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REVIEW

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The human anti-thyroid peroxidase autoantibody repertoire in Graves' and Hashimoto's autoimmune thyroid diseases

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Abstract Human anti-thyroid peroxidase (TPO) autoantibodies (aAb) are generated during autoimmune thyroid diseases (AITD). Within recent years, increasing knowledge of the TPO-specific aAb repertoire, gained mainly by the use of combinatorial library methodology, has led to the cloning and sequencing of around 180 human anti-TPO aAb. Analysis of the immunoglobulin (Ig) variable (V) genes encoding the TPO aAb in the ImMunoGeneTics database (IMGT) (http://imgt.cines.fr) reveals major features of the TPO-directed aAb repertoire during AITD. Heavy chain VH domains of TPOspecific aAb from Graves' disease patients preferentially use D proximal IGHV1 genes, whereas those from Hashimoto's thyroiditis are characterized more frequently by IGHV3 genes, mainly located in the middle of the IGH locus. A large proportion of the anti-TPO heavy chain VH domains is obtained following a VDJ recombination process that uses inverted D genes. J distal IGKV1 and IGLV1 genes are predominantly used in TPO aAb. In contrast to the numerous somatic hypermutations in the TPO-specific heavy chains, there is only limited amino acid replacement in most of the TPO-specific light chains, particularly in those encoded by J proximal IGLV or IGKV genes, suggesting that a defect in receptor edit-

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S. Péraldi-Roux Faculté de Pharmacie, CNRS UMR 5094, Institut de Biotechnologie et Pharmacologie, 15 avenue Charles Flahault, BP14491, 34093 Montpellier Cedex 5, France ing can occur during aAb generation in AITD. Among the predominant *IGHV1* or *IGKV1* TPO aAb, conserved somatic mutations are the hallmark of the TPO aAb repertoire. The aim of this review is to provide new insights into aAb generation against TPO, a major autoantigen involved in AITD.

Keywords Thyroid peroxidase · Autoantibody · Phage display · Variable gene · IMGT database

Introduction

The anti-thyroid peroxidase (TPO) autoantibodies (aAb) are the most frequently represented aAb in the sera of patients suffering from autoimmune thyroid disease (AITD); they are present in 90% of Hashimoto's thyroiditis and 75% of Graves' disease patients (Mariotti et al. 1990). In vitro cytotoxic effector functions mediated by TPO-specific aAb, such as C3 complement activation (Chiovato et al. 1993; Parkes et al. 1994; Wadeleux et al. 1989) and antibody-dependent cell cytotoxicity (Bogner et al. 1995; Guo et al. 1997; Metcalfe et al. 1997; Rodien et al. 1996; Weetman et al. 1989), trigger thyroid cell destruction. Moreover, it has been suggested that thyroidinfiltrating B lymphocytes as antigen-presenting cells through membrane-bound anti-TPO antibodies modulate antigen processing (Guo et al. 1996; McLachlan and Rapoport 1992; Rapoport et al. 1995).

Only one human anti-TPO antibody was obtained by cell immortalization (Horimoto et al. 1992). However, McLachlan and Rapoport's group pioneered the application of combinatorial libraries to the study of aAb in thyroid diseases (Portolano et al. 1991), and a large number of human anti-TPO aAb have since been isolated by this group and others (Chazenbalk et al. 1993; Hexham et al. 1994; Jaume et al. 1994a, b; Jaume et al. 1997; McIntosh et al. 1997; Portolano et al. 1992, 1993a, b; 1995; Prummel et al. 1994a, b). In the last 2 years, about 100 anti-TPO aAb directed against immunodominant or non-immunodominant epitopes have been described

(Chapal et al. 2000; 2001; Guo et al. 1999; Pichurin et al. 2001). Given this enlarged TPO-specific repertoire, and particularly the numerous Ig gene sequences published to date, we compiled and analyzed the genes encoding these aAb using the international ImMunoGeneTics database (IMGT) (http://imgt.cines.fr), an integrated information system devoted to the study of immunoglobulins, T-cell receptors, and major histocompatibility molecules of several vertebrate species (Giudicelli et al. 1997; Lefranc and Lefranc 2001).

TPO-specific heavy chain gene usage in AITD

Ig variable domain sequences encoding TPO aAb have been obtained from Fab and single chain variable fragment (scFv) combinatorial libraries, mainly derived from thyroid-infiltrating B cells of Graves' disease patients (Chapal et al. 2000; 2001; Chazenbalk et al. 1993; Jaume et al. 1994a, b, 1997; Portolano et al. 1992, 1993a, b, 1995; Prummel et al. 1994a, b). Only two libraries constructed from thyroid-infiltrating B cells or lymph node B lymphocytes of Hashimoto's patients have been described (Hexham et al. 1994; McIntosh et al. 1997). Although we cannot formally exclude that differences observed in IGV gene usage of TPO-specific aAb obtained from the libraries cited in Table 1 (consisting of parts a, b and c) are due to preferential primer amplification of certain IGV genes or gene families, we consider that the data reflect the reality in vivo since the analyses were carried out on more than 180 human anti-TPO aAb obtained from four laboratories that used different primers. Analysis of the heavy chain variable domains of the anti-TPO aAb shows a restriction in the IGHV gene usage in both Graves' and Hashimoto's AITD (Table 1, consisting of parts a, b and c) (McIntosh et al. 1998; McLachlan and Rapoport 2000). The heavy chains of the anti-TPO aAb are mainly encoded by genes of the IGHV1 (75.4%) and IGHV3 (21.2%) subgroups, with a large predominance of the IGHV1-3 gene in thyroid diseases.

Interestingly, IGHV gene analysis of anti-TPO aAb from patients with Graves' disease or with Hashimoto's hypothyroiditis clearly indicates a discrimination in IGHV subgroup usage (Table 2). In Graves' disease, the anti-TPO aAb mainly use IGHV1 subgroup genes (88.9%), with overrepresentation of IGHV1-3 (50.4%) and IGHV1-2 (25.5%). In Hashimoto's thyroiditis, the IGHV3 subgroup (71%) is dominant among the anti-TPO aAb, with a large predominance of IGHV3-21 (47.4%) and IGHV3-23 (18.4%) (Table 1 (consisting of parts a, b and c) and 2). Preferential use of IGHV4, IGHV5, and IGHV6 genes by aAb in autoimmune diseases was suggested by several studies (Dijk-Hard van et al. 1999; Melero et al. 1998; Pascual and Capra 1992; Pascual et al. 1992a, b, c; Roben et al. 1996). On the other hand, underexpression of the *IGHV1* subgroup in aAb is a very common feature in autoimmune diseases, as demonstrated for numerous autoantigens (Bona et al. 1993). The overexpression of the IGHV3 subgroup in Hashimoto's thyroiditis and that of the *IGHV1* subgroup in Graves' disease seems to be a characteristic of the anti-TPO aAb repertoire, and suggests that there is a skewing of *IGHV* gene usage in TPO-specific aAb in the sera of patients suffering from autoimmune thyroid diseases.

With regard to the organization of the human IGH locus (Fig. 1), TPO-specific aAb from patients with Graves' disease and from Hashimoto's hypothyroiditis preferentially use D proximal IGHV1 genes and D distal IGHV3 genes, respectively. Two different hypotheses can explain the preferential expression and/or selection of a particular IGHV gene: (1) selection derived from preferential rearrangement due to the gene position in the IGH locus and/or accessibility to the recombinase machinery and (2) functional selection based on the recognition of defined epitopes on the TPO molecule (Sasso et al. 1989). The preferential use of the D proximal IGHV5 subgroup gene previously designated 7183 is well documented in mice (Bona et al. 1993), but the fact that genes from IGHV subgroups are scattered throughout the IGH locus (Fig. 1) does not support the "position" hypothesis. On the other hand, the fact that non-IDR (immunodominant region) TPO-specific aAb show a restricted *IGHV1–69* gene usage (Pichurin et al. 2001) argues in favor of the second hypothesis.

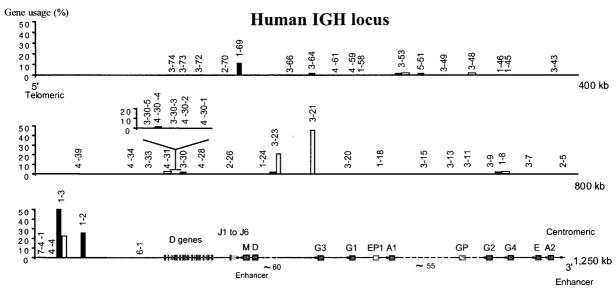
The D genes used by these aAb show a high diversity with a large number of genes in an inverted orientation of transcription (38%) (Table 1, consisting of parts a, b and c). Inverted D genes are rarely used by aAb, and this event seems to be a peculiarity of anti-TPO aAb. This observation suggests the possible involvement of particular mechanisms such as the use of D genes with irregular spacers (DIR elements) (Tuaillon and Capra 1998), preferential V-D rearrangements (Tuaillon and Capra 2000b), or modulation of terminal deoxynucleotidyltransferase activity (Tuaillon and Capra 2000a) to generate heavy chain diversity in the TPO repertoire. Analysis of D gene usage suggests that there is no apparent restriction in D gene use, whereas IGHJ4 (61.6%) and IGHJ6 (29.9%) are preferentially rearranged among the TPO-directed aAb (Tables 1 (consisting of parts a, b and c), 2) in Graves' disease.

TPO-specific light chain gene usage in AITD

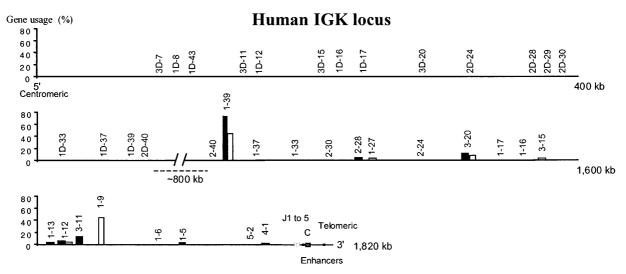
J distal *IGKV1* and *IGLV1* genes (Fig. 1) are preferentially rearranged in TPO-specific recombinant aAb (Tables 1 (consisting of parts a, b and c) and 2). Within

Fig. 1 Germline gene usage of human anti-thyroid peroxidase ► (TPO) antibodies in relation to their position on the immunoglobulin heavy (*IGH*), kappa (*IGK*), and lambda (*IGL*) variable gene loci. Percentage of anti-TPO clones derived from the corresponding germline gene of patients with Graves' disease (*solid bars*), and Hashimoto's thyroiditis (*open bars*). Genes *IGKVI-12* and *IGKVI-39* could not be differentiated from their duplicated genes *IGKVID-12* and *IGKID-39*, respectively. The loci representations were recovered and simplified from the IMGT database and the legend may be found at http://imgt.cines.fr

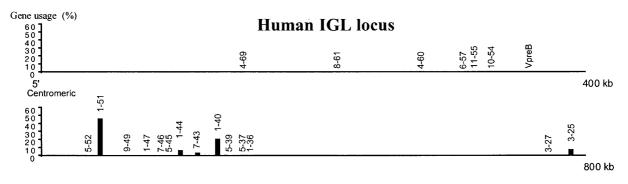
1a



1b



1c



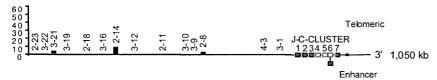


Table 1a Human anti-thyroid peroxidase (TPO) antibody fragments isolated from combinational libraries. Antibodies showing in-cell H/L associations are boxed

	Primer	Clone	Heavy chain ger	Y.		Light chain gene	b	Affinity*	TPO
	specificity		IGHV	IGHD ^d	IGHI	IGKV or IGLV	IGKU or IGLU	(nM)	domain
Lambda phage librar	les (λ>ZAP,)'							
F-1- 4 F	VIA CLEAN	001.0	1011111 0100	10	ICH HARD	IDIO II IID 30MI	ICH IDIO	0.00	100/4
Fab from Graves' thyroid pan B cells	$\gamma 1$ and κ	SP1.2 SP1.4	IGHV1-2*02	ND ND	IGHJ6*02	IGKV1/1D-39*01	IGKJ2*01 IGKJ3*01/4*01/5*01	0.08	IDR/A
(Partolana elf al., 1991; 1992)		SP1.5	IGHV1-2*02 IGHV1-2*02	ND	IGHJ6*02 IGHJ6*02	IGKV1/1D-39*01	IGKJ3*01/4*01/5*01	0.22	IDR/A1 IDR/A1
grandidid er at, 1991; 1992;	*	35.0	10HV1-2 02	NO	NOPEU UZ	IOKA131D/34 01	13133 0174 0173 01	0.00	Diches
SP1-2 IGHV x different	y) and k	SP1.12	id SP1.2	ld SP1.2	id SP1.2	IGKV1/1D-39*01	IGKJ1*01		
IGKV (Roulette)		SP1.13	ld SP1.2	ld SP1.2	ld SP1.2	IGKV1/1D-39*01	IGKJ2*01		
(Partolano et al., 1993b)		SP1.14	kd SP1.2	ld SP1.2	ld SP1.2	IGKV1/1D-39°01	IGKJ2*01		
		SP1.16	kd SP1.2	id SP1.2	ld SP1.2	IGKV1/1D-39*01	IGKJ2*01		
		SP1-17	id SP1.2	id SP1.2	id SP1.2	IGKV1/1D-39*01	IGKJ2*01		
		SP1.18	kd SP1.2	id SP1.2	id SP1.2	IGKV1/1D-39*01	IGKJ2*01	0.00	100.11
		SP1.20	kd SP1.2	id SP1.2	id SP1.2	IGKV1/1D-39*01	IGKJ1*01	0.09	IDR/A
SP1-2 IGKV x different	yl/yt and	sP4.6	IGHV1-2*02	IGHD2-2*01inv/02inv/03inv	IGHJ4*02	id SP1.2	ld SP1.2	0.15	IDR/A
IGHV (Roulette)		SP1.7	IGHV1-2*02	ND	IGHJ6*02	ld SP1.2	ld SP1.2		IDR/A
(Portolano et al., 1993b)		SP1.9	IGHV1-2*02	ND	IGHJ6*02	ld SP1.2	ld SP1.2		IDR/A
Fab from Graves'	yl and k	WR1.7	IGHV1-3*01	IGHD6-13*01	IGHJ4*02	IGKV1/1D-39*01c	IIGKJ1*01d	0.2	IDR/A2
thyroid pan B cells	365	WR1.9	IGHV1-3*01	IGHD6-13*01	IGHJ4*02	IGKV1/1D-39*01c			7.5
(Chazenbalk et al., 1993)									
Fab from Graves'	y4 and k	WR4.2	IGHV1-2*02	IGHD2-2*01lnv	IGH4 ⁹	IGKV1/1D-39*01	IGKJ2 ⁰		
thyroid pan B cells		WR4.3	IGHV1-2*02	IGHD2-2*01l/w	IGH4 ^a	IGKV1/1D-39*01	KGKJ2 ⁸		
(Chazenbalk et al., 1993)		WR4.4	IGHV1-2*02	IGHD2-2*01Inv	IGH4 ⁰	IGKV1/1D-39°01	IGKJ2*01		
		WR4.5	IGHV1-2*02	IGHD2-2*01inv	IGH4 ³	IGKV1/1D-39*01	IGKJ2*01	0.31	IDR/A
		WR4.7	_0	_0	-0	IGKV1/1D-39*01	IGKJ1*01		
		WR4.8	IGHV1-2*02	IGHD2-2*01Inv	IGH4 ^d	IGKV1/1D-39°01	KGKJ2*01		
		WR4. 9	_0	_0	_0	IGKV1/1D-39*01			
		WR4.10	IGHV1-2*02/4	IGHD2-2*01inv/2inv/3inv	IGHJ4*03	IGKV1/1D-39°01	IGKJ2*01		
		WR4.12	IGHV1-2*02/4	IGHD2-2*01inv/2inv/3inv	IGHJ4*02	IGKV1/1D-39*01	IGKJ2*01		
		WR4.21	IGHV1-2*02	IGHD2-2*01Inv	IGH4 ^d	IGKV1/1D-39*01			
		WR4.22	IGHV1-2*02	IGHD2-2*01inv	IGH4 ⁴	IGKV1/1D-39*01	IGKJ2*01 IGKJ2*01		
		WR4.25 WR4.27	IGHV1-2*02/4 IGHV1-2*02/4	IGHD2-2*01inv/2inv/3inv		IGKV1/1D-39*01 IGKV1/1D-39*010			
		Y 30 30 30 30 30	IGHV1-2*02/4	IGHD2-2*01lnv/2lnv/3lnv IGHD2-2*01lnv/2lnv/3lnv	IGHJ4*02 IGHJ4*02	IGKV1/1D-39*01	KGKJ2*01		
		WR4.28 WR4.31	IGHV1-2*02/4	IGHD2-2*01inv/2inv/3inv	IGHU4*02	IGKV1/1D-39*01	IGKJ2*01		
		WR4.32	IGHV1-2*02	IGHD2-2*01inv/2inv/3inv	IGHJ4*02	IGKV1/1D-39*01	IGKJ2*01		
		WR4.33	IGHV1-2*02	IGHD2-2*01inv	IGH4 ²	IGKV1/1D-39*01	IGKJ2*01		
		WR4.34	IGHV1-2*02	IGHD2-2*01inv/2inv/3inv	IGHU4*02	IGKV1/1D-39*01	IGKJ2*01		
		WR4.35	IGHV1-2*02	IGHD2-2*01inv/2inv/3inv	IGHJ4*02	IGKV1/1D-39*01	IGKJ2*01		
		WR4.36	IGHV1-2*02	IGHD2-2*01lnv	IGH4 ⁰	IGKV1/1D-39*01	IGKJ2*01		
		WR4.37	IGHV1-2*02	IGHD2-2*01l/w	IGH4 ³	IGKV1/1D-39*01	IGKJ2*01		
Each from Comme	and instantion	TD1 2	ICHVA FAMI	IGHD6-6*01inv	ICH MINS	1010/11/10 20:01	KCK II+01	0.51±0.01	IDR/A:B
Fab from Graves' thyroid pan 8 cells	γ 1 and κ	TR1.5	IGHV3-53*01 IGHV3-53*01	IGHD6-6*01inv	IGHJ6*03	IGKV1/1D-39*01 IGKV1/1D-39*01	IGKJ1*01 IGKJ2*01	0.0130.01	IDR/A:8
(Chazenbak et al. 1993)		TR1.6	IGHV1-69*06	IGHD6-13*01inv/5-12*01inv			IGKJ2*01		IDR/B1
(Citato Can of al., 1995)		TR1.8	IGHV1-69*06	IGHD3-16*01	IGHJ3*01	IGKV2/2D-28*01	IGKJ2*01	0.27±0.01	IDR/B1
		TR1.9	IGHV1-3*01	IGHD1-26*01	IGHJ4*02	IGKV1-13*02	IGKJ4*01	0.15±0.02	IDR/B2
		TR1.10	IGHV1-3*01	IGHD3-16*01inv/1-14*01	IGHJ4*02	IGKV1/1D-39*01	IGKJ1*01	0.15	IDR/A
			32 () () () () () () () () () (/3-3*01inv/2inv/1-20*01					
		TR1-13	IGHV1-3 ^a	_0	IGH,M ^o	IGKV1-13*02	IGKJ3*01		
Fab from Graves'	$\gamma 1$ and κ	JA1.9	IGHV1-2*02	ND	IGHJ6*02	IGKV1/1D-39*01	IGKJ4*01		
thyroid pan B cells									
(Charenbak et al., 1993) Faib from Graves'	yT and k/X	DESCRIPTION OF THE PROPERTY OF	IGHV3-30-3*01 ^g	IGHD5-5*01 ^g	IGHJ4°	IGKV4-1 ^g	IGKJ4 ^a	2.2	IDR/B
	YI GOOD K/A	WR1.223	IGHV3-23*01#	IGHD3-9*01inv ^g	IGHJ3 ⁰		IGKJ5 ⁸	0.81	IDR/B
thyroid pan B cells		WR1.223	IGHV3-23 UI*	IGHLU3-9 UTINY	IGHU3"	IGKV4-1 st	K-KJO"	0.81	IDR/B
(Jaume et al., 1997)	ul ond v	CANT	ICAJHI 28	ICHD3.3/3.39	ICH IKS	ICN/3-119	_0		
Fab from Graves'	γl and κ		IGVH1-2 ²	KGHD3-3/2-2 ⁹	IGHJ6°	IGKV3-11 ^g	_0		
Fab from Graves' thyroid pan B cells	yl and ĸ	G(N) 2	IGHV1-3 ²	ND	IGHJ4º	IGKV1/1D-39*01 ²			
Fab from Graves'	γ1 and κ	G(N) 2 G(N) 3	IGHV1-3 ²	ND ND	IGHJ4° IGHJ4°	IGKV1/1D-39*01 [©] IGKV1/1D-39*01 [©]	_0		
Fab from Graves' thyroid pan B cells	γl and κ	G(N) 2 G(N) 3 G(N) 4	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ²	ND	IGHJ4º IGHJ4º IGHJ6º	IGKV1/1D-39*01 ^g IGKV1/1D-39*01 ^g IGKV3-11 ^g	_9		
Fab from Graves' thyroid pan B cells	y) and κ	G(N) 2 G(N) 3	IGHV1-3 ²	ND ND	IGHJ4° IGHJ4°	IGKV1/1D-39°01° IGKV3-11° IGKV3-11° IGKV1/1D-39°01°	_0 _0		
Fab from Graves' thyroid pan B cells	yl and ĸ	G(N) 2 G(N) 3 G(N) 4	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ²	ND ND IGHD3-3/2-2 ⁹	IGHJ4º IGHJ4º IGHJ6º	IGKV1/1D-39*01 ^g IGKV1/1D-39*01 ^g IGKV3-11 ^g	_0 _0		
Fab from Graves' thyroid pan B cells	γ1 and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 5	IGHV1-3 ² IGHV1-3 ² IGHV1-3 ²	ND ND IGHD3-3/2-2 ⁹ IGHD1-26Inv/2-8inv ⁹	IGHJ4º IGHJ4º IGHJ6º IGHJ6º	IGKV1/1D-39°01° IGKV3-11° IGKV3-11° IGKV1/1D-39°01°	_0 _0 _0		
Fab from Graves' thyroid pan B cells	γl and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ⁹ IGHV1-3 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2 ⁹ IGHD1-26Inv/2-8inv ⁹ ND	IGHJ4º IGHJ6º IGHJ6º IGHJ4º IGHJ6º	IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01°	_9 _9 _0 _9		
Fab from Graves' thyroid pan B cells	γl and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ² IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8Inv ⁹ ND IGHD1-26Inv/2-8Inv ⁹ ND	IGHJ4° IGHJ6° IGHJ6° IGHJ6° IGHJ6° IGHJ6°	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01°	_9 _9 _0 _9		
Fab from Graves' thyroid pan B cells	γl and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁹ ND IGHD1-26Inv/2-8inv ⁹ ND IGHD3-3/2-2°	IGHJ4º IGHJ4º IGHJ6º IGHJ6º IGHJ6º IGHJ6º IGHJ6º IGHJ6º	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11°	_9 _9 _0 _0 _0 _0		
Fab from Graves' thyroid pan B cells	γl and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-2 ³ IGVH1-2 ³	ND ND IGHD3-3/2-2° IGHD1-26/nv/2-8/nv° ND IGHD1-26/nv/2-8/nv° ND IGHD3-3/2-2° IGHD3-3/2-2°	IGHU4º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11°	_9 _9 _9 _9 _0		
Fab from Graves' thyroid pan B cells	yl and ĸ	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17	IGHV1-3 ² IGHV1-3 ² IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁹ ND IGHD1-26Inv/2-8inv ⁹ ND IGHD3-3/2-2°	IGHJ4º IGHJ4º IGHJ6º IGHJ6º IGHJ6º IGHJ6º IGHJ6º IGHJ6º	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11°	_9 _9 _9 _9 _9 _9		
Fab from Graves' thyroid pan B cells		G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22	IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGVH1-2 ³ IGVH1-2 ³	ND ND IGHD3-3/2-2° IGHD1-26/nv/2-8/nv° ND IGHD1-26/nv/2-8/nv° ND IGHD3-3/2-2° IGHD3-3/2-2°	IGHU4º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11°	_9 _9 _9 _9 _9 _9		
Fab from Graves' thyroid pan B cels (Guo et al., 1999) Filamentous phage M	oranies (oho	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22	IGHV1-3 ^a IGHV1-3 ^a IGVH1-2 ^a IGHV1-3 ^a IGHV1-3 ^a IGHV1-3 ^a IGHV1-3 ^a IGHV1-3 ^a IGHV1-2 ^a IGVH1-2 ^a IGVH1-2 ^a IGVH1-2 ^a	ND ND IGHD3-3/2-2° IGHD1-26/nv/2-8/nv° ND IGHD1-26/nv/2-8/nv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2°	IGHU69 IGHU69 IGHU69 IGHU69 IGHU69 IGHU69 IGHU69 IGHU69 IGHU69	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV3-11"		0,35±0.11	
Fab from Graves' thyroid pan B cels (Super al., 1999)		G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22	IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGVH1-2 ³ IGVH1-2 ³	ND ND IGHD3-3/2-2° IGHD1-26/nv/2-8/nv° ND IGHD1-26/nv/2-8/nv° ND IGHD3-3/2-2° IGHD3-3/2-2°	IGHU4º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º IGHU6º	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11°	-9999 KSKL4*01	0.35±0.11	
Fab from Graves' thyroid pan B cels (Superal, 1999) Filamentous phage M Fab from Graves'	oranies (oho	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22	IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGVH1-2 ³ IGVH1-2 ³ IGVH1-2 ³ IGVH1-2 ⁴	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8Inv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2°	IGHJ4°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11"	-9999 KSKL4*01	0.35±0.11 0.54±0.15	IDR/A
Fab from Graves' thyroid pan B cels (Supertal, 1999) Filamentous phage lit Fab from Graves' thyroid pan B cels	oranies (oho	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 17 G(N) 22 TR1.21 TR1.22 TR1.23 TR1.32-1.33	IGHV1-3° IGHV1-3° IGVH1-2° IGHV1-3° IGHV1-3° IGHV1-3° IGVH1-2° IGVH1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-3° IGHV1-	ND ND IGHD3-3/2-2° IGHD1-26/mv/2-8/mv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01/mv IGHD5-18*01/mv/5-5*01/mv IGHD5-24*01 IGHD4-1*01/mv/4-4*01/mv	IGHU4° IGHU6°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01		0.54±0.15 0.57	IDR/A IDR/A:8
Fab from Graves' thyroid pan B cels (Supertal, 1999) Filamentous phage lit Fab from Graves' thyroid pan B cels	oranies (oho	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 17 G(N) 19 G(N) 12 G(N) 22 IRI.21 IRI.22 IRI.23	IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-2 ³ IGHV1-2 ³ IGVH1-2 ³ IGHV1-2 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2° IGHD1-26/nv/2-8/nv° ND IGHD1-26/nv/2-8/nv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01/nv IGHD5-18*01/nv/5-5*01/nv IGHD5-24*01	IGHU4° IGHU6°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11"		0.54±0.15	IDR/A
Fab from Graves' thyroid pan B cels (Superal, 1999) Filamentous phage M Fab from Graves' thyroid pan B cels (Parloino et al, 1993a)	oranies (oho 71 and k	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 7 G(N) 7 G(N) 19 G(N) 19 G(N) 19 G(N) 22 IRI.21 IRI.22 IRI.23 IRI.33-1.33	IGHV1-3° IGHV1-3° IGVH1-2° IGHV1-3° IGHV1-3° IGHV1-3° IGVH1-2° IGVH1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-2° IGHV1-3° IGHV1-	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-0*01/1-1*01 IGHD6-25*01/1inv/3-10*01	IGHU4° IGHU6°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01		0.54±0.15 0.57	IDR/A IDR/A:8 IDR/B
Fab from Graves' thyroid pan B cels (Sue et al., 1999) Filamentous phage Mit Fab from Graves' thyroid pan B cels (Pariotano et al., 1993a) Fab from Hashimoto's thyroid pan B cels	oranies (oho 71 and k	G(N) 2 G(N) 3 G(N) 4 G(N) 6 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22 IRI.21 IRI.22 IRI.23 IRI.33 GF	IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-2 ³ IGHV1-3 ³	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁹ ND IGHD1-26Inv/2-8inv ⁹ ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-20*01/1-1*01 IGHD6-25*01/1inv/3-10*01 /3-3*01/02	IGHU4° IGHU6°	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11° IGKV3-11° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01	-9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -	0.54±0.15 0.57 0.30 80	IDR/A-B IDR/A-B IDR/B Gs 2G4
Fab from Graves' thyroid pan B cels (Suc et al., 1999) Filamentous phage Mit Fab from Graves' thyroid pan B cels (Pantono et al., 1993a) Fab from Hashimoto's thyroid pan B cels	oranies (oho 71 and k	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 7 G(N) 7 G(N) 19 G(N) 19 G(N) 19 G(N) 22 IRI.21 IRI.22 IRI.23 IRI.33-1.33	IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-3 ³ IGHV1-2 ⁵ IGVH1-2 ⁵ IGVH1-2 ⁵ IGVH1-2 ⁵ IGVH1-2 ⁵ IGVH1-2 ⁵ IGVH1-2 ⁵ IGHV1-2 ⁵	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-0*01/1-1*01 IGHD6-25*01/1inv/3-10*01	IGHU4° IGHU6°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01 IGKV1/ID-39'01		0.54±0.15 0.57 0.30	IDR/AR IDR/AR IDR/B as 2G4
Fab from Graves' thyroid pan B cels (Sucertal, 1999) Filamentous phage Mit Fab from Graves' thyroid pan B cels (Paristano et al, 1993a) Fab from Hashimoto's thyroid pan B cels (Hasham et al, 1994	oraries (oha γ1 and κ γ1 and κ	G(N) 2 G(N) 3 G(N) 4 G(N) 6 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 12 IR1.21 IR1.22 IR1.23 IR1.37 6 F 7 F 101	IGHV1-3° IGHV1-3° IGVH1-2° IGHV1-3° IGHV1-3° IGHV1-3° IGHV1-3° IGVH1-2° IGVH1-2° IGHV1-2° IGHV1-2°02 IGHV1-2°02 IGHV1-3°01 IGHV1-8°01 IGHV1-8°01 IGHV3-3°01 IGHV3-3°01	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁹ ND IGHD1-26Inv/2-8inv ⁹ ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-10*01inv/4-4*01inv IGHD5-25*01/1inv/3-10*01 /3-3*01/02 IGHD3-10*01 IGHD3-10*01 IGHD3-10*01 IGHD3-3*01/2	IGHU4° IGHU6°	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11° IGKV3-11° IGKV3-11° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-20°01° IGKV1/ID-39°01°	GKU3*01 GKU3	0.54±0.15 0.57 0.30 80 80 9.3	IDR/As IDR/As IDR/B IDR/
Fab from Graves' thyroid pan B cells (Superal, 1999) Filamentous phage it Fab from Graves' thyroid pan B cells (Pensiano et al, 1993a) Fab from Hashimoto's thyroid pan B cells (Hesham et al, 1994 Fab from Graves'	oranies (oho 71 and k	G(N) 2 G(N) 3 G(N) 4 G(N) 6 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22 IR1.21 IR1.22 IR1.23 IR1.37 6 F 7 F 101 IR1.41	IGHV1-3 ³ IGHV1-2 ⁵ IGHV1-3 ⁵	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁰ ND IGHD1-26Inv/2-8inv ⁰ ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-10*01inv/4-4*01inv IGHD5-24*01 IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-3*01/12 IGHD3-3*01/2 IGHD3-10*01 IGHD3-10*01	IGHU4° IGHU6° IG	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11° IGKV3-11° IGKV3-11° IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV3-20°01 IGKV3-20°01 IGKV3-20°01	-9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -	0.54±0.15 0.57 0.30 80 80 9.3	IDR/AB IDR/BB IDR/B as 2G4 not 2G/ IDR/B
Fab from Graves' thyroid pan B cels (Guertal, 1999) Filamentous phage Mit Fab from Graves' thyroid pan B cels (Hantana et al, 1994) Fab from Hashimota's thyroid pan B cels (Hantana et al, 1994) Fab from Graves' thyroid pan B cels (Hantana et al, 1994)	oranies (otho y) and k y) and k y) and k	G(N) 2 G(N) 3 G(N) 4 G(N) 5 G(N) 6 G(N) 7 G(N) 9 G(N) 19 G(N) 19 G(N) 22 IRI.21 IRI.22 IRI.23 IRI.32-1.33 IRI.37 6 F 7 F 101 IRI.41 WRI.102	ICHV1-3 ³ ICHV1-3 ³ ICHV1-2 ³ ICHV1-3 ³ ICHV1-3 ³ ICHV1-3 ³ ICHV1-3 ³ ICHV1-2 ³ ICHV1-3 ³ ICHV1-6 ³	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv° ND IGHD1-26Inv/2-8inv° ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-11*01inv/4-4*01inv IGHD1-20*01/1-1*01 IGHD6-25*01/1-10*01 IGHD3-3*01/02 IGHD3-3*01/02 IGHD3-3*01/2 IGHD3-10*01	IGHU4° IGHU6°	IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV1/ID-39'01" IGKV3-11" IGKV3-11" IGKV3-11" IGKV3-11" IGKV1/ID-39'01	IGKL1*01 IGKL2*01 IGKL2*01 IGKL2*01 IGKL2*01 IGKL2*01 IGKL2*01	0.54±0.15 0.57 0.30 80 80 9.3 0.8 2	IDR/ABIDR/ABIDR/BB
Fab from Graves' thyroid pan B cells (Superal, 1999) Filamentous phage it Fab from Graves' thyroid pan B cells (Pensiano et al, 1993a) Fab from Hashimoto's thyroid pan B cells (Hesham et al, 1994 Fab from Graves'	oranies (otho y) and k y) and k y) and k	G(N) 2 G(N) 3 G(N) 4 G(N) 6 G(N) 6 G(N) 7 G(N) 9 G(N) 17 G(N) 19 G(N) 22 IR1.21 IR1.22 IR1.23 IR1.37 6 F 7 F 101 IR1.41	IGHV1-3 ³ IGHV1-2 ⁵ IGHV1-3 ⁵	ND ND IGHD3-3/2-2° IGHD1-26Inv/2-8inv ⁰ ND IGHD1-26Inv/2-8inv ⁰ ND IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-3/2-2° IGHD3-16*01inv IGHD5-18*01inv/5-5*01inv IGHD5-24*01 IGHD4-10*01inv/4-4*01inv IGHD5-24*01 IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-10*01inv/4-4*01inv IGHD4-3*01/12 IGHD3-3*01/2 IGHD3-10*01 IGHD3-10*01	IGHU4° IGHU6° IG	IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV1/ID-39°01° IGKV3-11° IGKV3-11° IGKV3-11° IGKV3-11° IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV1/ID-39°01 IGKV3-20°01 IGKV3-20°01 IGKV3-20°01	-9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -9 -	0.54±0.15 0.57 0.30 80 80 9.3	IDR/AB IDR/BB IDR/B as 2G4 not 2G4 IDR/B

Table 1b

Libraries	Primer	Clone	Heavy chain ge	ne	359	Light chain gene		Affinity	TPO
	specificity		IGHV	IGHD	IGH.J	IGKV or IGLV	IGKI or IGLI	(nM)	dama
Filamentous phage lib	raries (pha	ige alsplay))						
Fab from Hashimoto's	y) and κ/λ	126A	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*01/2	IGKV1-9*01	IGKJ4*01		
thyrold pan B cells		126B	IGHV3-21*01/2	IGHD5-12*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		IDR/8
(Mointosh et al., 1997)		126C	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
		126D	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*02	IGKV1-12*01/02 1D-12*02	IGKJ4*01	0.2	
		126 F	IGHV3-21*01/2	IGHD1-7*01/1-20*01	IGHJ5*01/2	IGKV1-9*01	IGKJ4*01		
		126G	IGHV3-21*01/2	IGHD1-1*01		IGKV1-9*01	IGKJ5*01	0.2-3.1	IDR/B
		126H	IGHV3-21*01/2	IGHD4-23*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01	0.2	IDR/B
		1261	IGHV3-21*01/2	IGHD5-12*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
		126J 126TO1	IGHV3-21*01/2 IGHV1-3*01	IGHD3-16*01 IGHD2-2*01iny/3iny	IGHJ6°02 IGHJ6°01	IGKV1-9*01 IGKV1/1D-39*01	IGKJ4*01	3.9	IDR/A
		126102	IGHV1-3*01	IGHD3-9*01inv	IGHJ6*01	IGKV1/1D-39*01		0.4-2.4	IDNO
		126TO3	IGHV1-3*01	IGHD3-9*01inv	IGHJ6*01	IGKV1/1D-39*01		0.4-2.4	
		126TO6	IGHV1-3*01	IGHD3-9*01Inv	IGHJ6*01	IGKV1/1D-39*01	IGKJ4*01	0.4-2.4	
		126108	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*02	IGKV1-9*01	IGKJ5*01	0.2-3.1	
		126TO9 126TO10	IGHV3-21*01/2 IGHV3-21*01/2	IGHD2-21*01 IGHD3-16*01	IGHJ5*02 IGHJ5*02	IGKV1-27*01 IGKV1-9*01	IGKJ4*01 IGKJ4*01	0.094-10	
		126TO15	IGHV3-21*01/2	IGHD5-12*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01	0.094-10	
Fab from Hashimoto's	al and ea	126TP1	IGHV3-21*01/2	IGHD3-16*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
lymph node pan B cel		126TP5	IGHV1-3*01	IGHD3-9*01inv	IGHJ6*02	IGKV1/1D-39*01	IGKJ4*01		IDR/A
(Mointosh et al., 1997)		126TP6	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
		126TP7	IGHV3-21*01/2	IGHD1-1*01		IGKV1-9*01	IGKJ4*01		
		126TP8	IGHV3-21*01/2	IGHD1-1*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
		126TP9	IGHV1-3*01	IGHD6-6*01inv/3-16*01 /3-10*01/2	IGHJ6*02	IGKV1/1D-39*01	1GKJ4*U1		
		126TP10	IGHV3-21*01/2	IGHD3-16*01	IGHJ5*02	IGKV1-9*01	IGKJ4*01		
		126TP13	IGHV1-3*01	IGHD2-2*01 inv/3inv	IGHJ6*02	IGKV1/1D-39*01	IGKJ4*01	2.8	
		126TP14	IGHV1-3*01	IGHD3-9*01inv	IGHJ6*02	IGKV1/1D-39*01	IGKJ4*01	9.1	
		126TP15 131TP2	IGHV1-3*01 IGHV3-23*01	IGHD3-9*01inv IGHD6-6*01inv/4-23*01inv	IGHJ6*02 IGHJ6*01	IGKV1/1D-39*01 IGKV1/1D-39*01	IGKJ4*01 IGKJ3*01	3.1 3.1-4.4	
		131TP5	IGHV3-23*01	/1-26*01inv IGHD6-6*01inv/4-23*01inv	IGHJ6°01	IGKV1/1D-39*01	IGKJ1*01	2.2-15	IDR/A
		131TP6	IGHV3-23*01	/1-26*01inv IGHD6-6*01inv/4-23*01inv	IGHJ6*01	IGKV1/1D-39*01	IGKJ3*01	3.1-4.4	IDR/A
		131TP7	IGHV3-23*01	/1-26*01inv IGHD6-6*01inv/4-23*01inv /1-26*01inv	IGHJ6*01	IGKV1/1D-39*01	IGKJ1*01	2.2-15	IDR/A
		131TP8	IGHV3-23*01	IGHD6-6*01inv/4-23*01inv	IGHJ6*01	IGKV1/1D-39*01	IGKJ1*01	2.2-15	
		131TP14	IGHV3-48*01	/1-26*01inv IGHD3-16*01inv /2-21*01inv/2inv	IGHJ6*01	IGKV3-15*01	IGKJ3*01	2.6	IDR/B
		131TP15	IGHV3-23*01	/2-8*01inv/2inv IGHD6-6*01inv/4-23*01inv /1-26*01irv	IGHJ6*01	IGKV1/1D-39*01	IGKJ1*01	2.2-15	
mAb from Hashimoto! thyroid pan B cells (Hor			GHV3-53°01/2	IGHD6-13*01/6-6*01	IGHJ4*02	IGKV3-20*01	IGKJ5*01	2.5	
			V21/25/02/25/02	170000000000000	12.770/227			05893	0.450000
Fab from Graves'	γl and κ		IGHV1-69*01/6	IGHD3-10*01	IGHJ6*02	IGKV1/1D-39*01		NM	non-ID
thyroid pan B cells		DN 7	IGHV1-3°	IGHD1-26inv/2-8inv ^G	IGHJ6°	IGKV1/1D-39*015			IDR
(select on denature TP		DN 8	IGHV1-3 ⁰	IGHD1-26inv/2-8inv ^a	IGHJ69	IGKV1/1D-39*01		0.15	IDB
(Guoetal, 1999, Pichurin et	al. 2001)	DN 14	IGHV1-3 ⁰	IGHD3-3/2-2 ⁰	IGHJ6 ⁰	IGKV3-11 ^{II}	-0	0.26	IDR
		DN 15	IGHV1-3 ⁰	IGHD1-26inv/2-8inv ^a	IGHJ6 ⁰	IGKV1/1D-39*019			IDB
		DN 16	IGHV1-30	IGHD1-26inv/2-8inv ⁰	IGHJ4º	IGKV1/1D-39*019		0.12	IDB
		DN 20	IGHV1-3 ⁰	ND	IGHJ4 ⁰	IGKV1/1D-39*01	1-9		IDR
Fab from Graves'	yl and κ	N2	IGHV1-3 ^d	ND	IGHJ4 ⁰	IGKV1/1D-39*01	-0		IDR
thyroid pan B cells	de la company	N 5	IGHV1-3 ⁰	ND	IGHJ4 ⁰	IGKV1/1D-39*015			IDR
(Guo et al., 1999)		Nó	IGHV1-3 ⁰	IGHD3-3/2-29	IGHJ6°	IGKV3-110	_9		IDR
		N 8	IGHV1-3 ⁰	ND	IGHJ4 ^G	IGKV1/1D-39*01	_9		IDR
		N 11	IGHV1-3 ^g	ND	IGHJ4 ⁰	IGKV1/1D-39*01			IDR
		N 12	IGHV1-3º	ND	IGHJ4º	IGKV3-20 ⁰	_0		IDR
In-cell scFv from Gravi	yl and k/k	ICA1	IGHV1-3*01	IGHD3-3*01inv/3-9*01inv	IGHJ4*02	IGLV1-51*01	IGLI1*01	4.17	7
thyroid CD19* B cells		ICA5	IGHV1-69*01	IGHD3-3*01	IGHJ4*02	IGLV1-40*02	IGLJ2*01/3*01	1.82	11
(Chapal et al., 2000)		ICB7	_IGHV1-3*01	IGHD3-3*01inv/3-9*01inv /4*03	IGHJ3*01/2	IGLV1-51*01	IGLJ1*01	1.20	111
scFv from Graves'	yl and κ/λ	. AT	IGHV1-3*01	IGHD3-16*01/5-24*01	IGHJ4*02/3	IGLV1-51*01	IGLJ2*01/3*01		
thyroid CD19" B cells	A CONTRACTOR	A2	IGHV1-3*01	IGHD3-16*01		IGLV1-51*01	IGLJ2*01/3*01	4.89	III
(Chapal et al., 2001)		A3	IGHV1-3*01	IGHD7-27*01	IGHJ4*02	IGLV1-51*01	IGLJ3*02		177
		A4	IGHV1-3*01	IGHD5-24*01 /3*01/2	IGHJ4*02	IGLV2-14*01	IGLJ2*01		
		A5	IGHV1-3*01	IGHD4-17*01/4-23*01		IGLV1-51*01	IGLJ2*01/3*01		
		A6 A7	IGHV1-3*02	IGHD7-27*01inv		IGLV1-40*02	IGL 12101 /3101		
		A7 A8	IGHV1-3*01 IGHV3-30*04	IGHD3-3*01Inv/3-9*01inv IGHD4-23*01	IGHJ4°02/3 IGHJ4°02	IGLV1-51*01 IGLV1-44*01	IGLJ2*01/3*01 IGLJ2*01/3*01		
		A9	IGHV3-64*01	IGHD2-15*01inv	IGHJ6*02	IGLV1-40*01	IGLJ2*01/3*01		
		A10	IGHV1-69*01	IGHD3-3*01	IGHJ4*02	IGLV1-40*02	IGLJ1*01	5.43	IV
		A11	IGHV1-3*01	IGHD3-16*01		IGLV1-51*01	IGLII*01	8.03	٧
		A12	IGHV3-64*01	IGHD2-15*01inv	IGHJ6*02	IGLV1-51*01	IGLJ2*01/3*01	1.21	VIII.
		A13 A14	IGHV1-3*01	IGHD3-3*01inv/3-9*01inv IGHD3-3*01inv/3-9*01inv	IGHJ4*02 IGHJ4*02	IGLV1-40*01 IGLV1-40*01	IGLJ1*01 IGLJ1*01		
		A15	IGHV1-3*01	IGHD3-16*01		IGLV1-51*01	IGLJ1*01		
		A16	IGHV1-3*01	IGHD3-3*01inv/3-9*01inv	IGHJ3*01/2	IGLV1-51*01	IGL/1*01		
		A17	IGHV1-69*01	IGHD3-3*01	/4*03 IGHJ4*02	IGLV1-44*01	IGLJ1*01		

Table 1c

Libraries	Primer	Clone	Heavy chain gen	x ·		Light chain gene		Affinity	TPO
	specificity		IGHV	IGHD	IGHJ	IGKV or IGLV	IGKJ or IGLI	(nM)	domair
Filamentous phage (ibraries (pha	nge display)						
scFv from Graves"	y) and k/X	B1	IGHV1-3*01	IGHD5-24*01	IGHJ4*02	IGLV1-40*02	IGLJ3*02		
thyroid pan B cells		B2	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ4*02	IGKV1/1D-39*01	IGKJ4*01	4.35	VI
(Chapal et al., 2001)		B3	IGHV1-3*01	IGHD5-24*01	IGHJ4*02	IGLV7-43*01	IGLJ3*02		
		B4	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ4*02	IGLV1-51*01	IGLJ3*02	2.83	VI
		B5	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ1*01	IGLV1-51*01	IGLJ2*01/3*01	1.99	VI
		B6	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ1*01	IGLV1-51*01	IGLJ1*01	3.54	VI
		B7	IGHV1-3*01	IGHD2-21*01/2/3-10*01/2 /3-22*01		IGKV1D-12*01	IGKJ5*01	2.17	VI/VIII
		88	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ4*02	IGLV1-51*01	IGLJ3*02	0.99	VI
		B9	IGHV1-3*01	KGHD2-21*01/2/3-10*01/2 /3-22*01		EIGLV1-51*01	IGLJ3*02		
		B10	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ4*02	IGLV2-14*01	IGLJ3*02	12.3	VII
		B11	IGHV5-51*01	IGHD3-16*01	IGHJ4*02	IGLV1-51*01	IGLJ2*01/3*01		
scFv from Graves'	yl and k/X	. 11	IGHV1-3*01	IGHD4-4*01/4-11*01	IGHJ4*02	IGLV1-51*01	ND		
thyroid TPO-purified		12	IGHV1-3*02	IGHD2-21*01	IGHJ4*03	IGKV3-11*02	ND	5.09	DX:
Bicells (Chaparetal, 2001)		T3	IGHV1-3*01	IGHD2-21*01/2/3-10*01/2 /3-22*01	IGHJ4*02/3	IGKV1/1D-39*01	IGKJ4*01	1.28	VI/VIII
		14	IGHV1-3*01	IGHD2-8*01inv/2inv /2-21*01inv/2inv	IGHJ4*02	IGLV2-8*01	IGW1*01		
		T5	IGHV1-8*01	IGHD3-3°02inv	IGHJ3*02	IGKV1-5*03	IGKJ2*01	0.77	VI/VIII
		T6	IGHV1-3*01	IGHD2-2*02	IGHJ4*02	IGKV1/1D-39*01	IGKJ2*01		
		17	IGHV1-3*01	ND	IGHJ6*02	IGLV1-40*01	IGLJ2*01/3*01	nesari	2000
		T8	IGHV1-3*01	IGHD2-21*01/2/3-10*01/2 /3-22*01		IGKV1/1D-39*01	IGK,14*01	4.50	VIII
		19	IGHV1-3*01	IGHD2-21*01/2/3-10*01/2 /3-22*01	IGHJ4*02	IGKV1/1D-39*01	ND		
		110	IGHV3-64*01	IGHD6-19*01	IGHJ6*02	IGKV3-11*01	IGKJ4*01	2.19	VI/VIII
		111	IGHV1-3*01	KGHD2-2*02		IGKV1/1D-39*01	IGKJ5*01		
		112	IGHV1-3*01	IGHD4-4*01/4-11*01		IGLV1-40*02	IGLJ3*02		
		113	IGHV1-3*01	ND	IGHJ6*02	IGLV1-40*01	IGLJ2*01/3*01/2	7.95	VIII
Faib from Graves'	yl and k	TF2.3	IGHV1-69°03	IGHD3-10°01	IGHJ6*02	IGKV3-20*01	IGKJ2*01		non-IDI
thyroid pan B cells (Pichuth et al., 2001)		TF2.4	IGHV1-69*04	IGHD3-10*01	IGHJ6*02	IGKV1-12*01/2 /1D-12*02	IGKJ1*01	2.0	non-IDF
		TF2.6	IGHV1-69°02/4/6	IGHD3-10*01	IGHJ6*02	IGKV1/1D-39*01	IGKJ1*01		non-IDF
		TF2.10	IGHV1-69*04	IGHD3-10*01	IGHJ6*02	IGKV1/1D-39*01	IGKJ1*01	2.7	non-IDF
		TF3.5	IGHV1-69*04/6	IGHD3-10°01	IGHJ6*02	IGKV3-20*01	ND	1.2	non-IDF
		TF3.12	IGHV1-69*04/6	IGHD3-10*01	IGHJ6*02	/1/1D-39*01/02	IGKJ2*01		non-IDI
		TF3.14	IGHV1-69*04/6	IGHD3-10*01	IGHJ6*02	IGKV3-20*01	IGKJ4*01		non-IDF
		TF3.19	IGHV1-69*04/6	IGHD3-10*01	IGHJ6*02	IGKV3-20*01	IGKJ2*01	10/200	non-IDI
		12.2	IGHV1-2*02	IGHD1-20*01Inv/1-1*01Inv /6-13*01/6-6*01	IGHJ6*02	IGKV3-11*01	IGKJ2*01	0.25	IDR
		12.5	IGHV5-51*01	IGHD5-18*01/5-5*01	IGHJ6*02	IGKV1D-39*01	IGKJ4*01	0.4	IDR
		T2.6	IGHV1-3*01	/5-18*01inv/5-12*01inv /5-5*01inv/3-22*01inv	IGHJ6*02	IGKV1D-39*01	IGKJ2*01	0.12	IDR
		T2.7	IGHV1-3*01	IGHD5-24*01inv	IGHJ4*02	IGKV1/1D-39*01	IGKJ1*01		IDR
		T2.11	IGHV1-3*01	IGHD3-10*01	IGHJ4*02	IGKV3-20°01	IGKJ2*01	1.6	IDR
		13.2	IGHV1-3*01	IGHD5-24*01inv /5-18*01inv/5-12*01inv	IGHJ6*02	IGKV1/1D-39*01	IGKJ4*01		IDR
		13.3	IGHV1-3*01	/5-5*01inv/3-22*01inv IGHD2-21*02inv /2-15*01inv	IGHJ6*02	IGKV1/1D-39*01	IGKJ1*01	0.2	IDR
		70.4	101 5 (1 040)	/2-2*01inv/2inv/3inv	1001111400	101011 10101 -	100114401	0.00	inc
		13.4	IGHV1-8*01	/6-19*01inv/6-13*01inv /6-6*01inv/5-24*01inv	IGHJ6*02	/ID-12*02	IGKJ4*01	0.22	IDR
		13.5	IGHV1-3*01	IGHD5-24*01inv	IGHJ4*02	IGKV1/1D-39*01	IGKJ1*01	0.12	IDR
		19.0	ROMA I-O DI	MARIDO-24 UTINV	19HJ4-02	IOVALLID-2A.01	noral ul	0.12	R.AK

 $^{^{\}circ}$ Each library was generated from a given single patient sample except those described by Chapal et al.

IGHV1-3*01

IGHV1-3*01

IGHV1-3°01

IGHJ4*02

IGHJ4*02

IGHJ4*02

IGHJ4*02

IGKV1/1D-39°01 IGKJ4°01

IGKV1/1D-39*01 IGKJ1*01

IGKV1/1D-39°01 IGKJ1°01

IGKV3-20*01

IDR

IDR

IDR

IDR

IGHD5-24*01inv

IGHD5-24*01inv

IGHD5-24*01Inv

KGHD3-10*01

13.7

T3.10

T3.13

T3.15

^b Putative closest germine genes determined with IMGT/V-QUEST sequence alignment software (http://imgt.cines.tr). The nomenclature is according to the IMGT (Letranc and Letranc, 2001) and HUGO (Human Genome Organization) nomenclature committee (http://www.gene.ucl.ac.uk/nomenclature). All the germine genes or affets presenting the same scare are presented

^o Affinity measurements were performed by various techniques (Scatchard analysis, Biacore, EUSA)

^dBecause of the short length of the D genes, several putative closest germline D genes have the same score of alignment.

^{*}TPO domains were defined by various methods (EUSA inhibition, Biacore inhibition), IDR characterized according to Chazenbalk et al. (1993) and regions I-X (Chapal et al. 2000, 2001) were determined independently.

All the human anti-TPO antibodies, except 2G4, were isolated from combinatorial libraries.

⁹ Nucleotide sequences not found in public databases. When available, information concerning the proposed germline genes is derived from the cited publications.

h The crystal structure of TR1.9 Fab has been solved (S. Chacko et al. 1996). Residue K713 has been identified to be involved in the TPO IDR epitope recognized by the TR1.9 autoantibody (Guo et al. 2001).

Sequence glanment by IMGT/V-QUEST and IMGT/JunctionAnalysis of ICA5 shows the same score for IGLV1-40°01 and for IGLV1-47°02.

ND: Not determined by IMGT/V-QUEST or IMGT/JunctionAnalysis.

inv: D genes in inverted orientation of transcription. id: identical to in the "roulette" studies.

NM: Not measurable IDR: Immunodominant region

Table 2 Germline genes used by the human TPO-specific autoantibody repertoire (ND not determined by IMGT/V-QUEST)

Thyroid	IG variable gene usagea	gene 1	usagea															
disease	IGHV gene	и	9%	IGHJ gene	и	9%	IGKV gene 1	u c	q%	IGKJ gene	и	9%	IGLV gene	и	9%	IGLJ gene	и	9%
Graves' disease ^c	diseasec																	
	IGHV1-2	35	25.5	IGHJI	2	1.4		_ (IGKJI	18	17.4	IGLVI-40	10	26.3	IGLJ1	13	34.2
	IGHV1-3	69	50.4	CHID	ľ	v	IGKVI-12		2.9	01451	00	0 76	IGLV1-44	C) 0	5.5	171	5	0 90
	1GHVI-69 IGHVI-69	16	11.6	CCHOI	C./	4.0		75	72.8	IGN 72	90	50.9	10-14701	10	4. 4.	19172	10	50.5
				IGHJ4	84.5 61.6	61.6				IGKJ3	α	2.9	IGLV2-8	1	2.6	IGLJ3	14	36.8
	IGHV3-23	7	1.4				IGKV2-28	α	2.7				IGLV2-14	\mathcal{C}	7.9			
	IGHV3-30	Ω,	1. c	1GHJ6	41	29.9	11 6/145/1	9	,	IGKJ4	15	14.5	10 271771	-	0	ND	1	2.6
	IGHV3-53 IGHV3-64	n m	2.2	p-	7	1.4		7	6.8	IGKJ5	8	2.9	IGLV3-21 IGLV3-25	7	5.2			
	IGHV4-30-4	4 1	0.7				IGKV4-1	2	1.9	ND	3	2.9	IGLV7-43	Τ	2.6			
	IGHV5-51	2	1.4							p_	23	22.3						
	p_	2	1.4															
Hashimo	Hashimoto's disease																	
	IGHV1-3	6	23.7	IGHJ4	2	5.2		16	42.1	IGKJI	2	13.1						
	IGHV I–8	_	7.0	IGHJ5	18	47.4	IGKVI–12 IGKVI–27		2.0 2.0	IGKJ2	1	2.6						
	IGHV3-21	18	47.4) i i i	9	1		17 4	44.7		-	į. C						
	IGHV3-23 IGHV3-48	\ -	18.4	IGHJ0	<u>8</u>	4./4	IGKV3-15		26	IGKJ3	4	5.01						
	IGHV3-53	- -	2.6				IGKV3-20	7	5.2	IGKJ4	24	63.0						
	<i>IGHV4–31</i>	\vdash	2.6							<i>IGKJ5</i>	4	10.5						
a IGHD g	a IGHD gene usage is not indicated since numerous anti-TPO	not inc	licated s	since numerou	ıs anti-	ТРО а	antibody gene sequences	ednen		: N=37 for <i>I</i> (GHV aı	nd for R	<i>3HJ</i> ; <i>N</i> =103 ¹	for IG	:KV and 1	$\stackrel{?}{\sim} N=37$ for IGHV and for IGHJ; N=103 for IGKV and for IGKJ; N=38 for IGLV and for	3 for I	GLV and for
present the by $6\% = n/N$	present the same alignment score with different germline genes $0 = m/N \times 100$ where $m = m + m + m = m + m + m + m + m + m = m + m +$	ent sc =numl	tore with	n different ger inti-TPO <i>IGH</i>	mline V_{V}	genes es in tl	present the same alignment score with different germline genes $0 = 100 \text{ JGHV}$ subgroun and	rolln		<i>IGL)</i> d Nucleotide	seamen	ces not	<i>IGLJ</i> d Nucleotide sequences not annotated by IMGT/V-OHEST	IMG	VV-OUE	LS		
N=total n	N=total number of anti-TPO IGHV genes studied	TPO /	GHV ge	nes studied	200		8000 11101 01	Jacar		N=35 for //	$\mathcal{F}HV$ an	nd for /G	e N=35 for IGHV and for IGHJ: N=38 for IGKV and for IGKL	r IGK	V and for	·IGKL		

N=total number of anti-TPO IGHV genes studied

e N=35 for IGHV and for IGHJ; N=38 for IGKV and for IGKL

the *IGKV1* subgroup, a strong restriction is observed: 72.8% of the κ anti-TPO aAb are encoded by genes derived from the IGKV1-39 (or IGKV1D-39) gene in Graves' disease (Tables 1 (consisting of parts a, b and c) and 2) (McIntosh et al. 1998; McLachlan and Rapoport 2000). Concerning the TPO-specific IGL repertoire, few anti-TPO recombinant Fab expressing a λ light chain have been characterized and sequenced. This is probably due to the fact that only a few libraries have been constructed using λ -specific amplification primers (Jaume et al. 1997; McIntosh et al. 1997; Prummel et al. 1994b). The decision by other authors to use only κ -specific amplification primers for library construction was based on the fact that κ-chain TPO aAb predominated in the sera of the thyroid disease patients from whom the library originated (Chazenbalk et al. 1993; Guo et al. 1999; Hexham et al. 1994; Pichurin et al. 2001; Portolano et al. 1991, 1992, 1993a, b). Using a mixture of κ - and λ - specific primers, we recently obtained numerous λ anti-TPO scFv by an in-cell library and random combinatorial libraries (Table 1, consisting of parts a, b and c) (Chapal et al. 2000; 2001). Analysis of this enlarged λ -derived TPO repertoire revealed a dominant use of the *IGLV1* subgroup in thyroid diseases, with two genes mainly found, IGLV1-51 (47.4%) and IGLV1-40 (26.3%) (Tables 1 (consisting of parts a, b and c), 2). Autoantibodies with λ light chains have been described in various autoimmune diseases (Cairns et al. 1989; Prummel et al. 1994a, b; Ravirajan et al. 1998; Serrano et al. 1994; Song et al. 1998); in particular, λ anti-TSHr aAb are involved in thyroid stimulation in patients with Graves' disease (Knight et al. 1986; Williams et al. 1988; Zakarija and McKenzie 1983). Moreover, five IGLV1-40- and one IGLV1-51-derived anti-Tg aAb have been isolated from a combinatorial library constructed from a patient with Hashimoto's thyroiditis (McIntosh et al. 1996, 1998).

H/L pairing of TPO aAb

Chain pairing in a TPO-selected random library can contain in vivo H/L combinations as suggested by "roulette" studies (Costante et al. 1994; Portolano et al. 1993a). This was demonstrated by comparison of H/L combinations obtained from an in-cell library with those obtained from various random libraries (Chapal et al. 2001). However, only TPO-directed aAb from an in-cell combinatorial library (Chapal et al. 2000) and clone 2G4 obtained from cell fusion (Horimoto et al. 1992) formally reflect the in vivo situation (Table 1, consisting of parts a, b and c).

Although a previous study described the lack of promiscuity between TPO-specific heavy and light chains (Portolano et al. 1993a), an extensive analysis of H/L rearrangements of anti-TPO aAb does not show apparent restriction in H/L pairing (Table 1, consisting of parts a, b and c). Indeed, the heavy chains encoded by the dominant *IGHV1–3* gene are associated with light chains encoded by 11 of 18 different *IGKV* or *IGLV* genes (Table 1,

consisting of parts a, b and c). Reciprocally, the most frequently used light chain genes, i.e., IGKV1-39, IGLV1-40, and IGLV1-51, are combined with around 50% of the IGHV genes used by TPO aAb. Overrepresentation of IGHV1-3/IGKV1-39, IGHV1-3/IGHLV1-51, and IGHV1-3/IGLV1-40 pairings probably reflects the predominance of the expressed IGHV, IGKV, and IGLV genes in the TPO antibody repertoire. The clones resulting from an in-cell library and from cell fusion show the *IGHV1–3/IGLV1–51*, *IGHV1–69/IGLV1–40*, and IGHV3-53/IGKV3-20 associations found respectively in 14, 1, and none of the anti-TPO aAb obtained from random combinatorial libraries (Table 1, consisting of parts a, b and c). These observations indicate the need to enlarge the number of in vivo clones to definitively conclude that there is a restricted H/L pairing in TPOspecific aAb, even though it is possible to obtain at least part of the in vivo anti-TPO repertoire with combinatorial libraries.

Amino acid multi-sequence alignment of TPO-specific aAb

Whereas numerous somatic hypermutations are observed in TPO-specific heavy chains whatever the library origin (Table 3, consisting of parts a, b and c)), there is no or only limited amino acid replacement in most TPO-specific light chains, particularly those encoded by the J proximal *IGLV2–14*, *IGKV1–9*, *IGKV3–11*, *IGKV3–15*, IGKV3-20, and IGKV4-1 genes (Tables 1 (consisting of parts a and b), 5). The pattern of mutations in IGHV genes from anti-TPO aAb is typical of an antigen-driven selection during AITD. On the other hand, preferential usage of J proximal IGLV or IGKV genes for some TPO aAb, with little or no residue mutations, strongly suggests a defect in receptor editing of the light chain during aAb generation in AITD, as demonstrated for lupus-associated anti-DNA aAb (Bensimon et al. 1994; Chen et al. 1997). In this case, certain TPO-specific B cells might have been blocked in their capacity to turn off their autoreactivity by light chain replacement, leading to the acquisition of a new specificity.

As previously suggested by others (McIntosh et al. 1997; Portolano et al. 1993b, 1995) and confirmed by our recent publications (Chapal et al. 2000; 2001), extensive analysis of somatic hypermutations among *IGHV1–3*, *IGHV1–2*, and *IGKV1–39* dominant-derived aAb indicate that certain residue replacements (e.g., Ile39 and Thr95 for *IGHV1* genes) are systematically found in the majority of TPO-specific aAb independently of the library, but other amino acid mutations are mostly library or patient specific (Tables 3 (consisting of parts a, b and c), 4 (consisting of parts a and b), and 5). These observations support the hypothesis that the hypermutation process could be the hallmark of the TPO aAb repertoire.

were obtained from databases except antibodies WR1.223, KMI, WR1.102, WR1.107, and WR1.112. Boxed amino acids at the N-terminus correspond to possible primer-de-**Table 3a** Amino acid sequences of human anti-TPO antibody *IGHV chains* aligned with the closest putative germline genes. Designation of the complementarity determining regions (CDR) and framework regions (FR) are according to IMGT (Lefranc and Lefranc 2001; Lefranc et al. 1999). Only substituted amino acids are shown. Antibody sequences

rived sequences

10.113 10.1	Antibody designation	FR1-IMGT	CDR1-IMGT	FR2-IMGT	CDR2-IMGT	FR3 - IMGT	CDR3-IMGT FR4-IMGT	 E 8
1		(1–26)	(27-38)	(39-55)	(29-95)	(66-104)	(/TT-COT)	(63)
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1.00 1.00	T2.6	VQL-E	1		V-GY	TDT		S SS
100 100	T3.2	-E	H-S	B	G	NL		SS
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14	A13		S-SI-N	IBI	-HTR			SS
	A14		S-SI-N	BI	-HTR			33
B	A16		S-SI-N	BI	-HTR.	LSL		SS
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Table 3

FR2-IMGT CDR2-IMGT FR3-IMGT CDR3-IMGT FR4-IMGT (105-117) (118-129)	40 50 60 70 80 90 100 111 112 118 120 130 130 1	MHWARQA FOGIEMMOR INPNSOCT NYAQKRQ GRVJSTRDTSISTAYMELSRLRSDDTVVYYC AR NFYGLIDVWGGGTTVTUSS NYAQKRQ GRVJSTRDTSISTAYMELSRLRSDDTVVYYC AR NFWELLOW NYAL NFYGLIDVWGGGTTVTUSS NYAL NFWELLOW NATURAL NFSER G-M - A - AT - TS - KA - A - F GLGVG NYGLIDVWGGGTLVTVSS NYGLIDVWGGGTLTVTVSS NYGLIDVWGGGTLTVTVSS NYGLIDVWGGGTLTVTVSS NYGLIDVWGGGTLTVTVSS NYGLIDVWGGGTLTVTVSS NYGLIDVWGGGTLTA NYGLIDVWGGG	ISWNRQAPGGLEWMGG IIPIFGTA NYAQKFQ GRVTITADESTSTAYMEI.SSILRSEDTAVYYC AR	INWIRGATION GYAQKFQ GRUTMTRATELSSLRSEDTAVYYC AR LYGLDWAGQCTTUTUSS CGNAG CANDON MIPRISCATUT CGNAG CGNAG
	50			GYTFTSYD. INWVRQATG2GLEWMGM MNP EF — H— - N-H— - N-H— - N-H— - N-H— - N-HTF -
FR1-IMGT (1-26)	1 10 20	QVQLVQSQA. BVKKPGASVKVSCKAS (VQL BE L. R. R. R. VKL EE L. R. R. R. R. VKL EE L. R. R. R. R. VKL EE L. R. R. R. P. VKL EE L. R. R. R. R.	QVQLVQSGA.EVKRPGSSVKVSCKAS. (. 0. E	OVOLVOSGA. EVKKPGASVKVSCKAS (VQL)-E
Antibody designation		X07448 IGHV1-2*01 A2506372 L12061 WR4.10 L12067 WR4.25 L12070 WR4.25 L12070 WR4.22 L12071 WR4.22 L12071 WR4.32 L12077 WR4.32 L12077 WR4.32 L12077 WR4.34 L12077 WR4.34 L12077 WR4.34 L12077 WR4.35 L12078 WR4.35 L12107 TR1.21 L12107 TR1.22 Z15084 SP1.6	122582 IGHV1-69*01 AF206350 TF2.13 AF306351 TF2.3 AF306353 TF2.4 AF306354 TF3.12 AF306354 TF3.12 AF306355 TF3.12 AF306357 TF3.14 AF306357 TF3.14 AF306327 DN4 AL23827 ICA5 AL2392810 AL1 AL239810 AL1 AL299810 AL1 AL299815 TF3.14 AL299816 TF3.14 AL299817 TF3.14 AL299817 TF3.14 AL299817 TF3.14 AL299817 TF3.14	M99637 IGHV1-8*01 A2399831 T3.4 A73856 6F X73856 6F X98933 IGHV3-21*01 X98934 1266 X98935 1266 X98936 1266 X98936 1266 X98937 1266 X98937 1266 X98937 1267 X98937 126708 X98944 126709 X98945 1267015 X98945 1267015 X98946 1267015

Antibody designation	FR1-IMGT (1-26)	CDR1-IMGT (27-38)	FR2-IMGT (39-55)	CDR2-IMGT (56-65)	FR3 (66	FR3-IMGT (66-104)	CD (1)	CDR3-IMGT (105-117)	FR4-IMGT (118-129)
	1 10 20	30	40 50	09	70 80	90 100	110 111	112 118	130
M99660 IGHV3-23*01 X98959 131TP2 X98950 131TP5 X98960 131TP6 X98961 131TP6 X98961 131TP7 X98964 131TP15 WR1.223 WR1.223	EVQLLESGG. GIVQPGGSLRLSCAAS GFT	GETTESSYA.	MSWVRQAPGKGLEWVSA I SGSG "T"		YYADSVK, GRPTISRDNSKNTJYLQMNSIRAED S-E	YYADSVK, GRFTISRDNSKATLVLQMASLRAEDTAVYYCC SEVRRSRS. SEVVB-K-SSSSSSSSSSSSS	AKAGRILGAVIARGILIGAVIARGPIPYYARGPIPYYARGPIPYYARGPIPYYARGPIPYYARGPIPYYARGPIPYYIESBEIESBE	WYSLYYGFDVMOQGTTVTVGS YYALDVLMQGTTVTVGS YYALDVLMQGTTVTVGS YYALDVLMQGTTVTVSS YYALDVLMQGTTVTVSS YYALDVLMQGTTVTVSS YYALDVLMQGTTVTVSS YYALDVLMQGTTVTVSS YALDVLMQGTTVTVSS YALDVLMQGTTVTVSS	GTTVTVSS FTTVTVSS FTTVTVSS FTTVTVSS FTTVTVSS FTTVTVSS
M83134 IGHV3-30*01 AJ399808 A8 KM1	QVQLVBSGG.GVVQPGRSLRLSCAAS GFTFSSYA. EA-DAFT. QVKLLLEM,S	GFTFSSYA	MHWVRQAPGKGLEWVAV	ISYDGSNKSA-TKT	YYADSVK.GRFTISRDNS -FC	YYADSVK, GRFTI SRDNSKNTLYLQMNSLRAEDTAVYYC -F	C AR DGDPI	GYYFDYWGPGTLVTVSS	CTLVTVSS
M99675 IGHV3-48*01 X98963 131TP14	EVQLVESGG.GLVQPGGSLRLSCAAS GFT	GFTFSSYS	MNWVRQAPGKGLEWVSY ISSSSSTI	ISSSSSTI	YYADSVK.GRFTISRDNAKNSIYLQMNSI	YYADSVK.GRFTISRDNAKNSLYLQMNSLRAEDTAVYYC AR RDF	C AR AFSLRFS	YYYGMDVWGPGTTVTVSS	CTTVTVSS
M99679 IGHV3-53*01 L12090 TR1.3 L12092 TR1.5 L12111 TR1.32 X73853 2G4	EVQLVESGS.GLIQPGGSLRLSCAAS GFTVSSNY [WK]-E	GFTVSSNYIN-KLIN-KLLN-KL	MSWVRQAPGKGLEWVSV -T	IYSGGST -FTD-NP T-TD-T T-TD-T	YYADSVK, GRPTISRDNSKNTIYLQMNSIRAED	YYADSVK. GRFTISRDNSKNTLYLQMNSLRAEDTAVYYC AR	- KYQGTRS KSQGTRS KSQGTRS	YYYDDWGKGTTVIVYYYDDWGKGTYYYDDWGKYYYDDWGKLAHWGQGTLVSVSS	GTTVIV GT SYLVSVSS
M99682 IGHV3-64*01 AJ399809 A9 AJ399811 A12 AJ399836 T10	EVQLVESCG. GLVQPCGSLRLSCAAS GFTFSSYA	GFTFSSYA	MHWVRQAPGKGLEYVSA VYG -YS	ISSNGGST GH GHT	YYANSVK GRFTISRDNSKNTLYL S	YYANSVK.GRFTISRDNSKNTLYLQNGSLRAEDMAVYYC AR S	C AREWQLPNFEWQLPNF	YSYGMDVWGQGTLVTVSS YSYGMDVWGQGTLVTVSS GGYFGLDVWGHGTLVTVSS	KILVIVSS KILVIVSS KILVIVSS
Z14238 IGHV4-30-4*01 WR1.112	QVQLQESGP.GLVKPSQTLSLTCTVS GGSISSGDYY.,	GGSISSGDYY	WSWIRQPPGKGLEWIGY IYYSGST	IYYSGST	YYNPSLK.SRVTISVDTS	YYNPSIK.SRVTISVDTSKNQFSIKLSSVTAADTAVYYC AR A,	C AR 		
L10098 IGHV4-31*01 X73857 7F	QVQLQESGP.GLVKPSQTUSLTCTVS GSSISSGSYY WSWIRQHPGKGLEWIGY IYYSGST	GGSISSGGYY	WSWIRQHPGKGLEWIGY L-H	IYYSGST	YYNPSLK.SLVTISVDTS	YYNPSLK, SLVTISVDTSKNOFSLKLSSVTAADTAVYYC AR T, GRIENFGRAALFG	C AR GRAALFG	SESYPLDHWGQGTLVTVSS	STLVTVSS
M99686 IGHV5-51*01 AF306373 T2.5 AJ399826 B11	EVQLVQSGA.EVRKPGESLKISCKGS GYSFTSYW [VQ]-B-E-ESSS	GYSFTSYW K-D	IGWVRQMPGKGLEWMGI IYPGDSDTA	IYPGDSDT		RYSPSFO. GQVTISADKSISTAYLQWSSLKASDTAMYYC AR K	HRDTAILTGQ. SFGAFRH	KNYYYYGMDVWGQGTTVTVSS	GTLVTVSS GTLVTVSS

2001; Lefranc et al. 1999). Only substituted amino acids are shown. Antibody sequences were obtained from databases except antibodies WR1.223 and KM1. Boxed amino acids at the N-terminus correspond to possible primer-derived sequences **Table 4a** Amino acid sequences of human anti-TPO antibody *IGKV* chains aligned with the closest putative germline genes. Designation of the complementarity determining regions (CDR) and framework regions (FR) are according to IMGT (Lefranc and Lefranc

T FR4-IMGT (118-129)	120	.WTFGQGTKVEIKR .LTFGGGTKVEIKR .LTFGGGTKVEIKR	.LTFGPGTKVDI .LTFGGGTKVEI	.LTFGGGTKSEIK	. YTFGQGTKLEIKR	LIPPGGTKVEIKR LIPPGGTKVEIKR LIPPGGTKVEIKR LIPPGGTKVEIKR LIPPGGTKVEIKR LIPPGGTKVEIK	VLSFGGGTRLEIKR	.YTFGQGTK .YTFGQGTKLE .YTFGHFTK	.LIYNFGQGTKLEIKR SFGGGTQLTVLS LTFGGGTKLEIKR	PFTFGPGTKVDIK	GAFQQTKLEIKR LTFCQCTKLEIKR LTFCGCTKVEIKR LTFCGCTKVEIKR LTFCGCTKVEIKR
CDR3-IMGT (105-117)	110	QQANSFP 	QQFNNYP S	QKYNSAP -QV-HY	QQYNSYS YT-P	VOLINSYP 	QQANSFP GY-S	MQALQTP P P	QQRSNWP NSL R	QQYNNWP	QQYGSSP L
(66-104)	90 100	SGTDFTLTISSLQPEDFATYYC	SLESGVP.SRESGSGSGTDFTLTISSLQPEDFATYYC T	TLQSGVP.SRFSGSGSGTDFTLTISSLQPEDVATYYC	SGTEFTLTISSLOPDDFATYYC		.SGTDFTLTISSLQPEDFATYYC	SGTDFTLKISRVEAEDVGVYYC	SGTDFTLTISSLEPEDFAVYYC	TRATGIP.ARFSGSGSGTEFTLTISSLQSEDFAVYYC	SGTDFTLTISRLEPEDPAVYYC
	70 80	SLOSGVP.SRFSGSG.	SLESGVP.SRFSGSC TS	TLQSGVP.SRFSGSC	SLESGVP.SRFSGSG. H-HD	TLQSGVP. SRFSGSG	SLQSGVP, SRFSGSG.	NRASGVP. DRFSGSG.	NRATGIP.ARFSGSG.	TRATGIP.ARFSGSC	SRATGIP DRESGSG
CDR2-IMGT (56-65)	09	AAS	DAS	AAS	DAS	AAAS.	AAS	LGS	DAS	GAS	GAS.
FR2-IMGT (39-55)	40 50	LAWYQQKPGKAPKLLIY	LA*YQQKPGKAPKLLIY W	LAWYQQKPGKVPKLLIY	LAWYQQKPGKAPKLLIY	LAMYQQKPGKAPKLLIY	LAWYQQKPGKAPKLLIY	LDWYLQKPGQSPQLLIY	LAWYQQKPGQAPRLLIY	LAWYQQKPGQAPRLLIY	LAWYQXREGAPRLLIY
CDR1-IMGT (27-38)	30	QGISSW -A-YT	QGISSA	QGI SNY	QSISSW	OSTSSY	QGISSW	QSLLHSNGYNY.	QSVSSY	OSVSSN	OSVSSSY
FR1-IMGT (1-26)	1 10 20	DIOMTQSPSSVSASVGDRVTITCRAS C	ALQLTQSPSSLSASVGDRVTITCRAS C ELVM	DIQMTQSPSSLSASVGDRVTITCRAS C	DIQMTQSPSTLSASVGDRVTITCRAS C E-VL-HPI	DIQLTQSPSFLSASVGDRVTITCRAS C	DIQMTQSPSSVSASVGDRVTITCRAS C	DIVMTQSPLSLPVTPGEPASISCRSS C EL	EIVLTQSPATLSLSFGERATLSCRAS (E	EIVMTQSPATLSVSPGERATLSCRAS (EIVLTQSPGTLSLSPGERATLSCRAS (E
Antibody designation		V01577 IGKVJ-12*01 AF306360 TF2.4 AF306389 T3.4 X98967 126D	Z00006 IGKV1-13*02(F) L12089 TR1.13 L12099 TR1.9	X63398 IGKV1-27*01 X98976 126T09	Z00001 IGKV1-5*01 AJ399874 T5	200013 IGKVJ-9*01 X98965 126C X98968 126F X98969 126G X98970 126H X98971 126TO X98977 126TOI X98978 126TPI X98978 126TPI X98981 126TPI X98981 126TPI X98981 126TPI X98981 126TPI	X17263 IGKV1D-12*01 AJ399871 B7	X12691 IGKVZD-28*01 L12095 TR1.6 L12097 TR1.8 L12114 TR1.37	X01668 IGKV3-11*01 AF306380 T2.2 AJ399872 T2 AJ399878 T10	M23090 IGKV3-15*01 X98990 131TP14	X12686 IGKV3-20*01 AF306359 TF2.3 AF906363 TF3.19 AF306364 TF3.14 AF306379 T2.11 AF306386 T3.15 AF30638 GRV4-1*01 RM1.223

Table 4b

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CDR3-IMGT (105-117)		OOSYSTP		:	: ,	N	N	F		:						:			:	:	N	FT	FT				· E		L		I		 	• E			G-D	I	D	A-T	A	T	N		:	0			T				•				I	1	1 1					:		·	D-V	L			:	-S	D	
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2001; Lefranc et al. 1999). Only substituted amino acids are shown. Antibody sequences were obtained from databases except antibodies WR1.102, WR1.107, and WR1.112. Boxed amino acids at the N-terminus correspond to possible primer-derived sequences Table 5 Amino acid sequences of human anti-TPO antibody IGLV chains aligned with the closest putative germline genes. Designation of the complementarity determining regions (CDR) and framework regions (FR) are according to IMGT (Lefranc and Lefranc

GT FR4-IMGT 7) (118-129)	120	VPGGGTKLTVLG	VFGTGTKVDIKSVFGTGTKVDIKRVFGGGTKVDIKRVFGGGTKVTULG I.VFGGGTKVTULG I.VFGGGTKVTULGVFGGGTKVTULGVFGGGTKVTULGVFGGGTKVTULGVFGGGTKVTULGVFGGGTKVTULGVFGGGTKVTULGVFGGGTKLTTULGVFGGGTKVDIKRVFGGGTKUTULGVFGGGTKUTULGGVFGGTKUTULGGVFGGTTKUTULGGVFGGTTKUTULGGVFGGTTKUTULGGVFGGTTKUTULGGVFGGTTKUTU	FOGGTKLEIKR VFGGGTKLIVLG	VFGSGTKLEIKR	VEGGGTOLITYLS
CDR3-IMGT (105-117)	OSYDSS P	C AAWDDSLNGSDSD AAWDDSLSG	C STADSSLSA C CSKAAGNTY 		C SSYAGSNNF FI- C QVWDSSSDH RN-	C QSADSSGTYYY C LLYYGGAQVHPR.
FR3-IMGT (66-104)	90 100 	QRPSGVP. DRFSGSK SGTSASLAI SGLQSEDEADYYC	SGTSATLGITGLOTGDEADYYC -E	NRPSGVS. NRFSGSK. SGNTASITI SGLQAEDEADYYC	. SGNTASLITVSGLQAEDEADYYC C . SGNTATLITSRVEAGDEADYYC 	ERPSGIP ERFSGSS . SGTTVTLTISGVQAEDEADYYC
	70 80	ORPSGVP. DRFSGSK	KRPSGIP DRFSGSK E	. NRPSGVS.NRFSGSK	. KRPSGVP. DRFSGSKNA DRPSGIP. ERFSGSN A	. ERPSGIP ERFSGSS
CDR2-IMGT (56-65)	60 60 60 60 60 60 60 60 60 60 60 60 60 6	SNIN GKS			EVS	7 KDS
FR2-IMGT (39-55)	VHWYQQLPGTAPKLLIY	VMMYQQLPGTAPKLLIYC	VSWYQQLPGTAPKLLITY	VSWYQQHPGKAPKLMIY	VSWYQQHPGKAPKLMIYYI VHWYQQKPGQAPVLVIY	AYMYQQKPQQAPVIJVIY -H
CDR1-IMGT (27-38)	30 	SGS SSNIGSNT	SGS SSNIGMNYS-K-KS-K-		TGT SSDVGGYNY TVD GGN NIGSKS	SGD ALPKQY
FR1-IMGT (1-26)	QSVLTQPPS.VSGAPQRVTISCC 	QSVJJQPPS.ASGTPGQRVTISC -PSSXB QSVJJQPPS.ASGTPGQRVTISCV	QSVLTQPPS . VSAAPQQKVT1SC	QSALTQPAS. VSGSPQQSITISCTGT	QSALTQPPS. ASGSPQSVTISCTGT -PV SYVLTQPPS.VSVAPGKTARITCGGN EL-VAQT-SD	SYELMOPPS VSVSPOGTARITCSGD
Antibody designation	M94116 IGIV1-40*01 AJ399845 A6 AJ399849 A10 AJ399852 A13 AJ399856 B1 AJ399867 T7 AJ399869 T12	Z73654 IGLV1-44*01 AJ399847 A8 AJ399855 A17 D87016 IGLV1-47*02 AJ238330 ICA5	273661 IGLV1-51*01 AJ238329 AJ398840 AJ399841 AJ399842 AJ399842 AJ399844 AJ399844 AJ399850 AJ399850 AJ399851 AJ399854 AJ399859 AJ399860 AJ399862 AJ399862 AJ399862 AJ399864	Z73664 IGLV2-14*01 AJ399843 AJ399863 B10 WRI.102	X97462 IGLV2-8*01 AJ399866 T4 X71966 IGLV3-21*01 U09085 TR1.41	X97474 IGLV3-25*01 WR1.107 WR1.112 X14614 IGLV7-43*01 AJ399857 B3

Correlation between Ig gene usage and TPO-specific antibody epitopes

Pairing of one defined heavy chain with different light chains does not alter antigen binding (Burton and Barbas 1992, 1994). This observation strongly suggests that the heavy chain initiates the formation of the antigen/antibody complex and thereby provides the specificity of the interaction, whereas its light chain counterpart stabilizes the interaction with subsequent affinity modulation (Noel et al. 1996). Such an effect of the anti-TPO aAb light chain on affinity is less conclusive, since neither IGKV nor IGLV gene usage of anti-TPO aAb has been shown to modulate antigen affinity (Chapal et al. 2000, 2001; McIntosh et al. 1997; Portolano et al. 1991, 1992, 1993b). On the other hand, several groups have pointed out that domain A of the TPO immunodominant region (IDR/A) is preferentially recognized by TPO-specific aAb with the IGKV1-39 light chain, whereas TPOspecific aAb showing other IGKV light chains map in domain B of the IDR (IDR/B) (Table 1, consisting of parts a, b and c) (Chazenbalk et al. 1993; Costante et al. 1994; Guo et al. 1998; Jaume et al. 1996, 1997; McIntosh et al. 1997; Portolano et al. 1995). This IDR/B has been at least partially identified even though the location of the IDR on the TPO molecule is still under debate. Region 713-721 is located on the C-terminal myeloperoxidase-like domain of the TPO molecule; this region, recognized by murine Mab 47/C21 antibody (Finke et al. 1991; Libert et al. 1991) and by serum polyclonal TPO aAb (Libert et al. 1991; Ruf et al. 1989), was initially thought to be outside the IDR (Chazenbalk et al. 1993). Furthermore, mutations in the 713–721 region do not affect the recognition of aAb directed against IDR (Nishikawa et al. 1996). On the other hand, high concentrations of IDR/B-specific aAb TR1.9 inhibited the binding of Mab47/C21 to TPO (Guo et al. 1998) and mapped an epitope comprising amino acid residue K713 (Guo et al. 2001), suggesting that region 713–721 is located on the fringe of an IDR. The crystal structure of the Fab TR1.9 has been solved (Chacko et al. 1996), but in the absence of the three-dimensional structure for the complex of TR1.9 with TPO, it is difficult to determine the structural details of the binding.

The role IGLV genes play in affecting anti-TPO specificity remains to be elucidated. The initially described λ -derived anti-TPO aAb had low affinity and were directed against TPO-IDR/B (Portolano et al. 1995; Prummel et al. 1994b). In contrast, some of our λ -derived aAb demonstrated high affinity to TPO and inhibited the binding of a majority of the serum aAb to TPO (Bresson et al. 2001; Chapal et al. 2001), suggesting that these aAb recognized the IDR (defined by epitope mapping using BIACORE as regions II, VI, and VIII) (Table 1, consisting of parts a, b and c). Future studies involving λ -derived aAb such as T13/VI, B4/VIII, or ICA5/II and Fab defining IDR/A and /B (WR1-7, SP1-4, TR1-8, and TR1-9) could shed new light on the epitope specificity and gene usage of these aAb that recognize IDR.

Recently, Pichurin et al. (2001) produced and characterized human recombinant aAb by phage display technology binding outside the TPO-IDR (defined as non-IDR). All these heavy chains are encoded by IGHV1–69, with an extremely long CDR3, and paired with different types of light chains, suggesting that non-IDR specificity is determined primarily by a common heavy chain. Interestingly, almost all IDR-specific aAb obtained in the same experiment use IGHV1-2 and IGHV1-3, as is also the case for a majority of the IDR aAb previously described (Table 1, consisting of parts a, b and c). Does IGHV1-2 or IGHV1-3 gene usage reflect a particular TPO-IDR specificity of recombinant aAb? Even though the methodologies used to define epitope recognition of anti-TPO recombinant aAb are different, these results reveal the difficulty of correlating gene usage with epitope recognition of TPO-specific aAb.

Conclusion

Several laboratories have produced and characterized numerous human anti-TPO aAb, leading to an enlarged autoantibody repertoire. Analysis of these antibodies using the IMGT database (Giudicelli et al. 1997; Lefranc 2001; Lefranc and Lefranc 2001; Lefranc et al. 1999) reveals several characteristics of the TPO-specific aAb repertoire: (1) a restriction in the IGV gene usage to generate anti-TPO aAb in AITD, (2) a VDJ recombination process using preferentially inverted D genes, (3) limited somatic mutations of J proximal light chain genes suggesting a defect in receptor editing in AITD, and (4) presence of certain somatic mutations systematically in the anti-TPO aAb repertoire. The annotations described in this paper and the protein display will soon be available as a new specialized IMGT page on human anti-TPO aAb genes. This page will evolve with time and integrate all the sequences devoted to autoantibodies that are published in the future.

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