Mice lacking cyclin D1 are small and show defects in eye and mammary gland development

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Using homologous recombination, mice lacking cyclin D1 were generated by replacing most of the first exon of the Cyl-1 gene with sequences encoding neomycin resistance. Cyl-1−/− mice were viable and fertile but consistently smaller than their heterozygous or wild-type littermates. The nullizygous animals also showed two distinctive abnormalities: a severe retinopathy caused by impaired development of all layers of the retina and, in the mammary gland during pregnancy, a marked reduction in acinar development accompanied by a failure to lactate. Approximately 50% of animals also had a malformation of the jaw that manifested itself as a misalignment of the incisor teeth. Mouse embryo fibroblasts isolated from 14 day nullizygous, heterozygous, or wild-type embryos and grown under standard conditions showed similar cell-cycle and growth characteristics. Thus although cyclin D1 kinase activity may facilitate G2 progression, it is not essential for the development of most tissues and organs, and only a few specialized cell lineages are demonstrably sensitive to its absence.

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The mammalian cell division cycle is regulated by the sequential activation of a series of cyclin-dependent protein kinases (cdks) whose function relies on association with a specific regulatory subunit, or cyclin (for review, see Norbury and Nurse 1992; Reed 1992; Pines 1993a; Hunter and Pines 1994). Although the activity of each holoenzyme is modulated further by specific phosphorylation or dephosphorylation, the temporal order of events is primarily dependent on the synthesis of the cyclin. Thus, cyclins D1, D2, and D3 in conjunction with their catalytic partners cdk4 and cdk6 appear to regulate the initial phases of the cell cycle, as cells exit G0 or progress through G1 (Matsushime et al. 1992, 1994; Xiong et al. 1992; Bates et al. 1994; Meyerson and Harlow 1994). The subsequent transitions are executed by complexes between cyclin E and cdk2, cyclin A and cdk2, and cyclin B and cdk1 (cdc2) (for review, see Pines 1993b; Sherr 1993).

The multiplicity of D-type cyclin/kinase complexes and their early appearance in cells exposed to growth factors, make them good candidates for coupling signal transduction pathways with the cell cycle machinery. The levels of these complexes reach a maximum in late G1 phase, at a time roughly coincident with the restriction point (Pardee 1989; Matsushime et al. 1991). This is an operationally defined transition, at which cells become irrevocably committed to the division cycle irrespective of external stimuli and correlates temporally with the hyperphosphorylation and functional inactivation of the retinoblastoma gene product (pRb) (Pardee 1989; for review see Hollingsworth et al. 1993; Weinberg 1995). Whether these two checkpoints are equivalent remains unproven, but it is clear that pRb and the related p107 protein are major substrates for the cyclin D-dependent kinases (Beijersbergen et al. 1995; for review, see Weinberg 1995).

The evidence favoring a functional link between D-type cyclins and pRb is persuasive. Microinjection of a monoclonal antibody that neutralizes cyclin D1 function causes normal cells to arrest in G1, but this arrest is not seen in cells that lack functional pRb (Baldin et al. 1993; Quelle et al. 1993; Lukas et al. 1994). Similarly, in cells synchronously released from G1 by addition of growth factors, the antibody will only have an effect if injected prior to a time in G1 corresponding roughly to the pRb checkpoint. A specific inhibitor of D-cyclin/kinase function, such as the product of the tumor suppressor gene p16, also causes a late G1 arrest that is dependent on the presence of functional pRb (for review, see Sherr and Roberts 1995).

Although all six complexes involving the D-type cyclins and either cdk4 or cdk6 appear capable of phosphorylating pRb in vitro, it is unlikely that in vivo they are functionally equivalent (Matsushime et al. 1992, Dowdy et al. 1993; Ewen et al. 1993; Kato et al. 1993; Meyerson and Harlow 1994). For example, it is well established...
that deregulated expression of cyclin D1 occurs in a number of human cancers, yet there is little direct evidence to implicate cyclins D2 and D3 similarly [for review, see Peters 1994]. In some tumors, such as mantle cell lymphomas carrying the t(11;14) translocation, cyclin D1 is activated in cell types that do not normally express the gene. In breast and squamous cell carcinomas, on the other hand, amplification of cyclin D1 simply increases the level of a gene that is already expressed. Attempts to confirm the oncogenicity of cyclin D1, by overexpressing the cDNA in transfected cells or in transgenic mice, have given confusing results [Ando et al. 1993; Jiang et al. 1993; Quelle et al. 1993; Bodrug et al. 1994; Lovec et al. 1994; Resnitzky et al. 1994; Wang et al. 1994]. In some systems, ectopic expression of cyclin D1 can accelerate the G1 phase, as predicted for a positive regulator of cdks, but in others, excessive cyclin D1 inhibits proliferation [Ando et al. 1993; Jiang et al. 1993; Quelle et al. 1993; Resnitzky et al. 1994]. In the mammary gland, ectopic expression of cyclin D1 leads to abnormal proliferation, whereas in B cells its oncogenic effect is only apparent when coexpressed with c-myc [Bodrug et al. 1994; Wang et al. 1994]. To try to gain further insights into the tissue specificity, functional redundancy and oncogenicity of the D-type cyclins, we have used homologous recombination in embryonic stem (ES) cells to generate mice lacking cyclin D1 protein.

**Results**

**Targeted disruption of Cycl-1 in ES cells**

The five coding exons of the mouse Cycl-1 gene extend over ~12 kb of genomic DNA [Fig. 1A; Smith et al. 1995]. The first exon of 336 bp encodes the LXCXE motif (Dowdy et al. 1993; Ewen et al. 1993), which is characteristic of proteins that interact with pRb, as well as several amino acids of the cyclin box motif. The targeting vector effectively replaced 200 bp of exon 1, between codon 2 and a Nhel site in the first intron, with 1.8 kb of a PGK-neo cassette (see Fig. 1A; Materials and methods). Electroporation of the vector into ES cells, using established procedures (see Materials and methods), gave rise to G418-resistant colonies that were screened by Southern blotting with probe 1 from exon 5 of Cycl-1 [Fig. 1A]. Of 400 G418-resistant colonies tested, four clones gave an additional band of the size predicted for a mutant allele arising from homologous recombination. Further analysis of two of these clones [D40 and E8] confirmed their status and they were used to generate mutant mice.

**Generation of mice with heterozygous and homozygous disruption of the Cycl-1 gene**

Chimeric mice were generated by injection of D40 or E8 recombinant ES cell clones into C57BL/6J blastocysts.
Transmission of the mutant gene was determined by PCR, and F1 mice carrying the disrupted allele were interbred. Two independent colonies of mice generated from clones D40 and E8, respectively, were maintained, and both gave rise to similar phenotypes. The genotype of F2 progeny carrying no, one, or two copies of the mutant or wild-type allele was established by PCR. Southern analysis of DNA from Cyl-1^+/+, Cyl-1^-/- and Cyl-1^-/- embryos confirmed the reliability of genotyping by PCR (Fig. 1C). Heterozygous crosses yielded a slightly lower proportion of Cyl-1^-/- mice than expected for the segregation of a single affected allele, suggesting that there was a significant degree of embryonic lethality for the Cyl-1^-/- genotype (Table 1). It was also apparent that virtually all mice null for Cyl-1 were abnormally small when compared with their heterozygous and wild-type littermates (Fig. 2A). During subsequent growth to adulthood, the Cyl-1^-/- animals remained proportionately smaller (between 10% and 40%), as illustrated by the growth curves of representative litters (Fig. 2B,C).

To determine whether there was any detectable expression of Cyl-1 transcripts in mice with both Cyl-1 alleles disrupted, total RNA for Northern blot analysis was isolated from several different tissues of Cyl-1^-/- mice and a wild-type littermate. Tissues were selected on the basis that they normally express readily detectable levels of Cyl-1 transcripts [V. Fantl, unpubl.]. A probe from exon 5 that hybridizes to the 3’ portion of the mRNA, downstream of the PGK-neo cassette, was used. A prominent transcript of 4.0 kb was observed in tissues from the Cyl-1^+/+ mouse, but no specific signal was discernible in RNA samples from Cyl-1^-/- mice (Fig. 3A). To determine whether there were any changes in expression of other D-type cyclins, the same blot was rehybridized sequentially with probes to Cyl-2, Cyl-3, and finally to β-actin to act as an RNA loading control (Fig. 3A). In some tissues from Cyl-1^-/- mice there appeared to be slightly elevated levels of Cyl-2 RNA compared to the β-actin controls, but Cyl-3 RNA levels were comparable.

Although there were no detectable Cyl-1 transcripts in mice with the disrupted allele, we sought to confirm these observations at the protein level. Extracts were prepared from wild-type and Cyl-1^-/- tissues and immunoblotted using a rabbit antiserum against the carboxy-terminal peptide of cyclin D1. The 36-kD cyclin D1 protein, which migrates as a doublet, was only detected in extracts from the wild-type animals (Fig. 3B; Matsushime et al. 1991; Bates et al. 1994).

Anatomical and histopathological abnormalities associated with Cyl-1^-/- mice

Several Cyl-1^-/-, Cyl-1^+/+, and wild-type littermates were autopsied to investigate the presence of developmental abnormalities. In general, the weights and sizes

| Table 1. Transmission of the mutant Cyl-1 gene from Cyl-1^-/- pairings |
|---------------------------|----------------|----------------|----------------|
| Mouse line | Total | Cyl-1^+/+ | Cyl-1^-/- | Cyl-1^-/- |
| D40 | 118 | 37 (31%) | 58 (49%) | 23 (19%) |
| E8 | 158 | 45 (28%) | 86 (54%) | 27 (17%) |
| Overall | 276 | 82 (30%) | 144 (52%) | 50 (18%) |

The percentage of Cyl-1^-/- progeny was lower than would be expected (25%) for the segregation of a single mutant allele (P = 0.007).
Mice deficient for Cyl-1

of the major organs of Cyl-1—/- mice were proportionately lower in keeping with their smaller size. Moreover, the histological appearance of these tissues was not discernibly different from the Cyl-1+/+ and wild-type animals, except for the abnormalities described below.

Direct observation showed ~50% of the Cyl-1—/— mice had misalignment of the incisor teeth, leading to their excessive growth (Fig. 4). In a few severe cases, the teeth needed to be reduced in length to allow the mice to feed properly. However, it should be noted that the small size of the Cyl-1—/— mice was not attributable to the misalignment affecting feeding, as mice without such defects were equally runted. To investigate the underlying cause of the misaligned incisors, selected mice were examined radiologically. Results indicated that while development of the teeth was normal, lateral distortion of the mandibles in the anterior part of the jaw was causing misalignment of the incisors. In one case, there was a marked distortion of the maxilla on one side (Fig. 4).

As part of a detailed histological analysis, several eyes from nullizygous and wild-type mice were examined. The stained sections revealed a striking reduction in thickness and organization of the retinal layers (Fig. 5). A reduction in thickness and organization of the surface ganglion cell layer was also seen. These features are consistent with a severe retinopathy, and in all cases examined it was bilateral. The maximal diameters of eyes from Cyl-1—/— mice (2.03±0.14 mm) were not significantly different from control mice (2.33±0.31 mm), taking into account the difference in size between the two groups.

Despite their reduced size, both male and female Cyl-1—/— mice proved to be fertile, and litters of up to 11 young were produced from such pairings. However, litters were invariably killed by their mothers, never surviving beyond the second day. Such litters could only be rescued when cross-fostered. This prompted us to examine the mid-pregnant and postpartum mammary glands to determine whether there was a problem with their growth or differentiation. At both stages of development, mammary glands of Cyl-1—/— mice showed poor acinar development and very little secretory activity (Fig. 6).

Cell cycle kinetics for embryo fibroblast lines from Cyl-1—/— mice

To investigate the growth of cells with and without cyclin D1, mouse embryo fibroblast (MEF) cultures were
prepared from a number of individual 14 day embryos derived from heterozygous crosses. The different embryo cell cultures were genotyped by PCR and their growth rates compared as described in Materials and methods. Most of the cell cultures showed similar growth rates, whether they were from Cyl-1^-/-, Cyl-1^+/-, or Cyl-1^+/+ mice, although two lines, one wild-type [not shown] and one mutant, have shown a slower rate of growth (Fig. 7). Following serum stimulation of quiescent MEFs, the proportion of cells in different phases of the division cycle (G0/G1, S, and G2/M) was monitored at intervals by flow cytometry. The results show that embryo cell cultures derived from Cyl-1^-/- and Cyl-1^+/+ mice displayed indistinguishable kinetics for entry into, and progression through, the cell division cycle (Fig. 8). Cells started to enter S phase between 15 and 18 hr, with a maximum proportion in S phase at 21 hr. At this time, there was a noticeable increase of cells entering G2/M, which peaks at 28 hr.

Table 2. Comparison of retinal tissue from Cyl-1^+/+, Cyl-1^-/-, and Cyl-1^-/- mice

| Cyl-1 status | Retinal layer thickness (µM) | Nuclear cell layer (no. of nuclei) |
|--------------|-----------------------------|-----------------------------------|
| +/-, +/-     | 14.8 ± 0.2                  | inner: 7.1 ± 0.8; outer: 12.9 ± 0.3 |
| +/-          | 6.7 ± 0.5                   | inner: 3.8 ± 0.2; outer: 5.2 ± 1.8 |

Observations were made on six eyes in each category. For individual eyes the retinal thickness and numbers of nuclei were measured at several points. Numbers refer to the mean values ± S.D.

no detectable cyclin D1 in MEFs from Cyl-1^-/- mice, but these cells showed normal patterns of expression for cyclins D2 and D3. Although in wild-type cells, cyclin D1 accumulation precedes that of cyclin D2, in the Cyl-1^-/- MEF there was no compensatory appearance of cyclin D2 at an earlier time (Fig. 9). Similarly, there was no significant difference in the time of cyclin D3 accumulation. We also examined cyclin A and a presumed target of cyclin D1/cdk activity, pRb. In both nullizygous and wild-type cell lines, cyclin A levels became detectable at 15 hr, concomitant with the onset of DNA synthesis, and reached a maximum in mid-G2/M. Similarly, pRb was detectable throughout the cell cycle, but its electrophoretic mobility became progressively retarded be-
Mice deficient for Cyl-1

determine whether there is a specific lack of growth hormone and prolactin-secreting cells in Cyl-1<sup>-/-</sup> mice.

Although the dwarfism and mammary gland defect may have a related cause originating in the pituitary, there is no obvious link between these defects and the retinal abnormality. All layers of the retina are present in the eyes of Cyl-1<sup>-/-</sup> mice, but there is a striking decrease in their thickness. Rather than affecting the differentiation of retinal cells, the absence of cyclin D1 appears to only affect their proliferation. Although transcripts for all three D-type cyclins are detectable in whole eyes (data not shown), this finding suggests that there is an inadequate compensatory mechanism for the loss of cyclin D1 in the retina. Whether this is attributable to a specific lack of cyclin D2 and/or D3 in this tissue remains to be determined.

As judged by the normal histological appearance of the majority of tissues in the Cyl-1<sup>-/-</sup> mice, most cells appear unaffected by the lack of cyclin D1. This contrasts
tween 12 and 18 hr. This change in migration correlates with increasing degrees of phosphorylation (Buchkovich et al. 1989; De Caprio et al. 1989). In Cyl-1<sup>-/-</sup> MEF, the pattern of phosphorylation was comparable to wild-type cells.

Discussion

The results presented here show that a functional Cyl-1 gene is not essential for the development of viable mice, although they are smaller than their Cyl-1<sup>+/+</sup> and Cyl-1<sup>+/+</sup> littermates [Fig. 2], and there is a detectable level of embryonic lethality (Table 1). Cyl-1<sup>-/-</sup> mice reach sexual maturity and are able to produce live offspring, but they cannot rear their young because of an inability to produce milk. As a result, the mothers invariably kill their litters within 2 days of birth. The litters can be rescued by cross-fostering, suggesting that lactational inactivity is the main and possibly only reason why the natural mothers fail to nurture their young.

The cause of this failure to lactate has yet to be established. It could be that an absence of cyclin D1 severely curtails the normal proliferation and differentiation of the mammary gland that is associated with pregnancy. Alternatively, it may reflect an inadequate lactotrophic response because of a deficiency of prolactin. Prolactin is secreted by the lactotrophic cells of the anterior pituitary, and it is an intriguing possibility that pituitary insufficiency could also explain the small size of Cyl-1<sup>-/-</sup> null mice. A proportional reduction in body size is a characteristic of pituitary dwarfism and is caused by inadequate levels of growth hormone. The latter is synthesized and stored in the somatotrophic cells of the anterior pituitary. However, other anterior pituitary hormones such as follicle-stimulating hormone and luteinizing hormone appear to be secreted at sufficient levels to maintain fertility in most Cyl-1<sup>-/-</sup> mice. A detailed investigation of pituitary function will be necessary to

Figure 8. Cell cycle kinetics of MEFs. (A) Cells derived from a wild-type mouse embryo [E8]; (B), cells from a nullizygous embryo [E6]. After serum depletion for 3 days, the quiescent cells were stimulated by the addition of medium containing 10% FBS, and samples taken for FACS analysis at the indicated times.
with the results reported for antibody microinjection and antisense experiments suggesting that abrogation of cyclin D1 function causes cells to arrest in G1 (Baldin et al. 1993; Quelle et al. 1993; Lukas et al. 1994). It is possible that antibodies produce a more severe effect by interfering with proteins related to, and associated with, cyclin D1, such as cyclin D2, cdk4, and cdk6. Alternatively, there might be a major difference between the effects observed in tissue culture with those occurring in the whole animal. However, MEFs originating from the Cyl-1−/− mice showed no significant differences in growth rates or in cell cycle kinetics, nor were we able to demonstrate a compensatory increase in the level or time of appearance of the two other D-type cyclins [Fig. 9]. Moreover, within the limits of the analysis afforded by immunoblotting, the phosphorylation of pRb also appeared unaffected. However, because resolution of the cell division cycle is always compromised by imperfect cell synchrony, small differences in the length of G1, or cyclin expression patterns might not be discernible. These preliminary studies were also conducted under optimal growth conditions, and it is possible that differences in the behavior of the nullizygous cells may only become apparent under more restrictive growth conditions when proliferation might be more dependent on signaling pathways that act preferentially through cyclin D1.

The observation that organs such as kidney, salivary gland, and seminal vesicles, which normally have readily detectable Cyl-1 transcript levels, are apparently unaffected in nullizygous mice indicates that during development, either cyclin D1 is not essential or there is some mechanism to compensate for its loss. This latter possibility suggests that either the D-type cyclins can substitute for one another or there are alternative signaling pathways. These alternative pathways may or may not involve the other D-type cyclins in the coupling of external signals with cell cycle control. However, in a few specialized tissues, such as the developing retina, there is an absolute requirement for cyclin D1. To distinguish between these possibilities it will be necessary to determine the normal pattern of D-type cyclin expression in the affected tissues during embryogenesis and also to assess the concerted effect of combined deletions of D-type cyclins in mice.

Materials and methods

Construction of targeting vector

The Cyl-1 genomic locus was isolated as a recombinant bacteriophage λ clone from a 129/Sv mouse genomic library using a mouse Cyl-1 cDNA probe (Smith et al. 1995). A 10-kb XbaI fragment was subcloned into Bluescript KS+ plasmid (Stratagene) for further manipulation. The targeting vector was generated by inserting a PGK-neo cassette between the NcoI and Nhel sites in the first exon of Cyl-1 by blunt end ligation [Adra et al. (1987)]. This was performed on a HindIII–EcoRI subclone and then reassembled into the equivalent of the 10-kb XbaI fragment (see Fig. 1).

Electroporation and selection of ES cells

GK129 ES cells derived from mouse strain 129/Ola were maintained on a monolayer of mitomycin C-inactivated, neomycin-resistant, STO fibroblast feeder cells, as described previously [Robertson 1987]. A suspension of trypsinized cells [5×10^6/ml in Dulbecco’s modified Eagle medium (DMEM)] was electroporated with 20 μg [1 mg/ml] of linearized DNA using a Bio-Rad gene pulser (190 V, 960 μF). Cells were plated immediately after transfection and allowed to recover for 24 hr before selection in medium containing G418 (300 μg/ml). Cells were fed daily and after 9 days the resulting colonies were divided equally between two 96-well plates. When cells were confluent, one plate was frozen [−80°C] while the duplicate was used to prepare DNA.

Analysis of targeted ES cell clones

The analysis of DNA from ES cell clones was initially carried out in 96-well plates according to the method of Ramirez-Solis et al. (1992). In brief, cells were washed with PBS, lysed in 50 μl of 10 mM Tris-HCl [pH 7.5] containing 10 mM EDTA, 10 mM NaCl, 0.5% sarcosyl, and 1 mg/ml of proteinase K. After overnight incubation at 60°C, 100 μl of a mix of NaCl and cold ethanol [15 μl of 5 M NaCl/ml] was added and DNA allowed to precipitate for 2 hr [or overnight] at 4°C. The supernatant was discarded and the DNA washed three times with 70% ethanol. Plates were left tilted to air-dry for 20 min before adding 35 μl of restriction enzyme mixture (1× restriction buffer, 100 μg/ml of BSA, 50 μg/ml of RNase, and 15 units of HindIII) to each sample. After thorough mixing, using a microtiter plate shaker, samples were incubated overnight at 37°C. Electrophoresis and Southern blotting of the digested DNA was performed as described previously [Fantl et al. 1990]. A 490-bp SalI–HindIII fragment [probe 1], which lies 3′ of the genomic sequences in the targeting vector, identified bands of ∼8.9 and 10.5 kb corre-
sponding to germ-line and homologous recombinant bands, respectively. Targeted clones were recovered from the duplicate plates and expanded in 35-mm dishes. DNA from these colonies was analyzed further by KpnI digestion, which produced an endogenous fragment of ~17 kb and a correspondingly larger (~16.6 kb) band from the mutant allele. These were identified using an 800-bp PstI–PstI fragment (probe 2) located upstream of exon 1.

Generation of mice carrying the disrupted Cyl-1 allele

Two different clones of targeted ES cells were used to generate chimeric mice according to procedures described previously (Bradley 1987). C57BL/6 blastocysts injected with 10–15 ES cells were transferred to pseudopregnant female mice. Chimeric mice, identifiable by agouti coat color, were mated with C57BL/6 mice. Offspring with agouti coat color were tested for the presence of the targeted locus by PCR. Heterozygotes were interbred, and PCR analysis was used to distinguish between offspring with no, one, or two copies of the mutant gene.

Screening mice for the targeted gene by PCR

DNA for PCR analysis was prepared from tissue obtained with an ear punch. Fresh tissue was incubated at 55°C in 40 μl of buffer [50 mM at pH 8.0 containing 20 mM NaCl, 0.1% SDS, and proteinase K at 1 mg/ml] for 2 hr. Following addition of water [four volumes], samples were boiled for 5 min, cooled on ice, and 2 μl used for PCR. For the targeted allele, a 384-bp product was generated using a sense oligonucleotide, 5′-GTAGCA-GAGAGCTACAGACT-3′ [nucleotide 17–36 in Cyl-1] [Smith et al. 1995], and antisense oligonucleotide, 5′-AGTGAGACGT-GCTACTTCCA-3′ [nucleotide -264' to -288' in the PGK promoter] (Adra et al. 1987). To detect the wild-type allele, the PGK oligonucleotide was replaced with the antisense oligonucleotide, 5′-CATCCAGGTGGCCACGATT-3′ [310′–329′], in Cyl-1 to give a product of 312 bp. As a positive control sense and antisense FGF3 oligonucleotides 5641–5660, and 6157′–6176′, respectively (Moore et al. 1986), were used in all reactions giving rise to a product 535 bp.

Northern blot analysis

Total tissue RNA was prepared by solubilization in guanidine isothiocyanate and centrifugation through a cushion of cesium trifluoroacetate (Pharmacia). Samples (20 μg) were fractionated by electrophoresis in formaldehyde-agarose gels. RNA was transferred to Hybond/ECL/nitrocellulose or Immobilon-PVDF membranes for 1 hr at 300 mA. Protein loading was checked by Ponceau-S staining. Membranes were processed for antigen detection by enhanced chemiluminescence according to the manufacturer’s, instructions (Amersham). Rabbit antibodies were used to detect cyclin D1 and D2 (Bates et al. 1994), and cyclin A [gift of Tim Hunt, ICRF, London, UK], whereas monoclonal antibodies were used to detect cyclin D3 (Lukas et al. 1995) and retinoblastoma protein [Pharmingen]. Anti-rabbit and anti-mouse horseradish peroxidase-conjugated secondary antibodies (Dokapet) were used appropriately.

Flow cytometry

For the analysis of cellular DNA content, 30-mm-diameter dishes of cells were harvested by trypsinization, washed once in PBS, and fixed in 70% EtOH/PBS. After multiple washes in PBS, the cells were resuspended in 100 μg/ml of RNase at 37°C for 15 min, washed in PBS, and stained with 50 μg/ml of propidium iodide. Fluorescence was measured above 600 nm on a Becton-Dickinson FACScan flow cytometer, and the number of cells plotted against DNA content.

Histology

For histological examination, dissected tissues fixed in 10% formal saline were processed and embedded in wax using routine procedures. Eyes were embedded in the vertical plane, and their maximum diameter measured by taking serial 4-μm sections until the central region was reached. Retinal thickness from the ganglion cell layer to the choroid layer was also measured at several points. An ocular micrometer in an Olympus BH-2 microscope was used for all these measurements. Replicate counts for the number of cells in the inner and outer nuclear layers of each retina were made. Observations were restricted to the orientated central zone, avoiding the fovea and peripheral region where the cell layers are thinner.

Radiology

The jaw and skull of selected mice were analyzed in a Siemens Orbix maxillofacial X-ray unit at 52 kV and 2 mA.

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