

Induction of medulloblastomas in *p53*-null mutant mice by somatic inactivation of *Rb* in the external granular layer cells of the cerebellum

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Medulloblastomas are among the most common malignancies in childhood, and they are associated with substantial mortality and morbidity. The molecular pathogenesis as well as the ontogeny of these neoplasms is still poorly understood. We have generated a mouse model for medulloblastoma by *Cre-LoxP*-mediated inactivation of *Rb* and *p53* tumor suppressor genes in the cerebellar external granular layer (EGL) cells. *GFAP-Cre*-mediated recombination was found both in astrocytes and in immature precursor cells of the EGL in the developing cerebellum. *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* or *LoxP/LoxP* mice developed highly aggressive embryonal tumors of the cerebellum with typical features of medulloblastoma. These tumors were identified as early as 7 weeks of age on the outer surface of the molecular layer, corresponding to the location of the EGL cells during development. Our results demonstrate that loss of function of RB is essential for medulloblastoma development in the mouse and strongly support the hypothesis that medulloblastomas arise from multipotent precursor cells located in the EGL.

[Key Words: Cre-LoxP system; medulloblastoma; *Rb*; GFAP; *p53*; astrocyte]

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Astrocytes are the most abundant non-neuronal cells in the central nervous system (CNS) and participate in a wide variety of physiologic and pathologic processes. In humans this cell type most frequently undergoes neoplastic transformation in the adult brain. Although some genetic events in the neoplastic transformation of astrocytes have been identified, relatively little is known about the pathways involved in the development and progression of glial tumors.

The glial fibrillary acidic protein (GFAP) gene encodes an intermediate filament cytoskeletal protein that is restricted primarily to astroglia in the adult CNS (Bignami et al. 1972). Mouse GFAP promoter sequences have been extensively characterized and used to express heterologous genes in transgenic mice (Toggas et al. 1994; Johnson et al. 1995). We reasoned that expressing the Cre recombinase of the *Cre-LoxP* system (for review, see Rajewsky et al. 1996) under control of this promoter would enable us to study the role of the tumor suppressor genes involved in the pathogenesis of gliomas.

There is evidence that loss of function of the tumor suppressor gene *P53* is one of the initiator events in the neoplastic transformation of astrocytes. Families with

germ line mutations of *P53* have an incidence of CNS tumors of 13%, most of which (73%) are astrocytomas (Kleihues et al. 1997). The pattern of mutations found in sporadic and inherited brain tumors is similar (Fulci et al. 1998). *p53* knockout mice have been generated and extensively characterized (Donehower et al. 1992; Jacks et al. 1994), but no increased incidence of glial tumors have been reported in the short life span of these highly tumor prone mice. Astrocytes derived from these *p53* knockout mice, however, undergo spontaneous immortalization (Bogler et al. 1995; Yahanda et al. 1995).

The retinoblastoma gene was the first tumor suppressor gene to be cloned (Dryja et al. 1986; Friend et al. 1986; Lee et al. 1987a,b) and has been found mutated in a variety of human sporadic and hereditary tumors (for review, see Weinberg 1995). The retinoblastoma gene product, RB, is not only required for regulating entry into the cell cycle but also for terminal differentiation of different cell types, including CNS precursors (Lee et al. 1994). There is evidence from *in vitro* studies for the involvement of RB in differentiation and maturation of neurons and glia. Retinoic acid treated p19 embryonal carcinoma cells failed to differentiate into neurons and glia upon functional inactivation of RB by E1A (Slack et al. 1995). Moreover, loss of RB has been implicated in glial malignancies. A proportion of children with the inherited

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form of retinoblastoma will develop brain tumors later in life, and alterations of chromosome 13 have been reported in glioma cell lines as well as in about one-third of malignant gliomas (Hamel et al. 1993).

Although inactivation of the *Rb* gene in the mouse germ line leads to embryonic lethality at day 12.5 of gestation (Clarke et al. 1992; Jacks et al. 1992; Lee et al. 1992) as a result of major defects in the hematopoietic and nervous system, mice heterozygous for *Rb* or mice chimeric for wild-type and *Rb*-deficient cells develop pituitary tumors early in life (Maandag et al. 1994; Williams et al. 1994). Therefore, the consequences of lack of RB on glial differentiation and neoplastic transformation cannot be studied in these mice.

We have chosen to investigate the role of RB in the neoplastic transformation of astrocytes using the *Cre-LoxP* system. To assess a possible cooperation in tumorigenesis between RB and p53, we examined the effects of *GFAP-Cre*-mediated *Rb* inactivation in a *p53*-null background. Moreover, to overcome the limitations linked to the short life span of the *p53* knockout mice, we used *p53* conditional mutant mice, and we studied the selective inactivation of this tumor suppressor gene in astrocytes either alone or in combination with *Rb*.

We report here that RB is not required for the normal maturation of astrocytes and that the independent inactivation of p53 or RB is not sufficient to cause the neoplastic transformation of these cells in vivo. However, we found that expression of the *GFAP-Cre* transgene is not restricted to mature astrocytes. Precursor cells located in the external granular layer (EGL) of the cerebellum also express Cre recombinase driven from the GFAP promoter. Lack of *Rb* in these cells in combination with either a somatic or a germ-line *p53* inactivation leads to medulloblastomas, highly aggressive embryonal tumors of the cerebellum. No glial tumors have been found in these mice.

Results

Generation and characterization of *GFAP-Cre* transgenic mice

Transgenic mice were generated in which Cre expression was under the control of the GFAP promoter (Johnson et al. 1995). Progeny resulting from the pronuclear injection of the *GFAP-Cre* construct (Fig. 1A) were screened for integration of the transgene. Transgenic lines were derived from all 12 founder mice (FVB inbred). Two lines, Tg2 and Tg6, expressing Cre mRNA in the brain (Fig. 1B) were selected for further analysis. To investigate the tissue specific expression and the functionality of the transgene, we crossed F₁ *GFAP-Cre* Tg2 and Tg6 mice with transgenic LacZ indicator mice (ACZL and/or R26R) in which LacZ expression is only present in Cre-expressing cells and their descendants (Fig. 1C) (Akagi et al. 1997; Soriano 1999). PCR analysis on double transgenic animals showed brain specific recombination of the indicator transgene in both lines (Fig. 1D). Testis, intestine, and peripheral nerves also showed recombination, in agreement with previous reports on GFAP expression outside the CNS (McLendon and Bigner 1994). In situ enzymatic staining for β -galactosidase in combination with immunostaining for GFAP on brains of adult double transgenic mice revealed specific staining of distinct subpopulations of astrocytes. Numerous fibrillary astrocytes located in the white matter (Fig. 2A), protoplasmic astrocytes located in different areas of the cerebral (Fig. 2B) and cerebellar cortex, the Bergmann glia of the cerebellum, subpial astrocytes, and ependymal cells (Fig. 2C) were positive for both LacZ and GFAP. Surprisingly, not only astrocytes in the regions described above but also certain subpopulations of neurons [CA3 and CA4 portion of the neuronal ribbon of the hippocampus (Fig. 2D), the dentate gyrus, some neuronal subpopulations of the cortical molecular layer, and scattered cerebellar granular cells] were β -galactosidase positive al-

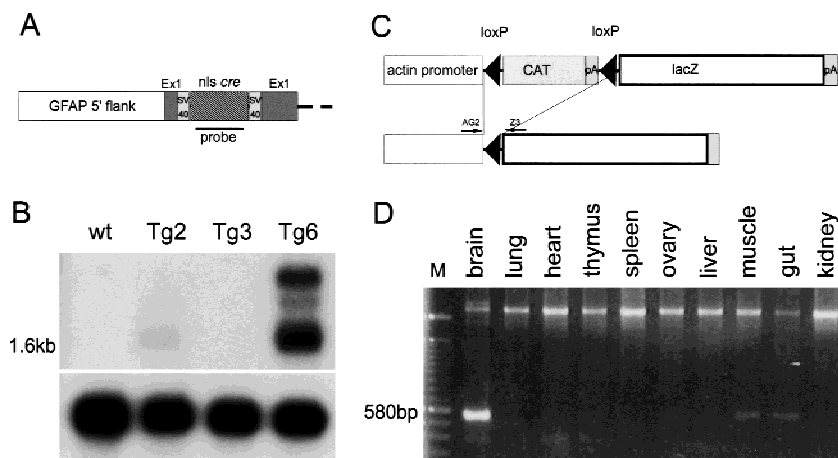
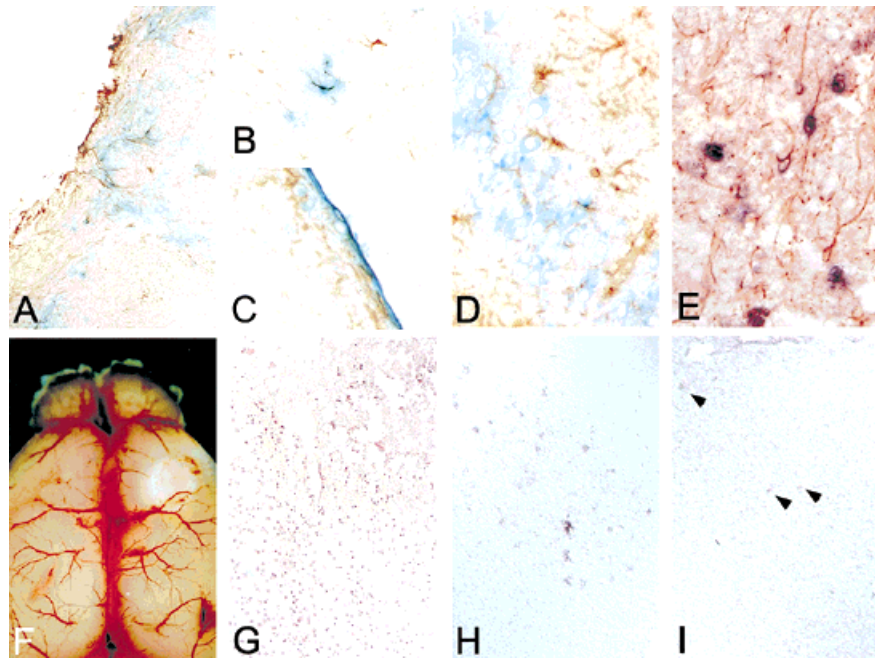


Figure 1. Structure of *GFAP-Cre* transgene and characterization of the transgenic mice. (A) Partial structure of the *GFAP-Cre* transgene. (B) Northern blot analysis of *Cre* expression on poly(A)⁺ RNA from three independent transgenic lines. A 1.6-kb mRNA corresponding to the predicted transcript from the *GFAP-Cre* transgene was detected in Tg2 and Tg6 mice. Control hybridization with hGAPDH probe shows an equal loading of RNA in each lane. (C) Genomic structure of the transgene in the ACZL mice. Cre-mediated recombination leads to juxtaposition of the *lacZ* gene to the chicken β -actin promoter. (D) PCR analysis of recombination on *GFAP-Cre;ACZL* double transgenic mice. A 580-bp product for the recombined allele is present in brain, muscle (likely originating from peripheral myelinated nerve fibers), and intestine (enteric glia). The upper band represents the nonrecombined allele. The location of the primers AG2 and Z3 is illustrated in C.

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Figure 2. In situ analysis of recombination in *GFAP-Cre;ACZL* double transgenic mice. (Upper row) Astrocytes double positive for LacZ staining (blue reaction product) and GFAP (brown stain) located in the brain stem (A) and in the cerebral cortex (B); double positive ependymal cells lining the ventricular wall (C) and neurons positive for LacZ staining and negative for GFAP in the CA4 portion of the neuronal ribbon of the hippocampus (D); double labeling of *Cre* mRNA (black) and GFAP protein (brown) in fibrillary white matter astrocytes. (E). (Lower row) Brain specimen 24 hr after cryolesioning. Macroscopical appearance of the lesioned brain (F) and histological analysis of the lesioned area on frontal sections stained with H&E (G) are shown. (H) Up-regulation of *Cre* expression in reactive astrocytes located in the necrotic area and surrounding the lesion as demonstrated by *Cre* mRNA in situ hybridization. In contrast, there is only little basal *Cre* expression in the cerebral cortex of an unlesioned animal (I).



beit GFAP negative. Both Tg2 and Tg6 lines showed this staining pattern. Endogenous β -galactosidase activity was ruled out by the absence of staining in single transgenic littermates.

In situ hybridization analysis revealed a specific expression of the *Cre* recombinase in astrocytes (Fig. 2E), subependymal cells, and scattered cells located in the CA4 layer of the hippocampus. This apparently ectopic expression might be explained by a transient *Cre* expression in multipotent precursor cells located in these regions (Palmer et al. 1997; Eriksson et al. 1998). Differentiation and maturation of these precursor cells would result in loss of *Cre* expression and sustained LacZ expression. To further assess the specificity of the *Cre* expression, we performed cryolesions to up-regulate GFAP-directed transgene expression (Johnson et al. 1995). In the mammalian CNS, GFAP expression is up-regulated in astrocytes reacting to pathologic processes. Cold lesions of the brain surface effectively induce astrocyte reaction and hence GFAP expression (Phillips et al. 1995). *Cre* in situ hybridization was performed on brain sections from *GFAP-Cre* mice, sacrificed 24 hr after cryolesioning (Fig. 2F,G). As shown in Figure 2H, reactive astrocytes in the vicinity of the lesion contained abundant *Cre* mRNA. Although the nontreated cortical gray matter contains only few *Cre* expressing astrocytes (Fig. 2I), their number was significantly increased after cryolesion. These findings suggest that *Cre* expression follows the specific expression behavior typical of the GFAP promoter.

Cell type-specific inactivation of *Rb* in the CNS

Mice carrying conditional *Rb* alleles were generated using an insertion type targeting vector in which two *LoxP*

sequences were inserted into the introns surrounding exon 19 of *Rb* (Fig. 3A). Excision of the region flanked by the *LoxP* sites results in truncation of the RB protein that is functionally equivalent to a null allele (Vooijs et al. 1998; Vooijs and Berns 1999).

We crossed *GFAP-Cre* Tg2 mice with homozygous *Rb^{LoxP/LoxP}* mice to obtain *GFAP-Cre;Rb^{LoxP/+}* mice. These mice were subsequently bred to *Rb^{LoxP/LoxP}* mice to obtain *GFAP-Cre;Rb^{LoxP/LoxP}* mice. These mice were healthy and fertile and showed no sign of CNS disease.

A cohort of 22 *GFAP-Cre;Rb^{LoxP/LoxP}* mice was kept under observation for up to 1 year of age (Table 1), and the brains were examined histologically at different time points (two mice at 1, 3, 5, and 7 months, respectively, and 14 mice at 1 year of age). PCR analysis for *Rb* recombination (Fig. 3B) and in situ-hybridization for *Cre* mRNA (data not shown) were performed on selected brain specimens at various time points and confirmed the presence of astrocytes devoid of RB. Histological analysis did not reveal any evidence for dysplastic astrocytes or small neoplastic lesions.

Medulloblastomas in mice with a somatic *Rb* inactivation in a p53-null background

In humans, *RB* mutations have been found in high-grade astrocytic tumors (Hamel et al. 1993), and the loss of its function is considered to be involved in the progression from the benign form to the highly malignant astrocytoma (Kleihues et al. 1993). Therefore, the lack of an initiation step might be responsible for the lack of neoplastic transformation of *Rb*-null mutant astrocytes. As there is evidence that loss of p53 function can act as an initiator event in low-grade astrocytoma pathogenesis (von Deimling et al. 1992; Ohgaki et al. 1993),

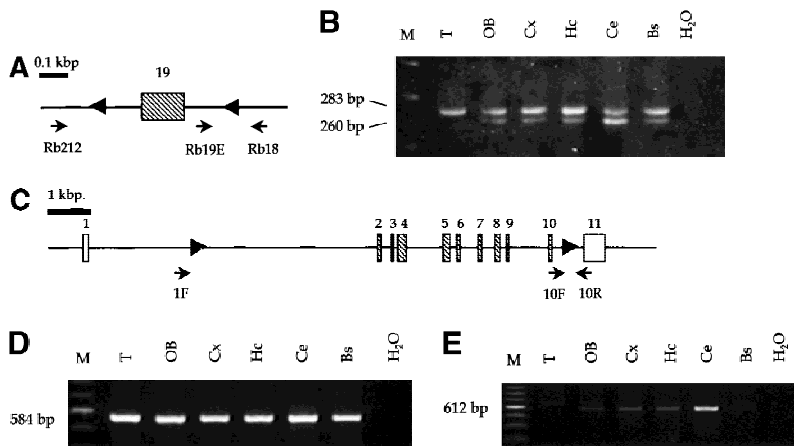
Medulloblastomas in *p53/Rb* compound mutant mice

Figure 3. Analysis of *Rb* and *p53* recombination in different areas of the adult mouse brain. (A) Structure of the *Rb* floxed allele: LoXP sites were inserted into the introns surrounding exon 19. (B) A 260-bp PCR product representing the recombined *Rb* allele (primers Rb212 and Rb18) is present in a subset of cells in all analyzed areas of the brain. The 283-bp product represents the floxed allele (primers Rb19E and Rb18). The location of the primers used for the analysis (Rb212, Rb19E, and Rb18) is illustrated in A. Structure of the *p53* floxed allele (C) and PCR analysis of recombination in different areas of the brain (D,E) are shown. The 584-bp band represents the floxed *p53* allele (primers 10F and 10R), and the 612-bp band represents the recombined *p53* allele (primers 1F and 10R). (M) Molecular marker; (T) tail; (OB) olfactory bulb; (Cx) cortex; (Hc) hippocampus; (Ce) cerebellum; (Bs) brain stem.

we asked whether there is cooperation between these two tumor suppressor genes in inducing neoplastic transformation of astrocytes. We therefore crossed *GFAP-Cre;Rb^{LoxP/LoxP}* mice with *p53*-deficient mice (Donehower et al. 1992). Thirteen mice with one intact copy of the *p53* allele (*GFAP-Cre;Rb^{LoxP/LoxP};p53^{+/-}*) were kept under observation for up to 8 months (Table 1), when they had to be sacrificed because of the development of lymphomas and different types of sarcomas (Harvey et al. 1995). Histological analysis of the brains of these mice showed no evidence for cellular atypia or neoplastic lesions of glial origin.

Seven *GFAP-Cre;Rb^{LoxP/LoxP}* mice in a *p53^{-/-}* background had to be sacrificed around 3–4 months of age (Table 1) because they developed similar tumors as mice with one inactivated *p53* allele. Three *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* mice, however, presented with symptoms of CNS disease. They circled, held their heads to one side, and showed tremor, gait disturbance with

ataxia, and loss of balance. These three mice and the additional four mice without neurological symptoms (all of them between 3 and 4 months of age) were sacrificed, and their brains were analyzed histologically. In all but one case, we detected tumors in the cerebellum (Fig. 4A,B). These tumors consisted of densely packed, polygonal cells with scant cytoplasm and hyperchromatic nuclei (Fig. 4C). The tumor cells were arranged in sheets and rows and occasionally formed pseudorosette structures. There was high mitotic activity throughout the tumor. These hypercellular tumors were located in the vermis or close to the cerebellar peduncles and showed an invasive growth pattern with ill-defined margins. Multifocal occurrence of these tumors was occasionally noted, although in these cases a secondary spread of the tumor could not be ruled out. Localization and morphological appearance of these tumors showed in all instances striking similarity to medulloblastomas, malignant tumors arising in the cerebellum of children.

Table 1. Occurrence of medulloblastoma in compound mutant mice

Genotype	Total no. of mice	Medulloblastomas in mice sacrificed at 8 months, 1 year, or when symptoms occurred	Medulloblastomas in nonsymptomatic mice sacrificed at earlier time point
<i>GFAP-Cre;Rb^{LoxP/LoxP}</i>	22	0/14 ^a	0/8
<i>GFAP-Cre;Rb^{LoxP/LoxP};p53^{+/-}</i>	13	0/13 ^b	N.D.
<i>GFAP-Cre;Rb^{LoxP/+};p53^{-/-}</i>	4	1/4 ^c (106d)	N.D.
<i>GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}</i>	7	5/6 ^c (76d–108d)	1/1
<i>GFAP-Cre;Rb^{LoxP/+};p53^{+/-}</i>	15	0/15 ^b	N.D.
<i>GFAP-Cre;p53^{LoxP/LoxP}</i>	11	0/10 ^b	0/1
<i>GFAP-Cre;Rb^{LoxP/LoxP};p53^{LoxP/+}</i>	5	0/5 ^b	N.D.
<i>GFAP-Cre;Rb^{LoxP/+};p53^{LoxP/LoxP}</i>	12	6/8 ^{b,c} (87d–196d)	0/4
<i>GFAP-Cre;Rb^{LoxP/LoxP};p53^{LoxP/LoxP}</i>	12	9/9 ^c (86d–118d)	1/3
<i>GFAP-Cre;Rb^{LoxP/+};p53^{LoxP/+}</i>	15	0/15 ^b	N.D.

The total number of mice analyzed for each genotype and the occurrence of medulloblastomas at different time points is presented.

^aMice sacrificed at 1 year of age.

^bMice sacrificed at 8 months of age.

^cMice sacrificed when symptoms occurred; time range in days is shown in parenthesis.

(N.D.) Not determined.

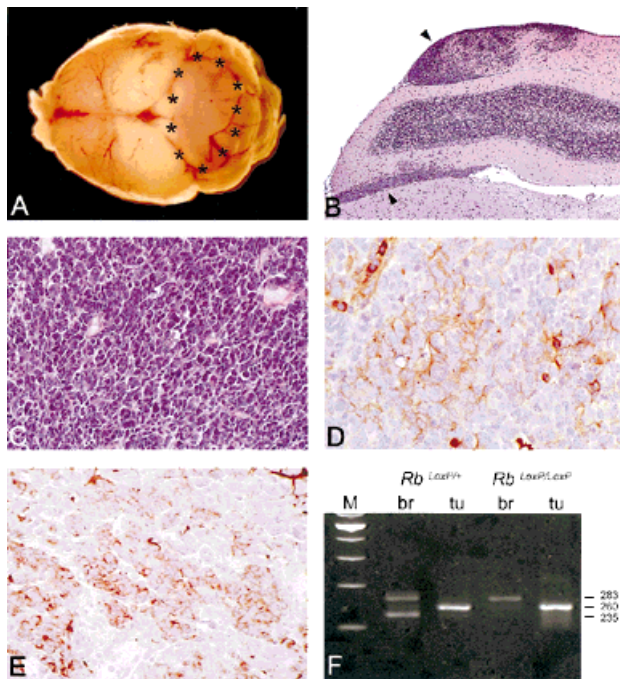


Figure 4. Histopathological analysis of medulloblastomas in *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* mice. (A) Macroscopic appearance of a medulloblastoma arising in the cerebellar vermis of a 16-week-old mouse. (B) Low-power magnification of a tumor diffusely infiltrating in a cerebellar folia and in the subarachnoidal space (upper and lower arrows, respectively; H&E). (C) High-power magnification of a hypercellular tumor consisting of densely packed, polygonal cells with scant cytoplasm and hyperchromatic nuclei (H&E). Areas of early neuronal (MAP-2; D) and glial (GFAP; E) differentiation are detectable. (F) PCR analysis of *Rb* recombination in tumors arising in *GFAP-Cre;Rb^{LoxP/+};p53^{-/-}* and *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* mice. The 283-, 260-, and 235-bp products correspond to the floxed (nonrecombined), the recombined, and the wild-type allele, respectively. The primers used for the analysis (Rb212, Rb19E, and Rb18) are the same as described in Fig. 3.

To find further support for their similarity to human medulloblastomas, we looked for the expression of glial and neuronal markers. Microtubule-associated protein (MAP-2), an early expressed neuronal antigen, was detected in the tumor cells (Fig. 4D), whereas other neuronal markers of advanced differentiation (neuron-specific enolase and synaptophysin) were not expressed (data not shown). GFAP expression was found in clusters of neoplastic cells, characterized by a thin rim of immunoreactivity around the nucleus (Fig. 4E). These cells were clearly discernible from reactive astrocytes with thin, elongated processes and a brighter, scattered nucleus. These findings indicate early neuronal and glial differentiation and suggest an origin from multipotential, uncommitted precursor cells.

To test whether the floxed *Rb* alleles were recombined in the tumors, we performed a PCR analysis on DNA extracted from tumor tissue and from adjacent normal CNS tissue. This analysis revealed that both floxed *Rb* alleles were recombined in the tumor tissue (Fig. 4F). In

addition to mice with both floxed *Rb* alleles in the *p53*-null background, there was one mouse with only one floxed *Rb* allele in the *p53*-null background (*GFAP-Cre;Rb^{LoxP/+};p53^{-/-}*) that also showed signs of CNS disease. Autopsy and histological examination of the brain showed a medulloblastoma morphologically indistinguishable from those described above. PCR analysis performed on neoplastic and normal tissue of this mouse showed that the floxed *Rb* allele was recombined and the wild-type allele was lost (Fig. 4F).

Histological examination of other brain areas of these mice revealed hydrocephalus in some cases (as a result of obstruction of the fourth ventricle) but no sign of other malignancies.

No medulloblastomas and no glial malignancies in GFAP-Cre;p53^{LoxP/LoxP} mice

Because the short lifespan of the *p53* conventional knockout mice might have masked the development of medulloblastomas and precluded the development of other CNS tumors such as gliomas, we crossed *GFAP-Cre* mice with mice carrying conditional *p53* alleles (Fig. 3C). These mice were generated by inserting *LoxP* sites in intron 1 and intron 10 of the *p53* gene using a subsequent targeting strategy. Mice homozygous for the floxed allele were monitored for up to 1 year of age and did not show increased incidence of diseases. On the other hand, *p53^{Δ2-10}* mice, obtained after crossing with mice expressing *Cre* in the germ-line (*Deleter-Cre*; Schwenk et al. 1995) were showing the same tumor spectrum of *p53* conventional knockout mice (J. Jonkers and A. Berns, in prep.).

GFAP-Cre;p53^{LoxP/LoxP} mice were healthy and fertile and were kept under observation for up to 8 months of age. The presence of astrocytes lacking *p53* was assessed on selected brain specimens with PCR-based analysis of recombination (Fig. 3D,E) and in situ hybridization for *Cre* mRNA (data not shown). The histological analysis on the brains of these animals reveals neither medulloblastomas nor malignancies of glial origin.

Conditional inactivation of both p53 and Rb leads to medulloblastomas

To ask whether *GFAP-Cre*-mediated somatic inactivation of both *p53* and *Rb* would lead to medulloblastomas, we crossed *p53^{LoxP/LoxP}* mice with *GFAP-Cre;Rb^{LoxP/LoxP}* to obtain *GFAP-Cre;Rb^{LoxP/+};p53^{LoxP/+}* mice. These mice were subsequently crossbred to obtain *GFAP-Cre;Rb^{LoxP/LoxP};p53^{LoxP/LoxP}* mice.

Between 3 and 4 months of age all compound mutants (9/9) developed medulloblastomas with the same histopathologic features described previously.

Six out of 8 mice with one floxed *Rb* allele and two floxed *p53* alleles (*GFAP-Cre;Rb^{LoxP/+};p53^{LoxP/LoxP}*), kept under observation for up to 8 months, developed medulloblastomas as well, although in some cases, after longer latency (Table 1). PCR analysis was performed on

four out of six tumors in which a good separation between neoplastic and adjacent normal tissue was present. In three out of four tumors, the floxed *Rb* allele was recombined and the wild type allele was lost. In one case, the wild-type and the floxed alleles were still detected in addition to a strong recombination band, most likely this was due to contaminating non-neoplastic cells.

EGL precursor cells are the cells of origin of medulloblastomas

One *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* mouse and three *GFAP-Cre;Rb^{LoxP/LoxP};p53^{LoxP/LoxP}* mice were sacrificed at 2 months of age without showing neurological disease, and the entire cerebella was analyzed by serial sections. In the *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* cerebellum, we identified three early neoplastic lesions located on the outer surface of the molecular layer (Fig. 5a), where the EGL cells are located during late embryonal and early postnatal development. RNA in situ hybridization showed strong *Cre* expression in the neoplastic cells (Fig. 5b). Similar lesions were found in one out of three *GFAP-Cre;Rb^{LoxP/LoxP};p53^{LoxP/LoxP}* cerebella analyzed.

Newborn mice double transgenic for *GFAP-Cre* and a

LacZ indicator construct (R26R) showed *LacZ* staining not only in the developing forebrain and in the spinal ganglia but also in the EGL cells of the cerebellum (Fig. 5c). Moreover, scattered precursor cells located in the EGL of the same animals are expressing *Cre*, as shown in Figure 5d. This finding suggests that precursor cells of the EGL activate the *GFAP* promoter and thus induce *Cre* expression and *Cre*-mediated recombination. An ectopic expression of the *GFAP-Cre* construct in the EGL precursor cells, however, cannot be excluded.

Math1 (Akazawa et al. 1995; Ben-Arie et al. 1996), the mouse homolog of the *Drosophila* gene *atonal*, encodes a basic helix-loop-helix transcription factor that is specifically expressed in the precursor cells of the EGL and is required in vivo for the genesis of granule cells (Ben-Arie et al. 1997).

We performed in situ hybridization for *Math1* on selected cases of medulloblastomas arising in *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* or *LoxP/LoxP* mice, and we observed strong *Math1* expression (Fig. 5e,f) in all tumors analyzed. The mature granule cells located in the IGL as well as the subependymal matrix cells did not express *Math1*, in agreement with previous reports (Ben-Arie et al. 1996, 1997).

These results strongly support our conclusions that these tumors arise from the granule cell precursors retained on the cerebellar surface.

Discussion

GFAP, an intermediate filament protein, is expressed in differentiated and mature astrocytes in the mammalian brain (Eng 1985; Bignami et al. 1992). A detailed expression analysis of neuronal and glial markers during the histogenesis of the human cerebellar cortex, however, described *GFAP* expression in scattered multipotent precursor cells located in the ventricular zone of the cerebellar anlage at 8 weeks of gestation (Yachnis et al. 1993). Furthermore, a recent study identified ependymal cells, known to express *GFAP*, as multipotent neural stem cells of the adult mammalian CNS (Johansson et al. 1999).

We report here that *GFAP*-driven expression of *Cre* is detectable in newborn mice not only in the developing forebrain and in spinal ganglia but also in precursor cells located in the EGL of the developing cerebellum. Although expression of the *Cre* transgene was mainly restricted to astrocytes in the brain of adult mice, *LacZ* expression was also detectable in other cell types. This could simply be due to a stochastic expression of the *GFAP-Cre* construct independently of the integration site, because we find the same expression pattern in two independently generated transgenic lines. On the other hand, a transient *GFAP* expression in precursor cells resulting in *Cre*-mediated recombination of the floxed transgene is an intriguing alternative hypothesis. In ACZL mice this would lead to *LacZ* expression in the lineage of the recombined precursor cell. After commitment of these precursor cells to a neuronal lineage, there

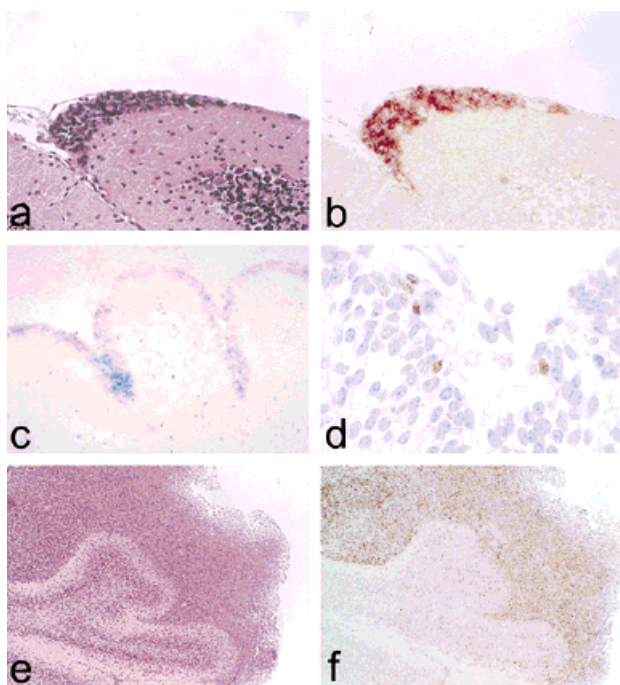


Figure 5. Analysis of the origin of medulloblastomas in *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}*. Early neoplastic lesions on the outer cerebellar surface of a 7-week-old mutant mouse (H&E; A) and in situ hybridization for *Cre* mRNA (B) are shown. *LacZ*-positive EGL cells in the cerebellum of a newborn *GFAP-Cre* mouse crossed to a *LacZ* indicator mouse show *LacZ* staining/Nuclear Red (C), and high-power magnification shows scattered *Cre*-expressing EGL precursor cells (D). Low-power magnification of a medulloblastoma surrounding cerebellar lamellae (H&E; E) and *Math-1* expression in the tumor cells RNA in situ hybridization (F) are shown.

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would be sustained LacZ expression albeit undetectable Cre expression.

GFAP expression has never been shown in the EGL of the developing cerebellum, although a low level of expression cannot be excluded. Delaney et al. (1996) generated transgenic mice expressing the thymidine kinase (*tk*) gene under the control of the *GFAP* promoter. Treatment of transgenic newborn mice with gancyclovir leads to a conversion of the drug into a toxic metabolite in cells expressing *tk*. These mice developed ataxia, and histological examination revealed a reduced size of the cerebellum and severe depletion of granule cells. This finding was explained by a deleterious effect on the radial glia (Bergmann glia) that normally guides external granule cells toward the internal granular layer. In the light of our experimental findings, however, this effect might be explained as a result of *tk*-mediated death of GFAP-expressing precursor cells of the EGL.

The developmental expression of *GFAP*-driven *Cre* recombinase in specific subpopulations of precursor cells renders *GFAP-Cre* mice particularly useful for analyzing the effects of tumor suppressor gene inactivation on the ontogeny of CNS tumors. In addition, *GFAP-Cre* mice may be instrumental in studying genes involved in the development and maturation of precursor cells or in the analysis of astrocyte-specific genes upon stimulation of mature astrocytes by pathologic stimuli.

Medulloblastoma, a malignant embryonal tumor of the cerebellum, accounts for ~20% of primary brain tumors in children. Little is known about the molecular genetic events that contribute to the development of this tumor. Mutations of the *P53* gene (Malkin et al. 1990; Cogen and McDonald 1996), amplification of *c-MYC* (Badiali et al. 1991) and *N-MYC* (Tomlinson et al. 1994), up-regulation of different *PAX* genes (Kozmik et al. 1995), and, more recently, lack of *PATCHED* (*PTC*) expression (Goodrich et al. 1997) have been implicated in the pathogenesis of these neoplasms. Mouse models for medulloblastomas are limited, a subset of mice heterozygous for *ptc* develop these tumors (Goodrich et al. 1997), and medulloblastomas have been induced in rats by transplantation of in vitro-transformed CNS precursor cells (Wiestler et al. 1992).

We describe here a new mouse model for medulloblastoma obtained by somatic inactivation of *Rb* in the EGL precursor cells of the developing cerebellum of *p53*-null mutant mice. The tumors observed in the cerebellum of *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* or *LoxP/LoxP* mice share a number of characteristics with the human tumors. These include onset in young animals, location in corresponding areas of the cerebellum, highly aggressive behavior with typical local infiltration, and a mixed neuronal–glial differentiation pattern.

The ontogeny of medulloblastomas has been a controversial issue since the original description by Bailey and Cushing in 1925. Two hypotheses have been discussed most extensively: The first postulates that these tumors arise from the subependymal matrix cells of the fourth ventricle, which are precursor cells giving rise to neurons and glia in the adult CNS (Yachnis et al. 1994). An

alternative hypothesis assumes an uncontrolled proliferation of precursor cells of the EGL (Vandenberg et al. 1987).

Because expression of *Cre* and recombination occur in scattered EGL cells before neoplastic transformation, we conclude that in our mouse model medulloblastomas arise from multipotent granule cell precursors retained on the cerebellar surface. The expression of *Math1*, a specific EGL cell marker, in the tumors provides an independent support to our conclusions.

While we observed medulloblastomas in *GFAP-Cre;Rb^{LoxP/LoxP};p53^{-/-}* or *LoxP/LoxP* mice as young as 7 weeks, we never found these tumors in *GFAP-Cre;Rb^{LoxP/LoxP}* or *GFAP-Cre;p53^{LoxP/LoxP}* or in *p53^{-/-}* mice. In line with the latter observation, only one medulloblastoma has been described in a series of 93 *p53^{-/-}* mice (Donehower et al. 1992), and no such tumors have been described in another series of 49 animals (Jacks et al. 1994).

Interestingly, not only recombination of the floxed *Rb* allele but also loss of the wild-type allele was observed in medulloblastomas arising in *GFAP-Cre;Rb^{LoxP/+};p53^{-/-}* or *LoxP/LoxP* mice. Hence, loss of function of RB is essential for medulloblastoma development in the mouse. The fact that these tumors are not found in *GFAP-Cre;Rb^{LoxP/LoxP};p53^{+/-}* mice suggests that *p53* deficiency may act as an initiator event in the ontogeny of these tumors. Alternatively, RB deficiency may stimulate excessive cell proliferation that, in normal conditions, might be counteracted by *p53*-induced apoptosis. Consequently, in a *p53*-null background, EGL precursor cells that have lost RB will fail to undergo apoptosis and ultimately give rise to neoplasms.

Although medulloblastomas represent 16% of brain tumors found in patients with germ line *P53* mutations (Li-Fraumeni syndrome), loss of *P53* is not commonly found in sporadic medulloblastomas. Additionally, RB has almost never been found mutated or lost in human medulloblastomas. Simultaneous mutations of both RB and *P53* thus appear to be rare events in these tumors. The lack of RB mutations in human medulloblastomas could be explained by mutations involving other components of the RB pathway, although a study on a small series of 16 medulloblastomas failed to detect loss of p16 and/or Cyclin D and CDK4 amplification (Sato et al. 1996).

Alternatively, *p53* and RB can be functionally inactivated and therefore contribute to the etiopathogenesis of medulloblastomas even if no mutations are found at the DNA level. SV40 Tag has been recently shown to be expressed (Huang et al. 1999) and to form specific complexes with *p53* and RB in 33% of medulloblastoma cases (Zhen et al. 1999).

We never detected gliomas in *GFAP-Cre;Rb^{LoxP/LoxP}* mice kept under observation for up to 1 year of age. Our findings in *GFAP-Cre;ACZL* double transgenic mice clearly show recombination in astrocytes; moreover, we reproducibly detected astrocytes expressing *Cre* in different brain areas of *GFAP-Cre;Rb^{LoxP/LoxP}* adult mice.

GFAP-Cre;p53^{LoxP/LoxP} mice, kept under observation

for up to 8 months of age, never developed gliomas either. On the other hand, malignant gliomas arise between 1 and 4 months of age in 20% of *GFAP-v-src* transgenic mice (Weissenberger et al. 1997), and well differentiated neoplasms between 4 and 5 months of age in 9% of *p19^{ARF}-/-* mice (Kamijo et al. 1999). Speculations on the possible causes of these differences between *GFAP-Cre;p53^{LoxP/LoxP}* and *p19^{ARF}-/-* mice are premature because the size of our cohort of mice is smaller and the genetic background different. A time course observation of *p19^{ARF}-/-* mice in the same genetic background as that of the *GFAP-Cre;p53^{LoxP/LoxP}* is currently under way.

We never observed glial malignancies in our *p53/Rb* combined mutants. These results might imply that mouse astrocytes are remarkably resistant to neoplastic transformation brought about by loss of p53 and RB. On the other hand, the shortened life span of the mice, which had to be sacrificed because of the development of medulloblastomas, might have precluded the study of late effects of p53 and RB combined deficiency in mature astrocytes.

Materials and methods

Construction of plasmids and generation of transgenic mice

The *GFAP-Cre* transgene was constructed from the *GFAP-LacZ* plasmid (Mucke et al. 1991). This construct contains not only a 2-kb fragment encompassing the *GFAP* promoter region but also positive and negative regulatory sequences located in the following introns (Miura et al. 1990).

The *Cre* open reading frame with an amino-terminal nuclear localization signal (Gu et al. 1993) was obtained as a *NotI* fragment after subcloning in a modified pBR vector. This fragment was cloned into the unique *NotI* cloning site of the *GFAP* construct. The final construct contained 2 kb of *GFAP* 5'-flanking sequences, 89 residues of *GFAP* exon 1, 0.2 kb of SV40 DNA with splice sites, a 1-kb *Cre* insert, and 0.2 kb of SV40 DNA with a poly(A) signal, followed by the rest of the *GFAP* gene. The construct was then released from the backbone, purified by agarose gel electrophoresis and electroelution, and injected into fertilized FVB/N mouse oocytes.

Screening of transgenic mice

Transgenic *GFAP-Cre* founder mice were identified by Southern blot analysis. DNA was extracted from tails according to standard protocols, digested with *EcoRV* and probed with a 1-kb *Cre*-containing fragment. After founders were established, genotyping was performed by PCR. Tail tip DNA was amplified with primers *Cre1* (5'-ACCAGCCAGCTATCAACTC-3') and *Cre2* (5'-TATACGCGTGCTAGCGAAGATCTCCATCTTCCAGCAG-3') yielding a 269-bp product. Thermocycling conditions consisted of 30 cycles of 30 sec at 94°C, 30 sec at 56°C, and 50 sec at 72°C. Reactions contained 200 ng of template DNA, 0.5 μM primers, 100 μM dNTPs, 9% glycerol, 2.5 units of *Taq* polymerase, 1.8 mM MgCl₂, 1× PCR buffer (GIBCO BRL) in a 20-μl volume.

ACZL and R26R mice were screened by PCR using the same conditions described above and primers LZ1 (5'-CGTCACACTACGTCTGAACG-3') and LZ2 (5'-CGACCAGATGATCACTACTCG-3'). Tail DNA from *Rb^{LoxP}* mice and *p53* mutant

mice was amplified for 30 cycles (30 sec at 94°C, 30 sec at 58°C, and 50 sec at 72°C) using primers Rb18 (5'-GGCGTGTGCCATCAATG-3') and Rb19 (5'-AACTCAAGGGAGACCTG-3'), p53a (5'-GCCTTCTATCGCCTTCTTGACGAGT-3'), p53b (5'-CGACCTCCGTTCTCTCTCTCTCTT-3'), p53c (5'-AGACCACAAAACAAAAACAAAATTACA-3') and p53-int10-fwd (5'-AAGGGGTATGAGGGACAAGG-3'), p53-int10-rev (5'-GAAGACAGAAAAGGGGAGGG-3'), respectively. The composition of the PCR mix was identical for all reactions.

Analysis of transgene transcription

Expression of the transgene was analyzed by Northern blotting. Total RNA was prepared from snap-frozen tissue using Trizol Reagent Total RNA isolation reagent (GIBCO BRL). The PolyA-Tract System (Promega) was used to enrich for poly(A)⁺ RNA. Northern blot was performed according to standard protocols using 5 μg of poly(A)⁺ RNA. The probes used were a 1-kb *Cre* fragment and a 1.3-kb hGAPDH fragment.

PCR analysis of recombination

PCR analysis of *Cre*-mediated recombination in the ACZL mice was performed on genomic DNA extracted from different organs according to published protocols and followed by phenol-chloroform extraction. Thermocycling conditions consisted of 35 cycles of 30 sec at 94°C, 50 sec at 57°C, and 50 sec at 72°C yielding a 580-bp product for the recombined allele. The primers used were AG2 (5'-CTGCTAACCATGTTCATGCC-3') and LZ3 (5'-GGCTCTTCGCTATTACG-3'); the composition of the PCR mix was as described above.

Microdissection of tumors and of adjacent normal brain tissue was performed using a laser capture microdissection system (Arcturus). Ten-micrometer-thick sections from formalin-fixed paraffin-embedded material were used after dewaxing in xylene and drying. The dissected material was transferred to 50 μl of proteinase K buffer and incubated at 37°C overnight, after which the proteinase K was inactivated for 10 min at 96°C. Five microliters were used for subsequent PCR amplification using primers Rb212 (5'-GAAAGGAAAGTCAGGGACATTGGG-3'), Rb18 (5'-GGCGTGTGCCATCAATG-3'), and Rb19E (5'-CTCAAGAGCTCAGACTCATGG-3') yielding a 283-bp product for the unrecombined *Rb^{LoxP}* allele, a 260-bp product for the recombined *RbΔ19* allele, and a 235-bp fragment for the wild-type *Rb* allele (Vooijs et al. 1998). The PCR cycling profile consisted of 35 cycles of 30 sec at 94°C, 30 sec at 58°C, and 50 sec at 72°C. The PCR mix consisted of 0.5 μM primers, 100 μM dNTPs, 9% glycerol, 2.5 units of *Taq* polymerase, and 1× PCR buffer (GIBCO BRL) in a 20-μl volume. A hot start was performed using Invitrogen Hot-wax MgCl₂ beads (1.5 mM). PCR products were analyzed on a 3% NuSieve agarose gel.

PCR analysis of *Cre*-mediated recombination in the *p53^{LoxP/LoxP}* mice was performed on genomic DNA extracted from different areas of the brain and followed by phenol-chloroform extraction. Thermocycling conditions consisted of 30 cycles of 30 sec at 94°C, 30 sec at 58°C, and 50 sec at 72°C yielding a 612-bp product for the recombined allele. The primers used were p53-int1-fwd (5'-CACAAAACAGGTTAAACCCA-3') and p53-int10-rev (5'-GAAGACAGAAAAGGGGAGGG-3'); the composition of the PCR mix was as described above.

Cryolesion

Mice were anesthetized, their scalps excised, and the parietal region of the head was exposed. A metal rod, cooled in liquid

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nitrogen, was applied to the skull for 45 sec. The skin wound was closed by interrupted sutures. Animals were sacrificed 24 hr or 3 days after cryolesioning; brains were dissected out, sliced coronally, and fixed in 4% PFA-PBS or frozen in OCT.

RNA in situ hybridization

In situ hybridization for Cre RNA was performed on selected brain sections. A 0.8-kb *NruI-XbaI* fragment of the *cre* coding region was directionally cloned in an *EcoRV-XbaI* cut pSP72 vector. Digoxigenin-labeled antisense probe was synthesized from the *BglIII* cleaved construct using the Sp6 polymerase. *Math1* antisense and sense probes were synthesized from the *XhoI* and *ERI* respectively cleaved ICTd construct (a gift of H. Zoghbi, HHMI, Baylor College of Medicine, Houston, TX) using T7 and T3 polymerase. The RNA in situ hybridization on frozen and on formalin-fixed paraffin-embedded material was essentially performed as described (Marino et al. 1995).

For subsequent GFAP immunolabeling, sections were processed as described below. The sections were mounted in glycerol gelatin.

Detection of β -galactosidase activity

Embryos were dissected into cold PBS and kept on ice. When all the embryos were collected, they were changed to cold 4% PFA-PBS and rocked at 4°C for 1–4 hr depending on the age of the embryo. Adult brains were dissected out, cooled at 4°C for 5 min, and sliced coronally into 2-mm slices. The slices were fixed in 4% PFA-PBS at 4°C for 2 hr. The β -galactosidase staining was performed according to standard procedure.

Selected brains were frozen and embedded in OCT compound, in a mixture of dry ice and 2-methylbutane. They were then cut with a cryostat in 6- and 12- μ m sections and stained according to standard procedures. For subsequent GFAP immunolabeling, sections were processed as described below. The sections were mounted in glycerol gelatin.

Histological analysis

Whole mouse brains were fixed for at least 24 hr in 4% paraformaldehyde in PBS. Coronal slices of ~2-mm thickness were dehydrated through graded alcohols and embedded in paraffin. Sections of 4- μ m nominal thickness were mounted on coated slides and routinely stained with hematoxylin and eosin (H&E).

Immunohistochemistry for GFAP (polyclonal, 1:500; DAKO), Synaptophysin (polyclonal, 1:50; DAKO), MAP-2 (monoclonal, 5 mg/ml; Boehringer Mannheim), NSE (monoclonal, 1:200; DAKO), and CRE (polyclonal, 1:14,000; Novagen) was performed on selected sections. Biotinylated secondary antibodies (goat anti-rabbit and rabbit anti-mouse; DAKO) were used at a dilution of 1:800. Visualization was achieved using biotin/avidin-peroxidase (DAKO) and diaminobenzidine as a chromogen.

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Induction of medulloblastomas in *p53*-null mutant mice by somatic inactivation of *Rb* in the external granular layer cells of the cerebellum

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The advertisement features a dark blue background with a glowing DNA double helix structure on the left. The 'horizon' logo is in white, with 'a PerkinElmer company' in smaller white text below it. To the right, the text 'Streamline your research with Horizon Discovery's ASO tool' is displayed in white, with 'Horizon Discovery's ASO tool' in a larger, bold font.