Supporting information for Hoyer *et al.* (2002) *Proc. Natl. Acad. Sci. USA*, 10.1073/pnas.212410199

Mouse	Ν	D_{H}	Ν	$ m J_{H}$	% clones mutated	J_H	D_H		
K7	g	TCTACTATGATTAC	ctacggctattggtgtg	TACTATGCTATGGACTACT	23.3	$J_{\rm H}4$	DSP2.2		
				GGGGTC <u>G</u> A <u>T</u> GAAC <u>AC</u> CA	(7/30)				
				G <u>C</u> CACCGTC <u>A</u> CCTCA					
P2	ttat	TACTA	cgatggtagctaccc	TACTGGTACTTCGATGTCT	NM	$J_{\rm H} 1$	DSP2.n		
				GGGGCGCAGGGACCAC	(2/2)				
				GGTCACCGT <u>C</u> TCCTCA					
K8	gttta	ttta ATTAC tacggtag TA		TA <u>G</u> TTTGACTACTGGGGC	20.0	$J_{\rm H}2$	DSP2.2		
				CAAGGCACCACTCTCACA	(4/20)				
				GTCTCC <u>A</u> CA					
K7 J (1) (2) (3)	H4: 7	FACTATGCTATGG	ACTACTGGGG	GTC <u>A</u> A <u>G</u> GAAC <u>CT</u> CAG <u>T</u> C G T	CACCGTC <u>T</u> CCT	СА			
(3)		A C							
(4)									
(5)				C					
(6)				С					
(7)					А				

Table 4. Somatic hyp	permutation of J _H I	region in DL	BCL, BLL,	and FBL
----------------------	---------------------------------	--------------	-----------	---------

Somatic mutations detected in BLL (K7), FBL (P2), and DLBCL (K8) tumor lines derived from *TCL1* transgenic mice. N-region additions flanking individual D_H gene segments are listed in lower case letters. The total variations detected from the germ-line J_H sequence in each tumor are underlined. Shown below the table are the positions of 7 mutations detected from 30 J_H region segments sequenced from tumor K7. All mutations were substitutions rather than insertions or deletions. Boldface underlined positions indicate transition-type mutations. The occurrence of four distinct transitions (Pyr \rightarrow Pyr and Pur \rightarrow Pur) and three independent transversions (Pyr \rightarrow Pur and Pur \rightarrow Pyr) is compatible with ongoing somatic hypermutation in this line. Mutations within

the $D_{\rm H}$ segments, 5' and 3' regions flanking the $DJ_{\rm H}$ segments were not observed. NM, not meaningful.