Adenosine Triphosphate–dependent Taurocholate Transport in Human Liver Plasma Membranes

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Abstract

Transport systems involved in uptake and biliary secretion of bile salts have been extensively studied in rat liver; however, little is known about these systems in the human liver. In this study, we investigated taurocholate (TC) transport in canalicular and basolateral plasma membrane vesicles isolated from 15 human livers (donor age 6–64 yr). ATP stimulated the uptake of TC into both canalicular and basolateral human liver plasma membrane vesicles (clLPM and blLPM, respectively). Considerable interindividual variations in the transport velocity were observed in the different membrane preparations used: 9.0±1.3 (mean±SEM, n=17; range 1.6–18.0) and 9.3±2.0 (range 1.1–29.8) pmol TC·mg protein−1·min−1 at 1.0 μM TC for clLPM and blLPM, respectively. TC transport was temperature sensitive and showed saturation kinetics with a high affinity for TC (Km 4.2±0.7 μM and 3.7±0.5 μM for clLPM and blLPM, respectively). Transport was dependent on the ATP concentration and saturable (Km 0.25±0.03 mM, n = 3). Neither nitrate, which reduces membrane potential, nor the protonophore FCCP strongly inhibited ATP-dependent TC transport, indicating that membrane potential and proton gradient are not involved in this process. TC transport was significantly inhibited by the classical anion transport inhibitor 4,4′-disothiocyanostilbene-2,2′-disulfonate (250 μM) and the glutathione conjugate S-(2,4-dinitrophenyl)glutathione (100 μM). In conclusion, high affinity ATP-dependent TC transport is present in human liver at both the canalicular and the basolateral sides of the hepatocyte. (J. Clin. Invest. 1992. 90:2321–2326.) Key words: canalicular plasma membrane vesicles • basolateral plasma membrane vesicles • primary active transport • DNP-SG

Introduction

Transport systems involved in vectorial hepatic transport of bile salts (i.e., uptake at the sinusoidal side and secretion at the canalicular pole of the hepatocyte) have been studied extensively in rat liver. However, little is known about these processes in the human liver. Vesicles are the system of choice to study these processes, since no interference by intracellular metabolism occurs and the media inside and outside the vesicles can be defined exactly and manipulated easily. We have recently developed a method to isolate canalicular (clLPM) and basolateral (blLPM) plasma membrane fractions simultaneously from human liver (1), which enables us to study transport processes in both types of membranes of the same liver.

Studies using membrane vesicles from rat liver indicate that uptake of taurocholate (TC) from the blood into the hepatocyte at the basolateral (sinusoidal) membrane is sodium dependent (2–6). Only a single study is known to us in which TC transport was investigated in human liver membranes: in basolateral membrane preparations isolated from two human livers, sodium-dependent TC transport could also be demonstrated (7).

The excretion of TC into bile has been shown to be sodium independent and membrane potential dependent in rat liver (5, 8, 9). A protein of ~ 100 kD appears to be involved in this transport process (10–12). The magnitude of the membrane potential, however, is too small to explain the actual bile salt concentration gradients that exist over the canalicular membrane (5). Recently, primary ATP-dependent TC transport was demonstrated in rat canalicular membranes (13–17), which apparently acts synergistically with potential-dependent TC transport (18). Photoaffinity labeling and reconstitution studies revealed that a 110-kD glycoprotein functions as carrier in the ATP-dependent TC transport (16).

Besides the bile salt transport system, the existence of two other ATP-dependent transporters has been demonstrated in rat canalicular membranes, namely, a transporter for non–bile acid organic anions, e.g., S-(2,4-dinitrophenyl)glutathione (DNP-SG), cysteinyl leukotrienes and glucuronides (19–25), and probably bile acid sulfates (25), and the P-glycoprotein, the product of the multidrug-resistance gene, which transports mainly hydrophobic cations (26). Available data indicate that the multidrug-resistance transporter is different from the anion carriers. There is also evidence from in vitro studies that TC transport and non–bile acid organic anion transport proceed via different systems (15, 16). This has previously been inferred from the existence of the Dubin-Johnson syndrome in man (27), and mutant Corridale sheep (28) and Wistar rats (29, 30), all showing impaired secretion of non–bile salt organic anions with normal bile salt secretion.

Since bile salt secretion is an important determinant of bile flow, information on mechanisms of hepatic bile salt transport in human liver may provide better insight into normal bile

1. Abbreviations used in this paper: blLPM, basolateral domains of liver plasma membranes; clLPM, canalicular domains of liver plasma membranes; DIDS, 4,4′-disothiocyanostilbene-2,2′-disulfonic acid; DNP-SG, S-(2,4-dinitrophenyl)glutathione; FCCP, carbonylcyanide-p-trifluoromethoxyphenylhydrazone; LAP, leucine aminopeptidase; RE, relative enrichment of enzyme activities; TC, taurocholate; UW, University of Wisconsin.
formation in man, and improve our understanding of mechanisms involved in the etiology of cholestasis. We therefore investigated TC transport in human liver plasma membranes. In this work we demonstrate the presence of an ATP-dependent TC transport system in human liver plasma membranes.

**Methods**

Materials. [3H]TC was purchased from DuPont/New England Nuclear (Boston, MA), FCCP (carbonyl cyanide-p-trifluoromethoxyphenyl-hydrazone) was from Fluka AG (Buchs, Switzerland), and valinomycin and DIDS (4,4'-diisothiocyanostilbene-2,2'-disulfonic acid) were purchased from Sigma Chemical Co. (St. Louis, MO). DNP-SG was kindly provided by Dr. P. Zimmniak (Division of Gastroenterology, University of Arkansas for Medical Sciences, Little Rock, AR). All other chemicals used were of analytical grade.

Human livers. Human liver tissue was obtained from livers harvested from multiorgan donors (six female and nine male, age 6–64 yr). Consent from legal authorities and family was obtained for the explantation of organs for transplantation purposes. The donor livers were split in order to be able to perform reduced-size liver transplants in children. The liver tissue remaining after the splitting procedure was used for the isolation of plasma membranes. Some livers discarded for transplantation after harvesting were used as well. Livers were perfused with University of Wisconsin (UW) solution (31) and stored in this buffer at 4°C until the start of the isolation procedure, which was performed within 48 h.

Isolation and characterization of plasma membrane fractions. cLPM and bLPM were isolated and characterized as described previously (1), with a small modification: 0.1 mM PMSF, a protease inhibitor, was added to the homogenization buffer. Membrane vesicles were stored in liquid nitrogen from 4 to 21 mo until use. No indication of loss of transport activity was observed during storage under these conditions. Marker enzymes, Na+ K+-ATPase (32) for basolateral and leucine aminopeptidase (33) for canalicular membranes, were determined as described. Protein was determined by the method of Lowry et al. (34) using BSA (Sigma Chemical Co.) as standard. Enrichments (specific enzyme activity of the membrane preparation relative to the specific enzyme activity in the total homogenate) of the canalicular membrane preparations were 59±8 for leucine aminopeptidase and 21±5 for Na+ K+-ATPase, and for basolateral preparations 14±2 and 38±6, respectively (17 preparations). We have previously shown that the subcellular organelles endoplasmic reticulum, mitochondria, Golgi membranes, and lysosomes, are only slightly enriched, if at all, in our membrane preparations (1).

Transport studies. TC transport was measured using a rapid filtration technique. Experiments were performed at 37°C in a final volume of 100 μl. Unless specified otherwise, the incubation medium contained various concentrations of [3H]TC, 1.0 mM ATP, an ATP regenerating system (10 mM creatine phosphate and 0.1 mg/ml creatine kinase), 1.0 mM ouabain, and 0.5 mg/ml BSA in 10 mM Tris·HCl/0.25 M sucrose/10 mM MgSO4/0.2 mM CaCl2, pH 7.4; final membrane protein concentration was 0.2 mg protein·ml⁻¹. Similar incubations without ATP served as blanks. Transport was stopped by adding 2.5 ml ice-cold buffer (10 mM Tris·HCl/0.25 M sucrose/10 mM MgSO4/0.2 mM CaCl2, pH 7.4). Vesicles were filtered through 0.45-μm mixed (nitrate/acetate) cellulose filters (Millipore HAWP; Millipore Corp., Bedford MA), which were prewetted with 1 ml 1.0 mM TC and washed twice with 2.5 ml buffer. Radioactivity on the filter was measured in a Wallac liquid scintillation counter (LKB, Turku, Finland). Experiments were performed in triplicate, with at least three different membrane preparations. K_m and V_max values were calculated using double reciprocal plots of 1/v as a function of 1/[substrate]; the intercept on the x-axis reflects –1/K_m, the intercept on the y-axis 1/V_max.

Values are expressed as mean±SEM unless indicated otherwise. In some studies least-squares linear regression analysis was applied.

**Results**

**ATP-dependent TC transport.** TC uptake into human liver plasma membrane vesicles at 37°C was stimulated by ATP (Fig. 1). Large differences in transport rate between different membrane preparations were observed (see below), but qualitatively the different membrane preparations showed the same time dependence: initial, rapid uptake was followed by slower uptake, probably a consequence of simultaneous leakage of TC out of the vesicles according to the TC gradient produced. Finally, probably because the ATP concentration in the system was reduced, release of TC occurred. This release was slower in the bLPM fraction shown in Fig. 1 than in the cLPM fraction, probably because in the latter fraction more Mg-ATPase was present (1), leading to faster consumption of ATP. No ATP-dependent transport of TC was observed when experiments were performed on ice (data not shown). To verify that uptake and not ATP-dependent binding was measured, studies were performed in the presence of different extravesicular concentrations of sucrose. With increasing osmolarity, less TC was taken up in the presence of ATP (Fig. 2), indicating that only a small fraction of the measured uptake reflects binding to the membranes. Binding of TC to the membrane vesicles (association at infinite medium osmolarity) amounted to 8 and 12% of the total uptake in normal medium in two different cLPM preparations, and to 8 and 9%, respectively, for bLPM fractions isolated from the same livers. Thus, the contribution of binding was similar for both types of membranes.

**Localization of ATP-dependent TC transport.** ATP-dependent TC transport was observed in both canalicular and basolateral membrane fractions. Large interindividual differences were observed: initial rates 1.6–18.0 pmol TC·mg⁻¹·min⁻¹ for cLPM and 1.1–29.8 for bLPM (medium concentration, 1.0 μM TC). In some of the preparations the transport rate (on protein basis) in basolateral membranes was higher than in the canalicular fraction, whereas in others this ratio was opposite (see below). Mean TC transport rate was 9.0±1.3 pmol TC·mg⁻¹·min⁻¹.

![Figure 1. TC uptake into human cLPM and bLPM: effect of ATP. Association of TC (medium concentration 1.0 μM) with membrane vesicles was determined at 37°C, in the absence (closed symbols) or presence (open symbols) of 1.0 mM ATP, cLPM, bLPM. Data are from a representative experiment (see text). The experiment was performed in triplicate; SD are not shown for clarity, they were <15% of the mean values.](http://www.jci.org)
Figure 2. Effect of medium osmolality on TC uptake in human liver plasma membrane vesicles. ATP-dependent uptake was determined after 30 min incubation in medium (TC concentration 1.0 μM) with different concentrations of sucrose. Data (±SD) are from a typical experiment in triplicate. Similar observations were made in four other membrane preparations. By extrapolation to the abscissa an estimate of the amount of TC bound to the membranes can be made.

pmol·mg⁻¹·min⁻¹ for cLPM and 9.3±2.0 pmol·mg⁻¹·min⁻¹ in bLPM (17 membrane preparations isolated from 15 different livers). It is theoretically possible that ATP-dependent TC transport is present only in human canalicular membranes, as described for rat liver (14, 15). Due to unavoidable contamination with canalicular membranes, ATP-dependent TC transport would in this case also be measurable in the bLPM fraction. This possibility could, however, be excluded. The relative amounts of cLPM present in the cLPM and bLPM fractions can be quantified by measuring the relative enrichment (RE) in leucine aminopeptidase (LAP) activity. If ATP-dependent TC transport were performed by canalicular membranes only, the transport rates corrected for the presence of cLPM (by dividing by LAP enrichment) should be identical in cLPM and bLPM. Transport rates thus calculated, however, yielded values of 0.19±0.04 pmol·min⁻¹·mg prot⁻¹·RE(LAP)⁻¹ for cLPM and 0.69±0.10 for bLPM preparations (P < 0.001), indicating that the ATP-dependent TC transport is not only located on the canalicular, but also on the basolateral side of the membrane. Similarly, to rule out the possibility that transport is located on bLPM only, transport rates were corrected for the presence of bLPM by dividing by the RE for Na⁺/K⁺-ATPase. Mean velocities were 0.56±0.08 for canalicular and 0.27±0.05 pmol·min⁻¹·mg prot⁻¹·RE(Na⁺/K⁺-ATPase)⁻¹ for basolateral membrane preparations (P < 0.01). These data thus show that the ATP-dependent TC transport activity is present in both bLPM and cLPM.

Age dependency of ATP-dependent TC transport. There seemed to be a tendency for a decrease in transport rate with donor age (Fig. 3). However, it appeared that the purity of the membrane preparations was also age dependent to some extent: with increasing age, less enrichment with Na⁺/K⁺-ATPase was observed in both cLPM and bLPM. Enrichments in LAP activity were not age dependent. Thus, it was essential to correct the uptake rate for the purity of the plasma membrane preparation. When this was done for the quantity of bLPM by dividing the transport rate by the enrichment of Na⁺/K⁺-ATPase activity in the membranes (Fig. 4), there was a tendency for a decrease in uptake rate with age for the basolateral preparation only. Transport rate corrected for LAP showed a decrease with age for both cLPM and bLPM, probably because of the occurrence of relatively less basolateral membranes in preparations of the older donors, as mentioned earlier.

No correlations between ATP-dependent TC transport and donor gender, preservation time of the liver in UW solution, or storage time of the membranes in liquid nitrogen were observed.

Kinetics of ATP-dependent TC transport. Transport of TC, measured in the presence of 1.0 mM ATP, showed saturation kinetics. Fig. 5 gives an example of the relationship between TC concentration and transport rate. The apparent Kₐ for TC was 4.2±0.7 μM for cLPM (four preparations), which is similar to the Kₐ determined for bLPM (3.7±0.5 μM; five preparations). As stated above, large differences in transport rates were observed for different preparations used. Vₘₐₓ values ranged from 6.5 to 188 pmol·mg⁻¹·prot⁻¹·min⁻¹ (five basolateral and four canalicular preparations isolated from six different livers). Fig. 6 shows the dependence of transport rate on ATP concentration in a typical experiment. Transport was saturable; the apparent Kₐ for ATP was 0.25±0.03 mM (n = 3). Vₘₐₓ for the three preparations (isolated from three different livers) studied ranged from 4.3 to 21 pmol·mg⁻¹·min⁻¹.

Role of membrane potential and pH gradients. To further characterize ATP-dependent TC transport, we studied the effects of several compounds. Since bLPM and cLPM showed

Figure 3. Age dependency of ATP-dependent TC transport in human liver membranes. Initial rapid TC uptake in medium containing 1.0 μM TC was determined as described in Methods. (A) cLPM (r = -0.567; P < 0.01); (B) bLPM (r = -0.479; P < 0.05).

Figure 4. Age dependency of ATP-dependent TC transport in human liver membranes. Transport rates, as presented in Fig. 3, were corrected for the amount of basolateral membranes actually present in the preparations by dividing the transport rate (pmol·mg⁻¹·min⁻¹) by the relative enrichment of the basolateral marker enzyme Na⁺/K⁺-ATPase. (A) cLPM (r = 0.069); (B) bLPM (r = -0.409). No significant correlation with age was observed.

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similar $K_m$ values for TC uptake, it is highly likely that the same transporter is responsible for transport in both vesicle preparations. We therefore did not study effects of the various compounds on cLPM and bLPM separately, but all studies were done at least once with a preparation of each membrane type. No qualitatively different effects on cLPM and bLPM were observed.

Initial rates of ATP-dependent TC transport were not dependent on pH gradient or membrane potential. The protonophore FCCP, which dissipates pH gradients, only slightly inhibited transport. The presence of nitrate, a permeable anion that dissipates a possible membrane potential, stimulated transport (Table 1). The K$^+$ ionophore valinomycin, which collapses membrane potential in the presence of KCl, inhibited TC transport, which might point to an involvement of membrane potential in the ATP-dependent transport. However, similar inhibition by valinomycin occurred in the absence of potassium, suggesting a nonspecific effect of valinomycin (Table 1).

### Table 1. Effects of Nitrate, Ionophores, DIDS, and DNP-SG on Plasma Membrane ATP-dependent TC Transport

<table>
<thead>
<tr>
<th>Relative transport %</th>
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<tbody>
<tr>
<td>Control</td>
<td>100</td>
</tr>
<tr>
<td>FCCP (5 μM)</td>
<td>81±9 (6)</td>
</tr>
<tr>
<td>Nitrate*</td>
<td>158±24 (4)</td>
</tr>
<tr>
<td>Valinomycin plus K$^{++}$</td>
<td>48±10 (4)</td>
</tr>
<tr>
<td>Valinomycin§</td>
<td>53±21 (3)</td>
</tr>
<tr>
<td>DIDS (250 μM)</td>
<td>6±5 (3)</td>
</tr>
<tr>
<td>DNP-SG (100 μM)</td>
<td>62±10 (4)</td>
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</tbody>
</table>

Initial rates of ATP-dependent TC uptake were measured in medium as described in Methods unless specified otherwise. TC concentration was 1.0 μM and ATP concentration was 1.0 mM. Data are given as percentage of control uptake rate ±SEM, with the number of different membrane preparations used in parentheses. The data presented were obtained from at least one cLPM and one bLPM preparation. No qualitative differences were observed between cLPM and bLPM.

* Medium inside as well as outside the vesicles contained 100 mM KNO$_3$ plus 50 mM sucrose instead of 250 mM sucrose in the uptake buffer. § Medium inside and outside the vesicles contained 100 mM KCl plus 50 mM sucrose instead of 250 mM sucrose in the uptake buffer; valinomycin concentration was 10 μg/mg protein.

### Discussion

Excretion of TC into bile was shown to be a sodium-independent, membrane potential–dependent process in rat liver (5, 8, 9). However, the magnitude of the membrane potential is insufficient to explain the gradient of bile salts across the canalicular membrane. The recent discovery of an ATP-dependent TC excretion system in rat liver may provide a clue for the existence of the extremely high biliary bile salt concentrations (13–17). This study clearly demonstrates the presence of an ATP-dependent transport system for TC in human liver plasma membranes as well. Transport was temperature dependent and showed a low $K_m$ value ($K_m = 4.2±0.7$ μM and 3.7±0.5 μM for cLPM and bLPM, respectively) for TC. In rat cLPM, $K_m$ values of 47 (14), 26 (15), 7.5 (16), and 2.1 μM (17) have been published for the ATP-dependent TC transport system. The $K_m$ for ATP was 0.25 mM, which is in the range of values reported in the literature for rat cLPM, i.e., 0.80 (16), 0.67 (15), and 0.064 mM (14), respectively. Experiments using the highly permeable anion nitrate and the protonophore FCCP indicated that neither membrane potential nor pH gradient was involved in the transport process. Similarly, it has been
reported that membrane potential is not involved in the ATP-dependent TC transport in rat canalicular membranes (14, 17). In our experiments, the ionophore valinomycin did inhibit transport. Since this inhibition was seen in both the absence and the presence of the transferred ion (potassium), this is probably attributable to a nonspecific effect of valinomycin. Similar to observations in rat liver cLPM (15, 17), the anion transport inhibitor DIDS very strongly inhibited ATP-dependent TC transport in human membrane vesicles.

The measured transport rates showed large interindividual variations. There are many factors that may influence the transport rate and may thus be responsible for this variation. First of all there are donor-dependent factors (genetic variations, age, gender, diet, previous medication, etc.). Furthermore, several factors may influence the quality of the isolated liver (quality of the perfusion, preservation time). Finally, the purity of the membranes and their orientation (in rat cLPM, ATP-dependent transport was shown in inside-out vesicles only [15]) may influence the measured transport rate. When corrected for the amount of basolateral membranes present in the membrane fractions, there was an insignificant tendency for an age-dependent decrease in TC transport in the bLPM fraction. It is known that in rat liver ouabain and TC uptake decline upon aging (35, 36). It is possible that a significant decrease of TC transport rate only takes place at higher ages than those studied here (maximum age 64 yr). For other parameters that might influence TC transport (donor gender, preservation time of the liver in UW, and storage time of the membranes in liquid nitrogen), we did not observe clear-cut effects.

ATP-dependent TC transport activity was present in both canalicular and basolateral human plasma membrane preparations, whereas in Sprague-Dawley rats it was demonstrated exclusively (14, 15), and in Wistar rats predominantly (16, and our own unpublished results) in canalicular membranes. In the latter rat membrane preparations only low rates of ATP-dependent TC transport were measured in bLPM fractions, possibly due to slight contamination of these fractions with cLPM. As was demonstrated in Results, in the human membranes transport activity in bLPM cannot be explained by the low level of contamination with cLPM. At the moment we do not know whether the presence of the transporter at the human basolateral membrane is physiologically relevant. Several explanations for the appearance of this transporter can be put forward. First, Bartles et al. (37) have shown in rat liver that, after synthesis, several canalicular proteins are first transported to the basolateral side of the cell and subsequently transferred to the canalicular membrane. It is conceivable, therefore, that the transporter is always present in the bLPM, although perhaps in a physiologically nonrelevant concentration. During the preservation of the liver in UW for a prolonged period of time (at least 10 h in our experiments) redistribution of canalicular and basolateral proteins might take place. Alternatively, the transport of the proteins to the bLPM may be less restrained than the redistribution of the proteins from the bLPM to the cLPM, leading to relatively high concentrations of the transporter at the bLPM. In rats, extrahepatic obstructive cholestasis causes a shift of the bile salt transport system from the canalicular to the basolateral side of the cell (38). The donor livers we studied might, to some extent, be in a “cholestatic state,” for instance due to medication of the donor or other factors inherent to the donor procedure. However, this possibility is not very likely, since no clinical signs of cholestasis were observed: plasma bilirubin and alkaline phosphatase were normal. Since we do not know whether TC is the physiological substrate for the ATP-dependent transport observed, it may be of physiological significance to transfer its natural substrate out of the cell and into the blood compartment. For several organic anions, transport from the liver to the blood (sinusoidal efflux) is known to occur (39).

There is some controversy in the literature as to whether in rat liver TC and non-bile acid organic anions share the same canalicular ATP-dependent transporter. In one study (13), ATP-dependent DNP-SG and TC transport showed mutual inhibition. Nishida et al. (15), however, did not observe inhibition of ATP-dependent TC transport by DNP-SG. It was also demonstrated (15, 16) that mutant rats lacking ATP-dependent DNP-SG transport retain ATP-dependent TC transport, suggesting different transport mechanisms. In our membrane preparations, DNP-SG, a substrate for the non-bile acid organic anion carrier in rats (13, 19), did inhibit ATP-dependent TC transport. This may indicate the use of a common transporter. However, more studies on both rat and human membranes are warranted to investigate the nature and kinetics of this inhibition.

In conclusion, we have demonstrated the presence of a high-affinity, temperature-sensitive, and saturable ATP-dependent TC transport system in human liver plasma membranes. Since this transport is independent of membrane potential and proton gradients, it apparently reflects direct ATP-dependent transport.

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