Unconjugated bilirubin in human bile: the nucleating factor in cholesterol cholelithiasis?

M K Dutt, G M Murphy, R P H Thompson

ORIGINAL ARTICLE

C holesterol is sparingly soluble in water and its transport in an aqueous medium such as bile is dependent on its aggregation with other biliary lipids. Consequently, studies of the pathogenesis of cholesterol gallstones have for many years concentrated on the relative stabilities of the different modes of cholesterol transport in bile. More recently, attention has been focused on the process of cholesterol crystal formation and, in particular, glycoproteins have been identified in human bile that may promote or inhibit cholesterol crystal precipitation. However, stones are rarely pure cholesterol; indeed, the “Western” cholesterol gallstone is often only 70% cholesterol by weight and almost invariably contains a cholesterol, bilirubin is also insoluble in water and its transport in human bile is bile acid dependent. Like cholesterol, bilirubin is also insoluble in water and its transport in bile from patients with stones than in bile from control patients, whether or not the bile from either group was saturated with cholesterol or not. Indeed, the mean concentration of cholesterol was significantly higher in control bile samples.

Conclusion: The presence of stones was more closely related to the proportion of unconjugated bilirubin than to the degree of saturation of bile with cholesterol. Bilirubin and its metabolites probably play an important part in the formation of cholesterol gallstones.

Aims: To investigate the concentrations of bilirubin, bilirubin conjugates, phospholipid, and cholesterol in the gall bladder bile obtained at surgery from patients with and without cholesterol gallstones.

Methods: Gall bladder bile was collected during surgery, by puncture, from 20 patients with gallstones undergoing routine cholecystectomy and from eight patients with normal liver blood tests. Concentrations of bilirubin, bilirubin conjugates, phospholipid, and cholesterol were measured using standard procedures.

Results: The proportion of total bilirubin that was unconjugated was significantly higher in the bile from patients with stones than in bile from control patients, whether or not the bile from either group was saturated with cholesterol or not. Indeed, the mean concentration of cholesterol was significantly higher in control bile samples.

Therefore, we compared the concentrations of bilirubin metabolites and biliary lipids in the bile of patients with and without cholesterol gallstones.

PATIENTS AND METHODS

Gall bladder bile was collected during surgery, by puncture, from 20 patients with gallstones undergoing routine cholecystectomy. Oesophagogastroduodenoscopy had been carried out in 14 of the 20 patients and was normal. All preoperative liver blood tests, including serum bilirubin, enzymes, and albumin were normal. Operative cholangiography revealed no bile duct stones.

Gall bladder bile was also obtained from eight patients with normal liver blood tests, six of whom were undergoing surgery for duodenal ulcer, one for adhesions, and one an exploratory laparotomy for unexplained abdominal pain in whom no abnormality was found. In each case the gall bladder was inspected and palpated and, being normal, the specimen of bile was aspirated before the surgical procedure.

The median age of the 20 patients with stones was 47 years (range, 21–73) and 12 were women. The median age in the eight patients without stones was 43 years (range, 21–66) and four were women. No patient was taking drugs. All stones were cholesterol in type (> 70%), as analysed by routine laboratory procedures. The concentrations of bilirubin and its conjugates in bile were determined using the base catalysed transesterification procedure of Blanckaert. Bilirubin standards were prepared in bile acid/lecithin solutions and in chloroform, to cover the concentration range of 0–200 µmol/litre. Recovery was monitored by the addition of 0.1 µCi 14C-bilirubin (specific activity 15.9 mCi/mmol; Amersham–Buchler, Germany) to 0.2 ml bile, and incubating for 15 minutes at 34°C before extraction; the mean recovery from two bile samples on five consecutive occasions was 97% (SD, 3%) and 96% (SD, 2%), respectively. The recovery from thin layer plates of bilirubin standards prepared in chloroform was indistinguishable from that of bilirubin standards prepared in the bile acid/lecithin solutions, and did not vary over the range of 2–30 µmol applications: mean, 69%; SD, 5% at 2 µmol; mean, 71%; SD, 2% at 30 µmol (n = 5).

Biliary bile acids, phospholipids, and cholesterol were measured as described previously, and the cholesterol saturation index (CSI) of each bile sample was calculated as described. The significance of the differences between the results was estimated with the Mann Whitney non-parametric method using “Minitab” software version 2.0.

All patients gave written permission for the bile to be obtained during surgery, and the procedure was approved by the St Thomas’s Hospital ethics committee.

RESULTS

One quarter of the 20 patients with predominantly cholesterol gallstones had bile unsaturated with cholesterol (CSI < 1.0), whereas half of the stone free bile samples were unsaturated; consequently, there was no significant difference with respect
Biliary bilirubin concentrations in patients with (triangle) and without (circle) predominantly cholesterol gallstones. Medians are shown.

**Figure 2** Unconjugated bilirubin [% total] in cholesterol saturated (cholesterol saturation index (CSI) > 1.0) and unsaturated (CSI < 1.0) bile samples from patients with and without gallstones.

Therefore the presence of stones correlated more with the proportion of unconjugated bilirubin than with the CSI. Indeed, in the 15 patients with cholesterol gallstones and bile saturated with cholesterol, the mean percentage of bilirubin present in the unconjugated fraction (mean, 3.3%; median, 2.84%; SD, 1.5%; n = 15) was higher than that in the four control subjects also with cholesterol saturated bile but without gallstones (mean, 1.63%; median, 1.51%; SD, 0.8%; n = 4; p < 0.02), and also higher, but not significantly different, from that in the other four control subjects whose bile was not saturated with cholesterol (mean, 1.85%; median, 1.72%; SD, 0.8%; n = 4; 0.05 > p > 0.10).

In the five patients with cholesterol gallstones but bile unsaturated with cholesterol, the percentage of bilirubin present in the unconjugated fraction (mean, 3.66%; median, 3.25%; SD, 1.4%; n = 5) was significantly higher than that seen in the eight subjects without gallstones, whether their bile was saturated with cholesterol (mean, 1.63%; median, 1.51%; SD, 0.8%; n = 4; p < 0.04) or not (mean, 1.85%; median, 1.72%; SD, 0.8%; n = 4; p < 0.04; fig 2).

**DISCUSSION**

Our results confirm that the CSI and lipids in the gall bladder bile of patients with gallstones are not consistently higher than in controls, although the bile of patients with cholesterol gallstones frequently contains a greatly increased proportion of unconjugated (and also probably of monoconjugated) bilirubin. Such observations are in accordance with the hypothesis that deconjugation of bilirubin glucuronides in bile may be an early event in the formation of cholesterol gallstones with a pigmented nidus, rather than a primary abnormality of lipid metabolism being the cause.

The major bilirubin metabolite in human bile is its diglucuronide, which, together with the monoglucuronide, accounts for more than 98% of total biliary bilirubin. Deconjugation may be catalysed by glucuronidases of bacterial or mucosal origin, and bilirubin monoglucuronide may also undergo non-enzymatic hydrolysis to form unconjugated bilirubin. Even in normal bile, the low concentration of unconjugated bilirubin still exceeds its aqueous solubility 100–1000-fold, so that bile has frequently been described as being supersaturated with calcium salts of bilirubin. Increased proportions of unconjugated bilirubin in gallbladder bile may be a consequence of impaired gallbladder motility. Prolonged storage of bile in the gallbladder, in addition to increasing the length of exposure of conjugated bilirubin to glucuronidases, may also decrease the biliary pH, although this has not been measured in humans. Endogenous β-glucuronidase activity can be detected at pH 7.5, but its optimum pH is 4.5–5.0, so that a lower pH would favour the formation of unconjugated bilirubin. Even in sterile gallstones, bacterial DNA can usually be detected, suggesting

| Table 1 Lipid concentrations in gall bladder bile |   |   |
| No stones (n=8) | With stones (n=20) |
| Cholesterol (mmol/l) | 14.03 (5.1) | 12.05 (4.9)* |
| Bile acids (mmol/l) | 119.7 (38.2) | 91.3 (27.5) |
| Phospholipids (mmol/l) | 54.2 (20.6) | 36.3 (15.5)* |
| Total lipids (g/l) | 114.8 (38) | 80.8 (26)* |
| Cholesterol saturation index | 0.99 (0.28) | 1.30 (0.48) |

Values are means (SD).

*p < 0.02, stones v no stones.
The presence of stones was more closely related to the proportion of unconjugated bilirubin than to the degree of saturation of bile with cholesterol. Bilirubin and its metabolites probably play an important part in the formation of cholesterol gallstones.

The concept that unconjugated bilirubin plays at least as an important role in the formation of cholesterol gallstones as does cholesterol saturation is not new, but has received little attention.

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The concept that unconjugated bilirubin plays at least as an important role in the formation of cholesterol gallstones as does cholesterol saturation is not new, but has received little attention. Our finding of an increased proportion of glucuronidase activity related to poor gall bladder motility, the nucleation of cholesterol, or that it is simply a marker of gated and/or monoconjugated bilirubin may be important in the possibility that the proportions of conjugated to unconjugated bilirubin is a feature of both cholesterol and pigment gallstones.

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