Deoxycholic acid in gall bladder bile does not account for the shortened nucleation time in patients with cholesterol gall stones

H Noshiro, K Chijiwa, I Makino, K Nakano, I Hirota

Abstract

The relations between the concentration of deoxycholic acid (DCA), the cholesterol saturation index, and the nucleation time in gall bladder bile were measured to determine the role of DCA in bile in the pathogenesis of cholesterol gall stone disease. Bile was obtained from patients with cholesterol gall stones (n=30), subjects without gall stones (n=35), and patients with pigment gall stones (n=9). Three of 30 cholesterol gall stone patients and 10 of 35 gall stone free subjects were treated with antibiotics by mouth to decrease the concentration of bile DCA and determine the effect of DCA on biliary lithogenicity. Both the percentage and concentration of DCA in bile were similar in patients with and without cholesterol gall stones despite significant differences in their cholesterol saturation indices and nucleation times. Neither the percentage nor the concentration of DCA in bile correlated with either the cholesterol saturation index or the nucleation time. Analysis of subgroups with matching cholesterol saturation indices showed no correlation between the proportion of DCA in the bile and the cholesterol nucleation time. The proportion of DCA in bile was decreased by antibiotic treatment, but this had no effect on the cholesterol saturation index or nucleation time. These results suggest that DCA in bile is not responsible for biliary cholesterol saturation or cholesterol nucleation time. (Gut 1995; 36: 121–125)

Keywords: gall stones, gall bladder bile, deoxycholic acid, nucleation time.

Deoxycholic acid (DCA), a metabolite of cholic acid, the precursor of DCA, has been considered a risk factor for cholesterol gall stone disease. An increased proportion of DCA in bile has been shown in patients with cholesterol gall stones compared with subjects without gall stones, and a significant correlation has been shown in patients with gall stones between the percentage of DCA in the bile acid pool and the cholesterol saturation index (CSI) of their bile. In normal subjects fed physiological amounts of DCA, the molar percentage of cholesterol in bile rose significantly along with the increased proportion of DCA in the bile. A similar result was also seen with cholic acid, the precursor of DCA. Moreover, when intestinal transit was slowed in healthy volunteers treated with lopamide, their circulating DCA pools expanded and their biliary CSI rose.

A decreased proportion of DCA in bile can be induced by the administration of metronidazole or ampicillin, by feeding bran, lactulose, or a preparation of Streptococcus faecium, or by treating hypothyroidism with thyroxine. Reduction of biliary CSI was reported to follow these manoeuvres.

Recent evidence has shown that nucleation time, the time to the appearance of cholesterol monohydrate crystal in vitro, seen under a polarising microscope, better predicts lithogenic bile than the CSI, as supersaturated bile is often seen in healthy subjects without gall stones. Mucous glycoprotein has been shown to promote the nucleation of cholesterol monohydrate crystals in bile, and hypersecretion of mucus glycoprotein into gall bladder can be induced through the arachidonic acid pathway or directly by DCA. In artificial bile, nucleation time is shortened when the predominant bile acid is the taurine conjugate of DCA, compared with the taurine conjugate of chenodeoxycholic acid or cholic acid predominates.

Based on this evidence, we hypothesised a close correlation between the concentration of DCA and the nucleation time in gall bladder bile. In this study, we determined the relation between the proportion of DCA and the nucleation time using human gall bladder bile samples. As a reduced biliary CSI was reported in bile samples with lowered DCA, we also examined the effect of decreasing the proportion of DCA by antibiotic treatment on the CSI and nucleation time.

Methods

Patients

Gall bladder bile samples were collected from 30 patients with cholesterol gall stones, nine patients with pigment gall stones, and 35 subjects without gall stones. Of these 74 subjects, 10 without gall stones and three patients with cholesterol gall stones were treated with antibiotics, kanamycin at a dose of 3 g/day and metronidazole 750 mg/day, for three days before intestinal surgery for colonic cancer, a treatment generally used for bowel preparation. There was no significant difference in age or sex among the five patient groups:
untreated cholesterol gall stone group (UCGS group), antibiotic treated cholesterol gall stone group (ACGS group), untreated gall stone free group (UGSF group), antibiotic treated gall stone free group (AGSF group), and pigment gall stone group (PGS group) (Table I). The patients in the ACGS group had colonic cancer in addition to their cholesterol gall stone disease. Samples from UGSF subjects were obtained during surgery for gastric cancer (n=20), malignant lymphoma of the stomach (n=1), colonic cancer (n=2), liver cyst (n=4), or adenoma of the papilla of Vater (n=1). No patient had biliary tract disorder or gastrointestinal obstruction. None of the patients were obese or suffered from hyperlipidaemia or liver dysfunction. None of the patients had previous bile acid treatment. Informed consent was obtained from all patients.

### BILE COLLECTION

A functional gall bladder was confirmed by preoperative cholangiography or ultrasonography, and bile samples with total lipid concentrations of less than 5 g/dl were excluded from the study after biliary lipid analysis.24 Bile samples were obtained by needle aspiration, withdrawing biliary gall bladder contents completely to avoid sampling errors caused by stratification.24 The fresh, sterile, samples were centrifuged immediately at 100 000×g for 120 minutes at 37°C in an ultracentrifuge (Hitachi 55P-62, Tokyo, Japan) and the isotropic, cholesterol crystal free, bile was collected from the interphase and subjected to nucleation time16 and biliary lipid analysis as previously described.25 Absence of gall stones was verified by preoperative ultrasound and by intraoperative palpation of the gall bladder. Any gall stones present were classified by gross inspection and chemical analysis in the laboratory.26 All cholesterol gall stones included more than 70% cholesterol by dry weight and pigment gall stones contained less than 30% cholesterol. All fresh bile samples used for the study were negative for bacterial contamination by standard aerobic and anaerobic bacterial culture. Histological examination of all gall bladder specimens obtained from the gall stone patients showed only mild cholecystitis.

### NUCLEATION TIME

Nucleation time was determined in our laboratory as previously reported,27 based on the method of Holan et al. Isotopic, cholesterol crystal free, bile was transferred to small sterile brown glass vials, and was stored in darkness in an incubator at 37°C without shaking. Small aliquots were examined daily under a polarising microscope (Nikon, XTP-II), to detect the formation of cholesterol monohydrate crystals, which were identified by their typical rhomboidal structure. Nucleation time was defined as the number of days until typical rhomboidal monohydrate crystals appeared during a 21 day observation period.

### BILARY LIPID ANALYSIS

Phospholipids were measured by the method of Bartlett.28 Cholesterol and individual bile acids were quantified simultaneously by gas-liquid chromatography using nordeoxycholic acid as an internal standard as previously described.29 Total bile acid concentration was calculated as the sum of the bile acids. The cholesterol saturation index (CSI) was calculated using a microcomputer program30 based on tables provided by Carey.

### STATISTICS

Values are expressed as means (SD). The statistical significance of differences was evaluated using the χ² test, analysis of variance (ANOVA) or the unpaired Student’s t test. Correlation between parameters was tested by linear regression analysis. p Values less than 0·05 were considered significant.

### Results

#### BILARY LIPID COMPOSITION AND NUCLEATION TIME

No significant difference was seen in the molar percentages of bile acids or phospholipids in gall bladder bile between the UCGS and the UGSF groups (Tables I and II). The proportion of cholesterol in the bile was significantly higher in the UCGS than in the UGSF group (Table I). The CSI was also significantly higher in the UCGS than the UGSF group (1·25 (0·29) vs 0·96 (0·26); p<0·01, Table I). There was no significant difference in CSI between the PGS and UGSF groups (0·87 (0·18) vs 0·96 (0·26), Table I). The nucleation time of the UCGS group was much shorter than those of the UGSF and PGS groups (2·9 (2·6), 12·0 (7·5), and 15·4 (6·2) days; p<0·01,
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TABLE II Bile acid composition in gall bladder bile

<table>
<thead>
<tr>
<th>Patient group</th>
<th>LCA</th>
<th>DCA</th>
<th>CDCA</th>
<th>UDCA</th>
<th>CA</th>
</tr>
</thead>
<tbody>
<tr>
<td>UGSF (n=27)</td>
<td>0.6 (0.7)</td>
<td>17.2 (11.7)</td>
<td>43.8 (10.6)</td>
<td>2.3 (2.0)</td>
<td>36.2 (7.5)</td>
</tr>
<tr>
<td>ACGS (n=3)</td>
<td>0.0 (0.1)</td>
<td>6.7 (8.3)</td>
<td>51.5 (14.5)</td>
<td>0.0 (0.6)</td>
<td>41.7 (6.5)</td>
</tr>
<tr>
<td>(0.1)</td>
<td>(9.5)</td>
<td>(7.7)</td>
<td>(75.2)</td>
<td>(0.0)</td>
<td>(31.1)</td>
</tr>
<tr>
<td>UGSF (n=25)</td>
<td>0.8 (0.8)</td>
<td>16.1 (8.6)</td>
<td>44.3 (7.6)</td>
<td>3.5 (3.2)</td>
<td>34.8 (10.4)</td>
</tr>
<tr>
<td>AGSF (n=10)</td>
<td>0.3 (0.7)</td>
<td>9.7 (7.0)</td>
<td>46.8 (11.2)</td>
<td>2.8 (2.9)</td>
<td>40.5 (11.1)</td>
</tr>
<tr>
<td>(0.5)</td>
<td>(10.1)</td>
<td>(13.2)</td>
<td>(27.9)</td>
<td>(3.9)</td>
<td>(22.1)</td>
</tr>
<tr>
<td>PGS (n=9)</td>
<td>1.2 (1.1)</td>
<td>14.1 (9.5)</td>
<td>48.0 (10.1)</td>
<td>2.8 (3.7)</td>
<td>33.9 (9.2)</td>
</tr>
<tr>
<td>(2.0)</td>
<td>(8.8)</td>
<td>(25.5)</td>
<td>(22.2)</td>
<td>(4.0)</td>
<td>(25.4)</td>
</tr>
</tbody>
</table>

Values are expressed as mean (SD) of: upper, the percentage of total bile acid; and, below, the concentrations (mM) of each bile acid in gall bladder bile. Significantly different from UGSF group; †p<0.05 and ‡p<0.01. Abbreviations used: LCA, lithocholic acid; DCA, deoxycholic acid; CDCA, chenodeoxycholic acid; UDCA, ursodeoxycholic acid; CA, cholic acid.

Other abbreviations as in Table I.

respectively, Table I). Despite the significant differences seen in CSI and nucleation time between patients with and without cholesterol gall stones, there was no significant difference in the percentage of DCA in the total bile acid or in the absolute concentration of DCA in gall bladder bile among the UCGS, UGSF, and PGS groups (17 (12), 16 (9), and 12 (10) for percentages and 23.9 (18.6), 31.6 (23.7), and 25.5 (22.2) mM for concentrations, respectively; Table II).

EFFECT OF ANTIBIOTIC TREATMENT ON THE DCA CONCENTRATION, CSI, AND NUCLEATION TIME IN SUBJECTS WITHOUT GALL STONES

In the 10 GSF patients treated with antibiotics, the proportion of DCA was significantly decreased compared with the untreated GSF group (10 (7)% v 16 (9); p<0.05, Table II). The molar per cent of cholesterol, and the CSI and nucleation time in the bile were not changed, however, by antibiotic treatment. The total biliary lipid concentration was significantly lower in the antibiotic treated GSF group than in the untreated GSF group (10 (9) (3.1) v 14.3 (4.6) g/dl; p<0.05, Table I). In the antibiotic treated CGS group, the proportion of DCA in bile was decreased and the CSI and nucleation time were not changed compared with the untreated CGS group, suggesting a similar response as that of the gall stone free subjects, although the sample number was insufficient for statistical comparison between the groups (Tables I and II).

RELATIONS BETWEEN DCA CONCENTRATION, TOTAL LIPID CONCENTRATION, CSI, AND NUCLEATION TIME

Nucleation time was significantly inversely correlated with the CSI (r=-0.621, p<0.01), but not with the total lipid concentration (r=0.077). No correlation was seen between the percentage of DCA and the molar percentage of cholesterol, phospholipids or total bile acid in bile (r=0.042, r=-0.076, and r=0.045, respectively). The percentage of DCA did not affect the CSI (r=0.112, Figure (A)). Similarly, neither the percentage nor the concentration of DCA in bile correlated with the nucleation time (r=-0.029 and r=0.075, Figure (B)). The total lipid concentration was not correlated with the percentage of DCA (r=0.005).

As no correlation was found between the proportion of DCA and the CSI, the bile samples were divided into four subgroups according to the CSI (CSI<0.75, 0.75<CSI<1.00, 1.00<CSI<1.25, and 1.25<CSI) to find out if the concentration of DCA was indeed independent of nucleation

Figure 1: Relations between the proportion of DCA and (A) CSI and (B) nucleation time in gall bladder bile.

There were no significant correlations. ○: untreated cholesterol gall stone patients; ♦: antibiotic treated cholesterol gall stone patients, ▲: antibiotic treated gall stone free patients, ◻: untreated gall stone free patients, △: pigment gall stone patients. NS: not significant.
time in bile samples with similar CSIs (Table III). The DCA concentration in bile was not increased in bile samples with rapid nucleation time (nucleation time ≤ 5 days) in any CSI subgroup.

**Discussion**

It has been suggested that an increased amount of biliary DCA may be a risk factor in cholelithiasis. Proposed mechanisms for the influence of DCA on the pathogenesis of cholelithiasis have included raising the cholesterol saturation of bile, and causing hypersecretion of mucous glycoprotein.  

The results in this paper show that neither the proportion nor the absolute concentration of DCA correlated with either the CSI or the nucleation time in human gall bladder bile. An acceleration by DCA of the cholesterol crystal nucleation time without an effect on the CSI was also not seen (Table III). It has been reported that DCA raises the cholesterol saturation of bile by increasing secretion of cholesterol. In this study, however, there was no correlation between the percentage or concentration of DCA and those of cholesterol in the bile. Moreover, both the percentage and concentration of DCA were very similar in untreated patients with cholesterol gall stones and in those without gall stones, while the CSI differed significantly between the two groups. These results support previous findings, and confirmed that the raised biliary cholesterol saturation indices in patients with cholesterol gall stones are independent of DCA in bile.

Our finding of no significant difference in biliary DCA between patients with and without gall stones differs from those of some previous studies. There are some possible reasons for the differing results. Firstly, to avoid confounding effects of cholecystitis, only patients with functioning gall bladders, as evidenced by bile with total lipid concentrations greater than 5 g/dl, were selected. Some bile samples in previous studies showed extremely low total lipid concentrations. In this study, no significant correlation was seen between the proportion of DCA and total lipid concentration of gall bladder bile (r = 0.005). Secondly, control bile samples were obtained from patients with few symptoms whose age and sex were similar to those of the gall stone group. Twenty of 27 untreated gall stone free patients in this study suffered from gastric cancer with few symptoms and normal food intake, while most control patients in previous studies had peptic ulcer or hiatial hernia with abdominal complaints. In contrast, studies using duodenal bile samples obtained from healthy volunteers and gall stone patients showed no significant difference in the proportion of DCA between the groups. Finally, this is a report of the relation of DCA and the pathogenesis of cholesterol gall stone in Japanese subjects. Racial differences seem to be unimportant, however, because both the CSI and nucleation time of gall bladder bile, which are considered to accurately reflect the tendency to cholesterol crystallisation, were similar in our subjects to those reported for Europeans.

To find out if DCA had an effect on nucleation time independent of the CSI, bile samples were divided into four subgroups according to their cholesterol saturation indices. This was based on previous findings that DCA accelerates hepatic secretion of arachidonic acid into bile. Arachidonic acid is a precursor of prostaglandins, which in turn mediate mucous glycoprotein secretion from the gall bladder epithelium and DCA directly stimulates mucous glycoprotein secretion from gall bladder epithelium. Mucous glycoprotein is a recognised nucleating factor. Bile samples belonging to the subgroups with CSI > 0.75 showed a wide range of nucleation times, suggesting the presence of nucleating or antinucleating factors in bile. DCA concentration showed no relation, however, to cholesterol crystallisation time in bile samples grouped by CSI (Table III).

Several mechanisms have been reported to reduce DCA in bile. Antibiotic administration is one of these, and metronidazole and ampicillin both reduce the bacterial flora in the colon and interfere with the dehydroxylation of cholic acid, decreasing DCA in bile. Although the proportion of DCA was significantly decreased after treatment with metronidazole and kanamycin in gall stone free patients (Tables I and II), neither a decrease of cholesterol saturation index nor a prolongation of nucleation time in gall bladder bile was seen, supporting previous reports. These results show that decreasing the concentration of DCA in bile by short term antibiotic treatment had no effect on cholesterol saturation or nucleation time in the bile.

In conclusion, the proportion or concentration of deoxycholic acid in bile did not correlate with either cholesterol saturation or nucleation time of gall bladder bile in patients.
Deoxycholic acid in gall bladder bile does not account for the shortened nucleation time in patients with cholelithiasis.