Vinculin strengthens the endothelial barrier during vascular development

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Abstract
Remodelling of cell-cell junctions is crucial for proper tissue development and barrier function. The Cadherin-based adherens junctions anchor via β-Catenin and α-Catenin to the actomyosin cytoskeleton, together forming a junctional mechanotransduction complex. Tension-induced conformational changes in the mechanosensitive α-Catenin protein induce junctional Vinculin recruitment. In endothelial cells, Vinculin protects the remodelling VE-Cadherin junctions. In this study, we have addressed the role of Vinculin in endothelial barrier function in the developing vasculature. In vitro experiments, using endothelial cells in which α-Catenin was replaced by a Vinculin-binding deficient mutant, showed that junctional recruitment of Vinculin promotes endothelial barrier function. To assess the role of Vinculin within blood vessels in vivo, we next investigated barrier function in the vasculature of Vinculin knockout zebrafish. In the absence of Vinculin, sprouting angiogenesis and vessel perfusion still occurred. Intriguingly, the absence of Vinculin made the blood vessels more permeable for 10 kDa dextran molecules, but not for larger tracers. Taken together, our findings demonstrate that Vinculin strengthens the endothelial barrier and prevents vascular leakage in developing vessels.
Introduction

The semi-permeable vascular barrier between the blood and the surrounding tissue is maintained by a monolayer of endothelial cells (1). The endothelial barrier regulates the extravasation of leukocytes and fluid (2,3). Changes in permeability induced by angiogenic growth factors or inflammatory cytokines are often temporal and reversible, ensuring recovery of the vascular barrier (4). Chronic disruptions of the endothelial barrier, however, perturb vascular homeostasis and contribute to a multitude of pathologies, including atherosclerosis, cancer and inflammatory diseases (1,5). Hence, maintaining a tight, yet adaptable, endothelial barrier is important.

The endothelium in the developing vasculature is exposed to multiple forces that are derived from the mural cells, blood pressure and hemodynamic forces from the blood stream (6–9). Endothelial cells sense and transmit such mechanical cues via their cell-cell contacts, which evoke proportional cellular responses to maintain the endothelial barrier (10). VE-Cadherin-based adherens junctions (AJs) are crucial adhesion structures that form endothelial cell-cell contacts (11,12). Endothelial junction remodelling is required for collective endothelial migration during sprouting angiogenesis (13,14). The cytoplasmic domain of the transmembrane VE-Cadherin protein binds to β-Catenin and α-Catenin, which in turn couples to the actomyosin cytoskeleton and forms the core junctional mechanotransduction complex (15–18). The interaction of the VE-Cadherin – Catenin complex with the actin cytoskeleton stabilizes the AJs and maintains endothelial monolayer integrity (19–21).

Tension on the VE-Cadherin complex results in unfolding of α-Catenin, which enhances its Actin-binding affinity and exposes a cryptic binding site for Vinculin (22–28). The Vinculin – α-Catenin interaction drives α-Catenin-mediated mechanotransduction and preserves junctional integrity during force-dependent remodelling in cultured endothelial and epithelial cells (22,25,29–32). Vinculin recruitment to AJs occurs to different extents during agonist-induced endothelial barrier enhancing and disrupting processes (22,33,34). In zebrafish embryos, Vinculin associates with endothelial junctions that are remodelled by changes in blood flow (35) and endothelial expression of Vinculin is important for angiogenesis in the postnatal mouse retina (36). Whether junctional Vinculin has a role in endothelial barrier function remains unclear.

In this study, we found that junctional Vinculin recruitment facilitates endothelial barrier function in vitro. To define the importance of Vinculin for endothelial tissue integrity in vivo we examined the vasculature of vinculin-knockout zebrafish. We found that the endothelial cells still generated a functional and perfused vasculature in the absence of Vinculin. Interestingly, the developed blood vessels in Vinculin mutant zebrafish were more permeable for small dextran molecules, whereas large dextran molecules did not extravasate. Taken together, these results point to a role for Vinculin in strengthening of the endothelial barrier in the vasculature.
Results

**Junctional Vinculin strengthens the endothelial barrier in vitro.**

Vinculin is recruited to tensile AJs by α-Catenin. In addition, Vinculin localizes at Integrin-based focal adhesions (FAs) through its force-dependent interaction with Talin (37). To specifically investigate the role of junctional Vinculin, we used lentiviral shRNA transductions that deplete endogenous α-Catenin from human umbilical vein endothelial cells (HUVECs). Subsequently, we lentivirally expressed mouse α-Catenin-GFP, or α-Catenin-∆VBS-GFP, a modified α-Catenin protein in which the binding to Vinculin is prevented (22). Western blot analysis confirmed the depletion of endogenous α-Catenin and expression of the α-Catenin-GFP and α-Catenin-∆VBS-GFP in the rescued cells (**Fig. 1A**). Immunofluorescent (IF) stainings for VE-Cadherin were performed to assess AJs in the different experimental conditions. Knockdown of α-Catenin led to disassembly of AJs. The expression of α-Catenin-GFP and α-Catenin-∆VBS-GFP restored the AJs in shα-Catenin HUVECs (**Fig. 1B**), as shown previously (22). Next, we examined junctional Vinculin recruitment by performing IF analysis. Vinculin localized at both FAs and AJs in wild type α-Catenin rescued HUVECs (**Fig. 1C, D**). Analysis of α-Catenin-∆VBS-based junctions showed efficient preclusion of Vinculin from the tensile AJs, while Vinculin localization at the FAs was maintained (**Fig. 1C, D**). This confirms the junction-specific depletion of Vinculin in the α-Catenin-∆VBS expressing cells. To examine whether junctional Vinculin controls endothelial barrier function, we next performed Electric Cell-substrate Impedance Sensing (ECIS) as readout for the tightness of the endothelial cell monolayers. Silencing of endogenous α-Catenin expression led to a decrease in transendothelial resistance, indicating an impairment of endothelial barrier function (**Fig. 1E**). Restoring α-Catenin levels by α-Catenin-GFP re-established endothelial monolayer integrity, however α-Catenin-∆VBS-GFP expression, the mutated form lacking the Vinculin binding site, did not fully restore barrier function (**Fig. 1E, F**). To investigate whether the junctional depletion of Vinculin might affect barrier loss upon physiological remodelling conditions we performed ECIS experiments following treatment with the permeability factor Thrombin. These experiments showed that thrombin-induced barrier loss occurs similarly in α-Catenin-GFP or α-Catenin-∆VBS-GFP rescued endothelial monolayers (**Fig. 1G**). Taken together, these results indicate that junctional Vinculin promotes basal endothelial barrier function.

**Vinculin mediates vascular barrier function for small molecules.**

Since junctional Vinculin strengthens the barrier function of cultured endothelial monolayers, we next investigated the consequence of Vinculin ablation on the vascular barrier in vivo. We recently showed that Vinculin is important for the formation of junctional fingers in response to blood flow in the developing vessels of zebrafish (35). The zebrafish genome encodes two Vinculin isoforms, vinculin a (vcla) and vinculin b (vclb) (38,39). The genetic ablation of both Vinculin isoforms delays sprouting angiogenesis during early vascular development (35). Nevertheless, the vcl-KO embryos develop functional blood vessels and no evident haemorrhages were observed (35). To assess the permeability of the endothelial barrier, the 10 kDa rhodamine-dextran tracer was injected into the duct of Cuvier at 48 hpf of control or vcl-KO Tg(fli1a:EGFP)y1 embryos, in which the fli1a promoter drives the endothelial specific expression of EGFP. One hour after dextran-microinjections, we examined vascular perfusion and leakage by imaging the blood vessels of the zebrafish trunk, namely the intersegmental vessels (ISVs), the dorsal longitudinal anastomotic vessel (DLAV) and their perivascular areas. The ISVs of control embryos were perfused, as shown by the presence of the 10 kDa rhodamine-dextran tracer in the lumen of blood vessels. In control zebrafish, the tracer was maintained in the
blood vessel lumen of the DLAV and the ISVs and did not extravasate into the perivascular regions (Fig. 2A-C), which indicates that endothelial junctions are sufficiently tight to prevent leakage of small molecular such as 10 kDa rhodamine-dextran. Conversely, in vcl mutants we observed extensive leakage of the tracer dye into the surrounding tissues (Fig. 2A, B). In vcla-/-;vclb+/- and in particular in vcl full-KO embryos, the 10 kDa rhodamine-dextran intensities were lower within the perfused ISVs, suggesting that dextran molecules extravasated from the circulation (Fig. 2C). Our analysis shows that vcla-/-;vclb+/- and vcl full-KO embryos exhibited increased perivascular dextran levels (Fig. 2A, B). To examine the requirement of Vinculin for vascular barrier function of larger molecules, we next injected 70 kDa rhodamine-dextran in fli1:EGFP control and vcl KO embryos. An hour after dextran-microinjections, no differences in vascular permeability and perfusion between control and vcl heterozygous or homozygous KO embryos were observed (Fig. 3A-C). Taken together, these experiments demonstrate that Vinculin is required for the strengthening of the endothelial barrier in newly developed blood vessels to prevent leakage of small molecules.
Discussion

Endothelial barrier function is tightly regulated through force-dependent remodelling of cell-cell contacts. Failure of the endothelial junctions to adapt to subjected forces, leads to vascular leakage and inflammation in cardiovascular disease (1). In this study, we examined the importance of the mechanotransduction protein Vinculin for the endothelial barrier using both in vitro and in vivo functional approaches. These results reveal that recruitment of Vinculin to AJs strengthens the endothelial cell-cell junctions in blood vessels.

Vinculin knockout mice are embryonically lethal due to neuronal and cardiovascular defects at E10.5 (40), demonstrating the importance of Vinculin for mammalian development. In addition, endothelial-specific Vinculin depletion constrains collective endothelial migration during retinal angiogenesis in mice (36), indicating that endothelial Vinculin contributes to vascular development. Endothelial Vinculin is recruited to both Integrin-based FAs and Cadherin-based AJs in a force-dependent manner (30,37). When tensile forces remodel AJs, Vinculin-mediated mechanotransduction occurs via the VE-Cadherin complex (22). By generating endothelial cells that form junctions through the Vinculin-binding deficient α-Catenin mutant (α-Catenin-ΔVBS), we now specifically showed that junctional Vinculin recruitment supports strengthening of monolayer integrity in cultured endothelial cells.

Single vcla or vclb mutant zebrafish display mild developmental defects (39,41). We find that the vcla-/-;vclb/- double knockout zebrafish exhibit mild defects in vascular morphogenesis and eventually the morphogenetic process gives rise to a functional vasculature (35). In line with the expectation that Vinculin modulates, rather than being needed for, endothelial cell-cell junctions (22,28,29), we observed vascular leakage specifically for small molecules, 10 kDa, in Vinculin knockout zebrafish, whereas the vasculature still acted as a barrier for larger molecules. This result is corroborated by the notion that recruitment of Vinculin to VE-Cadherin-based junctions does not affect histamine-induced vascular leakage of Evans Blue in mice, which is a measure for the extravasation of large molecules (28). Even though the measured resistance formed by cultured endothelial cells expressing α-Catenin-ΔVBS indicated minor barrier differences upon the junctional depletion of Vinculin, the knockout of Vinculin in vivo resulted in significant leakage of small molecules. This indicates that the endothelial dysfunction upon Vinculin depletion become aggravated in pressurised vascular conditions. Potentially, the mild phenotype of the Vinculin knockout zebrafish might be aggravated under pathological conditions that weaken the endothelial barrier, such as during inflammation or sepsis. Together, the data show that Vinculin tightens endothelial junctions in blood vessels.

We expect that the depletion of the junctional Vinculin pool underlies the vascular phenotype of the Vinculin KO zebrafish. Cultured endothelial cells that form junctions through α-Catenin-ΔVBS fail to sprout in collagen gels (data not shown). This suggests that the junctional pool of Vinculin controls endothelial dynamics within angiogenic sprouts, an effect that has also been observed in endothelial-specific knockout mice (36). Moreover, we recently observed differences in flow-induced endothelial junction dynamics in the ISVs of control and Vinculin KO zebrafish, whereas integrin-dependent filopodia still formed equally (35). Other groups showed that juvenile Vclb mutant zebrafish display epicardial defects and pericardial edema (39,41,42). Recruitment of vinculin was observed during the maturation of cell-cell junctions between cardiomyocytes in vivo (42). These findings suggest that in vivo, vinculin’s role at cell-cell junctions is prominent. Nevertheless, a potential contribution of Vinculin’s role at Integrin-based adhesion cannot be fully ruled out in this model system.
Finally, we surmise that junctional Vinculin recruitment fortifies the endothelial barrier for small molecules upon vascular remodelling. Future work targeting the protective function of Vinculin using pharmacological approaches, for instance by enhancing its interaction with the junctions, may provide a strategy to treat pathologies that entail vascular permeability.
Materials & Methods

Antibodies & reagents

Rabbit polyclonal anti-VE-Cadherin (Cat# 36-1900, diluted 1:200 for immunofluorescence (IF)) was from Thermo Fischer Scientific. Purified mouse anti-Vinculin (clone hVIN-1, Cat # V9131, diluted 1:400 for IF) was from Sigma Aldrich. Rabbit polyclonal anti-β-actin (Cat# 4867S, diluted 1:1000 for Western blot (WB)) and rabbit polyclonal anti-phospho-paxillin-Tyr118 (Cat# 69363, diluted 1:200 for IF) were from Cell Signaling. Mouse monoclonal anti-α-Catenin (Cat# 13-9700; diluted 1:1000 for WB) was from Invitrogen/Zymed and mouse monoclonal anti-GFP (B-2, Cat# sc-9996, diluted 1:1000 for WB) was from Santa Cruz Biotechnology. Promofluor 415 Phalloidin (Promokine, Cat# PK-PF415-7-01, diluted 1:200 for IF) was used for IF of F-actin. Alexa Fluor 488 or 594-coupled secondary antibodies were from Invitrogen (diluted 1:250 for IF). Secondary antibodies coupled to horseradish peroxidase (HRP) were form Bio-Rad (diluted 1:1000 for WB). Human plasma-derived thrombin (used at 1 U/ml) was purchased from Sigma-Aldrich.

Cell culture

Pooled primary Human Umbilical Vein Endothelial Cells (HUVECs) from different donors (Lonza) were cultured in Endothelial Cell Growth Medium 2 (EGM-2) culture medium supplemented with the Growth Medium 2 Supplement Pack (PromoCell) on gelatin coated tissue flasks. HEK293T cells (ATCC) were cultured in Dulbecco’s modified Eagle’s medium (DMEM) with L-glutamine and supplemented with 10% FCS and 1% pen/strep. Cells were recently authenticated and tested for contamination.

DNA plasmids & lentivirus production

To silence α-Catenin expression in HUVECs, pLKO.1-shRNA plasmid targeting human α-Catenin mRNA was used (TRCN0000062653). ShC002 was used as shRNA control (Sigma-Aldrich mission library). The mouse α-Catenin-GFP and α-Catenin-ΔVBS-GFP lentiviral plasmids were previously described (22). Lentivirus was generated by transfecting HEK293T cells with the lentiviral expression plasmids and 3rd generation packaging plasmids using Trans-IT-LTI transfection reagents (Mirus) as described previously (43). To transduce HUVECs, the supernatant containing the lentiviral particles were mixed at a 5:1 ratio with EGM-2 and incubated with the HUVECs for 16 hours. Subsequently, transduced HUVECs were selected for expression of the shRNA with 2.5 µg ml⁻¹ puromycin (Sigma). shRNA-based knockdown levels were analysed at least 72 hours after transduction.

ECIS

To measure endothelial barrier resistance, we used electric-cell-impedance-sensing as previously described (44). Gold electrode arrays (8W10E, Applied Biophysics) were treated with 10 mM L-cysteine (Sigma) for 15 minutes at room temperature. After washing with MQ water, the wells were coated with 5 µg ml⁻¹ fibronectin in MQ for 1 hour at 37°C and 5% CO₂. Subsequently, 120.000 cells per well were seeded on the arrays and the impedance was measured during monolayer formation at 4000 Hz using the ECIS model ZTheta (Applied BioPhysics).

Immunofluorescence stainings

For immunofluorescence stainings, HUVECs were cultured on 5 µg ml⁻¹ fibronectin coated coverslips and later fixed for 15 minutes at room temperature with 4% paraformaldehyde in PBS⁺⁺ (PBS supplemented with 1 mM CaCl₂ and 0.5 mM MgCl₂). The fixed cells were permeabilized for 5 minutes at room temperature with 0.5 % Triton X-100 in PBS, and blocked for 15 minutes in 2% bovine serum albumin (BSA) in PBS. Primary and secondary antibodies were diluted in 0.5% BSA in PBS and incubated
for 45 minutes. Between incubations, fixed cells were washed three times with 0.5% BSA in PBS. Coverslips were mounted in Mowiol4-88/DABCO solution (Sigma).

Immunoblot analysis

HUVECs were lysed using reduced sample buffer containing 4% β-mercaptoethanol. Samples were denatured at 95°C for 5 minutes and subsequently loaded on a 10% SDS page gel. Gel running was performed in SDS-page running buffer (25 mM Tris-HCl pH 8.3, 192 mM glycine and 0.1% SDS) and blotted on ethanol activated PVDF membranes using full-wet transfer blot buffer (25 mM Tris-HCl pH 8.3, 192 mM glycine and 20% (v/v) ethanol). Blots were blocked in 5% milk powder in Tris-buffered Saline (TBS) for 30 minutes and subsequently incubated with the primary antibodies in 5% milk powder in TBS supplemented with Tween-20 (TBS-t) overnight at 4°C. The secondary antibodies, coupled to HRP, were incubated for 45 minutes at room temperature. Between antibody incubations, blots were washed 3x with TBS-t. As a final step before visualization, blots were washed 1x with TBS. HRP signal was visualized using enhanced chemiluminescence (ECL) detection (Supersignal West Pico PLUS, ThermoFischer) with an ImageQuant LAS 4000 (GE Healthcare).

Zebrafish lines and maintenance

Zebrafish were maintained in standard housing conditions according to FELASA guidelines (45). All experiments were performed in accordance with federal guidelines and were approved by the Kantonales Veterinäramt of Kanton Basel-Stadt (1027H, 1014HE2, 1014G). The vcla\textsubscript{hu10818}; vclb\textsubscript{hu11202} zebrafish lines (39) were crossed into the transgenic Tg(fli1a:EGFP)\textsuperscript{y1} line, which labels all endothelial cells (46) as described in (35).

Genotyping of the Vinculin (vcl) mutant lines

For genotyping of the Vinculin mutant alleles, genomic DNA was extracted from adult fish fin biopsies or from whole embryos using a standard protocol (47) with addition of proteinase K to the sample. The extracted genomic DNA was then used to genotype the vcla and vclb loci. Genotyping protocol for vcla or vclb alleles was performed as described previously (35).

Microangiography

48 hpf zebrafish embryos were anaesthetized with 1x tricaine (0.08%, Sigma) and were injected with 250 µg/ml 10 kDa or 70 kDa rhodamine-dextran (Molecular Probes) in the duct of Cuvier using glass needles (Biomedical Instruments) and standard microinjection protocols (48,49). The injected embryos were transferred back to embryo media (5 mM NaCl, 0.17 mM KCl, 0.33 mM CaCl\textsubscript{2}, 0.33 mM MgSO\textsubscript{4}, pH 7.4) to recover and subsequently imaged 1 hour after microinjections. For visualization of vascular leakage, embryos were mounted in 0.7% low-melting-point agarose (Sigma) and imaged with the Zeiss Axioplan Airy (25X oil/0.8 NA objective, confocal mode).

Imaging and image analysis

Fixed HUVECs were imaged using widefield microscopy on a NIKON Eclipse TI, with a SOLA SE II light source, 60X 1.49 NA Apo TIRF (oil) objective and Andor Zyla 4.2 plus sCMOS camera and standard CFP, GFP or mCherry filter cubes (NIKON). For live-imaging of zebrafish, a Zeiss LSM880 Airyscan inverted confocal microscope, with a 25x 0.8 NA oil objective was used. First, live embryos were selected for fluorescence signal, and subsequently anaesthetized with 1x tricaine (0.08%) in E3 fish water and mounted in glass bottom Petri dishes (MatTek) using 0.7% low-melting-point agarose (Sigma) containing 1x tricaine. For live-imaging, E3 with 1x tricaine and 0.003% 1-phenyl-2-thiourea (PTU, Sigma) was added to avoid pigmentation. Images were acquired with a zoom of 1-1.6 and z-stack step.
size of 0.5 to 1.0 µm, with a time interval of 25 to 30 minutes. Vascular perfusion and vascular leakage were analysed based on the fluorescent dextran levels in zebrafish embryos at 48hpf. Vascular perfusion was defined as the ratio of fluorescent dextran levels in the ISVs to the levels of dextran inside the dorsal aorta (DA). Vascular leakage was defined as the ratio of the dextran fluorescent levels at the perivascular area of the vessels to the dextran levels inside the vessels. For this analysis, the dorsal longitudinal anastomotic vessel (DLAV) and its perivascular areas were analysed.

Statistical analysis

Graphpad Prism was used for the statistical analysis of the data. All violin plots represent data distribution, with the dashed line representing the quartiles and the straight line representing the median. When 2 groups were compared, a Wilcoxon test was used. When 2 or more groups were compared to the control, a One-way analysis of variance (ANOVA) was used, in combination with a Tukey’s or Dunnett’s post-hoc test for multiple comparisons and a D'Agostino-Pearson test for normality. Asterisks indicate p values, and are defined as n.s. Non-significant, * p<0.05, ** p <0.01, *** p<0.001.
Declaration of interest
The authors declare that there is no conflict of interest.

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Author contributions
M.v.d.S., M.K. and R.S. designed and performed experiments. S.H. and H.G.B. conceived and supervised the study. M.v.d.S., M.K., R.S., H.G.B. and S.H. analyzed the data. M.v.d.S., M.K., H.G.B. and S.H. wrote the manuscript. All authors reviewed the manuscript.

Data availability
Data are available from the authors upon reasonable request.
Figure Legends

Figure 1. Junctional Vinculin strengthens the endothelial barrier in vitro.

(A) Representative Western blot analysis of shControl and shα-Catenin transduced HUVECs rescued with lentiviral expression of α-Catenin-GFP or α-Catenin-ΔVBS-GFP. Blotted for α-Catenin, GFP or β-actin. Bar graphs indicate the average±S.E. expression levels of α-catenin-GFP and α-catenin-ΔVBS-GFP in shα-Catenin transduced HUVECs relative to endogenous α-catenin levels in shControl HUVECs. n.s. non-significant (non-parametric Wilcoxon matched-pairs signed rank test). Data is from 3 independent experiments. (B) Representative immunofluorescent images of shControl and shα-Catenin transduced HUVECs rescued with lentiviral expression of α-Catenin-GFP or α-Catenin-ΔVBS-GFP. Stained for F-actin (grey) and VE-Cadherin (magenta). (C) Representative IF images of shα-Catenin HUVECs rescued with α-Catenin-GFP or α-Catenin-ΔVBS-GFP (green) that were stained for Vinculin (magenta) and F-actin (grey). Colocalization of Vinculin with α-catenin–GFP or α-catenin–ΔVBS–GFP was analyzed by line scans displaying signal intensity (arbitrary units) across the AJs and FAs as indicated. Scale bar, 10 µm. (D) Representative IF images of shα-Catenin HUVECs rescued with α-Catenin-GFP or α-Catenin-ΔVBS-GFP (green) that were stained for Vinculin (magenta) and phosphor-Paxillin Tyr118 (grey). Colocalization of Vinculin with pPaxillin was analyzed by line scans displaying signal intensity (arbitrary units) across FAs as indicated. Scale bar, 10 µm. (E) Line graph showing the average resistance±S.E. measured with ECIS at 4000 Hz of indicated endothelial monolayers over time. Data is from 3 independent experiments. (F) Bar graphs representing the average resistance±S.E. measured with ECIS at 4000 Hz of indicated endothelial monolayers after 24 hours. Data is from 3 independent experiments. n.s. non-significant, *p<0.05, **p<0.01, ***p<0.001 (One-Way ANOVA with Tukey’s post-hoc test for multiple comparisons). (G) Line graph showing the average resistance±S.D. measured with ECIS at 4000 Hz of indicated endothelial monolayers over time following treatment with the permeability factor Thrombin. Data is normalized to the baseline values prior to Thrombin treatment and is derived from 2 independent experiments.

Figure 2. Vinculin ensures vascular barrier function for 10 kDa dextran.

(A) Images of ISVs from 48 hpf Tg(fli1:EGFP) vcla+/+;vclb+/+, vcla-/+;vclb+/+ or vcla-/+;vclb-- embryos injected with 10 kDa rhodamine-dextran (red). Lower panels are single channel images of the rhodamine signal. Scale bars, 50 µm. (A i-iii) Corresponding close-up images showing the dextran leakage in the perivascular area around the dorsal longitudinal anastomotic vessel (DLAV). (A j-jjj) Corresponding close-up images showing the dextran leakage in the perivascular area around an intersegmental vessel (ISV). Scale bars, 10 µm. Scale bars, 10 µm. (A i-III) Close-up images of an ISV and the dorsal aorta (DA) showing the rhodamine signal within these vessels. Scale bar, 20 µm. (B) Violin plots showing the average leakage±S.E. of 10 kDa dextran into the perivascular area of the DLAV or ISVs normalized to the dextran inside the DLAV or ISVs from 48 hpf embryos. The dotted lines represent the quartiles and the straight lines represent the median, n= 10 vcla+/+;vclb+/+, n=25 vcla+/--;vclb-- and n=13 vcla+/+;vclb-- embryos, n.s. non-significant, *p<0.05, **p<0.01, ***p<0.001 (One-way ANOVA and Dunnett’s post-test). PCV: posterior cardinal vein. (C) Violin plot showing the ISV perfusion determined as the ratio between fluorescent dextran levels inside the ISV±S.E. and dextran inside the dorsal aorta (DA) from 48 hpf embryos. The dotted lines represent the quartiles, the straight lines represent the median. n=10 vcla+/+;vclb+/+, n=25 vcla+/--;vclb-- and n=13 vcla+/+;vclb-- embryos.
Figure 3. Loss of Vinculin does not impair vascular barrier function for 70 kDa dextran.

(A) Images of ISVs from 48 hpf Tg(fli1:EGFP) vcla+/+;vclb+/+, vcla-/--;vclb+/-- or vcla-/--;vclb-/- embryos injected with 70 kDa rhodamine-dextran (red). Lower panels are single channel images of the rhodamine signal. Scale bars, 50 µm. (A i-iii) Corresponding close-up images showing the dextran levels in the perivascular area around the dorsal longitudinal anastomotic vessel (DLAV).

(A j-jjj) Corresponding close-up images showing the dextran levels in the perivascular area around an intersegmental vessel (ISV). Scale bars, 10 µm. (A I-III) Close-up images of an ISV and the dorsal aorta (DA) showing the rhodamine signal within these vessels. Scale bar, 20um. (B) Violin plot showing the average leakage±S.E. of 70 kDa dextran into the perivascular area of the DLAV or ISVs normalized to the dextran inside the DLAV or ISVs from 48 hpf embryos. The dotted lines represent the quartiles, the straight lines represent the median, n=7 vcla+/+;vclb+/+, n=16 vcla-/--;vclb+/-- and n=7 vcla-/--;vclb-/- embryos, n.s. non-significant, One-way ANOVA and Dunnett’s post-test. (C) Violin plot showing the ISV perfusion determined as the ratio between fluorescent dextran levels inside the ISV±S.E. and dextran inside the dorsal aorta (DA) from 48 hpf embryos. The dotted lines represent the quartiles, the straight lines represent the median, n=10 vcla+/+;vclb+/+, n=16 vcla-/--;vclb+/-- and n=8 vcla-/--;vclb-/- embryos.


16. Pannekoek W-J, de Rooij J, Gloerich M. Force transduction by cadherin adhesions in...


Figure 1.

A

Relative expression levels

kDa
α-catenin-GFP
α-catenin
GFP
β-actin
sha-catenin

+ + + +
A

Endothelial barrier (24h)

B

shControl

α-catenin

α-catenin-∆VBS

GFP

Vinculin

GFP

pPaxillin

GFP

Vinculin

pPaxillin

C

D

E

F

G

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Figure 2. Analysis of vascular endothelial barrier function in different genotypes.

A) 10 kDa Dextran filling with EGFP.

B) Leakage (10 kDa Dextran).

C) Perfusion (10 kDa Dextran).