A novel assay uncovers an unexpected role for SR-BI in LDL transcytosis

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Received 2 February 2015; revised 17 August 2015; accepted 24 August 2015; online publish-ahead-of-print 2 September 2015

Time for primary review: 39 days

Aims

Retention of low-density lipoprotein (LDL) cholesterol beneath the arterial endothelium initiates an inflammatory response culminating in atherosclerosis. Since the overlying endothelium is healthy and intact early on, it is likely that LDL passes through endothelial cells by transcytosis. However, technical challenges have made confirming this notion and elucidating the mechanisms of transcytosis difficult. We developed a novel assay for measuring LDL transcytosis in real time across coronary endothelial cell monolayers; we used this approach to identify the receptor involved.

Methods and results

Murine aortas were perfused *ex vivo* with LDL and dextran of a smaller molecular radius. LDL (but not dextran) accumulated under the endothelium, indicating that LDL transcytosis occurs in intact vessels. We then confirmed that LDL transcytosis occurs *in vitro* using human coronary artery endothelial cells. An assay was developed to quantify transcytosis of DiI-LDL in real time using total internal reflection fluorescence microscopy. DiI-LDL transcytosis was inhibited by excess unlabelled LDL, while degradation of the LDL receptor by PCSK9 had no effect. Instead, LDL colocalized partially with the scavenger receptor SR-BI and overexpression of SR-BI increased LDL transcytosis; knockdown by siRNA significantly reduced it. Excess HDL, the canonical SR-BI ligand, significantly decreased LDL transcytosis. Aortas from SR-BI-deficient mice were perfused *ex vivo* with LDL and accumulated significantly less sub-endothelial LDL compared with wild-type littermates.

Conclusion

We developed an assay to quantify LDL transcytosis across endothelial cells and discovered an unexpected role for SR-BI. Elucidating the mechanisms of LDL transcytosis may identify novel targets for the prevention or therapy of atherosclerosis.

Keywords

Endothelium • Transcytosis • LDL • SR-BI • Atherosclerosis

1. Introduction

One of the earliest stages in the pathogenesis of atherosclerosis is the deposition of low-density lipoprotein (LDL) cholesterol in the sub-endothelial intima. Remarkably, how circulating LDL traverses the endothelial cell layer remains poorly understood.

In principle, permeability across the endothelium could occur between cells (paracellular transport) or through the cytoplasm of individual cells (transcytosis).¹ Early electron microscopic studies of

rat arteries perfused with LDL demonstrated its internalization into cellular vesicles and its targeting to either lysosomes or the basolateral membrane of the cell.² Importantly, LDL was never observed to be penetrating intercellular junctions. Furthermore, the size of LDL³ precludes paracellular passage without the opening of inter-endothelial junctions,⁴ yet an intact endothelium is observed overlying early atherosclerotic lesions.⁵ Thus, a number of investigators have suggested that LDL crosses the endothelium by transcytosis.^{6–8} Others have instead suggested that LDL exits the vasculature at areas of damaged

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or dividing endothelium, implicating the paracellular route.^{9,10} However, the rate of endothelial mitosis is vanishingly low (<0.05%),¹¹ and more importantly, there is scant to no *in vivo* evidence of endothelial apoptosis or denudation early on in human atherosclerosis.⁵

The alternative route of endothelial transcytosis has been best characterized for the serum protein albumin; receptors on the apical plasmalemma mediate its internalization into membrane invaginations known as caveolae.¹² After scission from the surface by a process requiring the GTPase mechanoenzyme dynamin, albumin-bearing vesicles make their way through the cytoplasm and undergo exocytosis at the basolateral cell surface.¹³ A major constituent of caveolae is the protein caveolin-1, the depletion of which prevents the formation of caveolae and ablates albumin transcytosis.¹⁴ Thus, it is intriguing to note that aortic rings from caveolin-1 knockout mice demonstrate a >50% reduction in LDL internalization⁶ while *cav1*^{-/-}/*ApoE*^{-/-} mice are protected from atherosclerosis (compared with *ApoE*^{-/-} mice) despite having increased plasma LDL.¹⁵ This uncoupling of elevated plasma LDL levels and the development of atherosclerosis suggests a defect in the transit of LDL across the endothelial monolayer. This observation is all the more remarkable, because an increase in paracellular leak has been described in caveolin-1 knockout mice,¹⁶ implying that the paracellular route cannot account for the accumulation of sub-endothelial LDL. Indeed, overexpression of caveolin-1 specifically in the endothelium caused accelerated atherosclerosis,^{17,18} although transcytosis of LDL was not examined.

Thus, the route by which LDL crosses the endothelium is controversial, but the data favour transcytosis. It is an important issue since the transcellular transport of LDL is likely to be tightly regulated and amenable to therapeutic manipulation, in contrast to paracellular diffusion. However, there is a paucity of literature on LDL transcytosis and its regulation,^{7,19} likely due to technical difficulties in its study. Traditional experimental approaches using transwells are problematic, because manipulation of endothelial monolayers often introduces intercellular gaps, confounding analysis of transcytosis.¹ Even a basic question like which receptor or receptors are involved is unanswered.

In this work, we report a novel assay that allows LDL transcytosis by single coronary artery endothelial cells to be quantified in real time. Using this approach, we describe an unexpected role for SR-BI.

2. Methods

2.1 Cell culture

In vitro experiments were performed using primary human coronary artery endothelial cells (HCAECs; Lonza, Canada) used in passages 6–9 and serum-starved for 3 h prior to experiments. While different batches of primary cells were used over the course of the project, cells from the same batch were used in individual experiments (e.g. control vs. an experimental condition).

2.2 Transwell transcytosis assay

Biotinylated LDL (Intracel, MD) was added to the upper chamber (50 µg/mL) of a transwell containing confluent HCAECs. Transwells (0.4 µm pore polyester, Costar, Corning, NY, USA) were incubated at 4°C for 10 min to allow LDL binding but not internalization. The transwell was then transferred to a fresh plate and was washed twice with cold PBS. Fresh media were added and cells were incubated at 37°C for 2 h with or without platelet-activating factor (PAF, 10 µM; Sigma-Aldrich, ON, Canada) or fluorescein-Na (1 µg/mL, MW 376 Da; Sigma-Aldrich).

To measure the amount of LDL flux, 100 µL of media was taken from the lower chamber and analysed by ELISA.¹ The aliquot of media was added

along with 50 µL of 1 µg/mL of horseradish peroxidase–conjugated sheep anti-LDL (Bioss, Burlington, ON, Canada) to a 96-well plate coated in streptavidin (Thermo Scientific, Waltham, MA, USA) at room temperature for 2 h with shaking. The plate was washed with buffer (25 mmol/L Tris, pH 7.2, 150 mmol/L NaCl, 0.05% Tween-20, 0.1% gelatin), then 50 µL of 3,3',5,5'-tetramethylbenzidine (Thermo Scientific) was added to the well and incubated at room temperature for 15 min. Fifty microlitres of 2 mol/L sulfuric acid was added, and quantification of LDL was determined by measuring absorbance at 450 nm with a ThermoMax Microplate Reader (Molecular Devices, Sunnyvale, CA, USA). A standard curve using known quantities of LDL was used to determine the sensitivity of the assay.

2.3 Total internal reflection fluorescence transcytosis assay

Total internal reflection fluorescence (TIRF) microscopy images were acquired on an Olympus cell TIRF Motorized Multicolor TIRF module mounted on an Olympus IX81 microscope (Olympus, Hamburg, Germany). Samples were imaged using a ×150/1.45 objective with 561 nm excitation lasers and Volocity acquisition software. Unless otherwise indicated, the penetration depth, or the distance into the basal aspect of the cell that is visible by TIRF microscopy, was set at 110 nm. For each cell, 150 TIRF images were acquired at a frame rate of 6.67 per second for a constant duration of ~22 s. At least 10 randomly selected cells were imaged in each experimental replicate. Dil-labeled human native (non-oxidized) LDL (2 µg/mL) with or without unlabelled LDL or high-density lipoprotein (HDL; Intracel, MD) was added to confluent cells seeded on 25 mm glass coverslips. Cells were put at 4°C to allow binding; after 10 min, cells were rinsed in cold PBS and fresh media (HEPES-buffered RPMI, i.e. HPMI)²⁰ were added followed by immediate imaging with a heated (37°C) stage using a standard cell chamber (Life technologies, Catalog no. A7816). In some experiments, cells were pre-treated with Dyngo4A (30 µM; Abcam, Cambridge) for 30 min.

Quantification of the transcytotic events was performed in a blinded fashion using a vesicular detection and tracking algorithm custom-written and automated in Matlab.²¹ Using these scripts, images are filtered for noise and local background and individual vesicles identified based on their size (9–36 pixels²) and circularity (>0.2). The tracks of moving vesicles are then identified using a tracking algorithm that identifies vesicle tracks based on a maximum-probability assessment of how closely potential tracks resemble free and superdiffusive Brownian diffusion.²² The resulting tracks are analysed for the speed of vesicular movement, the duration the vesicle is present in the TIRF field, and the degree to which the particles' movement deviates from free Brownian diffusion (γ). Vesicles undergoing fusion with the plasma membrane are identified as those having a γ significantly less than that of an equivalent model population undergoing Brownian diffusion, typically $0 < \gamma < 0.873$,²³ and which undergo a decrease in fluorescence intensity over the last two time points of their tracks equivalent to a drop of at least 2.5 SDs of the vesicular intensity over the entire period the vesicle has been tracked. The assay overcomes the limitations of transwell and microscopy assays, as it differentiates actual fusion (transcytotic) events from both diffusion across endothelial junctions and transient interactions of vesicles with the basolateral plasma membrane.²³ Under control conditions, cells performed 10–30 transcytosis events during the period of observation.

2.4 Colocalization assay and immunostaining

Cells were incubated with Dil-LDL (2 µg/mL) at 37°C for 10 min and then washed twice with cold PBS. Cells were then fixed with 4% paraformaldehyde (PFA), permeabilized, and blocked. Cells were incubated with an antibody to SR-BI (Santa Cruz Biotechnology, Dallas, Texas, USA) for 1 h followed by incubation with a goat anti-rabbit secondary antibody tagged with Alexa Fluor 488 (Jackson ImmunoResearch, West Grove, PA, USA). Z-stack images were then acquired using a spinning disc confocal

microscope (Zeiss Axiovert 200 M microscope; $\times 60$ objective, numerical aperture 1.35). After setting a threshold to exclude background, colocalization of LDL with endogenous SR-BI was calculated using Volocity version 6.0.1 and reported using Manders' coefficient. Further colocalization analysis was performed using ImageJ software (NIH) with the JACoP and Bioformats plugins installed.^{24,25} The images were cropped to each individual cell and a Pearson correlation coefficient measured for each cell. To determine whether the degree of colocalization was significant, each cell's image was subject to Costes' analysis²⁴ in which the image is randomized and the Pearson coefficient was calculated for the randomized image. Costes' analysis was performed 100 times per image to produce a statistically representative sample of non-interacting molecules. In separate experiments, 50-fold excess unlabelled LDL was added at the same time as Dil-LDL to demonstrate the specificity of colocalization with SR-BI.

For immunostaining, cells were grown to confluency on coverslips and then fixed with 4% PFA, permeabilized, and blocked. Cells were incubated with antibodies to either occludin or VE-cadherin (Santa Cruz) for 1 h, followed by incubation with a goat anti-mouse Alexa Fluor 488 secondary (occludin) or a rabbit anti-goat Alexa Fluor 488 secondary (VE-Cadherin) (Jackson ImmunoResearch). Cells were stained with DAPI, mounted on slides, and imaged by spinning disk confocal microscopy on a Leica DMI6000B system (Wetzlar, Germany). Images were acquired with a z-stack interval of 0.2 μm .

For the time-lapse z-axis imaging, cells grown on coverslips were incubated with Dil-LDL (2 $\mu\text{g}/\text{mL}$) at 4°C for 10 min to allow membrane binding. A total of 20 $\mu\text{g}/\text{mL}$ fluorescein lectin (Vector Labs, Burlington, ON, Canada) was added to the media during membrane binding to identify the plasma membrane. At Time 0, cells were rinsed three times with PBS, and then a subset of coverslips was immediately fixed in 4% PFA. The other group was treated with serum-free media and placed in the incubator at 37°C for 30 min, then washed 3 $\times$ with PBS and fixed. For this time point, 20 $\mu\text{g}/\text{mL}$ lectin was added during the final 10 min of LDL uptake followed by fixation. Cells were stained with DAPI, mounted on slides, and imaged by spinning disk confocal microscopy (Zeiss Axiovert 200 M). Images were acquired with a z-stack interval of 0.3 μm . LDL localization was quantified using ImageJ by first measuring the total LDL signal (red) in the image after background subtraction. The proportion of LDL that was intracellular was then calculated by subtracting the LDL signal in contact with the apical membrane [as identified by the lectin (green)] and dividing the remainder (the intracellular signal) by the total signal. Images and quantification are representative of four independent experiments; in each experiment, 10 high-power fields were acquired per condition.

2.5 siRNA and overexpression

To overexpress SR-BI, cells were transfected with SR-BI-GFP²⁶ or GFP alone using Eugene HD (Promega, Madison, WI, USA) in accordance with the manufacturer's instructions. For transcytosis measurement, only cells expressing SR-BI-GFP or GFP alone were imaged by TIRF.

For SR-BI depletion, SR-BI siRNA (Qiagen, Valencia, CA, USA; Hs SCARB1 7, Cat. No. S102777201) or scrambled control siRNA (Qiagen) was transfected into cells using HiPerfect (Qiagen). Transcytosis was quantified 36 h later. Knockdown was confirmed by western blot. Lysates were analysed by SDS-PAGE using 12% polyacrylamide gel. Proteins were transferred to nitrocellulose membranes and then blocked and probed overnight at 4°C with the primary antibody. Blots were washed in PBS with Tween 20, and blots were incubated with horseradish peroxidase-conjugated secondary antibodies for 1 h, washed, and then visualized by enhanced chemiluminescence (Amersham, GE Healthcare, Mississauga, ON, Canada).

2.6 PCSK9 and LDLR

HEK293 cells grown in Dulbecco's Modified Eagle Medium (DMEM) were transiently transfected with a plasmid vector encoding human PCSK9²⁷

using FuGENE[®] HD Transfection Reagent according to manufacturer's instructions. DMEM was changed 12 h after transfection; 24 h later, cell culture supernatants were harvested and incubated with confluent primary HCAEC monolayers for a minimum of 6 h. Treated monolayers were then analysed for LDL transcytosis or harvested for immunoblotting as described above. Antibodies for LDLR and SR-BI were from Santa Cruz.

2.7 Ex vivo assay

Mouse work was performed in accordance with the Guide for the Care and Use of Laboratory Animals (Institute of Laboratory Animal Resources, seventh edition, 2011) and was approved by the Animal Care and Use Committee of the local government authorities (AUP# 896.18 and #542). Aortic arches were isolated from male C57BL/6 male mice (age 8–12 weeks, Figure 1), or male B6;129S2-Scarb1^{tm1Kri}/J (SR-BI^{-/-}) and wild-type (WT) littermate control mice (11–14 weeks old, Figure 7), all purchased from Jackson Laboratories (Bar Harbor, MN, USA). Mice were fed a standard chow diet and were given free access to food and water. Mice were sacrificed by isoflurane overdose, and then the chest cavity was opened. Three millilitres of 10 U/mL heparin were injected into the left ventricle, and the inferior vena cava was cut for exsanguination. The mouse was placed on a dissecting microscope stage, and the aorta was carefully cleaned and the branches off the arch were tied *in situ*. The aorta was then removed by cutting above the heart and before the renal branches and then mounted on a silicone-coated dish for cannulation. Vessels were cannulated with PE50 tubing and perfused for 3 h at a rate of 1 mL/h using MOPS [3-(N-morpholino)propanesulfonic acid] buffer containing biotinylated dextran (70 kDa, 100 $\mu\text{g}/\text{mL}$), as well as LDL (0.71 mg/mL) or PAF (50 nmol/L) (see Figure 1). In some experiments (Figure 7), perfusion was performed with biotinylated LDL (Alpha Diagnostics International, San Antonio, TX, USA) to increase the sensitivity of the assay and the ability to detect a difference between knockout and wild-type vessels. Vessels were then flushed with MOPS for 15 min, followed by perfusion with 4% PFA for 15 min. Vessels were subsequently immersed in 4% PFA for 15 min and then cryofixed.

For the perfusion experiment with LDL and dextran (Figure 1), the ascending aorta was sectioned sequentially and stained using Nile Red and streptavidin-Cy3 (Sigma); Nile Red was chosen as it detects both free and esterified cholesterol and is more sensitive than Oil Red O.²⁸ Vessels were then stained with DAPI and imaged by spinning disk confocal microscopy (Zeiss Axiovert 200 M microscope; Zeiss, Jena, Germany).

For the perfusion experiment using aortas from SR-BI^{-/-} and wild-type (WT) littermate animals (Figure 7), the aortic arch was sequentially sectioned between the second and third branches at a thickness of 5 μm . Slides were air-dried for 2 h before staining. OCT medium was removed by submerging slides in PBS for 20 min. Slides were stained with 20 μL of Streptavidin-Cy3 (1:1000) in PBS for 30 min followed by 3 $\times$ wash in PBS. The arch was then stained with DAPI followed by mounting with coverslips. Image acquisition was done by confocal microscopy on a Leica DMI6000 B microscope with settings kept constant between groups. A total of 30 images were randomly acquired from each vessel at a z-stack interval of 0.2 μm . All perfusion, sectioning, staining, and image analysis were performed as four separate experiments, each containing one WT and one SR-BI^{-/-} mouse. Images were quantified using ImageJ for Cy3 (red) fluorescence; all signal above background was measured. For each experiment, the same background correction was applied to images from both the knockout and matching wild-type aorta. Results are presented as the red fluorescence intensity normalized to wild type.

2.8 Statistics

Statistical analysis was performed by use of GraphPad Prism software (GraphPad Prism 5.0; GraphPad Software Inc., La Jolla, CA, USA), and a combination of Student's *t*-tests and one-sample *t*-tests was used. Data are presented as mean $\pm$ SEM. All experiments were performed at least on four different occasions.

Figure 1 LDL accumulates in the ascending aorta by transcytosis. (A) Aortas from C57BL/6 mice were perfused *ex vivo* with LDL simultaneously with biotinylated dextran of a smaller molecular radius. Sequential sections, which were stained with Nile Red or streptavidin-Cy3 (to detect cholesterol and dextran, respectively), demonstrate that LDL accumulates in the sub-endothelial intima in the absence of dextran (inset). Arrows indicate cholesterol deposits (in red); nuclei are stained with DAPI (blue). Size bars, 10 μm . (B) No cholesterol is detected in vessels not perfused with LDL. Size bar, 13 μm . (C) As a positive control, aortas were perfused with dextran alone (i.e. no LDL) in the absence (top panels) or presence (bottom panels) of 50 nM PAF, which induces inter-endothelial gaps. Dextran (in red) accumulated in PAF-treated vessels (inset); nuclei are in blue. Arrows indicate the endothelium; dotted lines indicate the vessel wall. Size bars, 80 μm .

3. Results

3.1 LDL crosses the endothelium by transcytosis *ex vivo*

We first wanted to establish that LDL transcytosis occurs in intact arteries. Aortas were isolated from C57BL/6 mice, perfused with LDL and biotinylated dextran (of a smaller molecular radius than LDL), and sequential sections were then stained with Nile Red and Streptavidin-Cy3 to detect accumulation of cholesterol and dextran, respectively. Perfusion with LDL for several hours resulted in sub-endothelial cholesterol deposits (Figure 1A) that were absent in control vessels perfused without LDL (Figure 1B). No intimal dextran was

observed despite its smaller size whether perfused at the same time as LDL (right-hand panel, Figure 1A) or when alone in the perfusate (Figure 1C). This was not due to impaired retention of the dextran, since aortas perfused with PAF, a known inducer of endothelial leak,²⁹ demonstrated an obvious increase in accumulation of intimal dextran compared with control (Figure 1C). Taken together, these data suggest that LDL reached the sub-endothelial intima by transcytosis.

3.2 LDL transcytosis is performed by primary HCAECs *in vitro*

Primary HCAECs were grown in culture and exhibited a typical cobblestone morphology under phase contrast microscopy. In addition, they

Figure 2 Primary HCAECs in culture retain endothelial characteristics and internalize LDL in a polarized fashion. (A) Coronary artery endothelial cells exhibit cobblestone morphology under phase contrast microscopy (left panel), as well as continuous rings of occludin (middle) and VE-cadherin (right panel) as determined by immunostaining. Images are z-stack projections and are representative of >5 independent experiments; nuclei are stained with DAPI (blue). Size bar is 19 μm . (B) Confluent HCAECs were pulsed with Dil-conjugated LDL and then rinsed to remove unbound ligand. After incubation at 37°C, monolayers were fixed at progressive time points and imaged by confocal microscopy. The apical membrane is labelled with a fluorescein-tagged lectin (green). Long images are side views (yz-axis) of separate cells with corresponding xy images shown at left for reference. xy images were taken 5 slices from the top of the cell in both cases, and white arrows denote vesicles containing LDL. Note the apical localization of LDL (red) at time zero (indicated by colocalization with the green lectin) followed by its movement towards the basal membrane (dotted line) 30 min later. Quantification is shown at right. Scale bar represents 3 μm , nuclei are stained with DAPI (blue); $n = 4$ independent experiments; $*P < 0.001$ by Student's *t*-test for 0 vs. 30 min.

displayed continuous rings of tight junctions, as demonstrated by immunofluorescence for occludin, as well as adherens junctions, as shown by plasmalemmal staining for VE-cadherin (Figure 2A); these suggest that cell polarity is maintained. When confluent monolayers of these cells were pulsed with fluorophore-tagged LDL, we observed the progression of the LDL from the apical membrane downwards over time (Figure 2B). To confirm that actual LDL transcytosis occurs in HCAECs, we adapted our published assay for albumin transcytosis.¹ Briefly, HCAECs seeded on transwells were exposed to biotin-LDL at 4°C, a temperature at which binding occurs but internalization is inhibited. Cells were rinsed to remove unbound LDL and placed at 37°C to allow transcytosis to begin; the binding and rinsing step is performed to decrease confounding from paracellular leak. Passage of LDL across the endothelial monolayer was measured using a highly sensitive ELISA (Figure 3A). As a control, we repeated the transcytosis assay in the presence of excess unlabelled LDL and found that permeability to biotin-LDL was significantly attenuated (Figure 3B) consistent with competition for binding sites. To prove that the assay is not affected by endothelial gaps, we added PAF to the cells. As expected, PAF significantly increased endothelial permeability to

fluorescein (Figure 3C); however, it had no effect on the flux of membrane-bound LDL (Figure 3D).

3.3 Development of a novel assay to study LDL transcytosis

While the transwell assay suggests that LDL transcytosis occurs *in vitro*, the method is cumbersome and not amenable to the molecular dissection of transcytosis. Transwell assays are affected by edge effects, low transfection efficiency (particularly in primary cells), and the potential of constant confounding from intercellular gaps. To overcome these issues, we developed a novel assay using TIRF microscopy.²¹ This technique selectively visualizes the basal (bottom) surface of cells (e.g. 100–150 nm or so), excluding overlying noise from the rest of the cytosol. Dil-labeled (fluorescent) LDL is added apically to confluent endothelial monolayers, and the fusion of LDL-containing vesicles with the basolateral membrane is observed in real time (see Supplementary material online, Movie S1 and Figure 4A and B). Quantification of fusion events is performed in a blinded and automated fashion. We validated this new

Figure 3 LDL transcytosis is performed by HCAECs *in vitro*. (A) Standard curve for ELISA for biotin-LDL showing linearity and high sensitivity; y-axis is optical density at 450 nm (see Methods). (B) Transwells coated with confluent HCAECs were allowed to bind biotin-LDL and then rinsed to remove unbound ligand. Two hours later, LDL in the lower chamber of the transwell was measured by ELISA. Addition of 50-fold excess unlabelled LDL (Comp) reduced biotin-LDL transcytosis, $*P < 0.05$ by Student's *t*-test. $n = 7$ experiments. (C and D) The transwell assay for LDL transcytosis is not affected by paracellular leak. Administration of PAF (10 μ M) increased the permeability of endothelial monolayers to Na-fluorescein (C) but did not affect LDL transcytosis (D) compared with untreated cells (Ctrl); $*P < 0.05$ by one-sample *t*-test, $n = 6$ experiments.

method in a number of ways. First, to prove that the assay is not confounded by endocytic traffic (i.e. vesicles moving in and out of the TIRF zone), we reduced the penetration depth of the TIRF laser to 75 nm; as expected, this had no effect on the number of transcytosis events detected (Figure 4C). As with the transwell method, the TIRF assay was not affected by an increase in paracellular leak induced by PAF (Figure 4D). In addition, an excess of unlabelled ligand caused an almost four-fold decrease in LDL transcytosis (Figure 4E), consistent with a receptor-mediated process rather than simple diffusion across paracellular gaps. Finally, incubation with Dyngo4A,³⁰ a specific chemical inhibitor of the GTPase dynamin (Figure 4F), essentially abrogated the process.

3.4 Degradation of the LDLR by PCSK9 does not decrease LDL transcytosis

We then wished to take advantage of this assay to identify which receptor was capable of mediating LDL transcytosis. To determine whether

the high-affinity LDL receptor (LDLR) is required, we exposed coronary endothelial cells to exogenous proprotein convertase subtilisin/kexin type 9 (PCSK9)²⁷ which mediates degradation of the receptor. As expected, PCSK9-treated cells demonstrated almost complete loss of LDLR (Figure 5A); of note, levels of another cell surface receptor, SR-BI, were unchanged. Under these conditions, LDL transcytosis was not affected (Figure 5B).

3.5 SR-BI mediates LDL transcytosis across HCAECs

Since LDLR does not mediate LDL transcytosis, we turned our attention elsewhere. The scavenger receptor SR-BI was described some time ago to bind native LDL,³¹ but the physiological significance of this interaction is unknown. Interestingly, SR-BI is expressed in coronary endothelium and is capable of bona fide endocytosis.³² We observed partial colocalization of SR-BI and Dil-LDL (Pearson's coefficient $R^2: 0.228 \pm 0.029$; Costes' $R^2: 0.0025 \pm 0.0009$; Manders' coefficient = 0.577 ± 0.073 ; Figure 6A), which was >100 -fold higher than that expected by chance. Addition of 50-fold excess unlabelled LDL essentially abrogated the colocalization. Overexpression of SR-BI resulted in a 50% increase in LDL transcytosis (Figure 6B), while even incomplete knockdown by siRNA had the opposite effect, resulting in a decrease in transcytosis by $\sim 40\%$ (Figure 6C and D). Lastly, addition of excess HDL, the canonical ligand for SR-BI, significantly attenuated transcytosis of LDL (Figure 6E), suggesting competition for the receptor.

3.6 Aortas from SR-BI-deficient mice accumulate less LDL during ex vivo perfusion

Finally, to validate our *in vitro* findings, we isolated the aortas from SR-BI-deficient mice and their wild-type littermates, followed by perfusion for 3 h with biotinylated LDL. Sequential sections of the vessels were then examined and quantified for intimal accumulation of LDL. Consistent with the cell culture experiments, deficiency of SR-BI caused a $>40\%$ reduction in aortic LDL accumulation (Figure 7). Thus, SR-BI contributes to LDL transcytosis both *in vitro* and *ex vivo*.

4. Discussion

Our study addresses the critical question of how LDL crosses the endothelium to accumulate in the intima, a necessary first step in the pathogenesis of atherosclerosis. We provide evidence in both intact vessels and *in vitro* that LDL can cross endothelial cells by transcytosis. This is consistent with the observation that early atherosclerotic lesions occur despite apparently healthy overlying endothelium.⁵

Elucidation of the molecular regulation of transcytosis is best determined *in vitro*, yet a major technical barrier is the difficulty in distinguishing it from paracellular leak, since manipulation of an endothelial monolayer by pharmacological or molecular approaches often perturbs barrier integrity.¹

Accordingly, we devised a means of quantifying transcytosis by single cells in real time, reasoning that this would avoid confounding by paracellular leak or low transfection efficiency in primary cells. After validating this technique, we then sought to identify the receptor involved in LDL transcytosis.

While the best characterized receptor for LDL is its high affinity receptor (LDLR), our data suggest that it is unlikely to be required for LDL transcytosis; specifically, almost total degradation of LDLR by PCSK9³³ did not impair transcytosis. The lack of involvement of

Figure 6 SR-BI mediates LDL transcytosis. (A) Coronary artery endothelial cells were incubated with Dil-LDL then fixed and immunostained for SR-BI. Z-stack images were then acquired using a spinning disk confocal microscope. Dil-LDL colocalized partially with SR-BI (arrows). Pearson coefficient R^2 : 0.228 ± 0.029 ; Costes' R^2 : 0.0025 ± 0.0009 ; Manders' coefficient = 0.577 ± 0.073 . Addition of 50-fold unlabelled LDL essentially ablated colocalization of Dil-LDL and SR-BI (Pearson R^2 = 0.125 ± 0.015 ; Manders' coefficient = 0.107 ± 0.025 ; $P < 0.001$ by Student's t -test for comparison of the Manders' coefficients, $P = 0.01$ by Student's t -test for Pearson's coefficients); $n = 8$ experiments with Dil-LDL alone and $n = 4$ with excess unlabelled LDL. White scale bar is $3 \mu\text{m}$. (B) Cells transfected with SR-BI-GFP exhibited increased LDL transcytosis compared with cells transfected with GFP alone (Control), as measured by TIRF, $*P < 0.01$ by one-sample t -test, $n = 4$ experiments. (C) Cells were transfected with siRNA to SR-BI or scrambled control (Control) followed by western blotting. Levels of LDLR were probed as a control for off-target effects while α -actinin was the loading control. (D) Even partial depletion of SR-BI by siRNA caused a significant drop in LDL transcytosis compared with scrambled control (scRNA) as measured by TIRF; $*P < 0.05$ by one-sample t -test, $n = 5$ experiments. (E) Addition of 50-fold unlabelled HDL at the same time as Dil-LDL reduced LDL transcytosis, $*P < 0.05$ by one-sample t -test, $n = 4$ experiments.

LDLR in transcytosis is perhaps unsurprising as it is known from human patients that mutations in LDLR causing marked elevations in circulating LDL nonetheless predispose to accelerated atherosclerosis.³⁴

Instead, we present multiple lines of evidence indicating that SR-BI mediates transcytosis of LDL, in addition to its previously established roles in HDL metabolism³⁵ and steroidogenesis.³⁶ This finding was unexpected given that deficiency of SR-BI has been reported to cause accelerated atherosclerosis in mice.^{36,37} However, the phenotype of these animals is dominated by expression of the receptor in the liver^{37,38} and macrophages.^{39,40} Our data suggest that endothelial SR-BI has a more complex role and may contribute to early atherosclerotic lesions. This apparent duality of SR-BI is not without precedent, as data suggest that in such early lesions, SR-BI expression in bone marrow-derived cells is also pathogenic.⁴¹ It is also possible that the pro-atherogenic contributions of SR-BI in the arterial endothelium are nullified by the receptor's beneficial role in the liver and macrophages. Our study used multiple approaches to implicate SR-BI *in vitro*, followed by validation of these results *ex vivo*; further work will be required to confirm these observations *in vivo*. However, it is reassuring that our observation that LDL and HDL can compete for binding to SR-BI is mirrored by an earlier report by Rohrer *et al.*,³⁵ who found that LDL could decrease binding of HDL to aortic endothelium.

These findings suggest the possibility that very low-density lipoprotein (VLDL), another reported ligand for SR-BI,⁴² could also inhibit LDL transcytosis, but this remains to be established experimentally.

Our data imply that in addition to SR-BI, other receptors are likely to contribute to LDL transcytosis. The endothelium expresses other scavenger receptors including CD36, SR-AI/II, LOX-1, and SREC.^{43,44} Of these,⁴⁵ only SR-BI and CD36 have been reported to bind native LDL^{31,46} and to be capable of performing bona fide receptor-mediated endocytosis.^{32,47} Intriguingly, the majority of both SR-BI and CD36 is found in lipid rafts or caveolae. Thus, the role of CD-36 in LDL transcytosis is the subject of ongoing work in our laboratory. Furthermore, although the receptor-mediated endocytosis of LDL by LDLR is well characterized,⁴⁸ the relationship, if any, of endothelial retention of cholesterol to cholesterol transcytosis remains uncertain. We believe that our development of a dynamic and facile assay for quantifying LDL transcytosis by individual cells will greatly facilitate the elucidation of this important issue.

In closing, given that the deposition of sub-endothelial LDL is the first step in atherosclerosis, it is startling how little is known about the process. We have devised a novel assay to quantify LDL transcytosis; our data implicate transcytosis as an important route and point to SR-BI as a contributing receptor. This assay is likely to prove useful in the study of

Figure 7 Deficiency of SR-BI reduces LDL transcytosis *ex vivo*. Aortas from *SR-BI*-deficient mice and wild-type (WT) littermates were perfused *ex vivo* with biotinylated LDL (25 $\mu\text{g/mL}$). Serial sections were stained with streptavidin-Cy3, and z-stack images were then acquired by confocal microscopy under identical microscope settings. Nuclei are stained with DAPI (blue). All perfusion, sectioning, staining, and image analysis were performed as four separate experiments, each containing one WT and one *SR-BI*^{-/-} mouse. Images were quantified using ImageJ for Cy3 (red) fluorescence; all signal above background was measured. For each experiment, the same background correction was applied to images from both the knockout and matching wild-type aorta. Results are presented as the red fluorescence intensity in a *SR-BI*^{-/-} aorta normalized to its WT equivalent. More punctae of LDL are visible in WT vs. *SR-BI*^{-/-} vessels (inset, arrowheads). Quantification is shown at right. * $P < 0.01$ by one-sample t-test, four mice per group.

transcytosis of other ligands, including modified forms of LDL and other lipoproteins such as HDL.³⁵ Further study into the mechanisms of LDL transcytosis may lead to new therapeutic approaches for the prevention and treatment of atherosclerosis.

Supplementary material

Supplementary material is available at *Cardiovascular Research* online.

Acknowledgements

The authors thank Johnny Su for assistance with densitometry and Michael Woodside and Paul Paroutis of the Imaging Facility at the Hospital for Sick Children for technical assistance with TIRF.

Conflict of interest: none declared.

Funding

This work was supported by the Canadian Institutes of Health Research (102741 to N.G.S.); Canada Research Chair (216684 to N.G.S.); Foundation Leducq (13 CVD 03 to N.G.S.). S.M.A received MD/PhD funding from the CIHR and the McLaughlin Centre for Molecular Medicine.

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