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Induction of pre-BCR expression on <i>in vitro</i> cultured pre-B cells ^a																
	Day 0				Day 1			Day 2			Day 3			Day 4		
SL^+ cells	77			79			71			<5			<5			
μH^+SL^+ cells	0			40			63			<5			<5			
% c-kit ⁺ cells	44			25			9			5			2			
% CD25 ⁺ cells	1			7			30		55			62				
large cells	45			45			58		46			5				
	RNA signals per nucleus ^b															
	1	2	4	1	2	4	1	2	4	1	2	4	1	2	4	
CD45																
PCNA ^{- c}	3	39	2	3	42	2	2	45	2	7	41	5	4	64	3	
\mathbf{PCNA}^+	0	4	15	0	3	18	1	7	27	2	7	15	0	1	8	
Total	3	43	17	3	45	20	3	52	29	9	48	20	4	65	11	
λ5																
PCNA ⁻	1	37	1	6	42	4	5	34	1	14	13	1	5	1	0	
$PCNA^+$	1	6	17	1	2	14	1	11	13	0	15	5	1	3	0	
Total	2	43	18	7	44	18	6	45	14	14	28	6	6	4	0	

Transcription pattern of the $\lambda 5$ gene after induction of pre-BCR expression

^a CD19⁺ BM cells from tet- μ H mice were purified and cultured *in vitro* for four days. Each day samples were analyzed by: FACS, to determine the percentages of surface SL⁺, μ H⁺, i.e. pre-BCR⁺ cells and cell size and by immuno-RNA-FISH to investigate the transcriptional status of the CD45 and λ 5 genes. ^b Nuclei contained either 1, 2 or 4 discrete RNA signals. ^c Percentages of the total nuclei with either 1, 2 or 4 signals staining as PCNA⁻ or PCNA⁺ (S phase) are shown. The data shown is from one representative experiment. The number of nuclei counted from each day and for each probe was >150.