Caveolin-1–eNOS signaling promotes p190RhoGAP-A nitration and endothelial permeability

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Endothelial barrier function is regulated by adherens junctions (AJs) and caveolae-mediated transcellular pathways. The opening of AJs that is observed in caveolin-1−/− (Cav-1−/−) endothelium suggests that Cav-1 is necessary for AJ assembly or maintenance. Here, using endothelial cells isolated from Cav-1−/− mice, we show that Cav-1 deficiency induced the activation of endothelial nitric oxide synthase (eNOS) and the generation of nitric oxide (NO) and peroxynitrite. We assessed S-nitrosylation and nitration of AJ-associated proteins to identify downstream NO redox signaling targets. We found that the GTPase-activating protein (GAP) p190RhoGAP-A was selectively nitrated at Tyr1105, resulting in impaired GAP activity and RhoA activation. Inhibition of eNOS or RhoA restored AJ integrity and diminished endothelial hyperpermeability in Cav-1−/− mice. Thrombin, a mediator of increased endothelial permeability, also induced nitration of p120-catenin–associated p190RhoGAP-A. Thus, eNOS-dependent nitration of p190RhoGAP-A represents a crucial mechanism for AJ disassembly and resultant increased endothelial permeability.

Introduction

The endothelium lining all blood vessels plays an essential role in the maintenance of tissue fluid homeostasis (Mehta and Malik, 2006). It finely controls the permeability of the vessel wall barrier to plasma proteins and small solutes via transcellular, caveolae-mediated transcytosis and paracellular, inter-endothelial junctional (IEJ) routes. Whereas IEJs are permeant to molecules with radii of 3 nm or less, caveolae-mediated vesicle trafficking transports different-sized plasma proteins across the vascular barrier (Michel, 1996). The interrelationships between paracellular and transcellular pathways are not well understood. The loss of caveolin-1 (Cav-1) results in not only disappearance of caveolar structures but also destabilization of IEJs and formation of inter-endothelial gaps (Schubert et al., 2002; Miyawaki-Shimizu et al., 2006). We surmised from these previous studies that opening of the paracellular route for albumin transport in Cav-1−/− mice might represent a general mechanism by which Cav-1 influences IEJs and regulates junctional integrity.

In addition to the transport of plasma proteins and lipids, caveolae compartmentalize signaling molecules within lipid raft microdomains (García-Cardeña et al., 1997). Cav-1 interacts with a variety of signaling molecules such as EGF receptor (EGFR), c-Src, phosphoinositide 3-kinase (PI3K), GTP-activating protein (GAP) p190RhoGAP-A was selectively nitrated at Tyr1105, resulting in impaired GAP activity and RhoA activation. Inhibition of eNOS or RhoA restored AJ integrity and diminished endothelial hyperpermeability in Cav-1−/− mice. Thrombin, a mediator of increased endothelial permeability, also induced nitration of p120-catenin–associated p190RhoGAP-A. Thus, eNOS-dependent nitration of p190RhoGAP-A represents a crucial mechanism for AJ disassembly and resultant increased endothelial permeability.
Cav-1 deficiency induces eNOS activation and AJ destabilization in endothelium. [a] Phosphorylation of eNOS at S\textsuperscript{1177} and Akt-1 at S\textsuperscript{473} in Wt and Cav-1\textsuperscript{−/−} MLVECs; n = 3. Molecular mass standards are indicated next to the gel blots in kilodaltons. [b] Basal NO generation was measured in the absence or presence of l-NNA or NPA and 1400W in combination. Arrow, l-arginine replenishment. Scale bars are shown for 100 nM NO and 10 min. Bar plot, NO accumulation in a media over 20 min; mean ± SEM (error bars); *, P < 0.05 as compared with Wt control; n = 6. [c] Wt and Cav-1\textsuperscript{−/−} endothelial monolayers stained for VE-cadherin, β-catenin (green in overlays), F-actin (red), and nuclei (blue). Inter-endothelial gaps in Cav-1\textsuperscript{−/−} monolayers are indicated by arrowheads. Bar, 10 µm. [d and e] Accumulation of β-catenin at AJs was expressed as a mean pixel intensity of threshold area shown in d (threshold above intracellular background is in orange). [f] Area of inter-endothelial gaps was determined using the same set of images as in e; n = 17. Black circles and error bars in e and f indicate mean and SEM, respectively; *, P < 0.01 as compared with Wt control.

Results and discussion

We first determined basal eNOS activity in murine lung vascular endothelial cells (MLVECs) isolated from wild-type (Wt) and Cav-1\textsuperscript{−/−} mice (Fig. S1). We observed an approximately two-fold increase in eNOS phosphorylation at S\textsuperscript{1177} (Fig. 1 a), the Akt-1 phosphorylation site known to regulate enzymatic activity of eNOS (Dimmeler et al., 1999). Phosphorylation of Akt-1 at S\textsuperscript{473} was also increased (Fig. 1 a), which indicates marked Akt-1 activation in Cav-1\textsuperscript{−/−} endothelium. Basal nitric oxide (NO) generation determined by l-arginine replenishment was fourfold greater in Cav-1\textsuperscript{−/−} MLVECs (Fig. 1 b). This finding signaling, which is normally held in abeyance by binding of eNOS to Cav-1 (Bucci et al., 2000), regulates IEJ integrity and thereby the permeability of the endothelial barrier.
is consistent with the reported increased plasma NO concentration in Cav-1–deficient mice (Miyawaki-Shimizu et al., 2006). Nitro-L-arginine (L-NNA) treatment at an inhibitory concentration for nNOS, and eNOS prevented NO generation, whereas treatment with 1400W and N-propyl-L-arginine (NPA) in combination (at inhibitory concentrations for iNOS and nNOS) had a marginal effect (Fig. 1 b). We concluded that Cav-1 deficiency in the endothelium resulted in Akt-1 and eNOS activation and an augmented NO production, which is in agreement with the role of Cav-1 in inhibiting eNOS activity in endothelium (García-Cardeña et al., 1997).

Next we determined the effects of Cav-1 deficiency on adherens junction (AJ) integrity, the primary adhesive complex in the continuous endothelium responsible for regulating junctional permeability. We did not observe any significant change in the expression of AJ proteins in Cav-1−/− MLVECs compared with Wt cells (Fig. S1 b), which is in contrast to the difference reported for the brain endothelial cells (Song et al., 2007). This inconsistency might reflect tissue specificity (i.e., continuous vs. the highly restrictive brain endothelial barrier). Whereas AJs play a primary role in the mechanism of AJ integrity in the continuous endothelium, they are secondary in maintaining brain–blood barrier function (Nitta et al., 2003). Integrity of AJs was assessed by immunofluorescent staining for VE-cadherin, the homotypic adhesive protein of AJs (Corada et al., 1999), and β-catenin. Both proteins accumulated at the cell border; however, clustering of VE-cadherin at the cell surface of Cav-1−/− endothelium was markedly reduced (Fig. 1 c). VE-cadherin formed discrete patches on the Cav-1−/− endothelial cell surface; in contrast, VE-cadherin was organized as bands encircling the Wt endothelial cells. Accumulation of β-catenin at AJs, also used to assess AJ integrity, was significantly reduced in Cav-1−/− endothelium (Fig. 1, c–e). Cav-1−/− monolayers were characterized by the presence of many inter-endothelial gaps (Fig. 1, c and f). L-NNA treatment partially restored the accumulation of β-catenin at AJs and resealed the gaps in Cav-1−/− monolayers (Fig. 1, c and e). Although AJs remained somewhat fragmented, they appeared to assemble as bands connecting adjacent cells. Interestingly, the PI3K p110α isoform, which acts upstream of the Akt-1–eNOS signaling node, was shown to negatively regulate the integrity of AJs during inflammation by inducing the association of VE-cadherin with Pyk2 (Cain et al., 2010). It is likely therefore that Cav-1 deficiency is also accompanied by PI3K activation, and the destabilization of VE-cadherin adhesion seen in the Cav-1−/− monolayers might result from independent parallel signaling events. Nevertheless, our data suggest that Cav-1 modulates permeability of endothelial barrier, in part, through its negative regulation of eNOS activity.

To address mechanisms of eNOS modulation of AJ integrity, we determined the effects of NO on modification of AJ proteins. NO can lead to S-nitrosylation of cysteine, the covalent attachment of a NO group to the thiol side chain (Lima et al., 2010), and thus influence protein function. S-nitrosylation of β-catenin was shown to facilitate destabilization of AJs in response to VEGF signaling (Thibeault et al., 2010). Therefore, we determined S-nitrosothiol (SNO) content of β- and p120-catenins and of p190RhoGAP-A (referred to as p190A), a binding partner of p120-catenin (Wildenberg et al., 2006). We reasoned that the apparent organization of actin cytoskeleton into stress fibers in Cav-1−/− MLVECs (Fig. 1 c) might result from insufficient control of RhoA activity. p190A, a GTPase-activating protein (GAP), facilitates RhoA-GTP hydrolysis (Tatsis et al., 1998) and is a major regulator of RhoA activity at the level of AJs (Mammoto et al., 2007). SNO content, determined by the biotin-switch assay (Jaffrey and Snyder, 2001), of AJ proteins was indistinguishable between Cav-1−/− and Wt endothelia, although in a control experiment we could detect SNO of β-catenin in VEGF-stimulated endothelial cells (Fig. 2 a). This finding raises the possibility that eNOS-redox signaling regulates integrity of AJs by other means.

When NO is accompanied by O2•− generation, it forms peroxynitrite. The diffusion-controlled reaction of NO with O2•− is a likely source of peroxynitrite in cells and in vivo (Beckman et al., 1990; Jouzdheuil et al., 2001). The resulting peroxynitrite rapidly decomposes in the presence of ubiquitous CO2 to produce nitrate (NO3−) as a stable end product as well as carbonate (CO3•−) and nitrogen dioxide (NO2•) radicals (Bonini et al., 1999; Augusto et al., 2002), whose concerted action redirects peroxynitrite reactivity toward nitration of protein tyrosine residues (Santos et al., 2000). A direct measurement of peroxynitrite in cells is precluded by the extremely short lifetime of peroxynitrite at physiological pH and by the presence of metals, reductants, and CO2. Because peroxynitrite formation depends on both O2•− and NO and produces NO3− as the only stable product after oxidant decomposition, the use of superoxide scavengers followed by measurement of nitrite (NO2−) accumulation provides an assessment of peroxynitrite formation (Bonini et al., 2002). Agents such as polyethylene glycol-superoxide dismutase (PEG-SOD) and 4-hydroxy-2,6,6-tetramethylpiperidinolxylo (TEMPOL) in adequate concentrations outcompete NO for O2•−, thus favoring NO autoxidation and NO2− accumulation (Hoffman et al., 2003). Here we used TEMPOL/SOD-dependent NO2− accumulation to assess peroxynitrite formation. Both scavengers had marginal effects on accumulation of nitrite resulting from basal NO production in WT cells (Fig. 2 b); in contrast, accumulation of nitrite was significantly higher in Cav-1−/− cells (Fig. 2 b). Consistent with this observation, Cav-1 deficiency was accompanied by a twofold increase in O2•− generation (Fig. 2 c), as measured by formation of 2-hydroxyethylidium (EOH), a product of dihydroethidium (DHE) oxidation. Furthermore, adding tetrahydrobiopterin (BH4) to these cells, but not L-arginine, restored O2•− generation to the level seen in WT cells (Fig. 2 c), which suggests that BH4 deprivation-mediated uncoupling eNOS from NO generation is the primary mechanism of O2•− generation in the Cav-1−/− endothelial cells. We thus concluded that up-regulation of eNOS activity in Cav-1−/− endothelium results in an augmented generation of both NO and O2•− and thereby the formation of peroxynitrite.

We next determined the nitration of AJ proteins by probing immunoprecipitated proteins with anti-nitrotyrosine antibody (3-N). We failed to detect any change in nitration of either β- or p120-catenins in Cav-1−/− endothelial monolayers (Fig. 2 d); however, nitration of p190A was significantly increased (Fig. 2 e).
We also found marked reduction in p190A nitration in Cav-1−/− endothelial cells pretreated with TEMPOL and in MLVECs isolated from lung vessels of Cav-1/eNOS double knockout mice (Fig. 2 e). Nitration of p190A also appeared to be reversible because scavenging of O2•− significantly reduced the nitrated protein levels, although the p190A expression level itself remained unchanged (Fig. 2 e). We speculate that nitration, which in some cases targets proteins for proteolytic degradation (Grune et al., 1998), does not regulate the metabolism and turnover of p190A. The dogma of irreversible nitration of proteins is under debate, and it is also now increasingly evident that the action of peroxynitrite might be specific (Aulak et al., 2004; Görg et al., 2007).

Nitrination compromises protein function by altering tyrosine phosphorylation (Gow et al., 1996). The ability of p190A to hydrolyze RhoA is regulated by c-Src and FAK phosphorylation of p190A (Chang et al., 1995; Holinstat et al., 2006). Therefore, we tested whether nitration alters tyrosine phosphorylation of p190A, which could thereby inhibit its GAP activity. We found a marked decrease in p190A phosphorylation in Cav-1−/− MLVECs (Fig. 3 a). We next addressed whether Y1105, the phosphorylation site critical for p190A activity (Roof et al., 1998), might be targeted for nitration. Mutation of Y1105, but not Y1087, to phenylalanine (Y→F) prevented the chemically induced nitration of p190A by 3-morpholinosydnonimine (SIN-1; Fig. 3 b). SIN-1 also induced nitration of endogenous and transiently expressed Wt p190A but not of p190RhoGAP-B (Fig. 3 b).

Down-regulation of p190A activity results in RhoA activation (Vincent and Settleman, 1999), which may increase junctional permeability (Holinstat et al., 2006). We thus determined whether nitration of p190A results in RhoA activation in the Cav-1−/− endothelium. For this, we assessed RhoA activity using the FRET-based RhoA biosensor (Pertz et al., 2006). Ratio images revealed a twofold increase in RhoA activity in Cav-1−/− monolayers compared with Wt (Fig. 3, c and d). We detected uniform RhoA activity from the edge of the cell inwards with
To demonstrate the causal link between RhoA activity and destabilization of AJs, we attempted to restore integrity of AJs in Cav-1−/− MLVECs by inhibiting RhoA. Treatment of Cav-1−/− endothelium with the Rho inhibitor C3-transferase significantly reduced phosphorylation of myosin light chain (MLC), a very little and infrequent accumulation at cell edges and along extensions (Fig. 3 c), which is consistent with the distribution of active RhoA in other nonmigrating cells (Pertz et al., 2006). Notably, we observed a fourfold increase in RhoA activity by the pull-down of RhoA-GTP with Rhotekin beads (Fig. 3 e).
To address the role of Cav-1–eNOS interaction in the mechanism of increased paracellular permeability in vivo, we recorded fluid shift in isolated-perfused murine lungs by the gravimetric method in response to a standard perturbation of the trans-endothelial oncotic pressure gradient ($\pi_c - \pi_i$; Fig. 4 d).

Elevating perfusate albumin from 0 to 10 g/100 ml caused significantly greater dehydration of lungs of Cav-1$^{-/-}$ mice than in Wt control (Fig. 4 d), which indicates much faster equilibration of fluid across the endothelium devoid of Cav-1 expression. The lung weight gain (during the 40–50th min of the experiment) was fit to a double exponential function characterizing the fast and slow filtration processes (Table S1). The filtration rate calculated downstream effector of RhoA/ROCK, in Cav-1$^{-/-}$ cells (Fig. S2). It also rescued $\beta$-catenin accumulation at AJs and restored permeability of Cav-1$^{-/-}$ monolayers to the Evans blue albumin (EBA) tracer (Fig. 4, a–c). Treatment of Cav-1$^{-/-}$ endothelium with Cav-1 scaffold domain (CSD)–antennapedia (AP) fusion peptide, which binds to and suppresses eNOS catalytic activity (Bucci et al., 2000), superoxide scavenger TEMPOL, and l-NNA as well as Cav-1/eNOS deficiency (DKO) also restored the integrity of AJs and the permeability of Cav-1$^{-/-}$ monolayers (Fig. 4, a–c). We concluded therefore that nitration of p190A provides a reversible switch mechanism regulating RhoA activity and thereby endothelial barrier integrity.

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The NO redox endothelial permeability-increasing mechanism described in this paper may contribute to deregulation of tissue fluid homeostasis during inflammation, which is known to be accompanied by loss of p190A activity and activation of RhoA (Mammoto et al., 2007). Therefore, we determined whether stimulation of endothelial cell monolayers with a pro-inflammatory mediator such as the serine protease thrombin induces p190A nitration. Thrombin proteolytically cleaves and from the initial slope of the slow exponential component was significantly greater in Cav-1−/− (5.5 ± 0.6 µl/min; n = 5) than in Wt (4.1 ± 0.3 µl/min; n = 9) lungs (Fig. 4 c). Perfusion with AP-CSD peptide that reanneals open IEJs (Fig. 4 a) restored transvascular fluid filtration in Cav-1−/− lungs to the Wt values (Fig. 4, d and e). We infer therefore that Cav-1 regulates the permeability of the paracellular or AJ pathways mainly by functioning to restrain the eNOS activity.

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activates the protease-activated receptor PAR-1 on the endothelial surface, leading to increased intracellular calcium concentration and activation of eNOS (Motley et al., 2007). Challenge of human pulmonary artery endothelial cells (HPAECs) with α-thrombin resulted in translocation of eNOS to AJs and a transient interaction between eNOS and p190A and nitration of p190A (Fig. 5, a–d). Nitrated p190A also remained bound to p120-catenin (Fig. 5 d). Thus, we suggest that the signaling mechanism responsible for increased vascular permeability described in this study may be important in inflammatory diseases, and hence we propose it represents a novel anti-inflammatory therapeutic target.

In conclusion, our study demonstrates that paracellular permeability of the endothelium is finely regulated by constitutive Cav-1–dependent inhibition of eNOS activity (Fig. 5 e). Deficiency in Cav-1 or activation of eNOS in conjunction with O•− production results in nitration of p190A and the concomitant activation of RhoA, which induces AJ destabilization and increased endothelial permeability. Because of the causal relationship of eNOS-specific signaling to the impairment of p190A activity, this mechanism induces RhoA activation in endothelial cells. In fibroblasts, however, Cav-1 depletion induces Src-dependent activation of p190A, resulting in loss of cell polarity and impaired wound healing (Grande-García et al., 2007). In contrast, in endothelial cells, Cav-1 knockdown increases directional migration (Gonzalez et al., 2004); thus, our findings help to inform the unique role of Cav-1 expression on p190A activity in endothelial cells and how Cav-1 regulates the permeability of the endothelial junctional barrier.

Materials and methods

Antibodies and reagents
We used anti-HA (clone 16B12; Covance), anti-phosphotyrosine (PY20), and PECA1-MAb (BD); anti-VE-cadherin, anti–β-actin, anti–β3, and anti-p120 catenin; anti-RhoA, anti-eNOS, and anti–Cav-1 (Santa Cruz Biotechnology, Inc.); anti-p190RhoGAP (Sigma-Aldrich); anti-nitrotyrosine, anti-Akt, and anti–Mlc 2; and phospho-eNOS-Ser 1177, phospho-Akt-Ser 473, and phospho–RhoA-Ser 199 (Cell Signaling Technology). Secondary Abs were HRP-conjugated donkey anti–mouse, anti–rabbit, and anti–goat (Jackson ImmunoResearch Laboratories, Inc.); TRITC, FITC, and Cy-5-conjugated donkey anti–mouse, anti–goat, and anti–rabbit (Jackson ImmunoResearch Laboratories, Inc.), Alexa Fluor 532 phalloidin, and DAPI (Sigma-Aldrich).

We used protein A/G beads (Santa Cruz Biotechnology, Inc.), DHE, and Dynabeads M-450 (Invitrogen). We also used a Rho activation assay kit and Rho inhibitor C3 transferase (Cytoskeleton), SIN-1 (EMD), NPA, i-NNA, and 1400W (Cayman Chemical), PEG-SOD, TEMPO, 2 µg/ml Rho inhibitor C3 transferase, and 6 µM AP-CPD peptide for 2 h. 70–80% confluent monolayers were transfected using FuGENE HD (Roche) and Lipofectamine 2000 (Invitrogen) transfection reagents in accordance with the manufacturer’s instructions.

Expression constructs, primers, and RT-PCR
HA-tagged rat p190A and p190B, Y1105F, and Y1087F mutants of p190A were gifts from S.J. Parsons (University of Virginia, Charlottesville, VA); the RhoA biosensor was from K. Hahn (University of North Carolina at Chapel Hill, Chapel Hill, NC). The mRNA was assessed by quantitative real-time PCR (Rajasingh et al., 2008). Primer and probe sequences were as follows: Cav-1 forward, 5′-CCAGCATCTGCTTGGGTAAT-3′; and reverse, 5′-TGTTCC- AGCCATCCCCGTA-3′; and reverse, 5′-ACCATAGGGAGGCTTCTC-3′; eNOS forward, 5′-TGCGCAGGTAGATCATG-3′; and reverse, 5′-CTTCAG- GCCCTCTGTGGT-3′; VE-cadherin forward, 5′-GTTGATGAGCCCCC- TGCTC-3′; and reverse, 5′-CAAGGCTTTCGCTGGTTCTT-3′; and forward, 5′-GGCTGT- CATGCTTGGACGCAC-3′. The relative mRNA expression was normalized to the 18S gene (Applied Biosystems).

Cell culture, treatments, and transfection
Cav-1-1/− and matching control mice of C57Bl/6 background were obtained from The Jackson Laboratory (JAX mice and services). Cav-1/eNOS double knockout was produced as described previously (Zhao et al., 2009). MVECs were isolated as described previously (Garrahan et al., 2006), with some modification. In brief, 3–4 wk-old mice were anesthetized by using isoflurane (2.5% in room air) according to the protocol approved by the University of Illinois Animal Care Committee. Depth of anesthesia was ascertained by loss of pain reflex to a vigorous paw pinch. After exsanguination, the lung was perfused with sterile HBSS supplemented with antibiotics. Lung was removed, minced, and digested with collagenase A (1.0 mg/ml in HBSS) for 60 min at 37°C on shaker. The released cells were centrifuged at 200 g for 10 min. The pellet was suspended in 10 ml of suspension buffer (Ca2+- and Mg2+-free PBS containing 0.5 g/100 ml bovine serum albumin, 2 mM EDTA, and 4.5 mg/ml l-glucose), and filtered through a 200-µm mesh sterile filter. The cell suspension was incubated with 10 µg of anti-mouse PECA1-MAb antibody on ice for 1 h, centrifuged to remove unbound antibody, and washed once with suspension buffer. The cells were incubated with Dynabeads M-450 (Sheep anti-IgG) for 60 min at 4°C. The cells attached to Dynabeads were trapped in a magnetic column to separate a subpopulation of PECA1-positive cells, washed, and resuspended in endothelial growth medium EGM-2 MV medium (Lonza), supplemented with 10% FBS (Invitrogen), and seeded in Matrigel-coated culture dishes (BD).

MVECs and human lung microvascular endothelial cells (HMVECs); Cav-1−/− were grown in EGM-2 MV medium (Lonza) supplemented with 15% FBS (Invitrogen). HPAECs (Lonza) were grown in EGM-2 Medium (Lonza) supplemented with 15% FBS (Invitrogen). Endothelial cells were used at passages 2–6. Human microvascular endothelial cell (HMECs; dermal; Lonza) were grown in MCDB 131 Medium (Invitrogen) supplemented with 10% FBS, 0.003 µg/ml h-EGF, 0.001 mg/ml hydrocortisone, and l-glutamine.

For measurement of basal NO generation, cells were incubated in l-arginine-free EBM media (Lonza) for 2 h. Cells were treated with 4 µM NPA, 4 µM 1400W, 4 µM i-NNA, and 100 µM SIN-1 for 30 min; and with 150 U/ml PEG-SOD, 100 µM TEMPO, 2 µg/ml Rho inhibitor C3 transferase, and 6 µM AP-CPD peptide for 2 h. 70–80% confluent monolayers were transfected using FuGENE HD (Roche) and Lipofectamine 2000 (Invitrogen) transfection reagents in accordance with the manufacturer’s instructions.

Co-immunoprecipitation and Western blotting
Cells were lysed in modified RIPA buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS, 1 mM NaVO4, 10 µg/ml leupeptin, 10 µg/ml aprotinin, and 0.1 mM PMSF). For co-immunoprecipitation experiments, 350 µg of total protein was incubated with 15 µg of antibody at 4°C, and precipitates were collected with protein A/G agarose beads. Densitometry of blots was performed using ImageJ software (National Institutes of Health).

RhoA-GTP pull-down assay
RhoA activity was determined by pull-down assays using GST-Rho-kinase beads (Knezevic et al., 2007). Cells were lysed in 50 mM Tris-HCl, pH 7.5, 10 mM MgCl2, 0.5 M NaCl, 1 mM EDTA, 1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS, 10 µg/ml leupeptin, 10 µg/ml aprotinin, and 0.1 mM PMSF. Cell lysates were incubated with GST-RBD beads at 4°C and bound proteins were analyzed by Western blotting. 5% of cell lysates were used to detect total RhoA.

NO measurement
NO formation was measured using porphyrinic NO electrodes (Brovkovich et al., 1999). Electodes were calibrated using a stock solution of NO-saturated water. The NO electrode was placed onto the cell surface and two additional electrodes were added into the solution to generate a 650-mV potential. The system was coupled to a FAS1 femtostat and a personal computer with electrochemical software (Gamry Instruments). The electrode current, which is proportional to NO concentration, was measured as a function of time.

Nitrre measurement
Nitrite accumulation in cell culture media was assessed by a chemiluminescence assay (Bonini et al., 2002) using an NO Analyzer (NOA 280; GE Healthcare). Authentic NaNO2 solutions of known concentrations were used as standards. All measurements were performed relative to cell-free controls.
measurement of intracellular superoxide

Cells plated in a 96-well plate were left untreated or incubated with 50 µM BH4 or 500 µM L-arginine for 30 min, then treated with 50 µM DHE for 1 h. Emission of EOH, a product of DHE oxidation, at λ = 580 nm, was measured using a SpectraMax M5e Multi-Mode Microplate Reader (Molecular Devices).

SNO measurement

SNO content of junctional proteins was detected by biotin-switch method using the S-nitrosylated Detection Assay kit (Cayman Chemical) according to the manufacturer’s instructions. The biotin-labeled proteins were precipitated with ice-cold acetone, and 350 µg of total protein was used for co-immunoprecipitation with specific Abs. SNO content was detected with SNO detection reagent I on Western blots. The same blot was reprobed for protein of interest.

Immunofluorescence staining and image analysis

Cell were fixed with 4% paraformaldehyde, permeabilized with 0.2% Triton X-100, and stained as described previously (Komarova et al., 2002). Samples were mounted with Prolong Gold antifade reagent (Invitrogen). Z-stack images were obtained using a confocal microscope (LSM 510 Meta; Carl Zeiss) equipped with a 63x, 1.2 NA water immersion objective lens and Ar ion and dual HeNe lasers, and driven by LSM software. 12-bit images were analyzed using MetaMorph software (Molecular Devices) and processed for illustration in Photoshop (Adobe). Projection images were generated by collecting the maximum pixel intensity from each image of the z stack and by projecting pixel intensity onto the single (projection) image. The relative accumulation of β-catenin (mean fluorescence intensity) at AJs and the area of gaps were measured on projected images. The 12-bit images were thresholded by subtracting intracellular background, and the mean fluorescence intensity at AJs was measured. For measurement of eNOS accumulation at AJs, the corresponding VE-cadherin image was used to generate a binary mask within 0 (outside of AJs) and 1 (AJs) area values after subtracting intracellular background, and the eNOS image was multiplied by a binary mask image for VE-cadherin. The mean fluorescence intensity of eNOS at AJs was then measured as described for β-catenin. Gap area was expressed as a percentage of the area outside of the cell.

Live cell imaging and image processing

Cells were imaged in phenol-free EBm media (Lonza) supplemented with 5% FBS at 37°C maintained by the stage heater (Tempcontrol-37; Carl Zeiss). Z-stack images were obtained using a confocal microscope (LSM 510 Meta; Carl Zeiss) equipped with a 63x, 1.2 NA water immersion objective lens and Ar ion and dual HeNe lasers, and driven by LSM software. 12-bit images were analyzed using MetaMorph software (Molecular Devices) and processed for illustration in Photoshop (Adobe). Projection images were generated by collecting the maximum pixel intensity from each image of the z stack and by projecting pixel intensity onto the single (projection) image. The relative accumulation of β-catenin (mean fluorescence intensity) at AJs and the area of gaps were measured on projected images. The 12-bit images were thresholded by subtracting intracellular background, and the mean fluorescence intensity at AJs was measured. For measurement of eNOS accumulation at AJs, the corresponding VE-cadherin image was used to generate a binary mask within 0 (outside of AJs) and 1 (AJs) area values after subtracting intracellular background, and the eNOS image was multiplied by a binary mask image for VE-cadherin. The mean fluorescence intensity of eNOS at AJs was then measured as described for β-catenin. Gap area was expressed as a percentage of the area outside of the cell.

Endothelial barrier permeability measurement

The permeability of MLVECs monolayer to EBA was determined as described previously (Patterson et al., 1992). Cells were grown to confluence on transwell inserts with a 0.4 µm pore size. Upper (luminal) and lower (abliminal) chambers were filled with HBSS containing 0.5% BSA and 20 mM HEPES. After a 30-min equilibration period, the luminal chamber was loaded with 0.057% EBA. Samples from the abluminal chamber were collected every 5 min for 1 h. EBA content was calculated from the optical density measured at 620 nm with the aid of the SpectraMax Plus microplate spectrophotometer (Molecular Devices). The rate of EBA clearance from luminal chamber was determined by linear regression between 5 and 60 min.

Perfusion of mouse lungs and measurement of transvascular fluid filtration

Murine lung preparations from WT and Cav-1−/− male mice (C57Bl/6 strain) were performed according to an approved protocol of the University of Illinois at Chicago Animal Care Committee (Vogel et al., 2000). In brief, isolated lung perfusions were performed with RPMI medium at constant flow (2 ml/min), temperature (37°C), and venous pressure (4 cm H2O).

The preparation was ventilated at a rate of 120 breaths per min, at constant peak inspiratory (~10 cm H2O) and end expiratory pressures (2 cm H2O). The lung weight change was recorded by a force-displacement transducer (Model FT03C; Grass Technologies). All lung preparations underwent a 10-min equilibration perfusion to establish isogravitometric conditions followed by a 30-min step change in perfusate albumin concentration (g/100 ml) of albumin from 0 to 10. In some cases, AP-CCD (final concentration of 10 µM) was added into the high-albumin perfusate between minute 20 and 40 from the beginning of perfusion. The fluid filtration rate was obtained by fitting a double exponential growth curve, y0 = α(1 - e−kx) + β(1 - e−60x), to the lung weight gain upon removal of albumin. The fast exponential component, which was of relatively low amplitude and coincident with the change of solution, most likely represented the effect of the shift in solution viscosity. The slow exponential component represented fluid filtration across the vessel wall induced by the change in oncotic pressure. The filtration rate was determined from the initial slope of this component extrapolated to zero time.

Statistical analysis

Data handling was performed using Sigma Plot software (SPSS). For statistical analysis, a paired two-tailed Student’s t-test was used to compare data between two groups.

Online supplemental material

Fig. S1 provides a characterization of primary MLVECs. Fig. S2 shows the effect of a Rho inhibitor on phosphorylation of MLC 2 in endothelial permeability studies. Table S1 summarizes the parameters of transvascular fluid filtration across the microvessel wall in isolated lungs. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201012129/DC1.

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