Vasculogenic mimicry: a novel target for glioma therapy

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Abstract

Anti-angiogenic therapy has shown promising but insufficient efficacy on gliomas. Recent studies suggest that vasculogenic mimicry (VM), or the formation of non-endothelial, tumor-cell–lined microvascular channels, occurs in aggressive tumors, including gliomas. There is also evidence of a physiological connection between the endothelial-lined vasculature and VM channels. Tumor cells, by virtue of their high plasticity, can form vessel-like structures themselves, which may function as blood supply networks. Our previous study on gliomas showed that microvessel density was comparably less in VM-positive tumors than in VM-negative tumors. Thus, VM may act as a complement to ensure tumor blood supply, especially in regions with less microvessel density. Patients with VM-positive gliomas survived a shorter period of time than did patients with VM-negative gliomas. Although the detailed molecular mechanisms for VM are not fully understood, glioma stem cells might play a key role, since they are involved in tumor tissue remodeling and contribute to neovascularization via transdifferentiation. In the future, successful treatment of gliomas should involve targeting both VM and angiogenesis. In this review, we summarize the progress and challenges of VM in gliomas.

Key words Glioma, vasculogenic mimicry, target therapy
Current Understanding of the Vascularization Process in Tumors

Angiogenesis and vasculogenesis are widely accepted processes of tumor vascularization, particularly for endothelium-dependent vessels. In both processes, tumor vascular endothelial cells develop from host cells located in normal tissues around the tumor or from endothelial progenitor cells. During the transition from endothelium-dependent vessels to mimicked vessels, mosaic vessels occur as a transitional type between endothelium-dependent vessels and VM channels, wherein both the host endothelium and tumor cells participate in tumor vascularization. VM is completely different from angiogenesis and vasculogenesis, in another word, the blood supply to tumors is proposed to involve three types: tumor-cell-lined vessels, mosaic vessels, and endothelium-dependent vessels. A study suggest that VM channels—the tumor-cell-lined vessels—could be the main source of blood supply in the early stage of tumor growth. Endothelium-dependent vessels could then replace VM channels via a transitional step as mosaic vessels to become the dominant blood supply pattern at the late stage of tumor growth.

Cancer stem cells (CSCs), or tumor-initiating cells, were identified as a unique subpopulation with stem cell features in many types of cancer. Current CSC studies provide novel insight into tumor angiogenesis and its interplay with the tumor microenvironment. CSCs have been shown to promote tumor angiogenesis by secreting vascular endothelial growth factor (VEGF) and via their potential for transdifferentiation into endothelial cells. Rebetz et al. showed that CD133+ cells could originate from either tumor blood vessels or gliomas. In another study, He et al. found that blood vessels near CD133+ or nestin niches as well as some CD31+ vessels co-expressed CD133 or nestin. Dong et al. reported that glioma stem cells (GSCs) were involved in tumor tissue remodeling in a xenograft model. Similarly, Ricci-Vitiani et al. and Wang et al. found that a subpopulation of cells within glioma can give rise to endothelial cells. In summary, the angiogenesis capacity of GSCs has been demonstrated; however, the detailed relationship of GSCs and the phenomenon VM in glioma is still unclear.

Biological Characteristics and Clinical Significance of VM in Gliomas

VM channels are negative for CD34 (and other endothelial blood vessel markers, such as CD31) and positive for periodic acid-schiff (PAS). VM can be identified histologically based on three elements: the plasticity of malignant tumor cells, the remodeling of the ECM, and the connection of VM channels to the host microcirculation system. VM has been observed in many human tumors, including melanoma, pancreatic cell carcinoma, breast cancer, ovarian carcinoma, primary gallbladder carcinoma, malignant esophageal stromal carcinoma, mesothelial sarcoma, and alveolar rhabdomyosarcoma, hepatocellular carcinoma, prostatic carcinoma, bladder carcinoma, osteosarcoma, and pheochromocytoma. We used CD34 staining to identify the endothelium in glioma tissue sections and PAS staining to determine the basement membrane of tumor blood vessels. Real tumor vessels stained positive for CD34 on their luminal surface and for PAS in their walls. However, in a subset of gliomas, we observed PAS-positive tubular structures that contained red blood cells but were lined by CD34-negative cells in the luminal surface, i.e., VM. VM was also found in human glioma cell line xenografts.

As described by Folberg et al., seven morphologic patterns of VM channels were identified in uveal melanoma: straight channels, arrangements of parallel straight channels, straight channels that cross-link, arcs (incompletely closed loops), arcs with branching, closed loops, and networks (networks were defined arbitrarily as at least three back-to-back, closed, PAS-positive loops). In our study, we found that VM channels can be sorted into two distinct types: tumor-cell-dependent (wherein CD34-negative tumor cells imitate the functions of the endothelium) and ECM-dependent (which involves only PAS-positive ECM without tumor cells). Neither type of tubing is dependent on the endothelium, but both contribute to microcirculation in gliomas.

Maniotis’ discovery of VM originally spawned controversy. However, different approaches have since confirmed that VM channels provide a mechanism of perfusion and a dissemination route within the tumor that functions either independently of or simultaneously with angiogenesis. Studies have demonstrated the functional role of VM channels in tumor circulation using several methods, including microinjection, Doppler ultrasonography, magnetic resonance imaging (MRI), laser scanning confocal angiography, and injection of absorbite particles. In our study, we found that the distribution of VM channels in gliomas looked patchy and abundant by the CD34-PAS double staining. Furthermore, VM channels were always located in regions where endothelium-dependent vessels were not found, and no necrosis or surrounding inflammatory cells were observed nearby. We investigated the potential association between VM and MVD in 48 glioblastomas and found that the MVD was comparably less in VM-positive tumors than in VM-negative tumors. This evidence supports that VM channels may be a complementary system to ensure tumor blood supply, especially in regions with less MVD.

For cancer patients, VM is associated with poor prognosis, as the unique structure of VM channels facilitates tumor cell metastasis. Tumor cells, which line the inner surface of VM channels, are directly exposed to blood flow, allowing them to leak out, migrate through the blood stream, and metastasize to other regions. Furthermore, tumor cells that line the VM channel are highly malignant, are poorly differentiated, and have high plasticity. These cells can degrade adjacent connective tissue and penetrate the basement membrane of blood vessels by secreting proteins that mediate tumor invasion and metastasis. This phenomenon has been confirmed in liver cancer, breast cancer, gastrointestinal stromal tumors, and glioblastomas. We performed a retrospective analysis on 101 glioma patients. Tumor samples were co-stained for CD34 and PAS. Then, the dual stained samples were stained for Ki-67, cyclooxygenase-2.
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Figure 1. Vasculogenic mimicry (VM) in human gliomas. These representative images show gliomas dual stained for CD34 and periodic acid-schiff (PAS) using immunohistochemistry. A, endothelial cells are detected with anti-CD34 (dark brown), and vascular basement membrane is detected with PAS (purple magenta) in normal tubular blood vessels. B, microvascular proliferation of glioblastoma. C–F, typical VM channels (denoted by black arrows). The channels are located in a viable area of the tumor, far from necrosis. C and E, seven morphologic patterns of PAS-positive channels. D and F, VM channels containing red blood cells positive for PAS but negative for CD34: large cross section (D) and longitudinal section (F). Magnification: A, B, D, E, and F, ×400; C, ×100. (Cited from Yue et al. [17]. The authors have got the permission to reprint this image.)

(COX-2), and matrix metalloproteinase-9 (MMP-9). VM was detected in 13 of 101 samples and was more frequent in high-grade gliomas than in low-grade gliomas. VM channels were also associated with expression of COX-2 and MMP-9. Patients with VM-positive tumors survived a shorter period of time than did patients with VM-negative tumors.

Targeting Vasculogenic Mimicry for Glioma Therapy

In 1971, Folkman [9] reported that tumors require a blood supply for survival, growth, and metastasis, and argued, for the first time, that anti-angiogenic therapy would have significant efficacy on
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Advances and Challenges

Abnormal, dysfunctional tumor vasculature and GSCs are believed to be major obstacles for effective glioma treatment. VM may represent an important tumor survival mechanism and may contribute to the failure of current anti-angiogenic therapy, which aims to completely deprive tumors of their blood supply. Targeting VM along with endothelium-dependent vessels may thus block the supply of oxygen and nutrition to tumor cells effectively and completely. Furthermore, the unique structure of VM channels directly exposes tumor cells, which line the channels’ inner surface, to blood vessels, thereby facilitating metastasis. VM is frequently seen in the regions between the tumor and surrounding normal tissues and is associated with poor prognosis. Therefore, therapies targeting VM channels have the potential to destroy the niche that maintains GSCs, block the passage through which tumor cells metastasize, and reduce cancer recurrence.

Nevertheless, tumor vascularization is a complex process that involves concomitant activity of several distinct pathways that may vary according to the patient, tumor type, tumor grade, and therapeutic effect. Successful treatment of gliomas should involve targeting one or more stages in the VM signaling cascade. Three factors affect VM channel formation: the plasticity of VM channel-associated tumor cells, the remodeling of extracellular matrix, and the connection of VM channels with the host microcirculation. Thus, anti-VM therapy should focus on inhibiting tumor cell plasticity as well as remodeling the ECM and tumor microenvironment by blocking the biochemical and molecular pathways underlying VM. However, further studies on the mechanisms of VM are needed to determine the potential for future translational studies and clinical applications.
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