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Antagonistic roles of ubiquitin ligase HEI10 and SUMO ligase RNF212 regulate meiotic recombination

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Abstract

Crossover recombination facilitates accurate segregation of homologous chromosomes during meiosis^{1,2}. In mammals, poorly characterized regulatory processes ensure every pair of chromosomes obtains at least one crossover, even though the majority of recombination sites yield non-crossovers³. Designation of crossovers involves selective localization of SUMO-ligase RNF212 to a minority of recombination sites where it stabilizes pertinent factors, such as MutS γ ⁴. Here we show ubiquitin-ligase HEI10/CCNB1IP1^{5,6} is essential for this crossover/non-crossover differentiation process. In *Hei10* mutant mice, RNF212 localizes to most recombination sites and dissociation of RNF212 and MutS γ from chromosomes is blocked. Consequently, recombination is impeded and crossing-over fails. In wild-type mice, HEI10 accumulates at designated crossover sites suggesting a late role to implement crossing-over. Like RNF212, dosage-sensitivity indicates

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AUTHOR CONTRIBUTIONS

H.Q., H.B.D.P.R., Y.Y., J.W. and N.H. conceived and designed the experiments. H.Q., H.B.D.P.R., Y.Y., J.H.F. J.C., D.D, K.N., R.S., E.S, J.K.H., J.W. and N.H. performed the experiments. H.Q., H.B.D.P.R., Y.Y., J.H.F., J.W. and N.H. analyzed the data. J.S., E.S. and P.E.C. contributed reagents, materials and/or analysis tools. H.Q., H.B.D.P.R., Y.Y., J.W. and N.H. wrote the manuscript.

COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

HEI10 is a limiting factor for crossing-over. We suggest SUMO and ubiquitin play antagonistic roles during meiotic recombination that are balanced to effect differential stabilization of recombination factors at crossover and non-crossover sites.

Variants of both *RNF212* and *HEI10/CCNB1IP1* are associated with heritable variation in the rate of crossing-over in humans⁷⁻¹⁰. Rnf212 and Hei10 also share structural and functional similarities. Both proteins have tripartite structures with RING, coiled-coil and tail domains, and are inferred to catalyze post-translational protein modification by ubiquitin-like proteins^{4-6,11,12}. Rnf212 is implicated as an E3 enzyme for SUMO modification while Hei10 has ubiquitin-ligase activity^{4,5,11} (Y.Y and N.H, unpublished observations). In both *Hei10^{mei4/mei4}* and *Rnf212^{-/-}* mutant mice, early stages of meiosis occur normally and full synapsis of homologous chromosomes (homologs) is achieved^{4,6}. However, crossover-specific recombination complexes, containing the MutL γ complex (MLH1 and MLH3) and cyclin-dependent kinase CDK2, fail to assemble^{4,6}. Consequently, crossing-over fails and the animals are sterile. These similarities prompted us to examine the relationship between these two pro-crossover factors.

Using immunofluorescence cytology, we previously described the dynamic localization pattern of RNF212 to synaptonemal complexes⁴, the meiosis-specific structures that connect homologous chromosomes (homologs) along their lengths during the pachytene stage of meiosis. As homologs undergo synapsis during zygonema, RNF212 localizes specifically to the central region of synaptonemal complexes forming a punctate pattern of immunostaining foci. Consistent with previous analysis⁴, in wild-type spermatocytes at early pachynema, when synapsis is complete, ~150 foci are observed per nucleus (Fig. 1a,k). However, by mid-pachynema, most staining has disappeared and RNF212 foci are retained only at sites where crossovers will form (Fig. 1b,c,k). These crossover-specific RNF212 foci are then lost by late pachynema, prior to the disassembly of synaptonemal complexes at diplonema (Fig. 1d,e,k).

In spermatocytes from *Hei10^{mei4/mei4}* mice, the early staining pattern of abundant RNF212 foci appears normal (155.9 ± 37.2 (s.d.), 20 early pachytene nuclei; versus 153.0 ± 42.8 in wild type, 20 nuclei; Fig. 1f,k). Strikingly, this pattern persists throughout pachynema and loss of RNF212 from the chromosomes is only seen when synaptonemal complexes are disassembled during diplonema (Fig. g,h,i,j,k). Moreover, the numbers of RNF212 foci are significantly higher than ever seen in wild-type spermatocytes ($P = 0.0003$, Mann-Whitney test). Thus, HEI10 is required for the post-synapsis turnover of RNF212 that culminates in its selective retention at future crossover sites.

To examine the consequences of persistent RNF212 for recombination in *Hei10^{mei4/mei4}* mutants, we examined chromosomal dynamics of the MutS γ complex (Fig. 2). MutS γ comprises MSH4 and MSH5, two meiosis-specific homologs of the bacterial DNA mismatch-binding factor MutS¹³. Evidence to date indicates that MutS γ binds and stabilizes DNA strand-exchange intermediates to promote both homolog synapsis and crossing-over^{14,15}. We previously showed that a minority of MutS γ foci present in early pachynema co-localizes with RNF212⁴. Analysis of *Rnf212^{-/-}* knock-out mice indicates that RNF212

acts to stabilize MutS γ and thereby designate a crossover fate to this subset of recombination sites.

In wild-type spermatocytes, chromosomal localization of MutS γ resembles that of RNF212: 82.9 ± 23.4 (s.d.) MSH4 immunostaining foci are observed in late zygonema and early pachynema; at mid pachynema only 39.4 ± 9.6 foci are retained; and by the onset of diplonema, MSH4 staining has essentially disappeared (Fig. 2a–e,k). In *Hei10^{mei4/mei4}* mutant spermatocytes, chromosomal dynamics of MutS γ are severely aberrant. Although normal numbers of MSH4 foci are formed, focus numbers remain high throughout pachynema and only decrease after homologs desynapse at diplonema (Fig. 2f–k).

Super-resolution structured illumination microscopy (SIM¹⁶) reveals that stabilization of MutS γ foci correlates with the degree of co-localization between MutS γ and RNF212 (Fig. 2l–p and Supplementary Fig. 1). Consistent with our previous analysis⁴, around a third of MSH4 foci colocalize with RNF212 in late zygotene and early pachytene spermatocytes (Fig. 2l,m,p; Supplementary Fig. 1). In sharp contrast, in *Hei10^{mei4/mei4}* mutant spermatocytes, MSH4 foci show a high level of colocalization (~90%) with RNF212 at all stages from early zygonema through early diplonema (Fig. 2l–p; Supplementary Fig. 1). These observations lend further support to our inference that RNF212 stabilizes association of MutS γ with recombination sites⁴. Thus, RNF212 and HEI10 have antagonistic functions with respect to the stabilization of MutS γ at recombination sites. We suggest that the balance of these two activities underpins the temporal dynamics and spatial patterning of RNF212 and MutS γ seen in wild-type spermatocytes.

Meiotic recombination is initiated by the programmed formation of DNA double-strand breaks (DSBs¹⁵). The persistence of MutS γ complexes implies that DSB-repair is delayed in *Hei10^{mei4/mei4}* spermatocytes. Immunostaining for DSB-induced H2AX phosphorylation (γ H2AX) supports this inference (Fig. 3). In wild-type nuclei, γ H2AX staining initially forms a pan-nuclear cloud, which diminishes to limited chromatin flares and foci as chromosome synapsis ensues, and finally disappears from autosomes around mid pachynema¹⁷ (Fig. 3a–e,k; note that γ H2AX accumulates as a large staining body on the chromatin of the sex chromosomes where it facilitates transcriptional silencing¹⁸). In *Hei10^{mei4/mei4}* spermatocytes, although pan-nuclear γ H2AX staining diminishes with synapsis, delayed DSB-repair is indicated by the persistence of γ H2AX foci throughout pachynema (Fig. 3f–j,k). Ultimately, however, DSBs appear to be repaired in *Hei10^{mei4/mei4}* spermatocytes because broken chromatids are not detected in late-stage nuclei⁶. Together, these data suggest that HEI10-dependent elimination of RNF212 from synaptonemal complexes is required for the timely removal of MutS γ from most recombination sites as pachynema progresses. This in turn allows the timely progression of recombination and repair of DSBs.

Despite their broadly similar phenotypes, we can now conclude that *Rnf212^{-/-}* and *Hei10^{mei4/mei4}* mutants have distinct defects with respect to the designation of crossover sites. In the absence of RNF212, designation of crossover sites fails because no MutS γ complexes are stabilized beyond early pachynema⁴; whereas absence of HEI10 causes most

or all MutS γ complexes to be stabilized, as though all sites have been designated a crossover fate. This raises the question, why don't *Hei10^{mei4/mei4}* mutants form lots of crossovers?

Insights into this question, and how HEI10 regulates the dynamics of RNF212 and MutS γ , came from examining the chromosomal localization patterns of HEI10. Although previous attempts to localize mouse HEI10 to meiotic chromosomes failed⁶, we were able to show that mouse HEI10 associates with synaptonemal complexes to form distinct immunostaining foci (Fig. 4a–k and Supplementary Figs. 2 and 3). Unlike RNF212, distinct foci of HEI10 were rarely detected along nascent synaptonemal complexes during zygonema (Fig. 4a). Given that *Hei10^{mei4/mei4}* phenotypes are already apparent at this time (e.g. Fig. 2p), we infer that cytologically undetectable HEI10 regulates early-stage dynamics of RNF212 and MutS γ . Indeed, the possibility this function of HEI10 does not involve association with meiotic chromosomes cannot be ruled out. However, by early pachynema, HEI10 foci could be detected (Fig. 4c,d) and their numbers peaked during mid pachynema with an average of 27 foci per nucleus (27.2 ± 8.8 (s.d.); 22 nuclei; Fig. 4e,f,k). At this stage, HEI10 focus numbers were quite variable, with as few as 15 and as many as 50 foci per nucleus. In late pachytene nuclei, the average HEI10 focus number was lower and less variable (21.1 ± 3.6 (s.d.); 20 nuclei; Fig. 4g,h,k). At the onset of diplonema, HEI10 foci were no longer detected (Fig. 4i,j).

As HEI10 promotes loss of RNF212 and MutS γ from synaptonemal complexes, we examined co-localization with these proteins (Fig. 4l–o). In mid-pachytene spermatocytes, only 23% of RNF212 foci and 29% of MSH4 foci colocalized with HEI10 (respectively, $22.6 \pm 7.7\%$ (s.d.) and $28.9 \pm 15.1\%$, 10 nuclei each). HEI10 is required for formation of crossover-specific complexes containing CDK2 and MutL γ (MLH1-MLH3). In contrast to RNF212 and MSH4, a high degree of co-localization was seen between these crossover-specific markers and HEI10 (Fig. 4p–s). $74.0 \pm 10.7\%$ (s.d.) of non-telomeric CDK2 foci and $94.3 \pm 6.9\%$ of MLH1 foci co-localized with HEI10 (10 and 19 nuclei respectively). These data are consistent with HEI10 foci superseding co-complexes of RNF212 and MutS γ at future crossover sites.

The genetic requirements for chromosomal localization of HEI10 were investigated using several mutant lines. SPO11 catalyzes DNA breakage to initiate recombination¹⁵. In *Spo11^{-/-}* spermatocytes, homolog pairing is defective but a significant fraction of nuclei assemble incomplete synaptonemal complexes, which generally involve non-homologous chromosomes^{19,20}. HEI10 foci were diminished, although not completely eliminated, in these nuclei (Fig. 5a,b,g). This dependency contrasts that of RNF212, which readily associates with synaptonemal complexes independently of recombination⁴. SYCP1 encodes a major component of the synaptonemal complex central region²¹. In the *Sycp1* knockout mice, meiotic recombination initiates normally and homologs closely align, but synapsis is precluded. Also, MutL γ foci are not detected in *Sycp1^{-/-}* nuclei indicating that designation or maturation of crossover sites fails in this mutant. Formation of HEI10 foci was greatly reduced in pachytene-like *Sycp1^{-/-}* spermatocytes, but most nuclei contained a few foci (4.5 ± 3.1 (s.d.) per nucleus, 20 nuclei; Fig. 5c,d,g). In *Rnf212^{-/-}* mutants, homologs undergo synapsis but designation of crossover sites is defective⁴. Similar to the *Sycp1^{-/-}* mutant, a few HEI10 foci were detected in *Rnf212^{-/-}* mutant spermatocytes ($6.7 \pm$

4.9 (s.d.), 23 nuclei; Fig. 5e,f,g). We infer that initiation of recombination, homolog synapsis and the designation of crossover sites are important for the normal formation and/or stabilization of HEI10 foci.

Finally, we examined HEI10 localization in mice lacking the MLH3 component of the crossover-specific factor, MutL γ ¹³. In *Mlh3*^{-/-} spermatocytes, homolog synapsis and initial designation of crossover sites appear normal, but implementation of crossing-over fails^{4,22}. In sharp contrast to the other mutants examined, high numbers of HEI10 foci were observed in *Mlh3*^{-/-} spermatocytes (Fig. 5h-m). Unlike wild-type, HEI10 foci were already detectable in zygonema (Fig. 5h,i,n), reached very high numbers during mid pachynema (89.9 \pm 24.5 (s.d.) foci per nucleus, 21 nuclei; Fig. 4j,k,n) and persisted into diplonema (Fig. 5l,m,n). At least half of the foci in mid pachynema were coincident with γ H2AX staining (53.0 \pm 12.5% (s.d.), 10 nuclei; Supplementary Figure 4), implying that the majority of HEI10 accumulates at sites of DSB repair in *Mlh3*^{-/-} cells.

These data suggest that chromosomal localization of HEI10 occurs in two phases: first, HEI10 is licensed to accumulate into foci associated with synaptonemal complexes. These, normally transient, complexes may generally promote progression of recombination. In wild type, the relatively high and variable numbers of HEI10 foci seen in mid pachynema may be a manifestation of this first phase. Subsequently, stable accumulation of HEI10 specifically at crossover sites is directed by MLH3 (and, presumably, MLH1). Notably, MLH3 restrains accumulation of HEI10 during zygonema, an early function that was not anticipated from the timing of crossover-specific MLH3 foci, which don't appear until early-mid pachynema¹³.

Taken together, our data imply that HEI10 functions during zygonema to limit the co-localization of RNF212 with MutS γ -associated recombination sites and thereby establish early differentiation of crossover and non-crossover sites. Later, HEI10 is directed by MutL γ (and perhaps CDK2) to stably accumulate at designated crossover sites. Here, we propose that HEI10 also promotes dissociation of RNF212 and MutS γ to allow progression of recombination and implementation of the final steps of crossing over. CNTD1 is a cyclin-B related protein and the mammalian homolog of the *C. elegans* pro-crossover factor, COSA-1²³. Intriguingly, the meiotic phenotypes of *Cntd1*^{-/-} and *Hei10*^{mei4/mei4} mice are remarkably similar and CNTD1 is required for chromosomal localization of HEI10, suggesting that the two proteins function together (P. Cohen, personal communication). The model in Supplementary Figure 5 synthesizes the key points of our analysis and those of previous studies.

Recently, recombination rate in humans has been associated with a variant in the untranslated 5' region of *HEI10/CNNB1IP1* (A. Kong, personal communication), which has the potential to alter expression level and suggests that Hei10 may be a dosage-sensitive regulator of crossing over. To determine whether the crossover function of mouse HEI10 is dosage sensitive, we analyzed spermatocytes from *Hei10*^{+/^{mei4}} heterozygotes (Fig. 6). Indeed, significant decreases in the numbers HEI10 foci (20.6%), MLH1 foci (13.5%) and chiasmata (10%) were detected ($P=0.0003$, < 0.0001 and 0.0088 , respectively; Mann-Whitney test). In spermatocytes, homologs not tethered by crossovers are detected by the

spindle checkpoint, which triggers apoptosis²⁴. Consistent with the reduced crossing-over seen in *Hei10*^{+/*mei4*} heterozygotes, we detected a significant increase in apoptotic cells (TUNEL positive) in testes sections from *Hei10*^{+/*mei4*} heterozygotes (Fig. 6j–m and Supplementary Fig. 6; $P < 0.0001$ Mann-Whitney test).

Intriguingly, the crossover function of RNF212 also shows dosage sensitivity in the mouse, and human *Rnf212* variants have been associated with changes in recombination rate^{4,7–9}. These observations are consistent with the idea that the balance of SUMO and ubiquitin is a key aspect of crossover regulation. It will be interesting to see if human alleles of *HEI10/CCNB1IP1*, *RNF212* and recombination factors such as MutS γ , interact to modulate recombination rate, fertility and the risk of aneuploidy.

ONLINE METHODS

Mice

All mice were congenic with the C57BL/6J background. Mice were maintained and used for experimentation according to the guidelines of the Institutional Animal Care and Use Committees of the University of California, Davis and the Middlebury College Animal Facility. The *Hei10*, *Mlh3*, *Rnf212*, *Spo11* and *Sycp1* mutant lines and primer sequences for genotyping were previously described^{4,6,19,21,22}. Male mice between 2–6 months were used for experimentation.

Protein blot analysis

Tissues from adult mice were sonicated in RIPA buffer, protein concentration was measured by the Bradford assay and 100–200 μ g of protein was separated by SDS-PAGE. After protein transfer to nitrocellulose membranes (Waterman), blots were incubated overnight with the following antibodies: mouse monoclonal anti-CCNB1IP1/HEI10 (ab118999 Abcam, 1:2000 dilution), rabbit polyclonal anti-CCNB1IP1/HEI10 (this study, 1:2000), or mouse anti-tubulin (BioLegend, 625902, 1:2,000). Secondary antibodies (1:10,000 dilution) were goat anti-rabbit or anti-mouse IgGs conjugated to horseradish peroxidase (HRP; SouthernBiotech, 4050-05 and 1031-05, respectively). HRP was detected using the ECL reagent (Pierce).

Antibody Production

A polyclonal antibody against mouse HEI10/CCNB1IP1 was raised in rabbits against a mixture of two C-terminal peptides. Antibodies were purified from serum using Protein A/G spin columns (GE Healthcare).

Cytology

Testes and ovaries were dissected from freshly killed animals and processed for surface spreading as described²⁵. For all quantification, images from at least two animals (2–5) were analyzed. Comparisons were made between animals that were either littermates or matched by age. All cytological analyses were performed by two observers; the second observer was blind to which group/genotype was being analyzed. Immunofluorescence staining was performed as described²⁶, using the following primary antibodies with incubation overnight

at room temperature: mouse anti-SYCP3 (sc-74568 Santa Cruz, 1:200 dilution), rabbit anti-SYCP3 (sc-33195 Santa Cruz, 1:300), guinea pig anti-SYCE1²⁷ (1:2000)(generously provided by Chist Höög, Karolinska Institutet), guinea pig anti-RNF212⁴ (1:50), rabbit anti-RNF212⁴ (1:200), rabbit anti-MSH4 (ab58666 Abcam, 1:100), mouse monoclonal anti-CCNB1IP1/HEI10 (ab118999 Abcam, 1:150), rabbit polyclonal anti-CCNB1IP1/HEI10 (this study), mouse anti-MLH1 (1:50, 550838 BD Pharmingen), mouse monoclonal anti- γ H2AX (05-636 Millipore, 1:500), mouse monoclonal anti-CDK2 (sc-6248 Santa Cruz, 1:200), guinea pig anti-H1t²⁸ (a gift from M.A. Handel, The Jackson Laboratory; 1:1,000 dilution). Slides were subsequently incubated with the following goat secondary antibodies for 1 h at 37 °C: anti-rabbit 488 (A11070 Molecular Probes, 1:1000 dilution), anti-rabbit 568 (A11036 Molecular Probes, 1:2000), anti-mouse 555 (A21425 Molecular Probes, 1:1000), anti-mouse 594 (A11020 Molecular Probes, 1:1000), anti-mouse 488 (A11029 Molecular Probes, 1:1000), and anti-guinea pig fluorescein isothiocyanate (106-096-006 FITC, Jackson Labs, 1:200). Coverslips were mounted with ProLong Gold antifade reagent (Molecular Probes). For chiasma counts, air-dried preparations of diakinesis/metaphase I-stage cells were prepared as described²⁹ and stained with DAPI.

TUNEL Assay

Testes were fixed in formalin, embedded in paraffin, sectioned and processed using the ApopTag Plus Peroxidase In Situ Apoptosis Detection kit (Chemicon).

Imaging

Immunolabeled chromosome spreads and DAPI-stained diakinesis/metaphase I nuclei were imaged using a Zeiss AxioPlan II microscope with 63 \times Plan Apochromat 1.4 objective and EXFO X-Cite metal halide light source. Images were captured by a Hamamatsu ORCA-ER CCD camera and processed using Volocity (Perkin Elmer) and Photoshop (Adobe) software packages. SIM analysis was performed using a Nikon N-SIM super-resolution microscope system and NIS-Elements 2 image processing software. MSH4-RNF212 colocalization was determined using NIS-Elements and co-foci were confirmed by visual inspection. Testes sections were imaged using an Axiovert 200 microscope and AxioCamMRc camera using AxioVision 4.4 software. Apoptotic cells were imaged and counted in representative fields of view.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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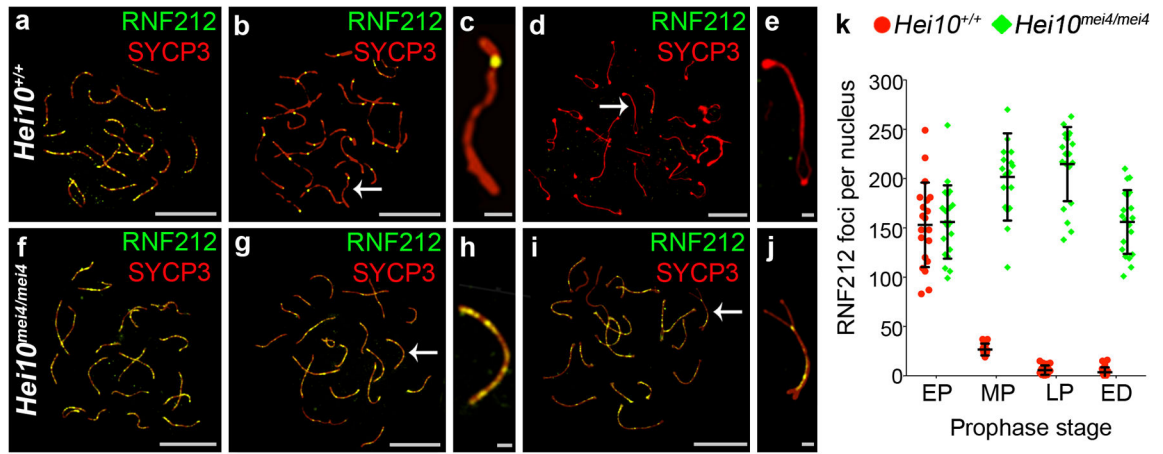


Figure 1.

RNF212 fails to dissociate from synaptonemal complexes in *Hei10*^{mei4/mei4} mutant spermatocytes. All nuclei were immunostained for RNF212 (green) and homolog axis component, SYCP3 (red). (a–e) Wild-type (*Hei10*^{+/+}) nuclei at (a) early pachynema, (b,c) mid pachynema, and (d,e) early diplonema. c and e show magnified views of chromosomes indicated by arrows in b and d, respectively. (f–j) *Hei10*^{mei4/mei4} mutant nuclei at (f) early pachynema, (g,h) mid pachynema, and (i,j) early diplonema. h and j show magnified views of chromosomes indicated by arrows in g and i, respectively. (k) Quantification of RNF212 foci (\pm s.d.) at successive prophase I stages. EP, early pachynema; MP, mid pachynema; LP, late pachynema; ED, early diplonema. 20 nuclei were analyzed for each stage. Scale bars, 10 μ m in a,b,d,f,g,i and 1 μ m in c,e,h,j.

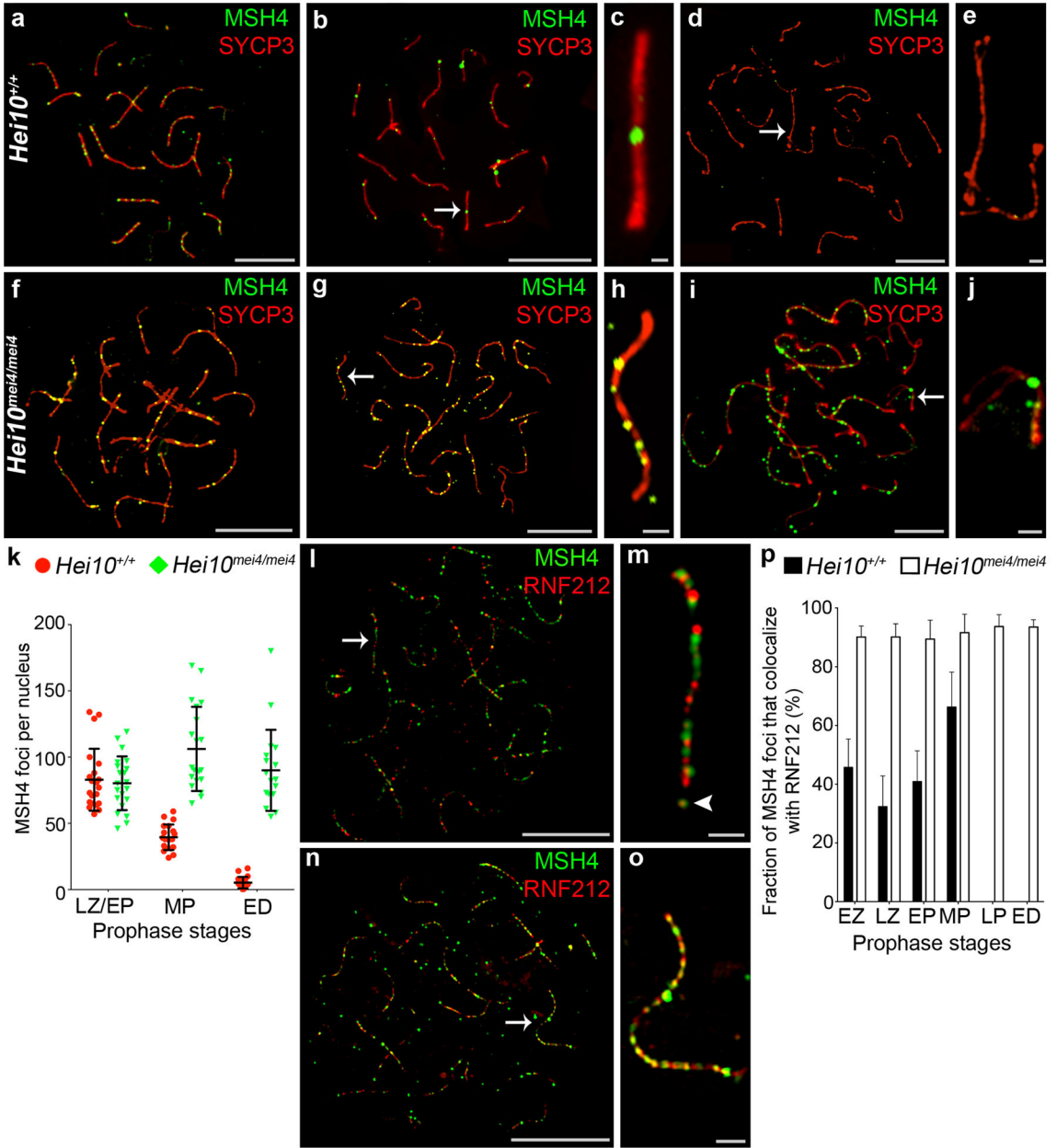


Figure 2. Persistence of MutS γ complexes in *Hei10^{mei4/mei4}* spermatocytes. (a–j) Spermatocyte nuclei immunostained for MSH4 (green) and SYCP3 (red). (a–e) Wild-type nuclei at (a) late zygonema, (b,c) mid pachynema, and (d,e) early diplonema. c and e show magnified views of chromosomes indicated by arrows in b and d, respectively. (f–j) *Hei10^{mei4/mei4}* mutant nuclei at (f) late zygonema, (g,h) mid pachynema, and (i,j) early diplonema. h and j show magnified views of chromosomes indicated by arrows in g and j, respectively. (k) Quantification of MSH4 foci (\pm s.d.) at successive prophase I stages. LZ, late zygonema; EP, early pachynema; ED, early diplonema. Number of nuclei analyzed at LZ/EP, MP and ED: 21, 20 and 20 for *Hei10^{+/+}*; and 21, 20 and 18 for *Hei10^{mei4/mei4}*. (l–o) SIM images of

early pachytene nuclei immunostained for MSH4 (red) and RNF212 (green). To allow accurate staging, nuclei were also stained for SYCP3 (not shown). **(l)** Wild-type nucleus. **(m)** Magnification of the chromosome highlighted by the arrow in **l**. **(n)** *Hei10^{me4/mei4}* mutant nucleus. **(o)** Magnification of the chromosome highlighted by the arrow in **n**. **(p)** Percentage of MSH4 foci (\pm s.d.) that colocalize with RNF212 at successive prophase substages determined by SIM analysis. EZ, early zygonema; LZ, late zygonema; EP, early pachynema; MP, mid pachynema; LP, late pachynema; ED, early diplonema. Number of *Hei10^{+/+}* nuclei analyzed at EZ, LZ, EP and MP, respectively: 10, 5, 10, and 7. Number of *Hei10^{mei4/mei4}* nuclei analyzed at EZ, LZ, EP, MP, LP and ED, respectively 5, 6, 25, 14, 9, 9.

Scale bars, 10 μ m in **a,b,d,f,g,i,l,n** and 1 μ m in **c,e,h,j,m,o**.

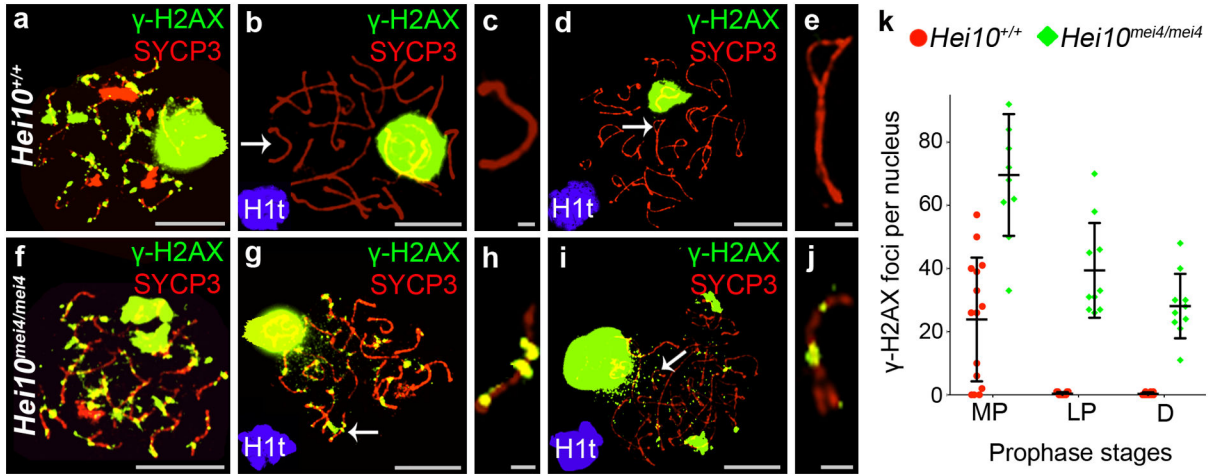


Figure 3.

Repair of DSBs is delayed in *Hei10*^{mei4/mei4} spermatocytes. All nuclei were immunostained for γ H2AX (green) and SYCP3 (red). (a–e) Wild-type nuclei at (a) early pachynema, (b,c) mid pachynema, and (d,e) diplonema. c and e show magnified views of chromosomes indicates by arrows in b and d, respectively. (f–j) *Hei10*^{mei4/mei4} mutant nuclei at (f) early pachynema, (g,h) mid pachynema, and (i,j) early diplonema. h and j show magnified views of chromosomes indicates by arrows in g and j, respectively. (k) Quantification of γ H2AX foci (\pm s.d.) at successive prophase I stages. MP, mid pachynema; LP, late pachynema; D, diplonema. Number of nuclei analyzed at MP, LP and D: 15, 11 and 10 for *Hei10*^{+/+}; and 10, 10 and 10 for *Hei10*^{mei4/mei4}.

Scale bars, 10 μ m in a,b,d,f,g,i and 1 μ m in c,e,h,j.

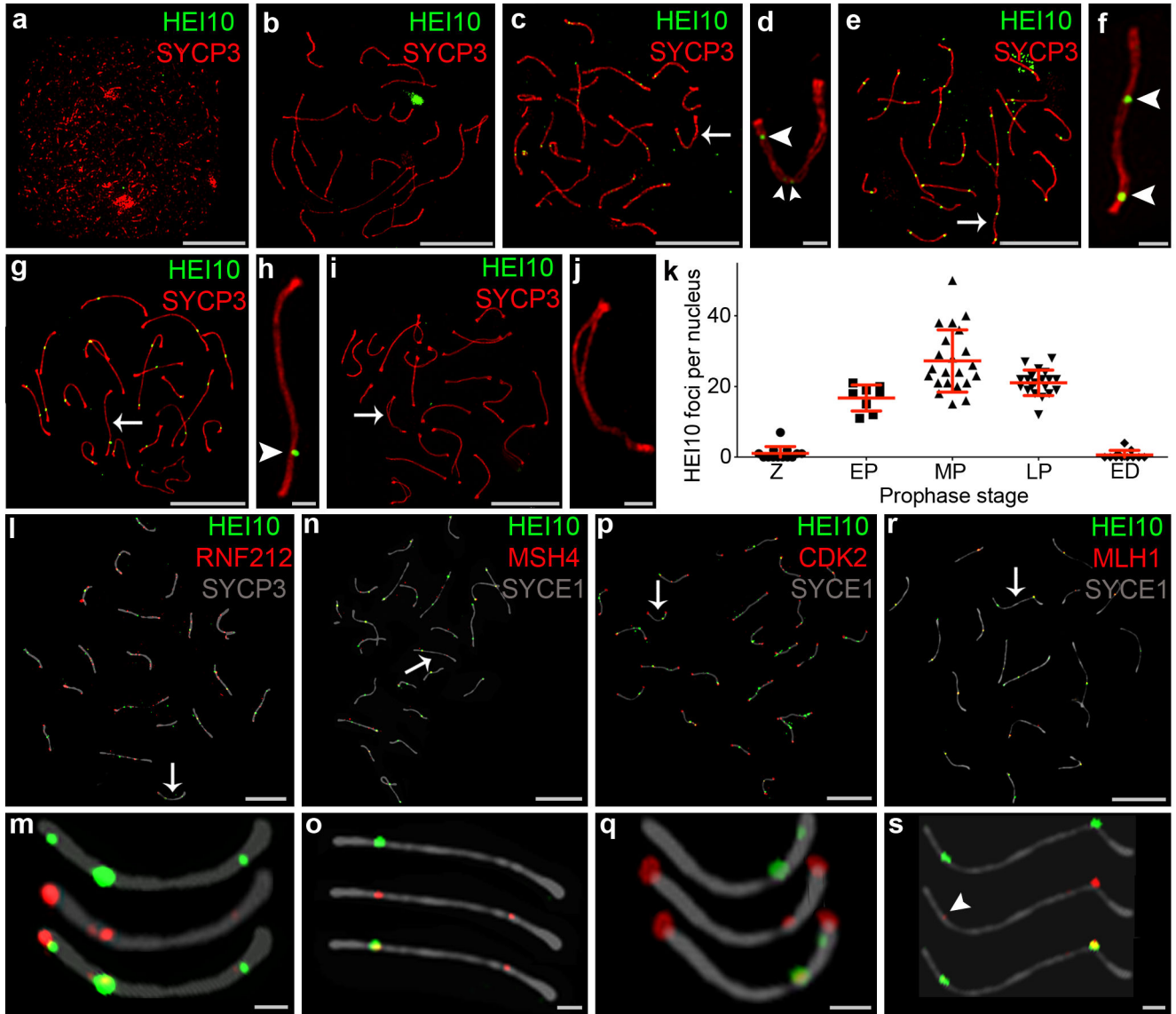


Figure 4. HEI10 localization to synaptonemal complexes and crossover sites. **(a–j)** Representative SIM images of wild-type spermatocyte nuclei immunostained for HEI10 (green) and SYCP3 (red) at **(a)** leptotene, **(b)** late zygotene, **(c,d)** early pachytene, **(e,f)** mid pachytene, **(g,h)** late pachytene, and **(i,j)** early diplotene. **(d, f, h, j)** Magnified views of the chromosomes indicated by arrows in **c, e, g, i**, respectively. Arrowheads highlight HEI10 foci. **(k)** Numbers of HEI10 foci per nucleus at successive prophase stages (Z, zygotene; EP, early pachytene; MP, mid pachytene; LP, late pachytene; ED, early diplotene). Horizontal bars represent means \pm s.d. Numbers of nuclei analyzed at Z, EP, MP, LP and ED, respectively: 13, 8, 22, 20 and 11. **(l,m)** Mid-pachytene spermatocyte immunostained for HEI10 (green), RNF212 (red), and SYCP3 (grey). **(m)** Magnified view of the chromosome indicated by the arrow in **l**. **(n,o)** Mid-pachytene spermatocyte immunostained for HEI10 (green), MSH4 (red) and synaptonemal complex central element protein, SYCE1

(grey). **(o)** Magnified view of the chromosome indicated by the arrow in **n**. **(p,q)** Mid-pachytene spermatocyte immunostained for HEI10 (green), CDK2 (red), and SYCE1 (grey). **(q)** Magnified view of the chromosome indicated by the arrow in **p**. **(r,s)** Mid-pachytene spermatocyte immunostained for HEI10 (green), MLH1 (red), and SYCE1 (grey). **(s)** Magnified view of the chromosome indicated by the arrow in **r**. The arrowhead highlights a small MLH1 focus that colocalizes with HEI10.

Scale bars, 10 μm in **a, b, c, e, g, i, l, n, p, r**; 1 μm in **d, f, h, j, m, o, q, s**.

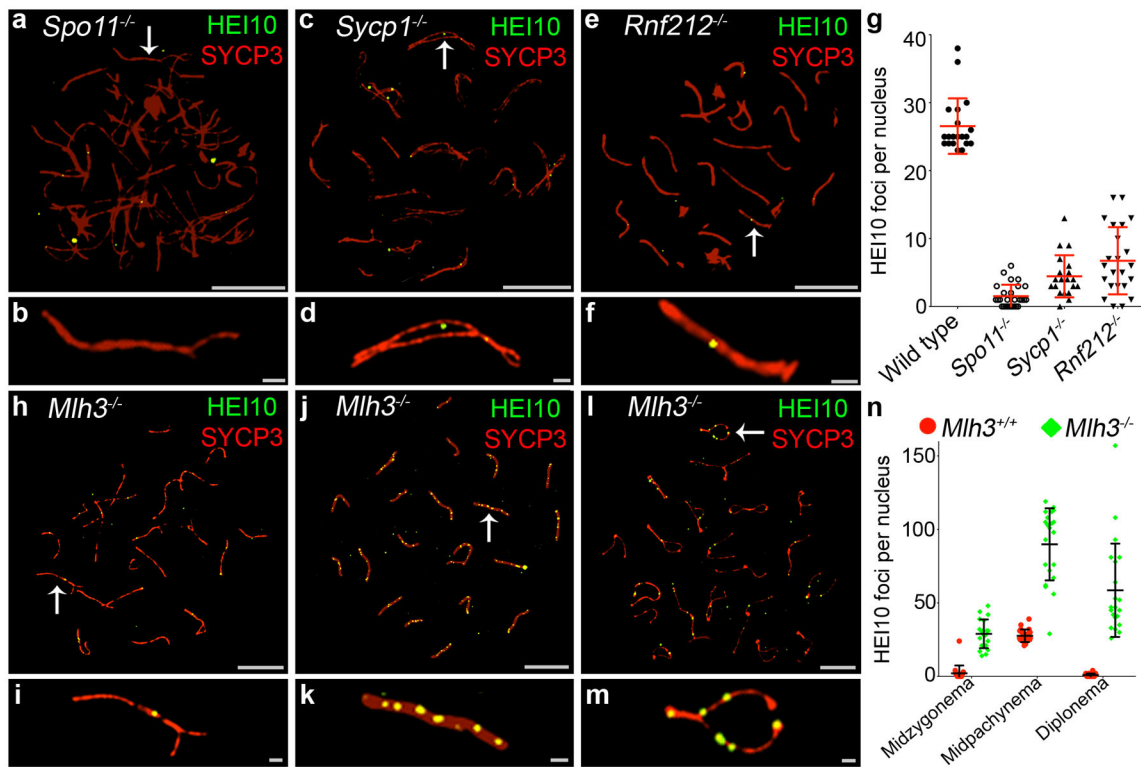


Figure 5.

Genetic requirements for HEI10 localization. **(a–m)** Representative spermatocyte nuclei, from indicated mutant lines, immunostained for HEI10 (green) and SYCP3 (red). **(a,b)** Spermatocyte nucleus from a *Spo11*^{-/-} mouse. **(b)** Magnification of the chromosome indicated by the arrow in **a**. **(c,d)** Spermatocyte nucleus from a *Sycp1*^{-/-} mouse. **(d)** Magnification of the chromosome indicated by the arrow in **c**. **(e,f)** Spermatocyte nucleus from a *Rnf212*^{-/-} mouse. **(f)** Magnification of the chromosome indicated by the arrow in **e**. **(g)** Numbers of HEI10 foci per nucleus in mid-late pachytene spermatocytes from wild-type, *Spo11*^{-/-}, *Sycp1*^{-/-} and *Rnf212*^{-/-} mice. Bars represent means \pm s.d. Numbers of nuclei analyzed: wild-type, 20; *Spo11*^{-/-}, 27; *Sycp1*^{-/-}, 20; *Rnf212*^{-/-}, 23. Analysis of *Spo11*^{-/-} was confined to nuclei with high levels of synapsis, predicted to be the equivalents of mid pachynema nuclei. Similarly, only *Sycp1*^{-/-} nuclei with a high degree of pseudo-synapsis were analyzed. However, true equivalents of wild-type mid pachynema nuclei may not be represented in *Spo11*^{-/-} and *Sycp1*^{-/-} testes due to apoptosis^{18,20}. **(h–m)** Representative spermatocyte nuclei from the *Mlh3*^{-/-} mutant. **(h)** Late zygotene nucleus. **(i)** Magnification of the chromosome indicated by the arrow in **h**. **(j)** Mid-pachytene nucleus. **(k)** Magnification of the chromosome indicated by the arrow in **j**. **(l)** Mid-diplotene nucleus. **(m)** Magnification of the chromosome indicated by the arrow in **l**. **(n)** Numbers of HEI10 foci per nucleus (\pm s.d.) at successive prophase stages in wild-type and *Mlh3*^{-/-} mice. MZ, mid-zygonema; MP, mid pachynema; D, diplonema. Number of nuclei analyzed at MZ, MP and D, respectively: 20, 22 and 20 for wild-type (*Mlh3*^{+/+}); and 20, 21 and 21 for *Mlh3*^{-/-}. Scale bars, 10 μ m in **a,c,e,h,j,l** and 1 μ m in **b,d,f,i,k,m**.

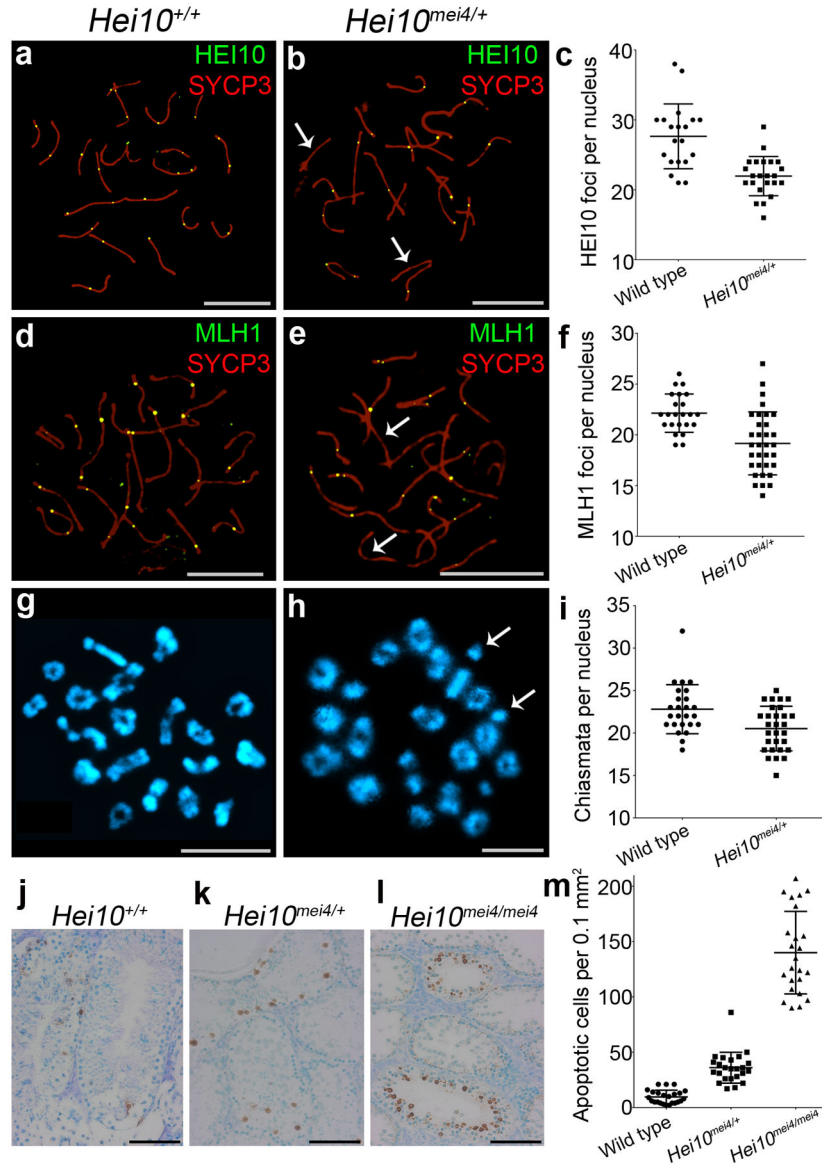


Figure 6. Dosage sensitivity of the HEI10 crossover function. (a,b) Mid-pachytene nuclei from (a) wild-type (*Hei10^{+/+}*) and (b) *Hei10^{+/mei4}* heterozygous mice immunostained for HEI10 (green) and SYCP3 (red). The arrows in b highlight chromosomes that lack HEI10 foci. (c) Numbers of HEI10 foci (\pm s.d.) per nucleus in mid-pachytene cells (23 wild-type and 33 *Hei10^{+/mei4}* nuclei). (d,e) Mid-pachytene nuclei from (d) wild-type and (e) *Hei10^{+/mei4}* heterozygous mice immunostained for MLH1 and SYCP3. The arrows in e highlight chromosomes that lack MLH1 foci. (f) Numbers of MLH1 foci per nucleus (\pm s.d.) in mid-pachytene cells (20 wild-type and 23 *Hei10^{+/mei4}* nuclei). (g, h) Chromosome spreads of diakinesis/metaphase I spermatocytes from (g) wild-type and (h) *Hei10^{+/mei4}* heterozygous mice stained with DAPI. (i) Numbers of chiasmata per nucleus (\pm s.d.) in diakinesis/metaphase I spermatocytes (25 wild-type and 29 *Hei10^{+/mei4}* nuclei). (j-l) TUNEL stained

testis sections from **(j)** *Hei10^{+/+}*, **(k)** *Hei10^{+/mei4}* and **(l)** *Hei10^{mei4mei4}* animals. **(m)** Quantification of TUNEL-positive apoptotic cells (\pm s.d.) in spermatocyte sections. Scale bars, 10 μ m in **a,b,d,e,g,h** and 100 μ m in **j,k,l**.

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