Bat3 deficiency accelerates the degradation of Hsp70-2/HspA2 during spermatogenesis

Toru Sasaki,1 Edyta Marcon,4 Tracy McQuire,1 Yoichi Arai,5 Peter B. Moens,4 and Hitoshi Okada1,2,3

1The Campbell Family Institute for Breast Cancer Research, 2Division of Signaling Biology, Ontario Cancer Institute, University Health Network, and 3Department of Medical Biophysics, University of Toronto, Toronto, Ontario M5G 2C1, Canada
4Department of Biology, York University, Toronto, Ontario M3J 1P3, Canada
5Department of Urology, Tohoku University, School of Medicine, Aoba-ku, Sendai 980-8574, Japan

Meiosis is critical for sexual reproduction. During meiosis, the dynamics and integrity of homologous chromosomes are tightly regulated. The genetic and molecular mechanisms governing these processes in vivo, however, remain largely unknown. In this study, we demonstrate that Bat3/Scythe is essential for survival and maintenance of male germ cells (GCs). Targeted inactivation of Bat3/Scythe in mice results in widespread apoptosis of meiotic male GCs and complete male infertility. Pachytene spermatocytes exhibit abnormal assembly and disassembly of synaptonemal complexes as demonstrated by abnormal SYCP3 staining and sustained γ-H2AX and Rad51/replication protein A foci. Further investigation revealed that a testis-specific protein, Hsp70-2/HspA2, is absent in Bat3-deficient male GCs at any stage of spermatogenesis; however, Hsp70-2 transcripts are expressed at normal levels. We found that Bat3 deficiency induces polyubiquitylation and subsequent degradation of Hsp70-2. Inhibition of proteasomal degradation restores Hsp70-2 protein levels. Our findings identify Bat3 as a critical regulator of Hsp70-2 in spermatogenesis, thereby providing a possible molecular target in idiopathic male infertility.

Introduction

Meiosis is a fundamental process for genetic exchange between maternal and paternal genomes in all eukaryotes. During prophase of the first meiotic division, homologous chromosomes undergo synopsis, genetic exchange, and gene conversion. Once paired, homologous chromosomes are connected by the synaptonemal complex (SC), a tripartite multiprotein structure. The SC consists of the central element, axial/lateral elements, and transverse filaments (Fawcett, 1956; Moses, 1956, 1969; Zickler and Kleckner, 1999). Formation of the fully synapsed autosomal SCs and the partially synapsed XY pair are essential for successful completion of DNA repair and recombination processes and subsequent desynapsis (Moens, 1994). Although the function and regulation of SC proteins are not fully understood, recent genome-wide screens and genetic studies have identified novel SC components (Wang et al., 2001; Maratou et al., 2004; Toure et al., 2005), including SYCE1, CESC1, and TEX12 (Costa et al., 2005; Hamer et al., 2006). These discoveries, combined with mouse genetics, have provided in-depth insight into the regulation of meiosis (Bolcun-Filas et al., 2007; Costa and Cooke, 2007).

Hsp70-2, another SC-interacting protein, is expressed exclusively in male germ cells (GCs) at specific stages of differentiation. Hsp70-2 is not expressed in spermatogonia but becomes detectable at leptotene and zygotene. Hsp70-2 is expressed highly in pachytene spermatocytes, where it has been found to associate with the lateral element of the SC (Allen et al., 1996). Consistent with this expression pattern, sperm development in Hsp70-2-deficient mice arrests in meiotic prophase I, and the majority of late pachytene spermatocytes are eliminated by apoptosis, resulting in complete male infertility (Dix et al., 1996). Recently, Hsp70-2 was reported to act as a chaperone of transition proteins TP1 and TP2 in spermatids (Govin et al., 2006).

Bat3 (also called Scythe; Thress et al., 1998) has been reported to regulate apoptosis in a variety of settings (Thress et al., 1998, 1999a, b; Desmots et al., 2005). Certain domains in Bat3...
also appear in the Bcl-2-associated athanogene (BAG) family of proteins that regulate Hsp70 function. Thus, Bat3 may be important for the folding and activity of apoptotic signaling molecules (Thress et al., 2001). Recently, p53 and apoptosis-inducing factor have been identified as Bat3 targets after DNA damage (Sasaki et al., 2007) and ER stress (Desmots et al., 2007), respectively. In addition, Bat3 has been shown to act as a ligand of a natural killer cell receptor (Pogge von Strandmann et al., 2007), suggesting that Bat3 may mediate signals from damaged cells to the immune system. However, although Bat3 is predominantly expressed in adult testes (Wang and Liew, 1994), the role of Bat3 in male GC development remains unknown.

In this study, we investigated male GC development in Bat3-deficient mice and found that most Bat3-deficient male GCs die of apoptosis at meiotic prophase I. Interestingly, the testis-specific Hsp70-2 protein was undetectable in Bat3-deficient cells even though Hsp70-2 transcript levels were normal. We found that Bat3 inactivation induced polyubiquitylation (poly-Ub) and subsequent degradation of Hsp70-2. Additional inactivation of proteasome activity restored Hsp70-2 protein levels. We conclude that Bat3 functions as a critical regulator of Hsp70-2 in spermatogenesis.

**Results and discussion**

We detected high levels of Bat3 mRNA in adult testes (Fig. 1 A), which is consistent with a previous study (Wang and Liew, 1994). Male but not female Bat3+/− mice were completely infertile. We observed that Bat3+/− testes at postnatal day 120 (P120) were significantly smaller (Fig. 1 B). The mean weight of Bat3+/− testes (40.0 ± 7.0 μg; n = 8) was one third of the weight of Bat3+/+ (125.0 ± 1.0 μg; n = 8) and Bat3−/− (115.0 ± 7.0 μg; n = 8) testes (Fig. 1 C). In contrast, no significant differences were observed in the size and weight of the epididymis for all three genotypes (Bat3+/+, 25.0 ± 2.0 μg; Bat3+/−, 24.0 ± 7.0 μg; Bat3−/−, 23.0 ± 1.0 μg; n = 8; Fig. 1 D). In addition, serum levels of follicle-stimulating hormone (FSH), luteinizing hormone (LH), and testosterone were not significantly different between Bat3+/+ and Bat3−/− mice (Table S1, available at http://www.jcb.org/cgi/content/full/jcb.200802113/DC1) relative to normal controls (Fig. 2 A and Fig. S1 A). Consistent with these morphological abnormalities, multiple γ-H2AX-positive foci were observed on paired and unpaired Bat3−/− chromosomes (Fig. 2 F and Fig. S1, E and F). The paired regions heavily coated with γ-H2AX (Fig. S1 F) suggest the existence of nonhomologously paired regions in Bat3−/− chromosomes. In contrast, only the partially paired XY was positive for γ-H2AX staining in Bat3+/− samples (Fig. 2 B and Fig. S1 D). We also detected abnormal SYCP3 and γ-H2AX staining in Bat3−/− cryosections (Fig. 2, O–T). These data indicate that multiple synaptic abnormalities occur in Bat3−/− pachytene spermatocytes.

Next, we analyzed the localization of the early recombination repair marker Rad51. Rad51 was located along synapsed and asynapsed portions of SCs in Bat3+/+ zygote spermatocytes and gradually disappeared in late zygote/early pachytene (Fig. 2 C). In contrast, Bat3−/− pachytene spermatocytes retained a significant number of Rad51 foci (Fig. 2 G) even though the number and localization of Rad51 recombine complexes appeared normal in meiotic nuclei during early prophase (not depicted). We also stained for replication protein A (RPA), which colocalizes with Rad51 (Sigurdsson et al., 2001; Govin et al., 2006). We detected a significant number of RPA foci in Bat3−/− (Fig. 2 H) but only trace levels in Bat3+/+ (Fig. 2 D) pachytene spermatocytes. These findings suggest that Bat3 is required for proper chromosomal pairing, recombination, and desynapsis.

Although most spermatocytes in Bat3−/− mice arrested and died in meiotic prophase I, we observed a small number of spermatids and sperm cells in chromosome spread preparations (Fig. 2 J). These elongated spermatids and sperm heads showed abnormal shape and mislocalization of TP-1 (Fig. 2, K–N). TP-1 functions in histone replacement within the sperm head (Sassone-Corsi, 2002). Thus, Bat3 may regulate histone replacement in postmeiotic GCs.

The phenotype observed in Bat3−/− males is characterized by the failure of SCs to disassemble, resulting in differentiation observed between Bat3+/+ and Bat3−/− testes, suggesting that Bat3 is not essential for the production of spermatogonia. Similarly, phosphoglycerate kinase-2 (a spermatocyte-specific marker; Boer et al., 1987) was expressed normally in the Bat3−/− background. In contrast, protamine 1, the mature haploid GC-specific marker expressed in postmeiotic GCs (Kleene et al., 1984), was undetectable in Bat3−/− testes, indicating a severe defect in GC differentiation beyond meiosis. We also found that GATA-1, a marker of Sertoli cells, was expressed normally in Bat3−/− testes.

We next investigated whether Bat3 deficiency induces apoptosis in male GCs at various developmental stages. Apoptotic cells were rarely detected at P7 (Fig. 1, P and Q). In contrast, the number of apoptotic cells increased significantly in Bat3−/− testes at P42 (Fig. 1, R–V). These data further support that spermatogenesis is impaired in the absence of Bat3.
arrest and apoptosis of spermatocytes, essentially leading to male infertility. Interestingly, this phenotype is similar to that seen in Hsp70-2−/− male mice (Dix et al., 1996). Hsp70-2 has also been found to regulate TP-1 and -2 (Govin et al., 2006). In addition, Bat3 has been shown to bind Hsp70/Hsc70 in a BAG domain–dependent manner and inhibits Hsp70-mediated protein refolding in vitro (Thress et al., 2001). Such mounting evidence prompted us to investigate a functional interaction between Hsp70-2 and Bat3.

Hsp70-2 was abundantly expressed in spermatocytes and round spermatids but not spermatogonia in Bat3−/− mice (Fig. 3, A and C). In contrast, Hsp70-2 was undetectable in Bat3−/− mice (Fig. 3, B and D). Hsp70-2 expression in spermatocytes correlated with the arrest and apoptosis observed in Bat3−/− mice (Fig. 3, E–L). The size (B) and weight of testes (C) and epididymides (D) were measured [n = 8]. Error bars represent SD. (E–L) Representative histological sections of hematoxylin and eosin–stained P7 (E and I), P14 [F and J], P42 [G and K], and P140 (H and L) testes and epididymides [M and N]. (O) Stage-specific expression of the indicated genes during spermatogenesis was analyzed by semiquantitative RT-PCR. GAPDH, loading control. Samples were prepared at the indicated developmental stages (P, postnatal day). (P–V) Increase in the number of TUNEL-positive Bat3−/− male GCs. TUNEL-positive cells were counted on P42 Bat3−/− and Bat3−/− sections (V). Bar graph represents mean ± SD (error bars; n = 20). Statistical significance was assessed using the unpaired t test. Bars: (B) 1 mm; (H and Q) 100 μm; [N and U] 20 μm.
Figure 2. **Bat3 is required for SC formation.** (A–N) Spermatocyte surface spreads were prepared from Bat3+/+ and Bat3−/− testes. SCs were visualized with SYCP3 antibody, and centromeres were stained with CEN antibodies (A and E). Analysis of SYCP3/γH2AX (B and F), SYCP3/Rad51 (C and G), SYCP3/RPA (D and H), SYCP3/TP1 (I and J), and TP1/DAPI (K–N). Arrows indicate abnormal SC formation in Bat3−/− samples (E). (O–T) Frozen sections from Bat3+/+ and Bat3−/− testes were stained with Bat3/SYCP3 (O and R) and SYCP3/γH2AX (P and S) antibodies. DNA was stained with DAPI (Q and T).

mice (Fig. 3, B and D). To eliminate the possibility that abnormal Hsp70-2 protein levels in Bat3−/− mice resulted from the loss of GCs at later developmental stages, we analyzed sections prepared at earlier stages. At P7, Hsp70-2 was not detected in either Bat3+/+ or Bat3−/− mice (Fig. 3, K and L). In contrast, Hsp70-2 became evident at P14 in Bat3+/+ but not Bat3−/− mice.
affected (Fig. 3 W), but Hsp70-2 protein levels were significantly lower in Bat3−/− GCs (Fig. 3 X). Therefore, we speculated that Bat3 deficiency might accelerate Hsp70-2 protein degradation.

We first examined whether Bat3 knockout or knockdown (KD) could reduce Hsp70-2 protein levels. Hsp70-2 levels were

(Fig. 3, Q and R). Nuclear staining indicated the presence of GCs in all of the Bat3−/− seminiferous tubules (Fig. 3 V).

We next investigated Hsp70-2 mRNA levels in Bat3+/+ and Bat3−/− testes. In Bat3+/+ testes, Hsp70-2 became expressed at P14, and Bat3 expression occurred at P7 (Fig. 3 W).

Interestingly, Hsp70-2 transcript levels were not significantly
We then examined whether Bat3 binds Hsp70-2. We transfected Flag–Hsp70-2 plus Myc-tagged full-length or deletion mutants of Bat3 into HEK293T cells (Fig. 5 A). Immunoprecipitation assays showed that Bat3 interacts with Hsp70-2 (Fig. 5 B) and that the BAG domain is essential for this interaction (Fig. 5 C), as shown previously for other members of the HSP70 family of proteins (Thress et al., 2001).

Next, we tested whether Bat3 KD could induce Hsp70-2 poly-Ub. We transfected U2OS cells with control or Bat3-targeted RNAi in combination with Flag–Hsp70-2 and HA-ubiquitin. Bat3 KD significantly increased the level of polyubiquitylated Hsp70-2 (Fig. 5 D). Hsp70-2 was also highly ubiquitylated in significantly lower in both Bat3−/− MEFs (Fig. 4 A) and multiple cell lines depleted for Bat3 (Fig. 4, B and C). We excluded potential off-target effects by introducing a Bat3 mutant resistant to RNAi (ΔRNAi-B3; Fig. 4 D; Sasaki et al., 2007). These data demonstrate that Bat3 is required for maintaining Hsp70-2 protein levels. We also found that Hsp70-2 protein levels were restored in Bat3−/− mouse primary GCs after treatment with MG132 (Fig. 4 E). Because Bat3 regulates the chaperone function of Hsp70 in vitro (Thress et al., 2001), we also examined Hsp70 protein levels. Notably, we found that Hsp70 protein levels were not significantly affected by Bat3 status (Fig. 4, A–C).

Figure 4. Bat3 is required for Hsp70-2 protein stability in multiple tissues. [A–C] Bat3, Hsp70-2, and Hsp70 protein levels were evaluated in Bat3+/+ and Bat3−/− MEFs [A], a human teratocarcinoma cell line, NTERA2 [B], and a mouse spermatocyte cell line, GC-2 [C], in the presence or absence of Bat3 with cycloheximide treatment for the indicated times (in hours). [D] Decreased Hsp70-2 protein stability induced by Bat3 KD was rescued by the RNAi-resistant Bat3 mutant (Bat3ΔRNAi). siCNT, control siRNA. (E) Hsp70-2 levels in primary Bat3-deficient male GCs were restored by proteasome inhibitor (MG132) treatment.
Bat3\(^{-/-}\) GCs (Fig. 5 E), suggesting that Bat3 negatively regulates the poly-Ub and subsequent degradation of Hsp70-2.

Because Hsp70 stability was not affected by Bat3 loss (Fig. 4, A–C), we speculated that the offending lysine (Lys) residues targeted by poly-Ub are specific to Hsp70-2. We aligned the amino acid sequences of Hsp70 and Hsp70-2 and identified unique Lys residues in Hsp70-2 (Fig. 5 G). Only the ΔN196 mutant lacking amino acids 1–196 could not be polyubiquitylated (Fig. 5 H). Further mutational analysis of Hsp70-2 identified Lys residues critical for poly-Ub (Fig. 5 I). These residues are conserved between human and mice (unpublished data). These data indicate that Bat3 binds and stabilizes Hsp70-2. Here, we propose a model in which the Bat3–Hsp70-2 interaction protects Hsp70-2 from ubiquitin ligase (Fig. 5 J).

We examined the expression of Bat3 and Hsp70-2 in Leydig and Sertoli cells. Hsp70-2 transcripts were undetectable in either cell type, whereas Bat3 transcripts were detectable in both (Fig. S2, available at http://www.jcb.org/cgi/content/full/jcb.200802113/DC1). The absence of Hsp70-2 in both cells was confirmed by immunostaining (Fig. S3). In addition, no significant differences in the staining pattern of calretinin (a Leydig cell marker) and Gata4 (a Sertoli cell marker) were observed in poly-Ub (Fig. 5 F). These results are conserved between human and mice (unpublished data). These data indicate that Bat3 binds and stabilizes Hsp70-2. Here, we propose a model in which the Bat3–Hsp70-2 interaction protects Hsp70-2 from ubiquitin ligase (Fig. 5 J).

Materials and methods

Histological analysis

For immunohistochemistry, paraffin-embedded sections were deparaffinized and rehydrated followed by antigen retrieval in boiled 10 mM sodium citrate buffer. Sections were blocked in 10% normal goat serum in PBS for 1 h at room temperature and incubated with primary antibody at 4°C overnight. After washing in PBS/0.3% Triton X-100 three times, the sections were incubated with secondary antibody for 1 h at room temperature. The sections were washed three times and mounted in ProLong Gold AntiFade Reagent (Invitrogen). TUNEL staining was performed using an In Situ Cell Death Detection kit (Roche) according to the manufacturer’s protocol. Primary antibodies used were polyclonal anti–mouse Bat3 antibody (Sasaki et al., 2007), Hsp70-2/HspA2 (M06; Abnova Corp.), calretinin (Santa Cruz Biotechnology, Inc.), and Gata-4 (Santa Cruz Biotechnology, Inc.). Secondary antibodies used were goat anti–rabbit AlexaFluor488 and goat anti–mouse AlexaFluor594 (Invitrogen). Hoechst33342 (Invitrogen) was used to visualize nuclei. Confocal images were acquired on a confocal microscope (LSM10; Carl Zeiss, Inc.) using 10×/0.5 NA, 20×/0.75 NA, and 63×/1.2 NA water immersion objectives. Three fluorophores (Hoechst33342, AlexaFluor488, and AlexaFluor594) were imaged sequentially using 351-, 488-, and 543-nm laser excitation and appropriate bandpass emission filters. LSM Image Browser (Carl Zeiss, Inc.) was used for capturing and overlaying images of different colors, and Photoshop CS2 (Adobe) was used to crop the images to proper size.

Immunostaining of surface-spread spermatocytes

Staining was performed as previously described (Dobson et al., 1994). The developmental stages of male GCs were determined based on the morphology and appearance of the XY bivalent and the localization of meiotic markers Rad51 and RPA. Chromosome cores and SCs were visualized using antibodies against SYCP3 and SYCP1 proteins (Dobson et al., 1994). Hsp70-2/RPA (provided by J. Ingle, University of Toronto, Toronto, Canada) and RAD51/DMC1 antibodies were previously described (Moens et al., 2002). Centromeres were visualized with human CREST serum (provided by S. Varmuza, University of Toronto, Toronto, Canada). TP-1 antibody was a gift from S. Kisler (University of South Carolina, Columbia, SC). γH2AX antibody was obtained from Abcam. Prolong Gold AntiFade reagent (Invitrogen) was used as a mounting agent to prevent fading of fluorescent signal. Rhodamine and FITC fluorochromes were used to visualize antibody-bound proteins, whereas DAPI fluorochrome allowed visualization of DNA. The images were acquired on a Polyvar microscope (model 602601; Carl Zeiss, Inc.) with oil immersion fluorescent objective lenses (Carl Zeiss, Inc.) of 40x/1.2 NA and 100x/1.32 NA magnification and were recorded with a cooled digital color camera (DP70; Olympus) using Image Pro 4.0 software (Media Cybernetics, Inc.). For image processing, Photoshop 7.0 was used. Using Photoshop, the brightness and contrast were enhanced, and the two images of different colors were overlapped and cropped to proper size.

RT-PCR

Sertoli and Leydig cells were isolated from three wild-type mice as previously described (Anway et al., 2003). Total RNA was extracted from the cells and tested using TRIZOL (Invitrogen). First-strand cDNA was synthesized using a SuperScript First-Strand kit (Invitrogen) according to the manufacturer’s instructions. Gene-specific primers are listed in Table S2 (available at http://www.jcb.org/cgi/content/full/jcb.200802113/DC1).

Cell lines, transfections, and RNAs

U2OS, 293T, NTERA2, and GC-2 cells were obtained from the American Type Culture Collection. Synthetic siRNA oligoduplexes for Bat3 as previously described (Sasaki et al., 2007) were purchased from Dharmacon. Cells were cultured to 60% confluency in Opti-MEM medium (Invitrogen) and transfected with siRNA oligoduplexes (40-nM final concentration) plus 3 µl Lipofectamine 2000 (Invitrogen). At 6–8 h after transfection, the medium was changed to complete medium. Transfected cells were assayed 48 h after transfection unless otherwise indicated. All control samples were transfected with control siRNA oligoduplex (Dharmacon).

Western blot analysis

Protein samples were prepared in lysis buffer (0.1% SDS, 0.1% sodium deoxycholate, 1% Triton X-100, 20 mM Tris-HCl, pH 7.5, 150 mM NaCl,
Figure 5. Bat3 binds and negatively regulates the poly-Ub of Hsp70-2. (A–C) The Bat3 BAG domain is required for interaction with Hsp70-2. 293T cells were transfected with Flag–Hsp70-2 and a series of Myc-Bat3 deletion mutants [A]. Lysates were immunoprecipitated with anti-Flag antibody and immuno-blotted with anti-Myc antibody [B and C]. (D) Negative regulation of Hsp70-2 poly-Ub by Bat3. Flag–Hsp70-2 and HA-Ub plus either siBat3 or siCNT (control siRNA) were transfected into U2OS cells. The lysates were immunoprecipitated and immunoblotted with anti-HA antibody. (E) Increased poly-Ub
1 mM EDTA, 10 mM NaF, and protease inhibitor cocktail [Roche]). 20 μg of lysates was immunoblotted according to standard procedures. Proteins were transferred to an Immobilon-P membrane [Millipore]. Blots were incubated with the primary antibody at 4°C overnight and with a horseradish peroxidase-conjugated secondary antibody at room temperature for 1 h. Signals were detected using Enhanced Chemiluminescence reagents [GE Healthcare]. Primary antibodies used were as follows: anti-mouse [Sasaki et al., 2007] and anti-human Bat3 (a gift from S. Karnbluth, Duke University, Durham, NC); anti-Hsp70 (K-20) and anti-HA (F-7; Santa Cruz Bio-technology, Inc.); and anti-Flag M2 and anti-actin [Sigma-Aldrich].

**Immunoprecipitations**

293T cells were transfected with Flag–Hsp70-2 plus pcDNA3 control vector (Invitrogen) or Myc-Bat3. Cells were lysed in buffer (50 mM Tris-HCl, 150 mM NaCl, and 1% NP-40) supplemented with a protease inhibitor cocktail [Roche] for 30 min on ice and cleared by centrifugation. Extracts were incubated with the appropriate antibodies at 4°C for 1 h with rotation and incubated for an additional 2–3 h with protein A/G agarose (Invitrogen) or Myc-Bat3. Cells were lysed in buffer (50 mM Tris-HCl, 150 mM NaCl, and 1% NP-40) supplemented with a protease inhibitor cocktail [Roche] for 30 min on ice and cleared by centrifugation. Extracts were incubated with the appropriate antibodies at 4°C for 1 h with rotation and incubated for an additional 2–3 h with protein A/G agarose beads (Santa Cruz Biotechnology, Inc.). Samples were fractionated by SDS-PAGE and subjected to immunoblotting as described in the previous section.

**Hormone measurement**

Serum levels of testosterone, LH, and FSH were measured in P40 Bat3+/− and Bat3−/− mice. LH and FSH were measured by the Center for Research in Reproduction Ligand Assay and Analysis Core of the University of Virginia. Testosterone was measured by an EUSA kit according to the manufacturer’s protocol [Assay Designs].

**Online supplemental material**

Fig. S1 shows additional pictures for SYCP3–/H2AX staining of SCs. Fig. S2 demonstrates that Hsp70-2 transcripts are undetectable in Leydig and Sertoli cells. Fig. S3 shows that Hsp70-2 protein is also undetectable in Leydig and Sertoli cells. Table S1 shows that Bat3 deficiency does not affect serum levels of testosterone, LH, or FSH. Table S2 lists RT-PCR primer sets used in this study. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200802113/DC1.

This paper is dedicated to Peter B. Moens and his outstanding contributions in the field of reproductive cell biology. We sincerely thank Sally Karnbluth for helpful discussion, human Bat3 antibody, and human myc-Bat3 plasmid; Steve Kistler for TP1 antibody; Andrew Vakeham for embryonic stem cell injection; Andrea Jurisicova and Andras Nagy for help in the initial stage of the project; BarbaraSpyropoulosforantibodypreparation;andEmmaGoolingandDavidSecundy for editing. H. Okada is supported by a grant from the Canadian Institutes of Health Research [MOP84353]. The University of Virginia Center for Research in Reproduction Ligand Assay and Analysis Core is supported by National Institute of Child Health and Human Development/National Institutes of Health [Specialized Cooperative Centers Program in Reproduction and Infertility Research] grant U54HD28934.

Submitted: 19 February 2008
Accepted: 7 July 2008

**References**


Table S1. *Bat3 deficiency does not affect hormone levels*

<table>
<thead>
<tr>
<th>Hormone/genotype</th>
<th>+/+ (n = 4)</th>
<th>--/-- (n = 4)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Testosterone (pg/ml)</td>
<td>1,838.1 ± 74</td>
<td>1,834.0 ± 51</td>
</tr>
<tr>
<td>LH (ng/ml)</td>
<td>0.22 ± 0.10</td>
<td>0.20 ± 0.04</td>
</tr>
<tr>
<td>FSH (ng/ml)</td>
<td>31.25 ± 7.8</td>
<td>33.56 ± 4.1</td>
</tr>
</tbody>
</table>

Data represent mean ± SEM.

Table S2. *List of RT-PCR primers*

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward (5'–3')</th>
<th>Reverse (5'–3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dazl</td>
<td>CCTCAACCATGATGAATCC</td>
<td>TCTGTATGCTTGGGCCAC</td>
</tr>
<tr>
<td>Plzf</td>
<td>TGAGATCTCTCCACCGAA</td>
<td>GTAGGACTCATGGGTGAGGA</td>
</tr>
<tr>
<td>Dmc1</td>
<td>GGAGGATCAAGTTGTGCAGGA</td>
<td>ATCCCTGCTCAGCTTAGA</td>
</tr>
<tr>
<td>Pgk2</td>
<td>CTGTTGCTGATGAGCTCAAG</td>
<td>ACTCCGACCATAAGACTGT</td>
</tr>
<tr>
<td>Prm1</td>
<td>ATGCTGCCGACGCAAAGCA</td>
<td>CACCTTATGGTATGAGCG</td>
</tr>
<tr>
<td>Gata-1</td>
<td>CGGTTTTCTCTTCTCTGGG</td>
<td>AAAGGACTGGGAAAATCG</td>
</tr>
<tr>
<td>Bat3</td>
<td>CCCATGAGGTCTAGAAGGATGAATGG</td>
<td>GGAGTCTGATAGGACATAGG</td>
</tr>
<tr>
<td>Hsp70-2</td>
<td>CTACGTGGGCTCCACTGACA</td>
<td>GTCAAGATGACACATCGA</td>
</tr>
<tr>
<td>Gapdh</td>
<td>ATGTCAGCAATGCACTCTG</td>
<td>TTTAGCTGGGATGACCTTGC</td>
</tr>
</tbody>
</table>
