

Nomenclature of the Human T Cell Receptor Genes

The human T cell receptors (TcR) α - β and γ - δ are the products of four sets of genes on two chromosomes: T cell receptors α (*TRA*) and δ (*TRD*) on chromosome 14 at 14q11.2, T cell receptor β (*TRB*) on chromosome 7 at 7q35, and T cell receptor γ (*TRG*) on chromosome 7 at 7p15-p14.

This appendix presents tabulated lists of the human TcR α , β , γ , and δ genes (Table A.10.1, Table A.10.2, Table A.10.3, and Table A.10.4, respectively) named in accordance with the International ImMuNoGeTics database (IMGT; see Internet Resources) and approved by the Human Genome Organization (HUGO) Nomenclature Committee in 1999. Two additional tables list corresponding nomenclatures for these genes (Table A.10.5 and A.10.6).

HUMAN T CELL RECEPTOR LOCI

TRA Locus

The human *TRA* locus at 14q11.2 spans 1000 kb (Fig. A.10.1; Table A.10.1). It consists of 54 *TRAV* genes belonging to 41 subgroups, 61 *TRAJ* segments localized on 71 kb, and a unique *TRAC* gene (Folch et al., 2000; Scaviner and Lefranc, 2000a,b). The most 5' *TRAV* genes occupy the most centromeric position, whereas the *TRAC* gene, 3' of the locus, is the most telomeric gene in the *TRA* locus. The organization of the *TRAJ* genes over a large area is quite unusual and has not been observed in the other immunoglobulin or T cell receptor loci (Lefranc, 1990b). Moreover, the *TRD* locus is nestled in the *TRA* locus between the *TRAV* and *TRAJ* genes. V-J-rearrangements in the *TRA* locus therefore result in deletion of the *TRD* genes localized on the same chromosome (Lefranc, 1990b). Deletion occurs in two steps, i.e., a deletion of the *TRD* genes, involving specific sequences located upstream from *TRDC* (sequence $\phi J\alpha$; De Villartray et al., 1988; Begley et al., 1989) would take place before the *TRAV*-J rearrangement. The potential repertoire consists of 45 to 47 functional *TRAV* genes belonging to 33 to 35 subgroups, 50 functional *TRAJ* genes, and the unique *TRAC* gene. Among the variable genes, five genes are included, designated as *TRAV/DV*, which belong to five different subgroups and which have been found rearranged either to *TRAJ* or to *TRDD* genes and

can therefore be used in the synthesis of α or δ chains.

The total number of human *TRA* genes per haploid genome is 116, of which 96 to 98 genes are functional. Enhancer sequences have been characterized 4.5 kb 3' from *TRAC* (Ho et al., 1989).

TRB Locus

The human *TRB* locus at 7q35 spans 620 kb (Fig. A.10.2; Table A.10.2). It consists of 64 to 67 *TRBV* genes belonging to 32 subgroups. Except for *TRBV30*, localized downstream of the *TRBC2* gene, in inverted orientation of transcription, all the other *TRBV* genes are located upstream of a duplicated D-J-C-cluster, which comprises, for the first part, one *TRBD* gene, six *TRBJ* genes, and the *TRBC1* gene, and for the second part, one *TRBD* gene, eight *TRBJ* genes, and the *TRBC2* gene (Folch and Lefranc, 2000a,b; Folch et al., 2000). The most 5' *TRBV* genes occupy the most centromeric position, whereas the *TRBV30* gene, 3' of the locus, is the most telomeric gene in the *TRB* locus. The potential repertoire consists of 40 to 48 functional *TRBV* genes belonging to 21 to 23 subgroups, the two *TRBD* genes, thirteen *TRBJ* genes (6 from the first cluster and 7 from the second cluster), and the two *TRBC* genes. Six *TRBV* orphans have been localized on chromosome 9 at 9p21. The total number of human *TRB* genes per haploid genome is 82 to 85, 88 to 91 if the orphans are included, of which 57 to 65 are functional. Enhancer sequences have been characterized 5.5 kb 3' from *TRBC2* (Gottschalk and Leiden, 1990).

TRG Locus

The human *TRG* locus at 7p15-p14 spans 160 kb (Fig. A.10.3; Table A.10.3). It consists of 12 to 15 *TRGV* genes belonging to 6 subgroups, upstream of a duplicated J-C-cluster, which comprises, for the first part, three *TRGJ* genes and the *TRGC1* gene, and for the second part, two *TRGJ* genes and the *TRGC2* gene (Lefranc and Rabbitts, 1989, 1990a,b; Lefranc et al., 1989; Folch et al., 2000). The most 5' *TRGV* genes occupy the most centromeric position, whereas the *TRGC2* gene, 3' of the locus, is the most telomeric in the *TRG* locus. The potential repertoire consists of 4 to 6 functional *TRGV* genes belonging to two subgroups, the

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Abbreviations
and Useful Data

A.10.1

Supplement 40

5 *TRGJ* genes and the 2 *TRGC* genes. Polymorphisms in the number of *TRGV* genes and in the exon number of the *TRGC2* gene have been described in different populations (Buresi et al., 1989; Ghanem et al., 1989, 1991). The total number of human *TRG* genes per haploid genome is 19 to 22 of which 11 to 13 are functional. Enhancer and silencer sequences have been characterized 6.5 kb downstream of the *TRGC2* gene (Lefranc and Alexandre, 1995).

TRD Locus

The human *TRD* locus (Fig. A.10.1; Table A.10.4) at 14q11.2 comprises a cluster of one *TRDV* gene (*TRDV2*), three *TRDD* genes, and four *TRDJ* genes, upstream of the unique *TRDC* gene (Table A.10.4); another *TRDV* gene (*TRDV3*) is localized downstream of the *TRDC* gene, in inverted orientation of transcription (Lefranc, 1990a; Folch et al., 2000). This cluster spans 60 kb and is localized inside the *TRA* locus, between the *TRAV* genes and the *TRAJ* genes (Fig. A.10.1). The *TRD* locus also consists of one *TRDV* (*TRDV1*) localized at 360 kb upstream of the *TRDC* gene, among the *TRAV* genes, and the five genes described above as *TRAV/DV*. The *TRDV* genes are unique members of different subgroups. All the *TRD* genes are functional, with the exception of one *TRAV/DV*, which has been found either functional or as a pseudogene.

The total number of human *TRD* genes per haploid genome is 11 (16 if the 5 *TRAV/DV* are included). All *TRD* genes are functional. Enhancer sequences have been described between the *TRDJ3* and the *TRDC* gene (Bories et al., 1990; Redondo et al., 1990).

TABLE GUIDE

Gene names are according to the IMGT gene name nomenclature for Ig and TcR of all vertebrates (IMGT Scientific chart; <http://imgt.cines.fr:8104>).

Criteria of functionality (F: functional, P: pseudogene, ORF: open reading frame) are described in the IMGT Scientific chart (Lefranc, 1998).

For each gene, an IMGT reference sequence accession number is given. For the functional or ORF genes, the IMGT reference sequence accession number is that corresponding to the allele *01. Although the IMGT accession numbers are the same as those from the EMBL/GenBank/DBJ generalist databases, the content of the IMGT/LIGM-DB flat files differs by the expert annotations added by IMGT. IMGT data

are available from IMGT/LIGM-DB, IMGT Repertoire, and from SRS sites (available from the IMGT Home page).

The number of alleles is according to the Tables of Alleles and Alignments of Alleles, in the IMGT Repertoire (<http://imgt.cines.fr:8104>), with a dash indicating that the allele polymorphism of the pseudogenes has not been studied.

Gene accession ID in the Genome Database (GDB; <http://www.gdb.org>), and in the NCBI LocusLink (<http://www.ncbi.nlm.nih.gov/LocusLink>; see Internet Resources) are provided.

IMGT gene names and IMGT gene definitions for the human Ig and TcR genes have been approved by the Human Genome Organization (HUGO) Nomenclature Committee in 1999. Note that in the HUGO symbols (<http://www.gene.ucl.ac.uk/nomenclature>) slashes and parentheses are omitted, and capital letters replace the lowercase letters found in some provisional IMGT gene names. Otherwise the gene symbols and all the full names (including slashes and parentheses) are identical in IMGT and HUGO nomenclatures.

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LITERATURE CITED

- Arden, B., Clark, S.P., Kabelitz, D., and Mak, T.W. 1995. Human T-cell receptor variable gene segment families. *Immunogenetics* 42:455-500.
- Baer, R., Lefranc, M.-P., Minowada, J., Forster, A., Stinson, M.A., and Rabbitts, T.H. 1986. Organization of the T-cell receptor α -chain gene and rearrangement in human T-cell leukemias. *Mol. Biol. Med.* 3:265-277.
- Begley, C.G., Aplan, P.D., Davey, M.P., de Villartay, J.P., Cohen, D.I., Waldmann, T.A., and Kirsch,

- I.R. 1989. Demonstration of delta rec-pseudo J alpha rearrangement with deletion of the delta locus in a human stem-cell leukemia. *J. Exp. Med.* 170:339-342.
- Bories, J.C., Loiseau, P., d'Auriol, L., Gontier, C., Bensussan, A., Degos, L., and Sigaux, F. 1990. Regulation of transcription of the human T cell antigen receptor delta chain gene. A T lineage-specific enhancer element is located in the J delta 3-C delta intron. *J. Exp. Med.* 171:75-83.
- Buresi, C., Ghanem, N., Huck, S., Lefranc, G., and Lefranc, M.-P. 1989. Exon duplication and triplication in the human T-cell receptor gamma constant region genes and RFLP in French, Lebanese, Tunisian, and black African populations. *Immunogenetics* 29:161-172. [Published erratum appears in *Immunogenetics* 30:148.]
- De Villartay, J.P., Hockett, R.D., Coran, D., Korsmeyer, S.J., and Cohen, D.I. 1988. Deletion of the human T-cell receptor delta-gene by a site-specific recombination. *Nature* 335:170-174.
- Folch, G. and Lefranc, M.-P. 2000a. The human T cell receptor beta variable (TRBV) genes. *Exp. Clin. Immunogenet.* 17:42-54.
- Folch, G. and Lefranc, M.-P. 2000b. The human T cell receptor beta diversity (TRBD) and beta joining (TRBJ) genes. *Exp. Clin. Immunogenet.* 17:107-114.
- Folch, G., Scaviner, D., Contet, V., and Lefranc, M.-P. 2000. Protein displays of the human T cell receptor alpha, beta, gamma and delta variable and joining regions. *Exp. Clin. Immunogenet.* In press.
- Forster, A., Huck, S., Ghanem, N., Lefranc, M.-P., and Rabbitts, T.H. 1987. New subgroups in the human T cell rearranging V gamma gene locus. *EMBO J.* 6:1945-1950.
- Ghanem, N., Buresi, C., Moisan, J.P., Bensmana, M., Chuchana, P., Huck, S., Lefranc, G., and Lefranc, M.-P. 1989. Deletion, insertion, and restriction site polymorphism of the T-cell receptor gamma variable locus in French, Lebanese, Tunisian, and black African populations. *Immunogenetics* 30:350-360.
- Ghanem, N., Soua, Z., Zhang, X.G., Zijun, M., Zhiwei, Y., Lefranc, G., and Lefranc, M.-P. 1991. Polymorphism of the T-cell receptor gamma variable and constant region genes in a Chinese population. *Hum. Genet.* 86:450-456.
- Giudicelli, V., Chaume, D., Bodmer, J., Müller, W., Busin, C., Marsh, S., Bontrop, R., Lemaitre, M., Malik, A., and Lefranc, M.-P. 1997. IMGT, the international ImMunoGeneTics database. *Nucl. Acids Res.* 25:206-211.
- Gottschalk, L.R. and Leiden, J.M. 1990. Identification and functional characterization of the human T-cell receptor beta gene transcriptional enhancer: Common nuclear proteins interact with the transcriptional regulatory elements of the T-cell receptor alpha and beta genes. *Mol. Cell. Biol.* 10:5486-5495.
- Ho, I.C., Yang, L.H., Morle, G., and Leiden, J.M. 1989. A T-cell-specific transcriptional enhancer element 3' of C alpha in the human T-cell receptor alpha locus. *Proc. Natl. Acad. Sci. U.S.A.* 86:6714-6718.
- Huck, S. and Lefranc, M.-P. 1987. Rearrangements to the JP1, JP and JP2 segments in the human T-cell rearranging gamma gene (TRG gamma) locus. *FEBS Lett.* 224:291-296.
- Huck, S., Dariavach, P., and Lefranc, M.-P. 1988. Variable region genes in the human T-cell rearranging gamma (TRG) locus: V-J junction and homology with the mouse genes. *EMBO J.* 7:719-726.
- Isobe, M., Russo, G., Haluska, F.G., and Croce, C.M. 1988. Cloning of the gene encoding the delta subunit of the human T-cell receptor reveals its physical organization within the alpha-subunit locus and its involvement in chromosome translocations in T-cell malignancy. *Proc. Natl. Acad. Sci. U.S.A.* 85:3933-3937.
- Koop, B.F., Rowen, L., Wang, K., Kuo, C.L., Seto, D., Lenstra, J.A., Howard, S., Shan, W., Deshpande, P., and Hood, L. 1994. The human T-cell receptor TCRAC/TCRDC (C alpha/C delta) region: Organization, sequence, and evolution of 97.6 kb of DNA. *Genomics* 19:478-493.
- Lefranc, M.-P. 1990a. The human T-cell receptor delta genes. *Res. Immunol.* 141:692-693.
- Lefranc, M.-P. 1990b. Organization of the human T-cell receptor genes. *European Cytokine Network* 1:121-130.
- Lefranc, M.-P. 1998. IMGT (ImMunoGeneTics) locus on focus. A new section of experimental and clinical immunogenetics. *Exp. Clin. Immunogenet.* 15:1-7.
- Lefranc, M.-P. and Alexandre, D. 1995. Gamma delta lineage-specific transcription of human T cell receptor gamma genes by a combination of a non-lineage-specific enhancer and silencers. *Eur. J. Immunol.* 25:617-622.
- Lefranc, M.-P. and Rabbitts, T.H. 1989. The human T-cell receptor gamma (TRG) genes. *Trends Biochem. Sci.* 14:214-218.
- Lefranc, M.-P. and Rabbitts, T.H. 1990a. A nomenclature to fit the organization of the human T cell receptor gamma and delta genes. *Res. Immunol.* 141:615-618.
- Lefranc, M.-P. and Rabbitts, T.H. 1990b. Genetic organization of the human T-cell receptor gamma and delta loci. *Res. Immunol.* 141:565-577.
- Lefranc, M.-P., Forster, A., Baer, R., Stinson, M.A., and Rabbitts, T.H. 1986a. Diversity and rearrangement of the human T cell rearranging gamma genes: Nine germ-line variable genes belonging to two subgroups. *Cell* 45:237-246.
- Lefranc, M.-P., Forster, A., and Rabbitts, T.H. 1986b. Genetic polymorphism and exon changes of the constant regions of the human T-cell rearranging gene gamma. *Proc. Natl. Acad. Sci. U.S.A.* 83:9596-9600.
- Lefranc, M.-P., Chuchana, P., Dariavach, P., Nguyen, C., Huck, S., Brockly, F., Jordan, B., and Lefranc, G. 1989. Molecular mapping of the

- human T cell receptor gamma (TRG) genes and linkage of the variable and constant regions. *Eur. J. Immunol.* 19:989-994.
- Loh, E.Y., Cwirla, S., Serafini, A.T., Phillips, J.H., and Lanier, L.L. 1988. Human T-cell-receptor delta chain: Genomic organization, diversity, and expression in populations of cells. *Proc. Natl. Acad. Sci. U.S.A.* 85:9714-9718.
- Posnett, D.N., Romagné, F., Necker, A., Kotzin, B.L., and Sekaly, R.-P. 1996. First human TcR monoclonal Antibody Workshop. *The Immunologist* 4:5-8.
- Redondo, J.M., Hata, S., Brocklehurst, C., and Krangel, M.S. 1990. A T cell-specific transcriptional enhancer within the human T cell receptor delta locus. *Science* 247:1225-1229.
- Rowen, L., Koop, B.F., and Hood, L. 1996. The complete 685-kilobase DNA sequence of the human beta T cell receptor locus. *Science* 272:1755-1762.
- Satyanarayana, K., Hata, S., Devlin, P., Roncarolo, M.G., De Vries, J.E., Spits, H., Strominger, J.L., and Krangel, M.S. 1988. Genomic organization of the human T-cell antigen-receptor alpha/delta locus. *Proc. Natl. Acad. Sci. U.S.A.* 85:8166-8170.
- Scaviner, D. and Lefranc, M.-P. 2000a. The human T cell receptor alpha variable (TRAV) genes. *Exp. Clin. Immunogenet.* 17:83-96.
- Scaviner, D. and Lefranc, M.-P. 2000b. The human T cell receptor alpha joining (TRAJ) genes. *Exp. Clin. Immunogenet.* 17:97-106.
- Slightom, J.L., Siemieniak, D.R., Sieu, L.C., Koop, B.F., and Hood, L. 1994. Nucleotide sequence analysis of 77.7 kb of the human V β T-cell receptor gene locus: Direct primer-walking using cosmid template DNAs. *Genomics* 20:149-168.
- Takahara, Y., Tkachuk, D., Michalopoulos, E., Champagne, E., Reimann, J., Minden, M., and Mak, T.W. 1988. Sequence and organization of the diversity, joining, and constant region genes of the human T-cell delta-chain locus. *Proc. Natl. Acad. Sci. U.S.A.* 85:6097-6101.
- Toyonaga, B., Yoshikai, Y., Vadasz, V., Chin, B., and Mak, T.W. 1985. Organization and sequences of the diversity, joining, and constant region genes of the human T-cell receptor beta chain. *Proc. Natl. Acad. Sci. U.S.A.* 82:8624-8628.
- Tunnacliffe, A., Kefford, R., Milstein, C., Forster, A., and Rabbitts, T.H. 1985. Sequence and evolution of the human T-cell antigen receptor beta-chain genes. *Proc. Natl. Acad. Sci. U.S.A.* 82:5068-5072.
- Wei, S., Charmley, P., Robinson, M.A., and Concannon, P. 1994. The extent of the human germline T-cell receptor V beta gene segment repertoire. *Immunogenetics* 40:27-36.
- Yoshikai, Y., Clark, S.P., Taylor, S., Sohn, U., Wilson, B.I., Minden, M.D., and Mak, T.W. 1985. Organization and sequences of the variable, joining and constant region genes of the human T-cell receptor alpha-chain. *Nature* 316:837-840.
- Zhang, X.M., Tonnel, C., Lefranc, M.-P., and Huck, S. 1994. T cell receptor gamma cDNA in human fetal liver and thymus: Variable regions of gamma chains are restricted to V gamma I or V9, due to the absence of splicing of the V10 and V11 leader intron. *Eur. J. Immunol.* 24:571-578.
- Zhang, X.M., Cathala, G., Soua, Z., Lefranc, M.-P., and Huck, S. 1996. The human T-cell receptor gamma variable pseudogene V10 is a distinctive marker of human speciation. *Immunogenetics* 43:196-203.

KEY REFERENCES

Giudicelli, V. and Lefranc, M.-P. 1999. Ontology for immunogenetics: IMGT-ONTOLOGY. *Bioinformatics* 15:1047-1054.

The immunoglobulin gene nomenclature is based on the CLASSIFICATION concept of the IMGT-ONTOLOGY.

Lefranc and Rabbitts, 1990a. See above.

This paper describes the rules for a standardized nomenclature of the T cell receptor genes.

Ruiz, M., Giudicelli, V., Ginestoux, C., Stoeck, P., Robinson, J., Bodmer, J., Marsh S.G.E., Bon-trop, R., Lemaitre, M., Lefranc, G., Chaume, D., and Lefranc, M.-P., 2000. IMGT, the international ImMunoGeneTics database. *Nucl. Acids Res.* 28:219-221.

The immunoglobulin gene nomenclature is part of the data standardization in the international ImMunoGeneTics database.

INTERNET RESOURCES

<http://www.gdb.org>

The Genome Database.

<http://www.gene.ucl.ac.uk/nomenclature>

HUGO Gene Nomenclature Committee web site.

<http://imgt.cines.fr:8104>

IMGT, the international ImMunoGeneTics database. IMGT/LIGM-DB contains the germline and rearranged sequences of the immunoglobulin and T cell receptor genes of human and other vertebrates.

<http://imgt.cines.fr:8104/textes/IMGTrepertoire.html>

IMGT Repertoire provides locus representations, germline gene tables, potential germline repertoires, correspondence between gene nomenclatures, protein displays, alignments of alleles, and tables of alleles for all the human TRA, TRB, TRG and TRD genes. 2D graphical representations designated as Colliers de Perles are provided for the variable genes.

<http://imgt.cines.fr:8104/textes/IMGTScientificChart.html>

IMGT Scientific chart describes the rules for a standardized immunoglobulin and T cell receptor gene and allele nomenclature.

<http://www.ncbi.nlm.nih.gov/LocusLink>

NCBI LocusLink.

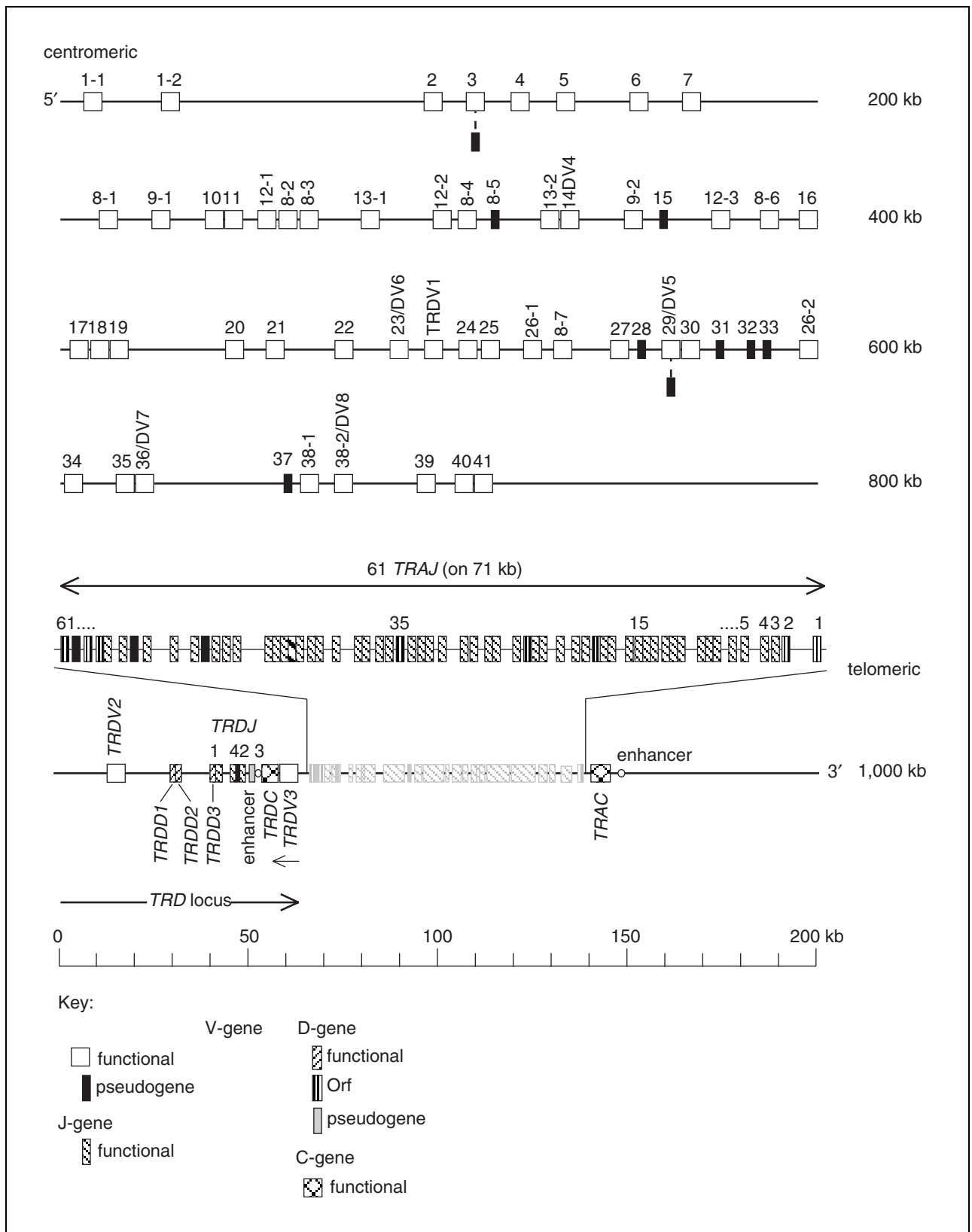


Figure A.10.1 Human *TRA/TRD* locus at 14q11.2. The boxes representing the genes are not to scale. The reader is referred to the following references for more information on *TRAV* genes (AE000658-AE000661): Boysen et al. (unpub. observ.); *TRAV* alignments: Folch et al. (2000), *TRAJ* genes: Koop et al. (1994); *TRAC* genes: Baer et al. (1985) and Yoshikai et al. (1985); *TRA* enhancer (4.5 kb 3' from *TRAC*): Ho et al. (1989); *TRDV* genes (AE000660, AE000661): Boysen et al. (unpub. observ.); *TRDD* genes: Loh et al. (1988) and Takihara et al. (1988); *TRDJ* genes: Isobe et al. (1988), Loh et al. (1988), and Satyanarayana et al. (1988); *TRDC* genes: Takihara et al. (1988); *TRD* enhancer (between *TRDJ3* and *TRDC*): Bories et al. (1990) and Redondo et al. (1990).

A.10.5

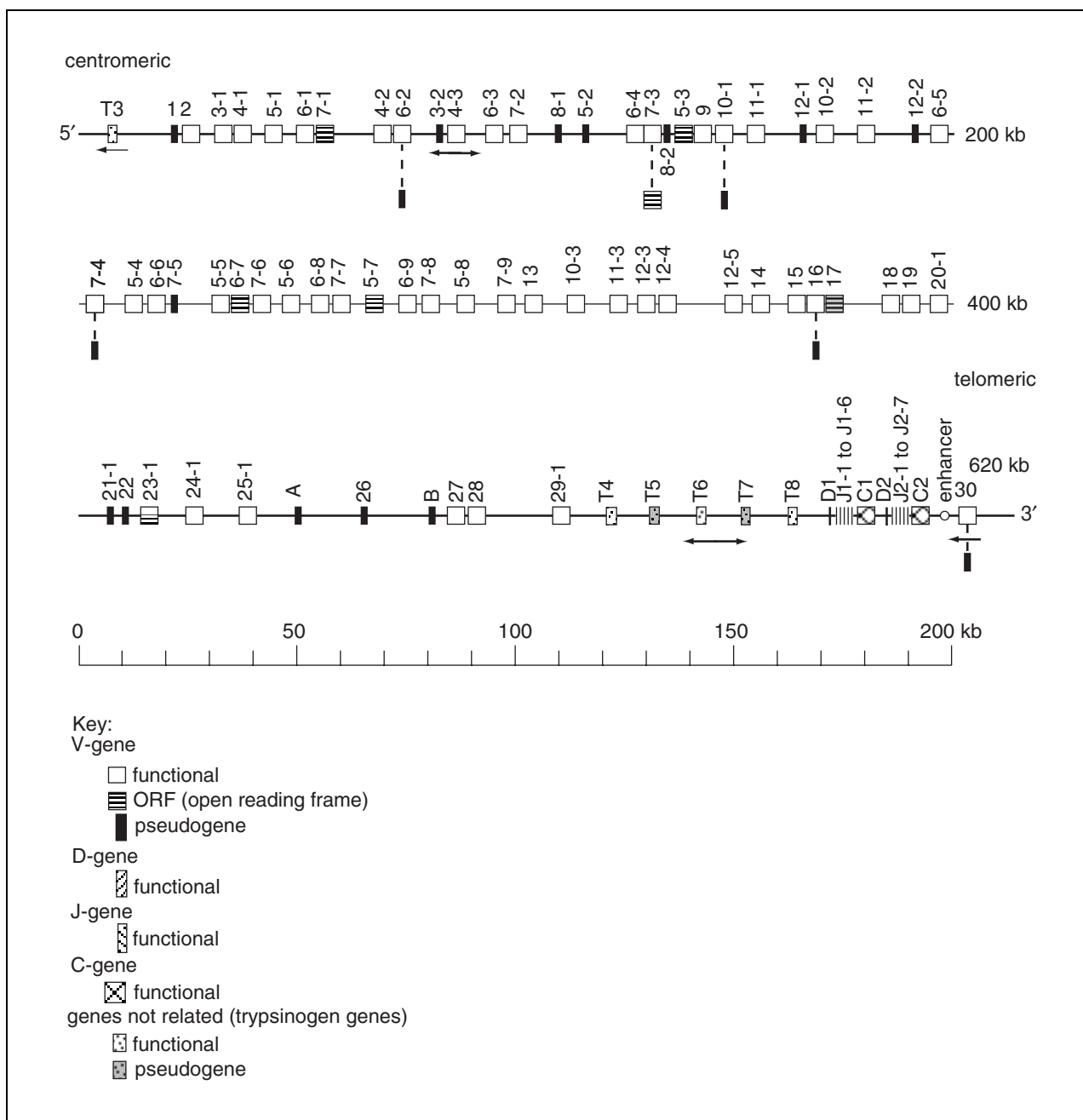


Figure A.10.2 Human *TRB* locus at 7q35. The boxes representing the genes are not to scale. Exons are not shown. *TRBV* gene designations according to Rowen et al. (1996). Nomenclature adopted by IMGT. Single arrows show genes whose polarity is opposite to that of the D-J-C-CLUSTER. Double arrows indicate insertion/deletion polymorphisms. *TRBV* genes: Wei et al. (1994), Posnett et al. (1996), and Rowen et al. (1996); *TRBD*-J-C-CLUSTER: Toyonaga et al. (1985) and Tunnacliffe et al. (1985); *TRBV* alignments: Folch et al. (2000); *TRBV* enhancer (5.5 kb 3' from TRBC2): Gottschalk and Leiden (1990).

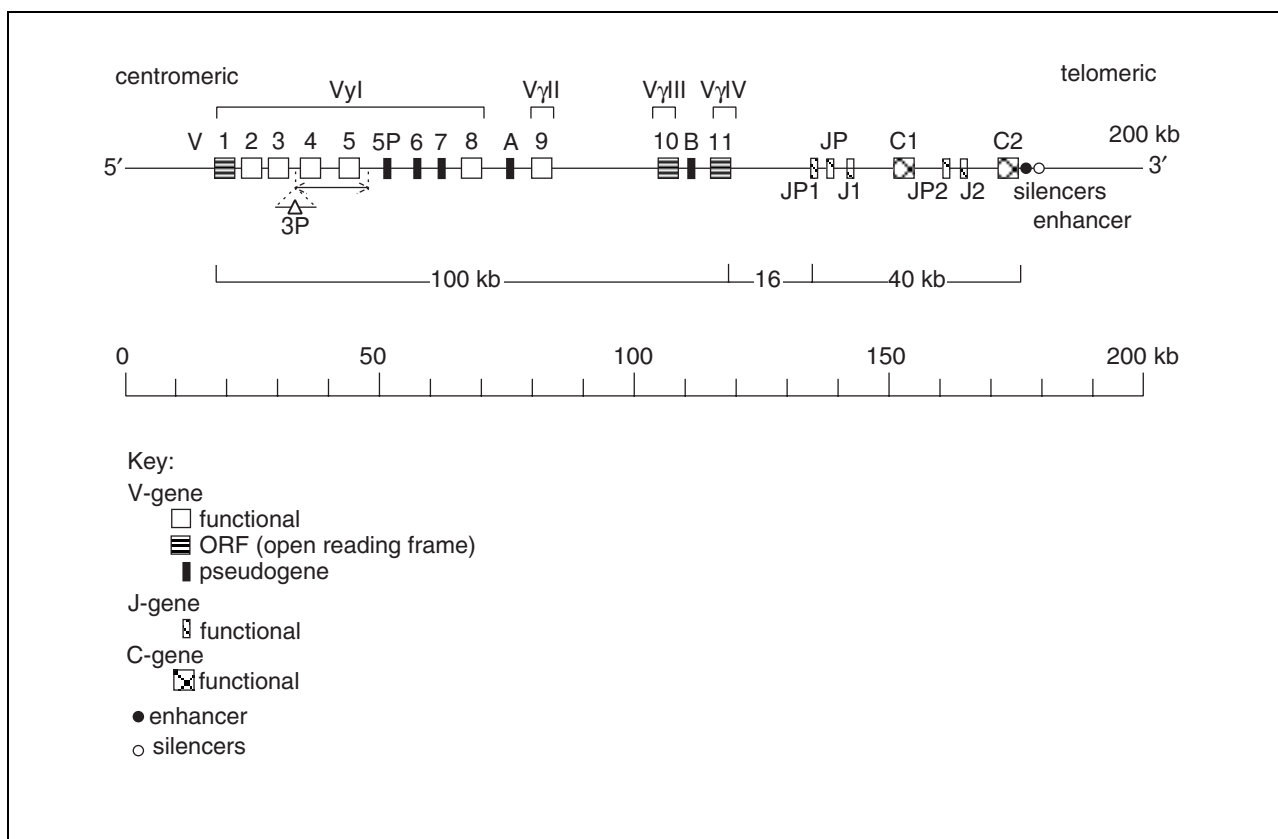


Figure A.10.3 Human *TRG* locus at 7p15-p14. The boxes representing the genes are not to scale. Exons are not shown. A double arrow indicates an insertion/deletion polymorphism. Lefranc and Rabbitts (1989, 1990a,b) and Lefranc et al. (1989); *TRGV* genes: Lefranc et al. (1986a), Forster et al. (1987), Huck et al. (1988), and Zhang et al. (1994, 1996); *TRGJ* genes: Huck and Lefranc (1987); *TRGC* genes: Lefranc et al. (1986b) and Buresi et al. (1989); Silencers and enhancers: Lefranc and Alexandre (1995); RFLP polymorphisms of the *TRGV* and *TRGC* genes: Buresi et al. (1989) and Ghanem et al. (1989,1991).

Table A.10.1 T Cell Receptor α (*TRA*) Genes^a

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
<i>TRA</i> locus on chromosome 14 at 14q11.2							
<i>TRAC</i>	<i>TRAC</i>	F	X02883	3	T cell receptor α constant	GDB:9953797	28755
<i>TRAJ</i> ^d	<i>TRAJ1</i>	ORF	X02884	1	T cell receptor α joining 1	GDB:9953799	28754
	<i>TRAJ2</i>	ORF	X02884	1	T cell receptor α joining 2	GDB:9953801	28753
	<i>TRAJ3</i>	F	X02884	1	T cell receptor α joining 3	GDB:9953803	28752
	<i>TRAJ4</i>	F	M94081	1	T cell receptor α joining 4	GDB:9953805	28751
	<i>TRAJ5</i>	F	M94081	1	T cell receptor α joining 5	GDB:9953807	28750
	<i>TRAJ6</i>	F	M16747	1	T cell receptor α joining 6	GDB:9953809	28749
	<i>TRAJ7</i>	F	M94081	1	T cell receptor α joining 7	GDB:9953811	28748
	<i>TRAJ8</i>	F	M94081	1	T cell receptor α joining 8	GDB:9953813	28747
	<i>TRAJ9</i>	F	M94081	1	T cell receptor α joining 9	GDB:9953815	28746
	<i>TRAJ10</i>	F	M94081	1	T cell receptor α joining 10	GDB:9953817	28745
	<i>TRAJ11</i>	F	M94081	1	T cell receptor α joining 11	GDB:9953819	28744
	<i>TRAJ12</i>	F	X02885	1	T cell receptor α joining 12	GDB:9953821	28743
	<i>TRAJ13</i>	F	M94081	1	T cell receptor α joining 13	GDB:9953823	28742
	<i>TRAJ14</i>	F	M94081	1	T cell receptor α joining 14	GDB:9953825	28741
	<i>TRAJ15</i>	F	X05775	2	T cell receptor α joining 15	GDB:9953827	28740
	<i>TRAJ16</i>	F	M94081	1	T cell receptor α joining 16	GDB:9953829	28739
	<i>TRAJ17</i>	F	X05773	1	T cell receptor α joining 17	GDB:9953831	28738
	<i>TRAJ18</i>	F	M94081	1	T cell receptor α joining 18	GDB:9953833	28737
	<i>TRAJ19</i>	ORF	M94081	1	T cell receptor α joining 19	GDB:9953835	28736
	<i>TRAJ20</i>	F	M94081	1	T cell receptor α joining 20	GDB:9953837	28735
	<i>TRAJ21</i>	F	M94081	1	T cell receptor α joining 21	GDB:9953839	28734
	<i>TRAJ22</i>	F	X02886	1	T cell receptor α joining 22	GDB:9953841	28733

*continued***A.10.8**

Table A.10.1 T Cell Receptor α (*TRA*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRAJ23</i>	F	M94081	1	T cell receptor α joining 23	GDB:9953843	28732
	<i>TRAJ24</i>	F	X02887	2	T cell receptor α joining 24	GDB:9953845	28731
	<i>TRAJ25</i>	ORF	X02888	1	T cell receptor α joining 25	GDB:9953847	28730
	<i>TRAJ26</i>	F	M94081	1	T cell receptor α joining 26	GDB:9953849	28729
	<i>TRAJ27</i>	F	M94081	1	T cell receptor α joining 27	GDB:9953851	28728
	<i>TRAJ28</i>	F	M94081	1	T cell receptor α joining 28	GDB:9953853	28727
	<i>TRAJ29</i>	F	X02889	1	T cell receptor α joining 29	GDB:9953855	28726
	<i>TRAJ30</i>	F	M94081	1	T cell receptor α joining 30	GDB:9953857	28725
	<i>TRAJ31</i>	F	M14905	1	T cell receptor α joining 31	GDB:9953859	28724
	<i>TRAJ32</i>	F	M94081	1	T cell receptor α joining 32	GDB:9953861	28723
	<i>TRAJ33</i>	F	M94081	1	T cell receptor α joining 33	GDB:9953863	28722
	<i>TRAJ34</i>	F	M35622	1	T cell receptor α joining 34	GDB:9953865	28721
	<i>TRAJ35</i>	ORF	M94081	1	T cell receptor α joining 35	GDB:9953867	28720
	<i>TRAJ36</i>	F	M94081	1	T cell receptor α joining 36	GDB:9953869	28719
	<i>TRAJ37</i>	F	M94081	1	T cell receptor α joining 37	GDB:9953871	28718
	<i>TRAJ38</i>	F	M94081	1	T cell receptor α joining 38	GDB:9953873	28717
	<i>TRAJ39</i>	F	M94081	1	T cell receptor α joining 39	GDB:9953875	28716
	<i>TRAJ40</i>	F	M35620	1	T cell receptor α joining 40	GDB:9953877	28715
	<i>TRAJ41</i>	F	M94081	1	T cell receptor α joining 41	GDB:9953879	28714
	<i>TRAJ42</i>	F	M94081	1	T cell receptor α joining 42	GDB:9953881	28713
	<i>TRAJ43</i>	F	M94081	1	T cell receptor α joining 43	GDB:9953883	28712
	<i>TRAJ44</i>	F	M35619	1	T cell receptor α joining 44	GDB:9953885	28711
	<i>TRAJ45</i>	F	M94081	1	T cell receptor α joining 45	GDB:9953887	28710

continued

Table A.10.1 T Cell Receptor α (*TRA*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRAJ46</i>	F	M94081	1	T cell receptor α joining 46	GDB:9953889	28709
	<i>TRAJ47</i>	F	M94081	1	T cell receptor α joining 47	GDB:9953891	28708
	<i>TRAJ48</i>	F	M94081	1	T cell receptor α joining 48	GDB:9953893	28707
	<i>TRAJ49</i>	F	M94081	1	T cell receptor α joining 49	GDB:9953895	28706
	<i>TRAJ50</i>	F	M94081	1	T cell receptor α joining 50	GDB:9953897	28705
	<i>TRAJ51</i>	P	M94081	—	T cell receptor α joining 51	GDB:9953899	28704
	<i>TRAJ52</i>	F	M94081	1	T cell receptor α joining 52	GDB:9953901	28703
	<i>TRAJ53</i>	F	M94081	1	T cell receptor α joining 53	GDB:9953903	28702
	<i>TRAJ54</i>	F	M94081	1	T cell receptor α joining 54	GDB:9953905	28701
	<i>TRAJ55</i>	P	M94081	—	T cell receptor α joining 55	GDB:9953907	28700
	<i>TRAJ56</i>	F	M94081	1	T cell receptor α joining 56	GDB:9953909	28699
	<i>TRAJ57</i>	F	M94081	1	T cell receptor α joining 57	GDB:9953911	28698
	<i>TRAJ58</i>	ORF	M94081	1	T cell receptor α joining 58	GDB:9953913	28697
	<i>TRAJ59</i>	ORF	M94081	1	T cell receptor α joining 59	GDB:9953915	28696
	<i>TRAJ60</i>	P	M94081	—	T cell receptor α joining 60	GDB:9953917	28695
	<i>TRAJ61</i>	ORF	M94081	1	T cell receptor α joining 61	GDB:9953919	28694
<i>TRAV^e</i>	<i>TRAV1-1</i>	F	AE000658	2	T cell receptor α variable 1-1	GDB:9953921	28693
	<i>TRAV1-2</i>	F	AE000658	2	T cell receptor α variable 1-2	GDB:9953923	28692
	<i>TRAV2</i>	F	AE000658	2	T cell receptor α variable 2	GDB:9953925	28691
	<i>TRAV3</i>	F, (P) ^f	AE000658	2	T cell receptor α variable 3	GDB:9953927	28690
	<i>TRAV4</i>	F	AE000658	1	T cell receptor α variable 4	GDB:9953929	28689
	<i>TRAV5</i>	F	AE000659	1	T cell receptor α variable 5	GDB:9953931	28688
	<i>TRAV6</i>	F	AE000659	6	T cell receptor α variable 6	GDB:9953933	28687

continued

**Nomenclature of
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Table A.10.1 T Cell Receptor α (*TRA*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRAV7</i>	F	AE000659	1	T cell receptor α variable 7	GDB:9953935	28686
	<i>TRAV8-1</i>	F	AE000659	2	T cell receptor α variable 8-1	GDB:9953937	28685
	<i>TRAV8-2</i>	F	AE000659	2	T cell receptor α variable 8-2	GDB:9953939	28684
	<i>TRAV8-3</i>	F	AE000659	3	T cell receptor α variable 8-3	GDB:9953941	28683
	<i>TRAV8-4</i>	F	AE000659	7	T cell receptor α variable 8-4	GDB:9953943	28682
	<i>TRAV8-5</i>	P	AE000659	—	T cell receptor α variable 8-5	GDB:9953945	28681
	<i>TRAV8-6</i>	F	X02850	2	T cell receptor α variable 8-6	GDB:9953947	28680
	<i>TRAV8-7</i>	F	AE000660	1	T cell receptor α variable 8-7	GDB:9953949	28679
	<i>TRAV9-1</i>	F	AE000659	1	T cell receptor α variable 9-1	GDB:9953951	28678
	<i>TRAV9-2</i>	F	AE000659	4	T cell receptor α variable 9-2	GDB:9953953	28677
	<i>TRAV10</i>	F	AE000659	1	T cell receptor α variable 10	GDB:9953955	28676
	<i>TRAV11</i>	F	AE000659	1	T cell receptor α variable 11	GDB:9953957	28675
	<i>TRAV12-1</i>	F	AE000659	2	T cell receptor α variable 12-1	GDB:9953959	28674
	<i>TRAV12-2</i>	F	AE000659	3	T cell receptor α variable 12-2	GDB:9953961	28673
	<i>TRAV12-3</i>	F	X06193	2	T cell receptor α variable 12-3	GDB:9953963	28672
	<i>TRAV13-1</i>	F	AE000659	3	T cell receptor α variable 13-1	GDB:9953965	28671
	<i>TRAV13-2</i>	F	AE000659	2	T cell receptor α variable 13-2	GDB:9953967	28670
	<i>TRAV14/DV4^g</i>	F	M21626	4	T cell receptor α variable 14/8 variable 4	GDB:9953969	28669
	<i>TRAV15</i>	P	AE000659	—	T cell receptor α variable 15	GDB:9953971	28668
	<i>TRAV16</i>	F	AE000659	1	T cell receptor α variable 16	GDB:9953973	28667
	<i>TRAV17</i>	F	AE000660	1	T cell receptor α variable 17	GDB:9953975	28666
	<i>TRAV18</i>	F	AE000660	1	T cell receptor α variable 18	GDB:9953977	28665

*continued***Abbreviations
and Useful Data****A.10.11**

Table A.10.1 T Cell Receptor α (*TRA*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRAV19</i>	F	AE000660	1	T cell receptor α variable 19	GDB:9953979	28664
	<i>TRAV20</i>	F	AE000660	4	T cell receptor α variable 20	GDB:9953981	28663
	<i>TRAV21</i>	F	AE000660	2	T cell receptor α variable 21	GDB:9953983	28662
	<i>TRAV22</i>	F	AE000660	1	T cell receptor α variable 22	GDB:9953985	28661
	<i>TRAV23/DV6^g</i>	F	AE000660	4	T cell receptor α variable 23/ δ variable 6	GDB:9953987	28660
	<i>TRAV24</i>	F	AE000660	2	T cell receptor α variable 24	GDB:9953989	28659
	<i>TRAV25</i>	F	AE000660	1	T cell receptor α variable 25	GDB:9953991	28658
	<i>TRAV26-1</i>	F	AE000660	3	T cell receptor α variable 26-1	GDB:9953993	28657
	<i>TRAV26-2</i>	F	AE000660	2	T cell receptor α variable 26-2	GDB:9953995	28656
	<i>TRAV27</i>	F	AE000660	3	T cell receptor α variable 27	GDB:9953997	28655
	<i>TRAV28</i>	P	AE000660	1	T cell receptor α variable 28	GDB:9953999	28654
	<i>TRAV29/DV5^g</i>	F, (P) ^f	AE000660	3	T cell receptor α variable 29/ δ variable 5	GDB:9954001	28653
	<i>TRAV30</i>	F	AE000660	4	T cell receptor α variable 30	GDB:9954003	28652
	<i>TRAV31</i>	P	AE000660	—	T cell receptor α variable 31	GDB:9954005	28651
	<i>TRAV32</i>	P	AE000660	—	T cell receptor α variable 32	GDB:9954007	28650
	<i>TRAV33</i>	P	AE000660	—	T cell receptor α variable 33	GDB:9954009	28649
	<i>TRAV34</i>	F	AE000660	1	T cell receptor α variable 34	GDB:9954011	28648
	<i>TRAV35</i>	F	AE000660	2	T cell receptor α variable 35	GDB:9954013	28647
	<i>TRAV36/DV7^g</i>	F	AE000660	4	T cell receptor α variable 36/ δ variable 7	GDB:9954015	28646
	<i>TRAV37</i>	P	AE000661	—	T cell receptor α variable 37	GDB:9954017	28645
	<i>TRAV38-1</i>	F	AE000661	4	T cell receptor α variable 38-1	GDB:9954019	28644

continued

Nomenclature of the Human T Cell Receptor Genes

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Table A.10.1 T Cell Receptor α (*TRA*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRAV38-2/DV8</i> ^g	F	AE000661	1	T cell receptor α variable 38-2/ δ variable 8	GDB:9954021	28643
	<i>TRAV39</i>	F	AE000661	1	T cell receptor α variable 39	GDB:9954023	28642
	<i>TRAV40</i>	F	X73521	1	T cell receptor α variable 40	GDB:9954025	28641
	<i>TRAV41</i>	F	AE000661	1	T cell receptor α variable 41	GDB:9954027	28640

^aSee Table Guide, above, for explanations of abbreviations and nomenclature; also see discussion of *TRA* Locus, above, Folch et al. (2000), and Scaviner and Lefranc (2000a,b) for more information.

^bIMGT *TRA* gene names have been approved by the HUGO Nomenclature Committee in 1999. Note that, in the HUGO symbols, slashes of the *TRAV/DV* gene names are omitted. Otherwise all the gene names (gene symbols) are identical in IMGT and HUGO nomenclatures.

^cGene definitions (full names) are identical (including slashes) in IMGT and HUGO nomenclatures. Note that in the databases, the Greek letters are written in full (e.g., α = alpha, δ = delta).

^d*TRAJ* genes are designated by a number for the localization from 3' to 5' in the locus.

^e*TRAV* genes are designated by a number for the subgroup followed, whenever there are several genes belonging to the same subgroup, by a hyphen and a number for their relative localization in the locus. Numbers increase from 5' to 3' in the locus.

^fFunctionality is shown between parentheses when the germline *TRAV* genes have not yet been isolated.

^gThe *TRAV14/DV4*, *TRAV23/DV6*, *TRAV29/DV5*, *TRAV36/DV7*, and *TRAV38-2/DV8* genes have been found rearranged to J genes of the *TRA* locus, and to D and J genes of the *TRD* locus.

Table A.10.2 T Cell Receptor β (*TRB*) Genes^a

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
<i>TRB</i> locus on chromosome 7 at 7q35							
<i>TRBC</i>	<i>TRBC1</i>	F	M12887	2	T cell receptor β constant 1	GDB:9954029	28639
	<i>TRBC2</i>	F	M12888	2	T cell receptor β constant 2	GDB:9954031	28638
<i>TRBD</i>	<i>TRBD1</i>	F	X00936	1	T cell receptor β diversity 1	GDB:9954033	28637
	<i>TRBD2</i>	F	X02987	2	T cell receptor β diversity 2	GDB:9954035	28636
<i>TRBJ^d</i>	<i>TRBJ1-1</i>	F	X00936	1	T cell receptor β joining 1-1	GDB:9954037	28635
	<i>TRBJ1-2</i>	F	X00936	1	T cell receptor β joining 1-2	GDB:9954039	28634
	<i>TRBJ1-3</i>	F	M14158	1	T cell receptor β joining 1-3	GDB:9954041	28633
	<i>TRBJ1-4</i>	F	M14158	1	T cell receptor β joining 1-4	GDB:9954043	28632
	<i>TRBJ1-5</i>	F	M14158	1	T cell receptor β joining 1-5	GDB:9954045	28631
	<i>TRBJ1-6</i>	F	M14158	1	T cell receptor β joining 1-6	GDB:9954047	28630
	<i>TRBJ2-1</i>	F	X02987	1	T cell receptor β joining 2-1	GDB:9954049	28629
	<i>TRBJ2-2</i>	F	X02987	1	T cell receptor β joining 2-2	GDB:9954051	28628
	<i>TRBJ2-2P</i>	ORF	X02987	1	T cell receptor β joining 2-2P	GDB:9954053	28627
	<i>TRBJ2-3</i>	F	X02987	1	T cell receptor β joining 2-3	GDB:9954055	28626
	<i>TRBJ2-4</i>	F	X02987	1	T cell receptor β joining 2-4	GDB:9954057	28625
	<i>TRBJ2-5</i>	F	X02987	1	T cell receptor β joining 2-5	GDB:9954059	28624
	<i>TRBJ2-6</i>	F	X02987	1	T cell receptor β joining 2-6	GDB:9954061	28623
	<i>TRBJ2-7</i>	F, ORF	M14159	2	T cell receptor β joining 2-7	GDB:9954063	28622
	<i>TRBV^e</i>						
	<i>TRBV1</i>	P	L36092	—	T cell receptor β variable 1	GDB:9954065	28621
	<i>TRBV2</i>	F	L36092	3	T cell receptor β variable 2	GDB:9954067	28620
	<i>TRBV3-1</i>	F	U07977	2	T cell receptor β variable 3-1	GDB:9954069	28619
	<i>TRBV3-2</i>	P	L36092	—	T cell receptor β variable 3-2	GDB:9954071	28618
	<i>TRBV4-1</i>	F	U07977	2	T cell receptor β variable 4-1	GDB:9954073	28617

continued

**Nomenclature of
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A.10.14

Table A.10.2 T Cell Receptor β (*TRB*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRBV4-2</i>	F	U07975	2	T cell receptor β variable 4-2	GDB:9954075	28616
	<i>TRBV4-3</i>	F	U07978	4	T cell receptor β variable 4-3	GDB:9954077	28615
	<i>TRBV5-1</i>	F	L36092	2	T cell receptor β variable 5-1	GDB:9954079	28614
	<i>TRBV5-2</i>	P	L36092	—	T cell receptor β variable 5-2	GDB:9954081	28613
	<i>TRBV5-3</i>	ORF	X61439	2	T cell receptor β variable 5-3	GDB:9954083	28612
	<i>TRBV5-4</i>	F	L36092	4	T cell receptor β variable 5-4	GDB:9954085	28611
	<i>TRBV5-5</i>	F	L36092	3	T cell receptor β variable 5-5	GDB:9954087	28610
	<i>TRBV5-6</i>	F	L36092	1	T cell receptor β variable 5-6	GDB:9954089	28609
	<i>TRBV5-7</i>	ORF	L36092	1	T cell receptor β variable 5-7	GDB:9954091	28608
	<i>TRBV5-8</i>	F	L36092	2	T cell receptor β variable 5-8	GDB:9954093	28607
	<i>TRBV6-1</i>	F	X61446	1	T cell receptor β variable 6-1	GDB:9954095	28606
	<i>TRBV6-2</i>	F, (P) ^f	X61445	3	T cell receptor β variable 6-2	GDB:9954097	28605
	<i>TRBV6-3</i>	F	U07978	1	T cell receptor β variable 6-3	GDB:9954099	28604
	<i>TRBV6-4</i>	F	X61653	2	T cell receptor β variable 6-4	GDB:9954101	28603
	<i>TRBV6-5</i>	F	L36092	1	T cell receptor β variable 6-5	GDB:9954103	28602
	<i>TRBV6-6</i>	F	L36092	5	T cell receptor β variable 6-6	GDB:9954105	28601
	<i>TRBV6-7</i>	ORF	L36092	1	T cell receptor β variable 6-7	GDB:9954107	28600
	<i>TRBV6-8</i>	F	L36092	1	T cell receptor β variable 6-8	GDB:9954109	28599
	<i>TRBV6-9</i>	F	X61447	1	T cell receptor β variable 6-9	GDB:9954111	28598
	<i>TRBV7-1</i>	ORF	X61444	1	T cell receptor β variable 7-1	GDB:9954113	28597
	<i>TRBV7-2</i>	F	X61442	4	T cell receptor β variable 7-2	GDB:9954115	28596
	<i>TRBV7-3</i>	F, ORF	X61440	5	T cell receptor β variable 7-3	GDB:9954117	28595
	<i>TRBV7-4</i>	F, (P) ^f	L36092	3	T cell receptor β variable 7-4	GDB:9954119	28594

continued

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Table A.10.2 T Cell Receptor β (*TRB*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRBV7-5</i>	P	L36092	—	T cell receptor β variable 7-5	GDB:9954121	28593
	<i>TRBV7-6</i>	F	L36092	2	T cell receptor β variable 7-6	GDB:9954123	28592
	<i>TRBV7-7</i>	F	L36092	2	T cell receptor β variable 7-7	GDB:9954125	28591
	<i>TRBV7-8</i>	F	M11953	3	T cell receptor β variable 7-8	GDB:9954127	28590
	<i>TRBV7-9</i>	F	L36092	7	T cell receptor β variable 7-9	GDB:9954129	28589
	<i>TRBV8-1</i>	P	L36092	—	T cell receptor β variable 8-1	GDB:9954131	28588
	<i>TRBV8-2</i>	P	L36092	—	T cell receptor β variable 8-2	GDB:9954133	28587
	<i>TRBV9</i>	F	L36092	3	T cell receptor β variable 9	GDB:9954135	28586
	<i>TRBV10-1</i>	F, (P) ^f	U17050	3	T cell receptor β variable 10-1	GDB:9954137	28585
	<i>TRBV10-2</i>	F	U17049	2	T cell receptor β variable 10-2	GDB:9954139	28584
	<i>TRBV10-3</i>	F	U03115	4	T cell receptor β variable 10-3	GDB:9954141	28583
	<i>TRBV11-1</i>	F	M33233	1	T cell receptor β variable 11-1	GDB:9954143	28582
	<i>TRBV11-2</i>	F	L36092	3	T cell receptor β variable 11-2	GDB:9954145	28581
	<i>TRBV11-3</i>	F	M33234	4	T cell receptor β variable 11-3	GDB:9954147	28580
	<i>TRBV12-1</i>	P	X07224	—	T cell receptor β variable 12-1	GDB:9954149	28579
	<i>TRBV12-2</i>	P	X06936	—	T cell receptor β variable 12-2	GDB:9954151	28578
	<i>TRBV12-3</i>	F	X07192	1	T cell receptor β variable 12-3	GDB:9954153	28577
	<i>TRBV12-4</i>	F	K02546	2	T cell receptor β variable 12-4	GDB:9954155	28576
	<i>TRBV12-5</i>	F	X07223	1	T cell receptor β variable 12-5	GDB:9954157	28575
	<i>TRBV13</i>	F	U03115	2	T cell receptor β variable 13	GDB:9954159	28574
	<i>TRBV14</i>	F	X06154	2	T cell receptor β variable 14	GDB:9954161	28573
	<i>TRBV15</i>	F	U03115	3	T cell receptor β variable 15	GDB:9954163	28572
	<i>TRBV16</i>	F, P	L26231	3	T cell receptor β variable 16	GDB:9954165	28571

continued

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Table A.10.2 T Cell Receptor β (*TRB*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRBV17</i>	ORF	U03115	1	T cell receptor β variable 17	GDB:9954167	28570
	<i>TRBV18</i>	F	L36092	1	T cell receptor β variable 18	GDB:9954169	28569
	<i>TRBV19</i>	F	U48260	3	T cell receptor β variable 19	GDB:9954171	28568
	<i>TRBV20-1^g</i>	F	M11955	7	T cell receptor β variable 20-1	GDB:9954173	28567
	<i>TRBV21-1^g</i>	P	L36092	—	T cell receptor β variable 21-1	GDB:9954175	28566
	<i>TRBV22^g</i>	P	L36092	—	T cell receptor β variable 22	GDB:9954177	28565
	<i>TRBV23-1^g</i>	ORF	L36092	1	T cell receptor β variable 23-1	GDB:9954179	28564
	<i>TRBV24-1^g</i>	F	M11951	1	T cell receptor β variable 24-1	GDB:9954181	28563
	<i>TRBV25-1^g</i>	F	L36092	1	T cell receptor β variable 25-1	GDB:9954183	28562
	<i>TRBV26</i>	P	L36092	—	T cell receptor β variable 26	GDB:9954185	28561
	<i>TRBV27</i>	F	L36092	1	T cell receptor β variable 27	GDB:9954187	28560
	<i>TRBV28</i>	F	U08314	1	T cell receptor β variable 28	GDB:9954189	28559
	<i>TRBV29-1^g</i>	F	L36092	3	T cell receptor β variable 29-1	GDB:9954191	28558
	<i>TRBV30</i>	F, P	L36092	5	T cell receptor β variable 30	GDB:9954193	28557
	<i>TRBVA</i>	P	L36092	—	T cell receptor β variable A	GDB:9954195	28556
	<i>TRBVB</i>	P	L36092	—	T cell receptor β variable B	GDB:9954197	28555
<i>TRBV orphans on chromosome 9 at 9p21</i>							
<i>TRBV</i>	<i>TRBV20/OR9-2</i>	ORF	L05149	2	T cell receptor β variable 20/OR9-2	GDB:9954199	6962
	<i>TRBV21/OR9-2</i>	ORF	L05151	1	T cell receptor β variable 21/OR9-2	GDB:9954201	6959
	<i>TRBV23/OR9-2</i>	ORF	L27615	1	T cell receptor β variable 23/OR9-2	GDB:9954203	28552
	<i>TRBV24/OR9-2</i>	ORF, P	L05153	2	T cell receptor β variable 24/OR9-2	GDB:9954205	6961

continued

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Table A.10.2 T Cell Receptor β (*TRB*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
	<i>TRBV25/OR9-2</i>	P	L05152	2	T cell receptor β variable 25/OR9-2	GDB:9954207	6960
	<i>TRBV29/OR9-2</i>	ORF	L05150	2	T cell receptor β variable 29/OR9-2	GDB:9954209	6958

^aSee Table Guide, above, for explanations of abbreviations and nomenclature; also see discussion of *TRB* Locus, above, Folch and Lefranc (2000a,b), and Folch et al. (2000) for more information.

^bIMGT *TRB* gene names have been approved by the HUGO Nomenclature Committee in 1999. Note that, in the HUGO symbols, slashes of the orphon names are omitted. Otherwise all the gene names (gene symbols) are identical in IMGT and HUGO nomenclatures.

^cGene definitions (full names) are identical (including slashes) in IMGT and HUGO nomenclatures. Note that in the databases, the Greek letters are written in full (β = beta).

^d*TRBJ* genes are designated by a number for the cluster followed by a hyphen and a number for their relative localization in the locus. Numbers increase from 5' to 3' in the locus.

^e*TRBV* genes are designated by a number for the subgroup followed, whenever there are several genes belonging to the same subgroup, by a hyphen and a number for their relative localization in the locus. Numbers increase from 5' to 3' in the locus.

^fFunctionality is shown between parentheses when the accession number refers to a rearranged sequence and the corresponding germline gene has not yet been isolated; brackets when the accession number refers to a DNA genomic sequence, but not known as being germline or rearranged.

^gSince orphans (OR) have been described for each of the following *TRBV* subgroups: 20, 21, 23, 24, 25, and 29 (see *TRBV* orphans), the single member gene in the main locus is designated by the subgroup number followed by a hyphen and the number 1. To date, no orphon has been reported which belongs to subgroup 22; therefore, the IMGT designation of the single member gene is *TRBV22*.

Table A.10.3 T Cell Receptor γ (*TRG*) Genes^a

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
<i>TRG</i> locus on chromosome 7 at 7p15-p14							
<i>TRGC</i>	<i>TRGC1</i>	F	M14996,97,98	2	T cell receptor γ constant 1	GDB:120408	6966
	<i>TRGC2</i> (2 \times)	F	M15002/M13231	3	T cell receptor γ constant 2 (2 \times)	GDB:120409	6967
	<i>TRGC2</i> (3 \times)	F	M17323/M25318	1	T cell receptor γ constant 2 (3 \times)	GDB:120409	6967
<i>TRGJ</i>	<i>TRGJ1</i>	F	M12960	2	T cell receptor γ joining 1	GDB:120410	6968
	<i>TRGJ2</i>	F	M12961	1	T cell receptor γ joining 2	GDB:120411	6969
	<i>TRGJP</i>	F	M12950	1	T cell receptor γ joining P	GDB:120412	6970
	<i>TRGJP1</i>	F	X08084	1	T cell receptor γ joining P1	GDB:120413	6971
	<i>TRGJP2</i>	F	M16016	1	T cell receptor γ joining P2	GDB:120414	6972

continued

Table A.10.3 T Cell Receptor γ (*TRG*) Genes^a, continued

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession number	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
<i>TRGV</i> ^{d,e}	<i>TRGV1</i>	ORF	M12949	1	T cell receptor γ variable 1	GDB:120415	6973
	<i>TRGV2</i>	F	M13429	1	T cell receptor γ variable 2	GDB:120418	6974
	<i>TRGV3</i>	F	M13430	1	T cell receptor γ variable 3	GDB:120419	6976
	<i>TRGV4</i>	F	X15272	2	T cell receptor γ variable 4	GDB:120420	6977
	<i>TRGV5</i>	F	X13555	1	T cell receptor γ variable 5	GDB:120421	6978
	<i>TRGV5P</i>	P	M13431	—	T cell receptor γ variable 5P	GDB:120422	6979
	<i>TRGV6</i>	P	M13432	—	T cell receptor γ variable 6	GDB:120423	6980
	<i>TRGV7</i>	P	M13433	—	T cell receptor γ variable 7	GDB:120424	6981
	<i>TRGV8</i>	F	M13434	1	T cell receptor γ variable 8	GDB:120425	6982
	<i>TRGV9</i>	F	X07205	2	T cell receptor γ variable 9	GDB:120426	6983
	<i>TRGV10</i>	ORF	X07206	2	T cell receptor γ variable 10	GDB:120416	6984
	<i>TRGV11</i>	ORF	Y11227	1	T cell receptor γ variable 11	GDB:120417	6985
	<i>TRGVA</i>	P	X07208	—	T cell receptor γ variable A	GDB:9953127	6986
	<i>TRGVB</i>	P	X07209	—	T cell receptor γ variable B	GDB:9953128	6987

^aSee Table Guide, above, for explanations of abbreviations and nomenclature; also see discussion of *TRG* Locus, above, Lefranc et al. (1989), Lefranc and Rabbits (1989, 1990a,b), and Folch et al. (2000) for more information.

^bIMGT *TRG* gene names have been approved by the HUGO Nomenclature Committee in 1999. All the gene names (gene symbols) are identical in IMGT and HUGO nomenclatures.

^cGene definitions (full names) are identical in IMGT and HUGO nomenclatures. Note that in the databases, the Greek letters are written in full (e.g., γ = gamma).

^d*TRGV* genes are designated by a number (or a letter, for pseudogenes that are single members of their subgroup) for their position from 5' to 3' in the locus (Lefranc et al., 1989; Lefranc and Rabbits, 1989, 1990a,b).

^eThe *IGHV3P* gene, a polymorphic gene by insertion, has been identified by Southern hybridization in a rare haplotype but has not been sequenced (Ghanem et al., 1991).

Abbreviations and Useful Data

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Table A.10.4 T cell Receptor δ (*TRD*) Genes^a

IMGT gene group	IMGT gene name ^b	Functionality	IMGT reference sequence accession numbers	Number of alleles	IMGT gene definition ^c	GDB accession ID	LocusLink number
<i>TRD</i> locus on chromosome 14 at 14q11.2							
<i>TRDC</i>	<i>TRDC</i>	F	M22148-M22151	1	T cell receptor δ constant	GDB:9954211	28526
<i>TRDD</i>	<i>TRDD1</i>	F	M23325	1	T cell receptor δ diversity 1	GDB:9954213	28525
	<i>TRDD2</i>	F	M22153	1	T cell receptor δ diversity 2	GDB:9954215	28524
	<i>TRDD3</i>	F	M22152	1	T cell receptor δ diversity 3	GDB:9954217	28523
<i>TRDJ</i>	<i>TRDJ1</i>	F	M20289	1	T cell receptor δ joining 1	GDB:9954219	28522
	<i>TRDJ2</i>	F	L36386	1	T cell receptor δ joining 2	GDB:9954221	28521
	<i>TRDJ3</i>	F	M21508	1	T cell receptor δ joining 3	GDB:9954223	28520
	<i>TRDJ4</i>	F	AJ249814	1	T cell receptor δ joining 4	GDB:9953677	28519
<i>TRDV^d</i>	<i>TRDV1</i>	F	M22198	1	T cell receptor δ variable 1	GDB:9953671	28518
	<i>TRDV2</i>	F	X15207	2	T cell receptor δ variable 2	GDB:9953287	28517
	<i>TRDV3</i>	F	M23326	2	T cell receptor δ variable 3	GDB:9953273	28516

^aSee Table Guide, above, for explanations of abbreviations and nomenclature; also see discussion of *TRD* Locus, above, Lefranc (1990a), and Folch et al. (2000) for more information.

^bIMGT *TRD* gene names have been approved by the HUGO Nomenclature Committee in 1999. All the gene names (gene symbols) are identical in IMGT and HUGO nomenclatures.

^cGene definitions (full names) are identical in IMGT and HUGO nomenclatures. Note that in the databases, the Greek letters are written in full (e.g., δ = delta).

^d*TRDV* genes are designated by a number for their position from 5' to 3' in the locus. The *TRAV14/DV4*, *TRAV23/DV6*, *TRAV29/DV5*, *TRAV36/DV7*, and *TRAV38-2/DV8* genes, which have been found rearranged to J genes of the *TRA* locus, and to D and J genes of the *TRD* locus, are displayed in the human *TRAV* table (Table A.10.1).

Table A.10.5 Correspondence Between *TRAV* Nomenclatures^{a,b}

IMGT <i>TRAV</i> gene name (Scaviner and Lefranc, 2000a)	Boysen et al. (unpub. observ.) ^c	Arden et al. (1995)
<i>TRAV41</i>	41S1	19S1
<i>TRAV40</i>	40S1	31S1
<i>TRAV39</i>	39S1	27S1
<i>TRAV38-2/DV8</i>	hADV38S2	14S1-ADV14S1
<i>TRAV38-1</i>	38S1	14S2
<i>TRAV37</i>	37S1	—
<i>TRAV36/DV7</i>	hADV36S1	28S1-DV28S1
<i>TRAV35</i>	35S1	25S1
<i>TRAV34</i>	34S1	26S1

continued

Table A.10.5 Correspondence Between *TRAV* Nomenclatures^{a,b}, continued

IMGT <i>TRAV</i> gene name (Scaviner and Lefranc, 2000a)	Boysen et al. (unpub. observ.) ^c	Arden et al. (1995)
<i>TRAV26-2</i>	26S2	4S1
<i>TRAV33</i>	33S1	—
<i>TRAV32</i>	32S1	—
<i>TRAV31</i>	31S1	—
<i>TRAV30</i>	30S1	29S1
<i>TRAV29/DV5</i>	hADV29S1	21S1-ADV21S1
<i>TRAV28</i>	28S1	—
<i>TRAV27</i>	27S1	10S1
<i>TRAV8-7</i>	8S7	—
<i>TRAV26-1</i>	26S1	4S2
<i>TRAV25</i>	25S1	32S1
<i>TRAV24</i>	24S1	18S1
<i>TRAV23/DV6</i>	hADV23S1	17S1-ADV17S1
<i>TRAV22</i>	22S1	13S1
<i>TRAV21</i>	21S1	23S1
<i>TRAV20</i>	20S1	30S1
<i>TRAV19</i>	19S1	12S1
<i>TRAV18</i>	18S1	—
<i>TRAV17</i>	17S1	3S1
<i>TRAV16</i>	16S1	9S1
<i>TRAV8-6</i>	8S6	1S3
<i>TRAV12-3</i>	12S3	2S2
<i>TRAV15</i>	15S1	—
<i>TRAV9-2</i>	9S2	22S1
<i>TRAV14/DV4</i>	hADV14S1	6S1-ADV6S1
<i>TRAV13-2</i>	13S2	8S2
<i>TRAV8-5</i>	8S5	—
<i>TRAV8-4</i>	8S4	1S2
<i>TRAV12-2</i>	12S2	2S1
<i>TRAV13-1</i>	13S1	8S1
<i>TRAV8-3</i>	8S3	1S4
<i>TRAV8-2</i>	8S2	1S5
<i>TRAV12-1</i>	12S1	2S3
<i>TRAV11</i>	11S1	—
<i>TRAV10</i>	10S1	24S1
<i>TRAV9-1</i>	9S1	—
<i>TRAV8-1</i>	8S1	1S1
<i>TRAV7</i>	7S1	—
<i>TRAV6</i>	6S1	5S1
<i>TRAV5</i>	5S1	15S1
<i>TRAV4</i>	4S1	20S1
<i>TRAV3</i>	3S1	16S1
<i>TRAV2</i>	2S1	11S1
<i>TRAV1-2</i>	1S2	7S2
<i>TRAV1-1</i>	1S1	7S1

^a*TRAV* genes are listed from 3' (top of the table) to 5' (bottom of the table). Cells with dashes indicate that no name exists for the gene in that system of nomenclature.

^bSee *TRA* Locus and Table A.10.1 for more information.

^cIMGT reference sequence accession numbers: AE000658-AE000661.

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Table A.10.6 Comparison of *TRBV* Gene Nomenclatures^{a,b}

IMGT <i>TRBV</i> gene name (Folch and Lefranc, 2000a)	Wei et al. (1994)	Arden et al. (1995)	Rowen et al. (1996)
<i>TRBV30</i>	20S1	20S1	30
<i>TRBV29-1</i>	4S1	4S1	29-1
<i>TRBV28</i>	3S1	3S1	28
<i>TRBV27</i>	14S1	14S1	27
<i>TRBVB</i>	—	34S1	—
<i>TRBV26</i>	—	28S1	26
<i>TRBVA</i>	—	33S1	—
<i>TRBV25-1</i>	11S1	11S1	25-1
<i>TRBV24-1</i>	15S1	15S1	24-1
<i>TRBV23-1</i>	19S1	19S1	23-1
<i>TRBV22</i>	—	29S1	22-1
<i>TRBV21-1</i>	10S1	10S1	21-1
<i>TRBV20-1</i>	2S1	2S1	20-1
<i>TRBV19</i>	17S1	17S1	19
<i>TRBV18</i>	18S1	18S1	18
<i>TRBV17</i>	26S1 ^c	26S1	17
<i>TRBV16</i>	25S1	25S1	16
<i>TRBV15</i>	24S1	24S1	15
<i>TRBV14</i>	16S1	16S1	14
<i>TRBV12-5</i>	8S3	8S3	12-5
<i>TRBV12-4</i>	8S2	8S2	12-4
<i>TRBV12-3</i>	8S1	8S1	12-3
<i>TRBV11-3</i>	21S4	21S2	11-3
<i>TRBV10-3</i>	12S2	12S1	10-3
<i>TRBV13</i>	23S1	23S1	13
<i>TRBV7-9</i>	6S5	6S4	7-9
<i>TRBV5-8</i>	5S8	5S4	5-8
<i>TRBV7-8</i>	6S3	6S2	7-8
<i>TRBV6-9</i>	13S4	13S4	6-9
<i>TRBV5-7</i>	5S7	5S7	5-7
<i>TRBV7-7</i>	6S14	6S6	7-7
<i>TRBV6-8</i>	13S7	13S7	6-8
<i>TRBV5-6</i>	5S2	5S2	5-6
<i>TRBV7-6</i>	6S4	6S3	7-6
<i>TRBV6-7</i>	13S8	13S8	6-7
<i>TRBV5-5</i>	5S3	5S3	5-5
<i>TRBV7-5</i>	6S12	6S9	7-5
<i>TRBV6-6</i>	13S6	13S6	6-6
<i>TRBV5-4</i>	5S6	5S6	5-4
<i>TRBV7-4</i>	6S11	6S8	7-4
<i>TRBV6-5</i>	13S1	13S1	6-5
<i>TRBV12-2</i>	8S5	8S5	12-2
<i>TRBV11-2</i>	21S3	21S3	11-2
<i>TRBV10-2</i>	12S3	12S3	10-2
<i>TRBV12-1</i>	8S4	8S4	12-1
<i>TRBV11-1</i>	21S1	21S1	11-1
<i>TRBV10-1</i>	12S4	12S2	10-1

continued

Table A.10.6 Comparison of *TRBV* Gene Nomenclatures^{a,b}, continued

IMGT <i>TRBV</i> gene name (Folch and Lefranc, 2000a)	Wei et al. (1994)	Arden et al. (1995)	Rowen et al. (1996)
<i>TRBV9</i>	1S1	1S1	9
<i>TRBV5-3</i>	5S5	5S5	5-3
<i>TRBV8-2</i>	—	32S1	8-2
<i>TRBV7-3</i>	6S1	6S1	7-3
<i>TRBV6-4</i>	13S5	13S5	6-4
<i>TRBV5-2</i>	—	31S1	5-2
<i>TRBV8-1</i>	—	30S1	8-1
<i>TRBV7-2</i>	6S7	6S5	7-2
<i>TRBV6-3</i>	13S2b	13S2b	6-3
<i>TRBV4-3</i>	7S2	7S2	4-3
<i>TRBV3-2</i>	9S2	9S2	3-2
<i>TRBV6-2</i>	13S2a	13S2a	6-2
<i>TRBV4-2</i>	7S3	7S3	4-2
<i>TRBV7-1</i>	6S10	6S7	7-1
<i>TRBV6-1</i>	13S3	13S3	6-1
<i>TRBV5-1</i>	5S1	5S1	5-1
<i>TRBV4-1</i>	7S1	7S1	4-1
<i>TRBV3-1</i>	9S1	9S1	3-1
<i>TRBV2</i>	22S1	22S1	2
<i>TRBV1</i>	—	27S1	1

^a*TRBV* genes are listed from 3' in the *TRB* locus (top of the table) to 5' (bottom of the table).

Blank cells indicate that no corresponding name exists.

^bSee *TRB* Locus and Table A.10.2 for more information.

^cIMGT note: 26S1 was defined in Slightom et al. (1994).

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