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Complete nucleotide sequence characterization of DRB5 alleles reveals a homogeneous allele group that is distinct from other DRB genes

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Abstract

Next Generation Sequencing allows for testing and typing of entire genes of the HLA region. A better and comprehensive sequence assessment can be achieved by the inclusion of full gene sequences of all the common alleles at a given locus. The common alleles of DRB5 are under-characterized with the full exon-intron sequence of two alleles available. In the present study the DRB5 genes from 18 subjects alleles were cloned and sequenced; haplotype analysis showed that 17 of them had a single copy of DRB5 and one consanguineous subject was homozygous at all HLA loci. Methodological approaches including robust and efficient long-range PCR amplification, molecular cloning, nucleotide sequencing and *de novo* sequence assembly were combined to characterize DRB5 alleles. DRB5 sequences covering from 5'UTR to the end of intron 5 were obtained for DRB5*01:01, 01:02 and 02:02; partial coverage including a segment

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Declaration of interest
none

Appendix A. Supplementary Data

Supplementary Materials and Methods: The HLA Database Construction protocol with evaluation and comparison with other algorithms.

Supplementary Table 1: Genomic sequences of cloned 5' fragment of DRB5 alleles aligned in reference to DRB5*01:01:01.

Supplementary Table 2: Genomic sequences of cloned 3' fragment of DRB5 alleles aligned in reference to DRB5*01:01:01.

Supplementary Table 3: Exon 2 to Intron 5 pairwise sequence alignment similarities between DRB5*01:01:01 with other DRB5 alleles and DRB1, DRB3, DRB4, DRB6 and DRB7 gene using Global alignment EMBOSS Needle.

Supplementary Table 4: Estimates of Evolutionary Divergence between DRB genes. The number of base substitutions per site from between sequences are shown. Analyses were conducted using the Jukes-Cantor model.

Supplementary Table 5: Global pairwise alignment analysis for each intron including human and non-human primates DRB5 genes

Supplementary Figure 1: Molecular Phylogenetic analysis of DRB5 intron 2 sequences obtained by the Maximum Likelihood method based on the Jukes-Cantor model.

Legend: Numbers on the branches indicate the bootstrap values for 500 repeats.

spanning exon 2 to exon 6 was obtained for DRB5*01:03, 01:08N and 02:03. Phylogenetic analysis of the generated sequences showed that the DRB5 alleles group together and have distinctive differences with other DRB loci. Novel intron variants of DRB5*01:01:01, 01:02 and 02:02 were identified. The newly characterized DRB5 intron variants of each DRB5 allele were found in subjects harboring distinct associations with alleles of DRB1, B and/or ethnicity. The new information provided by this study provides reference sequences for HLA typing methodologies. Extending sequence coverage may lead to identify the disease susceptibility factors of DRB5 containing haplotypes while the unexpected intron variations may shed light on understanding of the evolution of the DRB region.

Keywords

Next Generation Sequencing; *de novo* assembly; HLA-DRB5; Single Tandem Repeats; Gene conversion

1. Introduction

The major histocompatibility complex (MHC) was initially identified because differences in proteins from different individuals that are encoded in this genetic system play a major role in the rejection of tissues and organs. The class I and II MHC genes encode cell-surface heterodimers that play central roles in antigen presentation, tolerance, and self/non-self recognition [1–3]. The human histocompatibility class II genes encode for three cell-surface isotypes, designated HLA-DR, HLA-DQ, and HLA-DP. Each functional HLA class molecule is a heterodimer formed by an alpha and a beta subunit [4]. Molecular studies of the DR sub-region show one DRA gene, encoding the alpha chain, and multiple DRB genes, encoding the beta chains, on different haplotypes, which display, in addition, copy number variation. The DRA gene in humans is highly conserved while nine different HLA-DRB genes have been described. HLA-DRB1, -B3, -B4 and -B5 encode functional gene products, whereas -B2, -B6, -B7, -B8, and -B9 represent pseudogenes as manifested by various insertions/deletions (indels) and deleterious mutations [5]. Among the expressed HLA-DRB genes, DRB1 is the most polymorphic locus while DRB3, -B4 and -B5 have significantly less alleles as reported to the IMGT/HLA database [6]. The most common DRB5 alleles include DRB5*01:01:01, 01:02, 01:03, 01:08N and 02:02 [7–8]. Within the human population, five major region haplotype configurations have been described associated to serotypes DR1, DR51, DR52, DR8, and DR53, that are characterized by the presence of a unique combinations of DRB genes/pseudogenes [9–10]. For the chimpanzee (*Pan troglodytes*) and some macaque species 9 to more than 30 different DRB haplotypes have been described [11–12].

In the human MHC, the DRB5 locus is unique to the haplotypes bearing the DR51 serotype; DRB5 is adjacent to the DRB6 pseudogene. The DRB5 locus appears likely to derive from ancestral DRB1 alleles, and was generated more than 20 million yr; DRB5 is present in chimpanzees and gorillas [13–14]. The DRB gene organization with duplications and the extensive allelic polymorphism of expressed DR-beta molecules are striking features. These findings suggest that the DRB genes and the corresponding functional

molecules are under distinct selective pressures that resulted from episodic evolution involving both, population expansions and contractions with functional adaptations [14].

The evolutionary history of the HLA-DRB1 locus has been delineated thoroughly by the analysis of genomic full-length alleles (10–15 kb) of human and non-human primates [15]. In contrast, the full evolutionary history of the second expressed DRB loci of different haplotypes such as those bearing DRB5 has not been assessed because of incomplete intron sequence information. Significant information about the evolution and biological functions of DRB loci can be obtained through complete sequencing analysis in the second expressed DRB genes. These should include evaluation of both coding and non-coding regions such as the microsatellite (GT)_x(GA)_x repeats adjacent to exon 2 region [16]. This microsatellite in the MHC- DRB1 genes is interesting since not only in the exon/intron architecture is exactly conserved among all studied vertebrates but also has revealed that the exceptional polymorphism of exon 2 correlates with the variability of this microsatellite locus [17].

HLA matching for transplantation and mapping disease susceptibility and resistance factors, accurate and highly informative HLA allele assignment is desired. The application of whole gene next-generation sequencing (NGS) to the study of highly polymorphic and structurally complex regions of the human genome increases the throughput, accuracy, and resolution of genetic analysis by several orders of magnitude, presenting an opportunity to better understand the biological mechanisms underlying HLA disease associations [18]. For the evaluation of sequencing data, a comprehensive sequence reference database is needed in order to obtain accurate HLA assignments. Genomic references for some HLA allele lineages and loci are missing from the HLA sequences compiled by IMGT [6]. To address these limitations, we embarked in the characterization of genomic sequences of less studied or overlooked loci. In the present study we focused in the characterization of all common alleles of HLA-DRB5.

2. Materials and methods

2.1 Samples and DNA preparation

Samples from the international workshop cell lines (Research Cell Bank, Fred Hutchinson Cancer Research Center, Seattle, Washington) and from selected individuals previously typed by NGS were selected for this study. Genomic DNA was obtained using the QIAamp 96 DNA blood kit (Qiagen, Valencia, CA). Table 1 shows the samples and their HLA alleles. With exception of PGF, a cell line that is consanguineous and homozygous at all HLA loci [19], all the cell lines or subjects included in this study are heterozygous in DRB1 and carried one copy of DRB5.

2.2 HLA Database Construction Strategy

2.2.1 PCR Amplification—DRB alleles were amplified by long-range PCR (Long AMP polymerase, New England Biolabs) of genomic DNA. In order to determine the nucleotide sequences, two fragments overlapping in the highly polymorphic exon 2 region were amplified (Fig. 1). The primers were designed on the basis of examination of DRB sequences available in the IMGT database [6] and Ensembl [20] with the aid of Integrated

DNA Technologies OligoAnalyzer v3.1 software. 0.2 M Trehalose was introduced into PCR reaction to obtain reliable and efficient amplification [21].

2.2.2 Molecular Cloning of PCR products—The two PCR products were gel purified and cloned into the pCR-XL-TOPO vector using the TOPO® XL PCR Cloning Kit (ThermoFisher Scientific). The positively identified clones confirmed via gel agarose electrophoresis were further examined by Sanger sequencing (Elim Biopharmaceuticals, Inc).

2.2.3 Nucleotide Sequencing by NGS—The library construction protocol of the allelic clones for NGS was performed exactly as described by Wang et al. [22]. Sequencing was performed in a MiSeq sequencer using 250 bp paired-end reads run according to the manufacturer's instructions (Illumina, San Diego).

2.2.4 Processing of the NGS data and de-novo Assembly—The `fastq_barcode_splitter.pl` from Hannon/CSHL [23] was used to demultiplex of the raw fastq data generated by NGS sequencer. Read trimming was performed with `bwa -q 20` [24]. The fastq file produced for each sample was converted to a corresponding fasta, which was subsequently used as the input to a developed *de novo* assembly process. The developed algorithm was a blast based assembler [25] and performed the sequence assembly for each cloned amplicon. For the accurate determination of the STR and homopolymer lengths a localized sequence assembly, micro- assembly, of the reads mapping around these locations analyzed directly.

The exact HLA Database construction Strategy with evaluation and comparison with other available algorithms is included in the Supplementary Materials and Methods.

2.3 Phylogenetic Analysis

The full-length generated DRB sequences were deposited into the NIH [6] and IMGT HLA genetic database [26] (Table 1). The non-human primates DRB5 sequences of *Pan troglodytes* part-DRB5*01:02 and *Macaca mulatta* mamu-DRB5*03:01 were retrieved from the IPD-MHC-NHP database [27,28].

Pairwise comparisons of the DRB sequences were performed with the EMBOSS needle program using the Needleman-Wunsch algorithm [29]. Multiple sequence alignments executed by the Clustal Omega program [29]. The software MEGA 6.0 [30] was employed to calculate nucleotide diversity and to construct phylogenetic trees. The phylogenetic trees were reconstructed using the maximum likelihood method with Jukes-Cantor model [31]. The phylograms were a consensus of 500 bootstraps replicates.

3. Results

3.1

3.1.1 Description of HLA-DRB5 alleles—This report provides additional information regarding sequence variation at both, exons and introns of the most common DRB5 alleles. Out of the 87 samples genotyped, 18 samples harbored the DRB5 alleles (Table 1); the

resulting consensus sequences were analyzed and compared between them and with other alleles of the DRB gene families as well as DRB5 alleles of non-human primates.

In the present study we identified intron variants resulting from SNP or STR variation. We developed a local naming convention where the e suffix indicated sequence extension, the suffix v specified intron variants while the suffix _STR indicated differences in the STR length (Table 1).

The sequence analyses showed that seven individuals carried DRB5*01:01:01, six DRB5*02:02, two DRB5*01:02, and one of each, DRB5*01:03, DRB5*01:08N and DRB5*02:03. The sequence of DRB5*01:01:01 was confirmed in two samples (JS and PGF, Table 1). The CDS present in IMGT [6] for DRB5*01:02 and DRB5*02:02 was confirmed and extended from the 5'UTR up to the end of intron 5 including all intervening introns and exons with the total length of 12,681 and 12,638 bp respectively (Supplementary Tables 1 and 2). For DRB5*02:03, DRB5*01:03 and DRB5*01:08N alleles the cloning data extended the previously known sequence from the beginning of exon 2 to the end of intron 5 with lengths of 4,658, 4,693 and 4,672 bp respectively (Supplementary Table 2).

3.1.2 5' UTR sequences (PCR Fragment 1)—A segment of 173 bp upstream of exon 1 was identical for DRB5*01:01:01, 01:02 and 02:02. Examination of sequences from three alleles in which fragment 1 (5'UTR to exon 2) was analyzed showed that exon 1 of DRB5*01:02 is identical to that of DRB5*01:01:01; these alleles differ from DRB5*02:02 by two nucleotide substitutions resulting in a one amino acid change in codon -16 (K to V substitution) (Table 2). There is higher sequence divergence between DRB5*01 and DRB5*02 alleles in intron 1 (Fig. 2A,B, Supplementary Tables 1,2). The nucleotide differences between DRB5*01:01:01, DRB5*01:02 and DRB5*02:02 are shown in Figure 2A,B and Supplementary Tables 1,2. The location of two intron 1 homopolymers in reference to the first nucleotide of DRB5*01:01:01 in intron 1 is shown in Figures 2A,B. length variation (Table 2) and the sequences of these (T)_x (Fig. 2A) and (A)_x (Fig. 2B) homopolymers are distinctive among the described alleles. An estimation example of the length of the (A)_x and (T)_x intron 1 homopolymers is shown in Figures 3A and 3B.

3.1.3 Extended sequence coverage (PCR Fragment 2)—Fragment 2 (exon 2 to intron 5) was cloned and analyzed for all 18 samples bearing DRB5. In these samples, the length of this fragment ranged from 4,657 bp (DRB5*01:01:01v1) to 4,693 bp (DRB5*01:02_STR1) (Fig. 1). Interestingly, almost all the length variation between these alleles resulted from differences in the number of two dinucleotide STRs, (GT)_x and (GA)_x of intron 2 (50bp after exon 2) (Table 2).

As previously described [32–33], additional length variation results from the deletion of 2 and 19 nucleotides in exon 2 and exon 3, respectively of DRB5*01:10N and DRB5*01:08N (Supplementary Tables 1,2).

Three additional intron variants were identified among 7 subjects carrying DRB5*01:01:01; these are defined by differences in STR length in intron 2 (DRB5*01:01:01_STR1, DRB5*01:01:01v1, DRB5*01:01:01v1_STR1) and an intron 2 SNP variation (9990A/G)

(DRB5*01:01:01v1, DRB5*01:01:01v1_STR1) (Table 2). Figure 1 shows the location of two contiguous STRs located at the beginning of intron 2 in reference to DRB5*01:01:01. The sequences of both STRs of all DRB5 alleles described in the present study are shown in Figure 4; their length variation is shown in Table 2. Based on the analyses of SNPs in introns 2 and 3 (Table 2) three DRB5 lineages have been identified, namely DRB5*01:01:01, DRB5*01:02 (including also 01:03 and 01:08N) and DRB5*02. Table 1 shows the cell lines or subjects that carry the four variants of DRB5*01:01:01 also carry DRB1*15:01:01; these DRB5*01:01:01 intron variants are found in distinct haplotypes defined either by variations in the non-coding regions of the DRB1*15:01:01, in HLA-B or by distinguished by ethnic differences.

Two intron 2 STR variants of DRB5*01:02 were described in the present study (DRB5*01:02e1 and DRB5*01:02_STR1). These alleles were found in cells carrying DRB1*15:02:02 for DRB5*01:02e1 and DRB1*15:02:01 for DRB5*01:02e1_STR1 and they differ also in DQA1, DQB1 or at both DQ loci (Table 1).

The DRB5*01:02_STR1 and DRB5*01:03 alleles differ only by one nucleotide substitution in exon 2 and were identical in the gene segment spanning exon 2 to 5 (Table 2). The estimation of the STR lengths in DRB5*01:03 found in cell 119990 are shown in Figures 3C (GA)_x and 3D (GT)_x. The alleles DRB5*01:02_STR1 and DRB5*01:03 had high similarity with DRB5*01:08N in the assembled genomic region.

In this report two STR variants of DRB5*02:02 were identified (DRB5*02:02e1 and DRB5*02:02e1_STR1) (Fig. 4 and Table 2) and the genomic sequence of DRB5*02:03e1 was extended. The variants of DRB5*02:02 were found in haplotypes carrying different subtypes of DRB1*16. DRB5*02:02e1 and DRB5*02:03e1 were found in cells carrying DRB1*16:02:01 and DRB1*16:04 while the allele DRB5*02:02e1_STR1 was found in the cell line CHA, AJ that carries DRB1*16:01:01 (Table 1).

3.2

3.2.1 Comparison with other DRB genes—A global alignment (EMBOSS Needle) [29] and a phylogenetic tree was employed to determine fragment 2 homologies and phylogenetic relationships between DRB5*01:01:01 with other DRB5 alleles and DRB genes (DRB1/3/4/6/7). DRB5 alleles had very high similarity scores with distinctive differences with other DRB loci (Fig. 5A). The intra pairwise alignment displayed lower similarity scores between DRB5*01:01:01 and other DRB genes; the DRB5 gene had higher similarity with DRB4 followed by DRB3 and DRB1*15:01. The alignments between DRB5*01:01:01 and DRB pseudogenes (DRB6 and DRB7) showed the highest genetic dissimilarity (Supplementary Table 3). The pairwise comparisons between DRB5 alleles indicated that DRB5*01:02, DRB5*01:03 and DRB5*01:08N are highly homologous and as a group present the lowest distances with DRB5*01:01:01, DRB5*02:02 and DRB5*02:03 alleles (Supplementary Table 4). The phylogenetic analysis identified three clusters of alleles with the DRB5*01:02 group being in the middle and closer to the DRB5*01:01 group than to the DRB5*02 group (Fig. 5B).

3.2.2 Comparison with Non-Human primates DRB5 alleles—Phylogenetic analysis of exon 2 to intron 5 sequences and intron 2 only for human and non-human primate DRB genes and DRB5 alleles were performed. The results indicated that the hominid DRB5 alleles (*Homo sapiens* and *Pan troglodytes*) and the old world monkey mamu-DRB5*03:01 allele of *Macaca mulatta* form a separate clade in comparison to the other DRB genes (Fig. 5A, Supplementary Figure 1). In addition, a global alignment was conducted, using the Needleman-Wunsch algorithm [29], among HLA-DRB5 alleles, the common chimpanzee patr-DRB5*01:02 and Rhesus macaque mamu-DRB5*03:01. The results showed high homology of 92.6% between HLA-DRB5*01:01:01 and patr-DRB5*01:02 but lower with the other human DRB5 alleles. The mamu-DRB5*03:01 and HLADR5*01:01:01 sequences had 91.5% similarity which was higher than the alignment among other DRB genes (Supplementary Table 3). Further global pairwise alignment analysis was performed for separate introns between human and non-human primates of DRB5 alleles and can be found in Supplementary Table 5. Interestingly, SNPs that are informative and separate the human DRB5 families can be found also in non-human species (Table 2).

3.3 STR/Homopolymer Analysis of DRB5 alleles

In this study, the variations of simple repeat stretches with the basic structures poly(A), poly(T) and $(GT)_x(GA)_x$ in the different intronic areas of the HLA-DRB5 alleles were extensively investigated (Table 2, Fig. 3A–D). The NGS data showed an inverse correlation between the accuracy in the determination of STR length and the length of the repeats which is concordant with previous studies [34–37]. The short repeats $T_{(6,9)}$ in intron 1 of DRB5*01:01:01 were determined with high confidence because more than 90% of the reads in this area for each clone indicated the same poly(T) length. On the contrary, for longer homopolymers $(T)_{14}$, $(A)_{20}$ in intron 1 of DRB5*01:01:01 and STRs $GA_{(9)}$, $GT_{(21)}$ in intron 2 of DRB5*01:03e1, the reads showed a broader Gaussian like distribution; the highest frequency of the most representative reads for this area per clone were ranging from 22% to 60% (Fig. 3A–D). This broader distribution results in more difficult determination of the length of these STRs; more than 3 clones were required to be sequenced separately (instead of pooling them) for achieving accurate analysis.

The length variation and the basic structure of intron 2 STRs for all the DRB alleles for human and primates that has been examined in this report are shown in Table 2.

The $(GT)_x$ variations associate with specific haplotypes and ethnic background as shown in Table 1 while the intron $(GA)_x$ variants correlate with the three DRB5 lineages namely DRB5*01:01:01, DRB5*01:02 (including also 01:03 and 01:08N) and DRB5*02 (Table 2).

Comparing the human DRB5 intron 2 microsatellite structures with other DRB genes shows only similarity with DRB1*15/16 and DRB1*09 alleles (Table 2). The intron data indicates that DRB1*16:01:01 was generated through a gene conversion event involving DRB1*15:01:01:01 (recipient allele) and DRB5*01:01:01v1 (donor allele) alleles that are found on the same haplotype (Table 1); the exchange segment is shown in Figure 6.

Only three DRB5 alleles were examined in the first segment spanning intron 1; The poly(A) and poly(T) exhibited high variation in the number of repeats in both human and other primates DRB5 alleles (Table 2). More sequencing data is required to further characterize and determine the diversity and possible evolution of this region.

4. Discussion

In the present study, robust and improved methodological approaches were applied to characterize the gene structure and diversity of the second DRB locus expressed in haplotypes bearing DRB1*15 and DRB1*16 alleles. The novel strategy for full coverage and cost-effective allele sequencing took advantage of a modified long range PCR, highly efficient molecular cloning, clonal pooling, Illumina's NGS platform and a novel assembly algorithm specific to HLA genes [Supplementary Materials and Methods] [21, 38–39]. Long read sequencing technologies were not used in this study since may present major drawbacks such as PCR-chimera formation and biased reference alignment, which need to be considered when attempting to phase variants [38].

Because of the noise inherent to NGS protocols, cloning errors and sequence complexity in the HLA alleles, *de-novo* assemblers such as Velvet (v1.2.1) and SSAKE (v3.8.2) [40–41] could not generate the full length and error-free sequence of each selected allele. These limitations led us to design an in-house assembly algorithm [Supplementary Materials and Methods]. The addition of a local *de novo* assembly and the evaluation of isolated clones was able to resolve the low complexity and high error areas such as homopolymers and STR regions (Fig. 3A–D) [42–44]. This approach provided significant and valuable information that could have not been obtained otherwise. The novel variants reported here were not deemed to receive an official name by WHO Nomenclature Committee for Factors of the HLA System. Nevertheless, the information and publication of these sequences provides significant value for the application of NGS based methods to HLA typing and further delineate evolutionary relations of HLA alleles and haplotypes [45,46].

The SNP analysis of the most common DRB alleles shows that all the variation in DRB5 locus resides in the 5' side of the gene including exon 1 up to intron 3, whereas the region comprises exon 4 to intron 5 is completely conserved. Virtually all SNPs observed in introns 2 and 3 and the intron 2 (GA)_x correlate exactly with three DRB5 lineages (Fig 5B) also defined previously by exon analyses only (Table 2). The intron 2 (GT)_x appears to define recently generated variants in specific haplotypes and ethnic background as shown in Table 1. At the DRB5 locus, the intron 2 (GT)_x appears to evolve more rapidly in comparison to the intron 2 (GA)_x; the evolution rate of these STR is not similar at other DRB genes. This observation may be explained by: 1) different mutation rates among the DRB allele families and loci; and/or 2) different selective pressures in each DRB families. In the present study all alleles of the same DRB5 subfamily contain the same length (GA)_x in intron 2 while they may differ in the length of the (GT)_x. Both STRs are long; therefore, the mutation rate observed for these STRs indicates that the variation does not necessarily results from the repeat length. Interruptions in the perfect (GA)_x repeats may decrease the mutation rate; in addition differentially acting selective pressures such convergent evolution may decrease the exon variability to that of the intron microsatellites [47,48].

The observed intron variability of the DRB5 STRs could result in possible biological and functional effects that need to be further evaluated. Permanent coevolution with exons suggests a possible biological role of these composite intron microsatellites [49–53]. Despite the repeat length variations of the STRs, the basic structure of (GT)_x(GA)_x is highly conserved between different families of DRB5 gene (Table 2). Interestingly, similar structures can be seen in DRB1*09/15/16 alleles (Table 2) which led us to confirm a previous hypothesis regarding a reciprocal intergenic exchange between DRB loci [54]. This report shows that the DRB1*16:01:01 allele may have arisen by reciprocal intergenic exchange between DRB1*15:01:01:01 (recipient allele) and DRB5*01:01:01v1 (donor allele) in the DR51 haplotype and the postulated recombination sites are shown in Figure 6.

The phylogenetic analysis showed that the hominid DRB5 alleles (*Homo sapiens* and *Pan troglodytes*) and the old world monkey (OWM) mamu-DRB5*03:01 allele of *Macaca mulatta* form a separate clade in comparison to the other DRB genes (Fig. 5A). There were two major diversification events in the evolution of the HLA-DRB genes approximately 50 million years (my) ago. A DRB1*04 and an ancestor of the DRB1*03 cluster (DRB1*03, DRB1*15, and DRB3) diverged from each other approximately 50 million years (my) ago, and DRB5, DRB7, DRB8, and an ancestor of the DRB2 cluster (DRB2, DRB4, and DRB6) emerged by gene duplication [55]. These data confirm that the DRB5 locus is common to the Catarrhini's DRB region and the DRB5 gene originated before the OWM–HOM deviation (~25 My ago) as has been reported in other studies [27, 56].

The second expressed DRB loci (DRB3, DRB4, and DRB5) exhibit only limited allelic polymorphism in humans [57]. HLA typing of DRB5 alleles examining exon sequences shows only a few common alleles [6]. The structural protein sequence conservation in different ethnic groups is remarkable in spite of the rich haplotype variation identified when examining DRB1-DQA1-DQB1 haplotype blocks containing DRB5 (Table 1) [58,59]. The present study identifies additional non-coding variation that is haplotype specific and appears to indicate that there may be constraints for further diversification at the protein level compared to variations in the non-coding regions. It is proposed that DRB5 alleles may exert specific functions and may complement with molecules present in DRB1*15/16 and in the corresponding DQA1-DQB1 heterodimers [60–62]. Further work is needed to examine what immune responses are principally determined or restricted by the DRB5 alleles.

In all world populations that include haplotypes bearing in cis the genes encoding for DQA1*01:01 and DQB1*05:01, it has been observed that the DRB5 locus is absent in spite of the presence of the DRB6 pseudogene. These haplotypes include predominantly DRB1*01 alleles [59,63]. In addition to these haplotypes, Asian populations present frequently non-DRB1*01 haplotypes that include the DQ genes for the same DQ heterodimer (DQA1*01:01:01:01-DQB1*05:01:24); these haplotypes include the allele DRB1*15:02:01:03 and carry either DRB5*01:02 or DRB5*01:08N [32,64]. We speculate that the non-expressed DRB5*01:08N allele may have arisen recently in haplotypes bearing the latter DRB1-DQ alleles that also include DRB5*01:02. The putative mutational event with a deletion of 19 nucleotides in exon 3 of DRB5*01:02e1_STR1 may have resulted in the generation of DRB5*01:08N. This deletion produces a truncated non-membrane bound protein [32]. Given the high frequency of the allele DRB5*01:08N, it can be speculated that

the haplotypes bearing this allele may have been positively selected in Asian populations. A plausible explanation is that because the DRB expressed alleles may determine negative selection of some T cells in the thymus, the non-expression of a specific allele may allow for the generation of some T-cell clones that could be effective in responding pathogen antigens via presentation by other HLA class II molecules. The various haplotypes (bearing DRB1*01:01, DRB1*01:02, DRB1*15:02) that include genes in cis encoding DQA1*01:01 and DQB1*05:01 include alleles with identical protein sequences that differ by exon silent substitutions or intron variations; these observations suggest more distant origins with positive pressure for structural DQ conservation that may be related to absence of DRB5 expression.

The molecules encoded by the second expressed DRB genes DRB3, DRB4 and DRB5 appear to have lower expression than DRB1 [65–70]. It should be noted that DRB4*01:03:01:02N is another common null allele that is found frequently in subjects with European and Asian ancestry [59]. In Africans, approximately 8 percent of the haplotypes bearing DRB1*15:03 lack DRB5 gene [59]. Therefore, DRB5 alleles, as other DRB low expression alleles may play a role in both providing specific responses to pathogens as well as determining the size of T-cell repertoire of an individual.

The present study provides a novel strategy for generating extended allele sequences of high quality that are reliable sources for HLA genomic references. Additionally, the characterization of the most common DRB5 alleles may improve the NGS based HLA typing by providing valuable phasing information that can subsequently lead to greater precision in matching donor organs to transplant recipients.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations:

HLA	Human Leukocyte Antigens
MHC	Major Histocompatibility Complex
NGS	Next-Generation Sequencing
PCR	Polymerase Chain Reaction
STR	Single tandem repeats

UTR	Untranslated Region
indel	insertion or deletion

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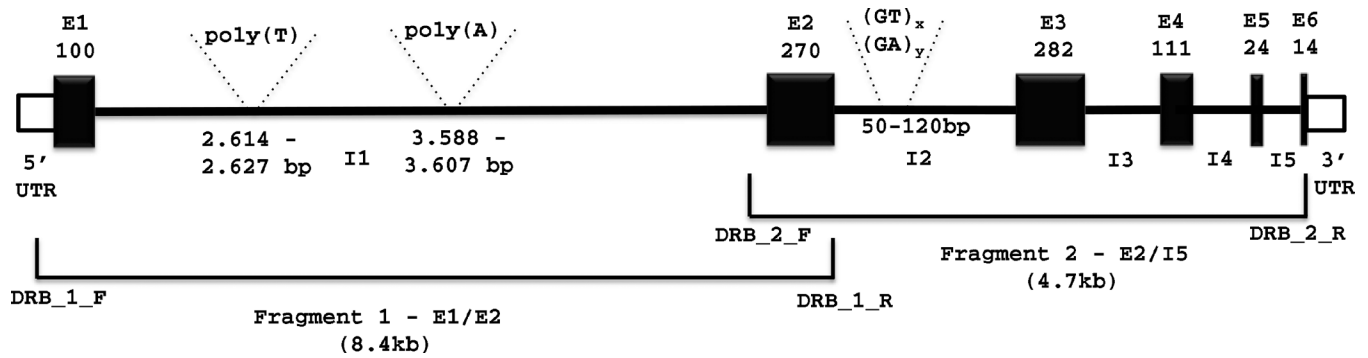


Figure 1:

Schematic illustration of amplified fragments of DRB5. The DRB5*01:01:01 is used as a reference for sequence annotation. Exons are represented by black boxes. E1–E6 denotes exons 1–6. I1–I5 denotes introns 1–5, 5'3'UTR, homopolymer (Poly Ts/As) and STRs locations. The numbers represent the exon length in base pairs. The two amplified fragments are showed at the bottom; they overlap in Exon 2 and flanking sequences. Primers DRB_1_F and DRB_1_R were used to amplify fragment 1 and primers DRB_2_F and DRB_2_R were utilized to amplify fragment 2.

A

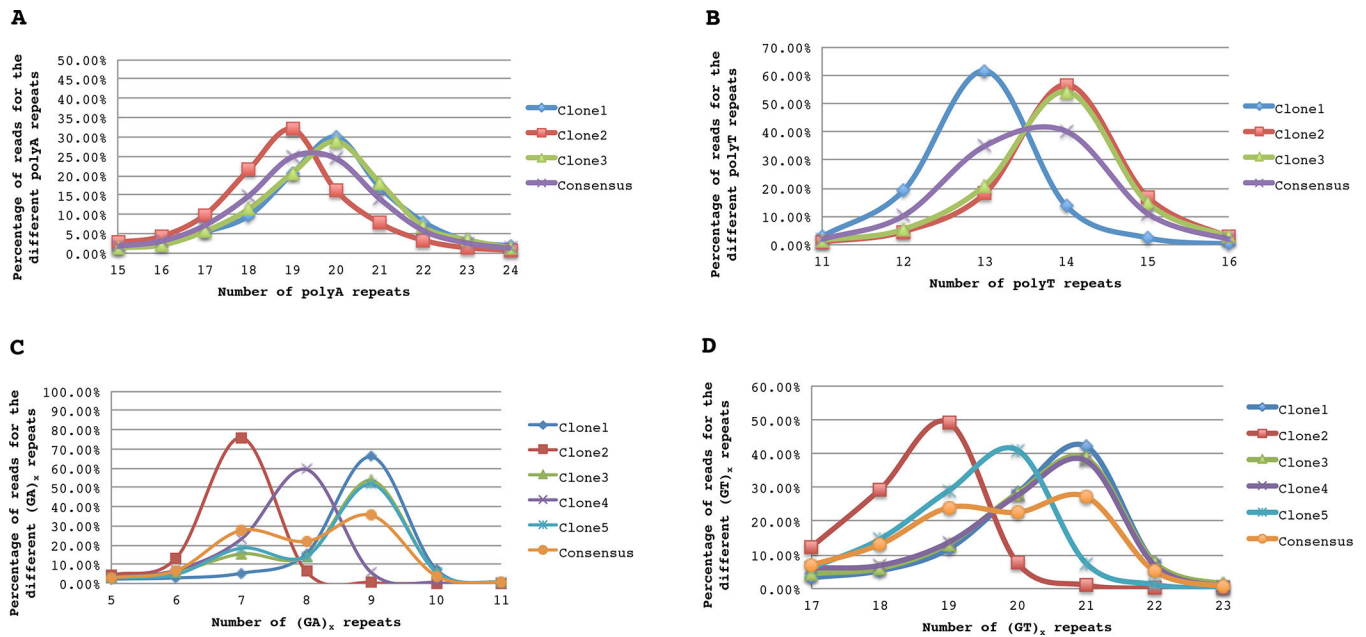
DRB5*02:02e1	ATTTTTTAATTTTTTTTTTTTTT...GAGATATAGTCTTGCTCTGTCACCAGGCTGGAGTG	2663
DRB5*01:01:01	ATTTTTTAATTTTTTTTTTTTTTT...GAGATATAGTCTTGCTCTGTCACCAGGCTGGAGTG	2662
DRB5*01:02e1	ATTTTTTAATTTTTTTTTTTTTTTTTGAGATATAGTCTTGCTCTGTCACCAGGCTGGAGTG	2664

B

DRB5*02:02e1	GGGTGACAGAGCAAGACTCCGTCTCAAAAAAAAAAAAAAAAAA...GAGACTCATGGTGAG	3619
DRB5*01:01:01	GGGTGACAGAGCAAGACTCCGTCTCAAAAAAAAAAAAAAAAAAAGAGACTCATGGTGAG	3622
DRB5*01:02e1	GGGTGACAGAGCAAGACTCCGTCTCAAAAAAAAAAAAAAAAAA.GAGACTCATGGTGAG	3623

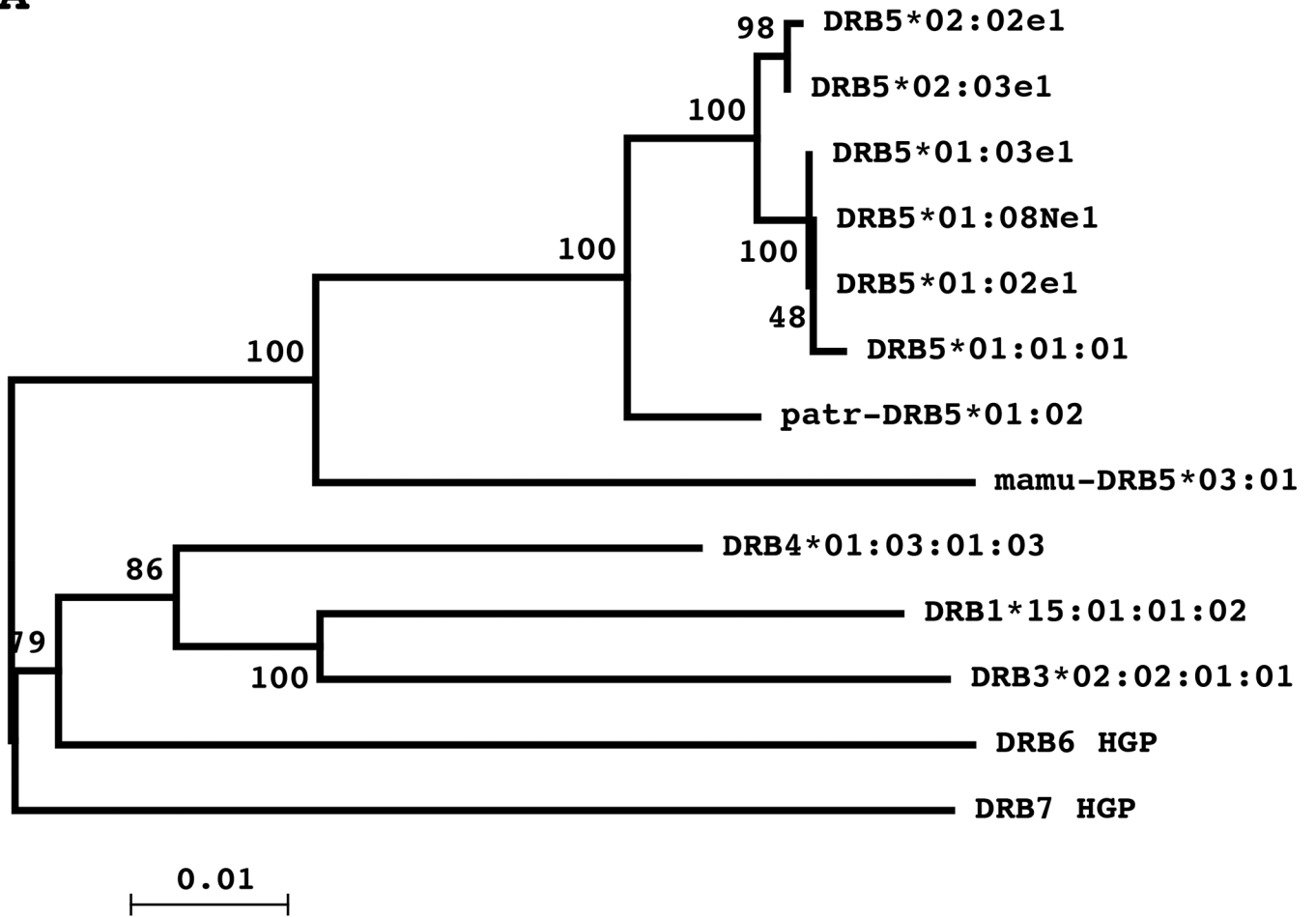
Figure 2.

A: CLUSTALW partial alignment of three *de novo* assembled DRB5 alleles within intron 1 shows the length variation of the poly(T) repeats. Numbering on the right indicates the relative nucleotide position at this intron for the corresponding allele. **B:** CLUSTALW partial alignment of three *de novo* assembled DRB5 alleles within intron 1 exhibits the length variation of the poly(A) repeats. Numbering on the right indicates the relative nucleotide position at this intron for the corresponding allele.

**Figure 3.**

A: Estimation of the length of intron 1 poly(A) in DRB5*01:01:01 from three clones isolated from cell line JS. The most representative STR length was selected for each clone according to percentages of reads; the length found in more clones was assigned as the final length for the tandem repeat. **B:** Estimation of the length of intron 1 poly(T) in DRB5*01:01:01 from three clones isolated from the cell line JS. The most representative STR length was selected for each clone according to percentages of reads; the length found in more clones was assigned as the final length for the tandem repeat. **C:** Estimation of the length of intron 2 (GA)_x STR in DRB5*01:03 found in cell 119990 from five clones of cell 119990. The most representative STR length was selected for each clone according to percentages of reads; the length found in more clones was assigned as the final length for the tandem repeat. **D:** Estimation of the length of intron 2 (GT)_x STR in DRB5*01:03 found in cell 119990 from five clones of cell 119990. The most representative STR length was selected for each clone according to percentages of reads; the length found in more clones was assigned as the final length for the tandem repeat.

A



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B

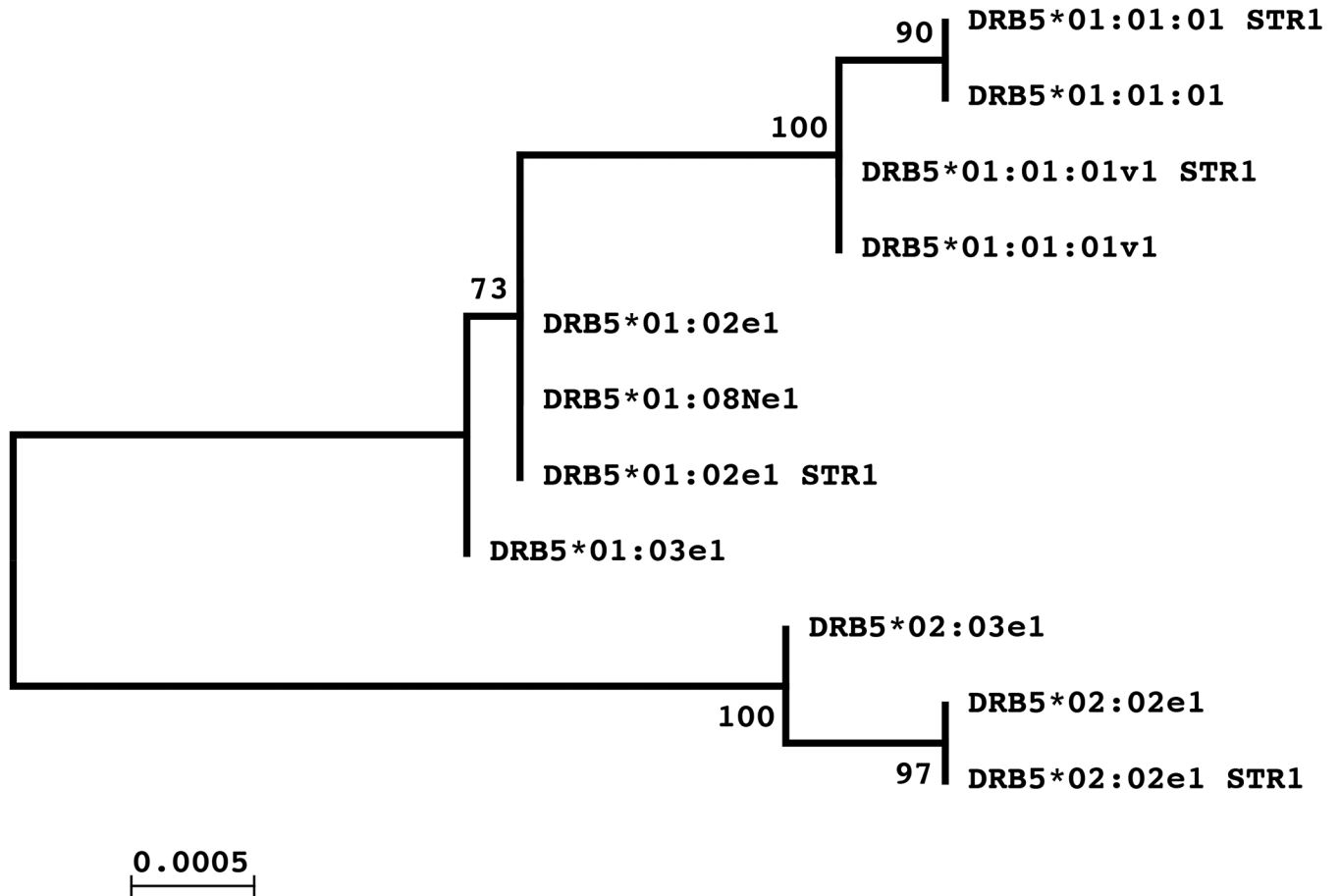


Figure 5.

A: Phylogenetic tree of DRB sequences spanning from exon 2 to exon 5 obtained by the Maximum Likelihood method based on the Jukes-Cantor model. Numbers on the branches indicate the bootstrap values for 500 repeats. **B:** Phylogenetic tree of DRB5 sequences spanning from exon 2 to exon 5 obtained by the Maximum Likelihood method based on the Jukes-Cantor model. Numbers on the branches indicate the bootstrap values for 500 repeats.

DRB1*16:01:01	CACGTTTCCTGTGGCAGCCTAAGAGGGAGTGTCAATTTCTTCAATGGGACGGAGCGGGTGC	60
DRB1*15:01:01:01	CACGTTTCCTGTGGCAGCCTAAGAGGGAGTGTCAATTTCTTCAATGGGACGGAGCGGGTGC	60
DRB5*01:01:01v1	CACGTTTCCTGTGGCAGCAGGATAAGTATGAGTGTCAATTTCTTCAACGGGACGGAGCGGGTGC	60
DRB1*16:02:01:01	CACGTTTCCTGTGGCAGCCTAAGAGGGAGTGTCAATTTCTTCAATGGGACGGAGCGGGTGC	60

	recombination site	
DRB1*16:01:01	GGTTCCTGGACAGATACTTCTATAAACCAGGAGGAGTCCGTGCGCTTCGACAGCGACGTGG	120
DRB1*15:01:01:01	GGTTCCTGGACAGATACTTCTATAAACCAGGAGGAGTCCGTGCGCTTCGACAGCGACGTGG	120
DRB5*01:01:01v1	GGTTCCTGGACAGACATCTATAACCAAGAGGAGGACTTGCCTTCGACAGCGACGTGG	120
DRB1*16:02:01:01	GGTTCCTGGACAGATACTTCTATAAACCAGGAGGAGTCCGTGCGCTTCGACAGCGACGTGG	120

DRB1*16:01:01	GGGAGTACCGGGCGGTGACGGAGCTGGGGCGCCTGACGCTGAGTACTGGAACAGCCAGA	180
DRB1*15:01:01:01	GGGAGTACCGGGCGGTGACGGAGCTGGGGCGCCTGACGCTGAGTACTGGAACAGCCAGA	180
DRB5*01:01:01v1	GGGAGTACCGGGCGGTGACGGAGCTGGGGCGCCTGACGCTGAGTACTGGAACAGCCAGA	180
DRB1*16:02:01:01	GGGAGTACCGGGCGGTGACGGAGCTGGGGCGCCTGACGCTGAGTACTGGAACAGCCAGA	180

DRB1*16:01:01	AGGACTTCTGGAAGACAGGCGCGCCGCGGTGGACACCTACTGCAGACACAACACTACGGGG	240
DRB1*15:01:01:01	AGGACTTCTGGAAGACAGGCGCGCCGCGGTGGACACCTACTGCAGACACAACACTACGGGG	240
DRB5*01:01:01v1	AGGACTTCTGGAAGACAGGCGCGCCGCGGTGGACACCTACTGCAGACACAACACTACGGGG	240
DRB1*16:02:01:01	AGGACTTCTGGAAGACAGGCGCGCCGCGGTGGACACCTACTGCAGACACAACACTACGGGG	240

	end of exon2	
DRB1*16:01:01	TTGGTGAGAGCTTACAGTGCAGCGCGAG GTGAGCATGGTGGGGGGCGGGGCTGGGT	299
DRB1*15:01:01:01	TTGGTGAGAGCTTACAGTGCAGCGCGAG GTGAGCATGGCCTGGGGGGCGGGGCTGAGT	299
DRB5*01:01:01v1	TTGGTGAGAGCTTACAGTGCAGCGCGAG GTGAGCATGGTGGGGGGCGGGGCTGGGT	299
DRB1*16:02:01:01	TTGGTGAGAGCTTACAGTGCAGCGCGAG GTGAGCATGGTGGGGGGCGGGGCTGGGT	299

	recombination site	
DRB1*16:01:01	CCTTGAGCTGGGAATCTGAGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT...	356
DRB1*15:01:01:01	CCCCGTGAGCTGGGAATCTGAGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT...	359
DRB5*01:01:01v1	CCTTGAGCTGGGAATCTGAGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT...	356
DRB1*16:02:01:01	CCTTGAGCTGGGAATCTGAGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT...	356

DRB1*16:01:01GAGAGAGAGAGAGA.....GGAAGAGAGAGAGAGAGCGCCATCTGTGAGC	403
DRB1*15:01:01:01	TGAGAGAGAGACAGAGAGAGACAGAGAGA.....GGAAGAGAGAGAGAGAGCGCCATCTGTGAGC	419
DRB5*01:01:01v1GAGAGAGAGAGAGAGAGGAAGAGA.....GAGACAGAGAGGGAGCGCCATCTGTGAGC	411
DRB1*16:02:01:01GAGAGAGAGAGAGAGA.....GGAAGAGAGAGAGAGAGCGCCATCTGTGAGC	403

DRB1*16:01:01	ATTTAGAATCCTCTCAGTCCGGAGCAAGCAGTTCTGAGAGCACAGGTGTGTGTGTAAAGT	463
DRB1*15:01:01:01	ATTTAGAATCCTCTCAGTCCGGAGCAAGCAGTTCTGAGAGCACAGGTGTGTGTGTAAAGT	479
DRB5*01:01:01v1	ATTTAGAATCCTCTCAATCTTGAGCAAGGACTTCTGAGGACACAGGTGTGTGTGTAGAGT	471
DRB1*16:02:01:01	ATTTAGAATCCTCTCAGTCCGGAGCAAGCAGTTCTGAGAGCACAGGTGTGTGTGTAAAGT	463

Figure 6:

Nucleotide sequences of a putative segmental exchange involved in the generation of DRB1*16:01:01. Boxes show the segments shared by DRB1*15:01 and DRB5*01:01 defining the boundaries of the segmental exchange.

Numbering on the right indicates the relative nucleotide position from the beginning of exon 2 for the corresponding allele.

Table 1:

HLA genotypes obtained by Next Generation Sequencing and reference sequences of 18 subjects and cell lines carrying DRB5.

International Workshop Cell line #	Local Sample ID	Ethnic Origin	Accession #	IMGT Submission #	CLASS II ALLELES										CLASS I ALLELES					
					DRB5	DRBI	DQA1	DQBI	DPA1	DPBI	HLA-A	HLA-B	HLA-C	DRB5	DRBI	DQA1	DQBI	DPA1	DPBI	HLA-A
IHW0 9228	JS	European - North America	AL713966		DRB5*01:01:01 DRB3*02:02:01:01	DRB1*15:01:01 DRB1*03:01:01:01	DQA1*01:02:01:01 DQA1*05:01:01:01	DQB1*06:02:01	DPA1*01:03:01:05 DPA1*01:03:01:01	DPB1*04:02:01:02 DPB1*02:02	A*03:01:01:01 A*30:03	B*07:02:01 B*18:01:01:01	C*07:02:01:03 C*05:01:01:01							
IHW0 9318	PGF	European - England, Europe	AL713966		DRB5*01:01:01	DRB1*15:01:01	DQA1*01:02:01	DQB1*06:02:01	DPA1*01:03:01:02	DPB1*04:01:01:01	A*03:01:01:01	B*07:02:01	C*07:02:01:03							
IHW0 9123	HAY, BD	Australian Aboriginal	KU593577	HWS10025809	DRB5*01:01:01 STR1 ^a DRB3*02:02:01:02	DRB1*15:01:01 DRB1*14:07:01	DQA1*01:02:01:01 DQA1*01:04:01	DQB1*06:02:01 DQB1*05:03:01:01	DPA1*01:03:01:02 DPA1*02:02:02	DPB1*04:01:01:01 DPB1*05:01:01	A*24:02:01:01	B*07:02:01 B*40:02:01	C*07:02:01:03 C*15:02:01:01							
IHW0 9394	BPT		KU593577	HWS10025809	DRB5*01:01:01 STR1 ^a	DRB1*15:01:01 DRB1*13:01:01	DQA1*01:02:01:01 DQA1*01:03:01:02	DQB1*06:02:01 DQB1*06:03:01	DPA1*01:03:01:04	DPB1*04:01:01:01	A*02:01:01:01	B*07:03 B*15:01:01:01	C*03:04:01:01 C*03:03:01							
	STA1001	Asian	KU593580	HWS10025817	DRB5*01:01:01 STR1 ^b	DRB1*15:01:01 DRB1*08:03:02	DQA1*01:02:01:03 DQA1*01:03:01	DQB1*06:02:01 DQB1*06:01:01	DPA1*01:03:01:01 DPA1*02:02:02	DPB1*05:01:01	A*02:01:01:01 A*32:01:01	B*54:01:01 B*52:01:01:02	C*01:02:01 C*12:02:02							
	STA1029	Asian	KU593580	HWS10025817	DRB5*01:01:01 STR1 ^b DRB4*01:03:01:02N	DRB1*15:01:01 DRB1*07:01:01	DQA1*01:02:01:03 DQA1*02:01	DQB1*06:02:01 DQB1*03:03:02:01	DPA1*02:01:01	DPB1*13:01:01	A*02:06:01:01 A*03:01:01:01	B*35:01:01:02 B*35:03:01	C*03:03:01 C*12:03:01:01							
IHW0 9368		Asian - Unknown	KU593576	HWS10025819	DRB5*01:01:01 STR1 ^c	DRB1*15:01:01 DRB1*09:01:02	DQA1*01:02:01:01 DQA1*03:02	DQB1*06:02:01 DQB1*03:03:02:02	DPA1*01:03:01:01 DPA1*02:02:02	DPB1*02:01:02 DPB1*05:01:01	A*24:02:01:01 A*26:04	B*38:02:01 B*39:01:03	C*07:02:01:01							
IHW0 9327	THA	European - India, Asia	KU593571	HWS10025805	DRB5*01:02:01 STR1 ^d DRB3*02:02:01	DRB1*15:02:02 DRB1*13:08	DQA1*01:03:01:01 DQA1*01:04:01	DQB1*06:01:01 DQB1*05:03:01	DPA1*01:03:01:04 DPA1*01:03:01:02	DPB1*04:01:01	A*31:01:02:01 A*03:01:01:01	B*35:03:01 B*15:18:01	C*04:01:01:01 C*07:04:01							
	STA1005	Asian	KU593573	HWS10025807	DRB5*01:02:01 STR1 ^e	DRB1*15:02:01 DRB1*10:01:01	DQA1*01:01:01 DQA1*01:05:01	DQB1*05:01:24 DQB1*05:01:01:02	DPA1*01:03:01:01 DPA1*02:02:02	DPB1*02:01:02 DPB1*13:01:01	A*11:01:01:01 A*29:01:01:01	B*38:02:01 B*07:05:01	C*15:05:02 C*07:02:01:01							
	STA1007	Asian	KU593574	HWS10025813	DRB5*01:03:01 STR1 ^f DRB3*03:01:03	DRB1*15:02:01 DRB1*12:02:01	DQA1*01:01:01 DQA1*06:01:01	DQB1*05:02:01 DQB1*03:01:01:01	DPA1*02:02:02 DPA1*01:03:01:03	DPB1*05:01:01 DPB1*21:01	A*02:03:01 A*24:02:01:01	B*40:02:01 B*15:02:01	C*03:04:01:01 C*08:01:01							
	STA1010	Asian	KU593575	HWS10025815	DRB5*01:08:01 STR1 ^g	DRB1*15:02:01 DRB1*10:01:01	DQA1*01:01:01 DQA1*01:05:01	DQB1*05:01:24 DQB1*05:01:01:02	DPA1*01:03:01:03 DPA1*02:02:02	DPB1*21:01 DPB1*05:01:01	A*11:01:01:01 A*29:01:01:01	B*38:02:01 B*07:05:01	C*07:02:01:01 C*15:05:02							
IHW0 9277	HS67	Asian - Japan	KU593572, KX687265	HWS10025811	DRB5*02:02:01 STR1 ^h DRB3*01:01:02	DRB1*16:02:01 STR1 ^k DRB1*08:03:02	DQA1*01:02:02 DQA1*01:03:01	DQB1*05:02:01 DQB1*06:01:01	DPA1*02:02:02	DPB1*05:01:01	A*11:01:01:01 A*24:02:01:01	B*67:01:01 B*48:01:01	C*07:02:01:01 C*08:03:01							
IHW0 9258	DAN72.3	American Indian - North America	KU593572, KX687282	HWS10025811	DRB5*02:02:01 STR1 ^h DRB3*01:01:02	DRB1*16:02:01 STR1 ^l DRB1*14:02:01	DQA1*05:05:01:03 DQA1*05:03	DQB1*03:01:01:01 DQB1*03:03:02:02	DPA1*02:01:01 DPA1*02:02:02	DPB1*14:01:01 DPB1*05:01:01	A*31:01:02:01	B*15:08:01 B*40:02:01	C*01:02:01 C*03:04:01:02							
IHW0 9317	FORE	European - France	KU593572, KX687283	HWS10025811, HWS09317	DRB5*02:02:01 STR1 ^h DRB4*01:03:01:03	DRB1*16:04 DRB1*04:04:01	DQA1*01:02:02 DQA1*03:01:01	DQB1*05:02:01 DQB1*03:02:01	DPA1*01:03:01:03	DPB1*03:01:01 DPB1*06:01	A*02:01:01:01 A*24:02:01:01	B*51:01:01:01 B*15:01:01:01	C*14:02:01 C*03:03:01							
IHW0 9365	GRC-138	American Indian - Brazil	KU593572	HWS10025811	DRB5*02:02:01 STR1 ^h DRB3*01:01:02	DRB1*16:02:01 DRB1*14:13	DQA1*05:05:01 DQA1*05:03	DQB1*03:01:01:01 DQB1*03:01:01:01	DPA1*01:03:01:05	DPB1*04:02:01:02	A*02:01:01:01 A*02:11:01	B*40:03 B*15:04:01	C*03:04:01:02 C*03:03:01							
	STA1016	European	KU593572	HWS10025811	DRB5*02:02:01 STR1 ^h DRB3*01:01:02	DRB1*16:02:01 DRB1*13:03	DQA1*01:02:02 DQA1*05:05:01:01	DQB1*03:01:01:03 DQB1*03:01:01:01	DPA1*01:03:01:05 DPA1*01:03:01:02	DPB1*04:02:01:02 DPB1*04:01:05	A*24:02:01:01 A*02:01:01:01	B*44:05:01 B*41:02:01	C*02:02:02:01 C*17:05							

International Workshop Cell line #	Local Sample ID	Ethnic Origin	Accession #	IMGIT Submission #	DRB5	CLASS II ALLELES							CLASS I ALLELES		
						DRB1	DQA1	DQB1	DPA1	DPB1	HLA-A	HLA-B	HLA-C		
IHW09112	CHA, AJ	European	KU593578,KX687264	HW810025821,HWS10026687	DRB5*02:02e1_STR1 ^f DRB5*02:02:01:01	DRB1*16:01:01e1 DRB1*03:01:01:02	DQA1*01:02:02 DQA1*05:01:01	DQB1*05:02:01 DQB1*02:01:01	DPA1*01:03:01:04	DPB1*04:01:01	A*03:01:01:01 A*24:03:01	B*39:06:02 B*50:01:01	C*12:03:01:01 C*15:04:01		
	STA1020	Asian	KU593579,KX687282	HWS10025823	DRB5*02:03e1 ^g DRB5*01:01:02	DRB1*16:02:01v1 ^h DRB1*13:02:01	DQA1*01:02:02 DQA1*01:03:01:02	DQB1*05:02:01 DQB1*06:03:01	DPA1*02:02:02	DPB1*02:02 DPB1*05:01:01	A*02:03:01 A*33:03:01	B*38:02:01 B*58:01:01:01	C*07:02:01:01 C*03:02:02:01		

^aDRB5*01:01:01_STR1 is an intronic variant of DRB5*01:01:01 with copy number variations in the intron 2 STR region

^bDRB5*01:01:01v1 is an intronic variant of DRB5*01:01:01

^cDRB5*01:01:01v1_STR1 is an intronic variant of DRB5*01:01:01 with copy number variations in the intron 2 STR region

^dDRB5*01:02e1 is an extended genomic sequence of DRB5*01:02

^eDRB5*01:02e1_STR1 is an intronic variant of DRB5*01:02 with copy number variations in the intron 2 STR region

^fDRB5*01:03e1 is an extended genomic sequence of DRB5*01:03

^gDRB5*01:08Ne1 is an extended genomic sequence of DRB5*01:08N

^hDRB5*02:02e1 is an extended genomic sequence of DRB5*02:02

ⁱDRB5*02:02e1_STR1 is an extended genomic sequence of DRB5*02:02

^jDRB5*02:03e1 is an extended genomic sequence of DRB5*02:03

^kDRB1*16:02:01v2 is an intronic variant of DRB1*16:02:01

^lDRB1*16:02:01v1 is an intronic variant of DRB1*16:02:01

SNPs outside of exon 2 and STRs variations in all DRB5 alleles examined.

Table 2:

Allele	Exon 1		Intron 1 Homopolymers				Intron 2 STR Structure					Intron 2 SNPs						Intron 3 SNPs								
	Codon16(aa)	poly(T)	poly(A)	poly(A)	(GA) ₅	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	(GA) ₄	CA	498	548	565	837	1480	1550	1670	1804	1890	1922	391	412	691	
DRB5*01:01:01	AAG(K)	14	20	(GT) ₂₁	(GA) ₅	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		T	G	T	T	G	T	A	T	C	A	A	A	C	T
DRB5*01:01:01 STR1 ^a				(GT) ₁₉	(GA) ₅	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		T	G	T	T	G	T	A	T	C	A	A	A	C	T
DRB5*01:01:01v1 ^b				(GT) ₁₈	(GA) ₈	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		T	G	T	T	G	T	G	T	C	A	A	A	C	T
DRB5*01:01:01v1 STR1 ^c				(GT) ₂₀	(GA) ₈	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		T	G	T	T	G	T	G	T	C	A	A	A	C	T
DRB5*01:02e1 ^d	AAG(K)	16	19	(GT) ₂₂	(GA) ₈	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA	(GA) ₇	CA	T	G	T	T	G	G	C	C	A	A	A	C	C
DRB5*01:02e1 STR1 ^e				(GT) ₂₁	(GA) ₉	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA	(GA) ₇	CA	T	G	T	T	G	G	C	C	A	A	A	C	C
DRB5*01:03e1 ^f				(GT) ₂₁	(GA) ₉	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA	(GA) ₇	CA	T	G	T	T	G	G	C	C	A	A	A	C	C
DRB5*01:08Ne1 ^g				(GT) ₂₀	(GA) ₉	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA	(GA) ₇	CA	T	G	T	T	G	G	C	C	A	A	A	C	C
DRB5*02:02e1 ^h	GTG(V)	12	16	(GT)	(GA) ₁₁	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		G	A	C	A	C	G	G	C	T	G	C	T	C	
DRB5*02:02e1 STR1 ⁱ				(GT) ₁₅	(GA) ₁₁	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		G	A	C	A	C	G	G	C	T	G	C	T	C	
DRB5*02:03e1 ^j				(GT) ₁₅	(GA) ₁₁	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA		G	A	C	A	C	G	G	C	T	G	C	T	C	
patr-DRB5*01:02	GTG(V)	13	18	(GT) ₄ GA(GT) ₇	(GA) ₇	(GA) ₄	CA	(GA) ₃	(GC AA)	(GA) ₄	CA	(GA) ₃	T	G	C	A	C	G	G	C	C	A	A	A	C	C
manmu-DRB5*03:01				(GT) ₂	(GA) ₂ AAGAAA(GA) ₂	(GA) ₄	(GC) ₄	(GC) ₄	AAGAAA	(GA) ₄	CA		T	G	C	A	C	G	G	T	C	A	A	A	C	C
DRB1*16:01:01:01				(GT) ₁₈	(GA) ₈	(GA) ₆		(GA) ₆	(GGAA)	(GA) ₆																
DRB1*16:02:01:01				(GT) ₁₈	(GA) ₈	(GA) ₆		(GA) ₆	(GGAA)	(GA) ₆																
DRB1*15:01:01:01				(GT) ₂₂	(GA) ₅	(GA) ₄	CA	(GA) ₃	CA	(GGAA)	(GA) ₄															
DRB1*09:01:02				(GT) ₁₀	(GA) ₉	(GA) ₄	CA	(GA) ₄	(GGAA)	(GA) ₄	CA															

^aDRB5*01:01:01_STR1 is an intronic variant of DRB5*01:01:01 with copy number variations in the intron 2 STR region

^bDRB5*01:01:01v1 is an intronic variant of DRB5*01:01:01

^cDRB5*01:01:01v1_STR1 is an intronic variant of DRB5*01:01:01 with copy number variations in the intron 2 STR region

^dDRB5*01:02e1 is an extended genomic sequence of DRB5*01:02

^eDRB5*01:02e1_STR1 is an intronic variant of DRB5*01:02 with copy number variations in the intron 2 STR region

^fDRB5*01:03e1 is an extended genomic sequence of DRB5*01:03

DRB1*16:02:01v1 is an intronic variant of DRB1*16:02:01

DRB1*16:02:01v2 is an intronic variant of DRB1*16:02:01

DRB5*02:03e1 is an extended genomic sequence of DRB5*02:03

DRB5*02:02e1_STR1 is an extended genomic sequence of DRB5*02:02

DRB5*02:02e1 is an extended genomic sequence of DRB5*02:02

DRB5*01:08Ne1 is an extended genomic sequence of DRB5*01:08N

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