When Does the Rate-Determining Step in the Hepatic Clearance of a Drug Switch from Sinusoidal Uptake to All Hepatobiliary Clearances? Implications for Predicting Drug-Drug Interactions[§]

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ABSTRACT

For dual transporter-enzyme substrate drugs, the extended clearance model can be used to predict the rate-determining step(s) (RDS) of a drug and hence predict its drug-drug interaction (DDI) liabilities (i.e., transport, metabolism, or both). If the RDS of the hepatic clearance of the drug is sinusoidal uptake clearance (CL_{in}^s), even if the drug is eliminated mainly by hepatic metabolism, its DDI liability (as viewed from changes to systemic drug concentrations) is expected to be inhibition or induction of uptake transporters but not hepatic enzymes; however, this is true only if the condition required to maintain CL_{in}^s as the RDS is maintained. Here, we illustrate through theoretical simulations that the RDS condition may be violated in the presence of a DDI. That is, the RDS of a drug can switch from CL_{in}^s to all hepatobiliary

clearances [i.e., metabolic/biliary clearance ($CL_{met} + bile$) and CL_{int}^s], leading to unexpected systemic DDIs, such as metabolic DDIs, when only transporter DDIs were anticipated. As expected, these analyses revealed that the RDS switch depends on the ratio of $CL_{met} + bile$ to sinusoidal efflux clearance (CL_{ef}^s). Additional analyses revealed that for intravenously administered drugs, the RDS switch also depends on the magnitude of CL_{in}^s . We analyzed published in vitro quantified hepatobiliary clearances and observed that most drugs have a $CL_{met} + bile/CL_{ef}^s$ ratio < 4; hence, in practice, the magnitude of CL_{in}^s must be considered when establishing the RDS. These analyses provide insights previously not appreciated and a theoretical framework to predict DDI liabilities for drugs that are dual transporter-enzyme substrates.

Introduction

Identifying liabilities with respect to drug-drug interactions (DDI) is important in drug development. In 2015, 25 of the 33 new drug applications contained in vitro transporter data, and of 20 clinical trials using the new molecular entities (NMEs) as victim drugs, only nine resulted in a significant area under the curve (AUC) change (Yu et al., 2017). These data acknowledge that drug transporters are important in determining drug disposition (Giacomini et al., 2010; Hillgren et al., 2013; Patel et al., 2016).

As shown by the hepatic extended clearance model (ECM), when a drug is both transported into and metabolized or biliary-excreted by the liver, the rate-determining step (RDS) in the systemic clearance of the drug can be its hepatic uptake clearance, metabolic clearance, biliary (canalicular efflux) clearance, or all hepatobiliary clearances (Miyauchi et al., 1987; Sirianni and Pang, 1997; Shitara et al., 2006; Kusuhara and Sugiyama, 2009; Li et al., 2014; Patilea-Vrana and Unadkat, 2016). The RDS of a drug can be identified using models such as the Extended Clearance Concept Classification System and the Extended Clearance Classification System, which use the drug's in vitro quantified hepatobiliary clearance values or the drug's physicochemical properties,

respectively (Camenisch and Umehara, 2012; Varma et al., 2015). Using such models is advantageous since the RDS of a drug helps identify where the DDI liabilities lie. Of note, unless indicated otherwise, all subsequent references to DDI should be interpreted as those DDIs that can be observed from measurement of the systemic concentrations of the victim drug. For example, if the RDS of a drug is its hepatic uptake clearance (RDS_{uptake}), then the focus of the DDI studies should be transporter-based [e.g., hepatic organic anion-transporting polypeptide (OATP)-mediated uptake of atorvastatin] (Maeda et al., 2011) or if the RDS is both hepatic uptake and metabolic or biliary clearance (RDS_{all}), the focus of DDI studies should be all hepatobiliary pathways (e.g., OATP and cytochrome P450-mediated clearance of cerivastatin) (Mück et al., 1999; Backman et al., 2002).

Here, we asked whether knowledge of the RDS of a drug is enough to predict DDI liabilities for drugs that are dual transporter-enzyme substrates. If not, the focus of DDI studies will be misdirected and will result in either a negative or unexpected DDI and therefore toxicity. Under the worst-case scenario, the latter will lead to discontinuation of drug development and the end result is that both outcomes will increase drug development cost (Paul et al., 2010). For these reasons, it is important to ask whether the RDS can switch from hepatic uptake clearance to all hepatobiliary clearance pathways thus resulting in unexpected systemic DDIs. Using the ECM theory and simulations, we aimed to: 1) provide a theoretical framework of when the RDS_{uptake} switches to RDS_{all} in the presence of a DDI and 2) apply the RDS

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ABBREVIATIONS: AUC, area under the curve; AUCR, area under the curve ratio; CL_{bile} , biliary (canalicular) efflux clearance; CL_{int} , intrinsic clearance; CL_{met} , metabolic clearance; CL_{in}^s , sinusoidal influx clearance; CL_{ef}^s , sinusoidal efflux clearance; DDI, drug-drug interaction; ECM, extended clearance model; IVIVE, in vitro to in vivo extrapolation; NME, new molecular entity; OATP, organic anion transporting polypeptide; PBPK, physiologically based pharmacokinetics; $PI_{met + bile}$, percent inhibition of $CL_{met} + bile$ necessary for RDS_{uptake} to switch to RDS_{all} ; RDS, rate-determining step.

framework to predict DDI liabilities through theoretical and practical examples. The resulting analyses and simulations provide novel insights, hitherto not appreciated, into factors that determine when a victim drug experiences the RDS_{uptake} switch to RDS_{all} and elucidate important considerations for predicting DDI liabilities for drugs that are substrates of both hepatic transporters and enzymes.

Materials and Methods

Theoretical Background. The ECM describes complex hepatobiliary clearance in terms of transport at the sinusoidal membrane via sinusoidal influx (CL_{in}^s) and efflux (CL_{ef}^s), transport at the canalicular membrane via biliary efflux (CL_{bile}), metabolism (CL_{met}), hepatic blood flow (Q_h), and fraction unbound in blood (f_u) (eq. 1). CL_{in}^s and CL_{ef}^s terms incorporate both transport-mediated plus passive diffusion clearance, whereas CL_{bile} describes active transport only. The interrelationships between the hepatobiliary clearances defined by the ECM create the RDS in the hepatic clearance of a drug. As described by us and others (Miyauchi et al., 1987; Sirianni and Pang, 1997; Shitara et al., 2006; Patilea-Vrana and Unadkat, 2016), these can be 1) $RDS_{\text{met} + \text{bile}}$ when the metabolic and biliary efflux clearances of the drug are much less than sinusoidal efflux clearance ($CL_{\text{met} + \text{bile}} < CL_{\text{ef}}^s$) and the drug is highly permeable (passive diffusion) $>>$ active transport, $CL_{\text{in}}^s \approx CL_{\text{ef}}^s$ and thus can rapidly distribute across the sinusoidal membrane; 2) RDS_{uptake} when the metabolic plus biliary efflux clearances are much greater than the sinusoidal efflux clearance ($CL_{\text{met} + \text{bile}} >> CL_{\text{ef}}^s$), or 3) RDS_{all} when a drug has both active transport and metabolism, but the preceding two extreme scenarios do not apply ($CL_{\text{in}}^s \neq CL_{\text{ef}}^s$):

$$CL_h = \frac{Q_h f_u CL_{\text{in}}^s (CL_{\text{met}} + CL_{\text{bile}})}{Q_h (CL_{\text{ef}}^s + CL_{\text{met}} + CL_{\text{bile}}) + f_u CL_{\text{in}}^s (CL_{\text{met}} + CL_{\text{bile}})} \quad (1)$$

Identifying the RDS of a drug can be used to predict the liability of transporter versus metabolic DDIs (see Patilea-Vrana and Unadkat, 2016, for simulations of systemic and hepatic AUC when hepatobiliary clearances are inhibited). For example, although a victim drug has RDS_{uptake} , inhibition of $CL_{\text{met} + \text{bile}}$ will not result in a significant increase in the systemic AUC, even though such DDI could result in significant drug accumulation in the liver and hence potentially enhance the hepatic efficacy or toxicity of the drug. That is, from the point of view of systemic (e.g., victim plasma concentrations) measurements, inhibition of $CL_{\text{met} + \text{bile}}$ will be incorrectly interpreted as negative because there will be no change in systemic concentrations of the drug. On the other hand, inhibition of CL_{in}^s will result in an increase in the drug's systemic AUC (and therefore potentially nonhepatic efficacy and toxicity of the drug) but will result in no changes in the hepatic AUC, provided the liver is the primary eliminating organ (for examples, see Patilea-Vrana and Unadkat, 2016). Less appreciated, however, is the fact that in the presence of metabolic or biliary efflux DDI, the RDS of a drug can switch from RDS_{uptake} to RDS_{all} and hence switch the DDI liability from uptake transporters to both metabolic or biliary and uptake pathways. Consequently, the drug's systemic AUC will significantly change owing to metabolic and biliary efflux DDIs, even though uptake was the RDS of the drug in the absence of a DDI, which would lead to unexpected DDIs as viewed from the systemic concentrations of the victim drug. Therefore, through MATLAB simulations (R2016a; MathWorks, Natick, MA), we illustrated when the RDS_{uptake} -to- RDS_{all} switch occurs for a victim drug in the presence of a DDI. We then applied our proposed RDS framework to published in vitro hepatobiliary clearances to determine whether in vivo observed DDI liabilities can be correctly predicted. Although the insights illustrated can be derived from analytical solutions of the ECM equation (eq. 1), for clarity, we chose to use simulations to illustrate the principles of these DDI liabilities within the RDS framework.

Simulation Assumptions. The hepatic ECM was simulated using the governing differential equations as previously described (Endres et al., 2009; Patilea-Vrana and Unadkat, 2016), and for simplicity, the following assumptions about the victim drug were made: 1) it was administered intravenously; 2) the fraction unbound (f_u) in blood and tissue (liver) was set to 1; 3) liver was the only eliminating organ; 4) Q_h was set to 1 liter/min. All references to systemic AUC are derived from drug concentrations in blood. Our conclusions regarding the RDS switch are generalizable to when victim drugs are administered orally, but our conclusions of the RDS dependence on CL_{in}^s apply only to intravenously

administered drugs (see text to follow). Furthermore, for oral drug administration, our findings apply only to changes to the hepatic clearance and bioavailability of the victim drug and do not address the intestinal availability of the victim drug. If there is significant nonhepatic clearance, our conclusions will stand except that the magnitude of the change observed in the systemic and/or hepatic AUC of the drug will differ (Patilea-Vrana and Unadkat, 2016).

Identifying When the RDS_{uptake} Switches to RDS_{all} and Factors that Influence this Switch. First, we determined when the RDS of a drug switches from uptake clearance to all hepatobiliary clearance pathways. This requires violating the condition $CL_{\text{met} + \text{bile}} >> CL_{\text{ef}}^s$, the condition necessary for uptake clearance to be the RDS in the hepatic clearance of drug. To illustrate this effect, for three theoretical victim drugs, where $CL_{\text{met} + \text{bile}} >> CL_{\text{ef}}^s$ ($CL_{\text{met} + \text{bile}} = 1, 10, 100$ liters/min, $CL_{\text{ef}}^s = 0.1$ liter/min, and $CL_{\text{in}}^s = 1Q_h$), the systemic AUC ratio (AUCR) of the victim drug in the absence and presence of 10%–99% inhibition of $CL_{\text{met} + \text{bile}}$ was simulated. In accordance with Food and Drug Administration guidelines, an AUCR of 1.25 was considered significant.

To illustrate that the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio, and not the absolute magnitude of $CL_{\text{met} + \text{bile}}$ and/or CL_{ef}^s , determines when the RDS_{uptake} switches to RDS_{all} , we conducted the following simulations: the systemic AUC of the drug was simulated for CL_{ef}^s values ranging from 0.1 to 10 liters/min (representing $0.1 \times$ to $10 \times Q_h$) with $CL_{\text{met} + \text{bile}}$ set to 1- to 20-fold the value of the corresponding CL_{ef}^s . The simulated systemic AUCs, when the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio was held constant, were compared with the simulated systemic AUCs when $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio varied.

Next, we defined the tipping point as the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio, at which RDS_{uptake} switches to RDS_{all} . Following the same strategy, we simulated the AUCR for various $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratios for victim drugs that originally had RDS_{uptake} to illustrate the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio at which AUCR = 1.25, thus signifying that RDS_{uptake} switched to RDS_{all} . The systemic AUC where the RDS is uptake was simulated such that the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio = 1000 (AUC_{ratio = control}: $CL_{\text{met} + \text{bile}} = 100$ liters/min, $CL_{\text{ef}}^s = 0.1$ liter/min). Then, systemic AUC was simulated for $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ “test” ratios ranging from 0.1 to 10 ($CL_{\text{met} + \text{bile}} = 0.01$ –1 liter/min, $CL_{\text{ef}}^s = 0.1$ liters/min), and the resulting AUC (AUC_{ratio = test}) was normalized to the control simulation (AUCR = AUC_{ratio = test}/AUC_{ratio = control}). The decrease in $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio is akin to inhibition of $CL_{\text{met} + \text{bile}}$ since CL_{ef}^s is held constant. The $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio, which resulted in a significant change to the systemic AUC (AUCR = 1.25) compared with control, was identified as the tipping point.

To illustrate that the magnitude of CL_{in}^s contributes to the tipping point, we simulated the tipping point for CL_{in}^s values ranging from $0.01 \times Q_h$ to $4 \times Q_h$ (henceforth, for simplicity, CL_{in}^s notation will be used instead of $f_u CL_{\text{in}}^s$ since $f_u = 1$). The tipping point can be explicitly derived from the ECM (eq. 1) by defining the RDS switch for any chosen AUCR as $AUCR = RDS_{\text{uptake}}/RDS_{\text{all}}$ and solving for the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio (eq. 2). This relationship (eq. 2 with AUCR = 1.25) was used later to identify DDI liabilities when considering CL_{in}^s magnitude and $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio of a drug:

$$\text{Tippingpoint} = \frac{1}{(AUCR - 1) \left(1 + CL_{\text{in}}^s / Q_h \right)} \quad (2)$$

Quantifying When a Drug with RDS_{uptake} Will Switch to RDS_{all} from Metabolic/Biliary Efflux DDIs. We defined the $PI_{\text{met} + \text{bile}}$ as the percent inhibition of $CL_{\text{met} + \text{bile}}$ required for RDS_{uptake} to switch to RDS_{all} . This quantifies when a significant DDI (AUCR ≥ 1.25) occurs from inhibition of $CL_{\text{met} + \text{bile}}$, even when uptake is the RDS in the absence of DDI. For $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratios ranging from 1 to 100, $CL_{\text{met} + \text{bile}}$ was inhibited 10%–99%. Simulations were conducted for CL_{in}^s values = 0.25, $1 \times$, $4 \times Q_h$. CL_{in}^s values were chosen to represent ER = 0.2, 0.5, and 0.8 [low-, mid-, and high-extraction ratio (ER), respectively] and were back-calculated from eq. 3 to eq. 4. The percent inhibition of $CL_{\text{met} + \text{bile}}$ at which the $CL_{\text{met} + \text{bile}}/CL_{\text{ef}}^s$ ratio reaches the tipping point (i.e., $PI_{\text{met} + \text{bile}}$) and thus causes the RDS_{uptake} to switch to RDS_{all} was calculated as shown in eq. 5:

$$CL_h = Q_h \cdot ER \quad (3)$$

$$CL_h = \frac{Q_h f_u CL_{\text{in}}^s}{Q_h + f_u CL_{\text{in}}^s} \quad (4)$$

$$PI_{\text{met} + \text{bile}}(\%) = \frac{CL_{\text{met} + \text{bile}} / CL_{\text{ef}}^s - \text{tippingpoint}}{CL_{\text{met} + \text{bile}} / CL_{\text{ef}}^s} \times 100 \quad (5)$$

Applying the RDS Framework to In Vitro and In Vivo Examples. Published data sets in which all hepatobiliary clearance pathways (CL_{in}^s , CL_{ef}^s , CL_{bile} , CL_{met}) were quantified in vitro were collected. The in vivo hepatobiliary clearances must be used to identify the RDS of a drug. As such, the provided in vitro to in vivo extrapolated (IVIVE) clearances were used; otherwise, in vitro hepatobiliary clearance values were scaled to in vivo using IVIVE scaling factors as provided by the authors. For all drugs, $fu_b CL_{in}^s/Q_h$ was used to calculate the tipping point using eq. 2 (see *Results* section to follow). RDS was labeled as RDS_{uptake} and RDS_{all} if the $CL_{met+bile}/CL_{ef}^s$ ratio was above and below the tipping point, respectively. For drugs with RDS_{uptake} , the $PI_{met+bile}$ was calculated using eq. 5. Finally, for selected drugs, the predicted DDI liabilities using the RDS and $PI_{met+bile}$ were compared with the observed in vivo data. To ensure that only the systemic clearance, and not the bioavailability of the victim drug, was affected, clinical DDI studies were included if the victim was a dual- transporter/enzyme substrate and coadministered with a selective enzyme inhibitor administered i.v. It should be noted that the availability of such studies was limited.

Results

Identifying the Tipping Point (i.e., When RDS_{uptake} Switches to RDS_{all}) and Factors that Influence this Switch. As described in the *Theoretical Background* section, RDS_{uptake} occurs when $CL_{met+bile} >>$

CL_{ef}^s , and, as such, inhibition of $CL_{met+bile}$ will not manifest in the systemic AUC of a victim drug. When this condition is violated owing to extensive inhibition of $CL_{met+bile}$, there will be a significant increase in the systemic AUC of the victim drug when $CL_{met+bile}$ is inhibited further. In other words, when $CL_{met+bile}$ is no longer $>> CL_{ef}^s$, then RDS_{uptake} switches to RDS_{all} . In Fig. 1A, 84%, 98%, and 99.8% inhibition of $CL_{met+bile}$ led to a clinically significant increase in the systemic AUC of the three theoretical victim drugs shown ($AUCR \geq 1.25$). Even though the victim drugs had different preinhibition $CL_{met+bile}$ values (1, 10, 100 liters/min), the postinhibition $CL_{met+bile}$ values were all the same (0.2 liters/min). Since CL_{ef}^s was kept constant (0.1 liters/min), an AUCR of 1.25 was observed when $CL_{met+bile}/CL_{ef}^s = 2$ for all three victim drugs. This simulation illustrates that the RDS_{uptake} switch to RDS_{all} depends on the $CL_{met+bile}/CL_{ef}^s$ ratio, not on the extent of $CL_{met+bile}$ inhibition.

To further emphasize the dependence on the $CL_{met+bile}/CL_{ef}^s$ ratio, we simulated the systemic AUC of the victim drug (in the absence of DDI) for different $CL_{met+bile}$ and CL_{ef}^s values while holding CL_{in}^s constant. The systemic AUC remained unchanged when the $CL_{met+bile}/CL_{ef}^s$ ratio remained fixed, even though the $CL_{met+bile}$ and CL_{ef}^s values varied,

Fig. 1. Identifying when RDS_{uptake} switches to RDS_{all} , that is, the tipping point. (A) Extensive inhibition of $CL_{met+bile}$ can lead to a significant increase in the systemic AUC for three theoretical victim drugs that have RDS_{uptake} (i.e., $CL_{met+bile} >> CL_{ef}^s$) in the absence of DDI. When inhibition of $CL_{met+bile}$ eventually violates the condition required for RDS_{uptake} , the RDS_{uptake} switches to RDS_{all} . An $AUCR \geq 1.25$ was observed when $CL_{met+bile}$ was inhibited $\geq 84\%$, $\geq 98\%$, and $\geq 99.8\%$ for $CL_{met+bile} = 1, 10, 100$ liters/h, respectively. For all three victim drugs, however, the $CL_{met+bile}$ value after such inhibition was similar (0.2 liters/min), as was the $CL_{met+bile}/CL_{ef}^s$ ratio ($= 2$). Simulations were performed as follows: $CL_{in}^s = 1 \times Q_h$, $CL_{met+bile} = 1, 10, 100$ liters/h, $CL_{ef}^s = 0.1$ liters/min. (B) The systemic AUC (in the absence of any DDI) of a theoretical drug remains unchanged when the $CL_{met+bile}/CL_{ef}^s$ ratio remains fixed (blue bars) but not when the $CL_{met+bile}/CL_{ef}^s$ ratio is varied (yellow bars), even though the absolute value of $CL_{met+bile}$ and CL_{ef}^s is varied in both scenarios. This trend was observed irrespective of the value of CL_{in}^s (Supplementary Fig. 1). Furthermore, this trend is true for when $CL_{met+bile} > CL_{ef}^s$ or $CL_{met+bile} < CL_{ef}^s$ (also refer to Supplementary Fig. 1). Thus, the $CL_{met+bile}/CL_{ef}^s$ ratio, irrespective of the magnitude of the absolute values of these clearances, is important for establishing the RDS and henceforth when the RDS switches from uptake to all hepatobiliary clearances. Simulations were performed as follows: $CL_{in}^s = 0.25 \times Q_h$, and the other input clearance values for scenarios A–E are shown in the table provided. (C) Since the RDS depends on the $CL_{met+bile}/CL_{ef}^s$ ratio, we define the tipping point as the $CL_{met+bile}/CL_{ef}^s$ ratio at which RDS_{uptake} switches to RDS_{all} . Similarly to (A), the RDS_{uptake} switch to RDS_{all} is represented by an AUCR of 1.25 and the decrease in $CL_{met+bile}/CL_{ef}^s$ ratio is akin to inhibition of $CL_{met+bile}$ when CL_{ef}^s is kept constant. As shown by the gray arrows, the tipping point for a low-, mid-, and high-ER drug is 3.2, 2, and 0.8, respectively. For example, if the $CL_{met+bile}/CL_{ef}^s$ ratio for the low ER drug is above the tipping point (i.e., $CL_{met+bile}/CL_{ef}^s > 3.2$), then the drug will have RDS_{uptake} ; therefore, DDIs from CL_{in}^s but not $CL_{met+bile}$ should be expected; however, inhibition of $CL_{met+bile}$ that makes the $CL_{met+bile}/CL_{ef}^s$ ratio lower than the tipping point (i.e., $CL_{met+bile}/CL_{ef}^s < 3.2$) will lead to a significant increase in the systemic AUC. Crossing the tipping point is indicative of the RDS_{uptake} switch to RDS_{all} . Simulations were performed as follows: systemic AUC were simulated for $CL_{in}^s = 0.25 \times, 1 \times, 4 \times Q_h$ (representing low, mid, and high ER, respectively) and $CL_{met+bile}/CL_{ef}^s$ ratios from 1 to 10 and then normalized to a control simulation where the $CL_{met+bile}/CL_{ef}^s$ ratio was set to 1000 (i.e., RDS_{uptake}).

demonstrating that the RDS in the hepatic clearance of a drug is dependent on the $CL_{met+bile}/CL_{ef}^s$ ratio, not on the absolute value of these clearances (Fig. 1B). This was true for both when $CL_{met+bile}$ was higher and lower than CL_{ef}^s (also see Supplementary Fig. 1). Since the systemic AUC decreased as the $CL_{met+bile}/CL_{ef}^s$ ratio increased, only the $CL_{met+bile}/CL_{ef}^s$ ratio needs to be considered when determining when the RDS_{uptake} switches to RDS_{all} for a victim drug.

Next, we identified the tipping point, defined here as the $CL_{met+bile}/CL_{ef}^s$ ratio when RDS_{uptake} switches to RDS_{all} . The RDS_{uptake} switch to RDS_{all} signifies when DDIs owing to inhibition of $CL_{met+bile}$ start to become significant for a victim drug that has RDS_{uptake} . As demonstrated already, the RDS_{uptake} switch to RDS_{all} depends on the $CL_{met+bile}/CL_{ef}^s$ ratio. As such, we identified the tipping point as the $CL_{met+bile}/CL_{ef}^s$ ratio at which the systemic AUC increases significantly ($AUCR = 1.25$) owing to a decrease in the $CL_{met+bile}/CL_{ef}^s$ ratio for a victim drug that has RDS_{uptake} (Fig. 1C). As demonstrated in Fig. 1C, the tipping point for a low, mid, and high ER drug was 3.2, 2, and 0.8, respectively.

Since the tipping point varied for a low, mid, and high ER, the magnitude of CL_{in}^s is also an important factor in determining when the RDS_{uptake} switches to RDS_{all} (Fig. 1C). Extending the simulations to identify the tipping point across a range of CL_{in}^s values, we established a theoretical (eq. 2) and practical (Fig. 2) relationship between CL_{in}^s/Q_h and the tipping point. The tipping point decreases as CL_{in}^s increases. In other words, as a drug's CL_{in}^s (and therefore its ER) increases, the drug is more likely to have RDS_{uptake} and a larger $PI_{met+bile}$, therefore making the drug more resistant to switching its RDS. In addition, as the influx across the sinusoidal membrane becomes large, hepatic clearance becomes limited by blood flow and therefore less likely to result in a change in AUCR when either CL_{in}^s (or for that matter $CL_{met+bile}$) is inhibited. On the other hand, when CL_{in}^s (or ER) is small and the hepatic clearance becomes proportional to CL_{in}^s , the victim drug becomes more susceptible to a change in RDS. This demonstrates that low ER drugs are more susceptible to RDS_{uptake} switching to RDS_{all} , whereas high ER drugs are more resistant to the RDS switch.

It should be noted that the relationship between CL_{in}^s/Q_h and the tipping point (eq. 2 and Fig. 2) depends on the chosen AUCR cutoff. Here, an AUCR of 1.25 was chosen based on Food and Drug Administration guidelines of what constitutes a positive DDI. If a higher AUCR cutoff were to be selected (Supplementary Fig. 2), this would lead to estimation of lower tipping points, thus making it more likely that drugs are labeled with RDS_{uptake} . Labeling a drug with RDS_{uptake} when in fact it has RDS_{all} can lead to underpredictions of DDI liabilities from metabolic enzymes and biliary transporters.

By understanding the relationship between CL_{in}^s and the tipping point, the RDS can be identified for any combination of a drug's hepatobiliary clearance values (Fig. 2). For example, a high-ER drug with a $CL_{met+bile}/CL_{ef}^s$ ratio of 3 will have RDS_{uptake} , but a low-ER drug with the same $CL_{met+bile}/CL_{ef}^s$ ratio will have RDS_{all} . Furthermore, a drug will always have RDS_{uptake} if the $CL_{met+bile}/CL_{ef}^s$ ratio is greater than 4, irrespective of the magnitude of CL_{in}^s . It should be noted that for orally administered drugs, the tipping point will no longer depend on the magnitude of CL_{in}^s and therefore will always be 4 because blood flow limitations from systemic clearance are cancelled out by blood flow limitations of hepatic bioavailability.

Quantifying the $PI_{met+bile}$ for Drugs with RDS_{uptake} . Identifying the RDS of a drug and when the RDS_{uptake} to RDS_{all} switch will happen identifies the drug's DDI liabilities. We quantified the $PI_{met+bile}$, defined here as the percent inhibition of $CL_{met+bile}$ needed to cause the RDS_{uptake} switch to RDS_{all} , to understand when inhibition of $CL_{met+bile}$ starts to become a DDI liability for victim drugs that have RDS_{uptake} . As the $CL_{met+bile}/CL_{ef}^s$ ratio of the victim drug (before inhibition) increases, the $PI_{met+bile}$ increases (Fig. 3A) because, as $CL_{met+bile}$ becomes $>>$

Fig. 2. The RDS framework helps identify DDI liabilities. The $CL_{met+bile}/CL_{ef}^s$ ratio and CL_{in}^s magnitude of a drug determine the RDS of the drug and when RDS_{uptake} switches to RDS_{all} . Combinations of hepatobiliary clearances found in the shaded area have RDS_{all} , whereas those in the nonshaded area have RDS_{uptake} . Any alterations in hepatobiliary clearances that cause a drug to switch from the nonshaded to the shaded area will cross the tipping point (dashed line, eq. 2) and therefore switch the RDS from uptake to all hepatobiliary clearances. The consequence of this switch is that DDIs from inhibition of $CL_{met+bile}$ will now manifest in the systemic AUC of a victim drug that originally had RDS_{uptake} . Consistent with Fig. 1C, the tipping point decreases as the magnitude of CL_{in}^s (and therefore the drug's ER) increases. This suggests that the greater the ER of the drug, the more likely it will have RDS_{uptake} and will be more resistant to switch to RDS_{all} . Furthermore, when $CL_{met+bile}/CL_{ef}^s > 4$, the RDS will always be uptake clearance, irrespective of the value of CL_{in}^s/Q_h ; however, when $CL_{met+bile}/CL_{ef}^s < 4$, the RDS can be either uptake or all hepatobiliary pathways, depending on the magnitude of CL_{in}^s . It should be noted that if a drug is administered orally, the tipping point will always be 4 because the blood flow limitations are no longer relevant. Simulations were performed as follows: the tipping point was simulated for CL_{in}^s values $(0.01 \times Q_h - 4 \times Q_h)$ using eq. 2.

than CL_{ef}^s , the victim drug become resistant to the RDS_{uptake} switch to RDS_{all} . High-ER drugs have a higher $PI_{met+bile}$ than low-ER drugs, demonstrating again that high-ER drugs are resistant to the RDS switch, whereas low-ER drugs are sensitive (Fig. 3A). Figure 3B illustrates that whereas a low-, mid-, and high-ER victim drug with $CL_{met+bile}/CL_{ef}^s$ ratio of 6 have RDS_{uptake} (before inhibition), inhibition of $CL_{met+bile} > 46\%$, 66% , and 87% , respectively, will cause the RDS_{uptake} to switch to RDS_{all} . This translates to observing a positive DDI owing to inhibition of $CL_{met+bile}$ for a victim drug that has been identified to have RDS_{uptake} (before inhibition). Without knowledge of the $PI_{met+bile}$, such a DDI may not be expected.

The purpose and conclusions of the simulations that have been used to establish the RDS framework up to this point are summarized in Fig. 4. As discussed, identifying the drug's RDS is not enough to predict correctly the drug's DDI liabilities. The tipping-point concept is an important consideration when identifying DDIs for victim drugs that are dual substrates of enzymes and transporters.

The flowchart in Fig. 5 can be used as a guide to identify the DDI liabilities for dual-transporter/enzyme substrates. All drugs with $CL_{met+bile}/CL_{ef}^s$ ratio > 4 will have RDS_{uptake} , whereas drugs with $CL_{met+bile}/CL_{ef}^s$ ratio < 4 will have RDS_{uptake} so long as this ratio is greater than the tipping point. Drugs with $CL_{met+bile}/CL_{ef}^s$ ratio less than the tipping point will have RDS_{all} . If the drug has RDS_{uptake} , then uptake transporters will become a DDI liability, whereas if the drug has RDS_{all} , then transporters and enzymes will be a DDI liability. Even for drugs that have RDS_{uptake} , however, $CL_{met+bile}$ can become a DDI liability if inhibition of $CL_{met+bile}$ is greater than the predicted $PI_{met+bile}$ and thus causes the RDS_{uptake} switch to RDS_{all} . The flowchart identifies the

Fig. 3. Identifying when drugs with RDS_{uptake} will start to experience a DDI from inhibition of CL_{met + bile}. (A) The PI_{met + bile}, defined as the % inhibition of CL_{met + bile} required for the RDS_{uptake} switch to RDS_{all}, depends on the CL_{met + bile}/CL_{ef}^s ratio (before inhibition) and the magnitude of CL_{in}^s (represented as low-, mid-, and high-ER drugs). The PI_{met + bile} identifies when a positive DDI from inhibition of CL_{met + bile} for a drug with RDS_{uptake} would be expected. Lower CL_{met + bile}/CL_{ef}^s ratios, as well as low ER drugs, are the most susceptible for the RDS_{uptake} switch to RDS_{all} owing to CL_{met + bile} inhibition. (B) For RDS_{uptake} to switch to RDS_{all} for a theoretical victim drug with CL_{met + bile}/CL_{ef}^s ratio of 6, CL_{met + bile} must be inhibited by >46%, >66%, or >87% if the drug is low-, mid-, and high-ER, respectively. Visually, the RDS_{uptake} switch to RDS_{all} happens when the theoretical victim drug crosses the dashed line (the tipping point) from the unshaded area (RDS_{uptake}) to the shaded area (RDS_{all}). Additional examples of PI_{met + bile} are given in the table provided. Simulations were performed as follows: PI_{met + bile} was calculated using eq. 5 for CL_{met + bile}/CL_{ef}^s ratios ranging from 1 to 40 and for CL_{in}^s = 0.25 ×, 1 ×, 4 × Q_h (representing low-, mid-, and high-ER, respectively).

CL_{met + bile}/CL_{ef}^s ratios at which 25%, 50%, 75%, and 95% expected inhibition of CL_{met + bile} is going to result in the RDS switch. This information can be used to assess when CL_{met + bile} starts to become a DDI liability for drugs with RDS_{uptake}. It also helps answer the question of how much larger CL_{met + bile} needs to be compared with CL_{ef}^s for sinusoidal uptake clearance to become (and maintain) the RDS in the hepatic clearance of any drug. Such information may be used during drug development to select drug candidates if a certain RDS is desired.

Applying the RDS Framework to In Vitro and In Vivo Examples. To provide context to the theoretical framework presented, examples from literature, where available, were used. For drugs with in vitro-quantified hepatobiliary clearances that were extrapolated to in vivo via IVIVE, the tipping point and the PI_{met + bile} were calculated using eq. 2 and eq. 5, and a subset of the analyzed data set, which includes primarily statin drugs, is shown in Fig. 6 (also see Supplementary Table 1) (Camenisch and Umehara, 2012; Jones et al., 2012; Varma et al., 2014; Kunze et al., 2015; Riede et al., 2017). If no empirical scaling factors (such as for active uptake clearance to match observed in vivo clearance) are included in the IVIVE process, then almost all drugs have RDS_{all}, except valsartan and pravastatin (Fig. 6A). This is because most CL_{met + bile}/CL_{ef}^s ratios are <4, and because the IVIVE CL_{in}^s magnitudes were small, most CL_{met + bile}/CL_{ef}^s ratios were less than the tipping point. Because many statins have been identified to have RDS_{uptake}, the trend in Fig. 6A suggests that CL_{in}^s was underestimated in vitro. When hepatobiliary clearances were adjusted by empirical scaling factors (Varma et al., 2014) or parameters were fitted from in vivo intravenous concentration-time profiles using a physiologically based pharmacokinetic (PBPK) model (Jones et al., 2012), the distribution of drugs is altered as low ER drugs (ER < 0.2) tended to have RDS_{all}, whereas mid- and high-ER drugs (ER > 0.2) were more likely to have RDS_{uptake} (Fig. 6B). This analysis of the published in vitro hepatobiliary clearances provides insight that drugs with RDS_{uptake} exist within the moderate RDS framework space, meaning that in general their CL_{met + bile}/CL_{ef}^s ratio is <4, and they are quite susceptible to the RDS switch (Supplementary Table 1). It further elucidates that current in vitro quantification techniques may underestimate CL_{in}^s, which can lead to erroneous labeling of the RDS and thus incorrect DDI liability predictions (Fig. 7; Supplementary Fig. 3).

To illustrate more fully the applicability of the RDS framework, predicted DDI liabilities using the RDS framework were compared with in vivo DDI examples. As indicated in Table 1, when empirical scaling factors are used during the IVIVE process or hepatobiliary clearances were estimated from in vivo via PBPK, atorvastatin and repaglinide have RDS_{uptake} and PI_{met + bile} of 10%–51% and 15%–40%, respectively, whereas bosentan has RDS_{all}. For atorvastatin and repaglinide, the in vitro data predicted that uptake transporters (OATPs) are the primary DDI liability, with the drugs' major metabolic enzymes (CYP3A and CYP2C8, respectively) becoming a potential liability only if the in vivo hepatic metabolic inhibition is greater than the PI_{met + bile}. For bosentan, the in vitro data predicted that both OATPs and CYP3A4 are potential DDI liabilities. Clinically, for atorvastatin, coadministration of rifampin (an OATP inhibitor) leads to an AUCR of 12, whereas 33% inhibition of CYP3A4 due to intravenous itraconazole (as measured using CYP3A4 probe midazolam) did not change atorvastatin systemic AUC, even though inhibition of atorvastatin metabolism was observed via a decrease in the 2-hydroxyatorvastatin concentrations (Maeda et al., 2011). In a similarly conducted experiment, coadministration of rifampin resulted in AUCR of 3.2 and 1.9 for bosentan or repaglinide, respectively, whereas 73% inhibition of CYP3A4 owing to intravenous itraconazole did not significantly change the systemic AUC of these drugs (Yoshikado et al., 2017). Furthermore, repaglinide coadministered with oral rifampin and trimethoprim (CYP2C8-selective inhibitor) resulted in AUCR 2.6 and 1.8, respectively (Kim et al., 2016). The in vivo DDI liability for OATPs was well predicted for all three victim drugs. The in vivo DDI liability for CYP3A4 was well predicted for atorvastatin. Since a probe was not used to assess the degree of CYP2C8 inhibition, it is difficult to determine whether the significant DDI when repaglinide was coadministered with trimethoprim was because RDS_{uptake} switched to RDS_{all} or because repaglinide truly has RDS_{all}. The in vitro metrics, as well as a whole-body PBPK DDI model, suggests that repaglinide has RDS_{uptake} (Varma et al., 2013); thus, the repaglinide-trimethoprim DDI is likely due to the RDS switch. Lastly, since bosentan was predicted to have RDS_{all}, a DDI was expected to result from CYP3A4 inhibition, but none was observed. It should be noted that the metabolic DDI liability prediction is assuming one main

Fig. 4. Summary of the purpose and conclusions for the simulations used to establish the RDS framework.

drug-metabolizing enzyme and no significant biliary efflux (e.g., $CL_{met+bile} = CL_{CYP3A4}$ for atorvastatin and bosentan). This assumption predicts the highest DDI risk owing to inhibition of $CL_{met+bile}$ and has a higher chance of predicting false-positive DDI results.

In the published in vitro data sets, discrepancies in the in vitro quantified values, particularly for CL_{in}^s , can be observed (Supplementary Table 1; Table 1). For example, in one report, the authors used empirical scaling factors for active sinusoidal uptake clearance to match hepatic clearance with clinically observed data that ranged from 1.1 to 101.8 with a geometric mean of 10.6 (Varma et al., 2014); however, the scaling factor used severely impacted the labeling of the RDS (e.g., fluvastatin, glyburide, pravastatin) or impacted the predicted $PI_{met+bile}$ of drugs (e.g., atorvastatin, rosuvastatin, fluvastatin, repaglinide) (Supplementary Table 1). Assumptions regarding CL_{ef}^s also caused discrepancies. In all reports, CL_{ef}^s was assumed to be equal to passive diffusion across the sinusoidal membrane, except in one report in which CL_{ef}^s was back-calculated from total sandwich cultured human hepatocytes CL_{int} (Camenisch and Umehara, 2012). The assumptions surrounding CL_{ef}^s impacted the $CL_{met+bile}/CL_{ef}^s$ ratio, which either changed how the RDS was labeled or the magnitude of the $PI_{met+bile}$ (e.g., aliskerin, ciprofloxacin, digoxin) (Supplementary Table 1). All in all,

mispredictions of any of the hepatobiliary clearances impact the RDS labeling, magnitude of the $PI_{met+bile}$, and DDI liability predictions.

Errors from in vitro quantification of hepatobiliary clearances can propagate when establishing the RDS and the predicted DDI liabilities. Underprediction of both CL_{in}^s and $CL_{met+bile}$ may erroneously label a drug with RDS_{all} when it is truly RDS_{uptake} (Fig. 7). $CL_{met+bile}$ is the more sensitive parameter for determining the RDS because underpredictions of CL_{in}^s may mislabel the RDS only for drugs with $CL_{met+bile}/CL_{ef}^s$ ratio < 4 (Supplementary Fig. 4). For such drugs, even moderate (e.g., 2- to 5-fold) underpredictions of either clearance pathway will lead to RDS mislabeling (Supplementary Fig. 4). Furthermore, underpredictions of both CL_{in}^s and $CL_{met+bile}$ leads to underprediction of $PI_{met+bile}$, resulting in predicting a larger DDI liability owing to $CL_{met+bile}$ inhibition for a drug with RDS_{uptake} (Fig. 7; Supplementary Fig. 4). Whereas underpredictions of hepatobiliary clearances will result in conservative DDI decisions, they also increase the chances of negative DDI studies.

Discussion

We built a theoretical RDS framework and identified important considerations when predicting DDI liabilities for dual transporter-enzyme

Fig. 5. Applying the RDS framework to identify DDI liabilities for dual transporter-enzyme substrate drugs. If $CL_{met+bile}/CL_{ef}^s > 4$, then the drug will have RDS_{uptake} , irrespective of the magnitude of CL_{in}^s . For drugs with RDS_{uptake} , DDIs due to inhibition of $CL_{met+bile}$ can become significant, depending on the drug's $CL_{met+bile}/CL_{ef}^s$ ratio and the expected inhibition of $CL_{met+bile}$. For example, 50% inhibition of $CL_{met+bile}$ may result in a significant DDI for a drug with RDS_{uptake} and $CL_{met+bile}/CL_{ef}^s$ ratio < 8 but no DDI will be observed if the drug has $CL_{met+bile}/CL_{ef}^s$ ratio > 8. The DDI liability owing to inhibition of $CL_{met+bile}$ increases as the $CL_{met+bile}/CL_{ef}^s$ ratio decrease and the expected $CL_{met+bile}$ inhibition increases.

Fig. 6. The distribution of drugs within the RDS framework using hepatobiliary clearance quantified in vitro and extrapolated to in vivo. Published in vitro hepatobiliary clearance values, when extrapolated to in vivo via IVIVE, can identify the RDS based on $fu_b CL_{in}^s/Q_h$ and $CL_{met+bile}/CL_{ef}^s$ ratio (eq. 2). (A) When no empirical scaling factors, such as to scale up active transport, are applied during the IVIVE process, all drugs except for valsartan and pravastatin have RDS_{all}. Most drugs had $CL_{met+bile}/CL_{ef}^s$ ratio <4 , indicating drugs primarily exist within the moderate RDS framework space. Furthermore, most drugs have $fu_b CL_{in}^s/Q_h < 0.4$, indicating severe underprediction of CL_{in}^s . (B) When empirical scaling factors are used or hepatobiliary clearances are estimates from in vivo data using PBPK modeling, the RDS of the drugs is altered severely. Now, RDS_{uptake} occurs more often for mid- and high-ER drugs with RDS_{all} primarily for low ER drugs (ER was calculated from in vivo hepatic clearance and blood flow). Furthermore, since all drugs have $CL_{met+bile}/CL_{ef}^s$ ratio <4 , information about both the magnitude of $fu_b CL_{in}^s$ and the $CL_{met+bile}/CL_{ef}^s$ ratio is necessary to correctly predict DDI liabilities. The dashed line represents the tipping point (eq. 2). The data shown are from Jones et al. (2012) and Varma et al. (2014) and represent a subset of the complete data set presented in Supplementary Table 1.

substrate drugs. First, inhibition of $CL_{met+bile}$ can cause the RDS of a victim drug to switch from RDS_{uptake} to RDS_{all} and hence result in an unexpected systemic DDI. Two metrics have been developed to identify when the RDS switch occurs: the tipping point, defined as the $CL_{met+bile}/CL_{ef}^s$ ratio at which RDS_{uptake} will switch to RDS_{all}, and the $PI_{met+bile}$, defined as the percent inhibition of $CL_{met+bile}$ at which a significant AUC change ($AUCR > 1.25$) for a drug with RDS_{uptake} will start to be observed. The tipping point depends on the drug's $CL_{met+bile}/CL_{ef}^s$ ratio and on the magnitude of CL_{in}^s . The former but not the latter condition is relevant when victim drugs are administered orally. Second, we showed that the $CL_{met+bile}/CL_{ef}^s$ ratio must be >4 for any drug to have RDS_{uptake}. Third, we applied the RDS framework to in vitro-quantified hepatobiliary clearances and observed that most drugs have $CL_{met+bile}/CL_{ef}^s$ ratio <4 ; hence, in practice, the magnitude of CL_{in}^s must be considered when establishing the RDS.

Our theoretical analysis demonstrates that the $CL_{met+bile}/CL_{ef}^s$ ratio, and not the absolute magnitudes of the clearances, determines the RDS in the hepatic clearance of a drug. Previous publications allude to this relationship. The authors of the ECCCS observed through experimental data that when $CL_{met+bile}$ is $2 \times CL_{ef}^s$, drugs that have RDS_{uptake} can be separated from those that do not (Riede et al., 2016). Furthermore, the β value [$\beta = CL_{met+bile}/(CL_{met+bile} + CL_{ef}^s)$] introduced by Yoshikado et al. (2016) can be used to differentiate the RDS, such as when β approaches unity (i.e., $CL_{met+bile} \gg CL_{ef}^s$), a drug has RDS_{uptake}. Our analyses corroborate and expand upon these results to provide a

quantitative definition of the demarcation point between RDS_{uptake} and RDS_{all} (i.e., the tipping point) and illustrate that the magnitude of CL_{in}^s , in addition to the $CL_{met+bile}/CL_{ef}^s$ ratio, is an important factor in determining the RDS of a drug. That is, as a drug's CL_{in}^s value increases, the drug is more likely to have RDS_{uptake} and to become resistant to the RDS_{uptake} switch to RDS_{all}.

We found good agreement for atorvastatin in vivo to predict DDI liabilities (Table 1). For bosentan, overprediction of expected DDI owing to inhibition of $CL_{met+bile}$ may be due to errors in the quantification of the hepatobiliary clearances. Indeed, a study in cynomolgus monkeys, in which bosentan plasma and liver drug concentrations were quantified, found that the in vitro scaled CL_{in}^s and CL_{met} were 28- and 13-fold underpredicted, whereas CL_{ef}^s (assumed equal to passive diffusion) was overpredicted by 2-fold compared with the in vivo-fitted values (Morse et al., 2017). Combining the in vitro metrics that identify RDS_{uptake} for repaglinide with in vivo repaglinide DDIs, it appears that CYP2C8 but not CYP3A4 inhibition may lead to RDS_{uptake} switch to RDS_{all}. Indeed, inhibition of repaglinide with gemfibrozil (CYP2C8 and OATP1B1 inhibitor) led to an 8-fold increase in systemic AUC; coadministration of itraconazole or cyclosporine (OATP1B1 and CYP3A4 inhibitor) led to much more modest 1.4- and 2.4-fold increases in systemic AUC (Niemi et al., 2003; Kajosaari et al., 2005).

Fig. 7. The impact of underpredictions of hepatobiliary clearance on DDI liability predictions. A representative 3-fold underprediction of either (A) CL_{in}^s or (B) $CL_{met+bile}$ can lead to erroneous labeling of the RDS for low-, mid-, and high-ER drugs (shown by the filled circles crossing from the nonshaded to shaded area (i.e., RDS_{uptake} switches to RDS_{all}). Mislabeling the RDS impacts the expected DDI risk from transporters versus enzymes. Furthermore, underpredictions of either CL_{in}^s or $CL_{met+bile}$ leads to identifying both transporters and enzymes as DDI liabilities when truly only uptake transporters are the true DDI liability. Please refer to Supplementary Fig. 4 for more detailed simulations.

TABLE 1

Comparison of predicted drug-drug interaction (DDI) liabilities from in vitro data to in vivo clinical studies

Hepatobiliary clearances, after in vitro to in vivo extrapolation (IVIVE), can be used to identify the rate-determining step (RDS) of a drug, such as if the $CL_{met} + bile/CL_{ef}^s$ ratio is $>$ or $<$ than the tipping point (eq. 2), then the drug will have RDS_{uptake} or RDS_{all} , respectively. If a drug has RDS_{uptake} , then the $PI_{met} + bile$ can be quantified (eq. 5) to predict when a significant DDI should be expected owing to inhibition of metabolic/biliary efflux clearance. An expanded analysis is shown in Supplementary Table 1.

Drug	Hepatobiliary Clearance (ml/min per kg)				$f_{ub}CL_{in}/Q_h^a$	$CL_{met} + bile/CL_{ef}^s$	Tipping Point	$PI_{met} + bile$	RDS In Vitro	RDS In Vivo	Reference
	CL_{in}^s	CL_{ef}^s	CL_{met}	CL_{bile}							
Atorvastatin	61 ^b	24.9	58.3	4.3	0.12	2.52	3.59	—	All	Uptake ^c	Varma et al. (2014)
	1194 ^d	24.9	58.3	4.3	2.27	2.52	1.22	$>51\%$	Uptake		Camenisch and Umehara (2012)
	405 ^e	24.9	58.3	4.3	0.77	2.52	2.26	$>10\%$	Uptake		Kunze et al. (2015)
	198 ^b	359	64.6	11.8	0.80	0.21	2.22	—	All		Maeda et al. (2011)
	198 ^b	57.7	64.6	11.8	0.80	1.32	2.22	—	All		
Bosentan	132 ^b	28.9	19.5	5.8	0.36	0.87	2.95	—	All	Uptake ^c	Varma et al. (2014)
	142 ^d	28.9	19.5	5.8	0.38	0.87	2.90	—	All		Jones et al. (2012)
	1117 ^e	28.9	19.5	5.8	3.02	0.87	0.99	—	All		Yoshikado et al. (2017)
	35 ^b	12.1	—	39 ^f	0.02	3.24	3.93	—	All		
	2035 ^g	14	—	5.0 ^f	1.09	0.36	1.91	—	All		
Repaglinide	166 ^b	63.6	128	0.3	0.19	2.01	3.35	—	All	Uptake ^c all ^h	Varma et al. (2014)
	1983 ^d	63.6	128	0.3	2.32	2.01	1.21	$>40\%$	Uptake		Jones et al. (2012)
	1151 ^e	63.6	128	0.3	1.35	2.01	1.71	$>15\%$	Uptake		Yoshikado et al. (2017)
	299 ^b	223	125	0.0	0.22	0.56	3.27	—	All		
	3671 ^g	352	125	0.0	2.73	0.35	1.07	—	All		

^aThe f_{ub} and Q_h values as noted in each reference were used for analysis; note that f_{ub} may vary for the same drug across different references.

^bIn vitro quantified + IVIVE.

^c RDS_{uptake} was established in vivo for atorvastatin and bosentan since there was no significant AUC change to victim drugs when coadministered with intravenous itraconazole (CYP3A inhibitor) which resulted in 33% and 73% CYP3A inhibition, respectively (Maeda et al., 2011; Yoshikado et al., 2017). Midazolam, a CYP3A probe, was used to assess magnitude of CYP3A inhibition.

^d RDS_{uptake} was established for repaglinide via whole-body physiologically based pharmacokinetic (PBPK) modeling of complex transporter- and enzyme-mediated DDIs (Varma et al., 2013).

^eIn vitro quantified + IVIVE + empirical scaling factor for active uptake transport (individual scaling factor).

^fIn vitro quantified + IVIVE + empirical scaling factor for active uptake transport (geometric mean scaling factor).

^gComposite $CL_{met} + bile$.

^hFitted parameters from in vivo using PBPK model.

ⁱ RDS_{all} was established in vivo for repaglinide even though there was no change to systemic AUC by intravenous itraconazole because CYP2C8 is the major hepatic drug-metabolizing enzyme (Yoshikado et al., 2017). In a different study, oral trimethoprim, a selective CYP2C8 inhibitor, increased repaglinide AUC by 1.8-fold (Kim et al., 2016).

The DDI liabilities discussed so far are relevant for systemic drug exposure but not necessarily for hepatic drug exposure, and thus efficacy/toxicity, if the site of action is in the liver. For example, the low-density lipoprotein cholesterol-lowering effect mediated by atorvastatin does not change for subjects with OATP1B1 polymorphism c.521T>C, even though there is a significant increase in atorvastatin systemic AUC (Maeda, 2015). This is because if the liver is the main eliminating organ, changes to sinusoidal uptake alter the hepatic concentration-time profile but not the hepatic AUC; however, a systemic increase of atorvastatin may lead to off-target toxicity, such as muscle myopathy. We refer the readers to our previous publication (Patilea-Vrana and Unadkat, 2016), which describes simulations that demonstrate the impact of inhibition of uptake or metabolism on both systemic and hepatic AUC when the liver is and is not the main eliminating organ.

The contrast between in vitro quantified CL_{in}^s with and without empirical scaling factors in Fig. 6 demonstrates that IVIVE of accurate transporter-mediated clearance remains challenging (Chu et al., 2013; Feng et al., 2014). The system used for in vitro quantification may be crucial since CL_{in}^s for statins quantified in sandwich cultured human hepatocytes appeared to be lower in magnitude than when quantified in suspended hepatocytes (Supplementary Table 1; Table 1). This may be mediated by significant intracellular localization of plasma membrane transporters (Kumar et al., 2017) or high interindividual variability when using individual donors (Vildhede et al., 2014). These reasons may also cause underpredictions of CL_{in}^s or CL_{bile} . For transporter IVIVE, we have previously recommended using a bottom-up proteomic approach and adjusting for in vitro activity via IVIVE transporter expression-based scaling factors (Prasad and Unadkat, 2014). We have recently demonstrated the successful prediction of hepatobiliary clearance of rosuvastatin in rats using the aforementioned approach (Ishida et al., 2018).

Special emphasis needs to be given to quantifying CL_{ef}^s along with $CL_{met} + bile$ since the $CL_{met} + bile/CL_{ef}^s$ ratio is one of the anchor points when establishing the RDS. Because CL_{ef}^s is a difficult parameter to

quantify in vitro, it is typically assumed to be equal to passive diffusion across the sinusoidal membrane; however, there are examples of active sinusoidal efflux transport, such as MRP3 efflux of rosuvastatin (Pfeifer et al., 2013). Active sinusoidal efflux would increase the magnitude of CL_{ef}^s and decrease the $CL_{met} + bile/CL_{ef}^s$ ratio, making a drug more likely to have RDS_{all} . One approach to measuring CL_{ef}^s is to use an integrative temporal modeling approach in sandwich cultured hepatocytes (Pfeifer et al., 2013; Ishida et al., 2018).

Errors in the quantification of CL_{in}^s and/or the $CL_{met} + bile/CL_{ef}^s$ ratio can impact DDI liability predictions. For example, patients with OATP1B1 polymorphism c.521T>C have about a 2-fold greater atorvastatin AUC compared with the wild-type allele (Maeda, 2015). Because of the lower CL_{in}^s , and therefore greater susceptibility to the RDS_{uptake} to RDS_{all} switch, patients with OATP1B1 polymorphism may experience a DDI attributable to inhibition of CYP3A, whereas patients with the wild-type allele may not. The same trend would be true for patients with polymorphic enzymes that result in lower $CL_{met} + bile$ and thus lower $CL_{met} + bile/CL_{ef}^s$ ratios. Polypharmacy use can also impact DDI liability predictions. For example, highly active antiretroviral therapy typically includes potent CYP3A4 and moderate OATP inhibitor ritonavir, among other drugs, which can impact the $CL_{met} + bile/CL_{ef}^s$ ratio more severely than if only one drug is administered. Indeed, the systemic AUC of atorvastatin increased by 3.9- and 9.4-fold when coadministered with saquinavir/ritonavir and tipranavir/ritonavir, respectively (Fichtenbaum et al., 2002; Pham et al., 2009). Lastly, the saturation of enzymes, leading to a lower CL_{met} with increased dose, may lower the $CL_{met} + bile/CL_{ef}^s$ ratio and cause DDIs owing to the RDS_{uptake} switch to RDS_{all} .

If a victim drug has RDS_{all} , but it has been mislabeled as RDS_{uptake} , then the DDI liability owing to inhibition of both transporter and metabolic activity could be underestimated. Considering potential DDI risks, it would be most conservative to assume a drug has RDS_{all} ; however, making such an assumption would lead to an increase in

negative DDI studies, particularly when conducting metabolic or biliary efflux DDI studies if the drug has RDS_{uptake} . An analysis of the DDIs performed for a cohort of NMEs in 2013 showed a modest return on investment because 57% ($n = 141$) of all in vivo DDIs were negative (Lesko and Lagishetty, 2016). Given the high prevalence of negative DDIs, it may be more appropriate to make mechanistic-based rather than conservative decisions regarding DDI liabilities.

The RDS framework presented here should be used as a guide for identifying the DDI liabilities, whereas PBPK models should be used to predict the direction and magnitude of complex transporter-enzyme DDIs. Several examples of such models (e.g., repaglinide, simvastatin, rosuvastatin) exist that predict complex interactions resulting from chemical inhibition or genetic polymorphism (Varma et al., 2013; Rose et al., 2014; Tsamandouras et al., 2015). Even with PBPK models, there are limitations. For example, when a drug has RDS_{uptake} , the $CL_{\text{met}} + \text{bile}$ is unidentifiable from plasma concentrations data since only CL_{in}^s plays a significant role in determining hepatic clearance. Focusing on capturing the correct $CL_{\text{met}} + \text{bile}$ magnitude, and not the $CL_{\text{met}} + \text{bile}/CL_{\text{ef}}^s$ ratio, can be misleading and will impact PBPK predictions. For instance, in an atorvastatin PBPK model, when cyclosporine CYP3A4 K_i was modulated 100-fold, a maximum 1.6-fold AUCR was achieved (Duan et al., 2017). Although the tendency is to run sensitivity analysis on the active components (transport and metabolism), a sensitivity analysis on CL_{ef}^s value (in the model, it was assumed to be equal to passive diffusion) should also be run as, for the specific example provided, it would likely have revealed a larger impact of cyclosporine on atorvastatin systemic AUC. Such an analysis may be helpful in consolidating in vitro K_i data with observed in vivo DDI data.

In summary, we introduced a theoretical RDS framework to predict more completely DDI liabilities for drugs that are dual transporter-enzyme substrates. We provide useful insights, such as the following: 1) the RDS_{uptake} switch to RDS_{all} depends on the ratio of $CL_{\text{met}} + \text{bile}/CL_{\text{ef}}^s$ and the magnitude of CL_{in}^s ; 2) $CL_{\text{met}} + \text{bile}/CL_{\text{ef}}^s > 4$ ensures RDS_{uptake} independent of CL_{in}^s magnitude or administration route; 3) there are existing drugs within a moderate space within the RDS framework that are susceptible to the RDS_{uptake} switch to RDS_{all} . Whereas these insights were obtained from the hepatic ECM, they can be equally applied to other organs, such as the kidneys, in which vectorial (basal-to-apical) transport of drugs is possible.

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Authorship Contributions

Participated in research design: Patilea-Vrana, Unadkat.

Conducted experiments: Patilea-Vrana.

Contributed new reagents or analytical tool: Patilea-Vrana.

Performed data analysis: Patilea-Vrana.

Wrote or contributed to the writing of the manuscript: Patilea-Vrana, Unadkat.

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