Expression and regulation of scavenger receptor class B type I (SR-BI) in gall bladder epithelium

J F Miquel, M Moreno, L Amigo, H Molina, P Mardones, I I Wistuba, A Rigotti

Background and aims: Biliary lipid absorption by the gall bladder mucosa and the cholesterol content of the gall bladder wall appear to play a role in cholesterol gall stone formation. As the scavenger receptor class B type I (SR-BI) regulates cellular cholesterol uptake, we studied its expression in human and murine gall bladders, its regulation by increased biliary lipid content, and its role in gall stone formation.

Methods and results: Using immunohistochemistry, SR-BI was found in the apical domain of human gall bladder epithelial cells. Immunoblotting of isolated membranes from gall bladder epithelial cells showed a specific signal for the 82 kDa SR-BI protein. In C57BL/6 mice, SR-BI was also found in the gall bladder epithelium. Using western blot analysis, an inverse relationship was observed between biliary cholesterol concentration and SR-BI expression in murine gall bladder mucosa. By comparing lithogenic diet fed wild-type and SR-BI deficient mice, gall bladder wall cholesterol content and gall stone formation were not found to be dependent on SR-BI expression.

Conclusions: (i) SR-BI is expressed in both human and murine gall bladder epithelium; (ii) biliary cholesterol hypersecretion is associated with decreased gall bladder SR-BI expression in mice; and (iii) murine SR-BI is not essential in controlling gall bladder wall cholesterol content and gall stone formation during diet induced cholelithiasis.

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medium consisting of a 1:1 (v/v) mixture of Dulbecco’s modified Eagle’s medium (DMEM) and Ham’s F12 (Life Technologies, Grand Island, New York, USA) containing antibiotics, as described by Auth et al with minor modifications. \(^\text{17}\) Isolation of human GB epithelial cells (GBEC) was performed between 30 minutes and one hour after surgery. The GB was incised longitudinally, the walls were reflected, and a small section of tissue was taken for histological analysis. Then, the mucosa was rinsed carefully with transport medium and wiped with gauze several times to remove adherent bile and mucus. The GB tunica mucosa was then placed in 0.125% collagenase solution (collagenase type IV; Sigma Chemicals, St Louis, Missouri, USA) for 20 minutes at 37°C. Every five minutes the mucosa was abraded thoroughly using a scalpel and flushed with DMEM medium. The resulting cell suspension was subjected twice to centrifugation at 85 g for five minutes at 20°C. An aliquot of freshly isolated GBEC was placed on a glass slide, fixed with ethanol, and stained with H&E for light microscopy. Semi-quantitative analysis demonstrated that more than 95% of cells had epithelial features. GBEC were kept at −80°C for further western blot analysis, as described below.

**Animals and diets**

C57BL/6 mice, purchased originally from the Jackson Laboratory (Bar Harbour, Maine, USA), were used to breed our own colony. Animals were housed in a humidity and temperature controlled room with reverse cycle lighting. All mice were obtained from Dr Monty Krieger (Massachusetts Institute of Technology, Cambridge, Massachusetts, USA) ad libitum, prior to the feeding experiments with cholesterol; Prolab RMH3000, PMI Feeds Inc, St Louis, Missouri, USA) and the abdominal cavity was exposed through a midline incision. Surgery was performed on mice that were fasted overnight and results were normalised to the signal generated by APN.

**Immunoblotting**

Crude membranes were prepared from isolated human GBEC, pools of murine GB and jejunal epithelium, and murine liver, as described elsewhere. \(^\text{18}\) Each sample (20 μg total membrane protein) was subjected to 10% sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) under reducing conditions and transferred to Hybond C membranes (Amersham, Arlington Heights, Illinois, USA). The blots were incubated with anti-SR-BI (1:3000 dilution) or anti-APN (1:400 dilution) polyclonal antibodies in Tris buffered saline/0.05% Tween 20 containing 5% non-fat dried milk powder. Visualisation of membrane bound antibodies was carried out with a standard chemiluminescence kit (Amersham) using horseradish peroxidase labelled mouse antirabbit IgG antibody (Sigma Chemicals) as a secondary antibody. Western blot signals were quantified by densitometric analysis (CS-9000 Dual-Wavelength Flying-Spot Scanner, Shimadzu), and results were normalised to the signal generated by APN.

**Analytical procedures**

Protein concentrations were determined with the Bradford assay, using bovine serum albumin as the standard curve. Biliary bile salts were determined by the 3α-hydroxysteroid dehydrogenase method, lecithin by the inorganic phosphorus procedure, and cholesterol by the cholesterol oxidase assay, as described previously. Unesterified and esterified cholesterol content was measured after lipid extraction \(^\text{19}\) in pools of GB samples (n=4–5) obtained from mice fed with chol or the lithogenic diet.

**Immunohistochemistry**

Tissue sections were deparaffinised and endogenous peroxidases were quenched with 0.3% H₂O₂ in methanol for 30 minutes at room temperature. Immunostaining was performed in a Nexes IHC staining system (Ventana Medical System, Inc., Tucson, Arizona, USA). Incubation with the primary antibody was carried out for 90 minutes in appropriate dilutions (anti-SR-BI 1:300). The specific signal was developed with an universal biotinylated secondary antibody and peroxidase activity was revealed with 3,3′-diaminobenzidine hydrochloride and H₂O₂, according to the reagents and instructions of the manufacturer’s detection system. Subsequently, sections were counterstained lightly with H&E, dehydrated, cleared, and mounted in Permount. Negative controls were obtained by omitting the primary antibody or by incubating sections with isotypic immunoglobulins.

**Collection of GB bile and tissues in mice**

Surgery was performed on mice that were fasted overnight with free access to water. Animals were anaesthetised with an intraperitoneal injection of pentobarbitial (4.5 mg/100 g body weight) and the abdominal cavity was exposed through a ventral incision; the cystic duct was ligated and the GB removed. Mice were then killed and the liver and 15 cm of the proximal jejunum were removed. The GB was examined visually for the presence of stones or sediment. Bile was aspirated by puncturing the GB with a fine needle and then stored at −20°C for further lipid analysis. \(^\text{19}\) GBs from animals of the same experimental group (n=4–5) were opened longitudinally, washed with phosphate buffered saline, and pooled. In some animals, following aspiration of bile as described, ethan-
Statistical analysis

Values are expressed as mean (SEM). The significance of differences between means was evaluated using the unpaired Student’s t test. The level of significance was set at p<0.05.

RESULTS

SR-BI is expressed in the apical domain of human and murine GB epithelial cells

Human GBs were obtained from patients with cholesterol gall stone disease after elective cholecystectomy and immunohistochemistry was performed to assess whether the scavenger receptor SR-BI was expressed in the GB epithelium. Routine H&E staining of the GB showed a well preserved monolayer of typical GB epithelial cells with an apical domain (arrows), and subepithelial capillary vessels (open arrowhead). Strong immunoperoxidase reaction for SR-BI protein was restricted to the plasma membrane of the apical domain of the GB epithelial cells (arrows). No staining was observed in the basolateral domain or in the tissues of the lamina propria. Original magnification: A and C, 200×; B and D, 400×.

![Figure 1](image1.png)

**Figure 1** Haematoxylin-eosin (H&E) staining [A, B] and immunohistochemical reaction for scavenger receptor class B type I (SR-BI) [C, D] in human gall bladder (GB) epithelium. Immunohistochemistry (performed as described in materials and methods) is based on a polyclonal anti-SR-BI as the primary antibody. [A, B] H&E staining revealed a well preserved monolayer of typical GB epithelial cells with an apical domain (arrows), and subepithelial capillary vessels (open arrowhead). [C, D] Strong immunoperoxidase reaction for SR-BI protein was restricted to the plasma membrane of the apical domain of the GB epithelial cells (arrows). No staining was observed in the basolateral domain or in the tissues of the lamina propria. Original magnification: A and C, 200×; B and D, 400×.

To evaluate whether SR-BI was also expressed in the GBEC of C57BL/6 mice, a well defined animal model used for diet induced cholesterol gall stone disease, GB of chow fed mice were also subjected to immunohistochemical analysis. Following ethanolic fixation, H&E staining showed a well preserved monolayer of murine GBEC (fig 3A, 3B) with identifiable apical (black arrows) and basal (white arrowheads) plasma membrane domains. Immunohistochemical analyses of murine GB with the anti-SR-BI serum are shown in figs 3C and 3D. Positive staining for SR-BI was restricted to the apical pole of the GBEC (arrows), similar to the signal found in human GB (fig 1C, 1D).

![Figure 2](image2.png)

**Figure 2** Immunoblot analysis of scavenger receptor class B type I (SR-BI) in crude membrane fractions from isolated human gall bladder epithelial cells (GBEC) and from male murine liver (chow diet). GBECs were harvested from a fresh human GB without cholelithiasis, and crude plasma membranes were isolated as described in materials and methods. Liver from a control C57BL/6 mouse was used for isolation of crude plasma membrane. Crude membranes (20 µg protein each) were submitted to sodium dodecyl sulphate-polyarylamide gel electrophoresis and electrophoretically transferred to a nitrocellulose filter. Immunoblot analysis was carried out using anti-human SR-BI polyclonal antibody. Bound antibodies were visualised as described in materials and methods.

In order to confirm whether the positive immunostaining in the GB corresponded to the well characterised SR-BI protein, GBEC were isolated from freshly harvested human GB and total cell membranes were prepared and subjected to immunoblot analysis. Figure 2 (left lane) shows the presence of the ≈82 kDa SR-BI protein in the total membrane fraction of human GBEC. Detection of the expected molecular weight of mature SR-BI protein in our GB samples is in clear contrast with the 57 kDa SR-BI protein recently reported in human GB (Johnson and colleagues, see discussion). It is noteworthy that in our GB samples, SR-BI expression levels were similar to those observed in a total membrane preparation from mouse liver (fig 2, right lane).

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The next part of the study was designed to determine whether expression of SR-BI in the GBEC could be modulated under conditions of increased GB biliary lipid content induced by dietary manipulations in mice. C57BL/6 mice were first fed for four weeks with a control diet, a diet supplemented with 2% cholesterol, or a high cholesterol/fat/bile salt lithogenic diet. As shown in table 1, GB biliary cholesterol and bile salts were increased significantly (by 2.7-fold and 1.5-fold, respectively) in animals fed the high cholesterol diet for four weeks compared with animals on the control diet. As we have shown previously, the increment in GB biliary lipids in animals fed a lithogenic diet for four weeks was even more dramatic. Compared with the control diet, GB biliary cholesterol, phospholipid, and bile salt concentrations increased by 6-, 3-, and 2.7-fold, respectively. All mice developed biliary sludge or gall stones when fed the lithogenic diet.

Figure 4 shows SR-BI protein expression from murine GB and proximal jejunum epithelia under the different experimental diets. It is worth noting that the level of expression of SR-BI in murine GB epithelia under the control diet was similar to the protein expression level in the liver (fig 2), and significantly higher than that in the proximal gut. It was evident that under either the cholesterol supplemented or lithogenic diet for four weeks, protein expression levels of SR-BI in the GB decreased dramatically, by 30% and 80%, respectively. In proximal intestine samples obtained from the same animals, only a low signal for SR-BI was visible, which was insufficient for quantifying differences in the level of intestinal SR-BI expression under the various experimental conditions.

Next we tested whether GB SR-BI expression could be regulated specifically in association with biliary cholesterol, but not bile salt, hypersecretion and whether downregulation of SR-BI in GB epithelium occurs before gall stone formation. The new experiment considered feeding of three groups of mice (n=4) for only 10 days, under the same experimental conditions and diets as described above, and an additional group of mice fed either with chow or chow supplemented with diosgenin (1% w/w) for 14 days. After 10 days, the lithogenic diet induced a 4- to 5-fold increase in GB biliary
subjected to sodium dodecyl sulphate alone, chow supplemented with 2% cholesterol (Chol), or a proximal jejunum epithelia of C57BL/6 male mice fed a diet of chow cant changes in bile salt or phospholipid content.

Figure 4 Immunoblot analysis of scavenger receptor class B type I (SR-BI) in crude membrane fractions from gall bladder (GB) and proximal jejunum epithelia of C57BL/6 male mice fed a diet of chow alone, chow supplemented with 2% cholesterol (Chol), or a lithogenic diet for four weeks or 10 days. Epithelium from GBs and jejunum from each group of mice were pooled (n=4 each), crude membranes were prepared, and aliquots (20 µg protein) were subjected to sodium dodecyl sulphate-polyarylamide gel electrophoresis analysis and transferred electrophoretically to a nitrocellulose filter, as described in the experimental procedures. Immunoblot analysis was carried out with rabbit anti-SR-BI as the primary antibody and bound antibodies were visualised as described in materials and methods. The degree of changes in SR-BI protein levels was compared with those of mice on the chow diet and are representative of two independent experiments. Quantification of bands was determined after correction for loading differences with APN, as described in materials and methods.

Cholesterol concentrations (1.08 (0.16) v 4.94 (0.86) mmol/l for control and lithogenic diets, respectively), without significant changes in bile salt or phospholipid content. Mice fed with 2% cholesterol showed no significant changes in GB biliary lipid composition after 10 days. Using polarised light microscopy, some cholesterol crystals, but not stones, were observed in GB bile of mice on the lithogenic diet. As previously described in rodents, diosgenin feeding increased GB biliary cholesterol content by 2.5-fold in C57BL6 mice (1.0 (0.2) for chow fed mice v. 2.45 (0.3) mM for diosgenin fed mice).

Pooled GB tissues of these cholesterol and diosgenin fed mice were subjected to western blot analysis, as described above, and protein expression levels for SR-BI and APN were quantified. After 10 days on the lithogenic diet, a 70% decrease in the expression level of SR-BI was observed. No changes in SR-BI expression were apparent after 10 days of feeding with the diet supplemented with cholesterol alone (fig 4). A 60% decrease in the expression level of SR-BI in the GB of diosgenin fed mice was also observed (data not shown). Taken together, these results indicate that increased GB biliary cholesterol content per se (without associated changes in any other biliary lipid components) correlated with decreased GB SR-BI expression in mice.

Lack of correlation between GB wall cholesterol content, gall stone formation, and SR-BI expression levels

Different animal models of cholesterol gall stone disease have demonstrated that a lithogenic diet induces an increment in the cholesterol content of the GB smooth muscle plasma membrane, generating a contractile defect of this organ leading to an increased residence time of lithogenic bile in the GB lumen contributing further to gall stone formation and growth. This excess of GB smooth muscle cholesterol may originate from the lithogenic GB bile itself. To explore if SR-BI was relevant in controlling the cholesterol content of the GB wall under conditions that dramatically increased biliary cholesterol concentration, we correlated GB SR-BI expression levels with cholesterol content in the GB wall. C57BL6 mice were sacrificed 3, 6, 12, and 30 days after feeding with chow or lithogenic diets. At each time point, GB bile and pooled GB walls were obtained for lipid analysis and SR-BI expression only during the first 12 days of administration of a lithogenic diet.

Figure 5 Correlation between gall bladder (GB) bile cholesterol content, GB wall total cholesterol content, and scavenger receptor class B type I (SR-BI) expression on administration of either a chow or lithogenic diet. C57BL6 mice were fed with a control or lithogenic diet for 3, 6, 12, or 30 days. Total cholesterol content of GB bile (A), protein expression of SR-BI in the GB wall, and total cholesterol content of GB tissue (B) were assessed as described in materials and methods over 30 days of diet administration. Total cholesterol content in bile increased progressively over time and correlated inversely with expression of SR-BI in the GB wall. Total cholesterol content of GB tissue correlated directly with SR-BI expression only during the first 12 days of administration of a lithogenic diet.

To further address the relevance of SR-BI expression in controlling GB wall cholesterol content and gall stone formation, mice with SR-BI deficiency in a mixed C57BL/6×129/SV background were studied. As shown in fig 6, GB cholesterol content of SR-BI knockout mice was similar to wild-type animals when fed a chow diet. After feeding the lithogenic diet...
for seven and 30 days, GB biliary cholesterol content was slightly, but not significantly, lower in SR-BI knockout mice compared with wild-type mice (2.9 (0.3) v 4.8 (1.9) mM, 3.8 (1.2) v 4.6 (1.8) mM, respectively; p>0.05). Furthermore, changes in GB wall cholesterol content of lithogenic diet fed SR-BI knockout mice were similar to those found in wild-type mice after feeding the same diet for 6, 12, or 30 days (fig 6). In addition, cumulative gall stone prevalence in SR-BI knockout mice after 30 days under a lithogenic diet was similar to that of wild-type mice (90 and 100%, n=10 and 12, respectively). These results indicate that SR-BI expression in the GB is not essential for controlling either GB cholesterol content or the development of cholesterol gall stones under a lithogenic diet.

**DISCUSSION**

In this study, we have demonstrated that SR-BI is highly expressed in GB epithelial cells. Immunohistochemical analysis of human and murine GB localises SR-BI to the apical domain of GBEC. SR-BI was markedly downregulated in mouse GBEC in association with increasing GB biliary cholesterol content. Finally, SR-BI expression was not essential in controlling either total GB wall cholesterol content under basal or lithogenic conditions or the incidence of gall stone formation.

SR-BI was initially discovered as a new class of receptor for modified lipoproteins. Later it was shown to be the first molecularly well defined cell surface high density lipoprotein receptor to be identified. Interestingly, SR-BI mediates a cellular process known as selective lipid uptake by which not only cholesteryl esters, but also various other lipid molecules (that is, free cholesterol, phospholipids) from receptor bound lipoproteins are directly transferred to the cell plasma membrane (reviewed by Rigotti and colleagues and Krieger). Furthermore, SR-BI can also facilitate both non-lipoprotein unesterified cholesterol uptake as well as free cholesterol efflux in cultured cells. Various genetic manipulations in vivo have established that SR-BI plays important physiological and pathophysiological roles in lipoprotein cholesterol metabolism in mice. More recently, SR-BI was proposed to play a role in intestinal cholesterol absorption based on SR-BI expression analysis and transport studies with intestinal brush border membranes and cells. Several studies have shown that SR-BI is indeed expressed in the apical membrane of enterocytes of rats and mice, mainly in proximal segments of the gut where cholesterol absorption take place.

More interestingly, Hauser et al used SR-BI ligands and anti-SR-BI antibodies to inhibit cholesterol uptake from bile salt mixed micelles and phospholipid vesicles into intestinal brush border preparations and CaCo2 cells. However, further studies aimed at evaluating more directly the potential role of SR-BI in intestinal cholesterol absorption in SR-BI knockout mice established that expression of SR-BI is not essential for this process.

Several studies have reported indirect evidence that GB cholesterol absorption may occur in vivo. In vitro experiments have shown that cholesterol is indeed efficiently absorbed by human GB mucosa. A recent study by Corradini et al, using an intraruminally perfused GB model, confirmed that the mucosa of normal human and pig GB is able to absorb 20–30% of biliary cholesterol over a period of five hours. As 800–1200 mg of free cholesterol are secreted daily from the liver into the biliary tract in humans, absorption of at least 20% of biliary cholesterol by the GB could be similar to the daily amount of dietary cholesterol absorbed in the intestine in subjects on a typical Western diet. This significant amount of biliary cholesterol uptake by the GB may have major implications for biliary cholesterol solubility, intestinal cholesterol absorption, and, ultimately, for body cholesterol homeostasis. In fact, this function is impaired in the GB of cholesterol gall stone patients suggesting that lack of this absorptive capacity may contribute to gall stone formation. Despite the potential physiological and pathophysiological relevance of GB function in cholesterol absorption, the detailed cellular and molecular mechanisms by which biliary lipids are actually absorbed by the GB epithelium remains unknown. A recent study has reported that apolipoprotein (apo) A-I, a normal component of human bile, stimulated cholesterol transfer from phospholipid vesicles on the mucosal side to the serosal side of cultured polarised GBEC. In addition, apo A-I, an amphipatic anionic polypeptide that exhibits immunological cross reactivity with apo A-I, could serve as a protein ligand for candidate apical GBEC receptors, which might be involved in GB cholesterol absorption. In this regard, the previously reported functional properties of SR-BI and the enterocyte cholesterol transport inhibition experiments together with its high level of expression in GBEC and its localization in the apical domain of these cells, as shown in this study, suggested a potential role of this cell surface receptor in GB cholesterol absorption.

While our western blot analysis showed the presence of a 82 kDa SR-BI protein in human and murine GB, a previous study has reported an immunoreactive SR-BI protein of 57 kDa, but not 82 kDa, in murine GB. It is interesting that this latter study also showed that most of the hepatic and ovarian protein detected by the anti-SR-BI antibodies exhibited a 57 kDa molecular weight. Johnson et al, who used SR-BI ligands and molecular mechanisms by which biliary lipids are actually absorbed by the GB epithelium remains unknown. A recent study has reported that apolipoprotein (apo) A-I, a normal component of human bile, stimulated cholesterol transfer from phospholipid vesicles on the mucosal side to the serosal side of cultured polarised GBEC. In addition, apo A-I, an amphipatic anionic polypeptide that exhibits immunological cross reactivity with apo A-I, could serve as a protein ligand for candidate apical GBEC receptors, which might be involved in GB cholesterol absorption. In this regard, the previously reported functional properties of SR-BI and the enterocyte cholesterol transport inhibition experiments together with its high level of expression in GBEC and its localization in the apical domain of these cells, as shown in this study, suggested a potential role of this cell surface receptor in GB cholesterol absorption.

Another important finding of this study was the dramatic downregulation of GB SR-BI expression under conditions of high biliary lipid content. Cholesterol feeding either in a high cholesterol diet or a high cholesterol/high fat/cholic acid lithogenic diet for four weeks resulted in lowered expression of SR-BI in murine GB (fig 4, 5), which correlated with an increased concentration of GB biliary lipids (table 1). Both the 2% cholesterol diet and the cholesterol rich lithogenic diet produced not only a dramatic increase in GB bile cholesterol concentration but also a significant increase in biliary phospholipids and/or bile salts. Therefore, this in vivo experimental model did not allow identification of exactly which increase in biliary lipid content was correlated with regulation
of SR-BI expression in GB epithelia. To determine whether a selective change in biliary cholesterol could be associated with regulation of GB SR-BI expression, a new set of experiments were carried out feeding mice with the same diets for a shorter time period. When C57BL/6J mice were fed a lithogenic diet for only specific biliary cholesterol concentration increased markedly, without significant changes in biliary bile salts or phospholipid content. Under this experimental condition, a clear downregulation of SR-BI was found (figs 4, 5). However, changes in bile salt composition are expected in lithogenic diet fed mice as dietary cholic acid is rapidly incorporated into the endogenous bile salt pool. Thus changes in bile acid composition apart from increased biliary cholesterol content may also explain downregulation of GB SR-BI. To gain more insight into the specific bile component associated with changes in GB SR-BI expression, we performed a diosgenin feeding experiment in mice. Diosgenin is a plant steroid that selectively increases biliary cholesterol secretion but does not affect biliary secretion of bile salts and phospholipids or bile salt pool composition. Increased GB bile cholesterol content under a diosgenin supplemented diet also correlated with significant downregulation of SR-BI expression in GB epithelial cells. If bile is the source of cholesterol that regulates SR-BI expression in GBEC, our results may be explained by cholesterol dependent modulation of the transcriptional rate of the target gene. In fact, it has been shown that the promoter region of the SR-BI gene contains sterol response elements through which sterol regulatory element binding proteins can bind and thereby regulate transcription of this gene. Additional studies are required using polarised monolayer GBEC in cultured conditions to gain more insights into the mechanisms by which SR-BI expression might be regulated by cholesterol in GBECs. This study was not aimed at directly evaluating the role of SR-BI in biliary lipid absorption in the GB. However, we performed additional experiments to test if GB SR-BI expression could be relevant in controlling GB wall cholesterol content and gall stone formation as an indirect measurement of a functional role of SR-BI expressed in the GB. Under lithogenic diet feeding, GB wall cholesterol content significantly decreased as early as three or six days and this change correlated with downregulation of SR-BI expression levels until day 12. However, after 30 days under a lithogenic diet, the cholesterol content of the GB wall intriguingly returned to basal levels while SR-BI expression levels remained low. These results suggest that the SR-BI expression is not always functionally linked to control of GB wall cholesterol content. Alternatively, SR-BI downregulation could be involved in controlling GB cholesterol content but compensatory mechanism(s) may overcome this effect when a lithogenic diet is maintained for longer periods of time. However, the fact that SR-BI knockout mice showed similar GB wall cholesterol content under chow or lithogenic diets in comparison with wild-type mice strongly suggests that SR-BI expression in the GB epithelium does not play a major role in controlling GB wall cholesterol content. Moreover, this study clearly demonstrated that SR-BI expression is not essential for cholesterol gallstone formation under exogenous cholesterol supply with a standard lithogenic diet. Recent data published during the preparation of this manuscript also indicated that SR-BI is not critical in controlling either the formation of lithogenic bile or formation of gall stones. In conclusion, we have demonstrated the presence of SR-BI expression in the apical membranes of human and mice GBECs. In the murine GBEC, SR-BI was downregulated in association with increased levels of GB biliary cholesterol. However, SR-BI expression in GB epithelial cells was not critical in controlling the cholesterol content of the GB wall or gall stone formation under a lithogenic diet. Further studies are required to elucidate the regulation of SR-BI in the GB and its function in physiology, particularly in transepithelial cholesterol transport as well as its potential role in prevalent GB disease conditions such as cholesterol gall stone and cholesterolosis.

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