Androgen promotes differentiation of PLZF⁺ spermatogonia pool via indirect regulatory pattern

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Abstract

Background: Androgen plays a pivotal role in spermatogenesis, accompanying a question how androgen acts on germ cells in testis since germ cells lack of androgen receptors (AR). Promyelocytic leukemia zinc-finger (PLZF) is essential for maintenance of undifferentiated spermatogonia population which is terminologically called spermatogonia progenitor cells (SPCs).

Aims: We aim to figure out the molecular connections between androgen and fates of PLZF⁺ SPCs population.

Method: Immunohistochemistry was conducted to confirm that postnatal testicular germ cells lacked endogenous AR. Subsequently, total cells were isolated from 5 dpp (day post partum) mouse testes, and dihydrotestosterone (DHT) and/or bicalutamide treatment manifested that Plzf was indirectly regulated by androgen. Then, Sertoli cells were purified to screen downstream targets of AR using ChIP-seq, and gene silence and overexpression were used to attest these interactions in Sertoli cells or SPCs-Sertoli cells co-culture system. Finally, these connections were further verified in vivo using androgen pharmacological deprivation mouse model.

Results: Gata2 is identified as a target of AR, and β1-integrin is a target of Wilms’ tumor 1 (WT1) in Sertoli cells. Androgen signal negatively regulate β1-integrin on Sertoli cells via Gata2 and WT1, and β1-integrin on Sertoli cells interacts with E-cadherin on SPCs to regulate SPCs fates.

Conclusion: Androgen promotes differentiation of PLZF⁺ spermatogonia pool via indirect regulatory pattern.

Keywords: Androgen receptor, β1-integrin, PLZF, Spermatogonial stem cells, WT1

Plain English summary

Androgen signaling plays a pivotal role in spermatogenesis, but the molecular mechanisms underlying androgen action in this process are unclear. Specifically, it is unknown if androgen receptor (AR) is expressed in germ cells. Thus, it’s interesting to reveal how androgen induces differentiation of spermatogonial stem cells (SSCs). Here we observed the AR is deficient in SSCs before spermatogenesis onset. Then we examined a regulatory role of the AR in spermatogenesis using a SSCs-Sertoli cells co-culture system, and demonstrated that androgen negatively regulated the key gene for SSCs stemness, Plzf. Additionally, we identified Gata2 as a target of AR in Sertoli cells, and demonstrated that Wilms tumor 1 (WT1) and β1-integrin as two putative intermediate molecules to transfer differentiation signals from Sertoli cells to SSCs, which was further verified using androgen pharmacological-deprivation mice model. These results demonstrate a regulatory pattern of androgen in SSCs niche in an indirect way via multiple steps of signal transduction: androgen activates AR in Sertoli cells to indirectly regulate ITGB1 on Sertoli cells via intermediate molecule GATA2 and WT-1, and ITGB1 on Sertoli cells interacts with E-cadherin on SSCs to regulate SSCs fates.
**Background**

Spermatogonial stem cells (SSCs) reside in the basement membrane and are subjected to control signal in the micro-environment to function as the original source for sperms in male testis. In the hierarchy of differentiation, spermatogonia consist of different subpopulations such as $A_s$, $A_{pp}$, $A_{ab}$, $A_{1}$,…,$A_{ab}$, $A_{pp}$, B etc. Of them, $A_s$, $A_{pp}$ and $A_{ab}$ spermatogonia are referred to as undifferentiated population, and are nominated as spermatogonial progenitor cells [1]. The mechanism of SSCs differentiation is not clear yet, especially the role of androgen in spermatogenesis is still a mystery. Androgen is produced by testicular Leydig cells and functions via binding and activation of its receptor AR in cytoplasm. Activated AR is subsequently imported into nucleus after release of heat shock protein and functions as a transcription factor [2]. Numerous target genes of AR have been identified, involving in many biological processes such as gonad development [3] and tumorigenesis [4]. However, the exact biological mechanism of AR in spermatogenesis is not fully understood, and whether AR is expressed in testicular germ cells is controversial. Some studies detected AR signal in germ cells of mouse testis [5, 6]. Other studies suggested that AR is presented only in the somatic cells of rodent testis [7]. It has been shown that germ cell specific $Ar$ knockout mice still had normal sperm [8] but conditional deletion of AR in Leydig or Sertoli cells caused spermatogenesis defects [9, 10]. These results suggest that AR expressed in Sertoli cells, Leydig cells and perivascular myoid cells may participate in spermatogenesis via interacting with surrounding spermatogonia [11]. However, SyceP1-driven Cre for $Ar$ deletion in germ cells was used in the study mentioned above [8], which only indicates AR is not required in germ cells since meiosis onset. Moreover, studies reported that androgen functions as a signal molecule in SSCs niche, namely androgen acts on peritubular myoid (PM) cells surrounding the seminiferous tubule to stimulate PM cells to produce GDNF, to promote self-renewal of SSCs [12, 13], indicating a complicated role of androgen in testicular niche. In all, the mechanism of spermatogenesis mediated by androgen still needs to be further investigated.

*Plzf* is a key transcription suppressor gene for SPCs maintenance. It was first discovered by its association with acute promyelocytic leukemia [14], and was subsequently characterized as an undifferentiated marker for SSCs in rodents [15] and primates [16]. Loss of *Plzf* did not affect spermatogonia formation, but led to progressive and significant deficiency of SSCs after neonatal life and finally caused infertility [15, 17], indicating its critical role in SSCs maintenance. Moreover, PLZF expression was detected in spermatogonia $A_s$, $A_{pp}$ and $A_{ab}$, not restricted in SSCs [18]. Thus, PLZF is a marker of SPCs, and PLZF is an important factor for maintenance of this pool [19]. Although the link of androgen and PLZF has not been reported in germ cells, much evidence from prostate tumorigenesis studies suggests the interaction between AR and PLZF. For example, *Plzf* represses prostate tumorigenesis and its expression can be inhibited by androgen antagonist, bicalutamide [20]. In prostate cancer cell line PCa cells, PLZF was identified as a repressor of AR as well as an activator of regulated in development and DNA damage responses 1 (REDD1), which suppressed mTORC1 [21]. AR was characterized as a critical transcriptional factor in prostate tumorigenesis [4], and mTORC1 has been found to participate in EMT (Epithelial mesenchymal transition) in prostate cancer [22]. Thus, PLZF functions as tumor suppressor and interacts with AR in prostate cancer system, but it’s unclear whether similar links exist in germ line.

In testsis, Sertoli cells in base membrane form niches to protect SSCs and regulate their fates [23], and many surface proteins, such as cadherins and integrins, are identified as functional components in the niche [24]. Many of these molecules are AR responsive and associated with the fate of SSCs [25], but the mechanism is largely unknown. Also, it’s necessary to focus on Wt-1 gene, which is specifically expressed in Sertoli cells and required for Sertoli cell lineage maintenance [26, 27]. Moreover, Wt1 functions as a suppressor of $Ar$ [28]. Thus, we ask whether Wt1 participates in the regulation of spermatogenesis mediated by androgen signal.

Here, we studied AR expression pattern in testis of postnatal mouse using a monoclonal antibody, and detected weak AR signal in pre-spermatogonia of 2 dpp testis, but found that this signal was absent in germ cells from 3 dpp, instead appeared exclusively in somatic cells. Spermatogenesis starts from about 5 dpp [29], so the possibility that germ cells need AR for spermatogenesis is eliminated. Thus, we investigated the indirect regulation pattern of androgen on SPCs differentiation via Sertoli cells using a SPCs-Sertoli cells co-culture system. Two transcription factors Gata2 and WT1 were identified as AR’s downstream targets in Sertoli cells, and a surface protein β1-integrin was characterized as a target of WT1 and may function as an intermediate molecule to transfer androgen signal from Sertoli cells to SPCs to regulate spermatogenesis. In all, we demonstrated the complicated regulation pattern of androgen in spermatogenesis in SSCs niche, which requires the cooperation of Sertoli cells and many intermediate molecules to regulate *Plzf* expression in undifferentiated spermatogonial populations.

**Methods**

**Animals**

CD-1 mice were supplied by Comparative Medicine Centre of Yangzhou University, and all animal experiments have been approved by Animal Protection Committee of Nanjing Agricultural University.
Isolation and culture of total testicular cells or SPCs

Total testicular cells extracted from testes of 5 dpp mice followed previous protocol [30] with minor modifications. Testes were cut into small particles and followed by collagenase IV and trypsin digestion. Undigested collagen was removed using micropipette after centrifugation, and cell pellet was resuspended in culture medium and then transferred to gelatin coated plates for culture. To identify the total testicular cells’ co-culture system, the expression of Sox9 and Wt1 (Sertoli cells marker), Coup-tf II (Leydig cells marker), Sma (Peritubular Myoid cells marker), Mvh (germ line marker), Ar, Ssp3 (meiosis marker), c-kit (SPCs differentiation marker), integrin-aV, Cd9, integrin-β1 (a6), Integrin-a6 and Plzf (SPCs marker) were detected using RT-PCR (Fig. 2u and v).

For SPCs sorting, resuspended cells after two-enzyme digestion were filtered with 70-μm filter and subsequently incubated with mouse Thy-1.2 antibody coated magnetic beads (BD, Cat.551518). Thy1+ fraction was collected and cultured on mitotically inactivated STO feeder layer at 37 °C with 5% CO2. Culture medium for total testicular cells or SPCs was composed of 90% MEMa, 10% FBS (Gibco), 1 mg/mL bFGF (Sino Bio Inc., 10,014-HNAE), 1 ng/mL EGF (Sino Bio Inc., 50,482-MNAY), 1 ng/mL GDNF (Sino Bio Inc., 10,561-HNCH), 10 ng/mL LIF (Santa Cruz, sc-4378), 20 μg/mL transferrin (sigma, T8158), and 5 μg/mL insulin (Aladdin, 1113907). For androgen stimulation or deprivation, dihydrotestosterone (sigma, cat. 31,573) or bicalutamide (sigma, PHR-1678) was added to the culture medium.

Testes of 5 dpp mice [31]. Briefly, testes were encapsulated and seminiferous tubules were pooled and digested with 0.5 mg/mL collagenase IV. After centrifugation, the supernatant containing the Leydig cells was pipetted out. The tubules were further digested with 1 mg/mL collagenase IV and 10 μg/mL DNase to detach peritubular cells. Digestion was monitored under microscopy. The supernatant containing peritubular cells after centrifugation was discarded. Finally, single cell suspension was obtained after trypsin digestion and blocked by FBS. The cell pellet was resuspended in DMEM/F12 media supplemented with 2.5% FBS, 5% horse serum, 2 mM L-Gln, 100 U/mL penicillin, 1μM sodium pyruvate, and then plated. After 48 h, hypotonic treatment was performed to remove germ cells as described [32], and the purity of Sertoli cells was assessed by morphology and immunofluorescent staining of WT1 (purity of Sertoli cells: 95%) [Fig. 4a-d and i].

Sertoli cell isolation

A modified method was used to isolate Sertoli cells from testes of 5 dpp mice [31]. Briefly, testes were encapsulated and seminiferous tubules were pooled and digested with 0.5 mg/mL collagenase IV. After centrifugation, the supernatant containing the Leydig cells was pipetted out. The tubules were further digested with 1 mg/mL collagenase IV and 10 μg/mL DNase to detach peritubular cells. Digestion was monitored under microscopy. The supernatant containing peritubular cells after centrifugation was discarded. Finally, single cell suspension was obtained after trypsin digestion and blocked by FBS. The cell pellet was resuspended in DMEM/F12 media supplemented with 2.5% FBS, 5% horse serum, 2 mM L-Gln, 100 U/mL penicillin, 1μM sodium pyruvate, and then plated. After 48 h, hypotonic treatment was performed to remove germ cells as described [32], and the purity of Sertoli cell was assessed by morphology and immunofluorescent staining of WT1 (purity of Sertoli cells: 95%) [Fig. 4a-d and i].

ChIP-qPCR

To characterize genome-wide binding patterns of AR and WT1 in Sertoli cells, ChIP and input DNA libraries were performed as previously described [33]. Briefly, cells were cross-linked with 1% formaldehyde for 10 min at room temperature and formaldehyde was then inactivated by the addition of 125 mM glycine for 5 min. Sonicated DNA fragments with 100-300 bp were pre-cleared and then immunoprecipitated with Protein A + G Magnetic beads coupled with anti-Androgen Receptor antibody (Millipore, 06-680) or anti-Wilms’ tumor 1 antibody (Abcam, ab69901). After reverse crosslinking, immunoprecipitated DNAs and input DNAs were end-repaired and ligated adapters to the DNA fragments using NEBNext Ultra End-Repair/ΔA-Tailing Module (E7442, NEB) and NEBNext Ultra Ligation Module (E7445, NEB). High-throughput sequencing of the ChIP fragments was performed using Illumina NextSeq 500 following the manufacturer’s protocols. The raw sequencing data were processed with trimmomatic (version 0.36) to filter low-quality reads [34]. The resulting data were mapped using bowtie2 (version 2.2.9) to the UCSC mm10 genome reference [35]. Peak detection was performed using the MACS peak finding algorithm (Model-based Analysis of ChIP-Seq; version 1.4.2) with parameters --nomodel --shift-size 25 and the p-value cutoff set to 0.05 [36].

Primers for ChIP-qPCR

GATA2-F: GCTAGAGAGTGCATTGGGGA GATA2-R: AGTTCCTGGGGCTGCGAG.

ITGB1-F: AGCTACATTCTTCTGAGCAATTATGGGA IT
GB1-R: CTGCTCTCCCAAGGACTCAA.

Control-F:TTAGGTGGCCTCAGATCCCT Control-R: CCTGCTCTCCTTTTGGACAG.

Bicalutamide injection

Bicalutamide injection was performed as described [37] with minor modifications: 6-week old CD1 male mice were intraperitoneally injected with bicalutamide (20 mg/kg) or DMSO (vehicle) once every other day for 4 times.
Immunohistochemistry (IHC) and immunofluorescence (IF)
Mouse testes were transfected and fixed in 4% neutral paraformaldehyde overnight, and subsequently dehydrated and embedded in paraffin. Histological sections were dewaxed and rehydrated in ethanol series, followed by microwave antigen retrieval in 0.01 M citrate (pH = 6.0) and methanol/H₂O₂ treatment at room temperature. After blocking with 5% goat serum, the slides were incubated with primary antibodies against PLZF (Santa Cruz, sc-22,839), AR (Abcam, ab133273), ITGB1 (Abcam, ab179471), c-Kit (Cell Signaling, 2074), and biotin labelled goat anti-mouse IgG (Vector, BA-9200) or goat anti-rabbit IgG (Vector, BA-1000), respectively. Streptavidin-HRP (Jackson Lab) and DAB kit (Vector, sk4100) were used for visualization. Cell IF was carried out as described [38], primary antibodies against MVH (Cell Signaling, 8761s), PLZF (Santa Cruz, sc-2831 and sc-22,839), WT1 (Abcam, ab89901), CDH1 (Abcam, ab76055), CD9 (Affinity, DF6565), ITGB1 (BD, 610467) and secondary antibodies Alexa Flour532 goat anti-rabbit (Invitrogen, A-11009), goat anti-mouse IgG FITC (ZSGB-BIO, ZF-0312) are used.

BrdU assay
Mice were intraperitoneally injected with BrdU (sigma, B5002, 50 mg/kg) 4 h before testis harvesting. Testes were fixed in 4% neutral PFA and embedded with paraffin. For BrdU detection, sections were incubated with 2 M HCl at 37 °C for 1 h, and washed with 0.1 M boric acid (pH = 8.5) for 3 times. IF was performed as aforementioned (BrdU, Santa Cruz, sc-32,323).

Gene silence and overexpression
The validated siRNA against PLZF (5′-CCAGCAAGAUGUUGAGAATT-3′) [39], ITGB1 (5′-UGAGAAAUGUUGGAACACUUUCGCUCC-3′) [40], WT1 (5′-GGCGAUUG-GUUAAGGCUCAUUU-3′) [41], gata2 (5′-CGGGCAC-CUGUUGUGCAAAUUGUCA-3′) [42], control (5′-UU CUCCGAACUGUGUCAGUTT-3′) and shRNA sequences against ITGB1 (5′-AAATCAAAAAGACAGATGTGATGATTTAGAATCTCCTGTGAATTCATCAATCCTGGG-3′) [43], control (5′-AAATCCAAA AACAACAGATGAGACCACCATCTGTGAAT TGGTGCTTCTTCATCTGGTGAG-3′) were synthesized by Shanghai GenePharma Company. AR expression plasmid was a gift from Prof. Zijie Sun (Stanford University). WT1 expression plasmid (EX-MM24822-02) was purchased from GeneCopoeia. Plasmids or siRNA oligos were transfected using lipofectamine 3000 (Invitrogen, L3000008).

Western blot and co-immunoprecipitation
For western blotting, tissue or cell lysates were separated in SDS PAGE gel and transferred to nitrocellulose membrane, and was subsequently blocked with 5% skim milk at 4 °C overnight and then incubated with primary antibodies Sycep3 (Santa Cruz, sc-74,569), Stra8 (Abcam, ab49602), claudin11 (Abcam, ab175236), nectin2 (Abcam, ab135246), GATA2 (Abclonal, A0677), GFRα1 (Affinity, DF7309), β-catenin (Cell Signaling, 8480p), β-tubulin (Anbo, P07437) and others mentioned before at room temperature for 1 h. The membranes were incubated with HRP-conjugated goat anti rabbit or mouse IgG (Santa Cruz, sc-2004, sc-2005) and ECL, and finally exposed the film. For Co-IP, lysates of adult mice testes or SPCs were prepared using a standard protocol (Beyotime, P0013), and 0.5 mg total proteins diluted with TBST were incubated with mouse anti-ITGB1 antibody (BD, 610467) and rabbit anti-CDH1 antibody (Santa Cruz, sc-7870) and rocked at 4 °C overnight. The pre-washed protein (A+G) sepharose beads (Beyotime, P2012) were added afterwards and incubated at 4 °C for 4 h. The beads were then washed three times with TBST, pelleted and boiled in 1XSDS loading buffer, and finally analyzed by western blot.

Statistical analysis
For cell counting, sections or immunofluorescent visual fields were selected randomly. Data was analyzed by SPSS statistics 17.0 and was presented as mean ± SD (standard deviation). Statistical significance was determined by t-test.

Results
PLZF+ spermatogonia pool keeps steady during testis development
The size of PLZF+ population in testis represents the capacity of self-renewal and spermatogenesis, thus the expression profile of PLZF was detected in neonatal (5 dpp), juvenile (10 dpp), puberty (20 dpp) and adult (42 dpp) testes using IHC staining. The results demonstrated that from neonatal to adult testes, all PLZF+ cells exclusively located at basement membrane with strong nucleus staining [Fig. 1a-d], or a few oblate pairs of cells in adult testes, which are Apr spermatogonia [Fig. 1d inset]. These PLZF+ cells were surrounded by Sertoli cells, which formed the microenvironment called niches. Strong PLZF intensity in PLZF+ spermatogonia is stable during testicular development.

AR expression in testis indicates an indirect regulation pattern in promoting spermatogenesis
Subsequently, we examined AR expression in 5, 10, 20 and 42 dpp testes using IHC, to investigate the role of
Fig. 1 Expression patterns of PLZF and AR during testis development. PLZF staining was exclusively detected in spermatogonia localized in basal membranes of 5 dpp (a), 10 dpp (b), 20 dpp (c) and 42 dpp (d) testes (inset in D, a representative of PLZF+ AR spermatogonia). AR staining in 5 dpp (e), 10 dpp (f), 20 dpp (g) and 42 dpp (h) testes was specifically detected in Sertoli cells. Weak AR staining was observed in pre-spermatogonia (red arrowheads) in the testicular lumen of 2 dpp testis, but not in Sertoli cells (i). In 3 dpp testis (j), pre-spermatogonia migrated to basement membranes and formed unique structures with surrounding Sertoli cells (red arrows in (j)), and identical structures were also found in 4 dpp and 5 dpp testes, red arrows in (e) and (k), and AR staining was detected in Sertoli cells but not in germ cells. k AR staining in 4 dpp testis, i a representative of niche in 5 dpp testis, in which SPCs were embraced by AR+ Sertoli cells (red frame). AR/PLZF dual IF exhibited no overlap of AR signals and PLZF signals in 4 dpp testis ((m), PLZF, (n), AR, (o), DAPI, (p), merge). From neonatal to adult stages, ratio of PLZF+ cells in seminiferous tubules decreased (q) but number of PLZF+ cells kept steady (r). Data represent means ± SD (**p < 0.01), n = 16, 4 of each group, bar = 20 μM
androgen in spermatogenesis. Intensive AR signals were detected in Sertoli cells, Leydig cells and myoid cells but not in germ cells [Fig. 1e-h], consistent with the conclusion that postnatal testicular germ cells lacked endogenous AR [7]. In 5 dpp testes, moderate AR staining was detected in Sertoli cells localized at basement membrane [Fig. 1e]. Both AR+ Sertoli cells number and AR signal intensity were obviously increased in 10 dpp testes, and some AR+ Sertoli cells moved to the second layers [Fig. 1f], and then formed AR+ layers in 20 dpp testes [Fig. 1g]. In 42 dpp testis, those AR+ second layers disappeared and AR+ cells distributed in different layers of seminiferous tubules [Fig. 1h]. In rodent Sertoli cells, AR expression was reported to start from 3 to 5 dpp [44], thereupon we examined AR expression in 2, 3 and 4 dpp testes, and detected AR signal in Sertoli cells only from 3 dpp testis and signal intensity grew stronger from 5 dpp [Fig. 1I-k]. We also noticed Sertoli cells did not express AR in 2 dpp testis as literature reported [44], but unexpectedly observed weak AR signal in pre-spermatogonia localized in the cord of testes [Fig. 1l]. Interestingly, AR signal was absent in the pre-spermatogonia of 3 dpp testes, at which time point some of these pre-spermatogonia had already migrated from cord to basal membrane [Fig. 1j]. Meanwhile, Sertoli cells began to express AR and two AR+Sertoli cells embraced one pre-spermatogonium to form a special structure nominated as ‘niche’ [Fig. 1j-l], implying AR in Sertoli cells commenced to function in microenvironment. Spermatogenesis starts from 3 to 5 dpp and differentiated spermatogonia can be observed at 6 dpp [29]. Dual immunofluorescent staining also exhibited no overlap of AR and PLZF signals in 4 dpp testes [Fig. 1m-p]. Collectively, AR expression pattern in Sertoli cells demonstrates their close connection with PLZF+ cells, but PLZF+ population is unable to be directly stimulated by androgen since they lack of endogenous AR when spermatogenesis starts.

Blockage of androgen with antagonist inhibits SPCs differentiation

To investigate the role of androgen in spermatogenesis, a co-culture system of testicular cells was established. Testes from 5 dpp mice (this time spermatogenesis just begins) were harvested and digested to single cells (composed of SPCs, differentiating spermatogonia and testicular somatic cells, and the markers of Sertoli cells (Sox9 and Wt-1), Leydig cells (Coup-tf II) and Peritubular Myoid cells (Sma) were detected using RT-PCR (Fig. 2u)) and cultured on gelatin or mitotically inactivated STO feeder layer (negative of AR expression) showed no obvious phenotype change after either DHT or bicalutamide treatment (data is not shown), suggesting that Sertoli cells were necessary for SPCs differentiation during androgen stimulation.

AR indirectly regulates Wt1 in Sertoli cells

To reveal regulatory mechanism of AR on spermatogenesis, we used ChIP-seq to screen the target genes of AR in purified Sertoli cells from 5 dpp mouse testis. In the candidates of targets, we noticed that Gata2 promoter was a binding region of AR (p-value < 0.01 by MACS [36]) [Fig. 4m]. This result provided an important cue, since Gata2 is identified as a factor to promote WT1 expression after DHT/bicalutamide treatment [Fig. 3g and h]. Additionally, we observed that DHT aggravated the down-regulation of PLZF expression induced by Plzf siRNA [Fig. 3i and j], confirming that DHT promoted SPCs differentiation. On the contrary, Thy1-MACS purified SPCs cultured on gelatin or mitotically inactivated STO feeder layer (negative of AR expression) showed no obvious phenotype change after either DHT or bicalutamide treatment (data is not shown), suggesting that Sertoli cells were necessary for SPCs differentiation during androgen stimulation.
Fig. 2 (See legend on next page.)
The β1-integrin gene is a direct target of WT1 in Sertoli cells
Since WT1 involves in multiple functions in Sertoli cells, we used ChIP-seq again to screen the target genes of WT1 regarding spermatogenesis in Sertoli cells. Notably, β1-integrin gene was identified as a direct binding target of WT1 [Fig. 5a]. Besides the role as a subunit of laminin receptor, β1-integrin is essential for SSCs homing since deletion of β1-integrin in SSCs or in Sertoli cells impaired SSCs homing [24], indicating its signal function in SSCs niche. After repeatedly validation of the binding of WT1 on β1-integrin gene using ChIP-qPCR [Fig. 5b], we further examined whether WT1 regulates β1-integrin expression at protein level. Knockdown of WT1 expression in Sertoli cells suppressed expression of β1-integrin [Fig. 5c], and WT1 overexpression strengthened

(See figure on previous page.)

Fig. 2 SPCs-somatic cells co-culture system was able to be maintained in vitro for short term. SPCs-somatic cells co-culture system stably expressed MVH (a) bright field, (b) MVH, (c) dapi, (d) merge, PLZF (e) bright field, (f) PLZF, (g) dapi, (h) merge, CD9 (i) bright filed, (j) WT1, (k) dapi, (l) merge, AR (m) bright filed, (n) AR, (o) dapi, (p) merge) and WT1 (q) bright filed, (r) WT1, (s) dapi, (t) merge. RT-PCR (n = 2) results exhibit the co-culture system expresses somatic markers (u) 1.Gapdh, 2.Sox9, 3. Sma, 4.Coup-tf II, 5. Wt1, 6.Mvh, Ar (V. 1) and germline markers (v) 2.Scp3, 3.kit, 4.Integrin-αV, 5.Cd9, 6.Integrin-β1, 7. Integrin-α6, 8. Pldf, 9. Mvh) after 3 passages. Western blot result confirmed this co-culture system was sensitive to DHT or bicalutamide stimulation, 1, DMSO; 2, 10nM DHT; 3, 10nM bicalutamide, n = 3 (w)
Fig. 4 (See legend on next page.)
β1-integrin’s expression level [Fig. 5d], suggesting that expression of β1-integrin was regulated by WT1. Moreover, we observed that androgen signal negatively regulated the expression of β1-integrin in Sertoli cells [Fig. 5e], and disturbance of Gata2 expression displayed similar impact on β1-integrin expression [Fig. 4p], which were consistent with aforementioned results that AR negatively regulated Gata2 via binding on its promoter region, and indirectly interfered in WT1 expression. Then we knocked down the expression of β1-integrin in co-culture system using siRNA and observed decreased expression level of PLZF [Fig. 5f], implying the indispensable role of β1-integrin in SPCs maintenance. Collectively, we concluded a cue from androgen signal to surface protein β1-integrin, via intermediate factor GATA2 and WT1 [Fig. 5g].

β1-integrin on Sertoli cells may function as an intermediate molecule to regulate SPCs differentiation

The important role of β1-integrin in vivo has been proven in Sertoli cells specific β1-integrin conditional knockout mice [24]. Here we used in vitro co-culture system to verify the role of β1-integrin on Sertoli cells for SPCs maintenance. Because primary Sertoli cells have limited prolifera-
tion ability of mice [24]. Here we used in vitro co-culture system to verify expression of β1-integrin before co-culture with SPCs. Morphology of Sertoli cells 72 h post infection [j], and infection efficiency was examined by expression of red fluorescence protein reporter [k]. Western blot revealed the relative expression level of β1-integrin was approximately decreased by 80% after β1-integrin shRNA interfering, 1, Igb1 siRNA, 2, control siRNA. Data represent means ± SD (*p < 0.05, **p < 0.01), n = 3 [l]. The representative view of AR binding sites (visualized by Integrative Genomics Viewer [http://software.broadinstitute.org/software/igenome]) at Gata2 promoter region [m]. ChIP-qPCR validated the binding of AR in the promoter region of Gata2 fold enrichment by antibodies against AR relative to control IgG was presented as mean value ± SD of two replicates [n]. Overexpression of AR or DHT treatment caused down-regulated expression levels of GATA2, WT1 and β1-integrin in Sertoli cells, n = 3 [o]. Knockdown of Gata2 using siRNA led to decreased expressions of WT1 and β1-integrin, n = 3 [p].

E-cadherin was validated in adult mouse testis lysates by immunoprecipitation. E-cadherin was pulled down by the antibody against β1-integrin, and vice versa [Fig. 6e], implying that E-cadherin on SPCs binds with β1-integrin in physiological condition. Moreover, E-cadherin siRNA was transfected into Thy1+ SPCs Sertoli cells co-culture system for 48 h, to detect the impact on SPCs fates. Unexpectedly, no obvious difference of growth condition or cell number was observed [Fig. 6f and g]. However, western blot revealed that E-cadherin knockdown reduced expression levels of SPCs markers, including PLZF, ITGB1, GFRA1, and up-regulated expression level of STRA8 [Fig. 6h], indicating that E-cadherin could interfere in SPCs at least at molecular level. Notably, E-cadherin is a specific undifferentiated spermatogonia marker, while β1-integrin is expressed on both spermatogonia and Sertoli cells [48, 49], which is consistent with our results from IF [Fig. 6i and j and Fig. 4e-h] and from western blot [Fig. 6k]. Thus, it’s not clear that E-cadherin on SPCs binds with β1-integrin on Sertoli cells or on SPCs themselves. Therefore, Thy1+ SPCs were used for IP [Fig. 6l], and the result eliminates the possibility that E-cadherin and β1-integrin on SPCs binds on themselves. These results suggest that β1-integrin on Sertoli cell is essential for SPCs maintenance, and probably interacts with E-cadherin on SPCs.

Pharmacologically androgen deprived mice are lack of differentiated germ cells but enriched with hyper-proliferated PLZF+ and β1-integrin+ spermatogonia

To further study the role of androgen on spermatogenesis, 33 6-week old male mice were intraperitoneally injected with bicalutamide, since mice at this age have mature Sertoli cells, stable AR expression and continuous spermatogenesis. Bicalutamide recipient mice exhibited retarded body weight growth and smaller testis size compared to control group [Fig. 7a and b]. Histological results revealed the severe deficiency of differentiated germ cells (spermatocytes, spermatids and sperms) in seminiferous tubules [Fig. 7e]. Moreover, number of AR+ cells [Fig. 7d] and expression level of AR [Fig. 7g and r] were reduced, but number of PLZF+ cells [Fig. 7c] and expression level of PLZF [Fig. 7i and r] were increased in bicalutamide recipient testes compared to
controls [Fig. 7h and j], indicating that androgen deprivation did not impair undifferentiated spermatogonia, but resulted in blockage of germ cell differentiation and expansion of undifferentiated spermatogonia population. Notably, in many seminiferous tubules of bicalutamide recipients, accumulated spermatogonia formed a second layer [Fig. 7k], which were identified as PLZF negative [Fig. 7m], β1-integrin and c-kit positive [Fig. 7n and o], and mitotically active [Fig. 7p] spermatogonia. These results indicated androgen deprivation blocked spermatogenesis at the step that ckit + β1-integrin + spermatogonia differentiate to more differentiated populations, and consequently led to accumulation of undifferentiated spermatogonia, including PLZF + population [Fig. 7s]. Western blot results showed the increased expressions of Gata2, WT1 in bicalutamide recipients [Fig. 7r], which was consistent with results from in vitro. Enhanced expression levels of PLZF, GFRA1 and β1-integrin resulted from accumulated spermatogonia, and decreased expression levels of SYCP3, Claudin-11 and Nectin-2 were the consequence of lost meiotic germ cells caused by androgen deprivation [Fig. 7r]. These results implied that blockage of androgen in adult mouse testis inhibited the differentiation from spermatogonia to spermatocytes, and meanwhile promoted SPCs accumulation.

**Fig. 5** WT1 is an intermediate molecule to transmit androgen signal to downstream target β1-integrin. The representative view of WT1 binding sites (visualized by Integrative Genomics Viewer [http://software.broadinstitute.org/software/igv/]) at β1-integrin promoter region (a). ChIP-qPCR validates the binding of WT1 in the promoter region of β1-integrin; fold enrichment by antibodies against WT1 relative to control IgG was presented as mean value ± SD of two replicates (b). Knockdown of WT1 in Sertoli cells decreased β1-integrin expression, 1, Wt-1 siRNA; 2, scrambled siRNA; 3, untransfected control, n = 5 (c), and transfection of WT1 cDNA enhanced β1-integrin expression, n = 3 (d). Androgen down-regulated expression of WT1 and β1-integrin in co-culture system, n = 4 (e). Knockdown of β1-integrin in the co-culture system suppressed expression level of PLZF, 1, Itgb1 siRNA; 2, scrambled siRNA; 3, untransfected control n = 3 (f). An illustration hypothesizes the potential signal pathway from AR to β1-integrin in Sertoli cells (g).
Fig. 6 (See legend on next page.)
### Discussion

Our results exhibit that SPCs are deficient of endogenous AR after 3 dpp, which is consistent with the dispensable role of AR in differentiated germ cells from AR germ cell conditional knockout mice. Interestingly, some studies detected AR expression in germ cells of fetal testis [50], and we examined weak AR signal in pre-spermatogonia of 2 dpp old testes. Therefore, we prefer the opinion that SPCs do not need endogenous AR for the initiation of spermatogenesis after neonatal stage, and the function of AR in embryonic and new born stages is still need to be revealed.

Testis Zinc Finger Protein (TZFP), a homolog of PLZF, could co-repress the activated AR [51] and participate in hormone signal pathways to regulate spermatogenesis [52]. However, IHC staining showed PLZF and AR staining were distributed in distinct cell types after neonatal stage, indicating no direct interaction between these two transcription factors during spermatogenesis. Based on our data, we hypothesize that transduction of androgen signal in spermatogenesis spatially can be summarized into three steps [Fig. 7t]: the first step is AR activation in Sertoli cells induced by androgen, including androgen-AR interaction and activation of AR’s downstream genes to produce signal molecules; second step is the communication between Sertoli cells and SPCs, involving in some surface or transmembrane proteins; the last one is signal transduction to SPCs nucleus to turn off PLZF and start differentiation. Our results suggest that AR inhibits Gata2 expression in Sertoli cells by binding on its promoter region, leading to a consequent decrease of WT1 expression, and WT1 binds to and regulates β1-integrin in Sertoli cells, which interacts with SPCs as a putative signal molecule to regulate SPCs differentiation. WT1 is able to bind the promoter of Ar to inhibit AR expression [28]. We revealed AR’s inhibitory role on WT1 expression via interacting with Gata2, indicating a feedback regulation on WT1 expression. Moreover, we proposed β1-integrin on Sertoli cells regulates SPCs differentiation. Routinely, β1-integrin combines with another subunit, i.e. α1-integrin, α6-integrin, to form a complex. On SPCs surface β1-integrin binds α6-integrin to form laminin receptor, which is essential for SPCs maintenance. Nevertheless, it’s not clear that α6-integrin binds with β1-integrin in Sertoli cells [53], thus we are interested in searching for the interactive molecule of β1-integrin on Sertoli cells in future study. Moreover, it’s not clear E-cadherin really functions as a signal molecule to regulate SPCs fate in responding to β1-integrin on Sertoli cells, since loss of E-cadherin in SPCs did not impair homing capacity. Our data also exhibited that the growth condition and cell number were unaffected after E-cadherin knockdown (Fig. 6f and g). However, we cannot conclude E-cadherin is redundant for SSCs, since SSCs enrich other types of cadherins which may compensate the role of E-cadherin, and some studies observed that E-cadherin mutation in SSCs led slightly reduced colony formation after transplantation [24]. Notably, our observations revealed the reduced expression levels of E-cadherin do have impact on expression levels of SPCs markers (Fig. 6h). A possibility is that the time for E-cadherin siRNA treatment is not long enough. Although other cadherins in SPCs may compensate the function of E-cadherin, loss of E-cadherin may regulate its own downstream targets or interactive molecules, which will impact on the fates of SPCs. For example, E-cadherin was reported to be functional in transformation of SSCs into pluripotent status [54]. Thus, next we will focus on the link of E-cadherin and SSCs fates, especially regarding β1-integrin’s binding partners on SSCs surface.

Two important junction proteins Claudin-11 and Nectin-2, which were reported to be significantly affected in testes of AR knockout mouse [25], declined in bicalutamide recipient testes. Claudin-11 is a key component of tight junction (TJ) barrier and closely related to spermatogenesis, and its deficiency caused infertility [55]. Ar deletion in Sertoli cells resulted in down-regulation of Claudin-3 (a homolog of Claudin-11 and a component of TJ junction) and a subsequent increase of BTB (blood-testis barrier) permeability [56]. Further study revealed that Claudin-11 replaced Claudin-3 during spermatoocyte translocation in spermatogenesis [57]. Therefore, we inferred that androgen deprivation caused the blockage of
Fig. 7 (See legend on next page.)
spERMATOGONIA differentiation and a consequent accumulation of SPCs. Reduced Claudin-11 expression after androgen deprivation indicated loose of TJ junction. Whether Claudin-11 interacts with β1-integrin is also an interesting question. Down-regulation of Nectin-2 after androgen deprivation from our experiments exhibited contradicting result to that from AR knockout mice [25]. As a anchor junction protein, Nectin-2 is mainly expressed in Sertoli cells and elongated spermatids, and Nectin-2 knockout mice had normal sperm titters but exhibited infertility phenotype due to serve spermatozoa malformation [58], suggesting the major role of Nectin-2 for sperm maturation. Thus, the dramatic decline of Nectin-2 expression in androgen deprivation assay was due to blockage of spermatogonia differentiation into spermatocytes and spermatids.

**Conclusions**

This study reveals androgen signal pathway in Sertoli cells which promotes differentiation of PLZF+ spermatogonia. GATA2, WT1 and β1-integrin were identified as pivotal intermediate molecules in this process. The take-home messages are listed below

1. The expression profile of androgen receptor (AR) in testes of postnatal male mice reveals that spermatogonia express AR before homing to their niche at 2 dpp, but do not express AR since spermatogenesis starts,
2. Sertoli cells are required for spermatogonia differentiation verified by a spermatogonia-Sertoli cells co-culture system.
3. Data from ChIP-seq and gene silence or overexpression assays conclude that AR is able to regulate WT-1 (a key factor for Sertoli cells) indirectly, and identify β1-integrin (a key surface molecule regulating SPCs homing) as a direct target of WT-1 in Sertoli cells,
4. Pharmacologically androgen deprived mice process accumulated undifferentiated spermatogonia and differentiating spermatogonia, but lack spermatocytes and sperm in testes.

In the future, we will focus on the communication pattern between Sertoli cells and SPCs, in particular β1-integrin’s role in regulation of SSCs fates.

**Additional file**

Additional file 1: Table S1, primer information for semi-quantitative and quantitative RT-PCR. (DOCX 28 kb)

**Abbreviations**

AR: Androgen receptor; Coup-tf II: Chicken ovalbumin upstream promoter-transcription factor; DHT: Dihydrotestosterone; dpp: Day post partum; GFRα1: Gial cell line derived neurotrophic factor family receptor alpha 1; ID4: Inhibitor of DNA binding 4; Mvh: Mouse vasa homologue; PLZF: Promyelocytic leukemia zinc-finger; SMC: A-smooth muscle actin; SPCs: Spermatogonia progenitor cells; SSCs: Spermatogonial stem cells; STO cells: SIM mouse embryo-derived thioguanine-and ouabain-resistant cells; STRA8: Stimulated by retinoic acid gene 8; SYCP3: Synaptonemal complex protein 3; TEX14: Testis expressed gene 14; WT1: Wilms tumour 1

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**Availability of data and materials**

The raw data of ChIP-seq assay in this study is available in the EMBL database. (http://www.ebi.ac.uk/arrayexpress/) under accession number E-MTAB-7129.

**Authors’ contributions**

JW Collection and/or assembly of data, data analysis and interpretation. JL Collection and/or assembly of data, data analysis and interpretation. WX Collection and/or assembly of data, data analysis. QX Collection and/or assembly of data, data analysis. YG Collection and/or assembly of data, data analysis. WS Collection and/or assembly of data, data analysis. XZ Collection and/or assembly of data. YY Collection and/or assembly of data. WW Data analysis and interpretation. HL Data analysis and interpretation. KZ
Conception and design, financial support, manuscript writing, final approval of manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate
All of the experimental procedures have been approved by the ethical committee at Nanjing Agricultural University.

Consent for publication
All of the authors agree to submit this manuscript to Cell Communication and Signaling.

Competing interests
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