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### Authors

Lei, Wen-Long  
Han, Feng  
Hu, Meng-Wen  
et al.

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# Protein phosphatase 6 is a key factor regulating spermatogenesis

Wen-Long Lei<sup>1,2</sup> · Feng Han<sup>1,3</sup> · Meng-Wen Hu<sup>1</sup> · Qiu-Xia Liang<sup>1,4</sup> · Tie-Gang Meng<sup>1,2</sup> · Qian Zhou<sup>1,2</sup> · Ying-Chun Ouyang<sup>1</sup> · Yi Hou<sup>1</sup> · Heide Schatten<sup>5</sup> · Zhen-Bo Wang<sup>1,2</sup> · Qing-Yuan Sun<sup>1,2</sup>

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## Abstract

Protein phosphatase 6 (PP6) is a member of the PP2A-like subfamily, which plays a critical role in many fundamental cellular processes. We recently reported that PP6 is essential for female fertility. Here, we report that PP6 is involved in meiotic recombination and that germ cell-specific deletion of PP6 by *Stra8-Cre* causes defective spermatogenesis. The PP6-deficient spermatocytes were arrested at the pachytene stage and defects in DSB repair and crossover formation were observed, indicating that PP6 facilitated meiotic double-stranded breaks (DSB) repair. Further investigations revealed that depletion of PP6 in the germ cells affected chromatin relaxation, which was dependent on MAPK pathway activity, consequently preventing programmed DSB repair factors from being recruited to proper positions on the chromatin. Taken together, our results demonstrate that PP6 has an important role in meiotic recombination and male fertility.

## Introduction

Spermatogenesis is a complex developmental process by which male germline stem cells divide and differentiate to

produce mature spermatozoa. In mammalian testes, this process occurs within seminiferous tubules and consists of three phases: mitosis, meiosis, and spermiogenesis [1]. In the second phase of spermatogenesis, meiosis is characterized by chromosome recognition, pairing, synapsis, recombination, and segregation of homologous chromosomes [2]. Meiosis I is different from meiosis II and mitosis in that it involves the exchange of genetic materials during prophase I [3]. This phase is subdivided into leptotene, zygotene, pachytene, diplotene, and diakinesis. Recombination is initiated at the leptotene stage by the formation of double-stranded breaks (DSBs) [4]. DSBs activate the ATM kinase, which performs an essential role in controlling DSB numbers via a negative feedback loop and plays an essential role in shaping DSB distribution across the genome [5, 6]. Any mistake in meiotic recombination can lead to meiotic arrest or chromosome segregation failure, resulting in severe consequences including infertility [7, 8].

Protein phosphorylation and dephosphorylation are essential for spermatogenesis [9]. Changes in the phosphorylation status are mediated by a conserved cohort of protein kinases and phosphatases. For various reasons, protein phosphatases have been much less studied than protein kinases. Among the phosphoprotein phosphatases, PP2A, PP4, and PP6 comprise the type 2A subfamily within the serine/threonine phosphatase gene family [10]. These enzymes play vital roles in multiple aspects of fundamental cellular signal processes [11]. Among type 2A

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✉ Zhen-Bo Wang  
wangzb@ioz.ac.cn

✉ Qing-Yuan Sun  
sunqy@ioz.ac.cn

<sup>1</sup> State Key Laboratory of Stem Cell and Reproductive Biology, Institute of Zoology, Chinese Academy of Sciences, Beijing 100101, China

<sup>2</sup> University of Chinese Academy of Sciences, Beijing 100101, China

<sup>3</sup> Key Laboratory for Major Obstetric Diseases of Guangdong Province, Key Laboratory of Reproduction and Genetics of Guangdong Higher Education Institutes, The Third Affiliated Hospital of Guangzhou Medical University, Guangzhou 510150, China

<sup>4</sup> Department of Reproductive Medicine, Peking University Shenzhen Hospital, Shenzhen, Guangdong 518036, China

<sup>5</sup> Department of Veterinary Pathobiology, University of Missouri, Columbia, MO 65211, USA

protein phosphatases, PP2A is the best known and is extensively studied. PP2A is involved in cell proliferation, signal transduction, DNA repair, and apoptosis [12]. We previously reported that PP2A has an indispensable role in oocyte meiotic maturation, though it is dispensable for folliculogenesis in the mouse ovary [13]. PP4 has been implicated in microtubule organization, histone modifications, DNA damage responses, and signal transduction [14–17]. Although abundantly expressed in cells with multiple roles in cellular signaling, PP6 has received less attention than its close relative PP2A and PP4.

Like other type 2 A Ser/Thr protein phosphatases, PP6 works as a holoenzyme, with regulatory subunits that are evolutionally conserved among eukaryotes, suggesting its fundamental importance. It has been found that mutations in PP6 exist in 9–12.4% melanomas surveyed and PP6 might act as a driver for melanoma development [18, 19]. More recent studies suggested an important role for PP6 in cell/organ size regulation through the Hippo pathway, inflammatory signaling and pre-mRNA splicing [20–22]. PP6 has been found to play a vital role in the G1–S transition and S phase arrest [23–25], as well as in regulating mitotic spindle formation and chromosome segregation by controlling the T-loop phosphorylation state of Aurora A bound to its activator TPX2 [26]. It is also suggested that PP6 is recruited to sites of DNA damage where PP6 may contribute to the dephosphorylation of  $\gamma$ -H2AX [27].

Our laboratory recently reported that a conditional knockout of PP6 in oocytes from growing follicles (by crossing *Ppp6c<sup>F/F</sup>* mice with *Zp3-Cre* mice) caused female subfertility by disrupting the second meiosis completion [28], and specific deletion of *Ppp6c* in *Ppp6c<sup>F/F</sup>;Gdf9-Cre* oocytes from the primordial follicle stage, caused female infertility owing to persistent phosphorylation of H2AX in oocytes, which leads to massive elimination of oocytes and premature ovarian failure [29]. Here, we crossed *Ppp6c<sup>F/F</sup>* mice with *Stra8-Cre* mice to generate mutant mice with specific deletion of the *Ppp6c* gene in male germ cells. We found that the PPP6c-deficient male mice were completely infertile and germ cells were arrested at the pachytene stage during spermatogenesis. Further investigation revealed that the depletion of PP6 in germ cells affected chromatin relaxation, which was dependent on MAPK pathway activity, thus preventing the recruitment of programmed DSB repair factors to the proper positions on chromosomes.

## Materials and methods

### Mice

Mice lacking *Ppp6c* in male germ cells (referred to as *Ppp6c<sup>CKO</sup>*) were generated by crossing *Ppp6c<sup>F/F</sup>* mice with

*Stra8-Cre* mice. Both transgenic mouse lines have C57BL/6J genomic background. The *Ppp6c<sup>WT</sup>* male mice were used as control group. Genotyping PCR for *Ppp6c* was performed using the following primers: forward: GCA-GAGGATGGGGTCACATAG, and reverse: ATCTCT-GAACCAATTCTGGAG. The PCR conditions were as follows: 94 °C for 5 min; 35 rounds of 94 °C for 30 sec, 56 °C for 30 sec, and 72 °C for 30 sec; and 72 °C for 5 min. Genotyping PCR for *Stra8-Cre* was performed using the following primers: forward: GTGCAAGCTGAACAA-CAGGA, and reverse: AGGGACACAGCATTGGAGTC. The PCR conditions were as follows: 94 °C for 5 min; 35 rounds of 94 °C for 30 sec, 66 °C for 30 sec, and 72 °C for 30 sec; and 72 °C for 5 min.

The mice were housed under controlled environmental conditions with free access to water and food. All animal operations were approved by the Animal Research Committee principles of the Institute of Zoology, Chinese Academy of Sciences.

### Antibodies

PPP6C (rabbit, A300-844A; Bethyl Laboratories, Inc.); SYCP3 (rabbit, NB300-231; Novus Biologicals);  $\beta$ -actin antibody (mouse, sc-47778; Santa Cruz); SYCP3 (mouse, sc-74569; Santa Cruz);  $\gamma$ H2AX (rabbit, 9718; Cell Signaling Technology, Inc.); Phospho-ATM (mouse, 4526; Cell Signaling Technology, Inc.); p44/42 MAPK (Erk1/2) (rabbit, 4695; Cell Signaling Technology, Inc.); Phospho-p44/42 MAPK (Erk1/2) (Thr202/Tyr204) (rabbit, 4377; Cell Signaling Technology, Inc.); MVH (mouse, ab27591; abcam); SYCP1 (rabbit, ab15090; abcam); MLH1 (mouse, ab14206; abcam); H3 dimethyl-K9 (rabbit, 07-212; Upstate); Histone H3K14ac (rabbit, BE3225; Easybio); Tak1 (phospho Thr187) (rabbit, YP0424; Immunoway); TAK1 Antibody (rabbit, 12330-2-AP; Proteintech); HMGA2 Antibody (rabbit, 20795-1-AP; Proteintech); HA-Tag mab (mouse, AE008; ABclonal); c-Myc antibody (mouse, m4439; sigma); MRE11 (rabbit) and DMC1 (rabbit) were gifted from Mengcheng Luo. Horseradish peroxidase-conjugated secondary antibodies were purchased from Zhongshan Golden Bridge Biotechnology Co, LTD (Beijing). Alexa Fluor 488-conjugated antibody and Alexa Fluor 594-conjugated antibody were purchased from Life Technologies.

### Breeding assay

Males of different genotypes (8–9 weeks) were used for the breeding assay. Each male mouse was caged with two wild-type C57BL/6J females (7–8 weeks), and their vaginal plugs were checked every morning. The number of pups in each cage was counted within a week of birth.

Each male underwent six cycles of the above breeding assay.

### Immunoblotting

To prepare protein extracts, testis extracts were prepared using a homogenizer in radioimmunoprecipitation assay (RIPA) buffer supplemented with protease and phosphatase inhibitor cocktail (Roche Diagnostics). Germ cell isolated from testes were resuspended in cold RIPA buffer as described above. After transient ultrasound, the testis lysates or germ cell lysates were incubated on ice for 30 min and then centrifuged at 4 °C, 12,000 rpm for 20 min. The supernatant was transferred to a new tube and equal volume loading buffer was added. After being boiled at 95 °C for 10 min, the protein lysates were used for immunoblotting analysis. Immunoblotting was performed as described previously [30]. In brief, the separated proteins in sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS-PAGE) were electrically transferred to a polyvinylidene fluoride membrane. After incubation with primary and secondary antibodies, the membranes were scanned with Bio-Rad ChemiDoc XRS+.

### Tissue collection and histological analysis

For histological examination, at least three adult mice for each genotype were analyzed. Testes and caudal epididymides were dissected immediately following euthanasia. The tissues were then fixed in Bouin's fixative overnight at room temperature, dehydrated in an ethanol series, and embedded in paraffin wax. Then, 5 µm sections were cut with a microtome. After 48 °C overnight drying, the sections were deparaffinized in xylene, hydrated by a graded alcohol series and stained with hematoxylin and eosin for histological analysis, or stained with periodic acid schiff (PAS)-hematoxylin for determining the seminiferous epithelia cycle stages. Images were collected with a Nikon inverted microscope with a charge coupled device (Nikon, Eclipse Ti-S, Tokyo, Japan).

### Immunofluorescence

Testes used for immunostaining were fixed in 4% paraformaldehyde (pH 7.4) overnight at 4 °C, dehydrated, and embedded in paraffin. Paraffin-embedded testes were cut into sections of 5-µm thickness. Then, the sections were deparaffinized, immersed in sodium citrate buffer (pH 6.0) and heated for 15 min in a microwave for antigen retrieval. After blocking with 5% donkey serum albumin, sections were incubated with primary antibodies at 4 °C overnight. Then the sections were incubated with an appropriate fluorescein isothiocyanate-conjugated secondary antibody.

The nuclei were stained with 4',6-diamidino-2-phenylindole. Images were captured using a laser scanning confocal microscope (Zeiss 780 META).

### TUNEL assay

Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay was carried out in accordance with the DeadEnd Fluorometric TUNEL System (Promega BioSciences, Madison, WI, USA). Images were captured using a laser scanning confocal microscope (Zeiss 780 META).

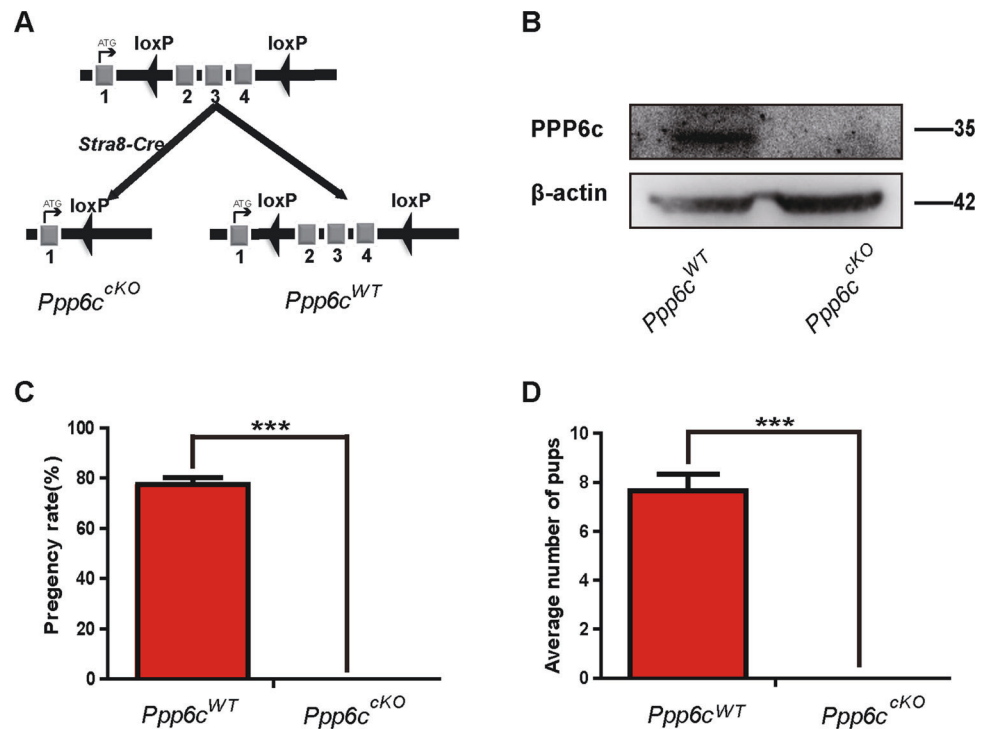
### Spermatocyte surface spreading

Spermatocyte surface spreading was performed according to the drying-down technique as previously described [31]. In brief, testes were dissected and the tubules were washed in phosphate-buffered saline (PBS) pH 7.4 at room temperature. Next, the tubules were submerged in a hypotonic extraction buffer (30 mM Tris pH 8.2, 50 mM sucrose, 17 mM trisodium citrate dihydrate, 5 mM ethylenediaminetetraacetic acid, 0.5 mM dithiothreitol, and 0.5 mM phenylmethylsulfonyl fluoride) for 30–45 min. Subsequently, the tubules were torn into pieces in 100 mM sucrose pH 8.2 on a clean glass slide and then pipetted gently to make a suspension. The cell suspensions were loaded on slides containing 1% paraformaldehyde pH 9.2 and 0.15% Triton X-100. The slides were dried overnight. Finally, the slides were washed with PBS for 10 min and immunostained with antibodies according to the standard protocols mentioned above.

### Isolation of mouse germ cells

Spermatogenic cells were isolated using a protocol previously described by Bellve with minor modification [32]. In brief, testes from adult *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>CKO</sup> mice were removed and decapsulated. The seminiferous tubules were torn into small pieces and incubated in 4.5 ml PBS containing 1 mg/ml collagenase (Sigma, C5138, St. Louis, MO, USA) and 1 mg/ml DNase I (Solarbio, D8071) at 37 °C for 5 min with gentle shaking. Then, the cells were collected by centrifugation at 200 × g for 1 min at 4 °C. Next, the cells were incubated in 4.5 ml 0.25% Trypsin and 1 mg/ml DNase I, and incubated at 37 °C for 5 min with gentle shaking. Thereafter, the cells were collected and washed with 800 µl FBS to cease digestion. Finally, the cells were collected by centrifugation at 600 × g for 6 min at 4 °C and used for immunoblotting analysis. For spermatocytes, after filtration through 40 µm Nylon Cell Strainer, the digested cells were separated by their different sedimentation velocities at unit gravity at 4 °C, using 2–4% bovine serum

**Fig. 1 PP6 is essential for male fertility.** **a** Schematic diagram of deletion of *Ppp6c* exons and creation of *Ppp6c*  $\Delta$  allele by *Stra8-Cre*-mediated recombination in male germ cells. **b** Western blotting analysis of PPP6C protein in *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> spermatocytes of 8-week-old mice.  $\beta$ -actin was detected as an internal control. **c** Pregnancy rates (%) of plugged wild-type females after mating with *Ppp6c*<sup>cKO</sup> or *Ppp6c*<sup>WT</sup> 8-week-old males. **d** Average litter size of plugged wild-type females after mating with *Ppp6c*<sup>cKO</sup> or *Ppp6c*<sup>WT</sup> 8-week-old males. For this part, at least five mice (8-week-old) of each genotype were used for analysis. Data are presented as the mean  $\pm$  SEM.  $P < 0.05$  (\*), 0.01 (\*\*), or 0.001 (\*\*\*)



albumin gradient in Dulbecco's Modified Eagle Medium. Finally, we could collect adequate spermatocytes.

### Plasmid construction and immunoprecipitation (IP)

Mouse *Ppp6c* gene (NM\_024209.3) was cloned into pCS2<sup>+</sup> vector and mouse *Tak1* gene (NM\_009316.1) was cloned into pCMV vector. The plasmids were transfected into 293 T cells using lipofectamine 2000 (11668-019, Invitrogen) according to the reagent protocol. After 36 h, the cells were washed by PBS and then were lysed in buffer for western and IP (P0013, Beyotime) supplemented with protease and phosphatase inhibitors (04693159001 and 04906845001; Roche). Next, the lysis buffer with cells were incubated on ice for 20 min and were centrifuged at 12,000 rpm for 15 min at 4 °C. The supernatant was collected into a new 1.5 ml Eppendorf tube. For IP, 5% of the lysates served as input and the rest were incubated with 1  $\mu$ l HA-Tag antibody and 30  $\mu$ l Protein A/G Magnetic beads (B23201, bimake) at 4 °C for 6 h. Finally, these samples were diluted with an SDS loading buffer, heated at 95 °C for protein denaturalization, and then analyzed by western blotting.

### Statistical analysis

All experiments were performed at least three times. Paired two-tailed Student's *t* test was used for statistical analysis. Data were presented as mean  $\pm$  SEM and  $P <$

0.05 (\*), 0.01 (\*\*), or 0.001 (\*\*\*) was considered statistically significant.

## Results

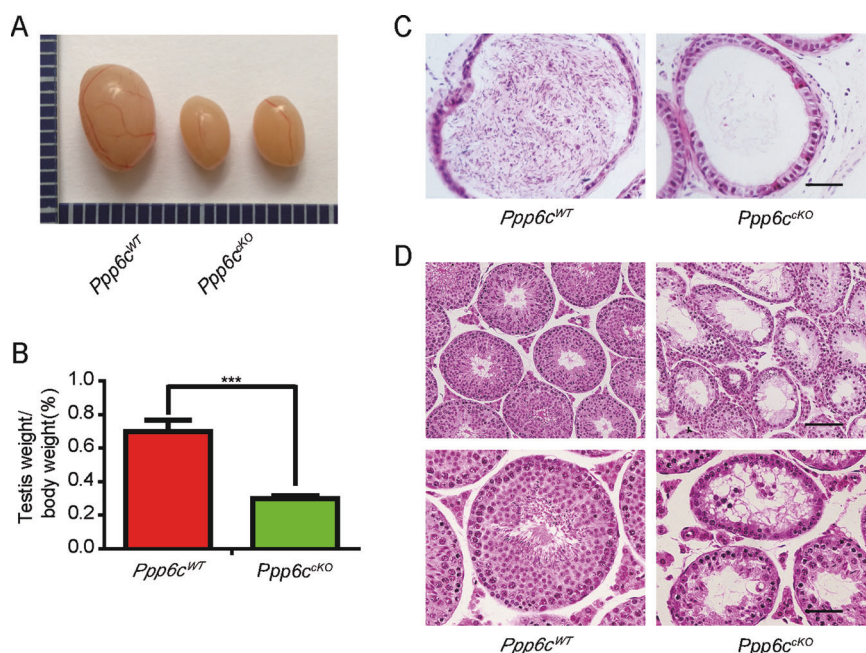
### Germ cell-specific deletion of *Ppp6c* gene causes male infertility

To investigate the function of PP6 in spermatogenesis (referred to as *Ppp6c*<sup>cKO</sup>), we first searched gene information about *Ppp6c* (NM\_024209.3) and found that START codon was located in exon I. So, we generated *Ppp6c*<sup>F/F</sup> mice in which exons II–IV of the *Ppp6c* gene were flanked with Loxp sites [28]. Then, PPP6c was specifically deleted from male germ cells by crossing *Ppp6c*<sup>F/F</sup> mice with *Stra8-Cre* transgenic mice (Fig. 1a). *Stra8-Cre* recombinase is specifically active in early-stage spermatogonia starting at 3 days after birth [33]. The results of western blotting (Fig. 1b) showed that PPP6c was absent in spermatocytes of *Ppp6c*<sup>cKO</sup> mice. Thus, we successfully established male germ cell-specific knockout mice for PP6.

The breeding assays showed that the *Ppp6c*<sup>cKO</sup> male mice were completely infertile (Fig. 1c, d). Although copulatory plugs were routinely observed, no pups were obtained when adult *Ppp6c*<sup>cKO</sup> males mated with normal fertile females. Compared with controls, the testes of *Ppp6c*<sup>cKO</sup> mice were much smaller (Fig. 2a) and the testis weight to body weight ratio of *Ppp6c*<sup>cKO</sup> was significantly



**Fig. 2 PPP6c is required for spermatogenesis.** **a** The testes of *Ppp6c<sup>CKO</sup>* were smaller than those of the *Ppp6c<sup>WT</sup>* (8-week-old, the same as below). **b** Testis weight to body weight ratio of *Ppp6c<sup>WT</sup>* and *Ppp6c<sup>CKO</sup>* mice ( $n = 7$ ). Data are presented as the mean  $\pm$  SEM.  $P < 0.05$  (\*),  $0.01$  (\*\*), or  $0.001$  (\*\*\*) **c** Histological analysis of the caudal epididymides of the *Ppp6c<sup>WT</sup>* and *Ppp6c<sup>CKO</sup>* mice. (Scale bar: 50  $\mu$ m) **d** Histological analysis of the seminiferous tubules of the *Ppp6c<sup>WT</sup>* and *Ppp6c<sup>CKO</sup>* mice. Scale bar: (top) 100  $\mu$ m; (bottom) 50  $\mu$ m.



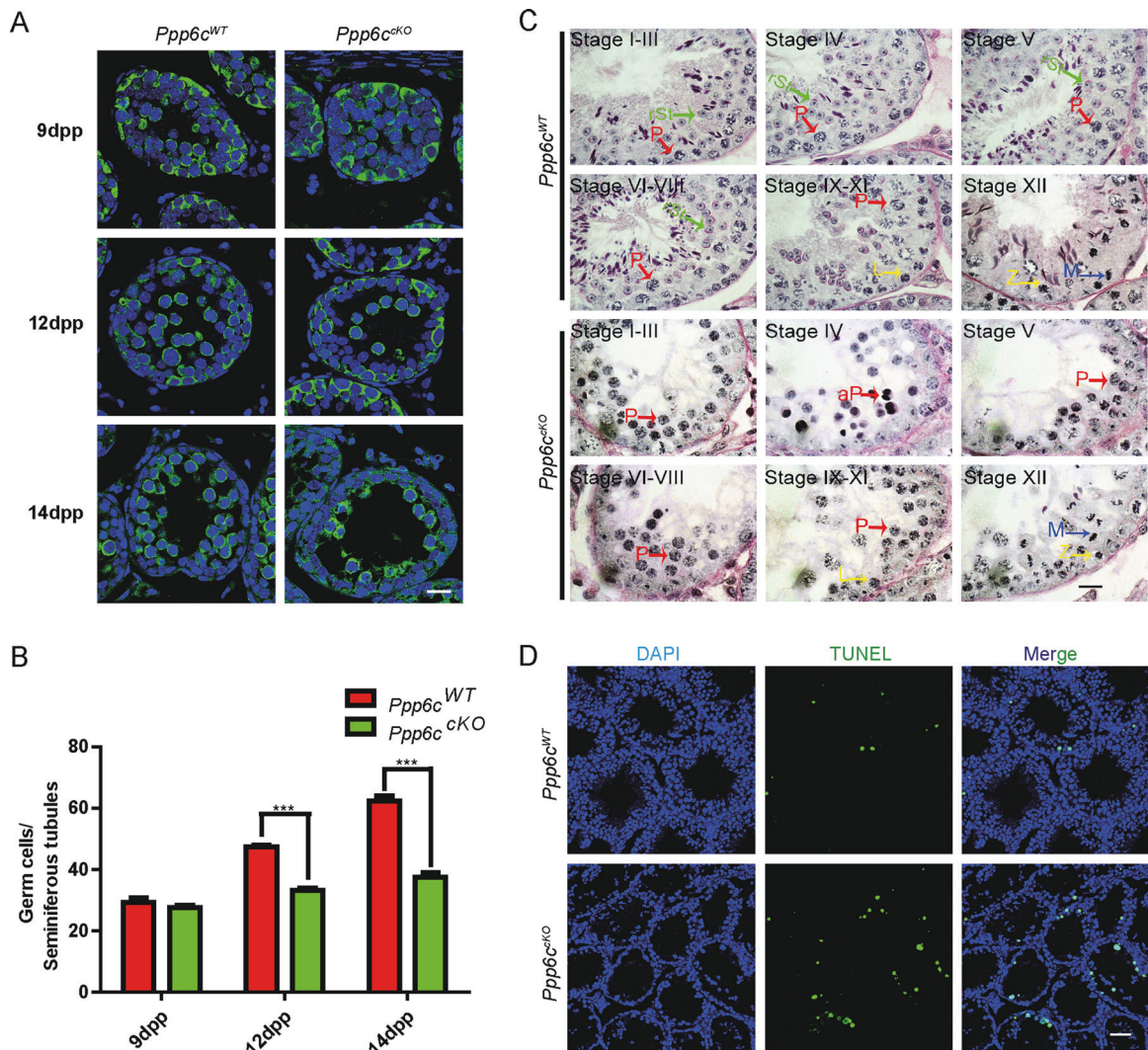
lower (Fig. 2b). We next analyzed the histology of the epididymides and testes by hematoxylin and eosin (H&E) staining. The results showed that almost no mature spermatozoa were found in the epididymal lumens of *Ppp6c<sup>CKO</sup>* mice (Fig. 2c). The seminiferous tubules of *Ppp6c<sup>WT</sup>* testes contained a basal population of spermatogonia, several types of spermatocytes and spermatids. However, germ cells were severely reduced in number, and nearly no round or elongated spermatids were observed in the testes of *Ppp6c<sup>CKO</sup>* males (Fig. 2d, Fig. S1A, and Fig. S1B). These results demonstrate that germ cell-specific PPP6c knockout results in spermatogenesis failure and thus male infertility.

### The PPP6c-deficient spermatocytes arrest at the pachytene stage

To identify exactly which stage of spermatogenesis was affected in PPP6c-deficient mice, we performed H&E and immunofluorescence staining of the germ cell marker MVH to characterize the first wave of spermatogenesis in mice at 9 dpp (days postpartum), when testes begin to generate leptotene spermatocytes, at 12 dpp, when testes begin to generate zygotene spermatocytes, and at 14 dpp, when testes begin to generate pachytene spermatocytes. The results revealed that the numbers of germ cells of *Ppp6c<sup>CKO</sup>* mice were similar to *Ppp6c<sup>WT</sup>* mice at 9 dpp, but these numbers decreased sharply at 12 dpp and 14 dpp (Fig. 3a, b and Fig S2A), suggesting that the prophase of meiosis I was affected in *Ppp6c<sup>CKO</sup>* mice. Spermatogenesis can be subdivided into 12 stages in mouse testes using PAS and hematoxylin staining of sections [34]. To identify which

stages of spermatogenesis were affected after PPP6c deletion, sections were stained with PAS and hematoxylin and we discovered that spermatogenesis of PPP6c-deficient mice was blocked at stages IV and most of the spermatocytes arrested at the pachytene stage (Fig. 3c). The percentage of germ cells was sharply decreased in *Ppp6c<sup>CKO</sup>* mice from 12dpp (Fig. 3b), probably because the pachytene checkpoint-arrested spermatocytes were undergoing apoptosis. To test this possibility, TUNEL assay was performed, and the results showed that germ cells underwent apoptosis in the *Ppp6c<sup>CKO</sup>* mice (Fig. 3d, Fig S2B, and Fig S2C).

Based on the above results, we found that PPP6C-deficient spermatocytes were abnormal. Therefore, its localization was first analyzed by spermatocyte surface spreading. During meiosis I prophase, the localization of PPP6c was consistent during different stages. From leptotene to diakinesis, PPP6c was always localized to the nucleus. PPP6c aggregated strongly throughout the nucleus during the zygotene stage to the diplotene stage (Fig. 4a). Moreover, we identified various stages of meiotic prophase I by staining for a component of the synaptonemal complex, SYCP3 [35]. All meiotic prophase I stages could be found in the spermatocyte nuclei of *Ppp6c<sup>WT</sup>* testes, whereas only spermatocytes from the leptotene to the pachytene stages were observed in the *Ppp6c<sup>CKO</sup>* mice testes (Fig. 4b). Further quantification of the meiotic prophase I stages in testes showed that the proportion of the early-pachytene cells was increased and that of mid-pachytene cells, diplotene cells and diakinesis cells was significantly decreased in the *Ppp6c<sup>CKO</sup>* mice (Fig. 4c). All these results indicate that the depletion of PPP6c leads to a pachytene stage arrest and thus spermatocyte death.



**Fig. 3 PPP6c depletion results in the abnormality and apoptotic death of spermatocytes.** **a** MVH immunofluorescence analysis of the juvenile *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>CKO</sup> mice showed that the amount of the germ cells began to drop from 12 dpp to 14 dpp. (Scale bar: 20  $\mu$ m) **b** Quantification of the germ cells in the seminiferous tubules of the juvenile *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>CKO</sup> mice. For each time point, at least three mice of each genotype were used for analysis. Data are presented as the mean  $\pm$  SEM.  $P < 0.05$  (\*), 0.01 (\*\*), or 0.001 (\*\*\*)

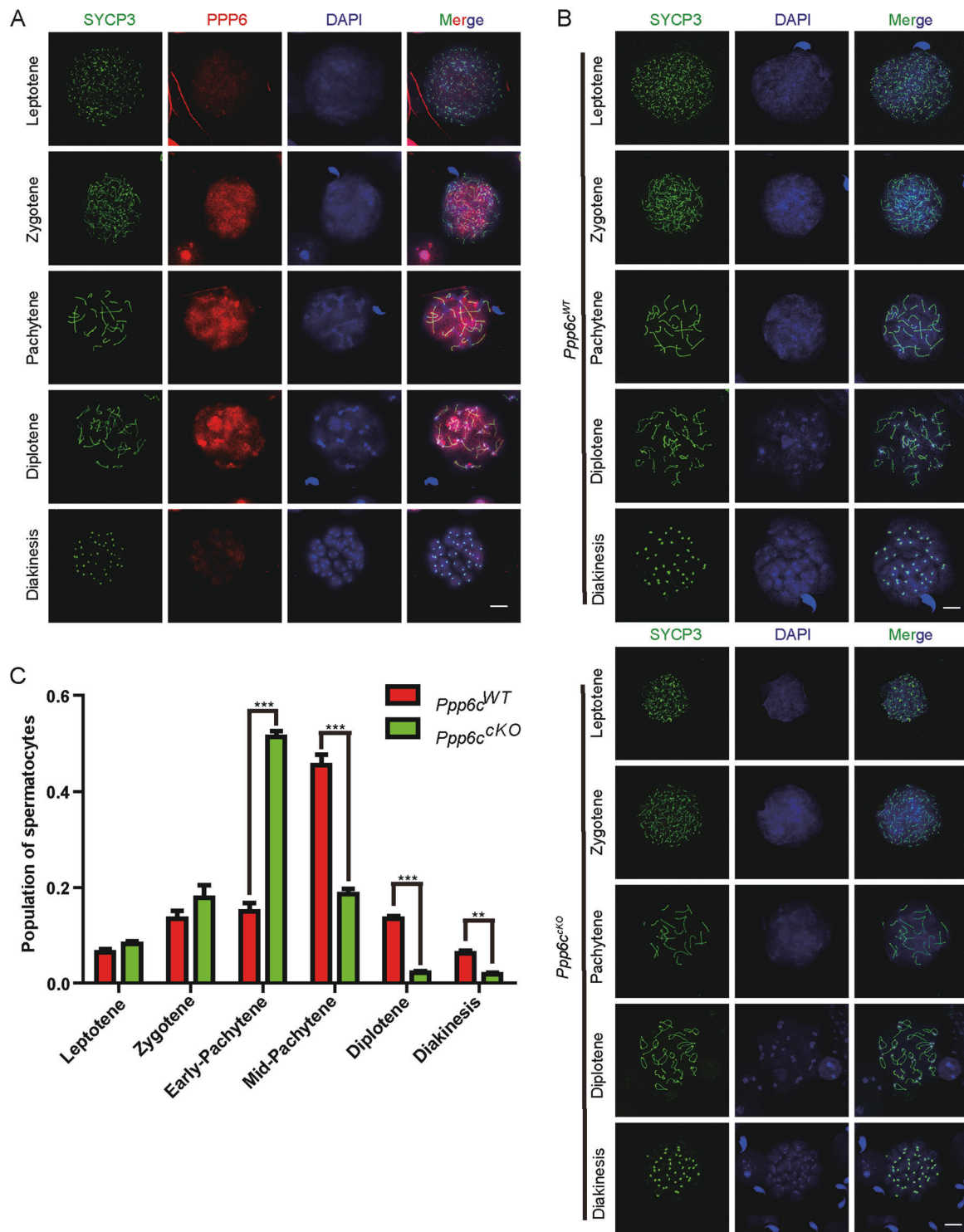
**c** Seminiferous tubules paraffin sections from *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>CKO</sup> testis (8-week-old) were stained with PAS-hematoxylin. L leptotene spermatocytes, Z zygotene spermatocytes, P pachytene spermatocytes, aP apoptotic pachytene spermatocytes, M meiotic divisions, rSt round spermatids. (Scale bar: 20  $\mu$ m) **d** TUNEL immunofluorescence staining of the testes of *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>CKO</sup>. (Scale bar: 50  $\mu$ m) Green: TUNEL positive signal; Blue: DAPI. At least three mice (8-week-old) of each genotype were used for analysis.

### PPP6c depletion causes unsuccessfully programmed DSB repair

According to the above-presented data, *Ppp6c*<sup>CKO</sup> mice displayed defects in the pachytene stage. Programmed DSB repair mainly occurs at this stage. Chromosome synapsis and recombination are facilitated by the introduction of DSB [36], which can be monitored by detecting  $\gamma$ H2AX-positive foci at the leptotene and zygotene stages and these are decreased on autosomes after DSB repair at the pachytene stage [37]. In the *Ppp6c*<sup>CKO</sup> spermatocytes, the  $\gamma$ H2AX signal was detected at the leptotene and zygotene

stages, which was similar to *Ppp6c*<sup>WT</sup> spermatocytes, indicating normal production of DSB (Fig. 5a). However, in the *Ppp6c*<sup>CKO</sup> spermatocytes, the  $\gamma$ H2AX signal was still diffused on the autosomes at the pachytene stage, rather than being accumulated only on the XY bodies (Figs. 5a, b). Consistent with the  $\gamma$ H2AX staining, pATM, which is integral to the regulation of programmed DSB formation [38], was also detected on the autosomes of *Ppp6c*<sup>CKO</sup> spermatocytes (Fig. 5c). These results suggested that PPP6c might be involved in programmed DSB formation and repair. PPP6c depletion caused failure in programmed DSB repair.



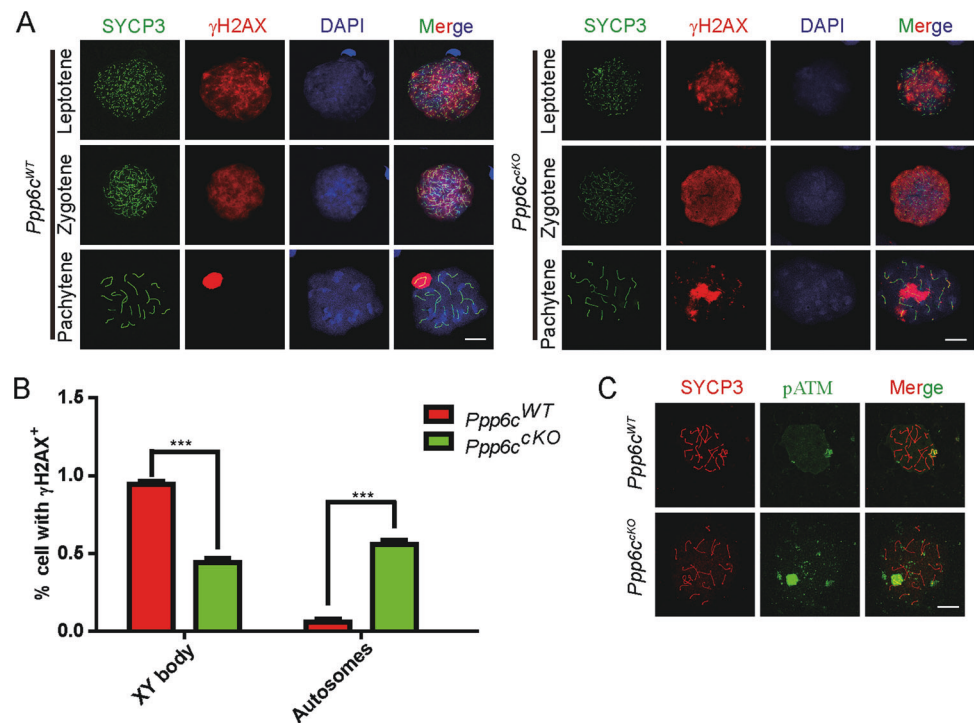


**Fig. 4 PPP6c-deficient spermatocytes arrest at pachytene stage.** **a** Representative images of subcellular localization of PPP6c during spermatogenesis. Spermatocytes (8-week-old) were double stained for PPP6c (red) and SYCP3 (green) at leptotene, zygotene, pachytene, diplotene, and diakinesis stages. (Scale bar: 10  $\mu$ m). **b** Spermatocyte stages in prophase in *Ppp6c<sup>WT</sup>* and *Ppp6c<sup>CKO</sup>* spermatocytes (8-week-

old). Chromosome spreads of spermatocytes were immunostained with antibodies against SYCP3 (green) and DAPI stain (blue). Scale bar: 10  $\mu$ m. (Meiotic stage frequencies in *Ppp6c<sup>WT</sup>* and *Ppp6c<sup>CKO</sup>* testes. For this part, at least three mice (8-week-old) of each genotype were used for analysis. Data are presented as the mean  $\pm$  SEM.  $P < 0.05$  (\*), 0.01 (\*\*), or 0.001 (\*\*\*)).



**Fig. 5 Programmed DSBs are not repaired efficiently due to PPP6c depletion.** **a** *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> chromosome spreads of spermatocytes were immunostained with antibodies against SYCP3 (green) and  $\gamma$ H2AX (red).  $\gamma$ H2AX marked DSB in spermatocytes (8-week-old). Scale bar: 10  $\mu$ m. **b** Percentage of cells retaining H2AX outside the sex chromosomes in *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> pachytene spermatocytes ( $n = 3$ ; 300 cells). **c** *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> chromosome spreads of spermatocytes (8-week-old) were immunostained with antibodies against SYCP3 (red) and pATM (green). Scale bar: 10  $\mu$ m.



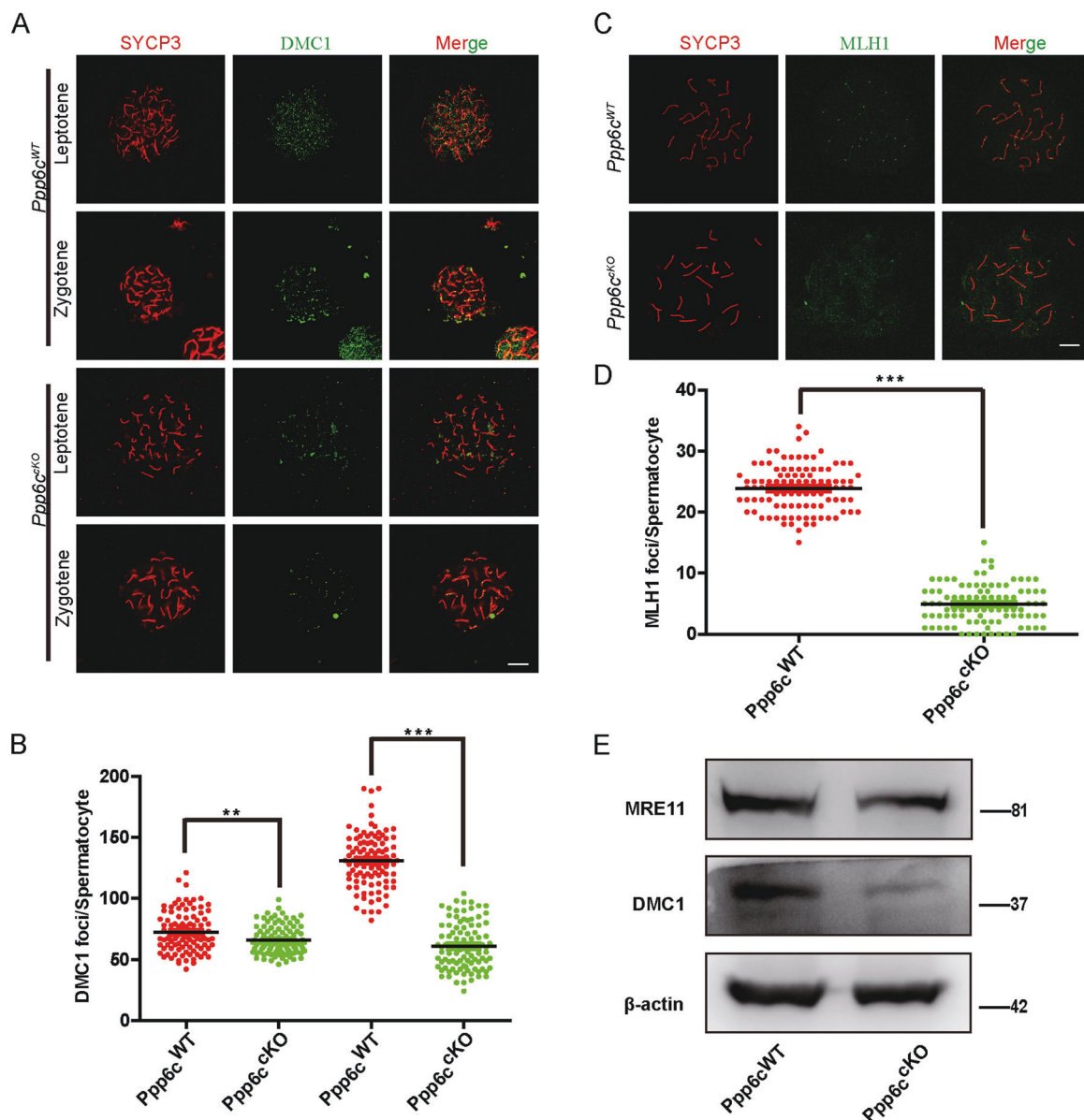
### PPP6c deletion results in impaired meiotic recombination in spermatocytes

In general, defects in meiotic recombination are significant causes for pachytene stage arrest [39]. Meiotic recombination is dependent on the successful repair of programmed DSB. We examined proteins involved in meiosis-specific programmed DSB formation and recombination, such as SYCP1, MRE11, DMC1, and MLH1. SYCP1 is one of the important components of the synaptonemal complexes and it is the central element that links two homologous chromosomes for pairing [35]. The MRN complex, consisting of MRE11, RAD50, and NBS1, is involved in DSB end resection [40]. RAD51 and DMC1 are important marker molecules for different stages of meiotic recombination. BRCA1 associates with RAD51 and RAD51/DMC1 facilitate the invasion of homologous chromosomes [41, 42]. MLH1 represents crossovers during meiotic recombination [43]. First, we examined chromosomal synapsis by using SYCP1 antibody staining and found that SYCP1 signals of *Ppp6c*<sup>cKO</sup> mice were similar to those of *Ppp6c*<sup>WT</sup> mice (Fig S3C). By western blotting, we found that MRE11 decreased dramatically in *Ppp6c*<sup>cKO</sup> spermatocytes (Fig. 6e), suggesting that the MRN complex might be insufficiently recruited to the programmed DSB sites in *Ppp6c*<sup>cKO</sup> mice. The number of DMC1 foci in *Ppp6c*<sup>cKO</sup> spermatocytes was also decreased compared with that of the *Ppp6c*<sup>WT</sup> mice (Fig. 6a, b). Moreover, we performed immunoblotting using germ cell lysates from *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> mice. As shown in Fig. 6e, the levels of DMC1 were all significantly

decreased in *Ppp6c*<sup>cKO</sup> mice, suggesting defects in single strand DNA invasion or generation. We also found that the number of MLH1 foci of *Ppp6c*<sup>cKO</sup> mice was significantly reduced compared with that of *Ppp6c*<sup>WT</sup> mice (Fig. 6c, d), suggesting the crossover formation was also impaired. All these results suggested that PPP6c deletion might influence recombination and crossover formation, finally resulting in a pachytene stage arrest.

### PPP6c promotes chromatin relaxation to regulate meiotic recombination

To investigate the mechanism underlying PPP6c deletion-caused pachytene stage arrest, we found that chromatin relaxation seems to be critical for meiotic recombination initiation and repair [44]. We proposed that PP6 might influence meiotic recombination by affecting chromatin structure. To test this possibility, we examined the acetylation status of H3K14 (H3K14ac), which is an active transcription histone marker, and dimethylation at histone H3K9 (H3K9me2), which is a heterochromatin-associated modification marker [45–47]. By spermatocyte surface spreading and western blotting, we found the level of H3K9me2 was significantly increased in *Ppp6c*<sup>cKO</sup> mice, whereas the level of H3K14ac was sharply decreased in *Ppp6c*<sup>cKO</sup> mice (Fig. 7a, b, and Fig S4), suggesting that PPP6c-deficient spermatocytes could not normally promote chromatin relaxation. Based on the above results, PPP6c depletion phenotypes could be explained by which PPP6c mediates chromatin structures and allows the



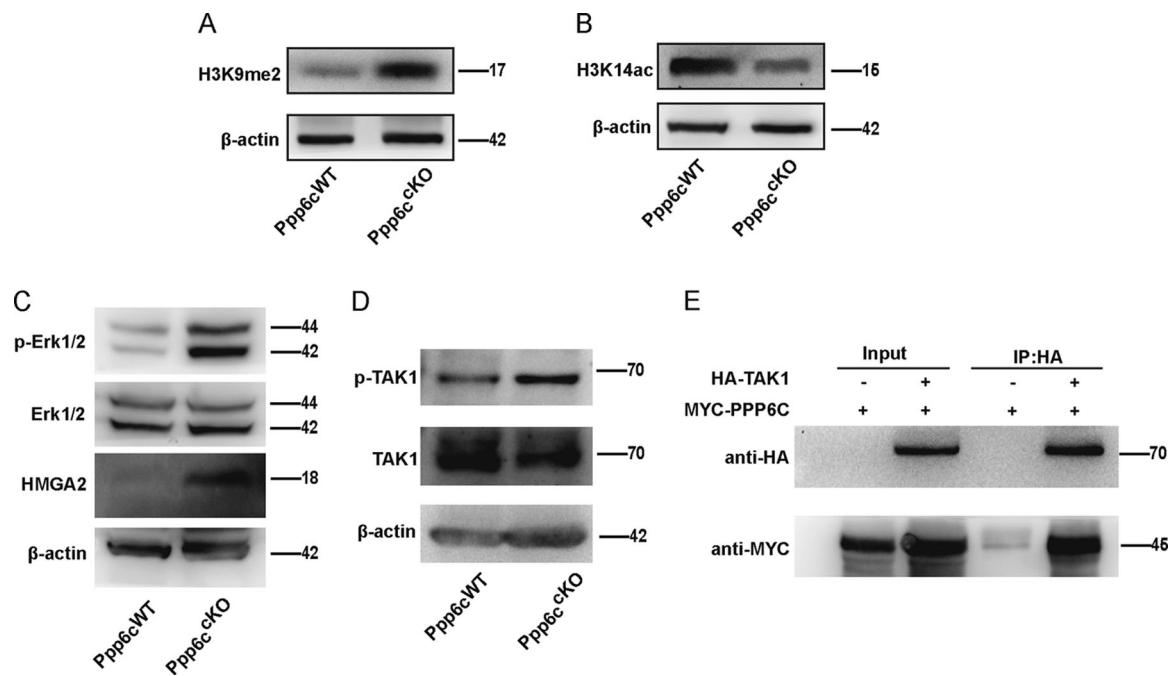
**Fig. 6 Strand invasion, meiotic crossover, and recombination are impaired in *Ppp6c*<sup>KO</sup> spermatocytes.** **a** Spermatocytes stained for SYCP3 (red) and DMC1 (green) in *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>KO</sup> 8-week-old mice. Scale bar: 10  $\mu$ m. **b** Quantification of DMC1 foci in leptotene stage and zygotene stage ( $n = 3$ ; 100 cells). **c** Spermatocytes stained for SYCP3 (red) and MLH1 (green) in *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>KO</sup> 8-week-old mice. MLH1 is a recombination marker in *Ppp6c*<sup>WT</sup>

spermatocytes and decreased MLH1 staining was observed in *Ppp6c*<sup>KO</sup> mice spermatocytes. Scale bar: 10  $\mu$ m. **d** MLH1 signals were counted in spreads. Quantification of the MLH1 foci numbers per cell in the *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>KO</sup> spermatocytes ( $n = 3$ ; 100 cells). **e** Immunoblotting analysis of MRE11, DMC1 in *Ppp6c*<sup>WT</sup>, and *Ppp6c*<sup>KO</sup> germ cells (12 dpp). Level of  $\beta$ -actin was detected as internal control. Each experiment was repeated at least three times.

recruitment of DSB repair factors to the proper position during meiosis.

However, how does the PPP6c mediate chromatin structures? It has reported that the mitogen-activated protein kinase (MAPK) pathway is required for maintaining the chromatin condensation during G2/M progression, whereas chromatin is relaxed in pachytene stage [48]. TAK1 (transforming growth factor-activated kinase 1) is a member of the mitogen-activated protein kinase kinase kinase (MAPKKK) family [49]. Moreover, ERK, as TAK1

downstream kinases, is a member of mitogen-activated protein kinase (MAPK) [50]. Upon activation of the MAPK pathway, p90Rsk2 interacts with and phosphorylates Nek2, which is bound to DNA through interaction with HMGA2. Phosphorylation of HMGA2 by Nek2 decreases its affinity for DNA and favors its release from the chromatin, possibly permitting condensation factors to enter the chromatin and to trigger chromosome condensation [48, 51]. Therefore, we proposed that there may have an interaction between PPP6c and TAK1. PPP6c may mediate chromatin structures



**Fig. 7 PP6 promotes chromatin relaxation to regulate meiotic recombination dependent on MAPK pathway activity.** **a** Immunoblotting detection of the expression of H3K9me2 in spermatocytes of *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> mice (8 week). **b** Immunoblotting detection of the expression of H3K14ac in spermatocytes of *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> mice (8 week). **c** Immunoblotting detection of the expression of p-ERK1/2, ERK1/2, and HMGA2 in spermatocytes of *Ppp6c*<sup>WT</sup> and

*Ppp6c*<sup>cKO</sup> mice (8 week). **d** Immunoblotting detection of the expression of p-TAK1 and TAK1 in spermatocytes of *Ppp6c*<sup>WT</sup> and *Ppp6c*<sup>cKO</sup> mice (8 week). **e** 293 cells were transfected with expression vectors for HA-Tak1 and MYC-PPP6c. The negative controls were transfected with expression vectors for HA and MYC-PPP6c. Proteins from the cell lysates were immunoprecipitated with anti-HA, and immunoblotted with anti-MYC and anti-HA. IP, immunoprecipitation.

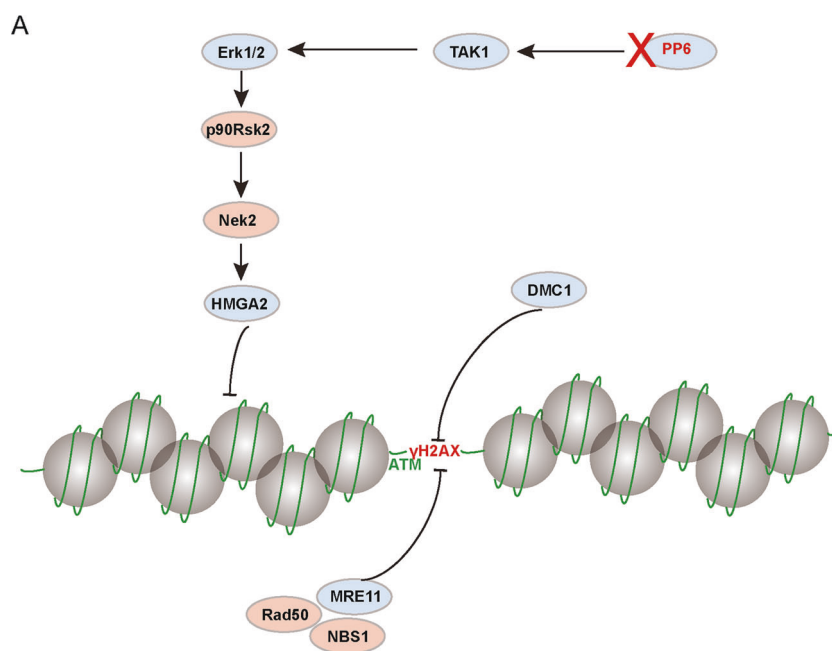
dependent on MAPK pathway activity. To test this possibility, we first performed immunoblotting using spermatocytes from *Ppp6c*<sup>WT</sup> mice and *Ppp6c*<sup>cKO</sup> mice to detect relevant phosphorylated protein levels of MAPK pathway. As expected, the level of p-ERK1/2 and p-TAK1 protein was markedly increased in *Ppp6c*<sup>cKO</sup> mice (Fig. 7c, d), suggesting that the absence of PPP6c affected MAPK pathway activity. Meanwhile, we found that the level of HMGA2 was increased (Fig. 7c), indicating that it was released from the chromatin, finally resulting in chromatin condensation. Moreover, the myc-PPP6c and HA-TAK1 were expressed in 293 T cells for IP experiments. As expected, PPP6c clearly immunoprecipitated with TAK1 (Fig. 7e). Therefore, we confirm that PPP6c interacts with TAK1 and deletion of PPP6c results in premature chromatin condensation.

## Discussion

As a member of the PP2A-like subfamily, PP6 is well known for its functional role in mitosis [26, 52]. Our recent study also revealed a critical role for PP6 in oocyte development [28, 29]. In this study, by crossing *Ppp6c*<sup>F/F</sup> mice with *Stra8-Cre* mice to generate mutant mice with a specific

deletion of *Ppp6c* in male germ cells, we were able to investigate the potential roles of PP6 in male germ cell meiosis and spermatogenesis. We found that PP6 is required for meiosis and fertility in males. Of particular note, in *Ppp6c*<sup>cKO</sup> mice, the spermatocytes arrested at the pachytene stage and programmed DSB repair failed.

Meiosis is a specific process for gamete formation in sexually reproducing organisms, which is generally considered to be the most crucial phase of spermatogenesis. Correct execution of meiosis is essential for fertility, for maintaining the integrity of the genome and for ensuring normal development of the offspring [53]. Meiotic prophase I is subdivided into four cytological stages: leptotene (chromatin condensation, initiation of DSB, and meiotic recombination), zygotene (initiation of synapsis of homologous chromosomes), pachytene (full synapsis, development of recombination sites into at least one crossover per homologous chromosome pair), and diplotene (de-synapsis and visible crossover sites, the so-called chiasmata) [54]. A specific event of this process is meiotic recombination between paternal and maternal chromosomes. Meiotic recombination involves several steps including formation of DSB, DSB end resection, and strand invasion into a chromatid of the homologous chromosome [55]. During meiotic recombination, there are specifically related proteins, such



**Fig. 8** A hypothetical model of the function of PP6 during meiotic recombination. **a** In the *Ppp6c<sup>CKO</sup>* spermatocytes, TAK1 cannot be normally dephosphorylated, resulting in persistent MAPK pathway activation. Upon activation of the MAPK pathway, p90Rsk2 interacts with and phosphorylates Nek2, which is bound to DNA through interaction with HMG A2. Phosphorylation of HMG A2 by Nek2 decreases its affinity for DNA and favors its release from the

chromatin, possibly permitting condensation factors to enter the chromatin and to trigger chromosome condensation. As such, meiotic chromatin cannot be relaxed, which affects the efficient recruitment of the MRN complex (MRE11/RAD50/NBS1) and other DNA repair factors to the DSB sites. Therefore, programmed DSBs cannot be repaired by meiotic recombination, resulting in spermatocyte cell apoptosis and finally male infertility in mice.

as SPO11, MRN complex, BRCA1, DMC1, RAD51, MLH1. In our study, we found that these related proteins decreased dramatically in *Ppp6c<sup>CKO</sup>* spermatocytes (Fig. 6). We speculated that the decreasing presence of these proteins is likely owing to failed recruitment to the programmed DSB sites. Finally, PPP6c depletion causes failure of programmed DSBs repair.

Several components of the chromatin remodeling protein, such as switching defective/sucrose non-fermenting (SWI/SNF) and RNF20, might be involved in chromatin relaxation during meiotic recombination [46, 56]. In our study, male germ cells-specific knockout of PPP6c resulted in a dramatic increase in MAPK pathway activity (Fig. 7). Upon activation of the MAPK pathway, p90Rsk2 interacts with and phosphorylates Nek2, which is bound to DNA through interaction with HMG A2. Phosphorylation of HMG A2 by Nek2 decreases its affinity for DNA and favors its release from the chromatin, permitting condensation factors to enter the chromatin and to trigger chromosome condensation [48]. Consequently, the MRN complex and other recombination protein factors could not be efficiently recruited to the programmed DSB sites, and those DSBs could not be repaired by meiotic recombination, resulting in spermatocyte cell apoptosis and finally male infertility in mice (Fig. 8). Infertility affects 10–15% couples in their reproductive life, and ~20–30% of infertility is caused by male factors [57]. In

humans, male infertility such as azoospermia or severe oligozoospermia is mostly characterized by decreased semen parameters [58]. Azoospermia or severe oligozoospermia is a multifactorial disorder caused by genetic, epigenetic, and environmental factors [59]. The majority (~75%) of male patients with spermatogenic failure are idiopathic, and a genetic factor is often considered to be one of the major causes [60, 61]. The PPP6c-deficient mouse model produces a phenotype that is very similar to some cases of human azoospermia or oligozoospermia. Although there are no reported PP6-associated male infertility patients, the existence of these types of patients should not be discounted. Many cases of male infertility remain undetermined. In summary, our study demonstrated for the first time that PP6 regulates meiotic recombination by promoting chromatin relaxation and facilitating the recruitment of some meiotic recombination-related factors to the chromatin through the MAPK pathway.

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## Compliance with ethical standards

**Conflict of interest** The authors declare that they have no conflict of interest.

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