An ENU-induced mutation in the mouse *Rnf212* gene is associated with male meiotic failure and infertility

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Abstract

The ENU-induced *repro57* mutation was identified in an unbiased screen for the discovery of novel genes for fertility. Male *repro57* homozygous mice are infertile and exhibit significantly reduced testis weight compared with WT mice. Histological examination of mutant testes revealed that spermatocytes degenerated during late prophase, and no mature spermatozoa were found in the seminiferous epithelium, suggesting that infertility is caused by the arrest of spermatogenesis at late meiotic prophase. Consistent with this hypothesis, the number of foci with MLH1, a protein essential for crossing over, is greatly reduced in *repro57* mutant spermatocytes, which also lack chiasmata between homologs and exhibit premature dissociation of XY chromosomes. In *repro57* mutant mice, we identified a mutation in the *Rnf212* gene, encoding Ring finger protein 212. The overall phenotype of *repro57* mice is consistent with the recently reported phenotype of the *Rnf212* knockout mice; slight differences may be due to genetic background effects. Thus, the *repro57* nonsense mutation provides a new allele of the mouse *Rnf212* gene.

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Introduction

During meiosis, homologous chromosomes undergo highly programmed and evolutionarily conserved processes, including pairing of homologous chromosomes, synapsis, recombination to form chiasmata between homologs, and chromosome segregation mediated by chiasmata resolution. At the onset of meiosis, recombination is initiated by formation of DNA double-strand breaks (DSBs) by SPO11 in most organisms (Phadnis *et al.* 2011). DNA ends at the DSB sites are loaded with the recombinases RAD51 and DMC1, promoting homolog interaction leading to synapsis, which is mediated by the formation of a unique structural scaffold, the synaptonemal complex (SC) (Cromie & Smith 2007). During recombination, DSBs are repaired by either crossing over (CO) or non-CO process. CO sites are created by exchange of non-sister chromatids between homologs, and formation of at least one CO on each chromosome is essential for the accurate segregation of homologous chromosomes (Baudat *et al.* 2013). However, the precise mechanisms and full repertoire of genes responsible for mammalian CO recombination are still unclear (Handel & Schimenti

2010, Bolcun-Filas & Schimenti 2012). ring finger protein 212 (*Rnf212*) gene has been shown to be essential for meiotic recombination in *Caenorhabditis elegans* (Jantsch *et al.* 2004, Bhalla *et al.* 2008), and, in humans, RNF212 has been associated with variation in the genome-wide recombination rate (Kong *et al.* 2008, Chowdhury *et al.* 2009). Moreover, the infertility and CO failure of mice bearing a targeted knockout of the mouse gene, *Rnf212*^{tm1Nhr} (herein referred to as *Rnf212* KO) provide evidence for an essential role of RNF212 in mammalian CO recombination (Reynolds *et al.* 2013).

To discover genes involved in meiosis and reproduction, an unbiased phenotype-driven approach using N-ethyl-N-nitrosourea (ENU)-induced mutagenesis and fertility screening was conducted by the Reproductive Genomics Program at The Jackson Laboratory (JAX) (Lessard *et al.* 2004, Handel *et al.* 2006). From this program, we identified homozygous *repro57* mice as infertile and report herein that the male *repro57* mutant mice exhibit meiotic arrest with defective CO. The *repro57* mice have a nonsense mutation in exon 4 of the *Rnf212* gene, encoding a, RNF212 (also known as Zip3 in budding yeast and ZHP3 in worms), thereby identifying a new allele of *Rnf212*.

Materials and methods

Mice

The *repro57* mice used in this study were produced by the NIH-supported Reproductive Genomics program at JAX (<http://reproductivegenomics.jax.org/>), where each mutant line was designated by a 'repro' number in the order of appearance. Mice of the JF1/Ms (JF1) strain were obtained from the National Institute of Genetics (Mishima, Japan), and C3HeB/FeJ (C3H) mice were obtained from CLEA Japan (Tokyo, Japan). The *repro57* mutation was induced in a C57BL/6J (B6) background and mutagenized mice were subsequently outcrossed to C3H (Handel *et al.* 2006), with creation of a *repro57*-C3H congenic line. For experimental analyses, *repro57* homozygous mice were obtained by mating of *repro57* heterozygous mice. Mice were killed using CO₂ at 10–12 weeks (or at 0–5 weeks of age to follow the first wave of spermatogenesis). All experiments using animals were approved by the animal care and use regulatory committees of both Okayama University and JAX.

Mapping the *repro57* mutation

Adult B6 mice were mutagenized with ENU and the *repro57* infertility phenotype was identified in the third generation by mating with normal mice for 3 weeks. More precise phenotypes were subsequently determined from IVF and histological examination of reproductive tissues (Handel *et al.* 2006). To establish the chromosomal linkage of the *repro57* mutation, genome scans using polymorphic microsatellite markers were performed on DNA obtained from affected and unaffected mice. For fine mapping analysis, 740 F2 mice were produced by mating the F1 mice derived from an intercross between the *repro57* and JF1 mice. The F2 offspring were phenotyped by histology and genotyped using polymorphic microsatellite markers to narrow the candidate region.

DNA sequencing was performed to identify the site of the *repro57* mutation. Genomic DNA was isolated from the liver of *repro57* mutants and founder strain B6 mice. DNA samples were re-extracted once with phenol–chloroform; the 260:280 ratio was ~2.0. Whole-genome libraries were prepared and

subjected to sequencing on the illumina HiSeq 2000 (illumina, CA, USA); sequence analysis was focused on the candidate region identified from mapping.

Bioinformatics analysis

The dataset was subjected to quality control using *NGSQCtoolkit* v2.3 and reads with base qualities greater ≥ 30 over 70% of total read length were used in the downstream analysis (Patel & Jain 2012). High-quality reads were mapped to the mouse genome (build-mm10) using *BWA* v0.5.10-tpx (Li & Durbin 2009). The resulting alignment was sorted by coordinates and further converted to a binary alignment map (BAM) format by *Picard* v1.8.4-*SortSam* utility (<http://picard.sourceforge.net>). *Picard*-*MarkDuplicates* module was used to remove duplicates from the data. Subsequently, *Genome Analysis tool kit* (GATK) v2.2-16 (McKenna *et al.* 2010, DePristo *et al.* 2011) modules *IndelRealigner* and *BaseRecalibrator* were used to pre-process the alignments. The realigned and recalibrated BAM alignment file was used as an input to *GATK-UnifiedGenotyper* (variant caller) at parameters, -stand_call_conf 50.0, -stand_emit_conf 30.0 and -dcov 200, and variant calls were restricted to the target region. Finally, variant calls were annotated by *snpEff* v2.0.5 (Cingolani *et al.* 2012).

Cytological preparations and immunostaining

Histology

Testes from mice were fixed in Bouin's solution and paraffin embedded. Sections (5 µm) were stained with hematoxylin and eosin (HE) for evaluation of spermatogenesis. Immunohistochemistry was performed as described previously (Fujiwara *et al.* 2013). Mouse monoclonal anti-human proliferating cell nuclear antigen (PCNA) primary antibody and anti-rat IgG HRP-labeled secondary antibody (Table 1) were used to label the histological sections. For fluorescence immunostaining, goat polyclonal anti-human DMC1 (Table 1) was used in PBS containing 5% skim milk for 3 h at 37 °C. A donkey anti-goat IgG-DyLight 488 secondary antibody

Table 1 Antibodies for fluorescence immunostaining.

Antibody	Host	Cat. number or reference	Dilution
Anti-βactin	Rabbit	Abcam, Cambridge, UK, ab8227	1:1000
Anti-DMC1	Goat	Santa Cruz, sc-8973	1:50
Anti-γH2AX	Mouse	Upstate, Billerica, MA, USA, JBW301	1:500
Anti-H1T	Guinea pig	Handel Lab, Bar Harbor, ME, USA (Cobb <i>et al.</i> 1999)	1:1000
Anti-MLH1	Mouse	BD Pharmingen, San Jose, CA, USA, 551091	1:50
Anti-PCNA	Mouse	Santa Cruz, sc-25280	1:500
Anti-SYCP1	Rabbit	Novous Biologicals, Littleton, CO, USA, NB300-229	1:100
Anti-SYCP3	Guinea pig	de Massy Lab, Montpellier, France	1:1500
Anti-SYCP3	Rabbit	Chuma Lab, Kyoto, Japan	1:1000
Anti-RNF212	Rabbit	Hunter Lab, Davis, CA, USA (Reynolds <i>et al.</i> 2013)	1:1000
Anti-goat IgG-DyLight 488	Donkey	Jackson Immuno Research, West Grove, PA, USA, 705-485-003	1:500
Anti-guinea pig IgG Alexa Fluor 594	Goat	Invitrogen, A11076	1:200
Anti-guinea pig IgG AMCA	Goat	Jackson Immuno Research, West Grove, PA, USA, 106-156-003	1:500
Anti-mouse IgG Alexa Fluor 488	Goat	Invitrogen, A21121	1:200
Anti-mouse IgG HRP	Goat	Santa Cruz, sc-2005	1:400
Anti-rabbit IgG Cy3	Donkey	Jackson Immuno Research, West Grove, PA, USA, 711-165-152	1:500
Anti-rabbit IgG HRP	Goat	Santa Cruz, sc-2004	1:20 000
anti-rabbit IgG Alexa Fluor 488	Goat	Invitrogen, A11008	1:200

(Table 1) was applied with the same buffer as the primary antibody and incubated for 1 h at RT. Nuclei were counterstained with DAPI, mounted using VECTASHIELD Mounting Medium (H-1200, Vector Laboratories, Inc., Burlingame, CA, USA), and observed using a ZEISS Axio Imager.A1 (ZEISS, Jena, Germany). A TUNEL assay was performed according to the manufacturer's instructions (11684817910, Roche Applied Science).

Surface-spread chromatin preparation

Meiotic cell spreads were prepared as described previously (Peters *et al.* 1997) with some modifications. Briefly, testes were immersed in PBS, and the retrieved seminiferous tubules were washed and immersed in ice-cold hypotonic extraction buffer (30 mmol/l TRIS, 50 mmol/l sucrose, 17 mmol/l trisodium citrate dihydrate, 5 mmol/l EDTA, 0.5 mmol/l DTT, and 0.5 mmol/l phenylmethylsulfonyl fluoride, pH 8.2) for 1 h. segments of the tubules were minced in 100 mmol/l sucrose solution (pH 8.2), and the cell suspension collected was spread onto glass slides, previously immersed in 1% paraformaldehyde containing 0.15% Triton X-100; the slides were placed in a moisture box for 2 h. The slides were washed twice for 2 min with PBS with 0.4% DRIWEL (Fuji Film, co., Ltd, Tokyo, Japan). After blocking with 5% skim milk in PBS, the preparation slides were incubated with primary antibodies diluted in the blocking buffer for 3 h at RT. Secondary antibodies were then incubated for 1 h at RT. The antibodies used were listed in Table 1. The slides were then mounted with VECTASHIELD Mounting Medium with DAPI (H-1200, Vector Laboratories, Inc.) and observed using a ZEISS Axio Imager.A1 (ZEISS).

Air-dried chromosome preparation

The procedure for preparation of chromosomes was as described previously (Fujiwara *et al.* 2013).

Western blot analysis

Testes removed from *repro57* mutant and control (WT) mice were homogenized in radioimmunoprecipitation assay buffer (RIPA: 150 mM NaCl, 10 mM Tris-HCl (pH 7.2), 0.1% SDS, 1.0% Triton X-100, 1% deoxycholate, and 5 mM EDTA). The protein extract was applied on 10% SDS-PAGE and transferred onto Immobilon-P Transfer Membranes (Millipore, Billerica, MA, USA). The membrane was incubated with an anti-RNF212 antibody and HRP-conjugated secondary antibody. Subsequently, the same membrane was incubated with anti-ACTIN antibody after the previous antibodies were removed using Stripping Solution (193-16375; Wako, Osaka, Japan). The antibodies used are listed in Table 1. The membrane was developed using an ECL Advance Western Blotting Detection Kit (RPN2135; Amersham Biosciences) and observed.

Gene expression analysis by RT-PCR

mRNAs were isolated from the testes of WT and *repro57* mutant mice at 10 weeks of age using TRIzol (Invitrogen) following the manufacturer's instructions. Genomic DNA in the RNA sample was digested using DNase I (Takara Bio, Inc., Shiga, Japan), and the RNA solutions were purified by phenol/chloroform

treatment. cDNAs were synthesized by RT reaction using Superscripts III reverse transcriptase (Invitrogen) and Oligo dT primer (Invitrogen). RNAs in the cDNA solution were then digested with RNase H (Toyobo, Osaka, Japan). Given evidence for splice variants of *Rnf212* (Reynolds *et al.* 2013) (http://www.ncbi.nlm.nih.gov/nuccore/XM_006535331.1, http://www.ncbi.nlm.nih.gov/nuccore/XM_006535332.1, and http://www.ncbi.nlm.nih.gov/nuccore/XM_001476621.5), the sequences of the variant-specific primers, how they align to the exonic structure, and the PCR cycles are shown in Supplementary Fig. 1., see section on supplementary data given at the end of this article.

Statistical analysis

Data were expressed as mean $\pm$ S.E.M. The statistical significance of differences in mean values was assessed by Student's *t*-test.

Results

Infertility and meiotic defects in the testes of *repro57* mutant mice

Both female and male *repro57* mutant mice on a congenic C3H background failed to produce offspring after natural mating with normal heterozygous or WT mice for more than 2 months. Heterozygotes were fully fertile with an average litter size of 6.9 ± 1.9 (Table 2). Adult homozygous *repro57* mice of both sexes showed no apparent morphological defects in non-reproductive organs, and there was no difference in body weight between the mutant and WT males. However, the testis weight of mutant mice was approximately half that of WT mice (Fig. 1A and B), suggesting germ cell depletion. Histological examination of testes from mutants at 12 weeks of age revealed many degenerating spermatocytes at late prophase stages, with no mature spermatozoa observed in the seminiferous tubules of *repro57* mice (Fig. 1D), compared with testes of WT mice (Fig. 1C). A TUNEL assay confirmed increased numbers of apoptotic germ cells in testes of *repro57* mice compared with WT mice (Fig. 1E), with cell death specific to metaphase spermatocytes (Fig. 1F).

To determine developmental onset of the phenotype, we examined juvenile WT and *repro57* testis histology. At 14 days *postpartum* (dpp), there were no abnormalities in either WT or *repro57* mice (Fig. 2A and B). However, by 21 dpp, the seminiferous tubules of WT mice were filled

Table 2 Mating results.

Female	Male	<i>n</i>	Number of pups
<i>repro57/repro57</i>	<i>+ / repro57</i>	4	0
<i>+ / repro57</i>	<i>repro57/repro57</i>	3	0
<i>+ / repro57</i>	<i>+ / repro57</i>	10	6.9 ± 1.9

n, number of mating pairs. Number of pups per litter is expressed as mean $\pm$ S.D.

Figure 1 Histology of the testis of WT and *repro57* mice. (A) Testes of WT (left) and *repro57* (right) mice. (B) The paired testis weights (TW in mg, white bars) and testis-body weight ratios (TW/BW, gray bars) were significantly reduced in *repro57* mice ($n=14$) compared with those of WT mice ($n=8$). Bars indicate s.e.m.; asterisk indicates $P < 0.005$. (C and D) HE-stained testis sections show degenerating spermatocytes in the seminiferous tubules of *repro57* mice (arrows in D). (E and F) Apoptotic cell death was observed only in metaphase spermatocytes in the testis of *repro57* mice. P, pachynema; D, diplonema; M, metaphase spermatocyte. Scale bars; 50 μ m in C, D, G, and H, and 200 μ m in E and F.

with round spermatids (Fig. 2C), while those of *repro57* mice exhibited degenerating germ cells and no spermatids (Fig. 2D). Reflecting this, testis weights of *repro57* mice were significantly lower than that of WT mice at 21 dpp (Fig. 2E). We examined PCNA, which plays an important role in both the DNA synthesis machinery of proliferating mouse spermatogonia (Yazawa *et al.* 2000) and meiotic progression (Roa *et al.* 2008). The distribution of PCNA was similar in the testes of WT and *repro57* mice (Fig. 2F and G).

Cytological CO in *repro57* spermatocytes

We assessed marker proteins of meiotic prophase progression to determine abnormalities of meiotic chromosomal dynamics in *repro57* mutant testes. The distribution of DMC1, an early recombination repair protein, is similar in WT and *repro57* germ cells (Fig. 3A and B). We determined the pattern of staining for γ H2AX (phosphorylated histone H2AFX), which accumulates at sites of meiotic DNA DSBs in early prophase and, later, is restricted to the XY body (Fig. 3C). The *repro57* mutant spermatocytes exhibited normal distribution of γ H2AX

during the pachytene stage (Fig. 3D). However, 30% of *repro57* mutant pachytene spermatocytes showed XY dissociation (Fig. 3D, $n=260$ cells), while only 3.7% of WT pachytene spermatocytes exhibited XY dissociation (Fig. 3C, $n=135$ cells).

To identify COs, we immunolabeled for MLH1 foci, which mark COs during mid- to late-pachytene substages and for the testis-specific histone H1 variant HIST1H1T (herein referred to by its common designation of H1t), which is expressed in spermatocytes at post-mid-pachytene stages (Cobb *et al.* 1999). H1t-positive spermatocytes in the WT mice contained normal numbers of MLH1 foci (1–2 foci on each chromosome) (Fig. 4A). However, almost all H1t-positive homozygous

Figure 2 Developmental analysis of testes in WT and *repro57* mice. (A, B, C, and D) HE-stained sections of the testis from WT (A and C, $n=5$ and 6 mice at 14 and 21 dpp respectively) and *repro57* (B and D, $n=4$ and 3 mice at 14 and 21 dpp respectively) mice during the first wave of spermatogenesis. G, spermatogonia; L/Z, leptoneuma/zygonema; P, pachynema; R, round spermatid. Arrows indicate degenerating germ cells. (E) Testis weights for WT and *repro57* mice during the peri-pubertal period of development. Bars indicate s.e.m.; asterisk indicates $P < 0.05$. (F and G) Immunolabeling for PCNA on testis sections from prepubertal WT (A) and *repro57* (B) mice at 14 dpp. Scale bar, 50 μ m.

Figure 3 Analysis of DNA damage repair and XY body dynamics in WT and *repro57* spermatocytes. (A and B) Sections from 14 dpp testes were immunolabeled for DMC1. Scale bars, 50 μ m. (C and D) Spread spermatocyte nuclei were immunolabeled for SYCP3 (red) and γ H2AX (green), with magnified views of the XY bodies (insets), revealing XY chromosome dissociation in *repro57* mutant spermatocytes. (original magnification of E, F = $\times 1000$).

mutant spermatocytes showed complete absence of MLH1 foci (Fig. 4B), although a few had 1–8 MLH1 foci per cell (Fig. 4C). We also scored COs by labeling diplotene spermatocytes for co-localization of SYCP3 and SYCP1 in partially desynapsed diplotene spermatocytes. In contrast to WT spermatocytes, the *repro57* mutant spermatocytes lacked SYCP1–SYCP3 co-localization (a mark of CO between homologs).

However, homologs in mutant germ cells were partially associated at centromere (Fig. 4E), as were homologs in WT diplotene spermatocytes (Qiao *et al.* 2012; Fig. 4D). At diakinesis/metaphase I, all 20 chromosomes exhibited CO in WT spermatocytes (Fig. 4F), but only a few chromosomes in *repro57* mutant spermatocytes exhibited chiasmata (Fig. 4G). Consistent with the lack of chiasmata, there were a significantly increased number of chromosomes (univalents) in mutant spermatocytes compared with WT spermatocytes (Fig. 4H).

A mutation in the *Rnf212* gene in *repro57* mice

The *repro57* mutation was mapped to ~ 3 Mb region of Chr. 5 between D5Mit338 and D5Mit117 by linkage analysis using 82 affected F2 male mice (Fig. 5A). This region includes 70 genes; among these, *Rnf212* was a strong candidate gene for harboring the *repro57* mutation. Similar to *repro57* mutant mice, mice with a targeted gene knockout of *Rnf212* exhibit male and female infertility, with male germ cells arrested at late meiotic prophase (Reynolds *et al.* 2013). High-throughput sequencing of DNA from homozygous *repro57* and WT mice revealed a C-to-T transversion at nucleotide 265 of the *Rnf212* mRNA (at 108757107 bp in Chr5, GRCm38.p2), creating a premature stop codon within the coiled-coil domain of RNF212 (Fig. 5B). This mutation was common to all mice with the *repro57* phenotype and, moreover, no other exonic mutation was detected within the *repro57* region. We assessed expression of *Rnf212* transcript variants in

Figure 4 Analysis of crossing-over in WT and *repro57* spermatocytes. (A, B, and C) Pachytene spermatocytes from WT (A) and *repro57* homozygous mice (B and C) immunolabeled with SYCP3 (red) and MLH1 (green). MLH1 foci are absent in most H1T-positive *repro57* spermatocytes (B); however, some mutant spermatocytes exhibited 1–8 weak MLH1 foci (C). Arrowheads indicate MLH1 focus. (D and E) Diplotene spermatocytes immunolabeled with SYCP3 (red) and SYCP1 (green) from (D) WT, with a magnified view of a chromosome with chiasma and (E) *repro57*, with a magnified view of a chromosome lacking CO. (F, G, and H) Giemsa-stained chromosome of diakinesis/metaphase I spermatocyte from WT (F) and *repro57* mice (G). Arrows in G indicate bivalent chromosomes with chiasma, but many lack chiasmata. Original magnification $\times 1000$. (H) The number of chromosomes, including both bivalent and univalent, was compared ($n=21$ cells for WT and $n=60$ cells for *repro57*). Bars indicate S.E.M.; asterisk indicates $P<0.005$. Data are expressed as mean $\pm$ S.E.M.

Figure 5 Fine mapping of *repro57* mutation and loss of RNF212 expression. (A) Segregation of alleles of seven microsatellite markers in 82 affected mice of the F2 progeny obtained from crosses between +/*repro57* and JF1 mice. The positions of the microsatellite markers in the critical region are indicated in cM according to the mouse genome sequence (NCBI Build 38) and the region that contains the *repro57* locus is indicated by a bar. (B) Schematic diagram of the *Rnf212* transcript indicating the position of the nonsense mutation in exon 4 of *Rnf212*. (C) Western blot analysis revealed no expression of RNF212 in the testis of adult *repro57* mice, and a single band was detected only in the testes of WT mice. Non-specific bands were detected in the testes of both WT and *repro57* mice. ACTIN was used as an internal control.

WT and *repro57* testes. Three variants contain the RING finger domain (Reynolds *et al.* 2013); bioinformatics analysis revealed that 5' end of two other transcripts contains variant-specific sequences, lacking the RING finger domain (GRCm38.p2, C57BL/6J). RT-PCR analysis of WT and *repro57* testis RNAs revealed expression of variants a, b, c, and e in both WT and *repro57* mice, while variant d was not detected in either WT or *repro57* mice (Supplementary Fig. 1).

To determine whether RNF212 protein is absent in *repro57* mice, we performed western blot analysis using whole testis lysates, with a polyclonal antibody raised against the full-length mouse RNF212. In WT testes, a single RNF212 band (correlating to splice variant a) was detected at ~34 kDa, while no RNF212, or even a truncated protein, was detected in *repro57* mutant testes (Fig. 5C). Non-specific bands at higher molecular sizes were observed in both WT and *repro57* lysates.

Discussion

A moderate-scale unbiased mutagenesis effort coupled with phenotype screening is a powerful tool to create animal models for human disease. We identified the *repro57* fertility model by a fertility screen in such an ENU mutagenesis program. We find that there is a nonsense mutation in the *Rnf212* gene in *repro57* mutant mice. Similar to the *Rnf212* knockout phenotype, the *repro57* mutant mice exhibit male infertility, arrest of spermatogenesis in meiosis, and defects in cytological markers of recombination and chiasma formation. These observations identify a new allele of the mouse *Rnf212* gene, *Rnf212^{repro57}*.

A nonsense mutation in *Rnf212*, encoding a meiotic recombination factor, in *repro57* mutants

Deep sequencing revealed a nonsense mutation in exon 4 of the *Rnf212* gene in *repro57* mutant. This mutation could

truncate the RNF212 protein, ablating the coiled-coil and serine-rich domains. Interestingly, both this study and the previous study (Reynolds *et al.* 2013) find that only a single RNF212 isoform is expressed in the testis, in spite of evidence for several splicing variants. This could suggest that the variants are not translationally competent.

There is considerable evidence for essential function of RNF212 in meiosis in a variety of species. *Rnf212* was shown to be essential for meiotic recombination in *Caenorhabditis elegans* (Jantsch *et al.* 2004, Bhalla *et al.* 2008). In human, variants of RNF212 have been associated with different genome-wide recombination rates (Kong *et al.* 2008, 2014, Chowdhury *et al.* 2009). Creation of a targeted mutation, *Rnf212^{tm1Nhr}*, revealed that mouse RNF212 was essential for CO formation; the *Rnf212* KO spermatocytes lack chiasmata and exhibit depletion of spermatids and mature spermatozoa (Reynolds *et al.* 2013). These phenotypes of *Rnf212* KO mutant mice resemble those of the *repro57* mutant mice, providing evidence that the new *Rnf212* allele, *Rnf212^{repro57}*, is responsible for infertility of the *repro57* mice. Final confirmation that the new *Rnf212^{repro57}* allele is solely causative of the *repro57* infertility phenotype awaits a formal genetic complementation test.

Spermatogenic and meiotic defects in *repro57* mice

Many aspects of the spermatogenesis and meiotic phenotype of *Rnf212^{repro57}* mutant males reflect those found in *Rnf212* KO males. Even during the juvenile onset of spermatogenesis, degenerating and apoptotic spermatocytes were observed (at 21 dpp). However, the normal expression pattern of PCNA in the mutant testes suggests that mitotic proliferation and meiotic initiation are normal in the absence of RNF212. Similar to mice with the *Rnf212* KO mutation (Reynolds *et al.* 2013), *Rnf212^{repro57}* spermatocytes form the SC and exhibit synapsis. Interestingly, another RNF212 homolog ZHP3

is also not required for SC formation (Jantsch *et al.* 2004); however, yeast *Zip3* mutants exhibited defective SC formation (Agarwal & Roeder 2000). Evidence that DNA DSBs are repaired in *Rnf212*^{repro57} spermatocytes includes apparently normal localization of DMC1, which is essential for DNA strand invasion (Moens *et al.* 2002) and redistribution of γ H2AX by mid-prophase in mutant spermatocytes. However, mutant *Rnf212*^{repro57} spermatocytes, similar to *Rnf212* KO spermatocytes, fail to form COs as marked by MLH1 foci, exhibiting lack of chiasmata in most homologous chromosome pairs. Together, observations on mutant spermatocytes for either of the two different *Rnf212* alleles suggest that mouse RNF212 plays a role in maturation of CO events in meiotic recombination, but the mechanism is still not clear. Recent evidence suggests that small ubiquitin-like protein (SUMO) plays an important role in assembly and disassembly of SC by regulating protein–protein interaction during meiosis (Watts & Hoffmann 2011). In this context, it is interesting that the mouse RNF212 mediates SUMOylation, with SUMO modification stabilizing the association of MutS γ with nascent CO intermediates (Reynolds *et al.* 2013). Human RNF212 and yeast homolog Zip3 contain a RING-finger domain and have ubiquitin-ligating activity for post-translational protein modification (Deshaies & Joazeiro 2009), and Zip3 has also been reported to be involved in SUMO modification (Perry *et al.* 2005, Cheng *et al.* 2006). In this context, it is relevant that the *repro57* mutation in the *Rnf212* gene may still lead to a protein with an intact RING domain. Loss of RING domain in the targeted *Rnf212* KO allele resulted in absence of MLH1 foci in germ cells (Reynolds *et al.* 2013), and although the majority of *Rnf212*^{repro57} spermatocytes also lacked MLH1 foci, a few had a markedly reduced number of MLH1 foci and chiasmata, suggesting that the *Rnf212*^{repro57} allele could be hypomorphic in this respect. As much of the value of different gene alleles lies in the ability to tease apart protein domain function, further experimental comparisons of the two mutated alleles are warranted.

***Rnf212* infertility models**

These results establish the *Rnf212*^{repro57} mice as a single-base change model for male infertility, suggesting the possibility that some human male infertility could be attributed to RNF212 mutation. Although the initial screening of the *repro57* mice suggested only male infertility (<http://reproductivegenomics.jax.org/mutants/G1-586-2.html>), female *repro57* homozygous mice on the congenic C3H background also exhibited infertility. Interestingly, the C3H congenic *Rnf212*^{repro57} males exhibited a remarkably higher rate of XY dissociation than the *Rnf212* KO mutants on a B6 background (Reynolds *et al.* 2013). Given also that recent studies have shown genomic variations between subspecies to cause alteration of phenotypes (Keane *et al.* 2011,

Yang *et al.* 2011), our findings indicate that phenotypic differences between *Rnf212* mutants could be due to not only the allelic difference but also genetic background effects.

In summary, this study identifies a new allele of the mouse *Rnf212* gene, *Rnf212*^{repro57}, and shows its association with male fertility. These results bolster genetic evidence that RNF212 is essential for CO and chiasma formation during meiosis. As COs promote the accurate segregation of homologs, thereby avoiding aneuploidy, the mouse *Rnf212* mutations provide important models for investigating the requirements for accurate gametic chromosome segregation. In humans, chromosome aneuploidy leads to infertility, especially in males (Egozcue *et al.* 2000), congenital defects such as down syndrome, or high rates of miscarriage due to defective embryo development (Ambartsumyan & Clark 2008). Thus, genetic models such as these continue to be valuable for human genetic disease.

Supplementary data

This is linked to the online version of the paper at <http://dx.doi.org/10.1530/REP-14-0122>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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