Prediction of Clinically Relevant Herb-Drug Clearance Interactions Using Sandwich-Cultured Human Hepatocytes: Schisandra spp. Case Study

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ABSTRACT

The Schisandraceae family is reported to have a range of pharmacological activities, including anti-inflammatory effects. As with all herbal preparations, extracts of Schisandra species are mixtures composed of >50 lignans, especially schizandrins, deoxyschizandrins, and gomisins. In China, Schisandra sphenanthera extract (SSE) is often coadministered with immunosuppressant treatment of transplant recipients. In cases of coadministration, the potential for herb-drug interactions (HDIs) increases. Clinical studies have been used to assess HDI potential of SSE. Results demonstrated that chronic SSE administration reduced midazolam (MDZ) clearance by 52% in healthy volunteers. Although clinical studies are definitive and considered the “gold standard,” these studies are impractical for routine HDI assessments. Alternatively, in vitro strategies can be used to reduce the need for clinical studies. Transporter-certified sandwich-cultured human hepatocytes (SCHHs) provide a fully integrated hepatic cell system that maintains drug clearance pathways (metabolism and transport) and key regulatory pathways constitutive active/androstan receptor and pregnane X receptor (CAR/PXR) necessary for quantitative assessment of HDI potential. Mechanistic studies conducted in SCHHs demonstrated that SSE and the more commonly used dietary supplement Schisandra chinensis extract (SCE) inhibited CYP3A4/5-mediated metabolism and induced CYP3A4 mRNA in a dose-dependent manner. SSE and SCE reduced MDZ clearance to 0.577- and 0.599-fold of solvent control, respectively, in chronically exposed SCHHs. These in vitro results agreed with SSE clinical findings and predicted a similar in vivo HDI effect with SCE exposure. These findings support the use of an SCHH system that maintains transport, metabolic, and regulatory functionality for routine HDI assessments to predict clinically relevant clearance interactions.

Introduction

Conventional medicines share the same metabolic enzymes and xenobiotic transporters (e.g., cytochrome P450 enzymes, glucuronosyltransferases, P-glycoprotein, and organic ion transporters) that evolved in response to early human exposure to plant alkaloids and other phytochemicals. Thus, combinations of herbal ingredients with modern drugs may result in herb-drug interactions (HDIs) mediated by those metabolic enzymes and transporters. Interactions that increase plasma drug levels are reduced below minimal effective concentrations. The occurrence and significance of any specific HDI depends on a variety of factors, such as dose, frequency and timing of herb/drug intake, dosing regimen, route of administration, and therapeutic range (Gurley, 2012; Sprouse and van Bremen, 2016).

In China, Schisandra sphenanthera extract (SSE) is prescribed as a tonic to improve hepatic function and is often coadministered with tacrolimus in treating renal and liver transplant recipients (Xin et al., 2007). In this case, any potential HDI may be life threatening; therefore, it is important to predict these interactions. In recent human clinical studies, administration of SSE altered the first-pass metabolism, bioavailability, and plasma pharmacokinetics of tacrolimus (Qin et al., 2010). In similar studies, SSE increased the bioavailability and decreased the clearance of midazolam, with differential effects at the level of the gastrointestinal tract and liver (Xin et al., 2009). These data indicated clinically significant inhibitory effects on CYP3A4/5 in the presence of SSE.

Although clinical studies are definitive and considered the “gold standard,” the number of combinations of herbal extracts and clinically used drugs is so large it is not feasible to routinely perform clinical-interaction studies (Roe et al., 2016). Alternatively, in vitro strategies easily scale and can be used to reduce the need for clinical studies (Food and Drug Administration, 2006, 2012; European Medicines Agency, 2012). However, the accuracy of in vivo prediction from translation of

ABBRévATIONS: DMSO, dimethylsulfoxide; HDI, herb-drug interaction; HPLC, high-pressure liquid chromatography; ICC, intracellular concentration; LC-MS/MS, liquid chromatography–tandem mass spectrometry; MDZ, midazolam; OHMDZ, 1-hydroxymidazolam; RIF, rifampicin; SCE, Schisandra chinensis extract; SCHH, sandwich-cultured human hepatocyte; SJW, St. John’s wort; SSE, Schisandra sphenanthera extract; TAO, trolleyandamycin.
in vitro results has been the “Achilles heel” of in vitro drug-interaction strategies. The choice of model system has been suggested to affect the prediction accuracy (McGinnity et al., 2005; Brown et al., 2007; Hewitt et al., 2007; Sprouse and van Breeemen, 2016). For example, hepatic microsomes lack cellular processes including transporter (uptake and efflux) and adaptive response (e.g., induction) function. Therefore, microsomal studies cannot account for mechanisms that may affect unbound cellular concentrations in the hepatocyte. The lack of these processes likely decreases the accuracy of microsomes as an in vitro drug-drug interaction model.

Primary human hepatocytes have become the preferred in vitro system for transport, metabolism, toxicity, and drug-interaction studies due to their biologic (e.g., relevant species) and physiologic (e.g., whole-cell model) relevance (Hewitt et al., 2007). Regulatory agencies, such as the Food and Drug Administration and European Medical Agency, have adopted primary human hepatocyte cultures as the standard in vitro model for predicting human cytochrome P450 enzyme induction by pharmaceuticals (European Medicines Agency, 2012; Food and Drug Administration, 2012). Under the proper culture conditions in conjunction with properly characterized human hepatocytes, a fully integrated hepatic cell system that maintains drug clearance pathways (metabolism and transport) and key regulatory pathways constitutive active/androstane receptor and pregnane X receptor (CAR/PXR) can be used to predictively screen for potential HDI.

Herein, we used a retrospective in vitro study design to evaluate the use of Transporter-Certified sandwich-cultured human hepatocytes (SCHHs) to predict in vivo HDI outcomes. Transporter-certified hepatocytes, which have been characterized for drug uptake and efflux transporter function while maintaining metabolism and nuclear receptor function, were used in this in vitro study to ensure hepatic physiologic relevance. SSE was used as a model herbal extract with proven clinical HDI while a second closely related and popular herbal dietary supplement was used in this in vitro study to ensure hepatic physiologic relevance. SSE was used as a model herbal extract with proven clinical HDI while a second closely related and popular herbal dietary supplement extract, *Schisandra chinensis* extract (SCE), with an uncharacterized HDI profile was evaluated in parallel. Both mechanistic (e.g., inhibition and induction) and net effect (e.g., intrinsic clearance) studies were performed in SCHHs to demonstrate the effectiveness of this model to predict in vivo HDI outcomes.

### Materials and Methods

**Materials**

Ketoconazole, aflatoxin, midazolam (MDZ), rifampicin (RIF), schisandrin A, gomisin A, schizandrin, d4-hydroxymidazolam, triazolam, dimethylsulfoxide (DMSO), and d5-atorvastatin were purchased from Toronto Research Chemicals (Toronto, Canada). Troleandomycin (TAO) was purchased from Cayman Chemical (Ann Arbor, MI). 1-Hydroxymidazolam (OHMDZ) was chased from Sigma-Aldrich (St. Louis, MO). 1-Hydroxymidazolam (OMHDZ) was purchased from Cayman Chemical (Ann Arbor, MI). Troleandomycin (TAO) and d5-atorvastatin were purchased from Toronto Research Chemicals (Toronto, ON, Canada). High-pressure liquid chromatography (HPLC)–grade methanol and acetonitrile were purchased from World Wide Medical Products (Bristol, PA).

Base cell culture media and other cell culture reagents were obtained from Thermo Fisher Scientific (Waltham, MA) and were of the highest purity available. St. John’s wort (SJW) extract was purchased from PureBulk (Roseburg, OR). CellTiter-Glo Luminescent Cell Viability Assay was purchased from Promega (Madison, WI). Qiagen RNasey kit and reagents were purchased from Qiagen (Germantown, MD). Quant-iT RiboGreen RNA Assay Kit, High Capacity cDNA Archive Kit, and TaqMan primer/probes were purchased from Thermo Fisher Scientific. Whatman 90-well Unifilter 25-mm melt blown polypropylene (MBPP) 0.45-μm polypropylene (PP) filter plates, Millipore MDRPNNP protein precipitation plate, and Millipore MSHVN45 0.45-μm filter plate were purchased from World Wide Medical Products.

Extracts of *Schisandra* fruit from both *S. schisandra* (SSE) and *S. chinensis* (SCE) were purchased from Flavax, Naturextracte GmbH (Rehlingen-Siersburg, Germany) and stored at room temperature prior to use. Extracts for both species were prepared from the dried fruit of *Schisandra spp.*, by supercritical fluid extraction with natural carbon dioxide, with no solvent residues, no inorganic salts, no heavy metals, and no reproducible microorganisms. Thus, SSE and SCE contain all CO₂-soluble lipophilic components, and both extracts have the appearance of a brown oily liquid with a weak aromatic smell. The SSE used in these studies contained 2.7% (g/100 g) deoxyschizandrin (also called schisandrin A) as detected by gas chromatography. The sum of schizandrin/gomisin (calculated as deoxyschizandrin) was 5.2%. The SCE used in these studies contained 0.65% deoxyschizandrin (schisandrin A), 1.8% schizandrin, and 0.13% gomisin A, as detected by the supplier using gas chromatography. The sum of schizandrin/gomisins (calculated as deoxyschizandrin) was 7.2%.

### Cell Culture

SCHHs were prepared using QTS transporter-certified cryopreserved human hepatocytes (Table 1) purchased from Triangle Research Laboratories (Qualyst Transporter Solutions; Durham, NC). Cryopreserved hepatocytes were thawed following the manufacturer’s thawing instructions. SCHHs were prepared by plating cryopreserved hepatocytes suspended in QualGro Seeding Medium, QTS propriety hepatocyte seeding medium, at a density of 0.8 million viable cells/ml onto BioCoat 24-well cell culture plates purchased from Corning (Corning, NY), and maintained under standard cell culture conditions (37°C, 5% CO₂, 95% humidity). Following plating, cells were allowed to attach for 2–4 hours, rinsed, and fed with warm (37°C) seeding medium. Eighteen to 24 hours later, cells were fed and overlaid with QualGro Hepatocyte Culture Induction Medium supplemented with 0.35 mg/ml Matrigel purchased from Corning. Cells were maintained in QualGro Hepatocyte Culture Induction Medium.

**SSE and SCE Stock and Dosing Solution Preparation.** SSE and SCE were provided as dark-yellow oils and diluted in DMSO to produce stock solutions, which were stored at room temperature. For chronic dosing studies (72 hours), SSE and SCE dosing solutions were prepared fresh daily in QualGro Hepatocyte Culture Induction Medium from these stock solutions, with final DMSO content ≤0.14%. Treatment of hepatocyte cultures began on day 2 and continued for 72 hours. Each day, treatment solutions were removed, and fresh treatment solutions were added to hepatocytes. On day 5, MDZ intrinsic clearance assays were conducted, and cells were harvested for RNA isolation and intracellular concentration determinations.

**Cytotoxicity Assessment.** On day 5 of culture, viability of hepatocytes was quantitatively assessed by determining the amount of adenosine triphosphate (ATP) content present following 24 and 72 hours of exposure to increasing concentrations of SSE or SCE (0.30, 3.0, 30, and 300 μg/ml and 0.32, 3.2, and 320 μg/ml, respectively). Aflatoxin (10 μM) and media plus solvent (0.14% DMSO) were included as positive and negative controls, respectively. Cellular ATP content was measured using CellTiter-Glo Luminescent Cell Viability Assay from Promega following the manufacturer’s instructions. Luminescence detection of samples was measured using a SpectraMax GeminiXS 96-well Plate Reader (Molecular Devices, Sunnyvale, CA). Each test condition was performed in triplicate. The percentage of control was calculated as the ratio of ATP content in treated samples compared with that in wells treated with vehicle control (0.14% DMSO) as described in eq. 1:

\[
\% \text{ of control} = \frac{\text{ATP content treatment}}{\text{ATP content 0.1% DMSO}} \times 100
\]

#### CYP3A4/5 Enzyme Inhibition

On day 5 of culture, direct and time-dependent inhibition of CYP3A4/5 in the presence and absence of SSE or SCE (0.30, 3.0, 30, and 300 μg/ml) was quantitatively assessed.

**TABLE 1**

<table>
<thead>
<tr>
<th>Hepatocyte Lot</th>
<th>Sex</th>
<th>Experiment</th>
<th>Figure</th>
</tr>
</thead>
<tbody>
<tr>
<td>HUM4059</td>
<td>Female</td>
<td>24-hour cytotoxicity assessment</td>
<td>Fig. 1A</td>
</tr>
<tr>
<td>HUM4075B</td>
<td>Male</td>
<td>72-hour cytotoxicity assessment</td>
<td>Fig. 1B</td>
</tr>
<tr>
<td>HUM4059</td>
<td>Female</td>
<td>CYP3A4/5 inhibition evaluation</td>
<td>Fig. 2</td>
</tr>
<tr>
<td>HUM4109</td>
<td>Female</td>
<td>CYP3A4 mRNA induction assessment</td>
<td>Fig. 3</td>
</tr>
<tr>
<td>HUM4122C</td>
<td>Male</td>
<td>Intracellular concentration evaluation</td>
<td>Fig. 4</td>
</tr>
<tr>
<td>HUM4122C</td>
<td>Male</td>
<td>MDZ intrinsic clearance studies</td>
<td>Figs. 5 and 6</td>
</tr>
</tbody>
</table>
32, and 320 µg/ml, respectively) was assessed by monitoring the formation of OHMDZ. In parallel incubations, direct (50 and 250 nM of ketonazole) and time-dependent (1 and 10 µM TAO) positive control inhibitors were evaluated as system controls. In brief, the medium of 5-day cultures of human hepatocytes was replaced with Williams’ E medium containing 5 µM MDZ or without SSE, SCE, or positive control inhibitor. Direct inhibition studies were conducted in situ at 37°C for 20 minutes. Following the conclusion of the incubation, medium was removed and stored at −80°C prior to sample analysis. To investigate time-dependent inhibition of CYP3A4, 5–day cultures of hepatocytes were incubated with SSE, SCE, or positive control for 60 minutes prior to addition of midazolam. After an additional 20-minute MDZ incubation period, medium was collected and stored at −80°C prior to sample analysis. Samples and matrix-matched calibration standards were extracted (100-µl sample volume) by the addition of 300 µl of internal standard solution (methanol containing 25 nM triazolam) using a protein precipitation plate stacked on a 96-deep-well block. The plates were shaken prior to centrifugation for collection of the filtered supernatant. The sample filtrate was evaporated to dryness, and the samples were reconstituted in 200-µl sample diluent, 40/60 methanol/10 mM ammonium acetate, and mixed for at least 20 minutes on a plate shaker. The reconstituted samples were transferred to a Millipore 0.45-µm filter plate, filtered into a 96-deep-well plate by centrifugation, and sealed with a silicone capmat. Liquid chromatography–tandem mass spectrometry (LC-MS/MS) analysis for the formation of the OHMDZ metabolite was performed using an API-3000 mass spectrometer (Applied Biosystems, Foster City, CA) operated in positive electrospray ionization mode. Chromatographic separation for OHMDZ was achieved using a binary HPLC system with LC-10ADvp pumps (Shimadzu, Columbia, MD) equipped with a Hypersil Gold 100 × 1.0 mm, 3 µm with matching guard and precolumn filter (Thermo Scientific, Bellefonte, PA) using a linear gradient of 10%–100% B at a flow rate of 0.1 ml/min, with mobile phase A as 10% methanol with 0.25 mM ammonium acetate and mobile phase B as 80% methanol with 0.25 mM ammonium acetate. Quantitation of OHMDZ was determined by HPLC-MS/MS monitoring for the mass spectrometry transition for OHMDZ (m/z 342→324). Calibration standards were prepared in blank media and extracted in parallel with study samples for quantitation of OHMDZ.

To quantify inhibition, percentage of remaining activity was calculated as shown in eq. 2:

\[
\% \text{ of remaining activity} = \frac{\text{OHMDZ formation of treated samples}}{\text{OHMDZ formation of solvent control}} \times 100
\]

(2)

Nonlinear regression analysis of the inhibition dose-response curves (e.g., percentage of remaining activity versus test article concentration) was conducted using GraphPad Prism Software (version 6.0; GraphPad Software, La Jolla, CA). Estimated IC₅₀ values and their respective 95% confidence intervals were determined using the nonlinear regression analysis functions within the software package. The strength of the parameter estimates was determined based upon the coefficient of determination (R²).

CYP3A4 mRNA Induction

SCHHs were treated with increasing concentrations of SSE or SCE (0.30, 3.0, 30, and 300 µg/ml and 0.32, 3.2, 32, and 320 µg/ml, respectively) or with a clinically relevant CYP3A4 inducer, RIF (10 µM), beginning on day 2 of culture and continuing for 72 hours. An extract of SJW (20 µg/ml), an herbal extract known to cause CYP3A4 induction, was also included for comparison. Following 72 hours of treatment (day 5 of culture), SCHHs were washed once with 1 volume of Hanks’ balanced salt solution and lysed by addition of 0.3 ml of Qiagen RLT Buffer supplemented with β-mercaptoethanol and frozen at −80°C. Lysates were thawed and total RNA was isolated from each treatment group, pooled from triplicate wells, and used to investigate hepatic SCHH gene expression after centrifugation. The sample filtrate was evaporated to dryness, and the samples were reconstituted in 150 µl of sample diluent, 60/40 methanol, and 10 mM ammonium acetate and mixed for at least 10 minutes on a plate shaker. The reconstituted samples were transferred to a Millipore 0.45-µm filter plate, filtered into a 96-deep-well plate by centrifugation, and sealed with a silicone capmat prior to LC-MS/MS analysis. LC-MS/MS analysis was performed using an Applied Biosystems API-3000 mass spectrometer operated in positive electrospray ionization mode. Chromatographic separation for schisandrin A, gomisin A, and schizandrin was achieved using a Shimadzu binary HPLC system with LC-10ADvp pumps equipped with a Thermo Scientific Hypersil Gold AQ column (100 × 1.0 mm, 3 µm) with matching guard and precolumn filter using a linear gradient of 20%–100% B, with mobile phase A as 0.2% formic acid in water and mobile phase B as 0.2% formic acid in acetonitrile. Quantitation was determined by HPLC-MS/MS monitoring for the mass spectrometry transition for schisandrin A (m/z 417→399), gomisin A (m/z 417→399), and schizandrin (m/z 435→415). Calibration standards were prepared using previously frozen 24-ml deep-well plates derived from SCHH spiked with known concentrations of each constituent and extracted in parallel with unknown samples. Sample concentrations were reported as peak-area ratios back calculated from the calibration curve. The ICC of each analyte was calculated by dividing the mass of analyte in the hepatocytes (cellular accumulation) by the hepatocyte intracellular fluid volume of 7.69 µl/mg protein for hepatocytes, as previously described (Lee and Brouwer, 2010). All mass values were generated from triplicate wells and were represented by the calculated mean and S.D. from these biologic replicates.

MDZ Intrinsic Clearance Assays

Intrinsic clearance of MDZ was evaluated in SCHH on day 5 of culture following 72 hours of exposure to RIF (10 µM). SJW (20 µg/mg), and SSE (3 and 30 µg/ml) and SCE (3 and 30 µg/ml). Twenty-four hours after last dosing, cell culture medium was aspirated and replaced with Williams’ E medium containing MDZ (1 µM). At multiple time points, incubations were terminated with the addition of 900 µl of methanol containing 100 µM α-hydroxymidazolam-d₄ (internal standard) and frozen at −80°C prior to analysis. The supernatants were removed, filtered, dried down under nitrogen, and reconstituted in 150 µl of 60/40 methanol/water with 10 mM ammonium acetate and analyzed for the loss of gene-expression analysis. Amplifications were performed on a ViiA 7 Real-Time PCR System (Thermo Fisher Scientific) in relative quantification mode for 45 amplification cycles using standard conditions for TaqMan-based assays. Threshold cycle (Cₚ) determinations were performed by the ViiA 7 system software for both target and endogenous control genes. Relative-fold mRNA content was determined for each treatment group relative to the endogenous control gene expression and the calibrator, DMSO vehicle control, using the ViiA 7 system software. Confidence intervals of 95% were calculated for each target gene relative quantitation mean by the ViiA 7 system software.

Relative-fold mRNA content that was less than 1-fold, indicating suppression of gene expression, was transformed using eq. 3 to enhance the graphical representation of these data:

\[
\text{relative fold suppression} = \frac{-1}{\text{relative fold mRNA content}}
\]

(3)

Intracellular Concentration of SSE and SCE Constituents

The intracellular concentration (ICC) of three constituents of SSE and SCE (schisandrin A, gomisin A, and schizandrin) was determined in SCHH dosed with SCE or SSE at final assay concentrations of 30 and 300 µg/ml for 72 hours. Following 72 hours of treatment, cell culture medium was removed, and the tight junctions of treated SCHH were modulated using B-CLEAR technology (QTS; Triangle Research Laboratories) as previously described (Liu et al., 1999; Chandra and Brouwer, 2004; Ghiebellini et al., 2007; Marion et al., 2012).

Cells were placed in frozen storage at −80°C overnight prior to extraction, and quantitation of the constituents (schisandrin A, gomisin A, and schizandrin) was conducted. To extract, a volume of 500 µl of lysis solution, 70:30 methanol/water (v:v) containing 100 nM d5-atorvastatin (internal standard), was added to each well. Plates were shaken for at least 15 minutes, and the cell lysates were transferred to a Whatman 96-well Unifilter 25-µM MBPP/0.45-µm PP filter plate stacked on a 96-deep-well plate. Lysate was filtered into the deep-well plate by centrifugation. The sample filtrate was evaporated to dryness, and the samples were reconstituted in 150 µl of sample diluent, 60/40 methanol, and 10 mM ammonium acetate and mixed for at least 10 minutes on a plate shaker. The reconstituted samples were transferred to a Millipore 0.45-µm filter plate, filtered into a 96-deep-well plate by centrifugation, and sealed with a silicone capmat prior to LC-MS/MS analysis. LC-MS/MS analysis was performed using an Applied Biosystems API-3000 mass spectrometer operated in positive electrospray ionization mode. Chromatographic separation for schisandrin A, gomisin A, and schizandrin was achieved using a Shimadzu binary HPLC system with LC-10ADvp pumps equipped with a Thermo Scientific Hypersil Gold AQ column (100 × 1.0 mm, 3 µm) with matching guard and precolumn filter using a linear gradient of 20%–100% B, with mobile phase A as 0.2% formic acid in water and mobile phase B as 0.2% formic acid in acetonitrile. Quantitation was determined by HPLC-MS/MS monitoring for the mass spectrometry transition for schisandrin A (m/z 417→399), gomisin A (m/z 417→399), and schizandrin (m/z 435→415). Calibration standards were prepared using previously frozen 24-ml deep-well plates derived from SCHH spiked with known concentrations of each constituent and extracted in parallel with unknown samples. Sample concentrations were reported as peak-area ratios back calculated from the calibration curve.
parent MDZ using an Applied Biosystems API-3000 mass spectrometer operated in positive electrospray ionization mode. Chromatographic separation for MDZ was achieved using a Shimadzu binary HPLC system with LC-10ADvp pumps equipped with a Thermo Scientific Hypersil Gold column (50 × 1.0 mm, 3 μm) with matching guard and precolumn filter using a linear gradient of 10%–100% B, at a flow rate of 0.075 ml/min, with mobile phase A as 0.2% formic acid in water and mobile phase B as 0.2% formic acid in methanol. Quantitation of MDZ was determined by HPLC-MS/MS monitoring for the mass spectrometry transition for MDZ (m/z 326 → 248).

The substrate-depletion approach was used to estimate metabolic intrinsic clearance. Linear regression analyses of the percentage of parent remaining (log-transformed) versus time profiles were used to estimate the elimination rate constant (k). Intrinsic clearance was estimated assuming monoexponential decay using eq. 4:

\[
\text{Clearance}_{\text{intrinsic}} = k \times V / N
\]

where \(V\) = total incubation volume, \(N\) = milligram of protein in the incubation, and \(k\) = elimination rate constant. The observed metabolic intrinsic clearance was scaled utilizing scaling factors derived from physiologic parameters for their respective species (Davies and Morris, 1993). To quantify the change in MDZ intrinsic clearance in response to treatment, percentage of control was calculated as the ratio of intrinsic clearance observed in treated samples to intrinsic clearance observed in vehicle control (0.14% DMSO) as described in eq. 5:

\[
\% \text{ of control} = \left( \frac{\text{MDZ intrinsic clearance}_{\text{treatment}}}{\text{MDZ intrinsic clearance}_{\text{0.14%DMSO}}} \right) 
\]

Statistical Analyses. All calculations were performed using Excel 2010 (Microsoft, Redmond, WA) unless otherwise stated. Statistical analysis was performed using GraphPad Prism Software (version 6.0). A one-way analysis of variance followed by a Dunnett’s test was performed for statistical comparisons of ATP content and MDZ intrinsic clearance. An * in figures denotes a \(P\) value \(\leq 0.05\).

Results

Cytotoxicity Assessment. Intracellular ATP content was evaluated following 24 and 72 hours of treatment to assess cytotoxicity of \textit{Schisandra} spp. extracts, SSE (0.30, 3.0, 30.0, and 300 μg/ml) and SCE (0.32, 3.2, 32, and 320 μg/ml). The cellular ATP content in SCHHs was significantly (\(P\) value \(\leq 0.05\)) reduced to 27.4% and 10.9% of solvent control following 24 and 72 hours, respectively, of exposure to aflatoxin (10 μM) (Fig. 1). These results demonstrated that the SCHHs were sensitive to known metabolism-dependent toxins, such as aflatoxin (Essigmann et al., 1977). No marked reduction in SCHH cell viability was observed in SCHHs treated for 24 or 72 hours with 0.30, 3.0, or 30 μg/ml SSE and 0.32, 3.2, or 32 μg/ml SCE (Fig. 1). In contrast, ATP content was significantly (\(P\) value \(\leq 0.05\)) reduced to 86.5% and 76.5% of solvent control in SCHHs following 24 hours of treatment with 300 μg/ml SSE and 320 μg/ml SCE (Fig. 1A). Following 72 hours of treatment with 300 μg/ml SSE and 320 μg/ml SCE, ATP content was significantly (\(P\) value \(\leq 0.05\)) reduced to 89.5% and 86.5% of solvent control in SCHHs (Fig. 1B). Overall, these results suggested that SSE and SCE were noncytotoxic at concentrations \(\leq 30\) and 32 μg/ml, respectively, following 24 or 72 hours of exposure.

CYP3A4/5 Inhibition Assessment. CYP3A4/5 activity was evaluated in the absence and presence of SSE (0.30, 3.0, 30.0, and 300 μg/ml) and SCE (0.32, 3.2, 32, and 320 μg/ml) with and without preincubation to assess the inhibition potential of \textit{Schisandra} spp. extracts. The rate of OHMDZ formation was significantly decreased approximately 28% or 72% when SCHHs were incubated in the presence of the known inhibitor ketoconazole (50 or 250 nM, respectively; data not shown). Similarly, inclusion of either SCE or SSE in the incubation medium significantly decreased the rate of OHMDZ formation in a concentration-dependent manner. In the absence of preincubation, CYP3A4 enzyme activity was reduced to 3.29% and 4.13% of solvent control at the highest SSE (300 μg/ml) and SCE (320 μg/ml) concentrations examined in SCHHs (Fig. 2). These results demonstrated that both \textit{Schisandra} spp. extracts have the potential to inhibit CYP3A4/5 activity as direct inhibitors. Nonlinear regression analysis of the inhibition plots (percentage of remaining activity vs. the log concentration of SCE or SSE) estimated IC\(_{50}\) values of 4.46 μg/ml for SCE and 1.23 μg/ml for SSE.

When TAO (1 μM) was added to wells 60 minutes prior to the addition of MDZ, the rate of OHMDZ formation by SCHH was reduced approximately 80% compared with a 20% reduction when TAO was not preincubated, thus demonstrating time-dependent inhibition by TAO (data not shown). SSE (3 μg/ml) and SCE (3.2 μg/ml) reduced the rate of OHMDZ formation to 18.6% and 24.6% of solvent control, respectively, in SCHH following a 60-minute preincubation (Fig. 2). Without preincubation, the OHMDZ formation rate by SCHH was reduced to 37.4% and 67.1% of solvent control (Fig. 2) in the presence of SSE (3 μg/ml) and SCE (3.2 μg/ml), correspondingly. Taken together, these results demonstrated that inhibition of OHMDZ formation in the
Inhibition of CYP3A4/5 by SSE (A) or SCE (B), without preincubation (●) and with preincubation (■), evaluated in SCHH by monitoring the formation of OFMDZ from DZ. SSE no preincubation IC$_{50}$ = 1.23 µg/ml, 95% confidence interval = 0.823–1.85, goodness of fit = 0.989; SSE preincubation IC$_{50}$ = 0.378 µg/ml, 95% confidence interval = 0.299–0.479, goodness of fit = 0.999. SCE no preincubation IC$_{50}$ = 4.46 µg/ml, 95% confidence interval = 4.44–4.48, goodness of fit = 1.0; SCE preincubation IC$_{50}$ = 1.45 µg/ml, 95% confidence interval = 1.44–1.46, goodness of fit = 1.0. Data are from a single experiment with treatments performed in triplicate wells. Mean values are reported ± S.D.

Fig. 2. Inhibition of CYP3A4/5 by SSE (A) or SCE (B), without preincubation (●) and with preincubation (■), evaluated in SCHH by monitoring the formation of OFMDZ from DZ. SSE no preincubation IC$_{50}$ = 1.23 µg/ml, 95% confidence interval = 0.823–1.85, goodness of fit = 0.989; SSE preincubation IC$_{50}$ = 0.378 µg/ml, 95% confidence interval = 0.299–0.479, goodness of fit = 0.999. SCE no preincubation IC$_{50}$ = 4.46 µg/ml, 95% confidence interval = 4.44–4.48, goodness of fit = 1.0; SCE preincubation IC$_{50}$ = 1.45 µg/ml, 95% confidence interval = 1.44–1.46, goodness of fit = 1.0. Data are from a single experiment with treatments performed in triplicate wells. Mean values are reported ± S.D.

presence of SSE (3 µg/ml) and SCE (3.2 µg/ml) in SCHH was time-dependent. Nonlinear regression analysis of the inhibition plots (with and without preincubation) estimated a ≥3-fold shift in the calculated IC$_{50}$ values for both Schisandra spp. extracts. The inhibition potency shift of both extracts was statistically significant (P value ≤0.05), since the calculated 95% confidence intervals of the IC$_{50}$ values (with and without preincubation) did not intersect. The observed inhibition potency shift confirmed that both extracts exhibited the potential to inhibit CYP3A4/5 activity in a time-dependent manner (Fig. 2).

CYP3A4 mRNA Induction Assessment. Following 72 hours of treatment, CYP3A4 mRNA content was evaluated to assess CYP3A4 induction potential of Schisandra spp. extracts, SSE (0.30, 3.0, 30.0, and 300 µg/ml) and SCE (0.32, 3.2, 32, and 320 µg/ml). CYP3A4 mRNA content was induced 8.56-fold and 5.54-fold greater than solvent control in SCHHs treated with RIF (10 µM) and SJW extract (20 µg/ml), respectively (Fig. 3). These results demonstrated that SJW, a well studied herbal extract, produced a CYP3A4 induction response that was

65% of the response elicited by RIF. These results demonstrated that the SCHH system was sensitive to both pharmaceutical and herbal CYP3A4-type inducers. Dose-related induction of CYP3A4 mRNA content was observed, ranging from 2.09- to 6.27-fold and from 2.08- to 7.48-fold in SCHH exposed to 3.0 and 30 µg/ml SSE and 3.2 and 32 µg/ml SCE, respectively (Fig. 3). CYP3A4 induction responses provoked by 72 hours of treatment with SSE (30 µg/ml) and SCE (32 µg/ml) were greater than the response produced by SJW extract treatment and were 73% and 87% of the RIF response. CYP3A4 induction responses were muted in SCHH treated with the highest concentrations of SSE and SCE examined (Fig. 3). Loss of ATP content was observed at the highest concentrations of SSE and SCE examined and may explain the reduced CYP3A4 induction response observed under these conditions. Overall, the results demonstrated that both SSE and SCE have the potential to strongly induce CYP3A4 mRNA content in SCHH under the conditions studied.

Intracellular Concentration of Schisandra spp. Constituents. B-CLEAR technology was used to determine the ICC of Schisandra spp. extract constituents, including schisandrin A, gomisin A, and schizandrin, in SCHH following 72 hours of treatment with SSE (30 and 300 µg/ml) or SCE (30 and 300 µg/ml). These constituents were chosen because they represent the more abundant lignans found across Schisandra spp. and, therefore, could be used to show that such lignans could accumulate in hepatocytes. ICCs of schisandrin A increased in a dose-related manner in hepatocytes treated with SSE, resulting in concentrations ranging from 28.2 to 1228 µM (Fig. 4). Gomisin A and schizandrin were not detected in SCHH lysates treated with SSE at any concentration examined. All three constituents were observed in hepatocytes treated with SCE, where ICCs of gomisin A and schisandrin A increased in a dose-related manner, ranging from 1.51 to 6.40 µM and 14.3 to 438 µM, respectively (Fig. 4). In contrast, schizandrin ICCs were not observed to increase in SCHH treated with increasing concentrations of SSE. Schisandrin A accumulation was 1.97-fold and 2.8-fold greater in SCHH treated with SSE at 30 and 300-µg/ml dose levels, respectively, compared with SCHH treated with SCE. In general, these data demonstrated that constituents of Schisandra spp. extracts were extensively present in hepatocytes, and each extract examined had a different intracellular concentration profile.
MDZ Intrinsic Clearance. SCHHs were treated for 72 hours prior to monitoring MDZ (1 μM) depletion over time to assess SSE (3 or 30 μg/ml) or SCE (3 or 30 μg/ml) treatment on intrinsic clearance. Percentage of parent drug remaining versus time profiles (Fig. 5) demonstrated that MDZ intrinsic clearance was clearly increased in SCHHs treated with known CYP3A4 inducers SJW (20 μg/ml) and RIF (10 μM). MDZ intrinsic clearance was increased to 2.36-fold or 11.8-fold above solvent control in SCHH treated with SJW or RIF, respectively (Fig. 6A). These results were in agreement with clinical studies that demonstrated a 2.0-fold (Backman et al., 1996) and 24-fold (Wang et al., 2001) increase in the mean MDZ clearance in humans following treatment with SJW or RIF, respectively. In contrast, MDZ intrinsic clearance in SCHH treated with either SSE (3 or 30 μg/ml) or SCE (3 or 30 μg/ml) was reduced (Fig. 5), ranging from 0.518- to 0.577-fold and 0.646- to 0.599-fold of solvent control, respectively, consistent with inhibition of metabolism (Fig. 6B). The in vitro results of SSE were in agreement with clinical findings that showed a 0.52-fold reduction of MDZ clearance across 12 healthy male subjects following a 7-day SSE treatment regimen (Xin et al., 2009). Similar in vitro results were observed for SCE, suggesting that SCE treatment has the potential to cause a similar in vivo effect.

Discussion

Dietary supplement sales have grown in the United States, reaching over $6 billion in 2015 (Smith et al., 2016). Consumer usage data for dietary supplements from 2007 to 2011 found that 64%–69% of respondents regularly use dietary supplements beyond simple multivitamins (Dickinson et al., 2014). In conjunction with increased dietary supplement usage, concurrent use with prescription medications is of concern, especially in older adults (Jou and Johnson, 2016). Thus, more routine assessments of herbal ingredients with prescription medications appear warranted.

Although clinical studies are definitive, a clinical strategy is unrealistic for routine HDI assessments. A more realistic approach would be to use a physiologic, relevant in vitro model system, such as SCHH. Using properly characterized hepatocytes (e.g., transporter-certified) under optimized culture conditions, SCHHs are particularly well suited to evaluate complex mixtures for the potential of HDI, as the hepatic whole-cell system maintains drug metabolism (phase I/I) and transporter (uptake/efflux) clearance pathways and vital drug clearance regulatory pathways (CAR/PXR) (Ghibellini et al., 2007; Hewitt et al., 2007; Jackson et al., 2009). Integration of these components into a whole-cell system is necessary in predicting HDI.

In this study, we evaluated the effectiveness of SCHH to predict in vivo HDI outcomes using an intrinsic clearance approach. Mechanistic inhibition and induction studies of CYP3A4 were performed to further elucidate the drug-interaction mechanisms in SCHH following exposure to herbal extracts. SSE was used as a model herbal extract with a known MDZ clinical HDI, whereas the closely related and increasingly popular herbal dietary supplement extract SCE, with an unknown clinical MDZ HDI profile, was evaluated in parallel.

Mechanistic inhibition studies in SCHH demonstrated that both SSE and SCE reduced the rate of OHMDZ formation in a dose-dependent manner with and without preincubation. Nonlinear regression analysis of the inhibition curves demonstrated that the inhibition potency (IC50) of both Schisandra spp. extracts increased ≥3-fold with preincubation, indicative of time-dependent inhibition. Toxicity did not appear to be the reason for the loss of OHMDZ formation since ATP content remained unchanged following 24-hour exposure to 30 μg/ml SSE and 32 μg/ml SCE. At these concentrations, inhibition of OHMDZ formation reached maximum effect, reducing activity to 99% and 98% of control, respectively, following a 60-minute preincubation. Taken together, these results demonstrated that loss of CYP3A4/5 activity was not attributable to cytotoxicity but was the result of enzyme time-dependent inhibition.

Using a standard in vitro approach recommended by the regulatory agencies (European Medicines Agency, 2012; Food and Drug Administration, 2012), induction of CYP3A4 mRNA was evaluated using transporter-certified hepatocytes in a sandwich-culture format following 72 hours of exposure to increasing concentrations of either Schisandra spp. extract. Interestingly, robust dose-related induction of CYP3A4 mRNA was observed in SCHH treated with either Schisandra spp. extract across the concentrations examined. A maximum CYP3A4 mRNA response 6.27-fold and 7.48-fold greater than vehicle control was observed in SCHH treated with 30 μg/ml SSE or 32 μg/ml SCE, respectively. Also, the maximum CYP3A4 induction observed in response to SSE or SCE was equivalent (113% and 135% of SJW) or almost equal (73.2% and 87.4% of RIF) to induction responses of known CYP3A4 inducers SJW and RIF. An induction response of 20%–40% of
positive control has previously been proposed as demonstrating clinically relevant induction potential (Bjornsson et al., 2003; Food and Drug Administration, 2006; Fahmi et al., 2010). Induction of CYP3A4 mRNA was reduced from maximum in SCHH treated at the highest concentrations of SSE or SCE evaluated. Following 72 hours of treatment at these high concentrations only, ATP content was significantly ($P$ value $\leq 0.05$) reduced, suggesting that the loss of CYP3A4 induction may have been the result of hepatotoxicity.

Our approach, using a fully integrated whole-cell system, demonstrated that both *Schisandra* spp. extracts significantly inhibited and induced CYP3A4 metabolism, a drug clearance pathway extensively involved in the clearance of $\approx 30\%$ of clinically prescribed medications (Zanger and Schwab, 2013). These results suggested that both *Schisandra* spp. extracts have the potential to elicit an HDI within a major drug clearance pathway; however, the net effect of *Schisandra* spp. extract exposure was difficult to estimate from these separate mechanistic studies. To evaluate the HDI net effect and predict an overall clinical effect, further experiments were performed using the same whole-cell system treated for 72 hours with either SSE or SCE (3.0 and 30 $\mu$g/ml) prior to monitoring the intrinsic clearance of MDZ.

Demonstrating that this system was responding appropriately, the positive controls, SJW and RIF, increased intrinsic clearance of MDZ 2.36-fold and 11.8-fold above solvent control, respectively. These data were consistent with reported clinical MDZ drug interactions. In clinical drug-interaction studies, SJW and RIF increased MDZ clearance 2.0-fold and 24-fold, correspondingly (Backman et al., 1996; Wang et al., 2001). These results demonstrated that an intrinsic clearance approach in SCHH using a sensitive probe substrate (MDZ) to assess the drug interaction net effect (e.g., induction) and the relative strength of the interaction was feasible. These in vitro results were in good agreement with clinical-interaction studies of well known perpetrators of in vivo drug interactions, SJW and RIF. Using our integrated SCHH system, intrinsic clearance of MDZ was reduced to 0.518-fold and 0.577-fold of solvent control treated with 3.0 and 30 $\mu$g/ml SSE, respectively. These results were in remarkable agreement with the previously mentioned clinical HDI study that demonstrated MDZ clearance was reduced 0.52-fold across 12 healthy volunteers treated with SSE twice daily for 7 days. Similarly, for the SCE extract, which had not been evaluated clinically, intrinsic clearance of MDZ was reduced to 0.646-fold and 0.599-fold of solvent control in SCHH treated with 3.0 and 30 $\mu$g/ml SCE, respectively. Although the changes in MDZ clearance were in close agreement with the clinical studies and support this screening approach, true translation to clinical data would require additional experiments, incorporation of other organ effects, and potentially in silico modeling efforts. The inhibition mechanism studies described earlier for both *Schisandra* spp. extracts demonstrated dose-related inhibition, with maximum inhibition achieved at $\approx 30 \mu$g/ml; however, no dose-related effects were observed with MDZ intrinsic clearance. The lack of dose-dependent inhibition of MDZ intrinsic clearance following 72 hours of either *Schisandra* spp. extract treatments (e.g., 3 and 30 $\mu$g/ml) may have been due to CYP3A4 induction. In support of this conclusion, mechanic induction studies demonstrated that SSE and SCE maximally induced CYP3A4 mRNA at $\approx 30 \mu$g/ml.

SCHHs cultured under optimized conditions have active transport and adaptive response mechanisms necessary to generate physiologically relevant intracellular concentrations. Thus, intracellular concentrations were determined for three of the more abundant constituents across *Schisandra* spp. extracts in our optimized cultures of SCHH following 72 hours of treatment with SSE or SCE at 30 and 300 $\mu$g/ml. All three constituents were observed in hepatocytes treated with SCE. Schisandrin A intracellular concentrations were 2.8-fold greater in SCHH treated with SSE at the 300-$\mu$g/ml dose level than in that treated with SCE. Interestingly, and consistent with these concentration differences, SSE was demonstrated to be a $\approx 3.8$-fold more-potent inhibitor of OHMDZ formation than SCE in the mechanistic inhibition studies. Intracellular concentration differences may explain these findings. In microsomal studies evaluating individual SCE constituents, the gomisinins (A, B, G, C, and N) were reported to be more-potent inhibitors of CYP3A4 activity than schisandrin (Iwata et al., 2004). HDI assessment strategies that interrogate each component separately for interaction potential may not account for synergistic effects among constituents; hence, a more integrative approach is required to evaluate complex mixtures and reproduce net drug-interaction effects.

In summary, we demonstrated that an intrinsic clearance approach in sandwich-cultured transporter-certified human hepatocytes, with integrated metabolism and transport functionality, was feasible to evaluate HDI net effect and the relative strength of the HDI. We believe this net-effect approach is an appropriate strategy to screen complex mixtures for drug interactions. It should be noted that additive or synergistic effects between mixture components were not explicitly evaluated, and additional studies would be required to explore those effects. The approach was remarkably consistent with the in vivo HDI effect.
observed in the clinic for two herbal extracts, SSE and SJW. The evaluation of SCE predicted an in vivo HDI with MDZ similar to that observed with the closely related SSE. Kinetic parameters (Kᵢ, Kᵢ, IC₅₀, EC₅₀) from mechanistic studies have been successful in evaluating drug-interaction potential; however, kinetic parameters are difficult to directly translate to an in vivo net effect. Although our results were in good agreement with clinical findings, improvements to the study design should be considered, including the use of multiple donors for intrinsic clearance studies and developing an in vitro assay to predict herbal extract bioavailability from gut absorption. Although many drug interactions occur in the hepatocyte, the gut is an additional site of interaction that has to be considered to fully evaluate the net effect and strength of drug interactions. Here, we present a strategy to efficiently predict the potential for clinically relevant HDI by using a fully integrated in vitro system that maintains key hepatic processes important to evaluating drug interactions and can provide intracellular concentrations of constituents to further elucidate mechanistic details.

**Authorship Contributions**

**Participated in research design:** Jackson, Black, Brouwer, Roe.

**Conducted experiments:** Jackson, Freeman, Friley.

**Performed data analysis:** Jackson, Freeman, Friley, Black, Brouwer, Roe.

**Wrote or contributed to the writing of the manuscript:** Jackson, Freeman, Herman, Friley, Black, Brouwer, Roe.

**References**


