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Intraluminal pressure triggers a rapid and persistent reinforcement of endothelial barriers†

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In response to mechanical cues, endothelial cells elicit highly sensitive cellular response pathways that contribute to the regulation of the physiology and disorders of the vascular system. However, it remains relatively unexplored how endothelial tissues process and integrate the intraluminal pressure, and in turn regulate the permeation flow across the vessel wall. Leveraging a tissue engineering approach to create microvessels (MVs), we measured real-time permeation flow induced by intraluminal pressures ranging from 0.1 to 2.0 kPa. Our findings reveal that mechanically stimulated MVs strengthen their barrier function within seconds of exposure to pressures below 1 kPa, with this enhanced barrier function persisting for 30 minutes. We demonstrate that this barrier reinforcement is linked to the closure of paracellular gaps. Additionally, we observe that it is associated with, and depends on, actin cytoskeleton reorganization, including the accumulation of stress fibers near intercellular junctions and the broadening of adherence junction protein localization. These findings provide insights into the ability of endothelial tissues to regulate interstitial fluid flow in response to sudden increases in blood pressure.

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Normal organ function relies on the precise regulation of blood vessel permeability, which governs the influx of molecular and macromolecular components from plasma.¹ This barrier function is actively maintained in quiescent vasculature,² primarily through organ-specific regulation of intercellular junction tightness.³ It is also dynamically responding to biochemical stimulations, *e.g.* thrombin⁴ or histamine,⁵ that destabilize the endothelial barrier function⁶ by disconnecting junctions and gradually forming intercellular separations. This destabilization is also associated with the reorganization of the actin cytoskeleton,⁷ a hallmark of mechanotransduction,⁸ wherein biochemical signals induce structural and physical changes within endothelial cells. Conversely, mechanotransduction triggered by shear stress^{9,10} prompts elongation and alignment of endothelial cells in the direction of flow¹¹ while enhancing

barrier integrity under shear stress below ~ 1 Pa.¹² Similar effects are observed in cyclic uniaxial stretch experiments,^{13–16} where elongating endothelial tissues induce cytoskeletal reorganization perpendicular to the strain direction and strengthen the barrier function at strains below $\sim 5\%$. At the molecular level, these adaptations involve the junctional mechanosensory complex,¹⁷ which orchestrates stress fiber assembly and regulates the distribution of junctional proteins.^{18,19} While these mechanisms have been extensively studied using fluid shear stress as a mechanical cue and monitoring barrier integrity through electrical resistance-based methods,²⁰ the dynamic permeability changes of stretched endothelial cell monolayers remain more elusive. Circumferential stress is nevertheless a physiological cue of blood flow, which is characterized by a typical intraluminal pressure of ~ 2 kPa in microvasculatures²¹ with abrupt increases observed during hypertension.²²

To address this knowledge gap, we developed a technology that enables real-time monitoring of permeation flows across stretched endothelial cell monolayers. This technology is based on a microvessel-on-a-chip (MV) platform,^{23–25} which allows dynamic regulation of intraluminal pressure within the range of 50 to 2000 Pa. Permeation flow rates are simultaneously recorded over time windows spanning minutes to hours using a time interval of 100 ms. We demonstrate that the endothelial barrier actively responds to intraluminal pressure, becoming significantly less permeable within tens of seconds. We show that this reinforcement of

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the barrier is explained by the closure of paracellular pores, and report that these closure events can be challenged by an excessive intraluminal pressure of 1500 Pa or more. In the low-pressure regime of barrier reinforcement, we then prove that intraluminal pressure stimulation induces and depends on the reorganization of the actin cytoskeleton with the accumulation of stress fibers at the vicinity of intercellular junctions. Barrier reinforcement triggers, but is independent of, the activation of the MAP kinase mechanosensory pathway. Conversely, inhibition of the rho kinase compromises barrier reinforcement, destabilizing intercellular junctions for low intraluminal pressures. We finally discuss these findings in the context of endothelial tissue response to bursts in blood pressure.

Results

MVs constitute dynamic barriers to *trans*-endothelial flow

Using human umbilical vein endothelial cells (HUVECs), which represent a standard for *in vitro* studies of vasculature and angiogenesis^{26,27} and for endothelial phenotypic response to mechanical stimulation,^{25,28,29} we formed MVs of

200 μm in diameter and 6 mm in length within a collagen gel scaffold (Fig. 1A; detailed fabrication methods in the Materials and methods section). After two days of culture in standard “static” conditions, we mounted a fluidic device onto the MV chip (Fig. 1B), enabling dynamic control of inlet pressures. By applying equal pressures to both inlets, we created a homogeneous intraluminal pressure with a fluid shear stress that was four orders of magnitude lower in amplitude (Fig. 1C and see the calculation in the Materials and methods). The fluidic device and pressure actuation approach were adapted from previously reported methods for collagen gel mechanical characterization.³⁰ To measure the permeation flow rate (Q) across the endothelial tissue and the collagen matrix (indicated by red arrows in Fig. 1B), we installed two flow meters along the injection tubing and applied mass conservation principles, which indicate that the fluid passing through these sensors equals the amount exiting across the MV and collagen slab. This setup hence allowed us to stimulate the MV with intraluminal pressure and simultaneously measure Q . Additionally, the system's integration with a wide-field microscope enabled cumulative observation of MV deformation over time.

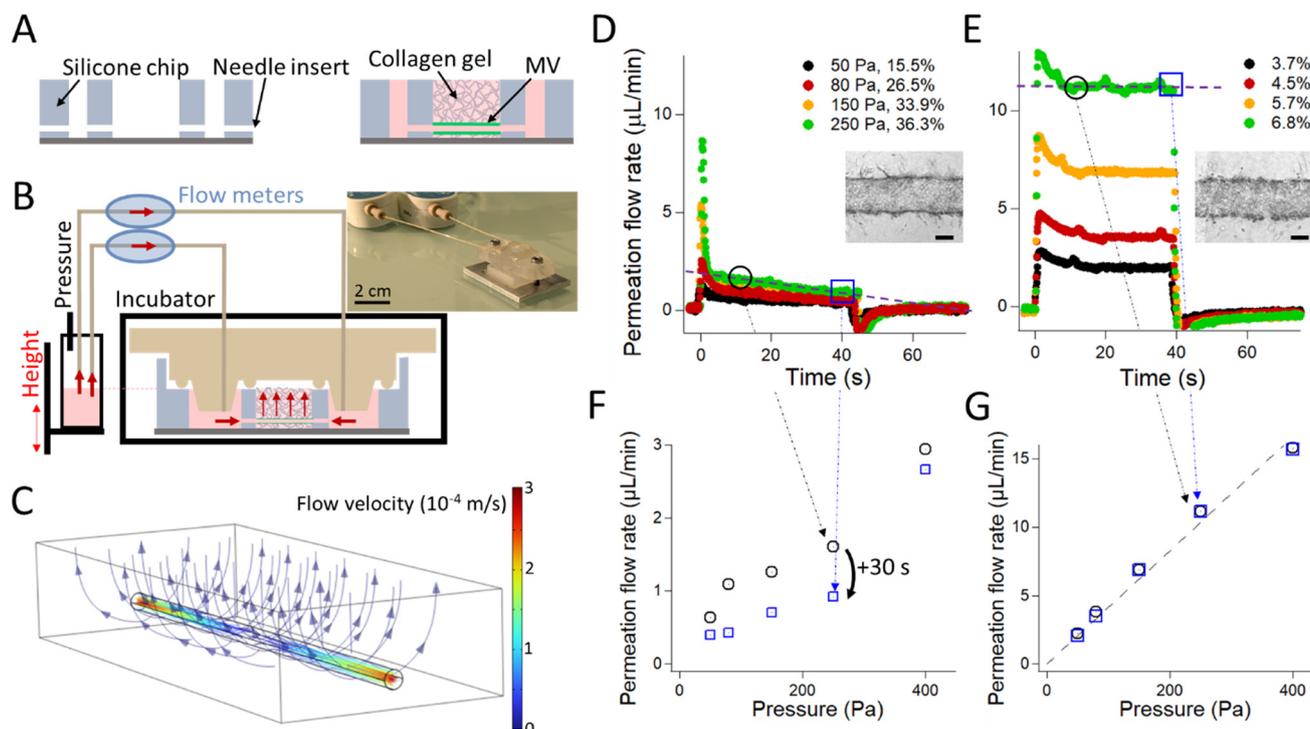


Fig. 1 Real-time recording of the permeation flow through MVs. (A) The sketch represents the silicone chip to fabricate a hollow lumen in a collagen gel, which is seeded with endothelial cells to form the MV (green lines). (B) Schematic of the system used to apply intraluminal pressure and measure the permeation flow. The inlet tubes are immersed in a media-filled vial (left), where the pressure is dynamically controlled. The tubes are connected to the MV chip at their opposite ends. The MV is semipermeable, allowing fluid to exit (indicated by red arrows) while passing through flow meters to quantify the flow rate. The accompanying photograph shows the MV chip, along with the tubing and flow meters. (C) Finite element simulation of the fluid flow velocity in the MV chip (see Fig. S1† for the pressure field and Materials and methods for the implementation). (D) The graph shows the permeation flow rate as a function of time for an intraluminal pressure spanning 50 to 250 Pa, as indicated in the legend. (E) The permeation flow rate is recorded after overnight treatment with sodium azide of the same MV as in (D). The micrographs in the inset of (D) and (E) show the MV and the lumen with cell residues, respectively. (F) The graph represents the flow rate as a function of the intraluminal pressure, as measured after 10 and 40 s of actuation (black and blue datasets, respectively). (G) Same as (F) after sodium azide treatment. The scale bars correspond to 100 μm .



We started by applying intraluminal pressures spanning 50 to 400 Pa, which is typically comparable to the stress induced by the pulsatile pressure in human capillaries of ~ 300 Pa, but not to its continuous component of 2000 Pa.³¹ The resulting permeation flow traversing the MV was on the order of $Q \sim 1 \mu\text{L min}^{-1}$, and the deformation of the MV in the range of 10 to 30% (Fig. 1D). The barrier function of the MV became evident when it was disrupted using the cytotoxic agent sodium azide, which caused a dramatic increase in flow through the collagen matrix—over tenfold for an intraluminal pressure of 250 Pa (Fig. 1E). Notably, we observed that the deformation of the MV and collagen matrix was fully reversible within this range of mechanical stimulation (Fig. S1).[†] In turn, these experiments enabled us to estimate the permeability of the MV, quantified by the Darcy coefficient.³² Considering the known permeability of collagen gels ($7 \times 10^{-14} \text{ m}^2$,³⁰) and the thickness of the collagen slab (2 mm) relative to the much thinner endothelial MV ($\sim 1 \mu\text{m}$,²⁵), the observed tenfold reduction in permeation flow suggested an endothelial tissue permeability of

approximately $5 \times 10^{-18} \text{ m}^2$ (see the details of the calculation in the Materials and methods section). This measurement was in agreement with published estimates,^{25,32} which modeled *trans*-endothelial fluid transport as convection through paracellular holes. To confirm that fluid transport occurred primarily *via* paracellular, rather than transcellular, pathways,³³ we fixed the MV with formaldehyde immediately after manipulation. The permeation flow rate Q remained similar in amplitude before and after the fixation (Fig. S2A[†]), indicating that fluid transport was passive and occurred through paracellular pores, rather than being actively mediated by endothelial cells.

We then investigated the temporal variation of the permeation flow rate. In experiments with dead cells (Fig. 1E), we observed a rapid reduction in the flow rate during the first 10 seconds, which was attributed to the deformation of a soft poroelastic material (Fig. S2B[†]). Following this transient phase, the flow rate stabilized, and we obtained the linear response in pressure expected for passive porous materials (dashed line in Fig. 1H). In contrast,

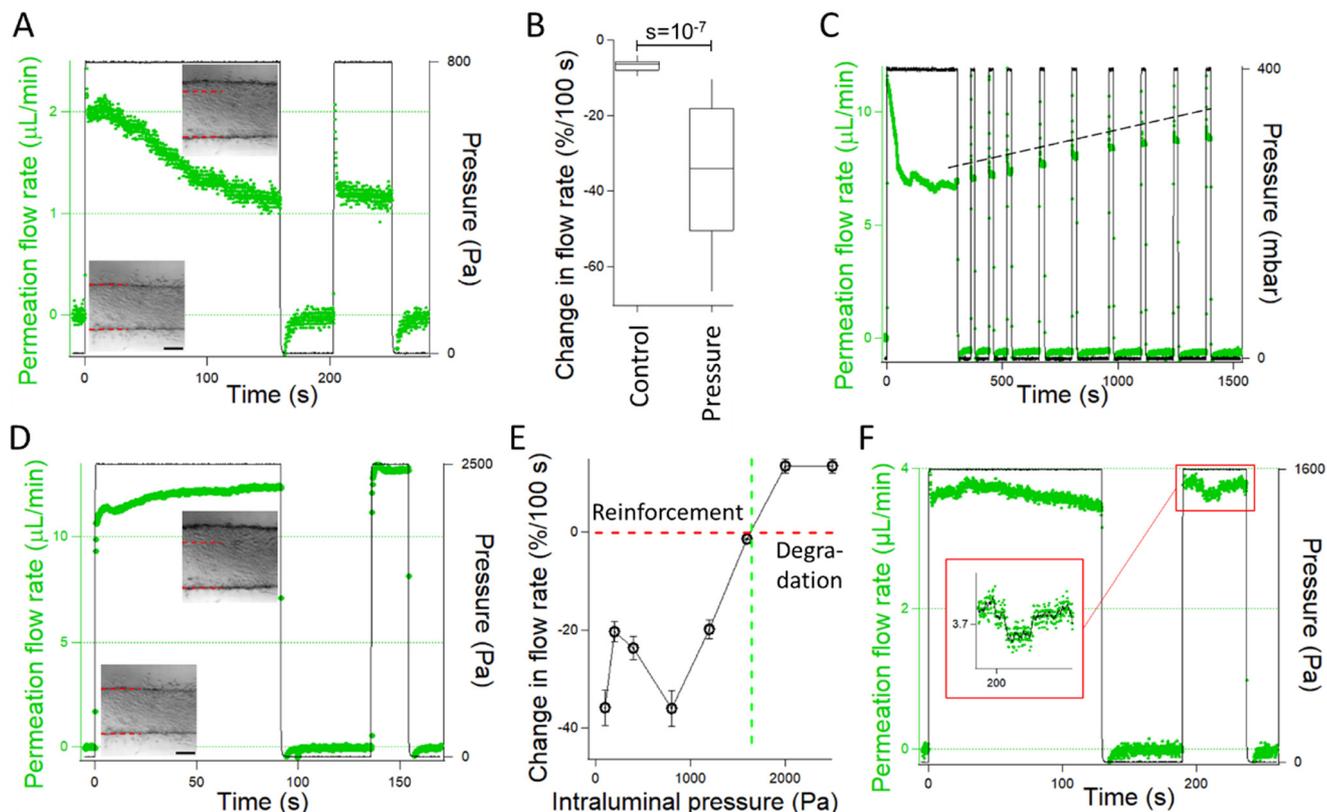


Fig. 2 Rapid and persistent barrier reinforcement under intraluminal pressure. (A) The graph shows the permeation flow rate as function of time (green dataset) using a temporal sequence of pressure actuation represented in black and reported on the right axis. Here, the intraluminal pressure is set to 800 Pa for 150 s, then stalled, and restored for 60 s. The two micrographs represent the MV in static and deformed by intraluminal pressure at the bottom and top, respectively. (B) The box plot reports the relative change in permeation flow rate in the low-pressure regime in live MVs or control based on fixed or dead cells. (C) Same experimental design as in (A) but the temporal sequence of ~ 30 min is ten times longer and the pressure is set to 400 Pa. (D) Using the same MV as in (A), we apply an intraluminal pressure of 2500 Pa for 90 s, arrest it for 45 s, and restore it for 20 s. (E) The graph reports the relative change in permeation flow rate after a time lag of 100 s as a function of the intraluminal pressure. The red dashed curve corresponds to a constant permeation flow over time. The green dashed line draws the limit between the regimes of reinforcement and degradation. (F) In the intermediate pressure regime of 1600 Pa, the permeation flow rate is recorded for 120 s. The zoom in the inset shows the fluctuations of the permeation flow. The scale bars correspond to 100 μm .



experiments with MVs exhibited a gradual decline in the flow rate over time (dashed line in Fig. 1D). To quantify this trend, we plotted the flow rate measured at 10 and 40 s that decreased across all pressure settings (Fig. 1F). This progressive decline in flow rate during stimulation suggests a change in the permeability of the endothelial cell monolayer, likely reflecting its active response to the mechanical cue.

Intraluminal pressure stably reinforces MV barrier function

We further characterized the active response of the stimulated MV to determine whether the modulation of the permeation flow rate by intraluminal pressure was transient or stable. To investigate this, we applied an intraluminal pressure of 800 Pa for 150 s, released the pressure for 60 s, and then re-applied the stimulation for an additional 50 s. The pressure is shown in the black dataset, with the y-axis on the right in Fig. 2A, and the resulting permeation flow rate in the green dataset on the left y-axis. This experiment confirmed the decrease of Q , which started from 2.0 and ended up at $1.1 \mu\text{L min}^{-1}$ in the first step of this experiment. The reduction was consistently observed in 24 MVs stimulated by an intraluminal pressure of 400 or 800 Pa and equal to $-36 \pm 20\%$ (Fig. 2B). On the contrary, it was significantly lower and equal to $-7 \pm 2\%$ in 14 controls obtained either by disrupting the tissue with sodium azide or by chemical fixation with paraformaldehyde (Fig. 2B). Then, Q returned to its initial null value as we stopped the pressure stimulation (Fig. 2A). More interesting was the fact that Q remained low during the second pressure pulse, with a value that precisely matched that at the end of the first stimulation of $1.2 \mu\text{L min}^{-1}$. Given that a decrease in permeation flow indicates an enhanced barrier, referred to here as 'reinforced', our data suggested a sustained barrier reinforcement following pressure stimulation. This result was strengthened by monitoring the permeation flow during ~ 30 minutes following an initial stimulation of five minutes at 400 Pa (Fig. 2C). Barrier reinforcement occurred within the first two minutes; however, we observed a slow relaxation phase over the ensuing 30 minutes while maintaining zero pressure and applying brief pulses of ~ 7 s to register Q (dashed line in Fig. 2C). We concluded that MV barrier reinforcement was rapid and persistent, in agreement with the swift activation of mechanosensitive pathways by fluid shear stress and the slow return to baseline.^{9,10}

High tensile forces can disrupt adhesion between endothelial cells.³⁴ Therefore, we anticipated that applying excessive intraluminal stress would destabilize the barrier. We repeated the experiment with the same MV as in Fig. 2A, but increased the intraluminal pressure to 2500 Pa (Fig. 2D). The permeation flow rate was observed to increase during mechanical stimulation, and the endothelial tissue remained degraded even after a 60 second pause. Structural analysis of the MVs using immunoconfocal microscopy confirmed that the stress-induced barrier degradation resulted in detectable gaps between adjacent cells (Fig. S3†). By scoring the relative

change in permeation flow rate during 100 s, we could then recapitulate a set of experiments performed on the same MV with two regimes of barrier reinforcement and barrier degradation (Fig. 2E). At the transition for an intraluminal pressure of 1500 Pa, the permeation flow rate remained roughly stable over time (Fig. 2F). However, a closer inspection of the signal revealed abrupt positive and negative fluctuations of Q . Using a Bayesian estimator to detect abrupt changes in temporal traces,³⁵ we registered a total of 35 jumps for a total recording time of 500 s and 5 different MVs. Their average amplitude was $95 \pm 81 \text{ nL min}^{-1}$ (positive and negative jumps being roughly equally frequent, Fig. S4†). At the threshold pressure, barrier reinforcement likely competed with the disruption of intercellular junctions, resulting in paracellular pore opening and closing events. Notably, this tentative mechanism could be used to estimate the typical size of these events. Indeed, the flow rate of $\sim 100 \text{ nL min}^{-1}$ for a pressure of 1500 Pa corresponded to a single pore approximately $1 \mu\text{m}$ in radius crossing an endothelial tissue of $1 \mu\text{m}$ in thickness (see the details of the calculation in the Materials and methods section). Consequently, analysis of the dynamics of barrier response to intraluminal pressure hints to a mechanism of reinforcement involving paracellular pore closure.

Paracellular pore closure events account for the reinforcement of the barrier

We aimed to gather further evidence on whether barrier reinforcement involved paracellular hole closure. We performed the macromolecular assay, which consisted in loading a fluorescent dextran molecule into the lumen of the microvessel (MV) and tracking its spatial redistribution within the surrounding extracellular matrix (Fig. 3A, see Materials and methods). The experiment was notably performed under constant intraluminal pressure, maintained at 100 Pa throughout the assay, as described in ref. 25 and 36. Leakage through paracellular gaps became immediately detectable following dye injection (upper left panel in Fig. 3A), because these gaps created an escape path for the rapid release of fluorescent molecules within the basal layer of the MV. These bright spots gradually spread over time due to diffusion, as illustrated in the difference micrograph between consecutive images (upper panel in Fig. 3B). Interestingly, this differential representation also revealed the appearance of localized dark spots at 15 and 38 s (Fig. 3B). These dark spots indicated an abrupt cessation of fluorescence convection, as expected for the closure of paracellular pores. These closure events occurred only once, as exemplified by the observation that the dark spot appearing after 15 s became undetectable at 38 s. Although spatial analysis beyond one minute was not possible with this assay, because the global fluorescence signal induced by leakage blurred out local intensity fluctuations, these experiments provided direct supporting evidence of localized rearrangements of the endothelial tissue architecture that reduce the strength of the permeation flow.



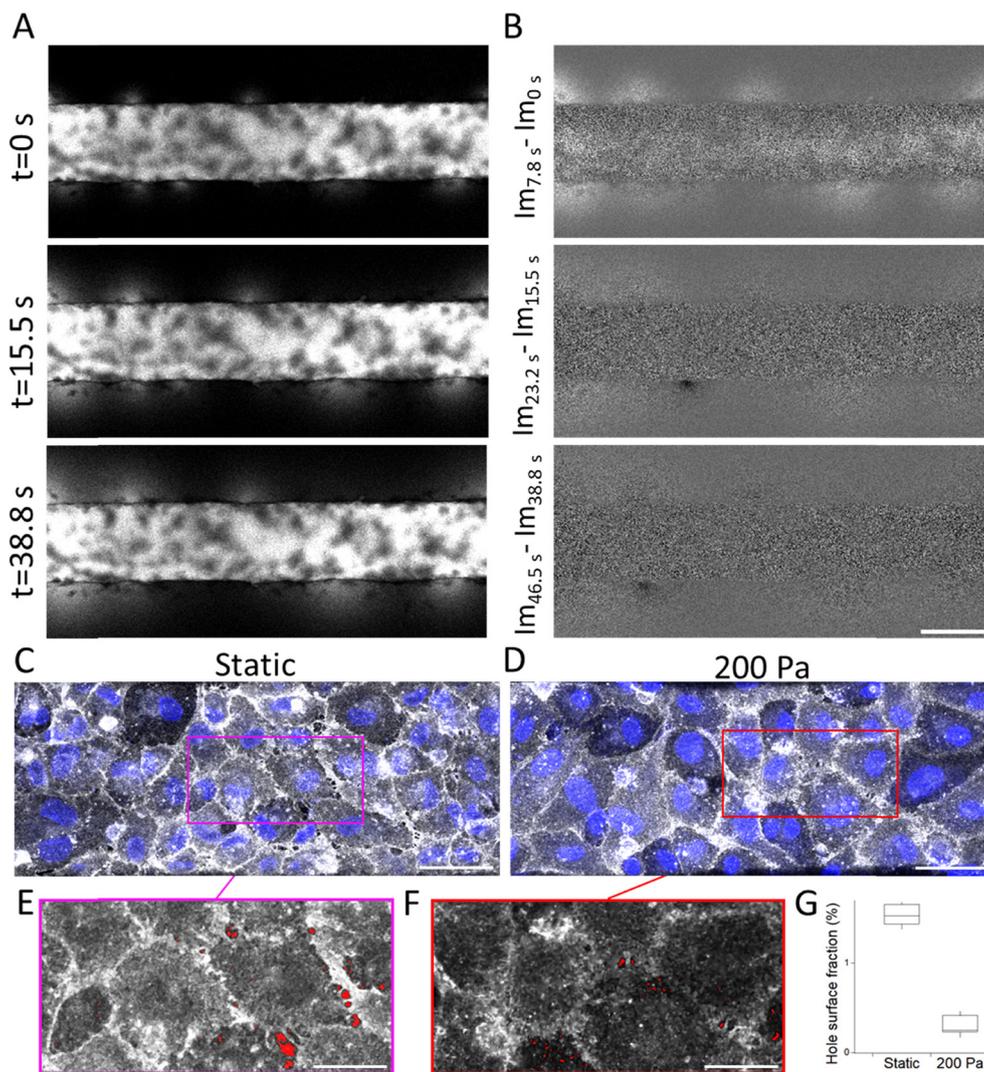


Fig. 3 Paracellular gap closure occurs under pressure. (A) Confocal time series of the macromolecular assay using the 4 kDa dextran (see Fig. S5† for 70 kDa dextran) performed with an intraluminal pressure of 100 Pa using the system described in ref. 36. The inter-frame interval is 7.8 s. (B) The micrographs correspond to the difference between two consecutive images of the time series in (A). (C) MIPs of confocal micrographs of a MV cultured in static conditions; labels: PECAM-1 in gray and nucleus in blue. (D) Same as (C) after the application of an intraluminal pressure of 200 Pa. Note that the spatial distribution of PECAM-1 appears more diffuse, as discussed in the following section. (E) and (F) are zoom-ins extracted from panels (C) and (D) with the detected paracellular gaps in red. (G) Surface fraction of gaps in static and after pressure stimulation. The scale bars correspond to 200 μm in A and B, 50 μm in C and D, and 25 μm in E and F.

Second, we aimed to validate the presence of paracellular pores by characterizing the structure of the endothelial cell monolayer with immunoconfocal microscopy performed in static conditions *vs.* under intraluminal pressure. We used cell-cell junction protein platelet endothelial cell adhesion molecule-1 (PECAM-1) to assess intercellular interactions. Qualitative inspection of MVs revealed occasional gaps, indicated by dark fluorescence signals under static conditions (Fig. 3C). These gaps were less frequent following intraluminal pressure stimulation at 200 Pa for 5 minutes (Fig. 3D). Although the frequency of these gaps varied across fabrication batches, we estimated the gap surface fraction using image analysis (see Materials and methods, Fig. 3E and F) across two separate fabrication batches,

including a total of 4 MVs in static conditions and 6 under mechanical stimulation. The average area fraction of gaps was estimated to be $1.55 \pm 0.12\%$ in static conditions, compared to $0.31 \pm 0.11\%$ after stimulation (Fig. 3G). Altogether, our data strongly support the conclusion that endothelial barriers respond and adapt to intraluminal pressure by closing paracellular gaps.

Intraluminal pressure stimulates actin cytoskeleton reorganization

The actin cytoskeleton plays a central role in regulating endothelial tissue function, as demonstrated by the morphological and orientational changes endothelial cells



undergo in response to mechanical stimulation from fluid shear stress or periodic stretching.^{17,37} We thus characterized the actin cytoskeleton and the patterns of cell–cell interaction proteins in order to clarify their structure after barrier reinforcement. We stained HUVECs with phalloidin and antibodies to the *adherens* junction (AJ) protein vascular endothelial cadherin (VE-Cad), PECAM-1 or the tight junction protein Claudin-5 (see Fig. S6† for VE-Cad and Claudin 5). In the control samples cultured in static conditions, we observed a well-defined cortical actin rim at the basolateral site, indicating a typical cytoskeletal organization in the

absence of pressure (Fig. 4A). We then stimulated MVs with 100 and 400 Pa of intraluminal pressure stimulation for 5 minutes, and fixed the samples immediately after. At 100 Pa, the strain was $17 \pm 2\%$, and this pressure load was sufficient to induce the formation of parallel actin stress fibers spanning the cell interior (Fig. 4B and S6A†). In some instances, we observed alignment of stress fibers between neighboring cells, accompanied by irregular intercellular junction patterns, an arrangement reminiscent of discontinuous AJs (ref. 38, orange arrows in Fig. 4B). Notably, these patterns closely resembled those observed upon

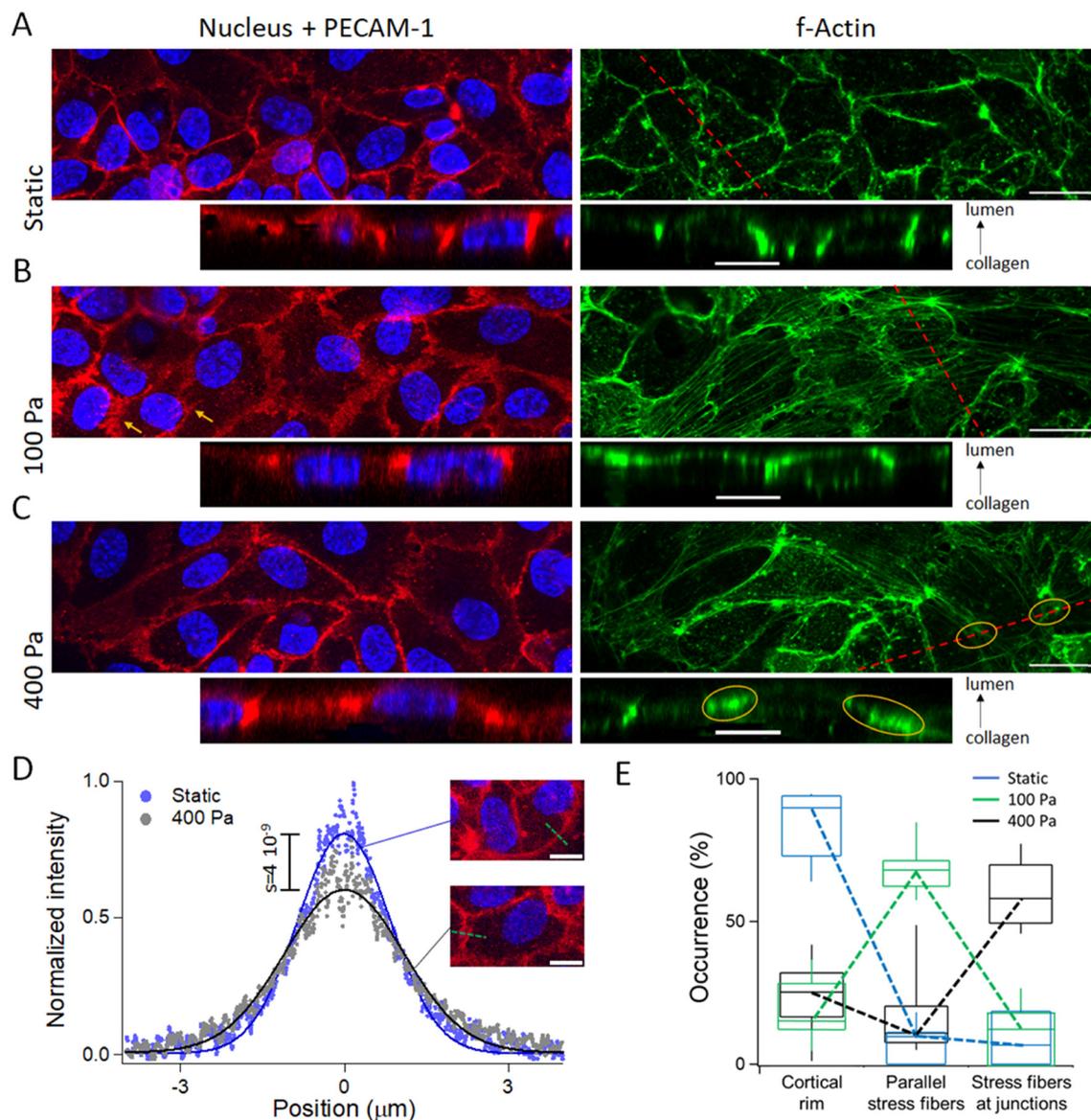


Fig. 4 Barrier enhancement involves actin cytoskeleton reorganization. MIPs and vertical cross-sections from the collagen gel to the lumen along the red dashed lines in static conditions (A) and after the application of 100 Pa (B) or 400 Pa (C); labels: F-actin in green, nucleus in blue, and PECAM-1 in red. The orange arrows indicate discontinuous AJs patterns. The scale bars correspond to 20 μm in the MIPs and 10 μm in the cross-sections (see Fig. S6† for VE-Cad data). (D) 30 integral-normalized intensity profile of the AJs extracted from MIP of PECAM-1 immunofluorescence micrographs are pooled in static and under 400 Pa of intraluminal pressure. The two fluorescence micrographs shown in inset are representative cell patterns, and intensity profiles are typically collected along the green dashed lines. (E) Enumeration of the number of HUVECs with a cortical rim, parallel stress fibers or accumulated stress fibers in static conditions or at 100 and 400 Pa (blue, green, black datasets, respectively). The dashed lines are guides to the eye.



stimulation of endothelial tissues with pro-inflammatory agents, such as tumor necrosis factor or histamine, both of which are associated with hyperpermeability.^{5,39,40} However, unlike these biochemical cues, mechanical stimulation-induced formation of aligned stress fibers and discontinuous AJs did not lead to a degradation of endothelial barrier function. Examination of stress fiber localization revealed their predominant accumulation on the apical surface of the tissue, with a few isolated fibers observed on the basal side (lower panel of Fig. 4B). When intraluminal pressure was increased to 400 Pa, actin stress fibers became concentrated near cell-cell junctions, as indicated by the orange-labeled regions in Fig. 4C. This pattern was also evident in VE-cadherin MIPs (Fig. S6A†). Notably, at 400 Pa, actin stress fibers were exclusively observed on the apical side (Fig. 4C), and discontinuous AJ patterns appeared less frequently. However, the intercellular junctions visualized in MIPs were broader under 400 Pa compared to static conditions, as quantified by the onset of 29% of the full width at half maximum of the Gaussian fit to the PECAM-1 signal (Fig. 4D).

Based on these observations, we identified three distinct cytoskeletal patterns: (i) a cortical rim, (ii) parallel stress fibers, and (iii) accumulated stress fibers. These patterns were counted under static conditions and at pressures of 100 and 400 Pa (blue, green, and black datasets in Fig. 4E), with data averaged across three microvessels, including ~120 cells, per condition. This analysis confirmed the phenotypic shift induced by intraluminal pressure because the prevalence of the three cytoskeletal patterns peaked at 84%, 55%, and 61% for the three different settings in intraluminal pressure. Thus, the intraluminal load stimulates cytoskeletal reorganization, which in turn stabilizes intercellular junctions.

Further, we tested whether barrier reinforcement was dependent on the mitogen-activated protein kinase (MAPK) pathway because cyclic-stretch experiments have shown that mechanical stretch activates this mechanosensory cascade independently of actin fiber reorientation.⁴¹ We used the selective MEK 1/2 inhibitor PD98059 at a concentration of 30 μM for 30 minutes, a dose that is non-cytotoxic but interferes with HUVEC migration,⁴² and then examined the response of MVs to intraluminal pressure. Immunostaining for the phosphorylated form of extracellular signal-regulated kinase (ERK) confirmed that the pressure stimulus activated the MAPK pathway and that this activation was inhibited by PD98059 (Fig. S7A and B†). We also monitored the permeation flow rate for 5 minutes with or without the inhibitor (Fig. S7C†), and did not detect any changes of the barrier reinforcement dynamics. Last, we observed that the density of holes in static conditions decreased similarly after pressure stimulation with or without drug treatment (Fig. S7D†). Consequently, the inhibitor did not affect the response of the endothelial barrier to intraluminal pressure, indicating that the activation of the MAPK pathway was coincident but not associated to the endothelial tissue reinforcement.

Barrier reinforcement is lost upon inhibition of ROCK

The reorganization of the actin cytoskeleton occurred concurrently with endothelial adaptation to intraluminal pressure. To further investigate this process, we aimed to disrupt actin reorganization by inhibiting ROCK—a kinase activated by RhoA that regulates myosin motor activity and cell contractility—with Y27632. This inhibitor has been shown to directly interfere with actin stress fiber formation without disrupting endothelial barrier function at concentrations below 10 μM ;⁴³ at these low concentrations, it even protects the barrier against thrombin-induced degradation.^{44,45} After one day of culture in static conditions, we supplemented the medium with 1 μM of Y27632 and maintained the MVs for another two days. This treatment did not induce any morphological change at the tissue level (Fig. 5A), and it also preserved the characteristic cortical rim pattern in mechanically unstimulated MVs (see more below). We then probed the response of MVs to intraluminal pressure by applying 400 Pa. The initial flow rate after a stimulation of ~1 s was comparable in the control and in the treated sample of 7.6 and 7 $\mu\text{L min}^{-1}$, respectively (gray and green datasets in Fig. 5B). These results demonstrated that the permeability of the barrier was not affected by Y27632 treatment. However, the response diverged noticeably after approximately 10 seconds of actuation with intraluminal pressure. Specifically, the control sample exhibited a characteristic rapid and sustained reinforcement of the barrier function, as shown by the red dashed trend line in Fig. 5B. In contrast, the permeation flow rate in Y27632-treated MVs remained consistently high and unchanged, even after 50 seconds (blue dashed line in Fig. 5B). This flat response indicated that the endothelial barrier lost its ability to actively respond to intraluminal pressure following treatment with Y27632. This conclusion was further supported by measuring the change in permeation flow rate after 100 seconds of actuation. Unlike the untreated MVs (Fig. 2B), the Y27632-treated MVs showed no significant differences in response when compared to dead or fixed cells (Fig. 5C). Taken together, these findings suggest that treatment with 1 μM Y27632 compromises the adaptive reinforcement of endothelial tissue in response to intraluminal pressure.

We next analyzed the actin cytoskeleton and PECAM-1 patterns in the presence of the ROCK inhibitor using immunoconfocal microscopy (Fig. 5D and E). Under static conditions, the actin cortical rim and linear AJs were visible in the MIPs (Fig. 5D), exhibiting patterns similar to the untreated control group (Fig. 4A). However, after exposure to 400 Pa of pressure, we observed significant alterations compared to the control. Specifically, discontinuous actin fibers were detected (Fig. 5E), predominantly distributed along both the basal and apical surfaces of the cells. The AJ patterns appeared irregular, with protrusions extending along typically thin actin fiber tracks (Fig. S8†). Additionally, ovoid gaps between adjacent cells were also detected, suggesting that intraluminal pressure disrupted intercellular junctions



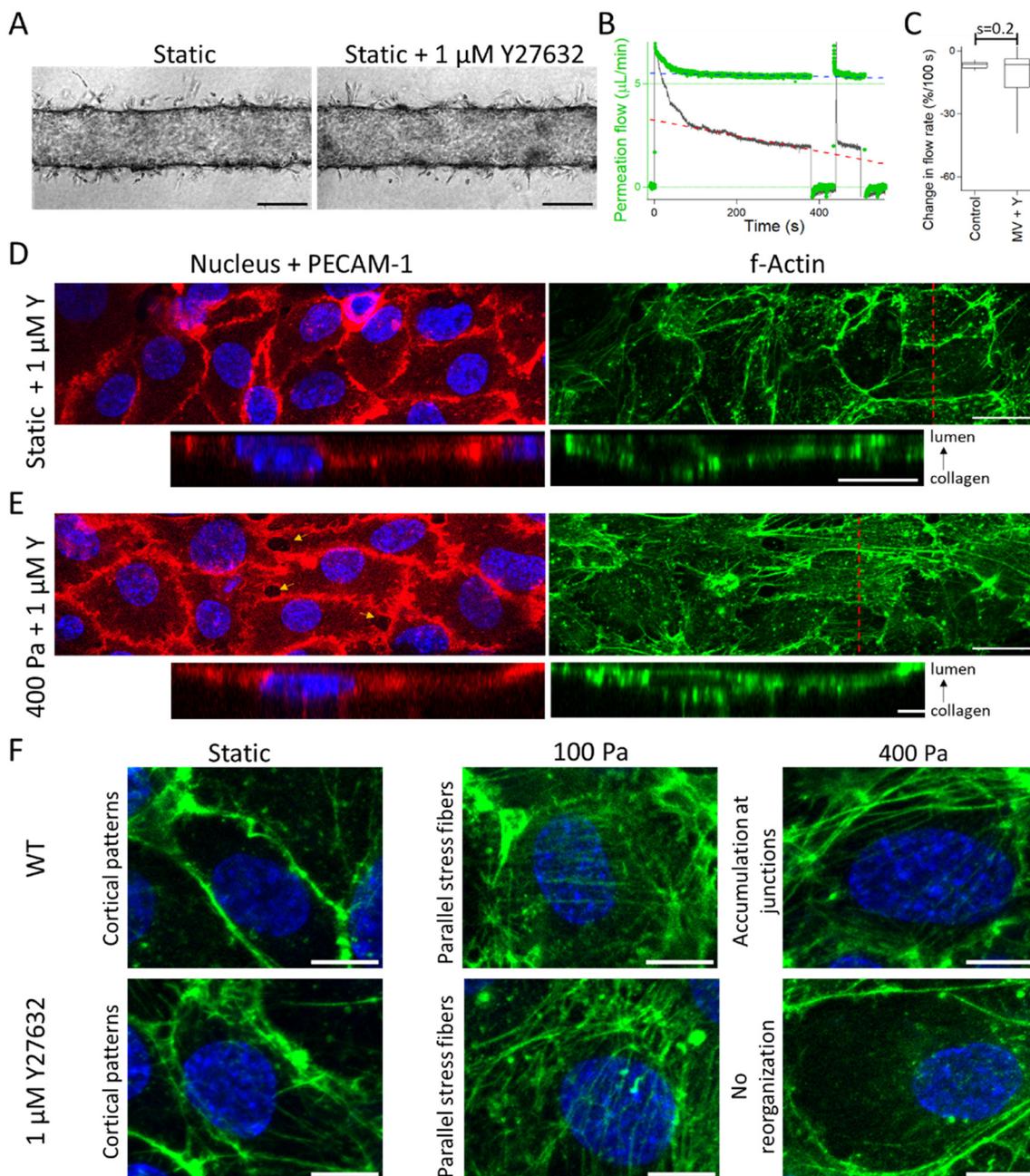


Fig. 5 Rho kinase inhibition disrupts barrier reinforcement. (A) Optical micrographs of MV treated or not with the drug Y27632 in static conditions. (B) The graph shows the permeation flow rate as function of time for a MV treated with Y27632 (green dots) and untreated (gray dots). The temporal sequence of pressure actuation is 400 Pa for 350 s, then 0 Pa for 50 s, and 400 Pa for 60 s. (C) The box plot reports the relative change in permeation flow rate after 100 s of stimulation with fixed or dead cells (same dataset as in Fig. 2B) or in treated conditions. MIPs of MVs treated with Y27632 and vertical cross-sections along the red dashed lines in static conditions (D) and after the application of 400 Pa (E); labels: F-actin in green, nucleus in blue, and PECAM-1 in red (see Fig. S8† for VE-Cad data). The scale bars correspond to 20 μm in the MIPs and 10 μm in the cross-sections. (F) MIPs of the apical surface of HUVECs, with an integration depth of 3 μm , under control and treated conditions. Labels: F-actin in green, nucleus in blue; the scale bars correspond to 10 μm .

(highlighted by orange arrows, Fig. 5E) and contributed to the loss of barrier integrity under treated conditions. In order to further document the loss of cytoskeleton reorganization induced by Y27632, we extracted the actin pattern on the apical side of the tissue in static conditions and after 100 and 400 Pa of pressure stimulation (Fig. 5F). While the actin

patterns under static conditions and at 100 Pa were comparable between the control and treated groups, the reorganization and accumulation of stress fibers at junctions observed at 400 Pa in controls was completely absent in Y27632-treated MVs. These findings collectively indicate that MVs with impaired cytoskeletal reorganization cannot



effectively withstand intraluminal stress, resulting in the formation of paracellular gaps and a compromised barrier function.

Conclusion

Using an assay to dynamically measure the permeation flow across endothelial cell monolayers, we report that HUVEC-based MVs respond to intraluminal pressure stimulation in seconds by reinforcing their barrier function. We show that this reinforcement is explained by the active closure of paracellular holes. We also demonstrate that this adaptive response can be challenged by intraluminal pressures greater than 1500 Pa, and uncover a dynamic equilibrium of opening and closing events at this threshold. Although the vasculature model used in this study is simplified regarding the extracellular matrix protein composition and perivascular cell integration, we note that this threshold is comparable to the physiological blood pressure in capillaries, approximately 2000 Pa.⁴⁶ This finding suggests that endothelial cells alone have a strong capacity to withstand the constraints imposed by blood pressure.

Our data further highlight the crucial role of actin cytoskeleton reorganization in reinforcing barrier function under mechanical stress. At a low intraluminal stress of 100 Pa, we observed parallel actin stress fibers spanning the cell interior, accompanied by zig-zag patterns of cell–cell adhesion proteins. These patterns resemble the perpendicular stress fibers at AJs commonly associated with inflammatory conditions.³⁸ While this cytoskeletal organization may be prone to destabilization, it undergoes a significant rearrangement at 400 Pa, where parallel actin fibers localize at cell junctions. Given that intercellular junctions in HUVECs are stabilized by the pushing forces of the branched actin network emerging from cortical actin bundles,⁴⁷ we propose that the accumulation of actin stress fibers at junctions enhances cell–cell association forces. This process effectively seals potential gaps in the tissue. Interestingly, this model suggests a counterintuitive mechanism in which increased intraluminal pressure, which stretches the tissue, triggers a corresponding rise in compressive tension between cells, thereby strengthening the barrier.

Finally, it is tempting to propose that the reinforcement of endothelial barriers plays a key role in regulating interstitial fluid flow and preventing the onset of edema. Increases in intraluminal pressure, commonly seen in hypertension or during physical exercise,⁴⁸ raise the pressure within blood vessels, including capillaries. When capillary pressure rises, permeation flow also increases, heightening the risk of fluid accumulation in tissues. Prolonged hypertension has been shown to induce structural changes in blood vessels, such as vessel wall thickening, which helps resist elevated pressure and limit interstitial fluid leakage.^{49,50} Aside from long-term vascular remodeling, barrier reinforcement offers a rapid and durable mechanism for regulating permeation flow,

potentially maintaining tissue homeostasis despite sudden changes in blood pressure.

Materials and methods

Cell culture and reagents

All chemicals were purchased from Sigma Aldrich, unless mentioned. Studies were performed with primary human umbilical vein endothelial cells (HUVEC; Catalog #C2519A, Lot #0000699241; Lonza, Basel, Switzerland) that were cultured in endothelial cell growth medium-2 BulletKit (EGM-2; Lonza). They were frozen in liquid nitrogen at passage 4 to 5, thawed and cultured for three days in culture dishes, and then used to load MV chips. The drug PD98059 was dissolved at 50 mM in anhydrous DMSO and diluted in culture medium to 30 μ M. Y27632 was purchased at 1 mM in H₂O and diluted in culture medium at 1 μ M.

MV fabrication

MV were fabricated in polydimethylsiloxane (PDMS)-based chips (25 mm \times 25 mm \times 5 mm: width \times length \times height), as developed by Matsunaga and collaborators.³⁶ The protocol includes an additional PDMS–collagen cross-linking step to avoid leaks at the PDMS/collagen interface during diffusion and pressure assays. The protocol started by O₂ plasma treatment of PDMS chips and acupuncture needles of 200 μ m (No. 08, J type; Seirin, Shizuoka, Japan) for one minute (basic plasma cleaner; Harrick Plasma, Ithaca, NY, USA). The PDMS chips and needles were then placed together in a vacuum reactor with 50 μ L of aminopropyl-triethoxysilane, and left at 0.1 mbar and room temperature for 30 minutes. Needles were then soaked in 1% (w/v) bovine serum albumin and dried. The chips were treated with 50 μ L of 2.5% glutaraldehyde for one minute, then thoroughly rinsed with water and dried. The collagen solution was subsequently prepared on ice by mixing Cellmatrix® type I-A collagen solution (Nitta Gelatin, Japan), 10 \times Hanks' buffer, and 10 \times collagen buffer (volume ratio 8 : 1 : 1) following manufacturer's protocol (final collagen concentration: 2.4 mg mL⁻¹). We poured 30 μ L of this ice-cold solution into the chip, and inserted the coated needle. The resulting devices were incubated at 37 $^{\circ}$ C for 15 min to induce collagen reticulation, and the needles were withdrawn to form a hollow channel. The chips were left in PBS at least overnight before cell seeding, and the holes for needle incorporation were sealed with unreticulated PDMS to prevent leaks.

Just prior to loading in the chips, HUVEC cells were harvested and resuspended at a density of 10⁷ cells per mL in the medium supplemented with 3% (m/v) dextran (500 kDa) in order to increase the viscosity and favor the attachment of HUVECs with the collagen gel. 50 000 cells were loaded at each opening of the channel, and let to attach to the collagen scaffold at 37 $^{\circ}$ C for 10 minutes. We eventually added 1 mL of fresh medium, and refreshed it every other day.



Microfluidics to monitor the permeation flow

MV were placed on an aluminum support of $30 \times 60 \text{ mm}^2$ with a set of tapped holes that enabled us to tightly hold 3D printed reservoirs fabricated by stereolithography (Expert Material Series, NSS, Japan). This system was connected to chromatography PEEK tubing of outer diameter $1/32''$ and inner diameter $380 \mu\text{m}$. They were connected to two microheater-based flow sensors (Fluigent, model S) through UNF 6–40 port, and then to pressurized reservoirs of 2 mL actuated with a 25 mbar pressure controller (MFCS, Fluigent). These reservoirs were mounted on a manually adjustable vertical stage, allowing precise control of fluid levels to eliminate hydrostatic flows. Prior to every experiment, the microfluidic system was first filled with 70% ethanol solutions, then extensively washed with sterile medium, before its assembly on the MV chip. The mounted MV chip was eventually placed in a miniature incubator controlled in temperature and CO_2 that was placed on an inverted bright field microscope (Zeiss AxioObserver). Note that the hydraulic resistance of the microfluidic set up of $15 \text{ Pa } \mu\text{L}^{-1} \text{ min}^{-1}$ was three and thirty times smaller than that of collagen gels and MV, respectively.

For an intraluminal pressure of 400 Pa, the permeation flow Q ranged from approximately 1 to $4 \mu\text{L min}^{-1}$, and it was associated to a Poiseuille flow within the MV lumen.⁵¹ Modeling the lumen as a cylindrical tube with a diameter of $200 \mu\text{m}$ and a length of 3 mm , we estimated the typical hydraulic resistance of the MV to be $5.5 \times 10^{10} \text{ Pa s m}^{-3}$. From this, we inferred that the pressure drop associated with the permeation flow was around 2 Pa, which is negligible compared to the intraluminal pressure of 400 Pa. This finding indicates that variations in permeation flow induced by the endothelium do not affect the strength of intraluminal pressure stimulation. Furthermore, this calculation enabled us to determine the residual shear stress induced by the permeation flow, which was 0.02 Pa (or 0.2 dyn cm^{-2}). This shear stress magnitude was four orders of magnitude lower than the intraluminal pressure and at least one order of magnitude smaller than the levels observed in blood vessels⁵² or 2D endothelial cell cultures.⁹

Macromolecular assay, fixation and immunostaining

For the macromolecular assay, we used the same aluminum support with a set of tapped holes and a different reservoir that maintained a constant intraluminal pressure through a 1 cm height difference between the inlet and outlet of the MV.³⁶ This reservoir was set up two minutes before introducing a solution containing 4 kDa and 70 kDa dextran, each at $5 \mu\text{g mL}^{-1}$ and labeled with fluorescein and rhodamine, respectively. We then acquired a confocal time series at 488 nm and 555 nm using the LSM 700 confocal microscope (Carl Zeiss) equipped with a $10\times$ objective (numerical aperture of 0.4) and a pinhole setting that recorded the signal over a $12 \mu\text{m}$ thickness. The inter-frame interval was 7.8 seconds.

The immunostaining protocol started by MV fixation using 4% (w/w) paraformaldehyde for 30 min, and then thoroughly rinsing with PBS. Permeabilization was performed with 0.5% Triton X-100 for 10 min at room temperature. Blocking with 1% BSA was performed overnight at $4 \text{ }^\circ\text{C}$. Cells were then incubated overnight at $4 \text{ }^\circ\text{C}$ with the primary antibodies VE-Cadh (rabbit mAb, D87F2, Cell Signaling Technology, 1:200), PECAM-1 (mouse mAb, GA610, Dako, 1:200), Claudin-5 (rabbit mAb, ab15106, Abcam, 1:200) or phosphorylated ERK1/2 (rabbit mAb, 4370T, Cell Signaling Technology, 1:200) diluted in blocking solution. After washing, cells were incubated for 2 h with the corresponding secondary antibodies (1:400), Alexa Fluor-488 Phalloidin (1:800), and Hoechst 33342 (1:1000). Labeled samples were washed and stored at $4 \text{ }^\circ\text{C}$ until imaging. Confocal images were captured with the LSM 700 confocal microscope equipped with a $40\times$ water immersion objective (numerical aperture of 1.2). We used a pinhole of 1 airy unit for the three lasers of 405, 488, 555 nm, a pixel size of $0.156 \mu\text{m}$, and set the vertical increment of confocal stacks to $0.5 \mu\text{m}$.

Image analysis and statistical test

Image analysis, including MIP and orthogonal view, was performed with ImageJ. Hole detection was performed by thresholding images (minimum method) after application of the “enhanced local contrast” function (CLAHE). All the plots were obtained with Igor Pro (version 5.0). Statistical significance was determined using the Student's t -test, that was reported with the parameter s (s -values < 0.05 were considered statistically significant; n.d., not different). Data were expressed as mean \pm standard error.

Qualitative estimation of MV permeability and fluid flow simulation methodology

The permeation flow velocity is related to the pressure gradient by the following relationship:

$$v = \frac{\kappa \text{d}P}{\mu \text{d}x} \quad (1)$$

with κ the permeability and μ the viscosity of the culture medium. We assume a 1D geometry and a linear pressure gradient, implying that the pressure gradient is P_{in}/H_c with H_c the collagen gel height of 1 mm. The endothelial barrier confines the pressure gradient in the cell layer h . Using the subscripts c and e for the collagen gel and the endothelial cell layer, respectively, we deduce that:

$$\kappa_e = \kappa_c \frac{Q_c h}{Q_c H} \quad (2)$$

We also performed simulations of the permeation flow using finite element simulations performed with COMSOL Multiphysics 6.0, following the approach in ref. 25. We used the Darcy law module to compute the flow velocity



field. We set the tube radius r_0 to 100 μm and placed it 280 μm above the glass coverslip. The barrier thickness δ was set to 2 μm . Three distinct domains were defined: (i) the collagen matrix with a porosity of 0.95, (ii) the central lumen free of collagen, and (iii) the cell barrier as characterized by a permeability of $5 \times 10^{-18} \text{ m}^2$. The permeability of collagen gels was set to $5 \times 10^{-14} \text{ m}^2$. Boundary conditions consisted of no flux at the edges of the gel except for an open boundary at the top surface and tube interface. The pressure was set to 100 Pa on the two lateral sections of the MV and to null at the top interface of the gel.

Data availability

All data supporting the findings of this study are included in the manuscript and its ESI.†

Conflicts of interest

The authors declare no conflict of interest.

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