A novel Amh-Treck transgenic mouse line allows toxin-dependent loss of supporting cells in gonads

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Abstract

Cell ablation technology is useful for studying specific cell lineages in a developing organ in vivo. Herein, we established a novel anti-Müllerian hormone (AMH)-toxin receptor-mediated cell knockout (Treck) mouse line, in which the diphtheria toxin (DT) receptor was specifically activated in Sertoli and granulosa cells in postnatal testes and ovaries respectively. In the postnatal testes of Amh-Treck transgenic (Tg) male mice, DT injection induced a specific loss of the Sertoli cells in a dose-dependent manner, as well as the specific degeneration of granulosa cells in the primary and secondary follicles caused by DT injection in Tg females. In the testes with depletion of Sertoli cell, germ cells appeared to survive for only several days after DT treatment and rapidly underwent cell degeneration, which led to the accumulation of a large amount of cell debris within the seminiferous tubules by day 10 after DT treatment. Transplantation of exogenous healthy Sertoli cells following DT treatment rescued the germ cell loss in the transplantation sites of the seminiferous epithelia, leading to a partial recovery of the spermatogenesis. These results provide not only in vivo evidence of the crucial role of Sertoli cells in the maintenance of germ cells, but also show that the Amh-Treck Tg line is a useful in vivo model of the function of the supporting cell lineage in developing mammalian gonads.

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Introduction

In mammalian embryos at early organogenic stages, undifferentiated genital ridges start to develop into the testis or ovary dependent on their genetic sex. As the first morphological sign of testis differentiation, germ cells are surrounded by differentiating Sertoli cells, leading to the formation of testis cords (future seminiferous tubules), a tubular structure of the germ and Sertoli cells packed within the basal lamina sheath (Harikae et al. 2013a, Svingen & Koopman 2013). In developing ovaries, both germ and pre-granulosa cells also aggregate to form the poorly defined ovarian cords (ovigerous cord), subsequently resulting in the formation of ovarian cysts, a closed sack-like structure of several oocytes and their surrounding pre-granulosa cells (Pepling 2006, Hummitzsch et al. 2013). After birth, the testis cords develop into the seminiferous tubules associated with active spermatogenesis, while in the ovary the ovarian cyst breaks down into each individual primordial follicle, an oocyte surrounded by a single layer of granulosa cells. A part of the primordial follicle continuously grows into primary and subsequent secondary follicle stages at the medullary side during the initial round of folliculogenesis (Mork et al. 2012, Harikae et al. 2013b).

From the fetal to adult stages, both Sertoli and granulosa cells secrete various growth factors to support the proliferation, differentiation and maintenance of the germ cells (Matsui 1998, Choi & Rajkovic 2006, Buratini & Price 2011, Zhang et al. 2011, Feng et al. 2014). Moreover, the germ cells physically contact and interact intimately with the supporting cells inside the basal lamina sheath. These histological and physiological findings indicate the crucial roles of the gonadal supporting cells in the maintenance and survival of germ cells in mammalian gonads. However, the duration...
for which the germ cells are maintained after the specific loss of supporting cells in vivo remains unclear.

A conditional in vivo depletion of lineage-specific cells can be achieved through administration of diphtheria toxin (DT) to toxin-receptor-mediated cell-knockout (Trek)-transgenic (Tg) mice that carry a human DT receptor (DTR; \(H\)BEGF-mut) cDNA transgene driven by a cell lineage-specific promoter (Saito et al. 2001). In this Trek system, the administration of DT to the Trek Tg mice causes depletion in only DTR-expressing cell lineages and specifically eliminates these target cells in vivo (Takada et al. 2008, Matsuoka et al. 2013).

In mice, anti-Müllerian hormone (\(Amh\)) is expressed specifically in the gonadal-supporting cell lineages: Sertoli and granulosa cells of postnatal immature gonads in both males and females (Lécureuil et al. 2002). Herein, we established three \(Amh\)-Trek Tg lines carrying the \(Amh\)-promoter-fused DTR cassette, in which conditional in vivo deletion of the gonadal-supporting cells was induced by DT injection at postnatal timepoints. Here we show a great potential in the \(Amh\)-Trek line as an in vivo functional assay system of the supporting cell lineage in mammalian gonads.

**Materials and methods**

### \(Amh\)-Trek Tg mouse line

The \(Amh\)-Trek transgene was prepared by inserting the entire 5′-flanking promoter region of the \(Amh\) gene (Guerrier et al. 1990) upstream of DT receptor (DTR; \(H117V/L148V\) mutant of human HB-EGF tagged with EGFP; Furukawa et al. 2006). We successfully established three \(Amh\)-Trek Tg lines (Takada et al. 2001, Takada et al. 2008, Matsuoka et al. 2013) carrying the \(Amh\)-promoter fused DTR cassette in which condition in vivo deletion of the gonadal-supporting cells was induced by DT injection at postnatal timepoints. Here we show a great potential in the \(Amh\)-Trek line as an in vivo functional assay system of the supporting cell lineage in mammalian gonads.

### Quantitative RT-PCR

Total RNA was reverse-transcribed using random primer with a Superscript-III cDNA Synthesis Kit (Invitrogen). PCR was carried out using an Applied Biosystems Step-One Real-Time PCR System with the primers and probes of \(Egfp\) (Mr04329676_mr) and \(Gapdh\) (Taqman control reagents).

**DT treatment**

\(Amh\)-Trek Tg mice (P3–21) were treated with DT (Sigma-Aldrich, 0.04–40.0 \(\mu\)g/kg, i.p.) as described previously (Saito et al. 2001, Takada et al. 2008, Matsuoka et al. 2013).

All tissues were subjected to histological and immunohistochemical analyses as described below.

## Transplantation of EGFP or Cherry-positive Sertoli cells

A single-cell suspension (including gonocytes and Sertoli cells) (1.0×10^6 cells/ml) was prepared from the testes of 1-week-old GFP/Cherry mice (C57BL6) by a two-step enzymatic digestion procedure (Shinozuka et al. 2003), and then transplanted into the testes of 3-week-old recipient Tg males pre-treated with DT (0.4–4.0 \(\mu\)g/kg). On days 10 and 45 after transplantation, all recipient Tg testes were removed from the tunica albuginea and roughly dissected in cold PBS under an Olympus fluorescence microscope (BX51N-34-FL2) and stereomicroscope (SZX16 plus U-LH100HG) (Nagai et al. 2012). The tissues were processed using the immunohistochemical techniques described below.

## Histology and immunohistochemistry

The tissues were fixed in 4% paraformaldehyde (PFA) or Bouin’s solution and routinely embedded in paraffin. The sections (4 \(\mu\)m thickness) were subjected to conventional histological (hematoxylin and eosin; H&E) and immunohistochemical staining as described previously (Hiramatsu et al. 2010, Harikae et al. 2013b).

In brief, for immunohistochemical staining, the sections were incubated with anti-\(AMH\) (1:100 dilution; sc-6886; Santa Cruz), anti-\(\beta\)-HSD (1:2000 dilution; sc-30821; Santa Cruz), anti-\(\beta\)-Gal (1:1000 dilution; sc-1237; Santa Cruz), anti-GFP (1:200 dilution; ab5096; Abcam), anti-GATA4 (1:250 dilution; sc-1237; Santa Cruz), anti-GFP (1:200 dilution; 598; MBL, Nagoya, Japan), anti-GFRα1 (1:100 dilution; AF560; R&D Systems, Minneapolis, MN, USA), anti-HuB/Elavl2 (1:1000 dilution; H1538; Sigma), anti-laminin (1:400 dilution; 10765; ICN Pharmaceuticals, Santa Ana, CA, USA), anti-MHJV/ \(\alpha\)-smooth muscle actin (anti-SMA) (1:500 dilution; A2547; Sigma), or anti-SOX9 (1:250 dilution; H2M Shinomura and others) antibody at 4°C for 12 h. The reaction was visualized using a biotin-conjugated secondary antibody in combination with the Elite ABC Kit (Vector Laboratories) or by secondary antibodies conjugated with Alexa-488/594. TUNEL assays were also performed by using the Apoptotic Detection Kit (MK500; TaKaRa, Shiga, Japan).

## Results and discussion

### Establishment of \(Amh\)-Trek Tg mouse lines

The transgene was constructed as the \(Amh\) promoter-driven DT receptor (DTR; \(H\)BEGF-mut) (\(H117V/L148V\) mutant of human HB-EGF) tagged with EGFP (Furukawa et al. 2006; Fig. 1A). We obtained three Tg lines (healthy and fertile in both sexes) by pronuclear injection of the plasmid-based transgene into the C57BL6 background. In Wt mice, endogenous \(Amh\) expression is found in the

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Figure 1 Characterization of AMH-Trecc Tg lines (#94). (A) Schematic representation showing the transgene of AMH promoter-driven diphtheria toxin receptor (DTR; HBEGF-mut) fused with EGFP. (B) H&E staining and anti-SOX9 immunostaining of the testes on day 4 after DT treatment, showing a dose-dependent degeneration of SOX9-positive Sertoli cells in Tg males treated with DT at 0, 0.04, 0.4, and 4.0 µg/kg. (C) H&E and TUNEL staining showing a time-course pattern of the Sertoli cell and germ cell degeneration in the Tg males treated with DT at 4.0 µg/kg. (D) Anti-GFRα1 and HuB immunostaining of the testes on day 4 (two left-handed rows) and day 10 (most right-handed row) after DT treatment, showing the dynamics of undifferentiated (GFRα1+ and HuB+) spermatogonia in the Wt and Tg littermates treated with DT at 4.0 µg/kg. Both GFRα1+ and HuB+ populations of the spermatogonia disappear by day 10 after DT treatment. Scale bars represent 20 µm.

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Sertoli cells of the testes from embryonic day (E) 12.5 to postnatal day (P) 14 (Supplementary Figure S2, see section on supplementary data given at the end of this article) and then almost repressed by P24 (Al-Attar et al. 1997). In the ovary, Amh expression is defined in the growing follicles after P3 and later stages (Lécureuil et al. 2002). The preliminary experiments carried out with DT treatment (4.0 μg/kg; Matsuoka et al. 2013) in the postnatal testes at P3–P10 showed degeneration in the majority of Sertoli cells in all of the three lines as well as no appreciable defects in other organs (Fig. 1 and Supplementary Figure S3). Among these three lines, the EGFP-tagged DTR expression was, although fluorescent signals are very weak, detectable in the seminiferous tubules and the primordial follicles of the testes and ovaries in the #94 line (Supplementary Figure S4), in addition to the gonad-specific high-level expression of the DTR transgene by qPCR (Supplementary Figure S1). Moreover, even a low-dose DT treatment (0.04 μg/kg) can affect the nuclear shape of Sertoli cells and their arrangement in the #94 line (see Fig. 1B), and thus we used mainly the #94 line for subsequent experiments.

DT injection induces Sertoli cell degeneration in postnatal testes of Tg males in a dose-dependent manner

First, we examined the dose-dependent effects of DT on the Sertoli cell degeneration in the Amh-Treck line. The Tg males and their Wt littermates at P21 (post-weaning) were treated with DT at various dosages, from 0.04 to 40.0 μg/kg, and their organs including testes were histopathologically examined on day 4 after DT treatment (Fig. 1B and Supplementary Figure S5, see section on supplementary data given at the end of this article). In the Tg males treated with 0.04 μg/kg DT, most of Sox9-positive Sertoli cells appeared morphologically healthy but the nucleus position was arranged irregularly in the seminiferous epithelia as compared with the non-treated control testes (Fig. 1B). DT treatment at either 0.4 or 4.0 μg/kg clearly induced the loss of the Sox9-positive Sertoli cells in the seminiferous tubules by day 4 after DT treatment (Fig. 1B). In contrast, no appreciable defects were found in the Wt mice treated with DT at 4.0–40 μg/kg (Supplementary Figure S5). In addition, no gross-anatomical and histological defects were observed in the other organs of both Wt and Tg males treated with 4.0–40 μg/kg DT (Supplementary Figure S5; data not shown).

Next, we examined the temporal changes in the testicular phenotype in DT-treated Tg testes (4.0 μg/kg) between days 1 and 10 after DT treatment (Fig. 1C). On day 1, DT treatment showed onset of Sertoli cell degeneration; however, histological loss of the nucleus was observed on day 4 after DT treatment. Moreover, histological analysis revealed that the germ cells appeared to undergo germ cell degeneration with visible accumulation of cell debris at the luminal side on day 4. Finally, by day 10 after DT treatment, all Sertoli and germ cells were degenerated in the seminiferous tubules and consequently, a large amount of cell debris had accumulated inside the seminiferous tubules (Fig. 1C). TUNEL staining also confirmed the appearance of apoptotic cell death in the germ cells at luminal side by day 4 after DT treatment (lower plates in Fig. 1C; Supplementary Figure S6, see section on supplementary data given at the end of this article). Anti-GFRα1 and anti-HuB immunostaining revealed that the GFRα1+ undifferentiated spermatogonia (including spermatogenic stem cells; Sato et al. 2011) and HuB+ differentiated spermatogonia (including pre-leptotene spermatocytes; Supplementary Figure S7) are detectable in the basal compartment of the seminiferous tubules on day 4 after DT treatment. However, both spermatogonial populations were depleted by day 10 after DT treatment (Fig. 1D). These findings, therefore, confirmed that loss of Sertoli cells led to rapid cell death of germ cells within several days. This is consistent with previous reports involving Sertoli cell-specific deletion of Wt1 (Gao et al. 2006) or Sox9/Sox8 (Barrionuevo et al. 2009) gene(s), which demonstrated a drastic reduction in germ cell number in the postnatal testes, together with defective Sertoli cell function.

Sertoli cell-specific loss of the seminiferous tubules in DT-treated Tg testes at P3 and P21

The specificity of the cell depletion in DT-treated testes was examined on day 4 after DT treatment. The male Tg pups and their Wt littermates at P3 and P21 were treated with DT (4.0 μg/kg) and the expression patterns of various somatic cell markers were examined immunohistochemically (Fig. 2A). In P3 testes treated with DT (upper plates in Fig. 2B), the almost complete depletion of the Amh-positive or Sox9-positive Sertoli cells was found throughout the entire testis area. This is in sharp contrast with E-cad-positive rete testis epithelia, which appeared histologically normal (Fig. 2B). Immunohistochemical analyses of P21 testes on day 4 after DT treatment also confirmed that GATA4-positive cells appeared intact in the peritubular and interstitial regions. However, no GATA4-positive signals were detectable in the seminiferous epithelia, leading to MVH-positive germ cells occupying the inside of the tubules (Fig. 2C). The laminin-positive basal lamina, SMA-positive peritubular myoid cells, and 3β-HSD-positive Leydig cells appeared intact in the interstitial region (Fig. 2C). The complete degeneration of cells inside the tubules by day 10 after DT treatment indicates that Sertoli cell-specific depletion occurs by day 4; germ cells then undergo rapid degeneration, which may result in the accumulation of a large amount of cell debris at the luminal side by day 10 after DT treatment.
Transplantation of healthy immature Sertoli cells can rescue the germ cell degeneration in DT-treated Tg testes, leading to a partial recovery of the spermatogenesis.

The intratubular injection method of the cell suspension prepared from immature testes previously showed that immature dividing Sertoli cells selectively colonize and support the recipient-derived spermatogenesis in the empty seminiferous tubules pre-treated with the Busulfan and cadmium (Shinohara et al. 2003). In order to rescue the germ cell degeneration, healthy immature Sertoli cells (around P7) labeled by cytoplasmic/nuclear EGFP...
Figure 3 A rescue experiment by transplantation of the healthy Sertoli cells into DT-treated Tg testes. (A) Schematic representation of Sertoli cell transplantation on day 4 and day 45 after DT injection (0.4–4.0 μg/kg). (B) Various degrees of contribution of donor-derived (EGFP/Cherry-positive) Sertoli cells in each testis on day 10 and day 45 after Sertoli cell transplantation. (C, D, E, and F) Donor-derived cytoplasmic/nuclear EGFP and nuclear Cherry signals in the transplanted tubules (lower magnified images; C and E) and EGFP/Cherry signals (green/red) and anti-SOX9 (brown in D, green in F) immunostaining images of the same testicular section (D and F), showing the contribution of donor-derived healthy Sertoli cells (arrowheads) in the DT-treated testis on day 10 after transplantation. In the most right-handed images of plates D and F, transplanted Sertoli cells directly support the host-derived germ cells (asterisks) in DT-treated Tg testes. (G and H) Anti-Cherry (red) and anti-SOX9 (green in H) double immunostaining images showing spermatogenic patches derived from both endogenous (asterisks) and exogenous (open arrow) germ cells in the transplanted DT-treated Tg testis on Day 45 after transplantation. Transplanted Sertoli cells directly support spermatogenesis of the host-derived germ cells (asterisks). In the host, plates C and D are the testis pre-treated with DT at 0.4 μg/kg, while plates E, F, G, and H are the testes pre-treated with DT at 4.0 μg/kg. Scale bars represent 200 μm in C, E, and G; 10 μm in D; 20 μm in F, H.
Figure 4 Specific depletion of the granulosa cells of AMH-positive presumptive primary and secondary follicles in developing ovaries. (A and B) Anti-AMH immunostaining and H&E staining images of the P7 ovaries of Tg and Wt littermates (DT, 40.0 µg/kg treatment at P3). In B, the ovarian surface region including an AMH-negative primordial follicle pool (indicated by bars) is shown on the left side. (C) Higher magnified images of H&E, anti-FOXL2, anti-MVH, anti-laminin and anti-SMA staining patterns in the presumptive primary and secondary follicles located in the centromedullary region of DT-treated ovaries (Tg: upper plates, Wt: lower plates). FOXL2-positive granulosa cells are missing in the affected follicles of the DT-treated Tg ovary, despite no differences in anti-laminin-/anti-SMA-immunostaining patterns between Tg and Wt ovaries. (D) Gross-anatomical, H&E staining and anti-3β-HSD/anti-AMH immunostaining images of the P28 ovaries of Tg and Wt littermates treated with DT (40.0 µg/kg) at P3 (day 25 after DT treatment). The new recruitment of AMH-positive primary and secondary follicles is shown in the DT-treated Tg ovary, and both the ovary size and the 3β-HSD-positive area are reduced in the Tg ovary. Nuclei are counterstained with hematoxylin in blue. In A and D, each inset shows a higher-magnification image of the area indicated by the broken rectangle. Scale bars represent 100 µm in A, B, and D; 20 µm in C.
Reproduction males, on day 4 after DT treatment (0.4–4.0 (CAG-EGFP) or nuclear Cherry (R26-H2B-mCherry) H8 testes); Fig. 3 B). Histological analysis revealed that a injected: 10/18 testes (EGFP, 4/8 testes; Cherry, 6/10 (number of testes with donor cells per total testes transplanted with exogenous healthy testis cells, the transplanted testes were examined on day 10 after Sertoli function. Indeed, this new Treck system is more simple transplantation studies in terms of evaluating Sertoli cell structural and endocrine roles of the supporting cell lineage at the postnatal stages, but also potentially for system is also useful not only for supporting cells at the postnatal stages using DT treatment. The present data confirm the essential role supporting cells at the postnatal stages using DT treatment. The present data confirm the essential role of Sertoli cells in germ cell survival in vivo. This model system is also useful not only for in vivo analyses of structural and endocrine roles of the supporting cell lineage at the postnatal stages, but also potentially for in vivo functional assay of defective supporting cells in stem cell niche and spermatogenesis.

**Supplementary data**

This is linked to the online version of the paper at http://dx.doi.org/10.1530/REP-14-0171.

**Declaration of interest**

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

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