Systems Pharmacology Modeling of Drug-induced Hyperbilirubinemia: Differentiating Hepatotoxicity and Inhibition of Enzymes/Transporters

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Elevations in serum bilirubin during drug treatment may indicate global liver dysfunction and a high risk of liver failure. However, drugs also can increase serum bilirubin in the absence of hepatic injury by inhibiting specific enzymes/transporters. We constructed a mechanistic model of bilirubin disposition based on known functional polymorphisms in bilirubin metabolism/transport. Using physiologically based pharmacokinetic (PBPK) model-predicted drug exposure and enzyme/transporter inhibition constants determined in vitro, our model correctly predicted indinavir-mediated hyperbilirubinemia in humans and rats. Nelfinavir was predicted not to cause hyperbilirubinemia, consistent with clinical observations. We next examined a new drug candidate that caused both elevations in serum bilirubin and biochemical evidence of liver injury in rats. Simulations suggest that bilirubin elevation primarily resulted from inhibition of transporters rather than global liver dysfunction. We conclude that mechanistic modeling of bilirubin can help elucidate underlying mechanisms of drug-induced hyperbilirubinemia, and thereby distinguish benign from clinically important elevations in serum bilirubin.

Study Highlights

WHAT IS THE CURRENT KNOWLEDGE ON THE TOPIC?
☑ Severe drug-induced liver injury increases serum bilirubin. However, drug-induced hyperbilirubinemia can also be induced by inhibition of enzymes/transporters that mediate bilirubin disposition.

WHAT QUESTION DID THIS STUDY ADDRESS?
☑ Can enzyme/transporter-mediated drug-induced hyperbilirubinemia be predicted by mechanistic modeling from in vitro inhibition data? Can a mechanistic model differentiate bilirubin increases due to overt liver injury vs. enzyme/transporter inhibition?

WHAT THIS STUDY ADDS TO OUR KNOWLEDGE
☑ The mechanistic bilirubin model, combined with PBPK model-predicted drug exposure and in vitro enzyme/transporter inhibition constants, correctly predicted indinavir-mediated unconjugated hyperbilirubinemia and minimal bilirubin changes by nelfinavir. CKA induced ALT and bilirubin elevations in rats, and simulations suggest that CKA-mediated bilirubin elevation was mostly due to inhibition of bilirubin transporters, rather than liver injury.

HOW THIS MIGHT CHANGE CLINICAL PHARMACOLOGY OR TRANSLATIONAL SCIENCE
☑ Mechanistic modeling that represents hepatotoxicity mechanisms and enzyme-mediated and transporter-mediated bilirubin disposition can be used to elucidate underlying mechanisms of drug-induced hyperbilirubinemia and may be useful in prospective prediction of bilirubin increases by drug candidates.

Bilirubin, the product of heme breakdown from red blood cells, is exclusively eliminated by the liver. Thus, circulating bilirubin is widely used as a diagnostic biomarker for liver function. Large postmarketing studies of patients with drug-induced liver injury (DILI) show that ~10% of subjects with hyperbilirubinemia or jaundice die or require a liver transplant.1–3 In the setting of a clinical trial of a new drug candidate, elevations in serum bilirubin may also indicate severe liver injury with global hepatic dysfunction. The 2009 US Food and Drug Administration guidance on assessing liver safety in clinical trials defines the “Hy’s Law Case” as a trial subject who experiences a hepatocellular injury with concomitant elevations in serum alanine aminotransferase (ALT) >3× the upper limit of normal (ULN) and serum total bilirubin (TB) elevation >2× ULN when there is no more likely cause than the study drug. The existence of Hy’s Law Cases in a clinical trial database is interpreted as indicating that the study drug is capable of causing liver failure, a conclusion that can lead to termination of the development program, requirement of

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glucuronosyltransferase (UGT)1A1. Liver-generated CB is metabolized to CB (glucuronide conjugate, primarily) by UDP and Dubin-Johnson Syndrome (DJS) exhibit elevated serum bilirubin; UGT, UDP glucuronosyltransferase. Dubin-Johnson Syndrome; GS, Gilbert's Syndrome; CNS, Crigler-Najjar Syndrome; HC, hepatocytes; MRP, multidrug resistance-associated protein; OATP, organic anion transporting polypeptide; RBC, red blood cell; RS, Rotor Syndrome; UB, unconjugated bilirubin; UGT, UDP glucuronosyltransferase.

Figure 1. Diagrams of hepatobiliary disposition of bilirubin (a) and the bilirubin submodel structure within DILIsym (b). CB, conjugated bilirubin; DJS, Dubin-Johnson Syndrome; GS, Gilbert’s Syndrome; HC, hepatocytes; MRP, multidrug resistance-associated protein; OATP, organic anion transporting polypeptide; RBC, red blood cell; RS, Rotor Syndrome; UB, unconjugated bilirubin; UGT, UDP glucuronosyltransferase.

extended clinical trials to assess risk management strategies, or restricted indications. Therefore, the observation of hyperbilirubinemia during clinical trials or postmarketing can raise concern about irreversible liver injury with serious regulatory consequences. However, drug-induced hyperbilirubinemia is not always indicative of hepatotoxicity. Drugs may also increase serum bilirubin in the absence of hepatic injury by inhibiting enzymes/transporters that are involved in hepatic elimination of bilirubin. Serum bilirubin consists of conjugated bilirubin (CB) and unconjugated bilirubin (UB). UB is released to the circulation with red blood cell degradation and is taken up into hepatocytes by active transport (organic anion transporting polypeptide (OATP)1B1/1B3) and passive diffusion.4–6 Once inside the hepatocytes, UB is metabolized to CB (glucuronide conjugate, primarily) by UDP glucuronosyltransferase (UGT)1A1.7 Liver-generated CB is excreted into bile by multidrug resistance-associated protein (MRP)2, or transported back to sinusoidal blood by the efflux transporter MRP3, and then taken up again by downstream hepatocytes by OATP1B1/1B3.6,8–10 Patients with inherited disorders of bilirubin metabolism and/or transport (e.g., Rotor Syndrome (RS), Gilbert’s Syndrome (GS), Crigler-Najjar Syndrome (CNS), and Dubin-Johnson Syndrome (DJS)) exhibit elevated serum bilirubin, suggesting that modulation of any of these enzyme-mediated and transport-mediated processes by drugs may increase serum bilirubin (Figure 1).8,11–14 Several studies have reported that drugs with potent in vitro inhibition for enzyme/transporters that mediate bilirubin disposition at clinical exposure levels are associated with hyperbilirubinemia.15–17

Although the inhibitory potential for bilirubin enzyme/transporters can be assessed using in vitro studies, it remains a challenge to translate these in vitro data to in vivo and to predict the net effects of inhibition of multiple steps involved in bilirubin disposition. In addition, interpretation is not straightforward when hyperbilirubinemia is observed with and without concomitant serum ALT elevations in clinical trials. This is because some drugs without hepatic liabilities can cause high and frequent elevations in serum ALT.1 In such cases, distinguishing hyperbilirubinemia that is mediated by liver injury from enzyme/transporter-mediated interaction is challenging.

Mechanistic modeling could provide a useful tool to investigate the underlying mechanisms of hyperbilirubinemia and/or predict this multifactorial event. DILIsym (the product of a public-private partnership involving scientists from industry, academia, and the US Food and Drug Administration) is a mechanistic, multiscale model of DILI that integrates pharmacokinetic and in vitro toxicity data to predict in vivo hepatotoxicity in humans and preclinical animals.18 DILIsym can leverage existing preclinical and clinical data to predict the risk of hepatocellular DILI and prioritize drug candidates, and help optimize dosing paradigms and monitoring. Construction of SimPops, simulated populations that include variability in toxicity mechanisms, enabled prediction of low-frequency injuries as well as characterization of patients at DILI risk.19,20 The current SimPops represent healthy subjects, and patients with diabetes or nonalcoholic fatty liver disease, but do not yet include ethnic differences or age-related changes nor represent variability in biomarker-related parameters. The bilirubin submodel within DILIsym was initially designed to predict hyperbilirubinemia due to severe liver injury.21 In the current study, the bilirubin submodels for human and preclinical species were updated to represent enzyme/transporter-mediated bilirubin elimination. Serum bilirubin levels in patients and animals with impaired function of bilirubin enzyme or transporter(s) were used to determine the impact of each pathway on the changes in serum UB and CB. The impact of three compounds, indinavir, nelfinavir, and an investigational drug (chemokine receptor antagonist (CKA), 1-(4-chloro-3-(trifluoromethyl)benzyl)-5-hydroxy-1H-indole-2-carboxylic acid),22 on bilirubin levels was simulated in humans and rats to validate the current model.

Indinavir and nelfinavir are protease inhibitors used to treat human immunodeficiency virus. Both drugs are known to inhibit UGT1A1 and OATP1B1,15 but only indinavir is associated with hyperbilirubinemia.23,24 Altered serum bilirubin levels by indinavir and nelfinavir were simulated with DILIsym by combining in vitro enzyme/transporter inhibition data with in vivo exposure predicted by physiologically based pharmacokinetic (PBPK) modeling. CKA was shown to inhibit multiple hepatic transporters.22 CKA showed minimal to modest ALT increase in humans; whereas it induced dose-dependent serum ALT increases in rats.
Previous modeling with DILIsym using *in vivo* exposure and *in vitro* mechanistic toxicity data correctly predicted species differences in hepatotoxicity.25 Interestingly, CKA induced hyperbilirubinemia in all the treated rats, whereas ALT elevations were observed only in a subset of treated rats, suggesting that factors other than liver injury might have contributed to the bilirubin increases. In the present study, CKA-mediated effects on serum bilirubin were simulated using *in vitro* bilirubin transporter inhibition data, PBPK-predicted exposure, and previously defined hepatotoxicity mechanistic parameters in simulated human and rat populations.

**RESULTS**

**Simulation of bilirubin levels in patients with impaired bilirubin metabolism and transport**

The normal range of serum TB is 0.1–1.2 mg/dL, and about 3.0–4.3% of serum TB exists as CB.14 Respective baseline values in the current bilirubin model are 0.55 mg/dL and 3.6%. The bilirubin model parameters were optimized to clinical bilirubin data in patients with impaired bilirubin metabolism and transport (Supplementary Table S2 online). RS is a genetic disorder characterized by near-complete loss of OATP1B1/1B3 expression.8 Patients with RS present with conjugated hyperbilirubinemia, suggesting that (1) there exist non-OATP-mediated hepatic uptake pathway(s) for UB, and (2) OATP1B1/1B3 is involved in blood-hepatocyte shuttling of CB. Therefore, hepatic uptake of UB was represented by both non-OATP and OATP-mediated uptake processes, and hepatic uptake of CB was represented as an OATP1B1/1B3-mediated process. The serum TB and percent serum CB were 2.28–6.91 mg/dL and 56.1–79%, respectively, in the simulated patients with RS (with 70–95% impaired function of OATP1B1/1B3), consistent with clinically observed values of 4.42 ± 2.18 mg/dL and 62.9 ± 12.1% (Figure 2).8

Patients with GS have ~30–50% of normal UGT1A1 function, whereas CNS is a rare genetic disorder with near-complete loss of UGT1A1 function. Patients with GS or CNS present with unconjugated hyperbilirubinemia; serum TB levels in patients with GS increase to 1.8–4 mg/dL, whereas patients with CNS manifest increased serum TB up to 6–44 mg/dL.12,14 Simulated patients with GS and patients with CNS had impaired function of UGT1A1 by 30–60% and 70–90%, respectively. Simulated serum TB in respective groups were 0.84–5.99 and 8.32–13.03 mg/dL and were mostly composed of UB, consistent with the clinical data (Figure 2).

Patients with DJS, characterized by conjugated hyperbilirubinemia, have mutations in genes encoding MRP2. Patients with DJS have decreased MRP2 function and adaptive increases in MRP3 expression.12 Thus, the simulated patients with DJS had impaired function of MRP2 by 50–90%, and MRP3 function increased threefold.26 Patients with DJS manifest with conjugated hyperbilirubinemia; serum TB often increases up to 2–5 mg/dL, and less often, up to 25 mg/dL.27 Simulated serum TB levels in the patients with DJS were 5.08–13.28 mg/dL, 48–88% of which is conjugated, consistent with the clinical data (Figure 2).

Overall, simulations reasonably recapitulated conjugated hyperbilirubinemia in patients with RS or DJS, and unconjugated hyperbilirubinemia in patients with GS or CNS.

**Simulation of bilirubin levels in rats and mice lacking enzyme or transporters**

The normal range of serum TB and percent serum CB in rats and mice were highly variable (Supplementary Table S1 online).
The baseline values of serum TB and percent serum CB in the current bilirubin model are 0.55 mg/dL and 3.6%, consistent with humans. The bilirubin model parameters were optimized to the bilirubin data in rats and mice lacking specific enzyme and/or transporter(s) that are involved in bilirubin disposition (Supplementary Table S2 online). Simulations recapitulated conjugated hyperbilirubinemia in Mrp2-deficient (TR−), Eisai hyperbilirubinemic rat (EHBR), and Mrp2-knockout rats, which have impaired function of Mrp2 (observed and simulated serum TB: 1.9–4.0 vs. 3.3–6.4 mg/dL), unconjugated hyperbilirubinemia in Gunn rats with impaired function of Ugt1a1 (observed vs. simulated serum TB: 4.7–8.0 vs. 1.6–4.8 mg/dL), and Ugt1a1 knockout mice (observed vs. simulated serum TB: 7.8 ± 1.3 vs. 1.3–3.0 mg/dL; Figure 3).

The mouse bilirubin submodel recapitulated altered serum bilirubin levels in mice lacking single or multiple transporters; knockout of Mrp2 caused a modest increase in serum TB (observed vs. simulated changes from the baseline: 1.7-fold vs. 3.4-fold, respectively). Knockout of Oatp1a/1b significantly increased serum TB (observed vs. simulated: 8.3-fold vs. 8.5-fold), which is reversed by additional knockout of Mrp3 (observed vs. simulated: 3.1-fold vs. 2.6-fold), but potentiated by additional knockout of Mrp2 (observed vs. simulated: 21.9-fold vs. 12.5-fold; Figure 4). Due to large variability in the mouse data, fold-

Figure 3  Simulated and observed bilirubin levels in rats lacking Mrp2 or Ugt1a1, and mice lacking Ugt1a1. Closed and open circles represent observed and simulated serum total bilirubin (TB), respectively. (a) Observed data represent mean serum TB in Mrp2 knockout rats, EHBR rats, and TR rats (top to bottom). Simulated rats have 50–100% impaired Mrp2 function. (b) Observed data represent mean serum TB in Ugt1a1 knockout rats obtained from multiple studies. Simulated rats have 50–100% impaired Ugt1a1 function. (c) Observed data represent mean serum TB in Ugt1a1(-/-) (top) and Ugt1a1 (+/-) mice (bottom). Simulated mice have 50–100% impaired Ugt1a1 function.

Figure 4  Simulated and observed bilirubin levels in mice lacking single or multiple transporters. Observed (closed bars) and simulated (open bars) serum total bilirubin (TB) (a), percent conjugated bilirubin (CB) in serum compared to serum TB (b), and biliary excretion of TB presented as percent of wild-type animals (c).
rats; peak plasma concentration (Cmax) and area under the curve (AUC) were within 1.5-fold of those from the mean observed plasma concentrations in humans.\textsuperscript{29,30} Nelfinavir was shown to inhibit UGT1A1 and OATP1B1 in \textit{in vitro} assays, but is not associated with drug-induced hyperbilirubinemia (Table 1). Simulation of clinical dosing of nelfinavir (1,250 mg b.i.d. for 1 month) did not increase serum TB, consistent with clinical data.

**Species differences in chemokine receptor antagonist-mediated hyperbilirubinemia**

In the human SimPops administered single doses of up to 900 mg CKA, serum bilirubin did not exceed the ULN for any simulated individual, consistent with the clinical data. Conversely, the rat SimPops predicted dose-dependent serum bilirubin elevations at 50–500 mg/kg doses, consistent with preclinical data, although the magnitude of increase was underestimated (Figure 6a). In the preclinical studies, the mean fold-increase in serum TB was 2.1, 3.9, and 5.6 at 50, 200, and 500 mg/kg doses, respectively. Simulations predicted average serum TB increase of 1.4, 1.7, and 2.1-fold at respective doses (Figure 6a). When the same dosing protocols were simulated without considering inhibition of bilirubin transporters, serum TB levels minimally changed; predicted fold-increase in serum TB was 1.0, 1.1, and 1.2 at respective doses (Figure 6b). On the other hand, simulations with only bilirubin transporter-inhibition constants predicted 1.4-fold, 1.7-fold, and 1.9-fold increase in serum TB at 50, 200, and 500 mg/kg doses, respectively.

**Table 1** Summary of inhibition constants of indinavir, nelfinavir, and chemokine receptor antagonist for enzymes and transporters involved in bilirubin disposition

| Enzyme/transporter | Indinavir | Nelfinavir | CKA |
|--------------------|-----------|------------|-----|
| OATP1B1 IC\textsubscript{50} (\mu M) | 4.1 ± 1.8\textsuperscript{a} | 2 ± 0.9\textsuperscript{a} | 0.90\textsuperscript{b} |
| OATP1B3 IC\textsubscript{50} (\mu M) | >100\textsuperscript{a} | >100\textsuperscript{a} | N/A |
| UGT1A1 IC\textsubscript{50} (\mu M) | 6.8 ± 1.0\textsuperscript{c} | 4.8 ± 0.1\textsuperscript{c} | N/A |
| MRP2 IC\textsubscript{50} (\mu M) | >100\textsuperscript{a} | >100\textsuperscript{a} | 68.5 ± 7.3\textsuperscript{a} |
| MRP3 IC\textsubscript{50} (\mu M) | N/A | N/A | 11.2\textsuperscript{a} |

CKA, chemokine receptor antagonist, N/A, not available.

\textsuperscript{a}Determined from HEK cells transfected with OATP1B1 or OATP1B3 using pitavastatin as a probe substrate.\textsuperscript{12,13} The average of IC\textsubscript{50} values from two independent studies (0.84 and 0.97 \mu M, respectively), each performed in triplicate. Inhibition of OATP1B1 activity was determined in a stably transfected HEK293 cell line that overexpressed the transporter using the probe substrate estradiol 17\beta-glucuronide (EG). CKA at varying concentrations was pre-incubated for 15 min at 37°C, followed by a 2-min incubation in presence of CKA and 5 \mu M EG at 37°C. EG was quantified via liquid chromatography mass spectrometry.\textsuperscript{14} Determined from the human liver microsome using \beta-glucuronidase as a probe substrate.\textsuperscript{25} Determined using MRP2 PREDIVEZ kit (Solvo) and (5/6)-carboxy-Z''-7''-dichlorofluorescein (CDF) as a probe substrate.\textsuperscript{15} Data represent geometric mean and SD of four separate test occasions, each performed in duplicate. Effects of CKA on Mrp2 activity were determined by quantifying inhibition of ATP-dependent uptake (5 min, 37°C) of 10 \mu M CDF into inverted plasma membrane vesicles prepared from Sf21 insect cells expressing Mrp2. CDF was quantified via fluorescence.\textsuperscript{26} Determined from one independent study performed in triplicate. Inhibition of MRP3 activity was determined from inverted membrane vesicles prepared from HEP293 cells expression MRP3 using EG as a probe substrate.

**Simulation of nelfinavir effects on serum bilirubin**

Simulated nelfinavir serum profiles reasonably described the observed data in humans (Supplementary Figure S1 online). Simulated plasma C\textsubscript{max} and AUC were within 1.4-fold of those from the mean observed plasma concentrations in humans.\textsuperscript{29,30}
change from baseline for increasing doses of CKA simulated with combined effects of hepatotoxicity and bilirubin transporter inhibition, hepatotoxicity only, or bilirubin transporter inhibition only.

DISCUSSION

Drug-induced hyperbilirubinemia may occur because of severe DILI. The bilirubin submodel within DILLSym, which was initially designed to predict impaired liver function, reasonably recapitulated hyperbilirubinemia induced by hepatotoxic compounds, such as troglitazone.

However, drugs may increase serum bilirubin by inhibition of enzymes and/or transporters involved in bilirubin disposition in the absence of overt liver injury. In the current study, the bilirubin submodels for human and preclinical species were updated to represent enzyme-mediated and transporter-mediated bilirubin disposition; simulated quantitative effects of impaired function of enzymes/transporters on serum bilirubin levels were optimized using in vivo bilirubin data from humans and animals with impaired bilirubin metabolism and transport. Next, the impact of multiple enzyme/transporter inhibitors, indinavir, nelfinavir, and CKA, on serum bilirubin was simulated by combining PBPK-predicted drug disposition, the bilirubin model constructed in this study, and inhibition constants for enzyme and/or transporters measured in vitro. CKA simulations also incorporated known hepatotoxicity mechanisms to simulate net effects of liver injury and inhibition of bilirubin disposition. For each compound, simulated bilirubin levels were compared to in vivo preclinical/clinical bilirubin data to assess the predictive performance of the current bilirubin model.

Depending on the main bilirubin species in serum, hyperbilirubinemia is classified as unconjugated or conjugated hyperbilirubinemia. Two main reasons for unconjugated hyperbilirubinemia are increased bilirubin production (e.g., hemolytic anemias, hemolysis) and defective hepatic bilirubin metabolism. The current bilirubin submodel reasonably recapitulated unconjugated hyperbilirubinemia in patients with GS and CNS, Gunn rats, and Ugt1a1 knockout mice (Figures 2 and 3). Conjugated hyperbilirubinemia can result from defective hepatobiliary transport of CB, such as cholestatic defects, biliary obstruction, and genetic disease, including DJS and RS. Patients with RS present near complete loss of OATP1B1/1B3 function. At first glance, it seems counterintuitive that impaired function of the uptake transporter increases serum CB because CB is generated in the liver and only excreted into bile via MRP2. The study by van de Steeg et al.

Indinavir and nelfinavir inhibit UGT1A1 and OATP1B1, but only indinavir has been associated with hyperbilirubinemia. Simulations reasonably predicted indinavir-mediated unconjugated hyperbilirubinemia in normal patients, which was more pronounced in individuals possessing GS alleles, consistent with the clinical data.

Rotger et al.
reported a similar extent of increase in median bilirubin levels between normal subjects and patients with GS treated with atazanavir or indinavir (0.75 vs. 0.7–1.0 mg/dL), suggesting that higher bilirubin increases in patients with GS result from additive effects of impaired UGT1A1 function and indinavir-mediated UGT1A1/OATP1B1 inhibition. On the other hand, Zucker et al.
reported greater bilirubin increases in patients with GS compared with normal subjects after indinavir treatment (1.36 ± 1.05 vs. 0.39 ± 0.3 mg/dL), suggesting synergistic effects. Simulations predict slightly greater bilirubin increase in patients with GS.

Figure 6  Simulated and observed peak serum total bilirubin (TB) levels in rats administered 50, 200, or 500 mg/kg chemokine receptor antagonist (CKA). (a) Maximum serum TB fold-change from baseline for increasing doses of CKA. Black dots represent data from preclinical trials and red asterisks denote simulation results for SimPops (n = 294) with combined effects of hepatotoxicity and bilirubin transporter inhibition. (b) Maximum serum TB fold-change from baseline for increasing doses of CKA simulated with combined effects of hepatotoxicity and bilirubin transporter inhibition, hepatotoxicity only, or bilirubin transporter inhibition only.

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compared with normal subjects (0.21–0.30 vs. 0.14 mg/dL), which is more consistent with the Rotger et al.24 data. Underlying reasons for the seemingly synergistic increase in patients with GS reported by Zucker et al.23 still remain to be investigated. The current model also correctly predicted the minimal impact of nelfinavir on serum bilirubin. Although the inhibition potency and systemic exposure were comparable, nelfinavir showed higher protein binding (fraction unbound in plasma: 0.001 vs. 0.36), which led to lower unbound concentrations in the liver and sinusoidal blood, which mainly contributed to differential effects on serum bilirubin.15

CKA induced dose-dependent serum ALT and bilirubin elevations in rats, whereas only modest increases in ALT and no bilirubin increases were observed in humans. In vitro assays showed that CKA inhibited multiple bile acid and bilirubin transporters, inhibited mitochondrial electron transport chain (ETC) function, and induced oxidative stress.22,25 Our previous modeling using DILIsym correctly predicted minimal ALT elevations in humans up to 900 mg, and dose-dependent, reversible ALT elevations in rats administered single oral doses of 50–500 mg/kg CKA.25 CKA modeling with its hepatotoxic mechanistic data combined with bilirubin transporter inhibition correctly predicted minimal bilirubin increases in humans up to 900 mg, and dose-dependent bilirubin increases in rats at 50–500 mg/kg, although fold-changes were underestimated. To determine the underlying mechanism of bilirubin increase in rats, simulations were performed either only with hepatotoxic mechanistic inputs or only with bilirubin transporter inhibition. Simulation results indicate that, although ALT and bilirubin were concomitantly increased, bilirubin increases were mostly due to OATP1B1 and MRP2 inhibition, with minor contributions from liver injury. A previous study22 showed no histological features indicative of overt hepatotoxicity in rats administered up to 500 mg/kg CKA, supporting the conclusion of the current study. Reasonable prediction of serum ALT, a biomarker of liver injury, also validates the quantitative contribution of liver injury assessed by the current model.25

Differences between observed and simulated fold-change bilirubin levels in rats administered CKA are likely due to multiple reasons. The main reason is a lack of metabolite effects in the simulations. CKA glucuronide was also shown to be a potent inhibitor of MRP2, but disposition of CKA glucuronide was not represented in the PBPK framework due to a lack of available pharmacokinetic data. If the metabolite had been included, it is very likely that bilirubin levels would have further increased in the simulations. Another reason could be lack of variability in parameters surrounding the bilirubin model. As currently modeled, all individuals in the SimPops have the same initial bilirubin parameter values and variability in bilirubin metabolism and transport processes were not represented, which leads to minimal bilirubin variability in the simulation results. By design, SimPops were constructed to represent variability in a large number of parameters related to toxicity mechanisms. If parameters related to biomarkers (e.g., bilirubin disposition) are also varied, it would require much larger populations (thousands instead of hundreds) to cover all the parameter spaces, which would significantly increase computation time.

Not representing metabolite effects and interindividual variability in bilirubin disposition are limitations of the current study. Assessment of metabolite effects on hepatotoxicity and enzyme/transporter function could provide reliable predictions, but identification and synthesis of stable metabolites are not feasible especially during early stages of drug development. Also, future efforts will be needed to incorporate variability in baseline bilirubin levels, variability in multiple bilirubin disposition pathways, as well as environmental effects, such as caloric intake,32,33 which may potentially impact serum bilirubin levels. Accurate prediction of the drug concentration in the liver is important in predicting inhibition of hepatic enzyme/transporters, but clinical measurement of liver concentration is challenging and there exists uncertainty in PBPK-predicted liver concentrations of the hepatic uptake transporter substrates based on the plasma profile.34 In the current study, nelfinavir liver concentrations were predicted using a PBPK model trained to the plasma concentrations, which is another limitation. To explore the potential impact of the uncertainty in liver concentration, simulations were performed with up to 10-fold increases in nelfinavir liver concentrations. Nelfinavir still did not induce hyperbilirubinemia, supporting our overall conclusion.

Drugs may induce hyperbilirubinemia due to liver injury, inhibition of bilirubin enzymes and transporters, or a combination of both. Observations of drug-induced hyperbilirubinemia in clinical trials often require thorough assessments of hepatotoxic potential and investigation of underlying mechanisms. This would involve measurement of in vivo liver safety biomarkers and mechanistic in vitro studies. Given the complexity of underlying mechanisms and multifactorial nature of drug-induced hyperbilirubinemia, interpretation and translation of various data is not always straightforward. In the current study, a quantitative systems pharmacology model that incorporates drug disposition, the hepatocyte life cycle, multiple hepatotoxic mechanisms, and mechanistic biomarker models was used to assess underlying mechanisms of drug-induced hyperbilirubinemia. Differential effects of indinavir and nelfinavir on serum bilirubin were correctly predicted, and species differences in CKA-mediated hyperbilirubinemia were reasonably reproduced. Simulations also suggested that bilirubin increases in rats administered CKA were mainly due to bilirubin transporter inhibition with some contribution of liver injury, demonstrating that the current approach may be useful in predicting potential hyperbilirubinemia and assess underlying mechanisms for observed bilirubin increase in vivo.

**METHODS**

**DILIsym bilirubin submodel update**

Representation of bilirubin disposition in DILIsym prior to version 5A was published elsewhere and also described in the Supplementary Methods online.21 In the current study, the bilirubin submodel was updated explicitly to represent the enzyme-mediated and transporter-mediated hepatobiliary disposition of bilirubin. In the bilirubin submodel, both serum and liver bilirubin pools consist of UB and CB. OATP1B1/1B3 facilitates hepatic uptake of UB and CB.4–6 UB can also enter the liver.
by passive diffusion. Conversion of UB to CB in the liver is mediated by UGT1A1. CB is excreted into bile via MRP2 or transported to sinusoidal blood by MRP3. Each of the enzyme-mediated or transporter-mediated processes can be competitively inhibited by drugs or metabolites. Inhibition of OATP1B1/1B3 is dependent on the unbound drug concentration in the liver. hepatic artery concentrations. Inhibition of UGT1A1, MRP2, and MRP3 is dependent on the unbound drug concentration in the liver. Renal excretion and intestinal secretion pathways are added as compensatory excretion pathways for CB and UB, respectively, under hyperbilirubinemia. MRP3 is dependent on the unbound drug concentration in the liver. Hepatic uptake and efflux processes represented by Michaelis-Menten kinetics. Metabolite disposition was not tracked because metabolite effects on bilirubin disposition remain unknown. Thus, it was assumed that only the parent modulated bilirubin disposition. Drug-specific parameters, obtained from the literature or estimated using available plasma and liver PK profiles of indinavir and nelfinavir, are listed in Supplementary Table S2 online. Detailed methods for bilirubin submodel construction can be found in the Supplementary Method online.

Development of physiologically based pharmacokinetic models
PBPK models were developed in DILIsym to describe the disposition of indinavir, nelfinavir, and CKA in humans and rats. Briefly, the PBPK model consisted of a central compartment representing the blood, three liver compartments representing three zones of the liver, and extrahepatic tissues (i.e., muscle, gut, and other tissues). Blood and tissue compartments were linked by blood flow. Tissue distribution of indinavir, nelfinavir, and CKA was represented with tissue partition coefficients; hepatic distribution of nelfinavir was transporter-mediated. Indinavir and nelfinavir are mainly eliminated by hepatic metabolism, which is represented by Michaelis-Menten kinetics. Metabolite disposition was not tracked because metabolite effects on bilirubin disposition remain unknown. Thus, it was assumed that only the parent modulated bilirubin disposition. Drug-specific parameters, obtained from the literature or estimated using available plasma and liver PK profiles of indinavir and nelfinavir, are listed in Supplementary Table S3 online. PBPK parameters for CKA were previously reported. Physiological parameters for tissue volumes and blood flows in the PBPK submodel of DILIsym were published elsewhere.

Simulation of drug-induced hyperbilirubinemia
Indinavir-mediated hyperbilirubinemia was simulated in the baseline human and the baseline rat using PBPK model-predicted exposure and in vitro inhibition constants for UGT1A1, OATP1B1, and MRP2 (Table 1). In human simulations, the baseline human and a simulated patient with GS and CNS and transport (RS and DJs), Bilirubin submodels for the rat, mouse, and dog were calibrated using bilirubin levels in preclinical species lacking specific enzymes or transporters (i.e., TR, ECHR, MRP2-knockout, and Gunn rats; mice lacking Mrp2, Mrp3, Oatp, or Ugt1a1; and dogs with biliary obstruction). Affinities of UB and CB for enzymes and transporters were obtained from literature or optimized, if unavailable. Parameters in the bilirubin submodel, obtained from the literature or estimated using serum bilirubin levels, are listed in Supplementary Table S2 online. Detailed methods for bilirubin submodel construction can be found in the Supplementary Method online.

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CONFLICT OF INTEREST
K.Y., C.B., J.L.W., S.Q.S., and B.A.H. are employees and stockholders of DILIsym Services Inc., which serves as the coordinating member of the DILI-sim Initiative. P.B.W. directs the DILI-sim Initiative and owns equity in DILIsym Services Inc. S.H.S. and J.T.M. are employees of AstraZeneca.

AUTHOR CONTRIBUTIONS
K.Y., C.B., J.L.W., S.H.S., J.T.M., P.B.W., S.Q.S., and B.A.H. wrote the manuscript. K.Y., C.B., J.L.W., S.H.S., J.T.M., P.B.W., S.Q.S., and B.A.H. designed the research. K.Y., C.B., J.L.W., S.H.S., and J.T.M. performed the research. K.Y., C.B., J.L.W., S.H.S., J.T.M., P.B.W., S.Q.S., and B.A.H. analyzed the data.

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