

Tumorigenesis and Neoplastic Progression

Up-Regulation of Angiopoietin-2, Matrix Metalloprotease-2, Membrane Type 1 Metalloprotease, and Laminin 5 γ 2 Correlates with the Invasiveness of Human Glioma

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Diffuse infiltration of malignant human glioma cells into surrounding brain structures occurs through the activation of multigenic programs. We recently showed that angiopoietin-2 (Ang2) induces glioma invasion through the activation of matrix metalloprotease-2 (MMP-2). Here, we report that up-regulation of Ang2, MMP-2, membrane type 1-MMP (MT1-MMP), and laminin 5 γ 2 (LN 5 γ 2) in tumor cells correlates with glioma invasion. Analyses of 57 clinical human glioma biopsies of World Health Organization grade I to IV tumors displaying a distinct invasive edge and 39 glioma specimens that only contain the central region of the tumor showed that Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 were co-overexpressed in invasive areas but not in the central regions of the glioma tissues. Statistical analyses revealed a significant link between the preferential expression of these molecules and invasiveness. Protein analyses of microdissected primary glioma tissue showed up-regulation and activation of MT1-MMP and LN 5 γ 2 at the invasive edge of the tumors, supporting this observation. Concordantly, in human U87MG glioma xenografts engineered to express Ang2, increased expression of MT1-MMP and LN 5 γ 2, along with MMP-2 up-regulation, in actively invading glioma cells was also evident. In cell culture, stimulation of glioma cells by overexpressing Ang2 or exposure to exogenous Ang2 promoted the expression and activation of MMP-2, MT1-MMP, and LN 5 γ 2. These results suggest that up-regulation of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 is

associated with the invasiveness displayed by human gliomas and that induction of these molecules by Ang2 may be essential for glioma invasion. (*Am J Pathol* 2005, 166:877–890)

A hallmark of highly malignant human gliomas is their rapid invasive growth into surrounding brain parenchyma. The infiltration by glioma cells throughout the brain renders these tumors incurable even by the combined approaches of surgery, radiotherapy, chemotherapy, and immunotherapy.^{1–3} Studies from both *in vitro* and *in vivo* tumor model systems have suggested a link between increased expression and activation of several matrix metalloproteases (MMPs) such as MMP-2 and membrane type 1 (MT1)-MMP and malignant glioma invasiveness.⁴ These MMPs produced by tumor, endothelial, and/or stromal cells degrade extracellular matrix at the invasive fronts of the glioma cells, thus removing the extracellular matrix barrier allowing subsequent tumor cell migration into newly created, more permissive spaces in the adjacent brain structures.^{5,6} Additionally, increased expression and activation of laminin (LN) 5 γ 2, a basement membrane protein, has been linked to the invasiveness of malignant human cancers such as breast, lung, pancreatic, colon, esophageal, and cervical cancers.⁷ Accumulating evidence demonstrates that interaction between MMP-2 and MT1-MMP leads to the activation of MMP-2⁵ and interactions between MMP-2,

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MT1-MMP, and LN 5 γ 2 results in the activation of LN 5 γ 2.^{8,9} However, the involvement and mechanisms of these proteins in the initiation and maintenance of human glioma cell invasion are not well understood.

Angiopoietin-2 (Ang2) is an angiogenic factor that plays critical roles in angiogenesis and tumor progression. During angiogenesis, Ang2 antagonizes Ang1 activity by competitively inhibiting the binding of Ang1 to their cognate endothelial receptor, Tie2, causing destabilization of the vasculature. Ang2 also acts in concert with vascular endothelial growth factor to regulate vessel growth.¹⁰ In human cancers, increased expression of Ang2 in tumor cells is closely correlated to the progression, invasiveness, and metastases of lung, gastric, colon, and breast cancers.^{11–15} Stable overexpression of Ang2 by human tumor cells promotes tumor growth, angiogenesis, and metastases in animals.^{12,16} We recently have shown that Ang2 induces human glioma cell invasion through the activation of MMP-2 both *in vivo* and *in vitro*. Up-regulated expression of Ang2 and MMP-2 were found in the invasive regions of primary human glioma specimens and in intracranial xenografts of U87MG glioma cells that stably overexpress Ang2.¹⁷ Here, we report that co-expression of Ang2, MT1-MMP, and LN 5 γ 2 were found in the invasive areas, but not in the central regions of primary human glioma specimens. Furthermore, Ang2 induces the expression of MT1-MMP and LN 5 γ 2 in invasive glioma xenografts formed by human U87MG Ang2-expressing glioma cells in the murine brain and in cell culture. These data strongly suggest that Ang2 induces glioma cell invasion through the stimulation of MMP-2, MT-1 MMP, and LN 5 γ 2 in Tie2-deficient glioma cells.

Materials and Methods

Cell Lines and Reagents

Human U87MG glioma cells were obtained from American Type Culture Collection (Rockville, MD) and their culture was previously described.¹⁸ Fetal bovine serum was from HyClone Inc., Salt Lake City, UT. The following reagents were used in this study: anti-Myc-tag antibody (1 μ g/ml) from Medical & Biological Laboratories, Ltd., Nagoya, Japan; anti-Ang2 (C-19, 1:50) and anti-LN 5 γ 2 antibodies (C-20, 1:100 or 1 μ g/ml) from Santa Cruz Biotechnology, Inc., Santa Cruz, CA; anti-mouse CD31 antibody (MEC 13.3, 1:1000) from BD PharMingen, San Diego, CA; anti-MMP-2 (AB809, 1:100), anti-MT1-MMP antibodies (AB8012, 1:500; AB805, 1:500; and AB815, 1 μ g/ml), anti-TIMP-2 antibody (mAb 3310, 1:200) from Chemicon Int., Temecula, CA; anti-MMP-9 antibody (1:1000) from CalBiochem., San Diego, CA; recombinant human Ang2 protein (623-AN-025), anti-Ang2 antibody (AF623, 1:200) from R&D Systems, Minneapolis, MN; and anti-von Willebrand factor (vWF) antibody (1:1000) from DAKO, Carpinteria, CA. The secondary and tertiary antibodies were from Vector Laboratories (Burlingame, CA) or Jackson ImmunoResearch Laboratories (West Grove, PA). A DAB elite kit was from DAKO, Aqua block was

from East Coast Biologics, Inc. (North Berwick, ME). Cell culture media and other reagents were from Invitrogen/BRL, Grand Island, New York; Sigma Chemicals, St. Louis, MO; or Fisher Scientific, Hanover Park, IL.

Immunohistochemical (IHC) Analyses of Primary Human Glioma Specimens

Of the 96 human glioma specimens investigated (Table 1 and data not shown), there were 8 pilocytic astrocytomas (PA) [World Health Organization (WHO) grade I], 8 diffuse astrocytomas (DA) (WHO grade II), 3 oligoastrocytomas (OA) (WHO grade II), 7 oligodendrogliomas (OD) (WHO grade II), 10 anaplastic astrocytomas (AA) (WHO grade III), 6 anaplastic oligodendrogliomas (AOD) (WHO grade III), 3 anaplastic oligoastrocytomas (AOA) (WHO grade III), and 51 glioblastoma multiforme (GBM) (WHO grade IV). Among these grade I to IV glioma specimens, 57 samples display an invasive edge and 39 samples contain only the central area of the tumors identified by hematoxylin and eosin (H&E) staining. In addition, four normal human brain specimens were also included as normal controls. All of the tumor samples were surgical specimens obtained during the last 7 years at the Department of Neurosurgery, Saitama Medical School, Saitama, Japan. The normal brain samples were from the cerebral hemisphere or cortex of the brain and obtained through autopsies from patients who did not have any brain lesions. The specimens were fixed in 10% formaldehyde and embedded in paraffin. The WHO glioma grade and the presence of invasive areas in the primary glioma biopsies were verified using H&E-stained tissue sections by Dr. Takanori Hirose, a neuropathologist at the Department of Pathology, Saitama Medical School, Saitama, Japan. The diagnosis of the WHO tumor grade and invasiveness was further confirmed by Dr. Ronald Hamilton, a neuropathologist at the Department of Pathology, University of Pittsburgh, Pittsburgh, PA. The 5- μ m sections were deparaffinized in prewarmed xylene (55°C) for 5 minutes followed by dehydration. After washing with Tris-buffered saline, the antigen was retrieved by boiling the sections in a citrate buffer (pH 6.0) twice for 5 minutes. The IHC analyses using diaminobenzidine as a chromogen were performed using anti-Ang2 (C-19), anti-MMP-2 (AB809), anti-MT1-MMP (AB8012), anti-LN 5 γ 2 (C-20), and anti-vWF antibodies as previously described.¹⁷ Quantitative analyses of blood vessel densities were performed on serial-cut human glioma specimens that were stained with the anti-vWF antibody as previously described.¹⁹

Statistical Analyses

A χ^2 test for trend was performed to examine the association between positive staining for Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 and each area of the glioma specimens (center area, border area, and invasive area) using the StatView Version 5.0 software (SAS Institute Inc., Cary, NC). A two-sided *P* value was calculated on the basis of the χ^2 test. *P* values less than 0.05 were considered significant.

Table 1. Immunohistochemical Staining for Ang2, MMP-2, MT1-MMP, and LN5 γ 2

Case	Histology (WHO grade)	Ang2			MMP-2			MT1-MMP			LN5 γ 2		
		Center	Border	Invasion	Center	Border	Invasion	Center	Border	Invasion	Center	Border	Invasion
J140#	Normal	±			—			—			—		
J141#	Normal	±			—			—			—		
J142#	Normal	1+			—			—			—		
J143#	Normal	±			—			—			—		
J39	P.A. (I)	±	2+	1+	2+	2+	2+	—	1+	1+	1+	2+	1+
J40*	P.A. (I)	2+	2+	2+	±	1+	1+	±	1+	±	—	2+	2+
J42	P.A. (I)	—	2+	1+	—	1+	1+	—	1+	1+	—	1+	1+
J43	P.A. (I)	1+	2+	1+	2+	2+	2+	—	1+	1+	1+	2+	2+
J49	P.A. (I)	—	1+	±	±	3+	3+	—	—	—	—	1+	±
J51	P.A. (I)	±	1+	2+	±	1+	2+	—	1+	1+	—	1+	±
J182	P.A. (I)	—	±	±	1+	2+	2+	1+	2+	2+	1+	2+	1+
J13	D.A. (II)	—	3+	1+	—	3+	2+	—	3+	2+	—	3+	1+
J63	D.A. (II)	1+	1+	2+	—	1+	1+	1+	2+	2+	—	1+	1+
J173	D.A. (II)	1+	2+	1+	±	2+	2+	1+	2+	1+	2+	3+	2+
J178	O.A. (II)	1+	2+	1+	±	2+	1+	1+	2+	1+	2+	3+	2+
J19	O.D. (II)	1+	2+	2+	±	2+	1+	±	1+	1+	±	±	±
J48	O.D. (II)	1+	1+	1+	±	1+	2+	±	1+	1+	—	1+	1+
J181	O.D. (II)	±	1+	1+	1+	2+	1+	1+	1+	1+	1+	2+	2+
J10	A.A. (III)	1+	3+	2+	—	1+	1+	±	2+	2+	1+	3+	2+
J11	A.A. (III)	1+	2+	2+	—	1+	1+	—	—	—	2+	2+	2+
J47	A.A. (III)	2+	2+	2+	1+	1+	1+	1+	1+	1+	—	—	±
J60	A.A. (III)	2+	2+	2+	—	±	±	±	±	1+	—	1+	1+
J62	A.A. (III)	1+	2+	2+	1+	1+	2+	±	1+	2+	1+	2+	2+
J165	A.A. (III)	1+	2+	2+	1+	2+	2+	1+	2+	1+	1+	2+	1+
J16	A.O.D. (III)	1+	2+	2+	1+	2+	1+	—	±	—	1+	2+	1+
J17	A.O.D. (III)	2+	3+	3+	1+	2+	1+	±	1+	1+	1+	2+	2+
J95	A.O.D. (III)	1+	2+	2+	±	1+	3+	±	1+	1+	2+	2+	2+
J96	A.O.D. (III)	1+	2+	2+	±	1+	2+	—	1+	±	±	3+	3+
J18	A.O.A. (III)	2+	2+	2+	—	±	±	±	1+	—	2+	2+	3+
J97	A.O.A. (III)	1+	3+	3+	—	1+	1+	±	1+	1+	1+	2+	2+
J1	GBM (IV)	—	2+	2+	—	3+	2+	—	3+	2+	—	3+	2+
J2	GBM (IV)	1+	2+	2+	2+	2+	2+	1+	2+	2+	2+	2+	3+
J3	GBM (IV)	2+	3+	3+	—	1+	2+	±	2+	2+	1+	2+	2+
J5	GBM (IV)	1+	2+	2+	1+	1+	1+	—	±	±	2+	2+	2+
J6	GBM (IV)	2+	3+	2+	2+	2+	2+	±	1+	2+	2+	1+	2+
J7	GBM (IV)	±	2+	2+	1+	2+	2+	1+	1+	2+	1+	2+	2+
J58	GBM (IV)	1+	2+	2+	±	2+	2+	±	2+	1+	—	2+	2+
J64	GBM (IV)	±	2+	±	—	1+	1+	—	1+	—	±	2+	1+
J66	GBM (IV)	1+	2+	2+	2+	2+	2+	—	1+	1+	1+	1+	2+
J68	GBM (IV)	1+	2+	2+	2+	2+	2+	±	1+	1+	1+	2+	2+
J71	GBM (IV)	2+	3+	3+	±	2+	2+	—	1+	1+	±	±	1+
J72	GBM (IV)	±	2+	2+	1+	2+	1+	—	1+	1+	1+	1+	1+
J78	GBM (IV)	2+	2+	3+	1+	2+	1+	—	1+	1+	—	1+	1+
J79	GBM (IV)	2+	3+	3+	—	—	—	—	1+	1+	1+	2+	2+
J80	GBM (IV)	1+	2+	2+	±	2+	1+	—	±	±	1+	2+	2+
J81	GBM (IV)	1+	3+	3+	1+	2+	2+	—	1+	1+	1+	2+	2+
J82	GBM (IV)	2+	3+	3+	±	2+	1+	—	1+	1+	—	1+	1+
J83	GBM (IV)	±	2+	2+	±	1+	2+	—	1+	1+	2+	2+	2+
J85	GBM (IV)	2+	3+	2+	—	1+	1+	1+	1+	2+	1+	2+	1+
J86	GBM (IV)	2+	3+	3+	1+	2+	2+	±	1+	1+	1+	2+	1+
J87	GBM (IV)	1+	1+	2+	±	2+	1+	1+	2+	2+	2+	2+	±
J89	GBM (IV)	1+	2+	1+	1+	2+	2+	1+	2+	2+	±	1+	1+
J94	GBM (IV)	1+	2+	2+	1+	2+	2+	—	1+	1+	±	2+	2+
J98	GBM (IV)	1+	2+	2+	1+	1+	2+	1+	1+	2+	1+	1+	1+
J151	GBM (IV)	2+	2+	2+	—	2+	2+	—	1+	1+	1+	2+	2+
J152	GBM (IV)	1+	2+	2+	1+	2+	1+	1+	2+	2+	1+	2+	2+
J153	GBM (IV)	1+	3+	2+	±	2+	1+	—	1+	1+	2+	3+	3+
J154	GBM (IV)	1+	2+	2+	1+	3+	2+	1+	2+	2+	1+	2+	2+
J155	GBM (IV)	1+	2+	2+	1+	2+	1+	±	2+	2+	±	2+	1+
J156	GBM (IV)	2+	3+	3+	2+	2+	3+	1+	2+	2+	1+	2+	2+
J164	GBM (IV)	1+	2+	2+	1+	3+	2+	±	1+	1+	±	2+	1+

*Contains neurofibroma 1.

#Scores of normal brain were indicated in the columns of "Center" for convenience.

P.A., pilocytic astrocytoma; D.A., diffuse astrocytoma; O.A., oligoastrocytoma; O.D., oligodendroglioma; A.A., anaplastic astrocytoma; A.O.D., anaplastic oligodendroglioma; A.O.A., anaplastic oligoastrocytoma; GBM, glioblastoma multiforme.

Microdissection of Paraffin-Embedded Tissue Sections of Human Glioma Specimens

Microdissection of paraffin-embedded primary human glioma tissues was performed as previously described.²⁰ Briefly, paraffin-embedded glioma specimens were sectioned separately at 5 μm and 50 μm thickness and mounted onto plain glass slides. To identify the center, border, and invasion regions within the glioma specimen, 5- μm -thick sections were stained by H&E. Three 50- μm -thick sister sections of each sample were then deparaffinized in xylenes, rehydrated in graded ethanol, immersed in distilled water, and air-dried. To exclusively collect center-tumor or border-tumor tissues, the targeted areas were cut microscopically under an Olympus SZ-ST5 stereomicroscope (Melville, NY) with a fine needle referring to the microscopic observation of the morphology of serial H&E-stained sections.

Extraction and Western Blot Analyses of Proteins from Microdissected Human Primary Glioma Tissues

Protein extraction from microdissected formalin-fixed, paraffin-embedded tissues was performed following protocols that were previously described.²¹ Briefly, the collected tumor tissues were suspended in lysis buffer containing 25 mmol/L sodium phosphate buffer (pH 7.6), 150 mmol/L sodium chloride, 1% Triton X-100, 2% sodium dodecyl sulfate, 12 mmol/L sodium deoxycholate, 0.2% sodium azide, 0.95 mmol/L fluoride, 2 mmol/L phenylmethyl sulfonyl fluoride, 50 mg/ml aprotinin, and 50 mmol/L leupeptin and incubated at 100°C for 20 minutes followed by incubation at 60°C for 2 hours. After incubation, the tissue lysates were centrifuged at 15,000 $\times g$ for 20 minutes at 4°C, and the supernatants containing extracted proteins were collected. The protein extracts were then subjected to Western blot analyses using a sensitive protocol.²² Briefly, 10 μg of total proteins were separated by NuPAGE 4 to 12% Bis-Tris polyacrylamide gels (Invitrogen) and transferred to Immobilon-P transfer membranes (pore size 0.45 μm ; Millipore, Bedford, MA) according to the manufacturer's standard procedure. The membrane was blocked and separately probed with anti-MT1-MMP (AB8012, Chemicon), anti-LN 5 γ 2 (C-20, Santa Cruz Biotechnology), and anti- β -actin (Sigma) antibodies. Antibody detection was accomplished with SuperSignal West Femto maximum sensitivity substrate (Pierce).

Generation of U87MG Cell Lines That Stably Express Ang2 or LacZ

Transfected U87MG cell clones that stably express Ang2 were generated by transfecting U87MG cells with a cDNA for Ang2 in the pSecTagB/Myc-His(+) expression vector (Invitrogen, San Diego, CA). The clones that expressed exogenous biologically active Ang2 or LacZ were expanded and characterized as previously described.^{17,18}

Tumorigenicity, Glioma Invasion, Mouse Brain Tissue Processing, and IHC

U87MG LacZ- or U87MG Ang2-expressing cell clones (5×10^5) were stereotactically implanted into individual nude mouse brains with five mice per group. When mice developed neurological symptoms because of disturbance of their central nervous system, mice were sacrificed and their brains were removed, processed, and analyzed by IHC as previously described.^{17,18}

Quantification Analyses of IHC Data

To quantify microvascular density or the degree of staining by antibodies (for mouse xenograft tissues), five to seven serial-cut sections were stained with the anti-CD31, anti-vWF, anti-Ang2, anti-MMP-2, anti-MT1-MMP, or anti-LN 5 γ 2 antibodies. The stained images (10 or more random areas per section) were captured using an Olympus BX51 microscope equipped with a digital camera, imported into the Image Pro Plus program (Version 4.1; Media Cybernetics, Silver Spring, MD) and analyzed. The mean values of microvessel density or relative intensity of antibody staining from serial brain sections (five or more individual mice per group) in each group were used. Microvessel density was expressed as the ratio of positively stained areas to the total area of the image (object areas/ mm^2).¹⁹ The degree of antibody staining in U87MG/Ang2 gliomas was shown as fold of increase to that in U87MG/LacZ tumors.¹⁷

To quantify the intensities of IHC staining on primary human glioma specimens using anti-Ang2, anti-MMP-2, anti-MT1-MMP, and anti-LN 5 γ 2 antibodies, two experienced researchers (P.G. and Y.I.) independently examined all of the aforementioned sections stained with the four antibodies under $\times 200$ and $\times 400$ magnification. In each stained section, 5 to 10 identified areas (center area, border area, and invasive area) in each type were examined. The intensities of IHC staining with each antibody were defined as no stain (–), weakest (\pm), low (1+), medium (2+), and strong (3+) stains. The immunoreactivities of each antibody on the paraffin-embedded sections were different. The anti-Ang2 and anti-MMP-2 antibodies produced excellent staining without nonspecific background staining. The anti-MT1-MMP had the weakest activity whereas the anti-LN 5 γ 2 antibody had nonspecific staining in some tissue areas. These variations were considered during the scoring. Greater than 95% of the scoring by the two examiners determined in a double-blinded manner was comparable. Discrepancies between the scores were resolved by re-examining the stained sections side-by-side followed by consultation between the examiners.

Western Blot and Zymography Analyses

Western blot and zymography analyses were performed as previously reported.¹⁷ Briefly, six-well plates were coated with or without 10 $\mu\text{g}/\text{ml}$ of recombinant Ang2 or bovine serum albumin in phosphate-buffered saline

(PBS) at 4°C overnight, washed three times with PBS, and then dried. U87MG cells or U87MG Ang2-expressing cells were seeded onto the plates and incubated in Dulbecco's modified Eagle's medium with 10% heat-inactivated cosmic calf serum for 4 hours at 37°C. The medium was then replaced by serum-free Dulbecco's modified Eagle's medium, and the cells were maintained in the medium for 24 hours at 37°C. Whole cell lysates containing 30 μ g of total protein from the various cells were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions and transferred onto a nitrocellulose membrane. The membrane was blocked and probed with various antibodies. The reacted proteins were visualized by enhanced chemiluminescence reaction. Quantification of the expression of MT1-MMP, TIMP-2, and LN 5 γ 2 was performed by importing and analyzing the scanned images using the Image Pro Plus program.

The zymography analysis for proteolytic activities of MMP-2 toward gelatin was performed as previously described.¹⁷ Briefly, serum-free conditioned medium from a 48-hour culture of various U87MG cells plated onto six-well plates coated with Ang2, bovine serum albumin, or noncoating was collected. Conditioned medium containing 20 μ g of total proteins was separated at 4°C in a 7.5% sodium dodecyl sulfate polyacrylamide gel containing 0.2% gelatin. The sodium dodecyl sulfate in the gel was removed by washing the gel three times with a Tris-HCl buffer containing 2% Triton X-100. The gel was rinsed and developed in a gel-developing buffer containing 1% Triton X-100 and 5 mmol/L of CaCl₂ at 37°C for 16 hours. The gel was then stained by a Coomassie Blue solution, destained, and photographed. The proteolytic activities of MMP-2 toward gelatin appear as unstained areas.

Results

Preferential Expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in Invading Glioma Cells and Neovessels of Human Primary Glioma Specimens

We recently reported that Ang2 and MMP-2 are co-overexpressed in the invasive areas including tumor borders, but not in the central regions of primary human glioma biopsies.¹⁷ To further demonstrate whether there is a significant association between up-regulated expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 and tumor invasiveness displayed by various human gliomas in all four WHO grade stages, we conducted a comprehensive survey by IHC analyses. The IHC analyses were performed on a total of 96 human glioma specimens and 4 normal human brain biopsies in our collection. We tested several well-characterized antibodies against each of these molecules that have been reported in similar clinical investigations on human tumor specimens and chose goat anti-Ang2 (C-19),^{11,12,15} rabbit anti-MMP-2 (AB809),^{9,23} rabbit anti-MT1-MMP (AB8012),^{9,23} and goat anti-LN 5 γ 2 (C-20) antibodies as described in the

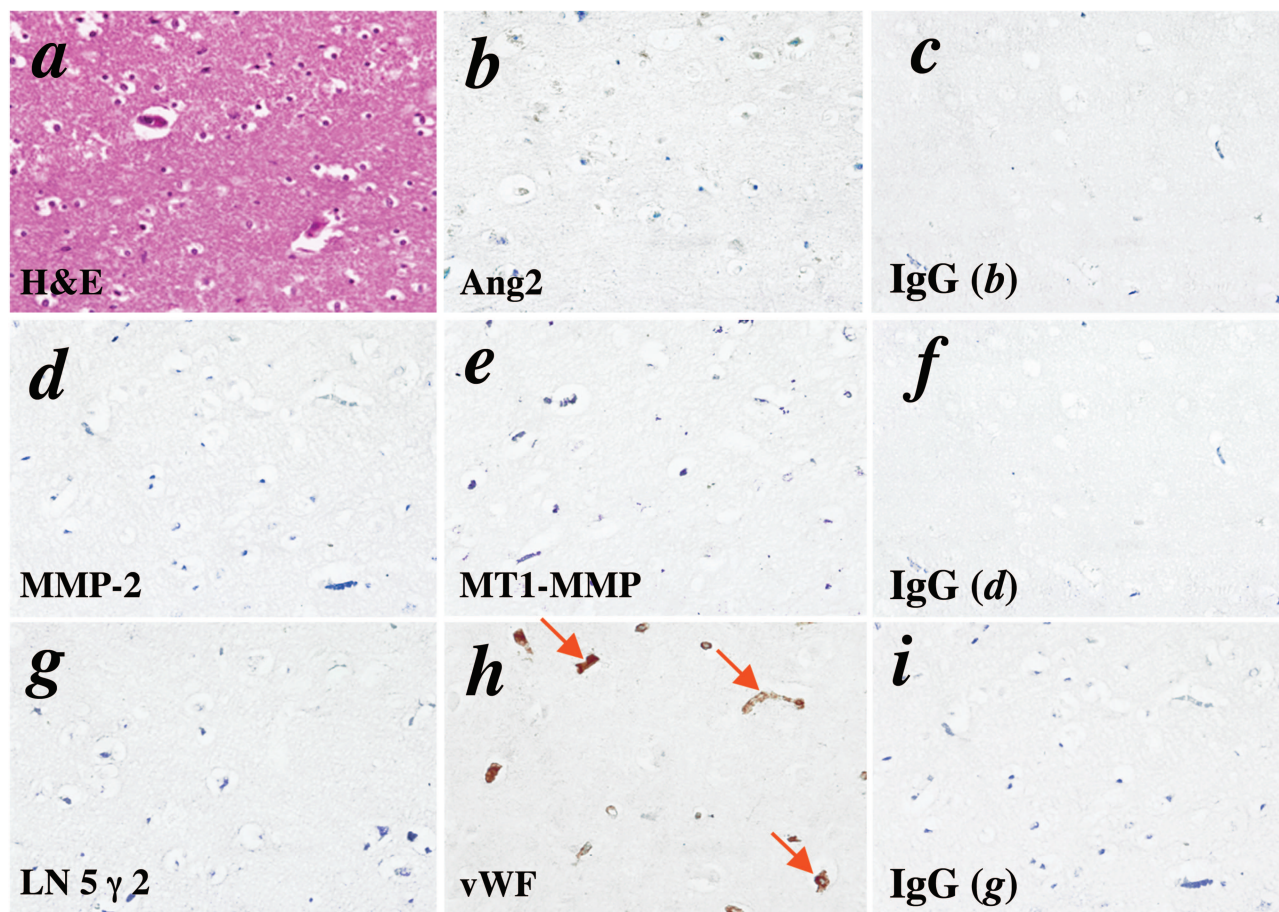
Materials and Methods section. For the negative controls, we applied isotype-matched IgG at the same concentration onto the sister sections of each specimen during the IHC staining, which all showed negative stains (see the insets in Figure 2). The specificity of each of these antibodies on detecting the corresponding protein products were verified by Western blot analyses using these antibodies on U87MG cells that express endogenous MMP-2, MT1-MMP, and LN 5 γ 2 proteins (Figure 4, and data not shown) and U87MG Ang2-expressing cells (for Ang2).¹⁷ Table 1 summarizes our findings of the relative immunoreactivities of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in 57 paraffin-embedded biopsies of human gliomas that display invasiveness or normal human brain specimens. The relative intensities of the immunostaining by each antibody were quantified as described in the Materials and Methods section.

Normal Human Brain Tissues

We performed IHC analyses on four normal brain samples that were from autopsied patients who did not have brain lesions (specimens J140 to J143, Table 1). All four specimens were mostly from the cerebral hemisphere or cortex of the brain where gliomas occur most frequently.³ These specimens were homogenous normal brain tissue with similar morphology judged by H&E staining (Figure 1a). As shown in Figure 1 (specimen J141) and Table 1, no or very weak immunoreactivities of the antibodies to Ang2 (Figure 1b), MMP-2 (Figure 1d), MT1-MMP (Figure 1e), or LN 5 γ 2 (Figure 1g) were detected. These data indicate that under normal physiological conditions, the expression of these molecules is below or near the detectable levels by IHC analyses.

Pilocytic Astrocytomas (WHO Grade I)

Eight PA biopsies were included in this study (specimens J39, J40, J42, J43, J49, J51, J182, and J183). By H&E staining, we found distinct invasive areas in seven of eight PA biopsies (87.5%; Table 1, data not shown). In addition, intratumoral hemorrhage is common in most of these grade I tumors (Figure 2A, b, data not shown). Figure 2A shows the IHC staining of specimen J42 using H&E and various antibodies. In the center regions of this PA, low expression of Ang2 (Figure 2A, d), MMP-2 (Figure 2A, g), MT1-MMP (Figure 2A, j), and LN 5 γ 2 (Figure 2A, m) was detected (Table 1). In the tumor border (Figure 2A, b), increased expression levels of Ang2 (Figure 2A, e), MMP-2 (Figure 2A, h), and LN 5 γ 2 (Figure 2A, n) are found while a low level of MT1-MMP expression was seen (Figure 2A, k). Interestingly, high levels of Ang2 and MMP-2, but not MT1-MMP and LN 5 γ 2, were seen in neovessels (arrowheads in Figure 2A, e and h), suggesting the role of Ang2 and MMP-2 in vascular growth. In invasive areas (0.25 mm away from the border, Figure 2A, c) where glioma cells are actively invading into the brain structure, only tumor cells express Ang2 (Figure 2A, f, arrows), MMP-2 (Figure 2A, i, arrows), MT1-MMP (Figure 2A, l, arrows), and LN 5 γ 2 (Figure 2A, o, arrows).



Normal human brain (specimen J141)

Figure 1. Expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in normal human brain tissue. IHC on serial sections of a normal human brain biopsy (specimen J141) using H&E (a), goat polyclonal anti-Ang2 (b), rabbit polyclonal anti-MMP-2 (d), rabbit polyclonal anti-MT1-MMP (e), goat anti-LN 5 γ 2 (g), and rabbit polyclonal anti-vWF (h) antibodies. The isotype-matched IgG controls (c, f, and i) are the identical areas shown in b, d, and g. Arrows in h indicate blood vessels. A total of four individual normal human brain tissues were analyzed. The experiments were repeated two additional times with similar results. Original magnifications, $\times 400$.

Thus, co-expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 are detected in the invading glioma cells and growing neovessels in tumor borders, but not in the central regions of PA tumors. These results suggest that at the early stage of glioma progression, up-regulation of these molecules correlates with the initial invasion displayed by PA.

Diffuse Astrocytomas (DA, Grade II), Oligoastrogliomas (OA, Grade II), and Oligodendrogliomas (OD, Grade II)

There were 8 DA, 3 OA, and 7 OD biopsies that are WHO grade II tumors in our collection. In the DA specimens, diffuse morphology of tumor cells is evident. In some tumors, diffusive tumor cells formed a cellular gradient toward the normal brain structure without identifiable borders (data not shown). Among the 18 grade II tumors, 7 tumor specimens (38.8%) contain identifiable tumor borders and clear invasive regions (at least 0.25 mm away from the tumor border, Table 1). Figure 2B shows the IHC

staining of specimen J13 (a DA) using H&E and various antibodies. Consistent with the expression profiles in the WHO grade I gliomas (Figure 2A), strong expression of Ang2 (Figure 2B, e), MMP-2 (Figure 2B, h), MT1-MMP (Figure 2B, k), and LN 5 γ 2 (Figure 2B, n) were evident in tumor cells (arrows) and in neovessels (arrowheads) along the tumor border, but not in the center region of the tumors (Figure 2B; d, g, j, and m). Increased expression of Ang2 (Figure 2B, f), MMP-2 (Figure 2B, i), and LN 5 γ 2 (Figure 2B, o), and strong expression of MT1-MMP (Figure 2B, l) were detected in cells (arrows in Figure 2B; f, i, l, and o) and in vessels (arrowheads) in the invasive areas that are 0.25 mm away from the tumor border.

Anaplastic Astrocytomas (AA, Grade III), Anaplastic Oligodendrogliomas (AOD, Grade III), and Anaplastic Oligoastrocytomas (AOA, Grade III)

There were 10 AA, 6 AOD, and 3 AOA gliomas in our collection. All of the grade III gliomas displayed in-

creased cellularity and malignancy, especially in the central region (data not shown). Among the 19 grade III glioma biopsies, 12 tumors contain tumor borders and invasive areas (63.2%, Table 1). In contrast to the grade I and II glioma specimens, the tumor borders were distinct and with sharp edges (data not shown). Similar to that in the grade II gliomas (Figure 2B), up-regulated Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 were found in the tumor borders, but not in the central regions of these tumors (data not shown). In addition, increased expression of all four molecules were detected in cells in the invasive region 0.25 mm away from the tumor borders in the grade III gliomas (data not shown).

GBM (WHO Grade IV)

There were 51 GBM specimens in our collection. These grade IV glioma biopsies display the highest cellularity and contain evident necrotic areas and intratumoral hemorrhage (data not shown). Among the 51 GBM, 31 specimens contain tumor borders and invasive areas (60.8%, Table 1). The tumor borders are also sharp and easily distinguishable (Figure 2C, a). Figure 2C shows the IHC staining of specimen J1 (a GBM) using H&E and various antibodies. We found that protein expression profiles of Ang2 (Figure 2C, e), MMP-2 (Figure 2C, h), MT1-MMP (Figure 2C, k), and LN 5 γ 2 (Figure 2C, n) were located in the tumor borders similar to IHC staining of the grade II and III gliomas (Figure 2B and data not shown). In addition, increased levels of expression of Ang2 (Figure 2C, f), MMP-2 (Figure 2C, i), MT1-MMP (Figure 2C, l), and LN 5 γ 2 (Figure 2C, o) were also found in the invasive areas of the GBM tumors. The IHC results of WHO grade III and IV gliomas demonstrate that increased expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 are further elevated in the tumor borders and invasive regions of high-grade human glioma specimens.

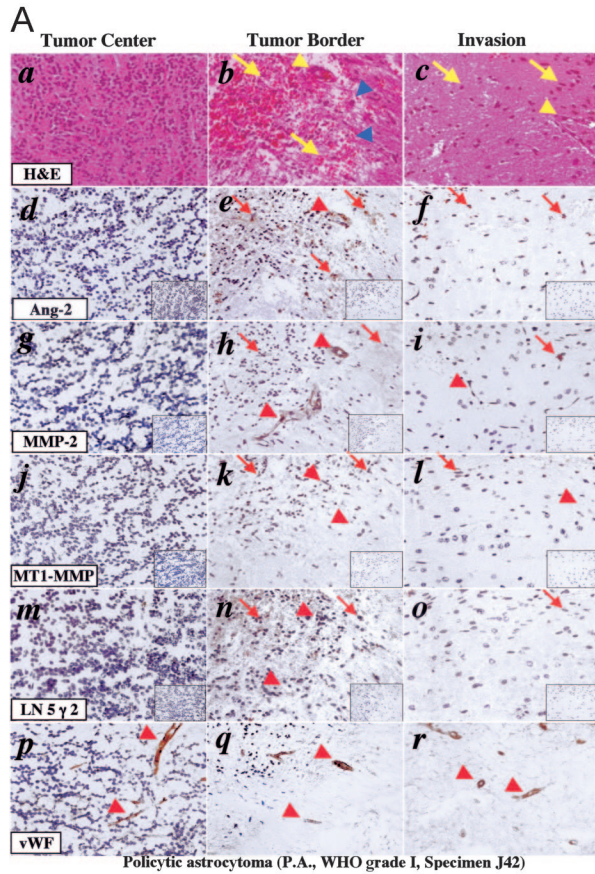
Glioma Angiogenesis

As shown in Figures 1 and 2, we stained all of the normal brain and glioma tissue specimens using an anti-vWF antibody that stains tumor endothelium. We evaluated glioma angiogenesis by quantifying the microvessel density in all of the stained tissue sections. In normal human brain tissues, vessels were apparent (Figure 1, h). In WHO grade I (Figure 2A, p and q) and II (Figure 2B, p and q) tumors, when compared with the MCD in normal brain tissues, a 1.8-fold increase of MCD was found. In WHO grade III gliomas, a 3.2-fold increase of MCD was seen. In WHO grade IV gliomas (Figure 2C, p and q), a further increase of MCD (7.1-fold) was found (data not shown). These data corroborate with previous studies showing that glioma angiogenesis is not evident in low-grade (I and II) gliomas, but significantly increased in high-grade (III and IV) tumors.²⁴

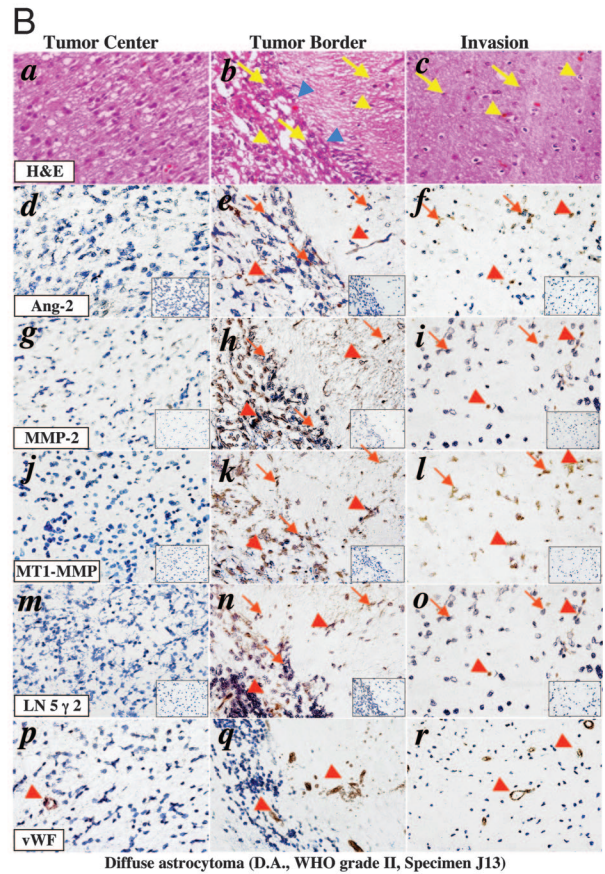
Up-Regulated Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 Are Preferentially Associated with Tumor Invasiveness among the Clinical Glioma Specimens

As shown in Table 1, we have found that 57 glioma specimens contain identifiable tumor borders and nearby invasive regions. There are 39 additional glioma specimens that only contain the central region of the tumors. We hypothesize that there is a significant association between up-regulated expressions of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 and tumor invasiveness displayed by all four WHO grades of human gliomas. To test our hypothesis, we examined the IHC analyses data on the cases that contain all three areas within the same sample: tumor center, border, and invasive areas, and determined the number of the cases analyzed that meet these criteria. As shown in Table 1, in seven grade I gliomas, there are five to six cases of seven samples (71 to 85.7%) that have expression scores in the areas both in the border and invasive areas higher than that of the center area. Similarly, the scores of protein expression level in the majority of cases in grade II to IV specimens meet the same criteria: grade II, 5 to 6 cases of 7 (71 to 85.7%); grade III, 7 to 9 cases of 12 (58 to 75%), and grade IV, 23 to 28 cases of 31 (74.1 to 90%). Together, in a total of 50 glioma grade II to IV specimens that contain all three areas (center, border, and invasion), 35 to 43 cases (70 to 86%) are available for testing our hypothesis (Table 2).

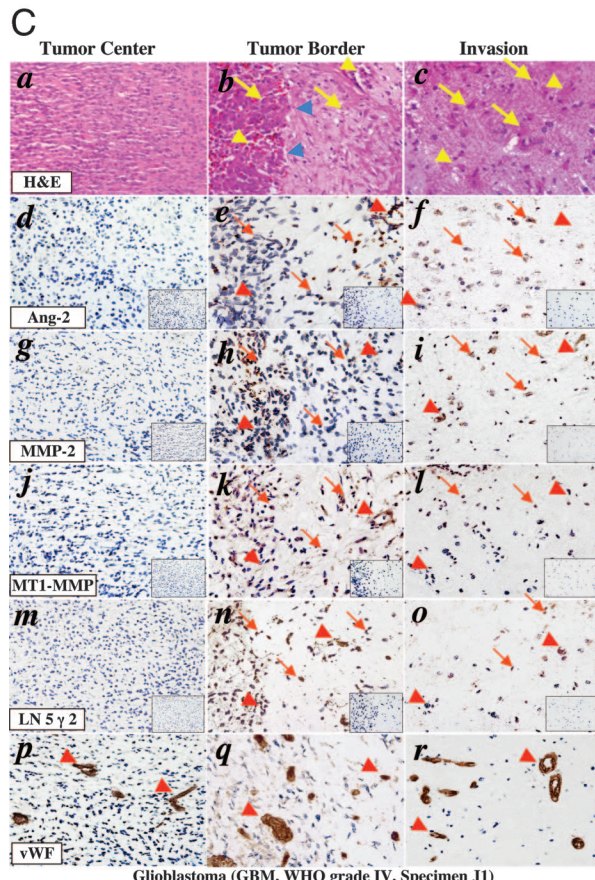
Next, we determined whether there is a distinct link between the up-regulation of the four molecules that are shown to be vital in the invasiveness of human cancers. A χ^2 test for trend was performed to examine the association between positive staining for Ang2, MMP-2, MT1-MMP, LN 5 γ 2, and each area of grade II to IV glioma specimens (center area, border area, and invasion area). A two-sided P value was calculated on the basis of the χ^2 test. As shown in Table 2, significant links are found between the positive immunoactivities of Ang2 ($P < 0.0001$ for both comparisons), MMP-2 ($P < 0.0001$ for border versus center and $P = 0.0001$ for invasion versus center, respectively), MT1-MMP ($P < 0.0001$ for both comparisons), and LN 5 γ 2 ($P < 0.0001$ for border versus center and $P = 0.0002$ for invasion versus center, respectively), and the invasiveness displayed by these gliomas. In contrast, there are no differences in the expression profiles of these four proteins between the tumor borders and invasive regions (between $P = 0.5120$ to $P = 0.7880$). Because the pathogenesis of PAs is different from that of grade II to IV gliomas,³ we separately performed the χ^2 test for trend on the IHC data for PA gliomas. No significant association of the expression levels of these four molecules and tumor invasiveness could be found, possibly because of the small sample size analyzed (data not shown). In addition, because the majority of the glioma specimens analyzed in Table 2 are grade IV GBM (31 of 50, 62%), we separately performed the χ^2 test for trend analyses on the IHC data of these 31 grade IV GBM samples. A statistically significant link



Polycystic astrocytoma (P.A., WHO grade I, Specimen J42)



Diffuse astrocytoma (D.A., WHO grade II, Specimen J13)



Glioblastoma (GBM, WHO grade IV, Specimen J1)

Figure 2. Co-overexpression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in the invasive areas of primary human glioma tissues. IHC on serial sections of PA (**A**: specimen J42, WHO grade I), DA (**B**: specimen J13, grade II), and GBM (**C**: specimen J1, grade IV) tissues using H&E (**a** to **c**), goat polyclonal anti-Ang2 (**d** to **f**), rabbit polyclonal anti-MMP-2 (**g** to **i**), rabbit polyclonal anti-MT1-MMP (**j** to **l**), goat anti-LN 5 γ 2 (**m** to **o**), and rabbit polyclonal anti-vWF (**p** to **r**) antibodies. **Insets** in **d** to **o** are the isotype-matched IgG controls that are the identical areas shown in each panel. **a, d, g, j, m, and p**: Central regions of the tumor mass; **b, e, h, k, n, and q**: tumor borders; and **c, f, i, l, o, and r**: areas that are 0.25 mm away from the tumor edge shown in **b, e, h, k, n, and q**. **Blue arrowheads** in **b** indicate the border between the tumor mass (left, high cellular density) and normal brain tissue (low cellular density). **Red or yellow arrowheads** in **b** to **r** indicate tumor vessels. **Arrows** in **b** to **o** are glioma cells or activated neural cells that were stained by anti-Ang2 (**e, f**), anti-MMP-2 (**h, i**), anti-MT1-MMP (**k, l**), and anti-LN 5 γ 2 (**n, o**) antibodies, respectively. A total of 96 individual primary tumor specimens (WHO grades I to IV) were analyzed. The experiments were repeated two additional times for each section with similar results. Original magnifications, $\times 400$.

Table 2. Scores and Correlation between Positivity for Staining and Each Area of Tumor (Grade II–IV, $n = 50$)

	Ang2			MMP-2			MT1-MMP			LN5 γ 2		
	Center	Border	Invasion	Center	Border	Invasion	Center	Border	Invasion	Center	Border	Invasion
3+	0	14	11	0	2	3	0	0	0	0	5	4
2+	14	32	33	5	29	22	0	16	16	11	31	27
1+	31	4	5	19	16	22	17	29	27	22	11	16
\pm	5	0	1	15	2	2	16	4	3	9	2	3
–	0	0	0	11	1	1	17	1	4	8	1	0
<i>P</i> value of correlation between positivity and area of tumor*												
Border vs center	<0.0001			<0.0001			<0.0001			<0.0001		
Invasion vs center	<0.0001			0.0001			<0.0001			0.0002		
Border vs invasion	N.S.			N.S.			N.S.			N.S.		

*Analyzed by χ^2 test for trend based on the distribution of the scores of each area. N.S., not significant.

between the expression profiles of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 and invasiveness of these GBM was also found ($P < 0.0001$ to $P = 0.0159$ for border versus center or invasion versus center). Similarly, there is no difference in the immunoactivities of these four molecules between the border and invasive areas ($P = 0.3939$ to $P = 0.9996$, data not shown).

Lastly, we assessed the protein expression levels in the 39 cases of grade II to IV gliomas that only have the center area, but no tumor border and invasive area. Among these 39 samples, 25 to 33 cases (64.1 to 84.6%) have low levels of protein expression of the four molecules that were examined (scores $\leq 1+$, data not shown). Because we do not have portions of these glioma biopsies containing tumor border and invasion areas, we cannot evaluate the correlation of the expression levels of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 with the invasive edge in these 39 human glioma specimens.

MT1-MMP and LN 5 γ 2 Are Overexpressed and Activated in the Invasive Region of Primary Human Glioma Specimens

To support our observations from the IHC analyses, we identified five primary human glioma specimens from our collection that have shown increased co-expression of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in the invasive regions (Table 1) with sufficient tumor material: J181, WHO grade II OD; J165, a grade III AA; J152, a grade IV GBM; J152, a grade IV GBM; and J155, a grade IV GBM. We microdissected the tissues of the central region and the invasive edge of these primary glioma specimens (Figure 3A).²⁰ Then, we performed Western blot analyses on the total proteins extracted from the microdissected glioma tissue samples using anti-MT1-MMP and anti-LN 5 γ 2 antibodies according to the protocols that were previously reported.^{21,22} As shown in Figure 3B, a twofold to threefold increase in protein expression of intact MT1-MMP (70 kd) was found in the samples from the tumor invasive edge of all of the five primary glioma specimens when compared with the central regions in the same sample. Furthermore, significant proteolytic activation of LN 5 γ 2 (155 kd) was evident by the appearance of γ 2'

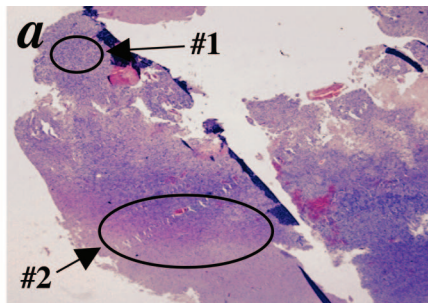
(105 kd) and γ 2x (90 kd) fragments⁸ in the samples of the tumor invasive edge of all five specimens. Autocleaved MT1-MMP protein fragments (45 kd) were also detected, albeit to a lesser degree, indicating that activation of LN 5 γ 2 and MT1-MMP occurred in these regions of the gliomas. These findings corroborate with the IHC data shown in Figure 2 and Table 1 that MT1-MMP and LN 5 γ 2 are co-up-regulated and co-activated with Ang2 and MMP-2 in the invasive edge of primary human glioma grade II to IV specimens. Taken together, our results (Tables 1 and 2 and Figures 1 to 3) demonstrate that up-regulation of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 are preferentially associated with the invasive edge of WHO grade I to IV gliomas.

MT1-MMP and LN 5 γ 2 Are Overexpressed at the Invasive Fronts and Disseminated U87MG Ang2-Expressing Gliomas in the Brain

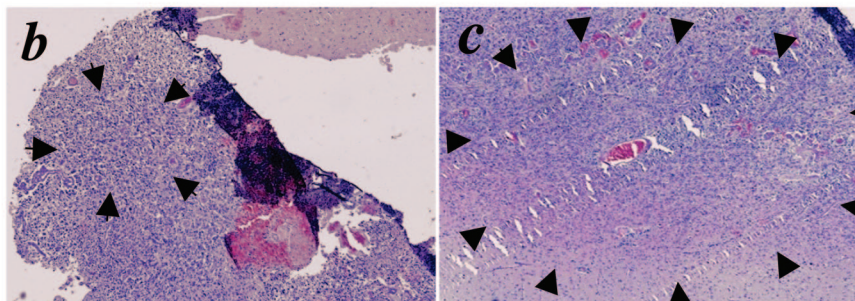
We recently reported that overexpression of Ang2 by human U87MG glioma cells caused aggressive glioma invasion compared to isogenic control tumors. We found that Ang2 and MMP-2 were co-overexpressed in the actively invading regions of U87MG Ang2-expressing tumors.¹⁷ To determine whether the expressions of MT1-MMP and LN 5 γ 2 were also up-regulated in the invasive regions of the Ang2-expressing U87MG gliomas, we performed IHC on various U87MG gliomas. As shown in Figure 4, the expression of MT1-MMP (Figure 4, f) and LN 5 γ 2 (Figure 4, h) was co-overexpressed with Ang2 (Figure 4, d) in the Ang2-expressing gliomas, whereas neither of these two proteins was detected in the isogenic control tumors (LacZ-expressing tumors; Figure 4, e and g). Because both MT1-MMP (Figure 4, f) and LN 5 γ 2 (Figure 4, h) are extracellular matrix proteins, the protein stains detected in the invasive Ang2 tumors were relatively diffusive. The highest expressions of MT1-MMP and LN 5 γ 2 (a fourfold to fivefold increase compared to that in LacZ tumors) were found at the invasive fronts as well as in the disseminated tumor clusters of the U87MG/Ang2 gliomas (arrows in Figure 4, f and h). Importantly, high expression of MT1-MMP (Figure 4, f) and LN 5 γ 2

A

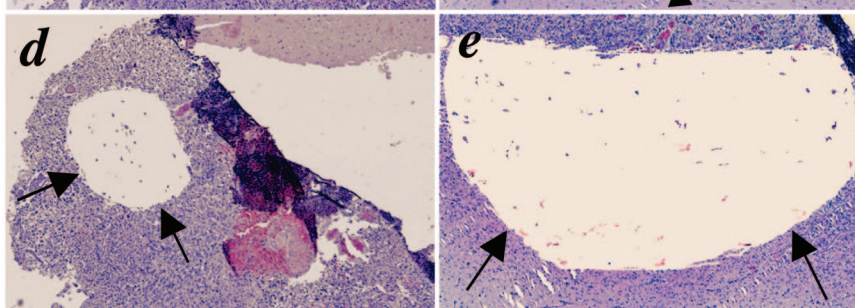
Whole Specimen (GBM)



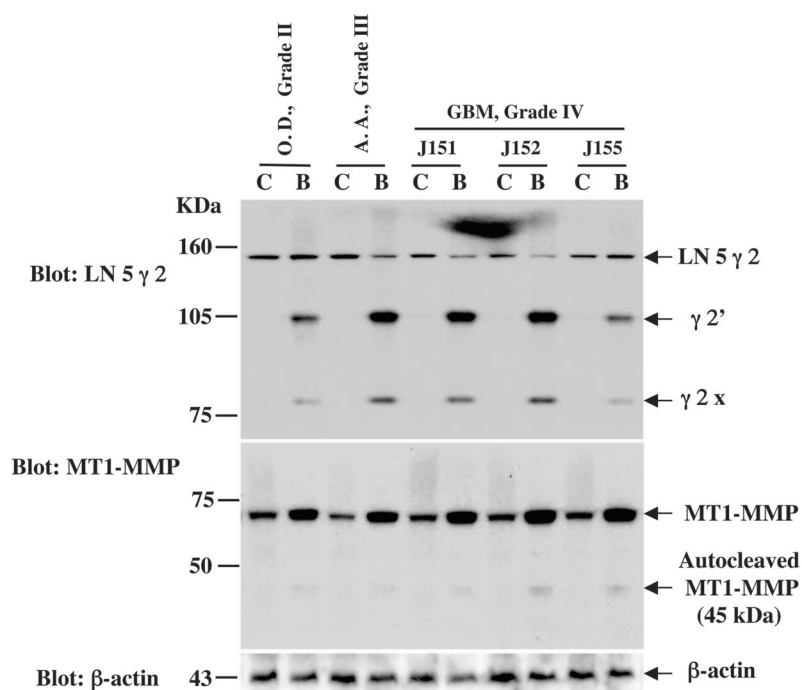
Before Microdissection



After Microdissection



B



(Figure 4, h) are not only co-localized in the invading malignant cells or disseminated tumor clusters, but also co-expressed in the same cell population where Ang2 (Figure 4, d) and MMP-2 are overexpressed.¹⁷ Co-overexpression of these four molecules in the regions where Ang2-induced gliomas invasion occurs suggests that Ang2 induces human glioma invasion through the activation of the downstream effectors MMP-2, MT1-MMP, and LN 5 γ 2.

Ang2 Induces the Stimulation of MT1-MMP and LN 5 γ 2 in Vitro

Increased expression and activation of both MT1-MMP and LN 5 γ 2 are strongly associated with tumor invasion in various types of human cancers.^{7,25} To determine whether Ang2 can directly stimulate the expression and activation of MT1-MMP and LN 5 γ 2 in glioma cells, we assessed the expression and proteolysis of these two molecules in various conditions. As shown in Figure 5, overexpression of Ang2 by U87MG cells or exposure to recombinant Ang2 by U87MG, U251MG, and T98G cells (data not shown) results in a 2.2- to 3.4-fold increase in expression and activation of LN 5 γ 2 (Figure 5, top) and MT1-MMP (Figure 5, the second panel) in comparison to untreated or bovine serum albumin-treated U87MG cells. Importantly, the stimulation of both MT1-MMP and LN 5 γ 2 under these conditions correlates with increased expression and activation of MMP-2 in the glioma cells (Figure 5, bottom). Because MT1-MMP is responsible for the activation of MMP-2 by interacting with tissue inhibitor-2 of MMP (TIMP-2),⁵ we also determined the expression of TIMP-2 in the glioma cells under these conditions and in the invasive U87MG/Ang2 and noninvasive U87MG/LacZ gliomas. We found that there was no alteration in the expression levels of TIMP-2 in Ang2-stimulated or nonstimulated glioma cells (Figure 5, the third panel), nor in the invasive or noninvasive U87MG gliomas established in the mouse brain (data not shown). Therefore, similar to that of MMP-2,¹⁷ the stimulation of MT1-MMP and LN 5 γ 2 is involved in human glioma invasion induced by Ang2 *in vitro* and *in vivo*.

Discussion

The involvement of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in the invasiveness of various human cancers including gliomas has been intensively investigated using clinical tumor specimens, xenografts of human tumor cells, and cell culture models.^{5,7,12,15,25,26} However, most of the

studies focused on the roles of only one or two of these molecules in human tumor invasion and metastases. In this report, we use three distinct model systems to determine whether there is a significant link between the expression profiles and four important molecules, Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 and human glioma invasion. First, we performed a comprehensive survey of the expression profiles of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 by analyzing a total of 96 primary human glioma specimens that included 57 samples containing a distinct invasive edge and 39 glioma biopsies containing only central regions of the tumor, and 4 normal brain biopsies. Statistical analysis of our results establishes that there is a significant association between the up-regulated expression of these molecules and the invasiveness of the various gliomas in all four WHO grades. Additionally, low levels of protein expression of these four molecules were detected in the central areas of the majority of these clinical glioma samples. Second, Western blot analyses of microdissected primary glioma specimens showed that MT1-MMP and LN 5 γ 2 proteins were not only co-up-regulated but also co-activated in the cells consisting of the invasive edge of the tumor when compared with the center regions of the same sample, thus supporting our findings of IHC analyses. Third, we further analyzed our recently established glioma invasion animal model¹⁷ and showed that the aggressively invasive U87MG Ang2-expressing gliomas were accompanied by up-regulated MMP-2, MT1-MMP, and LN 5 γ 2 in tumor cells that are actively invading the brain. Finally, stimulation of human glioma cells by overexpressing Ang2 or exposure to exogenous Ang2 induced the expression and activation of MT1-MMP, MMP-2, and LN 5 γ 2 in glioma cells *in vitro*. Together, our data provide evidence suggesting that up-regulation of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 in actively invading tumor cells correlates with human glioma invasion and stimulation of MMP-2, MT1-MMP, and LN 5 γ 2 are involved in Ang2-induced glioma invasion.

MMP-2, MT1-MMP, and LN 5 γ 2 are important in tumor invasion. In addition to the preferential expression in the invading tumor fronts found in various types of primary human cancers,^{4,7,25} these three molecules interact with each other in promoting tumor cell invasion. It has been shown that MMP-2, MT1-MMP, and TIMP-2 form a complex on the cell surface, where MT1-MMP binds to and activates pro-MMP-2 by proteolysis.⁵ Both MMP-2 and MT1-MMP also cleave LN 5 γ 2 within the γ 2 chain (155 kd) and generate proteolytic fragments, γ 2' (110 kd) and γ 2' X (80 kd). The released γ 2' X fragment is capable of promoting tumor cell invasion^{8,9} and preventing tumor

Figure 3. Co-overexpression and co-activation of MT1-MMP and LN 5 γ 2 in the invasive edge of primary human glioma specimens. **A:** An example of performing microdissection on the center region (**a**, circled area 1) and invasive edge of the tumor (**a**, circled area 2). **a:** An overview of the specimen (J151, GBM, grade IV). **b** and **c:** Photographs of the central areas (**1** in **a**) and invasive edge (**2** in **a**) before microdissection. **d** and **e:** Photographs of the central areas (**1** in **a**) and invasive edge (**2** in **a**) after microdissection. **b** and **c:** Enlarged photos of tumor center (circled area 1 in **a**). **c** and **e:** Tumor invasive edge (circled area 2 in **a**). **B:** Western blot analyses of total protein extracted from microdissected glioma tissues of tumor center (**C**, illustrated as area 1 in **A**; **a**, **b**, and **d**) and tumor invasive edge (**B**, illustrated as area 2 in **A**; **a**, **c**, and **e**). OD grade II, specimen J181, an oligodendroastrocytoma; AA, grade III, specimen J165, an astrocytic astrocytoma; and J151, J152, and J155, GBM, grade IV, GBM. **Top:** Expression and activation of LN 5 γ 2; **middle:** expression and activation of MT1-MMP (the second panel), and **bottom:** β -actin (a loading control). Mature LN 5 γ 2 runs at 155 kd and cleaved γ 2' and γ 2x run at 105 kd and 80 kd, respectively. Mature and autocleaved MT1-MMP run at 68 kd and 45 kd, respectively. β -Actin runs at 43 kd. The experiments were performed two independent times and similar results were obtained. Original magnifications: $\times 12.5$ (**Aa**); $\times 40$ (**Ab-Ae**).

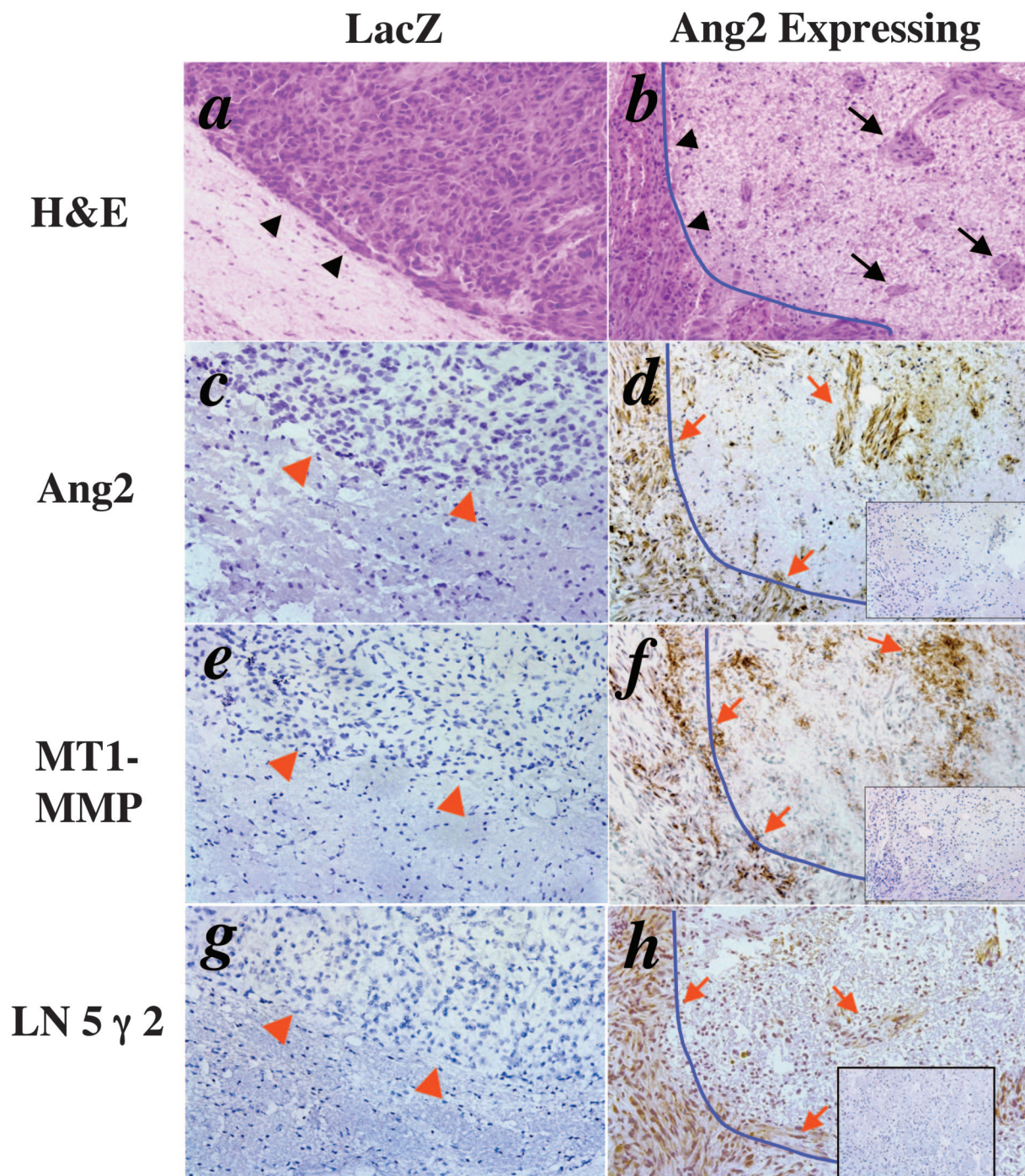


Figure 4. Overexpression of Ang2 by U87MG cells induces glioma invasion and promotes the expression of MT1-MMP and LN 5 γ 2 in the murine brain. IHC of various U87MG gliomas established in the murine brain. **a** to **d** were previously published in Hu and colleagues.¹⁷ The only purpose that we present here again is to serve as controls showing that co-overexpression of Ang2, MT1-MMP, and LN 5 γ 2 are localized in the same invading glioma cells in the brain. **a**, **c**, **e**, and **g**: A glioma established by control U87MG/LacZ cells. **b**, **d**, **f**, and **h**: An invasive glioma formed by U87MG/Ang2 cells. **a** and **b**: H&E staining. **Arrowheads** in **a** indicate the clean edge of U87MG/LacZ tumor spheroid. **Arrowheads** in **b** indicate spikes extended from tumor mass of the U87MG/Ang2 glioma. **Arrows** in **b** show disseminated tumor clusters of U87MG/Ang2 gliomas. **c** to **h**: IHC on serial sections of U87MG/LacZ or U87MG/Ang2 gliomas using anti-Ang2 (**c**, **d**), anti-MT1-MMP (**e**, **f**), and anti-LN 5 γ 2 (**g**, **h**) antibodies. **Arrowheads** in **c**, **e**, and **g** indicate the clean edges of U87MG/LacZ tumor spheroid. **Arrows** in **d**, **f**, and **h** show the invasive tumor-spikes and disseminated tumor clusters that expressed Ang2 (**d**), MT1-MMP (**f**), or LN 5 γ 2 (**h**). **Insets** in **d**, **f**, and **h** show isotype-matched IgG controls. Ten to twelve individual samples in each class were analyzed. The experiments were repeated two additional times with similar results. Original magnifications, $\times 200$.

cell apoptosis through the interaction with the epidermal growth factor receptor.²⁷ LN 5 γ 2 and MT1-MMP are found to be co-localized in invasive fronts of human breast cancers.⁹ Indeed, cooperative interactions of LN 5

γ 2, MMP-2, and MT1-MMP are required for tumor vascular mimicry vasculogenesis.^{28,29} Our results corroborate these observations. We found that up-regulated expression of MMP-2, MT1-MMP, and LN 5 γ 2, but not

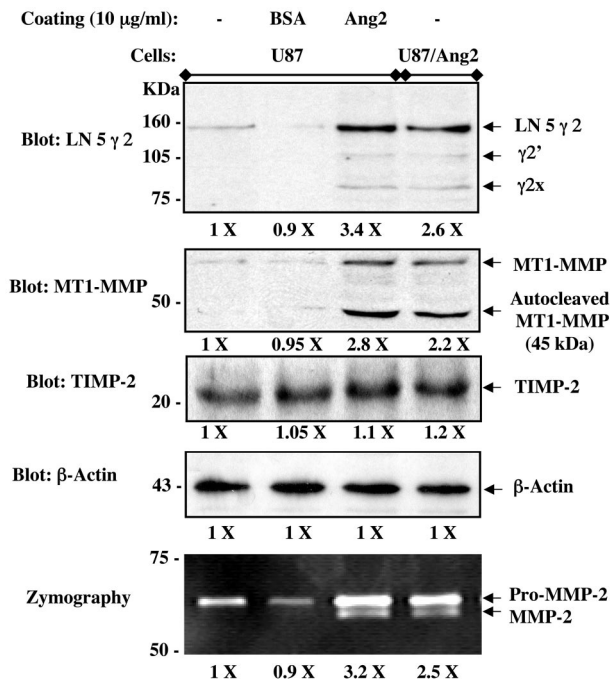


Figure 5. Stimulation of human glioma cells by overexpressing Ang2 or exogenous Ang2 promotes the expression and activation of MMP-2, MT1-MMP, and LN 5 γ 2, but not TIMP-2 *in vitro*. **Top** four panels: Western blot analyses for expression and activation of LN 5 γ 2 (the first panel from the **top**), MT1-MMP (the second panel), TIMP-2 (the third panel), and β -actin (the fourth panel, a loading control). Mature LN 5 γ 2 runs at 155 kd and cleaved γ 2' and γ 2x run at 105 kd and 80 kd, respectively. Mature and autocleaved MT1-MMP run at 68 kd and 45 kd, respectively. TIMP-2 runs at 22 kd. β -Actin runs at 43 kd. **Bottom:** Zymography assays for the activation of MMP-2. The pro-MMP-2 and the mature form of MMP-2 are shown at 72 kd and 64 kd, respectively.

TIMP-2, co-localized with the increased Ang2 in the invasive areas of primary human glioma specimens, in U87MG Ang2-expressing gliomas *in vivo* as well as in Ang2-stimulated glioma cells *in vitro*. Interestingly, stronger proteolytic co-activation of LN 5 γ 2 and MT1-MMP were found in the proteins extracts from microdissected primary glioma invasive tissues (Figure 3B) than the Ang2 stimulation (by overexpression or exogenous treatment with recombinant Ang2) of U87MG glioma cells *in vitro*, suggesting an optimal tumor microenvironment *in vivo* was more potent in stimulating the activities of these two extracellular matrix-modifying enzymes critical for glioma cell invasion. Thus, our data indicate that co-overexpression of these molecules is involved in glioma invasion.

The ability of glioma cells to diffusively infiltrate normal brain tissues without disrupting brain cytoarchitecture or neuronal function is a hallmark of malignant human gliomas.¹⁻³ Glioma invasion does not correlate with glioma grade because low-grade astrocytomas invade brain parenchyma with high frequency. However, the degree of tumor invasiveness is correlated with glioma grade.¹⁻³ In this study, seven of eight PAs displayed an infiltrative phenotype (Table 1) whereas grade II, III, and IV glioma specimens in our collection contain distinct invasiveness with identifiable tumor border and invasive areas. Expression profiles of Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 are correlated with the invasive edge of human gliomas, especially in the invasive areas that are 0.25 mm away

from the tumor border. Increased levels of these proteins were evident in the invasive regions in these grade I to IV tumors (Tables 1 and 2; Figures 2A, 2B, and 2C, f, i, l, o; and data not shown) and in the tumor borders (Tables 1 and 2; Figures 2A, 2B, and 2C, e, h, k, n; and data not shown). When compared with the other three molecules, the expression profiles of LN 5 γ 2 were relatively increased in both invasive and noninvasive regions of grade IV GBM tumors (Table 1). Strong activation of LN 5 γ 2 at the invasive edge of the tumor (Figure 3B) supports our IHC data that this tumor invasion marker is involved in the progressive growth of malignant GBM tumors. These data show that up-regulation of these four molecules, particularly Ang2, MMP-2, and MT1-MMP, is correlated with the invasiveness of malignant human gliomas.

In summary, our data demonstrate that co-localization of up-regulated Ang2, MMP-2, MT1-MMP, and LN 5 γ 2 is significantly associated with human glioma invasion. Our results from the engineered human U87MG glioma (Ang2-overexpressing, Figure 4) xenografts and cell culture model (stimulation of glioma cells by overexpressing Ang2 or exposure to exogenous Ang2, Figure 5), and from previous reports³⁰⁻³³ suggest that Ang2 plays a critical role in promoting human glioma invasion through the activation of MMP-2, MT1-MMP, and LN 5 γ 2 in Tie2-independent-pathways. The determination of the mechanisms by which Ang2 induces glioma invasion could provide critical information with regard to the potential of Ang2 and its effectors as new therapeutic targets in glioma treatments.

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