**ABSTRACT**

Venlafaxine (VEN), a first-line antidepressant, and Zuojin Pill (ZJP), a common Chinese herbal medicine consisting of *Rhizoma Coptidis* and *Fructus Evodiae*, have a high likelihood of combination usage in patients with depression with gastrointestinal complications. ZJP exhibits inhibitory effects on recombinant human cytochrome P450 isoenzymes (rhP450s), especially on CYP2D6, whereas VEN undergoes extensive metabolism by CYP2D6. From this perspective, we investigated the influence of ZJP on the metabolism of VEN in vitro and in rats for the first time. In this study, ZJP significantly inhibited the metabolism of VEN in both rat liver microsomes (RLM) and human liver microsomes (HLM); meanwhile, it inhibited the O-demethylation catalytic activity of RLM, HLM, rhCYP2D6*1/*1, and rhCYP2D6*10/*10, primarily through CYP2D6, with IC$_{50}$ values of 129.9, 30.5, 15.4, and 2.3 μg/ml, respectively. Furthermore, the inhibitory effects of ZJP on hepatic metabolism and pharmacokinetics of VEN could also be observed in the pharmacokinetic study of rats. The area under drug concentration-time curve (AUC) of VEN and its major metabolite O-desmethylvenlafaxine (ODV) increased by 39.6% and 22.8%, respectively. The hepatic exposure of ODV decreased by 57.2% 2 hours after administration ($P=0.014$). In conclusion, ZJP displayed inhibitory effects on hepatic metabolism and pharmacokinetics of VEN in vitro and in rats mainly through inhibition of CYP2D6 activity. The human pharmacokinetic interaction between ZJP and VEN and its associated clinical significance needed to be seriously considered.

**SIGNIFICANCE STATEMENT**

Zuojin Pill, a commonly used Chinese herbal medicine, demonstrates significant inhibitory effects on hepatic metabolism and pharmacokinetics of venlafaxine in vitro and in rats mainly through suppression of CYP2D6 activity. The human pharmacokinetic interaction between Zuojin Pill and venlafaxine and its associated clinical significance needs to be seriously considered.

---

**Introduction**

Venlafaxine (VEN), a selective serotonin and noradrenaline reuptake inhibitor, is one of the most efficacious and commonly prescribed antidepressants (Cipriani et al., 2018). VEN undergoes extensive metabolism by cytochrome P450 (CYP450) isoenzymes. Its major metabolite O-desmethylvenlafaxine (ODV) is predominantly catalyzed by CYP2D6, which has similar efficacy as VEN, whereas minor metabolites with little pharmacological effect, including N-desmethylvenlafaxine (NDV) and N, O-didesmethylvenlafaxine (NODV), are catalyzed by CYP2C9, CYP2C19, and CYP3A4 (Fig. 1) (Otton et al., 1996; Fogelman et al., 1999). The blood (including plasma) concentrations of VEN + ODV (active moiety) are considered clinically relevant to a relatively narrow therapeutic reference range (100–400 ng/ml) (Hiemke et al., 2018). The factors that may influence the pharmacokinetics of VEN, such as age, disease state, genetic polymorphism of metabolic enzymes, and drug combination, should also be considered seriously (Magalhães et al., 2015).

Recently, increasing evidence has suggested that concomitant administration of CYP2D6 inhibitors may incur pharmacokinetic drug-drug interaction (DDI) of VEN (Jiang et al., 2015; Magalhães et al., 2015).

Zuojin Pill (ZJP) is a traditional Chinese herbal formula recorded in the Chinese Pharmacopoeia for treating gastrointestinal disorders (Chinese Pharmacopoeia Commission, 2015). It consists of two commonly used herbs, *Rhizoma Coptidis* and *Fructus Evodiae*, following a 6:1 (w/w) ratio. The primary active ingredients in ZJP are supposed to be alkaloid compounds such as *Rhizoma Coptidis* alkaloids berberine and coptisine and *Evodiae* alkaloids evodiamine and rutaecarpine (Gao et al., 2010; Wang et al., 2013). In previous studies, ZJP
and its bioactive components, such as berberine and coptisine, exhibited inhibitory effects on CYP2D6, CYP1A2, and CYP3A4 in vitro (Han et al., 2011; Liu et al., 2014). Moreover, when ZJP is coadministered with dextromethorphan, a specific CYP2D6 probe substrate, the area under drug metabolism-time curve (AUC)$_{0-24}$ hour of dextromethorphan in healthy CYP2D6*1/*1 carriers increases 3.0-fold (Qiu et al., 2016). Because of the concomitant symptoms of depression itself and the adverse reactions of antidepressants, patients with depression are usually accompanied by several gastrointestinal disorders, and there exists a high likelihood of the combination usage of ZJP and VEN in patients with depression with gastrointestinal complications (Qiu et al., 2015; Wang et al., 2020).

Herbs and natural products (including botanical dietary supplements and foods) can produce clinically significant pharmacokinetic interactions with conventional drugs (Johnson et al., 2018). Abnormally increased or decreased drug exposure caused by herbs and natural products will generate potential risks in clinical treatment. In the classic cases, St. John’s wort and grapefruit juice can significantly influence the pharmacokinetics of cyclosporine and felodipine, respectively, which will further interfere with the therapeutic efficacy of both drugs (Paine and Oberlies, 2007; Niculusi et al., 2020).

Therefore, in the present study, we investigated the influence of ZJP on the metabolism of VEN in human liver microsomes (HLM), rat liver microsomes (RLM), and recombinant human P450 enzymes (rhP450s). Moreover, the potential pharmacokinetic interaction between ZJP and VEN was evaluated by assessing the effects of ZJP on VEN and its metabolites in rats.

Materials and Methods

Chemicals and Materials. Venlafaxine hydrochloride (purity >99%) and ODV (purity >99%) were obtained from Selleck Chemicals (Houston, TX). NDV (purity >98%) and NODV (purity >98%) were purchased from Toronto Research Chemicals (Toronto, Canada). Berberine hydrochloride (purity ≥95%), coptisine hydrochloride (purity ≥98%), evodiamine (purity ≥99%), and rutaecarpine (purity ≥98%) were obtained from Aladdin (Shanghai, China). The pooled liver microsomes of rats (a total of 75 rats of the same sex) and humans (a total of 25 adult donors of the same sex) were obtained from the Research Institute for Liver Diseases Co., Ltd. (Shanghai, China) and BiodVT (Westbury, NY), respectively. Recombinant human CYP2D6*1 and CYP2D6*10 expressed in Escherichia coli were obtained from Cyrex Ltd. (Scotland, UK). Reduced NADPH was obtained from Solarbio (Beijing, China). Diphenhydramine was purchased from Sigma-Aldrich (St. Louis, MO). All other reagents were of the highest quality commercially available.

Zuojin Pill (ZJP, batch number: 20180304) was purchased from Hubei Xianglian pharmaceutical Co., Ltd. (Wuhan, Hubei, China). The ZJP dosing solutions were prepared as follows: briefly, Zuojin Pill was ground into powder and dissolved in DMSO for in vitro study, whereas for in vivo study, 10-fold volumes of deionized water were subsequently added. The method for determining the concentration of major alkaloids in ZJP solution was similar to the liquid chromatography–tandem mass spectrometry (LC-MS/MS) method in a biologic matrix described below. The amounts of four major alkaloids in ZJP, namely berberine, coptisine, evodiamine, and rutaecarpine, were calculated as 25.5, 3.1, 0.31, and 0.48 mg/g, respectively (Supplemental Table 1).

Animals. Wistar rats (male, 180–220 g) were purchased from Charles River laboratory animal Co., Ltd. (Beijing, China). Rats were housed in a room at 16–26°C, with a light/dark cycle of 12/12 hours and humidity of 40–70%. All animal experiments were approved by the Animal Care and Use Committee of Shanghai University of Traditional Chinese Medicine and followed the Guidance for the Care and Use of Laboratory Animals in China.

Metabolic Stability Assessment of VEN in HLM and RLM. The total volume of the incubation mixture was 200 μl, which included VEN (1 μM), HLM (1.0 mg/ml) or RLM (0.5 mg/ml), and NADPH (10 mM), in 100 mM potassium phosphate-buffered solution (PBS) (pH 7.4). The reaction was conducted in a shaking water bath at 37°C for 60 minutes. To quench the reaction, 200 μl of ice-cold methanol containing diphenhydramine (100 ng/ml) as an internal standard was added. The mixture was vortexed for 1 minute and centrifuged at 15,000×g for 5 minutes. Finally, the supernatant was collected and injected into a high-performance LC-MS/MS analysis. The VEN depletion data were used for calculating enzymatic parameters.

Enzymatic Kinetics Assessment of VEN in RLM, HLM, and rhP450s. Preliminary experiments were conducted to determine the optimal conditions for linear product formation, such as protein concentration and incubation time with RLM, HLM, and rhCYPs, respectively. Subsequent experiments were conducted under the conditions of linear product formation. For both HLM and RLM experiments, the reaction mixture containing 0.2 mg/ml of microsomes and VEN (0.2–750 μM) in 100 mM PBS buffer (pH 7.4) was preincubated at 37°C for 5 minutes. Then, the mixture was incubated with 10 mM NADPH at 37°C for 15 minutes. For rhCYP2D6*1*1 or *10*10, the reaction mixture containing 8 pmol/ml of each enzyme and VEN (0.04–150 μM for CYP2D6*1*1 and 2–1500 μM for CYP2D6*10*10) was preincubated at 37°C for 5 minutes, followed by another 15-minute incubation after the addition of 10 mM NADPH. As described above, all reactions were quenched and treated with ice-cold methanol, and the supernatant was collected for LC-MS/MS analysis. The ODV formation data were used for calculating enzymatic parameters.

Evaluation of the Influences of ZJP and Its Major Components on the Metabolism of VEN In Vitro. The influences of ZJP and its major components,
administration, after 12 hours of fasting but free access to water, the rats in ZJP
dissolved in saline) once daily for 9 consecutive days. On the 9th day of
rat was intragastrically administered normal saline or ZJP solution (2.52 g/kg,
after administration, rats were sacrificed, and liver samples were collected immediately
and ZJP group. Twelve hours before the experiment, all rats were forbidden from
allocated into two groups (n = 6 in each group), which were blank control group and ZJP group. Each
rat was intragastrically administered normal saline or ZJP solution (2.52 g/kg,
dissolved in saline) once daily for 9 consecutive days. On the 9th day of
administration, after 12 hours of fasting but free access to water, the rats in ZJP
group received VEN (2.63 mg/kg) as well as ZJP, whereas the rats in blank control
group received the same volume of normal saline. Blood samples were collected in
heparinized tubes at 0, 0.5, 1, 1.5, 2, 3, 4, 6, 8, 12, 24, 36, and 48 hours after
VEN administration and then centrifuged at 4000 g 4°C for 10 minutes, and
the plasma was collected and stored at –80°C for LC-MS/MS analysis.

**Determination of Major Bioactive Components of ZJP in Rat Livers.**
Four major alkaloids (berberine, coptisine, evodiamine, and rutacarpine) of ZJP
in rat livers were quantitatively determined by a validated LC-MS/MS method.
The LC-MS/MS system contained an LC20AD liquid chromatography (Shi-
madzu, Kyoto, Japan) and an API 4000 QTRAP triple quadrupole mass
spectrometer (Applied Biosystems, Foster City, CA) equipped with an electrospray ionization source. Chromatographic separation was performed on an
Agilent Eclipse XDB C18 reversed-phase column (150 × 4.6 mm, 5 μm) at
room temperature. The gradient of the mobile phase consisting of water
(0.08% formic acid and 4 mM ammonium acetate) (A) and acetonitrile (B) with
a flow rate of 0.8 ml/min was set as follows: 0–3 minutes, 65% A; 3–4.5 minutes,
65%–20% A; 4.5–9 minutes, 20% A; 9–11 minutes, 20%–65% A; and 11–15
minutes, 65% A. Positive ionization mode and multiple reaction monitoring mode
were selected for quantification. The parameters of mass spectrometry for each
analyte were shown in Supplemental Table 2. The intra- and interday precision
and accuracy of the quality control samples for each analyte were<15% (Supplemental Table 3). All analytes were stable in the measurement circumstances (Supplemental Table 4).

**Determination of VEN and Its Three Metabolites in In Vitro and In Vivo
Samples.** VEN and its three metabolites (ODV, NDV, and NODV) from in vitro
samples, such as HLMs, RLMs, or rhCYP2D6 reaction samples, and in vivo
samples, such as rat plasma and liver samples, were quantitatively determined
by a validated LC-MS/MS method. The LC-MS/MS system contained an Agilent-
1260 liquid chromatograph (Agilent Technologies, Santa Clara, CA) and an API
4000 triple quadrupole mass spectrometer (Applied Biosystems) equipped with an
electrospray ionization source. Chromatographic separation was performed on a
Phenomenex Kinetex XB-C18 reversed-phase column (100 × 2.1 mm, 2.6 μm) at
40°C. The gradient of the mobile phase consisting of water (0.1% formic acid and
5 mM ammonium acetate) (A) and methanol (B) with a flow rate of 0.5 ml/min
was set as follows: 0–1 minutes, 60% A; 1–4 minutes, 60%–20% A; 4.5–5.5
minutes, 20% A; 5.5–6.5 minutes, 20%–60% A; and 5.6–8 minutes, 60% A. Positive
ionization mode and multiple reaction monitoring mode were selected for
quantification. The parameters of mass spectrometry for each analyte were shown
in Table 1. Linear calibration curves for VEN, ODV, NDV, and NODV were
constructed in relative matrices with concentrations ranging 2.5–2000, 0.35–280,

### Table 1

<table>
<thead>
<tr>
<th>Analytes</th>
<th>Transition (m/z)</th>
<th>Spray Voltage (V)</th>
<th>Temp. (°C)</th>
<th>DP (V)</th>
<th>CE (V)</th>
<th>CXP (V)</th>
<th>EP (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td>VEN</td>
<td>278.2 &gt; 58.1</td>
<td>5500</td>
<td>400</td>
<td>46</td>
<td>46</td>
<td>12</td>
<td>10</td>
</tr>
<tr>
<td>ODV</td>
<td>264.2 &gt; 58.1</td>
<td>5500</td>
<td>400</td>
<td>46</td>
<td>46</td>
<td>12</td>
<td>10</td>
</tr>
<tr>
<td>NDV</td>
<td>264.2 &gt; 246.1</td>
<td>5500</td>
<td>400</td>
<td>35</td>
<td>14</td>
<td>12</td>
<td>10</td>
</tr>
<tr>
<td>NODV</td>
<td>250.2 &gt; 232.1</td>
<td>5500</td>
<td>400</td>
<td>30</td>
<td>14</td>
<td>12</td>
<td>10</td>
</tr>
<tr>
<td>Diphenhydramine</td>
<td>256.0 &gt; 167.1</td>
<td>5500</td>
<td>400</td>
<td>46</td>
<td>25</td>
<td>12</td>
<td>10</td>
</tr>
</tbody>
</table>

| Temp. | DP, declustering potential. CE, collision energy. CXP, CXP, cell exit potential. EP, entrance potential. |  |  |  |  |  |  |

**Fig. 2.** The remaining VEN and generated demethylation metabolites, ODV, NDV, and NODV, after 60-minute incubation of VEN in HLM (A) or RLM (B). Data for each component were normalized to spiked VEN concentration in reaction mixtures and represented as means ± S.D. in triplicates.
Results

Metabolic Stability of VEN in HLM and RLM. After a 60-minute incubation, the remaining VEN in RLM and HLM were 15.6% and 78.3%, respectively, indicating a faster phase I metabolism of VEN in rats than in humans (Fig. 2). In both RLM and HLM, ODV was the primary metabolite (30.9% vs. 31.0%), whereas NDV was 6.3-fold higher in RLM than HLM (14.1% vs. 2.2%), indicating a species difference in the metabolism of VEN.

Enzymatic kinetics of VEN in HLM, RLM, and rhP450s. Because ODV was the dominant metabolite of VEN, we explored its generating rate to reflect the enzymatic kinetics of VEN in RLM, HLM, and rhP450s, respectively (Fig. 3; Table 2). In the RLM and HLM experiments, although $K_m$ was very similar, the $V_{max}$ of RLM was 6.6-fold higher than that of HLM. Hence, there was a 2.1-fold increase comparing $CL_{int}$ of VEN in RLM to that in HLM (Fig. 3, A and B).

rhP450 experiments, the two rhCYP2D6s with different alleles, *1/*1 and *10/*10, showed similar $V_{max}$ values (0.014 vs. 0.013 nmol/min per picomoles P450), whereas the CYP2D6*10/*10 showed a 13.1-fold higher $K_m$ value than that of CYP2D6 *1/*1, resulting in a 14.0-fold lower $CL_{int}$ than CYP2D6*1/*1 (Fig. 3, C and D).

Influence of ZJP and Its Major Components on the Metabolism of VEN in HLM and RLM. To explore whether ZJP could influence the demethylation process of VEN, ZJP and its major components, berberine and coptisine, were incubated with VEN (1 μM) in HLM and RLM, respectively (Fig. 4). The results showed that ZJP significantly inhibited VEN depletion (94.5%) and the formation of its demethylation products, ODV, NDV, and NODV (91.0%, 69.4%, and 66.5%, respectively), compared with the control group in HLM (all P values <0.001) (Fig. 4, A–D). Coptisine and berberine demonstrated a similar tendency as ZJP. CYP2D6 was primarily responsible for the metabolism of VEN and the generation of ODV in HLM, which were inhibited by QND at inhibitory degrees of 96.5% and 95.6%, respectively. Meanwhile, NDV increased by 33.7% compared with the control group after QND treatment. In contrast, KTZ significantly inhibited the production of NDV and NODV (both P <0.001) but without obvious effect on ODV (P = 0.557).

Similarly, the VEN depletion and formation of ODV and NDV significantly decreased after ZJP (48.3%, 33.0%, and 78.9%, respectively) and coptisine (19.8%, 13.5%, and 53.8%, respectively) treatment in RLM (all P values <0.05) (Fig. 4, E–H). Meanwhile, NDV increased by 73.5%, 66.8%, and 86.3% in the presence of ZJP, coptisine, and berberine, respectively. Interestingly, ZJP exhibited very similar tendencies in either VEN depletion or its metabolite generation as that of QND, a specific inhibitor of rat CYP2D2 and KTZ, indicating that CYP3A4 could also play an important role in the metabolism of VEN in rats.

IC$_{50}$ of ZJP and Its Major Components on the Metabolism of VEN in HLM, RLM, and rhCYP2D6s. To further disclose the inhibitory potency of ZJP and its major bioactive components on the metabolism of VEN, the inhibitory effects, or IC$_{50}$, of ZJP, coptisine, or berberine on ODV production in microsomes or on rhCYP2D6s were investigated subsequently (Fig. 5). The individual IC$_{50}$ values of ZJP, berberine, and coptisine on the metabolism of VEN are shown in Table 2.

<table>
<thead>
<tr>
<th>Subject</th>
<th>$K_m$ (μM)</th>
<th>$V_{max}$ (nmol/min per Milligram Protein or nmol/min per Picomoles P450)</th>
<th>$CL_{int}$ (μl/min per Milligram Protein or μl/min per Picomoles P450)</th>
</tr>
</thead>
<tbody>
<tr>
<td>RLM</td>
<td>41.2 ± 23.4</td>
<td>0.480 ± 0.013</td>
<td>11.7 ± 5.5</td>
</tr>
<tr>
<td>HLM</td>
<td>39.9 ± 1.1</td>
<td>0.182 ± 0.004</td>
<td>5.7 ± 0.2</td>
</tr>
<tr>
<td>rhCYP2D6*1/ *1</td>
<td>11.3 ± 3.2</td>
<td>0.014 ± 0.003</td>
<td>1.2 ± 0.3</td>
</tr>
<tr>
<td>rhCYP2D6*10/ *10</td>
<td>148.2 ± 26.5</td>
<td>0.013 ± 0.002</td>
<td>0.088 ± 0.016</td>
</tr>
</tbody>
</table>
Table 3. Those results showed that ZJP, coptisine, and berberine could inhibit the O-demethylation of VEN while showing little effect (IC$_{50}$ > 280 μg/ml or 100 μM) on the N-demethylation process of VEN. In addition, ZJP and its two bioactive components exhibited higher inhibitory potency in HLM compared with RLM. Meanwhile, coptisine was more potent than berberine in inhibiting the formation of ODV and NODV, as it displayed lower IC$_{50}$ values both in HLM and RLM (7.5% and 41.9%, respectively) than those of berberine. Both coptisine and berberine were potent inhibitors of rhCYP2D6*1/*1 and rhCYP2D6*10/*10, as IC$_{50}$ values did not exceed 2.5 μM. As NDV and NODV were not

Fig. 4. Inhibitory effects of ZJP (150 μg/ml), coptisine (CPS, 30 μM), berberine (BBR, 30 μM), quinine (QNN, 2 μM), quinidine (QND, 2 μM), and ketoconazole (KTZ, 2 μM) on the metabolism of VEN and formation of ODV, NDV, and NODV in HLM (A–D) and RLM (E–H). Each column represented the remaining VEN (% control) or product formation (% spiked VEN). Data were presented as means ± S.D. in triplicates. *P < 0.05 compared with the control. Cntl, control.

Fig. 5. Inhibitory effects (IC$_{50}$) of ZJP (A–D), coptisine (E–H), and berberine (I–L) on ODV formation in HLM, RLM, CYP2D6*1/*1, and CYP2D6*10/*10, respectively. Data were presented as means ± S.D. in triplicates. The curve represented the fitting of the observed ODV formation rate (% control) (y) vs. the inhibitor concentration (x). Conc., concentration.
TABLE 3

<table>
<thead>
<tr>
<th>IC&lt;sub&gt;50&lt;/sub&gt; of ZJP, berberine, and coptisine toward VEN metabolism in microsomes and rhCYP2D6</th>
<th>Inhibitors</th>
<th>ODV</th>
<th>IC&lt;sub&gt;50&lt;/sub&gt; value</th>
<th>NDV</th>
<th>NODV</th>
</tr>
</thead>
<tbody>
<tr>
<td>RLM</td>
<td>ZJP (µg/ml)</td>
<td>129.9 ± 2.6</td>
<td>&gt;575</td>
<td>42.8 ± 8.0</td>
<td></td>
</tr>
<tr>
<td>Berberine (µM)</td>
<td>64.5 ± 11.5</td>
<td>&gt;1000</td>
<td>59.5 ± 18.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coptisine (µM)</td>
<td>27.0 ± 6.2</td>
<td>&gt;200</td>
<td>16.9 ± 0.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HLM</td>
<td>ZJP (µg/ml)</td>
<td>30.5 ± 5.5</td>
<td>&gt;288</td>
<td>11.5 ± 10.2</td>
<td></td>
</tr>
<tr>
<td>Berberine (µM)</td>
<td>30.5 ± 9.6</td>
<td>&gt;500</td>
<td>&gt;500</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coptisine (µM)</td>
<td>2.3 ± 1.3</td>
<td>&gt;100</td>
<td>&gt;100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>rhCYP2D6*1/*1</td>
<td>ZJP (µg/ml)</td>
<td>15.4 ± 4.5</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
</tr>
<tr>
<td>Berberine (µM)</td>
<td>2.5 ± 0.9</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coptisine (µM)</td>
<td>0.7 ± 0.2</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
<td></td>
</tr>
<tr>
<td>rhCYP2D6*10/*10</td>
<td>ZJP (µg/ml)</td>
<td>2.3 ± 1.0</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
</tr>
<tr>
<td>Berberine (µM)</td>
<td>1.5 ± 0.2</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coptisine (µM)</td>
<td>2.2 ± 0.2</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

N/A, not applicable.

detected in rhCYP2D6s but in microsomes, other P450s than CYP2D6 were responsible for the N-demethylation process of VEN.

Inhibitory Effects of ZJP on the Metabolism of VEN and Pharmacokinetics in Rats. VEN and its major metabolites, ODV and NDV, were detected in rat plasma after intragastric administration of VEN alone or coadministration with ZJP (VEN+ZJP). The mean plasma concentration-time curves are plotted in Fig. 6. The pharmacokinetic parameters of VEN and its metabolites were presented in Table 4. Compared with the VEN group, the AUC<sub>0–24</sub> for VEN and ODV increased obviously by 39.6% and 22.8%, respectively, in the VEN+ZJP group (Fig. 7). The ratio of ODV/VEN in the liver decreased significantly by 57.2% (P = 0.014), whereas NDV exposure increased significantly by 72.7% in VEN+ZJP group (P = 0.014). Meanwhile, the ratio of ODV/VEN in the liver also decreased significantly by 37.7% (P = 0.018) upon ZJP administration.

Discussion

Depression is the leading cause of psychiatric disorders, with an estimated 350 million people affected worldwide (Cipriani et al., 2018). Depression can be long-lasting or recurrent, and it can dramatically affect people’s ability to live a rewarding life. VEN, approved in 1995, is a classic antidepressant, but it still stands as one of the most efficacious and prescribed antidepressants, particularly for the treatment of selective serotonin reuptake inhibitor-resistant depression (Cipriani et al., 2018). VEN undergoes extensive phase I metabolism and is primarily catalyzed by CYP2D6 to generate an active metabolite, ODV. The recommended therapeutic reference range for VEN is 100–400 ng/ml of active moiety, which is calculated from the total amount of VEN and ODV (Hiemke et al., 2018). The metabolic ratio ODV/VEN is usually used as a tool to distinguish patients as poor metabolizers or extensive metabolizers of CYP2D6 (Lobello et al., 2010). When VEN is coadministered with a CYP2D6 inhibitor such as paroxetine, both C<sub>max</sub> and AUC of VEN are significantly elevated in patients, which might explain VEN responses and adverse events (Jiang et al., 2015). Although the discrepancy on the clinical implication of exposure changes to VEN and/or ODV is still great, knowledge of the mechanisms of potential DDI between VEN and a drug with CYP2D6 inhibition is necessary for assessing and minimizing clinical risks (Schoretsanitis et al., 2019).

Traditional Chinese medicines (TCMs) belong to herbal medicines and have a long history of usage in clinical practice. In Asia, TCMs are prescribed concomitantly with conventional medications (prescription and nonprescription) for treatments. However, the risk assessment of the pharmacokinetic interaction between TCMs and conventional drugs is insufficient, leading to safety and efficacy issues in contemporary pharmacotherapy. ZJP is a widely used traditional Chinese herbal formula recorded in the Chinese Pharmacopoeia for routine treatments of gastrointestinal disorders (Chinese Pharmacopoeia Commission, 2015). It is commonly prescribed to patients with indications of gastritis, such as gastric ulcer, pyloric obstruction, gastroesophageal reflux disease, etc. It is also beneficial for depression with maladjusted gastrointestinal function (Wang et al., 2020). ZJP consists of two commonly used herbs, *Rhizoma Coptidis* and *Fructus Evodiae*, following a 6:1 (w/w) ratio. *Rhizoma Coptidis* is reported to inhibit
CYP2D6 activity with an IC$_{50}$ value of 5.8 µg/ml of its extracted powder (approximately equal to 30.4 µg/ml of crude herbs) in HLM (Han et al., 2011). The inhibitory effect might be mostly attributable to Coptis alkaloid components, especially coptisine, with an IC$_{50}$ value of 4.4 µM (Han et al., 2011). Moreover, when ZJP is coadministered with dextromethorphan, a known CYP2D6 substrate, the AUC$_{0–24}$ hour of dextromethorphan in healthy volunteers with dominant CYP2D6 phenotype (CYP2D6*1/*1) increases 3.0-fold higher than those administered dextromethorphan only (Qiu et al., 2016). With the high likelihood of combination usage of the two conventional prescriptions ZJP and VEN for treating depression patients with gastrointestinal disorders in China and potential herb-drug interaction (Qiu et al., 2015; Wang et al., 2020), we investigated the pharmacokinetic influences of ZJP on VEN for the first time. In this study, we found that ZJP significantly inhibits the metabolism of VEN in vitro. After coincubation of VEN and ZJP for 1 hour, the VEN depletion and ODV formation significantly decreased by 94.5% and 91.0%, respectively, compared with the blank group in RLM. It was supported by some DDI data in rats. In addition, it was worth mentioning that because of focuses on the plasma exposure of VEN (between 100 and 400 ng/ml) in clinical practice that was far below the $K_m$ values obtained both in HLM and RLM, we selected a clinically relevant VEN concentration, e.g., 0.5 or 1 µM, for IC$_{50}$ experiments.

Species difference is probably the reason for weak evidence of DDI potential between ZJP and VEN in rats. CYP2D isoforms between rats and humans share a good homology (>70%) (Venhorst et al., 2003). Rat CYP2D1 is known as an ortholog (approximately 83% homology) of human CYP2D6 (Martignoni et al., 2006). However, when comparing the metabolic capability of VEN, the two species are obviously different. In literatures, the C$_{max}$ and AUC$_{0–24}$ of VEN in poor metabolizers increased by approximately 70%–80% and 200%–300%, respectively, compared with those of extensive metabolizers (Lessard et al., 1999; Lobello et al., 2010). In contrast, the C$_{max}$ and AUC$_{0–24}$ of VEN in CYP2D1-null rats only increased by 24% and 59%, respectively, compared with those of wild-type rats, indicating an apparently weaker inhibitory effect involved in CYP2D6 inhibition in rats than humans (Zhou et al., 2019). Similarly, we observed that the capability and magnitude of inhibitory effects of ZJP were not comparable between HLM and RLM. First, the metabolic characteristics of VEN were different between the two species. CYP2D6 was the sole principle P450 involved in CYP2D6 inhibition in rats than humans compared with those of wild-type rats, indicating an apparently weaker inhibitory effect involved in CYP2D6 inhibition in rats than humans (Zhou et al., 2019). Similarly, we observed that the capability and magnitude of inhibitory effects of ZJP were not comparable between HLM and RLM. First, the metabolic characteristics of VEN were different between the two species. CYP2D6 was the sole principle P450.

**TABLE 4**

Pharmacokinetic parameters of VEN, ODV, and NDV in rats after oral administration of VEN alone or VEN combined with ZJP

<table>
<thead>
<tr>
<th>Analyte</th>
<th>Groups</th>
<th>$C_{max}$ (ng/ml)</th>
<th>$T_{max}$ (h)</th>
<th>$t_{1/2}$ (h)</th>
<th>CLz/F (l/h per kilogram)</th>
<th>AUC$_{0–24}$ (µg/l × h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>VEN</td>
<td>VEN</td>
<td>3.47 ± 0.80</td>
<td>1.0 ± 0.6</td>
<td>2.4 ± 2.2</td>
<td>231.7 ± 18.6</td>
<td>11.1 ± 0.9</td>
</tr>
<tr>
<td>VEN+ZJP</td>
<td>VEN+ZJP</td>
<td>2.01 ± 0.85*</td>
<td>3.1 ± 1.8*</td>
<td>4.0 ± 2.2</td>
<td>198.7 ± 151.3</td>
<td>15.5 ± 7.6</td>
</tr>
<tr>
<td>ODV</td>
<td>VEN</td>
<td>4.82 ± 4.18</td>
<td>1.1 ± 0.9</td>
<td>6.2 ± 3.3</td>
<td>172.7 ± 123.5</td>
<td>19.3 ± 11.3</td>
</tr>
<tr>
<td>NDV</td>
<td>VEN</td>
<td>3.41 ± 2.00</td>
<td>2.1 ± 1.4</td>
<td>10.9 ± 6.5</td>
<td>116.8 ± 66.8</td>
<td>23.7 ± 12.8</td>
</tr>
<tr>
<td></td>
<td>VEN</td>
<td>1.24 ± 0.90</td>
<td>2.0 ± 1.8</td>
<td>5.1 ± 2.9</td>
<td>468.0 ± 208.7</td>
<td>6.45 ± 4.34</td>
</tr>
</tbody>
</table>

$P < 0.05$ compared with VEN group. CLz/F: apparent oral clearance.

**Fig. 7.** Hepatic exposures of VEN (A), ODV (B), NDV (C), NODV (D), and ODV/VEN ratio (E) at 2 hours after intragastric administration of VEN alone or coadministration with ZJP (VEN+ZJP) in rats ($n = 6$). Data were presented as means ± S.D. *$P < 0.05$ compared with the VEN group. Conc., concentration.
enzyme responsible for VEN metabolism in HLM but not in RLM. After 1 hour of incubation of VEN, ODV (generated primarily by CYP2D6) was the major metabolite (31.0% of spilled VEN), and NDV (generated primarily by CYP3A4 and CYP2C19) was the minor metabolite (2.2% of spilled VEN) in HLM. In contrast, OD and NDV were both major metabolites in RLM, with 30.9% and 14.1% of formation rate percentages, respectively. Secondly, the inhibitory effects of CYP2D6 inhibitor on ODV formation in rats were weaker than those in humans. Quinidine, a known human CYP2D6 inhibitor, inhibited VEN depletion (96.5%) and ODV formation (95.6%) completely in HLM. In contrast, quinine, the quinidine enantiomer and a known rat CYP2D6 inhibitor, inhibited VEN depletion (58.4%) and ODV formation (50.6%) partly in RLM. The inhibitory efficacy (\(IC_{50}\) values obtained from ODV formation) of the potent CYP2D6 inhibitor in HLM was greater than that in RLM (10.0 μM vs. 31.5 μM) (Supplemental Fig. 1). Taken together, CYP2D6 (in fact CYP2D1 in rats) did not play a dominant role in systemic exposures for the four major alkaloid constituents, berberine, coptisine, evodiamine, and rutaecarpine in rats. A sensitive index substrate of CYP2D6, dextromethorphan, was also assessed in healthy volunteers (which contain 20 mg/kg of berberine and are coadministered drugs. The underlying mechanism may be multiple principles involving one or more processes of absorption, disposition, metabolism, and excretion. In this study, we found that the concentrations of Coptis alkaloids berberine and coptisine in liver tissue were much higher (>10.0-fold higher) than those in plasma, which was in accordance with previous literature (Liu et al., 2010). Although the concentration of single alkaloids in the rat liver (e.g., 25.5 and 3.1 mg/g for berberine and coptisine, respectively) for CYP2D6-mediated metabolism of VEN was slightly lower than the \(IC_{50}\) (e.g., 21.7 and 8.6 mg/g for berberine and coptisine, respectively), considering the structural and mechanistic similarities of the alkaloids in ZIP, whether structural and mechanistic similarities of the alkaloids in ZIP, whether structur


Address correspondence to: Bo Tan, Laboratory of Clinical Pharmacokinetics, Shuguang Hospital Affiliated to Shanghai University of TCM, 528 Zhangheng Rd., Shanghai 201203, China. E-mail: tbot@163.com; or Furong Qiu, Laboratory of Clinical Pharmacokinetics, Shuguang Hospital Affiliated to Shanghai University of TCM, 528 Zhangheng Rd., Shanghai 201203, China. E-mail: furong_qiu@126.com