Identification of a Recombinogenic Major Histocompatibility Complex Q Gene with Diverse Alleles

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Summary

Structural diversity enables class Ia molecules to present a diverse repertoire of peptides to the T cell receptor. This diversity is thought to be generated by recombinations between class I genes. We have found that two class Ib Q2 alleles exhibit extremely high sequence diversity, even higher than class Ia alleles. Clustered nucleotide differences between Q2 b and Q2 k suggest that this sequence diversity was generated by microrecombinations between Q2 genes and other class I genes. The relatively high expression of Q2 b in the thymus may be significant and perhaps suggests a novel role for a Q2 b product in the education and selection of the T cell repertoire.

Materials and Methods

Genes. The Q1 b and Q2 genes from cosmid clones LSHT36 and H26, respectively, were obtained from Dr. R. Flavell (Yale University, New Haven, CT) (8), and were digested with BamHI, subcloned into pBLUESCRIPT (Stratagene, La Jolla, CA), andideoxy sequenced (Sequenase; United States Biochem. Corp., Cleveland, OH) using gene- and vector-specific primers.

RNA Preparation and PCR Analysis. The preparation of whole cellular RNA was as previously described (9). cDNA was prepared by incubating 2 μg of RNA with 100 ng of Q2 3’ primer (CCCTGAGCTTCTGCTCACCAGC, positions 150-157) in 20 μl of 1 x PCR buffer (PEXPRESS, Norwalk, CT) with additional MgCl₂ to 6.5 mM and 20 U reverse transcriptase (RT) (Life Sciences, Inc., St. Petersburg, FL) for 30 min at 50°C, followed by 95°C for 5 min. Nucleotide sequence in lower-case letters at 5’ of the oligonucleotides represents added restriction enzyme sites. After the addition of 400 ng 3’ primers, 500 ng of Q2 b 5’ primer (tctggagaatCCCTGAAACCTGACTGAGACA, positions -3 to -9) and K b 5’ primer (ttcaagaattCCCTGGCTCCGACTCAGA, positions -3 to -9) 2 U Taq polymerase (PEXPRESS), and 1 x PCR buffer to 100 μl, amplification proceeded by incubation at 94°C for 30 s, 55°C for 30 s, and 72°C for 1 min. After 15 cycles, 5 μl was removed and used as input DNA for an additional 25 cycles. Fractionation by agarose gel electrophoresis, transfer to GenScreen (New England Nuclear, Boston, MA), and hybridization analysis with specific probes was as previously described (9). Q2 b- and Kb-specific RT-PCR were carried out in the same or in different amplification tubes with similar results.

Results and Discussion

Q1 Alleles Are Conserved. Sequence analysis of the Q genes of the C57BL/6 mouse indicated that Q1 b, while potentially coding for an intact class Ib molecule, is one of the most divergent Q genes (10). Sequence analysis of exons 1-3 of the Q1 b gene demonstrates that the two alleles are almost identical, with only two nucleotide differences (Fig. 1 and Table 1). This finding extends the previous observation of allelic sequence conservation of Q loci (4, 11).

Q2 b and Q2 k Are Alleles and Exhibit Sequence Diversity. In contrast to Q1, a nucleotide sequence comparison between Q2 b and Q2 k reveals that the two alleles differ from each...
The asterisk at position 338 of Q2 b indicates the position of a single base insertion in Q2 b. These can be seen in the sequence data presented in Figure 1. The number of differences between listed alleles in the 57 amino acids that make up the ARS in domains 1 and 2 (1).

Table 1. Sequence Differences between Allelic Genes in the H-2^k and H-2^k Haplotypes

| Alleles | No. of nucleotide differences in exons 2 and 3* | No. of amino acid differences in: |
|---------|-----------------------------------------------|---------------------------------|
| K^k/K^k | 51                                            | Domains 1 and 2: 27 14 13        |
| D^k/D^k | 63                                            | ARS: 30 0 0 17                  |
| Q4b/Q4k | 1                                             | Framework: 19                  |
| Q10b/Q10b | 1                                      |                                  |
| Q1b/Q1k | 2                                             |                                  |
| Q2b/Q2k | 97                                            |                                  |

DNA sequences other than Q1b and Q2b are from Kuhner and Goodenow (13). Amino acid sequences other than Q1b and Q2b are from Watts et al. (18).

* The number of differences between listed alleles in the 546 nucleotides that make up exons 2 and 3.
† The number of differences between listed alleles in the 182 amino acids that make up domains 1 and 2.
§ The number of differences between listed alleles in the 57 amino acids that make up the ARS in domains 1 and 2 (1).
†† The number of differences between listed alleles in the 125 amino acids that make up the non-ARS positions in domains 1 and 2.

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other by 97 of the 546 nucleotides in exons 2 and 3 (comprising the α1 and α2 domains), resulting in 46 amino acid replacements (Fig. 1 and Table 1). This level of allelic diversity is even greater than that observed at H-2 loci (Table 1).

The allelic relationship between Q2b and Q2k is supported by both physical mapping studies and critical sequence homologies. Both genes have been cloned in cosmids that are part of large, defined contiguous genomic segments that encompass the entire Q region of their respective haplotypes (8, 10). Q2b and Q2k map as the first genes telomeric (~20 kb) to the Q1 genes in each haplotype (8, 10). Given the conserved nature of Q1 alleles as a benchmark, the mapping data support the idea that Q2b and Q2k are alleles.

Striking sequence homologies further support the allelic relationship between Q2b and Q2k. Flanking Q2k (and Q1k) are a unique 5′ sequence of ~650 nucleotides (beginning 82 nucleotides upstream of the initiation codon and continuing in the 5′ direction) and a unique 3′ sequence of ~500 nucleotides (beginning ~100 nucleotides downstream of the termination codon) (10). Neither of these sequences has a counterpart in the GenBank data library (10). Q2b contains the almost identical unique sequences at precisely the same positions, suggesting an allelic relationship between the two genes (Lapierre, L. A., and J. Geliebter, manuscript in preparation).

Further, coding and intron sequence homologies also suggest an allelic relationship between Q2b and Q2k. It has previously been shown that class I alleles exhibit sequence conservation in the 3′ portion of their genes, while nonallelic genes often display sequence diversity (12, 13). For example, in exon 5 of the H-2b and H-2k haplotypes, K, D, Q4, and Q10 loci display >95% allelic sequence conservation, but <86% sequence identity between nonallelic genes (13). Q2b and Q2k are ~95% identical in exons 5–8, and introns 5–7, which is in line with expectations for alleles (Fig. 1, and unpublished data). Therefore, based on sequence conservation and physical mapping it can be inferred that Q2b and Q2k are alleles.

With the exception of CCAAC (instead of CCAAT) and TATAA sequences, none of the usual H-2 and Q regulatory sequences have been found in Q2b or Q2k, a situation also found in TL region genes (10, 11, and Lapierre, L. A., and J. Geliebter, manuscript in preparation). Splice donor and acceptor sites are intact in all seven introns (data not shown), and Q2b would be expected to code for a 339-amino acid integral membrane glycoprotein. It is also possible that Q2b could be anchored to the membrane by a phosphatidylinositol (PI) linkage as with the Qa-2 protein (gene QT) (11).

Due to a single base insertion in exon 7, Q2b translation would terminate at the TAA stop codon. 3′ acceptor sites are intact in all seven introns (data not shown), and Q2b would be expected to code for a 339-amino acid integral membrane glycoprotein. It is also possible that Q2b could be anchored to the membrane by a phosphatidylinositol (PI) linkage as with the Qa-2 protein (gene QT) (11).

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Figure 2. Potential microrecombination sites in the Q2 gene. Numbers on top of nucleotide sequences indicate residue number according to the K8 sequence. Dashes beneath the top sequence indicate identity to that sequence. Sequences other than Q2b are from Kuhner and Goodenow (13) and Bronson et al. (25). The asterisk in the intron 2 sequence of Q2k indicates the position of a single base insertion in Q2b. Data are not meant to imply that all or any of the differences between Q2b and Q2k were generated since the establishment of inbred mouse strains, rather that potential donor sequences, are available.

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that Q2 may be expressed on the cell surface and function as a recognition structure.

Q2 Appears to Have Undergone Multiple Microrecombinations. Many of the nucleotide differences between Q2 and Q2 are clustered and can be found at homologous positions in other class I genes (Figs. 1 and 2). These characteristics are taken to reflect past microrecombination or gene conversion-like events with nonallelic class I genes (20). Kuhner et al. (21) analyzed sequences from 23 class I genes using a statistical algorithm and detected 25 past conversion events among all the genes analyzed. Visual examination of Q2, Q2, and six class I sequences of the H-2 haplotype revealed evidence for more than six events (Fig. 2).

While most examples of past Q2 microrecombination events are in exons 2 and 3, examples are seen in introns and the transmembrane exon 5 (Fig. 2). It is not always certain which Q2 gene was the recipient of genetic information. However, in exon 5, Q2 differs from the published sequences of both Q2 and Q2 by the same nucleotides that are found in the D gene. This implies that Q2 was probably the recipient of genetic information in a microrecombination event with a D-like gene.

Q2 Sequence Diversity Appears to Be the Result of Selection. The reassortment of class I sequences into H-2 genes has been suggested to provide the raw material for diversification, to be acted upon by natural selection (22). The lack of diversity and polymorphism of Q loci may reflect a unidirectional bias in conversion from Q genes to H-2 genes (22). Alternatively, since most conversion events are lost in the absence of selection for the novel products, Q gene sequence conservation may simply reflect the lack of strong selection pressure for diversification (20). If the latter is correct, Q2 gene diversity and apparent maintenance of microrecombination products may reflect selection and imply a function more closely related to class Ia molecules than class Ib molecules. This notion is supported by two related observations; a very high percentage of amino acid replacements between the two alleles are in the antigen recognition site (ARS), and an unusually high percentage of nucleotide substitutions in the ARS result in amino acid replacements (see below).

The 3 and 2 domains of class Ia molecules contain 182 amino acids. Of these, 125 are in framework positions, and 57 are designated ARS positions (1). Of the ~30 amino acid differences between R and R or D and D, 50% are in the ARS, despite ARS positions constituting only 31% of positions (Table 1). Likewise, of the 49 amino acid differences between Q2 and Q2 in the 3 and 2 domains, 57% are in the ARS (Table 1). Thus, allelic differences between H-2 and Q2 molecules are predominantly in the ARS, where they affect peptide binding and interactions with the TCR.

A high ratio of nonsynonymous to synonymous nucleotide substitutions in the ARS is consistent with positive selection for sequence diversity. For class Ia, in the ARS, the rate of nonsynonymous substitutions per nonsynonymous site (ds) is 21.4 ± 2.4, while the rate of synonymous substitutions per synonymous site (dS) is 13.8 ± 3.2, indicating positive selection for diversification (23). Comparing Q2 and Q2 under the same analysis yields a dS of 38.2 ± 6.8 and a ds of 22.15 ± 8.2, also indicating selection for diversity between the two alleles (Lorraine Flaherty, personal communication).

These data indicate that like class Ia, structural differences between Q2 alleles could result in functional diversity, and that these differences may be maintained by natural selection.

Q2 Is Transcribed in Various Tissues. cDNA synthesis followed by RT-PCR of Q2 mRNA indicates that Q2 transcription is highest in the thymus (Fig. 3). Lower expression was also observed in the lung and kidney, and with prolonged exposure of the hybridized filter to film, most tissues showed a band of the appropriate size (PCR controls remained negative). Since RT-PCR is not quantitative and tissues were not perfused, weak expression in some tissues may reflect a disproportionate contribution of certain cell types. Yet, the above data do clearly show that Q2 is transcribed and may be translated. The relatively high expression of Q2 in the thymus may be significant and perhaps suggests a novel role for a Q2 product in the education of and selection of the T cell repertoire.

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