LETTER

The Complex Mutation Pattern of a Microsatellite

Claudia Macaubas,^{1,5} Li Jin,² Joachim Hallmayer,^{2,4} Akinori Kimura,³ and Emmanuel Mignot¹

¹Center for Narcolepsy, Department of Psychiatry, Stanford University, Palo Alto, California 94304;

²Department of Genetics, Stanford University, Palo Alto, California 94304; ³Department of Tissue Physiology, Medical Research Institute, Tokyo Medical and Dental University, Tokyo 101, Japan

DQCAR is a (CA)_n microsatellite located in the HLA class II region and tightly linked to *HLA-DQB1*. Previous studies showed a strikingly low level of size variation in *DQCAR* alleles within an extensive subfamily of *HLA-DQ* subtypes (*DQI*). *DQCAR* alleles in non-*DQI* subtypes showed a higher degree of size polymorphism. In this study sequence analysis demonstrates that *DQI*-associated *DQCAR* alleles have a single C → A nucleotide substitution interrupting the CA repeat array. Frequent CA → GA mutations are also observed in *DQI*-associated microsatellites with identical allele sizes. In contrast, *DQCAR* alleles associated with non-*DQI* haplotypes display a perfect CA repeat sequence and the variation in allele size is attributable only to differences in the number of CA repeats. Our results imply that several mutational mechanisms are involved in the generation of allelic diversity within the same microsatellite locus. The possibility of different mutation rates in the same locus should be taken into account when using these markers in evolutionary and disease studies.

[The sequence data described in this paper have been submitted to the GenBank data library under accession nos. U96944–U96962 and S87165.]

Microsatellites are abundant in mammalian genomes (Weber 1990; Stallings et al. 1991) and have attracted an enormous amount of interest in recent years. Instability of trinucleotide repeats is involved with certain neurodegenerative conditions (for review, see Rosenberg 1996), suggesting that underlying mutational mechanisms may be responsible for genetic morbidity. Polymorphic dinucleotide repeats have been used extensively for gene mapping (Dib et al. 1996), forensic identification (Edwards et al. 1992) and studying the history of human population (Bowcock et al. 1994; Deka et al. 1995).

In spite of their popularity as a genetic tool, the mechanisms underlying microsatellite allelic diversity are still poorly understood. Estimations of mutation rate for microsatellites range between 10⁻² and 10⁻⁵ per generation (Edwards et al. 1992; Mahtani and Willard 1993), with Weber and Wong (1993) observing that dinucleotide repeats have a lower average mutation rate than tetranucleotides (5.6 × 10⁻⁴ and 2.1 × 10⁻³, respectively). Most of

the observed changes in microsatellite allele sizes are attributable to deletion or addition of a few repetitive units (Straub et al. 1993; Deka et al. 1995). In principle, this pattern of mutation fits with a simple stepwise mutation model (Ohta and Kimura 1973) but might not account for all the variation observed (Shriver et al. 1993; Valdes et al. 1993; Deka et al. 1995). A two-phase model that allows for bigger differences in repetitive units has also been proposed (Di Rienzo et al. 1994). Strand slippage during replication has been suggested as the most likely mechanism involved in the generation of microsatellite mutations (Levinson and Gutman 1987; Schlötterer and Tautz 1992; Strand et al. 1993).

Most of the studies involving microsatellites have analyzed a large number of loci. Detailed sequence analyses of mutation patterns at individual loci have been performed especially for trinucleotides associated with diseases (Rosenberg 1996) and some tetranucleotides (Mahtani and Willard 1993; Talbot et al. 1995). Minisatellites have also been analyzed (Jeffreys et al. 1994). Dinucleotides, however, have received less attention.

Previously, we studied a (CA)_n microsatellite (*DQCAR*) located in the human leukocyte antigen (HLA) class II region, ~12 kb centromeric to *HLA-DQA1* and 1–2 kb telomeric to the *HLA-DQB1* gene

⁴Present address: Graylands/UWA Centre for Clinical Research in Neuropsychiatry, Gascoyne House, Graylands Hospital, WA 6010, Australia.

⁵Corresponding author. Present address: TVW Telethon Institute for Child Health Research, West Perth, WA 6872, Australia. E-MAIL macaubas@icmr.uwa.edu.au; FAX 61-9-388-3414.

(Satyanarayana and Strominger 1992). *DQA1* and *DQB1* encode for the respective α and β chains, which form the heterodimeric protein *HLA-DQ* expressed in the surface of cells involved in the immune response. Fourteen different *DQCAR* alleles were identified and these were found to be in tight linkage disequilibrium with *HLA-DQ* (Macaubas et al. 1995; Mignot et al. 1995; Jin et al. 1996). Extremely low size polymorphism was found within *DQCAR* alleles associated with *DQ1*, the predominant *HLA-DQ* subtype in several populations (Imanishi et al. 1992). The *DQCAR* repeat associated with the other DQ subtypes (*DQ2*, *DQ3*, and *DQ4*) showed both larger numbers and sizes of alleles (Macaubas et al. 1995). In this study we carried out a sequence analysis of *DQCAR* alleles to investigate the basis for the observed allelic differences between *DQ1* and non-*DQ1* haplotypes.

RESULTS

The phylogeny of *DQB1* alleles is shown in Figure 1

(left). There are two groups of alleles—the *DQ1* subgroup (*DQB1*05* and *DQB1*06*) and non-*DQ1* (*DQB1*02*, *DQB1*03*, and *DBQ1*04*). The typing of 2140 chromosomes (Fig. 1, column 2), showed that in the non-*DQ1* subfamily, more *DQCAR* alleles per *DQB1* subtypes were found (Fig. 1, column 3) in comparison with the *DQ1*-associated subgroup. Non-*DQ1*-associated *DQCAR* alleles also have longer microsatellite sizes (Macaubas et al. 1995; Jin et al. 1996). The exceptions were *DQCAR* alleles associated with *DQB1*0201*, with only two short microsatellite alleles detected in >150 chromosomes typed.

At least one *DQCAR* allele from 15 *DQB1* lineages was then subcloned and sequenced. All *DQCAR* alleles associated with the *DQ1* subfamily showed a C $\rightarrow$ A nucleotide substitution in the fifteenth base pair, resulting in a CAAA motif interrupting the CA repeat array (Table 1; Fig. 2). In contrast, the microsatellite allele sequences associated with non-*DQ1* subfamilies showed an uninterrupted CA repeat motif (Table 1; Fig. 2).

Figure 1 Phylogenetic relationships of *DQB1* alleles (left) and *DQCAR* alleles (right) based on their sequence data. Phylogenies of 14 *DQB1* and 15 *DQCAR* alleles were reconstructed as detailed in Methods. (a) Number of chromosomes typed for *DQB1* and *DQCAR* in a mixed population. (b) Number of *DQCAR* alleles associated with each *DQB1* allele in the mixed population. (c) Number of uninterrupted CA repeats for each *DQCAR* allele observed in the mixed population, deduced from the sequenced alleles on Table 1. (d) The *DQCAR* alleles represented in the tree are the sequenced from Table 1. Areas with high *DQCAR* size variation (more than two *DQCAR* alleles found in association with a lineage) are boxed.

Table 1. *DQCAR* Sequences Observed in Association with Various *HLA-DQB1* Alleles

DQ	DQB1 line	Cell	DQCAR allele ^a	Sequence						Repeat motif
				10	20	30	40	50	60	
1(6)	*0602	9010	103	CAGACACACA	CACAAACACA	CACACACACA	CA-----	-----	GAGAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 8 (GA) 4
1(6)	*0602	9083	103	CAGACACACA	CACAAACACA	CACACACACA	CA-----	-----	GAGAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 8 (GA) 4
1(6)	*0603	9062	103	CAGACACACA	CACAAACACA	CACACACACA	CACA-----	-----	--GAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 9 (GA) 3
1(6)	*0604	9063	103	CAGACACACA	CACAAACACA	CACACACACA	CACA-----	-----	--GAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 9 (GA) 3
1(6)	*0609	9055	103	CAGACACACA	CACAAACACA	CACACACACA	CACA-----	-----	--GAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 9 (GA) 3
1(5)	*0502	9012	103	CAGACACACA	CACAAACACA	CACACACACA	CACA-----	-----	--GAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 9 (GA) 3
1(5)	*0501	9080	103	CAGACACACA	CACAAACACA	CACACACACA	CACACA----	-----	---GAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 10 (GA) 2
1(5)	*0501	9004	103	CAGACACACA	CACAAACACA	CACACACACA	CACACA----	-----	---GAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 10 (GA) 2
1(5)	*05031	9054	107	CAGACACACA	CACAAACACA	CACACACACA	CACACACA--	-----	--GAGAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 11 (GA) 3
1(6)	*0601	9011	107	CAGACACACA	CACAAACACA	CACACACACA	CACACACACA	-----	---GAGATG AGGTAAGA	CAGA (CA) 5AA (CA) 12 (GA) 2
2	*0201	9019	99	CAGACACACA	CACACACACA	CA-----	-----	----GCAAGA	GAGAGAGATG AGATAAGA	CAGA (CA) 9 GCAA (GA) 5
3(8)	*0302	9029	111	CAGACACACA	CACACACACA	CACACACACA	CACA-----	----GCAAGA	GAGAGAGATG AGATAAGA	CAGA (CA) 15 GCAA (GA) 5
3(8)	*0302	9034	111	CAGACACACA	CACACACACA	CACACACACA	CACA-----	----GCAAGA	GAGAGAGATG AGATAAGA	CAGA (CA) 15 GCAA (GA) 5
3(7)	*0301	9070	113	CAGACACACA	CACACACACA	CACACACACA	CACAGA----	----GCAAGA	GAGAGAGATG AGATAAATA	CAGA (CA) 15GA GCAA (GA) 5
4	*0401	N*106	113	CAGACACACA	CACACACACA	CACACACACA	CACACA----	----GCAAGA	GAGAGAGATG AGATAAGA	CAGA (CA) 16 GCAA (GA) 5
4	*0402	9072	115 ¹	CAGACACACA	CACACACACA	CACACACACA	CACACACA--	----GCAAGA	GAGAGAGATG AGATAAGA	(CAGA) 2 (CA) 15 GCAA (GA) 5
3(9)	*03032	9052	117 ¹	CAGACACACA	CACACACACA	CACACACACA	CACACACACA	----GCAAGA	GAGAGAGATG AGATAAGA	CAGA (CA) 18 GCAA (GA) 5
3(7)	*0301	9035	119 ¹	CAGACACACA	CACACACACA	CACACACACA	CACACACACA	CA--GCAAGA	GAGAGAGATG AGATAAATA	CAGA (CA) 19 GCAA (GA) 5
2	*02	MANN	121	CACACACACA	CACACACACA	CACACACACA	CACACACACA	CACAGCAAGA	GAGAGAGATG AGATAAATA	(CA) 22 GCAA (GA) 5
2	*0202	9051	121	CACACACACA	CACACACACA	CACACACACA	CACACACACA	CACAGCAAGA	GAGAGAGATG AGATAAATA	(CA) 22 GCAA (GA) 5

Sequences were obtained as described in Methods.

^a(1) Actual size of the fragment sequenced (2 bp less than the allele defined initially; Macaubas et al. 1995). *DQCAR* primers sequence are also included in the size of the allele.

Non-*DQ1*-associated alleles differed in size as a result of varying numbers of CA repeats. *DQCAR* 99 (the shortest allele sequenced) has a repeat array of 9 uninterrupted CA units, and *DQCAR* 121 (the longest) has a stretch of 22 CA units. Some changes from C → G into the repeat array were also observed in some alleles on both sides of the CA repeat (Table 1).

Each *DQCAR* allele is composed of a CA repeat, followed by a GA motif. Non-*DQ1*-associated alleles always show an intervening tetramer GCAA between CA and GA motifs. In *DQ1*-associated alleles the GA repeat structure lies immediately next to the CA repeat array (Table 1; Fig. 2). The GA repeat motif in non-*DQ1*-associated alleles is monomorphic, with five repeats of GA (Table 1; Fig. 2). The number of GA repeats varies for *DQ1*-associated alleles, even when the sizes of two alleles, are identical. For example, *DQCAR* 103 alleles associated with *DQB1**0602 and with *DQB1**0603 have the motif (CA)8(GA)4 and (CA)9(GA)3, respectively. The total number of repetitive units for those alleles remains constant (i.e., 12 units) while their composition varies. Using routine genotyping methods, this variation is missed.

The sequence flanking the 3' end of *DQCAR* showed subtype-specific variations. *DQ1*-associated alleles showed a G nucleotide in the fifth nucleotide after the GA run (Table 1; Fig. 2). Non-*DQ1* alleles have an A in this position. Sequences from *DQB1**0202 and *DQB1**0301 reveal a T in the ninth nucleotide after the GA stretch. All of the other sequences (including *DQ1* sequences) have a G in this position (Table 1).

A phylogenetic tree of *DQCAR* alleles, based on their flanking sequence, was then constructed (Fig. 1, right). As for *DQB1* alleles, two groups were observed: *DQCAR* alleles associated with *DQ1* subtypes formed one group, whereas microsatellite alleles observed with non-*DQ1* subtypes clustered in a separate group. The number of uninterrupted CA repeats for each *DQCAR* allele observed in the typed population was deduced from the sequence data (Fig. 1, column 4). The non-*DQ1*-associated *DQCAR* alleles showed a higher number of uninterrupted CA repeats (13–24 CA repeats), compared to *DQCAR* associated with *DQ1* haplotypes (between 8 and 12 CA repeats). The only exceptions were *DQCAR* alleles associated with *DQB1**0201, with 9 and 10 repeats.

Figure 2 Sequence from two representative cell lines. (Left) Cell line 9072 (*DQ4*, *DQB1*0402*, *DQCAR117*); (right) cell line 9062 (*DQ1*, *DQB1*0603*, *DQCAR103*). The left arrow points to the A nucleotide interrupting the CA repeat sequence in the *DQ1* cell line. Note the tetramer GCAA between the (CA)_n region and the (GA)₅ in the *DQ4* cell line. The right arrow points to the G substitution in the flanking region, characteristic of *DQCAR103* and *DQCAR107*. Sequence of opposite strand confirmed the results in both cases (not shown). Sequence was performed as detailed in Methods.

DISCUSSION

In the *DQCAR* locus, three forms of mutations are responsible for the generation of new alleles: (1) simple gain or loss of CA units; (2) a single C → A substitution in the CA repeat array; and (3) GA → CA or CA → GA mutations in the region immediately following the CA repeat array. Only the first mutation mechanism resulted in a change of allele size.

We observed that for *DQCAR* alleles with perfect CA-repeat runs, the change in allele size is attributable to small differences in the number of repetitive units, mostly by only 2 bp. This pattern of mutation seems to fit with a model of small changes in repeat unit number via a strand-slippage mechanism (Strand et al. 1993). In addition, the increase was more remarkable above 13 repeat units, suggesting, as pointed out by Weber (1990), that informativeness tends to increase with the number of repeats.

The interruption of the perfect repeat array had

a dramatic effect on the stability of the *DQCAR* alleles. It has been shown that imperfect repeats are less polymorphic (Weber 1990; Garza et al. 1995), and, interestingly, stabilization of repetitive sequences has also been described for certain trinucleotide repeats associated with diseases. In spinocerebellar ataxia type I (SCA1), Chung et al. (1993) showed that expanded alleles had a continuous run of (CAG)_n repeats in the SCA1 region, whereas unexpanded alleles had CAT interruptions. The gene for spinocerebellar ataxia 2 (SCA2) has been identified recently and also contains a stretch of (CAG)_n repeats uninterrupted in mutated alleles and interrupted by one to three CAAs in normal ones (Imbert et al. 1996). Similarly, fragile X syndrome is associated with an expansion of an almost uninterrupted run of triple repeat (CGG)_n at the 5'-untranslated region of the *FMR1* gene with stable alleles showing interspersed AGGs (Eichler et al. 1994; Zhong et al. 1995).

It is not totally understood how interruptions stabilize repetitive sequences. Interruptions may create more matching points during replication, and a misalignment could be detected more easily by mismatch repair enzymes, as suggested by Heale and Petes (1995). They observed that in yeast, the interruption of a repetitive run by one different repeat increased by 100 times its stability, and this stabilization was dependent on a functional DNA mismatch repair system.

Interestingly, the phylogeny of *DQB1/DQCAR* haplotypes suggests that the C → A mutation interrupting the CA repeat is monophyletic and therefore occurred only once in the evolution of this locus. The interruption of the CA repeat array in *DQ1*-associated *DQCAR* alleles must have been present before the divergence of this subtype, as it was found in all *DQCAR* alleles associated with *DQ1* haplotypes and in none of the non-*DQ1* haplotypes. The four major DQ subfamilies (*DQ1*, *DQ2*, *DQ3*, and *DQ4*) can be identified in nonhuman primates. They have persisted for at least 5 million years, and diversity within the subfamilies seems to have arisen after speciation (Gyllenstein et al. 1990).

Despite the low level of size variations in the *DQCAR* alleles associated with *DQ1* haplotypes, additional sequence diversity was observed between alleles of identical size because of CA → GA changes at the end of the repeat. As the number of CA units increases, a concomitant decrease in the number of GA units keeps the overall allele size constant. Additional polymorphism is thus obtained for alleles of similar size, as observed previously at other microsatellite loci (Saha et al. 1993; Estoup et al. 1995).

Sequence variation has also been observed in minisatellites (Jeffreys et al. 1994). The variation in the repetitive sequence is not restricted to *DQCAR* alleles associated with *DQ1* alleles but is also found in alleles associated with non-*DQ1* haplotypes (e.g., *DQB1*0301* and *DQB1*0402*). Nevertheless, the frequent change of allelic composition of CA and GA repeat numbers in alleles of identical size suggests a new mutational mechanism that has not been reported previously in microsatellites.

In conclusion, sequence analysis shows a complex mutational pattern in a single (CA)_n microsatellite. This complexity could also occur in other microsatellite loci and must be considered when using these markers for evolutionary and gene mapping studies.

METHODS

Tree Construction

A phylogeny of 14 frequent *DQB1* alleles was reconstructed on the basis of the aligned coding sequences of these alleles using the neighbor-joining method (Saitou and Nei 1987). Genetic distances were estimated using Kimura's two-parameter model (Kimura 1980). Several other measures of genetic distances were used, and all gave identical results in terms of the topology of the phylogeny obtained using the neighbor-joining method (Jin et al. 1996). The maximum parsimony tree displayed a slightly different topology, which became identical with the neighbor-joining tree once peptide-binding sites were removed from the analysis. The tree was rooted by *DQB* sequences of pig and horse. A maximum parsimony tree of *DQCAR* locus based on the flanking sequences of *DQCAR* alleles was also constructed.

Microsatellite Amplification and Sequence

Genomic DNA (200 ng) was used as a template, in a total volume of 25 μ l of 10 mM Tris-HCl, 50 mM KCl, 1.5 mM MgCl₂, 50 pmoles of each primer (see below), 125 μ M of each dNTP, and 1 unit of AmpliTaq DNA polymerase (Perkin Elmer). After an initial incubation for 5 min at 95°C, the amplification was carried for 30 cycles, with each cycle consisting of 30 sec at 94°C, 1 min at 57°C, and 1 min at 72°C, with a final extension at 72°C for 15 min. The primers used were CAR1, 5'-GAAACATATATTAACAGAGACAGACAAA-3'; and CAR2, 5'-CATTCTCTTCTTATCACTTCATA-3' (Satyanarayana and Strominger 1992; Macaubas et al. 1995). The presence of PCR product was confirmed by gel electrophoresis, followed by extraction with chloroform and ligation to pT7Blue T-vector (Novagen), as detailed in the product manual. Both strands of plasmid DNA from several clones were sequenced with Sequenase version 2.0 T7 DNA Polymerase (U.S. Biochemical), using T7 and U19 primers, following the manufacturer's instructions.

Cell Lines

Twenty HLA class II homozygous B cell lines were analyzed.

Eighteen B cell lines were obtained from the Tenth Histocompatibility Workshop panel (Yang et al. 1987; Kimura et al. 1992; Aldener and Olerup 1993). The other two B cell lines were cell line MANN (donated by Dr. George Blanck, Harvard University, Cambridge, MA; Blanck and Strominger 1988) and B cell line no. 106, acquired from the UCLA DNA reference panel (Dr. P.I. Terasaki, University of California, Los Angeles).

HLA Typing

Chromosomes (2140) from several populations were typed for *HLA-DQB1*, *HLA-DQA1* and *DQCAR* loci, and haplotypes were deduced as described in Kimura et al. (1992) and Jin et al. (1996).

ACKNOWLEDGMENTS

This work was funded by National Institutes of Health (NIH) grants NS23724 and 33797 (to E.M.) and by NIH training grant T32-GS08404 (to L.J.) We thank Drs. P. Underhill and P. Oefner for suggestions regarding cloning and sequencing strategies, and Professor Jonathan Majer for reviewing the manuscript.

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Received October 7, 1996; accepted in revised form April 4, 1997.

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Claudia Macaubas, Li Jin, Joachim Hallmayer, et al.

Genome Res. 1997 7: 635-641

Access the most recent version at doi:[10.1101/gr.7.6.635](https://doi.org/10.1101/gr.7.6.635)

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