Transient loss of venous integrity during developmental vascular remodeling leads to red blood cell extravasation and clearance by lymphatic vessels

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Summary statement

This study uncovers previously unrecognized roles of platelets and lymphatic vessels during developmental vascular remodeling and provides mechanistic insight into how vascular abnormality characterized by blood-filled lymphatic vessels arises.

Abstract

Maintenance of blood vessel integrity is critical for vascular homeostasis and mainly controlled at the level of endothelial cell (EC) junctions. Regulation of endothelial integrity has largely been investigated in the mature quiescent vasculature. Less is known how integrity is maintained during vascular growth and remodeling involving extensive junctional reorganization. Here we show that embryonic mesenteric blood vascular remodeling is associated with a transient loss of venous integrity and concomitant extravasation of red blood cells (RBC), followed by their clearance by the developing lymphatic vessels. In wild type mouse embryos, we observed activated platelets extending filopodia at sites of inter-EC gaps. In contrast, embryos lacking the activatory C-type lectin-like receptor-2 (CLEC2) showed extravascular platelets and excessive number of RBCs associated with and engulfed by the first lymphatic EC clusters that subsequently form lumenized blood-filled vessels connecting to the lymphatic system. These results uncover novel functions of platelets in maintaining venous integrity and lymphatic vessels in clearing extravascular RBCs during developmental remodeling of the mesenteric vasculature. They further provide insight into how vascular abnormality characterized by blood-filled lymphatic vessels arises.
Introduction

Maintenance of blood vessel integrity is critical to vascular homeostasis, and its disruption can lead to hemorrhage, edema, inflammation and tissue ischemia (Murakami and Simons, 2009). Endothelial integrity and barrier properties are mainly controlled at the level of endothelial cell-cell junctions. Associated mural cells and the basement membrane (BM) stabilize the vessel wall and form an additional barrier preventing leakage. Disruption of cell-cell junctions may be accompanied by exposure of the thrombogenic extracellular matrix of the vessel wall, which triggers thrombus formation. Thereby, platelets also play an important role in vascular homeostasis by sealing gaps in the injured endothelium. They can additionally promote barrier function in resting endothelium by releasing a variety of soluble factors (Ho-Tin-Noé et al., 2011).

Most studies have focused on the role of platelets in the maintenance of the mature vasculature, but their role in promoting neo-angiogenesis and maintaining integrity of angiogenic vessels in adult tissues has also been reported (Ho-Tin-Noé et al., 2011). Platelets are instead thought to be dispensable for embryonic vessel integrity with the exception of the cerebral vasculature, where platelet C-type lectin-like receptor-2 (CLEC2, encoded by Clec1b) interaction with podoplanin (PDPN) on the neuroepithelium is required for platelet adhesion, aggregation and secretion (Lowe et al., 2015). While CLEC2 appears to have a minimal role in physiological hemostasis (Bender et al., 2013), it is important for platelet function in preventing inflammation induced hemorrhaging (Boulaftali et al., 2013) and mediating initiation of deep vein thrombus formation (Payne et al., 2017). In addition, platelet dysfunction caused by Clec1b deficiency in mice has been associated with abnormal blood filling of lymphatic vessels (Welsh et al., 2016). However, this phenotype was shown to be caused by back filling of lymphatic vessels with blood due to defective platelet aggregation and thrombus formation at the lymphovenous junction (Hess et al., 2014), rather than defects in the blood vasculature.

Here we studied the regulation of vessel integrity during embryonic vascular morphogenesis. We found that remodeling of the mesenteric blood vasculature is associated with a transient disruption of venous endothelial integrity. We further identify previously unrecognized roles of platelets and lymphatic vessels during developmental vessel remodeling in maintaining endothelial integrity and clearing of red blood cells (RBC), respectively.
Results and discussion

Transient extravasation of RBCs and their engulfment by lymphatic vessels during mesenteric vascular development

Most lymphatic vessels in mammals have been described to form through lymphangiogenic sprouting from embryonic veins (Srinivasan et al., 2007). In contrast, lymphatic vessels in the mesentery form through lymphvasculogenic assembly of lymphatic EC (LEC) progenitors into clusters that further coalescence to lumenized vessels between embryonic day (E)13 and E14 (Stanczuk et al., 2015). Immunofluorescence combined with differential interference contrast (DIC) imaging of E14 mesenteries from wild type embryos unexpectedly showed the presence of cells characteristic of RBCs inside the developing lymphatic vessels (Fig. 1A). TER-119 staining confirmed RBC identity and revealed both nucleated (TER-119\textsuperscript{low}) and enucleated (TER-119\textsuperscript{high}) RBCs inside lymphatic vessels (Fig. 1B).

To investigate the significance of this phenomenon, we first determined the frequency of extravascular RBCs in the developing mesentery based on whole-mount immunofluorescence for markers of RBCs as well as blood and lymphatic ECs (Fig. 1C, Movie S1, Fig. S1A). Rare extravascular RBCs were observed at E13-E13.5 (0.55 ± 0.57% (n=5) and 2.05 ± 3.07% (n=23) of total RBCs, respectively; Fig. 1D), but the frequency increased to 7.38 ± 7.33% (n=39) at E14 (Fig. 1C,D). Notably, the majority of extravascular RBCs interacted with or were captured within LEC membrane protrusions, or engulfed by LEC clusters at E14 (Fig. 1C,E, Movie S1). At E15, RBCs were frequently observed in the lumen of the developing lymphatic vessels (Fig. 1C), but were no longer present after E16 (Fig. 1D; 1.26% ± 2.00% extravascular RBCs (n=37)), once lymphatic drainage is initiated (Sabine et al., 2012).

To better understand the mechanisms leading to RBC interaction with and engulfment by LECs, we analyzed mesenteries at E13-E13.5 when LEC clusters first appear. Consistent with the low number of extravascular RBCs, most LEC clusters did not interact with RBCs at this stage (Fig. S1B), suggesting that LEC emergence and cluster formation is not controlled by extravascular RBCs. To further investigate LEC-RBC interactions in high resolution, we mosaically labelled individual LECs by Cre-activated expression of a membrane bound GFP in \textit{R26-mTmG;Vegfr3-CreER\textsuperscript{T2}} embryos (Martinez-Corral et al., 2016) using a suboptimal dose of 4-hydroxytamoxifen. Autofluorescence signal in combination with PROX1 staining allowed visualization of RBCs and LEC nuclei, respectively. Deconvolved confocal images showed close association of RBCs with mesenteric LECs at E13.5 (Fig. 1F, Movie S2).
Imaging of whole mount mesenteries with structured illumination microscopy (SIM) and 3D reconstruction confirmed direct contacts between LEC protrusions and RBCs (Fig. 1G, Movie S3). Despite the observed interaction of RBCs with LEC clusters at E14 (Fig. 1E), LECs were frequently found to initially extend protrusions in a random manner (Fig. S1C), arguing against a specific RBC-derived chemoattractant in driving the formation of LEC protrusions.

Taken together, these results indicate transient RBC extravasation during the development of the mesenteric vasculature. Selective association of RBCs with LECs further suggests a role for developing lymphatic vessels in the capture and clearance of extravasated RBCs.

**Remodeling of the developing mesenteric blood vasculature is associated with a transient loss of endothelial integrity**

In the tumor vasculature, extravasation of RBCs leading to hemorrhage is associated with disruption of EC integrity (Hashizume et al., 2000). To analyze the integrity of the developing mesenteric blood vessels we first visualized the morphological changes in the vasculature at the critical stages of development using the *Cldn5-GFP* reporter that allows visualization of all endothelial cells (ECs) by strong GFP fluorescence (Stanczuk et al., 2015). As previously reported (Hatch and Mukouyama, 2015), at embryonic day (E)13 mesenteric blood vessels form a primary plexus that remodels by E13.5 into a segmentally organized pattern of veins and arteries running in parallel (Fig. 2A). Arterial-venous identity was established prior to remodeling, as indicated by expression of the venous EC marker Nrp2 in only a subset of vessels within the primitive plexus (Fig. 2A). At E14, arteries had a continuous basement membrane (BM) and extensive mural cell coverage (Fig. 2B), typical of mature vessels. In contrast, E14 mesenteric veins had only a fragmented BM and few mural cells (Fig. 2B).

ECs of E14 arteries were aligned along the longitudinal axis of the vessels in the direction of flow and showed continuous cell-cell junctions (Fig. 2B and data not shown). In contrast, E14 mesenteric veins unexpectedly showed disrupted cell-cell junctions between adjacent venous ECs, characterized by large intercellular gaps and filopodial extensions (Fig. 3A,B, Movie S4, 5). Intercellular gaps were observed by staining with both Nrp2 and PECAM1 antibodies (Fig. 3A). Co-staining with TER-119 antibodies showed the presence of RBCs in the intercellular gaps at the level of EC layer and protrusion into the extravascular space (Fig. 3B). Most endothelial gaps were 6-10 μm in diameter (Fig. 3C), with an average diameter of 9.5 ± 6.2 μm (n=294), and thus permissive for extravasation of both enucleated (6 μm) and nucleated (8 μm in diameter) RBCs. Concomitant with increased mural cells
recruitment and BM deposition (Fig. 2B), intercellular gaps were no longer detected in E15 mesenteric veins (data not shown). These observations suggest that normal vascular remodeling in the mesentery involves a transient loss of venous endothelial integrity, which correlates with extravasation of RBCs.

**Platelets maintain venous integrity and prevent excessive RBC extravasation during mesenteric vascular remodeling**

Platelets form aggregates to limit blood loss and plasma leakage in primary homeostasis and upon vascular injury (Ho-Tin-Noé et al., 2011). We hypothesized that they may also be involved in maintaining vessel integrity during mesenteric vascular remodeling. Staining for the CD41 marker revealed the presence of platelets at endothelial gaps, but also at the endothelium in areas where no gaps were detected (Fig. 4A). Although large aggregates were not observed, filopodia extension was indicative of platelet activation (Fig. 4A). Notably, unlike RBCs, platelets were neither found outside of the blood vessels, nor in association with LECs (Fig. 4A,B), suggesting that their activation prevents extravasation. Lack of endothelial association at E15 further suggests that platelets adhere to the endothelial layer only at the stage when intercellular gaps are present (Fig. 4A).

To investigate the functional importance of platelets in the maintenance of venous integrity we used Clec1b (encoding CLEC2) deficient mice that show impaired platelet function leading to decreased thrombus stability (May et al., 2009; Suzuki-Inoue et al., 2010), and, intriguingly, blood filled lymphatic vessels (Bertozzi et al., 2010; Finney et al., 2012; Suzuki-Inoue et al., 2010). The latter phenotype was initially described as a failure in the developmental separation of the two vascular systems following the formation of the first lymphatic vessels via sprouting from major veins. More recently, the underlying mechanism was assigned to back filling of lymphatic vessels with blood from the thoracic duct due to defective platelet aggregation and thrombus formation at the lymphovenous junction (Hess et al., 2014; Welsh et al., 2016). In agreement with previous studies, we found that mesenteric lymphatic vessels of E14 Clec1b−/− embryos were filled with RBCs (Fig. 4C). Notably, Clec1b−/− mesenteries also showed extravascular platelets that were associated with LECs (Fig. 4B,C). Both platelets and RBCs were found in the interstitial space and associated with LEC clusters at E14 but prior to establishment of lumenized lymphatic vessels in the mesentery (Fig. 4C) and skin of Clec1b−/− mutants (Fig. S2). The observation that Clec1b−/− embryos show extravascular RBCs and platelets prior to establishment of functional lumenized lymphatic vessels that are connected to the circulation argues that their
filling with blood occurs secondary to disruption of blood vessel integrity due to platelet dysfunction.

These findings raise a question on the mechanism of CLEC2 dependent activation of platelets at venous endothelial gaps. Podoplanin is so far the only known endogenous ligand of CLEC2 (Navarro-Núñez et al., 2015; Pollitt et al., 2014), and Pdpn knock-out mice also show blood-filling of lymphatic vessels (Fu et al., 2008; Uhrin et al., 2010). Podoplanin is highly expressed in mesenteric LEC clusters at E14 (Fig. S3A,B; (Stanczuk et al., 2015)), but also in non-ECs (Fig. S3A,B). This is in agreement with previous reports showing expression of podoplanin in multiple cell types, of which mesothelial cells (Schacht et al., 2005), leukocytes (Kerrigan et al., 2012; Lee et al., 2010) and stromal fibroblasts (Kawase et al., 2008; Christer Betsholtz, personal communication) may be relevant in the context of the developing mesentery. To further characterize the PDPN+ cells, we utilized embryos carrying the Pdgfrb-eGFP transgene that labels mural cells (He et al., 2016) and a large stromal cell population in E14 mesenteries (Fig. S3B). FACS analysis of E14 mesenteries showed three distinct podoplanin expressing cell populations; PECAM1+PDPNhighGFP-LECs as well as PECAM1+stromal cells of which PDPNIintermedGFP- cells likely represent mesothelial cells and PDPNlowGFP+ fibroblasts (Fig. S3C). Thus, at least three populations of PDPN expressing cells; LECs, mesothelial cells and fibroblasts, are in the immediate vicinity of the developing mesenteric veins and may be involved in establishing endothelial integrity through interaction with platelet CLEC2. Generation of mice lacking podoplanin in specific cell types during embryonic development will be important for addressing this question in the future. Notably, postnatal LEC-specific deletion of Pdpn leads to progressive blood filling of lymphatic vessels over a few weeks (Bianchi et al., 2017). The mechanism is however different from the embryonic process and attributed in both Pdpn and Clec1b deficient mice to back filling of the thoracic duct and lymph nodes with blood through the lymphovenous connection (Bianchi et al., 2017; Hess et al., 2014). Yet another CLEC2-PDPN mediated mechanism safeguarding the lymphatic vasculature has been described in the lymph node, where the integrity of high endothelial venules (HEVs) was shown to depend on the interaction between PDPN in fibroblastic reticular cells, surrounding HEVs, and platelet CLEC2 (Herzog et al., 2013). Similarly, platelet CLEC2 interaction with PDPN on the neuroepithelium is required for the maintenance of cerebrovascular integrity (Lowe et al., 2015).

Taken together, our study demonstrates that embryonic remodeling of the mesenteric blood vasculature is associated with a transient loss of venous integrity due to formation of large intercellular gaps in the endothelium and consequent RBC extravasation. Such a disruption
of vascular integrity has been observed in pathological conditions (Hashizume et al., 2000; Mazzone et al., 2009), but our results show that it also occurs as part of a normal developmental process. We further show that platelets are pivotal to maintain venous integrity and prevent excessive RBC extravasation during mesenteric vascular remodeling, while lymphatic vessels clear the extravasated RBCs from the tissue (Fig. 4D). It is appealing to speculate that CLEC2 represents an embryonic mechanism of platelet activation when BM collagens, which provide a major platelet activation signal in the mature vasculature (Dütting et al., 2012), are scarce. Finally, our data provide mechanistic insight into how vascular abnormality characterized by blood-filled lymphatic vessels arises, by showing that it can occur secondary to loss of venous integrity. Notably, the blood-filled lymphatic vessel phenotype has been observed in various genetic models and has been considered as direct evidence of primary lymphatic vascular defects. In light of our findings, defects in the blood vasculature may also need to be considered.

Materials and methods

Mice

Vegf3-CreER<sup>T2</sup> (Martinez-Corral et al., 2016), R26-mTmG (Muzumdar et al., 2007), Cldn5-GFP (a gift from C. Betsholtz; Barbara Lavina and Christer Betsholtz, unpublished), Clec1b<sup>flox</sup> (Acton et al., 2014) Pdgfrb-eGFP (He et al., 2016) and wild type mice were analyzed on a C57BL/6J background. Clec1b<sup>flox</sup> were crossed with PGK-Cre mice to generate germline heterozygous mice that were further crossed to generate germline homozygous embryos. The morning of vaginal plug detection was considered as embryonic day 0. Mosaic labeling of individual LECs by membrane-bound GFP in E13 R26-mTmG;Vegf3-CreER<sup>T2</sup> embryos was induced by administration of 0.5 mg of 4-hydroxytamoxifen to pregnant females at E12. Experimental procedures were approved by the Uppsala Laboratory Animal Ethical Committee.

Whole-mount immunofluorescence

Mesenteries were fixed in 4% paraformaldehyde at room temperature for two hours and stained as previously described (Stanczuk et al., 2015). The following primary antibodies were used: mouse anti-α-Smooth Muscle Actin-Cy3 (Sigma-Aldrich C6198) (1:250), rat anti-Endomucin (V.7C7) (Santa Cruz SC-65495) (1:200), rat anti-CD41-FITC (eBioscience 11-0411-81) (1:50), rabbit anti-Collagen IV (BioRad 2150-1470) (1:500), chicken anti-GFP (Abcam ab13970) (1:200), goat anti-Neuropilin2 (R&D Systems AF567) (1:200), hamster
anti-PECAM1 (Millipore MAB1398Z) (1:1000), rat anti-PECAM1 (BD Pharmingen 553370) (1:1000), rat anti-PECAM1-AF594 (BioLegend, 102520) (1:100), rabbit anti-human PROX1 (Stanczuk et al., 2015) (1:200), rat anti-TER-119 (eBioscience, 145921) (1:200), rat anti-TER-119-AF647 (Biolegend 116218) (1:50), goat anti-VE-Cadherin (Santa Cruz SC-6458) (1:200), and anti-podoplanin (Developmental Studies Hybridoma Bank at the University of Iowa, clone 8.1.1) (1:800). Autofluorescence signal at 550-600 nm wavelength was used for visualization of RBC. Secondary antibodies conjugated to AF488, AF594, AF647 or Cy3 were from Jackson ImmunoResearch, and all were used at 1:300 dilution.

**Image acquisition and quantification**

Confocal image stacks were acquired using Leica SP8 confocal microscope and LAS X software. Images represent maximum intensity projections of tiled z stacks. Images were processed with Fiji or Adobe Photoshop software. Deconvolution (Fig. 1F) was carried out in Huygens Essential v16.5 (Scientific Volume Imaging) using a theoretical PSF and automatic background estimation. Stopping criteria were set to 40 iterations and a signal to noise ratio of 10. Structural illumination raw image stacks were acquired on a Zeiss Elyra S.1 LSM710 together with a Plan-Apochromat 63x/1.4 Oil DIC M27 objective. GFP and Cy3 were imaged sequentially as separate tracks and excited at 488 and 561 nm respectively using the same grid size of 28 µm. For detection beam splitters BP 495-550 + LP 750 and BP 570-620 + LP 750 were used. Tracks were switched for each z stack. Reconstruction of structural illumination raw images was done using the SIM module of ZEN software (Carl Zeiss). The noise filter was manually set to -4.0. Same parameters were set for both channels. Maximum intensity projections of deconvolved or reconstructed image stacks were generated using Fiji. Movies S1, 4, 5 were generated in Fiji and recorded at 3 fps. Movies S2, 3 were generated in Imaris v8.4 (Bitplane) and recorded at 25 fps.

For quantification of intra/extravascular RBCs, mesenteries were stained for TER-119, PROX1 and Nrp2 (all samples), and PECAM1 (E13, E13.5 and E14 samples) to label RBCs, LECs, and (venous) blood ECs respectively. Images of mesenteries were acquired as multiple tile scans using HC FLUOTAR L 25x/0.95 W VISIR objective in resonant scan mode on a Leica SP8 confocal microscope. For each image, a z-stack was acquired at 5 µm intervals through the entire thickness of tissue containing blood and lymphatic vessels. RBCs were counted from individual stacks of images of whole mesenteries (E13; n=5 embryos) or along segmentally organized blood vessels covering the entire length from the mesenteric root to the intestinal wall (E13.5 (n=4 embryos and 5-7 vessels per embryo; total 23 vessels), E14 (n=7 embryos and 4-7 vessels per embryo; total 39 vessels), E16 (n=5
embryos and 6-9 vessels per embryo; total 37 vessels) using the Cell Counter plugin for Fiji.

Data summarized in Fig. 1D represent values for individual embryos (E13) or vessels (13.5, E14, E16). Different categories of RBCs were assigned based on localization: intravascular (inside the blood vessels) or extravascular (outside of the blood vessels). The latter were further categorized as associated with LECs or their protrusions, or not associated. If RBCs co-localized with LEC cell bodies and/or protrusion in the same optical section(s), they were defined as associated with LECs/protrusions; otherwise they were defined as not associated.

For quantification of openings within the endothelial layer, gaps were extracted from individual single Z stacks from which they could be clearly observed, and their areas (A) were measured using the Measure tool of Fiji. The gaps were assumed to be circular, and diameters (d) were calculated from measured areas (d = 2√A/π), as described previously (Hashizume et al., 2000). 294 gaps from 40 images acquired from 14 wild type E14 mesenteries were measured.

For quantification of platelet areas, mesenteries from E14 embryos (Clec1b+/− (n=3) and Clec1b−/− (n=3)) were stained for CD41 to label platelets, and Nrp2 to label LECs and venous blood ECs. Single-tile images were acquired using a HC PL APO CS2 63x/1.30 Glyc CoRR CS3 objective and Leica SP8 confocal microscope. For each sample, three single-tile images were taken from three different regions of the mesentery. The images were acquired at 2 μm intervals through the entire thickness of tissue containing blood and lymphatic vessels. CD41 positive areas, both total and LEC-associated, were measured in maximum intensity projection images after threshold adjustment using Fiji. Optical sections to be included for each measurement (e.g. covering the thickness of a lymphatic vessels) were defined individually from each z-stack.

The data were calculated using Microsoft Excel for Mac 2011, and summarized and graphed using GraphPad Prism 6 for Mac.

Flow cytometry

Mesenteries of E14 Pdgfrb-eGFP embryos were dissected and dissociated at 37°C and 550 rpm for 5 to 10 minutes with 2 mg/mL Collagenase IV (Life Technologies) and 0.2 mg/mL DNase I (Roche) in PBS supplemented with 0.2% FBS (Gibco). Digests were quenched by adding 2 mM EDTA, filtered through a 70 μm nylon filter (BD Biosciences) and washed twice.
with FACS buffer (0.5% FBS and 2 mM EDTA in PBS). Digests were incubated for 15 min with 5 µg/mL rat anti-mouse CD16/CD32 IgG (clone 93, eBioscience) to block Fc-receptor binding and subsequently stained with rat anti-PECAM1/CD31-PE-Cy7 (0.67 µg/mL clone 390, eBioscience), hamster anti-PDPN-eF660 (2 µg/mL clone eBio8.1.1., eBiosciences), rat anti-CD11b-eF450 (4 µg/mL clone M1/70, eBioscience), and rat anti-CD45-eF450 (4 µg/mL clone 30-F11, eBiosciences) antibodies for 30 min on ice. 1 mM Sytox-Blue (Life Technologies) was used to assess cell viability. Single stained samples were used for compensation. Samples were analysed on a BD Cytoflex S flow cytometer equipped with CytExpert software (BD Biosciences) and processed using FlowJo 10.3 software (FlowJo, LLC).

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Author contributions: YZ designed, performed and analyzed most of the experiments. ND designed, performed and analyzed experiments shown in Fig. 1F,G, S1C. SS designed, performed and analyzed experiments in Fig. S3B,C. TM designed experiments and analyzed data, supervised the research and wrote the manuscript with YZ. All authors discussed the results and commented on the manuscript.

Competing interests

No competing interests declared.

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References


Figures

A E14 wild type mesentery

B E14 wild type mesentery

C extravascular RBC - LEC interactions

D Extravascular RBC

E RBC association with LEC

F R26-mTmG;Vegfr3-CreERT2

G R26-mTmG;Vegfr3-CreERT2

3D-rendered
Figure 1. Transient RBC extravasation during the development of the mesenteric vasculature

(A, B) Whole-mount immunofluorescence (A, B) and differential interference contrast (DIC) imaging (A) of E14 mesenteries showing RBCs inside lymphatic vessels (arrowheads). Both nucleated (TER-119$^{\text{low}}$, yellow arrowheads in B) and enucleated (TER-119$^{\text{high}}$, white arrowhead in B) were present. V, mesenteric vein.

(C) Whole-mount immunofluorescence of mesenteric vessels for the indicated antibodies showing TER-119$^+$ RBC interactions with Nrp2$^{\text{high}}$ LEC protrusions (E14; on the left), and engulfment by forming (E14; in the middle) and lumenized lymphatic vessels (E15; on the right). Z-views at the indicated positions are shown below and on the right of each image.

(D) The percentage of extravascular RBCs versus total (intravascular and extravascular) RBCs in the mesentery at the indicated stages of development. Dots represent individual embryos (E13 n=5) or vessels (13.5 n=23, E14 n=39, E16 n=37) and horizontal lines represent mean values (see materials and methods for details).

(E) The percentage of extravascular RBCs at E14 showing association with LEC protrusions (light grey), the LEC cell body (in most cases engulfment of RBCs by a cluster of LECs, and less frequently localization next to a cluster but associated with the cell body of an LEC; dark grey), or no association with LECs (white). Dots represent individual embryos (n=7), mean ± SD.

(F) Deconvolved confocal image showing contact (arrowhead) between a protrusion from a mesenteric LEC (visualized by Nrp2 and PROX1 staining and GFP expression in mosaically induced R26-mTmG;Vegfr3-CreERT2 embryo) and extravascular RBC (visualized by autofluorescence signal). Boxed area is shown on the right as a surface-rendered 3D image from angles that reveal association between RBC and LEC protrusion.

(G) Reconstructed SIM images showing direct contacts between LEC (PROX1$^+$GFP$^+$) protrusions and RBC (autofluorescence signal) in E13.5 R26-mTmG;Vegfr3-CreERT2 mesentery. Boxed area is shown below as a digital zoom-in of maximum intensity projection of central part of the image stack and as surface-rendered 3D reconstructions of whole image stack viewed from opposite angles illustrating how LEC protrusions align with the surface of the RBC.

Scale bars: 20 μm (A-C), 10 μm (F,G upper panel), 2 μm (G lower panel).
Figure 2. Mesenteric blood vasculature undergoes extensive remodeling and maturation between E13-E15.

(A) Visualization of the mesenteric vasculature in Cldn5-GFP embryos showing extensive remodeling from a primary plexus into segmentally organized pattern of veins and arteries between E13-E14. Single channel images of boxed areas show co-staining for the venous EC marker Nrp2 in only a subset of vessels.

(B) Whole-mount immunofluorescence of E14 (left) and E15 (right) mesenteric vessels for markers of ECs (VE-cad), mural cells (αSMA) and basement membrane (Collagen IV). Single channel images of indicated stainings are shown. Note poor EC alignment as well as mural cell and BM coverage in E14 vein (V) when compared to E15 vein (arrowheads), or the artery (A).

Scale bars: 100 μm (A) and 20 μm (B).
Figure 3. Remodeling of the mesenteric blood vasculature is associated with a transient loss of venous endothelial integrity.

(A, B) Whole-mount staining of E14 mesenteries for the indicated antibodies, showing intercellular gaps in the veins. Z-projections of confocal stacks are shown and boxed areas are magnified on the right. Z-views at the indicated positions are shown below. Arrowheads indicate gaps in the endothelial layer.

(C) Size distribution of intercellular gaps in wild type E14 mesenteric veins (n=294 gaps from 14 embryos).

Scale bars: 20 μm.
Figure 4. Platelets are critical for maintaining integrity of remodeling veins.

(A) Adherence of platelets (CD41⁺) to venous ECs (Nrp2low) and gaps at E14, but not at E15. Z-views of the stack at the indicated positions are shown and boxed area is magnified on the right. Note filopodia extension (arrows in the magnified image showing CD41 staining alone), indicative of platelet activation, at E14.
(B) Quantification of extravascular (LEC-associated) platelets in E14 wild type and Clec1b−/− mesenteries. Dots represent individual embryos (n=3, mean of three areas imaged for each embryo) and the horizontal lines represent mean values.

(C) Whole-mount immunofluorescence of E14 wild type and Clec1b−/− mesenteries for Ter119 (RBC), CD41 (platelets), Nrp2 (lymphatic vessels (high) and veins (low)). Note the presence of RBCs and platelets in association with LEC clusters (E14-clusters panel, arrowhead) and in the extravascular space (arrows), as well as in lumenized lymphatic vessels (E14-vessels panel, arrowheads) in Clec1b−/− but not wild type mesenteries. RBC and platelet staining of the boxed areas are shown to the right (E14-vessels panels). Z-views of the stack at the indicated positions are shown below and on the right of the bottom panel E14-clusters.

(D) Proposed model of platelet function in mesenteric vascular development and the mechanism underlying blood filling of lymphatic vessels in Clec1b−/− embryos.

Scale bars: 20 μm.
**Supplementary figures**

**Figure S1. Analysis of extravascular RBCs in the mesentery.**

(A) Representative maximum intensity projections of whole-mount stained E13, E13.5 and E14 mesenteries used for quantification of intra/extravascular RBCs. Antibodies are indicated. Boxed areas are magnified in Fig. 1C.
(B) Representative maximum intensity projection of whole-mount stained E13 mesentery showing rare extravascular RBCs (arrows) and their locations relative to LEC clusters. Many LEC clusters are not associated with RBCs at this stage (arrowheads). Antibodies are indicated.

(C) Maximum intensity projection of a LEC cluster in E13 mesentery extending membranous protrusions of which some are in direct contact with nucleated (TER-119\textsuperscript{low}) extravascular RBC. Antibodies are indicated.

Scale bars represent 100 μm (A), 50 μm (B) and 10 μm (C).
Figure S2. Association of RBCs and platelet with dermal LEC clusters in Clec1b<sup>−/−</sup> skin.

Whole-mount immunofluorescence staining of E14 wild type and Clec1b<sup>−/−</sup> dorsal skins using the indicated antibodies. TER-119<sup>+</sup> RBCs and CD41<sup>+</sup> platelets are found outside of Nrp2<sup>low</sup> (left panel) and PECAM1<sup>+</sup> (right panel) blood vessels (arrowheads) and engulfed inside Nrp2<sup>high</sup> LEC clusters (left panel; arrow) in Clec1b<sup>−/−</sup> but not in wild type skins. Z-views at the indicated position are shown. Scale bars represent 20 μm.
Figure S3. Podoplanin expression in the developing mesentery.

(A) Whole-mount staining of E14 mesentery for the indicated antibodies showing podoplanin^{+}\text{PROX1}^{+} LECs and podoplanin^{+}\text{PROX1}^{-} stromal cells. Z-projection of a confocal
stack and Z-views at the indicated positions are shown on the left. The distance between the podoplanin⁺PROX1⁻ cell layer (shown on the right as a single plane image) and the Endomucin⁺ vein is about 12 μm.

(B) Whole mount staining of an E14 Pdgfrb-eGFP mesentery for the indicated antibodies showing broad expression of GFP in mural cells and mesenteric stromal cells but not in ECs (single plane, arrowhead; LV, lymphatic vessel; V, vein). The maximum intensity projection of a tiled z-stack over a distance of 32 μm reveals the presence of podoplanin expressing stromal cells (arrows). Boxed area is magnified on the right.

(C) Flow cytometric analysis of podoplanin expression in PECAM1 positive endothelial cells and PECAM1 negative stromal cells from E14 Pdgfrb-eGFP mesenteries. FACS plots are representative of 12 mesenteries.

Scale bars represent 20 μm (A) and 50 μm (B).
Supplementary Table S1. Image acquisition details.

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Movies

Movie S1. Z-stack of confocal images of E14 mesenteric vessels stained for markers of venous/lymphatic ECs (Nrp2) and RBCs (TER-119).
Series of confocal images of E14 mesenteric vessels stained for Nrp2, PROX1 and TER-119, showing interactions between mesenteric LECs and extravascular RBCs. The movie goes through 7.28 μm distance. Images within the series were acquired at 0.28 μm intervals and the movie is displayed at 3 frames per second.
Movie S2. 3D reconstruction of E13.5 mesenteric vessels showing close association between LEC and extravascular RBC.

A movie showing 3D reconstruction of deconvolved confocal images of an E13.5 R26-mTmG;Vegfr3-CreERT2 mesentery stained for GFP (green), Nrp2 (red), PROX1 (grey; LEC nuclei) or visualized by autofluorescence (grey; RBC). The image stack was acquired at 0.20 μm intervals covering a z distance of 23.2 μm. 3D surface rendering was applied based on signal intensity to highlight close association of a LEC protrusion and RBC. A corresponding z-projection is shown in Fig. 1F. The movie is displayed at 25 frames per second.
Movie S3. 3D reconstruction of structural illumination image stack showing close association between LEC cluster and extravascular RBC in an E13.5 mesentery.

A movie showing 3D reconstruction of an image stack acquired by structural illumination microscopy of whole mount E13.5 R26-mTmG;Vegfr3-CreERT2 mesentery stained for GFP (green), PROX1 (grey; LEC nuclei) or visualized by autofluorescence (grey; RBC). The image stack was acquired at 0.11 μm intervals covering a z distance of 10.1 μm. 3D surface rendering was applied based on signal intensity. A corresponding z-projection is shown in Fig. 1G. The movie is displayed at 25 frames per second.
Movie S4. Z-stack of confocal images of E14 mesenteric vessels stained for the pan-EC marker PECAM1.

Series of confocal images of E14 mesenteric vessels stained for PECAM1, showing intercellular gaps in the venous endothelium. The movie goes through 28.28 μm distance covering the entire thickness of tissue containing blood vessels (Movie S4A), or 4.2 μm distance covering the venous endothelial layer at the marked position (yellow arrow) to highlight an intercellular gap (Movie S4B). Corresponding z-projections are shown in Fig. 3A (left panels). Images within the series were acquired at 0.28 μm intervals and the movie is displayed at 3 frames per second.
Movie S5. Z-stack of confocal images of E14 mesenteric vessels stained for markers of venous/lymphatic ECs (Nrp2) and RBCs (TER-119).

Series of confocal images of E14 mesenteric vessels stained for Nrp2 and TER-119, showing intercellular gaps in the venous endothelium. The movie goes through 13.72 μm distance covering the entire thickness of tissue containing blood and lymphatic vessels (Movie S5A), or 1.96 μm distance covering the venous endothelial layer at the marked position (yellow arrow) to highlight an intercellular gap (Movie S5B). Corresponding z-projections are shown in Fig. 3B (right panels). Images within the series were acquired at 0.28 μm intervals and the movie is displayed at 3 frames per second.