## **Supplemental materials**

Supplemental FIG. S1. Overexpression of *FEN-1* enhanced I-*Sce*I-induced gene conversion in wild-type cells. The recombination frequency in the SCneo reporter construct is shown as the number of G418-resistant colonies derived from 10<sup>7</sup> cells transfected with the indicated plasmid. Expression vector of chicken *FEN-1* was cotransfected with the I-*Sce*I expression plasmid into wild-type cells. The experiments were done more than four times.

Supplemental FIG. S2. Generation of  $XPG^{-/-}$  DT40 cells. (A) Schematic representation of targeted disruption of the chicken XPG gene. The chicken XPG locus, the two targeting constructs and the resulting targeted locus are shown. The black boxes indicate the exons of the XPG gene. The triangles flanking the blasticidin S resistance (*bsr*) and histidinol dehydrogenase (*hisD*) genes designate the loxP sequences. The italic B denotes a *Bam*HI site. The figure is not drawn to scale. (B) Southern blot analysis of targeted cells. *Bam*HI-digested genomic DNA of wild-type (+/+), heterozygous (+/-) and homozygous mutant (-/-) cells was hybridized with the probe shown in (A).

Name	Sequence				
NeoS	GGATCGGCCATTGAACAAGATGGATTGCAC				
NeoE	CCTCAGAAGAACTCGTCAAGAAGGCGATAG				
Mneo-1 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCAGGGATAACAGG				
	GTAATCGATGCCTGCTTGCCGAATATCATGGTGGA				
Mneo-1 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGATTACCCTGTTATC				
	CCTGGTCACGACGAGATCCTCGCCGTCGGGCAT				
Mneo-2 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCAGGGACAATAGG				
	GTAATCGATGCCTGCTTGCCGAATATCATGGTGGA				
Mneo-2 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGATTACCCTATTGTC				
	CCTGGTCACGACGAGATCCTCGCCGTCGGGCAT				
Mneo-3 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCAGAGACAATAGA				
	GTAATCGATGCCTGCTTGCCGAATATCATGGTGGA				
Mneo-3 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGATTACTCTATTGTC				
	TCTGGTCACGACGAGATCCTCGCCGTCGGGCAT				
Mneo-4 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCAGGGTAATCGATG				
	CCTGCTTGCCGAATATCATGGTGGA				
Mneo-4 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGATTACCCTGGTCA				
	CGACGAGATCCTCGCCGTCGGGCAT				
Mneo-5 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCATCGATGCCTGCT				
	TGCCGAATATCATGGTGGA				
Mneo-5 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGATGGTCACGACGA				
	GATCCTCGCCGTCGGGCAT				
Mneo-6 (+)	ATGCCCGACGGCGAGGATCTCGTCGTGACCGATGCCTGCTTGC				
	CGAATATCATGGTGGA				
Mneo-6 (-)	TCCACCATGATATTCGGCAAGCAGGCATCGGTCACGACGAGAT				
	CCTCGCCGTCGGGCAT				

Supplemental table S1. Primers used in I-SceI-induced gene targeting assay

	OVA	RAD54	CENP-H	β-ACTIN
Wild-type	29/41 (70.7%)	8/40 (20%)	40/48 (83.3%)	26/63 (41%)
FEN-1 <sup>-/-</sup>	19/72 (26.4%)	2/51 (3.92%)	24/47 (51.1%)	37/87 (43%)
FEN-1 <sup>-/-</sup> +FEN-1	15/34 (44.1%)	5/46 (10.9%)	N.D.	N.D.

Supplemental table S2. The gene targeting efficiencies at various loci in  $FEN-1^{-/-}$  cells.

Shown are the number of targeted clones over the number of clones analyzed. N.D., not determined.

**S**1



