

SUPPLEMENTAL MATERIAL

Helmink et al., <http://www.jem.org/cgi/content/full/jem.20081326/DC1>

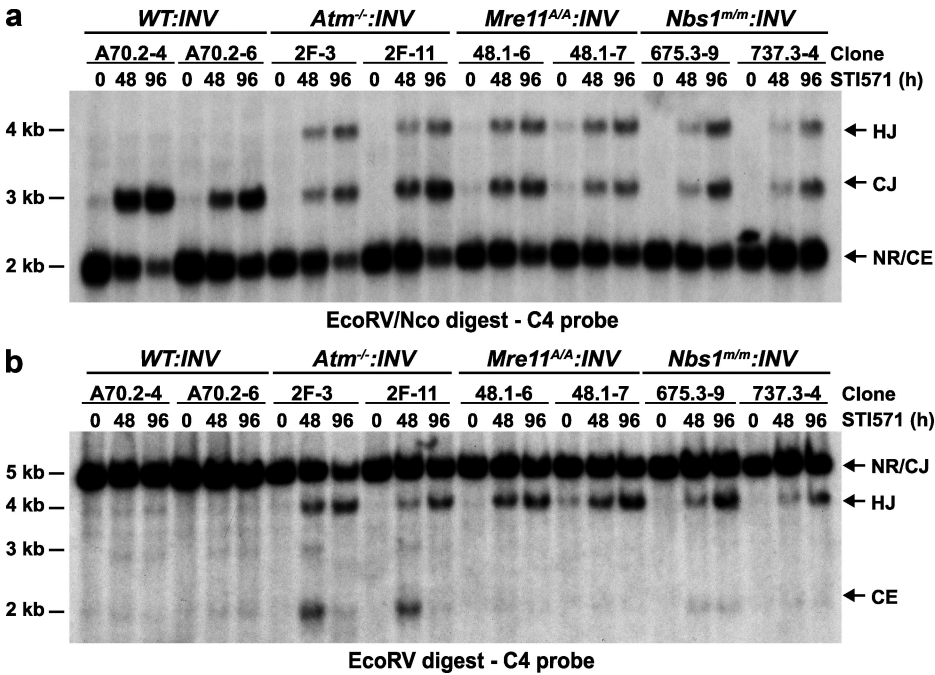


Figure S1. Southern blot analyses of pMX-INV rearrangement on additional abl pre-B cell clones. Analyses were carried out as described in Fig. 1 b (a) and Fig. 1 c (b) on additional abl pre-B cell clones not shown in Fig. 1.

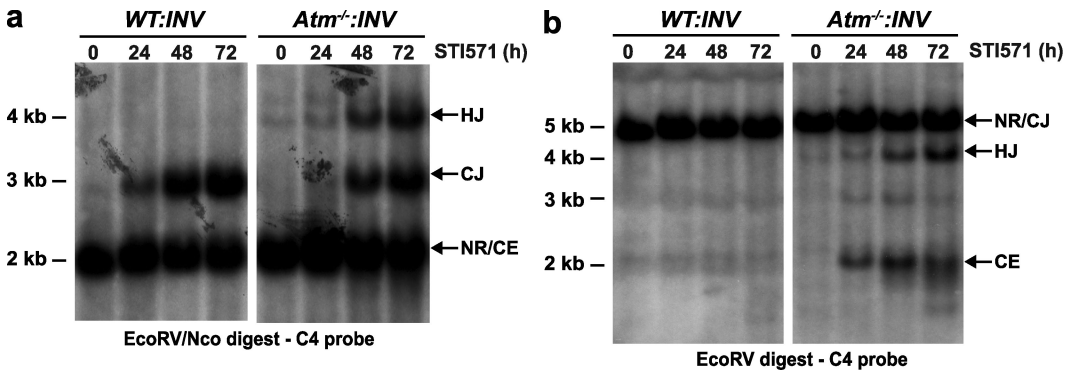


Figure S2. Longer exposure of the *WT:INV* and *Atm<sup>-/-</sup>:INV* Southern blot analyses. The lighter exposure is shown in Fig. 1 d (a) and Fig. 1 e (b).

[illegible]

Figure S3. Sequences of pMX-INV coding joints from *WT:INV*, *Atm<sup>-/-</sup>:INV*, *Mre11<sup>A/A</sup>:INV*, and *Nbs1<sup>m/m</sup>:INV* abl pre-B cells. N and P nucleotides are indicated, as are potential microhomologies (red). Complete coding end sequence is noted at the top of the figure.

pMX-DEL <sup>SJ</sup> 5' SE		pMX-DEL <sup>SJ</sup> 3' SE	
ttttgttccagtcctgtagcactgtg		cacagtggtagtactccactgtctggc	
<b>a <i>wt</i></b>			
pMX-DEL <sup>SJ</sup> 5' SE	N	pMX-DEL <sup>SJ</sup> 3' SE	Frequency
ttttgttccagtcctgtagcactgtg	0	cacagtggtagtactccactgtctggc	56
ttttgttccagtcctgtagcactgtg	GAG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	AG	cagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	GGCACAGTG	ggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	TGTG	ggtagtactccactgtctggc	1
			60
<b>b <i>Atm</i><sup>-/-</sup></b>			
pMX-DEL <sup>SJ</sup> 5' SE	N	pMX-DEL <sup>SJ</sup> 3' SE	Frequency
ttttgttccagtcctgtagcactgtg	0	cacagtggtagtactccactgtctggc	69
ttttgttccagtcctgtagcactgtg	GGG	acagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	G	cacagtggtagtactccactgtctggc	1
			71
<b>C <i>Mre11</i><sup>A/A</sup></b>			
pMX-DEL <sup>SJ</sup> 5' SE	N	pMX-DEL <sup>SJ</sup> 3' SE	Frequency
ttttgttccagtcctgtagcactgtg	0	cacagtggtagtactccactgtctggc	84
ttttgttccagtcctgtagcactgtg	AG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcac	CGTG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	0	agtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	0	cagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactg	CG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	GA	cacagtggtagtactccactgtctggc	1
			90
<b>d <i>Nbs</i><sup>m/m</sup></b>			
pMX-DEL <sup>SJ</sup> 5' SE	N	pMX-DEL <sup>SJ</sup> 3' SE	Frequency
ttttgttccagtcctgtagcactgtg		cacagtggtagtactccactgtctggc	61
ttttgttccagtcctgtagcactgtg	A	agtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	GCC	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactg	AGCA	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	GCCTG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	GTCCACG	tggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	G	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	CC	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	AT	cacagtggtagtactccactgtctggc	1
-67	CTCTGTTCCACTGTG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	A	gtctggc	1
ttttgttccagtcctgtagcactgt	CC	cacagtggtagtactccactgtctggc	1
-67	TCCGCCCCCCCCCTGTG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	AAGGG	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	CTC	cacagtggtagtactccactgtctggc	1
ttttgttccagtcctgtagcactgtg	CGCAC	tggtagtactccactgtctggc	1
			76

Figure S4. Sequences of pMX-DEL<sup>SJ</sup> signal joints from *WT:DEL<sup>SJ</sup>*, *Atm*<sup>-/-</sup>:*DEL<sup>SJ</sup>*, *Mre11*<sup>A/A</sup>:*DEL<sup>SJ</sup>*, and *Nbs1*<sup>m/m</sup>:*DEL<sup>SJ</sup>* abl pre-B cells. The complete signal end sequences are noted at the top of the figure.

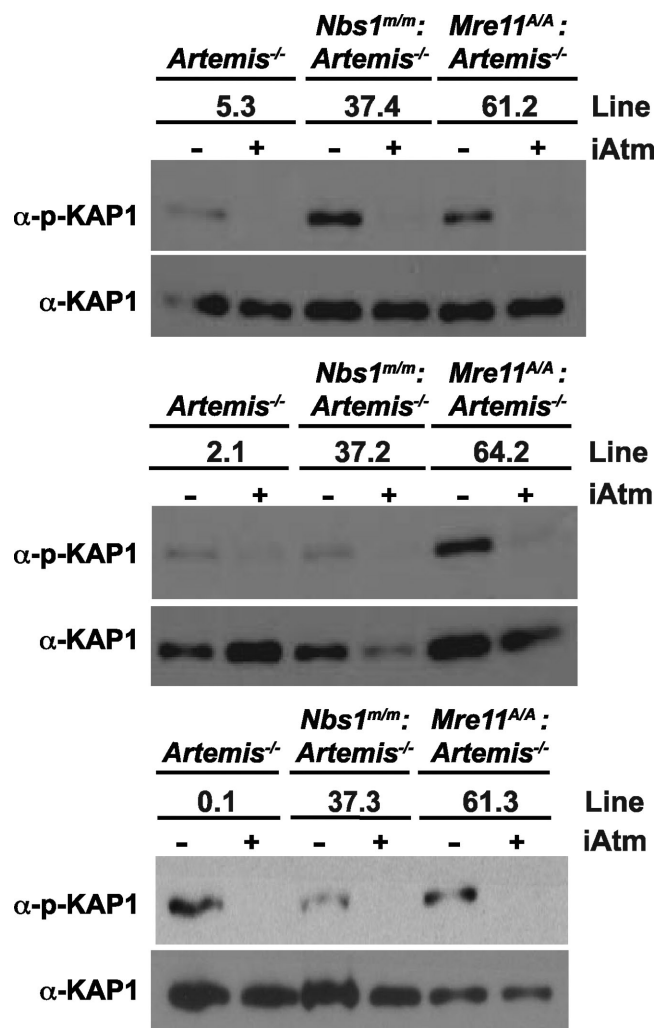


Figure S5. Analysis of KAP-1 phosphorylation carried out as described in Fig. 4 a on additional *Artemis*<sup>-/-</sup>, *Artemis*<sup>-/-</sup>:*Nbs1*<sup>m/m</sup>, and *Artemis*<sup>-/-</sup>:*Mre11*<sup>A/A</sup> abl pre-B cell lines.

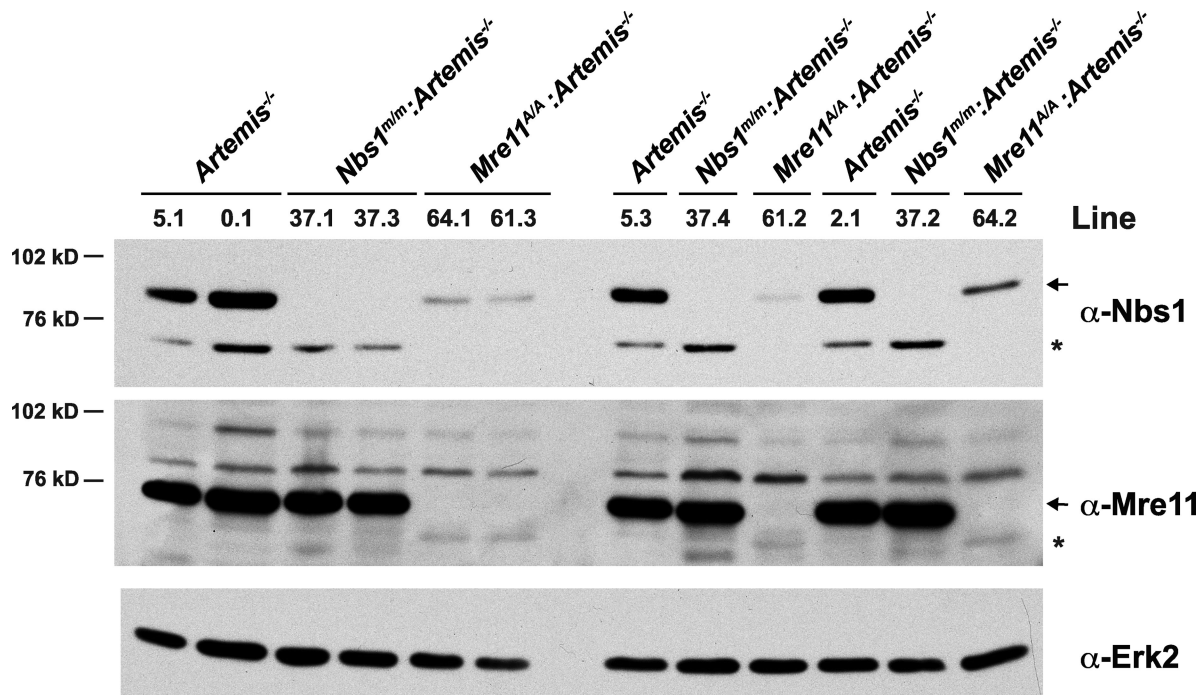


Figure S6. Nbs1 and Mre11 western blot analyses in the different *Artemis*<sup>-/-</sup>, *Artemis*<sup>-/-</sup>;*Nbs1*<sup>m/m</sup>, and *Artemis*<sup>-/-</sup>;*Mre11*<sup>A/A</sup> abl pre-B cells analyzed. Erk2 is shown as a protein loading control. The arrows indicate the native WT Nbs1 and Mre11 proteins. The asterisks indicate bands of expected size for the hypomorphic Nbs1 and Mre11 proteins. The identity of the band in the *Artemis*<sup>-/-</sup> abl pre-B cells that is of similar size to that expected for the *Nbs1*<sup>m</sup> hypomorphic protein is not known.

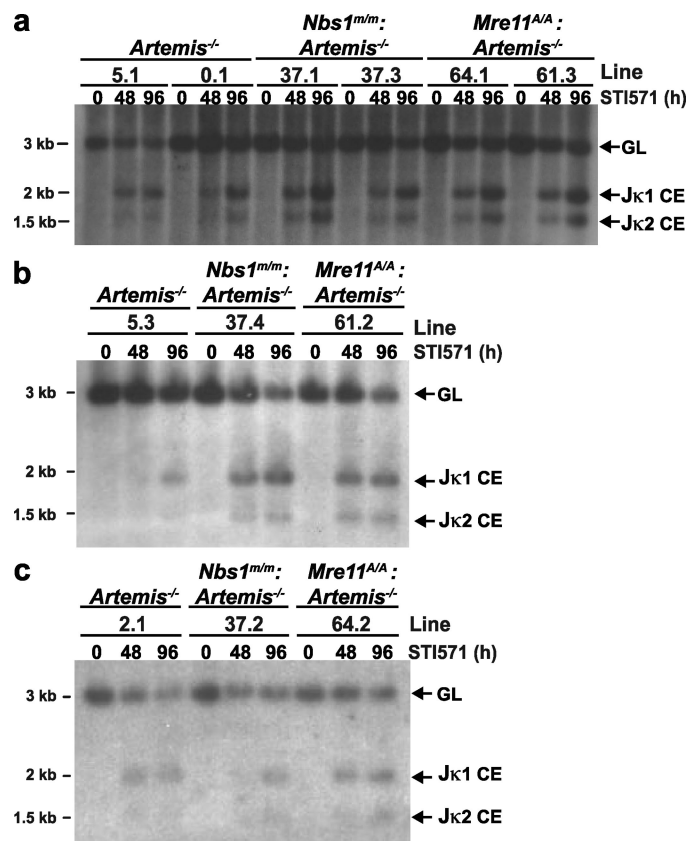


Figure S7. Southern blot analyses of IgL- $\kappa$  locus J $\kappa$  coding ends after Rag induction in the different *Artemis*<sup>-/-</sup>, *Artemis*<sup>-/-</sup>:*Nbs1*<sup>m/m</sup>, and *Artemis*<sup>-/-</sup>:*Mre11*<sup>A/A</sup> abl pre-B cell lines. Analyses were carried out as described in Fig. 3 a.

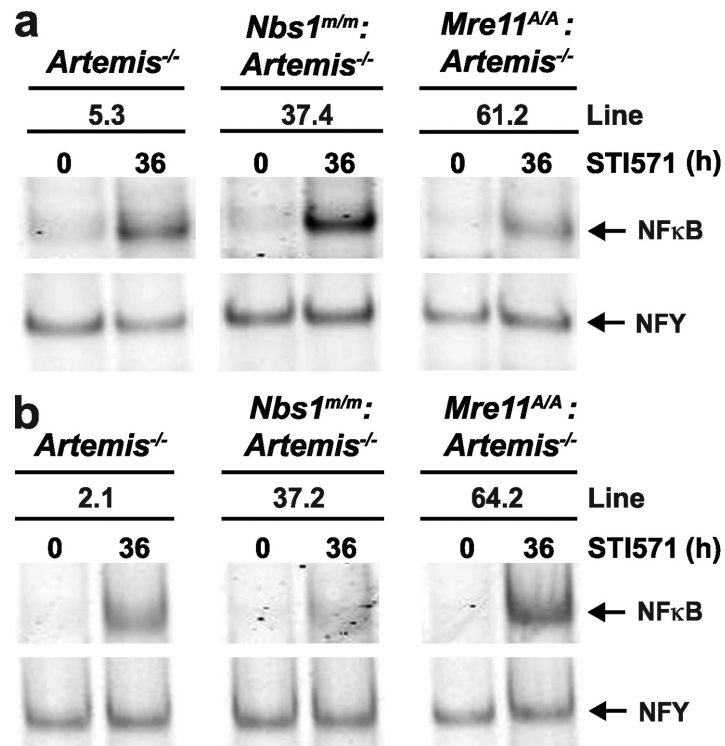


Figure S8. EMSA analyses of NF-κB translocation to the nucleus in the additional *Artemis*<sup>-/-</sup>, *Artemis*<sup>-/-</sup>:*Nbs1*<sup>m/m</sup>, and *Artemis*<sup>-/-</sup>:*Mre11*<sup>A/A</sup> abl pre-B cells not analyzed in Fig. 4 c. Analyses were carried out as described in Fig. 4 c.