Abnormal Expression of REST/NRSF and Myc in Neural Stem/Progenitor Cells Causes Cerebellar Tumors by Blocking Neuronal Differentiation

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Medulloblastoma, one of the most malignant brain tumors in children, is thought to arise from undifferentiated neural stem/progenitor cells (NSCs) present in the external granule layer of the cerebellum. However, the mechanism of tumorigenesis remains unknown for the majority of medulloblastomas. In this study, we found that many human medulloblastomas express significantly elevated levels of both myc oncogenes, regulators of neural progenitor proliferation, and REST/NRSF, a transcriptional repressor of neuronal differentiation genes. Previous studies have shown that neither c-Myc nor REST/NRSF alone could cause tumor formation. To determine whether c-Myc and REST/NRSF act together to cause medulloblastomas, we used a previously established cell line derived from external granule layer stem cells transduced with activated c-myc (NSC-M). These immortalized NSCs were able to differentiate into neurons in vitro. In contrast, when the cells were engineered to express a doxycycline-regulated REST/NRSF transgene (NSC-M-R), they no longer underwent terminal neuronal differentiation in vitro. When injected into intracranial locations in mice, the NSC-M cells did not form tumors either in the cerebellum or in the cerebral cortex. In contrast, the NSC-M-R cells did produce tumors in the cerebellum, the site of human medulloblastoma formation, but not when injected into the cerebral cortex. Furthermore, the NSC-M-R tumors were blocked from terminal neuronal differentiation. In addition, counteracting REST/NRSF function blocked the tumorigenic potential of NSC-M-R cells. To our knowledge, this is the first study in which abnormal expression of a sequence-specific DNA-binding transcriptional repressor has been shown to contribute directly to brain tumor formation. Our findings indicate that abnormal expression of REST/NRSF and Myc in NSCs causes cerebellum-specific tumors by blocking neuronal differentiation and thus maintaining the “stemness” of these cells. Furthermore, these results suggest that such a mechanism plays a role in the formation of human medulloblastoma.

Most medulloblastomas are believed to originate from undifferentiated neural stem/progenitor cells (NSCs) present in the external granule layer cells of the cerebellum (15, 18, 22, 41, 50). The primitive “embryonal” appearance of medulloblastoma cells, as well as their capacity for divergent differentiation, has led to the suggestion that they have a neural stem cell-like phenotype (18, 22). In addition to arising from stem cells, human medulloblastomas appear to contain stem-like cells required for tumor propagation (59). Induction of neuronal and glial markers in medulloblastoma cells has been documented as a response to several proposed chemotherapeutic agents (6, 42, 64), supporting the hypothesis that these tumors show lack of terminal differentiation and suggesting regulation of differentiation status as a promising treatment avenue.

Pathways regulating cerebellar development, such as Hedgehog and Wnt, have been found to be activated by genetic alterations during medulloblastoma tumorigenesis (15, 21, 44, 50, 66, 68, 69). Both Hedgehog and Wnt are thought to regulate proliferation and differentiation of neural stem cells and may play a similar role in medulloblastoma. However, mutations activating these pathways have been documented in only a modest percentage of human tumors. The receptor gene PTCH is the member of the Hedgehog pathway most commonly altered in medulloblastoma, but sensitive techniques such as direct sequencing have identified mutations in no more than 10% of cases (14). Similarly, mutations in CTNNB1 and Axin activating the Wnt pathway have been identified in less than 20% of sporadic medulloblastomas (13, 32, 70). Thus, the mechanism of tumorigenesis for the majority of medulloblastomas is still unknown.

The Myc oncogenes are also important in medulloblastoma pathogenesis (1, 7, 26). c-myc and N-myc are commonly amplified in the biologically aggressive large cell/anaplastic...
REST/NRSF and Myc in Medulloblastoma

MATERIALS AND METHODS

Histological and immunohistochemical assays. For the histological studies, 21 surgically excised human brain tumor tissue and adjacent normal tissue samples fixed in 10% buffered formalin and embedded in paraffin were obtained from the Brain Tumor Center tissue bank at The University of Texas M. D. Anderson Cancer Center, stained with hematoxylin and eosin (H&E), and examined under a light microscope, as previously described (19, 65). These experiments were done with institutional review board approval. All specimens had been previously determined to be medulloblastomas on the basis of histological findings. Six normal cerebellar tissue samples adjacent to tumors, which were histopathologically similar in structure to the tumors, were included in the array as negative controls. For the immunohistochemical assays, the brain sections were stained with an anti-REST antibody (a gift of Gail Mandel), an anti-c-Myc antibody (Santa Cruz), and an anti-N-Myc antibody (Santa Cruz), and this was done essentially as previously described (19, 22, 65). The staining intensity was graded as negative (−), weakly positive (+), or strongly positive (+++) and was determined on the basis of the staining intensity of 15 to 20 representative fields from each section.

Real-time RT-PCR. Real-time reverse transcription-PCR (RT-PCR) was performed as follows. Medulloblastoma samples were obtained from The Johns Hopkins Hospital and were snap-frozen at the time of resection. Diagnoses were confirmed by a neuropathologist (C.G.E.). These studies were approved by the local institutional review board. Total RNA was prepared using QIAGEN RNAeasy tissue kits according to the manufacturer’s instructions (QIAGEN Corp., Valencia, CA). An RNase-free DNase (QIAGEN) was used to reduce genomic DNA contamination. Quantification of mRNA levels was performed using a two-step real-time PCR method on a Bio-Rad iCycler (Bio-Rad). Approximately 1,000 ng of total RNA was reverse transcribed to cDNA using random hexamers and Moloney murine leukemia virus reverse transcriptase (Applied Biosystems, Foster City, CA) and then used to determine mRNA levels of β-actin, c-myc, N-myc, and REST in separate reactions. Each unknown was run at least three times, and expression values were generated with lecser software from a standard curve created using 0.1 to 50 ng of cDNA from the medulloblastoma cell line D283 (American Type Culture Collection). All values were then normalized to β-actin content in the cDNA sample and are shown as means with standard error. β-Actin and c-myc levels were determined using commercially available Taqman primers and probes (Applied Biosystems). The sequences of the forward and reverse primers used for SYBR green quantitative PCR were 5′-GAGGAGGAAGGCTGTTTACC-3′ and 5′-TCACAGCACTTGGCATTCA-3′, respectively. For N-myc, the forward and reverse primers were 5′-TGAAGGAAGATGAAAGAAGAAGA-3′ and 5′-GTGACAGCTTGGTGTGGA-3′, respectively. Single melt peaks were obtained from both REST and N-myc SYBR green primer pairs, and sequence analysis confirmed the identity of the products. Statistical analysis of the correlation ( Spearman test) and categorical association (two-sided Fisher’s exact test) between c-Myc and REST was performed using GraphPAD Prism4 (GraphPAD Software, San Diego, CA).

Plasmids and stable transfection. The Nehl/Xhol fragment of pcDNA3.1-REST (28) was subcloned into the Nehl/Xhol-digested plasmid pBl2gr (62). The clone obtained was confirmed by sequencing the junction region. The mouse multipotent C17.2 (NSC-M) cells were derived from neonatal mouse cerebellum transduced with v-Myc. Because v-Myc represents the activated form of the c-Myc oncogene (38), these cells provide an excellent model for our experiments (60). Construction and characterization of stable clonal cell lines NSC-M-V and NSC-M-R were performed as described previously (63).

Cell culture conditions. NSC-M cells were cultured at 5% CO2 at 37°C in either proliferation medium ( Dulbecco’s modified Eagle’s medium containing 10% fetal bovine serum, 5% horse serum, and 2 mM glutamine, and antibiotics/antimycotics [all from Gibco]) or differentiation medium (Neurobasal medium plus B27, antibiotics/antimycotics, and 2 mM glutamine [all from Gibco], plus nerve growth factor [Chemicon], at a final concentration of 200 ng/ml). For NSC-M-V or NSC-M-R stable clonal cells, the media were supplemented with 200 μg/ml hygromycin (Roche) to maintain the presence of the transgene and.
when needed, 20 μg/ml doxycycline (Sigma) to repress the human REST transgene expression. NSC-M, NSC-M-V, and NSC-M-R cells were propagated in proliferation medium on uncoated tissue culture dishes. For the assay of cell proliferation or differentiation, the cells were seeded on polylysine (Sigma)- and laminin (Invitrogen)-coated dishes in the presence of proliferation medium. Once the cells attached to the dish (approximately 2 h), they were processed either immediately (proliferation condition, day 0) or at different time points after the medium was replaced by differentiation medium (differentiation conditions, day 2, 4, or 6).

RT-PCR assay. The RNaseasy kit from QIAGEN was used to prepare total RNA from the cells. RNAs were quantitated spectrophotometrically, and 200 ng of total RNA was used for each sample to detect REST/NRSF in the RT-PCR analysis. Primers used in these reactions had the following sequences: mActin F 5'-CCAGGAGCTTCTTCCAT-3'; mActin R 816-841, 5'-CCCTCGGTTCGCAAGTGTGAC-3'; hREST F 3311–3333, 5'-GACGACACAGCTGCTGCTG-3'; hREST R 3334–3354, 5'-GAGACACAGCTGCTGCTG-3'; and mREST 2817-2792R, 5'-CAGATACGTCTTGGTCATAGTG-3'. An RT-PCR kit (QIAGEN) was used per the manufacturer's instructions to perform the initial reverse transcription. All three primer sets for actin, hREST, and mREST were used in the same tube at the time of reaction. The cDNA was amplified for 20 or 25 cycles under the following conditions: melting at 94°C for 1 min, annealing at 55°C for 1 min, and extension at 72°C for 1 min. The PCR product was separated by electrophoresis on a 2.0% agarose gel at 80 V for 1.5 h.

Reporter gene assays. The REST-dependent promoter repression was determined in NSC-M, NSC-M-V, and NSC-M-R cells growing either in proliferation medium (day 0) or in differentiation medium for 4 days (day 4) by transfection with reporter plasmids pNaCh (+RE) and pNaCh (-RE). Transfected cells were further incubated for 24 h in the same medium, and the assay was performed as described previously (28, 37). The plasmid pSTIuc was used as an internal control (37, 53). The average chloramphenicol acetyltransferase (CAT) activities from three experiments, which were normalized to luciferase activity, were calculated as the percentage of REST activity: 100 - [100 × pNaCh (+RE)/pNaCh (-RE)], where the REST activity for NSC-M at day 0 was taken as 100%.

Immunofluorescence assay. Immunofluorescence assay experiments were performed as described previously (63, 65). We used the following antibodies: anti-pVp6 (1:100) (Clontech), anti-unique β-tubulin (1:500) (Tuj1; Covance Research Products), anti-MAP2 (1:1,000, HM-2; Sigma), anti-synaptophysin 1 (1:200, mab506; Chemicon), anti-gial fibrillary acidic protein (anti-GFAP; 1:500; DAKO), horseradish peroxidase-conjugated anti-mouse or anti-rabbit immunoglobulin G (IgG; H + L, 1:20,000; Amersham), and Cy3-labeled anti-mouse or anti-rabbit IgG (H + L, 1:1,000; Amersham). A Leica epifluorescence microscope was used to examine the staining of cells. Further analysis was carried out using Metamorph software.

Cell proliferation assay. The cells growing either in proliferation medium (day 0) or in differentiation medium for various time points (day 2 or 4) were transfected with 10 μM bromodeoxyuridine (BrDU; Sigma) and then washed twice with 1× PBS and fixed with 10% buffered formalin for 20 min at room temperature. The immunofluorescence of BrDU-labeled cells was elicited by first blocking the cells with blocking buffer (1× PBS containing 10% normal goat serum, 10% powdered dry milk, and 0.2% Triton X-100) for 1 h at room temperature. The blocking buffer was removed, and the cells were incubated with 1× PBS containing mouse anti-BrDU antibody (containing nucleases) (Roche) at a dilution of 1:15 for 1 h at 37°C. The cells were washed three times for 5 min each with 1× PBS with gentle agitation. This was followed by incubation with Cy3-conjugated goat anti-mouse antibodies (Amersham) at a dilution of 1:1,000 for 1 h at 37°C. The cells were washed three times with 1× PBS with gentle agitation and allowed to air dry. The stained cells were covered with slow-fade antifade (Molecular Probes) containing 1 μg/ml 4',6'-diamidino-2-phenylindole (DAPI; Molecular Probes). The cells were analyzed by immunofluorescence assay as described above. The average values from five different fields were used for each data point. Each experiment was repeated four times.

Western blotting assay. Cells were seeded at a density of 5 × 10⁵ cells per 15-cm petri dish coated with 0.1 mg/ml polylysine and 0.1 mg/ml of laminin and grown in 5 ml Neurobasal medium (GIBCO, Bethesda, MD) supplemented with B27 and nerve growth factor (Chemicon) at a final concentration of 200 ng/ml and 1% (v/v) of 5% CO2 tension. In some experiments, replication incompetent adenovirus expressing either green fluorescent protein (GFP) or REST-VP16 (Ad.GFP and Ad.REST-VP16, respectively) was used to infect these cells at a multiplicity of infection of 100 for 4 h. Cells were either immediately harvested (day 0), or the volume of medium was then raised to 15 ml and cells were incubated in the differentiation medium further for 3 days postinfection. Infection efficiency was determined by counting the number of GFP-positive cells and was found to be >50%. Cells, adenovirus infected or uninfected, were harvested and lysed by addition of Laemmli buffer, and cell extracts were subjected to Western blot analysis using mouse anti-PARP antibody (BD Pharmingen) at a dilution of 1:500. The antibody recognizes an 85-kDa product from mouse cells. In some experiments, anti-neuronal β-tubulin (Tuj1, 1:500 dilution; Covance) or anti-α-tubulin (1:500 dilution; Covance) was also used.

Intracranial inoculation of cells into mice and assay for tumor formation. The intracranial inoculation of cells into nude mice was performed following "M. D. Anderson Institutional Animal Care and Use Guidelines." Cells (2 × 10⁵ cells in 5 μl of cell growth medium) were inoculated into groups of mice by using an implantable guide-screw system we have described elsewhere (19, 35, 65). The mice were sacrificed by CO2 inhalation 6 weeks later, and their brains were fixed with formalin and embedded in paraffin. 4- to 5-μm brain sections were examined in the same way as the human medulloblastoma brain specimens.

RESULTS

Many primary human medulloblastomas express both REST/NRSF and c-Myc proteins. As described above, many human medulloblastoma tumor specimens and cell lines have been found to express c-Myc or REST/NRSF. However, because neither protein by itself can cause tumor formation, we wondered whether human medulloblastoma tumor samples expressed both Myc and REST/NRSF proteins. We used immunohistochemical analysis to examine 21 surgically excised, formalin-fixed, paraffin-embedded tumor tissue specimens organized in a tissue array with anti-REST, anti-c-Myc, and anti-N-Myc antibodies. The results are shown in Fig. 1 and summarized in Table 1. As expected, normal cerebellar cortex tissue was composed of granular cell neurons with uniform, round nuclei and interspersed pale neuropil islands (glomeruli), whereas medulloblastoma tissue was composed of densely packed cytologically atypical tumor cells with enlarged, irregular, hyperchromatic nuclei (Fig. 1). All six normal control cerebellar tissues stained negatively for anti-REST/NRSF, anti-c-Myc, and anti-N-Myc antibodies. However, 17 of 21 medulloblastoma tumor samples stained positively for REST/NRSF, with strong signals (+ +) in 6 and weak signals (+) in 11. The remaining four samples showed no REST/NRSF signal. In contrast, 14 of the 21 tumor samples stained positively for c-Myc, with strong signals (+ +) in 9 and weak signals (+) in 5. The remaining five samples showed no c-Myc positivity. Similarly, 16 of the 21 tumor samples stained positively for N-Myc, with strong signals (+ +) in 6 and weak signals (+) in 10. The remaining seven samples showed no c-Myc positivity. High-magnification (×1,000) photomicrographs showed the nuclear staining for REST, c-Myc, and N-Myc (Fig. 1). Thus, whereas 52% samples showed coexpression of both N-Myc and REST/NRSF proteins at various levels (11 of 21), strong signals for both c-Myc and REST/NRSF were observed in only 14% (3 of 21) of samples. Likewise, 48% of samples (10 of 21) showed coexpression of REST/NRSF, c-Myc, and N-Myc proteins, with strong signals in 14% (3 of 21) of samples. Finally, 48% of the samples expressed REST/NRSF, c-Myc, and N-Myc proteins. It is worth noting that in two cases on Table 1, REST and N-myc are coexpressed in the absence of detectable c-Myc protein. This suggested the possibility that either c- or N-Myc is capable of cooperating with REST and was supported by our data shown below (Fig. 2). Although double-labeling studies were not performed, the fact that the majority of cells in a given
tumor were immunopositive for both REST and Myc suggests that the proteins were being coexpressed in individual nuclei. To obtain more quantitative data, we used real-time RT-PCR to investigate the association between REST and Myc transcript levels. Expression was analyzed in mRNA extracted from 40 snap-frozen primary medulloblastomas for which we had previously measured c-Myc and N-Myc expression levels (D. Stearns and C. G. Eberhart, unpublished data). REST was detected in all samples, and many of the tumors with high REST levels were also among those with elevated c-Myc expression (Fig. 2). REST and c-Myc mRNA levels showed a statistically significant positive correlation overall (Spearman $r = 0.36; P = 0.02$). However, several cases with high REST levels contained relatively little c-Myc. We were curious whether N-myc might substitute for c-Myc in these cases, as N-myc gene amplification has also been associated with aggressive medulloblastoma biology (16). Interestingly, four cases with high REST and low c-Myc had above-median levels of N-Myc (Fig. 2, inset). Thus, high-level REST expression appears to be commonly associated with elevated expression of one of these two myc oncogenes. As a final measure of the association between c-Myc and REST, we used Fisher’s exact test to determine whether tumors with very high (top third) REST expression also had very high (top third) c-Myc expression. The combined high-

\[ \text{TABLE 1. Status of immunoreactivity in medulloblastoma patient samples} \]

<table>
<thead>
<tr>
<th>Age (yr)</th>
<th>REST/NRSF</th>
<th>c-Myc</th>
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<tr>
<td>7</td>
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\[ ^a \text{–, negative; +, moderately positive; ++, strongly positive.} \]
REST/high-c-Myc category (33%) was significantly more common than would be predicted by chance association ($P = 0.04$), supporting the concept that these genes might act together in medulloblastoma pathogenesis.

Construction of NSC clones stably expressing Myc and doxycycline-regulated human REST/NRSF transgene. We used cerebellum-derived, neonatal mouse brain NSCs stably expressing an activated form of c-Myc (NSC-M cells) to further study the functional roles of Myc and REST in medulloblastoma tumorigenesis (60, 61). We generated stable clonal cells expressing either the vector (NSC-M-V cells) or human REST transgene (NSC-M-R cells) by using the bidirectional doxycycline-regulated pBig2r vector system by a method we described previously (63, 65). Selected clones were assayed using PCR to verify the appropriate clonal cells (data not shown). The pBig2r construct efficiently expressed the encoded transgene in the absence of doxycycline ($Dox$), and the cells of the clonal cell line NSC-M-R stably expressed high levels of the human REST transgene mRNA when grown without doxycycline ($-Dox$) (data not shown). The pBig2r construct efficiently expressed the encoded transgene in the absence of doxycycline ($-Dox$), and the cells of the clonal cell line NSC-M-R stably expressed high levels of the human REST transgene mRNA when grown without doxycycline ($-Dox$) (data not shown). As expected for normal NSCs (2, 3), NSC-M cells and NSC-M-V cells expressed the endogenous mouse REST (mREST) transcripts only under proliferation conditions ($d0$), and the expression ceased upon neuronal differentiation ($d2, d4, and d6$) as assayed by RT-PCR (Fig. 3A). NSC-M-R cells followed the same expression pattern for mREST but, in contrast, expressed the human REST transgene (hREST) under both proliferation and differentiation conditions (Fig. 3A).

To confirm that the expression pattern of REST/NRSF transcript levels during neuronal differentiation shown in Fig. 3A was also reflected in its transcriptional repression activity, we used a reporter gene system that is well established to detect REST/NRSF-specific activity in the cell (11, 28, 37, 65). The system involves comparison of reporter gene activities from two plasmids. The first, pNaCh (+RE), contains the bacterial CAT reporter gene under the control of sodium channel type II promoter/enhancer (NaCh) elements. NaCh elements are mammalian native neuronal promoter/enhancer sequences that contain the REST/NRSF binding site (RE) and are acted on by functional REST/NRSF. The second reporter plasmid, pNaCh (–RE), contains the CAT reporter gene under the control of the “minimal” NaCh elements without the RE1 sequence (–RE). Functional REST/NRSF does not repress the promoter activity of this plasmid. Thus, when functional REST/NRSF is present in the cell, the CAT activity generated from pNaCh (+RE) is lower than that from pNaCh (–RE) because REST/NRSF causes promoter repression from pNaCh (+RE), and not pNaCh (–RE).

The NSC-M and NSC-M-V cells, grown under either cycling conditions (day 0) or differentiation conditions (day 4), were transiently cotransfected with the reporter plasmid pNaCh (+RE) or pNaCh (–RE) and an internal control plasmid, pSTluc, which contains the luciferase reporter gene under the control of an Sp1-containing basic promoter (36, 43). The average CAT activities from three experiments, which were normalized to luciferase activity, were rendered as the percentage of REST activity, as described in Materials and Methods. As shown in Fig. 3B, whereas undifferentiated cycling NSC-M and NSC-M-V cells grown with or without doxycycline showed high levels of REST/NRSF activity (day 0), the differentiation
conditions (day 4) resulted in severely reduced REST/NRSF activity, indicating that REST/NRSF activity is lost during the neuronal differentiation of NSCs.

When NSC-M-R cells were cultured with doxycycline (Fig. 3B), they showed a pattern of REST/NRSF activity similar to that of NSC-M and NSC-M-V cells; that is, activity was high in cycling cells (day 0) and diminished in differentiating cells (day 4). In contrast, when NSC-M-R cells were cultured without doxycycline, they showed higher REST/NRSF activity than did NSC-M cells and NSC-M-V cells under cycling conditions (day 0) and continued showing high REST activity under differentiation conditions (day 4), suggesting that the human REST transgene was responsible for the continued REST activity in these cells (Fig. 3B). Thus, whereas all cells expressed REST/NRSF activity under proliferation conditions, only the NSC-M-R cells (Dox) expressed the REST/NRSF activity under differentiation conditions.

**Forced expression of REST/NRSF in NSCs blocks neuronal differentiation.** To determine the effect of continued REST/NRSF expression in NSC-M-R cells during neuronal differentiation (day 4), we examined whether the terminal neuronal differentiation markers neuronal β-tubulin and MAP2 were expressed in clonal cells growing under differentiation conditions with and without doxycycline. As shown in Fig. 4, the NSC-M-V cells, like the NSC-M cells, expressed both markers irrespective of whether they were cultured with or without doxycycline. NSC-M-R cells cultured with doxycycline also expressed these markers. In contrast, NSC-M-R cells cultured without doxycycline did not express either marker (Fig. 4). The pattern of expression of other neuronal differentiation genes, synapsin and secretogranin II, was similar (data not shown). Another REST/NRSF clone, NSC-M-R’, behaved very similarly to the NSC-M-R cells in all our experiments (data not shown), indicating that the effect of REST/NRSF was independent of its genomic integration site. Taken together, these results indicate that proliferating NSCs normally express endogenous REST/NRSF, but its activity is rapidly blocked as the cells enter neuronal differentiation. These cells then become mature neurons. In contrast, NSC-M-R cells continue to express REST/NRSF as they enter neuronal differentiation and, as a result, cannot become terminally differentiated neurons.

**REST/NRSF-mediated blockade of neuronal differentiation confers a proliferation advantage in vitro.** Because forced expression of REST/NRSF in NSCs blocked neuronal differentiation, to determine whether expression of REST/NRSF in NSC-M-R cells also confers a proliferative advantage, we measured BrdU incorporation into NSC-M cells and NSC-M-R cells under in vitro conditions during cycling (day 0) and differentiation (days 2 and 4). By culturing the cells either with or without doxycycline in the medium, we also performed these experiments in the presence and absence of exogenously expressed REST/NRSF. Figure 5A shows the results of the immunocytochemical assay for BrdU uptake, and Fig. 5B shows the percentages of BrdU-positive cells. As expected, the rate of BrdU incorporation into NSC-M cells cultured either with or without doxycycline, decreased drastically as they differentiated. The rate at which BrdU was incorporated into NSC-M-R cells assumed the same pattern when the expression of the
exogenous REST/NRSF was prevented by culturing the cells with doxycycline. In contrast, the rate at which BrdU was incorporated into NSC-M-R cells remained high when the expression of the exogenous REST/NRSF was kept steady by culturing the cells without doxycycline. Thus, although c-Myc is known to cause neuronal progenitor cell proliferation, these experiments indicated that, by blocking differentiation, the additional REST/NRSF expression conferred an enormous proliferative advantage on NSC-M-R cells compared with NSC-M and NSC-M-V cells, but only when they traveled down the neuronal differentiation pathway.

**FIG. 4.** Forced expression of REST/NRSF in NSCs blocks neuronal differentiation. NSC-M, NSC-M-V, and NSC-M-R cells were cultured with (+Dox) or without (−Dox) doxycycline under neuronal differentiation conditions for 4 days, and their level of neuronal differentiation was determined in an immunofluorescence analysis in which anti-neuronal β-tubulin (Tuj1) and anti-MAP2 antibodies were used.

REST/NRSF-mediated blockade of neuronal differentiation in vivo causes tumors in the cerebellum, but not in the cerebral cortex. Because the cerebellum is the major site of medulloblastoma tumorigenesis (31, 52), we wanted to determine the properties of NSC-M-R cells in mouse cerebellum. We injected NSC-M, NSC-M-V, or NSC-M-R cells into the cerebellums of nude mice (10 mice per group). We sacrificed the mice 6 weeks later and performed histological analysis of their paraffin-embedded brains. As shown in Fig. 6 and 7, the NSC-M and NSC-M-V cells injected into the cerebellum integrated into the cerebellum within about 4 weeks in a nontumorigenic, cytoarchitecturally appropriate manner (10 out of 10 mice for both NSC-M and NSC-M-V cells). Histological analysis of the brains of these mice showed that they had a normal cerebellar architecture with an appropriate vermiculiform histology, including normal molecular, Purkinje cell, and granule cell layers, with granule cell neurons showing the characteristic monotonous, regular, round nuclei. In contrast, large cerebellar tumors with irregular, pleomorphic, and hyperchromatic nuclei developed in mice inoculated with NSC-M-R cells (8 out of 10 mice), and these tumors were morphologically similar to human medulloblastomas (see Fig. 1). In most cases, the vermiculiform cerebellar cortex, including the granule cell layer, was compressed as a result of tumor expansion. Thus, NSCs that overexpressed activated c-Myc alone (NSC-M and NSC-M-V cells) did not produce cerebellar tumors in mice but did produce tumors when they also overexpressed REST/NRSF, indicating that the overexpression of both activated c-Myc and REST/NRSF in NSCs produces cerebellar tumors.

To determine whether tumor formation is location specific, we injected NSC-M, NSC-M-V, or NSC-M-R cells into the cerebral cortex of nude mice as described above. As shown in Fig. 6, none of the cells formed tumors, suggesting that the cerebellar environment facilitates c-Myc-REST/NRSF-mediated tumor formation.

REST/NRSF-mediated cerebellar tumors are blocked from terminal neuronal differentiation. To confirm that the cells that made up the cerebellar tumors produced by NSC-M-R cells did not differentiate in the glial pathway, we examined the expression of GFAP in paraffin-embedded sections. As shown in Fig. 8, NSC-M-V cells did not produce any GFAP. These results indicate that the abnormal expression of REST/NRSF in these tumors prevented the cells from undergoing terminal neuronal differentiation.

The gene encoding synaptophysin, another marker of terminal neuronal differentiation, is also a direct target of REST/NRSF (63, 65). To further confirm that the tumor-producing cells did not express REST/NRSF target genes, we examined paraffin-embedded sections for immunohistochemical expression of synaptophysin (SYP). NSC-M-V cells produced synaptophysin at levels indistinguishable from those of background brain structures. In contrast, NSC-M-R tumors did not produce any MAP2. These results indicate that the abnormal expression of REST/NRSF in these tumors prevented the cells from undergoing terminal neuronal differentiation.

To confirm that the tumor-producing cells did not differentiate in the glial pathway, we examined the expression of GFAP in paraffin-embedded sections with anti-GFAP antibo-
ies. NSC-M-V cells did not show any difference in GFAP staining from those of background brain structures (data not shown). As shown in Fig. 8, strong cytoplasmic reactivity for GFAP was seen in astrocytes but the tumor cells did not express GFAP, suggesting that NSC-M-R cells did not undergo glial differentiation.

**Countering REST/NRSF function blocks tumorigenicity.** Previously we found that REST-VP16 counters the effects of endogenous REST/NRSF in human medulloblastoma cells overexpressing REST/NRSF, induces apoptosis, and counteracts the cells’ tumorigenicity (19, 37). To determine whether the NSC-M-R cells respond to REST-VP16 in the same fashion, we infected NSC-M-R cells grown in the absence of doxycycline in vitro with Ad.GFP (control) and Ad.REST-VP16 at a multiplicity of infection of 100. We then looked for the active apoptosis of these cells on the basis of the detection of the anti-PARP antibody, which recognizes only the cleaved form of PARP. As shown in Fig. 9A, REST-VP16 specifically caused

![Diagram](image)

**FIG. 5.** REST/NRSF-mediated neuronal differentiation block confers a proliferation advantage in vitro. The amount of BrdU incorporation was determined by immunocytochemical analysis (A) and was represented as the percentage of cells (NSC-M or NSC-M-R) showing BrdU labeling (B) growing in proliferation medium (d0) or differentiation medium for day 2 (d2) or day 4 (d4) with (+Dox) or without (−Dox), as described in the text.

![Images](image)

**FIG. 6.** REST/NRSF-mediated neuronal differentiation block causes tumors in the cerebellums but not cerebral cortex of mice. NSC-M, NSC-M-R, and NSC-M-R’ (a second clone of NSC-M-R) cells were injected into the cerebellums and cerebral cortex of mice. The mice were sacrificed 6 weeks later, and their paraffin-embedded brain sections were histologically analyzed. Only the NSC-M-R and NSC-M-R’ cells produced cerebellar tumors (arrows).
high levels of apoptosis in NSC-M-R cells but not in NSC-M or NSC-M-V cells. Thus, these experiments showed that NSC-M-R cells followed the same fate in vitro in response to REST-VP16 as human medulloblastoma cells do.

We then inoculated REST-VP16-infected NSC-M-R cells into the cerebellums of mice, sacrificed the mice 6 weeks later, and examined paraffin-embedded sections of their brains. As shown in Fig. 9B, Ad.REST-VP16, but not Ad.GFP, blocked the tumorigenicity of NSC-M-R cells, indicating that, similar to human REST/NRSF-containing human medulloblastoma cells, countering REST/NRSF in NSC-M-R cells can eliminate their cerebellar tumorigenic potential.

The mechanism of REST-VP16-mediated induction of apoptosis in human medulloblastoma cells (19, 37) and NSC-M-R cells is unknown. One possible mechanism is the persistent presence of the opposing REST/NRSF and REST-VP16 activities in these cells, leading to the generation of conflicting signals of stem cell properties and differentiation properties, respectively. To confirm that REST-VP16-mediated apoptosis in NSC-M-R cells is caused by REST-VP16 activity and not by abnormal production of toxic levels of the protein, we examined the expression of the REST-VP16 target gene neuronal β-tubulin in these cells by performing an immunofluorescence assay. Although neither NSC-M nor NSC-M-R cells produced neuronal β-tubulin when infected with Ad.GFP (data not shown), they did produce it when infected with Ad.REST-VP16 (Fig. 9C). To further confirm this observation, we investigated the expression of neuronal β-tubulin in NSC-M and NSC-M-R cells, uninfected or infected with Ad.REST-VP16, by performing the Western blotting assay. As shown in Fig. 9D, only the Ad.REST-VP16-infected cells showed the expression of neuronal β-tubulin. These results indicate that REST-VP16 expressed in NSC-M and NSC-M-R cells as a result of adenoviral infection produced functionally active protein.

**DISCUSSION**

Our results indicate that many human medulloblastomas coexpress abnormally high levels of Myc and REST/NRSF. We also observed that NSCs overexpressing activated Myc and REST/NRSF proteins are blocked in neuronal differentiation and give rise to cerebellar tumors morphologically similar to human medulloblastoma. These cells did not produce tumors in the cortex, indicating a role of the local brain environment in formation of the tumors. Although the environmental difference between the cerebellum and the cortex very likely is because of the difference in trophic factors, the exact nature of such factors is not yet known. We further found that countering the effects of REST/NRSF in NSC-M-R cells caused apoptosis and counteracted the tumorigenic potential of the cells. On the basis of these findings, we propose a model in which efficient medulloblastoma tumorigenesis occurs when two conditions occur in NSCs: overall increased proliferation as a result of Myc expression and lack of differentiation (maintenance of “stemness”) as a result of REST/NRSF expression (Fig. 10).
Normally, when NSCs are signaled to travel down the neuronal differentiation pathway, they rapidly stop transcribing the REST/NRSF gene. Because the REST/NRSF protein is no longer present to block its target terminal differentiation genes, the cells can successfully differentiate into neurons. In medulloblastoma, the Myc and REST/NRSF proteins are abnormally overexpressed. When NSCs are induced to proliferate rapidly by Myc, then travel down the neuronal differentiation pathway, the deregulated REST/NRSF expression cannot be extinguished, leading to an ongoing block of neuronal differentiation. These still-dividing cells abnormally accumulate, thereby initiating medulloblastoma formation.

Our hypothesis is not at odds with the possibility that tumor maintenance and progression then result from the deregulation of other genes, caused by the abnormal physiology of the tumor cells. This hypothesis of tumor formation resulting from the capture of dividing cells at a stage before full differentiation may be a new paradigm explaining the initiation of tumors believed to originate from and resemble undifferentiated or partially differentiated cells, such as medulloblastomas, neuroblastomas, and rhabdomyosarcomas (48). Indeed, such tumors usually occur in children and adolescents, in whom differentiation pathways are probably more active than they are in adults.

Our model of efficient medulloblastoma formation from NSCs as the result of the cooperation of both proliferation advantage and differentiation-block is not at odds with the less-efficient medulloblastoma formation that occurs in the presence of only one of these steps. For example, activation of the Shh pathway as the result of a PTCH gene mutation, which
increases NSC proliferation, is known to occur in approximately 10% of medulloblastoma patients (50, 66). The severity of this disease can be enhanced by addition of c-Myc, another factor promoting NSC proliferation (52). Thus, it is conceivable that medulloblastomas develop as a result of a factor or factors favoring either the proliferation step or the differentiation-block step but that additional factors reinforcing the same step and/or the other step augment tumorigenesis.

Previously, we found that many human medulloblastoma cell lines and human medulloblastoma patient samples abnormally overexpress REST/NRSF as compared with neuronal cells or normal brain cells (19, 37). As expected, the REST/NRSF-positive tumor cells do not express the REST/NRSF target genes, such as synapsin, which, in turn, can be expressed by REST-VP16 (19, 37). That the overexpression of REST/NRSF plays such an oncogenic role in neuronal cells is further supported by studies from other laboratories indicating overexpression of REST/NRSF in several neuroblastoma cells with concomitant repression of neuronal differentiation genes (27, 46). Furthermore, when the neuroblastoma cells were forced to differentiate, they showed a decrease in REST/NRSF expression and an increase in neuronal markers. These studies suggested that the abnormal overexpression of REST/NRSF in neuronal cells, in which REST/NRSF is normally not present, blocks these cells from terminal neuronal differentiation and produces the cancerous phenotype, perhaps by forcing the cells to persist in a stem/progenitor state.

In contrast, several studies also showed that whereas normal bronchial epithelium expressed REST/NRSF activity, several established small cell lung cancer cell lines as well as primary samples showed reduced REST/NRSF activity with concomitant abnormal expression of REST/NRSF target genes, such as the glycine receptor α1 subunit or vasopressin (12, 25, 45). Such a tumor-suppressor function of REST/NRSF was also seen in colorectal cancers (47, 67). These studies suggested that the abnormal lack of REST/NRSF activity in lung and colon epithelial cells, which normally express REST/NRSF and repress expression of neuronal genes, leads to oncogenic conversion of these cells, perhaps by aberrant expression of REST/NRSF target genes resulting in the abnormal neuroendocrine phenotype sometimes observed in these tumors. Thus, REST may have both tumorigenic and tumor-suppressor effects, de-

FIG. 9. Countering REST/NRSF function blocks cerebellar tumorigenicity. (A) Cell extracts prepared from Ad.GFP (G)- or Ad.REST-VP16 (RV)-infected NSC-M, NSC-M-V, and NSC-M-R cells were subjected to Western blot analysis in which antibodies were used that detect cleaved PARP (85 kDa) and α-tubulin (55 kDa). (B) NSC-M-R cells infected with Ad.GFP or Ad.REST-VP16 were inoculated into the mouse cerebellum. The mice were sacrificed 6 weeks later, and their paraffin-embedded brain sections were histologically analyzed. The tumor produced by NSC-M-R cells infected with Ad.GFP is indicated by arrows. (C) NSC-M and NSC-M-R cells infected with Ad.REST-VP16 were examined by immunofluorescence using anti-neuronal β-tubulin (red). The cell nuclei were also labeled with DAPI (blue). (D) NSC-M and NSC-M-R cells, uninfected (-) or infected (+) with Ad.REST-VP16, were examined by Western blotting analysis using anti-neuronal β-tubulin. Anti-α-tubulin was used as an internal control.

FIG. 10. A model for medulloblastoma tumorigenesis triggered by the transcriptional repressor REST/NRSF and Myc. Please see the text for details.
pending on the cell type, which would indicate that REST is a major biological regulator of normal and abnormal development.

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