Cell Autonomous and Nonautonomous Function of CUL4B in Mouse Spermatogenesis* **

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CUL4B ubiquitin ligase belongs to the cullin-RING ubiquitin ligase family. Although sharing many sequence and structural similarities, CUL4B plays distinct roles in spermatogenesis from its homologous protein CUL4A. We previously reported that genetic ablation of Cul4a in mice led to male infertility because of aberrant meiotic progression. In the present study, we generated Cul4b germ cell-specific conditional knockout (Cul4blovlox), as well as Cul4b global knock-out (Cul4blox/lox) mouse, to investigate its roles in spermatogenesis. Germ cell-specific deletion of Cul4b led to male infertility, despite normal testicular morphology and comparable numbers of spermatozoa. Notably, significantly impaired sperm motility caused by reduced mitochondrial activity and glycolysis level were observed in the majority of the mutant spermatozoa, manifested by low, if any, sperm ATP production. Furthermore, Cul4blovlox spermatozoa exhibited defective arrangement of axonemal microtubules and flagella outer dense fibers. Our mass spectrometry analysis identified INSL6 as a novel CUL4B substrate in male germ cells, evidenced by its direct polyubiquitination and degradation by CUL4B E3 ligase. Nevertheless, Cul4blovlox global knock-out males lost their germ cells in an age-dependent manner, implying failure of maintaining the spermatogonial stem cell niche in somatic cells. Taken together, our results show that CUL4B is indispensable to spermatogenesis, and it functions cell autonomously in male germ cells to ensure spermatozooa motility, whereas it functions non-cell-autonomously in somatic cells to maintain spermatogonial stemness. Thus, CUL4B links two distinct spermatogentic processes to a single E3 ligase, highlighting the significance of ubiquitin modification during spermatogenesis.

Protein degradation via the ubiquitin-proteasome system plays a critical role during mammalian spermatogenesis. Timely removal of outlived proteins is crucial to ensure progression through different phases of spermatogenesis including mitotic, meiotic divisions, and postmeiotic morphogenesis. The ubiquitin–proteasome system selects its targets via a group of E3 ubiquitin ligases, which upon interaction with E2 ubiquitin-conjugating enzymes tag substrates for proteasomal degradation by the 26S proteasome. The largest mammalian E3 ligase family is the CRL (cullin-RING finger ubiquitin ligase) family, which includes eight members. Each cullin family member serves as a molecular scaffold and forms an E3 ligase complex with substrate-recruiting receptors and a linker protein (1). The Cul4b gene, located on the X chromosome, shares extensive sequence homology and functional redundancy with the other CRL4 family member, Cul4a. Both proteins employ DDB1 (DNA damage-binding protein 1) as the linker protein, which in turn recruits common or distinct DDB1-CUL4 associated factors for substrate binding. The DDB1-CUL4-mediated protein modification/degradation is involved in critical cellular processes including DNA replication, DNA repair, cell cycle control, and histone modifications (2).

Male infertility accounts for ~40–50% of infertility cases in humans (3, 4). Many factors contribute to male infertility; however, the vast majority of male infertility cases are associated with oligozoospermia (decreased sperm number), asthenozoospermia (reduced sperm motility), and/or teratozoospermia (abnormal sperm morphology). In the mammalian testis, a single pluripotent spermatogonial stem cell (SSC) undergoes several rounds of mitotic divisions followed by meiotic divisions and complex postmeiotic morphogenesis, eventually giving rise to a cohort of mature spermatozoa. We have previously reported that the CRL4 proteins exhibited complementary expression patterns in adult mouse testis, where CUL4A was predominantly detected in primary spermatocytes, whereas CUL4B was highly expressed in Sertoli cells, spermatagonia, and spermatids (5). The absence of CUL4B in spermatocytes is likely due to meiotic sex chromosome inactivation, a transient X chromosome inactivation caused by sex chromosome condensation and subsequent gene silencing (6). However, the

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‡ This article contains supplemental Table S1.

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absence of Cul4a gene expression in Cul4b-expressing cells suggests that these two family members play distinct and non-overlapping/nonredundant roles in specific testicular cell populations. We reported previously that loss of Cul4a in null mutant mice led to male infertility because of meiotic defects (5). Here we investigate the role of Cul4b in murine spermatogenesis. Our previous studies showed that Cul4b mutant mouse embryos died in midgestation because of a G2/M arrest in overlapping/nonredundant roles in specific testicular cell populations. We reported previously that loss of Cul4a in null mutant mice led to male infertility because of meiotic defects (5). Here we investigate the role of Cul4b in murine spermatogenesis. Our previous studies showed that Cul4b mutant mouse embryos died in midgestation because of a G2/M arrest in

Experimental Procedures

**Mice—**Generation of conditional Cul4b mice was described previously (7). The Vasa(Ddx4)-Cre (stock no. 006954), Amh-Cre (stock no. 007915), and Sox2-Cre (stock no. 008454) transgenic mice were purchased from the Jackson Laboratory (Bar Harbor, ME). All mouse colonies were maintained in a barrier facility at Washington University in St. Louis, and all animal experiments were performed in accordance with the institution’s regulations under an approved protocol.

**Electron Microscopy, Histology, and Immunofluorescence (IF)** of the Testis Tissues and Epididymal Spermatozoa—Preparations of testis sections and sperm smears, as well as histological analyses, were performed following protocols described previously (5). Antibodies and dilutions used were: 1:100 for CUL4B (Proteinintech Group, Chicago, IL), PLZF (CalBiotech, Gibbstown, NJ), CUL4A (Bethyl Laboratories, Montgomery, TX), and 1:1000 for Alexa 594 goat anti-rabbit and Alexa 488 goat anti-mouse (Life Technologies, Inc.). Images were captured under a Zeiss AxioSkop 2 epifluorescence microscope. Transmission electron microscopy (TEM) was performed on ultrathin cauda epididymal sections as previously described (5).

**Sperm Motility Assay—**Cauda epididymides were dissected, pierced, and incubated in EmbryoMax® human tubal fluid (EMD Millipore) for 10 min at 37 °C in 5% CO2 to disperse spermatozoa. Sperm motility was determined using a computer-assisted sperm analysis system (Hamilton-Thorne Research, Beverly, MA).

**Flow Cytometric Analysis—**Single testicular cell suspensions and epididymal spermatozoa were prepared as previously described (5). Live spermatozoa were stained with JC1 dye (Life Technologies, Inc.) according to the manufacturer’s instructions, followed by flow cytometry on the BD FACScan™ system (BD Biosciences, San Jose, CA). To prepare splenocyte suspensions, spleen removed from one mouse was meshed through a 70-μm cell strainer (BD Biosciences) with a syringe plunger and washed with cold PBS (calcium- and magnesium-free). After a brief spin, splenocyte pellets were resuspended and incubated in 5 ml of red blood cell lysis buffer (Sigma-Aldrich). The remaining cells were washed once in PBS, centrifuged, and processed for marker staining and FACS analysis.

Isolated splenocytes and testicular cells were incubated in blocking solution (PBS with 1% bovine serum albumin and 1% mouse Fc receptor block (BioLegend, San Diego, CA)) for 15 min, followed by incubation with antibodies diluted in blocking solution for 30 min on ice, protected from light. Dilutions for antibodies and fluorescent dyes used were: 1:100 for anti-mouse GR1-FITC, anti-mouse F4/80-PE, anti-mouse CD11b-APC, and anti-mouse CD45-Pacific Blue, and 1:500 for 7AAD (all antibodies were from AnaSpec Inc., Fremont, CA). Following antibody incubation, the cells were fixed in 4% paraformaldehyde with actinomycin (ACROS Organics, Geel, Belgium) overnight at 4 °C and analyzed on a BD LSRFortessa™ cytometer the following day. The data analyses were completed by using FlowJo V10 software (TreeStar Inc., Ashland, OR).

**Determination of Sperm ATP Level—**Sperm extracts were prepared as previously described (8). The ATP levels were quantified by using the Molecular Probes® ATP determination kit (Life Technologies, Inc.) following the manufacturer’s instructions. Bioluminescence was measured on a GloMax® microplate reader (Promega, Madison, WI), and a standard curve for a series of ATP standard solutions with concentrations ranging from 10 pM to 2.5 μM was generated. The ATP concentration of each sample was determined according to the standard curve, and sperm mass (total protein content) was determined by standard Bradford assay. Normalized ATP levels were calculated as ATP (micromoles)/sperm mass (gram), n = 4 for each genotype in two replicates.

**Determination of Sperm Phosphoenolpyruvate (PEP) Level—**A commercial fluorometric assay kit was used to determine sperm intracellular PEP level (Cayman Chemical Company, Ann Arbor, MI). Epididymal spermatozoa were collected, lysed, and deproteinated in the metaphosphoric acid buffer provided in the kit. Protein pellets were reconstituted, and concentration was determined using a bicinchoninic acid assay (Pierce, Rockford, IL). PEP concentration in each sample was determined by using a standard curve with concentrations ranging from 0 to 100 μM, and the PEP levels of individual samples were determined accordingly. Normalized PEP level was calculated as PEP concentration (μM)/sperm mass (mg). Four biological replicates were used for each genotype assayed in duplicate.

**Affinity Purification and Identification of Ubiquitinated Testicular Proteins—**Seminiferous tubules from control or Cul4bVasa-/- mice were minced in PBS, and the suspension was pipetted repetitively for 1 min. Large fragments were allowed to settle for 15 min, and the remaining cell suspension was centrifuged at 600 x g for 5 min. Cell pellets were resuspended in 1 ml of PBS. To enrich for germ cells, cell suspension was carefully layered on top of a Percoll gradient in a 15-ml conical tube, composed of 1 ml each of 45, 30, 22, and 15% Percoll solutions. After centrifugation at 600 x g for 25 min, cell population in the lower 22% Percoll fraction was recovered by slow pipetting. This fraction contained largely round and elongated spermatids, with some remaining spermatoocytes. Cell pellets were washed three times in PBS and resuspended in 0.2–0.5 ml immunoprecipitation assay buffer (1% Triton X-100, 150 mM
NaCl, 20 mM Tris-Cl, pH 7.4, 1 mM EGTA, 1× protease inhibitor mixture; Roche Applied Science). To affinity-purify ubiquitinated proteins, 10–15 μl of p62-derived ubiquitin-associated (UBA) agarose beads (agarose immobilized recombinant ubiquitin-binding domain of p62/SQSTM1 protein; ENZO, catalog no. UW9010) were added to cell lysates and incubated overnight at 4 °C on a shaking platform. After removing supernatant containing unbound proteins, affinity-purified complexes were pelleted, washed in PBS three times, and submitted to the Proteomics and Mass Spectrometry Program core facility at Washington University School of Medicine for mass spectrometry analysis (detailed method available upon request). Scaffold (Proteome Software, Portland, OR) was used to validate protein identifications using PeptideProphet and to derive corresponding protein ID probability scores (9). The mass spectrometry data processing pipeline and detailed method for peptide quantification were described previously (10).

Cell Culture, Transfection, Immunoprecipitation, and Western Blotting—HEK 293 cells were maintained in DMEM supplemented with 10% fetal bovine serum and 1% penicillin and streptomycin. For the His-tagged ubiquitin-expressing plasmid ubiquitination assay, 80% confluent HEK293 cells were transiently transfected with 2 μg of His-tagged ubiquitin-expressing plasmid (11, 12), HA-CUL4B (generated by Dr. Zhou’s lab), and/or MYC-INS6 (purchased pENTR223-INS6 plasmid from Harvard PlasmID Database, then subcloned INS6 ORF into pCS2-Myc vector in Dr. Zhou’s lab) as specified, and treated with 20 μM MG132 2 h prior to harvesting. Trypsinized cells were pelleted and washed in PBS, and 1/20 of the cell aliquots were boiled in 2× SDS-PAGE buffer for Western blot analysis. The remaining cells were resuspended in immunoprecipitation buffer (0.1 mM NaH2PO4, 10 mM Tris-HCl, pH 8.0, 6 M guanidine HCl, 10 mM imidazole) and sonicated. After brief centrifugation, supernatants were collected and incubated with nickel-nitrotriacetic acid-agarose beads (Qiagen) for 2 h. nickel-nitrotriacetic acid-precipitated proteins were released by boiling the agarose beads in 2× SDS-PAGE buffer, and analyzed by Western blotting. For substrate accumulation assay, 80% confluent HEK293 cells were transfected with 6 μg of INS6-V5-TOPO (purchased pENTR223-INS6 plasmid from Harvard PlasmID Database then subcloned INS6 ORF into pCMV-V5 vector by Gateway system in Dr. Zhou’s lab) alone or with 0, 0.5, 1 or 1.5 μg of HA-CUL4B plasmid, respectively. After 48 h, cells were harvested with Nonidet P-40 lysis buffer containing protease inhibitor mixture (Roche), and 15 μg of cell lysates were directly subjected to SDS-PAGE and immunoblotting. HEK293 cells were transfected with siRNA oligonucleotides (GE Healthcare Dharmacon Inc., Lafayette, CO) using Lipofectamine RNAiMAX reagent (Life Technologies, Inc.). The siRNA sequences are as follows: RBX1, 5’-GACTTTCCC-TGCTGTATCTAA-3’ and for DDB1, 5’-CGTGTGATGG-CAAAAAC-3’. Co-immunoprecipitation was performed following routine procedure after co-transfection of HEK293 with 5 μg of MYC-INS6 and 5 μg of HA-DDB1 (generated by Dr. Zhou’s lab). Antibody dilutions used are: 1:8000 for anti-α-tubulin (ProteinTech), 1:1000 for anti-MYC (Cell Signaling Technology, Danvers, MA), 1:1000 for anti-HA (Cell Signaling Technology), and 1:4000 for anti-V5 (Abcam) 1:500 for β-actin (Santa Cruz).

Results

Germ Cell- but Not Sertoli Cell-specific Cul4b-null Males Are Infertile—We previously demonstrated dynamic and complimentary expression patterns for the two Cul4 genes in the mouse testis and reported that loss of Cul4a led to male infertility caused by defects in meiotic progression (5). Because Cul4b is highly expressed in both somatic and germ cells in the testis, we first used the well established Vasa-Cre line (13) to generate germ cell-specific Cul4b mutants. These germ cell-specific mutants are hereafter referred to as Cul4bVasa, as compared with their wild type congeners, termed Cul4bFl/+. As shown in Fig. 1C and our previous study (5), CUL4B protein is predominantly detected in Sertoli cells, spermatogonia, and spermatids in adult wild type mouse testes. Close examination of wild type seminiferous tubule sections revealed that as spermiogenesis progresses, CUL4B protein gradually localizes to the cytoplasm adjacent to the postacrosomal segment of elongating spermatids, coinciding with the location of the caudal manchette, a transient microtubule-based spermatid structure (Fig. 1B, compare with Fig. 1A, arrows). In the Cul4bVasa testis, however, CUL4B protein was no longer detectable in germ cells, even though its expression was unaffected in Sertoli cells (Fig. 1D). To further confirm this finding, double immunofluorescence staining was performed using antibodies against CUL4B and an SSC marker, PLZF (14). High magnification epifluorescence microscopy showed complete removal of CUL4B from PLZF-positive spermatogonia, as well as round spermatids residing in the same seminiferous tubules (Fig. 1, E–J), demonstrating successful deletion of Cul4b in male germ cells.

To address whether in the absence of CUL4B, spermatocytes turn on Cul4a expression to compensate for the CRL4 function, we performed double immunofluorescence staining of CUL4A and PLZF proteins in the Cul4bVasa testis. No CUL4A staining was detected in PLZF-positive spermatogonia or spermatids of the Cul4bVasa testis (Fig. 1, K–N), suggesting that loss of CUL4B protein does not lead to compensatory Cul4a gene expression in these germ cell populations.

Cul4bVasa males exhibit no defects during development and have a normal life span. However, mutant males are completely infertile. Fertility records showed regular and successful copulatory activities of mutant males, evidenced by the presence of vaginal plugs in females housed in the same cages. However, none of the females mated with Cul4bVasa males produced any pups, in contrast to those mated with control males, averaging 11.2 pups per litter (Table 1).

To address the function of CUL4B in Sertoli cells, we generated Sertoli cell-specific Cul4b conditional knock-out mice using the Amh-Cre mouse (15), hereafter referred to as Cul4bAmh. Interestingly, despite its high expression in Sertoli cells (Figs. 1E and 2A), abolishing Cul4b in these somatic cells did not cause any reproductive phenotype. Immunofluorescent staining on Cul4bAmh testis sections showed complete absence of CUL4B in their Sertoli cells (Fig. 2B); however, these males exhibited normal fertility and seminiferous epithelium histol-
ogy (Fig. 2F). These results suggest that cell autonomous CUL4B function is critical to male germ cell lineage but is dispensable in Sertoli cells.

**Spermatozoa in Cul4b\textsuperscript{Vasa} Testis Are Immotile—**Anatomical examination of the Cul4b\textsuperscript{Vasa} mice revealed normal internal male reproductive organs, including the testes, epididymides, vas deferens, and seminal vesicles. No significant differences in size, weight, or morphology of the testes were noted between the two genotypes. As shown in Fig. 1 (O–R), seminiferous tubule histology of the two genotypes is indistinguishable, and germ cells of all phases are present in the Cul4b\textsuperscript{Vasa} testis.

![Image](image-url)

**TABLE 1**

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<td>Cul4bf/Y\textsuperscript{+/+} (5)</td>
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To investigate the cause of infertility in Cul4b\textsuperscript{Vasa} males, we harvested epididymal spermatozoa from these mice and used the computer-assisted sperm analysis to evaluate the quality of the mutant spermatozoa. As shown in Fig. 3A, Cul4b\textsuperscript{Vasa} produced equivalent numbers of spermatozoa as compared with their wild type counterparts (54.5 ± 5.1 M/ml, n = 4, versus 54.8 ± 5.6 M/ml, n = 5 in controls; p = 0.925). In contrast, sperm motility in the mutants was dramatically compromised: only 6.0 ± 1.4% mutant spermatozoa were progressive motile (n = 4), as compared with 87.2 ± 9.3% in control mice (n = 5, p < 0.0001). We next examined the morphology of epididymal
spermatozoa using epifluorescence microscopy after staining with sperm structure-specific dyes. In typical wild type spermatozoa, the sickle-shaped heads contained highly condensed DNA, which stained positive for DAPI (Fig. 3B, blue), and were covered by a crescent-shaped acrosome stained by peanut agglutinin-lectin (green). The midpiece of the spermatozoa is covered by a helical mitochondrial sheath, and active mitochondria were stained bright red after live spermatozoa were incubated with a cell-permeable fluorescent dye, MitoTracker Red CMXRos (Fig. 3B). Most Cul4bVasa sperms exhibited normal morphology when examined under light microscopy; however, more than half of the mutant spermatozoa stained very weakly or negatively for MitoTracker CMXRos (Fig. 3D, G, and H), suggesting a defect in mitochondrial membrane potential. Intriguingly, a small percentage (6.6 ± 3.4%) of Cul4bVasa spermatozoa exhibited a double-headed phenotype (Fig. 3E and G, asterisk). The individual heads of these spermatozoa were morphologically normal, with properly condensed nuclei and visible acrosomes. Nevertheless, they were often joined at the connecting piece and shared a common flagellum.

To further characterize the impaired mitochondrial activity observed in the Cul4bVasa spermatozoa, we stained epididymal spermatozoa with a vital carbocyanine dye, JC1, which fluoresces at different wavelengths when binds to mitochondria with differential membrane potentials, and analyzed JC1 staining profiles by FACS (16). Fig. 3I showed the FACS results of JC1-stained spermatozoa from representative control and Cul4bVasa mice. On average, 90.2 ± 6.2% of the control spermatozoa (n = 5) exhibited high mitochondrial membrane potential, evidenced by the orange JC1 staining (Fig. 3I and J). However, this population in Cul4bVasa mice was significantly reduced (62.3 ± 10.8%, n = 3, p = 0.045), accompanied by an increase in the population of spermatozoa with low mitochondrial membrane potential stained green with JC1 (Cul4bVasa, 16.8 ± 3.3%, n = 3; control, 6.2 ± 1.1%, n = 5; p = 0.024). These results were consistent with our findings with MitoTracker staining and statistically demonstrated that lack of Cul4b in mouse germ cells led to defective mitochondrial activity in some spermatozoa.

ATP is the energy source that is hydrolyzed by the dynein ATPase to generate the propelling and coordinated force along
the axoneme that enables proper sperm movement (17, 18). The immotile phenotype of Cul4bVasa prompted us to examine ATP production in the mutant spermatozoa by analyzing intracellular ATP level using a previously described method (8). Not surprisingly, we observed low ATP production in Cul4bVasa spermatozoa (2.4 ± 0.5 × 10^3 μmoles/g versus 6.7 ± 1.3 × 10^4 μmoles/g in controls, n = 4, p = 0.002) determined by a bioluminescence assay (Fig. 3K). There is continuous debate on the source of sperm ATP, and two metabolic pathways are believed to be involved in this process. Mitochondria, localized in the sperm tail midpiece, have long been considered to be a major source of ATP production because of their high efficiency of energy production via oxidative respiration (19, 20). Recent evidence, however, suggests that locally produced ATP via less efficient, but high throughput glycolysis in the principle piece of the sperm tail is the major energy source for sperm motility in the mouse (21–23). In our Cul4bVasa mutants, the reduction in mitochondrial membrane potential in spermatozoa may contribute to lowered ATP production. However, loss of ATP production in mutant spermatozoa may also reflect a defective glycolytic pathway. To test this hypothesis, we measured the spermatozoa intracellular PEP level using a fluorometric assay kit. PEP is the intermediate produced during glycolysis that is metabolized to the final product, pyruvate, by pyruvate kinase. We chose PEP instead of pyruvate as the indicator of glycolysis because pyruvate could also come from other metabolic pathways including amino acid metabolism (serine, alanine, and glycine), as well as conversion from lactate by lactate dehydrogenase. As shown in Fig. 3L, a significant decrease in PEP levels was detected in the mutant spermatozoa (Cul4bVasa, 374.8 ± 68.0, n = 3; Cul4bVasa, 116.4 ± 18.8, n = 3, p = 0.017), indicating perturbation of glycolysis in the mutant spermatozoa. Taken together, these results suggest that a lack of ATP production resulting from impaired mitochondrial activity and glycolysis was a major cause of low motility/asthenozoospermic phenotype in Cul4bVasa spermatozoa.

We next examined the ultrastructure of mutant spermatozoa by TEM because a loss of sperm motility is often associated with defective axonemal structure. The TEM analysis revealed that Cul4bVasa mutant sperm axonemes frequently displayed missing, supernumerary, or displaced microtubule doublets (Fig. 3, Q–T). These defects were observed mostly in peripheral microtubule doublets, as opposed to the central microtubule doublet, and it was not clear whether the displaced microtubule doublets were still associated with dynein arms or radial spokes. The mitochondrial sheaths of the mutant spermatozoa appeared morphologically normal (Fig. 3P). These data collectively demonstrated that deleting Cul4b from mouse spermatogonia, and subsequent germ cell populations did not affect the mitotic and meiotic phase of spermatogenesis but compromised spermiogenesis and biogenesis of the sperm flagellum in particular. Lack of Cul4b in these cells resulted in abnormal axonemal microtubule structure and reduced mitochondrial activity in the midpiece, which rendered Cul4bVasa spermatozoa immotile and subsequently male infertile.

**INSL6 Is a Direct Cul4B Substrate in Mouse Testes**—Given the function of Cul4B as an E3 ubiquitin ligase, it is reasonable to hypothesize that in the Cul4b-null germ cells, certain Cul4B substrates may not be efficiently ubiquitinated, which led to defective spermiogenesis. In an attempt to identify potential Cul4B ligase substrates, we performed a mass spectrometry analysis on ubiquitinated proteins purified from enriched spermatids of control and Cul4bVasa mice. Ubiquitinated proteins from pooled spermatid lysates were affinity purified through binding to p62-UBA immobilized agarose beads, and Cul4B target candidates were expected to be under-represented in Cul4bVasa samples. The identity of potential targets were predicted and visualized by Scaffold 4 following mass spectrometry (9). In total, 2504 proteins were identified with a minimum protein threshold of 95.0% and a false discovery rate of 0.6% or lower (supplemental Table S1). Among these, an insulin family member, INS6L, showed high spectrum count in the control testis but was absent in the p62-UBA-enriched testis lysates from Cul4bVasa, suggesting INS6L as a potential Cul4B substrate in the testis. INS6L was previously reported to be highly expressed in germ cells, particularly in spermatids of both rodent and human testes (24) (HPA021364 SIGMA), although its function in spermatogenesis remains unknown. To test the ability of Cul4B to directly mediate polyubiquitination of INS6L, a plasmid encoding MYC-labeled full-length INS6L was co-transfected into HEK293 cells along with His-tagged ubiquitin-expressing plasmid, with or without HA-CUL4B-expressing plasmid. As shown in Fig. 4A, the presence of Cul4B evidently increased polyubiquitinated INS6L, demonstrating that INS6L is a direct substrate for Cul4B. Next HEK293 cells were co-transfected with INS6L-coding plasmid (INS6L-V5) along with various amount of HA–CUL4B plasmids as indicated in Fig. 4B, and a clear Cul4B dose-dependent decrease in INS6L protein level was detected, indicating that INS6L is directly degraded through the Cul4B–mediated ubiquitin-pro-
teasome system mechanism. To further investigate the role of CRL complex in INSL6 polyubiquitination, we tested interactions between INSL6 and two key components of CRL4, the N-terminal binding DDB1 and the C-terminal interacting RING finger protein RBX1 (Rbx1/ROC1/Hrt1) (25). As shown in Fig. 4C, attenuating DDB1 and RBX1 functions, respectively, by RNA silencing in HEK293 cells resulted in reduced levels of INSL6 polyubiquitination. Moreover, DDB1 co-immunoprecipitated with INSL6 after co-transfection of INSL6-V5 and indicated amount of HA-CUL4B plasmid. C, gene silencing by RNAi of DDB1 and RBX1 resulted in reduction of polyubiquitinated INSL6 in HEK293 cells. D, co-immunoprecipitation of DDB1 and INSL6 in HEK293. β-Actin was used as loading control.

**FIGURE 4.** CRL promotes polyubiquitination and degradation of INSL6. A, co-expression of CUL4B resulted in increased amount of multiple slower migrating forms of polyubiquitinated INSL6 in HEK293 cells. Immunoblotting against MYC (Myc-INSL6), HA (HA-CUL4B), and α-tubulin, respectively, were performed on lysates prior to nickel-nitrilotriacetic acid immunoprecipitation. B, immunoblotting against V5 (INSL6-V5), HA (HA-CUL4B), and α-tubulin, respectively, on HEK293 cell lysates, co-transfected with INSL6-V5 and indicated amount of HA-CUL4B plasmid. C, gene silencing by RNAi of DDB1 and RBX1 resulted in reduction of polyubiquitinated INSL6 in HEK293 cells. D, co-immunoprecipitation of DDB1 and INSL6 in HEK293. β-Actin was used as loading control.

IB, immunoblotting; IP, immunoprecipitation; Ni-NTA, nickel-nitrilotriacetic acid.

deletion of Cul4b via Sox2-Cre gave rise to viable Cul4b-null mice (Cul4b<sup>Sox2</sup>) (7). Because Cul4b is expressed in many somatic cell types, we characterized the fertility phenotype of Cul4b<sup>Sox2</sup> males and compared with that of Cul4b<sup>Vasa</sup> to determine whether Cul4b also functions in somatic cells to support spermatogenesis. Cul4b<sup>Sox2</sup> males were completely infertile as expected; however, they exhibited a different and more severe age-dependent testicular/sperm phenotype when compared with their Cul4b<sup>Vasa</sup> counterparts. Younger adult Cul4b<sup>Sox2</sup> (4-month-old) testes exhibited no distinct defects (Fig. 5B), and their sperm phenotype resembled that in the Cul4b<sup>Vasa</sup> mice (data not shown), suggesting relatively normal mitotic and meiotic progression despite impaired spermiogenesis. However, as Cul4b<sup>Sox2</sup> mutants aged (8-month-old), their testes shrank, and many seminiferous tubules started to lose their germ cell populations (Fig. 5D). Such a germ cell depletion phenotype varied between individuals (Fig. 5D, inset), but was not detected in older Cul4b<sup>Vasa</sup> mice (Fig. 5C, inset). As a result, oligoastheno-
CUL4B Regulates Mammalian Spermatogenesis

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sperm smear

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testicular cells

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<td>CD45&lt;sup&gt;+&lt;/sup&gt;CD11b&lt;sup&gt;+&lt;/sup&gt;</td>
<td>CD45&lt;sup&gt;+&lt;/sup&gt;CD11b&lt;sup&gt;+&lt;/sup&gt;</td>
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<td>Cul4b&lt;sup&gt;FF&lt;/sup&gt;</td>
<td>Cul4b&lt;sup&gt;Sox2&lt;/sup&gt;</td>
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zooospermia was often observed in older Cul4bSox2 mutants (Fig. 5F; 54.8 ± 5.6 m/ml, n = 5 in controls; 16.0 ± 13.6 m/ml, n = 4 in Cul4bSox2; p = 0.007). Accompanied by low sperm counts, abnormal morphological phenotypes including deformed sperm heads, double- or triple-headed spermatozoa, and sperm heads detached from the flagella were common in Cul4bSox2 mutants (Fig. 5F, insets). This germ cell depletion phenotype may have resulted from a failure to maintain the SSC niche.

Sertoli cells play a crucial role in SSC niche maintenance (26). We tested the hypothesis whether Cul4b expression in Sertoli cells is required to maintain the SSC niche. As mentioned earlier, knocking out Cul4b in Sertoli cells alone did not cause an infertility phenotype (Fig. 2). We followed these mice as they aged (up to 10 months) but did not observe the germ cell depletion phenotype described above. These data argue against a role for Cul4b in Sertoli cells to maintain the SSC niche and suggest that other somatic populations are responsible for this phenotype. One such somatic cell population could be macrophages. It was estimated that macrophages represent as much as one quarter of the interstitial cell population in rodent testes (27, 28), and they play essential roles in regulating the SSC niche (29). In humans, increased number of testicular macrophages was often reported in patients with male infertility (30). Because the germ cell depletion phenotype in aging Cul4bSox2 mutants was often accompanied by splenomegaly (data not shown), we hypothesized that loss of Cul4b in the spleen may have caused a systemic perturbation of macrophages, which may in turn adversely affected spermatogonial maintenance and self-renewal. To address this possibility, surface marker staining followed by FACS analysis was performed on dissociated spermatocytes and testicular cells of control and Cul4bSox2 animals at 9 month of age. We employed four-color FACS to distinguish different hematopoietic populations: CD45 as a marker for nucleated hematopoietic cells (31), CD11b as a marker for macrophage/microglia (32), F4/80 as a mature macrophage marker (33), and GR1 as a neutrophil marker in peripheral organs (34). The splenocytes isolated from control animals contained 11.05 ± 1.96% CD45−CD11bhigh macrophages/granulocytes of all CD45+ lymphocytes (Fig. 5G), among which 5.81 ± 1.61% were F4/80high GR1low macrophages (n = 5; Fig. 4F). No statistically significant changes were observed in the Cul4bSox2 splenocytes (CD45−CD11bhigh: 12.02 ± 1.65%, n = 6, p = 0.41; F4/80high GR1low: 5.50 ± 1.66%, n = 6, p = 0.76; Fig. 5, H and J). The isolated testicular cells from control animals comprised of 21.12 ± 4.54% CD45−CD11bhigh macrophages/granulocytes of all lymphocytes, and the vast majority of them were macrophages (F4/80high GR1low: 91.84 ± 2.32%, n = 5; Fig. 5, K and M). CD45−CD11bhigh population remained largely unchanged in Cul4bSox2 testes (17.20 ± 5.30%, n = 6, p = 0.22; Fig. 5L); no statistically significant difference in macrophages was noted in these mutants (88.81 ± 3.36%, n = 6, p = 0.11; Fig. 5N). Comparison between the lymphocyte compositions in the spleen and testis of old (10–12 months old) and young (4 months old) adult Cul4bSox2 mice revealed no significant difference. Altogether, these data demonstrated that lack of Cul4b in both germ cells and somatic cells led to male infertility caused by spermiogenesis defects similar to that found in the Cul4bVasa mice and age-dependent germ cell depletion. The cause of the latter phenotype is yet to be determined but is unlikely to be associated with abnormal macrophage infiltration to the mutant testes.

Discussion

In the present study, we assessed the function of the E3 ubiquitin ligase CUL4B in spermatogenesis using genetically modified mouse models. We discovered a male infertility phenotype in both germ cell-specific conditional Cul4b knock-outs, as well as global Cul4b-null mutants.

Abolishing Cul4b in male germ cells did not cause any apparent morphological changes to the testis, and the progression of spermatogenesis appeared to be normal. The seminiferous tubules of Cul4bVasa conditional knock-outs contained male germ cell populations of all phases, and a normal number of spermatozoa were recovered from these animals. The cause of male infertility in the mutants is low sperm motility (93% reduction compared with controls). Several factors could contribute to this decrease in motility. We detected a varied level of decreased mitochondrial activity and loss of mitochondrial membrane potential, evidenced by MitoTracker Red CMX Ros and JC1 staining (Fig. 3). The mitochondrial sheath, tightly wrapped around the axoneme of the sperm tail midpiece, may provide energy in the form of ATP required for flagella movement. In humans, numerous cases of idiopathic and varicocele-related asthenozoospermia have been reported to be associated with defective mitochondrial respiratory functionalities, with or without phenotypic abnormalities of the mitochondrial sheath (35–37). Furthermore, a positive correlation was found between high mitochondrial membrane potential and increased sperm motility in a human male infertility study (37). Oxidative phosphorylation relies on the H+ concentration gradient generated and maintained by the electron transport chain, which requires the impermeability of inner mitochondrial membrane. A decrease in mitochondrial membrane potential therefore indicates an inner membrane breach that could potentially lead to impaired ATP production. Indeed, measurement of sperm ATP level revealed an almost complete loss of ATP in the Cul4bVasa spermatozoa. However, the partial reduction in mitochondrial activity cannot fully account for the near total loss of ATP production in mutant spermatozoa. Therefore, the alternative pathway for ATP production, glycolysis, could also be impaired. We measured spermatozoa intracellular levels of PEP, the second to last intermediate of glycolysis, and revealed a significant decrease compared with the controls (Fig. 3). Therefore, loss of CUL4B simultaneously caused reduction of mitochondrial activity in the midpiece and...
compromised glycolysis, which resulted in lack of energy supply to mutant spermatozoa, ultimately leading to the immotile phenotype. In addition, we observed structural defects in the mutant spermatozoa. Specifically, the axonemes of many mutant spermatozoa showed microtubule anomalies, including supernumerary, missing, or misplaced outer microtubule doublets (Fig. 3). During murine spermiogenesis, the distal centriole nucleates microtubule polymerization to form the axoneme and the proximal centriole organizes the anlage of the caput. Abnormal microtubule arrangements found in Cul4bVasa mutant sperm flagella suggest potential structural defects in the distal centriole or failure, at least in part, of microtubule nucleation.

To uncover the molecular mechanisms leading to these defects, we performed proteomic analysis comparing ubiquitinated proteins isolated from control and Cul4bVasa spermatozoa by affinity purification using p62-UBA domain-immobilized beads. Among the proteins identified as potential CUL4B targets, INSL6 showed relatively high spectrum count in control samples but was completely absent in Cul4bVasa samples. INSL6 is an insulin-like peptide that shows significant sequence similarity to the relaxin B chain (38). Intriguingly, high INSL6 expression has been reported in both mouse and human male germ cells, specifically in spermatids (24) (HPA021364 SIGMA), and INSL6 mutation in mouse led to compromised fertility in males only (39). Rearrangement of INSL6-containing chromosome 9q24.1 region in humans has been reported to be associated with testicular development failure (40), even though its function in spermatogenesis remains ambiguous.

In this study, we performed in vitro experiments and provided solid evidence that CUL4B mediates both INSL6 polyubiquitination, as well as its degradation, implicating INSL6 as a direct substrate of the CUL4B E3 ubiquitin ligase (Fig. 4). Direct interaction between INSL6 and DDB1 was also demonstrated. In addition, knocking down either DDB1 or RBX1 resulted in a reduction in INSL6 polyubiquitination (Fig. 4). It is unfortunate that there is currently no commercially available antibody against mouse INSL6, which prevented us from analyzing INSL6 expression and accumulation in mouse germ cells. Further investigations aiming at revealing the function of INSL6 is underway, which will shed light on understanding the underlying mechanisms of its involvement in spermatogenesis.

Interestingly, Cul4b-null mutation by Sox2-Cre resulted in a different male fertility phenotype, in that the aged mutant mice exhibited a germ cell depletion phenotype. This led us to hypothesize that CUL4B is required to maintain the SSC niche. It is believed that the SSCs are not randomly distributed along the basement membrane of seminiferous tubules but rather preferably located to "SSC niches" that are adjacent to the interstitial tissues rich in blood vessels (41, 42). Several types of somatic cells are responsible for maintaining the SSC niche. Inside the seminiferous tubules, Sertoli cells are thought to provide growth factors regulating SSC self-renewal/proliferation, including FGF2, GDNF (glial cell line-derived neurotrophic factor), and SCF (stem cell factor) (43–45). Because Cul4b was found to be highly expressed in Sertoli cells and loss of Cul4b was not compensated by Cul4a, we first speculated that it played a cell autonomous role in the Sertoli cells to maintain the SSC niche. However, we found no germ cell depletion phenotype in Sertoli-specific Cul4b mutants (Cul4bAlox; Fig. 2). This result suggests that Cul4b is dispensable in Sertoli cells. The interstitial Leydig cells are thought to regulate Sertoli cell function through growth factor production via testosterone signaling. There is a possibility that loss of CUL4B in Leydig cells altered this signaling, causing disturbance in Sertoli cell functions. However, measurements of circulating testosterone in Cul4bSox2 males revealed no significant difference compared with controls (data not shown). It was also proposed that interstitial macrophages play essential roles in maintaining SSC niche, and infiltration of macrophages in the testis was reported in patients with male infertility (30). High expression of Cul4b was reported in macrophages, and loss of CUL4B protein by shRNA knockdown significantly reduced TNFα protein in the activated macrophages (46). This led us to test whether Cul4b loss caused changes in testicular macrophage populations, which may in turn cause SSC loss. We tested this hypothesis by analyzing immunocytochemistry of the spleen and testis tissues of control and Cul4bSox2 animals. We used F4/80 as a macrophage marker for FACS analysis, and no noticeable differences were discovered between the two genotypes or mutants of different ages (Fig. 5). Thus, loss of SSCs in Cul4bSox2 mutant testis is not the result of a change in testicular macrophage number. However, our study does not exclude the possibility that loss of Cul4b may change the properties of testicular macrophages, thereby affecting their ability to maintain the SSC niche. One final cell population that could contribute to the germ cell loss phenotype is the peritubular myoid cell. However, we did not detect either CUL4B or CUL4A expression in this cell type; hence it is unlikely that CUL4B is required in these cells to maintain the stem cell niches. It is noteworthy that our global mutants manifest a male germ cell depletion phenotype with a rather late adult onset, in contrast to most other germ cell depletion mouse models exhibiting early onset. This makes our animals an excellent model for studying long term maintenance of the SSC niche. In conclusion, CUL4B is required in the male germ cells for spermiogenesis and in somatic cells to maintain the SSC population of the testis.

Author Contributions—L. M. and P. Z. designed the study, and Y. Y., L. L., C. Y., C. L., G. M. V., C. W., and P. S. contributed to data acquisition and analysis. Y. Y., L. M., and P. Z. wrote the paper.

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CUL4B Regulates Mammalian Spermatogenesis

References


