TBX20 and the PROK2-PROKR1 pathway—new kid on the block in angiogenesis research

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Introduction

The term angiogenesis stems from the Greek expression for “angeíon” (vessel) and “genesis” (formation, development) and describes the formation of new blood vessels. Technically speaking, angiogenesis means the formation and sprouting of blood vessels from pre-existing vascular structures. By comparison, vasculogenesis is the de novo development of blood vessels from angioblasts (1). During embryogenesis, initial vascular structures are formed through vasculogenesis, which is followed by angiogenesis for further organ development and also blood vessel formation under physiological and pathophysiological conditions (such as tissue growth, wound healing, the female reproductive cycle, but also in cancer and ischemia). Angiogenesis is considered a pivotal process in almost all tissues in health and disease (2). The first step to induce angiogenesis is associated in most cases with pathological events and/or activating stimuli. Angiogenesis is a dynamic process that requires endothelial cells to transition from a resting to an activated state. This step involves the formation of a new vascular sprout and, once this is accomplished, the newly formed vascular structure is elongating. The subsequent process of further sprouting is a tightly regulated action of proliferation (3), cell migration (4), degradation of basement membranes to facilitate further expansiveness, vessel lumen formation, and last but not least also tissue stabilization (5). However, endothelial cells are just a part of the whole picture, as angiogenesis is further regulated by growth factors (e.g., vascular endothelial growth factor, VEGF), cytokines, chemokines, metalloproteinases, and on an intracellular level by activation of signaling kinases and gene transcription (6).

Here in this editorial, we want to focus on the TBX20-Prokineticin 2 (PROK2)-prokineticin receptor (PROKR1) signaling pathway.

The study by Meng et al.

In the recently published study by Meng et al., the authors sought to analyse the function of the key transcription factor T-box 20 (TBX20) and the signaling pathway which is activated during angiogenesis (7). TBX20 is considered a pivotal transcription factor during embryogenesis. Furthermore, it has been shown that its deficiency is associated with congenital heart disease. However, very little is yet known about its role in angiogenesis and vessel formation. Meng et al. determined the role of TBX20 in angiogenesis in a very elaborate approach using loss- and gain-of-function assays by use of in vitro and in vivo experiments. TBX20 was shown to be an important regulator in cardiac development and knockout of Tbx20 in mice resulted in cardiac defects and embryonic lethality, whereas in human, mutations in the TBX20 gene are associated with different cardiac abnormalities (septation
defects, altered chamber formation and cardiomyopathy) (8). A gene array was used to find key downstream targets of TBX20, which identified TBX20 as a novel transcription factor for signaling through the PROK2-PROKR1 pathway in the regulation of angiogenesis. Furthermore, the consequences of TBX20 activation or blocking were analysed in three different in vivo models. By application of a murine model of angiogenesis using subcutaneously implanted matrigel plugs saturated with VEGF, the authors observed that TBX20 deficiency led to reduced expression of PROK2 and also hindered angiogenesis. Additionally, a murine model of hind limb ischemia (induced by ligation of the femoral artery) was utilized to assess the effects of recombinant PROK2 administration. Here, enhanced angiogenesis and increased blood flow was documented in ischemic legs using Laser Doppler perfusion measurements. In a zebrafish model, the transient knockdown of TBX20 also resulted in impairment of angiogenesis.

In these experimental settings, the authors determined new and yet unknown functions of TBX20 and PROK2. PROK2, a secreted protein has been shown to influence angiogenesis, circadian rhythm, reproductive function and neurogenesis (9-11). Here in this study, the authors analyzed the whole TBX20-PROK2-PROKR1 axis and gained broad knowledge on its function and regulation in angiogenesis. Furthermore, they showed that TBX20 controls endothelial cell migration but not their proliferation. VEGF and its receptor VEGFR2 are known to be important regulators in angiogenesis. VEGFR2 activation by VEGF triggers downstream signaling cascades such as PI3K-Akt, PLC-γ-ERK1/2, SRC-Small GTPase and p38 (12,13). In this concise experimental setup by Meng et al. the TBX20-PROK2-PROKR1 axis, which can be activated by VEGF, joined these aforementioned pathways, as relevant alternative mediators in angiogenesis and can help to further sustain the pro-angiogenic effects of VEGF (see Figure 1).

**Future implications**

Angiogenesis is a fundamental process in both health and disease (14). Whereas considered beneficial as in wound healing and in ischemic diseases, overshooting angiogenesis can lead to cancer progression. Therefore, as angiogenesis itself is such a double-edged sword, a thorough knowledge of how it is regulated, initiated and deactivated is of paramount importance in order to better understand (patho-)physiological processes in many diseases.

Apart from angiogenesis, the understanding of intracellular signaling and its consequences are of utmost importance. As for angiogenesis, the same cellular signaling pathways (e.g., PI3K-Akt) are involved in cell activation...
and proliferation, which are beneficial effects in ischemia and tissue damage, and laterally reversed, of detrimental effects in oncogenesis. Yellon, Hausenloy and others have determined that activation of the ERK, Akt and PI3K kinases are involved in conditioning mechanisms in ischemia and can reduce the extent of tissue damage in myocardial infarction (15-18). On the other hand, the activation of the same signaling axes can lead to tumor progression in cancer (19,20). Targeting single signaling factors of these pathways is thought to be an attractive strategy for the development of new drug compounds and novel therapeutic strategies to treat different forms of cancer. A fundamental knowledge of all these cross-linked signaling axes is of importance as intra- and inter-pathway compensatory loops exist in these pathways that can lead to the reactivation of trigger factors in the same cascade (upstream or downstream of the point of blockade). Moreover, as also underpinned by the data by Meng et al., activation of alternate signaling pathways can occur, which can hinder the effectiveness of cancer therapies.

Over recent years, our knowledge of molecular mechanisms in signaling axes has shown advances with a rapid increasing speed. Still, our understanding of how signaling factors interact and different pathways are interlocked with each other requires further research in order to provide the scientific background for the development of new drug agonist or antagonists to be applied in different disease conditions.

The data provided in this current experimental study provides a new jigsaw piece in the whole complex picture of intracellular signaling in angiogenesis. The TBX20-PROK2-PROKR1 pathway described by Meng et al. could also be a therapeutic target to treat disease conditions that are associated with dysregulated angiogenesis. It could provide therapeutic angiogenesis in patients with ischemic heart failure, stroke or peripheral occlusive artery disease as PROK2 has been shown to improve blood perfusion and functional recovery.

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**Footnote**

*Conflicts of Interest:* The authors have no conflicts of interest to declare.

**References**


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