Targeted disruption of the murine cholecystokinin-1 receptor promotes intestinal cholesterol absorption and susceptibility to cholesterol cholelithiasis

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Cholecystokinin (CCK) modulates contractility of the gallbladder, the sphincter of Oddi, and the stomach. These effects are mediated through activation of gastrointestinal smooth muscle as well as enteric neuron CCK-1 receptors (CCK-1R). To investigate the potential physiological and pathophysiological functions linked to CCK-1R-mediated signaling, we compared male WT and CCK-1R−/− mice (129/SvEv). After 12 weeks on either a standard mouse chow or a lithogenic diet (containing 1% cholesterol, 0.5% cholic acid, and 15% dairy fat), small-intestinal transit time, intestinal cholesterol absorption, biliary cholesterol secretion, and cholesterol gallstone prevalence were compared in knockout versus WT animals. Analysis of mice on either the chow or the lithogenic diet revealed that CCK-1R−/− animals had larger gallbladder volumes (predisposing to bile stasis), significant retardation of small-intestinal transit times (resulting in increased cholesterol absorption), and increased biliary cholesterol secretion rates. The elevation in bile cholesterol, coupled with a tendency toward gallbladder stasis (due to the absence of CCK-induced contraction), facilitates nucleation, growth, and agglomeration of cholesterol monohydrate crystals; this sequence of events in turn results in a significantly higher prevalence of cholesterol gallstones in the CCK-1R−/− null mice.

Introduction
Cholecystokinin (CCK) is a neuroendocrine peptide hormone that exhibits a variety of effects on many target tissues (1). Molecular, cellular, and physiologic approaches have been used to establish a functional role of the CCK-1 receptor (CCK-1R) on gallbladder smooth muscle, on pancreatic acini, at many levels in the gastrointestinal tract, and in the enteric nervous system and brain (2–4). Gallbladder contractile dysfunction in response to exogenously administered CCK or CCK octapeptide (CCK-8) has been observed in cholesterol gallstone patients (5, 6) and to a lesser extent (7) in pig gut. Administration of CCK or CCK octapeptide (CCK-8) to normal humans results in increased bile flow (= 37), significantly (P < 0.01) greater than the volume in WT mice of 26 ± 18 μl (n = 26). Furthermore, gallbladder volumes and prevalence rates, and characteristics of gallstones.

Nonstandard abbreviations used: body weight (BW); CCK octapeptide (CCK-8); CCK-1 receptor (CCK-1R); cholecystokinin (CCK); cholesterol saturation index (CSI).

Conflict of interest: The authors have declared that no conflict of interest exists.

Figure 1

Relative lipid compositions of bile specimens from CCK-1R–/– and WT mice. Relative lipid compositions (mol per 100 mol) are plotted on partial condensed phase diagrams according to the approximate total lipid concentrations of the bile samples (A, 10.0 g/dl for gallbladder bile; B, 2.0 g/dl for hepatic bile; see Table 1). The one-phase micellar zone (at bottom) is enclosed by a solid curved line, and two solid and two dashed lines divide the phase diagram into regions a–e with different crystallization sequences (see ref. 24). (A) Lipid compositions of pooled gallbladder bile specimens (n = 20 per group) from the CCK-1R–/– (circle) and WT mice (square) fed the lithogenic diet for 12 weeks are located in a central three-phase area where, at equilibrium, bile samples are composed of cholesterol-saturated mixed micelles, solid cholesterol crystals, and liquid crystals, as observed by microscopy. (B) Analogous regions of the condensed phase diagram exhibit the same physical states at equilibrium as those in the phase diagram shown in A; however, with decreases in total lipid concentration all crystallization pathways are shifted to the left and the micellar zone becomes smaller. These alterations generate a new condensed phase diagram with an enlarged region e. Lipid compositions of individual hepatic bile specimens (n = 6 per group) from both WT (squares) and CCK-1R–/– mice (circles) locate in region e, where, at equilibrium, the bile samples are composed of liquid crystals and saturated micelles, but no solid cholesterol monohydrate crystals are present.

Table 1

Biliary lipid compositions of pooled gallbladder and individual hepatic bile specimens

<table>
<thead>
<tr>
<th>Mouse</th>
<th>Mol% Ch</th>
<th>Mol% L</th>
<th>Mol% BS</th>
<th>L/(L + BS)</th>
<th>[TL] (g/dl)</th>
<th>CSI</th>
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<tr>
<td>WT</td>
<td>7.84</td>
<td>19.07</td>
<td>73.09</td>
<td>0.21</td>
<td>9.35</td>
<td>1.21</td>
</tr>
<tr>
<td>CCK-1R–/–</td>
<td>9.84</td>
<td>18.55</td>
<td>71.61</td>
<td>0.21</td>
<td>9.99</td>
<td>1.50</td>
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Individual hepatic bile specimens

WT

<table>
<thead>
<tr>
<th>n</th>
<th>Mol% Ch</th>
<th>Mol% L</th>
<th>Mol% BS</th>
<th>L/(L + BS)</th>
<th>[TL] (g/dl)</th>
<th>CSI</th>
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</thead>
<tbody>
<tr>
<td>1</td>
<td>6.66</td>
<td>16.53</td>
<td>76.81</td>
<td>0.18</td>
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<td>1.81</td>
</tr>
<tr>
<td>2</td>
<td>4.29</td>
<td>14.13</td>
<td>81.58</td>
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</tr>
<tr>
<td>3</td>
<td>3.86</td>
<td>12.29</td>
<td>83.84</td>
<td>0.13</td>
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<tr>
<td>4</td>
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<td>16.99</td>
<td>77.75</td>
<td>0.18</td>
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<td>5</td>
<td>7.08</td>
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<td>71.74</td>
<td>0.23</td>
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<tr>
<td>6</td>
<td>5.36</td>
<td>17.13</td>
<td>77.51</td>
<td>0.18</td>
<td>1.65</td>
<td>1.34</td>
</tr>
</tbody>
</table>

Mean ± SD 5.42 ± 1.27 16.37 ± 3.02 78.21 ± 4.19 0.18 ± 0.03 1.96 ± 0.47 1.35 ± 0.26

CCK-1R–/–

<table>
<thead>
<tr>
<th>n</th>
<th>Mol% Ch</th>
<th>Mol% L</th>
<th>Mol% BS</th>
<th>L/(L + BS)</th>
<th>[TL] (g/dl)</th>
<th>CSI</th>
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<td>7.09</td>
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<td>73.82</td>
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<td>1.61</td>
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Mean ± SD 6.91 ± 0.85 15.76 ± 2.08 77.33 ± 1.93 0.17 ± 0.02 1.92 ± 0.20 1.76 ± 0.31

Values were measured from pooled gallbladder bile specimens (n = 20 per group) and individual hepatic bile specimens (total n = 6 in each group). The CSIs of pooled gallbladder and six hepatic bile samples were calculated from the critical tables (23). ¥P < 0.05 compared with hepatic bile specimens of WT mice. CCK-1R, cholecystokinin-1 receptor; Ch, cholesterol; L, lecithin; BS, bile salt; [TL], total lipid concentration.

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see Table 1) (24). Relative lipid compositions of gallbladder bile samples from CCK-1R–/– mice fed the lithogenic diet plot in a central three-phase area denoted region e. By phase analysis, these bile samples are predicted to be composed of cholesterol monohydrate crystals, liquid crystals, and cholesterol-saturated mixed micelles at equilibrium (24, 25), as was observed experimentally. Gallbladder bile specimens of the WT mice fed the lithogenic diet also plotted in the central three-phase area; however, compared with the CCK-1R–/– mice they contained appreciably lower amounts of cholesterol.

We plot in Figure 1B the relative lipid compositions of individual hepatic bile samples (from Table 1) in relation to an appropriate micellar phase boundary (24, 25) and cholesterol crystallization pathways for dilute (2.0 g/dl) taurocholate-rich bile (rounded-off value for means of both sets of hepatic bile specimens in Table 1) (25). Relative lipid compositions plot within the crystallization pathway denoted region e, where only liquid crystals but never solid crystals phase separate at equilibrium (24). As observed in model (24) and native human bile samples (26), when total lipid concentrations decrease, equilibrium zones for all physical states and crystallization pathways shift markedly to the left, that is, to lower lecithin contents, and, in addition, the boundary of the micellar zone contracts downward. Our results show that relative lipid compositions from CCK-1R–/– mice and WT controls plot in region e, with relative lipid compositions of hepatic bile samples from CCK-1R–/– mice shifting upward significantly (P < 0.05), indicating increases in relative cholesterol contents (Table 1).

When gallbladder and hepatic bile samples from mice on a lithogenic diet were analyzed, all bile salt compositions from both WT and CCK-1R–/– mice were found to be taurine conjugated and displayed similar molecular species distributions. Because of the 0.5% cholic acid included in the lithogenic diet, taurocholate was, as expected (27, 28), the major bile salt (75.4–76.9%), followed by diminishing relative proportions of tauro-β-muricholate (10.3–10.6%) and taurodeoxycholate (2.0–2.1%), taurochenodeoxycholate (1.6–1.7%), and tauro-α-muricholate (0.8–0.9%) were also found. Hydrophobicity indexes (29) of gallbladder (0.03–0.06) and hepatic bile specimens (0.03–0.05) were identical between WT and CCK-1R–/– mice.

Bile flow and biliary lipid secretion rates. Bile flow rates after 12 weeks of feeding the lithogenic diet (see Methods) were measured during the first hour after interruption of the enterohepatic circulation, thereby avoiding appreciable perturbation of the enterohepatic circulation of bile salts (27). In CCK-1R–/– mice, the bile flow rate (69 ± 10 μl/min/100 g BW) was similar to that of the WT mice (64 ± 16 μl/min/100 g BW). Figure 2 plots the individual biliary lipid outputs in WT and CCK-1R–/– mice during the first hour of acute fistulation. A significant (P < 0.05) difference was observed only in biliary cholesterol secretion rates, which were 9.7 ± 1.7 μmol/h/kg BW in CCK-1R–/– mice, and 6.8 ± 2.1 μmol/h/kg BW in WT mice. No appreciable differences in lecithin or bile salt outputs (Figure 2) were noted; in CCK-1R–/– mice, these values were 21.9 ± 3.0 and 106.2 ± 3.4 μmol/h/kg BW, respectively, whereas in WT mice the values were 20.8 ± 4.9 and 113.0 ± 4.7 μmol/h/kg BW, respectively.

Volumes of the circulating bile salt pool were also identical between CCK-1R–/– (2.4 ± 0.3 μmol) and WT mice (2.3 ± 0.2 μmol). However, after factoring in each bile salt mass measured in pooled gallbladder bile samples, total bile salt pool sizes became appreciably greater in CCK-1R–/– mice (7.2 μmol) compared with WT mice (5.4 μmol). This finding is consistent with the significantly enlarged fasting gallbladder volumes in CCK-1R–/– mice compared with those of WT mice.

Intestinal cholesterol absorption by dual-isotope and cholesterol balance methods. Displayed in Figure 3 are values for percentage cholesterol absorption in mice ingesting chow as calculated from the plasma ratios of [1H]cholesterol and [3H]cholesterol 3 days after dosing (30, 31). Percentage cholesterol absorption values were significantly (P < 0.01) larger in CCK-1R–/– mice (38% ± 7%) compared with WT mice (30% ± 5%). Since chow contains trace cholesterol (<0.02%), and since both CCK-1R–/– and WT mice ate similar amounts of food (4.3–4.5 g/day), we calculated total cholesterol mass absorbed from the small intestine to be approximately 0.33 mg/day in CCK-1R–/– mice and approximately 0.26 mg/day in WT mice. To demonstrate that this observation was sustained during feeding the lithogenic

Figure 2
Total biliary lipid outputs (μmol/h/kg BW) during the first hour of interruption of the enterohepatic circulation in WT and CCK-1R–/– mice (n = 6 per group) after feeding the lithogenic diet for 12 weeks. (A) The CCK-1R–/– mice displayed significantly higher biliary cholesterol outputs (P < 0.05) compared with the WT mice. However, biliary outputs of (B) lecithin and (C) bile salts were similar between the two sets of mice.
diet, it became necessary to study cholesterol absorption in mice ingesting the diet high in cholesterol, cholic acid, and triglycerides, as used for the cholesterol gallstone prevalence study.

Table 2 summarizes the per diem results for cholesterol intake, biliary cholesterol outputs, fecal total neutral steroid excretion, absorbed cholesterol mass, and percentage cholesterol absorption in CCK-1R/- and WT mice in the metabolic steady state while on the lithogenic diet or ingesting chow. Basically, we found that CCK-1R/- and WT mice ingested similar amounts of food whether they were fed the chow or the lithogenic diet. On the chow diet (P < 0.02% cholesterol), biliary cholesterol secretion rates in CCK-1R/- mice (1.65 ± 0.27 mg/day) were significantly (P < 0.01) higher than in the WT mice (1.24 ± 0.16 mg/day). Because of their higher biliary cholesterol outputs, CCK-1R/- mice showed significantly (P < 0.05) augmented daily fecal neutral steroid excretion (2.15 ± 0.27 mg/day) compared with WT mice (1.81 ± 0.16 mg/day). Nonetheless, an input–output analysis revealed that the absorbed cholesterol mass in CCK-1R/- mice (0.31 ± 0.06 mg/day) was significantly (P < 0.05) larger than in the WT mice (0.24 ± 0.04 mg/day). The calculated percentage cholesterol absorption in CCK-1R/- mice was 38.2% ± 5.5%, a value that is significantly (P < 0.01) greater than in WT mice (29.3% ± 4.2%), and both are essentially identical to the results obtained with the dual-isotope method (see earlier). While ingesting the lithogenic diet, CCK-1R/- mice also secreted significantly (P < 0.01) more biliary cholesterol than the WT mice (Table 2). Furthermore, cholesterol mass absorbed from the small intestine was significantly (P < 0.001) greater in CCK-1R/- mice than in the WT mice, consistent with the latter excreting appreciably more neutral steroids in feces. As was clarified elsewhere (27), we note that compared with chow, cholesterol absorption efficiency measured by the mass balance method was increased significantly (P < 0.001) in both strains of mice fed the lithogenic diet, mostly as a result of the dietary cholic acid (27). Furthermore, with lithogenic-diet feeding, CCK-1R/- mice continued to display significantly (P < 0.001) higher percentage cholesterol absorption (48.4% ± 4.8%) compared with the WT mice (34.9% ± 4.0%). Taken together, these results suggest that CCK-1R/- mice absorb more cholesterol from the small intestine than do WT mice, whether they are fed trace or high levels of dietary cholesterol (and cholic acid).

Small-intestinal transit times. Figure 4 depicts the distributions of radioactivity administered as [3H]sitostanol along the length of the small intestines of CCK-1R/- and WT mice while each strain was being fed chow (Figure 4A) or the lithogenic diet (Figure 4B) for 14 days. These data were obtained at exactly 30 minutes after intraduodenal instillation of the radioisotope dissolved in medium-chain triglyceride (see Methods and references therein). On the chow diet, distributions of radioactivity lengthwise throughout the small intestine were significantly (P < 0.01) different between the WT and CCK-1R/- mice, with peaks occurring between segments 8 and 15 in WT mice compared with segments 4 and 11 in CCK-1R/- mice. The geometric center (see Methods) of the [3H]sitostanol distribution profiles in the small

### Table 2

<table>
<thead>
<tr>
<th>Mouse</th>
<th>Cholesterol intake (mg/day)</th>
<th>Biliary cholesterol (mg/day)</th>
<th>Steroid excretion (mg/day)</th>
<th>Absorbed cholesterolA (mg/day)</th>
<th>Cholesterol absorptionB (%)</th>
</tr>
</thead>
<tbody>
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<td>Chow diet</td>
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<td></td>
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</tr>
<tr>
<td>WT</td>
<td>1.80 ± 0.09</td>
<td>1.29 ± 0.13</td>
<td>2.08 ± 0.29</td>
<td>35.1 ± 4.0</td>
<td>43.9% ± 7.2</td>
</tr>
<tr>
<td>CCK-1R/-</td>
<td>1.65 ± 0.27</td>
<td>2.15 ± 0.27</td>
<td>30.88 ± 1.6</td>
<td>29.3% ± 4.2</td>
<td>38.2% ± 5.5</td>
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<tr>
<td>Lithogenic diet</td>
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</tr>
<tr>
<td>WT</td>
<td>1.24 ± 0.16</td>
<td>2.08 ± 0.16</td>
<td>35.27 ± 1.33</td>
<td>31.2% ± 4.0</td>
<td>34.9% ± 4.0</td>
</tr>
<tr>
<td>CCK-1R/-</td>
<td>1.99 ± 0.61</td>
<td>29.13 ± 4.70</td>
<td>29.3% ± 4.70</td>
<td>31.2% ± 4.0</td>
<td>34.9% ± 4.0</td>
</tr>
</tbody>
</table>

A: Absorbed cholesterol was determined by subtracting the daily fecal neutral steroid output from the daily cholesterol intake and the daily biliary cholesterol output as measured by HPLC (31). B: The percentage cholesterol absorption was determined by the cholesterol balance analysis according to published methods (31).

P < 0.01 compared with WT mice on chow. P < 0.001 compared with WT mice on chow. P < 0.0001 compared with WT mice on chow. P < 0.001 compared with WT mice fed the lithogenic diet. P < 0.001 compared with WT mice fed the lithogenic diet.
Cholesterol gallstone formation is associated with biliary cholesterol supersaturation, rapid cholesterol crystallization through several intermediate steps, and gallbladder stasis (9). Recent progress in understanding the molecular genetic (32) and physicochemical (28, 33) basis of biliary cholesterol hypersecretion and cholesterol crystallization has provided many new insights into the complex pathophysiological mechanisms involved in murine cholesterol gallstone formation. The present study in a new murine model highlights two important issues: (a) gallbladder size and small-intestinal motility are, in part, mediated by CCK-1R–induced signaling, and (b) the CCK-1R control of small-intestinal transit times is a physiological response that seems to be an important factor for the putative risk factor for murine cholesterol gallstone formation.

Discussion

Cholesterol gallstone formation is associated with biliary cholesterol supersaturation, rapid cholesterol crystallization through several intermediate steps, and gallbladder stasis (9). Recent progress in understanding the molecular genetic (32) and physicochemical (28, 33) basis of biliary cholesterol hypersecretion and cholesterol crystallization has provided many new insights into the complex pathophysiological mechanisms involved in murine cholesterol gallstone formation. The present study in a new murine model highlights two important issues: (a) gallbladder size and small-intestinal motility are, in part, mediated by CCK-1R–induced signaling, and (b) the CCK-1R control of small-intestinal transit times is a physiological response that seems to be an important factor for the putative risk factor for murine cholesterol gallstone formation.

It has been observed in rodents that CCK inhibits gastric emptying (34). However, the physiological response of the small intestine to CCK has remained unclear until now. Expression of the Cck-1r gene has been found not only on smooth muscle of gallbladder and stomach, but also on smooth muscle of the small intestine (35). We observed that CCK-1R−/− mice displayed significantly slower small-intestinal transit times compared with WT mice, whether ingesting chow or the lithogenic diet. Compared with the chow diet, the lithogenic diet accelerated small-intestinal transit times slightly in both strains of mice. Because small-intestinal transit time in mice is not influenced by feeding large amounts of dietary cholesterol or cholic acid (36), such an alteration could only be induced by fatty acids derived from digestion of dietary triglycerides in the lithogenic diet. Nonetheless, our results indicate that a physiologically relevant mechanism mediated by CCK-1R regulates small-intestinal motility appreciably. Furthermore, a retardation of small-intestinal transit time enhances cholesterol absorption from the intestine (Table 2, Figure 3), most likely because of a longer residence time of the sterol in the small-intestinal lumen. This, in turn, would increase cholesterol's incorporation into mixed micelles and also would promote partitioning of cholesterol monomers out of micelles, rendering them available for intestinal capture by cholesterol influx transporters(s) on apical membranes of small-intestinal enterocytes (37). This suggests that the transit rate of cholesterol molecules throughout the small intestine is a crucial and, in health, a normal physiological step in the regulation of cholesterol absorption. Surprisingly, this concept has not been readily appreciated, although it was confirmed in humans over two decades ago (21). In the present study, we prove that CCK and its CCK-1R regulate intestinal cholesterol absorption physiologically in an animal model. Moreover, dysfunction of this receptor has deleterious effects on biliary cholesterol solubility because it leads to the secretion into bile of larger amounts of cholesterol. In earlier work we found that the percentage of cholesterol absorption decreased progressively with increases in dietary cholesterol content over the examined range of 0.02–2% by weight (31). This significant negative relationship was secondary to the pronounced effect of dietary cholesterol diluting the trace mass of radioisotope in the upper small-intestinal lumen (27, 38). Therefore, in the present study we used a highly precise and validated cholesterol balance method (31) to examine cholesterol absorption efficiency in both CCK-1R−/− and WT mice while they were ingesting the lithogenic diet. We found that CCK-1R−/− mice displayed significantly higher intestinal cholesterol absorption efficiency compared with WT mice, whether mice were ingesting chow or the lithogenic diet (see Table 2). Moreover, we show that the cholesterol mass absorbed from the intestine, which reaches the liver through the chylomicron remnant pathway (39), must have in turn enhanced biliary cholesterol secretion and thereby induced biliary cholesterol supersaturation in our CCK-1R−/− mice. We reported in abstract form (40) that efficiency of intestinal cholesterol absorption actually correlates positively with prevalence of cholesterol gallstones in a large series of inbred mouse strains fed the lithogenic diet. These concepts suggest, therefore, that high intestinal cholesterol absorption efficiency is an independent pathophysiological risk factor for murine cholesterol gallstone formation.

Although it has been suggested in the past (41–44) that motility of the small bowel as well as the large bowel is sluggish in cholesterol gallstone patients, the pathogenetic mechanisms for the putative role of these conditions in cholelithogenesis are not known. In these earlier studies the authors focused on the role of the large intestine.

Figure 4
Small-intestinal transit rates in WT mice (top panels) and CCK-1R−/− mice (bottom panels) ingesting chow (A) or the lithogenic diet (B) for 14 days. Data were determined by the distribution of radioactivity at 30 minutes along the entire length of the small intestine after intraduodenal instillation of [3H]sitostanol dissolved in medium-chain triglyceride (30). Each bar shows the percent of radioactivity in each segment for n = 13 mice per group. Segments 1–20 represent evenly divided portions from the most proximal to the most distal parts of the small intestine placed on a 50-cm ruler (see Methods). (A) Arrows indicate the geometric center, which is significantly (P < 0.001) shorter in chow-fed CCK-1R−/− mice, indicating significantly slower small-intestinal transit times (geometric center = 7.8 ± 0.8) compared with the WT mice (geometric center = 10.8 ± 1.0). (B) Upon ingesting the lithogenic diet, CCK-1R−/− mice continue to display significantly slower small-intestinal transit times (geometric center = 9.1 ± 1.5) compared with the WT mice (geometric center = 13.3 ± 2.0).


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in cholelithogenesis, and so their hypothesis differs fundamentally from our observations in the CCK-1R−/− mouse model. The authors of these human studies (41–44) proposed that prolonged large-bowel transit time, together with greater numbers of anaerobes displaying 7α-dehydroxylation activity, promote increased formation and absorption of deoxycholic acid from the large intestine (41–44). Increased amounts of deoxycholate conjugates in bile are believed to be contributory to cholesterol gallstone formation (45) because they promote biliary cholesterol hypersecretion as well as accelerating cholesterol crystallization in gallbladder bile. However, in our studies we failed to find any differences in composition of biliary bile salt species between CCK-1R−/− and WT mice (see Results). Moreover, we reported elsewhere (46) that feeding 0.5% deoxycholic acid increased biliary deoxycholate pools in male C57L/J mice by 27% compared with the typical levels on chow (3.4%), but small-intestinal motility was not affected (46). Interestingly, in none of the aforementioned human studies was the possibility of increased cholesterol absorption by a sluggish small-intestinal transit addressed, nor was any allusion made to the possibility that increased cholesterol absorption from the intestine promotes the formation of lithogenic bile.

Impairment of gallbladder emptying in cholesterol gallstone patients (8, 9) is not the result of the physical presence of gallstones per se, since the defect does not correlate with the size or number of stones, and lithotripsy-induced gallbladder ablation does not reverse the gallbladder motility defect (47). Gallbladder contraction in response to i.v. CCK administration in cholesterol gallstone patients is diminished compared with the response of normal subjects to the same treatment (48), suggesting that some gallstone patients might have impaired gallbladder CCK-1R function as well as reduced CCK-1R number and/or CCK-binding capacity to CCK-1Rs (49, 50). Because there is impaired smooth muscle contractility in cell isolates from human gallbladders and from animal models with cholesterol gallstones, Behar and colleagues (51–53) have proposed that the gallbladder contractile mechanisms involve CCK-1R-mediated activation of phospholipase C, leading to signal-transduction decoupling when the sarcoclemmas of the smooth muscle cells of the gallbladder are highly enriched in cholesterol. This may be attributed to a cholesterol-dependent decrease in the number of or dysfunction in CCK-1Rs of gallbladder smooth muscle (52), leading to impaired contractility. A likely possibility is that this results from insertion of cholesterol molecules absorbed from bile within caveolin rafts of sarcosomal membranes. It is apparent from the present work that a similar phenomenon may be active in the CCK-1R−/− mice. As can be inferred from Table 1, there is a 0.26-unit difference in mean CSI between hepatic and gallbladder bile specimens in CCK-1R−/− mice compared with a smaller difference in mean CSI of 0.14 in WT mice. This is consistent with the possibility that, because of gallbladder stasis, biliary cholesterol molecules are absorbed in excess from gallbladders of CCK-1R−/− compared with the situation in WT mice. Our results therefore support the concept (51–53) that the crucial CCK-1R signal-transduction pathways may become yet more dysfunctional via the excess absorbed biliary cholesterol molecules in CCK-1R−/− mice, thus compromising gallbladder contractility and promoting cholelithogenesis. In parallel with the findings in humans, our results substantiate the concepts that the absence of CCK-1R in mice perturbs the physiological control of gallbladder contraction as well as small-intestinal motility, leading to organ dysfunction with absorption of excess cholesterol from the lumina of both organs. The scenario for cholesterol gallstone formation in CCK-1R−/− is propagated by a vicious cycle, since bile is supersaturated with cholesterol as a result of cholesterol absorption from the small intestine, and therefore gallbladder function is in turn further compromised by absorbed cholesterol molecules from cholesterol-supersaturated bile within its lumen. Hence, the CCK-1R–null mouse provides an excellent animal model for the study of cholesterol gallstone formation as promoted by these myodysfunctional mechanisms. Furthermore, the model can be systematically evaluated to investigate principles underlyng gallbladder and small-intestinal contractile functions in cholelithogenesis, at both the molecular and the cellular levels.

In summary, ablation of the Cck-1r gene impairs small-intestinal transit in mice whether chow or a lithogenic diet is ingested; this impairment in turn leads to increased intestinal cholesterol absorption, which results in enhanced biliary cholesterol secretion. Absence of functional CCK-1Rs also increases gallbladder size and cholesterol absorption from bile, which further impairs muscle contractility and probably results in profound gallbladder stasis (54). These dysfunctions of gastrointestinal motility contribute to cholesterol gallstone formation by facilitating more efficient intestinal cholesterol absorption and biliary cholesterol hypersecretion, leading to crystallization and growth, as well as agglomeration of solid-cholesterol crystals in gallbladders that have become “large, lax and lazy” (55). These findings, taken together, indicate that prolonged small-intestinal transit times, increased cholesterol absorption, and gallbladder hypomotility are important interdependent and conflating risk factors for cholesterol gallstone formation in this animal model.

**Methods**

**Chemicals.** Intralipid (20%, wt/vol) was purchased from Pharmacia (Clayton, North Carolina, USA), and medium-chain triglyceride was obtained from Mead Johnson Nutritionalscs (Evansville, Indiana, USA). Radioisotopes [1,2-3H]cholesterol and [4-14C]cholesterol were purchased from NEN Life Science Products Inc. (Boston, Massachusetts, USA), and [5,6-3H]sitostanol was obtained from American Radiolabeled Chemicals Inc. (St. Louis, Missouri, USA). For HPLC of biliary lipids, all reagents were spectra-analyzed HPLC grade from Fisher Scientific Co. (Fair Lawn, New Jersey, USA).

**Animals and diets.** Generation of CCK-1R−/− mice and general phenotypic properties have been reported previously (22). The CCK-A receptor is now referred to as CCK-1R (official gene symbol is Cck-1r) (56). We studied male homozygous CCK-1R knockout mice and WT mice of the same 129/SvEv background (Charles River Laboratories, Wilmington, Massachusetts, USA) at 3–6 months of age. All animals were maintained in a temperature-controlled room (22 ± 1°C) with 12-hour light (6 am–6 pm) cycles. Mice were allowed free access to water and standard Purina rodent chow (Purina Mills Inc., St. Louis, Missouri, USA), which contains trace quantites (<0.02%) of cholesterol (27). During gallstone-induction experiments, animals were fed a semi-synthetic lithogenic diet (28) containing 1% cholesteral, 0.5% cholic acid, and 15% butterfat for 12 weeks. All experiments were executed according to accepted criteria for the care and experimental use of laboratory animals, and euthanasia was consistent with recommendations of the American Veterinary Medical Association. Ethical protocols for animal experimentation were approved by the Institutional Animal Care and Research Committees of Harvard Medical School and Tufts–New England Medical Center, both of Boston, Massachusetts, USA.

**Collection and microscopic analysis of gallbladder bile specimens and gallstones.** After 12 weeks on the lithogenic diet, the WT mice (n = 26) and CCK-1R−/− mice (n = 37) were fasted overnight but allowed free access to water. Animals were weighed and anesthetized with an intraperitoneal injection of 35 mg/kg pentobarbital (Abbott Laboratories, North Chicago, Illinois, USA). After cholecystectomy (28), gallbladder volume was measured by weighing the whole gallbladder and numerically estimating gallbladder weight with
gallbladder volume (28). Gallbladders were then opened at the fundus, and 5 μl of fresh gallbladder bile were digitally expressed and examined by direct and polarized light microscopy for mucin gel, liquid crystals, cholesterol crystals, and stones according to previously established criteria (28). Pooled gallbladder bile samples were ultracentrifuged at 100,000 g for 30 minutes at 37°C and filtered through a preheated (37°C) Swinnex-GS filter (0.22 μm) assembly (Millipore Corp., Bedford, Massachusetts, USA). Samples were then frozen and stored at −20°C for further lipid analysis (see later).

Collection of hepatic bile specimens and measurement of circulating bile salt pool sizes. Biliary lipid secretory studies (33) were carried out on additional groups of CCK-1R−/− and WT mice (n = 6 per group) fed the lithogenic diet for 12 weeks. In brief, after caudal ligation, the common bile duct was cannulated with a polyethylene catheter (PE-10). After observation of bile fistula flow, the cystic duct was doubly ligated and a cholecystectomy was performed. Hepatic bile was collected under gravity drainage every hour for 8 hours. Each collection was examined by direct and polarized light microscopy, volumes were determined, and samples were frozen and stored at −20°C for further lipid analyses (see later). For measurement of the circulating bile salt pool sizes, 8-hour biliary washout studies in mice were performed according to published methods (33). During surgery, mouse body temperature was maintained at 37 ± 0.5°C with a heating lamp and monitored with a thermometer. Animals were kept lightly anesthetized with an intraperitoneal injection of 17 mg/kg pentobarbital every 2 hours, and 100 μl of 0.9% NaCl was given hourly via the abdominal cavity to maintain hydration.

Determination of intestinal cholesterol absorption by the plasma dual-isotope ratio method. Cholesterol absorption was measured by a plasma dual-isotope ratio method (27, 30, 31) in chow-fed CCK-1R−/− and WT mice (n = 14 per group). Each animal was injected i.v. with 100 μl of Intralipid containing 2.5 μCi [3H]cholesterol; immediately thereafter we administered an intragastric dose of 150 μl of medium-chain triglyceride in which 1 μCi of [14C]cholesterol was dissolved. To determine the ratio of [14C]cholesterol and [3H]sitostanol — as a nonabsorbable reference marker, dissolved in 100 μl of medium-chain triglyceride — was instilled into the small intestine of mice via a previously fitted in situ externalized duodenal catheter (30). Exactly 30 minutes after instillation, mice were again anesthetized with an intraperitoneal injection of 35 mg/kg pentobarbital. The abdomen was opened, and stomach, small and large intestines, and cecum were removed rapidly while avoiding digital or instrumental compression. The small intestine was frozen promptly in liquid N2, placed on a 50-cm ruled template, and cut into 20 equal segments with a scalpel blade. Individual segments were placed in tubes containing 10 ml of CHCl3-CH3OH (2:1, vol/vol), homogenized, and centrifuged at 10,000 g for 30 minutes. The samples were then stored at 4°C for 48 hours. Well-mixed portions (1 ml) were pipetted into counting vials, and the solvent was evaporated under N2. EcoLite (7 ml) was then added, and radioactivity was determined by liquid scintillation counting. Samples of stomach, cecum, and large intestine were also analyzed, but none showed appreciable radioactivity above background. Using these data, a calculation of small-intestinal transit time was carried out by two arithmetic methods (30): (a) the percentages of total [3H]sitostanol radioactivity in each of the 20 small-intestinal segments were transformed to cumulative percentages passing each segment; (b) the geometric center for the distribution of radioactivity within the small intestine was derived from the sum of the proportions of [3H]sitostanol per segment multiplied by segment number.

Biliary lipid analyses. Biliary phospholipids were analyzed by an inorganic phosphorus method (57). Total and individual bile salt concentrations were measured by HPLC (28). Biliary cholesterol as well as cholesterol content in chow and gallstones was determined by HPLC (28). CSIs of gallbladder and hepatic bile samples were calculated from critical tables (23). Relative lipid compositions of mouse gallbladder and hepatic bile samples were plotted on triangular phase diagrams according to their rounded-off mean total lipid concentrations (24). Phase boundaries and crystallization pathways were extrapolated from model bile systems based on sodium taurocholate at 37°C (24, 25). Hydrophobicity indexes of hepatic bile were calculated according to Heuman’s method (29).

Statistical analyses. All data are expressed as mean ± SD. Differences among groups of mice were assessed for statistical significance by Student’s t test, Mann-Whitney U test, or χ2 test. Analyses were performed with SuperANOVA software (Abacus Concepts Inc., Berkeley, California, USA). Statistically significant differences were defined as a two-tailed probability of less than 0.05.

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