“Splitting the matrix”: intussusceptive angiogenesis meets MT1-MMP

Gabriela D’Amico†, José M Muñoz-Félix†, Ana Rita Pedrosa† & Kairbaan M Hodivala-Dilke‡

Pathological angiogenesis contributes to tumour progression as well as to chronic inflammatory diseases. In this issue of EMBO Molecular Medicine, Esteban and co-workers identify endothelial cell MT1-MMP as a key regulator of intussusceptive angiogenesis (IA) in inflammatory colitis. Thrombospondin 1 (TSP1) cleavage by MT1-MMP results in the binding of the c-terminal fragment of TSP1 to αvβ3 integrin, which induces nitric oxide (NO) production, vasodilation and further initiation of IA. This novel control mechanism of inflammatory IA points towards promising new therapeutic targets for inflammatory bowel disease.

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See also: S Esteban et al (February 2020)

Angiogenesis, the formation and growth of blood vessels, plays an important role in normal development and ischaemic diseases. Increasing evidence supports the existence of different mechanisms of blood vessel growth, including sprouting and intussusceptive angiogenesis (IA). Both sprouting and IA occur under normal physiological and pathological conditions, including tumour growth (De Spiegelaere et al., 2012).

Sprouting angiogenesis, where blood vessel endothelial cells locally proliferate, bud and form branching vessels off a main vessel, is an invasive process initiated by multiple matrix metalloproteinasises (MMPs). These enzymes degrade extracellular matrix proteins, allowing vascular endothelial cells to invade the tissue and generate vessel sprouts. In contrast, IA is the expansion of the microvasculature through the formation of intraluminal pillars via invagination of the vessel wall into the vascular lumen, ultimately resulting in vessel “splitting”. Blood flow dynamics appear to be crucial in shaping this process, and neither pillar formation nor IA itself appear to be invasive processes (Paku et al., 2011). However, little is known about the cellular and molecular control of IA, highlighting the need for improved in vivo and in vitro models (De Spiegelaere et al., 2012).

Sprouting angiogenesis often precedes IA, especially in the tumour setting (Karthik et al., 2018). Importantly, since IA is stimulated after initiation of anti-angiogenic therapy, it constitutes an alternative target to combat vascular disease or resistance to anti-angiogenic therapy.

In this issue of EMBO Molecular Medicine, Esteban et al report a previously unrecognised role for endothelial MT1-MMP in IA (Esteban et al., 2020). The expression of the membrane protease MMP family member MT1-MMP, also known as MMP-14, is upregulated in angiogenesis and in inflammatory diseases; however, its potential implications in either IA or colitis were unknown.

Here, in a DSS-induced model of colitis, mice lacking endothelial cell-MT1-MMP had reduced number of IA capillary events, resulting in ameliorating colitis symptoms. Using a new whole-mount imaging method to overcome previous limitations in identifying and analysing IA (Hlushchuk et al., 2011; Nowak-Sliwinska et al., 2018), the authors showed that MT1-MMP expression levels were separately controlled in arterioles and capillaries, leading to a novel bimodal role of action for this enzyme. Constitutive MT1-MMP expression in arterioles drove NO production, increasing blood flow in the downstream capillary plexus. However, in the capillary y-junctions, DSS-induced upregulation of MT1-MMP drove NO production, which led to endothelial cell remodelling and pillar formation. Although not confirmed in the current study, the authors previously showed that NO regulates MT1-MMP activity during endothelial cell migration, therefore suggesting that, by a positive feed-back loop, NO could further reinforce MT1-MMP upregulation (Genis et al., 2007). Moreover, by combining a Cleavpredict search with in silico protein modelling, the authors identified that MT1-MMP drives TSP1 cleavage, allowing c-terminal TSP1 fragment release and interaction with CD47/αvβ3 integrin in the induction of NO during IA (Fig 1). This elegant in silico analysis could potentially be expanded to aid identification of novel substrates and the subsequent development of cell-specific strategies.

Targeting αvβ3 integrin has been extensively studied in angiogenesis (Robinson & Hodivala-Dilke, 2011), and some inhibitors such as the RGD-mimetic Cilengitide underwent clinical trials for cancer treatment. Unfortunately, Cilengitide did not meet the primary endpoints and therefore failed in randomised clinical trials (Demircioglu & Hodivala-Dilke, 2016). This present study suggests that specifically targeting αvβ3-TSP1 c-terminal fragment interactions may allow IA targeting. Another important finding of this study is the proposed use of serum cleaved TSP-1 as a biomarker of ulcerative colitis or related Crohn’s disease. TSP-1 was proposed as a biomarker in different pathologies such as pulmonary hypertension and other
cardiovascular diseases, and it would be interesting to investigate whether it could be further used as a biomarker in cancers that undergo IA and could benefit from anti-IA rather than anti-angiogenic therapies.

While the clinical and translational applications of this work remain to be established, the present results from Arroyo’s laboratory (summarised in Fig 1) provide exciting new opportunities for inflammatory disease control via the specific regulation of IA. All the players in this exquisitely balanced pathway of MT1-MMP-TSP1-αvβ3 have been therapeutic targets in the past, suggesting that drug repurposing may be of value in this context. But should we consider endothelial-specific MT1-MMP genetic editing as a real option for improved colitis control? Only the future will tell.

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