Loss of p53 but not ARF Accelerates Medulloblastoma in Mice Heterozygous for patched1

Cynthia Wetmore, Derek E. Eberhart2 and Tom Curran3

Departments of Developmental Neurobiology [C. W., D. E. E., T. C.] and Hematology/Oncology [C. W.]. St. Jude Children’s Research Hospital, Memphis, Tennessee 38105

Abstract

Brain malignancies represent the most common solid tumors in children, and they are responsible for significant mortality and morbidity. The molecular basis of the most common malignant pediatric brain tumor, medulloblastoma, is poorly understood. Mutations in several genes including the human homologue of the Drosophila segment polarity gene, patched (PTCH), the adenomatous polyposis coli gene (APC), β-catenin, and p53 have been reported in subsets of hereditary and sporadic medulloblastoma. Inactivation of one Ptc allele in mice results in a 14% incidence of medulloblastoma. Here, we report a dramatic increase in the incidence of medulloblastoma in mice heterozygous for Ptc that lack p53. The acceleration of tumorigenesis in Ptc+/− mice is specific for loss of p53, because no change in tumor incidence was observed in Ptc+/− mice carrying a mutation in APC (Min+/−) or in Ptc+/− mice deficient in p19ARF. Thus, there is a specific interaction between p53 loss and heterozygosity of Ptc that results in medulloblastoma. These findings may enhance the rate of acquisition of secondary mutations. Ptc+/−;P53−/− mice provide a useful model for investigation of the molecular bases of medulloblastoma and for evaluation of the efficacy of therapeutics in medulloblastoma.

Introduction

Medulloblastoma arises from the primitive neuroectoderm in the posterior fossa of children generally between the ages of 3 and 9. The tumors are derived from cerebellar granule precursor cells that undergo a dramatic proliferative expansion during the early phases of postnatal brain development. Several gene mutations have been described in medulloblastoma, although they occur in small subsets of tumors (1). A low frequency of mutations in patched (PTCH; Ref. 2), p53 (3), the adenomatous polyposis coli (APC) gene (4), and β-catenin (5, 6) have been reported in subsets of sporadic medulloblastoma. In addition, brain tumors, including medulloblastoma, have been reported to occur more frequently in patients carrying germ-line mutations in PTCH, APC, or p53 (7–9). In the case of heterozygous loss of PTCH, which is associated with Gorlin syndrome, also known as basal cell nevus syndrome (OMIM 109400), the incidence of medulloblastoma increases from 2 per million to 4 per hundred in children <18 years of age (10, 11). Thus, it is likely that there are several unknown prevalent mutations in these tumors that contribute to disease progression.

Progress in understanding the etiology of medulloblastoma has been hampered by the lack of an appropriate animal model. Recently, a mouse strain was generated in which Ptc was mutated by targeted disruption (12, 13). Heterozygous deletion of p53 results in embryonic lethality, whereas mice heterozygous for Ptc exhibit several features of Gorlin syndrome, including an increased propensity to develop tumors in the brain and soft tissues. Histological analysis of the brain tumors showed that they closely resemble human medulloblastoma (12, 14). However, only 14% of mice heterozygous for Ptc develop medulloblastoma over a period of 10 months, indicating that it is likely that additional genetic lesions are required for oncogenic transformation.

Ptc functions as a component of the receptor complex that transduces a signal from Hedgehog (Hh) through a complex pathway that was first described in Drosophila (7). The interaction of Shh, the mammalian orthologue of Hh, with Ptc relieves suppression of smoothed (Smo), resulting in increased transcription of Gli1 and other target genes (7). During cerebellar development, Shh, produced by Purkinje cells, functions as a mitogen to stimulate proliferation of granule cell precursors (15). Ptc does not function as a classic tumor suppressor gene in medulloblastomas in Ptc+/− mice because the normal allele is not lost, and it continues to be expressed in tumors (14, 15). p53 functions as a transcription factor that transduces signals elicited by physiological stress and DNA damage to regulate cell proliferation and apoptosis. Abrogation of p53 function attenuates both of these responses (17). The mouse tumor suppressor gene p19ARF (p14ARF in humans) is the product of an alternative reading frame encoded by the INK4a-ARF locus. ARF functions as a sensor of normal proliferative signals upstream of p53 by interfering with Mdm2, a negative regulator of p53 function (18). Thus, loss of p19ARF diminishes p53 activity and promotes tumor formation (19). Mice deficient in p53 do not develop brain tumors, although they are predisposed to develop tumors in several other tissues by 5 months of age (20, 21). Approximately 10% of ARF-null mice develop glial tumors by 6 months of age (19).

To address the possible involvement of tumor suppressor genes in medulloblastoma and to accelerate the incidence of these tumors, we crossed Ptc+/− mice with mice carrying mutations in other tumor suppressor genes. We selected APC because it regulates the levels of β-catenin, which functions in the Wnt signaling pathway (22). Humans with brain tumor-polyposis, or Turcot’s syndrome, carry germline mutations in APC, and they have an increased incidence of tumors arising in colon and brain (8). In addition, mutations in β-catenin have been reported in spontaneous medulloblastoma, albeit at a low frequency (5, 6). We also crossed the Ptc+/− mice with mice carrying inactivating mutations in two major tumor suppressor genes that are defective in more than half of all human cancer, p53 and ARF (18). These genes serve critical functions in the repression of cell proliferation, apoptosis, and response to DNA damage (18, 23).
Materials and Methods

Animals. The Ptc+/− mice used in this study were generated and maintained on a mixed C57BL/6 × 129Sv background, as described previously (14), and crossed with mice carrying targeted disruptions in p53, APC (C57BL/6J-Min+/−; Jackson Laboratories, Bar Harbor, ME), and ARF (19) to generate the following cohorts of mice: Ptc+/− p53+/+ (n = 440), Ptc+/− p53+/− (n = 68), Ptc+/− p53−/− mice (n = 40), Ptc+/− ARF−/− (n = 40), and Ptc+/− Min−/− (n = 16). Cohorts of mice were observed for tumor formation for a minimum of 6 months after birth. All mice were observed daily for signs of increased intracranial pressure and for evidence of enlarged occipital prominence three times weekly for at least 24 weeks. Animals were euthanized when they were moribund according to NIH-approved institutional guidelines or when they showed signs of increased intracranial pressure or when extracranial tumors were evident. Brains were removed from the surrounding calvarium, and tumor tissue was carefully separated from surrounding brain parenchyma under a dissecting microscope. In every mouse, the presence of tumor was confirmed by gross examination of the brain. If the mouse was not available for examination or if no tumor was detected, the cause of death was attributed to “unknown causes.” Fresh tissue was snap frozen and stored at −80°C for later extraction of RNA, DNA, and protein. For histochemical analyses, animals were deeply anesthetized and perfused transcardially with 4% paraformaldehyde in 0.1 M PBS and processed for immunohistochemical analyses as described previously (14).

RNA Isolation and Northern Analysis. Total cellular RNA was isolated from 11 mouse medulloblastomas using Trizol (Ambion, Inc., Austin, TX) according to the manufacturer’s directions. Five to 10 μg of total RNA were electrophoresed on a 1.8% agarose-formaldehyde gel, transferred to a nitrocellulose filter (Hybond N +; Amersham Pharmacia; Buckinghamshire, United Kingdom), and hybridized under stringent conditions (18 h at 68°C in 5 × SSPE, 50% formamide, 5 × Denhardt’s solution, 1% SDS, and 0.1 mg/ml denatured salmon sperm DNA) with a 32P-labeled RNA probe. Filters were washed (twice × 20 min in 0.1 SSC, 0.1% SDS at 68°C) and exposed to film (Eastman Kodak) for 12–72 h at −80°C. Control and tumor tissues were analyzed by hybridization with 32P-labeled RNA probes specific for mouse Ptc (12), Gli1 (mouse EST clone 38654), and mdm2.

Immunoblot Analysis. Protein extracts were prepared by Dounce homogenization of 80–100 mg of snap-frozen tumor or normal tissue as described (14). Extracts were clarified by microcentrifugation at 14,000 rpm for 30 min. Protein lysates (200 μg) from medulloblastoma arising in Ptc+/− mice (tumor nos. 185, 199, 241, 448, 530, and 574), mouse leukemia cells known to express mutated (CR246) or wild-type P53 (CR205), medulloblastoma from a Ptc+/− p53−/− mouse (1138), and normal adult mouse brain were separated on 10% polyacrylamide gels and transferred to nitrocellulose membranes. The membranes were incubated with anti-p53 antisera (Ab7; Oncogene; 1:5000), followed by donkey antiserum IgG-horseradish peroxidase (Chemicon; 1:2500) diluted in 5% evaporated milk powder in 1% TBST [50 mM Tris-Cl, 0.15 M NaCl (pH 8.0) with 1% Tween]. The signal was detected by enhanced chemiluminescence. The membranes were stripped and incubated with antibodies directed against Ref-1 and β-tubulin to control for protein loading and transfer efficiency.

RT-PCR. Two-step RT-PCR was carried out to maximize uniformity of PCR templates for all reactions. cDNA was derived in 20-μl volumes with random hexamers, oligo dT, and gene-specific priming using SuperScript reverse transcriptase (Life Technologies, Inc., Rockville, MD). The reverse transcriptase first-strand cDNA synthesis reactions were carried out using 3 μg of total RNA prepared from adult C57BL/6 mouse cerebellum and from seven tumor samples (tumor nos. 185, 199, 241, 448, 530, 574, and 646) according to the manufacturer’s directions. Gene-specific oligonucleotides corresponding to sequences within the open reading frame of p53 were synthesized, and PCR amplification of overlapping regions was performed to generate templates for nucleotide sequencing. Sequence analysis of PCR products generated from both the sense and antisense strands of p53 were analyzed from two separate cDNA templates and from multiple PCR reactions.

Nucleotide Sequencing. Sequencing reactions were performed by the Hartwell Center for Biotechnology at the St. Jude Children’s Research Hospital on template DNA using rhodamine or dRhodamine dye terminator cycle sequenc-
Gli1 is normally repressed by Ptc, and in adult, nonproliferating tissues, Gli1 expression is not readily detected. However, during development, when cells are rapidly proliferating, repression of Gli1 transcription is abrogated by the interaction of Shh with the Ptc/Smo receptor complex (7). When Ptc is mutated or absent, intrinsic signaling by Smo is not suppressed, resulting in increased transcription of Gli1 and other downstream genes. A single copy of Ptc was sufficient to maintain repression of Gli1 in the non-tumor containing Ptc+/− cerebellum, because no increases in Gli1 levels were detected in control brain tissues compared with the dramatic increases in Gli1 mRNA seen in the tumors (Fig. 2A). However, in medulloblastomas that arise in Ptc−/− mice, there is persistent expression of the wild-type Ptc allele, suggesting that derepression of Gli1 expression does not require complete loss of Ptc (14). These data indicate that genes other than Ptc may influence Gli1 expression in the cerebellum. Expression of p53 mRNA was found to be elevated in all tumors (Fig. 2B). This is consistent with prior reports of elevated p53 expression in populations of rapidly dividing cells during development and tumorigenesis (23, 24).

No Acceleration of Medulloblastoma Formation Was Observed in Ptc+/Min+/− Mice. Only 1 of 16 of these mice (6%) developed a posterior fossa tumor by 22 weeks of age. This tumor was phenotypically similar to the medulloblastomas found in Ptc+/− mice. These data suggest that genetic lesions in the PTCH and APC pathways found in subsets of sporadic human medulloblastoma do not act synergistically in mice.

p53 Loss Is Not Required for Medulloblastoma Formation in Ptc+/− Mice. The dramatic acceleration of medulloblastoma formation in Ptc+/− p53−/− mice prompted us to investigate the status of p53 in tumors arising in Ptc−/− mice in which there is no germ-line mutation of p53. Interestingly, these tumors contained high levels of p53 mRNA compared with control tissues. In contrast, there was no consistent difference in Mdm2 mRNA levels between normal and tumor tissues (Fig. 2B). The Mdm2 gene product acts to repress p53 activity, and amplification of Mdm2 inactivates p53 in a subset of astrocytomas (25, 26). Wild-type p53 protein has a short half-life, and it is not readily detected in populations of nonproliferating cells unless it has been stabilized by mutation (23). Therefore, we performed immunoblotting analysis to look for evidence of p53 inactivation in tumors from Ptc−/− mice. As shown in Fig. 3, despite the increase in p53 mRNA, p53 protein was present at significantly lower levels in medulloblastomas from Ptc−/− mice, compared with those observed in a mouse lymphoma with a E254G substitution mutation in p53 (CR246). However, expression of p53 protein and mRNA were higher in the tumors than in control brain tissue, which contains relatively few proliferating cells (Figs. 2A and 3). Nucleotide sequence analysis of p53 revealed no mutations in any of the seven tumor mRNAs examined. Thus, although germ-line loss of p53 accelerates tumorigenesis in Ptc−/− mice, mutation of p53 is not required for medulloblastoma formation. This contrasts with a report of increased medulloblastoma formation in mice carrying homozygous mutations in both the retinoblastoma (Rb) and p53 genes. In this mouse model, brain tumors were not detected in mice in which only one of these genes was disrupted (27).

No Accelerated Tumor Formation Was Noted in Ptc+/−p53+/− or in Ptc+/−ARF−/− Mice. The lack of acceleration of medulloblastoma in Ptc+/− p53+/− mice may be attributed to the very limited time window in which the presumed tumor precursor cells are proliferating. Granule cell precursors undergo rapid expansion in the external germinal layer of the cerebellum during the first 2 weeks of postnatal life.
In mice, these cells differentiate and migrate to their mature positions in the internal granular layer by the third week of postnatal development (28). In humans, this process is completed by the ninth postnatal month (29). Thus, Ptc\(^{+/−}\) precursor cells have a limited number of cell divisions in which to acquire the additional mutation(s) that contribute to medulloblastoma formation.

Loss of \(p53\) leads to accumulation of cytogenetic abnormalities (23, 30). Indeed, we observed a much higher incidence of random chromosome loss in tumors from Ptc\(^{+/−}\)/p53\(^{−/−}\) mice compared with those from Ptc\(^{−/−}\)/p53\(^{−/+}\) mice. In p53\(^{+/−}\) mice, there may be an insufficient number of cell generations to lose the remaining p53 allele and to acquire other genetic changes. Additionally, no acceleration in tumorigenesis was noted in Ptc\(^{+/−}\)/ARF\(^{−/−}\) mice. This may be because ARF does not increase genomic instability, and therefore, the tumor precursor cells may be less prone to sustain DNA damage than cells deficient in p53. It is likely that the genomic instability associated with complete loss of p53 function accelerates the mutation rate in granule cell precursors. This may synergize with the effects of reduced Ptc expression in these mice to increase the incidence of medulloblastoma.

Survivors of pediatric brain tumors have significant morbidity as a direct consequence of the therapy required to eradicate tumor cells from the developing brain of a child. Genetic mutations have been detected only in small subsets of medulloblastoma, and the molecular basis of the majority of these tumors remains to be elucidated. The high frequency and rapid onset of tumors in Ptc\(^{+/−}\)/p53\(^{−/+}\) mice provide a useful model to investigate other molecules that influence the balance between proliferation and cell death in the nervous system.

Acknowledgments

We thank M. Scott and L. Goodrich for the Ptc\(^{+/−}\) mice and Ptc plasmids (617 and M2-3); G. Zambetti for mdm2 plasmid; C. Sherr for the ARF\(^{−/−}\) mice; S. Mathew and J. Dalton for karyotype analysis; M. Connelly for assistance with tumor cell culture; and C. Eischen for mouse lymphoma cell lysates.

References

Loss of \textit{p53} but not \textit{ARF} Accelerates Medulloblastoma in Mice Heterozygous for \textit{patched}

Cynthia Wetmore, Derek E. Eberhart and Tom Curran


Updated version
Access the most recent version of this article at:
http://cancerres.aacrjournals.org/content/61/2/513

Cited articles
This article cites 27 articles, 12 of which you can access for free at:
http://cancerres.aacrjournals.org/content/61/2/513.full.html#ref-list-1

Citing articles
This article has been cited by 65 HighWire-hosted articles. Access the articles at:
/content/61/2/513.full.html#related-urls

E-mail alerts
Sign up to receive free email-alerts related to this article or journal.

Reprints and Subscriptions
To order reprints of this article or to subscribe to the journal, contact the AACR Publications Department at pubs@aacr.org.

Permissions
To request permission to re-use all or part of this article, contact the AACR Publications Department at permissions@aacr.org.