Serine/threonine phosphatase 5 (PP5C/PPP5C) regulates the ISOC channel through a PP5C-FKBP51 axis

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Abstract
Pulmonary endothelial cells express a store-operated calcium entry current (I\textsubscript{soc}), which contributes to inter-endothelial cell gap formation. I\textsubscript{soc} is regulated by a heterocomplex of proteins that includes the immunophilin FKBP51. FKBP51 inhibits I\textsubscript{soc} by mechanisms that are not fully understood. In pulmonary artery endothelial cells (PAECs) we have shown that FKBP51 increases microtubule polymerization, an event that is critical for I\textsubscript{soc} inhibition by FKBP51. In neurons, FKBP51 promotes microtubule stability through facilitation of tau dephosphorylation. However, FKBP51 does not possess phosphatase activity. Protein phosphatase 5 (PP5C/PPP5C) can dephosphorylate tau, and similar to FKBP51, PP5C possesses tetratricopeptide repeats (TPR) that mediate interaction with heat shock protein-90 (HSP90) chaperone/scaffolding complexes. We therefore tested whether PP5C contributes to FKBP51-mediated inhibition of I\textsubscript{soc}. Both siRNA-mediated suppression of PP5C expression in PAECs and genetic disruption of PP5C in HEK293 cells attenuate FKBP51-mediated inhibition of I\textsubscript{soc}. Reintroduction of catalytically competent, but not catalytically inactive PP5C, restored FKBP51-mediated inhibition of I\textsubscript{soc}. PAEC cell fractionation studies identified both PP5C and the ISOC heterocomplex in the same membrane fractions. Further, PP5C co-precipitates with TRPC4, an essential subunit of ISOC channel. Finally, to determine if PP5C is required for FKBP51-mediated inhibition of calcium entry-induced inter-endothelial cell gap formation, we measured gap area by wide-field microscopy and performed biotin gap quantification assay and electric cell-substrate impedance sensing (ECIS\textsuperscript{6}). Collectively, the data presented indicate that suppression of PP5C expression negates the protective effect of FKBP51. These observations identify PP5C as a novel member of the ISOC heterocomplex that is required for FKBP51-mediated inhibition of I\textsubscript{soc}.

Keywords
phosphatase, PP5C, FKBP51, ISOC calcium channel, endothelial barrier

Disruption of the endothelial monolayer leads to increased vascular permeability, increasing deposition of fluid, solutes, and macromolecules into perivascular tissue. Specifically, inter-endothelial cell gap formation is one mechanism of endothelial dysfunction that contributes to increased permeability in the pulmonary circulation.\textsuperscript{1} Increases in cytosolic calcium contribute to inter-endothelial cell gap formation in pulmonary endothelial cells,\textsuperscript{2} and transient receptor potential canonical (TRPC) proteins contribute to increased cytosolic calcium through mediating calcium entry.\textsuperscript{3} TRPC4 and TRPC1 are necessary for the calcium conductance of

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the moderately calcium-selective current. $I_{soc}$ a small inwardly rectifying current with a reversal potential around $+32 \text{mV}$.\textsuperscript{4,5} Additionally, $I_{soc}$ is a thapsigargin-activated current. When activated, $I_{soc}$ leads to inter-endothelial cell gap formation within the pulmonary circulation.\textsuperscript{6}

Our laboratory has shown that a large molecular weight immunophilin, FKBP51, is part of the ISOC heterocomplex. Furthermore, we have shown that FKBP51 regulates calcium entry in pulmonary artery endothelial cells (PAECs).\textsuperscript{7} FKBP51 inhibits thapsigargin-induced $I_{soc}$ in both PAECs and HEK293 cells, without affecting other thapsigargin-induced calcium entry pathways. This suggests that FKBP51 is a specific inhibitor of $I_{soc}$. However, the mechanism by which FKBP51 regulates $I_{soc}$ is poorly understood. FKBP51 promotes microtubule polymerization in neurons through dephosphorylation events of microtubule-associated proteins\textsuperscript{8} and we have recently demonstrated that inhibition of $I_{soc}$ by FKBP51 is dependent upon an FKBP51-mediated increase in microtubule polymerization.\textsuperscript{9} However, because FKBP51 lacks phosphatase activity the link between FKBP51 and dephosphorylation remains unknown. Serine/threonine protein phosphatase five (PP5C; encoded by PPP5C) dephosphorylates microtubule-associated proteins\textsuperscript{8} and we have recently demonstrated that PP5C plays a role in FKBP51-mediated inhibition of $I_{soc}$.\textsuperscript{9}

We found that neither the suppression nor genetic disruption of PP5C expression significantly altered thapsigargin-induced $I_{soc}$; however, without PP5C FKBP51-mediated inhibition of $I_{soc}$ was abolished. Reintroduction of wild-type (WT) PP5C, but not catalytically deficient PP5C, to PP5C$^{-/-}$ cells restored the FKBP51-mediated inhibition of $I_{soc}$. Further, a subpopulation of PP5C localizes and interacts with the ISOC heterocomplex. Since the inhibition of $I_{soc}$ by FKBP51 leads to reduced inter-endothelial cell gap formation,\textsuperscript{9} and we observed that PP5C is required for the FKBP51-mediated inhibition of $I_{soc}$, we next investigated the role of PP5C in inter-endothelial cell gap formation. Indeed, the FKBP51-mediated protection of the endothelial barrier against calcium entry-induced gap formation was lost upon PP5C small interfering RNA (siRNA) treatment. Cumulatively, our observations reveal a novel role for PP5C in ion channel regulation and endothelial barrier function.

**Materials and methods**

**Reagents**

All reagents were obtained from Sigma–Aldrich (St. Louis, MO, USA) unless otherwise indicated. Cell culture medium was obtained from Thermo Fisher Scientific (Waltham, MA, USA) or Santa Cruz Biotechnology (Santa Cruz, CA, USA). Penicillin/streptomycin was obtained from Thermo Fisher Scientific. Fetal bovine serum (FBS) was purchased from Thermo Fisher Scientific. EquaFETAL\textsuperscript{®} bovine serum (EBS) was purchased from Atlas Biologicals, Inc. (Fort Collins, CO, USA). Hank’s Balanced Salt Solution (HBSS) was purchased from Thermo Fisher Scientific. Monoclonal antibodies to FKBP51 were described previously.\textsuperscript{11} TRPC4 antibody was obtained from Alomone Labs (Jerusalem, Israel). Polyclonal antibody for PP5C was previously described.\textsuperscript{12} β-actin antibody was purchased from Santa Cruz Biotechnology.

**Cell culture**

PAECs were isolated from Sprague–Dawley rats as previously described.\textsuperscript{13} Cell culture media of PAECs and HEK293 cells contained high glucose DMEM supplemented with 10% FBS or EBS, penicillin G (50 U/mL), and streptomycin (0.05 mg/mL). All animal work was approved by the University of South Alabama Institutional Animal Care and Use Committee in compliance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

**Western blotting**

Cell lysis was achieved by scraping and sonication (10 s at 12% output; Branson Digital Sonifier model S-450D; Branson, Danbury, CT, USA) in RIPA buffer (Boston BioProducts; Ashland, MA, USA) with 1% protease inhibitor cocktail. Whole cell lysates were centrifuged at 12,000 x g for 20 min. Lysates were electrophoresed on 4–12% bis-Tris gels (Thermo Fisher Scientific) and proteins transferred to nitrocellulose membranes at 100 V. Membranes were blocked for 1 h with milk (5% non-fat dry milk/0.2% BSA in PBS supplemented with 0.1% Tween-20) at room temperature. Membranes were then incubated with primary antibodies overnight at 4°C and with secondary antibodies for 1 h at room temperature. The dilutions for primary antibodies were as follows: 1:700 for FKBP51 and 1:1000 for TRPC4, PP5C and β-Actin. Dilutions for secondary antibodies were: 1:5000 anti-mouse for FKBP51, 1:30,000 anti-mouse for β-actin, and 1:10,000 anti-rabbit for PP5C and TRPC4. Protein visualization was achieved with Supersignal West Pico or West Femto chemiluminescent substrates (Thermo Fisher Scientific). Densitometry of visualized westerns was achieved using Image J software.\textsuperscript{15}

**Genetic suppression of PP5C expression with siRNA**

The suppression of PP5C expression was achieved with siRNA targeted to the sequence

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5’-AATGGGCGATGGCGAGGGCGA-3’
\] (Qiagen; Hilden, Germany). Transfection of PAECs was initiated when monolayers had obtained 50–60% confluence. PAEC monolayers were transfected with a 20 nM final siRNA concentration in lipofectamine 3000 (Thermo Fisher Scientific). Complete cell culture media was added to the transfected monolayers at 4 h and siRNA incubation continued to 48 h.
Construction of PP5C expression plasmid

Complementary DNA (cDNA) comprising the complete coding region of human PP5C\textsuperscript{16} was subcloned into pBlueScript (Stratagene; San Diego, CA, USA; Genbank X52328). A prokaryotic expression vector was constructed by polymerase chain reaction (PCR) amplification with primers incorporating a 5' EcoRI restriction site, a methionine start codon followed by a sequence encoding a hexahistidine affinity tag (MGHHHHHHHG: his-tag), and a 3' PstI restriction site. The amplified fragment was subcloned into pKK223-3 (Pharmacia; New York City, NY, USA; Genbank M77749). A peptide NLYFQGA (which forms a part of a Tobacco Etch Virus [TEV] protease recognition sequence) was added to the N-terminal and subcloned into pMalc2E (New England Biolabs; Ipswich, MA, USA; NEB #N8066) using conventional methods. Expression constructs with the indicated mutations were made using a QuikChange Site-Directed Mutagenesis Kit (Stratagene). These expression constructs encoded an N-terminal Maltose Binding Protein (MBP) fusion with the catalytic domain of PP5C (residue Ser169 to residue Met499; designated PP5C), plus a linker sequence (MGHHHHHHHG SVVDSLDIENLYFQGA) between the fusion partners containing a hexahistidine affinity tag, a spacer sequence, and a TEV protease cleavage site. The N-terminal MBP aids the folding of PP5C when expressed in \textit{E. coli} and the His-tag aids purification. Primer sequences used in the generation of these constructs are provided in Supplementary Table 1. Following purification, the MBP and hexahistidine tags were removed by treatment with TEV protease. The catalytically active proteins were then further purified using ion exchange (monoQ; Pharmacia) chromatography. Additional details are described in Ni et al.\textsuperscript{17}

Phosphatase assays

Protein phosphatase activity was measured by the dephosphorylation of DiFMUP in assay buffer (50 mM MOPS, 5 mM β-ME, 1 mM MnCl\textsubscript{2}) according to established methods.\textsuperscript{16–18}

PP5C\textsuperscript{−/−} and PP5C catalytically inactive cell lines

Disruption of PP5C expression in HEK293 cells was achieved using CRISPR/Cas9-mediated cleavage of exon one, adapting essentially the same strategy used to generate PP5C knockout mice.\textsuperscript{19} Briefly, following the X-tremeGENE protocol, HEK293 cells were transfected with 1 μg of a Cas9 expressing plasmid into which a RNA-guide template targeting PP5C was subcloned after the U6-promoter (Addgene; Cambridge, MA, USA; #42230) and allowed to grow for three days. On day 3 using flow cytometry, cells were individually sorted into 96 well plates. The clonally derived cell lines were then screened using western analysis to identify lines with no PP5C protein. Genomic DNA sequencing was used to validate the disruption of PP5C expression, which occurred due to stochastic insertion that induces frame shift.

Reintroduction of WT (PP5C-WT) and catalytically inactive (PP5C-CatΔ) expression constructs. Human PP5C was amplified by PCR incorporating EcoRI and NotI sites by adding the appropriate sequence into the synthetic primers. The construct was then subcloned into pcDNA3. Plasmids allowing the expression of cDNA encoding PP5C (WT) were transfected into PP5C\textsuperscript{−/−} cells with X-tremeGENE (0.5 μg) for 24 h following the manufacturer’s protocol. To generate catalytically inactive PP5C, we performed a comprehensive site-directed mutagenesis of PP5C, in which all ten of conserved amino acids within the catalytic site identified by the high-resolution crystal structure of PP5C\textsuperscript{16} were systematically mutated using Stratagene QuikChange II Site-Directed Mutagenesis. Oligonucleotide sequences used for the introduction of the mutations are provided in Supplementary Table 2. All plasmids were sequenced to verify the fidelity of the constructs. Each of the mutant forms of PP5C were then subcloned into pKK223-3. After assessing purity via SDS-PAGE and Coomassie blue staining (Supplementary Fig. 1), the purified proteins were tested for phosphatase activity using identical buffers and substrates, as described for WT PP5C above. His304 was confirmed as the proton donor in the catalytic mechanism and the catalytically inactive construct (PP5C-H304N) subcloned into P-lenti6-V5-D-topo (Addgene). Catalytically dead PP5C (PP5C-CatΔ) or WT PP5C was then reintroduced into HEK293 PP5C\textsuperscript{−/−} cell line as described previously.\textsuperscript{20} Following blastomycin selection, expression of the reintroduced constructs was confirmed by western analysis.

Overexpression of FKBP51 in PAECs

Production of lentivirus encoding human FKBP51 (lv3899) and stable transduction of PAECs were as previously described.\textsuperscript{9} For acute overexpression of FKBP51, HEK293 cells, and PAECs were treated with a 1:1 ratio of cell culture media and lv3899 in the presence of polybrene (8 μg/mL) for 24 h and cells were subjected to patch clamp electrophysiology or lysis for western blot analysis after the 24-h lentiviral infection.

Patch-clamp electrophysiology

Cells were seeded onto glass coverslips and grown to confluence in 35-mm dishes. Patch-clamp electrophysiology recordings were performed in whole-cell configuration on electrically isolated cells as described.\textsuperscript{7,14} Non-enzymatic cell dissociation solution in PBS (Sigma, cat. no. C5789) was utilized for the isolation of single cells. Transmembrane current evoked by step depolarization in 20 mV increments in the range of −100 to +80 mV was measured with an Axopatch 200B Amplifier (Molecular Devices; Sunnyvale, CA, USA). PClamp10 software was used to measure current as the mean value of the current amplitude during the last
20 ms of each step. Standard pipette and bath solutions were composed as previously described, and all solutions were adjusted to 290–300 mOsm/L with sucrose. Hemo capillaries (A-M Systems; Sequim, WA, USA) were pulled by a micro-pipette puller (P-97; Sutter Instruments; Novato, CA, USA) to produce recording pipettes. Pipettes were heat-polished by a microforge (MF-830; Narishige; Tokyo, Japan) and filled with standard pipette solution to a final resistance of 3–5 megohms. All experiments were performed at room temperature. Representative patch tracings are shown for each newly introduced cell type in the respective figure corresponding to the current-voltage (IV) plot in which electrophysiological recordings of the cell line are first depicted. Representative tracings are indicative of the electrical recording of one cell within the corresponding IV plot. Patch tracings for subsequent treatments of the cell lines are shown in the supplemental material (Supplementary Fig. 2).

Membrane-cytoskeletal preparation

Cell fractionation of PAECs was performed in the cold room (4°C) using the membrane-cytoskeletal preparation as previously described.14 PAEC lysates were pelleted and resuspended in ice-cold sucrose buffer and homogenized with a dounce homogenizer. The homogenate was centrifuged at 3000 × g (17 min, 4°C) and the supernatant collected. The supernatant was subsequently centrifuged at 50,000 × g (30 min, 4°C). The pellet was re-suspended in octyl-D-glucopyranoside containing extraction buffer with potassium iodide (KI). Samples were incubated for 40 min in the cold room and centrifuged at 145,000 × g (50 min, 4°C). The membrane containing supernatant was collected and the cytoskeletal containing pellet was solubilized with sonication.

Co-precipitations

Co-precipitation of TRPC4 with PP5C was achieved with SureBeads™ IgG conjugated magnetic beads (Bio-Rad; Hercules, CA, USA) following manufacturer’s instructions. Following a 10 min incubation of the magnetic beads with the precipitating antibody, whole cell lysates of PAECs were incubated with the beads for 1 h under agitation. After washing, protein was eluted from the beads with 40 μL of 1 × Laemlli buffer (diluted from 6 × Laemlli buffer; Boston BioProducts, Ashland, MA, USA) for 10 min at 70°C.

Wide-field gap formation studies

PAECs were seeded onto 35-mm coverslips and grown to confluence. PAEC monolayers were imaged (20×) at 15 s intervals for 2 min using a Zeiss Observer.D1 wide-field microscope and AxioVision Rel 4.8 Software. Thapsigargin (1 μM) was then added and monolayers imaged for 30 min. Cell gap size was measured by outlining gap areas and using the pixel-to-micron conversion feature of the AxioVision Rel 4.8 Software.

Biotin gap assay

Protocol was performed as previously described.21 PAECs were grown to confluence in biotin-coated plates (Thermo Fisher Scientific; PI15151). Control PAECs were treated with either vehicle (ethanol, 0.05%) or dexamethasone (10 nM) for 48 h. To activate calcium entry-induced gap formation, cells were treated with thapsigargin (1 μM) for a total of 10 min, or DMSO (0.05%). Seven minutes after thapsigargin treatment, cells were incubated with streptavidin-488 (5 μg/mL) for 3 min. Monolayers were then washed twice with PBS (200 μL/well). A final volume of 100 μL PBS was added to each well and fluorescence of biotin-conjugated streptavidin-488 was measured on a BioTek (Winooski, VT) Synergy 2 plate reader and the data were recorded with BioTek Gen5 software.

Electric cell-substrate impedance sensing (ECIS®)

ECIS® experiments were performed as previously described.22 A stable baseline resistance was recorded for at least 30 min before administration of thapsigargin (1 μM). Resistance changes were recorded until a new constant baseline was established.

Statistical analysis

GraphPad version 5.0 software (San Diego, CA, USA) was utilized for statistical analysis. A two-way ANOVA and Bonferroni post hoc test was used to assess comparisons between multiple groups. Data are presented as mean ± SEM and values were considered significantly different when P < 0.05.

Results

PPSC is required for the FKBPS1-mediated inhibition of I_{soc}

The moderately calcium-selective I_{soc} is activated by the plant alkaloid thapsigargin, which results in depletion of endoplasmic reticulum stores through inhibition of the sarcoplasmic/endoplasmic reticular Ca²⁺/ATPase.23 We have previously shown that two immunophilins, FKBPS1 and FKBPS2, regulate thapsigargin-induced calcium entry.7 Calcium entry through I_{soc} is inhibited by FKBPS1 overexpression in both HEK293 cells and PAECs via a mechanism dependent upon increased microtubule polymerization.7,9 FKBPS1 has been implicated in promoting microtubule polymerization through the isomerization and dephosphorylation of tau.8 Since FKBPS1 lacks phosphatase activity, a phosphatase must be involved, and PP5C is a phosphatase reported to dephosphorylate tau.10 Like FKBPS1, PP5C contains TPR domains through which it interacts with HSP90.24 We therefore wanted to study the role of PP5C in the FKBPS1-mediated inhibition of I_{soc}.
FKBP51 overexpression was achieved by lentiviral transduction of human FKBP51 in HEK293 cells (Fig. 1a, b). Consistent with our previous observations in PAECs, FKBP51 overexpression significantly inhibited Isoc in HEK293 cells (Fig. 1c). To determine whether PP5C is required for the inhibition of Isoc by FKBP51, FKBP51 was over-expressed in HEK293 cells in which PP5C expression was genetically disrupted. Although FKBP51 over-expression inhibited Isoc in control HEK293 cells, over-expression of FKBP51 was unable to inhibit Isoc in the PP5C−/− cells (Fig. 1d–f). Reintroduction of full-length PP5C into PP5C−/− cells restored the FKBP51-mediated inhibition of Isoc.

The catalytic activity of PP5C is required for the FKBP51-mediated inhibition of Isoc

We have previously reported the high-resolution (1.6 Å) crystal structure of PP5C in complex with phosphate (mimicking the leaving group phosphate; PDB code 1S95), which allowed us to define the PP5C catalytic mechanism. The PP5C active-site comprises ten amino acids—D242, XH244(X)26–27, D271, XXD274, R275(X)28, N303, H304(X)48, H352(X)48–54, R400, (X)27, H427—that are 100% conserved in PP1C, PP2AC, and PP2BC, which share the same catalytic mechanism. Six of the conserved active site amino acids act as metal-coordinating residues (Asp242, His244, Asp271, Asp274, His304, His352).
Asn$^{303}$, His$^{352}$, and His$^{427}$) and four (Arg$^{275}$, Asn$^{303}$, His$^{304}$, and Arg$^{400}$) position the phosphate ion through strong hydrogen bonds to the phosphoryl oxygens of the substrate (Supplementary Fig. 3). The structure of the PP5C catalytic domain with bound phosphate provided a model of substrate-enzyme interactions in a near attack configuration. Analysis of the substrate analog complex revealed that His$^{304}$ forms a short (2.6 Å), strong hydrogen bond with O$^4$ of phosphate (the substrate leaving group oxygen). Asp$^{274}$ hydrogen bonds with the imidazole of His$^{304}$ (increasing the imidazole pKa). This stabilizes the histidine side chain in a cationic state that is capable of protonating the leaving group. Thus, His$^{304}$ acts as a general acid in the hydrolysis reaction. In addition, positively charged His$^{304}$, along with Arg$^{275}$, and Arg$^{400}$, contribute to electrostatic stabilization of the transition state by neutralization of the negative charge developing on the leaving group oxygen. Therefore, mutation of His$^{304}$, Asp$^{274}$, or the key metal coordinating amino acids were predicted to generate a catalytically inactive enzyme.

To validate the proposed catalytic mechanism and to determine the best amino acid to mutate in order to generate a catalytically inactive enzyme, we performed a comprehensive site-directed mutagenesis of PP5C, in which all the conserved amino acids within the catalytic site were systematically mutated (Supplementary Fig. 4). Each mutant form of PP5C was then expressed and purified using methods developed for the production of PP5C-Cat$^+$ (Supplementary Experimental Procedures). We then reintroduced PP5C-Cat$^+$ into the HEK293 cell line in which we previously used CRISPR/Cas9 to disrupt PP5C expression. In the PP5C-Cat$^+$ cells, FKBP51 overexpression was unable to inhibit $I_{soc}$ (Fig. 1i) indicating that the phosphatase activity of PP5C is needed for the FKBP51-mediated inhibition of $I_{soc}$. However, as shown above (Fig. 1f), when native PP5C was reintroduced, FKBP51-mediated inhibition of $I_{soc}$ was restored. Together, these observations indicate that PP5C catalytic activity is needed for the FKBP51-mediated inhibition of $I_{soc}$.

### PP5C is also required for the FKBP51-mediated inhibition of $I_{soc}$ in PAECs

PAECs and HEK293 cells express similar $I_{soc}$ electrophysiology. However, because mechanisms of $I_{soc}$ regulation are so poorly understood it was essential to verify PP5C is also needed for the FKBP51-mediated inhibition of $I_{soc}$ in PAECs before exploring physiological significance. To address this, we increased FKBP51 expression using two independent methods and decreased PP5C expression using siRNA. We first utilized dexamethasone (Dex) to increase the expression of FKBP51 in PAECs (Fig. 2a, b), which our laboratory previously showed inhibits $I_{soc}$ in PAECs. To verify that the inhibition of $I_{soc}$ was due to FKBP51 and not off-target effects of dexamethasone or glucocorticoid signaling, a PAEC line was generated to constitutively overexpress FKBP51 (cFKBP51 Fig. 2d, e). Dexamethasone treatment (Fig. 2c) and constitutive FKBP51 overexpression (Fig. 2f) both significantly inhibited $I_{soc}$ in PAECs in the presence of scrambled siRNA. In contrast, the inhibition of $I_{soc}$ by Dex (Fig. 2c) or with FKBP51 overexpression (Fig. 2f) was negated with even a modest decrease in PP5C expression by siRNA. These data confirm that PP5C contributes to the FKBP51-mediated inhibition of $I_{soc}$ in PAECs.

### PP5C associates with the ISOC channel heterocomplex

The ISOC channel heterocomplex comprises both channel pore-forming proteins and proteins that associate with the intracellular N- and C-terminal domains of the channel proteins. A spectrin-protein 4.1-TRPC4 interaction is required for the activation of $I_{soc}$ in PAECs and TRPC4 knockout mice do not express $I_{soc}$. These observations indicate that TRPC4 is an essential channel protein in the ISOC heterocomplex. Our laboratory has also shown that the cochaperones FKBP51 and FKBP52 co-purify with TRPC4 in PAECs. As FKBP51 has been shown to serve as a scaffold independent of HSP90 for serine/threonine phosphatase activity.
phosphatases, we wanted to determine whether PP5C also associates with TRPC4 of the ISOC heterocomplex. We first tested to determine whether PP5C is found in plasma membranes where the ISOC channel is localized. To detect PP5C in cell membranes of PAECs, a membrane-cytoskeletal preparation was utilized to isolate TRPC4-containing membrane fractions of PAECs. We observed that some PP5C is located in the TRPC4-containing membrane fraction of PAECs (Fig. 3a). The cytoskeletal protein actin was resolved only in the cytoskeletal fraction. To address whether PP5C associates with TRPC4 in PAECs, co-precipitations were performed (Fig. 3b). PP5C co-precipitated with TRPC4 in PAECs and reciprocally, TRPC4 co-precipitated with PP5C. Co-precipitation of PP5C and TRPC4 was also observed in HEK293 cells (data not shown). Collectively, these data reveal that a population of PP5C resides at the plasma membrane and interacts with the ISOC channel.

Fig. 2. PP5C is required for the FKBP51-mediated inhibition of $I_{\text{soc}}$ in PAECs. (a) Genetic suppression of PP5C expression was achieved by 48 h siRNA transfection of PAECs. As analyzed by western blot and (b) densitometry (calculated as band intensity of the indicated protein relative to that of $\beta$-actin and normalized to one for control PAECs), dexamethasone (Dex; 10 nM) treatment resulted in the overexpression of FKBP51 in PAECs transfected with PP5C siRNA or a scrambled control. $^aP < 0.05$ vs. no Dex treatment; $n = 3$. (c) $I_{\text{soc}}$ in Dex (10 nM)-treated PAECs (red) was significantly decreased from control PAECs (gray). PP5C siRNA in PAECs expressing endogenous FKBP51 (vehicle control) did not change $I_{\text{soc}}$, but in PAECs expressing increased FKBP51, genetic suppression of PP5C prevented inhibition of $I_{\text{soc}}$ ($^aP < 0.05$; n = 6). (d) PP5C siRNA treatment decreases PP5C expression in constitutive FKBP51 overexpressing PAECs (cFKBP51) as shown by western blot and densitometry (calculated as band intensity of the indicated protein relative to that of $\beta$-actin and normalized to one for control cFKBP51 PAECs. $^aP < 0.05$ vs. scrambled siRNA; n = 3). (e) cFKBP51 cells treated with scrambled siRNA (red) demonstrated reduced $I_{\text{soc}}$ relative to control PAECs (gray) ($^P < 0.05$; n = 5). FKBP51 overexpressing PAECs treated with PP5C siRNA (blue) resulted in significantly increased $I_{\text{soc}}$ compared to scrambled control (red) ($^P < 0.05$; n = 5). cFKBP51 cells treated with PP5C siRNA (blue) do not have a significantly different $I_{\text{soc}}$ than control PAECs (gray). Data for control PAECs (gray) are those shown in (c) and are included here for comparison.

Suppression of PP5C expression attenuates the FKBP51-mediated inhibition of calcium entry-induced inter-endothelial cell gap formation

Activation of $I_{\text{soc}}$ contributes to inter-endothelial cell gap formation in endothelial cells of the pulmonary circulation. We have recently shown that FKBP51-mediated inhibition of $I_{\text{soc}}$ is sufficient to decrease calcium-induced inter-endothelial cell gap formation in PAECs. Having demonstrated that PP5C is required for the FKBP51-mediated inhibition of $I_{\text{soc}}$, we wanted to determine whether PP5C plays a biological role needed for the protective effect of FKBP51 on the endothelial barrier. In these studies, dexamethasone or lentiviral infection were used to increase the expression of FKBP51, and siRNA was used to decrease PP5C expression. We then measured thapsigargin-induced inter-endothelial cell gap formation in confluent PAECs using wide-field microscopy. In the presence of scrambled
siRNA, FKBP51 overexpression resulted in significant decreased gap area 10 min following thapsigargin treatment compared to control PAECs (Fig. 4a). In PAECs treated with PP5C siRNA without FKBP51 overexpression, gap formation was comparable to control cells. To determine whether PP5C is needed for the FKBP51-mediated inhibition of inter-endothelial cell gap formation in PAECs, suppression of PP5C expression with siRNA was again utilized. PP5C suppression attenuated the dexamethasone-mediated inhibition of gap formation, and significantly increased gap formation in the constitutive FKBP51 overexpressing PAECs (Supplementary Fig. 5). Due to the substantial difference in initial gap area between groups, relative area is depicted in which initial gap area at t = 0 min is normalized to 1 for each group (Fig. 4b). Normalization of gap area reveals a significant increase in inter-endothelial cell gap formation due to PP5C siRNA-treated PAECs in which FKBP51 was overexpressed.

To confirm that PP5C is needed for the FKBP51-mediated inhibition of inter-endothelial cell gap formation, and ensure our observations are not subject to bias due to the inherent subjectivity of wide-field gap measurements, a 96-well fluorescent assay to assess gap formation first described by Dubrovskyi et al. was utilized. In this assay, gap formation is assessed by quantifying fluorescence of desmosomes described by Dubrovskyi et al. was utilized. In this assay, the inherent subjectivity of wide-field gap measurements, a

**Discussion**

Activation of the endothelial $l_{oc}$ leads to inter-endothelial cell gap formation, which increases vascular permeability in the pulmonary circulation. Our laboratory has shown that moderate increases in expression of FKBP51 inhibits $l_{oc}$ and protects the endothelium from calcium entry-induced disruption. However, upregulation of FKBP51 may not be an effective therapeutic strategy against endothelial barrier disruption as it is also a potent inhibitor of glucocorticoid receptor (GR) signaling. GRs regulate many cellular functions through their regulation of nuclear transcription factors. Specifically, nuclear translocation of GR and the resulting transcriptional regulation are needed for the downregulation of inflammatory cytokines. As many instances of inter-endothelial cell gap formation are initiated by inflammatory events, such as in atherosclerosis and acute respiratory distress syndrome, the ability of GR to suppress inflammatory cytokines in the endothelium is essential to maintain homeostasis.
However, when FKBP51 is present in mature GR it prevents the translocation of GR to the nucleus thus inhibiting GR-mediated transcription. Therefore, it is essential that we unravel the mechanisms by which FKBP51 inhibits Isoc to optimize its efficacy in inflammatory diseases.

In this study, we identify PP5C as a novel member of the ISOC heterocomplex that is required for inhibitory action of FKBP51 on both thapsigargin-induced Isoc and inter-endothelial cell gap formation. PP5C is a serine/threonine phosphatase that is known to interact with other proteins via its N-terminal TPR domains. FKBP51 has a similar TPR domain, and the TPR domains of PP5C and FKBP51 dynamically compete for a consensus binding to HSP90 within mature GRs. Through this interaction, PP5C also regulates GR activity in cells via mechanisms that are independent of its catalytic activity. Like FKBP51, most studies indicate that PP5C is a negative regulator of GR signaling. Specifically, Zou et al. demonstrated that siRNA-mediated suppression of PP5C expression resulted in an increase in basal and dexamethasone-stimulated GR activity in A549 lung carcinoma cells. Furthermore, it has been determined that dephosphorylation of GR by PP5C at Ser211 inhibits GR translocation into the nuclei in MCF-7 cells. Dephosphorylation of Ser211 by PP5C has also been...
shown to inhibit steroid responsiveness and GR activity in airway smooth muscle cells. Together, these findings indicate that PP5C possesses similar pharmacological limitations to FKBP51 as a standalone treatment of inflammatory-mediated endothelial barrier disruption, as both PP5C and FKBP51 are negative regulators of GR.

Despite the inhibitory roles of FKBP51 and PP5C on GR activity, the PP5C-FKBP51 axis remains a promising pharmacologic target in the regulation of $I_{soc}$ and inter-endothelial cell gap formation. Specifically, as FKBP51 is such a highly GR-regulated protein that infers glucocorticoid resistance, it would be beneficial to determine how to prime the FKBP51-mediated inhibition of $I_{soc}$ without altering expression levels of FKBP51. As PP5C is essential for the effect of FKBP51 on $I_{soc}$ targeted activation of PP5C may allow for the optimization of the FKBP51-mediated inhibition of $I_{soc}$. Our current study shows that a population of PP5C resides at the membrane of PAECs where it interacts with the ISOC heterocomplex. Chatterjee et al. demonstrated that the TPR domain of PP5C binds to Rac1 GTPase, which when activated, increases the translocation of PP5C to HeLa cell membranes. As GR resides in the cytosolic and nuclear fractions of cells, increasing the translocation of PP5C to the plasma membrane could allow for an increase in the FKBP51-mediated inhibition of $I_{soc}$ without analogous inhibitory effects of GR. As other serine/threonine phosphatases including PP2A have been shown to serve a protective role in endothelial junctions, it is possible that PP5C also would serve a similar effect if a larger population was present at the membrane.

In addition to promoting translocation of PP5C to the plasma membrane as a potential therapeutic strategy, many physiological activators of PP5C have been identified. Ramsey et al. identified that long-chain fatty acyl-CoA esters and the C-terminal domain of HSP90 increase substrate access to the catalytic domain of purified PP5C. Furthermore, Yamaguchi et al. demonstrated that members of the S100 protein family greatly increase the activity of PP5C in COS-7 cells in a calcium-dependent manner. As calcium entry through the ISOC channel results in inter-endothelial cell gap formation, S100 proteins may serve as novel targets in the calcium-dependent activation of PP5C and protection against calcium-mediated gap formation. Unfortunately, while much progress has been made in the development of specific pharmacologic inhibitors of PP5C, little work has been dedicated to developing PP5C agonists.

Lastly, as FKBP51 and PP5C both mediate their effects in neurons via dephosphorylation of the microtubule-associated protein tau, it is possible that the effect of PP5C on the FKBP51-mediated inhibition of $I_{soc}$ may be microtubule-dependent. Endothelial cells of the pulmonary circulation express many microtubule-associated proteins, including tau. As microtubule stabilization with taxol has been shown to inhibit $I_{soc}$ in pulmonary microvascular endothelial cells, it is possible that FKBP51 and PP5C may be working together to dephosphorylate tau.
resulting in increased microtubule stability in PAECs. Indeed, we observed an increase in microtubule stability in FKBP51 overexpressing PAECs and demonstrated that this increased microtubule stability was critical for the FKBP51-mediated inhibition of $I_{soc}$. Future studies will determine if PP5C is required for the FKBP51-mediated promotion of microtubule stability and tau dephosphorylation in PAECs.

Collectively, our observations demonstrate that PP5C associates with the ISOC heterocomplex and is required for the FKBP51-mediated inhibition of $I_{soc}$ and interendothelial cell gap formation. The data indicate specifically that the catalytic activity of PP5C is important for the FKBP51-mediated inhibition of $I_{soc}$. Future studies will be directed towards better understanding the mechanisms by which PP5C regulates FKBP51 function.

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Conflict of Interest
None declared.

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