Vascular cell-matrix adhesion in development and cancer

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Title: Vascular cell-matrix adhesion in development and cancer

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Abstract:
The development and homeostasis of vertebrate organisms depend on the “tree of life”, that is the intricate network of vascular tubes composed by endothelial cells attached to the basement membrane and surrounded by perivascular cells. Although many studies have revealed the fundamental role of cytokines, growth factors and Notch signalling in vascular morphogenesis, we still lack sufficient understanding of the molecular mechanisms controlling the various steps of the angiogenic processes. Emerging data highlight that cell adhesions are key players in vascular morphogenesis. In this review, we focus on endothelial cells and we present the current state of knowledge regarding the role of cell-matrix adhesions in developmental and tumour angiogenesis, attained mainly from genetic studies and animal models.

Sort title: Endothelial adhesome in angiogenesis

Keywords: endothelial, angiogenesis, adhesion, migration, cancer

Abbreviations:
3D, three-dimension; E, embryonic day; EC, endothelial cell; ECM, extracellular matrix; FAK, focal adhesion kinase; HUVEC, human umbilical vein endothelial cell; ILK, integrin-linked kinase; PI3K, phosphoinositide-3-kinase; PINCH, particularly interesting new cysteine-histidine-rich protein; siRNA, small interference RNA; VEGF/VEGFR, vascular endothelial growth factor/ receptor
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Introduction

Blood vessels are crucial for the life of all vertebrates. Besides delivering nutrients and oxygen and removing waste metabolites, blood vessels provide instructive cues to the developing organs, control immune responses and form specific niches for stem cells, facilitating tissue homeostasis and regeneration. Therefore, it is not surprising that the vascular system is the first system to develop in the vertebrate embryo (Rissau, 1997). The vascular system consists of blood vessels - further divided into arteries, veins and capillaries - and lymphatics - arranged into ducts, large collecting vessels and smaller capillaries. The basic unit of the vascular system is the endothelium, a monolayer of endothelial cells (ECs) attached to each other and to the extracellular matrix (ECM), named basement membrane (Kostourou and Papalazarou, 2014). Depending on the vessel type, other perivascular cells (pericytes and smooth muscle cells) are firmly or more loosely attached to ECs (Potente et al., 2011). EC adhesion is key to blood vessel structure and thus critically influence vascular development, homeostasis and function. This review will focus on the significance of cell-matrix adhesion in regulating vascular morphogenesis and present recent advances and in vivo studies that reveal the key factors governing EC adhesion.

Fundamentals of vascular development

The vascular system forms in the embryo, initially by the process of vasculogenesis, namely the de novo coalescence of mesoderm-derived endothelial progenitors, termed angioblasts. In the yolk sac, angioblasts form the blood islands that give rise to the primary plexus and in the embryo proper, angioblasts arrange into primitive tubes that will differentiate into the dorsal aorta and the cardinal vein (Rissau, 1997). The intricate network of blood vessels then is produced by the growth and the remodelling of pre-existing vessels, a process known as angiogenesis. There are two main types of angiogenesis, the formation of new vascular branches through sprouting angiogenesis (Potente et al. 2011) and the splitting of vessels by intussusception (Makanya et al. 2009). In certain pathological cases, circulating precursor endothelial cells could be integrated into fast-growing vessels (Pearson, 2010). Also, tumours
exploit additional ways to fuel their exponential growth and obtain routes for metastasis, co-opting the host vasculature or forming tube networks by vascular mimicry (Lugano et al. 2019).

**Sprouting angiogenesis**

Very often in the literature the term “angiogenesis” is used to denote angiogenic sprouting, namely, the development of new branches from pre-existing vessels (Figure 1). This is a multistep process that starts with the activation of quiescent ECs caused by changes in tissue energy and oxygen demands. In pathological situations such as cancer, EC activation is triggered by a shift in the balance of pro-angiogenic and anti-angiogenic factors that govern vessel homeostasis. The activation of ECs results in extensive remodelling of extracellular matrix and EC differentiation into migrating tip cells and proliferating stalk cells (Potente et al. 2011). Tip cells extend actin-based protrusions called filopodia, to receive and translate the guidance cues of the microenvironment and migrate to form the new sprout. Stalk cells proliferate to provide additional ECs that will expand the sprout. Subsequently, ECs in the extended sprout form the vascular lumen. Different processes have been proposed for lumen formation, including cord hollowing (Iruela-Arispe and Beitel, 2013). During luminisation, ECs establish apical-basal polarity and separate apically at their contact surfaces by fusing intracellular vessels. In a similar process of cell hollowing, ECs generate vacuoles that fuse in the cytoplasm and create an intracellular lumen which expands inside the cell before fusing to the plasma membrane and connecting to a neighbouring intracellular lumen (Betz et al. 2016). Studies in zebrafish unravelled the generation of intracellular lumens by hemodynamic forces which create a plasma membrane invagination that extends towards the cytoplasm of EC. Cell rearrangements and lumen expansion connects adjacent lumens and create a continuous blood-filled channel (Betz et al. 2016). The new sprout eventually fuses with another new sprout (head-to-head sprout anastomosis) or an established vessel (sprout-to-side anastomosis) to enable vessel perfusion and participation in blood circulation (Potente et al. 2011).

**Intussusceptive angiogenesis**
The intussusceptive angiogenesis is less understood and the role of cell adhesion is still elusive because of the lack of specific in vivo and in vitro studying models (Nowak-Sliwinska et al. 2018) and the difficulty in visualising and detecting the process in tissues. So far, intussusceptive angiogenesis has been mainly studied in the chicken chorioallantoic membrane (CAM) and in vascular “corrosion” casts from the kidney (Makanya et al. 2009). Intussusception involves the formation of an intraluminal pilar composed of extracellular matrix components in the middle of a dilated vessel that is subsequently divided into two by the action of pericytes and matrix-metalloproteinases (Burri and Djonov, 2002). It is believed to be induced by increased hemodynamic flow and regulated by potent angiogenic factors, such as VEGF and hypoxia. The recent demonstration that intussusception follows angiogenic sprouting in vivo during the development of the zebrafish vasculature (Karthik et al. 2018), and the evidence for the initiation of vessel intussusception after anti-angiogenic treatment to sustain tumour growth are expected to spark new investigations aiming to clarify the unknown molecular mechanisms driving intussusceptive vascular growth. It would be interesting to delineate the contribution of vascular adhesion in this process.

**Vascular patterning**

Following the formation of new blood vessels, blood flow dynamics will fine-tune the newly formed vascular network by processes of pruning and stabilisation (Figure 1). Newly-formed blood vessels that exhibit very low or no blood flow will regress while others will get stabilised to create a functional network. Vascular remodelling is followed by blood vessel maturation and patterning which allows the hierarchical organisation into arteries, veins and capillaries (Fonseca et al. 2020). Deposition of ECM in the basement membrane and recruitment of mural cells (pericytes and smooth muscle cells) trigger the quiescence of ECs and stabilise the neovessel, safeguarding vessel integrity (Potente et al. 2011). Additionally, the expression of specific proteins by ECs generates organ-distinct vascular beds and confers arterial-venous identity.

**Tumour neovascularisation**
Since the initial studies of Judah Folkman and his team in 1971, it is now well-established that tumours are unable to grow and form metastasis without the formation of a vascular network (Folkman, 1971; Potente et al. 2011). Indeed, angiogenesis is a clinical marker for cancer stratification, intimately linked to tumour malignancy. Research advances have moved from a cancer cell-centric focus to the significance of the tumour microenvironment in malignancy and metastasis, opening new avenues for therapeutic interventions. It is well-appreciated the contribution of different types of tumour stromal cells in cancer progression, where the vascular system is at the central stage, regulating immune cell infiltration, tumour cell survival, proliferation and invasion (Nikolopoulou et al. 2021). Even at pre-malignant stages, an imbalance in growth-stimulating factors and suppressors triggers the “angiogenic switch” and initiate the involvement of the vascular system (Hanahan and Folkman, 1996). In the tumour microenvironment, hypoxia and pro-angiogenic factors secreted by cancer and stromal cells induce mainly sprouting angiogenesis (Potente et al. 2011). Intussusceptive angiogenesis has been reported in certain cancers and as an alternative way of vascular re-growth upon inhibition of angiogenic factors, such as vascular endothelial growth factor (VEGF) (Burri and Djonov, 2002).

**Alternative ways of tumour vascularisation**

Aggressive cancers, including melanoma, glioma and lung cancer are also using vascular mimicry to propel their growth. Tumour cells can organise themselves in matrix-rich channels that can be joined to tumour vascular circulation, mimicking blood vessels (Lugano et al. 2019). In a process related to vascular mimicry, tumour cells or cancer stem cells can express at their cell surface endothelial markers that facilitate their integration in the developing vascular network (Fernández-Cortés et al. 2019). Both vascular mimicry and the acquisition of an endothelial-like phenotype by cancer cells, hinder the efficacy of anti-angiogenic treatments.

Furthermore, both at initial tumour formation and later stages in organs rich in blood vessels such as lung and liver, cancer can exploit the normal tissue vascular plexus and progress by vessel co-option. Tumour cells expand in the vicinity of the host vasculature, alongside blood vessels that modify accordingly to support tumour
exponential growth and obtain routes for metastasis (Bridgeman et al. 2016; Lugano et al. 2019).

The outcome of the above processes is a highly tortuous, chaotic and disorganised tumour vasculature. The vessels themselves are highly leaky and the vascular patterning is severely disrupted, giving rise to a dysfunctional network that hinders drug delivery and facilitates metastasis.

**Molecular pathways in angiogenesis.**

Developmental angiogenesis and tumour neovascularisation have been the focus of extensive research for the past 50 years, which led to the identification of essential growth factors and critical molecular pathways (Potente et al. 2011; Fonseca et al. 2020). For example, the family of VEGF and their receptors (VEGFRs), as well as Notch-Delta signalling, along with other cytokines, angiopoietins, hypoxia-inducible factors (HIF) and WNT/b-catenin signalling have been shown to regulate many aspects of the angiogenic processes both in physiology and pathology (Potente et al. 2011; Lugano et al. 2019). Many intracellular signal transduction pathways, including phosphoinositide-3-kinase (PI3K), mitogen-activated protein kinases (MAPKs) and protein-tyrosine phosphatases (PTPs) are also critically involved in regulating vascular morphogenesis (Ramjaun and and Hodivala-Dilke 2009; Graupera and Potente 2013; Hale et al. 2017). Blood flow through the function of transcription factors such as KLF2 and the interplay between ECs and perivascular cells through placental derived growth factors (PDGFs) and their receptors (PDGFRs), Angiopoietins, Notch, and Ephrins/EphBs signalling participate in stabilising the neovessels and maintain vascular homeostasis (Betsholtz, 2018).

Although the biochemical signals that are involved in the formation and maintenance of the vascular system have been well-studied, the significance of cell adhesion is still relatively unexplored. Both cell-to-cell and cell-to-ECM adhesions need to get remodelled and re-organised to enable cell shape changes, migration, invasion, proliferation and lumen formation, all key angiogenic steps. Emerging data also
indicate that vascular patterning is adhesion-dependent and the subsequent stabilisation of the vascular network relies on direct interactions of ECs with the ECM and perivascular cells (Ramjaun and Hodivala-Dilke 2009; Fonseca et al. 2020). Additionally, it is becoming increasingly evident that besides chemical cues, the mechanical properties of tissues play a central role in organ physiology and pathology. Indeed, in tumours, it is well-established that the mechanical properties of the tumour microenvironment define both the malignancy and the therapeutic outcome (Mohammadi and Sahai, 2018). More specifically, the tumour vasculature contributes and is affected by cancer mechanics (Zanotelli and Reinhart-King, 2018). Cell-matrix adhesions pose as one of the main mechanical hubs, where the rigidity of the microenvironment and tensional forces are perceived and translated to biochemical signals governing cellular responses (Weinberg et al. 2017).

Since there are many recent excellent reviews on the role of cell-cell junctions in angiogenesis and blood vessel remodelling (Dorland and Huveneers 2016; Claesson-Welsh et al. 2020) here we discuss the current understanding of how endothelial cell-matrix adhesions control vascular development and tumour angiogenesis.

**Fundamentals of cell-matrix adhesion**

To delineate the involvement of cell-matrix adhesions in vessel morphogenesis, we will first provide a brief overview of the molecular composition and architecture of cellular adhesions (Figure 2). Cell-matrix adhesions are highly dynamic structures located basally at the specific sites where the cell interacts with the ECM. They can be classified into different types characterised by distinct morphology, localisation, lifespan and protein composition. Nascent adhesions are the smallest adhesive structures formed at the leading edges of membrane protrusions. They are very short-lived and either mature into focal complexes or dissolve. Focal complexes are also highly dynamic with a fast turnover, located mainly at the cell periphery. Focal adhesions are more complex and stable structures with a half-life of minutes and can form both at the periphery and central parts of the cell (Gardel et al. 2010).
Adhesome

All types of cell-matrix adhesions have a common structure, consisting of a network of scaffolding and signalling proteins, called “adhesome” which links integrin transmembrane receptors to cell cytoskeleton. The molecular composition and organisation of the adhesome differ in time and space, making it a highly dynamic network, adept to regulate all-important cellular functions, ranging from cell adhesion and migration to proliferation and differentiation (Zaidel-Bar and Geiger, 2010). Indeed, the adhesome physical anchor cells to the surrounding environment, sense and respond to mechanical cues and receive signals from growth factors and various molecular pathways, posing as a key integrating hub to orchestrate cellular responses. The adhesome protein network encompasses more than 232 members, including i) the family of integrins cell surface receptors, ii) structural/adaptor proteins: talin, vinculin, ILK-PINCH-parvin, paxillin, zyxin, tensin, KANK, etc), many of which have also a mechanosensing ability (such as talin, vinculin, zyxin, etc) iii) signalling proteins: kinases (such as FAK, Src family, PI3Ks, myosin light chain kinase), phosphatases (PTPs, PTEN, SHP2) and iv) cell cytoskeletal-regulating proteins: Rho GTPase family, and actin-associating proteins, α-actinin, filamin, Arp2/3 etc. These proteins are intrinsic components of cell-matrix adhesions or transiently associate with the adhesion network (Geiger and Yamada, 2011). Furthermore, adhesome proteins facilitate the cross-talk between actin structures (stress fibres, cortical actin, lamellipodia, filopodia) and microtubules, to stabilise cell shape and facilitate cellular homeostasis (Horton et al. 2015). Given the critical role of cell-matrix adhesions in regulating cell behaviour, current research focuses on elucidating the requirement of adhesome components in angiogenesis. Here, we discuss evidence that supports the important role of adhesome proteins in the regulation of developmental and tumour angiogenesis.

Cell-matrix adhesions and developmental angiogenesis

Distinct approaches, including global and EC-specific genetic deletions, various cellular assays and inhibition studies have revealed the critical involvement of adhesome components in all steps of blood vessel formation. Many angiogenic
studies have focused on the role of integrin cell-surface receptors that directly interact with various ECM components (Avraamides et al. 2008). Integrins are heterodimers composed of an α and a β subunit. From the 24 unique combinations of α and β pairs, EC expresses α1β1, α2β1, α3β1, α4β1, α5β1, α6β1, α9β1, ανβ3, ανβ5, that can bind, among other ECM components, collagens, laminins and fibronectins, found in the basement membrane and the provisional matrix of blood vessels (Hynes, 2002). Many integrins, such as α6β1, α5β1 and ανβ3 are upregulated in angiogenic endothelium, suggesting a major role in angiogenesis (Niland and Eble, 2012). Consistent with this, genetic deletion of integrin β1, the most common subunit of integrin dimers, specifically in ECs led to embryonic lethality at embryonic data (E)9.5-10.5 with severe vascular defects, indicating an important function of integrins in developmental angiogenesis (Carlson et al. 2008; Lei et al. 2008; Tanjore et al. 2008). Nevertheless, global genetic deletions of the majority of integrin α subunits, including αν, α1, α2, α4, α3, and α6, although affect the viability of the embryo because of defects in other organs, do not impede the formation of embryonic blood vessels (Niland and Eble, 2012). These findings suggest redundant and compensatory functions of different integrins during embryonic angiogenesis. A striking exception poses integrin α5. Mice with global genetic ablation of integrin α5 die at E10-11 with defective vascular development of the yolk sac and the embryo proper, indicating an essential function for this integrin in physiological angiogenesis (Yang et al. 1993; Francis et al. 2002). Other integrins, like integrin α9β1 are highly expressed in lymphatic endothelial cells and have a role in lymphangiogenesis without affecting blood vessel development (Huang et al. 2000; Bazigou et al. 2011).

Although it is difficult to disentangle the involvement of integrins in embryonic angiogenesis from their developmental role in other organs, there is genetic evidence for the requirement of integrin-associated proteins in physiological angiogenesis. For example, talin binds directly to integrins and is required for integrin activation and clustering (Critchley and Gingras, 2008). Endothelial deletion of talin1 results in early embryonic lethality due to vascular defects (Monkley et al. 2011). Similarly, integrin-linked kinase (ILK) has a major role in embryonic blood vessel development, through regulation of EC survival and co-ordination of angiogenic growth factor signalling
ILK is found in mammalian cells in complex with particularly interesting new cysteine-histidine-rich protein (PINCH) and parvin family proteins, α- and β- parvins. Whereas the vascular role of PINCH is unknown, endothelial deletion of both α- and β- parvins led to embryonic lethality at E11.5 with abnormal vascular network formation (Pitter et al. 2018).

Furthermore, a wealth of evidence supports a vital role for integrin-mediating signalling in developmental angiogenesis, with focal adhesion kinase (FAK) being a key player. Endothelial deletion of FAK impaired EC proliferation and migration causing severe vascular defects, haemorrhaging and embryonic lethality by E11.5 (Shen et al. 2005; Braren et al. 2006). Interestingly, the kinase function of FAK was required for EC migration and cell-cell junctional stability but not EC survival, since an endothelial-specific knock-in mouse expressing kinase-deficient FAK displayed no apoptotic defects. However, these mice also died at E13.5 due to defective angiogenesis and vascular barrier function (Zhao et al. 2010). Taken together these findings demonstrate kinase-dependent and -independent functions of FAK, highlighting the importance of thorough analysis of FAK function in the whole organism.

Besides structural and signalling adhesome components, the connection to the cell cytoskeleton is a key function of cell-matrix adhesions. Regulation of actin cytoskeleton by the Rho-GTPases family is essential for organising cell morphology and cell migration and thus, is involved in angiogenic processes. Although endothelial deletion of RhoA-GTPase exhibits normal blood vessel development, other members of the Rho-family affected embryonic angiogenesis (Zahra et al. 2019). Specifically, genetic studies demonstrated that cdc42 is required for EC survival and embryonic angiogenesis (Jin et al. 2013; Barry et al. 2015). Furthermore, early endothelial deletion of Rac-1 led to embryonic lethality by E9.5 and reduced vessel formation, whereas, loss of endothelial Rac at later stages of embryonic development caused defects in lymphatic-blood vessel separation and later embryonic lethality (Tan et al. 2008; D'Amico et al. 2009).

**Cell-matrix adhesions and EC migration**
Cell migration is a highly coordinated action of cell-matrix adhesion re-arrangements and cell cytoskeletal remodelling. Actin polymerisation drives membrane extensions in the form of filopodia and lamellipodia that are stabilised by the formation of nascent adhesions and focal complexes (Figure 2). Stable focal adhesions are re-organised or resolved and actomyosin contractility generates appropriate tension to facilitate the translocation of the cell body. During angiogenesis, ECs migrate to generate the new vessel. Furthermore, endothelial tip cells extend filopodia protrusions to explore the angiogenic microenvironment and migrate along with the stalk cells to form the new sprout. Therefore, it is evident that adhesome members will have a central role in this process. Indeed, reduction in cell migration along with decreased cell survival is the main reason for developmental angiogenic defects and embryonic lethality. Various experimental approaches, ranging from in vitro cell migration and invasion assays, ex vivo aortic ring assays to in vivo models of vessel formation in zebrafish and the postnatal development of retina vasculature have been exploited to shed light on the requirement of specific adhesome members for angiogenic sprouting.

Indeed, cell migration is impaired in mice lacking endothelial expression of integrin β1 and integrin α5 is upregulated in angiogenic endothelium and activates intracellular signalling pathways to enhance EC migration (Francis et al. 2002; Carlson et al. 2008). Similarly, ILK regulates EC spreading, migration and capillary formation in response to VEGF and FGF angiogenic factors (Tan et al. 2004; Vouret-Craviari et al. 2004; Malan et al. 2013). Loss of EC-ILK function in the developing retina severely impaired angiogenic sprouting, decreased vessel branching and compromised endothelial tip cell invasion and deeper vascular layer formation (Park et al. 2019). Defective lamellipodia formation, cell spreading and migration exhibit also ECs completely deficient in FAK expression, whereas increase migration is observed in FAK heterozygous EC which express half the amount of FAK (Braren et al. 2006; Tavora et al. 2010; Kostourou et al. 2013). Detailed examination of the developing retina vasculature in mice with inducible endothelial deletion of FAK postnatally, revealed decreased angiogenic sprouting without affecting endothelial tip cell specification (Tavora et al. 2010). Similarly, defective sprout formation and radial outgrowth of the retina vasculature was observed in mice lacking endothelial expression of other key
adhesome members including talin, vinculin, and parvin. Additional to the defective migration during the superficial retinal layer, ECs deficient for these proteins were unable to sprout perpendicular and invade into the deeper retina to form other capillary layers (Fraccaroli et al. 2015; Carvalho et al. 2019; Pulous et al. 2020). Remarkably, EC tip selection was unaffected and filopodia were present albeit in aberrant numbers, suggesting compromised cell-matrix adhesion re-organisation rather than defects in cytoskeletal remodelling. Changes in cell-matrix adhesions, however, are strictly controlled, as exemplified by studies of paxillin, an intrinsic adhesome component. Paxillin can regulate adhesion assembly and disassembly, along with Rho-GTPase dynamics (Nishiya et al. 2005). Depletion of endothelial paxillin, using siRNA, increased EC migration and invasion and caused excessive angiogenic sprouting in the developing retina vasculature by interfering with growth factor signalling (German et al. 2014).

Genetic studies of Rho-family GTPases, also have highlighted the central role of actin regulation in EC migration and vessel formation. Endothelial cdc42 deletion postnatally reduced EC filopodia formation and migration during retinal sprouting angiogenesis (Barry et al. 2015). Lack of lamellipodia formation and reduced EC motility in response to pro-migratory stimuli is also observed in Rac1 deficient ECs (Tan et al. 2008). In developing retina vasculature, endothelial deletion of Rac1 decreased vessel branching and EC invasion necessary to form the deeper plexus (Nohata et al. 2016). Also, deletion of the endothelial-specific RhoJ impaired EC migration and reduced vessel branching and sprouting angiogenesis in the retina, by interfering with α5 integrin trafficking, further supporting the functional link of cell-matrix adhesions with cell cytoskeleton (Fukushima et al. 2020; Sundararaman et al. 2020).

**Cell-matrix adhesions and lumen formation**

The formation of the endothelial lumen is essential for vascular morphogenesis and involves re-arrangement of stalk cells and changes in EC polarity, contractility and shape. The successful completion of these steps requires the remodelling of both cell adhesions and the actin cytoskeleton. Therefore, it is not surprising that integrins and
Rho-GTPases have been shown to regulate vessel lumenisation. Initial studies culturing human umbilical vein endothelial cells (HUVECs) in a 3D matrix revealed the formation of lumens from the coalescence of intracellular vacuoles. Using integrin inhibitors, it was shown that both fibronectin-binding integrins such as \(\alpha v \beta 3\) and \(\alpha 5\beta 1\) and collagen-binding integrins such as \(\alpha 2\beta 1\) participate in lumen formation (Davis and Camarillo 1996; Bayless et al. 2010). In the same 3D cellular system, the important role of Rac1 and cdc42 small GTPases in vacuole coalescence and lumen expansion was unravelled using dominant-negative forms of these proteins (Bayless and Davis, 2002). Similarly, knockdown of RhoJ decreased lumen formation by downregulating Rac1 and cdc42 activity and upregulated RhoA (Yuan et al. 2011).

Direct evidence, though, for the involvement of integrins and Rho-GTPases in vascular lumen formation was attained much later from in vivo studies. Endothelial-specific deletion of cdc42, and mosaic gene ablation studies in the developing retina, all confirm a central role of cdc42 in controlling vascular lumen formation by regulating apical-basal and axial polarity (Barry et al. 2015; Laviña et al. 2018). Regulation of small Rho GTPases, Rac1 and cdc42 by Ras interacting protein 1 (Rasip1) and Arhgap 29 were also required for lumenisation in the mouse embryo (Xu et al. 2011). Further studies demonstrated that Rasip1 regulates locally cdc42 and actomyosin contractility to reorganise adherens junctions and induce apical membrane separation and lumen formation (Barry et al. 2016).

Zebrafish has also proven to be a very useful model to study lumen formation in real-time. Visualising lumen formation in zebrafish intersomitic vessels, it became evident the contribution of blood flow in the process of cell and cord hollowing. Forces from blood flow deform the EC membrane to create inverse blebs that develop into transcellular invaginations. Later studies revealed that actin polymerising and bundling proteins such as formins and myristoylated alanine-rich C-kinase substrate like 1 (MARCKSL1) respectively, reorganise actin and regulate lumen formation and expansion (Phng et al. 2015; Kondrychyn et al. 2020).

Exploiting an inducible endothelial-specific system that deleted integrin \(\beta 1\) later in embryonic development, it was demonstrated that integrin \(\beta 1\) is critical for the apical-
basal polarisation of ECs via controlling Par-3 localisation (Zovein et al. 2010). The significant contribution of core adhesome members in lumen formation is poorly-defined. The major vascular defects caused by the elimination of the function of key adhesome members poses a significant challenge in ascertaining their contribution in EC migration and survival versus lumen formation. Further genetic tools and cellular assays will need to be developed to shed light on the molecular mechanisms governing vessel lumenisation and vascular morphogenesis.

**Cell-matrix adhesions and vascular patterning**

Following the formation of the initial network by sprouting anastomosis, angiogenic remodelling will define an optimised vascular plexus with a hierarchical organisation of arteries, veins and capillary. New vascular branches will be stabilised and others will regress. Hemodynamic forces generated by blood flow often determine which vessel connections are strengthened and which are removed. Superfluous vascular structures are pruned by a tightly regulated process that does not involve EC apoptosis but rather cell movement (Franco et al. 2015). It has been proposed that ECs polarise axially (front to rare) against the direction of blood flow and migrate away from regions with low or no flow. The mechanisms regulating these movements are not well-understood. The axial polarisation of ECs and the re-organisation of AJs to facilitate collective cell migration are thought to be involved (Fonseca et al. 2020). Several molecular pathways have been implicated, including non-canonical wnt signalling, sphingosine 1-phosphate S1P and its receptor S1PR1 along with signalling effectors such as PI3K, small RhoGTPases and actomyosin contractility (Fonseca et al. 2020).

Studies using mice with heterozygous integrin β1 endothelial deletion which display 40% of integrin β1 expression in ECs, have abnormal vascular remodelling responses to blood flow, suggesting a role of this integrin in vascular patterning (Lei et al. 2008). Reduced vascular density is also a common phenotype of many adhesome -deficient vasculatures, including integrin α5, ILK, talin, parvin (Francis et al. 2002; Monkley et al. 2011; Park et al. 2019; Pitter et al. 2018). Similarly, endothelial deletion of FAK or expression of FAK defective in the kinase activity or phosphorylation on tyrosine-397 (FAK-Y397F) exhibit reduced vessel density and abnormal vascular remodelling
(Corsi et al. 2009; Lim et al. 2010; Tavora et al. 2010). However, it is difficult to discern whether the reduced vessel density is caused by decreased development of vascular branches or increased regression of newly formed vessels. Interestingly, vinculin, an adhesome component that is localised both at cell-matrix and adherens junctions, is involved in vessel pruning. Mice with endothelial deletion of vinculin develop normally with no obvious embryonic vascular defects. Detailed examination of the retina vasculature in these mice showed reduced radial outgrowth and increased numbers of regressing vessels because of defects in collective EC polarity (Carvalho et al. 2019).

**Cell-matrix adhesions and vessel maturation**

Vessel regression is also associated with vessel instability due to defective vascular maturation. Recruitment and association of perivascular cells, namely pericytes and smooth muscle cells, and the establishment of a quiescent ECM that comprise the vascular basement membrane are both critical for the stability of blood vessels. RhoJ is involved in vessel maturation by regulating the deposition of fibronectin around new blood vessels (Sundararaman et al. 2020). Although integrins are involved in organising the ECM, the role of adhesome members in the remodelling of vascular basement membrane remains still elusive.

Haemorrhaging and oedema are common manifestations of defective sprouting angiogenesis caused by adhesome members, whether are associated with reduced pericyte coverage is not always explored. Therefore, little evidence exists about the role of endothelial cell-matrix adhesions in mediating EC-perivascular cell interactions during developmental angiogenesis. Endothelial expressing integrin α4β1 has been shown to mediate EC-perivascular cell interactions by binding to VCAM1-expressing smooth muscle cells (Garmy-Susini et al. 2005). Furthermore, besides the sprouting defects caused by endothelial deletion of α-parvin, newly-formed embryonic blood vessels displayed also impaired coverage by smooth muscle cells (Montanez et al. 2009). Similarly, heterozygous mice for kindlin2, an integrin-associated protein, exhibited thinner vascular basement membrane and reduced pericyte coverage, leading to vessel instability and leakage (Pluskota et al. 2011). Further studies are
required to decipher how adhesome components regulate cellular interaction during vascular morphogenesis.

**Cell-matrix adhesions and tumour angiogenesis**

Despite the pivotal role of cell-matrix adhesions in cancer cell motility and survival, the significance of endothelial adhesome in tumour neovascularisation remains largely unexplored (Nikolopoulou et al. 2021). Genetic deletion approaches and inhibitor studies have been used to delineate the contribution of integrins in tumour angiogenesis. In contrast to normal vascular development, mice deficient in many integrin subunits, including $\alpha_1$, $\alpha_2$, $\alpha_3$, $\alpha_6$, $\beta_4$, $\alpha_v\beta_3$ exhibit aberrant and opposing angiogenic responses in cancer. For example, endothelial deletion of integrin $\alpha_3$ did not impact embryonic vascularisation but increased tumour growth and angiogenesis in the adult mice by regulating VEGFR2 function (da Silva et al. 2010). Conversely, antagonists of integrin $\alpha_4\beta_1$ blocked tumour growth and angiogenesis (Garmy-Susini et al. 2005). Cellular studies using antibody inhibitors or small interfering RNA for integrin $\alpha_6$, showed impaired EC migration and tube formation, in vitro (Lee et al. 2006). Global genetic deletion of integrin $\beta_4$ subunit, that form a dimer with integrin $\alpha_6$ subunit, decreased postnatal angiogenesis and tumour neovascularisation (Nikolopoulos et al. 2004). Contrary to these studies, endothelial-specific ablation of integrin $\alpha_6$ increased tumour growth and angiogenesis, suggesting that integrin $\alpha_6$ is not a good target for anti-angiogenic therapies (Germain et al. 2010). Furthermore, some integrins although share the same ECM ligand, differentially regulate tumour angiogenesis, as exemplified by the collagen-binding integrins $\alpha_1\beta_1$ and $\alpha_2\beta_1$. Mice deficient in integrin $\alpha_1$ display reduced tumour angiogenesis, while mice deficient for integrin $\alpha_2$ exhibit increased tumour angiogenesis (Pozzi et al. 2000; Zhang et al. 2008). These contrasting effects could be explained by cross-regulation of integrin subunits, which has been described for many integrin pairs. Integrin $\alpha_v\beta_3$ is poorly expressed in quiescent ECs and is upregulated in tumour endothelium, making it an interesting therapeutic target. Indeed, pharmacological inhibition of integrin $\alpha_v\beta_3$ decreased tumour angiogenesis (Brooks et al. 1995). Similarly, mice expressing integrin $\beta_3$ deficient in the phosphorylation of its cytoplasmic tail exhibited impaired
tumour angiogenesis (Mahabeleshwar et al. 2006). In contrast, genetic deletion of integrin β3 increased tumour angiogenesis and growth (Reynolds et al. 2002). Accordingly, low concentrations of RGD-mimetics binding at the ectodomain of integrin αvβ3 increased tumour angiogenesis (Reynolds et al. 2009). Taken together these studies suggest that inhibition of integrin β3 is context-dependent and therefore targeting the ectodomain of integrin β3 may not be an effective therapeutic approach. Consistent with the embryonic studies, integrin α5β1 is upregulated in activated ECs during tumour angiogenesis and antagonists for α5β1 impaired tumour growth and neovascularisation (Kim et al. 2010). Furthermore, regulation of integrin activation by talin has been suggested to affect tumour development and angiogenesis (Pulous et al. 2020). The role of another regulator of integrin, kindlin2 in cancer was studied using heterozygous mice. Prostate tumours grew less efficiently and displayed reduced vessel density in these mice compared to control littermates (Pluskota et al. 2011).

Besides integrins, the contribution of FAK in tumour angiogenesis has been the focus of intensive research and inhibitors of FAK have participated in clinical trials. According to its important function in developmental angiogenesis, pharmacological inhibition of FAK or inducible endothelial deletion of FAK impeded tumour growth and angiogenesis by inhibiting EC migration and proliferation and enhancing EC apoptosis (Tavora et al. 2010; Nikolopoulou et al. 2021). Also, in a glioma tumour model, endothelial deletion of FAK, also, impaired tumour growth by decreasing tumour vessel diameter and increasing EC barrier function (Lee et al. 2010). FAK heterozygous mice, however, that express reduced levels of FAK, or suboptimal FAK inhibition increased tumour formation and angiogenesis, indicating a dose-dependent effect of FAK inhibition and calling for a thorough characterisation of FAK function before clinical application (Kostourou et al. 2013). Accordingly, several functional domains of FAK, including the FAK kinase domain, the autophosphorylation tyrosine site Y397 (blocking Y397F or mimic Y397E phosphorylation), and the Src-phosphorylation site Y861 (blocking Y861F) have been targeted and the corresponding EC-specific mutant mice have been used in tumour studies (Tavora et al. 2014). These studies revealed that both the FAK kinase activity and the autophosphorylation site promote tumour angiogenesis and vascular leakage,
facilitating cancer growth and metastasis (Alexopoulou et al. 2017; Pedrosa et al. 2019). The requirement of FAK-phosphorylation at tyrosine -861 is context-dependent, with early stages of tumour development being more susceptible than established tumours. Although these findings suggest that a delicate balance of FAK regulation is a prerequisite for anti-angiogenic therapy, recent studies have unravelled a paracrine role of FAK in sensitising tumours to DNA-damaging agents, opening new avenues for combinatorial anticancer treatment (Nikolopoulou et al. 2021).

Consistent with an essential function of ILK in developmental angiogenesis, pharmacological inhibition of ILK decreased tumour growth and vessel density by inducing apoptosis both in ECs and tumour cells (Nikolopoulou et al. 2021). Similarly, siRNA depletion of ILK in prostate cancer impaired tumour growth and angiogenesis by inhibiting hypoxia-driven VEGF expression (Tan et al. 2004). In a pancreatic cancer model, however, pharmacological inhibition of ILK impeded tumour development without affecting tumour angiogenesis (Yau et al. 2005). Genetic studies of endothelial-specific ILK deletion are necessary to ascertain cancer-intrinsic versus angiogenic effects of ILK in cancer development.

The impact of actin regulation in tumour angiogenesis has been explored in mice deficient in RhoJ. Both global and endothelial-specific deletion of RhoJ disrupted tumour blood vessel formation and integrity, resulting in impaired cancer growth. Similar defective tumour angiogenesis and growth was observed following siRNA-mediated depletion of Rac1 from the whole tumour environment (Nikolopoulou et al. 2021). However, targeting specifically Rac1 in tumour endothelial cells did not affect tumour growth and angiogenesis in wild-type mice but restricted the elevated tumour development and neovascularisation induced by integrin-β3 deficiency (D’Amico et al. 2010).

Concluding remarks

A wealth of studies has advanced our knowledge about the cellular events and the molecular pathways that regulate angiogenesis. While the contribution of growth factors and Notch signalling has been established, we still lack a good understanding of the key players and the signals controlling the different angiogenic steps during
development and in cancer. Open questions still remain regarding the molecular events regulating vascular lumen formation, remodelling and patterning. Also, it is becoming increasingly evident the involvement of mechanical forces derived from blood flow and the rigidity of the basement membrane in determining blood vessel formation and function. Cell-matrix adhesions, although understudied, have emerged as critical regulators of angiogenesis, as discussed in this review. Besides controlling cell migration and proliferation, we still lack sufficient understanding of the impact of adhesome members in other steps of the angiogenic process. Given that cell-matrix adhesions present the main hub for cell mechanotransduction, it is envisaged that adhesome members critically determine blood vessel responses to mechanical cues and control every angiogenic step. Most importantly, our understanding of blood vessel development in cancer is limited to sprouting angiogenesis. Next to nothing is known about the signals regulating intussusception, vessel co-option and mimicry and the role of endothelial cell-matrix adhesions have not yet been explored. The latter is particularly important for the efficient therapeutic manipulation of angiogenesis and the development of novel anti-cancer therapies.

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Figure legends

**Figure 1**: Schematic presentation of the cellular events during the multistep angiogenic process. ECs are activated by pro-angiogenic cues, and acquire a tip cell migratory or a stalk cell proliferating identity. ECM is remodelled and new sprouts invade the surrounding environment, form a lumen and connect with other sprouting vessels. The primary vascular plexus matures into a patterned network of arteries, veins and capillaries. Further stabilisation of blood vessels is achieved by association with pericytes and smooth muscle cells and establishment of the basement membrane.

**Figure 2**: Schematic organisation of blood vessels, cell-matrix adhesions and adhesome members with a known role in angiogenesis. Blood vessels consist of ECs forming a monolayer attached to the basement membrane and surrounding by perivascular cells. During cell migration, membrane protrusions are stabilised by nascent adhesions and focal complexes that further mature in stable focal adhesions. Adhesome is a dynamic network of cytoskeletal signalling and structural proteins that connect integrins to the cell cytoskeleton.
