Claudin 11 deficiency in mice results in loss of the Sertoli cell epithelial phenotype in the testis.
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Claudin 11 Deficiency in Mice Results in Loss of the Sertoli Cell Epithelial Phenotype in the Testis

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INTRODUCTION

The integrity of epithelial cell layers is maintained by intercellular junctional complexes composed of adhesive (adherens junction, desmosomes) and occluding (tight) junctions, and gap junctions promote intercellular communication. This is an Open Access article, freely available through Biology of Reproduction’s Authors’ Choice option.

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202
virus and adenovirus receptor (CAR; official symbol CXADR) [15, 16]. The phenotypes of mice deficient in various components of these TJAs vary from normal (no apparent phenotype), as revealed in $F11r^{-/-}$ mice [17], to slowly degenerative, as for $Occh^{-/-}$ mice [18], to sterility in $Cldn11^{-/-}$ mice [19]. $Cldn11^{-/-}$ mice exhibit neurologic, auditory, and reproductive deficits, including slowed central nervous system nerve conduction, conspicuous hind limb weakness, profound sensorineural deafness, and male sterility [19–21].

In the testis of $Cldn11^{-/-}$ mice, spermatogenesis does not proceed beyond the spermatocyte stage, and cell clusters are observed in the seminiferous lumen. To understand the relationship between claudin 11 loss and seminiferous tubule disorganization, we have determined the etiology of this phenotype, in particular during the period when the BTB forms. Our comprehensive survey reveals that the absence of a mature BTB in Sertoli cells lacking claudin 11 [19] is linked to a spermatogenesis defect in neighboring germ cells. Furthermore, Sertoli cells lose polarity, detach from the basement membrane, undergo an epithelial-to-fibroblastic cell shape transformation, and proliferate while maintaining expression of differentiation markers. These changes are associated with TJ regulation as well as actin-related and cell cycle gene expression.

### MATERIALS AND METHODS

#### Animal Handling, Tissue Collection, and Processing

Claudin 11-null mice [19] were maintained on a mixed genetic background comprising 129SvEv and C57BL/6J strains and have been brother-sister mated for more than 10 generations. The testicular phenotype has been stable in this colony for 10 yr. Males were injected intraperitoneally with 50 mg/kg bromodeoxyuridine (BrdU) dissolved in saline 3 h before killing. Testes and epididymides were collected at 2–8 wk of age in PBS, and depending on the primary antibody used, sections were incubated for 2 h with either horseradish peroxidase-conjugated anti-rabbit antibody (Envision + system-HRP; Dako), biotinylated anti-goat antibody (1:500 dilution; Vector Laboratories Canada, Burlington, ON, Canada), or anti-rat antibody (1:200 dilution; Vector Laboratories), and finally for 30 min with a peroxidase-conjugated streptavidin-horseradish complex (LSAB + Kit; Dako). The reaction product was developed using 3′,3′-diaminobenzidine tetrahydrochloride (DAB; Sigma-Aldrich). Sections were counterstained with hematoxylin and mounted with Eukitt (Sigma-Aldrich). For negative controls, primary antibody was omitted. Slides were analyzed with a Zeiss Axioskop II and Axioshot microscopes (Carl Zeiss, New York, NY) connected to a digital camera (Spot RT Slider; Diagnostic Instruments, Sterling Heights, MI).

### TUNEL Assays

Detection of apoptotic cells was performed on paraffin sections using an in situ cell death detection kit (Roche). After rehydration, sections were boiled in 0.1 M citrate buffer (pH 6.0) and were incubated for 1 h at 37°C with the TUNEL reaction mixture containing terminal transferase to label free 3′-hydroxy ends of genomic DNA with fluorescein-labeled deoxy-UTP. After washing, sections were incubated overnight at 4°C with propidium iodide (Roche). Apoptotic cells were revealed with DAB and sections were counterstained with hematoxylin. Negative controls with the reaction mixture without the enzyme were applied to serial sections and revealed similarly.

### Morphometric Analysis

Apoptotic cells and germ cells in the division phases were quantified in transverse seminiferous tubule sections. Fragmented DNA in nucleus of apoptotic cells was stained with TUNEL, and nuclei of dividing cells were labeled with phosphorylated histone H3. For each animal, at least three nonserial testicular sections were used, and in each section, all transverse section tubules were quantified, for a total of at least 100 tubules per animal (with a mean of 330 and 300 tubules per animal). In addition, the number of phosphorylated histone H3-labeled cells was counted in at least 50 tubules per animal. Sertoli cell nuclei were quantified in transverse seminiferous tubule sections stained for GATA4. For each animal, at least three nonserial testicular sections were used, and in each section, all transverse section tubules were quantified, for a total of at least 50 tubules per animal.

### Microarray Analysis

Total RNA was prepared from P20 testes by using an RNeasy minikit (Qiagen, Courtaboeuf, France). RNA integrity was determined with the Agilent 2100 Bioanalyzer and RNA 6000 Nano Kit (Agilent Technologies, Massy, Mitsubishi Kagaku Institute of Life Sciences, Tokyo, Japan). Cln (TRA-369 antibody; diluted 1:1000; kindly provided by Dr. H. Tanaka, Osaka University, Osaka, Japan); Gat4 (diluted 1:100; Santa Cruz Biotechnologies), phosphorylated serine 10 (ser10) of histone H3 (diluted 1:500; Upstate Biotechnology/ Euromedex, Mundolsheim, France), BrdU (diluted 1:100; Roche), and vimentin (LN-6 clone; diluted 1:100; DakoCytomation, Trappes, France).

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**TABLE 1. List of primer sequences used for RT-qPCR.**

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<th>5′→3′ Reverse primer</th>
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A pool of P20 $Cldn11^{-/-}$ testes originating from three different males was used as a common reference and compared to three testes originating from three different $Cldn11^{-/-}$ males. One microgram of total RNA was amplified with the Amino Allyl MessageAmp II aRNA kit (Ambion, Austin, TX) according to the manufacturer’s instructions. This mRNA amplification procedure is well validated, and it has been demonstrated that it does not distort the relative abundance of individual mRNAs within a RNA population [22].

Fluorescent probes were synthesized by chemical coupling of 5 μg of aminoaallyl aRNA with cyanine 3 (Cy3) or Cy5 dyes (GE Healthcare Biosciences, Orsay, France). After purification with an RNeasy Mini Kit (Qiagen), probes were fragmented with 25× RNA Fragmentation Reagents (Agilent Technologies) and hybridized with 2× Agilent Hybridization Buffer (Agilent Technologies) to Mouse opArray (Operon Biotechnologies GmbH, Cologne, Germany) in an Agilent oven at 67°C for 16 h, following a dye swap experimental procedure to correct for gene-specific dye bias [23]. Microarrays were washed and scanned with a GenePix 4000B scanner (Molecular Devices).

FIG. 1. Chronology of the testis phenotype in $Cldn11^{-/-}$ mice. Testicular sections of P13 (A–C), P20 (D–F), P28 (G–I), and P60 (J and K) from control (A, D, G, J, C, F, and L) and $Cldn11^{-/-}$ (B, E, H, and I) mice were processed for PAS histological staining (A, D, G, J, C, F, and L) and $Cldn11^{-/-}$ (B, E, H, and I) claudin 11 immunohistochemistry (C, F, and I). L) Control (C; black) and $Cldn11^{-/-}$ (KO; white) testis weight (n = 3). Arrows point to the round cell clusters. Asterisks indicate statistical significance ($P < 0.001$) by one-way ANOVA followed by Tukey posthoc testing. Bars = 100 μm.

Quantitative PCR

Quantitative PCRs were done on RNA from the same three $Cldn11^{-/-}$ and three $Cldn11^{-/-}$ testes, assessed individually. First-strand cDNAs were synthesized from 1 μg of total RNA in the presence of 100 units of Superscript II (Invitrogen, Egany, France) and a mixture of random hexamers and oligo(dT) primers (Promega, Charbonnieres, France). Real-time PCR assays were performed in duplicates for each sample with a Rotor-Gene 6000 (Corbett Research, Mortlake, Australia). The PCR primers are listed in Table 1.
RESULTS

Chronology of the Cldn11−/− Phenotype

To characterize the beginning of the testicular phenotype in Cldn11−/− mice, we first analyzed testis histological sections throughout postnatal testicular development (Fig. 1). During normal testis development, at 2 wk of age, seminiferous tubules contained Sertoli cells and early pachytene spermatocytes, which are the most mature cells of the germ cell lineage. One week later, early spermatids largely populated the tubules, and differentiating elongated spermatids were first detected at P28. At P60, when the animals are adults, spermatogenesis was cyclic and could be divided into 12 stages (I–XII) based on the morphological transformation of spermatids into spermatozoa in a process referred as spermiogenesis [26].

The first signs of disorganization in Cldn11−/− testes appeared at P20 (Fig. 1, A, B, D, and E) and were obvious by P28. Indeed, when compared to the well-organized epithelium in the first wave of spermatogenesis from control testes (Fig. 1G), germ cells in Cldn11−/− testes appear abnormally localized. Round clusters of cells were observed closely apposed to the base of the seminiferous epithelium, and the testis tubular lumen was poorly defined and was filled with round spermatids, with the acrosome labeled with PAS or cell clusters (Fig. 1H, arrows). Moreover, although elongated spermatids appeared in some tubules of P28 control testes (Fig. 1G), only scarce ectopically localized round spermatids were present in Cldn11−/− testes (Fig. 1H). In addition, elongated spermatids were never observed in Cldn11−/− testes at P28 or P60, and spermatogenesis did not proceed through the complete process of germ cell differentiation, which is consistent with published findings [19]. The phenotype was accentuated at P60 (Fig. 1, J and K), and PAS-positive material was observed inside the cell clusters (Fig. 1K, arrows), indicative of the presence of glycoproteins.

We next compared the temporal appearance of the Cldn11−/− testis phenotype with that of protein expression in control testes (Fig. 1, C, F, and I). Although claudin 11 was not detected by immunohistochemistry at P10 (data not shown), it was obvious in some tubules at P13 in testes from wild-type animals, with labeling extending from the basal membrane to the lumen of Cldn11−/− testes (Fig. 1C). Such staining is consistent with the localization of claudin 11 to plasma membrane and the polarization of Sertoli cells from P20 onward (Fig. 1, F and I). In time-matched Cldn11−/− testes, no labeling was observed, thus confirming the specificity of the antibody (Supplemental Fig. S1; all Supplemental Data are available online at www.biolreprod.org). Testis weight in Cldn11−/− mice was normal up to P28 (Fig. 1L).

Cell Clusters Comprise Sertoli Cells

Histological sections reveal that nuclei from the vast majority of cells in adluminal cell clusters are clear and contain one to three nucleoli, which is reminiscent of normal Sertoli cell morphology (data not shown). However, in P60 Cldn11−/− testes, mixed cell nuclei are observed in the clusters (data not shown and Gow et al. [19]). To identify the cells comprising these clusters, we performed immunohistochemis-

try for several germ cell markers (Fig. 2): the DDX4 protein (mouse vasa homolog) is expressed from zygotene spermatocytes to round spermatids (Fig. 2A) [27–29]; Calmegin (CLGN) is expressed in early pachytene spermatocytes to step 14 spermatids (Fig. 2D) [30, 31]; and vitronectin [32] and claudin 1 (data not shown) are expressed in the acrosomes of spermatids. The absence of these markers demonstrates that cell clusters in Cldn11−/− seminiferous tubules do not contain germ cells (Fig. 2, B, C, E, F, and I), and in P60–P180 mice (data not shown), immunolabeling of GATA4, which identifies Sertoli cells inside the seminiferous epithelium (Fig. 2G) [33, 34], clearly indicated that clusters comprise Sertoli cells (Fig. 2, H and I).

Increased Apoptosis in Testis of Cldn11−/− Mice

Although claudin 11 expression in testis is Sertoli cell specific, the primary defect in Cldn11−/− mice is that of a failure of spermatogenesis. Apoptosis is the dominant pathway for eliminating germ cells whenever the supporting Sertoli cells are unable to provide a supportive environment for their development [35]; thus, we characterized the fate of germ cells by using TUNEL labeling (Fig. 3). Quantification of TUNEL-labeled cells in Cldn11−/− testis showed a significant (P < 0.05) increase in the incidence of apoptosis at P20 and P28 compared with age-matched controls (Fig. 3). The proportion of round tubules containing one or more TUNEL-positive cells was increased 1.5-fold (P < 0.05) in Cldn11−/− testes at P15 (Fig. 3E), suggesting an early impact of the absence of claudin 11 on germ cell development. Apoptosis reached a peak at P20 and was maintained through P28, but it remained elevated at
P60 and was not statistically different from levels at P20. In addition, abundant vesicles of fragmented TUNEL-positive DNA were detected in the cytoplasm of sloughing Sertoli cells (arrows in C) and absent in control TUNEL analysis (D). In E, counts of round seminiferous tubules containing at least one TUNEL-positive cell. Data show that first significant increase was seen at P15. Values are mean ± SEM of three to six animals. Asterisks indicate statistical significance ($P < 0.001$) by one-way ANOVA followed by Holm-Sidak posthoc testing. Bars = 100 μm (A and B) and 50 μm (C and D).

FIG. 3 High incidence of germ cell apoptosis in Cldn11−/− testes. TUNEL analysis of control (A) and Cldn11−/− (B and D) testes at P20 (A and B) shows increased germ cell apoptosis in Cldn11−/− testes. In addition to TUNEL-positive germ cells localized in the seminiferous epithelium, small vesicles of fragmented TUNEL-positive DNA reminiscent of phagocytosis are abundant in cytoplasm of sloughing Sertoli cells (arrows in C) and absent in control TUNEL analysis (D). In E, counts of round seminiferous tubules containing at least one TUNEL-positive cell. Data show that first significant increase was seen at P15. Values are mean ± SEM of three to six animals. Asterisks indicate statistical significance ($P < 0.001$) by one-way ANOVA followed by Holm-Sidak posthoc testing. Bars = 100 μm (A and B) and 50 μm (C and D).

Dynamics of Sertoli Cell Sloughing in Cldn11−/− Testes

To determine how a lack of claudin 11 might have an impact on Sertoli cell topography within the seminiferous epithelium, sections were immunolabeled with GATA4 (Fig. 5). In control testes, Sertoli cell nuclei were positioned at the basement membrane of the seminiferous epithelium, as expected. In Cldn11−/− testes, the nuclei were localized toward the center of the tubule as early as P13 (Fig. 5, A and B), which coincided with the beginning of claudin 11 expression in control testes (Fig. 1C). At P28 (Fig. 5, C–F)—also seen at P20 (data not shown)—basally located Sertoli cells were observed, as were several abnormal Sertoli cell arrangements. These include groupings of Sertoli cells adjacent to the base of the tubule (Fig. 5D), round clusters with few cells attached to the basement membrane (Fig. 5E), and completely detached Sertoli cell clusters filling the tubule lumen (Fig. 5F). These data are probably indicative of dynamic Sertoli cell sloughing from their basal sites into the lumen of the tubule. Consistent with our proposal that Sertoli cells are sloughed from the epithelium into the duct system is our observation that GATA4-positive cell clusters occur in the lumen of the epididymis of P60 Cldn11−/− mice (Fig. 5G).

In this light, we hypothesize that Sertoli cells may migrate along the basement membrane to form small groups, which
then detach, or they may form clusters by cell division and thereafter detach. To distinguish between these possibilities, we examined changes in the number of Gata4-labeled Sertoli cells from P15 to P60 (Fig. 6, A–C). Consistent with the temporal development of the phenotype, the number of Sertoli cells per cluster rose from P15 to P28 and was maintained at this level through P60 in \( \text{Cldn11}^{-/-} \) testes (Fig. 6A). In addition, the proportion of round tubules containing detached Sertoli cells almost doubled between P15 and P60 (\( P, 0.05 \)). Indeed, more than 80% of the tubules contained Sertoli cell clusters at P60 (Fig. 6B).

Although the number of Sertoli cells per round tubule decreased from P15 to P60 in control testes (probably as a result of dilution due to a massive increase of the germ cell population within the tubules as the animals mature), the size of this population remained almost constant in \( \text{Cldn11}^{-/-} \) testes (Fig. 6C). The number of Sertoli cells per round tubule in control and \( \text{Cldn11}^{-/-} \) testes was similar at P15 and at P28 if both peripherally located and cluster-located Sertoli cells are considered. At P60, the number of peripherally located Sertoli cells was similar in control and \( \text{Cldn11}^{-/-} \) testes but higher in \( \text{Cldn11}^{-/-} \) testes when considering total Sertoli cell numbers. Together, these data suggest a continuous renewal of Sertoli cells to compensate for their losses from detachment and sloughing into the lumen.

**Sertoli Cells Undergo Cell Division in \( \text{Cldn11}^{-/-} \) Testes**

Sertoli cells cease dividing in immature animals during the first 2 wk after birth, which is concomitant with the appearance of meiotic germ cells and formation of the lumen [37, 38]. To determine the cell cycle status of Sertoli cells, we performed an
in-depth survey of GATA4-labeled Sertoli cells (Fig. 6, D–F). In control testes, GATA4 labeling of Sertoli cell nuclei was homogenous, with the exception of nucleoli (Fig. 6D, arrows), and similarly labeled Sertoli cells were found in P28 (Fig. 6, E and F) and P60 (data not shown) Cldn11−/− testes. In addition, heterogeneous chromatin in GATA4-positive nuclei was also observed (Fig. 6, E and F, arrowheads), together with the characteristic meiotic figures of germ cells (Fig. 6E, arrows).

Juxtaposed Sertoli cells in the plane parallel to the basement membrane in Cldn11−/− testes (Fig. 6E) suggest that this cell population may be dividing. To identify dividing Sertoli cells, we used in vivo BrdU incorporation prior to the fixation and tissue processing (Fig. 6, G–J). Comparison of the Gata4-positive Sertoli cell and BrdU-positive dividing cell populations in wild type testes reveals the canonical complementary pattern of Sertoli cell and spermatogonia (Fig. 6G). This is a hallmark of stage VIII tubules, wherein spermatogonia proliferate and leptotene spermatocytes traverse the BTB into the adluminal compartment for further development [12, 27, 29, 32]. In Cldn11−/− testes, stage VIII tubules contained Gata4-positive Sertoli cells (Fig. 6, H–J, arrowheads) and BrdU-labeled spermatogonia (Fig. 6, H and I, solid arrows), indicating that the lack of claudin 11 expression does not hamper the spermatogenic cycle, or at least the cyclical entry of spermatogonia into mitosis at stage VIII [26]. However, we also observed double-labeled cells at the periphery of some tubules (Fig. 6, H and J, open arrows), which demonstrates that Sertoli cells are proliferating in Cldn11−/− testes. These double-labeled Sertoli cells, however, were found at a frequency of one to three per testis section, mostly grouped in one tubule, suggesting that the rate of Sertoli cell proliferation is probably low. Importantly, we did not observe BrdU labeling of cell clusters. It may indicate that Sertoli cell division occurs prior to cluster formation and shedding into the lumen.

*Loss of Sertoli Cell Polarity but Maintenance of Differentiation Markers*

In Figure 5, we observe that Sertoli cells in Cldn11−/− testes changed shape in the course of cell cluster sloughing. Thus, the nuclei of basal Gata4-labeled Sertoli cells were round or triangular, whereas Sertoli cells located at the periphery of clusters had round nuclei, and those within clusters had smaller and elongated or comma-shaped nuclei. Such shapes are reminiscent of fibroblasts (Fig. 7A) and indicate a loss of cell polarity, and possibly differentiation status. To examine these possibilities, we analyzed several Sertoli cell differentiation markers. Vimentin, androgen receptor, and N-cadherin labeling (Fig. 7, B–E, H, and I; and data not shown) demonstrated that Cldn11−/− Sertoli cells retain mature markers, even in luminal cell clusters from P180 testes when the number of germ cells and, consequently, testis weight is in decline. We also observed vimentin-positive Sertoli cell clusters in the epididymides of Cldn11−/− mice (Fig. 7, F and G), probably suggesting that they are cleared from the testes in a fashion similar to sperm cells. Consistent with the absence of sperm, the lumen of most epididymides were empty (Fig. 7G).
contrast, epididymides from control animals were filled with sperm cells (Fig. 7F).

**Molecular Characterization of Cldn11−/− Testes Using Microarrays**

To better understand the molecular consequences of the absence of claudin 11 on Sertoli cells, we performed a microarray analysis of whole-testis mRNA from P20 Cldn11−/− mice. Littermate Cldn11+/+ mice were used as controls, which are fertile and otherwise indistinguishable from wild-type mice in all studies we have performed. The choice of P20 stems from two main considerations; testis weight and germ cell composition are similar in Cldn11−/− and control groups, and Sertoli cells detached from the basement membrane can be observed. The strategy was based on a subtractive approach following a dye swap experimental procedure to catch the most differentially upregulated and downregulated genes. Microarray analysis identified 108 genes significantly upregulated and 98 downregulated by more than 1.3-fold in Cldn11−/− compared with Cldn11+/+ testes (SAM procedure with a false discovery rate <5%). Importantly, Cldn11 was the most downregulated gene (Table 2), which accords with the genotype of Cldn11−/− mice. In addition, we selected six upregulated genes and one downregulated gene on the basis of the amount of expression and their possible biological significance and confirmed differences in the expression levels by quantitative PCR analysis (Table 2).

In agreement with the histological data (Fig. 1, D and E), among the upregulated genes, 22 (20.4%) were inferred to Sertoli cells and four (3.7%) to germ cells; among the downregulated genes, 28 (29.6%) were inferred to germ cells, most often spermatocytes (Supplemental Table S1). Also consistent with the immunohistochemical data, the Vim gene was already upregulated in P20 Cldn11−/− testes. Interestingly, typical Sertoli cell markers, such as ETV5 (also known as ERM), SOX8, SOX9, Desert hedgehog, transferrin, GJA1 (connexin 43), or Kit ligand, displayed a similar expression in control and Cldn11−/− testes (data not shown). Notably, WebGestalt analysis of microarray data revealed a statistical enrichment of TJ (P = 1.36 × 10−8), regulation of actin cytoskeleton (P = 2.72 × 10−3), cell cycle (P = 1.14 × 10−4), glutathione metabolism (P = 2.75 × 10−3), and adherens junction (P = 1.84 × 10−3) KEGG pathways. Interestingly, expression of genes of the tight and adherens junctions and actin and intermediate filament cytoskeleton pathways were coordinately upregulated (16 of 18 genes and 14 of 15 genes, respectively).

**DISCUSSION**

The present data highlight the critical role of claudin 11 in the maintenance of Sertoli cell epithelial differentiation. Indeed, in the absence of claudin 11, Sertoli cells differentiate, can proliferate, and are not permanently arrested as nonmitotic quiescent cells. Thus, Sertoli cells can exhibit one feature of an immature cell without reverting to the state of immaturity. They form homogeneous clusters, lose their contact with basement membrane, detach from the seminiferous epithelium, and acquire a fibroblastlike cell shape. At the molecular level, the absence of claudin 11 expression is associated with selective changes in several cell junction-related genes, specifically TJ genes, as well as cytoskeleton-related genes.

A growing number of genes associated with male infertility have been generated using homologous recombination in embryonic stem cells [39, 40]. A frequent feature of mouse models of infertility is germ cell defect consecutive to either depletion of stem cells, meiosis arrest, or spermiogenesis default. Disruption of meiosis notably induces massive germ cell apoptosis and elimination by their phagocytosis by Sertoli cells, resulting in such histological characteristics as vacuoles in Sertoli cell cytoplasm and multinucleated spermatids. As well, the primary impact of somatic cell physiology disruption (very often through impairment of hormonal action) is germ cell development damage. By comparison, ablation of genes encoding TJ proteins, such as occludin (Ocln), ZO-1 (Tjp1),
and F11r, displays variable phenotypes, ranging from Sertoli cell-only syndrome of the old adult to the apparent absence of a testicular phenotype [15, 18, 41]. Together, our results demonstrate that claudin 11 plays a key role in both Sertoli cell physiology and spermatogenesis.

Sertoli cell clusters have been described previously in several mouse models of infertility consecutive to germ cell loss [42, 43]. Among them, very few involve a primary defect in the Sertoli cell maturation process (i.e., proliferation and differentiation), and none of them result in the uncoupling between these two states. For example, delayed Sertoli cell differentiation), and none of them result in the uncoupling

TABLE 2. List of selected up- and down-regulated genes in P20 Cldn11−/− testes.

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<td>NM_011733</td>
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<td>Actin or intermediate filament cytoskeleton</td>
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<td>Cytosolic FMR1 interacting protein 1 (Sra1)</td>
<td>Cygp1</td>
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<td>LIM and SH3 protein 1</td>
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<td>NM_018688</td>
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<td>+1.59*</td>
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<td>Act7b</td>
<td>NM_025271</td>
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* Statistical significance (p < 0.005) in P20 Cldn11−/− testes versus time-matched wild-type testes.
Sertoli cell number in proliferate, albeit at a low proliferation rate. Nevertheless, BrdU at P28 and P60, thereby demonstrating that they can incorporate some Sertoli cells lacking claudin 11 in vivo. In the current study, we observe that be arrested proliferative cells rather than terminally differentiated somatic cells [59, 60]. In vitro, recent studies show that P7–P8 and adult mouse Sertoli cells cease to divide, which suggests that BTB formation could be of several mature Sertoli cell markers, including Gata1 and Fshr [65], at P20. Therefore, we find no evidence that Sertoli cells are committed to a dedifferentiation process in the absence of TJs.

Alternatively, Sertoli cells could become committed to a preneoplastic process in the absence of TJs. In addition to Sertoli cell proliferation, Cldn11−/− testes express high levels of Mtdh, which is a known tumor cell marker [66]. However, dividing Sertoli cells are scarce and only observed at the periphery of tubules, and not in luminal cell clusters, which would indicate that these cells become quiescent before they slough. Moreover, instead of invading the entire testis, Sertoli cell clusters are cleared via the normal conducts and are found in the epididymis. At 6 mo, testis histology is not significantly different from that at 2 mo, but it is dramatically different from typical Sertoli cell tumors [67, 68]. Finally, testicular tumors have not been observed in our mouse colony in more than 10 yr, despite maintaining mice beyond 12 mo of age (Gow, unpublished results). Although several studies have demonstrated the induction or suppression of various claudin genes in different cancers, in support of a relationship between TJ-based barrier function and cell proliferation [51–53], and our data strongly suggest that the absence of claudin 11 in Sertoli cells does not commit these cells to a neoplastic transformation. In addition, although more than one claudin family member has been shown at the BTB [69], we demonstrate a lack of functional redundancy at the BTB.

It is interesting that Clu was the most highly stimulated gene in our microarray study. Indeed, its encoded protein is known to cause aggregation of Sertoli cells from immature rats and TM-4 cells from mouse testis [70]. Aggregation of these cells is the first sign of fetal testis differentiation [71], but this behavior is probably independent of claudin 11 expression, because testis development and morphology at P10 are normal in Cldn11−/− mice. Although mechanisms regulating homophilic recognition and cell aggregation in Sertoli cells are unclear, changes in expression of intercellular junction and cytoskeletal genes from our microarrays suggest a role for claudin 11 in intracellular signaling, including a feedback loop for the cell to compensate for excessive TJ permeability. Regulation of adherens junction proteins, such as N-cadherin (data not shown), may be sufficient for Sertoli cell aggregation, with claudin 11 mediating polarization. In addition, induction of the actin and intermediate filament genes may be related to the Sertoli cell shape changes and their sloughing into the lumen.

Among mouse models of male infertility, Cldn11−/− mice display a unique testis phenotype in which the primary defect is epithelial disorganization, mitosis, and detachment of Sertoli cells from the basement membrane in the face of adult differentiation marker expression. Components of TJs directly influence signaling pathways and regulate epithelial differentiation and proliferation [72, 73]. To further dissect the role of claudin 11 at TJs between Sertoli cells, it would be of great interest to track the incidence of Cldn11 knockdown on purified Sertoli cells, both at the phenotypic and transcriptome levels. Then, it would be possible to identify the genes that

Conceivably, the induction of these genes may be related to a germ cell apoptosis-related shift in the composition of the epithelium; however, several canonical Sertoli cell markers are unchanged, which argues against a significant change in the Sertoli cell:germ cell ratio. Although dedifferentiation of Sertoli cells might be an expected outcome of persistent proliferation and reorganization of the vimentin network, our microarray data show that these cells maintain the expression of several mature Sertoli cell markers, including Gata1 and Fshr [65], at P20. Therefore, we find no evidence that Sertoli cells are committed to a dedifferentiation process in the absence of TJs.

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show the highest degree of upregulation or downregulation, and the putative canonical pathways that might be affected by claudin 11.

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