The protein ENH is a cytoplasmic sequestration factor for Id2 in normal and tumor cells from the nervous system

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Id2 is a natural inhibitor of the basic helix–loop–helix transcription factors and the retinoblastoma tumor suppressor protein. Active Id2 prevents differentiation and promotes cell-cycle progression and tumorigenesis in the nervous system. A key event that regulates Id2 activity during differentiation is translocation from the nucleus to the cytoplasm. Here we show that the actin-associated protein enigma homolog (ENH) is a cytoplasmic retention factor for Id2. ENH contains three LIM domains, which bind to the helix–loop–helix domain of Id proteins in vitro and in vivo. ENH is up-regulated during neural differentiation, and its ectopic expression in neuroblastoma cells leads to translocation of Id2 from the nucleus to the cytoplasm, with consequent inactivation of transcriptional and cell-cycle-promoting functions of Id2. Conversely, silencing of ENH by RNA interference prevents cytoplasmic relocation of Id2 in neuroblastoma cells differentiated with retinoic acid. Finally, the differentiated neural crest-derived tumor ganglioneuroblastoma coexpresses Id2 and ENH in the cytoplasm of ganglionic cells. These data indicate that ENH contributes to differentiation of the nervous system through cytoplasmic sequestration of Id2. They also suggest that ENH is a restraining factor of the oncogenic activity of Id proteins in neural tumors.

Results

The LIM Domains of ENH Bind to the Helix–Loop–Helix (HLH) Domain of Id Proteins. To identify new interactors of Id2 from the nervous system, we performed yeast two-hybrid screening from a human fetal brain cDNA library using full length Id2 as bait. This screening yielded 47 validated cDNA clones corresponding to four different Id2-associated proteins. Among them, 24 clones code for Id2, 13 clones code for the bHLH transcription factor E2-2, eight clones code for the bHLH transcription factor HEB, and two clones code for the PDZ-LIM protein ENH. All Id2 and ENH clones contain an intact HLH domain. This finding is consistent with the essential role of the HLH domain for heterodimerization. The presence of endogenous Id2 is explained by the strong homodimerization ability of Id2 and its abundant expression in the fetal brain (16, 17). The identification of two E proteins, E2-2 and HEB, demonstrated that our screening was capable of identifying specific Id2 interactors. The only two clones that did not contain a HLH domain code for ENH, a member of the Enigma family of LIM domain proteins, a class of proteins associated with the actin cytoskeleton (18–21). Proteins of the Enigma family possess an N-terminal PDZ domain and three LIM domains at the C terminus (Fig. 1A). All members of the PDZ-LIM Enigma family, including ENH, are cytoplasmic proteins that bind to the actin cytoskeleton through direct interaction between the PDZ domain and α-actinin (19, 20). Sequence analysis of the two ENH clones identified in our two-hybrid assay established that both clones retained a C-terminal fragment of ENH (amino acids 461–596) that includes part of the last two LIM domains but lacked the N-terminal region with the basic helix–loop–helix domain of Id proteins (Fig. 1B). To validate the specificity of the binding between ENH and Id2 and identify the domains that mediate this interaction, we used GST fusion proteins and in vitro-translated proteins in pull-down assays. GST-Id2 bound efficiently to in vitro-translated, 35S-labeled full-length Id2.

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Abbreviations: RA, retinoic acid; GNB, ganglioneuroblastoma; sRNA, small interfering RNA; HLH, helix–loop–helix; bHLH, basic HLH; EN, embryonic day n; VZ, ventricular zone; MZ, mantle zone.

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length ENH and to C-terminal ENH deletion constructs that retain two (ENH LIM1–2) or one (ENH LIM1) LIM domains. However, GST-Id2 did not bind to an ENH polypeptide lacking all LIM domains (ENH6LIM) (Fig. 1B Left and Center). GST-Id2 fusion protein carrying a deletion of the HLH domain (GST-ΔHLHId2) failed to bind ENH (Fig. 1B Right). Given the high homology among the HLH domains of the three Id proteins, we asked whether Id1 and Id3 could also bind ENH in this assay. Indeed, both GST-Id1 and GST-Id3 bound to full-length ENH (Fig. 1B Right). To determine whether Id2 binds ENH in vivo, we performed coimmunoprecipitation experiments after transfecting Id2 and Flag-tagged ENH in Cos-1. Anti-Id2 antibodies precipitated full-length Flag-ENH but not Flag-ENH6LIM. Bound proteins were analyzed by autoradiography. (Right) The lane “input” shows in vitro-translated 35S-labeled full-length ENH. Cos-1 cells were transfected with the indicated expression plasmids. Lysates were analyzed directly (Input) or immunoprecipitated with antibodies against Id2 (C) or Flag (D). Immunoprecipitated proteins were analyzed by Western blot for Flag (C) or Id2 (D). NRlg, normal rabbit immunoglobulins.

ENH is Expressed in the Nervous System and Is Induced During Differentiation. The role of ENH has been mostly studied in cardiac and skeletal muscle cells, where ENH has been proposed as a key factor for the integrity of the actin cytoskeleton in differentiated myocytes (19). However, recent reports show that ENH binds to the N-type calcium channel and suggest that a PKC–ENH–calcium channel complex regulates channel activity in neurons (22). Our isolation of ENH from a fetal brain cDNA library suggests that this protein may be implicated in neural development as well. To conduct our study, we raised a polyclonal antibody against a peptide shared by human and mouse ENH. From Western blot experiments, we confirmed that the antibody interacts specifically with exogenously expressed Flag-ENH and endogenous ENH from RA-treated SK-N-SH neuroblastoma cells (Fig. 2A). To test whether ENH is expressed during mouse development, we stained sections from embryonic day 15.5 (E15.5) mouse embryo and RA-treated SK-N-SH neuroblastoma cells treated with RA. (A) Western blot analysis shows specificity of the ENH antibody for endogenous ENH in RA-treated SK-N-SH cells and ectopically expressed Flag-ENH. The ENH band is lost after treatment of the cells with siRNA oligonucleotides to ENH (siENH). SiCtr is a smart-pool siRNA mixture to luciferase (Dharmacon). (B) ENH immunohistochemistry from E15.5 mouse embryo shows expression in neural and muscle tissues. [Magnification: ×20 (Upper) and ×100 (Lower).] (C) Northern Blot analysis of ENH and Id2 in SH-F and SH-N cells treated with RA for the indicated times. 28S RNA is lost as a loading control. (D) Lysates from parallel cultures were analyzed by Western blot by using ENH and Id2 antibodies. The asterisk indicates a nonspecific band.
Human neuroblastoma cells are frequently used as in vitro models to recapitulate differentiation of the nervous system (23, 24). To ask whether ENH expression is regulated during differentiation of the nervous system we used clonal derivatives of the human neuroblastoma cell line SK-N-SH, the SK-N-SH-N (SH-N) and SK-N-SH-F (SH-F) cells. These cells, which lack N-myc gene amplification, have been used to characterize the cell-cycle exit associated with differentiation of neural cells (25). When treated with a low concentration of RA (0.1 μM) SH-N cells undergo differentiation along the neuronal lineage, whereas SH-F cells acquire an epithelioid morphology and rapidly enter into a senescent-like state. Both cell types arrest in the G₁ phase of the cell cycle within 48 h of treatment with RA (25). Remarkably, RA induced progressive elevation of ENH mRNA and protein in SH-N and SH-F cells, suggesting that ENH may play a role in multiple differentiation pathways in the nervous system (Fig. 2 B and C). Although higher concentrations of RA led to marked inhibition of N-myc and Id2 gene expression in N-myc-amplified neuroblastoma cells (13), we noted that RA at the concentration of 0.1 μM caused little change of Id2 expression in the SK-N-SH derivatives. However, a late decrease of Id2 protein was evident in RA-treated SH-N cells (Fig. 2D).

ENH Is Essential for Cytoplasmic Relocation of Id2 in Neuroblastoma Cells Treated with RA. We sought to ask whether elevation of ENH in RA-treated neuroblastoma cells leads to sequestration of Id2 in the cytoplasm by two independent experimental approaches. First, we examined the subcellular localization of Flag-Id2 after treatment with RA of SK-N-SH using double immunofluorescence staining of endogenous ENH and Flag. Flag-Id2 was predominantly nuclear in untreated cells (Fig. 3A Top Left). In agreement with results shown in Fig. 2, logarithmically growing neuroblastoma cells showed minimal ENH staining (Fig. 3A Middle Left). After treatment with RA, Flag-Id2 relocated to the cytoplasm in cells that had acquired high ENH expression (Fig. 3A Top Right, arrows; see also Middle for expression of ENH in the same cells) but remained nuclear in ENH-negative cells (Fig. 3A Top Right, arrowheads). Quantitative analysis of the subcellular localization of Id2 and ENH from three independent experiments demonstrated that, after treatment with RA, Flag-Id2 relocated to the cytoplasm in ~60% of the ENH-positive cells compared with 10% of the ENH-negative cells and 15% of untreated cells (Fig. 3C). Next we introduced ectopic Flag-ENH in SK-N-SH and examined the subcellular localization of endogenous Id2. Ectopic ENH localized to the cytoplasm with a pattern compatible with actin stress fibers (Fig. 3B Lower Right). As expected, Id2 was mainly nuclear in cells transfected with empty vector (Fig. 3B Left). However, expression of ENH caused translocation of Id2 to the cytoplasm (Fig. 3B Upper Right).

To establish the functional significance of endogenous ENH for the cytoplasmic relocation of Id2 induced by RA, we took advantage of a loss-of-function approach using small interfering RNA (siRNA) oligonucleotides directed to ENH. Transfection of RA-treated SK-N-SH with siRNA targeting ENH resulted in efficient depletion of ENH from these cells (Fig. 2A). Flag-Id2 translocated to the cytoplasm of RA-treated SK-N-SH in the presence of scrambled siRNA oligonucleotides, but the siRNA-mediated silencing of ENH prevented entirely the RA-induced relocation of Flag-Id2 to the cytoplasm (Fig. 4A; see also Fig. 4B for the quantitative analysis of subcellular localization of Flag-Id2 in the experiment shown in A (at least 300 cells were scored for each sample).

![Fig. 3. ENH relocates Id2 to the cytoplasm. (A) SK-N-SH cells expressing Flag-id2 were treated with RA or vehicle control for 48 h. Cells were double-immunostained for Flag (green) and ENH (red). Nuclei were counterstained with DAPI (blue). Arrows indicate cells showing coexpression of cytoplasmic Flag-Id2 and ENH. Arrowheads indicate cells with nuclear Flag-Id2 that lack ENH. (B) SK-N-SH cells were transiently transfected with Flag-ENH and immunostained for endogenous Id2 (red) and Flag (green). Nuclei were counterstained with DAPI. (C) Quantitative analysis of SK-N-SH cells displaying cytoplasmic Flag-Id2 from the experiment shown in A (at least 300 cells were scored for each sample).](image-url)

![Fig. 4. ENH knockdown prevents translocation of Id2 to the cytoplasm in neuroblastoma cells treated with RA. (A) Control (scrambled) or ENH-specific siRNA oligonucleotides were introduced in SK-N-SH expressing Flag-id2 before treatment with RA or vehicle control for 72 h. Cells were immunostained for Flag-id2 (red) and counterstained with DAPI. Arrowheads indicate cells displaying full relocation of Flag-id2 to the cytoplasm after treatment with RA. (B) Quantitative analysis of cells displaying predominant nuclear Flag-id2 (at least 500 cells were scored for each sample).](image-url)
Id2). Together, these results indicate that activation of ENH expression by RA is essential for cytoplasmic sequestration of Id2 in neuroblastoma.

**ENH Counters Id2 Activity and Is an Inhibitor of Proliferation and Cell-Cycle Progression.** To test the hypothesis that ENH restrains the inhibitory effects of Id2 on bHLH-mediated transcription by acting as a cytoplasmic retention factor for Id2, we performed luciferase reporter assays with five immunoregulated E-boxes driving expression of luciferase (E-box-luc). We transfected the E-box-luc plasmid in the presence of mammalian expression vectors for the ubiquitously expressed bHLH protein E47, Id2, and increasing amounts of ENH. Id2 inhibition of E47-mediated transcription was relieved by coexpression of ENH in a dose-dependent manner (Fig. 5 A). A well-known function of Id2 is the ability to enhance cell proliferation by promoting the transition from G1 to S phase of the cell cycle (6, 7). Therefore, we asked whether ENH inhibited cell proliferation and opposed Id2-mediated entry into S phase. Expression of ENH in three human neuroectodermal cell lines (the glioma cell line SF188 and the neuroblastoma cell lines IMR-32 and SK-N-SH) markedly inhibited colony formation, suggesting that ENH has antiproliferative effects (Fig. 5B). Next we transfected SK-N-SH with ENH and Id2 in the presence of a GFP expression plasmid and measured the rate of DNA synthesis by incorporation of BrdU of the successfully transfected, GFP-positive cells. Ectopic ENH strongly inhibited S phase entry and abrogated the Id2-mediated stimulation of DNA synthesis (Fig. 5C). These results suggest that, through its ability to sequester Id2 in the cytoplasm, ENH can efficiently suppress the functions of Id2 requiring nuclear localization, including the stimulation of cell-cycle progression.

**The ENH-Id2 Pathway in Development and Cancer from the Nervous System.** Taken together, the above findings indicate that, even when ectopically expressed, Id2 may be efficiently inactivated through cytoplasmic relocation implemented by differentiation signals that converge upon up-regulation of ENH. To test this hypothesis in a genetic mouse model in vivo, we generated transgenic mice expressing Flag-Id2 from the neural-specific promoter Nestin. We established six independent Nestin-Flag-Id2 mouse transgenic lines. We confirmed that Flag-Id2 is expressed in the telencephalon of hemizygous embryos by Western blot (Fig. 6A) and immunohistochemistry (Fig. 6C Upper). The older transgenic mice of this colony are >1 year old. We did not observe any abnormality in growth and differentiation of the nervous system during embryogenesis or postnatal life of Nestin-Flag-Id2 transgenics. Thus, we took advantage of this transgenic system to ask whether normal differentiation in the nervous system requires ENH-mediated relocation of Id2 to the cytoplasm. First, we used coimmunoprecipitation experiments to show that Flag-Id2 interacted specifically with endogenous ENH in Nestin-Flag-Id2 transgenic brains (Fig. 6B). Next, we performed immunohistochemistry for Flag and ENH on adjacent sections of the telencephalon at E15.5. At this developmental age, active proliferation of neural precursors is present in the periventricular, germinal layer [ventricular zone (VZ)], whereas differentiated neurons migrate radially and enter the mantle zone (MZ), which contains postmitotic cells. Flag-Id2 was predominantly nuclear in the neural precursors of the VZ but relocated to the cytoplasm in the differentiating neurons migrating toward the MZ (Fig. 6C Lower). Interestingly, ENH was barely detectable in the proliferating and undifferentiated precursors of the VZ but was coexpressed with Id2 in the cytoplasm of differentiated neurons (Fig. 6C Lower). These findings suggest that ENH is a component of the physiologic neural differentiation machinery that promotes cytoplasm relocation of Id2 in the developing brain.

Our earlier work established that Id2 displays predominant nuclear expression in aggressive neuroblastoma, an undifferenti-
that the LIM–HLH interaction is also used by ENH to inhibit the position of particular transcription complexes. Our findings suggest these associations occur in the nucleus and determine the compartmentalization of Id2, the process of differentiation, asso-

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captured form of pediatric tumor derived from the neural crest (6). A more differentiated form of these tumors, the ganglioneuroblas-toma (GBN), is characterized by the presence of a differentiated cellular component interspersed with a predominant, undiffer-

cellular compartmentalization of Id2 in primary neural tumors, we compared the expression of ENH and Id2 in four GNBs by immunostaining tumor sections with antibodies against Id2 and ENH. Most of the tumor cells stained negative for both ENH and Id2. However, we detected cytoplasmic accumulation of Id2 in the mature ganglionic cells of these tumors. Interestingly, the cytoplasm of the same cells was marked positively for ENH. Representative images are shown in Fig. 7.

The tumor expression data support our findings in cell culture and embryonic mouse brain and further strengthen the hypothesis that differentiation of neural cells requires ENH to sequester Id2 in the cytoplasm.

Discussion

Regulation of transcription factors by subcellular compartmental-

ization has been demonstrated in a number of cases (27). A common mechanism elicited in this process is sequestration of the factor into inactive compartments, usually through direct or indirect association with the cytoskeleton (28–33). The cellular localization of Id2 was recently proposed to be critical for the regulation of Id2 function (10, 14, 15). Id2 activity is primarily executed in the nucleus, where the Id2 protein antagonizes the function of DNA-binding proteins and pocket proteins of Rb family (4, 34). Although other biological conditions may regulate subcellular compartmentalization of Id2, the process of differentiation, associated with the state of proliferative quiescence, requires nuclear exclusion of Id2 (10, 14, 15). Here we have found that the cytoskeleton-associated protein ENH binds and sequesters Id2 in the cytoplasm, thus preventing its nuclear actions.

ENH belongs to a growing family of adaptor proteins that are anchored to the actin cytoskeleton through the PDZ domain and direct LIM-associated partners to actin filaments (18). The LIM domains of ENH are cysteine-rich double zinc finger motifs, which are known to mediate protein–protein interactions (35). They contact the HLH domain of Id2. Interestingly, there are previous examples of interactions between the HLH and the LIM domains. These include binding between the HLH protein TAL1 and the LIM transcription factor LMO2, as well as the interaction of MyoD, MRF4, and myogenin with MLP, another LIM protein (36, 37). These associations occur in the nucleus and determine the composition of particular transcription complexes. Our findings suggest that the LIM–HLH interaction is also used by ENH to inhibit nuclear shuttling of Id2 and drive differentiation. Knockdown of ENH had marked consequences on the cytoplasmic translocation of Id2 promoted by treatment of neuroblastoma cells with RA, a powerful inhibitor of cell proliferation and inducer of multiple pathways of differentiation. By anchoring itself to the actin cyto-

skeleton through the N-terminal PDZ domain, ENH tethers Id2 to the cytoskeleton. This mechanism recapitulates that ascribed to other cytoskeleton-associated proteins for their ability to sequester transcription factors in the cytoplasm (28–33). Although we have been focused primarily on the functional interaction between ENH and Id2 in neural cells, the ability of ENH to bind other Id proteins combined with its participation in differentiation of other tissue types (e.g., muscle) suggests that ENH may be a general inducer of differentiation through binding and cytoplasmic sequestration of Id proteins. Recently, additional isoforms of ENH (ENH2, ENH3, and ENH4) have been identified in human and mouse muscle tissues (19, 38). These isoforms lack the three LIM domains and resemble the ENHΔLIM mutant tested by us in Fig. 1 A and B. Based on our results, we conclude that the alternative ENH isoforms are unable to bind Id2, a property that might contribute to a potential dominant-negative activity toward full-length ENH (ENH1) in vivo (19, 38). Id proteins are aberrantly accumulated in various forms of human cancer, where they drive multiple hallmarks of neoplasia (34). The most common mechanism selected by tumor cells to activate Id function is to elevate the expression of Id genes through oncogenic activation of the upstream transcriptional enhancers. Now we suggest that tumor cells may also target another level of regulation of the Id biology. Our findings in GNB implicate that, by limiting the access of Id2 to the nuclear targets, expression of ENH may be a crucial safeguard against full-blown anaplasia in more differentiated tumors. As we have shown for ENH, other members of the PDZ-LIM domain family of proteins are more abundantly expressed in nontransformed cells and suppress growth of tumor cells (39–41). It is likely that this is a general attribute of this family of proteins. It will be interesting to test whether the most aggressive forms of human neoplasm select genetic and/or epigenetic alterations of the genes coding for PDZ-LIM proteins.

Materials and Methods

Yeast Two-Hybrid Screening. The Proquest system (Life Technolo-

gies) was used for yeast two-hybrid screening. The entire coding se-
quence of human Id2 was subcloned into the bait plasmid pDBLeu. A human fetal brain cdNA library in pPC86 (Life Technologies) was transformed into MaV203 yeast cells and screened for interactors with the bait plasmid according to the manufacturer’s protocol.

Cell Culture, Colony-Forming Assay, and Transfection. Neuroblastoma cell lines SK-N-SH, IMR-32, and LAN-1, the glialoma cell line SF188, and COS-1 cells were maintained in 10% FBS (Sigma) in DMEM (Cambrex). For colony-forming assay cells were transfected with pcDNA3-ENH or vector control. Cells were selected in G418 for 14 days, and colonies were scored in triplicate cultures. Cells were transfected by using Lipofectamine 2000 according to the manufacturer’s instructions.

Northern Blot. RNA was isolated by the TRIZol (Invitrogen) method. Twenty micrograms of total RNA was electrophoresed on an agarose-formaldehyde gel and transferred to nylon membrane (Nytran SPC; Schleicher & Schuell). cDNA of human ENH was used as a probe.

GST Pull-Down Assay, Western Blot, and Coimmunoprecipitation. GST fusion proteins were purified from BL21 Star (Invitrogen). GST pull-down assay was performed as described (42, 43). For Western blot analysis cellular pellets were lysed in ice-cold RIPA buffer (50 mM Tris, pH 7.5/150 mM NaCl/1% Nonidet P-40/0.5% sodium...
deoxynucleotide/0.1% SDS) containing Complete Mini protease inhibitor pellet (Roche) and 1 mM PMSF. Lysates were electrophoresed on SDS/PAGE gels and transferred to nitrocellulose membrane (Amersham Pharmacia Biotech). Membranes were stained with antibodies against ENH, Id2, (Santa Cruz Biotechnology), and α-tubulin (Sigma), and blots were developed by using ECL Western Blotting Detection System (Amersham Pharmacia Biotech). The anti-ENH antibody is a rabbit polyclonal that was produced in collaboration with Zymed against a peptide that is fully conserved in the mouse sequence (KOQNGPPRKHI). Immunoprecipitation of Id2 and ENH from cells transfected with pcDNA.3-Id2 and p3XFlag-ENH was performed as described (44).

Luciferase Assay. The luciferase reporter construct 5xE-box-luciferase (43) was cotransfected with pcDNA.3-E47 and pcDNA.3 vector or pcDNA.3-Id2 and pcDNA.3-ENH into SK-N-SH cells. pCMV-β-gal was cotransfected for normalization. Twenty-four hours later luciferase and β-galactosidase activities were measured as described (7).

BRD IIucorporation Study. SK-N-SH cells were plated in Lab-Tek Chamber Slides (Nalge Nunc). Cells were transfected with plasmids expressing the empty vector, Id2, ENH, or both and an EGFP expression vector to identify transfected cells. After 24 h, cells were labeled with 10 μM BrdU for 6 h and 14 h, fixed, and stained with anti-BrdU antibody (Roche) for1 h at room temperature. Secondary antibody was donkey anti-mouse, Cy3-conjugated (Jackson Immunoresearch). Nuclei were counterstained with DAPI. Cells were examined on an Olympus epifluorescence microscope. BRD II-positive cells were scored by counting at least 500 GFP-positive cells in three independent experiments.

Quantitative Analysis of Subcellular Localization of Id2 and siRNA Experiments. SK-N-SH-Flag-Id2 cells untreated or treated with RA for 48 h were fixed in 4% paraformaldehyde. Flag-Id2 and endogenous ENH were immunostained by using Flag-M2 (Sigma) and ENH antibodies, respectively. SK-N-SH cells were transfected with vector or p3XFlag-ENH and immunostained by using Flag-M2 and Id2 (Zymed) antibodies. For silencing of ENH, ENH siRNA (sGenome Smartpool reagent M-006950-00) and control, nontargeting Negative control Smartpool reagent D-0026-13) siRNA mixtures were purchased from Dharmacon. SK-N-SH cells stably expressing Flag-Id2 were treated with vehicle control or RA for 48 h and transfected with 60 nM siRNAs by using Lipofectamine 2000 (Invitrogen). Thirty-six hours after transfection cells were fixed with 4% paraformaldehyde and immunostained by using Flag-M2 antibodies. Parallel cultures were analyzed by Western blot. Secondary antibodies were FITC- or Cy3-conjugated anti-rabbit and Cy3-conjugated anti-mouse (Jackson Immunoresearch). Nuclei were counterstained with DAPI. Slides were mounted in 90% glycerol in PBS and analyzed on an Olympus epifluorescence microscope. The percentage of cells displaying Id2 staining in the nucleus was scored by counting at least 500 cells from triplicate samples.

Transgene Construction, Generation, and Screening of Mice. To direct transgenic expression of Id2, Flag-tagged Id2 cDNA was driven by the enhancer element contained in the second intron of the Nestin gene coupled with the thymidine kinase minimal promoter (45). The second intron of the Nestin gene directs expression in central nervous system progenitor cells. The first intron from the rat insulin II gene was included to enhance expression levels (46). The transgene fragment was microinjected at a concentration of 6 ng/μl into fertilized mouse eggs. Transgenic mice were identified by PCR analysis of DNA samples prepared from tail biopsies.

Immunohistochemistry. GNB sections were from anonymous tumor stores in the Columbia University tumor bank. Sections from E15.5 mouse brain or primary tumors were deparaffinized in xylene and rehydrated in a graded series of ethyl alcohol. Primary antibodies were Flag-M2 (Sigma), Id2, and ENH (Zymed). Avidin–biotin–peroxidase complex technique was used for primary antibody detection (Vectastain kit; Vector Laboratories). Staining was developed by using diaminobenzidine (brown precipitate). Sections were counterstained with hematoxylin. Rabbit or mouse IgG (Vector Laboratories) and tissue from Id2−/− mice were used as controls for specificity of the staining.

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