## Supplemental Fig. S1: Short Tandem Repeat (STR) Analysis

	Xenografts: 736 777 9216R 9218R 8R and 8L		<b>22R</b> v1			CWR-R1	
locus	alleles	frequency, 1 in	alleles	frequency, 1 in	alleles	frequency, 1 in	
D5S818*	11, 12	3.91	11, <mark>13</mark>	9.50	11, 12	3.91	
D7S820 <sup>†</sup>	9,10,11	11.60	9,10,11	11.60	9,10, <mark>12</mark>	11.60	
D13S317	8, 12	15.94	<mark>9</mark> , 12	26.39	8, 12	15.94	
D16S539	12, 12	9.77	12, 12	9.77	12, 12	9.77	
TH01	6, 9.3	7.39	6, 9.3	7.39	6, 9.3	7.39	
TPOX	8, 8	4.11	8, 8	4.11	8, 8	4.11	
CSF1PO	10, 11	5.86	10, 11	5.86	10, 11	5.86	
Random Match Probability, 1 in :		1,259,877		5,062,112		1,259,877	
Random chance that the xenografts and 22Rv1 or R1 would have the same allele pattern:1.6 X 10 <sup>-13</sup> 6.3 X 10 <sup>-13</sup>							

\*STR analysis at locus D5S818 was not determined for sample 736. <sup>†</sup>Frequency for D7S820 was determined using only the 9 and 10 allele frequencies.

## Supplemental Fig. S2: Primers used for PCR



Primer:	Sequence 5	i' nucleotide position#	1	2	Х
U3f	5'-GTTTAATTAAAGAATAAGGCTGAATAAC-3'	7826	+	_	+
13r	5'-ATGTCTTCTAACAGCTTTTTGGACACG-3'	5108	+	-	+
18f	5'-GGCAGAGGATGAGCAGAGAGAGAG-3'	1974	- !	+	+
24f	5'-AAGAAAAGGGACACTGGGCTAAGG-3'	2129	+	+	+
lbf	5'-AGGCATTCCCGACCAAGCG-3'	5048	+	_	+
8r	5'-CTGGATGCTACCGGAGCCC-3'	6284	+	-	+
8f	5'-GGGCTCCGGTAGCATCCAG-3'	6266	+	-	+
U3r	5'-CCCCTTTTTTATAGGGCTAGGAC-3'	8096	- !	_	+
8fsa	5'-CCTGTTTTGATTCCTCAGTGGG-3'	6247	+	_	+
U5rsa	5'-TCTGAGGAGACCCTCCCAAGG-3'	112	+	+	+
Flf	5'-GCGCCAGTCATCCGATAGACTGAGTCGCCCGGGTAC	ч -			
	CCGTGTTCCCAATAAAGCC-3'	1	-	+	+
Flr	5'-GCCGACGCCAAGGTCCCAGTTTTTGCGTTAGGACGC	ч -			
	CTTTGGCGTAGCCCTGCTTCTCGTCGACAAAGAGTTC	-3′ 3754	_	+	+
F2f	5'-GAACTCTTTGTCGACGAGAAGCAGGGCTACGCCAAA	7			
	GGCGTCCTAACGCAAAAACTGGGACCTTGGCGTCGGC	-3′ 3682	-	+	+
F2r	5'-TTGCAAACAGCAAAAGGCTTTATTGGGAACACGGGI	1			
	ACCCGGGCGACTCAGTCTATCGGATGACTGGC-3'	8185	+	-	+
XmU3f	5'-GTCCTAGCCCTATAAAAAGGGGG-3'	8074	-	-	+
G2f	5'-CCCTTATACCCGCTCACCAAGAC-3'	3550	+	_	_
G3r	5'-TGGAGCTGCTCAAATTGTTGGG-3'	7204	+	_	_
Xelf	5'-GTGGCCCAATCAGTAAGTCCGAG-3'	410	+	-	-
Xe2f	5'-CACTCCCTTGAGTCTGACCCTTG-3'	630	+	_	_
GAGr	5'-TCCCCCAACAAAGCCACTCCA-3'	473	_	+	+
midX1f	5'-TTGTCGACGAGAAGCAGGGC-3'	3689	-	+	+
midX2r	5'-TGCGTTAGGACGCCTTTGGC-3'	3731	_	+	+
envOUT1f	5'-CTGACCCAACAGTATCACCAACTC-3'	7629	+	+	+
C12 1f	5'-TGCTGGACAGAATCTCTGGTCTCT-3'	Ch12	_	_	_
C12 4r	5'-GATACTCAAGTGGTTCCCACCC-3'	Ch12	_	_	_
129 <sup>_</sup> 1r	5'-GCGGTTTCGGCGTAAAACCGAAAGCA-3'	537	_	+	+
4847r	5'-CTTTGCTGGCATTTACTTGGGCA-3'	4909	+	- !	+
4257f	5'-GATGGCAGAAGGTAAGAAGCTAAATGTTTA-3'	4296	+	- !	+
M19f	5'-TGGCCTTACTGAAAGCTCTCTTCC-3'	4436	_	+	_

#: 5'nucleotide position relative to XMRV-22Rv1 (Acc. FN692043).

-!: primer has mismatches with the proviral genome, but was able to amplify the proviral sequence.

## SUPPLEMENTAL FIG. S3: Location and Numbers of Cloned PCR Products Used for Sequencing Analysis





## Supplemental Figure S4A: Phylogenetic analysis of full-length genomes

Supplemental Figure S4B: Phylogenetic analysis of XMRV and PreXMRV-1 homology



Supplemental Figure 4C: Phylogenetic analysis of XMRV and PreXMRV-2 homology



Supplemental Figure 4D: Phylogenetic analysis of the U3-R region



Supplemental Fig. S5: Predicted Regions of RT Template Switching Events and XMRV Sequence Diversity

Α

B

**Predicted Regions of RT Template Switching Events** 



**XMRV Sequence Diversity** 

