Cellular Accumulation of Cholyl-Glycylamido-Fluorescein in Sandwich-Cultured Rat Hepatocytes: Kinetic Characterization, Transport Mechanisms, and Effect of Human Immunodeficiency Virus Protease Inhibitors

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ABSTRACT:
The present study was aimed at characterizing the in vitro cellular uptake mechanism and kinetics of the bile salt analog cholylglycylamido-fluorescein (CGamF) in sandwich-cultured rat hepatocytes (SCRHs). Concentration-dependent inhibition of active CGamF accumulation by seven human immunodeficiency virus (HIV) protease inhibitors (PIs) was also determined and compared with inhibition data obtained with taurocholate (TC) as a substrate. A \( K_m \) value of 9.3 ± 2.6 \( \mu M \) was obtained for saturable CGamF accumulation in SCRHs. The organic anion-transporting polypeptide (Oatp) inhibitor rifampicin (100 \( \mu M \)) inhibited CGamF (1 \( \mu M \)) accumulation in SCRHs by 72%; sodium depletion did not further reduce CGamF accumulation. In contrast, TC uptake was reduced by only 25% in the presence of rifampicin, whereas additional sodium depletion resulted in a complete loss of TC accumulation. These data imply that Oatp(s) and sodium taurocholate-cotransporting polypeptide preferentially mediate hepatic uptake of CGamF and TC, respectively. Coincubation of CGamF with HIV PIs (amprenavir, atazanavir, darunavir, indinavir, nelfinavir, ritonavir, saquinavir) revealed that five of them had a concentration-dependent inhibitory effect on CGamF accumulation in SCRHs, with IC\(_{50}\) values between 0.25 ± 0.07 and 43 ± 12 \( \mu M \). The rank order for inhibition of CGamF accumulation in SCRHs was: ritonavir \( \gg \) saquinavir \( \gg \) atazanavir \( \gg \) darunavir \( \gg \) amprenavir. Indinavir (up to 100 \( \mu M \)) did not alter CGamF accumulation, whereas nelfinavir solubility was limited to 10 \( \mu M \). Taken together, these findings illustrate the utility of CGamF as a suitable probe (complementary to TC) for rapid in vitro determination of interaction potential with sodium-independent uptake mechanisms (likely Oatps) in rat liver.

In the past decade, the key transport proteins mediating hepatic uptake and biliary excretion of endogenous compounds as well as xenobiotics have been identified, and their major roles have been recognized (Annaert et al., 2007). In the liver, the sodium taurocholate-cotransporting polypeptide [NTCP (human), Ntcp (rat), \( SLC10A1 \)] mediates sodium-dependent transport of conjugated and unconjugated bile salts across the basolateral (sinusoidal) membrane from blood to hepatocyte. Members of the organic anion-transporting polypeptide family [OATP (human), Oapt (rat); \( SLC21A1 \) gene family] also support hepatic uptake of bile salts (in addition to a variety of organic anions, neutral compounds, and cations) but in a sodium-independent manner. At the apical (canalicular) membrane of the hepatocyte, the bile salt export pump [BSEP (human), Bsep (rat); \( ABCB11/Abcb11 \)] and the multidrug resistance-associated protein-2 [MRP2 (human), Mrp2 (rat); ABCCC2/Abcc2] are responsible for bile salt excretion from hepatocytes to bile. As also stated in the European Federation for Pharmaceutical Sciences report on the 2005 Copenhagen Drug Transporters conference (Meijer and Lennernas, 2005), knowledge in this field has increased rapidly, yet quantitative insights into the exact impact of various transport proteins on absorption, distribution, metabolism, and excretion processes and toxicity in vivo remain limited. Nevertheless, interference with hepatobiliary transport of endogenous compounds (i.e., bile salts) is now clearly established as a possible mechanism of hepatotoxicity (Funk et al., 2001; Kostrubsky et al., 2003; Mita et al., 2006; Marion et al., 2007). Compounds that inhibit one or more of the hepatic transporters may cause toxicity following alterations in the levels of various endogenous compounds, such as bile salts, but also thyroid hormones and bilirubin. The detergent effects exerted by elevated intracellular concentrations of bile salts on hepatocyte membranes have been put forward as a mechanism to explain hepatotoxicity (Spivey et al., 1993; Funk et al., 2001).

This emphasizes the need for a better mechanistic understanding of transport processes through the use of new and existing in vitro models. Therefore, thorough characterization of in vitro models, not only with respect to transporter/enzyme expression at the protein and

ABBREVIATIONS: Ntcp, sodium taurocholate-cotransporting polypeptide; Oatp, organic anion-transporting polypeptide; MRP, multidrug resistance-associated protein; CGamF, cholyl-glycylamido-fluorescein; SCRH, sandwich-cultured rat hepatocyte; TC, taurocholate; HIV, human immunodeficiency virus; PI, protease inhibitor; FBS, fetal bovine serum; ANOVA, analysis of variance; RFP, rifampicin.
mRNA level, but also at the level of transporter function and activity, is crucial. In addition, a sufficiently diverse panel of model substrates (and inhibitors) needs to be identified and profiled with respect to transporter affinity and interaction. This will ultimately lead to well-characterized in vitro models enabling rapid and reliable screening for drug interaction and cellular toxicity potential.

In this context, the present study was aimed at characterizing the hepatic uptake mechanism and in vitro cellular kinetics of the conjugated bile salt cholyglycylamido-fluorescein (CGamF) in sandwich-cultured rat hepatocytes (SCRHs). The utility of CGamF as a convenient fluorescent probe simulating bile salt transport in various in vitro and in vivo models of different species has been demonstrated previously (Maglova et al., 1995; Bravo et al., 1998; Holzinger et al., 1998; Mita et al., 2006). Among the amido-fluorescein-tagged compounds, the cholyglycyl derivative (CGamF) was transported most efficiently. Its transport rate exceeded that of cholylamido-fluorescein and ursodeoxycholylamido-fluorescein, both of which were transported at the same rate (Holzinger et al., 1997). CGamF offers the advantages that no radiolabeled compound is needed and that it is amenable to fluorescence microscopy studies. However, the affinity profile of CGamF for human and rodent bile salt transporters appears to differ from the widely used model compound taurocholate (TC) (Mita et al., 2006). Hepatic TC transport is well characterized: its hepatic uptake is mediated by NTCP and OATPs for about 80 and 20% of total uptake, respectively (Meier et al., 1997; Kouzuki et al., 2000). Therefore, inhibition of OATPs alone would not be expected to affect TC accumulation kinetics to a large extent. The relative role of these transporters has not been reported in the case of the bile salt analog CGamF, potentially confounding interpretation of the actual impact of interference with its transport in vitro.

In addition to characterization of cellular CGamF kinetics in rat hepatocytes, the results of the present study illustrate the utility of this probe substrate for in vitro screening of interactions with hepatic uptake of endogenous and/or exogenous compounds by coinubations with seven HIV protease inhibitors (PIs). A study by McRae et al. (2006) already demonstrated that the antiretroviral drugs ritonavir, saquinavir, and efavirenz inhibit TC transport in sandwich-cultured human and rat hepatocytes. The results obtained in the present study with CGamF in sandwich-cultured rat hepatocytes support significant interaction of various HIV PIs with sodium-independent uptake transporters (most likely Oatps) in rat liver.

Materials and Methods

Chemicals. Amprenavir was kindly provided by GlaxoSmithKline (Uxbridge, Middlesex, UK). Ritonavir, indinavir sulfate, saquinavir mesylate, and nelfinavir mesylate were donated by Hetero Drugs Limited (Hyderabad, India). Atazanavir was obtained from Bristol-Myers Squibb Co. (Stamford, CT), darunavir was obtained from Cilag AG (Switzerland). 3-H-Taurocholic acid (5 Ci/mmol) and 3H-estrone-3-sulfate (57.3 Ci/mmol) were obtained from PerkinElmer Life and Analytical Sciences (Waltham, MA). Williams’ medium E, l-glutamine, penicillin-streptomycin mixture (contains 10,000 units of penicillin and 10,000 µg streptomycin sulfate/ml in 0.85% saline), fetal bovine serum (FBS), and Hanks’ balanced salt solution were purchased from Lonza Verviers SPRL (Verviers, Belgium). HEPES was purchased from Sigma-Aldrich (St. Louis, MO). Cholyl-glycylamido-fluorescein was kindly provided by Prof. Alan Hofmann (University of California, San Diego). All other chemicals and reagents were of analytical grade and were readily available from commercial sources. Standard buffer consisted of Hanks’ balanced salt solution, containing 10 mM HEPES and adjusted to pH 7.4. For experiments in which the effects on substrate accumulation were investigated, sodium-containing buffer and sodium-free buffer were prepared as described by Su et al. (2004). Sterile collagen was prepared in-house according to established procedures.

Animals. Male Wistar rats (240–300 g) were used for hepatocyte isolation from whole liver. The rats were housed according to the Belgian and European laws, guidelines, and policies for animal experiments, housing, and care in the Central Animal Facilities of the University. These facilities have the obligatory accreditation of the authorized Belgian Ministry and are registered under license number LA1210261. Approval for this project was granted by the Institutional Ethical Committee for Animal Experimentation.

Isolation of Rat Hepatocytes. Rat hepatocytes were isolated from male Wistar rats using a collagenase perfusion as described previously (Annaert et al., 2001), with slight modification; the recirculation system volume was 50 ml, and no trypsin inhibitor was added. The typical perfusion times were 10 min with Ca2+-free buffer, followed by about 6 min with buffer containing collagenase. Viability was determined using the trypan blue exclusion method. Viability was always >90% with a yield of 4 to 6 x 10⁸ cells. Cells were resuspended in Williams’ medium E and diluted to a final concentration of 1.33 x 10⁶ cells/ml.

Culture of Sandwich-Cultured Rat Hepatocytes. Six-well plates were coated with ice-cold collagen solution (~1.5 mg/ml final concentration; 0.1 ml/well) prepared by neutralizing a mixture of 4 ml of rat-tail collagen, 4 ml of deionized water, and 1 ml of 10X Dulbecco’s modified Eagle’s medium with 1 ml of 0.2 N NaOH solution (final pH, ~7.4). Each well was hydrated with 1.5 ml of phosphate-buffered saline and placed overnight at 37°C in a humidified incubator. Two million cells were added to each well and cultured for 1 to 2 h, and then unattached cells were removed by aspirating the medium. The cells were overlaid with 100 µl of rat tail collagen solution (1.5 mg/ml, pH 7.4) to obtain a “sandwich” configuration (“day 0”). One hour later, Williams’ medium E (1.5 ml/well, containing 5% FBS, 100 U/ml penicillin, 100 µg/ml streptomycin, 4 mg/l insulin, and 0.1 µm dexamethasone) was added onto the cultures. The medium was changed every day with FBS-containing medium.

Transport Studies with CGamF, H-Tc, and H-Estrone-3-Sulfate in Sandwich-Cultured Hepatocytes. SCRHs were rinsed twice with 1.5 ml/well standard buffer and preincubated for 10 min at 37°C. For experiments in which the effect of inhibitors was investigated, cells were preincubated for 10 min with the inhibitor at desired concentration. Subsequently, cells were coincubated in 1.5 ml of 1 µM substrate with or without inhibitors at desired concentration in standard buffer for 10 min (at 37°C unless mentioned otherwise) and subsequently rinsed four times with 1.5 ml of ice-cold standard buffer. Hepatocytes were lysed with 1 ml of 0.5% Triton X-100 solution (in phosphate-buffered saline) by placing plates on a shaker for 20 min at room temperature. Cell lysates were analyzed by fluorescence spectroscopy (excitation, 490 nm; emission, 524 nm) in a Tecan Infinite 200 plate reader (Tecan, Grödig, Austria) for determination of CGamF concentrations and by liquid scintillation spectroscopy (Wallac 1410; PerkinElmer Life and Analytical Sciences-Wallac Oy, Turku, Finland) for H-Tc or H-Estrone-3-sulfate. Accumulation was normalized to the protein content of the hepatocytes in each well and was measured by using the BCA Protein Assay kit (Pierce Chemical, Rockford, IL). All accumulation data were corrected for nonspecific binding to collagen-coated, hepatocyte-free culture plates.

Data Analysis. For the characterization of CGamF accumulation kinetics into SCRHs, the following equation was used:

\[ V = \frac{K_a \cdot C}{K_a + C} + V_{\text{max}} \cdot \frac{C}{K_c + C} \]

with \( K_a \) representing the rate constant for the nonsaturable accumulation and \( K_c \) and \( V_{\text{max}} \) representing the kinetic parameters for the saturable (Michaelis-Menten) accumulation. The best fit of the equation to the individual accumulation data sets was obtained by Pharsight WinNonlin software version 5.2 (Pharsight, Mountain View, CA).

In addition, the sigmoid inhibitory effect model was used to describe concentration-dependent inhibition of CGamF accumulation by taurocholate, rifampicin, and the various HIV PIs:

\[ E = \frac{E_{\text{max}} - (E_{\text{max}} - E_0) \cdot C^7}{C^7 + E \cdot C_{50}^7} \]
Results

Determination of CGamF Accumulation Kinetics in SCRHs.

Time-dependent cellular accumulation profiles of CGamF (1–50 μM) in SCRHs were linear for incubation times up to 90 s. Initial accumulation rates were concentration-dependent and could be described by a model incorporating both a Michaelis-Menten (saturable) and a linear (non-saturable) component. Average estimates for $V_{\text{max}}$ and $K_m$ were $462 \pm 30$ pmol/mg protein/min and $9.7 \pm 0.4$ μM/mg protein/min, respectively. Kinetic analysis was performed using the WinNonlin software version 5.2 (Pharsight).

Statistics. Two-tailed Student’s t test and ANOVA were used to evaluate statistical differences (SPSS version 11.0 for Windows; SPSS Inc., Chicago, IL) at an α level of 0.05.

Effect of Temperature on Accumulation of CGamF and Estrone-3-Sulfate in Day 1 SCRHs.

To confirm the prominent role of temperature in the primary transport mechanisms mediating CGamF accumulation, time course experiments were conducted at 4°C and 37°C. CGamF accumulation was clearly greater with rifampicin compared with TC, whereas the IC50 values (as determined in three independent experiments, i.e., three batches of hepatocytes) corresponded to inhibition of CGamF accumulation by rifampicin (3.2 ± 1.0 μM) and TC (9 ± 4 μM) were rather comparable.

Inhibition of CGamF Accumulation in SCRHs by HIV PIs.

The effect of preincubation and coinhibition of different HIV PIs [100 μM, except for nelfinavir (10 μM), ritonavir (50 μM), and saquinavir (50 μM), due to the solubility limitations] on CGamF and TC accumulation in SCRHs is shown in Fig. 5. The initial accumulation rate of CGamF was inhibited by amprenavir (100 μM), atazanavir (100 μM), saquinavir (50 μM), ritonavir (50 μM), and darunavir (100 μM) both in sodium-containing or sodium-free buffer. The accumulation rates were decreased to 144 ± 13, 130 ± 7, 109 ± 5, 144 ± 9, and

\[ \frac{\text{Passive rate}}{\text{Measured rate}} = \frac{0.98 \pm 0.06}{0.75 \pm 0.07} = 1.31 \]

\[ \frac{\text{Active rate}}{\text{Calculated rate}} = \frac{0.98 \pm 0.06}{0.75 \pm 0.07} = 1.31 \]
116 ± 10 pmol/mg protein/10 min, respectively, compared with control values of 323 ± 25 pmol/mg protein/10 min in sodium-containing buffer. The inhibition profiles obtained for these five PIs in sodium-free buffer only were only slightly lower than those obtained in the presence of sodium. In the case of darunavir, sodium depletion did not have any additive effect on CGamF accumulation. For nelfinavir (10 μM), there was no influence on CGamF accumulation in SCRHs and indinavir at 100 μM caused only slight inhibition of the CGamF accumulation. Except for the ~50% reduction of TC accumulation by saquinavir, the effects of the HIV PIs on TC accumulation remained rather limited in comparison with the effects on CGamF. In addition, although the effect of sodium depletion on CGamF accumulation was relatively small, TC accumulation fell to ~15% of control values when a combination of sodium-free buffer and HIV PIs was applied.

Figure 6 shows the concentration-dependent inhibition of CGamF accumulation in SCRHs by five HIV PIs. In this model system and under the present conditions, ritonavir, with an IC50 value of 0.25 ± 0.07 μM, was by far the most potent inhibitor. For the other four HIV PIs, more comparable inhibitory profiles were obtained. The rank order of inhibition of the CGamF accumulation in SCRHs is ritonavir > saquinavir > atazanavir > darunavir > amprenavir. The IC50 values along with other in vitro kinetic parameters are listed in Table 2. Interestingly, the maximum extent of inhibition of CGamF accumulation that was predicted for ritonavir was lower than the values obtained for the other HIV PIs. The in vitro data obtained in the present study were obtained in rat hepatocytes, and species differences
SCRHs offer the advantage that incubations can be done at higher temperature and for a more extended period of time. Data supporting the maintained expression and activity of the hepatic uptake transporters Ntcp and Oatp in rat hepatocytes early after isolation as well as in day 1 SCRHs have been generated previously. Liu et al. (1998) indeed showed superior expression of Ntcp in days 1 to 5 SCRHs compared with conventionally cultured rat hepatocytes as well as decreased Ntcp activity with increasing culture time. The data generated for TC accumulation (see Fig. 2) in day 1 SCRHs in the present study are consistent with these previous findings. More recently, Bow et al. published immunofluorescence images showing that Oatp1a1 remained expressed in the sinusoidal membrane of isolated rat hepatocytes, which was in contrast to the cytosolic internalization of efflux transporters (Mdr1, MRP2). In the present study, we also showed rifampicin- and temperature-inhibitable accumulation of the Oatp substrate estrone-3-sulfate, supporting expression of fully functional transporters in the plasma membrane of days 1 and 0 SCRHs. Furthermore, as shown by the data presented in Table 1, the use of day 1 versus day 4 SCRHs appears to offer the advantage of higher CGamF accumulation. A similar phenomenon was observed by Hoffmaster et al. in transporter-based interactions are likely to exist. Nevertheless, reported literature values for peak plasma concentration ranges of PIs observed in patients are also included in Table 2. Evaluation of these in vivo and in vitro data shows that the IC_{50} value for ritonavir is comparable with the plasma level of unbound ritonavir observed in patients. For the other protease inhibitors (amprenavir, darunavir, atazanavir, saquinavir), plasma levels are lower than IC_{50} values observed in vitro in rat hepatocytes.

**Discussion**

The aim of the present study was to characterize the bile salt analog CGamF as a probe substrate for in vitro hepatic uptake in rat hepatocytes and to demonstrate its utility for mechanistic investigation of drug interactions at the hepatic uptake level. A series of HIV protease inhibitors were selected as interacting model drugs. In addition, a comparison was made with the established probe substrate taurocholate.

As compared with freshly isolated suspended hepatocyte preparations, which have to be used within 4 to 6 h after isolation, day 1 SCRHs offer the advantage that incubations can be done at higher

![Fig. 5. Effect of different HIV PIs on the accumulation of CGamF (top) and TC (bottom) in SCRHs. Day 1 SCRHs were preincubated with inhibitors (APV, ATV, SQV, RTV, DRV) on the accumulation of 1 μM CGamF in SCRHs. Day 1 SCRHs were preincubated with inhibitors for 10 min, then coincubated with 1 μM CGamF and inhibitors at desired concentration for 10 min in standard buffer and intracellular accumulation measured as described under Materials and Methods. Points represent average ± S.D., n = 3 CGamF accumulation. Lines represent best fit of E_{max} – IC_{50} model to the accumulation data.](image-url)

![Fig. 6. Concentration-dependent inhibitory effect of different HIV PIs (APV, ATV, SQV, RTV, DRV) on the accumulation of 1 μM CGamF in SCRHs.](image-url)

**TABLE 2**

<table>
<thead>
<tr>
<th>Inhibitors</th>
<th>E_{max} – E_{o}</th>
<th>IC_{50}</th>
<th>Plasma Protein Binding*</th>
<th>C_{max} Corrected for Plasma Protein Binding**</th>
</tr>
</thead>
<tbody>
<tr>
<td>APV</td>
<td>188 ± 16</td>
<td>64 ± 6</td>
<td>43 ± 12</td>
<td>90</td>
</tr>
<tr>
<td>DRV</td>
<td>238 ± 45</td>
<td>81 ± 16</td>
<td>29 ± 7</td>
<td>93</td>
</tr>
<tr>
<td>ATV</td>
<td>196 ± 23</td>
<td>67 ± 7</td>
<td>13 ± 4</td>
<td>86</td>
</tr>
<tr>
<td>SQV</td>
<td>203 ± 24</td>
<td>69 ± 8</td>
<td>6 ± 2</td>
<td>98</td>
</tr>
<tr>
<td>RTV</td>
<td>158 ± 27</td>
<td>52 ± 10</td>
<td>0.25 ± 0.07</td>
<td>99</td>
</tr>
</tbody>
</table>

* From Swainston Harrison and Scott, 2005; Rittweger and Arasteh, 2007.
** From Williams and Sinko, 1999; Hoetelmans et al., 2003; Swainston Harrison and Scott, 2005; Marin-Niebla et al., 2007; Ruane et al., 2007.
Rat hepatocyte incubation were noted when taurocholate was used as a substrate; the gest inhibition of Ntcp-mediated CGamF accumulation by rifampicin. sodium depletion alone on CGamF accumulation, these results sug-
buffer. Together with the slight (but statistically significant) effect of did not differ between sodium-containing buffer and sodium-free
inhibition of Ntcp activity by rifampicin. In the present study, how-
motors in the presence of various concentrations of either rifampicin or Ntcp further confirmed the above findings (Fig. 3). Indeed, the maximum extent of inhibition of CGamF accumulation was higher for rifampicin (mainly affecting Oatps) compared with TC (mainly affecting Ntcp). Comparison of the dis-
eral uptake of these substrates (for a schematic summary, see also Fig. 7). First of all, sodium depletion resulted in only a slight reduction in CGamF accumu-
lations (Fig. 2), suggesting a limited role of Ntcp in active CGamF accumulation. Furthermore, in the presence of the known Oatp inhibitor rifampicin, CGamF accumulation was substantially reduced to ~30% of control levels. This represents a complete loss of active CGamF uptake since about 70% of CGamF uptake was calculated to result from an active process. Rifampicin has indeed been shown to be a potent inhibitor of Oatp2 (Oatp1a4) in LLC-PK1 cells (Shitara et al., 2002), rat hepatocytes (Fattinger et al., 2000), and of various OATP/SLCO isoforms in human hepatocytes (Vavricka et al., 2002). At a concentration of 100 μM, rifampicin was also shown to inhibit Oatp1a4-mediated transport of digoxin by 60% (Shitara et al., 2002). To date, no reports have been published on the possible inhibition of Ntcp activity by rifampicin. In the present study, however, CGamF accumulation in SCRHs in the presence of rifampicin did not differ between sodium-containing buffer and sodium-free buffer. Together with the slight (but statistically significant) effect of sodium depletion alone on CGamF accumulation, these results suggest inhibition of Ntcp-mediated CGamF accumulation by rifampicin. Completely different effects of sodium depletion and rifampicin co-
percentage of overall TC accumulation in day 4 SCRHs. The extent of inhibition of TC accumulation in sodium-containing buffer (and reflecting mainly Ntcp-mediated processes) remains rather limited, except for saquina-
vir (50% inhibition). On the other hand, the relative inhibition (up to 80%) of TC accumulation by sodium-independent processes (likely Oatps) is much more pronounced. It is of interest to compare our data regarding inhibition of TC accumulation with those obtained previously for ritonavir (RTV) and saquinavir (SQV) in day 4 SCRHs (McRae et al., 2006). First of all, overall TC uptake in day 1 SCRHs observed in the present study (about 130 pmol/mg protein/min) was substantially higher than the TC uptake observed in day 4 SCRHs in the earlier study (35 pmol/mg protein/min). This is consistent with declining expression and activity of uptake transporters (including Ntcp) with increasing culture times in SCRHs (Liu et al., 1998). Interestingly, the maximal extent of inhibition (about 80%) observed in the study by McRae et al. at the highest SQV and RTV concentrations corresponds to the relative inhibition percentages of sodium-independent TC accumulation by several HIV PIs in our study. This may reflect relatively higher contribution of sodium-independent processes to overall TC accumulation in day 4 SCRHs.

The results of this study have provided a direct comparison of the inhibitory potential of seven clinically used HIV protease inhibitors toward hepatic uptake transport of CGamF in rat liver. These data may aid in understanding relative differences between these compounds with respect to hepatotoxicity in the rat as the main toxicological species. Clearly, comparable in vitro data in human hepatocytes are

To further establish the usefulness of CGamF as a probe substrate for detecting Oatp-mediated interactions with active hepatic uptake of endogenous and exogenous substrates, a comparative in vitro study was conducted with HIV PIs as interacting drugs. Several HIV PIs are widely used in the clinic, mostly in combinations with other drugs. In addition, a few in vitro studies have provided evidence that HIV PIs display affinity for members of the Oatp family. Su et al. (2004) demonstrated for the first time that OATP-A (OATP1A2) was capable of transporting saquinavir in vitro. Concentration-dependent inhibition of [1H]estradiol-17ß-d-glucuronide uptake by indinavir, nelfinavir, saquinavir, and ritonavir was observed in OATP-C (OATP1B1)-transfected HeLa cells (Tirona et al., 2003). In the present study, concentration-dependent inhibition of (Oatp-mediated) CGamF accumulation in SCRHs by several HIV PIs (ritonavir, saquina-
vir, atazanavir, darunavir, amprenavir) was observed. For indinavir, only a slight effect was observed at 100 μM, whereas incubations with nelfinavir were only possible up to 10 μM due to solubility limitations. The most remarkable result with respect to inhibition of active CGamF accumulation in SCRHs is the substantially higher potency of ritonavir (IC50 = 0.25 μM), compared with the other HIV PIs. Indeed, the IC50 value for saquinavir, which appeared to be the second most potent HIV PI tested in this study, was almost 25 times higher.

The preferential interference of HIV PIs with sodium-independent (likely Oatps) transport mechanisms is also reflected in the inhibition data for TC accumulation (Fig. 5). The extent of inhibition of TC accumulation in sodium-containing buffer (and reflecting mainly Ntcp-mediated processes) remains rather limited, except for saquina-
vir (50% inhibition). On the other hand, the relative inhibition (up to 80%) of TC accumulation by sodium-independent processes (likely Oatps) is much more pronounced. It is of interest to compare our data regarding inhibition of TC accumulation with those obtained previously for ritonavir (RTV) and saquinavir (SQV) in day 4 SCRHs (McRae et al., 2006). First of all, overall TC uptake in day 1 SCRHs observed in the present study (about 130 pmol/mg protein/min) was substantially higher than the TC uptake observed in day 4 SCRHs in the earlier study (35 pmol/mg protein/min). This is consistent with declining expression and activity of uptake transporters (including Ntcp) with increasing culture times in SCRHs (Liu et al., 1998). Interestingly, the maximal extent of inhibition (about 80%) observed in the study by McRae et al. at the highest SQV and RTV concentrations corresponds to the relative inhibition percentages of sodium-independent TC accumulation by several HIV PIs in our study. This may reflect relatively higher contribution of sodium-independent processes to overall TC accumulation in day 4 SCRHs.

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needed to determine clinical relevance of the present data as well as potential species differences in drug interactions. Nevertheless, it is interesting to note that free plasma concentrations observed clinically for ritonavir in patients (Table 2) range around the IC_{50} value obtained for this PI in the present study.

Inhibition of one or more Oatp isoforms by HIV protease inhibitors can potentially result in drug interactions with Oatp substrates. In addition, because certain endogenous substrates also rely on Oatp transporters (e.g., thyroid hormones), (hepato)toxicity may result. Hepatotoxicity has been reported as an adverse effect in the prescribing information for the U.S. Food and Drug Administration-approved HIV PIs, nelfinavir, indinavir, atazanavir, ritonavir, lopinavir/ritonavir, saquinavir, ampranavir, and fosamprenavir. The incidence values ranged from one to two cases per 100 patients for nelfinavir, which was the lowest among this group, and from 2.2 to 9.5 for lopinavir/ritonavir (Bruno et al., 2006). Sulkowski et al. (2000) also observed that the relative risk of severe hepatotoxicity in patients receiving a PI-containing regimen was more than twice that seen in patients who were prescribed regimens with dual nucleoside analogs. The possible role of interference with Ntcp/NTCP-mediated bile salt transport as a mechanism of hepatotoxicity associated with antiviral drugs has already been investigated in the aforementioned in vitro study by McRae et al. However, a simple relationship between inhibition of NTCP-mediated bile salt uptake in human hepatocytes and the risk for hepatotoxicity in the clinic could not be established (McRae et al., 2006).

In the light of the obtained results in the present study, inhibition of Oatp-mediated substrate transport (in addition to NTCP-mediated transport) by HIV PIs may be considered as another potential mechanism in PI-associated hepatotoxicity. Although the present data were obtained in rat hepatocytes, and species differences are expected to exist, it is interesting to note that ritonavir exhibits the highest risk for hepatotoxicity in clinical therapy. This finding may be especially important given the widespread combination of ritonavir as a pharmacokinetic booster with other HIV PIs to improve pharmacokinetics of these PI.

In conclusion, our data have illustrated that thorough characterization and careful use of complementary probe substrates are important to get better insights in relative contributions of various transport mechanisms in drug interaction potential and risk for hepatotoxicity. Our data confirm that compared with hepatocytes cultured in a conventional configuration, SCRHs already after 1 day in culture represented a model system for in vitro evaluation of inhibition of hepatic uptake transporters by drugs. Quantitative in vitro data on inhibition of hepatic uptake transporters may help to understand mechanisms underlying side effects of certain HIV PIs.

References


