

Supplementary Table 4

Rearrangement	Representative Accession	Coordinates		Repeat Element at Breakpoint (BP)		Species with identical BP <sup>†</sup> in available sequence	State of element at breakpoint: Crossing or Truncated	Original (vs. Derived) State	Rearrangement Type	Size <sup>‡</sup> (bp)	Inferred Mechanism	
		Left	Right	Left	Right							
1	PAB_AC188481.3	120780	124140	3360	L1M1	L1M1	PAB, GGO	C	C	0		
	MMU_AC150715.2	33597			L1M1/L1M1*		MMU	T/T				
2	PAB_AC188481.3	128077			L1M5		PAB, GGO	C	O			
	MMU_AC148535.2	36394	42556	6162	L1M3	L1M3	MMU	T	T	Insertion	5466 LINE RT	
3	HSA_HG18	100212678			L1M3		HSA, PTR 15q	C	O			
	MMU_AC148535.2	36394	42556	6162	L1M3	L1M3	MMU	T	T	Insertion	6162 LINE RT	
4	HSA_HG18	100213018	100222547	9520	LOR1-int	L2	HSA, PTR 15q	C	C	O		
	MMU_AC150715.2	134811			LOR1-int	L2	MMU	T/T		Deletion	9531 NHEJ	
5	HSA_HG18	100239257	100242320	3063	L1PA7	L1PA7	HSA, PTR 15q, PTR 4q	C	C	O		
	MMU_AC148535.2	61701			L1PA7/L1PA7*		MMU	T/T		Deletion	3063 NHEJ	
6	HSA_HG18	100242452	100245066	2614	L1PA7	***	HSA, PTR 15q, PTR 4q	C	(C)	O		
	MMU_AC148535.2	61888			L1PA7/THEID-int		MMU	T/T		Deletion	≥2614 NHEJ	
7	HSA_HG18	100253145	100258937	5792	-	L1P4	HSA, PTR 4q	-	C	O		
	MMU_AC148535.2	67357			-	L1P4	MMU	-	T		Deletion	≥5792 NHEJ
8	PAB_AC18330.3	29985			L2C		PAB, HSA, PTR 4q	C	O			
	MMU_AC148535.2	70211	74131	3920	L2C	L2C	MMU	T	T	Insertion	3920 LINE RT	
9	PTR4q_AC183669.3	63635	68394	4859	L2	L1PA3	PTR4q	C	C	O		
	MMU_AC148535.2	145306			L2/L1PA3		MMU	T/T		Deletion	4859 NHEJ	
10	PAB_AC18330.3	165419	169603	4184	-	-	PAB					
	MMU_AC148620.2	73570			-	-	MMU			Indel	4184 NHEJ	
11	PAB_AC18330.3	176355	181239	4884	-	L1P4	PAB		C	O		
	MMU_AC148620.2	81628			-	L1P4	MMU	-	T		Deletion	4884 NHEJ
12	MMU_AC150715.2	52580	55392	2812	-	Tigger3	MMU		C	O		
	PAB_AC188481.3	137368			-	Tigger3	PAB, GGO	-	T		Deletion	2812 NHEJ
13	HSA_HG18	100200440	100213892	13252	L1M1	LTR57-int	HSA, PTR 15q, PTR 4q	C	C	O		
	PAB_AC18330.3	136240			L1M1/LTR57-int		PAB	T/T			Deletion	13252 NHEJ
14	HSA_HG18	100219570	100245066	25496	MLTH1	***	HSA, PTR 15q, PTR 4q	C	(C)	O		
	PAB_AC18330.3	18965			MLTH1/THEID-int		PAB	T/T			Deletion	≥25496 NHEJ
15	MMU_AC148535.2	67051			AluY		MMU	C	O			
	HSA_HG18	100248092	100252855	4763	AluY	AluY	HSA, PAB <sup>†</sup> , PTR 15q, PTR 4q	T	T		Insertion	4763 LINE RT
16	HSA_HG18	100256005	100260556	3651	L1MAB/L1ME3B		HSA, PTR 4q	C	C	O		
	PAB_AC18330.3	28542			L1MAB/L1ME3B		PAB	T/T			Deletion	3651 NHEJ
17	MMU_AC149242.2	36929	57410	20481	L1M4C	-	MMU, HSA, PTR 4q	C	O			
	PAB_AC18330.3	33511			L1M4C	-	PAB	T/-			Deletion	20480 NHEJ
18	Inferred	PAB_AC18330.3	33511	39774	6263	L1M4C	-	PAB	C	O		
	PTR4q_AC183669.3	62153	72625	10472	-	AluSq	PTR4q, MMU	T			Insertion	6263 LINE RT
19	PAB_AC18330.3	88115			-AluSq		PAB	-	C	O		
	MMU_AC148620.2	52576	58771	6195	L1M1C1	-	MMU	C	O			
20	PAB_AC18330.3	149153			L1MC1/-		PAB, PTR 4q	T/-			Deletion	6195 NHEJ
	MMU_AC150715.2	16098	68983	52885	-	-	MMU, PAB, GGO			O		
21	HSA_HG18	100155912			-	-	HSA, PTR 15q				Deletion	52885 NHEJ
	PAB_AC18330.3	100155912			-	-	MMU, PAB			O		
22	MMU_AC148535.2	†††	64830		-	-	HSA, PTR 15q, PTR 4q			O		
	HSA_HG18	100245068			-	-	MMU, PAB				Deletion	≥2995 NHEJ
23	MMU_AC149242.2	42064	50763	8699	-	L1PB	MMU		C	O		
	HSA_HG18	100259229			-	L1PB	HSA	-	T		Deletion	8700 NHEJ
24	MMU_AC148535.2	94830			L1MCA4		MMU	C	O			
	HSA_HG18	100273152	100276019	2867	L1MCA4	L1MC4A	HSA, PTR 4q	T	T		Insertion	2867 LINE RT
25	MMU_AC148535.2	127784	††††		LTR16C	AluSq/x	MMU, PAB 15q, PTR 4q	C	C	O		
	HSA_HG18	100307958			LTR16C/AluSq/x	HSA	T/T				Translocation-Mediated Deletion	-160000 NHEJ
26	HSA_HG18	100258107			AT-rich		HSA			O		
	PTR4q_AC150448.2	94792	98005	3213	AT-rich	AT-rich	PTR4q				Insertion	3213 LINE RT
27	MMU_AC148535.2	83068	89013	5945	-	L1PA5	MMU		C	O		
	PTR4q_AC183669.3	1661			-L1PA5		PTR4q	T/T			Deletion	5945 NHEJ
28	PAB_AC18330.3	104672			-	-	MMU, PAB			O		
	PTR4q_AC183669.3	89614	92759	3145	-	-	PTR4q				Insertion	3145 LINE RT
29	MMU_AC148620.2	33295			-	-	MMU, PAB			O		
	PTR4q_AC183669.3	112843	117305	4462	-	-	PTR4q				Insertion	4462 SUZ12 RT

<sup>†</sup> Chromosome locations are MMU 7q, PAB 15q, HSA 15q, and GGO 15q unless denoted otherwise.<sup>‡</sup> When a deletion size is ≥ a value, this deletion encompasses or overlaps deletion(s) in all genomes and its true size cannot be determined.<sup>\*</sup> These two Line elements are joined at nonhomologous sites<sup>\*\*</sup> Deletion is internal to L1PA7.<sup>\*\*\*</sup> THEID-int is truncated or missing from all genomes analyzed, but it can be inferred to have crossed the righthand breakpoint. It is missing from HSA due to rearrangement #22.<sup>††</sup> Repeat masker incorrectly identifies the portion of the AluY to the right of this insertion as an AluSq. PAB lacks a portion of the insertion and the AluY sequence to the right of the breakpoint due to a sequencing gap.<sup>†††</sup> Insertion is at the right edge of the truncated L1M4C created by rearrangement #17; or, alternatively, rearrangement #17 occurred subsequent to this insertion and deleted part of it.<sup>††††</sup> MMU & PAB are missing sequence for the lefthand breakpoint due to additional deletions (rearrangements 6 & 14).<sup>†††††</sup> Several human subtelomeres and the 2q fusion site retain block 2 sequence as an intact Alu element crossing the other translocation breakpoint healed by NHEJ to create this join (Linardopoulou et al., 2005).

There is ~160 kb from the breakpoint of HSA15q and MMU7 sequence homology to the fusion site.