Tumor suppressor gene Rb is required for self-renewal of spermatogonial stem cells in mice

The MIT Faculty has made this article openly available. Please share how this access benefits you. Your story matters.

<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>As Published</td>
<td><a href="http://dx.doi.org/10.1073/pnas.1311548110">http://dx.doi.org/10.1073/pnas.1311548110</a></td>
</tr>
<tr>
<td>Publisher</td>
<td>National Academy of Sciences (U.S.)</td>
</tr>
<tr>
<td>Version</td>
<td>Final published version</td>
</tr>
<tr>
<td>Accessed</td>
<td>Tue Mar 19 03:41:28 EDT 2019</td>
</tr>
<tr>
<td>Citable Link</td>
<td><a href="http://hdl.handle.net/1721.1/85906">http://hdl.handle.net/1721.1/85906</a></td>
</tr>
<tr>
<td>Terms of Use</td>
<td>Article is made available in accordance with the publisher’s policy and may be subject to US copyright law. Please refer to the publisher’s site for terms of use.</td>
</tr>
<tr>
<td>Detailed Terms</td>
<td></td>
</tr>
</tbody>
</table>
The retinoblastoma tumor suppressor gene \( Rb \) encodes the multifunctional protein RB, which actively controls multiple cellular processes, including cell-cycle progression and differentiation (1–3). Somatic stem cells, which possess the ability to self-renew and differentiate into specialized cells, are critical for maintaining tissue homeostasis as well as for repairing tissues after injury. It has been shown that somatic stem cells are largely held in quiescence through a process that involves \( Rb \) (3, 4). Mice that are conditionally deficient in \( Rb \) or \( Rb \) gene family members in the stem cell compartments of somatic tissues—such as blood (5), liver (6), muscle (7), and skin (8)—show a common phenotype: stem cells exit quiescence and proliferate. However, this defect does not appear to affect the stem cells’ self-renewal capacity (5, 9).

In addition to maintaining the quiescence of somatic stem cells, \( Rb \) also plays critical roles in their differentiation (see review in ref. 3). Somatic stem/progenitor cells that lack RB are unable to undergo terminal differentiation, reflecting RB’s function in controlling somatic cell fate through modulating the transcriptional activity of master differentiation regulators (10–12). Cell death resulting from \( Rb \) deficiency also contributes to the absence of terminally differentiated somatic cells (5, 13). Moreover, \( Rb \) deficiency in stem/progenitor cells can lead to tumor formation in somatic tissues (6, 13, 14).

\( Rb \)’s important function in somatic stem cells raises the question of whether it plays a similar role in the regulation of germ-line stem cells. However, this question has remained unexplored. Male germ-line stem cells, which are located in the testis, are also known as spermatogonial stem cells (SSCs). As with somatic stem cells, SSCs must undergo both self-renewal to sustain the stem cell pool and differentiation to give rise to terminally differentiated cells: spermatozoa (sperm). Spermatogenesis follows a differentiation scheme similar to that for somatic cell lineages. SSCs undergo mitotic divisions to generate progenitor (transit-amplifying) spermatogonia, followed by a series of differentiation events, including meiosis and spermiogenesis, to form highly specialized sperm cells. Despite the similarities between somatic stem cells and SSCs, they differ fundamentally in their cell-cycle status. Somatic stem cells are largely quiescent (for example, ~95% of adult hematopoietic stem cells in bone marrow), whereas SSCs are actively cycling throughout an animal’s reproductive life (15–17). This difference poses interesting questions about how RB functions in SSCs and how this function compares with its role in somatic stem cells. We therefore decided to explore RB function in the germ line throughout the various stages of spermatogenesis.

Spermatogenesis normally begins from single, isolated germ cells called \( A_{single} \) or \( A_{s} \) spermatogonia, the population of which is thought to be, or at least contain, SSCs (18–23). \( A_{s} \) spermatogonia divide, with incomplete cytokinesis, to form chains of 2 (\( A_{paired} \) or \( A_{pa} \)) and then 4, 8, 16, or even 32 (\( A_{aligned} \) or \( A_{al} \)) spermatogonia. \( A_{pa} \) spermatogonia then differentiate into spermatogonia in a wave-like manner (once every 8.6 d), moving synchronously through several phases: differentiating spermatogonia, spermatocytes (meiotic), and spermatids (postmeiotic) (24). In the mouse, the synchronized passage of spermatogonic cells through these phases results in 12 recognizable associations, known as seminiferous epithelial stages 1–XII (24).

Hereafter we use the term \( A_{pa} \) spermatogonia to refer, collectively, to \( A_{s} \), \( A_{pa} \), and \( A_{al} \) spermatogonia (23). \( A_{al} \) spermatogonia consist of both SSCs and progenitor spermatogonia, express specific protein markers [for example, promyelocytic leukemia zinc finger (PLZF)], and maintain the potential to self-renew or revert to self-renewing cells (21, 25, 26).

To explore the function of RB in the mouse germ line, we used a Cre recombinase-\(loxP \) conditional knockout strategy to remove \( Rb \) in germ cells before birth. Loss of \( Rb \) in germ cells resulted in rapid exhaustion of the SSC pool. Specifically, GFRa1-protein-expressing \( A_{single} \) (GFRa1\(^{+} \) \( A_{s} \)) spermatogonia, which are thought to be the major source of SSCs (19–21, 27), were defective in self-renewal, and production of germ cells in the testis was quickly depleted. Thus, our study indicates that \( Rb \) is required for stem cell maintenance in the male germ line.

The authors declare no conflict of interest.

Author contributions: Y.-C.H., D.G.d.R., and D.C.P. designed research; Y.-C.H. and D.G.d.R. performed research; Y.-C.H. and D.G.d.R. analyzed data; and Y.-C.H. and D.C.P. wrote the paper.

The authors declare no conflict of interest.

1To whom correspondence should be addressed: E-mail: dcpage@wi.mit.edu.

This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1311548110/-/DCSupplemental.
Results and Discussion

Germ Cell Expression of RB Protein Ends Before Birth in Conditional Knockout Males. We crossed Rb<sup>fl/fl</sup> conditional mutant mice (28) with mice carrying a newly generated mouse vasa homolog (Mvh<sup>Cre-mOrange<sup>+</sup></sup>; also known as Ddx4) allele (Fig. S1) in which the endogenous Mvh promoter drives expression of a Cre-mOrange fusion protein. The Mvh<sup>Cre-mOrange<sup>+</sup></sup> line showed Cre recombination activity in the germ line starting from ~embryonic day (E) 15.5, as assessed by crossing into the ROSA26-LacZ reporter strain (29) (Fig. S1F). To assay whether RB expression was subsequently lost in germ cells of homozygous Rb<sup>fl/fl</sup>; Mvh<sup>Cre-mOrange<sup>+</sup></sup> male mice (hereafter referred to as Rb cKO), we examined RB expression in testes. We found that, at postnatal day 0 (P0), RB was expressed in control (Rb<sup>fl/+</sup>) but not in Rb cKO germ cells (Fig. 1). Therefore, RB expression ceases in Rb cKO germ cells before birth.

Sterility by 2 mo of Age. Mature Rb cKO mice were grossly normal except that the testes were small (Fig. 2A). During the first 3–4 wk of age, testis weight in Rb cKO animals increased much like controls (including Rb<sup>fl/+</sup> and Rb<sup>fl/fl</sup>; Mvh<sup>Cre-mOrange<sup>+</sup></sup>), but testis weight declined thereafter and remained low throughout adulthood (Fig. 2A). To assess the fertility of Rb cKO mice, each male was housed with a single wild-type female for 21 wk, beginning at 4 wk of age. Each control male sired four to six litters throughout the test period. By contrast, each Rb cKO male sired only one litter, born when the males were 9–11 wk of age (Fig. 2B); impregnation would have occurred 3 wk earlier, when the males were 6–8 wk of age. The Rb deletion allele was transmitted at normal Mendelian ratios. The average size of litters sired by Rb cKO males was modestly but not significantly reduced compared with controls (6.4 ± 2.9 vs. 8.1 ± 1.4 pups, respectively; P > 0.05). Taken together, these data suggest that RB-deficient germ cells can develop into spermatozoan and produce offspring, but Rb cKO males become sterile by about 2 mo of age.

Limited Rounds of Spermatogenesis. As wild-type males develop, their germ cells undergo a G1/G0 cycle arrest from E14.5 until 1–2 d after birth (30). In the experiments reported here, RB was ablated in the germ cells while they were arrested, but we observed no difference in germ cell numbers between control and Rb cKO testes at P0 (Fig. 3A). We also observed that both control and Rb cKO germ cells remained quiescent at P0, as determined by BrdU incorporation assay (>95% germ cells negative for BrdU in both groups). These results suggest that RB is not required to maintain the G1/G0 arrest, consistent with a previous report (31).

To seek explanations for the loss of fertility in early adulthood in Rb cKO males, we examined the testicular histology of these mice from P7 to 2 mo of age. To ensure consistency in our analysis, we focused on seminiferous tubule cross-sections at stage VII in both control and Rb cKO testes (except at P7, when seminiferous tubule staging is not possible). By P28, Rb cKO tubules displayed a striking depletion of “early” (less advanced differentiating) spermatogenic cell types (Fig. S2B). In P28 control tubules, we observed four types of germ cells (spermatagonia, preleptotene spermatocytes, pachytene spermatocytes, and round spermatids), corresponding to four rounds of spermatogenesis. By contrast, many Rb cKO tubules contained only the more advanced differentiating cell types (pachytene spermatocytes and round spermatids) and lacked the earlier cell types (spermatagonia and preleptotene spermatocytes). In quantitative terms, only 28% of Rb cKO stage VII tubule cross-sections contained spermatagonia, compared with 91% of controls (Fig. 3C). Consistent with these findings, the number of preleptotene spermatocytes per 1,000 Sertoli cells was dramatically reduced in Rb cKO testes by P40, compared with controls (Fig. S3). By 2 mo of age, most tubules in Rb cKO testes were devoid of germ cells, and the few germ cells that remained were mostly elongated spermatids (highly advanced differentiating cells; Fig. 3B, Inset). These results demonstrate that Rb cKO testes initiate a very limited number of rounds of spermatogenesis, but that these limited rounds are histologically like those observed in controls. These findings suggest that the principal defect in Rb cKO testes is premature exhaustion of the SSC pool.

Transient Increase in Number of Spermatagonia. Next, we examined the kinetics of germ cell depletion and were surprised to find that, in Rb cKO tubules, germ cell depletion (resulting from SSC exhaustion) is preceded by a transient increase in the number
of spermatogonia (particularly progenitor spermatogonia). We discovered this by examining the number of spermatogonia per 1,000 Sertoli cells in stage VII tubule cross-sections, where nearly all spermatogonia are progenitor spermatogonia (32). In control tubules, the number of spermatogonia was relatively constant across the time points examined (Fig. 3D). In contrast, the number of spermatogonia in Rb cKO testes was markedly increased at P10 and P14, compared with controls, followed by the previously described reductions at P28 and P40 (Fig. 3D). We postulated that this transient increase in the number of spermatogonia in Rb cKO testes might reflect a second role for Rb (in addition to that of maintaining SSCs). Specifically, Rb might limit proliferation of progenitor spermatogonia. If so, we might expect that, in Rb cKO testes, progenitor spermatogonia would continue to overproliferate at P28 and P40, even as the total numbers of spermatogonia declined because of SSC exhaustion. To test this prediction, we measured the number of spermatogonia per 1,000 Sertoli cells in the stage VII tubule cross-sections that still retained spermatogonia at P28 and P40. We observed that the number of spermatogonia per 1,000 Sertoli cells was increased in these tubule cross-sections of Rb cKO testes compared with their counterparts in controls (Fig. S4). This result corroborates the hypothesis that Rb restricts the number of progenitor spermatogonia. Taken together, the histological analyses allow us to identify two germ cell phenotypes in Rb cKO testes: premature SSC exhaustion and increased expansion of progenitor spermatogonia.

**Progressive Loss of GFRα1+ A<sub>c</sub> Spermatogonia with Proliferation Unaffected.** To substantiate the finding that SSCs are prematurely exhausted in Rb cKO testes, we measured the abundance of GFRα1<sup>+</sup> A<sub>c</sub> spermatogonia between P5 and P28 by staining whole-mount tubules for PLZF and GFRα1. PLZF is expressed in all A<sub>spa</sub> spermatogonia, whereas GFRα1 is expressed mainly in A<sub>c</sub> and A<sub>ger</sub> spermatogonia (21, 33, 34). To ensure that Rb deficiency does not affect the GFRα1 expression pattern, we examined PLZF<sup>+</sup> A<sub>c</sub> spermatogonia in control and Rb cKO testes.
spermatogonia that retained BrdU label at time of testis recovery. 

GFRα1 spermatogonia in SD, testes at P10 (mean +/−). Consistently, we observed the absence of BrdU+ GFRα1+ spermatogonia in Rb cKO testes when mice were injected with BrdU at P10/P11 and killed 3–5 d later (Fig. 5B). Accordingly, we surmise that upon cell division, Rb-deficient GFRα1+ spermatogonia do not self-renew to form new GFRα1+ spermatogonia; each GFRα1+ cell can therefore initiate only one round of spermatogenesis.

Extended Proliferation of Progenitor Spermatogonia. The histological analysis revealed that progenitor (transit-amplifying) spermatogonia might overproliferate in Rb cKO testes (Fig. 3 and Fig. S4). To better understand this phenomenon, we closely

that Rb deficiency does not affect the proliferative activity of GFRα1+ A spermatogonia.

To determine whether increased cell death contributed to the progressive loss of GFRα1+ A spermatogonia in Rb cKO testes, we assayed cell apoptosis by staining whole-mount tubules for GFRα1 and active caspase-3. We found no GFRα1+ A spermatogonia that were positive for active caspase-3 in either control or Rb cKO testes at P10 (>45 GFRα1+ A spermatogonia examined for each genotype). These results suggest that the loss of GFRα1+ A spermatogonia in Rb cKO testes is not due to increased apoptosis.

No Newly Generated GFRα1+ A Spermatogonia. Upon mitotic division, an A spermatogonium either self-renews by forming two new A spermatogonia or takes a step toward differentiation by forming an A pr two-cell chain (Fig. 5A). Having found that Rb-deficient GFRα1+ A spermatogonia proliferate at a rate comparable with controls (Fig. 4C), we wondered whether they fail to self-renew, forming A pr spermatogonia only, leading to their depletion. To test this hypothesis, we performed a BrdU-tracing experiment to trace the mitotic progeny of A spermatogonia in both control and Rb cKO testes. In this experiment, mice were given a single injection of BrdU at P5/P6 and killed 5 d later. If Rb-deficient GFRα1+ A spermatogonia failed to self-renew, we would expect to detect only BrdU+ GFRα1+ A pr spermatogonia, but no BrdU+ GFRα1+ A spermatogonia, in Rb cKO mice at the time of testis recovery.

In control testes, we readily detected both BrdU+ GFRα1+ A and BrdU+ GFRα1+ A pr spermatogonia in seminiferous tubules, as expected given that wild-type A spermatogonia undergo both self-renewal and differentiation (Fig. 5B). In Rb cKO testes, by contrast, we found BrdU+ GFRα1+ A spermatogonia but no BrdU+ GFRα1+ A pr spermatogonia (Fig. 5B), suggesting that Rb-deficient GFRα1+ A spermatogonia were not capable of self-renewal. Similarly, we observed the absence of BrdU+ GFRα1+ A spermatogonia in Rb cKO testes when mice were injected with BrdU at P10/P11 and killed 3–5 d later (Fig. 5B). Accordingly, we surmise that upon cell division, Rb-deficient GFRα1+ A spermatogonia exclusively form A pr spermatogonia and do not self-renew to form new GFRα1+ A spermatogonia; each GFRα1+ A cell can therefore initiate only one round of spermatogenesis.
examined whole-mount seminiferous tubules at P14. In controls, the density of A\textsubscript{ps} spermatogonia (particularly A\textsubscript{ps} progenitor spermatogonia) rises and falls periodically along the length of the seminiferous tubules, peaking in epithelial stages II–VII (Fig. 4B, yellow brackets) and reaching its lowest point in stages VIII–IX (30, 32, 33). This periodic variation occurs largely because active proliferation of A\textsubscript{ps} spermatogonia is limited to stages IX to III, coupled with the fact that they transition to becoming differentiating spermatogonia in stages VII–VIII. In the Rb cKO tests, we observed stretches of seminiferous tubules that did not contain any A\textsubscript{ps} spermatogonia because of premature SSC exhaustion. However, in the areas that still retained A\textsubscript{ps} spermatogonia, the chains of these cells were larger and more densely packed than in controls (Fig. 4B, yellow brackets). These overly accumulated cells were nearly all A\textsubscript{al} progenitor spermatogonia (GFRA1\textsuperscript{+} PLZF\textsuperscript{+} chains as well as GFRA1\textsuperscript{+} PLZF\textsuperscript{+} chains of four or more cells). These results confirm that, in Rb cKO tests, progenitor spermatogonia undergo a marked expansion—albeit transiently—because of the eventual exhaustion of the SSCs from which the progenitor spermatogonia derive.

To investigate the mechanism underlying this expansion of progenitor spermatogonia in Rb cKO tubules, we examined the proliferation status of A\textsubscript{ps} (PLZF\textsuperscript{+}) spermatogonia using the mitotic marker phospho-(Ser10)-histone H3 (PH3) (PH3). We studied these spermatogonia in tubule cross-sections at stages VII and X–XII, respectively, the mitotically inactive and active phases. We found no significant difference in the PH3 labeling index in PLZF\textsuperscript{+} spermatogonia between control and Rb cKO tests during the mitotically active phase (stages X–XII). However, during the mitotically inactive phase (stage VII), when only ∼7% of control PLZF\textsuperscript{+} spermatogonia were positive for PH3, ∼59% were positive in Rb cKO tests (Fig. 6). Taken together with the findings that GFRA1\textsuperscript{+} A\textsubscript{ps} spermatogonia proliferate at a normal rate and are progressively lost in Rb cKO tests (Fig. 4), these results indicate that Rb-deficient progenitor spermatogonia (non-SSC A\textsubscript{ps} spermatogonia) fail to arrest at a time when control spermatogonia are mitotically quiescent, leading to their increased accumulation.

In wild-type seminiferous tubules, excess spermatogonia are subject to density control regulation, undergoing cell death at the differentiating spermatogonial stage (36, 37). We expected that the overproduced progenitor spermatogonia in Rb cKO tests would be culled when they progress to become differentiating spermatogonia. To test this prediction, we counted apoptotic cells in control and Rb cKO testis sections at P10, at which time spermatogonia (and, in some tubules, early spermatocytes) comprise the majority of testicular germ cells. Indeed, we observed more apoptotic cells in Rb cKO than in control tubules (Fig. S5). As a consequence of this density-control regulation, the increased number of progenitor spermatogonia at P10 and P14 (Fig. 3D) did not lead to any significant increase in the number of preleptotene spermatocytes at the same times and up to P28 in Rb cKO tests (Fig. S3). We concluded that spermatogonial density control is unaffected in Rb cKO tubules, and thus that Rb is dispensable for this process.

Although Rb deficiency resulted in extended proliferation of progenitor spermatogonia, our histological analysis indicated that these cells subsequently differentiated into spermatozoa in a stage-dependent manner similar to that observed in controls. Given the known functional redundancy among the Rb family genes (5, 38), functional compensation from other Rb family genes may contribute to the absence of a significant differentiation phenotype in spermatogenesis of Rb cKO mice.

Regulation of self-renewal and differentiation of stem cells depends on both intrinsic and extrinsic factors (23, 25, 26, 39). One possible explanation for the diversion of SSC fate from self-renewal to differentiation in Rb cKO tests is that RB is involved in the intracellular signaling essential for SSC entry into the self-renewal pathway. Alternatively, Rb-deficient germ cells may fail to acquire self-renewal capacity in response to cues from stem-cell niches in the postnatal testis. Because a definitive marker for self-renewing SSCs is lacking at present, the question of whether the GFRA1\textsuperscript{+} A\textsubscript{ps} spermatogonia seen in young Rb cKO tests had properly transitioned to the SSC state awaits further investigation.

Interestingly, a phenotype similar to that reported here in Rb cKO mice has been observed in Id4-null mice. Id4-null SSCs were shown to lose their self-renewal capacity, albeit much more gradually than in Rb cKO mice, without affecting their differentiation to form functional sperm (40). It will be important to determine whether RB and ID4 interact directly in regulating SSC self-renewal, as has been shown for RB and ID2 in other systems (41, 42). Other factors known to regulate mouse SSC maintenance intrinsically include NANOS2, PLZF, GFRA1, RET, and ETV5. NANOS2 functions in SSCs to maintain their undifferentiated state (22, 27). PLZF, GFRA1, RET, and ETV5 regulate proliferation of cells within the A\textsubscript{ps} population, and, as assayed by transplantation, the absence of any one of these factors eliminates or reduces the number of SSCs (22, 43–46). It will be of interest to determine whether RB governs SSC self-renewal through interaction with these known factors or others yet to be identified.

In summary, our data demonstrate that Rb plays an unexpected role in maintaining the germ-line stem cell pool in the mouse testis by regulating SSC self-renewal. Although Rb deficiency results in overexpansion of many somatic stem-cell populations, it leads to rapid exhaustion of SSCs. In addition to its role in SSC self-renewal, we also find that Rb plays a separate role, as a cell-cycle regulator, in progenitor spermatogonia. Rb is required for these cells to become quiescent during the mitotically inactive phase (stages III–VIII) in the seminiferous tubules. Despite these defects, however, Rb deficiency does not impair the ability of SSCs to differentiate into mature spermatzoa, which is in contrast to RB’s general roles in fate decision and differentiation of somatic stem cells. Rb’s regulatory roles in spermatogenesis—including its essential function in male germ-line stem cell renewal—are remarkably different from its known roles in somatic lineages.

Materials and Methods

Mice. Mice carrying the Rb conditional allele (28) were a gift from Tyler Jacks and Jacqueline Lees (Massachusetts Institute of Technology). Mvh\textsuperscript{Cre-mOrange} mice were generated by gene targeting in C57BL/6 ES cells (Fig. S1). The targeting vector replaced a portion of exon 1 and intron 1 of the Mvh locus with the Cre-mOrange gene. The mOrange plasmid (47) used to generate the targeting vector was a gift from Roger Tsien (University of California, San Diego, CA). Rb conditional mutants and littermate controls were obtained by crossing Rb\textsuperscript{fl/fl} and Rb\textsuperscript{fl/+}: Mvh\textsuperscript{Cre-mOrange}\textsuperscript{+} mice. All experiments involving mice were approved by the Committee on Animal Care at the Massachusetts Institute of Technology.

Histological Analysis and Seminiferous Tubule Staging. Tests were fixed overnight in Bouin’s solution, embedded in paraffin, and sectioned. Slides were then dewaxed, rehydrated, and stained with periodic acid-Schiff–hematoxylin. Stage-identifying criteria were described previously (24). Spermatogonia and preleptotene spermatocytes were counted in stage VII tubule cross-sections and normalized to the number of Sertoli cells (a supporting somatic lineage) to correct for variation in tubule size.

Immunohistochemistry. Immunohistochemical staining of testis sections was carried out as described previously (48). Primary antibodies against active caspase-3 (ab13847, Abcam), BrdU (OBT0030, Accurate Chemical and Scientific), GFRA1 (AF560, R&D Systems), histone H3 (phospho S10, ab5176, Abcam), MVH (AF2030, R&D Systems), PLZF (OP128, EMD Biosciences), and RB (sc-7905, Santa Cruz Biotechnology) were used in the study.
Whole-Mount Seminiferous Tubule Staining. Mouse testes were dissected to remove the tunica albuginea, and seminiferous tubules were untangled using forceps. Whole-mount immunohistochemical staining was carried out as described previously (49). Stained tubules were spread on glass slides and imaged. Primary antibodies used are listed in the previous paragraph.

For BrdU incorporation experiments, mice were injected intraperitoneally with 100 mg/kg body weight of BrdU 4 h before sacrifice. For BrdU-tracing experiments, BrdU was given 3–5 d before mice were killed. Testes were then removed and processed for whole-mount immunohistochemical staining, following the procedure described previously, except that the tubules were denatured with 3.5 N HCl for 2 min before blocking. The percentages of GFRa1+ A, and A+ spermatogonia positive for BrdU label were determined using an LSM710 confocal microscope (Zeiss). In each animal, we counted all GFRa1+ A, and A+ cells positive or negative for BrdU in at least two seminiferous tubules, each at least 10 mm in length.

To determine the density of GFRa1+ A spermatogonia, we counted all GFRa1+ A cells in at least two seminiferous tubules, each at least 10 mm in length, in each animal. The density was obtained by dividing the total number of GFRa1+ A spermatogonia by the length of the tubule.

ACKNOWLEDGMENTS. We thank J. Jacks and J. A. Lees for Ref60 mice; R. Y. Tisen for the mOrange plasmid; M. Goodheart for performing blastocyst injections; D. S. Pearson and Y.-H. Lee for experimental support; and R. Desgraz, R. D. George, J. F. Hughes, B. J. Lesch, K. A. Romer, S. Y. Q. Soh, and R. A. Weinberg for critical reading of the manuscript. This work was supported by the Howard Hughes Medical Institute.