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ABCA7 Downregulation Modifies Cellular Cholesterol Homeostasis and Decreases Amyloid- β Peptide Efflux in an *in vitro* Model of the Blood-Brain Barrier

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Abstract. The role of ABCA7 in brain homeostasis and Alzheimer's disease (AD) is currently under intense scrutiny, since it has been reported that polymorphisms in the *Abca7* gene and a loss of function of the protein are closely linked to excessive accumulation of amyloid peptides and disturbed cholesterol homeostasis. The blood-brain barrier (BBB), which isolates the brain from the blood compartment, is involved in both of these processes. We therefore hypothesized that ABCA7 downregulation might affect cholesterol and amyloid exchanges at the BBB. Using siRNA and primary cultures of mouse endothelial cells purified from brain microvessels and seeded on Transwell[®] inserts, we investigated the role of ABCA7 in cholesterol and amyloid exchanges across the BBB. Our results showed that a decrease in ABCA7 expression at the BBB provokes *in vitro* a reduction in ABCA1 expression and a decrease in APOE secretion. *In vitro*, these decreases reduce cholesterol exchange across the BBB, particularly for high-density lipoproteins and ApoA-I particles. When ABCA7 was absent, we observed a reduction in A β peptide basolateral-to-apical transport in the presence of ApoA-I, with non-significant changes in the expression levels of *Rage*, *Lrp1*, *Abcb1*, *Abcc1*, and *Abcg2*. Our study in murine BBB model highlighted a putative new role for ABCA7 in AD *via* the protein's involvement in cholesterol metabolism and amyloid clearance at the BBB.

Keywords: ABCA7, A β peptides, Alzheimer's disease, blood-brain barrier, cholesterol metabolism

INTRODUCTION

In developed countries, Alzheimer's disease (AD) is the most prevalent form of dementia in the

elderly [1]. One of the pathological hallmarks of this neurodegenerative disease is the accumulation of amyloid- β peptides (A β) in brain parenchyma and cerebral blood vessel walls [2]. It has been clearly demonstrated that the majority of late-onset cases of AD are associated with several risk factors linked to cholesterol metabolism and transport [3, 4]. For example, the apolipoprotein E $\varepsilon 4$ (*APOE* $\varepsilon 4$) allele is the strongest genetic risk factor for sporadic,

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late-onset AD. The ApoE4 protein is the main cholesterol carrier in brain tissue and is thus involved in regulation of cholesterol metabolism in the brain. The *APOE* ϵ 4 allele is also known to promote A β aggregation [5]. In contrast, the *APOE* ϵ 2 allele is associated with protection against AD. Moreover, a high-cholesterol diet and thus hypercholesterolemia are associated with AD. In preclinical studies of murine AD models, the hypercholesterolemia induced by a cholesterol-enriched diet was associated with an elevated level of A β peptides in the brain [6], whereas A β deposition was lower when mice were treated with cholesterol synthesis inhibitors [7]. Thus, the level of cholesterol in the blood can modify A β peptide levels in the brain [6, 8, 9]. These effects are seen despite the presence of the blood-brain barrier (BBB), which restricts the passage of several molecules (including cholesterol) from the circulation into the brain. The key structural components of the BBB are the brain endothelial cells (BECs), which are sealed together by tight junctions. Therefore, the passage of molecules across the BBB is facilitated by transporters that are expressed at the BECs' luminal and/or abluminal membranes.

These transporters include members of the ATP-binding cassette (ABC) superfamily, which use ATP hydrolysis to transport a wide range of molecules against their concentration gradients and thus participate in the maintenance of the brain's microenvironment [10]. At the BBB, ABCA1 contributes to cholesterol efflux [11, 12] by transferring lipids to apolipoproteins and initiating high-density lipoprotein (HDL) formation [13]. Other ABC transporters involved in the lipidation/maturation of lipoproteins are also expressed at the BBB [11, 14], although their exact roles require further characterization [15, 16]. Thus, ABC transporters are major players in cholesterol efflux and lipoprotein metabolism [13, 17], and confer the BBB with an important role in the regulation of brain cholesterol levels [18].

Interestingly, genome-wide association studies (GWASs) of AD have never identified single-nucleotide polymorphisms in members of the ABC transporter superfamily as possible genetic factors – albeit with the exception of *ABCA7*, for which loss of function is associated with an increased risk of developing AD [19, 20].

In the periphery, *ABCA7* participates in cholesterol efflux to ApoE discs in kidney cells [21], although cholesterol release is not affected in macrophages from *Abca7* knockout (KO) mice [22]. In the brain, *Abca7* is strongly expressed in several

different cell types but is particularly present in microglia [23]. It has been demonstrated that *ABCA7* is involved in phagocytosis and in A β clearance by microglia [24]. In neurons, *ABCA7* deficiency alters the lipid profile and promotes A β peptide production through increased BACE 1 expression [25]. Despite the fact that *ABCA7* expression has been detected in an *in vitro* model of the BBB [14], its function at this barrier has not previously been studied.

By using RNA silencing, a mouse-based *in vitro* model of the BBB [26–28], the objective of the present study was to characterize the effect of *ABCA7* deficiency on cholesterol and amyloid exchanges at the BBB.

Our *in vitro* results show that *ABCA7* deficiency induced a decrease in *ABCA1* and *APOE* expression, as well as a decrease in cholesterol efflux towards ApoA-I and HDL particles. When *ABCA7* is downregulated, a decrease in A β peptide efflux across the BBB is observed in the presence of ApoA-I. Taken as a whole, our results highlight a role for *ABCA7* in brain cholesterol metabolism and in A β peptide clearance across the BBB. The results may also explain (at least in part) the association between *ABCA7* and AD.

MATERIALS AND METHODS

Chemicals

siRNA and 5X siRNA buffer were purchased from Dharmacon (Velizy-Villacoublay, France). Lyophilized siRNA was reconstituted in 1X siRNA buffer at a concentration of 5 μ M. ApoA-I and HDL were purchased from VWR (Fontenay-sous-Bois, France). Transfection reagent Ribocellin, ApoE2 and ApoE4 were purchased from Clinisciences (Nanterre, France). Lyophilized ApoE2 and E4 were reconstituted in sterile water at a concentration of 1 mg/mL. Lucifer yellow (LY), FITC-inulin and bovine serum albumin (BSA) were purchased from Sigma-Aldrich (Lyon, France). [3 H]-cholesterol (40–60 μ Ci/mmol) was purchased from Biotrend (Köln, Germany) and [14 C]-sucrose was bought from Perkin Elmer Life and Analytical Sciences (Waltham, MA, USA). Fluorescent A β ₄₀-Cy5 labeled was purchased from Phoenix Pharmaceuticals (Strasbourg, France).

Cell culture reagents

Most reagents were purchased from Sigma-Aldrich: HEPES 1 M, amino acids 50X, vitamins

100X, tosyl-lysine-chloromethyl-ketone (TLCK), Hanks balanced saline solution (HBSS) 1X, dextran and basic fibroblast growth factor (bFGF). L-glutamine was obtained from Merck (Fontenay-sous-Bois, France). Dulbecco's modified Eagle's medium (DMEM) was purchased from Life Technologies (Illkirch, France). DNase I and collagenase/dispase were purchased from Roche Diagnostics (Meylan, France). Matrigel was purchased from BD Biosciences (Le Pont de Claix, France) and Transwell® polyester membrane inserts (pore size: 0.4 µm for all experiments) were obtained from Corning (Corning, NY, USA). Newborn calf serum (CS) was obtained from Integro B.V. (Zaandam, The Netherlands).

Animal experiments

In accordance with French legislation, the animal facility at the University of Artois has been approved by the local authorities (reference: B62-498-5). In compliance with European Union legislation (Directive 2010/63/EU), all procedures were approved by the local Animal Care and Use Committee (Comité d'Éthique en Expérimentation Animale du Nord-Pas-De-Calais; reference: C2EA 75) and the French Ministry of Research (reference: 2015090115412152). For *in vitro* experiments, 4- to 6-week-old C57BL/6 mice were provided by Janvier Labs (Le Genest-Saint-Isle, France).

Cell culture

The *in vitro* BBB model consisted of a primary monoculture of mouse brain capillary endothelial cells (MBCECs) seeded on filters coated with Matrigel® [26–28]. Briefly, after removing the meninges and the outer vessels, cortices were homogenized in HBSS containing 10 mM HEPES and 0.1% BSA (WBB buffer) using a Dounce homogenizer. The resulting homogenate was mixed with 30% dextran (v/v, molecular weight 100,000–200,000) in HBSS and then centrifuged to discard neural cells. The pellet containing the vascular component was resuspended in HBSS and then filtered through a 59 µm nylon mesh, in order to retain larger vessels on the mesh surface while capillaries passed through. The resulting capillary-enriched fraction was digested with collagenase/dispase (2 mg/mL) for 33 min at 37°C in WBB by supplemented with 10 µg/mL DNase and 0.147 µg/mL TLCK. Next, the resulting capillary suspensions were seeded on

Matrigel®-coated inserts (51,000 digested capillaries/cm²) and cultured in DMEM supplemented with 15% CS, 2 mM glutamine, 1X amino acids, 1X vitamins, 50 µg/mL gentamycin and 1 ng/mL bFGF. Remaining cellular debris and red blood cells were removed by washing with medium 24 h after seeding.

siRNA treatment

Twenty-four hours after seeding, endothelial cells were transiently transfected with 40 nM mouse *Abca7* siRNA (treated condition, siABCA7 cells) or untargeted siRNA (the control condition) according to the manufacturer's instruction (protocol D-044433-03-0005, GE Healthcare, Velizy-Villacoublay, France). First, the siRNA was incubated with 1X siRNA buffer and the transfection reagent Ribocellin for 15 min at room temperature (RT). Next, warmed cell culture medium (with no added antibiotic) was added to the solution. Lastly, cells were incubated with siRNA cell culture medium supplemented with 1 ng/mL bFGF for 60 h at 37°C in a 5% CO₂ incubator. The medium was refreshed, and experiments were performed 24 h after.

BBB permeability measurements

Permeability to LY was used to check the MBCECs' tightness, as previously described [29]. Filter inserts were transferred in a 12-well plate containing warm, HEPES-buffered Ringer's solution (RH buffer, pH 7.4, HEPES 5 mM; NaHCO₃ 6 mM; NaCl 150 mM; KCl 5.2 mM; CaCl₂ 2.2 mM; MgCl₂ + 6 H₂O 1.2 mM), and a solution of LY (50 µM) was added on the luminal side of the filter. Every 20 min for 1 h, the filter inserts were transferred into another well containing fresh RH. LY fluorescence was measured with a Synergy H1 multiplate reader (Biotek, Winooski, VT, USA). Lastly, the permeability coefficient was calculated as described by Cecchelli et al.

Immunostaining

MBCECs were rinsed twice with warmed CMF-PBS and then fixed for 1 min in cold methanol/acetone (v/v). After three washes with CMF-PBS, the filters were cut from the plastic insert and (to avoid nonspecific binding) incubated for 30 min in CMF-PBS with 10% normal goat serum. The cells were then incubated with the primary antibody (anti-ZO-1; dilution: 1/200; anti-claudin 5 (dilution: 1/100; both

from Invitrogen, Cergy-Pontoise, France) diluted in CMF-PBS supplemented with 2% NGS for 1 h. After three washes in CMF-PBS with 2% NGS, the cells were incubated in the diluted secondary antibody (AlexaFluor 488/568; Invitrogen) for 60 min. Nuclei were stained with Hoechst 33258 reagent. Lastly, after a wash with CMF-PBS with 2% NGS followed by three washes with CMF-PBS, the cells were mounted using Mowiol reagent containing Dabco (Sigma-Aldrich). A Cool SNAP RS Photometrics camera (Leica Microsystems) was used to acquire the images, which were then processed using Adobe Photoshop software (Adobe Systems, San Jose, CA, USA).

mRNA extraction and reverse transcription-quantitative polymerase chain reactions

Twenty-four hours after the end of siRNA treatment and refreshment of the medium, MBCECs were rinsed twice in sterile cold phosphate buffered saline buffer and lysed using RLT lysis buffer (Qiagen, Valencia, CA, USA). Three filters were pooled for each condition. Total RNA was purified using an RNeasy total RNA extraction kit (Qiagen), according to the manufacturer's instructions. Next, each sample's purity and concentration were assessed by measuring absorbance at 260/280 nm using a Take 3 microplate reader (SynergyTM H1, Biotek). Reverse transcription was performed with 250 ng of mRNA and iScriptTM Reverse Transcription Supermix (Bio-Rad Laboratories, Marnes-la-Coquette, France), according to the manufacturer's instructions. cDNA from each sample was amplified by real-time PCR using the SsoFastTM EvaGreen Master Mix kit (Bio-Rad Laboratories) and custom-designed primers. The sequences used for primer design and primer sequences are listed in Table 1. The amplification consisted of 40 cycles with an annealing temperature of 60°C, using a CFX96 thermocycler (Bio-Rad Laboratories). The specificity of amplification was assessed by melting curve analysis. mRNA levels were quantified using the comparative CT method and normalized against the reference genes *Actb* and *Hprt*.

Protein extraction and immunoblots

MBCECs were rinsed twice in cold CMF-PBS. The cells were scraped off with UT4 lysis buffer (urea 7 M, thiourea 4 M, 4% 3-[(3-cholamidopropyl)

dimethylammonio]-1-propanesulfonate) supplemented with a protease and phosphatase inhibitor cocktail (Sigma). After a freeze/thaw cycle, the cell lysate was centrifuged to eliminate cellular debris. The protein concentration was evaluated using Bradford's method (Bio-Rad Laboratories) after dilution of the samples. Electrophoresis of 25 µg (ABCA1) or 35 µg (other proteins) of protein was performed in a 4–15% SDS-PAGE gel (Bio-Rad), followed by electrotransfer onto nitrocellulose membranes (GE Healthcare). Next, membranes were blocked in Tris-buffered saline with 0.1% Tween and 5% skimmed milk for 90 min at RT. Immunoblotting was performed by incubating membranes with dilutions of the appropriate primary antibody (anti-ABCA7; dilution: 1/1000; Proteintech, Manchester, UK; anti-ABCA1 [11, 12, 30, 31]; dilution: 1/1000; Abcam; anti-ABCG4 [32, 33]; dilution: 1/1000; Alpha Diagnostic International, San Antonio, TX, USA; anti-RAGE [34, 35], dilution: 1/1000, Abcam; anti-ABCB1 (C219) [12, 35], dilution: 1/400, Gene-Tex, Irvine, CA, USA; anti-ABCG2 (BXP-53) [32], dilution: 1/80, Santa Cruz Biotechnology, Dallas, TX, USA; anti-ABCC1 [36], dilution 1/20, Enzo Life Sciences, Farmingdale, NY, USA; anti-LRP1 [35, 37], dilution 1/200, Santa Cruz Biotechnology and anti-β-actin, dilution: 1/20000, Sigma) overnight at 4°C with slight agitation.

After rinsing, the membranes were incubated with horseradish-peroxidase-conjugated secondary antibodies (Dako and Millipore) for 1 h at RT. Immunoreactions were developed using an enhanced chemiluminescence kit (GE Healthcare) and revealed on chemiluminescence-sensitive films (GE Healthcare). The bands' relative densities were measured using TotalLab TL 100 1D Gel Analysis software (Nonlinear Dynamics, Newcastle, UK). In these experiments, each deposit represented a pool of at least eight inserts per condition. Each experiment was performed at least four times.

ApoE detection

After 60 h of incubation with siRNA, cells were incubated at 37°C with serum-free medium. After 24 h, medium was collected and then centrifuged to eliminate cellular debris. Secreted proteins were precipitated with 20% trichloroacetic acid (Sigma). Cold acetone was added and the solution thus obtained was centrifuged. Protein pellets were washed twice with cold acetone, and the remaining acetone was air-dried. Proteins were resuspended in RIPA lysis buffer

Table 1
DNA primers and accession numbers used for the qPCR experiments

mRNA	F/R	Sequence	Accession number
Abca7	F	5'-TCTGCTGTGGAAAAATTACA-3'	NM_013850.1
	R	5'-GACAGCCACTAGGATGAAGA-3'	
Abca1	F	5'-CAAGCATCTTCAGTTCATCAGC-3'	NM_013454.3
	R	5'-GAGTGTAGCAGGGACCACATAA-3'	
ApoE	F	5'-CTCCAAGTCACACAAGAAGT-3'	NM_009696.3
	R	5'-TCCTCCAGCTCCTTTTTGTAAG-3'	
Abcg4	F	5'-CTGAGGACCTTCGCAAGATGT-3'	NM_138955
	R	5'-GCTTCAGGTTGGCAGAGACCAT-3'	
Abcg1	F	5'-GGGACTCGGTCTGACACATCT-3'	NM_009593
	R	5'-AGGACCTTCTTGCTTCGTTC-3'	
Abcb1	F	5'-TGAAATGAAGCTGTTGTCTG-3'	NM_011075.2
	R	5'-AATAGTGCAGGAAAGTTTCAA-3'	
Abcg2	F	5'-GGTGTCTGTAGCAACACTTCTC-3'	NM_011920
	R	5'-TAAAGTACTGAAGCCAGGACAG-3'	
Lrp1	F	5'-CTTGATGGCACAGAGAGAGT-3'	NM_008512.2
	R	5'-CATCACACCAGTAGAGCTTG-3'	
Rage	F	5'-CCCACCTTATTAGGGACACT-3'	NM_001271424.1
	R	5'-TCAAGTCCAACACTACCGAGTC-3'	
Abcc1	F	5'-CCCTGTGTAACACTTCAAT-3'	NM_008576.3
	R	5'-CTACTTTCAGGTCACACTCTGGTG-3'	
Actb	F	5'-TGACGTTGACATCCGTAAGAC-3'	NM_007393.5
	R	5'-GCAGTAATCTCTTCTGCATCC-3'	
Hprt	F	5'-TCTGGCCTGTATCCAACACTTCG-3'	NM_013556.2
	R	5'-AAGCAGTACAGCCCCAAAATGG-3'	

supplemented with protease/phosphatase inhibitors (Sigma). The protein concentration was evaluated using Bradford's method (Bio-Rad Laboratories). Equal amounts of total secreted proteins (20 µg) were loaded onto 4–15% SDS-PAGE gel (Bio-Rad Laboratories). Immunoblots were performed as described above, using anti-ApoE primary antibodies (dilution: 1/3000, Abcam). Each experiment was performed three times.

Cholesterol efflux studies

[³H]-cholesterol (0.5 µCi/mL) was incubated with serum for 6 h at 37°C. Next, 2 mM glutamine, 1X amino acids, 1X vitamins and DMEM were added in order to obtain radiolabeled medium. Cells were incubated with this radioactive medium supplemented with siRNA solution and 1 ng/mL bFGF for 60 h. After two washes with DMEM/0.05% BSA, cells were then incubated for 24 h at 37°C with DMEM/0.05% BSA/bFGF. After this equilibration step, cells were rinsed twice, and cholesterol efflux was assessed for 7 h in the presence or absence of acceptors (20 µg/mL HDL, 35 µg/mL ApoA-I, ApoE2 and ApoE4). At the end of efflux experiments, media were collected and centrifuged (4 min, 4000 rpm, 4°C) in order to remove cellular debris. Lastly, after three washes with cold PBS, cells were lysed with PBS/1% Triton X-100. Radioactivity in

the cell lysates and media was measured in a liquid scintillation counter (Tri-Carb 2100TR). The percentage of cellular cholesterol efflux was calculated from the disintegrations per minute (DPM) using the following equations:

$$\begin{aligned} &\text{percentage of luminal efflux} \\ &= (\text{DPM}_{\text{luminal}}) \times 100 / (\text{DPM}_{\text{luminal}} \\ &+ \text{DPM}_{\text{abluminal}} + \text{DPM}_{\text{cell lysate}}) \end{aligned}$$

$$\begin{aligned} &\text{percentage of abluminal efflux} \\ &= (\text{DPM}_{\text{abluminal}}) \times 100 / (\text{DPM}_{\text{luminal}} \\ &+ \text{DPM}_{\text{abluminal}} + \text{DPM}_{\text{cell lysate}}) \end{aligned}$$

$$\begin{aligned} &\text{percentage of total efflux} \\ &= (\text{DPM}_{\text{luminal}} + \text{DPM}_{\text{abluminal}}) \\ &\times 100 / (\text{DPM}_{\text{luminal}} + \text{DPM}_{\text{abluminal}} \\ &+ \text{DPM}_{\text{cell lysate}}) \end{aligned}$$

Each assay was performed using three filters, and each experiment was run at least three times.

Quantification of cell cholesterol

After 60 h of incubation with siRNA, cells were incubated for 24 h at 37°C with serum-free cell culture medium. After this equilibration step, cells were rinsed with cold RH buffer and then lysed with 1 N NaOH. One aliquot of the cell lysate (from a single

insert) was used for the protein concentration assay using Bradford's method (Bio-Rad), and another was homogenized in chloroform/isopropanol/NP-40 (7:11:0.1) for 5 min with vigorous agitation, in order to extract lipids. After 15 min of centrifugation at 14000 rpm and 4°C, the organic phase was collected and then air-dried at 50°C for 2 h. The dried lipids were dissolved in the Cholesterol Reaction Buffer provided with a Cholesterol/Cholesteryl Ester Quantitation Kit (Calbiochem) by vortexing vigorously for 5 min. The reaction mixture was prepared according to the manufacturer instructions. Equal volumes of sample and reaction mixture were placed in a 96-well plate and incubated at 37°C for 1 h. Next, fluorescence was measured with a Synergy H1 multiplate reader (Biotek) at an excitation wavelength of 535 nm and an emission wavelength of 587 nm. The amount of cholesterol in each sample was determined with reference to a standard curve and then adjusted as a function of the protein level in each sample. We used three filters for each assay, and each experiment was run three times.

Amyloid-β peptide (Aβ) transport studies

Twenty-four hours after medium refreshment (i.e., following the end of siRNA treatment), MBCECs were rinsed with prewarmed RH buffer. Next, for apical-to-basolateral transport studies, filter inserts were transferred into 12-well plates containing 1.5 mL of RH/BSA 0.1% per well. A solution of RH/BSA 0.1% supplemented with Cy5-Aβ₄₀ (10 nM), [¹⁴C]-sucrose (Perkin Elmer Life and Analytical Sciences, Waltham, MA, USA), FITC-inulin and (or not) ApoA-I (200 nM) was then added (or not) to the apical (luminal) compartment. For basolateral-to-apical transport studies, filter inserts were transferred into 12-well plates containing 1.5 mL of Cy5-Aβ₄₀ FITC-inulin in RH/BSA 0.1%, and a solution of [¹⁴C]-sucrose supplemented (or not) with ApoA-I (200 nM) was added in the luminal compartment.

Cells were incubated at 37°C with slight agitation. Transport was measured after 3 h. BBB permeability and non-receptor-mediated transport were respectively evaluated by measuring [¹⁴C]-sucrose permeability and FITC-inulin transport. Radioactivity was measured using a liquid scintillation counter, whereas fluorescence was measured using a Synergy H1 spectrophotometer (Biotek). The percentage passage was calculated in the presence and absence of cells, in order to determine the influence of the filter and the coating on transport. The results for inulin

were used to normalize the Aβ results, as previously described [31]. The mass balance was also calculated, and the value had to be between 90% and 110%. Each assay was performed using three filters, and each experiment was run at least three times.

Statistical analysis

Results were expressed as the mean ± SEM and analyzed using a Student's T-test or a one-way analysis of variance. All statistical analyses were performed using Prism Software (GraphPad Software Inc., San Diego, CA, USA).

RESULTS

Abca7 knock-down does not affect the BBB's integrity in vitro

First, we used an RNA silencing technique to downregulate *Abca7* expression in our murine BBB model previously described [26–28]. *In vitro*, *Abca7* knock-down resulted in a 58.7 ± 4.7% reduction in mRNA expression and a 40 ± 18% reduction in protein expression, relative to the control condition (Fig. 1A–C).

We then evaluated the impact of *Abca7* downregulation on the BBB's integrity. As shown in Fig. 2A, the MBCECs' paracellular permeability to LY was similar in the siABCA7 and untargeted siRNA conditions ($0.41 \pm 0.08 \times 10^{-3} \text{ cm} \cdot \text{min}^{-1}$ and $0.40 \pm 0.07 \times 10^{-3} \text{ cm} \cdot \text{min}^{-1}$, respectively). The value in non-treated cells was $0.34 \pm 0.07 \times 10^{-3} \text{ cm} \cdot \text{min}^{-1}$. Immunofluorescent labeling of the tight junction proteins claudin-5 and ZO-1 showed strong, intact signals at the cell periphery (Fig. 2B) – suggesting that siRNA treatments did not affect the MBCECs' integrity.

Downregulation of Abca7 decreases expression of the transporters involved in cholesterol metabolism and lipid exchanges at the BBB

We then decided to investigate the expression of transporters and acceptors involved in cholesterol metabolism and lipid exchange at the BBB in siRNA-treated cells. In *Abca7*-depleted cells, mRNA expression levels of *Abca1*, *Abcg4*, and *ApoE* were low (by 53 ± 5%, 34 ± 4%, and 51 ± 4%, respectively), whereas *Abcg1* expression was not significantly different (Fig. 3A). The ABCA1 protein expression levels correlated with the mRNA

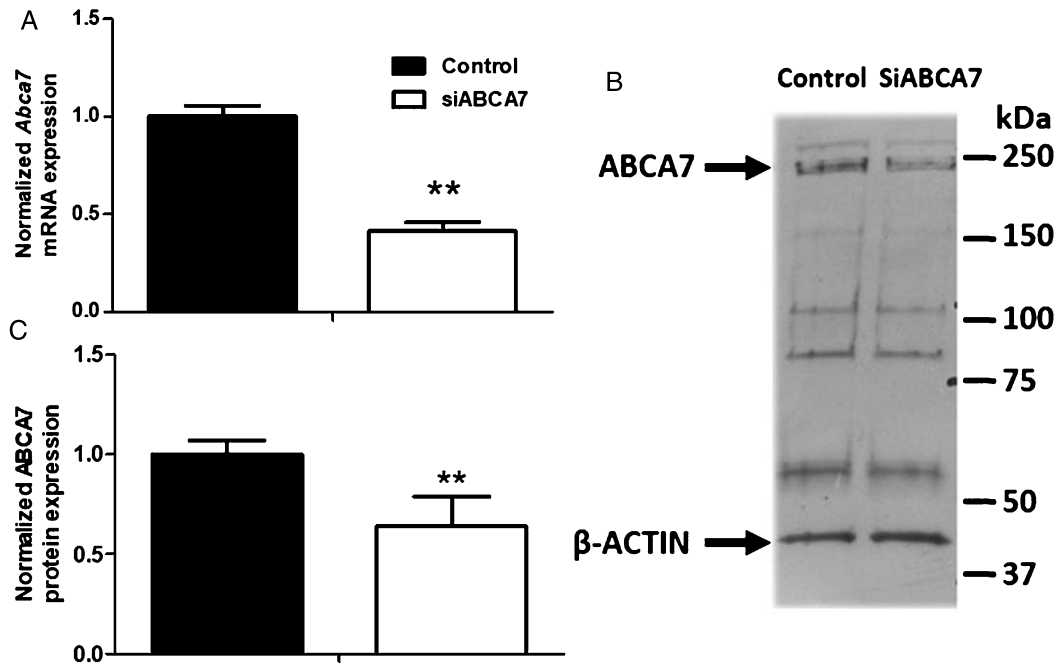


Fig. 1. Effect of siRNA treatment on ABCA7 mRNA and protein expression. A) Transcriptional expression of *Abca7* was assessed in MBCECs using the primers listed in Table 1. Each bar represents the level of mRNA expression normalized against the housekeeping genes *Actb* and *Hprt*, relative to the control condition (untargeted siRNA). Data are quoted as the mean \pm SEM of three experiments pooled from three filters. Statistical analysis: *t*-test, ***p* < 0.01. B) Protein levels were assessed by immunoblotting. ABCA7 detection gives unspecific bands as previously reported in mouse brain total extracts [24]. C) Quantification of ABCA7 expression. β -Actin was used as a loading control. Data are quoted as the mean \pm SEM of at least six experiments pooled from eleven filters. Statistical analysis: *t*-test, ***p* < 0.01.

expression levels (Fig. 3B). Indeed, a statistically significant $38 \pm 13\%$ relative reduction in the ABCA1 protein level was observed. The ABCG4 protein level was not significantly different (Fig. 3C). Furthermore, the total secreted ApoE fraction was $45 \pm 27\%$ lower but did not reach statistical significance (Fig. 3D). However, we observed that ApoE secretion was significantly lower in the abluminal media (by $63 \pm 17\%$) but not in the luminal compartment ($18 \pm 44\%$).

Thus, ABCA7 deficiency is associated with abnormally low expression of key players in HDL genesis. We therefore next assessed the ability of MBCECs to transfer cholesterol to apolipoproteins and lipoproteins.

Abca7 knock-down is associated with lower cholesterol release from the MBCEC membrane and greater cholesterol accumulation

The literature data indicate that human, porcine and bovine BBB cells release membrane cholesterol to ApoA-I, ApoE2, ApoE4, and HDL [12, 31, 38].

We therefore investigated the MBCECs' ability to efflux membrane cholesterol in basal conditions in the presence or absence of lipid acceptors. As shown in Fig. 4A, the presence of HDL, ApoE2, and ApoE4 was associated with significantly greater cholesterol release (6.8%, 6.3% and 7.9% respectively; i.e., up to 8-fold greater than the control value (0.8%) in the abluminal (brain side) compartment). The same trend was observed for cholesterol efflux to the luminal (blood side) compartment (Fig. 4B), which was promoted in the presence of each of the three lipid acceptors (11.8%, 14.6%, and 15.8% for HDL, ApoE2, and ApoE4, respectively, compared with the control value of 3.4%).

As a consequence, total cholesterol efflux was promoted in the presence of HDL, ApoE2, and ApoE4 (Fig. 4C). No significant differences between the three cholesterol acceptors were observed.

To investigate the impact of *Abca7* downregulation on cholesterol efflux, we then investigated cholesterol release in siABCA7-treated cells. As shown in Fig. 5A and B, we observed significantly lower total cholesterol efflux in the presence of ApoA-I and HDL

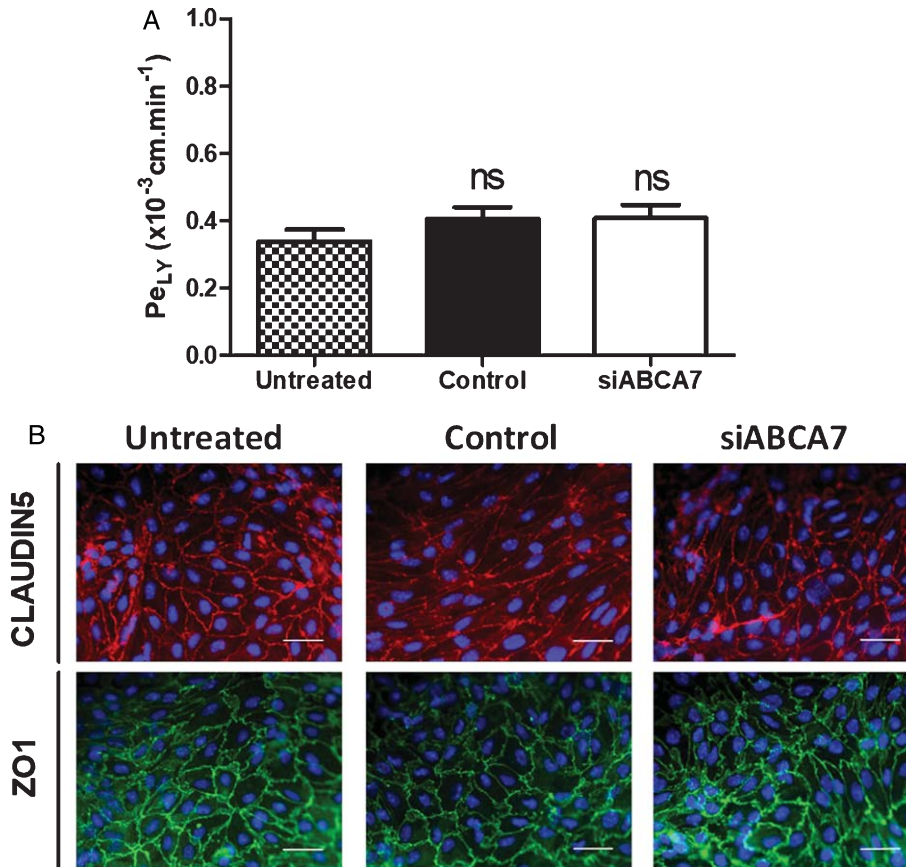


Fig. 2. Effect of siRNA treatment on permeability and tight junction integrity in a monolayer of MBCECs. Endothelial cells were incubated with medium alone (Untreated), untargeted siRNA (Control) or siRNA targeting *Abca7* mRNA (siABCA7) for 60 h. Twenty-four hours after the medium had been refreshed, permeability to the paracellular marker LY was evaluated (A). The bars represent the mean \pm SEM of five experiments, each performed with three filters per condition. Statistical analysis was performed using a one-way ANOVA followed by Dunnett's correction for multiple comparisons. ns: non-significant. B) Tight junction integrity was also assessed by immunostaining of claudin 5 (red) and ZO-1 (green). Nuclei were stained with Hoechst 33258 reagent (in blue). Scale bar: 25 μ m.

(32% and 20%, respectively, relative to the control). Cholesterol release to the luminal side was lower in the presence of ApoA-I, whereas cholesterol release to both sides was lower in the presence of HDL. These results are in line with those obtained in mouse macrophages in which *Abca1* and *Abca7* have been deleted [39].

In the presence of ApoE4 and ApoE2, total cholesterol release did not significantly differ when comparing siABCA7 cells and the control (Fig. 5C and D, respectively). However, when considering only the abluminal side, ApoE4-mediated cholesterol efflux was reduced by $16 \pm 2\%$ in siABCA7 relative to the control, whereas ApoE2-mediated cholesterol release was similar in both conditions. Hence, *Abca7* knock-down is associated with a relative decrease in cellular cholesterol efflux; this suggests that lipids accumulate in siABCA7 cells, relative to controls.

We thus evaluated the level of cholesterol accumulation in cells. As expected, we found that ABCA7 knock-down promoted a $26 \pm 9\%$ increase in cholesterol levels in MBCECs, relative to the control condition (Fig. 5E).

ABCA7 knock-down decreases basolateral-to-apical transport of A β peptide mediated by ApoA-I

Given that *Abca7*^{-/-} mice show elevated levels of amyloid deposition in the brain [24, 40, 41], we hypothesized that *Abca7* downregulation in MBCECs might impact amyloid transport across the BBB. Therefore, apical-to-basolateral and basolateral-to-apical transport assays of A β peptides were performed in our *in vitro* BBB models, as previously described [12, 31].

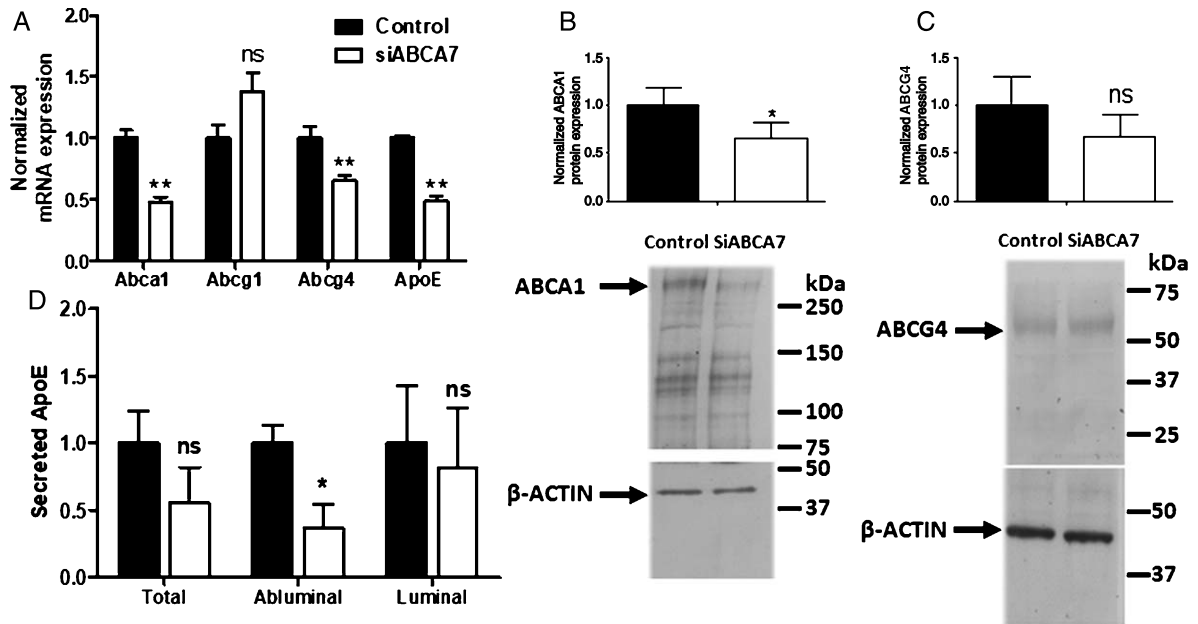


Fig. 3. Effect of *Abca7* silencing on the expression of transporters involved in cholesterol release at the BBB. A) For transcriptional expression, each bar represents the level of mRNA expression normalized against the housekeeping genes *Actb* and *Hprt*, relative to the control condition. Primers are listed in Table 1. Data are quoted as the mean \pm SEM of three experiments pooled from three filters. Protein levels of ABCA1 (B) and ABCG4 (C) were also evaluated. *Actb* was used as a loading control. Data are quoted as the mean \pm SEM of at least four independent experiments pooled from eight filters. Secreted APOE protein was measured in media recovered from luminal and abluminal compartments (D). Data are quoted as the mean \pm SEM of three independent experiments pooled from nine filters. From A to D, statistical analysis: *t*-test, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, ns, non-significant.

As shown in Fig. 6A, ABCA7 downregulation did not directly impact the influx (upper graph) or the efflux of $A\beta_{1-40}$ peptides (lower graph).

We previously reported that the presence of ApoA-I in the luminal compartment promotes $A\beta$ peptide efflux [42] and had observed in the present study that luminal ApoA-I lipidation is lower when ABCA7 is downregulated (Fig. 5A). Therefore, we next investigated $A\beta$ peptide transport in siABCA7 cells in the presence of ApoA-I in the luminal compartment. When ApoA-I was present, we observed a slight but statistically significant relative decrease in basolateral-to-apical transport of $A\beta$ (Fig. 6B, lower graph; $86.5 \pm 3.4\%$ compared with the control condition in the absence of ApoA-I). No significant differences in apical-to-basolateral transport were observed (Fig. 6B, upper graph).

To obtain additional data on the effect of *Abca7* depletion on the expression of receptors and transporters found to be involved in $A\beta$ clearance by us and others [12, 31, 37, 43–46], we used qRT-PCR and immunoblot assays to measure the mRNA and protein expression of *Abcb1*, *Abcc1*, *Abcg2*, *Lrp1*, and *Rage*. Downregulation of *Abca7* was not associated

with a difference in expression for any of these genes and proteins (Fig. 7). Despite our efforts, LRP1 and ABCC1 proteins were not detected, probably because they are expressed at very low level at the endothelial cells of the BBB as previously demonstrated [37, 47, 48].

Taken as a whole, these results suggest that ABCA7 downregulation could alter BBB cholesterol metabolism that in turn could affect amyloid clearance across this barrier.

DISCUSSION

We and others have demonstrated that the ABCB1 (also known as P-gp), ABCG2 (BCRP) and ABCC1 (MRP1) proteins expressed by endothelial cells of the BBB can restrict $A\beta$ entry into the brain and can also mediate $A\beta$ efflux into the blood stream [34, 49, 50]. ABCA1, ABCG1, and ABCG4 are more involved in cholesterol release from cellular membrane, and control cholesterol homeostasis in the brain and lipid exchanges across the BBB. These observations suggested that the modulation of ABC transporter expression might restore cholesterol metabolism and

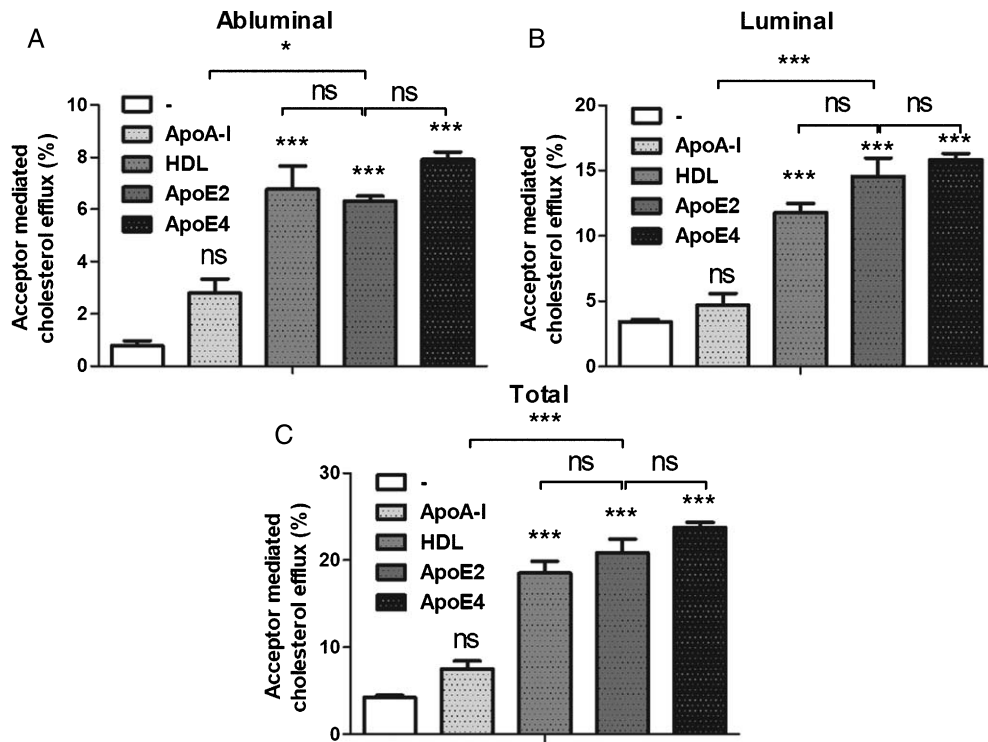


Fig. 4. Basal cholesterol efflux from MBCECs. Cholesterol efflux was assessed for 7 h in the absence of cholesterol acceptor (DMEM/BSA 0.05%, the control condition) or in the presence of the acceptor ApoA-I (35 $\mu\text{g}/\text{mL}$), HDL (20 $\mu\text{g}/\text{mL}$), ApoE2 (35 $\mu\text{g}/\text{mL}$) or ApoE4 (35 $\mu\text{g}/\text{mL}$) in the abluminal (A) and luminal (B) compartments. By summing the levels of cholesterol efflux in the two compartments, we obtained the total cholesterol efflux (C). Data are quoted as the mean \pm SEM of 3 independent experiments representing 7 inserts. Statistical analysis: a one-way ANOVA followed by Bonferroni's correction for multiple comparison, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

decrease amyloid deposition in AD patients. Promising results were obtained in AD transgenic mice in which the elevation of ABCB1 expression (by administering pregnenolone-16 α -carbonitrile [51] or caffeine [52]) promoted A β efflux across the BBB. Regarding cholesterol homeostasis, elevated ABCA1 expression in the AD mouse was associated with less amyloid deposition and less aberrant behavior [53–55]. Furthermore, we recently demonstrated that oxysterols (natural agonists of liver X receptors) were able to enhance both ABCA1 and ABCB1 expression—thus promoting cholesterol exchange and restricting A β influx across the BBB [31, 56]. Despite these interesting and promising results, however, little is known about the processes that lead to amyloid deposition and cholesterol imbalance in the central nervous system (CNS) of patients with AD. Interestingly, the only ABC transporter found to be involved in AD in GWASs is ABCA7; very few studies had focused on this protein until the publication of the GWAS results in 2011 [19]. More recent studies have tried to elucidate ABCA7's role in the

brain in health and during AD. For example, it was shown that ABCA7 does not have a significant role in the regulation of cell proliferation or neurogenesis in the adult mouse [57], and that the polymorphism identified in AD patients leads to a loss of function of ABCA7 [20]. Although *Abca7*^{-/-} mice exhibit normal sensory abilities, neurological reflexes, and motor functions [58], it is now clear that *Abca7*^{-/-} in AD transgenic mice promotes amyloid deposition. However, some researchers have explained this accumulation by the overproduction of A β peptides [21, 25, 40] whereas others have suggested that a decrease in microglial phagocytosis was involved [24, 41]. Despite the fact that *Abca7* is known to be expressed at the BBB [14], the effect of a decrease in *Abca7* expression on amyloid and cholesterol exchanges across the BBB had not previously been investigated.

Here, we used RNA silencing to decrease *Abca7* expression in an *in vitro* BBB model composed of primary MBCECs [26–28]. We did not observe any change in BBB integrity when *Abca7* mRNA and protein expression were downregulated. Given that

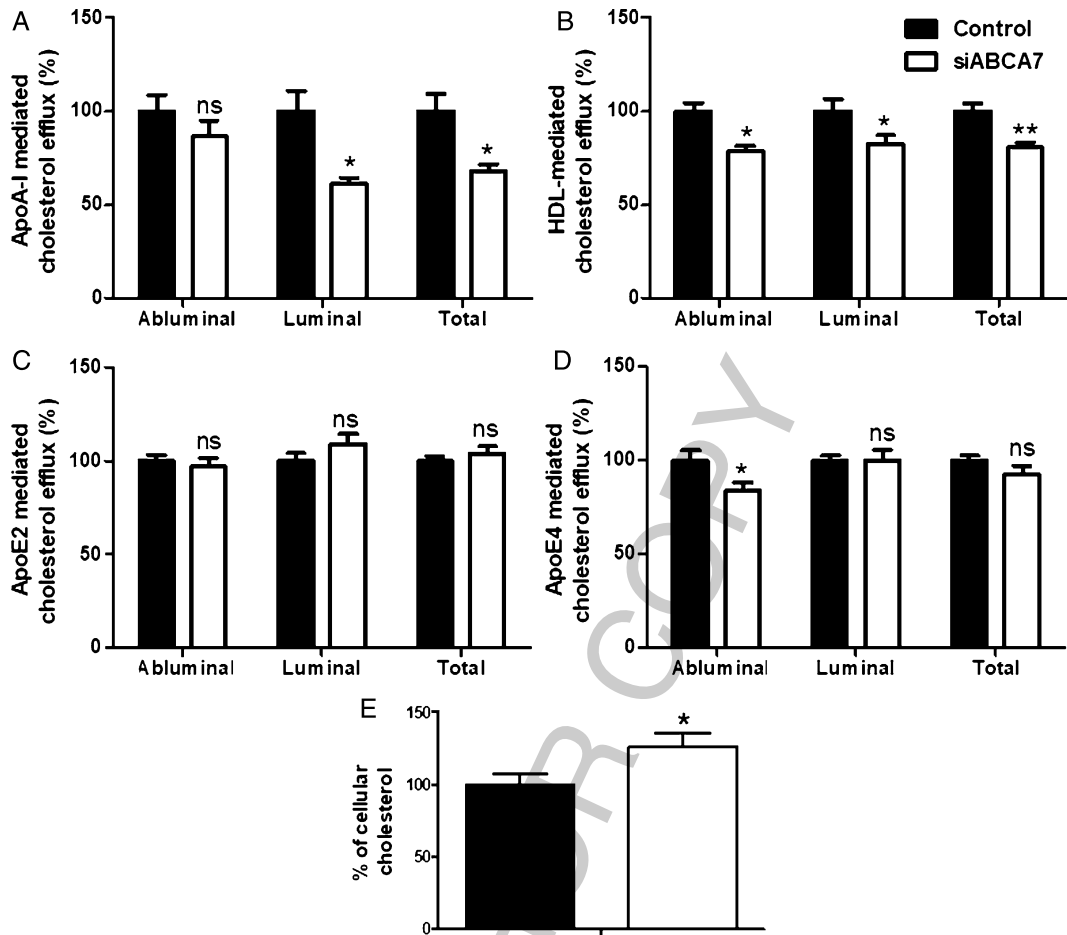


Fig. 5. Effect of *Abca7* knock-down on cholesterol release from MBCECs. As described above, cholesterol efflux was assessed for 7 h in the presence of various acceptors [ApoA-I (A), HDL (B), ApoE2 (C) and ApoE4 (D)] in control and SiABCA7 cells. Acceptors were added to the luminal and abluminal compartments. Data are quoted as the mean \pm SEM for at least 7 filters from 2 independent experiments. E) Effect of *Abca7* knock-down on the cellular cholesterol level. Cholesterol content was assessed in MBCECs after siRNA treatment. Data are quoted as the mean \pm SEM cellular cholesterol level from 9 filters per condition (3 independent experiments). Statistical analysis: *t*-test, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, ns, non-significant.

ABCA1 and ABCA7 display around 54% amino acid homology [42] and that ABCA7 generates HDL [21, 59, 60], we hypothesized that cholesterol exchange across the BBB could be impacted by *Abca7* down-regulation. Indeed, we observed a relative decrease in cholesterol release (and particularly for HDL and ApoA-I) in this context. However, we cannot be absolutely sure that this relative decrease in cholesterol efflux is directly linked to ABCA7 downregulation because 1) ABCA1 mRNA and protein levels are also lower and 2) we previously demonstrated that ABCA1 is involved in cholesterol efflux to HDL and ApoA-I in human and bovine BBB models [12, 31, 56]. We found no change in the expression of ABCG1, which has been recently identified as a key player in cholesterol metabolism at the BBB

[38]. *Abcg4* expression was significantly reduced *in vitro* when *Abca7* is deleted but protein level was not affected. ABCG4 is exclusively expressed in the CNS and generates (along with ABCA1 and ABCG1) cholesterol rich-HDL [61]. However, this transporter is also involved (along with ABCG1) in the regulation of the intracellular pool of cholesterol in astrocytes and neurons [62]. Therefore, the downregulation of ABCA1 observed in the present study might also explain the decrease in cholesterol release to ApoA-I and HDL and thus the intracellular accumulation of cholesterol.

Interestingly, we also observed lower *APOE* expression and lower APOE secretion into the abluminal compartment (mimicking the CNS) of our BBB model. In the CNS, ApoE is required for lipoprotein

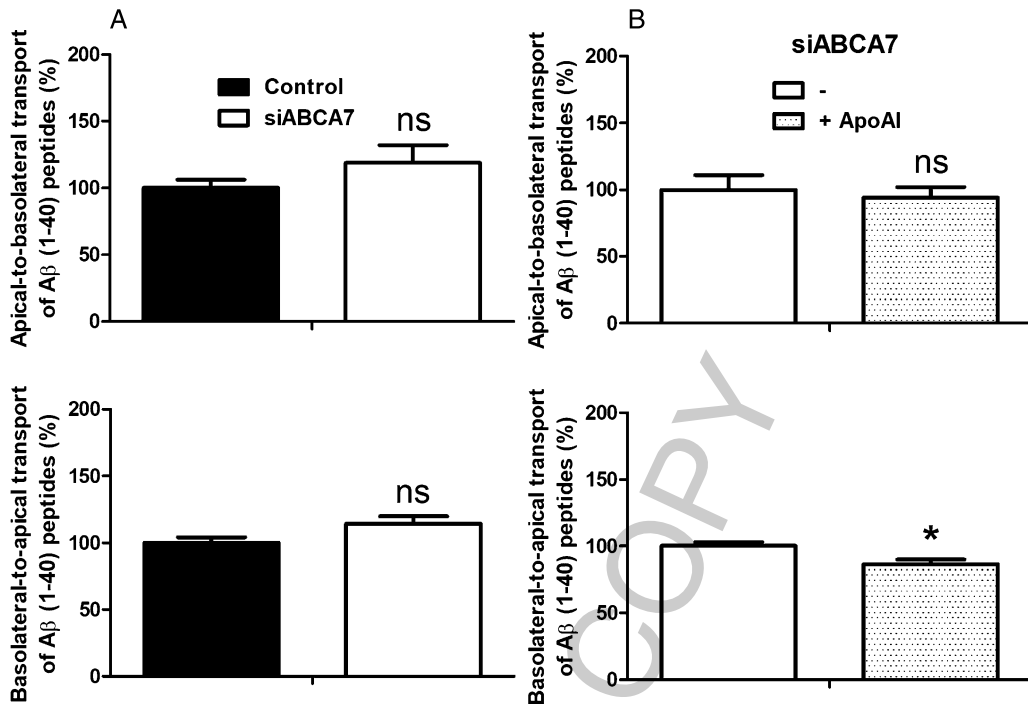


Fig. 6. Effect of *Abca7* knock-down and ApoA-I on A β peptide transport across the MBCEC monolayer. A) A β peptide transport was assessed for 3 h in MBCECs treated with untargeted siRNA (Control) or siABCA7. B) The impact of *Abca7* downregulation was then investigated in the presence or absence of ApoA-I (200 nM, in the luminal compartment). Inulin was used as a marker of non-specific vesicular transport. Inulin passage was assessed in parallel, and the data were used to normalize the A β results. In addition to inulin, paracellular integrity was assessed using radiolabeled sucrose (data not shown) for each experiment. Data are quoted as the mean \pm SEM at least seven filters per condition (2 independent experiments). Statistical analysis: *t*-test, **p* < 0.05, ns, non-significant.

uptake into neurons *via* the LDL family of receptors [63]. ApoE polymorphisms constitute the most common genetic factor involved in AD [64], and APOE mediates A β peptide elimination across the BBB. Downregulation of APOE secretion might lead to a reduction in A β transport across the BBB and therefore to greater accumulation of these peptides in the brain [64]. Noteworthy, ABCA1 deletion in AD mouse models is also associated with significant reductions in apoE level [65–67].

Our study is not the only one to investigate links between APOE, ABCA1, and ABCA7 regulations. Indeed, in *Abca7*^{-/-} mice, Kim et al. reported that ApoE levels were not affected when compared with wild-type mice [22]. ABCA1 expression was not modulated in brain. However, *in vitro*, an upregulation of ABCA7 has been reported in *Abca1*-deficient mouse fibroblasts [68] whereas *Abca7*-deficiency in mouse macrophages increases ABCA1 expression [39]. Therefore, all these data suggest that links exist between ABCA7, ABCA1, and APOE but the molecular processes remain largely unknown; they might involve sterol regulatory element binding proteins or

liver X receptors, which are key molecules regulating cholesterol and lipid metabolism [69, 70].

Taken as a whole, these data indicate that by decreasing APOE secretion and ABCA1 expression at the BBB, ABCA7 loss of function can significantly alter cholesterol exchange across the BBB and thus promote AD. This is consistent with a previous study reporting the contribution of ABCA7 in the cerebral lipid metabolism [25].

In addition to lipid metabolism, we also investigated the role of ABCA7 in amyloid transport across the BBB. When ABCA7 was downregulated, we did not observe any difference in amyloid exchange through the BBB in the absence of ApoA-I. However, as discussed above, we found that ABCA7 depletion was associated with a reduction in cholesterol release to ApoA-I; the latter is known to promote A β efflux across MBCECs [71]. Indeed, after adding ApoA-I to the luminal compartment of our experimental set-up, we observed a reduction in the basolateral-to-apical transport of A β peptide. Given that the expression of receptors and transporters known to mediate A β exchanges across the BBB did not change when

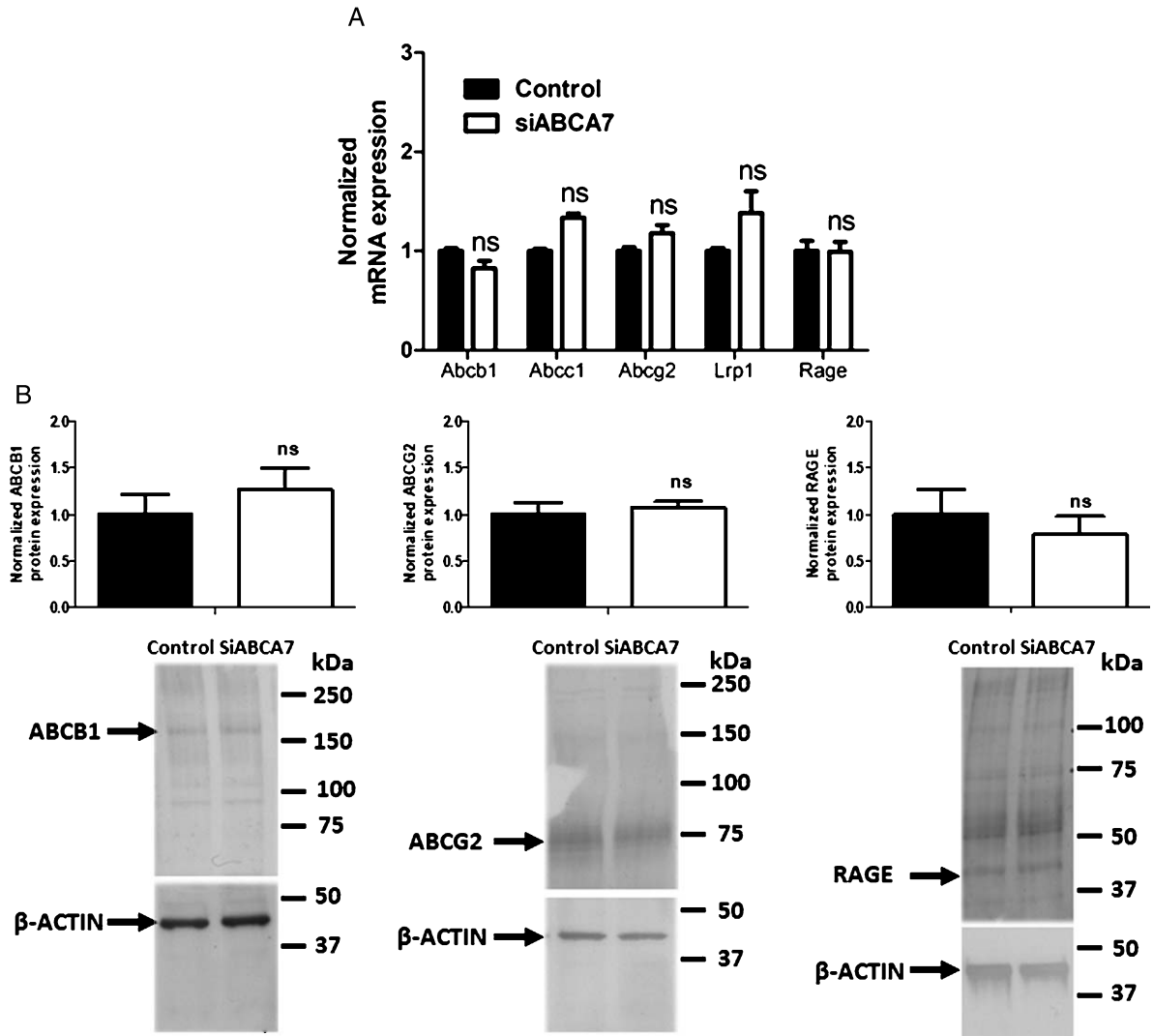


Fig. 7. Effect of *Abca7* silencing on the expression of receptors and transporters involved in A β peptide transport at the BBB. A) mRNA levels of key players of amyloid transport were measured by qRT-PCR. Each bar represents the level of mRNA expression normalized against the housekeeping genes *Actb* and *Hprt*, relative to the control condition. The primers used are listed in Table 1. Data are quoted as the mean \pm SEM of three independent experiments pooled from three filters. Statistical analysis: *t*-test; ns, non-significant. B) Protein levels were also evaluated and quantified. B-actin was used as a loading control. Data are quoted as the mean \pm SEM of at least four experiments pooled from eight filters.

ABCA7 expression was reduced, we suggest that the decrease in ABCA7 reported provokes a reduction of ABCA1 and APOE expression which in turn decreases lipoproteins lipidation and thus, reduces amyloid elimination across the BBB.

Lastly, by linking ABCA7 to cholesterol and amyloid exchanges across the BBB, our results 1) revealed novel pathways underlying the association between ABCA7 dysfunction and AD pathogenesis, and 2) suggest that this transporter may be a potential therapeutic target for curing or preventing AD.

All these experiments were performed using an *in vitro* BBB model, therefore further *in vivo* experiments using wild-type and *Abca7*^{-/-} mice should be performed to confirm the relevance of these data and to confirm the links between ABCA7, BBB, and AD.

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