Sensing of Dietary Lipids by Enterocytes: A New Role for SR-BI/CLA-1

Olivier Béaslas1,2,3, Carine Cueille1,2,3, François Delers1,2,3, Danielle Chateau1,2,3, Jean Chambaz1,2,3, Monique Rousset1,2,3, Véronique Carrière1,2,3*

1 Université Pierre et Marie Curie - Paris 6, UMR S 872, Les Cordeliers, Paris, France, 2 INSERM, U 872, Paris, France, 3 Université Paris Descartes, UMR S 872, Paris, France

Abstract

Background: The intestine is responsible for absorbing dietary lipids and delivering them to the organism as triglyceride-rich lipoproteins (TRL). It is important to determine how this process is regulated in enterocytes, the absorptive cells of the intestine, as prolonged postprandial hypertriglyceridemia is a known risk factor for atherosclerosis. During the postprandial period, dietary lipids, mostly triglycerides (TG) hydrolyzed by pancreatic enzymes, are combined with bile products and reach the apical membrane of enterocytes as postprandial micelles (PPM). Our aim was to determine whether these micelles induce, in enterocytes, specific early cell signaling events that could control the processes leading to TRL secretion.

Methodology/Principal Findings: The effects of supplying PPM to the apex of Caco-2/TC7 enterocytes were analyzed. Micelles devoid of TG hydrolysis products, like those present in the intestinal lumen in the interprandial period, were used as controls. The apical delivery of PPM specifically induced a number of cellular events that are not induced by interprandial micelles. These early events included the trafficking of apolipoprotein B, a structural component of TRL, from apical towards secretory domains, and the rapid, dose-dependent activation of ERK and p38MAPK. PPM supply induced the scavenger receptor SR-BI/CLA-1 to cluster at the apical brush border membrane and to move from non-raft to raft domains. Competition, inhibition or knockdown of SR-BI/CLA-1 impaired the PPM-dependent apoB trafficking and ERK activation.

Conclusions/Significance: These results are the first evidence that enterocytes specifically sense postprandial dietary lipid-containing micelles. SR-BI/CLA-1 is involved in this process and could be a target for further study with a view to modifying intestinal TRL secretion early in the control pathway.

Introduction

The increased incidence of metabolic disorders (obesity, metabolic syndromes and diabetes) and the ensuing atherosclerosis and cardiovascular diseases are linked to the significant changes in dietary habits that have occurred in recent decades, among which is an increase in fat intake [1]. Marked and prolonged postprandial hypertriglyceridemia, characterized by the accumulation of apolipoprotein B-containing triglyceride-rich lipoproteins (TRL), is a significant contributor to the development of dyslipidemia and a known risk factor for atherosclerosis [2]. Enterocytes in the intestine, the first organ to come into contact with digestion products, transfer dietary lipids to the organism and largely contribute to the production of TRL. It is thus important to characterize the mechanisms involved in the control of lipid absorption in these cells, especially those modulating the synthesis and secretion of TRL, as this could lead to the development of drugs acting on the early control steps in the intestinal transfer of dietary lipids, which could be used to reduce postprandial hypertriglyceridemia.

The absorption of lipids through the intestine is no longer considered a passive process but rather an active regulation of highly polarized mechanisms [3]. During the postprandial period, dietary lipids, mostly triglycerides (TG), after being hydrolyzed by pancreatic enzymes into fatty acids (FA) and monoglycerides (MG) and solubilized by bile salts and lipids in the intestinal lumen, are supplied to enterocytes as postprandial micelles (PPM). After absorption of FA and MG by enterocytes, TG must be re-synthesized in the endoplasmic reticulum and associate with the structural apolipoprotein (apo) B48, and apoA-I and apoA-IV to produce chylomicrons, the intestinal postprandial form of TRL that are secreted into lymph and then into the general circulation [3,4]. The small intestine must adapt to the variations in lipid load and composition that occur daily between postprandial and interprandial periods (for review [5]). The adaptation of enterocyte function has mainly been studied in terms of the effect of dietary lipids on gene expression and the action of transcription factors (for review [6]).

At the same time, the small intestine signals nutrient abundance to the organism and contributes to satiety by the release of hormones and enteropeptides from enteroendocrine cells and by the secretion of chylomicrons and apoA-IV, a satiety signal [7], by enterocytes (for reviews [8,9]). However, most studies on the
effects of dietary lipids in the small intestine do not relate the structural aspects of lipid supply to enterocytes to the specificity of the effects. More precisely, they do not address whether the ‘physiological’ mode of delivering dietary lipids to the apical pole of enterocytes as complex micelles, known to be critical for intestinal lipid intake [10], is required to induce the effects reported. While some mechanisms by which enteroendocrine cells sense lipids have been described [11], it is not known whether enterocytes sense dietary lipids. Previous works from our group indicated that there is some sensing of micellar dietary lipids by enterocytes as they demonstrated that the apical supply of postprandial micelles (mimicking those present in the intestinal lumen in vivo after a meal) induced specific effects in Caco-2/TC7 cells, a model reproducing in culture most of the morphological and functional characteristics of enterocytes [12,13]. These effects involved the rapid displacement of the apical brush-border-associated pool of apoB towards intracellular secretory compartments [14] and the intracellular neosynthesis of TG, leading to the secretion of TRL [15]. In addition, the apical supply of PPM induced the activation of apoA-IV transcription via the transcription factor HNF-4 [16], a process also recently reported to occur induced the activation of apoA-IV transcription via the transcription factor HNF-4 [16], a process also recently reported to occur induced the activation of apoA-IV transcription via the transcription factor HNF-4 [16], a process also recently reported to occur.

Materials and Methods

Chemicals and antibodies

Unless indicated, all chemicals were purchased from Sigma France. Antibodies used were: mouse polyclonal anti-Cla-1 (BD Transduction Laboratories), rabbit polyclonal anti-SR-BI (Novus 400-104), sheep polyclonal anti-PLAP (ABDSerotec 0300-1999), rabbit polyclonal anti-ERK (Cell Signaling), rabbit polyclonal anti-P-ERK (Thr202/Tyr204, Cell Signaling), rabbit polyclonal anti-p38 MAPK (Cell Signaling), rabbit polyclonal anti-P-p38 MAPK (Thr180/Tyr182, Cell Signaling), mouse monoclonal anti-E-cadherin (BD transduction laboratories), goat polyclonal anti-EEA1 (Santa Cruz Biotechnology), mouse monoclonal anti-flotillin-1 (BD Transduction Laboratories), mouse monoclonal anti-sucrase-isomaltase (provided by Dr. Swallow, University College London) and goat polyclonal anti-apoB (Immuno France). Secondary Cy3 and Cy2 dye-labeled antibodies were from Jackson Immunoresearch Laboratories. Peroxidase-conjugated immunoglobulins were from Bissy France. Alexa 488-conjugated Concanavalin A was from Molecular Probes.

Cell culture and stable knockdown of SR-BI/CLA-1 gene expression by shRNA

Caco-2/TC7 cells [12] were cultured on semi-permeable filters as described [14]. All experiments were performed 21 days after seeding, i.e. when the cells have acquired their optimal capacity of TRL secretion [15]. In some experiments, cycloheximide was added 1 hour before the treatment of the cells and during all the time of experiment, as already reported [14]. For stable SR-BI/CLA-1 knockdown experiments, lentiviral particles containing short hairpin RNA (shRNA) sequences targeted to human SR-BI/CLA-1 (GenBank accession number NM_005503) were purchased from Sigma MISSION Lentiviral Transduction Particles. Two lentiviral vectors containing shRNA sequences identified in The RNAi Consortium shRNA library as TRCN0000056963 (shRNA 65) and TRCN0000056964 (shRNA 64) were used and their transfection in Caco-2/TC7 cells gave Cell populations 63 and 64, respectively. One day after seeding on 24-well culture plates (Corning, France), dividing Caco-2/TC7 cells were incubated overnight with 0 μg/ml hexadecamethylene bromide and 200 μl of shRNA lentiviral particles containing 105 lentiviral transducing particles/ml in DMEM-FBS. The medium was then changed daily. The SR-BI/CLA-1 knockdown cells were able to grow and to differentiate as well as untransfected Caco-2/TC7 cells (data not shown). Stably transfected cells were selected with 10 μg/ml puromycin. Total puromycin-resistant cell populations were analyzed. Stock SR-BI/CLA-1 knockdown cell populations were cultured in the presence of puromycin to sustain the selection. Whatever the culture passages that were analyzed, all cell populations remained resistant to puromycin. For maintenance of infected cell populations, a 6-day passage frequency was used. The experiments were performed after 21 days of culture on semi-permeable filters, without puromycin.

Preparation of lipid micelles-containing media

Preparation of lipid micelles was performed as described [15]. The composition of postprandial lipid micelles (PPM) was similar to that of human duodenum lumen [10] after a lipid-rich meal (0.6 mM oleic acid, 0.2 mM L-α-lysophosphatidylcholine, 0.05 mM cholesterol, 0.2 mM 2-monooeleoylglcerol, 2 mM taurocholic acid). Inter-prandial lipid micelles (IPM) were prepared as PPM but, according to physiological conditions [19], were devoid of oleic acid and 2-monooeleoylglcerol. PPM or IPM were added to the medium of the upper compartment for the last day of culture, for the indicated times.

Competitors and inhibitors of SR-BI/CLA-1 and of protein kinases

HDL3 were isolated by ultracentrifugation as described [20], from EDTA-plasma obtained from human volunteers, supplemented with 0.005% gentamycin, 0.04% sodium azide and protease inhibitors. HDL3 (100 μg/ml) were added in the apical medium at the same time as PPM. For studies with inhibitors, Caco-2/TC7 cells were pre-incubated 1 hour in the presence of 20 μM of the selective inhibitor of P38 MAPK, SB203580 (Calbiochem), or 10 μM of BLT-1 (Chembridge corporation, USA), an inhibitor of SR-BI-dependent lipid transport, in both apical and basal compartments. Fresh solutions of inhibitors were then added in the apical medium at the same time as PPM. Both inhibitors were diluted in DMSO.

Confocal Fluorescence Microscopy

Immunofluorescence and confocal analyses were performed as described [14]. Cells were fixed with 4% paraformaldehyde (wt/vol) and permeabilized with 0.2% Triton X-100 for apoB or 0.01% saponin for SR-BI/CLA-1, at 4°C. In some experiments, alexa 488-conjugated concanavalin A was added at the same time as primary antibodies to visualize the apical brush-border. Immunofluorescence was examined by confocal fluorescence microscopy (LSM510 microscope Zeiss Germany, with a 63x water lens, 80 μm pinhole). Images were treated using Photoshop and Illustrator software (Adobe).
Cell surface biotinylation assay

Caco-2/TC7 cells cultured on filters were incubated without or with PPM, and washed with ice-cold PBS containing 1 mM CaCl₂ and 0.5 mM MgCl₂. Cells were then incubated (30 min, 4°C) with 0.9 mg/ml sulfo-NHS-SS-Biotin, added only to the apical compartment. Apical membrane expression of SR-BI/CLA-1 was analyzed by biotin labeling of surface proteins with EZ-Link Sulfo-NHS-SS-Biotin, followed by immunoprecipitation with immobilized streptavidin using the Cell Surface Protein Biotinylation and Purification Kit (Pierce), according to manufacturer’s instructions. Total cell lysates obtained before immunoprecipitation (total), immunoprecipitated fractions containing apical biotinylated proteins (apical fraction) and non-immunoprecipitated fractions containing unlabeled proteins (non apical fraction), were subjected to immunoblot analyses. In some experiments sulfo-NHS-SS-Biotin was added only in the basal compartment and the basal fraction containing biotinylated proteins was recovered as described above.

Sucrose gradient

For sucrose density gradients, 10⁸ Caco-2/TC7 cells were homogenized on ice in 2 ml of 10 mM Tris-HCl, pH 8, 150 mM NaCl buffer (TBS) containing 1% Triton X-100 and protease and phosphatase inhibitor cocktails. Homogenate was adjusted to 40% sucrose by 2 ml of 80% sucrose/TBS. The resulting 4 ml were covered with 4 ml of 30% sucrose and 4 ml of 5% sucrose and centrifuged (SW41, L8 Beckman, 18h, 39,000 rpm, 4°C). Sequential 1 ml fractions were collected from the top of the tube. Eleven fractions were obtained, to which NP-40 was added (1% final concentration).

Ligand blotting assay with micelles

Homogenates from differentiated Caco-2/TC7 cells were prepared in Tris/mannitol buffer (2 mM Tris-HCl pH 7.4, 50 mM mannitol, 2% protease inhibitor cocktail) and centrifuged for 15 min (5,000 rpm, 4°C). Supernatants were subjected to SDS-PAGE. Proteins were transferred onto Bio-Rad nitrocellulose membranes, which were saturated for 30 min with 6% defatted milk in PBS containing 0.25 M NaCl (PBS/NaCl). For lipid overlay, 0.05 mM biotinylated phosphatidylethanolamine was incorporated to the preparation of PPM or IPM with a 1/200 molar ratio toward L-α-lysophosphatidylcholine. Micelles were mixed with coupled horseradish-streptavidin peroxydase (1 μg/ml, Jackson ImmunoResearch) and incubated 10 min at room temperature. Membranes containing blotted proteins were incubated with this solution for 2 hours at room temperature, washed with PBS/NaCl and revealed by ECL system.

For immunoprecipitation of SR-BI/CLA-1, 5 μl of rabbit anti-SR-BI antibodies (Novus NB400-104) or rabbit non-immune serum were mixed with 25 μl of 400 mg/ml protein A-sepharose (Healthcare Biosciences) in Tris buffer (20 mM Tris-HCl, pH 7.4, 5 mM EDTA, 0.15 M NaCl, 5 mM lactate and 0.1% ovalbumin) for 1 hour at room temperature. After several washings with Tris buffer, SR-BI antibodies combined to protein A-sepharose were incubated with 90 μg of Caco-2/TC7 cell homogenates, for 2 hours at 4°C. After several washings with Tris buffer, immunoprecipitated material was recovered with Laemmli buffer, subjected to SDS-PAGE and then to lipid overlay.

Western blot analysis

Caco-2/TC7 cells were scraped into lysis buffer (20 mM Tris-HCl, pH 7.4, 5 mM EDTA, 0.15 M NaCl, 1% Triton X-100, 0.5% sodium deoxycholate) containing protease and phosphatase inhibitor cocktails. Aliquots of cell lysates (40-80 μg) were subjected to SDS-PAGE under reducing conditions in a 7.5% polyacrylamide gel. Proteins were transferred onto nitrocellulose membranes, developed using ECL Western blotting reagents according to the manufacturer’s instructions (Amersham Biosciences, Orsay, France) and quantified using Image J software. The levels of phospho-ERK and phospho-p38MAPK were normalized with total ERK and p38MAPK, respectively.

Figure 1. Trafficking of apoB is specifically induced by alimentary lipid-containing micelles. Confocal image of apolipoprotein B (apoB, red channel) and sucrase-isomaltase (SI, used as a brush-border marker, green channel) localizations in differentiated Caco-2/TC7 cells cultured in the absence (control) or presence of postprandial (PPM) or inter-prandial (IPM) micelles added at the apical pole of the cells for 20 min. Top panels show XY acquisitions at the apical level and at 3 μm and 5 μm below the apical plane. Bottom panels show the corresponding XZ projections (zoom ×2). (bar, 20 μm).

doi:10.1371/journal.pone.0004278.g001
Figure 2. Signaling pathways specifically induced by postprandial micelles. (A) Kinetworks™ phosphoprotein immunoblots from lysates of Caco-2/TC7 cells cultured on filters without micelles (control) or with PPM or IPM for 5 min. Duplicate immunoblots using specific antibodies directed against phosphoproteins were analyzed by Kinexus. Bands corresponding to phosphoproteins specifically up-regulated by PPM are indicated by arrows. The antibodies recognized the phosphosites: S144/S141/S154 for PAK1/2/3, T180/Y182 for p38 MAPK, S129/S133 for CREB1, T202/Y204 for ERK1, T185/Y187 for ERK2 and S338 for PKA Cβ. The phosphosites analyzed for MEK1 are indicated in the figure. Histograms show the amounts of phosphoproteins expressed as the ratio of phosphorylation in the PPM versus the IPM sample (from three independent experiments). (B) Caco-2/TC7 cells cultured on semi-permeable filters were incubated in the absence or presence of PPM or IPM in the apical compartment for the indicated times. Cell lysates were analyzed by immunoblot with antibodies against phospho-ERK1/2 (P-ERK), and phospho-p38 MAPK (P-p38 MAPK). Total ERK, P-ERK, and P-p38 MAPK were quantified using densitometry.
Phosphoprotein screens
Caco-2/TC7 cell lysates were analyzed by Kinetworks™ KPSS-9.0 phosphorylation screens (Kinexus Bioinformatics Corp., Canada). Kinetworks analysis, performed by Kinexus, involved resolution of a single lysate sample by SDS-PAGE with 20-lane multi-immunoblotters and subsequent immunoblotting using specific antibodies against 37 phosphoproteins. Kinexus performed all the normalization and statistical analyses on the data.

Immunoelectron microscopy
Differentiated Caco-2/TC7 cells were fixed with 4% PFA plus 0.1% glutaraldehyde. After alcohol-graded dehydration, sections were embedded in LR White, and ultra-thin sections were incubated with rabbit anti SR-BI/CLA-1 antibody (Novus, NB400-104), gold (12 nm particles)-labeled donkey anti-rabbit IgGs (Jackson Immunoresearch) being used as secondary antibodies. Double labeling of SR-BI/CLA-1 and alkaline phosphatase (PLAP) was performed by using secondary antibodies labeled with 10 nm and 12 nm gold particles, respectively. Sections were analyzed in a Jeol 100 CX II electron microscope.

Statistical analyses
Statistical analyses were performed using Student’s t test for unpaired data.

Results
Cell signaling events are specifically induced in enterocytes by postprandial micelles
We previously showed that addition of PPM to Caco-2 enterocytes caused the chase of apoB from the apical brush-border towards basolateral secretory domains within 15 min [14]. To evaluate the specific effect of alimentary lipids supplied as ‘physiological’ complex PPM, we compared this effect on apoB trafficking to that induced by interprandial micelles (IPM) devoid of digested TG products [Figure 1]. In the absence of micelles from the apical compartment, apoB co-localized with the brush-border marker sucrase-isomaltase (SI). After PPM were delivered to the apical pole, apoB was chased towards intracellular compartments, as previously reported [14]. This delocalization, which was not observed after IPM supply [Figure 1], indicates that the mobilization of apical apoB depends specifically on the supply of micellar alimentary lipids leading us to hypothesize that PPM can induce specific signaling pathways in enterocytes. We therefore tested whether any of multiple signaling proteins, mainly kinases, were activated using a specific, semi-quantitative method to identify them with Kinetworks™ screens. Time-course experiments were performed 5, 10 and 15 min after the apical supply of PPM or IPM. As early as 5 min after PPM supply, we observed an increase in the phosphorylation of five protein kinases, namely PAK1/2/3, MEK1, p38 MAPK, ERK1/2 and PKA Cβ (Figure 2A). Of these, MEK1, responsible for the activation of ERK1/2 in the canonical ERK pathway, was phosphorylated at three phosphosites (T202/S201, T297, and T298). There was also an increase in the phosphorylation of the CREB1 protein, a target transcription factor of MAPK pathway (for review [21]). None of these effects were observed when IPM was supplied to cells (Figure 2A) even after 15 min (data not shown). As most of the kinases that were activated after PPM supply belong to the MAPK signaling pathway, we focused on ERK1/2 and p38 MAPK, two kinases downstream in this pathway. PPM supply specifically increased the phosphorylation of ERK1/2 and p38 MAPK by 5 min and 2 min respectively [Figure 2B and 2C], in a dose-dependent manner [Figure 2D] as determined by immunoblotting.

The supply of PPM specifically regulates the subcellular distribution of SR-BI/CLA-1
We hypothesized that a protein located at the apical membrane of enterocytes could trigger the observed specific cell signaling events through an interaction with PPM. To identify proteins that specifically interact with PPM, we tested the interaction of cell extracts with labelled phospholipid-containing PPM or IPM in a ligand blotting assay. While both types of micelles interacted with several proteins, these interactions were enhanced with PPM; PPM also interacted with one 75 kDa protein that IPM did not interact with [Figure 3A]. Of the lipid receptors known to be localized at the apical membrane of enterocytes with a similar molecular weight is the scavenger receptor SR-BI (CLA-1 in Human) [22], which is expressed in the brush border membrane of enterocytes of the duodenum and jejunum and in Caco-2 cells [23-25]. Other candidates NPC1L1 and FabP1 are 150 kDa [26] and 40 kDa [27] respectively, and FAT1/CD36 [28] is not expressed in Caco-2 cells [25,29] and Carrière et al. unpublished results). To test whether SR-BI/CLA-1 interacts with PPM, Caco-2/TC7 cell homogenates were immunoprecipitated with antibody against SR-BI/CLA-1 and then assayed by ligand-blotting. Results presented in Figure 3B.
confirm that there is an interaction between PPM and SR-BI/CLA-1 in the immunoprecipitate.

We then analyzed the effects of PPM on the expression of SR-BI/CLA-1. Immunoelectron microscopy showed that in control Caco-2/TC7 cells SR-BI/CLA-1 was predominantly localized at the apical pole of the cells, in microvilli and in the terminal web (Figure 4A), as described previously for pig and human enterocytes [24,30]. Immunofluorescence analysis of SR-BI/CLA-1 localisation confirmed that SR-BI/CLA-1 co-localized with the brush-border marker sucrase-isomaltase in control Caco-2/TC7 cells (Figure 4B). SR-BI/CLA-1 formed clusters at the apical membrane of enterocytes as soon as 5 min after PPM supply (Figure 4B and 4D). These clusters, which were still present after 20 min, were not observed in IPM-treated cells (Figure 4C). To determine whether SR-BI/CLA-1 clustering resulted from an increased recruitment at the apical membrane, cell surface biotinylation assay for apical SR-BI/CLA-1 was used to detect the amount of SR-BI/CLA-1 at the apical membrane. Biotinylated fractions were isolated from the apical and basolateral cell membranes using a biotin-streptavidin complex. The amount of SR-BI/CLA-1 in total cell homogenates, in biotinylated fractions and in non-biotinylated fractions was compared by immunoblotting. There was much more SR-BI/CLA-1 at the apical membrane than at the basolateral membrane of enterocytes (Figure 4E, compare left and right panels). No increase in the apical

**Figure 4. Subcellular localization of SR-BI/CLA-1 after the supply of postprandial micelles.** (A) Immunoelectron micrograph of SR-BI/CLA-1 in untreated differentiated Caco-2/TC7 cells. MV, microvilli; TW, terminal web (bar, 0.5 μm). Note the significant amount of intracellular trafficking of SR-BI/CLA-1 in addition to its main apical localization (arrowheads). (B) Immunolocalization of SR-BI/CLA-1 (green channel) and sucrase isomaltase (SI, red channel) in differentiated Caco-2/TC7 cells before (T0) and after 5, 10 and 15 min of apical PPM supply. Panels represent XY acquisitions at the apical level (bar, 10 μm). Arrowheads show clusters of SR-BI/CLA-1. (C) Immunolocalization of SR-BI/CLA-1 in differentiated Caco-2/TC7 cells in the absence (control) or presence of PPM or IPM for 20 min (bar, 20 μm). Arrowheads indicate clusters of SR-BI/CLA-1 formed clusters at the apical membrane of enterocytes as soon as 5 min after PPM supply (Figure 4B and 4D). These clusters, which were still present after 20 min, were not observed in IPM-treated cells (Figure 4C). To determine whether SR-BI/CLA-1 clustering resulted from an increased recruitment at the apical membrane, cell surface biotinylation assay for apical SR-BI/CLA-1 was used to detect the amount of SR-BI/CLA-1 at the apical membrane. Biotinylated fractions were isolated from the apical and basolateral cell membranes using a biotin-streptavidin complex. The amount of SR-BI/CLA-1 in total cell homogenates, in biotinylated fractions and in non-biotinylated fractions was compared by immunoblotting. There was much more SR-BI/CLA-1 at the apical membrane than at the basolateral membrane of enterocytes (Figure 4E, compare left and right panels). No increase in the apical

**Figure 4. Subcellular localization of SR-BI/CLA-1 after the supply of postprandial micelles.** (A) Immunoelectron micrograph of SR-BI/CLA-1 in untreated differentiated Caco-2/TC7 cells. MV, microvilli; TW, terminal web (bar, 0.5 μm). Note the significant amount of intracellular trafficking of SR-BI/CLA-1 in addition to its main apical localization (arrowheads). (B) Immunolocalization of SR-BI/CLA-1 (green channel) and sucrase isomaltase (SI, red channel) in differentiated Caco-2/TC7 cells before (T0) and after 5, 10 and 15 min of apical PPM supply. Panels represent XY acquisitions at the apical level (bar, 10 μm). Arrowheads show clusters of SR-BI/CLA-1. (C) Immunolocalization of SR-BI/CLA-1 in differentiated Caco-2/TC7 cells in the absence (control) or presence of PPM or IPM for 20 min (bar, 20 μm). Arrowheads indicate clusters of SR-BI/CLA-1 formed clusters at the apical membrane of enterocytes as soon as 5 min after PPM supply (Figure 4B and 4D). These clusters, which were still present after 20 min, were not observed in IPM-treated cells (Figure 4C). To determine whether SR-BI/CLA-1 clustering resulted from an increased recruitment at the apical membrane, cell surface biotinylation assay for apical SR-BI/CLA-1 was used to detect the amount of SR-BI/CLA-1 at the apical membrane. Biotinylated fractions were isolated from the apical and basolateral cell membranes using a biotin-streptavidin complex. The amount of SR-BI/CLA-1 in total cell homogenates, in biotinylated fractions and in non-biotinylated fractions was compared by immunoblotting. There was much more SR-BI/CLA-1 at the apical membrane than at the basolateral membrane of enterocytes (Figure 4E, compare left and right panels). No increase in the apical
amount of SR-BI/CLA-1 was observed after PPM supply (Figure 4E, left panel), indicating that PPM, while they did induce clusters of apical SR-BI/CLA-1, did not modify its targeting to the apical membrane of enterocytes.

It has been reported that SR-BI function may depend on its localization in raft microdomains [30–32]. The effect of PPM or IPM supply on the association of SR-BI/CLA-1 with rafts was assessed by fractionating cell contents through a sucrose gradient (Figure 5). In cells that received no micelles in the apical medium, a large amount of trafficking SR-BI/CLA-1 was found in non-raft/high-sucrose density cell fractions (fractions 7–11), which contained the early endosome marker EEA1. However, SR-BI/CLA-1 was also localized in the low-sucrose density fractions (fractions 4–6) in raft microdomains identified by the expression of flotillin-1 (Figure 5A). When PPM were supplied to cells, more SR-BI/CLA-1 was observed in raft fractions (fractions 4–6) (Figure 5B). This localization was confirmed by immunoelectron microscopy (Figure 5C), which showed that after PPM supply SR-BI/CLA-1 was in the vicinity of the raft-associated alkaline phosphatase [33,34]. No difference in the partitioning of SR-BI/CLA-1 was observed in cells supplied with IPM and control cells. PPM and IPM supply did not modify the subcellular distribution of flotillin-1 indicating that they did not alter properties of the sucrose gradient.

SR-BI/CLA-1 is involved in PPM-induced cell signaling in enterocytes

To determine how SR-BI/CLA-1 acts in PPM-induced cell signaling, we first analyzed the effect of SR-BI/CLA-1 knockdown in Caco-2/TC7 cells. Two different cell populations were obtained by stable lentivirus-mediated transfection of two different SR-BI/CLA-1 shRNA sequences. The expression of SR-BI/CLA-1 protein was 80% lower in Cell population 63 than in control Caco-2/TC7 cells, while Cell population 64 expressed SR-BI/CLA-1 protein at nearly the same level as control Caco-2/TC7 cells and was thus used as a negative control (Figure 6A). PPM supply did not modify the amount of SR-BI/CLA-1 protein in these transfected cell populations. After several passages, we observed that SR-BI/CLA-1 expression in Cell population 63 rose to 60% of that in control cells, whereas there was no change in the level of SR-BI/CLA-1 expression in Cell population 64 over the same period (Figure 6B). We used these differences in SR-BI/CLA-1 protein levels to analyze the relationship between PPM-dependent ERK1/2 phosphorylation and the amount of SR-BI/CLA-1 protein, i.e., by comparing Cell population 63 at early (20% expression) and late passages (60% expression) with control cells (100% expression, untransfected) and Cell population 64 (100% expression, transfected). As shown in Figure 6B, the PPM-induced ERK1/2 phosphorylation level was positively correlated with the amount of SR-BI/CLA-1, with the least ERK1/2 activation in the cell population expressing the least SR-BI/CLA-1. The PPM-dependent delocalization of the apical pool of apoB was compared in controls and Cell population 63 (at early passages). Cells were incubated in the presence of cycloheximide, which inhibits protein synthesis, so the disappearance of the preexisting apical pool of apoB (in the presence of PPM) resulting from its chase towards intracellular compartments could be visualized without confusion with the trafficking of newly synthesized apoB targeted to the apical membrane [14]. As shown in Figure 6C, in SR-BI/CLA-1-knockdown cells PPM supply did not stimulate the chase of apoB, which was maintained at the
apical pole unlike control cells incubated with PPM, in which apoB disappeared from the brush-border domain.

To further assess the involvement of SR-BI/CLA-1 in the cell signaling events that are specifically induced by PPM, we checked whether a ligand of SR-BI/CLA-1 could compete with PPM and interfere with their downstream effects, namely the phosphorylation of MAP Kinases and the apoB trafficking. HDL₃, a strong SR-BI/CLA-1 ligand, was chosen as it has already been used as a competitor in other studies [35] including those using purified brush-border membrane vesicles of enterocytes [36]. The PPM-induced movement of the apical pool of apoB towards intracellular compartments (Figure 7A) and the phosphorylation of ERK1/2 (Figure 7B) were impaired in the presence of HDL₃ at the apical pole of Caco-2/TC7 cells. By contrast, the addition of HDL₃ in the absence of PPM had no effect compared to the controls. That SR-BI/CLA-1 has a role in PPM-dependent signaling is also reinforced by the fact that apoB trafficking is impaired in the presence of the inhibitor of SR-BI-dependent lipid transport BLT-1 [29,37] (Figure 7C).

Finally, specific inhibition of p38 MAPK phosphorylation with SB203580 (Figure 7D) impaired the PPM-dependent apoB trafficking (Figure 7C), suggesting that the activation of p38 MAPK would be one of the mechanisms involved in the first step of TRL secretion.

Discussion

Our previous studies led us to hypothesize that dietary lipids are sensed in their physiological form of postprandial micelles (PPM) by enterocytes [14–16,38], a process that would trigger specific
events leading enterocytes to secrete TG as TRL. Here we show that such a sensing system does exist in enterocytes and it involves the scavenger receptor class B type 1. We demonstrate that the enterocyte sensing of postprandial lipid micelles is characterized by the rapid delocalization of apoB, the structural protein necessary for the assembly of TRL, and by the rapid activation of ERK and p38 MAPK. In the absence of SR-BI/CLA-1, or in the presence of SR-BI/CLA-1 inhibitors or ligand competitors, these processes are impaired. This sensing step involves the clustering of SR-BI/CLA-1 at the apical membrane, a process associated with SR-BI/CLA-1

Figure 7. PPM-induced effects are impaired in the presence of competitors or inhibitors of SR-BI/CLA-1 and of the MAP kinase pathway. (A) Caco-2/TC7 cells were analyzed before and after a 10-min incubation with PPM with or without apical HDL₃ (100 µg/ml). ApoB distribution was analyzed by confocal microscopy. Upper panels show XY acquisitions at the apical brush-border (BB) level and 6 µm below the apical plane. Lower panels show the corresponding XZ projections. Dotted lines indicate the base of the cells (bar, 20 µm). (B) Caco-2/TC7 cells were incubated with PPM and HDL₃ as in (A). Cell lysates were analyzed in phospho-ERK1/2 (P-ERK) immunoblots with total ERK (ERK) and E-cadherin (E-cadh) levels used as loading controls. Lower panel, P-ERK levels normalized to total ERK expression. Results from two independent experiments are expressed as the mean±SEM, *P<0.05. (C) Caco-2/TC7 cells were pre-incubated for 1 hour with cycloheximide and 10 µM BLT-1 or 20 µM SB203580 or DMSO in both the apical and basal compartments. Inhibitors or DMSO were added again when PPM or DMEM were added in the apical compartment for 20 min. Immunolocalization of apoB in these cells using conA to visualize the apical membrane. Upper panels show the dual labeling of conA (green) and apoB (blue) in the apical plane (XY planes; bar, 50 µm), lower panels show the same apical XY planes with apoB alone (white). (D) Caco-2/TC7 cells were analyzed before or after a 10-min incubation with PPM with or without SB203580, a p38 MAPK inhibitor, added at the same time as PPM or DMEM. Cell lysates were analyzed in immunoblots of phospho-p38 MAPK (P-p38) and total p38 MAPK (p38). Lower panel, P-p38 MAPK levels normalized to total p38 MAPK levels. Results from two independent experiments are expressed as the means±SEM. *P<0.05, **P<0.01. doi:10.1371/journal.pone.0004278.g007
function in other cell models [39], and its recruitment into raft domains, that are known to contain various signaling molecules, including MAP kinases [40]. The localization of SR-BI/CLA-1 in raft domains differs from the observation of Hansen et al. (2003) [30]. This could be explained by differences in the experimental conditions used; these authors studied the lipid raft association of SR-BI in membrane preparations from intestinal mucosa scrapings of fasted pigs. One hypothesis is that the raft association of SR-BI depends on the nutritional status or on other environmental factors. It should be noted that the subcellular localization of SR-BI varies in different nutritional and hormonal conditions in adipocytes [41,42].

SR-BI/CLA-1 has been extensively studied in terms of its involvement in cholesterol transport in the liver and in macrophages [43]. Recently, a role for SR-BI in lipid sensing in vascular endothelial cells has been described. In these cells, where SR-BI acts as a cholesterol sensor of LDL at the plasma membrane, the interaction with its ligand leads to a sequential activation of protein kinases that stimulate eNOS enzymatic activity [44,45] and Rac GTPase activation [46]. The role of SR-BI/CLA-1 in the intestine remains controversial [47–50]. It was previously thought that SR-BI/CLA-1 was responsible for the absorption of micellar cholesterol from the intestinal lumen [24,51,52]. However, deficiency of intestinal cholesterol absorption is not observed in SR-BI knockout mice [53] and NPC1L1 was discovered to be the major intestinal transporter of cholesterol [54] together suggesting that SR-BI/CLA-1 may play another role in enterocytes. The newly assigned role of SR-BI/CLA-1 as a sensor of lipid micelles in enterocytes, described here, is in accordance with results showing that 1) SR-BI/CLA-1 binds not only high-density lipoproteins but also lipid structures devoid of apolipoproteins like PPM. 2) The three-dimensional structures of SR-BI/CLA-1 ligands are important in modulating its activity [51] and 3) various SR-BI/CLA-1 ligands trigger MAPK [35,55] and PI3K/Akt [45] signaling pathways. Furthermore, the overexpression of SR-BI in the intestine leads to increases in cholesterol absorption and triglyceride secretion indicating a functional role for SR-BI in triglyceride lipid metabolism [50].

Our observations suggest that interaction between PPM and SR-BI/CLA-1 induces the establishment of a signaling platform in the brush border membrane of enterocytes that leads to the activation of cellular events, namely the activation of MAPK kinases and the trafficking of apoB, the structural protein involved in TRL, assembly and secretion, from apical towards basolateral domains. A link has been made between apoB secretion and MAPK activation [60,61] and in Caco-2 cells [15,62].

In conclusion, our results provide evidence for an enterocyte sensing of alimentary lipid-containing micelles, which could prepare these absorptive cells for the massive postprandial availability of lipids, and that SR-BI/CLA-1 participates early in this process at the apical membrane of enterocytes. Changes in PPM sensing, and in the resulting secretion of TRL, could be caused by factors such as diet, hormones or pharmaceutical compounds, which modify the expression and/or activity of SR-BI/CLA-1, a new target in studying how the early stages of the intestinal transfer of dietary lipids is controlled.

Acknowledgments

We thank D. Pastier for HDL₄ isolation and M. Danielsen for the choice of raft markers. Confocal and electron microscopy analyses have benefit from facilities of the Cordeliers Research Centre.

Author Contributions

Conceived and designed the experiments: OB FD VC. Performed the experiments: OB CC FD JC MR VC. Contributed reagents/materials/analysis tools: VC. Wrote the paper: OB JC MR VC.

References

1. Shikany JM, White GI, Jr (2000) Dietary guidelines for chronic disease prevention. South Med J 93: 1138–1151.
2. Tomkin GH, Owens D (2001) Abnormalities in apo B-containing lipoproteins in diabetes and atherosclerosis. Diabetes Metab Rev 17: 27–43.
3. Mu H, Hoy C-E (2004) The digestion of dietary triacylglycerols. Progress in Lipid Research 43: 105–133.
4. Davidson NO, Shelness GS (2000) APOLIPROTEIN B: mRNA editing, lipoprotein assembly, and presecretory degradation. Annu Rev Nutr 20: 169–193.
5. Drozdowski L, Thomson AB (2006) Intestinal mucosal adaptation. World J Gastroenterol 12: 4614–4627.
6. Sanderson BR, Naik S (2008) Dietary regulation of intestinal gene expression. Annu Rev Nutr 29: 311–330.
7. Qin X, Tso P (2005) The role of apolipoprotein AV on the control of food intake. Curr Drug Targets 6: 145–151.
8. Raybould HE (1999) Nutrient tasting and signaling mechanisms in the gut. I. Sensing of lipid by the intestinal mucosa. Am J Physiol 277: G731–755.
9. Cummings DE, Overhult J (2007) Gastrointestinal regulation of food intake. J Clin Invest 117: 13–23.
10. Nordskog BK, Phan CT, Nutting DF, Tso P (2001) An examination of the factors affecting intestinal lymphatic transport of dietary lipids. Adv Drug Deliv Rev 50: 21–44.
11. McLaughlin J (2007) Long-chain fatty acid sensing in the gastrointestinal tract. Biochem Soc Trans 35: 1199–1202.
12. Chantrat I, Redollesos A, Barbat A, Dussaulx E, Brot-Laroche E, et al. (1994) Differential expression of sucrase-isomaltase in clones isolated from early and
late passages of the cell line Caco-2: evidence for glucose-dependent negative regulation. J Cell Sci 107(Pt 1): 213–225.

13. Zucco F, Batto AF, Bises G, Chambaz J, Chiusolo A, et al. (2005) An interlaboratory study to evaluate the effects of medium composition on the differentiation and barrier function of Caco-2 cells. Altern Lab Anim 33: 603–618.

14. Morel E, Demignot S, Chateau D, Chambaz J, Rousset M, et al. (2004) Lipid-dependent bidirectional traffic of apolipoprotein B is polarized enterocytes. Mol Biol Cell 15: 1322–1334.

15. Chateau D, Pasquai T, Delers F, Rousset M, Chambaz J, et al. (2005) Lipid micelles stimulate the secretion of triglyceride-enriched apolipoprotein B48-containing lipoproteins by Caco-2 cells. J Cell Physiol 202: 767–776.

16. Cattaneo V, Vidal R, Lazou K, Luciani M, Delers F, et al. (2005) HNF4-dependent induction of apolipoprotein A-V gene transcription by an apical supply of lipid micelles in intestinal cells. J Biol Chem 280: 5406–5413.

17. Leng X, Su A, Yao Y, Kan Z, Morris GS, et al. (2007) Hepatocyte nuclear factor-4 mediates apolipoprotein A-V transcriptional regulation by fatty acid in newborn swine enterocytes. Am J Physiol Gastrointest Liver Physiol 293: G475–483.

18. Helquist S, Staggers JE, Carey MC (1996) Physical-chemical behavior of dietary and biliary lipids during intestinal digestion and absorption. 2. Phase analysis and aggregation states of luminal lipids during duodenal fat digestion in healthy adult human beings. Biochemistry 29: 2041–2056.

19. Porter CJ, Trevaskis NL, Charman WN (2007) Lipids and lipid-based formulations: optimizing the oral delivery of lipidic drugs. Nat Rev Drug Discov 6: 231–246.

20. Bosier E, Lambert G, Aitger V, Tran NQ, Pasteur D, et al. (1999) Overexpression of human apoprotein A-II in mice induces hypertriglyceridemia due to defective very low density lipoprotein hydrolysis. J Biol Chem 274: 11564–11572.

21. Lang R, Lambert M, Mages J (2006) DUSP meet immunology: dual specificity MAPK phosphatases in control of the inflammatory response. J Immunol 177: 7497–7504.

22. Voshol PJ, Schwarz M, Rigotti A, Krieger M, Groen AK, et al. (2001) Down-regulation of intestinal scavenger receptor class B, type I (SR-BI) expression in rodents under conditions of deficient bile delivery to the intestine. Biochem J 356: 317–325.

23. Cai SF, Kirby RJ, Howes PN, Hui DY (2001) Differentiation-dependent expression and localization of the class B type I scavenger receptor in intestine. J Lipid Res 42: 902–909.

24. Levy E, Menard D, Sic I, Delvin E, Marcil V, et al. (2004) Ontogeny, immunolocalisation, distribution and function of SR-BI in the human intestine. J Cell Sci 117: 327–337.

25. Lobo MV, Huerta L, Ruiz-Velasco N, Teixeiro E, de la Cueva P, et al. (2001) Localization of the lipid receptors CD36 and CLA-1 (CD36 and LIMPII analogous-1) in normal rat jejunal mucosa. Chem Phys Lipids 117: 135–145.

26. Davies JP, Scott C, Olohi K, Lisip K, Amaou Y. (2003) Inactivation of NPC1L1 causes multiple lipid transport defects and protects against diet-induced hypercholesterolemia. J Biol Chem 280: 12710–12720.

27. Potter BJ, Sorrentino D, Berk PD (1989) Mechanisms of cellular uptake of free long-chain fatty acids. Annu Rev Nutr 9: 253–270.

28. Assanasen C, Mineo C, Seetharam D, Yuhanna IS, Marcel YL, et al. (2003) Cholesterol binding, efflux, and a PDZ-interacting domain of scavenger receptor-BI mediate HDL-initiated signaling. J Clin Invest 115: 969–977.

29. Mineo C, Shaul PW (2003) HDL stimulation of endothelial nitric oxide synthase: a novel mechanism of HDL action. Trends Cardiovasc Med 13: 229–231.

30. Seetharam D, Mineo C, Gormley AK, Gibson LL, Vongpatanasin W, et al. (2004) High-density lipoprotein promotes endothelial cell migration and inhibits proliferation via scavenger receptor BI. J Cell Biochem 90: 63–72.

31. Labonte ED, Howes PN, Granholm CA, Rojas JC, Davies JP, et al. (2007) Class B type I scavenger receptor is responsible for the high affinity cholesterol binding activity of intestinal brush border membrane vesicles. Biochim Biophys Acta 1771: 1132–1139.

32. Levy E, Spahs S, Sinnett D, Peretti N, Maupas-Schwalm F, et al. (2007) Intestinal cholesterol transport proteins: an update and beyond. Curr Opin Lipidol 18: 310–318.

33. Hui DY, Howes PN (2005) Molecular mechanisms of cholesterol absorption and transport in the intestine. Semin Cell Dev Biol 16: 183–192.

34. Bietrix F, Yan D, Nauze M, Rolland C, Bertrand-Michel J, et al. (2006) Accelerated lipid absorption in mice overexpressing intestinal SR-BI. J Biol Chem 281: 7124–7129.

35. Rinaldi D, Bruseau L (2004) The role of scavenger receptor class B type I (SR-BI) in lipid trafficking, defining the rules for lipid transporters. Int J Biochem Cell Biol 36: 39–77.

36. Hausner H, Dyer JH, Nandy A, Vega MA, Werder M, et al. (1989) Identification of a receptor mediating absorption of cholesterol in the intestine. Biochemistry 28: 17843–17850.

37. Mardones P, Quinones V, Amigo L, Moreno M, Mique Jr, et al. (2001) Hepatic cholesterol and bile acid metabolism and intestinal cholesterol absorption in scavenger receptor class B type I-deficient mice. J Biol Chem 276: 13522–13527.

38. Atmann SW, Davis HR Jr, Zhu LJ, Yao X, Hoo LM, et al. (2004) Niemann-Pick C1 Like 1 protein is critical for intestinal cholesterol absorption. Science 303: 1201–1204.

39. Grewal R, de Diego I, Kirchoff MF, Tebar F, Hirem J, et al. (2003) High density lipoprotein-induced signaling of the MAPK pathway involves scavenger receptor type BI-mediated activation of Ras. J Biol Chem 278: 16478–16481.

40. Federico LM, Naples M, Taylor D, Adeli K (2006) Intestinal insulin resistance and hyperlipidemia via scavenger receptor class B type I. J Cell Biochem 98: 170–180.

41. Altmann SW, Davis HR Jr, Zhu LJ, Yao X, Hoo LM, et al. (2004) Niemann-Pick C1 Like 1 protein is critical for intestinal cholesterol absorption. Science 303: 1201–1204.

42. Yesilaltay A, Kocher O, Pal R, Leiva A, Quinones V, et al. (2006) PDZK1 is disruption of the PDZK1 gene in mice causes tissue-specific depletion of the high density lipoprotein receptor scavenger receptor class B type I and altered lipoprotein metabolism. J Biol Chem 281: 50280–50285.

43. Robertson MD, Parkes M, Warren BF, Ferguson DJ, Jackson KG, et al. (2003) Adipose differentiation related protein (ADRP) and the secretion of triglyceride-rich lipoproteins by Caco-2 cells. J Biol Chem 278: 25998–25970.

44. Altmann SW, Davis HR Jr, Zhu LJ, Yao X, Hoo LM, et al. (2004) Niemann-Pick C1 Like 1 protein is critical for intestinal cholesterol absorption. Science 303: 1201–1204.

45. Quinones V, Amigo L, Moreno M, Mique Jr, et al. (2001) Hepatic cholesterol and bile acid metabolism and intestinal cholesterol absorption in scavenger receptor class B type I-deficient mice. J Biol Chem 276: 13522–13527.

46. Davis HR Jr, Zhu LJ, Yao X, Hoo LM, et al. (2004) Niemann-Pick C1 Like 1 protein is critical for intestinal cholesterol absorption. Science 303: 1201–1204.