, et al. (2005) *Plasmodium falciparum* variant surface antigen expression patterns during malaria. *PLoS Pathog* 1: e26. doi:10.1371/journal.ppat.0010026
 32. Kaestli M, Cortes A, Lagom M, Ott M, Beck HP (2004) Longitudinal assessment of *Plasmodium falciparum* var gene transcription in naturally infected asymptomatic children in Papua New Guinea. *J Infect Dis* 189: 1942–1951.
 33. Taylor HM, Kyes SA, Harris D, Kriek N, Newbold CI (2000) A study of var gene transcription in vitro using universal var gene primers. *Mol Biochem Parasitol* 105: 13–23.
 34. Gardner MJ, Hall N, Fung E, White O, Berriman M, et al. (2002) Genome sequence of the human malaria parasite *Plasmodium falciparum*. *Nature* 419: 498–511.
 35. Walther BA, Morand S (1998) Comparative performance of species richness estimation methods. *Parasitology* 116 (Pt 4): 395–405.
 36. Lewontin RC (1967) An estimate of average heterozygosity in man. *Am J Hum Genet* 19: 681–685.
 37. Schierup MH, Hein J (2000) Consequences of recombination on traditional phylogenetic analysis. *Genetics* 156: 879–891.
 38. Bull PC, Kortok M, Kai O, Ndungu F, Ross A, et al. (2000) *Plasmodium falciparum*-infected erythrocytes: Agglutination by diverse Kenyan plasma is associated with severe disease and young host age. *J Infect Dis* 182: 252–259.
 39. Kaestli M, Cockburn IA, Cortes A, Baea K, Rowe JA, et al. (2006) Virulence of malaria is associated with differential expression of *Plasmodium falciparum* var gene subgroups in a case-control study. *J Infect Dis* 193: 1567–1574.
 40. Kyriacou HM, Stone GN, Challis RJ, Raza A, Lyke KE, et al. (2006) Differential var gene transcription in *Plasmodium falciparum* isolates from patients with cerebral malaria compared to hyperparasitaemia. *Mol Biochem Parasitol* 150: 211–218.
 41. Rottmann M, Lavstsen T, Mugasa JP, Kaestli M, Jensen A T, et al. (2006) Differential expression of var gene groups is associated with morbidity caused by *Plasmodium falciparum* infection in Tanzanian children. *Infect Immun* 74: 3904–3911.
 42. Rowe JA, Kyes SA, Rogerson SJ, Babiker HA, Raza A (2002) Identification of a conserved *Plasmodium falciparum* var gene implicated in malaria in pregnancy. *J Infect Dis* 185: 1207–1211.
 43. Salanti A, Jensen AT, Zornig HD, Staalsoe T, Joergensen L, et al. (2002) A sub-family of common and highly conserved *Plasmodium falciparum* var genes. *Mol Biochem Parasitol* 122: 111–115.
 44. Winter G, Chen Q, Flick K, Kremsner P, Fernandez V, et al. (2003) The 3D7var5.2 (var COMMON) type var gene family is commonly expressed in non-placental *Plasmodium falciparum* malaria. *Mol Biochem Parasitol* 127: 179–191.
 45. Salanti A, Staalsoe T, Lavstsen T, Jensen AT, Sowa MP, et al. (2003) Selective upregulation of a single distinctly structured var gene in chondroitin sulphate A-adhering *Plasmodium falciparum* involved in pregnancy-associated malaria. *Mol Microbiol* 49: 179–191.
 46. Rowe JA, Kyes SA (2004) The role of *Plasmodium falciparum* var genes in malaria in pregnancy. *Mol Microbiol* 53: 1011–1019.
 47. Trimmell AR, Kraemer SM, Mukherjee S, Phippard DJ, James JH, et al. (2006) Global genetic diversity and evolution of var genes associated with placental and severe childhood malaria. *Mol Biochem Parasitol* 148: 169–180.
 48. Bruce MC, Galinski MR, Barnwell JW, Donnelly CA, Walmsley M, et al. (2000) Genetic diversity and dynamics of *Plasmodium falciparum* and *P. vivax* populations in multiply infected children with asymptomatic malaria infections in Papua New Guinea. *Parasitology* 121 (Pt 3): 257–272.
 49. Owusu-Agyei S, Smith T, Beck HP, Amenga-Etego L, Felger I (2002) Molecular epidemiology of *Plasmodium falciparum* infections among asymptomatic inhabitants of a holoendemic malarious area in northern Ghana. *Trop Med Int Health* 7: 421–428.
 50. Paul RE, Brockman A, Price RN, Luxemburger C, White NJ, et al. (1999) Genetic analysis of *Plasmodium falciparum* infections on the north-western border of Thailand. *Trans R Soc Trop Med Hyg* 93: 587–593.
 51. Berriman M, Ghedin E, Hertz-Fowler C, Blandin G, Renaud H, et al. (2005) The genome of the African trypanosome *Trypanosoma brucei*. *Science* 309: 416–422.
 52. Anderson TJ, Haubold B, Williams JT, Estrada-Franco JG, Richardson L, et al. (2000) Microsatellite markers reveal a spectrum of population structures in the malaria parasite *Plasmodium falciparum*. *Mol Biol Evol* 17: 1467–1482.
 53. Mu J, Awadalla P, Duan J, McGee KM, Joy DA, et al. (2005) Recombination hotspots and population structure in *Plasmodium falciparum*. *PLoS Biol* 3: e335. doi:10.1371/journal.pbio.0030335
 54. Joy DA, Feng X, Mu J, Furuya T, Chotivanich K, et al. (2003) Early origin and recent expansion of *Plasmodium falciparum*. *Science* 300: 318–321.
 55. Jensen AT, Magistrado P, Sharp S, Joergensen L, Lavstsen T, et al. (2004) *Plasmodium falciparum* associated with severe childhood malaria preferentially expresses PfEMP1 encoded by group A var genes. *J Exp Med* 199: 1179–1190.
 56. Baruch DI, Ma XC, Singh HB, Bi X, Pasloske B, et al. (1997) Identification of a region of PfEMP1 that mediates adherence of *Plasmodium falciparum*-infected erythrocytes to CD36: Conserved function with variant sequence. *Blood* 90: 3766–3775.
 57. Chen Q, Heddiini A, Barragan A, Fernandez V, Pearce SF, et al. (2000) The semiconserved head structure of *Plasmodium falciparum* erythrocyte membrane protein 1 mediates binding to multiple independent host receptors. *J Exp Med* 192: 1–10.
 58. Smith JD, Craig AG, Kriek N, Hudson-Taylor D, Kyes S, et al. (2000) Identification of a *Plasmodium falciparum* intercellular adhesion molecule-1 binding domain: A parasite adhesion trait implicated in cerebral malaria. *Proc Natl Acad Sci U S A* 97: 1766–1771.
 59. Oduola AM, Milhous WK, Weatherly NF, Bowdre JH, Desjardins RE (1988) *Plasmodium falciparum*: Induction of resistance to mefloquine in cloned strains by continuous drug exposure in vitro. *Exp Parasitol* 67: 354–360.
 60. Cox MJ, Kum DE, Tavul L, Narara A, Raiko A, et al. (1994) Dynamics of malaria parasitaemia associated with febrile illness in children from a rural area of Madang, Papua New Guinea. *Trans R Soc Trop Med Hyg* 88: 191–197.
 61. Bhasin VK, Trager W (1984) Gametocyte-forming and non-gametocyte-forming clones of *Plasmodium falciparum*. *Am J Trop Med Hyg* 33: 534–537.
 62. Burkot TR, Williams JL, Schneider I (1984) Infectivity to mosquitoes of *Plasmodium falciparum* clones grown in vitro from the same isolate. *Trans R Soc Trop Med Hyg* 78: 339–341.
 63. Chattopadhyay R, Sharma A, Srivastava VK, Pati SS, Sharma SK, et al. (2003) *Plasmodium falciparum* infection elicits both variant-specific and cross-reactive antibodies against variant surface antigens. *Infect Immun* 71: 597–604.
 64. Imrie H, Fowkes F, Michon P, Tavul L, Hume J, et al. (2006) Haptoglobin levels are associated with haptoglobin genotype and alpha-thalassaemia in a malaria endemic area. *Am J Trop Med Hyg* 74: 965–971.
 65. Cattani JA, Tulloch JL, Vrbova H, Jolley D, Gibson FD, et al. (1986) The epidemiology of malaria in a population surrounding Madang, Papua New Guinea. *Am J Trop Med Hyg* 35: 3–15.
 66. Burkot TR, Graves PM, Paru R, Wirtz RA, Heywood PF (1988) Human malaria transmission studies in the *Anopheles punctulatus* complex in Papua New Guinea: Sporozoite rates, inoculation rates, and sporozoite densities. *Am J Trop Med Hyg* 39: 135–144.
 67. Ljungstrom I, Perlmann H, Schlichtherle M, Scherf A, Wahlgren M, editors (2004) *Methods in malaria research*. 4th edition. Manassas (Virginia): Malaria Research and Reference Reagent Resource Center. 248 p.
 68. Anderson TJ, Su XZ, Bockarie M, Lagom M, Day KP (1999) Twelve microsatellite markers for characterization of *Plasmodium falciparum* from finger-prick blood samples. *Parasitology* 119 (Pt 2): 113–125.
 69. Felger I, Tavul L, Beck HP (1993) *Plasmodium falciparum*: A rapid technique for genotyping the merozoite surface protein 2. *Exp Parasitol* 77: 372–375.
 70. Thompson JD, Higgins DG, Gibson TJ (1994) CLUSTAL W: Improving the sensitivity of progressive multiple sequence alignment through sequence weighting, position-specific gap penalties, and weight matrix choice. *Nucleic Acids Res* 22: 4673–4780.
 71. Kumar S, Tamura K, Nei M (2004) MEGA3: Integrated software for molecular evolutionary genetics analysis and sequence alignment. *Brief Bioinform* 5: 150–163.
 72. Henikoff S, Henikoff JG (1992) Amino acid substitution matrices from protein blocks. *Proc Natl Acad Sci U S A* 89: 10915–10919.