Interaction of Ethambutol with Human Organic Cation Transporters of the SLC22 Family Indicates Potential for Drug-Drug Interactions during Antituberculosis Therapy

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According to the 2012 WHO global tuberculosis (TB) report (http://apps.who.int/iris/bitstream/10665/75938/1/9789241564502 _eng.pdf), the death rate for tuberculosis was over 1.4 million patients in 2011, with ~9 million new cases diagnosed. Moreover, the frequency of comorbidity with human immunodeficiency virus (HIV) and with diabetes is on the rise, increasing the risk of these patients for experiencing drug-drug interactions (DDIs) due to polypharmacy. Ethambutol is considered a first-line antituberculosis drug. Ethambutol is an organic cation at physiological pH, and its major metabolite, 2,2’-(ethylenediamino) dibutyric acid (EDA), is zwitterionic. Therefore, we assessed the effects of ethambutol and EDA on the function of human organic cation transporter 1 (hOCT1), hOCT2, and hOCT3 and that of EDA on organic anion transporter 1 (hOAT1) and hOAT3. Potent inhibition of hOCT1- and hOCT2-mediated transport by ethambutol (50% inhibitory concentration $IC_{50} = 92.6 \pm 10.9$ and $253.8 \pm 90.8 \mu M$, respectively) was observed. Ethambutol exhibited much weaker inhibition of hOCT3 ($IC_{50} = 4.1 \pm 1.6 \mu M$); however, significant inhibition ($>80\%$) was observed at physiologically relevant concentrations in the gastrointestinal (GI) tract after oral dosing. EDA failed to exhibit any inhibitory effects that warranted further investigation. DDI analysis indicated a strong potential for ethambutol interaction on hOCT1 expressed in enterocytes and hepatocytes and on hOCT3 in enterocytes, which would alter absorption, distribution, and excretion of coadministered cationic drugs, suggesting that in vivo pharmacokinetic studies are necessary to confirm drug safety and efficacy. In particular, TB patients with coexisting HIV or diabetes might experience significant DDIs in situations of coadministration of ethambutol and clinical therapeutics known to be hOCT1/hOCT3 substrates (e.g., lamivudine or metformin).

Ethambutol dihydrochloride (EMB) (Fig. 1) is a potent antmycobacterial agent employed in the treatment of tuberculosis (TB) and Mycobacterium avium complex infections. The main pharmacological effect of EMB is to inhibit arabinosyl transferase, preventing the synthesis of arabinogalactan, which is a vital component of the mycobacterial cell wall. EMB, pyrazinamide, isoniazid, and rifampin comprise the four first-line antituberculosis drugs recommended by the World Health Organization (WHO) (1). EMB is used in the treatment of multidrug-resistant tuberculosis to prevent the emergence of isoniazid- or rifampin-resistant mycobacteria (2). In clinical practice, EMB use is associated with many adverse effects, including optic neuritis (reported in 1% to 5% of patients) (3) and reduced renal clearance of urate (reported in about 66% of patients) (4), the incidence of which is higher when EMB is concomitantly administered with pyrazinamide (5).

The in vivo pharmacokinetic properties of EMB have been well defined (6–8). In humans, EMB showed high bioavailability (~80%) and low plasma protein binding (20 to 30%) (9). A peak serum concentration ($C_{\text{max}}$) of ~4.5 μg/ml was reported after oral dosing of 25 mg/kg (6). Only a small portion of EMB (8% to 15%) is metabolized in the liver to form the final dicarboxylic metabolite, 2,2’-(ethylenediamino) dibutyric acid (EDA) (Fig. 1) (10). Renal elimination is the major clearance mechanism for EMB, with 70 to 84% of an intravenous dose being excreted in the urine as the unchanged parent compound (9). The renal clearance rate (~417 ml/min) indicates that active net tubular secretion, as well as glomerular filtration (120 ml/min), is involved (6). While EMB has been identified as a substrate of P-glycoprotein (11), based upon their chemical structures (Fig. 1) and physicochemical properties, EMB and EDA have the potential to be a substrate and/or inhibitor of organic cation transporters (OCTs) and/or organic anion transporters (OATs).

The organic cation/anion/zwitterion transporters belong to the solute carrier 22 (SLC22) family and in vivo mediate the absorption, distribution, and elimination of a broad variety of charged endogenous and exogenous organic substances (12–14). OATs and OCTs are widely expressed in many barrier organs, such as the intestine, kidney, liver, and brain (12–14). In the intestine (Fig. 2), human OCT1 (hOCT1) (SLC22A1) is expressed in the basolateral membrane and hOCT3 (SLC22A3) is targeted to the brush border membrane of enterocytes, where they mediate the cellular entry of cationic compounds (12). hOCT1 and hOCT3 are both expressed in the sinusoidal membrane (blood side) of hepatocytes (Fig. 2) and represent the first step in the hepatic excretion of many substances (12, 15). In the kidney (Fig. 2), hOCT2 (SLC22A2), hOCT3, hOAT1 (SLC22A6), and hOAT3 (SLC22A8) are each expressed on the basolateral sides of proximal tubule cells and extract drugs from the blood. hOCT1 mRNA has been detected; however, the protein has not been immunolocal-
ized (12, 13). Hundreds of important clinical therapeutics are known substrates and/or inhibitors of OATs and OCTs, such as chemotherapeutics (e.g., cisplatin and paclitaxel), analgesics (e.g., morphine), cholesterol-lowering drugs (e.g., atorvastatin and pravastatin), antivirals (e.g., cidofovir and lamivudine), and antidiabetic agents (e.g., metformin) (12–14).

Recently, it was reported that TB infection is closely linked to both human immunodeficiency virus (HIV) and diabetes. According to the WHO, 10% of TB patients also suffer from diabetes and 13% of the ~9 million TB patients newly diagnosed in 2011 were coinfected with HIV (1). Thus, the complexity of these disease treatment regimens (TB, TB-diabetes, or TB-HIV), in terms of polypharmacy, raises concerns regarding the potential for drug-drug interactions (DDIs) involving OCTs and/or OATs. The aim of the present study was to explore the inhibitory effects of EMB and its metabolite EDA on hOCT1, hOCT2, hOCT3, hOAT1, and hOAT3. Potent inhibition was further characterized by kinetic investigations to estimate IC50s, which were used to quantitatively evaluate the clinical DDI potential on these transporters during antituberculosis therapy.

MATERIALS AND METHODS

Chemicals. Tritiated p-aminohippuric acid ([3H]PAH), estrone sulfate ([3H]ES), and 1-methyl-4-phenylpyridinium ([3H]MPP⁺) were purchased from PerkinElmer Life and Analytical Science (Waltham, MA). Unlabeled ES, MPP⁺, PAH, probenecid, and ethambutol dihydrochloride (EMB, ≥95% purity) were obtained from Sigma-Aldrich (St. Louis, MO). Quinine monohydrochloride dihydrate was purchased from Acros Organics (Fair Lawn, NJ). 2,2′-(Ethylenediamine)dibutyric acid (EDA; ≥99% purity) was purchased from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA).

Tissue culture. Derivation of stably transfected Chinese hamster ovary (CHO) cell lines expressing hOAT1 (CHO-hOAT1) and stably transfected human embryonic kidney 293 (HEK) cells expressing hOAT3 (HEK-hOAT3), hOCT1 (HEK-hOCT1), hOCT2 (HEK-hOCT2), or hOCT3 (HEK-hOCT3), as well as their corresponding empty vector-transfected background control cell lines, has been described previously (16–19). CHO-hOAT1 cells were maintained at 37°C with 5% CO2 in phenol red-free RPMI 1640 medium (Gibco-Invitrogen, Grand Island, NY) containing 10% serum, 1% penicillin/streptomycin, and 1 mg/ml G418. HEK-hOAT3 cells were maintained at 37°C with 5% CO2 in high-glucose Dulbecco’s modified Eagle medium (DMEM; Mediatech, Inc., Herndon, VA) containing 10% serum, 1% penicillin/streptomycin, and 125 µg/ml hygromycin B. The HEK-hOCT1, HEK-hOCT2, and HEK-hOCT3 cell lines were maintained at 37°C with 5% CO2 in high-glucose DMEM containing 10% serum, 1% penicillin/streptomycin, and 600 µg/ml G418.

Cell accumulation assay. The procedure for the cell accumulation assay has been described previously (20, 21). Briefly, cells were seeded into 24-well tissue culture plates at a density of 2 × 10⁵ cells/well (without antibiotics) for 48 h. On the day of the cell transport experiment, the cells were equilibrated to serum-free conditions with transport buffer for 10 min (500 µl of Hanks’ balanced salt solution containing 10 mM HEPES, pH 7.4). Equilibration buffer was replaced with 500 µl of fresh transport buffer containing 1 µM unlabeled substrate spiked with [3H]PAH (0.5 µCi/ml), [3H]ES (0.25 µCi/ml), or [3H]MPP⁺ (0.25 µCi/ml) in the presence or absence of test compounds. At the end of the incubation, the cells were quickly rinsed three times with ice-cold transport buffer and lysed. The radioactivity of cell lysate was quantified by liquid scintillation count-
Drug-drug interaction (DDI) index calculation. The DDI index (22) was estimated as the ratio of the maximal unbound EMB concentration in biofluid (unbound $C_{max}$) after therapeutic dosing of EMB divided by transporter-specific $IC_{50}$s estimated by in vitro assay. A cutoff value of $\geq 0.1$ is thought to indicate the need for a prospective in vivo pharmacokinetic study (22). Specifically, the predicted intestinal lumen concentration of EMB after oral dosing (25-mg/kg dose, 70-kg patient body weight, 250-ml volume) was used to estimate the DDI index for hOCT3 located in the apical (luminal) membranes of enterocytes, while the portal venous plasma concentration (presystemic circulation) of EMB was used to estimate DDI index values for hOCTs in the basolateral membranes of hepatocytes (hOCT1 and hOCT3) and/or enterocytes (hOCT1 and hOCT2), and the plasma concentration (systemic circulation) of EMB was used to estimate DDI index values for hOCTs in the basolateral membranes of renal proximal tubule cells (Fig. 2). The concentration of EMB in the intestinal lumen was estimated to be $\sim 34 \mu M$, assuming that patients (70 kg) take EMB with 250 ml of water. From clinical reports, after oral dosing (25 mg/kg), the plasma $C_{max}$ of EMB was 22.02 $\mu M$, with about 20% being protein bound (6, 9, 23). Accordingly, the method of Ito et al. can be used to estimate drug concentrations in the portal vein (24): $C_{port,vn} = C_{max} + (k_d \times D)/(Q_s \times F_p)$, where $C_{port,vn}$, $k_d$, and $F_p$ represent concentration in the portal vein, absorption rate constant, dose, hepatic blood flow rate ($\sim 1.200 \, \text{ml/min}$), and fraction absorbed, respectively. For EMB, $k_d$ and $F_p$ were reported as 0.48 h$^{-1}$ and 80% (9, 23). Using an average patient body weight of 70 kg, the predicted portal venous blood concentration for EMB was $\sim 67 \, \mu M$.

Statistical analysis. The data in the figures and the raw uptake scores are expressed as means $\pm$ standard deviations (SD), while $IC_{50}$ estimates are means $\pm$ standard errors of the means (SEM). Statistical differences were assessed using one-way analysis of variance (ANOVA) followed by post hoc analysis with Dunnett’s t test ($\alpha = 0.05$).

RESULTS

Inhibitory effects of EMB on hOCT1-, hOCT2-, and hOCT3-mediated MPP$^\mathbf{\ast}$ uptake. The inhibitory effects of EMB were examined on three OCT paralogs, hOCT1, hOCT2, and hOCT3, using MPP$^\mathbf{\ast}$ as a prototypical substrate. Significant accumulation of MPP$^\mathbf{\ast}$ ($\sim 33$-fold) was observed in stably transfected hOCT1-expressing cells relative to empty-vector-transfected background control cells (132.6 $\pm$ 9.9 versus 4.0 $\pm$ 0.2 pmol/mg protein/10 min, respectively). The known hOCT inhibitor, quinine (200 $\mu M$), showed virtually complete inhibition of hOCT1-mediated MPP$^\mathbf{\ast}$ uptake ($\sim 90\%$ inhibition) (Fig. 3A). Significant inhibition (66%) of hOCT1-mediated MPP$^\mathbf{\ast}$ transport by EMB (at 100 $\mu M$ versus 1 $\mu M$ MPP$^\mathbf{\ast}$) was observed (Fig. 3A). Subsequent dose-response ($10^{-7}$ to $10^{-3}$ M EMB) studies were performed to derive the $IC_{50}$ for EMB on hOCT1 (Fig. 4, top). The $IC_{50}$ of EMB for hOCT1 was estimated as 92.6 $\pm$ 10.9 $\mu M$. Therefore, the DDI indices were calculated to gauge DDI potency in enterocytes, hepatocytes, and proximal tubule cells, using the EMB concentration in gastrointestinal (GI) biofluid, portal venous blood, or the systemic circulation, as appropriate (Fig. 2 and Table 1).

We next examined the inhibitory effect of EMB on hOCT2-mediated transport. HEK-hOCT2 cells exhibited marked accumulation of MPP$^\mathbf{\ast}$ (93.0 $\pm$ 5.2 pmol/mg protein/10 min) compared to background control cells (4.0 $\pm$ 0.2 pmol/mg protein/10 min). Quinine (200 $\mu M$) completely blocked ($> 99\%$) hOCT2-mediated MPP$^\mathbf{\ast}$ uptake (Fig. 3B). The cell accumulation assay demonstrated that EMB at 100 $\mu M$ significantly inhibited hOCT2-mediated MPP$^\mathbf{\ast}$ uptake (21%), and complete inhibition of hOCT2 transport activity (96%) was observed at 1 mM EMB. Kinetic studies were conducted to estimate the $IC_{50}$ for EMB on hOCT2. Using EMB concentrations ranging from $10^{-6}$ to $10^{-3}$ M, the $IC_{50}$ was estimated as 253.8 $\pm$ 90.8 $\mu M$ (Fig. 4, middle). As shown in Table 1, the DDI index for hOCT2 expressed in kidney was estimated as 0.1.

Stably transfected hOCT3-expressing (HEK-hOCT3) cells also showed marked accumulation of MPP$^\mathbf{\ast}$ ($\sim 30$-fold) compared to empty-vector-transfected background control cells (80.3 $\pm$ 11.3 versus 2.7 $\pm$ 0.1 pmol/mg protein/10 min). Such active transport was completely (99%) blocked by quinine at 200 $\mu M$ (Fig. 3C).
Though EMB failed to show significant inhibition on hOCT3 transport activity at 100 μM, marked inhibition was observed with increasing concentrations (Fig. 3C). EMB showed ~50% and ~80% inhibition at 1 and 10 mM, respectively. Further dose-response (10⁻³ to 10⁻¹ M EMB) studies yielded an estimated IC₅₀ of 4.1 ± 1.6 mM (Fig. 4, middle). Due to low EMB concentration in the presystemic (portal) and systemic circulation, DDI indices resulting from EMB were not performed.

**Inhibitory effects of EDA on hOAT (hOAT1 and hOAT3) and hOCT (hOCT1, hOCT2, and hOCT3) transport activity.** The zwitterionic dicarboxylic acid metabolite of EMB, EDA, was examined for interactions with hOAT1, hOAT3, hOCT1, hOCT2, and hOCT3 (Fig. 5). The standard substrate used for hOAT1 was PAH, and that for hOCT3 was ES. Stably transfected hOAT1-expressing (CHO-hOAT1) and hOAT3-expressing (HEK-hOAT3) cells showed 5-fold- and 3-fold-greater substrate accumulation than empty-vector-transfected background control cells, respectively (7.5 ± 1.6 versus 1.6 ± 0.2 pmol/mg protein/10 min for hOAT1 and 4.1 ± 0.2 versus 1.6 ± 0.1 pmol/mg protein/10 min for hOAT3). The known OAT inhibitor probenecid completely blocked (>99%) hOAT1- and hOAT3-mediated substrate uptake (Fig. 5). With the exception of hOAT3 (~17% inhibition), EDA (100 μM) failed to produce any significant inhibitory effects. Despite the modest inhibition of hOAT3, for all tested transporters IC₅₀ of EDA were expected to be greater than 100 μM, which greatly exceeds the EDA concentration in the systemic circulation, and further kinetic analysis was not performed.

**DISCUSSION**

Although numerous antitubercular drugs have been developed since the 1940s, TB is still a prevalent disease and a common cause of death worldwide. Recently, the spread of multidrug-resistant TB strains (MDR-TB; estimated to account for 3.7% of new cases and 20% of previously treated cases) and the emergence of an extensively drug-resistant strain (XDR-TB; ~9% of MDR-TB cases) that has proven virtually impossible to successfully treat has become new challenges to disease management (1, 25). The appearance of XDR-TB has been reported in 84 countries, and the threat of its spread is viewed as a significant enough health issue that the United Kingdom has enacted regulations obligating people traveling from regions where TB is prevalent to have a current chest X-ray in their possession upon arrival in order to be issued a visa (26). Isolation and therapy may be required for up to 2 years and can cost upwards of one million U.S. dollars per patient (26).

A further complication in TB therapy arises from the increasing clinical association of TB with diabetes and HIV, with TB becoming one of the leading causes of mortality among HIV-infected patients (27, 28). Indeed, of the ~9 million new TB patients diagnosed in 2011, 13% were reported to be coinfectcd with HIV, and of the 1.4 million TB deaths, 430,000 were HIV associated (1). As a result, more TB patients need to be prescribed combination therapies to treat comorbidities or to minimize the development of drug resistance, making therapy management and compliance extremely complex and increasing the risk of patients’ experiencing unintended drug-drug interactions. Indeed, there are a number of case reports regarding adverse events, including nephrotoxicity, associated with EMB use during antituberculosis therapy (29–32). In these patients, discontinuation of EMB resulted in improvement in renal function.

Competition for binding to membrane transporters is one mechanism resulting in DDIs, and while there is currently insufficient clinical data to conclusively determine transporter-mediated DDI involvement, it is possible that hOCT-mediated DDIs

**TABLE 1 Estimated DDI index values for EMB on hOCT-mediated transport after an oral dose of 25 mg/kg**

<table>
<thead>
<tr>
<th>hOCT</th>
<th>GI tract</th>
<th>Liver</th>
<th>Kidney</th>
</tr>
</thead>
<tbody>
<tr>
<td>hOCT1</td>
<td>0.6</td>
<td>0.6</td>
<td>0.2</td>
</tr>
<tr>
<td>hOCT2</td>
<td>NE</td>
<td>NE</td>
<td>0.1</td>
</tr>
<tr>
<td>hOCT3</td>
<td>8.3 &lt;0.1</td>
<td>&lt;0.1</td>
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*a Drug-drug interaction index is defined as the unbound concentration of drug divided by the drug IC₅₀ for the transporter of interest. A DDI index value of >0.1 is thought to indicate the potential for clinically relevant DDIs.

*b Assuming basolateral targeting for hOCT1.

NE, transporter not expressed in this tissue.
Role of hOCTs and hOATs in Antituberculosis Therapy

In summary, our findings demonstrate the inhibitory effect of EDA on hOATs and hOCTs. (A) Inhibition of hOAT1-mediated PAH uptake by EDA (100 μM) and probenecid (1,000 μM). (B) Inhibition of hOAT3-mediated ES uptake by EDA (100 μM) and probenecid (1,000 μM). (C) Inhibition of hOCT1-mediated MPP⁺ uptake by EDA (100 μM) and probenecid (1,000 μM). (D) Inhibition of hOCT2-mediated MPP⁺ uptake by EDA (100 μM) and quinine (200 μM). (E) Inhibition of hOCT3-mediated MPP⁺ uptake by EDA (100 μM) and quinine (200 μM). EMB concentrations in hepatocytes were estimated to be 0.6 μM, and in enterocytes, 0.2 μM; all the DDI indexes are above the 0.1 threshold. The IC₅₀ for hOCT3 was higher (4.1 μM), indicating that 38% of tubule cells (0.2 μM) might be inhibited in enterocytes and potentially renal proximal tubule cells. Furthermore, the estimated DDI index of 8.3 (Fig. 4, bottom, and Table 1) is 83-fold higher than the DDI threshold value and is indicative of a marked potential for EMB to interfere (89% inhibition) with intestinal absorption of coadministered drugs that are hOCT3 substrates, including metformin.

As many anti-HIV agents are also hOCT substrates, the same DDI potential exists for polypharmacy in TB-HIV patients. Recently, expression of hOCT1 and hOCT2 mRNA was detected in CD4 cells isolated from patients with HIV (42). Therefore, EMB present in the systemic circulation also might reduce accumulation of anti-HIV agents in CD4 target cells that serve as a viral reservoir, resulting in loss of antiviral efficacy via inhibition of hOCT1 and/or hOCT2 function. Similarly, systemic EMB might impact hOCT1-mediated (DDI index = 0.2; ~20% inhibition) and/or hOCT2-mediated (DDI index = 0.1; ~9% inhibition) renal secretion.

hOCTs are also involved in the pharmacological action of platinum antineoplasics (e.g., cisplatin and oxaliplatin), raising the possibility for significant DDIs during chemotherapy in TB patients (43–46). Patients receiving cisplatin therapy frequently experience cumulative dose-dependent nephrotoxicity involving the proximal tubule (47–49). Moreover, treatment with inhibitors of hOCTs (e.g., tetraethylammonium and cimetidine) prevented accumulation of cisplatin in renal proximal tubule cells, and subsequently, cisplatin was identified as a competitive inhibitor of rat Oct2 (43, 50–53). Thus, coadministration of EMB and cisplatin or oxaliplatin might alter their oral absorption, as well as their associated hepatic and renal accumulation and toxicity.

In summary, our findings demonstrate the inhibitory effect of EMB on hOCT1, hOCT2, and hOCT3, as well as in the wildtype ferromorphs. hOCTs in TB-diabetes patients.
Human OCT3 appear to be necessary in order to optimize clinical safety between EMB and known substrates for hOCT1, hOCT2, and hOCT3 to appear to be necessary in order to optimize clinical safety and efficacy in these complex patient populations.

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REFERENCES