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FOXC2 marks and maintains the primitive spermatogonial stem cells subpopulation in the adult testis

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Abstract

In adult mammals, spermatogenesis embodies the complex transition from spermatogonial stem cells (SSCs) to spermatozoa. This process is initiated by the dynamic transition among a series of SSCs subpopulations. However, it remains elusive and controversial for the identity of the primitive adult SSCs at the top of this developmental hierarchy. Using single-cell analysis and lineage tracing, we identified forkhead box protein C2 (FOXC2) as a specific marker for the primitive SSCs subpopulation in adult mice and humans. During homeostasis, $FOXC2^{+}$-SSCs can initiate spermatogenesis, and through which give rise to all sets of spermatogenic progenies. Specific ablation of the $FOXC2^{+}$-SSC results in depletion of the undifferentiated spermatogonia pool. During germline regeneration, spermatogenesis can be completely restored by $FOXC2^{+}$-SSCs. Germ cell-specific *Foxc2* knockout resulted in accelerated exhaustion of SSCs and eventually led to male infertility. Mechanistically, FOXC2 is required for maintaining the quiescent state of the primitive SSCs by promoting the expression of negative regulators of cell cycle phase transition. Overall, this work proposed $FOXC2^{+}$-SSCs as an indispensable and primitive subgroup during homeostasis and regeneration in the adult testis.

eLife assessment

This **important** study reports Foxc2+ cells in the testis might be the true spermatogonial stem cells (SSCs). The data supporting this claim are **solid** and the finding, if proven true, would have a great impact on reproductive biology and stem cell biology as the genes responsible for maintaining the quiescent state of SSCs during spermatogenesis remain elusive.

Introduction

Through spermatogenesis, spermatozoa are generated from spermatogenic cells that are originated from spermatogonial stem cells (SSCs). It is critical for this process to be continuous and successful that SSCs are maintained in a homeostatic balance between self-renewal and differentiation (1). The SSCs, as the least differentiated spermatogonia, belong to a subgroup of undifferentiated spermatogonia (uSPG) that are morphologically categorized into three subtypes, i.e., A_{single} (As), A_{paired} (Apr), and $A_{aligned}$ (A_{al}) cells (2). So far, three models have been proposed for the mechanism underlying SSCs' self-renew based on the dynamic transitions among subgroups. In the 'A_s model', A_s spermatogonia serve as SSCs that are capable of both self-renew and further transformation into A_{pr} and A_{al} that eventually give rise to spermatogonia (3; 4). Later, based on the discovery of Ngn3 and Gfrα1 as SSCs markers, the 'fragmentation model' suggests all three subgroups with stem cell potential and the SSCs renewal is achieved through the fragmentation of pairs and chains (5). Further work on SSCs markers such as ID4 and PAX7 inspired the 'hierarchical A_s model', in which only specific A_s spermatogonia possess the potential for long-term self-renewal whereas the majority are restricted in their capacity (6; 7). Though their standing points of view differ, each model seems well supported by the respective collection of evidence, which to some extent reflects the nature of heterogeneity and dynamics among SSCs subpopulations.

In recent years, great insights into SSCs behaviors and regulations have been provided by a body of pioneer works, especially with recent advances in single-cell gene-expression profiling, highlighting great heterogeneity of SSCs and focusing on characterizing the nature of SSCs states especially for seeking the primitive subgroup among them. Within the population of uSPG, a number of genes relatively higher expressed in primitive subfractions have been identified and well investigated, i.e., Gfra1, ID4, Ret, Eomes, Pax7, Nanos2, Shisa6, T, Pdx1, Lhx1, Egr2 and Plvap ((5)–(15)). Particularly, Gfra1, ID4, Eomes, Pax7, Nanos2, and Plvap are further validated as the SSCs markers through lineage tracing experiment, which is considered to be a reliable method to study the origin and development of stem cells. However, some essential and primitive sub-populations remain undiscovered, and the identification of which is of great significance for elucidating the developmental process of SSCs renewal and its behavior in testis.

Adult stem cells (ASCs), as the undifferentiated primitive cells that can be found in nearly all types of tissues in mammals, are characteristic for a unique quiescent status reflected by both reversible cell cycle arrest and specific metabolic alterations (16). Putative the primitive SSCs subgroups appear to share this characteristic, as revealed in recent single-cell RNA-sequencing (scRNA-seq) analysis in humans and mice, being largely non-proliferative while capable of reciprocating between the quiescent and activated status ((17)–(21)). However, rigorous biological validation of these populations is lacking through live imaging or genetic lineage tracing, or other means. On the other hand, cells in a quiescent state are supposed to be more resilient to genotoxic insults, which shall enable the primitive SSCs to sufficiently restore spermatogenesis upon such disturbance.

Here, we identified a subpopulation of adult SSCs specifically marked by forkhead box protein C2 (FOXC2). In adult mice, spermatogenic cells derived from the $FOXC2^+$ population were able to complete the whole spermatogenesis. Upon the loss of this specific subpopulation of SSCs, the undifferentiated spermatogonia pool was exhausted, eventually leading to defective spermatogenesis. Specifically, FOXC2 is required for maintaining SSCs quiescence by promoting the expression of negative regulators of cell cycle phase transition, thus symbolizing the primitive state of these adult SSCs. Moreover, the $FOXC2^+$ population endured the chemical insult with busulfan and effectively restored spermatogenesis, thereby critical for keeping the reproductive homeostasis in male adult mice. Thus, our

results demonstrate that FOXC2 marks the primitive SSCs subpopulation in the adult testis, and is also required for the homeostasis and regeneration of SSCs.

Results

Identification of FOXC2^{+}-SSCs as the quiescent and developmental starting point of adult uSPG

We performed single-cell RNA-seq (10x genomics) of the uSPG from adult mice testes marked by THY1, a widely recognized surface marker for uSPG with self-renewing and transplantable state (22; 23), to dissect the heterogeneity and developmental trajectory (Fig.1A, Fig. S1A, B). Among 5 distinct clusters identified, Cluster1 was characterized by the high expression of stemness markers whereas other clusters were featured by progenitor or differentiating spermatogonia (dSPG) markers (Fig.1B, Fig. S1C, D). Primarily mapped to the extreme early point of the developmental trajectory, Cluster1 cells appeared quiescent and likely represented the primitive state of uSPG populations (Fig.1B, Fig. S1E-G). The top10 differentially expressed genes (DEGs) associated with Cluster1 are featured by SSCs markers such as *Mcam* (24), *Gfra1* (5), *Tcl1* and *Egr2* (12; 18) (Fig.1C, Fig. S2A, Supplemental Table S1) in addition to six others expressed in different stages of germ cells and/or somatic cells, in which only FOXC2 was exclusively localized in the nucleus of a subgroup of ZBTB16^{+} uSPG (25; 26) in mice (Fig. 1D, Fig. S2B). More specifically, in adult mice, FOXC2 displayed differential expressions among various subtypes of uSPG, being more specific in As (59.9%) than other subtypes including A_{pr} (5.2%), A_{pr-} 1 (4.1%), A_{al4-1} (1.83%), A_{al8-1} (1.5%), and A_{al16-1} (1.67%) (Fig. 1E). There was only a small fraction (5.1%) was active in proliferation as indicated by MKI67 (Fig. 1F), suggesting that FOXC2^{+} cells are primarily quiescent. Additionally, when examining the SSCs markers validated previously by lineage tracing (27), we found that FOXC2 displays a higher level of co-localization with GFRA1 and EOMES than PAX7 and NEUROG3 (28), indicating the FOXC2^{+} cells contain but differ from the known SSCs subsets (Fig. 1G).

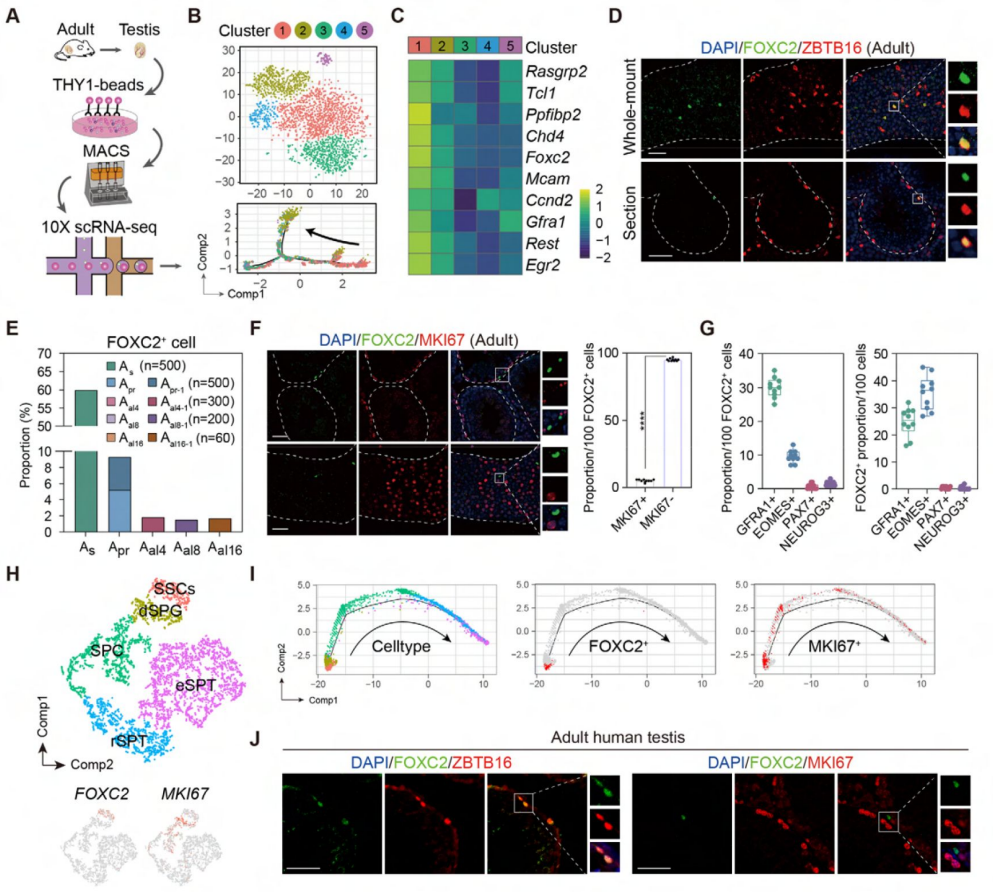

Figure 1.

Identification of the FOXC2$^+$-SSCs in adult mouse and human testis.

(**A**) Schematic illustration of the single-cell analysis workflow. (**B**) t-SNE plot and developmental trajectory of all uSPG, colored by cluster. (**C**) Heatmap of the Top10 DEGs in Cluster1. (**D**) Immunostaining for ZBTB16 (red), FOXC2 (green), and DAPI (blue) in testicular paraffin sections from wild-type adult C57 mice. Scale bar, 50 μm; C57, C57BL/6J. (**E**) The proportion of FOXC2$^+$ cells in different uSPG subtypes. (**F**) Immunostainings for MKI67 (red), FOXC2 (green), and DAPI (blue) in adult mice testis and the proportion of MKI67$^+$ cells in FOXC2$^+$ population (n=10). Scale bar, 50 μm; values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (****p-value < 0.0001). (**G**) The co-expression proportion between the FOXC2 and differential known SSCs makers (n=10). (**H**) t-SNE plot of germ cells in adult human testis (GSE112013), colored by germ cell type. Feature plot showing the expression patterns of *FOXC2* and *MKI67* in human germ cells. (**I**) The developmental trajectory of the human germ cells, colored by germ cell type, FOXC2 expression cells (red), or MKI67 expression cells (red). (**J**) Immunostaining for ZBTB16/MKI67 (red), FOXC2 (green), and DAPI (blue) in testicular paraffin sections from adult humans.

We next analyzed the expression of FOXC2 in adult human testis using the published scRNA-seq dataset (17) (GSE112013). As expected, FOXC2 was also specifically expressed in the human SSCs, most of which were MKI67$^-$ (Fig. 1H, Fig.S2C). Pseudotime analysis showed that the FOXC2$^+$ cells located at the start of the developmental trajectory with a proportion of about 90% that were MKI67$^-$ (Fig. 1I). Immunofluorescence staining confirmed that FOXC2$^+$ cells were a subset of ZBTB16$^+$ spermatogonia in adult human testis, and most of them were MKI67$^-$ (Fig. 1J), possibly representing the A_{dark} SSCs also known as the reserve stem cells or 'true SSC' in human testis((29)–(33)). These results suggested that FOXC2 was similarly expressed in the SSCs of adult human and mouse testis and may possess a conserved function.

FOXC2^{+}-SSCs can sufficiently initiate and sustain spermatogenesis

We generated *Foxc2*$^{CRE/+}$;*R26T/G*$^{f/f}$ mice in which FOXC2^{+} cells were specifically labeled with GFP to enable the progeny tracing after tamoxifen treatment (Fig. S3A) (34). Tamoxifen was introduced at 2-month of age, after which the FOXC2-expressing lineage (GFP^{+}) was tracked at d3 (day3), w1 (week1), w2, w4, w6, m4 (month4), m7, and m12 respectively (Fig. 2A). At d3, the tracked cells were both GFP^{+} and FOXC2^{+} (Fig. 2B) and constituted 0.027% of the total testicular cells as indicated by the fluorescence-activated cell sorting (FACS) analysis (Fig. 2C). FACS-sorted GFP^{+} cells were then transplanted into testes of recipient mice pre-treated with busulfan, in parallel to THY1^{+} cells derived from *eGFP*$^{Tg/+}$ mice as control. Two months after transplantation, FOXC2^{+} cells generated 5 times greater number of colonies than the THY1^{+} control (Fig. 6D, E), indicating that the FOXC2^{+} cells possess higher stemness as convinced by stronger transplantable viability.

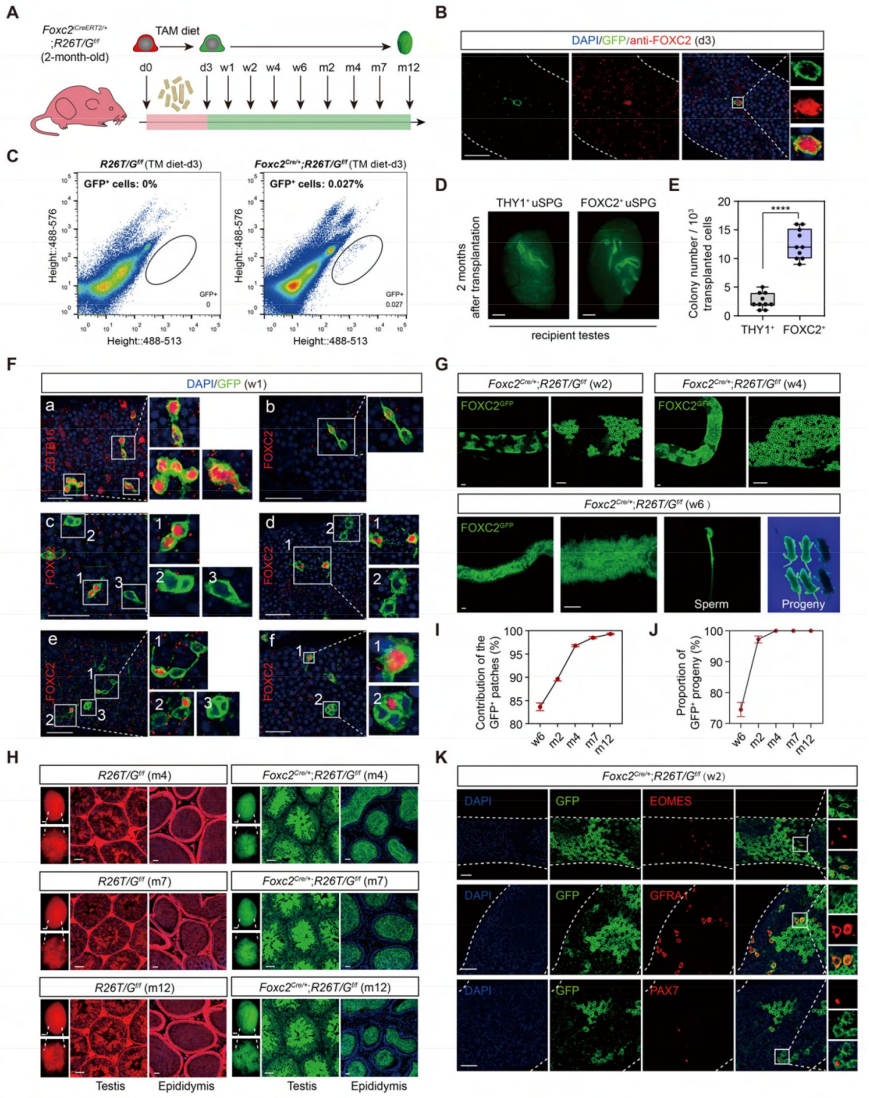

Figure 2.

Lineage tracing and functional validation of FOXC2^{+}-SSCs in *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice.

(**A**) Schematic illustration of the lineage tracing workflow for FOXC2^{+} cells. (**B**) Immunostainings for DAPI (blue) and FOXC2 (red) at day 3 post TAM induction. Scale bar, 50 μm; d, day. (**C**) FACS analysis of GFP^{+} populations derived from *R26T/G$^{f/f}$* or *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice at day 3 post TAM induction. (**D**, **E**) The recipient mice testes (**D**) and colony numbers (**E**) 2 months after transplantation (n=10) of the FACS-sorted GFP^{+} cells from the *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice 3 days after TAM diet and the MACS-sorted THY1^{+} cells from adult mice. Scale bar, 1 mm; values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (****p-value < 0.0001). (**F**) Immunostaining for DAPI (blue), ZBTB16/FOXC2 (red), and GFP (green) at week 1 post TAM induction (scale bar, 50 μm). (**G**) Seminiferous tubules of *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice 2, 4, and 6 weeks post TAM induction. Scale bar, 50 μm. (**H**) Testes (scale bar, 1 mm), seminiferous tubules, and epididymis (scale bar, 50 μm) at month 4, 7, and 12 post TAM induction in *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice. (**I, J**) The GFP^{+} patches (**I**) and progeny (**J**) population dynamics (n=10). Values, mean ± s.e.m. (**K**) Immunostainings for DAPI (blue), EOMES (red), GFRA1 (red), or PAX7 (red) in GFP^{+} population at week 2 post TAM induction. Scale bar, 50 μm.

At w1, all GFP^{+} cells were identified as uSPGs, encompassing A_s, A_{pr}, and $A_{al\text{-}4}$ (Fig. 2F$_a$). Specifically, FOXC2^{+} A_s gave rise to 3 types of A_{pr}, i.e., FOXC2^{+}/FOXC2^{+}, FOXC2^{+}/FOXC2^{-}, and FOXC2^{-}/FOXC2^{-} (Fig. 2F$_{c1, b, c2, d2}$), which then either produced FOXC2^{+} or FOXC2^{-} A_s through symmetric or asymmetric division (Fig. 2F$_{c3, d1, f1}$), or developed into A_{al} with no more than one FOXC2^{+} cell in the chains (Fig. 2F$_{e, f2}$). These results confirmed that FOXC2^{+} cells were capable of self-renewal to sustain the population as well as replenishing the uSPG pool by producing downstream progenies, thereby serving as primitive SSCs. In the following 2-6 weeks, GFP^{+} colonies further expanded and produced GFP^{+} sperms in the epididymis, from which healthy GFP^{+} offspring were given birth by C57 female recipients (Fig. 2G). The GFP^{+} colonies constituted 83.67%, 90.48%, 96.78%, 98.55%, and 99.31% of the total length of the seminiferous tubules at w6, m2, m4, m7, and m12 respectively (Fig. 2H, I). All offspring were GFP^{+} from m4 onwards (Fig. 2J). Additionally, the EOMES^{+}, GFRA1^{+} and PAX7^{+} cells were all GFP^{+} at w2, further confirming these progenies were derived from the FOXC2^{+} cells (Fig. 2K).

Overall, FOXC2^{+}-SSCs can produce all subtypes of uSPG, thus initiating spermatogenesis in adult mice.

Specific ablation of the FOXC2^{+}-SSC results in depletion of the uSPG pool

We then prepared *Foxc2*$^{Cre/+}$;*R26*$^{DTA/+}$ mice to investigate the physiological requirement of FOXC2^{+}-SSCs in spermatogenesis (34). FOXC2^{+} population in 2-month-old mice was specifically ablated with tamoxifen-induced diphtheria toxin (DTA). The testes of these mice were examined at day3, day7, and day14 post tamoxifen induction (Fig. 3A). Gradual loss of weight in testes coincided with the reduction in the size of testes in all the mice while body weight was maintained (Fig. 3B, C). Specifically, at d3, there were no detectable FOXC2^{+} cells in addition to the decrease in the number of GFRA1^{+}, LIN28A^{+} (35) and ZBTB16^{+} uSPG at the basement membrane of seminiferous tubules; at d14, all GFRA1^{+}, LIN28A^{+} and ZBTB16^{+} uSPG disappeared while vacuoles formed at the basement membrane with remaining spermatocytes and spermatids in the seminiferous lumen (Fig. 3D-F, Fig. S3B). Meanwhile, the expression of DDX4 (36) and DAZL (37) as germ cell markers was significantly reduced along with nearly undetectable expression of uSPG markers such as ZBTB16, LIN28A, GFRA1, RET, and NEUROG3 (28) (Fig. 3G). These results indicate an uSPG exhaustion as the result of the FOXC2^{+}-SSCs ablation, therefore supporting the critical role in spermatogenesis played by FOXC2^{+} population.

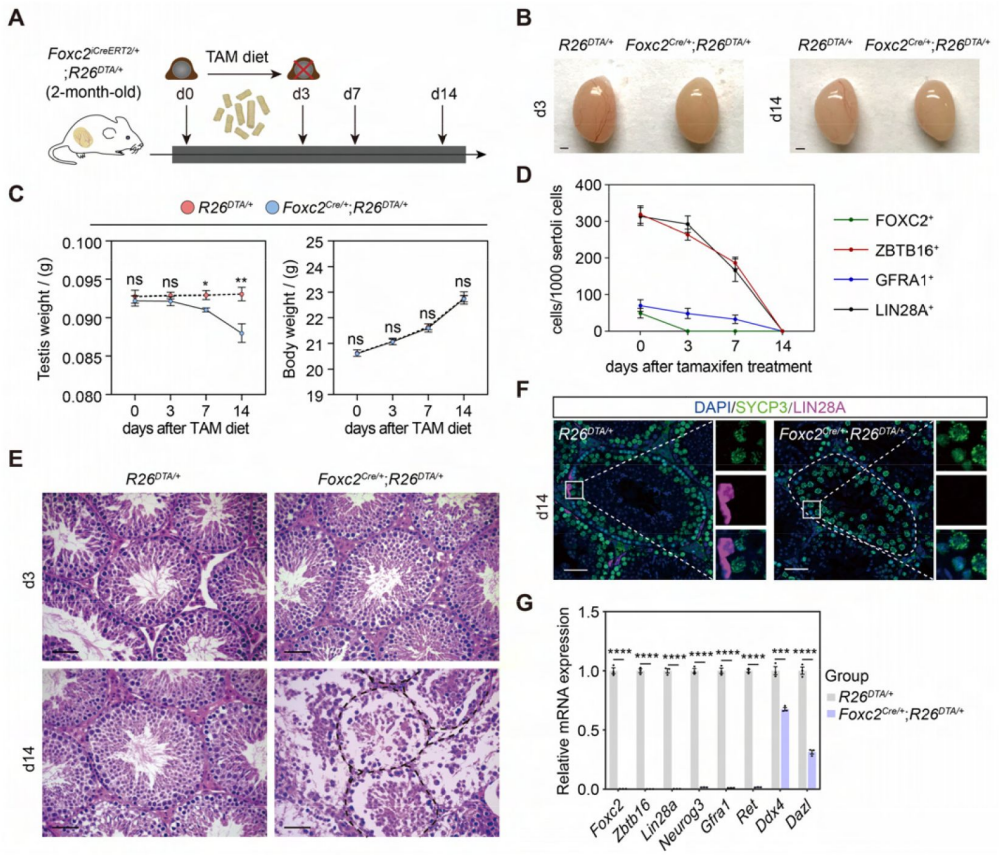

Figure 3.

Specific ablation of FOXC2$^+$-SSCs and phenotypic validation in *Foxc2$^{Cre/+}$;R26$^{DTA/+}$* mice.

(**A**) Schematic illustration of the lineage tracing workflow for FOXC2$^+$ cells. (**B-D**) Phenotypic validation of the *R26$^{DTA/+}$* and *Foxc2$^{Cre/+}$;R26$^{DTA/+}$* mice (n=5) for testes size (**B**), testis weight and body weight (**C**), and HE-staining of the testes (**D**). Scale bars in (**B**), 1 mm; in (**D**), 50 μm; d, day; values were mean ± s.e.m.; p-values were obtained using two-tailed t-tests (ns > 0.05, *p-value < 0.05, **p-value < 0.01). (**E**) ZBTB16$^+$, GFRA1$^+$, LIN28A$^+$, and FOXC2$^+$ SPG populations dynamics. Values, mean ± s.e.m. (n=10); p-values were obtained using one-way ANOVA followed by Tukey test (ns > 0.05, *p-value < 0.05, **p-value < 0.01, ****p-value < 0.0001). (**F**) Immunostainings for DAPI (blue), SYCP3 (green), and LIN28A (magenta) at day 14 post TAM induction. d, day; scale bar, 50 μm. (**G**) Quantitative RT-PCR analysis of SPG markers expression in the testes of the *R26$^{DTA/+}$* and *Foxc2$^{Cre/+}$;R26$^{DTA/+}$* mice (n=3). Values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (***p-value < 0.001, ****p-value < 0.0001).

FOXC2$^+$-SSCs are resilient to genotoxin and indispensable for germline regeneration

Next, we examined the regenerative viability of FOXC2$^+$-SSCs. At d20 post busulfan treatment (20mg/kg), FOXC2$^+$ cells constituted the majority of uSPGs (Fig. 4A). Following a sharp decrease in cell number in the first five days, ZBTB16$^+$ and GFRA1$^+$ cells began to recover from d25 while the number of FOXC2$^+$ cells remained stable (Fig. 4B), indicating that this population is insensitive to busulfan. We then checked changes in the proportion of MKI67$^+$ cells, active in proliferation, in FOXC2$^+$ population after busulfan treatment (Fig. 4C, D). At d30, the MKI67$^+$ proportion rose to 15.92%, indicating a higher level of proliferation, albeit the total cell number stayed static (Fig. 4B, D), thereby becoming the driving force in restoring spermatogenesis. Up to d120, the MKI67$^+$ proportion had settled gradually back to the pre-treatment level, accompanied by the full recovery of spermatogenesis (Fig. 4D). Further details of this process were revealed during lineage tracing (Fig. 4E). Three days after tamoxifen induction, the 2-month-old *Foxc2$^{CRE/+}$;R26T/G$^{f/f}$* mice were treated with busulfan. Consistent with the results above, at d20, the survived uSPG were predominantly GFP$^+$ (Fig. 4F). Over 68.5% of the total length of the seminiferous tubules were GFP$^+$ at m2,

and this proportion rose to 95.43%, 98.41%, and 99.27% at m4, m7, and m12 respectively (Fig. 4G, H), which was comparable to the proportion by tamoxifen induction alone (Fig. 2I). From m4 onwards, nearly all germ cells, spermatids, and their offspring were GFP$^+$ (Fig. 4G, I). Together, these results confirmed that FOXC2$^+$-SSCs are indispensable for germline regeneration that is central to spermatogenesis recovery from interruptions.

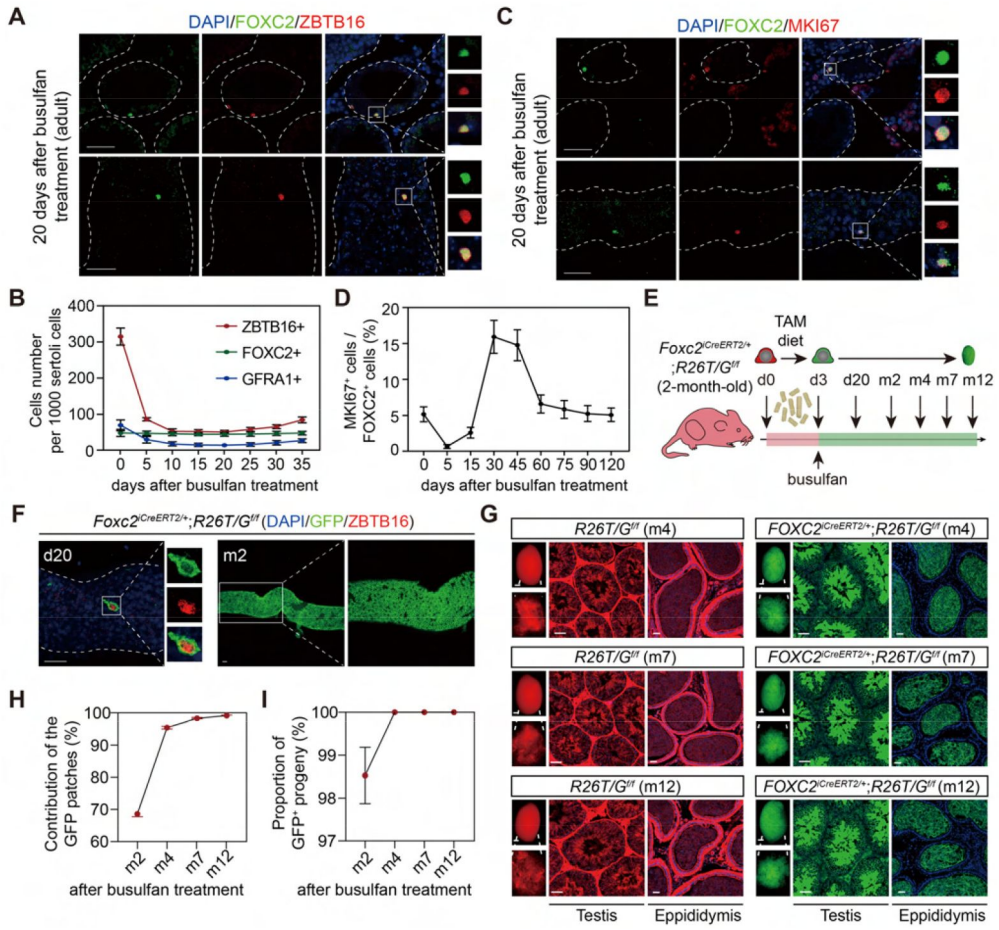

Figure 4.

FOXC2$^+$-SSCs are critical for germline regeneration.

(**A**) Co-immunostaining of FOXC2 (green) with ZBTB16 (red) in seminiferous tubules of the adult testes at day 20 post busulfan treatment. Scale bar, 50 μm. (**B**) ZBTB16$^+$, GFRA1$^+$, and FOXC2$^+$ population dynamics after busulfan treatment (20 mg/kg, n=10). (**C**) Co-immunostaining of FOXC2 (green) with MKI67 (red) in seminiferous tubules of the adult testes at day 20 post busulfan treatment. Scale bar, 50 μm. (**D**) MKI67$^+$FOXC2$^+$ proportions in relation to the whole FOXC2$^+$ population at different time points after busulfan treatment (n=4). (**E**) Schematic illustration for lineage tracing of FOXC2$^+$ cell after busulfan treatment. (**F**) Lineage tracing of the GFP$^+$ cells at day 20 and month 2 after busulfan treatment (scale bar, 50 μm). (**G**) The testes (scale bar, 1 mm), seminiferous tubules, and epididymis (scale bar, 50 μm) at month 4, 7, and 12 post TAM induction and busulfan injection. m, month. (**H**, **I**) The proportion dynamics of GFP patches (**H**) and GFP$^+$ progenies (**I**). Values, mean ± s.e.m. (n=10). w, week; m, month.

FOXC2 is essential for SSCs maintenance in adult mice

We then focused on dissecting FOXC2's role in the SSCs maintenance using *Foxc2$^{f/-}$;Ddx4-cre* mice (38) (Fig. 5A). No significant difference was observed in the expressions of various uSPG markers, including ZBTB16 and LIN28A, between *Foxc2$^{f/-}$; Ddx4-cre* and *Foxc2$^{f/+}$* mice at the age of 1 week (Fig. S4B). However, adult *Foxc2$^{f/-}$;Ddx4-cre* mice displayed clear testis weight loss without significant body weight loss (Fig. 5B, C). Moreover, in these mice, we observed severe degeneration of seminiferous tubules, reduced number of spermatids in the epididymis, and decreased size of the uSPG population with age (Fig. 5D–G) but without apparent signs of apoptosis (Fig. S5B). The 6-month-old *Foxc2$^{f/-}$;Ddx4-* cre mice were infertile, in which over 95% seminiferous tubules were Sertoli-only with hardly detectable

expressions of DAZL, DDX4, LIN28A, and ZBTB16 (Fig. 5D–F, H). Therefore, FOXC2 is essential for maintaining the SSCs homeostasis and normal spermatogenesis in adult mice.

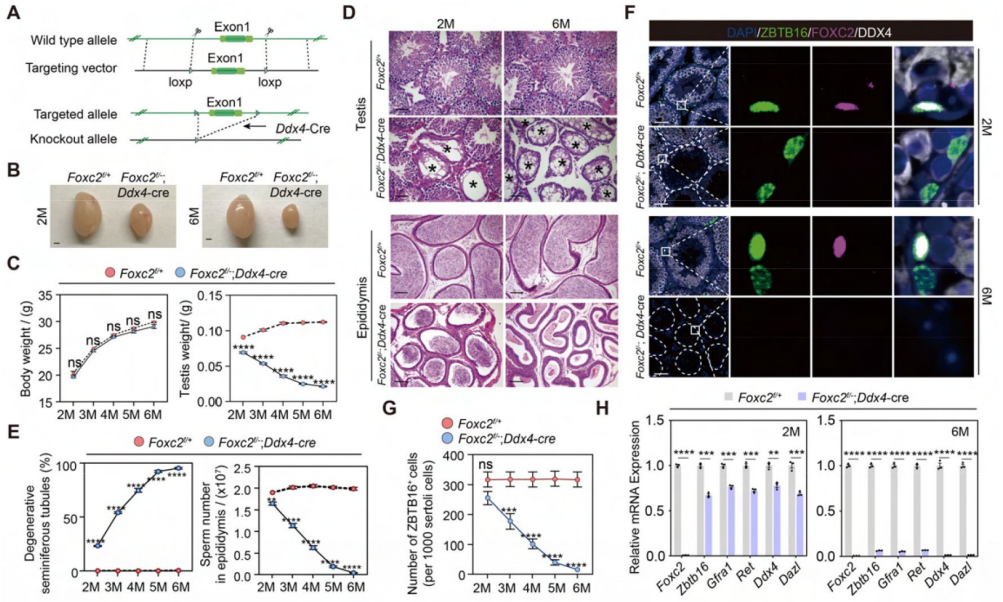

Figure 5.

Spermatogenesis exhaustion in the adult *Foxc2$^{f/-}$;Ddx4-cre* mice.

(**A**) Construction of the *Foxc2$^{f/-}$;Ddx4-cre* mice. (**B**) The testes size of the *Foxc2$^{f/-}$;Ddx4-cre* mice. Scale bar, 1 mm; M, month. (**C**) Body weight and testis weight of the *Foxc2$^{f/-}$;Ddx4-cre* mice at different age (n=5). M, month; values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (ns > 0.05, ****p-value < 0.0001). (**D**) HE-staining of the testis and epididymis. Scale bar, 50 μm; M, month. (**E**) Estimation of degenerative tubules and sperm counts in cauda epididymis of the *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$;Ddx4-cre* mice with age (n=5). Values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (**p-value < 0.01, ****p-value < 0.0001). (**F**) Immunostainings for DAPI (blue), ZBTB16 (green), FOXC2 (magenta), and DDX4 (white) in the seminiferous tubules of the *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$; Ddx4-cre* mice. Scale bar, 50 μm. (**G**) Estimation of ZBTB16^{+} uSPG number in the *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$;Ddx4-cre* mice with age (n=5). Values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (ns > 0.05, ***p-value < 0.001, ****p-value < 0.0001). (**H**) Quantitative RT-PCR analysis of the uSPG and germ cell markers expressed in the testis of the *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$; Ddx4-cre* mice (n=3). M, month; values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (**p-value < 0.01, ***p-value < 0.001, ****p-value < 0.0001).

FOXC2 maintains the SSCs homeostasis via negative regulation of cell cycle

We collected THY1^{+} uSPGs from 4-month-old *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$; Ddx4-cre* mice and compared their transcriptome signatures revealed from scRNA-seq (Fig. 6A). The pseudotime analysis identified Cluster1, which represented the FOXC2-expressing SSCs in *Foxc2$^{f/+}$* mice corresponding to the FOXC2-deleting SSCs in the *Foxc2$^{f/-}$; Ddx4-cre* mice, was specifically assigned to the extremely early stage of the development trajectory in respective samples, which was validated by the expression of corresponding markers (Fig. 6B, Fig. S5A, B). Aggregated analysis of the overall uSPG populations showed that cells derived from *Foxc2$^{f/-}$; Ddx4-cre* mice were specifically associated with the late stage of the development trajectory, as opposed to *Foxc2$^{f/+}$* mice where nearly all the cells derived were concentrated at the early stage of development (Fig. 6C, Fig. S5C). This implies that the loss of Foxc2 prompts the SSCs to progress into a more differentiated stage with defection in maintaining the primitive identity of SSCs. Further analysis of the cells in Cluster1 revealed two distinct subclusters, i.e., Subclusters0 and Subclusters1 (Fig. S6A). Formed primarily by the Cluster1 cells derived from *Foxc2$^{f/+}$* mice, Subclusters0 was featured by stemness markers, while Subcluster1,

representing the majority of Cluster1 cells from *Foxc2*$^{f/-}$*; Ddx4-cre* mice, was featured by progenitor markers (Fig. S6B, C). Consistently, pseudotime analysis showed that Cluster1 cells from *Foxc2*$^{f/+}$ mice projected a forward stage of the developmental trajectory indicated by stemness markers, whereas Cluster1 cells from *Foxc2*$^{f/-}$*; Ddx4-cre* mice were associated with a later stage of the developmental trajectory (Fig.6D, Fig. S6D, E). More specifically, less number of cells were found at the starting state1 in Cluster1 from *Foxc2*$^{f/-}$*; Ddx4-cre* mice than in *Foxc2*$^{f/+}$ mice, with rather more cells in the developmental progression (from state1 to state5), especially at the advanced state5 (Fig. 6E). Thus, FOXC2 deletion caused defective SSCs maintenance and committed the primitive SSCs to a differentiation destiny. Further, there were 932 genes down-regulated in Cluster1 cells derived from *Foxc2*$^{f/-}$*; Ddx4-cre* mice in comparison to *Foxc2*$^{f/+}$ mice (Fig. 6F, Supplemental Table S2), which were functionally associated with both stem cell population maintenance and mitotic cell cycle (Fig. 6G). Consistently, the GSEA analysis revealed a more progressive cell cycle in Cluster1 upon *Foxc2*-knockout (Fig. 6H), confirming the role of FOXC2 in regulating the cell cycle of the primitive SSCs.

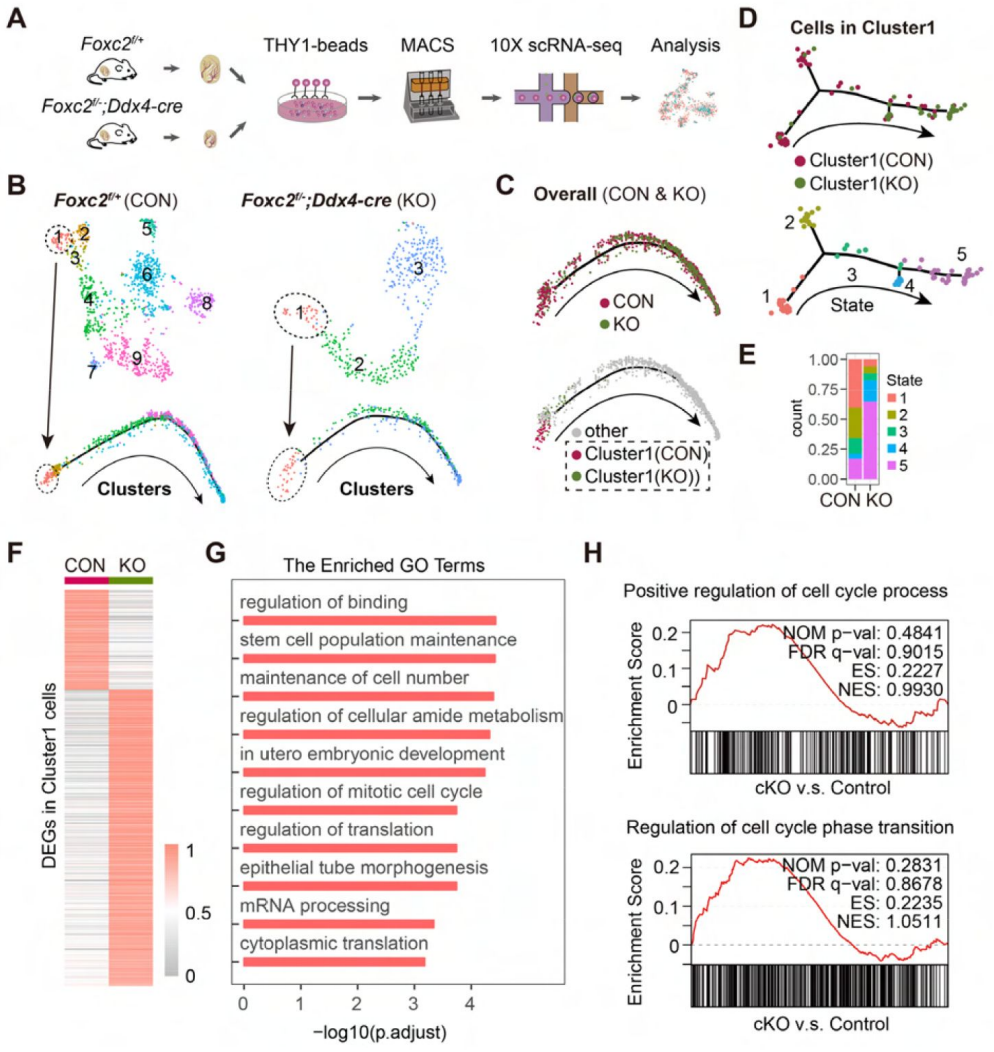

Figure 6.

scRNA-seq analysis of THY1$^+$ uSPG in *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*;Ddx4-cre* mice.

(**A**) Schematic illustration of the scRNA-seq workflow. (**B**) t-SNE plot and developmental trajectory of uSPG from *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*;Ddx4-cre* mice respectively, colored by cluster. (**C**) Developmental trajectories of uSPG from *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*;Ddx4-cre* mice, colored by sample or derivation. (**D**) Developmental trajectories of the cells in Cluster1 from *Foxc2*$^{f/+}$ (CON) and *Foxc2*$^{f/-}$*;Ddx4-cre* (KO) mice, colored by derivation or developmental state. (**E**) The Cluster1 cells proportion of each state in CON and KO mice. (**F**) Heatmap showing the DEGs in the Cluster1 cells from the *Foxc2*$^{f/-}$*;Ddx4-* cre mice compared with the *Foxc2*$^{f/+}$ mice. (**G**) Top GO terms enrichment by the down-regulated DEGs in KO mice. (**H**) Gene set enrichment analysis (GSEA) of the Cluster1 cells (*Foxc2*$^{f/-}$*;Ddx4-* cre v.s. *Foxc2*$^{f/+}$ mice). NOM, nominal; FDR, false discovery rate; ES, enrichment score; NES, normalized enrichment score.

We then performed Cleavage Under Targets and Tagmentation (CUT&Tag) sequencing to explore the underlying mechanism (39; 40), for which GFP$^+$ SSCs from *Foxc2C$^{RE/+}$;R26T/G$^{f/f}$* mice 3 days after tamoxifen induction, representing the FOXC2$^+$-SSCs, were isolated for CUT&Tag sequencing (Fig. 7A). Specific peaks enriched in the promoter region of 3629 genes (Fig. 7B, C; Supplemental Table S2) showed functional enrichment in biological processes such as DNA repair and mitotic cell cycle regulation (Fig. 7D). By overlapping with the 932 genes down-regulated in Cluster1 cells from *Foxc2$^{f/-}$; Ddx4-cre* mice, we obtained 306 genes as the candidates subjective to the regulation by FOXC2 (Fig. 7E; Supplemental Table S2). Further, GO enrichment analysis of these genes highlighted a distinctive functional cluster (11 genes) focusing on the negative regulation of cell cycle (Fig. 7F; Supplemental Table S3) ((41)–(50)). More specifically, significant peaks enrichment at the promoter region were observed for these candidate genes (Fig. 7G). Meanwhile, as predicted using the JASPAR Scan function (binding potential >0.8), there showed strong binding potential of FOXC2 towards these candidate genes (Fig. 7I) via the binding motif of FOXC2 (Fig. 7H), which was further confirmed by the results from the CUT&Tag qPCR (Fig. 7J). Overall results implied that FOXC2 may function as a gatekeeper that ensures the quiescent state of the primitive SSCs by impeding cell cycle progression.

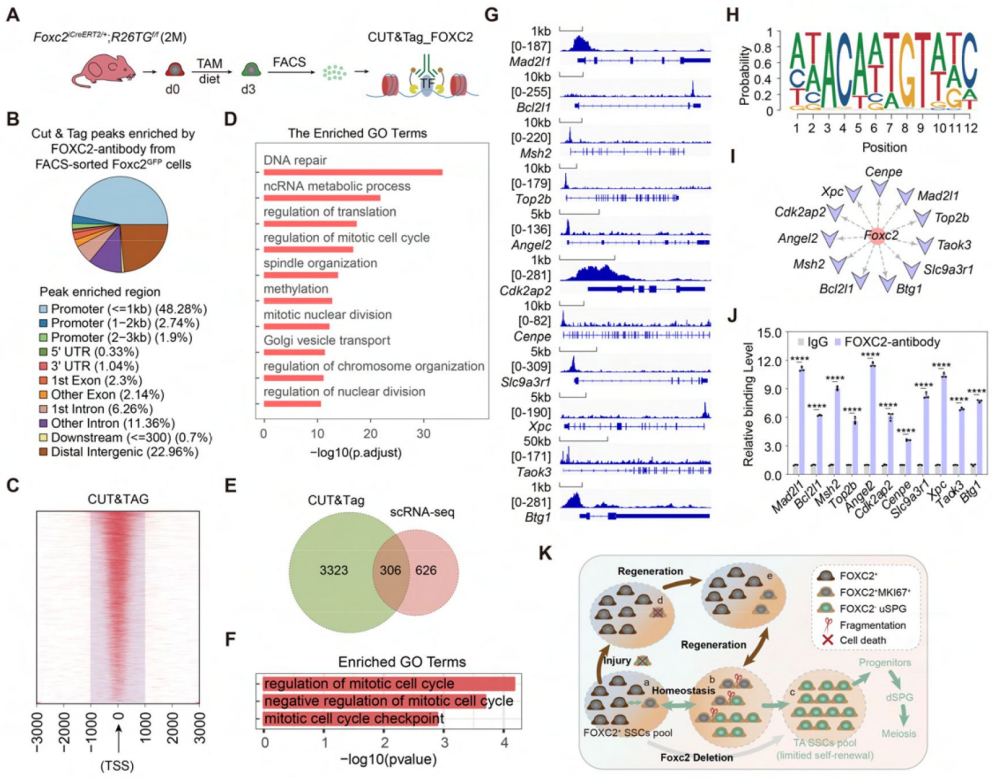

Figure 7.

FOXC2 is essential for sustaining the primitive SSCs via regulating cell cycle.

(A) Workflow schematic illustration of the CUT&Tag_FOXC2 analysis on the FACS-sorted FOXC2$^+$ cells. (B) Pie chart for CUT&Tag_FOXC2 peaks genome distribution. (C) Profiling of CUT&Tag_FOXC2 peaks in proximity to transcriptional starting site (TSS). The distance to TSS within 1000 was highlighted in the purple box. (D) Top GO terms enrichment by genes annotated by CUT&Tag_FOXC2 peaks. (E) Venn diagram of FOXC2 target genes defined by overlapping the CUT&Tag sequencing and scRNA-seq datasets. (F) GO terms enrichment by the FOXC2 target genes related to cell cycle regulation. (G) Chromatin landscapes of CUT&Tag_FOXC2 peaks of the candidates associated with negative cell cycle regulation. (H) The DNA-binding motif for FOXC2 (predicted with HOMER). (I) The cell cycle-related candidates possessing high binding potential (>0.8, predicted with JASPAR SCAN). (J) CUT&Tag-qPCR validation of the cell cycle arrest regulatory genes. (n=3). Values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (****p-value < 0.0001). (K) The model for the maintenance of the FOXC2+ SSCs subpopulation in adult testis.

Discussion

In this work, a comprehensive analysis of uSPG populations with scRNA-seq and the following lineage tracing study by whole-mount immunofluorescence assay led to the identification of FOXC2-expressing SSCs as an important and primitive SSCs subpopulation in adult mice. Further investigation through functionality analysis confirmed FOXC2 is essential for SSCs self-renewal and stemness, thereby is required for maintaining the SSCs population that is critical for continuous spermatogenesis. Importantly, our data demonstrated that the colonies formed by FOXC2$^+$ cells constituted nearly the total length of the seminiferous tubules (99.31 %), implying that the FOXC2$^+$- SSCs can support the complete spermatogenesis in adult mice.

GFRA1$^+$ A_{pr} and A_{al} cells were found to break randomly and a portion of them can return to the stem cell state (5). Interestingly, our findings showed FOXC2 appeared in one of the A_{pr} or A_{al} cells at times, therefore raising a possibility that the subset of GFRA1$^+$ cells that return to stem cell state after intercellular bridge break, maybe FOXC2$^+$ due to different cell cycle state. If so, based on both findings, GFRA1$^+$FOXC2$^+$ could represent a quiescent state whereas GFRA1$^+$FOXC2$^-$ is proliferate active, which certainly requires further validation possibly through multiple lineage tracing and live imaging.

We observed that the FOXC2$^+$-SSCs were almost all in a non-proliferative state (~94.9%), and further revealed that FOXC2 functioned in the negative regulation of cell cycle progression, thus confirming that FOXC2-expressing SSCs are quiescent SSCs population in adult mice. The finding that FOXC2 inhibited cell cycle and differentiation of SSCs in testis is consistent with that reported in other tissues (51; 52). In general, the quiescent state is a protective mechanism for stem cell storage and prevents stem cells from damage or depletion under genotoxic stresses ((1), (53)–(55)). In our work, after the busulfan treatment, the quantity of FOXC2$^+$ cells remained stable and the survived uSPGs were predominantly FOXC2$^+$, indicating its insensitivity to cytotoxic agents. However, the proportion of MKI67$^+$FOXC2$^+$ cells increased by 15.92% after 30 days of the busulfan treatment and decreased back to the pre-treatment level (5.08%) at 120 days, implying that the quiescent FOXC2$^+$ cells were able to transform into the proliferative FOXC2$^+$ cells to replenish the SSCs pool to maintain the SSCs homeostasis and normal spermatogenesis. We further confirmed by lineage tracing analysis that FOXC2-expressing cells were the only remaining SSCs population and were responsible for germline regeneration after the busulfan treatment, indicating that FOXC2$^+$-SSCs represent a functionally important stem cell population with regenerative ability. In the future, more insights into the unique regulation of SSCs can be drawn from studying and comparing the transition between the quiescent and proliferative states in FOXC2$^+$ and other SSCs subpopulations.

According to our findings, we proposed a model for the maintenance of the FOXC2+ SSCs subpopulation (Fig. 7K). Under physiological conditions, FOXC2$^+$ A_s cells (including FOXC2$^+$GFRA1$^+$, FOXC2$^+$EOMES$^+$ cells, etc.) constitute the primitive population of SSCs, of which only a small proportion (~5.1%) cells are proliferative while the majority remains quiescent (Fig. 7Ka). This primitive population can divide symmetrically or asymmetrically into different A_{pr} and A_{al} (Fig. 7Kb). Then FOXC2$^+$ cells (Fig. 7Kb) may break from the syncytial and return to A_s state (Fig. 7Ka) to maintain the stable number of the primitive SSCs. FOXC2$^-$ progenies, derived from the FOXC2$^+$ primitive population, form a transit amplification (TA) SSCs pool (Fig. 7Kc) to support spermatogenesis. However, it requires continuous supply from the FOXC2$^+$ population and is subject to exhaustion when the supply is disrupted. In the context of regeneration conditions, the FOXC2$^+$MKI67$^-$ cells can survive and set out the recovery process (Fig. 7Kd). At the early stage, increasing proportions of

MKI67^{+}/MKI67^{-} ratio returns to the physiological level in FOXC2^{+} population (Fig. 7Ka), leaving the total number of FOXC2^{+} cells stable therefore maintaining the SSCs homeostasis. However, it is necessary to perform more investigation to further improve and modify this model to gain a complete understanding of the connections between different primitive SSCs subpopulations via lineage tracing assays in the testes of adult mice.

Based on our observation, FOXC2 seems nonessential for the transformation from gonocytes to SSCs in infant mice, in contrast to its requirement for adult spermatogenesis. A recent study showed that FOXC2 was present in a fraction of A8 and A16 cells in the postnatal mouse testis (<5 weeks), however, this FOXC2^{+} subpopulation appeared more active in proliferation than the adult counterpart (56). Such differential functionality might reflect the difference in the physical nature of spermatogenesis between developmental stages. For example, the maturity of spermatogenesis is still under development during the juvenile period with a focus on expanding the SSCs pool. Therefore, it would be interesting to explore differences in individual functional contexts as well as the underlying regulatory mechanisms. Meanwhile, FOXC2, highly conserved between mice and humans with 94% identity in amino acid sequence (57), is also expressed in a subset of human adult SSCs, raising the possibility of an evolutionarily conserved mechanism governing SSCs homeostasis in humans. Further work following this direction might be of great clinical significance specifically to patients who suffer from infertility. Moreover, the developmental correlation between FOXC2^{+}-SSCs and other SSCs subpopulations proposed previously should be revealed via biological methods such as multiple lineage tracing and live imaging. Collectively, our work here provides new insights into the investigation of adult SSCs and serves as a reference for studying the homeostasis and regeneration of other stem-cell systems.

Materials and Methods

Data Availability

All data are available in the main text or supplementary materials. The scRNA-seq and CUT&Tag sequencing data have been uploaded to the GEO with accession codes GSE183163, GSE180729, and GSE180926. All of the R packages were available online and the code was used according to respective R packages documentation as described in the Methods. The MSigDB (v.7.0) used in this study is available at https://www.gsea-msigdb.org/gsea/msigdb.

Additional Experimental Procedures

The procedures for mice, magnetic-activated cell sorting (MACS), single-cell RNA-seq, single-cell RNA-seq data processing, CUT & TAG sequencing and analysis, enrichment analyses, transplantation assay, fluorescence-activated cell sorting (FACS), immunofluorescence, RNA isolation and quantitative RT-PCR analysis, tamoxifen inducible, analyses of cell density, sperm counts, histology, evaluation of degenerating tubules, and statistical analysis are presented in the Supplemental Materials and Methods.

Acknowledgements

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Supplemental Figures

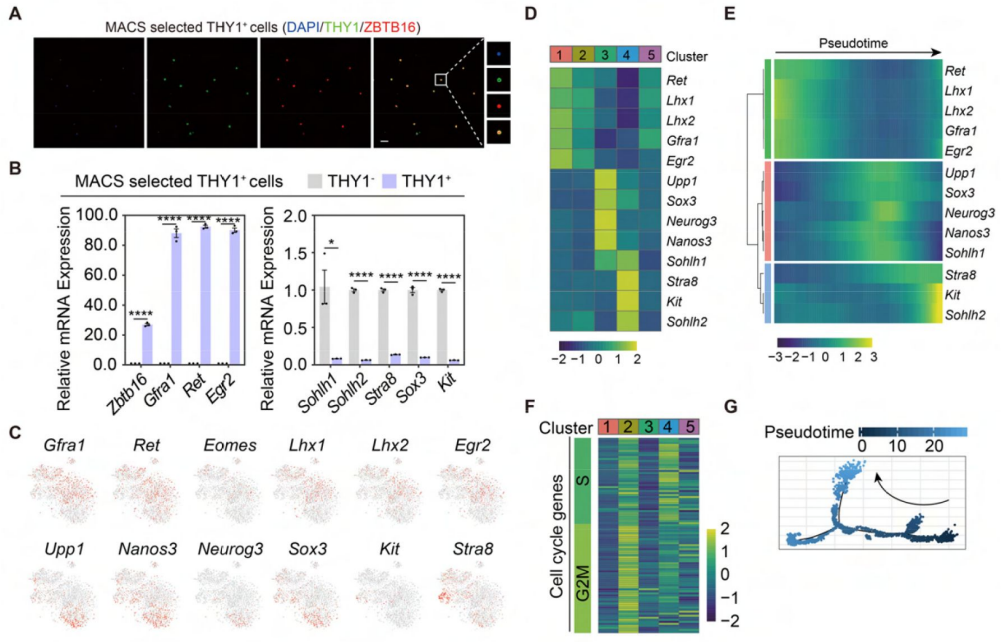

Figure S1.

Validation and characterization of the MACS-sorted THY1$^+$ uSPG from wild-type adult C57 mice.

(**A**) Immunostainings of DAPI (blue), THY1 (green), and ZBTB16 (red) in the MACS-sorted THY1$^+$ cells (n=5). Scale bar, 50 μm. (**B**) Quantitative RT-PCR analysis of uSPG and dSPG markers expressed in the MACS-sorted THY1$^+$ cells (n=3). Values, mean ± s.e.m.; p-values were obtained using two-tailed t-tests (ns > 0.05, *p-value < 0.05, **p-value < 0.01, ***p-value < 0.001, ****p-value < 0.0001). (**C**) Feature plots showing the expression pattern of classic SPG markers (stemness and differentiation). (**D**) Heatmap showing the expression pattern of markers for SPG in different clusters. (**E**) Expression pattern dynamics of the SPG markers with pseudotime progression. (**F**) Heatmap showing the expression pattern of markers for cell cycle phase in different clusters. (**G**) The developmental trajectory of the overall SPG, colored by pseudotime.

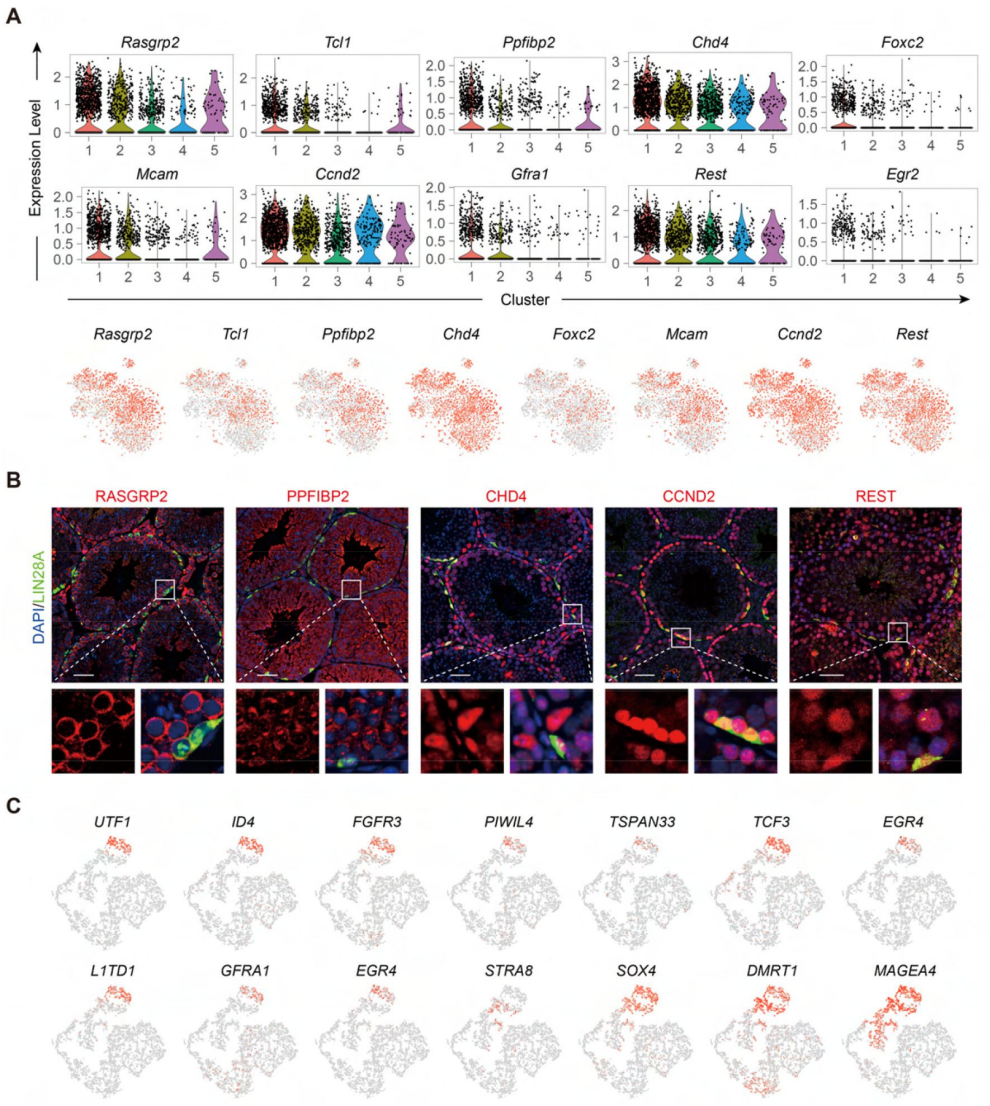

Figure S2.

Expression of top10 DEGs of Cluster1 in Figure 1B and classic SSC and SPG markers in adult human germ cells.

(**A**) Feature plots and violin plots of the Top10 DEGs of Cluster1. (**B**) Immunostainings for LIN28A (red), DAPI (blue), and newly-found markers (green) in testicular paraffin sections from adult mice. Scale bar, 50 μm. (**C**) Feature plots showing the expression pattern of classic SSCs and SPG markers in adult human germ cells.

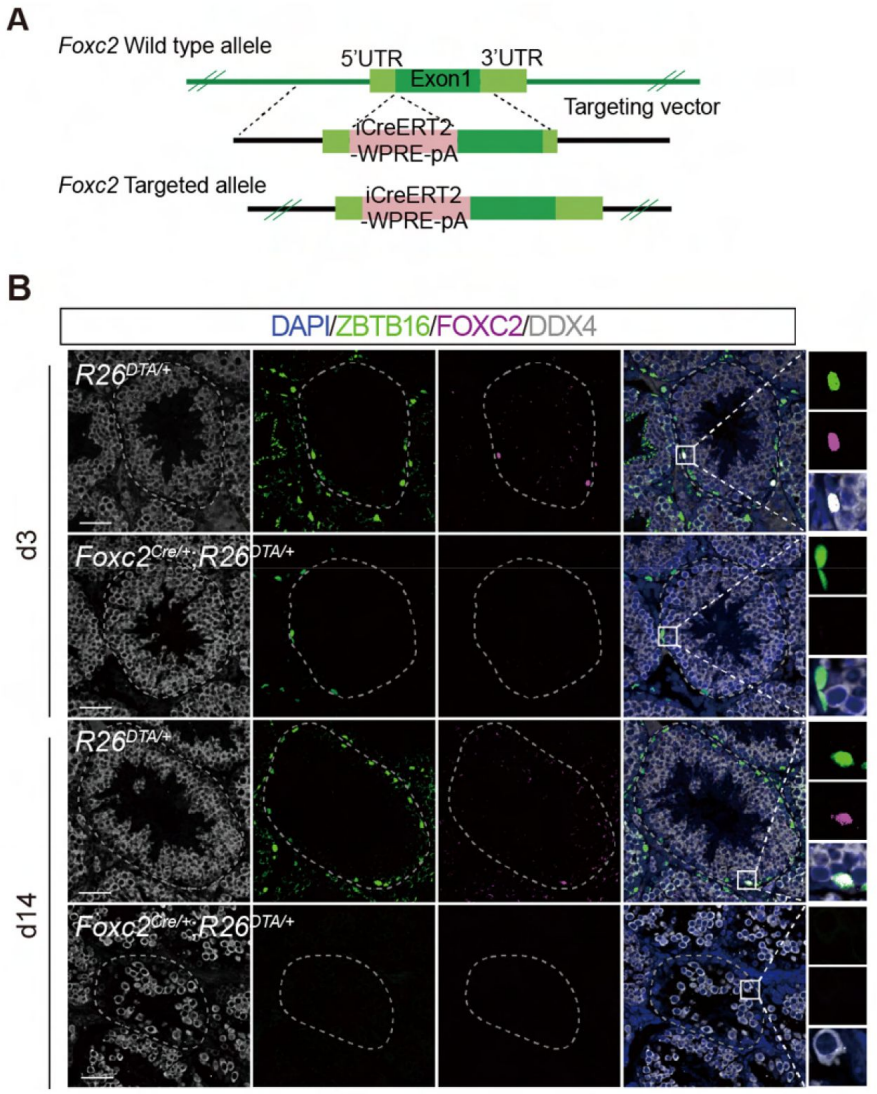

Figure S3.

Construction of the *Foxc2*^{iCreERt2} mice and depletion of uSPG pool in *Foxc2*$^{Cre/+}$;*R26*$^{DTA/+}$ mice 14 days after specific ablation of FOXC2^{+}-SSCs.

(**A**) Construction of the *Foxc2*^{iCreERt2} mice. (**B**) Immunostainings for DAPI (blue), DDX4 (white), ZBTB16 (green), and FOXC2 (magenta) at day 3 and day 14 post TAM induction in *Foxc2C*$^{re/+}$;*R26*$^{DTA/+}$ mice (scale bar, 50 μm).

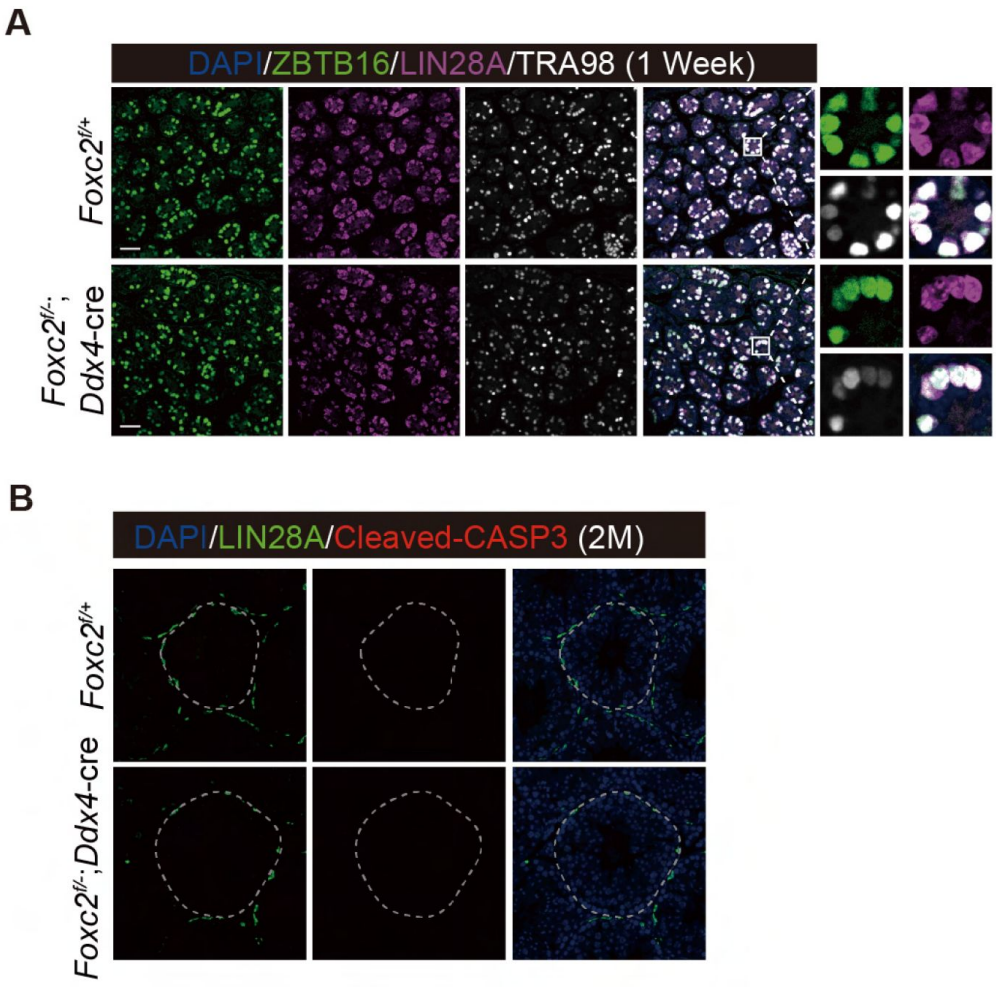

Figure S4.

Phenotypic validation of the *Foxc2$^{f/-}$;Ddx4-cre* mice.

(**A**) Immunostainings for DAPI (blue), ZBTB16 (green), LIN28A (magenta), and TRA98 (white) in seminiferous tubules of 1-week-old *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$;Ddx4-cre* mice. Scale bar, 50 μm. (**B**) Immunostainings for DAPI (blue), LIN28A (green), and Cleaved-CASP3 (red) in seminiferous of the *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$; Ddx4-cre* mice (2-month-old). M, month; scale bar, 50 μm.

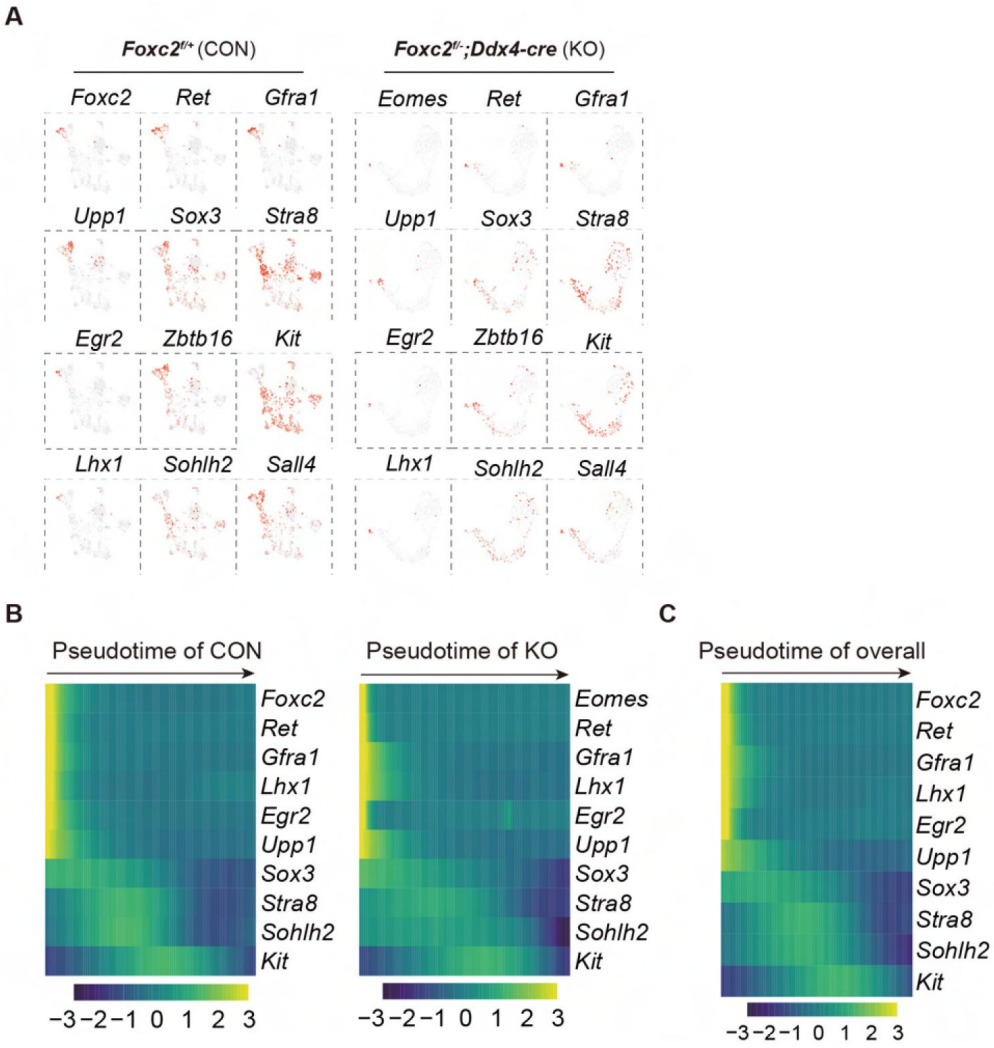

Figure S5.

scRNA-seq analysis of THY1^{+} uSPG in adult *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$;Ddx4-cre* mice.

(**A**) Feature plots of classic SPG markers for uSPG in adult *Foxc2$^{f/+}$* or *Foxc2$^{f/-}$;Ddx4-cre* mice. (**B**) Expression dynamics of SPG markers with pseudotime progression for uSPG from *Foxc2$^{f/+}$* or *Foxc2$^{f/-}$;Ddx4-cre* mice respectively. (**C**) Expression dynamics of SPG markers with pseudotime progression for overall uSPG from *Foxc2$^{f/+}$* and *Foxc2$^{f/-}$;Ddx4-cre* mice.

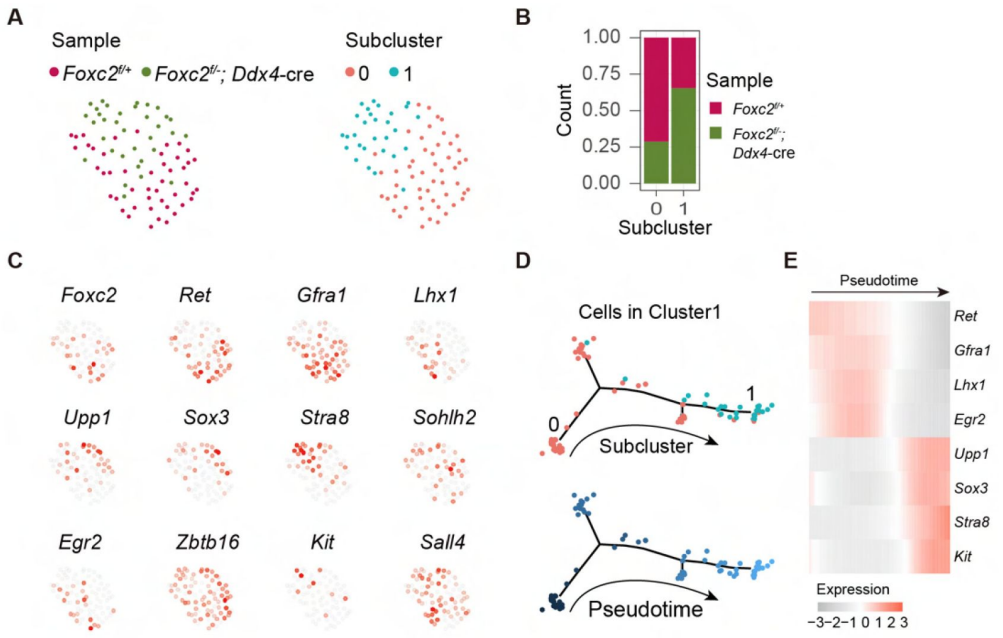

Figure S6.

Re-cluster and developmental trajectory analysis of cells in Cluster1 derived from adult *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*;Ddx4-cre* mice.

(A) The t-SNE plot of the Cluster1 cells aggregated from the *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*;Ddx4-cre* mice colored by sample or subcluster. (B) The cell proportion of each sample in each subcluster. (C) Feature plots of SPG markers expression. (D) Developmental trajectory of the aggregated Cluster1 cells colored by subcluster or pseudotime. (E) Expression dynamics of SPG markers with pseudotime progression.

Supplemental Tables (separate files)

Supplemental Table S1. List of the top30 differentially expressed genes of different clusters.

Supplemental Table S2. List of the differentially expressed genes found by CUT&Tag sequencing and scRNA-seq respectively and their respective enriched Gene Ontology terms.

Supplemental Table S3. List of the Gene Ontology terms of the 306 crossed candidates.

Supplemental Table S4. Primers and antibodies used in this study.

Supplemental Materials and Methods

Mice

Animal experiments were approved by the Committee on Animal Care of the Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences and Peking Union Medical College. The 8-week-old C57BL/6J wild-type mice were used for magnetic-activated cell sorting. The Rosa26mTmG^{flox} mice (stock no. 007676), *Ddx4*-Cre mice (stock no. 000692) and EGFP$^{Tg/+}$ mice (stock no. 021930) were bought from the Jackson Laboratory. The *Foxc2*^{iCreERt2} mice and the *Foxc2*$^{flox/flox}$ (*Foxc2*$^{f/f}$) mice were constructed and bought from the Biocytogen. The Rosa-eGFP-DTA (R26$^{DTA/+}$) mice were bought from GemPharmatech. All mice were housed and bred under specific pathogen-free conditions (temperature: 22-26°C, humidity: 40-55%, 12-h light/dark cycle) in the animal facility at the Institute of Basic Medical Sciences. DNA was isolated from the tails, and the genotypes of the mice were checked using

PCR with specific primers (Supplemental Table S4). All mice were randomly assigned to experiments and no statistical methods were used to predetermine sample size. The person performing the experiments did not know the sample identity until after data analysis. No data were excluded from analyses and the data displayed included a minimum of three independent experiments and a minimum of three biological replicates for each independent experiment. The 8-week-old C57BL/6J WT mice were treated with busulfan (40 mg/kg) and used as recipient mice 1 month later.

Magnetic-activated cell sorting (MACS)

The testes from 8-week-old C57BL/6J wild-type mice or 4-month-old *Foxc2*$^{f/+}$ and *Foxc2*$^{f/-}$*; Ddx4*-cre mice (n=4) were minced and digested in the collagenase type IV (1mg/mL, Sigma) and DNase I (500μg/mL, Sigma) at 37°C for 15 min. The cell suspension was pipetted up and down once every 5 minutes and the digestion process was stopped with DMEM (containing 10% FBS). The cell suspension was filtered through a 40-μm nylon mesh, and after centrifugation, the cells were resuspended in 8mL PBS. The 15 mL conical centrifuge tubes were slowly overlayed with 2 mL of 70% Percoll solution, 2 mL of 30% Percoll solution, and then 2 mL of testicular cell suspension and centrifuge at 600 × g for 10 min at 4 °C without using the centrifuge brake. After centrifugation, the cells at the interface between the 70% and the 30% Percoll solution were carefully removed into the new conical centrifuge tubes, washed with PBS, and then centrifuge at 600 × g for 10 min at 4 °C. After centrifugation, the cells were resuspended in 360μL MACS buffer, added with 40μL of magnetic microbeads conjugated with anti-Thy-1 antibody (Miltenyi Biotec 130–049–101, Auburn, CA), and mixed well. Incubate the cell suspension containing Thy-1 microbeads for 20 min at 4 °C. Mix gently by tapping every 10 min. Add 20 mL of MACS buffer to the tube to dilute Thy-1 microbeads and centrifuge at 300 ×g for 10 min at 4 °C. Remove the supernatant completely and resuspend in 2 mL of MACS buffer. Place the separation columns (MS Column; Miltenyi Biotec 130-042-201) in the magnetic field of the mini MACS Separation Unit (Miltenyi Biotec 130-142-102) and rinse with 0.5 mL of MACS buffer. Apply the cell suspension to the columns (500μL/ column). After the cell suspension has passed through the column and the column reservoir is empty, wash the column with 0.5mL of MACS buffer three times. Remove the column from the MACS Separation Unit and elute the magnetically retained cells slowly into a 50 mL conical centrifuge tube with 1mL of MACS buffer using the plunger supplied with the column. Centrifuge the tube containing the cells at 600 × g for 10 min at 4 °C and resuspend the cell pellet with 10mL of MACS buffer for rinsing. Repeat this step once. After the final rinsing step, resuspend cells in 0.04% BSA and count the cell number.

Single-cell RNA-seq

The MACS-sorted Thy1^{+} cells were used for loading onto the Chromium Single Cell 3’ Chip kit v2 (10x Genomics, PN-120236) according to the instructions. Cell capturing and library preparation was performed following the kit instructions of the Chromium Single Cell 3’ v2 Library and Gel Bead Kit (10x Genomics, PN-120237). In brief, 5000 cells were targeted for capture, and after cDNA synthesis, 10-12 cycles were used for library amplification. The libraries were then size-selected, pooled, and sequenced on a Novaseq 6000 (Illumina). Shallow sequencing was performed to access the library quality and to adjust the subsequent sequencing depth based on the capture rate and the detected unique molecular indices (UMI).

Single-cell RNA-seq data processing

Raw sequencing reads were processed using the Cell Ranger v.3.0.1 pipeline of the 10x Genomics platform. In brief, reads from each sample were demultiplexed and aligned to the

mouse mm10 genome, and UMI counts were quantified for each gene per cell to generate a gene-barcode matrix. Default parameters were used. The UMI counts were analyzed using the Seurat R Package (58) (v.3.0.1) following the Seurat pipeline. Cells with more than 200 detected genes or less than 10% mitochondria reads were retained. Genes not detected in at least 10 cells were removed from subsequent analysis. The resulting matrix was normalized, and the most variable genes were found using Seurat's default settings, then the matrix was scaled with regression against the mitochondria reads. The top 2000 variable genes were used to perform PCA, and Jackstraw was performed using Seurat's default settings. Variation in the cells was visualized by UMAP for the top principal components. Cell types were determined using marker genes identified from the literature (59). We used the Seurat function CellCycleScoring to determine the cell cycle phase, as this program determines the relative expression of a large set of G2-M and S-phase genes. After removing the undefined cells, the spermatogonia were used for trajectory analysis, and the single-cell pseudotime trajectory was constructed with the Monocle 2 package (v2.12.0) ((60)–(62)) according to the provided documentation. The Monocle function clusterCells was used to detect cell clusters between clusters. The Seurat function FindAllMarkers with default settings was used to find DEGs upregulated in each cluster compared to the other cells. The top200 DEGs of cluster1 were used for ordering cells, and the discriminative dimensionality reduction with trees (DDRTree) method was used to reduce the data to two dimensions. The dynamic expression patterns with the spermatogonial developmental trajectory of specific genes were visualized using the Monocle function plot_genes_in_pseudotime and plot_pseudotime_heatmap. The procession data of the adult human single-cell dataset was downloaded from Gene Expression Omnibus (GEO): GSE112013 and the UMI counts were analyzed using the Seurat R Package (v.3.0.1) following the Seurat pipeline with the same parameters and functions as mentioned previously. According to the known markers, the germ cells characterized was used for trajectory analysis, and the single-cell pseudotime trajectory was constructed with the Monocle 2 package (v2.12.0) as mentioned previously.

CUT & Tag sequencing and analysis

CUT&Tag assay was performed using CUT&Tag 2.0 High-Sensitivity Kit (Novoprotein scientific Inc., Cat# N259-YH01). The detailed procedures were described in (40; 63). In brief, cells were harvested by trypsin and enriched by ConA-magnetic beads. 10,000 cells were re-suspended in 100 mL Dig-wash Buffer (20 mM HEPES pH 7.5; 150 mM NaCl; 0.5 mM Spermidine; 13 Protease inhibitor cocktail; 0.05% Digitonin) containing 2 mM EDTA and a 1:100 dilution of primary FOXC2 antibody. The primary antibody was incubated overnight at 4°C. Beads were washed in Dig-wash Buffer 3 times and incubated with secondary antibody for 1 hour at a dilution of 1:200. After incubation, the beads were washed 3 times in Dig-Hisalt Buffer (0.05% Digitonin, 20 mM HEPES, pH 7.5, 300 mM NaCl, 0.5 mM Spermidine, 13 Protease inhibitor cocktail). Cells were incubated with proteinA-Tn5 transposome at 25°C for 1 h and washed 3 times in Dig-Hisalt buffer to remove unbound proteinA-Tn5. Next, cells were re-suspended in 100mL Tagmentation buffer (10 mM MgCl2 in Dig-Hisalt Buffer) and incubated at 37°C for 1 h. The tagmentation was terminated by adding 2.25 mL of 0.5 M EDTA, 2.75 mL of 10% SDS and 0.5 mL of 20 mg/mL Proteinase K at 55°C for 1 hour. The DNA fragments were extracted by phenol chloroform and used for sequencing on an Illumina HiSeq instrument (Illumina NovaSeq 6000) to generate 2 × 150-bp paired-end reads following the manufacturer's instructions.

Raw reads were analyzed by removing low-quality or adaptor sequences using Trim_galore (v0.5.0) and cleaned reads were mapped to the reference genome mm10 using Bowtie2 (v2.2.5). We used MACS2 (v2.1.2) to call peaks found in different groups. Homer (v4.11.1) de novo motif discovery tool was used for finding the binding motifs of Foxc2 with the findMotifsGenome.pl command. The binding potential of candidate target genes at the binding motif was predicated using the JASPAR Scan function (binding potential >0.8). The

peaks filtered by fold change more than 5 and transcription start site (TSS) less than 3000 bp were annotated by R package Chip Seeker for gene category analysis. R package Cluster profiler was used for gene function annotation such as KEGG and GO analysis.

Enrichment analyses

Gene Ontology (GO) and KEGG pathway enrichment analyses were conducted using the ClusterProfiler package (v3.12.0) (Yu et al., 2012) and the ClueGO app (v2.5.7) in Cytoscape (v3.8.1) with default settings and a p-value cut-off of 0.05. GSEA enrichment analysis was assessed using the GSEA (v4.0.2) algorithm with MSigDB (v7.0) with default settings. The signaling pathways enriched by niche-derived paracrine factors and undifferentiated SPG-derived membrane proteins in the DEGs of the four samples were characterized. Then for each niche cell type, the niche-derived signaling pathways in all four samples were crossed with the SSC-derived signaling pathways to identify the candidate signaling pathways pivotal to SSCs maintenance.

Transplantation assay

The 8-week-old C57BL/6J WT mice were treated with busulfan (40 mg/kg) and used as recipient mice 1 month later. SSCs were transplanted into the testis of recipient mice (1×10^3 cells/testis), and two months after transplantation, the testes were harvested and observed under a fluorescence microscope.

Fluorescence-activated cell sorting (FACS)

Single-cell suspensions were generated from testes or *in vitro* cultured SSCs. FACS was performed using an SH800 machine (Sony Biotechnology) to isolate the GFP^+ cells. Briefly, the GFP^+ gating area was based on the point of the fluorescence intensity axis where cells were considered as being GFP^+, set based on the background fluorescence intensity of a non-transgenic control testis cell population.

Immunofluorescence

Mouse testes were fixed in 4% Paraformaldehyde (PFA) at 4°C overnight, dehydrated, embedded in paraffin, and cut into 5-μm thick sections. The rehydrated mouse or human testis sections were subjected to antigen retrieval, blocked in 5% BSA with 0.1% Triton X-100, and incubated with primary antibody (Supplemental Table S4) at 4°C overnight, including the germ cell marker DDX4, undifferentiated spermatogonia markers ZBTB16, LIN28A, ECAD (64), GFRA1, EOMES, PAX7, progenitor marker NEUROG3, and spermatocyte marker SYCP3 (65). After three 5-min washes in PBS, the sections were incubated with secondary antibodies (Supplemental Table S4) and DAPI (Sigma) at 37°C for 1 h. After three 5-min washes in PBS, coverslips were then mounted on glass slides using anti-quencher fluorescence decay (Solarbio). Images were captured using a Zeiss 780 laser-scanning confocal microscope. Whole-mount immunofluorescence of seminiferous tubules was performed as previously described (66). Briefly, seminiferous tubules were disentangled from testicular biopsies and immediately fixed in 4% PFA at 4°C for 12 h. After fixation, the seminiferous tubules were permeabilized with 0.5% Triton X-100 in PBS and treated with 5% BSA in PBS overnight at 4°C. After three 30-min washes, the seminiferous tubules were incubated with primary antibody (Supplemental Table S4) overnight at 4°C. After three 30-min washes, the seminiferous tubules were incubated with species-specific secondary antibodies and DAPI at 4°C for 12 h. After three 30-min washes, the seminiferous tubules

were mounted on slides with anti-quencher fluorescence decay (Solarbio) and observed with a Zeiss 780 laser-scanning confocal microscope.

RNA isolation and quantitative RT-PCR analysis

Total RNA was extracted from the testes or cultured cells using the RNeasy kit (Qiagen), reverse-transcribed using RevertAid First Strand cDNA Synthesis kit (Thermo), and processed for qRT-PCR using PowerUp SYBR Green Master Mix (Applied Biosystems) and a LightCycler 480 system (Roche) with gene-specific primers (Supplemental Table S4). Reactions were run in triplicate and the mRNA levels were normalized to Gapdh and quantified using the delta-delta Ct method. The values shown are mean ± s.e.m. from three biological replicates.

Tamoxifen inducible

According to a previous report for activation of iCre (9), the mice were fed with TD.130859 (TAM diet) for three days. The food was formulated for 400 mg tamoxifen citrate per kg diet, which would provide ~40 mg tamoxifen per kg body weight per day.

Analyses of cyst length

The cyst length was obtained according to the previous report (67). Briefly, to determine the cyst length, after immunofluorescence staining with anti-E-CAD antibody, the whole mount seminiferous tubule specimens were observed under a fluorescence microscope. The E-CAD staining coupled with staining for FOXC2 enabled us to reliably identify syncytial cysts of $FOXC2^{+}$ cells.

Analyses of cell density

The cell density was counted according to a previous report (68). Briefly, the densities of the $ZBTB16^{+}$, $GFRA1^{+}$, $LIN28A^{+}$, or $FOXC2^{+}$ cells were measured on the seminiferous tubules with whole-mount staining, the numbers of which per 1000 Sertoli cells were determined.

Sperm counts

Total sperm counts were obtained according to the previous report (69). Briefly, epididymal caput and cauda were minced and incubated in prewarmed M16 medium (Sigma-Aldrich) at 37°C in air containing 5% CO2 for 30 min to allow the sperm to swim out. Then, the sperm were diluted in water and counted using a hemocytometer.

Histology, evaluation of degenerating tubules

Testes of WT and mutant mice were fixed with PFA fixative and processed for paraffin-embedded section preparation (5 μm thick) and hematoxylin and eosin staining, according to standard procedures. The percentage of degenerating seminiferous tubules was calculated based on the cross-sections of seminiferous tubules (n > 200) that appeared on one transverse section for each testis. In normal (WT) mouse testes, four generations of germ cells, each synchronously progressing through spermatogenesis, form cellular associations of fixed composition (called seminiferous epithelial stages). In the testes of $Foxc2^{flox/-}$; Ddx4-cre mice, a few tubule cross-sections lacked one or more out of the four germ cell layers, which was defined as “degenerative tubules” in this study.

Statistical analysis

All statistical analyses were performed using GraphPad Prism (v7.0). All experiments were repeated at least three times, and data for evaluated parameters are reported as mean ± s.e.m. The p-values were obtained using two-tailed unpaired Student's t-tests or one-way ANOVA followed by Tukey test (ns represents p-value > 0.05, * represents p-value < 0.05, ** represents p-value < 0.01, *** represents p-value < 0.001, and **** represents p-value < 0.0001).

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Reviewer #1 (Public Review):

The expression and localization of Foxc2 strongly suggest that its role is mainly confined to As undifferentiated spermatogonia (uSPGs). Lineage tracing demonstrated that all germ cells were derived from the FOXC2+ uSPGs. Specific ablation of the FOXC2+ uSPGs led to the depletion of all uSPG populations. Full spermatogenesis can be achieved through the transplantation of Foxc2+ uSPGs. Male germ cell-specific ablation of Foxc2 caused Sertoli-only testes in mice. CUT&Tag sequencing revealed that FOXC2 regulates the factors that inhibit the mitotic cell cycle, consistent with its potential role in maintaining a quiescent state in As spermatogonia. These data made the authors conclude that the FOXC2+ uSPG may be the true SSCs, essential for maintaining spermatogenesis. The conclusion is largely supported by the data presented, but two concerns should be addressed: 1) terminology used is confusing: primitive SSCs, primitive uSPGs, transit amplifying SSCs... 2) the GFP+ cells used for germ cell transplantation should be better controlled using THY1+ cells.

Reviewer #2 (Public Review):

The authors found FOXC2 is mainly expressed in As of mouse undifferentiated spermatogonia (uSPG). About 60% of As uSPG were FOXC2+ MKI67-, indicating that FOXC2 uSPG were quiescent. Similar spermatogonia (ZBTB16+ FOXC2+ MKI67-) were also found in human testis.

The lineage tracing experiment using *Foxc2CRE/+;R26T/Gf/f* mice demonstrated that all germ cells were derived from the FOXC2+ uSPG. Furthermore, specific ablation of the FOXC2+ uSPGs using *Foxc2Cre/+;R26DTA/+* mice resulted in the depletion of all uSPG population. In the regenerative condition created by busulfan injection, all FOXC2+ uSPG survived and began to proliferate at around 30 days after busulfan injection. The survived FOXC2+ uSPGs generated all germ cells eventually. To examine the role of FOXC2 in the adult testis, spermatogenesis of Foxc2f/-;Ddx4-cre mice was analyzed. From a 2-month-old, the degenerative seminiferous tubules were increased and became Sertoli cell-only

seminiferous tubules, indicating FOXC2 is required to maintain normal spermatogenesis in adult testes. To get insight into the role of FOXC2 in the uSPG, CUT&Tag sequencing was performed in sorted FOXC2+ uSPG from *Foxc2CRE/+;R26T/Gf/f* mice 3 days after TAM diet feeding. The results showed some unique biological processes, including negative regulation of the mitotic cell cycle, were enriched, suggesting the FOXC2 maintains a quiescent state in spermatogonia.

Lineage tracing experiments using transgenic mice of the TAM-inducing system was well-designed and demonstrated interesting results. Based on all data presented, the authors concluded that the FOXC2+ uSPG are primitive SSCs, an indispensable subpopulation to maintain adult spermatogenesis.

The conclusion of the mouse study is mostly supported by the data presented, but to accept some of the authors' claims needs additional information and explanation. Several terminologies define cell populations used in the paper may mislead readers.

1. "primitive spermatogonial stem cell (SSC)" is confusing. SSCs are considered the most immature subpopulation of uSPG. Thus, primitive uSPGs are likely SSCs. The naming, primitive SSCs, and transit-amplifying SSCs (Fig. 7K) are weird. In general, the transit-amplifying cell is progenitor, not stem cell. In human and even mouse, there are several models for the classification of uSPG and SSCs, such as reserved stem cells and active stem cells. The area is highly controversial. The authors' definition of stem cells and progenitor cells should be clarified rigorously and should compare to existing models.

2. scRNA seq data analysis and an image of FOXC2+ ZBTB16+ MKI67- cells by fluorescent immunohistochemistry are not sufficient to conclude that they are human primitive SSCs as described in the Abstract. The identity of human SSCs is controversial. Although Adark spermatogonia are a candidate population of human SSCs, the molecular profile of the Adark spermatogonia seems to be heterogeneous. None of the molecular profiles was defined by a specific cell cycle phase. Thus, more rigorous analysis is required to demonstrate the identity of FOXC2+ ZBTB16+ MKI67- cells and Adark spermatogonia.

3. FACS-sorted GFP+ cells and MACS-THY1 cells were used for functional transplantation assay to evaluate SSC activity. In general, the purity of MACS is significantly lower than that of FACS. Therefore, FACS-sorted THY1 cells must be used for the comparative analysis. As uSPGs in adult testes express THY1, the percentage of GFP+ cells in THY1+ cells determined by flow cytometry is important information to support the transplantation data.

4. The lineage tracing experiments of FOXC2+-SSCs in Foxc2CRE/+;R26T/Gf/f showed ~95% of spermatogenic cells and 100% progeny were derived from the FOXC2+ (GFP+) spermatogonia (Fig. 2I, J) at month 4 post-TAM induction, although FOXC2+ uSPG were quiescent and a very small subpopulation (~ 60% of As, ~0.03% in all cells). This means that 40% of As spermatogonia and most of Apr/Aal spermatogonia, which were FOXC2 negative, did not contribute to spermatogenesis at all eventually. This is a striking result. There is a possibility that FOXC2CRE expresses more widely in the uSPG population although immunohistochemistry could not detect them.

5. The CUT&Tag_FOXC2 analysis on the FACS-sorted FOXC2+ showed functional enrichment in biological processes such as DNA repair and mitotic cell cycle regulation (Fig.7D). The cells sorted were induced Cre recombinase expression by TAM diet and cut the tdTomato cassette out. DNA repair process and negative regulation of the mitotic cell cycle could be induced by the Cre/lox recombination process. The cells analyzed were not FOXC2+ uSPG in a normal physiological state.

6. Wei et al (Stem Cells Dev 27, 624-636) have published that FOXC2 is expressed predominately in As and Apr spermatogonia and requires self-renewal of mouse SSCs; however, the authors did not mention this study in Introduction, but referred shortly this at the end of Discussion. Their finding should be referred to and evaluated in advance in the Introduction.

Reviewer #3 (Public Review):

By popular single-cell RNA-seq, the authors identified FOXC2 as an undifferentiated spermatogonia-specific expressed gene. The FOXC2+-SSCs can sufficiently initiate and sustain spermatogenesis, the ablation of this subgroup results in the depletion of the uSPG pool. The authors provide further evidence to show that this gene is essential for SSCs maintenance by negatively regulating the cell cycle in adult mice, thus well-established FOXC2 as a key regulator of SSCs quiescent state.

The experiments are well-designed and conducted, the overall conclusions are convincing. This work will be of interest to stem cell and reproductive biologists.