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Coprs inactivation leads to a derepression of LINE1 transposons in spermatocytes

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Repression of retrotransposons is essential for genome integrity during germ cell development and is tightly controlled through epigenetic mechanisms. In primordial germ cells, protein arginine N-methyltransferase (Prmt5) is involved in retrotransposon repression by methylating Piwi proteins, which is part of the piRNA pathway. Here, we show that in mice, genetic inactivation of coprs (which is highly expressed in testis and encodes a histone-binding protein required for the targeting of Prmt5 activity) affects the maturation of spermatogonia to spermatids. Mass spectrometry analysis revealed the presence of Miwi in testis protein lysates immunoprecipitated with an anti-Coprs antibody. The observed deregulation of Miwi and pachytene pre-piRNAs levels and the derepression of LINE1 repetitive sequences observed in coprs-/- mice suggest that Coprs is implicated in genome surveillance mechanisms.

Spermatogenesis is a multistep process that takes place in the seminiferous tubules whereby mature spermatozoa are continuously produced during adulthood reproduction lifetime. Specifically, primordial germ cells (PGCs) generate spermatogonia that either self-renew to maintain a pool of stem cells or undergo differentiation. After two meiotic divisions, secondary spermatocytes are produced. Then, during spermiogenesis, they differentiate into elongating spermatids and finally spermatozoa [1–5]. Importantly, these changes are associated with chromatin compaction, transcription arrest, and specific epigenetic modifications that contribute to the production of mature sperm. Concomitantly, to maintain genome integrity, PGCs need to repress repetitive DNA elements because their spread can lead to inherited diseases. These repetitive sequences (for instance, long interspersed nuclear elements 1 (LINEs), short interspersed nuclear element (SINEs), and intracisternal A-particle (IAP) sequences) derive mostly from transposable elements and represent

Abbreviations
Coprs, coordinator of Prmt5 and differentiation stimulator; IAP, intracisternal A-particle; LINE1, long interspersed nuclear element 1; piRNA, Piwi-interacting RNA; Prmt5, protein arginine N-methyltransferase; SINE, short interspersed nuclear element.
about 40% of the human genome [3,6–8]. Thus, their repression is challenging and is tightly controlled in germ cells by the Piwi-interacting RNA (piRNA) pathway and through epigenetic mechanisms [9,10]. In the mouse, this pathway involves the three Piwi proteins Miwi, Mili, and Miwi2 (PIWIL1, PIWIL2, and PIWIL4 in humans, respectively) that show different developmental expression patterns in testis [11,12]. Miwi is expressed in fetal germ cells up to the round spermatid stage, while Miwi2 expression is restricted to fetal and perinatal germ cells. In adult testes, Miwi is expressed in pachytene spermatocytes up to round spermatid testes [13]. Mili and Miwi bind to both piRNAs in pachytenes spermatocytes and postmeiotic spermatids [14]. Deletion of the Miwi gene arrests cells at the round spermatid stage, and Mili deficiency leads to male sterility [14]. Chromatin-modifying enzymes also participate in the regulation of these repetitive elements. For instance, protein arginine N-methyltransferase 5 (Prmt5) is implicated in the piRNA pathway in PGCs via a Prmt5-dependent methylation of Piwi proteins. In somatic cells, Prmt5 modulates cell proliferation and differentiation, for instance, during myogenesis and adipogenesis, by interacting with various proteins, including Coprs (previously termed Copr5), a histone-binding protein [15–24]. In addition, Prmt5-methylated arginine residues can be bound by proteins harboring a Tudor domain. In PGCs, many of these proteins (e.g., Piwi proteins) are essential for male fertility [25]. It was suggested that Prmt5 maintains DNA integrity during global epigenetic reprogramming and that its subcellular localization affects the piRNA pathway, at least in part, and promotes transposon silencing [26]. In addition, conditional loss of Prmt5 in early mouse PGCs causes complete male and female sterility that was attributed to global impairment of DNA demethylation in the genome [26].

Here, we show that in coprs-/- mice, maturation of spermatogonia to spermatids is affected, although animals are fertile. Moreover, the mRNA levels of CenAl (cyclin A1), Prml (sperm protamine P1), Miwi, pachyteny piRNA precursors, and LINE1 are deregulated in coprs-/- testes, suggesting that Coprs contributes to genome surveillance mechanisms via the Piwi-piRNA pathway.

**Materials and methods**

**Flow cytometry**

Extraction of cells in testis was performed essentially as described in Mays-Hoopes et al. [27]. Testes from wild-type (WT n = 3) and knockout (KO n = 4) mice were dissected and decapsulated to release the tubules. Seminiferous tubules were incubated with 0.25 mg·mL⁻¹ collagenase type IV (Sigma) at 37 °C under rapid agitation for 5 min and washed to release Leydig cells and interstitial cells. Dispersed tubules were allowed to settle and washed twice to remove peritubular cells. Washed tubules were then incubated with 0.5% trypsin/EDTA (Gibco) and 1 µg·mL⁻¹ DNase RQ1 (Promega) at 37 °C for 5 min. Trypsin digestion was stopped by adding DMEM with 10% FBS. Suspensions were washed and disaggregated into single-cell suspensions by trituration before filtration through a 50-µm cell strainer. For analysis, cells were fixed in 0.4 M citrate buffer (pH 4.5) overnight and resuspended in 1 mL of cold propidium iodide (PI) staining solution (10 mM Tris/HCl (pH 8.0), 1 mM NaCl, 0.1% Nonidet P-40, 50 µg·mL⁻¹ PI, 10 µg·mL⁻¹ RNase A), vortexed, and incubated on ice for 10 min to lyse the plasma membrane and stain nuclear DNA. DNA content was assessed on a FACSCalibur II (Becton Dickinson) equipped with the CELQuest software.

**Western blotting**

Anti-Prmt5 (Millipore) and anti-Miwi (Abcam) antibodies were used according to the manufacturer’s instructions. The anti-Coprs antibody (AGRO-BIO, Clermont-Ferrand, France) was against the last 20 amino acids of Coprs C terminus [21].

**Mass spectrometry analysis**

In-gel digestion of bands excised from Colloidal Blue-stained gel was done before LC/MS/MS analysis that was performed at the Taplin Mass Spectrometry Facility (Harvard Medical School, Boston).

**Immunohistochemistry (IHC)**

Tissues were fixed in Bouin’s fixative and embedded in paraffin. Then, 4-µm-thick sections were cut and processed for IHC staining. IHC was performed using the same anti-Coprs and anti-Miwi (Abcam) antibodies employed for western blotting, followed by a biotinylated secondary antibody coupled to the streptavidin–peroxidase complex (ABC Vectastain Kit; Vector Laboratories). Revelation was performed with the peroxidase substrate DAB (3,3′-diaminobenzidine) from Vector Laboratories.

**RNA isolation, cDNA synthesis, and RT-qPCR amplification from mice**

Total semen RNA was isolated with the TRIzol Reagent (Life Technologies) according to the manufacturer’s instructions. Briefly, 800 µL of TRIzol and 200 µL of chloroform were added to 200 µL of sperm. The mixture was mixed for...
15 s and left at room temperature for 5 min. After centrifugation at 12 000 g at 4 °C for 15 min, supernatants were transferred to fresh tubes containing 1 volume equivalent of 70% ethanol. Then, total RNA was purified using the miR-Neasy Serum/Plasma Kit (Qiagen) according to the manufacturer’s recommendations. RNA was quantified with a NanoDrop ND-1000 spectrophotometer (NanoDrop Technologies Inc., DE, USA). RNA isolation from mouse testes and RT-qPCR were performed as described [18]. Briefly, testes were lysed in TRIzol reagent (Invitrogen), and total RNA was isolated according to the manufacturer’s recommendations. cDNA was synthesized from 1 l of total RNA using random hexamers and SuperScript III Reverse Transcriptase (Invitrogen). Real-time qPCR was performed on a LightCycler 480 SW 1.5 apparatus (Roche) with Platinum Taq DNA Polymerase (Invitrogen) in the presence and the SYBR Green Mix.

Oligonucleotide sequences

Oct4 f CAATGAGAACCTTCAGGAGATATGC, r TCA ATGCTAGTTCCGTTTCTTC;
Amh f TAGTCTACATCTGGCTGAAGTGATATG, r CCAGGTGAGGCTCTTGGA;
Cca1 f CACTTCGTGGATTTCAAC, r CGATGAGATCTCCTCTGCATAC;
Prm1 f GCCGCTCATACACCATAAGG, r CAAGATGTGGCGAGATGC;
Prm2 f GCAGAAGATCCCGAAGGAG, r CTCCAGCGAATGGACAG;
H1t2 f AGGGAAGAGAAGGGACAGGA, r CTTGGAGGCCCATATGGAAAA;
Hils f GTCCCAAGCCAGAGTGAGAG, r CTTGAGCGCCAGGTGTTAT;
Miwi f ATGATCGTGGGCAT, r AGGCCACTGCTGTCTACATA;
pre-piR1 f GTTAGCGAAGGACATTATTCTAACC, r TGACATGAACACAGGTGCTCAGAT;
pre-piR2 f CTATGCTTATGATGGCATTGGAGAG, r TTCCAGTTCAACAGGGACACGGGAC;
pre-piR3 f GTTCTCACTTTATCAGCTCTCAAG, r TGAGAGTGGCATCTAAATGTTTAG;
Line1 f GGAGGGACATTTCATTCTCATCA, r GCTGCTCTTGTATTGGGACATAGA;
SineB1 f GTGGCGCACGCCTTTAATC, r GACAGGTGTTCTCTGTGTAG;
IAP f ATTGTTCCCTCACTGGCACA, r ATTGTTCCCTCAGTGGCACA;
Actin f AGAAGAGCTATGAGCTGCCT, r TCATCGTACTCCTGCTGTCT;
hCOPRS f TCCGCGTCCACTGTGATACC, r GGCCCGCCACCTCACA;
hHPRT f TGACACTGGCAAAACATGCA, r GTGCTTTTCACCCAGCAAGTCT.

Mice and animal care

Animal experiments were approved by the Ethics Committee of the Languedoc-Roussillon Region (France).

Patient’s characterization

Nine patients were recruited from our IVF/ICSI program at the ART-PGD Department of Montpellier University Hospital, France, after signature of the written informed consent. Each patient was classified according to the fraction (%) of typical spermatozoa forms (TF) evaluated using one of three classical methods [28–30]. The study methodologies were approved by the local ethics committee and conformed to the standards set by the Declaration of Helsinki.

Results

Coprs is localized in spermatogonia and its genetic inactivation leads to male germ cell accumulation at the pachytene stage

Our previous studies [21] and the available Gene Expression Omnibus (GEO) data on Coprs expression profile in various human tissues confirmed that it was strongly expressed in testis (Fig. S1). To investigate Coprs role in testis, we used coprs KO mouse [22]. Morphological analysis of 8- and 36-week-old coprs KO males did not show testicular atrophy or reduced body weight (data not shown). IHC analysis of paraffin-embedded testis sections prepared from coprs KO animals and control littermates showed that Coprs was expressed only in germ cells and was mostly localized in the nucleus, although the anti-Coprs antibody stained also faintly the cytoplasmic compartment (Fig. 1A; note the absence of expression in coprs KO testis). A similar nuclear localization was previously observed in different culture cell types [21]. As we and others previously reported that Coprs is implicated in cell differentiation [16–18,22], we analyzed germ cell maturation and the distribution of the different cell types in WT and KO testes by flow cytometry. The proportion of germ cells that progressed through meiosis in the leptotene and zygotene stages was reduced in KO mice, and cells accumulated at the pachytene stage (Fig. 1B). Moreover, the proportion of round and elongated spermatids was lower in coprs KO than WT testes (Fig. 1B). These data suggest that Coprs absence perturbs spermatogenesis between the end of meiosis and spermiogenesis.

To further document at what stage coprs inactivation impaired the maturation of spermatogonia, we measured by RT-qPCR the expression level of various differentiation markers in testes of 8-week-old mice.
The mRNA levels of Oct4 and Amh, two markers of undifferentiated spermatogonia and Sertoli cells, respectively, were not significantly different between WT and KO samples (Fig. 1C). This suggested that the initial pool of undifferentiated spermatogonia was comparable in WT and KO animals. In agreement with the flow cytometry data, the mRNA levels of the premeiotic marker Ccnal were downregulated, whereas the level of the histone variant H112, which is usually expressed in pachytene spermatocytes and persists in early spermatids, was upregulated in coprs KO testes (Fig. 1C). Similarly, Prm1 mRNA level was significantly increased in coprs KO testis cells (Fig. 1C). In contrast, spermatid-specific linker histone Hils1 and Prm2 levels did not vary significantly between WT and KO in elongated spermatids (Fig. 1C). Consequently, the Prm1/Prm2 ratio, an indicator of chromatin compaction [31], was significantly higher in coprs KO testes than in controls (1.34 in coprs KO mice vs 0.92 in WT animals), suggesting that Coprs deficiency results in perturbations of chromatin compaction.

These data indicated that coprs is implicated in spermatocyte maturation.
Coprs gene ablation leads to deregulation of the piRNA pathway and derepression of LINE1 repetitive elements

To explore how Coprs regulates spermatocyte maturation, we next characterized Coprs protein interactors in testis by proteomics. After immunoprecipitation of Coprs in cell lysates prepared from testes of WT and coprs KO mice, and immunoprecipitate separation by SDS/PAGE and Colloidal Blue staining, we excised bands that were present in WT but not in KO extracts for mass spectrometry analysis (Fig. 2). Although Prmt5 and histone H4 are well-characterized direct partners of Coprs [21], we did not identify out of the background any specific band in the range corresponding to Prmt5 molecular weight. Conversely and as expected, histones were differentially enriched in Coprs immunoprecipitates, validating the efficiency of the immunopurification procedure. Besides histones, Miwi was the most differentially enriched protein in Coprs immunoprecipitates, suggesting that Coprs is part of a Piwi complex (Fig. 2). Of note, this analysis identified also Stk31 (Tdrd8), a Tudor domain containing protein that interacts with Miwi and Mili upon Prmt5-dependent methylation, but not critical for male germ cell development. IHC analyses of testis tissue sections showed that Miwi staining was lower in samples from coprs KO mice than WT littermates (Fig. 3A). Similarly, Miwi protein (western blotting) and mRNA (RT-qPCR) levels were significantly reduced in testes from coprs KO mice compared with WT littermates (Fig. 3B,C).

As Miwi and piRNAs associate in a functional complex that controls genome integrity, we evaluated whether piRNAs expression was altered upon coprs gene ablation. RT-qPCR analysis showed that pachytene piRNA precursors (pre-piRNAs) 1, 2, and 3 were less abundant in coprs KO than WT testes (Fig. 4A). Although 75% of piRNAs expressed at the pachytene stage are not yet annotated, approximately 17% of them play a key role in inhibiting retrotransposon expression [32]. Therefore, we used the RNA expression levels of the most abundant retrotransposons, namely LINE1, SINE B1, and IAP, as readout of potential dysfunction in the piRNA pathway. The transcript level of LINE1, but not of SINE B1 and IAP, was upregulated in coprs KO compared to WT testis samples (Fig. 4B).

Altogether, these results suggest that Coprs is involved in the piRNA pathway and that its deficiency in male germ cells perturbs LINE1-related genome surveillance mechanisms.

Coprs and teratozoospermia

To evaluate whether the molecular alterations identified in coprs KO testes influenced reproduction, we first measured the mating efficiency of coprs KO males and control littermates. Several independent crosses of coprs heterozygous animals identified a non-Mendelian inheritance of coprs-floxed allele in males (Fig. 5A). Matings of coprs KO males with WT females were successful without noticeable reduction of litter size (data not shown), indicating that spermatocyte maturation impairment in these mice was not severe enough to impair fertility. Considering that several knockout mice with reduced sperm count or altered spermatogenesis, but normal fertility, have been described previously [33], we evaluated the number, motility, and morphology of spermatozoa in 11-week-old mice. We did not find any significant difference between WT and coprs KO mice (Fig. 5B,C). However, probing GEO data for biological situation of differential Coprs expression, low level of Coprs mRNAs correlated strongly with teratozoospermia, a human pathology linked to male infertility (Fig. 6A). Therefore, we conducted a pilot experiment to evaluate Coprs mRNA level by RT-PCR in patients with teratozoospermia associated with fertility defects and confirmed that it was downregulated compared with patients without teratozoospermia (Fig. 6B,C).

Discussion

Here, we show in mouse that Coprs is expressed in spermatogonia and that its absence, maturation of spermatogonia during meiosis and spermiogenesis is slightly impaired. This leads to a significant accumulation of spermatocytes at the pachytene stage and a decrease of round/elongated spermatids. Many changes occur within the cell nucleus during spermatocyte maturation, including the programmed replacement of histones by a set of basic nuclear proteins such as histone variants and protamines to increase chromatin compaction, an essential event for male fertility. Consistent with an accumulation of cells in pachytene, the mRNA level of H1t2, a histone variant that is specifically expressed in pachytene spermatocytes, was increased in coprs KO cells. Moreover, the Prm1/Prm2 ratio was increased in KO spermatocytes, a parameter that was reported as being associated with male infertility in humans [34–36]. Our proteomic analysis identified Miwi as a potential Coprs interactor, although at this stage our data do not address whether this interaction is direct or through the presence of these two proteins within a common complex. Moreover, both Miwi
Fig. 2. Identification of Miwi as a major protein immunoprecipitated with Coprs. Protein extracts from WT and coprs KO testes were immunoprecipitated with an anti-Coprs antibody. Bands (*) present only in the WT lane were excised from the Colloidal Blue-stained gel (left) and analyzed by mass spectrometry (right).

| Unique | Total | AVG | Gene Symbols | Avg Precursor Intensity |
|--------|-------|-----|--------------|-------------------------|
| 48     | 49    | 2.87| Pwitm1       | 5.57e+06                |
| 30     | 36    | 3.23| Trin28       | 4.35e+06                |
| 32     | 32    | 3.01| Psmd2        | 3.18e+06                |
| 20     | 20    | 2.95| Pwitm2       | 1.77e+06                |
| 29     | 29    | 2.74| Nup93        | 1.72e+06                |
| 22     | 23    | 3.74| Uba1        | 1.71e+06                |
| 21     | 21    | 3.3 | Usp5        | 1.48e+06                |
| 21     | 25    | 3.64| Psmd1       | 1.13e+06                |
| 20     | 21    | 3.65| Xpo1       | 1.10e+06                |
| 20     | 21    | 3.85| Stk31      | 1.05e+06                |

Fig. 3. coprs KO affects Miwi expression level. (A) IHC analysis of WT and KO mouse testis sections with or without (negative control, -) an anti-Miwi antibody; bar = 50 μm. (B) Left panel: western blot detection of Miwi expression in whole-cell extracts from WT and KO testes with anti-Miwi and anti-Gapdh antibodies; right panel: quantification of Miwi expression with ImageJ after normalization to Gapdh expression. Values are expressed in arbitrary units (a.u.) and are the mean ± SEM of two independent experiments. (C) Miwi expression was assessed by qPCR. Data were normalized to actin level and expressed in arbitrary units (a.u.) (mean ± SEM of three independent mice/group). **P < 0.05 (Student’s t-test).
protein and mRNA levels were significantly decreased in \textit{coprs} KO mice, as well as the expression level of pachytene piRNAs. Altogether, these results suggest that the Miwi-piRNA pathway is altered in \textit{coprs} KO testis. How Coprs controls Miwi and piRNA levels remains unknown, and future studies will address this point. It can be hypothesized that the stability of Miwi protein or Miwi complexes requires Coprs. In addition, the reduced Miwi level in \textit{coprs} KO testes, resulting \textit{de facto} in a lower Miwi slicer activity, might impact on piRNAs that target active retrotransposons, such as LINE1, thus weakening the maintenance of transposon silencing [12].

In contrast to \textit{Miwi} KO mice in which spermatogenesis is arrested at the round spermatid stage with no retrotransposon upregulation [14], \textit{coprs} KO testes present only a slower spermatogenesis progression but showed a significant derepression of \textit{LINE1} expression. This suggests that the low but detectable level of Miwi present in \textit{coprs} KO animals is sufficient to ensure spermatogenesis and that Coprs could contribute to the control of \textit{LINE1} expression and genome stability beyond its impact on Miwi expression. A similar mild phenotype, that is, normal fertility and viability associated with a derepression of \textit{LINE1} transcripts), was described in mice deficient for \textit{Exd1} which encodes for a partner of Miwi2 piRNA biogenesis factor \textit{TDRD12} [37]. Importantly, the functional consequences of \textit{Exd1} loss in testis, that is, massive derepression of \textit{LINE1} elements and an arrest in spermatogenesis, are revealed only in the \textit{Exd1} \textsuperscript{-/-};\textit{Tdrd12} \textsuperscript{+/-} genetic background [38]. As well, we can make the assumption that a stronger impact of Coprs depletion on spermatogenesis might be observed by breeding \textit{coprs} mice with animals in which other Piwi-associated protein-encoding genes have been invalidated. However, we cannot exclude on the basis of the non-Mendelian inheritance of the \textit{coprs}-floxed allele that the more severe \textit{coprs} KO phenotypes might affect early embryo viability. This
could explain how coprs absence led to an apparent mild phenotype compared with that of Miwi and Miwi-associated proteins. Thus, it is possible that KO mice upon F2 and/or the following generations might affect more strongly the reproduction parameters and give rise to a phenotype reminiscent of human teratozoospermia, a pathology associated with low Coprs expression level. Although this conclusion requires confirmation in a larger cohort of patients, these data support the hypothesis that Coprs is a candidate diagnostic marker for teratozoospermia.

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**Conflict of interest**

The authors declare no conflict of interest.

**Authors contributions**

PC executed and analyzed FACS experiments; HD carried out immunohistochemistry experiments; DH and SH provided human samples and critical discussions of the manuscript; SC provided critical discussions of the manuscript; and FE carried out PCR data analysis, designed the study, and drafted the manuscript.

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Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Fig. S1. (A) Western blot detection of whole cell extracts from WT (*n* = 4) and *coprs* KO (*n* = 3) testes with antibodies against the indicated proteins. The expression of Prmt5, a *Coprs* partner, was comparable in KO and WT samples. (B) *Coprs* RNA expression level from RNA-seq data in different human tissues (GEO data) is expressed as TPM (transcripts per million), i.e., the mean values of the different samples from each tissue. Sample color-coding is based on tissue groups with common functional features.