Doxycycline Suppresses Cerebral Matrix Metalloproteinase-9 and Angiogenesis Induced by Focal Hyperstimulation of Vascular Endothelial Growth Factor in a Mouse Model

Chanhung Z. Lee, MD, PhD; Bin Xu, MD; Tomoki Hashimoto, MD; Charles E. McCulloch, PhD; Guo-Yuan Yang, MD, PhD; William L. Young, MD

Background and Purpose—A number of central nervous system (CNS) disorders are associated with abnormalities in or activation of angiogenesis, including vascular malformations. To test the hypothesis that the nonspecific matrix metalloproteinase (MMP) inhibitor, doxycycline, suppresses vascular endothelial growth factor (VEGF)-induced cerebral angiogenesis through inhibition of MMPs, we used a mouse model with enhanced cerebral angiogenesis induced by focal hyperstimulation of VEGF from adenovirus DNA (AdVEGF) transduction.

Methods—The time course study of MMP activity was performed at 7 and 14 days after AdVEGF transduction. MMP activity and expression were examined by zymography and immunohistochemistry, respectively. As an index of cerebral angiogenesis, microvessel counting was performed in the brains of 3 groups of mice (n=110): (1) control; (2) AdVEGF only; and (3) AdVEGF plus doxycycline (30 mg/kg per day).

Results—Brain MMP-9 activities increased 4-fold (883±137 versus 179±179; 1-sided P<0.001) at 7 days after AdVEGF transduction. VEGF transduction increased vessel counts by 19% (255±27 versus 215±15, 1-sided P<0.01). Doxycycline treatment decreased MMP-9 activity (89±72 versus 883±137; 1-sided P<0.001) and cerebral microvessel number (231±17 versus 255±27; 1-sided P<0.05).

Conclusions—Doxycycline is effective in decreasing stimulated cerebral MMP-9 activity and parenchymal angiogenesis. The decrease in MMP-9 activity is associated with decreased microvessel counts. Brain pathophysiological processes that involve abnormally enhanced angiogenesis may be amenable to manipulation by MMP inhibitors, including tetracycline derivatives. (Stroke. 2004;35:1715-1719.)

Key Words: drug therapy ■ metalloproteinases ■ angiogenesis

There is growing appreciation that a number of central nervous system (CNS) disorders are associated with abnormalities in or activation of angiogenesis,1,2 including ischemic stroke3,4 and brain arteriovenous malformations (BAVMs). Little is known about the pathogenesis of BAVMs, in part because there is no good animal model that mimics the human disease of recurrent spontaneous intracranial hemorrhage. Recent data suggests that BAVMs are associated with excessive vascular remodeling caused by pathologically increased angiogenesis.5,6,7

VEGF and MMPs are among the most potent regulators of angiogenesis. VEGF is the specific growth factor for endothelial cells and major regulator of blood vessel formation, ie, angiogenesis. VEGF165, typically expressed as a 46-kDa homodimer, is the most biologically active form in in vitro studies.8 Angiogenesis also requires degradation of vascular matrix proteins. MMPs degrade extracellular matrix proteins, cell surface molecules, and other pericellular substances.9 Recent findings have indicated that gelatinases, including A (MMP-2, constitutive) and B (MMP-9, inducible), in particular, play a central role during angiogenesis. Studies have shown that VEGF and MMP also influence each other in the process of angiogenesis,10–12

Altered expression of VEGF ligand and VEGF receptors have been described in surgical BAVM specimens13 and increased VEGF expression has been linked to recurrent BAVMs.14 We have recently described increased levels of MMP-9 relative to tissue inhibitors of metalloproteinase in surgical BAVM specimens.15 Taken together, evidence from clinical studies suggests that VEGF and MMPs may contribute to the development or maintenance of the diseased vascular phenotype.

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From the Departments of Anesthesia and Perioperative Care (C.Z.L., B.X., T.H., G.-Y.Y., W.L.Y.), Epidemiology and Biostatistics (C.E.M.), Neurological Surgery (G.-Y.Y., W.L.Y.), and Neurology (W.L.Y.), and the Center for Cerebrovascular Research (C.Z.L., B.X., T.H., G.-Y.Y., W.L.Y.), University of California, San Francisco.

Correspondence to Dr William L. Young, Center for Cerebrovascular Research, 1001 Potrero Ave, Room 3C-38, San Francisco, CA 94110. E-mail ccr@anesthesia.ucsf.edu

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There is growing evidence that MMP inhibition may be useful in the management of vascular diseases.16,17 Tetracycline derivatives, including doxycycline, have nonspecific MMP inhibitory effects that are distinct from their antimicrobial action.18 To study the effect of MMP inhibition on states of enhanced angiogenesis, we used a murine model newly developed in our laboratory. The model consists of focal hyperstimulation by VEGF<sub>ad</sub> in the brain using adenovirus-mediated DNA transduction, resulting in enhanced formation of cerebral microvessels.19,20 Our model is not a model of BAVM, but it is characterized by some of the phenotypic features of BAVM lesional tissue. Therefore, use of this model is a first step to mechanistically examine key angiogenic pathways and their response to pharmacological manipulation.

We hypothesized that the nonspecific MMP inhibitor, doxycycline, can suppress AdVEGF-induced cerebral angiogenesis and the suppression is mediated through inhibition of MMP-9 expression and activity. To test this hypothesis, we examined the effects of doxycycline on MMP activities and microvessel formation in the mouse brain after adenovirus-mediated VEGF transduction.

**Materials and Methods**

**Animals and Treatment**

Animal use was approved by the University of California San Francisco Committee of Animal Research. Male CD-1 mice weighing 30 to 35 g were purchased from Charles River Laboratory (Wilmington, Mass). The mice were allowed free access to food and water with a 12-hour alternating light–dark cycle.

Our previous data have shown that VEGF expression increases at day 5 in the mouse brain after AdVEGF transduction compared with the control group, with the peak of microvessel counts occurring later, at ~3 weeks.19 This finding suggests that there is significant lag time between the activation of angiogenic factors and the actual formation of new vessels. At day 3, mild inflammatory responses around the needle track was detected in both Ad<i>lac</i> and Ad<i>VEGF</i> groups to a similar degree.19 A time course study at 7 and 14 days after AdVEGF transfection was performed to determine the appropriate time point for assessing MMP expression change induced by VEGF overexpression.

To study the effect of doxycycline treatment on cerebral angiogenesis induced by AdVEGF transfection, the mice were divided into 3 groups: control, Ad<i>VEGF</i>, and Ad<i>VEGF</i> with doxycycline treatment. The control and Ad<i>VEGF</i> groups received Ad<i>lac</i> and Ad<i>VEGF</i> injection, respectively. In the treatment group, doxycycline (Sigma) was administered starting on the day of Ad<i>VEGF</i> injection, at 30 mg/kg per day via drinking water, a dose shown to inhibit VEGF administration and a reduction with the added administration of doxycycline, 1-sided <i>P</i> values were used for those comparisons; <i>P</i>&lt;0.05 is considered statistically significant.

**Statistical Analysis**

Data are expressed as mean±standard deviation. Parameters between different groups in the MMP expression time course study and doxycycline treatment study were analyzed using 2-way ANOVA and Student t test. Because theory and previous results distinctly predict an increase of MMP activity and microvessel formation with VEGF administration and a reduction with the added administration of doxycycline, 1-sided <i>P</i> values were used for those comparisons; <i>P</i>&lt;0.05 is considered statistically significant.

**Results**

To determine the appropriate time point to examine MMP expression induced by VEGF, the time course of MMP expression after AdVEGF transduction was studied (Figure 1). At 7 days, acute inflammatory response was minimal in all of the animals by H&E staining (data not shown). At day 7 after gene transfer, MMP-9 activities were increased near 4-fold in the AdVEGF transducted mouse brain compared with the ones with Ad<i>lac</i> (883±137 versus 179±179, arbitrary units [AU] 1-sided <i>P</i>&lt;0.001). At day 14, there was little MMP-9 detected in either of the groups (73±18 versus 165±94 AU, 1-sided <i>P</i>&gt;0.05), and the differences in MMP-9 levels were greater at 7 days than at 14 days (2-way factorial test for interaction, <i>P</i>&lt;0.01). Therefore, day 7 after gene transfer appeared to be an appropriate time for the MMP study. Unlike MMP-9, there was no significant change in MPP-2 expression in response to VEGF either at day 7 (3363±1204 versus 2997±998 AU, 1-sided <i>P</i>&gt;0.2) or at day
Immunohistochemistry staining was performed to compare the distribution of MMP-9 expression at day 7 after gene transfer (Figure 2). Diffuse positive staining of MMP-9 was distributed in surrounding areas of the needle track after AdVEGF transduction, as illustrated in Figure 2B, but not in the AdlacZ mouse brain (Figure 2A).

Doxycycline suppressed MMP-9 expression at 7 days after AdVEGF transduction in mouse brain (Figure 3). MMP-9 activities were much lower in doxycycline-treated mice than those that did not receive the drug (88 ± 72 versus 88 ± 137 AU, 1-sided P < 0.001). In contrast, the AdlacZ group was not statistically significantly different from the AdVEGF plus doxycycline group (179 ± 179 versus 89 ± 72 AU, 2-sided P > 0.4). In other words, the VEGF-induced MMP-9 activities were completely diminished by doxycycline treatment. Again, unlike MMP-9, there was no change of MMP-2 activity in response to doxycycline treatment (2997 ± 998 versus 2684 ± 821 AU, 1-sided P > 0.3) (Figure 3A).

To determine the effect of doxycycline on cerebral angiogenesis, 3 groups of mice (n=6 in each group) at 3 weeks after adenoviral DNA transduction were used for comparison. Mice with AdlacZ transfer in the brain were used as the control group. Our results showed that doxycycline decreased microvessel counts induced by focal VEGF hyperstimulation in the mouse brain (Figure 4). VEGF transduction increased microvessel formation by 19% in the mouse brain in comparison with the lacZ group (255 ± 27 versus 215 ± 15, number of microvessels; 1-sided P < 0.01). The number of microvessels was lower in the doxycycline group than in the VEGF group (231 ± 17 versus 255 ± 27, number of microvessels; 1-sided P < 0.05).

Discussion
Our results have demonstrated for the first time to our knowledge that doxycycline suppresses VEGF-induced cerebral MMP-9 activity in vivo. The doxycycline-induced changes in MMP-9 activity were associated with decreased regional angiogenesis, as evidenced by decreased microvessel counts.

Previous reports have shown that tetracycline derivatives, including doxycycline, influence many aspects of the angiogenesis process. Many of those data, however, were obtained from in vitro systems or from large blood vessels in animals. First, there have been conflicting results of the effect of doxycycline...
on cell growth between in vitro and in vivo studies, suggesting the possibility of different mechanisms. For example, in vivo studies on smooth muscle cell proliferation showed that effects of doxycycline differ from known MMP inhibitors, including GM6001 \(^{22}\) and BB94. \(^{23}\) However, in vitro studies using tissue culture showed contradictory results. One reported that doxycycline inhibited angiogenesis whereas GM6001 did not. \(^{24}\) The other one reported that the antiangiogenesis effects of tetracycline derivatives were associated with inhibition of MMP activities. \(^{25}\) Secondly, vascular endothelium in the central nervous system may be functionally distinct from the endothelium of other organ systems. Doxycycline has been shown to inhibit MMP-9 in aorta homogenate cultures and in human and animal aortic aneurysm studies. \(^{26,27}\) However, there are significant differences in baseline endothelial MMP activities between the brain microvessel and the aorta. For example, MMP-9 levels increased in the brain microvessel endothelial cells in response to stimulation by inflammatory cytokines, whereas no change was observed in the aortic endothelial cells. \(^{28}\) Our finding that doxycycline inhibits MMP-9 activity and the formation of capillaries in the mouse brain provides evidence that doxycycline can influence cerebral angiogenesis.

VEGF and MMPs are considered as potent regulators of angiogenesis. Our findings have shown that focal VEGF hyper-stimulation is associated with increased MMP-9 expression in the mouse brain. Similarly, Lamoreaux et al reported that VEGF increases the release of another gelatinase, MMP-2, and decreases the release of tissue inhibitors of metalloproteinases by microvascular endothelial cells in vitro. \(^{10}\) Further, MMP-9 can facilitate the availability of tissue-bound VEGF, \(^{11,12}\) which in turn may potentiate angiogenic activities.

There is somewhat contradictory evidence of MMP-9 and MMP-2 in response to stimulatory or inhibitory factors. \(^{26,28,29}\) In our study, there was no change in MMP-2 levels with VEGF stimulation or after doxycycline treatment, in contrast to changes in microvessels and MMP-9 levels. In the brain, in addition to endothelial cells and smooth muscle cells, astrocytes constitutively produce MMP-2. \(^{30}\) MMP-2 activities that we detected from all groups of mice regardless of the treatment could be a reflection of constitutive expression of MMP-2, which, in this case, does not respond to either the stimulation from VEGF or the inhibition from doxycycline.

There are several limitations to our study. As pointed out, this model is not a specific model for any particular disease but can...
allow mechanistic investigations of various potential interventions in the angiogenic process. Ideally, we would have had a dose–response study, but the dose used appears to be sufficient to provide proof-of-concept for the hypothesized effects of doxycycline, taken together with other information in the literature. An inflammatory tissue response from adenosivel transduction may confound the direct effects of VEGF overexpression, but our previous studies have demonstrated a minimal degree of acute inflammation with this model.10,11 Finally, we have only demonstrated an association between decreased MMP-9 activity and diminished ability of VEGF to induce capillary angiogenesis; further studies can better characterize the causal relationship of the 2 observations.

In conclusion, the present study has demonstrated that doxycycline can reduce MMP-9 activity and angiogenesis induced by focal VEGF hyperstimulation in the mouse brain. The ability to manipulate angiogenesis may have importance in the study of various CNS disorders, and in particular may be of interest in developing models to study the pathogenesis of brain vascular malformations. The mechanism of the effect of doxycycline on brain angiogenesis in relation to its anti-MMP activity remains to be further clarified.

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