Genomewide Association Study of Simvastatin Pharmacokinetics

Anssi J. H. Mykkänen1,2, Suvi Taskinen1,2, Mikko Neuvonen1,2, Maria Paile-Hyvärinen1,2, E. Katriina Tarkkainen1,2, Tuomas Lilius1,2, Tuija Tapaninen1,2, Janne T. Backman1,2, Aleksi Tornio1,2 and Mikko Niemi1,2,*

We investigated genetic determinants of single-dose simvastatin pharmacokinetics in a prospective study of 170 subjects and a retrospective cohort of 59 healthy volunteers. In a microarray-based genomewide association study with the prospective data, the $\text{SLCO1B1}$ c.521T>C (p.Val174Ala, rs4149056) single nucleotide variation showed the strongest, genomewide significant association with the area under the plasma simvastatin acid concentration-time curve (AUC; $P = 6.0 \times 10^{-10}$). Meta-analysis with the retrospective cohort strengthened the association ($P = 1.6 \times 10^{-17}$). In a stepwise linear regression candidate gene analysis among all 229 participants, $\text{SLCO1B1}$ c.521T>C ($P = 1.9 \times 10^{-15}$) and $\text{CYP3A4}$ c.664T>C (p.Ser222Pro, rs55785340, $\text{CYP3A4}^*2$, $P = 0.023$) were associated with increased simvastatin acid AUC. Moreover, the $\text{SLCO1B1}$ c.463C>A (p.Pro155Thr, rs11045819, $P = 7.2 \times 10^{-6}$) and c.1929A>C (p.Leu643Phe, rs34671512, $P = 5.3 \times 10^{-4}$) variants associated with decreased simvastatin acid AUC. Based on these results and the literature, we classified the volunteers into genotype-predicted $\text{OATP1B1}$ and $\text{CYP3A4}$ phenotype groups. Compared with the normal $\text{OATP1B1}$ function group, simvastatin acid AUC was 273% larger in the poor (90% confidence interval (CI), 137%, 488%; $P = 3.1 \times 10^{-6}$), 40% larger in the decreased (90% CI, 8%, 83%; $P = 0.036$), and 67% smaller in the highly increased function group (90% CI, 46%, 80%; $P = 2.4 \times 10^{-4}$). Intermediate $\text{CYP3A4}$ metabolizers (i.e., heterozygous carriers of either $\text{CYP3A4}^*2$ or $\text{CYP3A4}^*22$ (rs35599367)), had 87% (90% CI, 39%, 152%, $P = 6.4 \times 10^{-4}$) larger simvastatin acid AUC than normal metabolizers.

These data suggest that in addition to no function $\text{SLCO1B1}$ variants, increased function $\text{SLCO1B1}$ variants and reduced function $\text{CYP3A4}$ variants may affect the pharmacokinetics, efficacy, and safety of simvastatin. Care is warranted if simvastatin is prescribed to patients carrying decreased function $\text{SLCO1B1}$ or $\text{CYP3A4}$ alleles.

Study Highlights

**WHAT IS THE CURRENT KNOWLEDGE ON THE TOPIC?**

- The $\text{SLCO1B1}$ c.521T>C variant increases simvastatin exposure and risk for muscle-related adverse effects. However, marked unexplained interindividual variability exists in simvastatin pharmacokinetics.

**WHAT QUESTION DID THIS STUDY ADDRESS?**

- With our genomewide association study of simvastatin pharmacokinetics, we comprehensively assessed the effect of genetic variation on simvastatin pharmacokinetics in 229 healthy volunteers.

**WHAT DOES THIS STUDY ADD TO OUR KNOWLEDGE?**

- $\text{OATP1B1}$ no function variants $\text{SLCO1B1}^*5$ and $\text{SLCO1B1}^*15$ associate with increased simvastatin acid exposure and $\text{OATP1B1}$ increased function variants $\text{SLCO1B1}^*14$ and $\text{SLCO1B1}^*20$ associate with reduced simvastatin acid exposure. In addition, $\text{CYP3A4}^*2$ and $\text{CYP3A4}^*22$ variants associate with increased simvastatin exposure. This is the first study to demonstrate the effects of $\text{SLCO1B1}^*14$, $\text{SLCO1B1}^*20$, and $\text{CYP3A4}^*2$ on simvastatin pharmacokinetics.

**HOW MIGHT THIS CHANGE CLINICAL PHARMACOLOGY OR TRANSLATIONAL SCIENCE?**

- Our findings indicate that $\text{SLCO1B1}$ and $\text{CYP3A4}$ genetic variants affect simvastatin exposure. Simvastatin is probably best avoided in individuals with genetically poor $\text{OATP1B1}$ or $\text{CYP3A4}$ activity, or decreased activity of both $\text{OATP1B1}$ and $\text{CYP3A4}$. Lowering of the simvastatin dose should be considered in individuals with decreased $\text{OATP1B1}$ or $\text{CYP3A4}$ activity.

1Department of Clinical Pharmacology, University of Helsinki and Helsinki University Hospital, Helsinki, Finland; 2Individualized Drug Therapy Research Program, University of Helsinki, Helsinki, Finland. *Correspondence: Mikko Niemi (mikko.niemi@helsinki.fi)

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Simvastatin is a 3-hydroxy-3-methylglutaryl coenzyme A (HMG-CoA) reductase inhibitor widely prescribed for hypercholesterolemia. Simvastatin is usually effective and well-tolerated, but, for some patients, it can cause muscle symptoms ranging from mild myalgia to potentially lethal rhabdomyolysis. Factors increasing simvastatin exposure, such as drug interactions or genetic variation, heighten the risk for adverse effects. Factors increasing simvastatin exposure, such as drug interactions or genetic variation, heighten the risk for adverse effects. In simvastatin pharmacokinetics, marked interindividual variability exists. Simvastatin is an inactive lactone prodrug, which converts to pharmacologically active simvastatin acid. Simvastatin lactone and acid are extensively metabolized by cytochrome P450 (CYP) enzymes, with a minor contribution from CYP2C8 and uridine diphosphate-glucuronosyltransferases (UGT). The CYP3A4*22 (rs3559367) intron variant reduces CYP3A4 expression and associates with increased simvastatin exposure and cholesterol-lowering efficacy. Furthermore, the CYP3A5*3 (rs776746) splice site variant causing loss-of-function for CYP3A5, may elevate simvastatin concentrations.

Simvastatin acid is also a substrate for the organic anion transporting polypeptide 1B1 (OATP1B1), a hepatic influx transporter encoded by the SLCO1B1 gene. The SLCO1B1 c.521T>C (p.Val174Ala, rs4149056) missense variant raises simvastatin acid exposure and increases the risk for adverse effects. Some studies have suggested that simvastatin acid is also a substrate of the P-glycoprotein, an efflux transporter encoded by the ABCB1 gene, although the evidence is contradictory. Evidence as to the effect of genetic variation in ABCB1 on simvastatin pharmacokinetics is unclear.

As the risk for muscle-related adverse effects increases along with simvastatin exposure, understanding the genetic variants affecting simvastatin pharmacokinetics is important. Studies on simvastatin pharmacokinetics have identified several important pharmacogenetic variants, however, these studies have investigated candidate-gene associations, and yet-unrecognized variants may exist. The aim of this study was to comprehensively characterize the effect of genetic variation on simvastatin pharmacokinetics by use of genomewide methods in a relatively large cohort.

METHODS

Subjects and study design
The study sample comprises a prospective simvastatin pharmacokinetic study cohort and a retrospective cohort with earlier published pharmacokinetic data on simvastatin. A total of 170 (83 men and 87 women, age 24.6 ± 4.2 years (mean ± SD), height 174.4 ± 9.4 cm, weight 68.2 ± 11.1 kg, and body mass index 22.3 ± 2.4 kg/m²) who had participated in earlier single-dose pharmacokinetic studies on either 20 mg (24 participants) or 40 mg (35 participants) simvastatin. The volunteers ingested simvastatin tablets after an overnight fast and received a standardized warm meal 4 hours and standardized light meals 7 and 10 hours after simvastatin intake. Competent ethics committees and national authority (Fimea) approved the study protocols, including genetic analyses. Each participant had given written informed consent.

Determination of drug concentrations
Simvastatin lactone, simvastatin acid, 3′,5′-dihydrodiol simvastatin lactone, 3′,5′-dihydrodiol simvastatin acid, simvastatin-D6, and simvastatin acid-D6 were purchased from Toronto Research Chemicals (Toronto, Ontario, Canada).

Prior to analysis, plasma samples were pre-treated using a VersaPlate supported liquid extraction system in 96-Well format (Agilent Technologies, Waldbronn, Germany). A volume of 150 μL of plasma was mixed with 100 μL of iced cold ammonium acetate (100 mM, pH 4.5), containing the internal standards (10 ng/mL broth). The sample mixture was then transferred to extraction plate, incubated for 10 minutes, and eluted three times with 0.5 mL of methyl tert-butyl ether. The sample extract was evaporated and reconstituted in 80 μL of 40% acetonitrile, and a volume of 5 μL was delivered to the analytical column. The chromatographic separation of analytes was carried out on a Luna Omega polar C18 column (100 × 2.1 mm I.D., 1.6 μm particle size; Phenomenex, Torrance, CA) using 5 mM ammonium formate (pH 3.9, adjusted with 98% formic acid) as mobile phase A and acetonitrile as mobile phase B. The flow rate and the column temperature were maintained at 300 μL/min and 40°C. The mobile phase gradient conditions were as follows: 1 minute at 20% B on hold, then a linear ramp from 20% B to 40% B over 3 minutes followed by a second linear ramp to 90% B over 2 minutes, and 1 minute at 90% B before a re-equilibration step back to the initial conditions (20% B). The drug concentrations were determined using a Sciex 5500Qtrap mass spectrometer (ABSciex, Toronto, Ontario, Canada) interfaced with an electrospray ion source. The characteristic multiple reaction monitoring mass-to-charge (m/z) ion transitions were applied for each analyte. Simvastatin lactone (436–285), simvastatin acid (437–303), and 3′,5′-dihydrodiol simvastatin lactone (470–319) were detected in positive mode, and 3′-hydroxy simvastatin lactone (451–335) and 3′,5′-dihydrodiol simvastatin acid (469–353) were detected using negative mode. Isotope labeled simvastatin lactone served as an internal standard for simvastatin lactone and isotope labeled simvastatin acid for all other analytes. The limits of quantification (ng/mL) for simvastatin lactone, simvastatin acid, 3′-hydroxy simvastatin acid, 3′,5′-dihydrodiol simvastatin lactone, and 3′,5′-dihydrodiol simvastatin acid were 0.05, 0.05, 0.25, 0.25, and 0.5, respectively. The day-to-day precision values for all compounds were below 15% (expressed as percent coefficient of variation) and the accuracy within ±15%, except for the lower limit of quantification, for which both precision and accuracy were within ±20%.

Pharmacokinetics
In the prospective study, we calculated the area under the plasma concentration-time curve (AUC), peak plasma concentration (Cmax), and elimination half-life (t1/2) values for simvastatin, simvastatin acid, 3′,5′-dihydrodiol simvastatin lactone, 3′,5′-dihydrodiol simvastatin lactone, simvastatin acid, 3′,5′-dihydrodiol simvastatin lactone, and 3′,5′-dihydrodiol simvastatin acid.
acid, and 3′-hydroxy simvastatin by standard noncompartmental methods (Phoenix WinNonlin, version 6.4; Certara, Princeton, NJ). We calculated the AUC value from 0 hours to infinity (AUC_{0-∞}) for all metabolites, with the exception of 3′,5′-dihydrodiol simvastatin acid and 3′-hydroxy simvastatin acid, for which AUC from 0 to 24 hours (AUC_{0-24hours}) was determined instead, because the t_{1/2} could not be reliably determined for all participants. In our retrospective cohort, concentration data were available only for simvastatin lactone and simvastatin acid, and pharmacokinetic variables were recalculated from the original concentration data.

**Genotyping**

We extracted genomic DNA from whole blood samples using the Maxwell 16 LEV Blood DNA Kit on a Maxwell 16 Research automated nucleic acid extraction system (Promega, Madison, WI). The Institute for Molecular Medicine Finland (FIMM) Technology Centre, University of Helsinki, performed the genomewide genotyping. The volunteers in the prospective study were genotyped with the Illumina HumanCoreExome-24v1-1_A BeadChip (Illumina, San Diego, CA). Quality thresholds for including genotype data in statistical analysis were Hardy–Weinberg equilibrium P value >10^{-5} and proportion of missing ≤0.03. The genomewide statistical analysis included 261,851 single-nucleotide variants (SNVs) with minorallele frequency (MAF) of ≥0.05. The retrospective cohort was genotyped with Illumina GSAMD-24v2-0_A1 beadchip (Illumina). Data for the genomewide association study meta-analysis included 134,431 SNVs present in both genotyping chips. In order to supplement missing genotypes for candidate-gene analysis, we used TaqMan genotyping assays on a QuantStudio 12 K Flex Real-Time PCR system (Thermo Fisher Scientific, Waltham, MA) to genotype the participants for selected variants in ABCB1, CYP3A4, CYP3A5, SLCO1B1, SLCO1B3, UGT1A1, and UGT1A3. For the candidate-gene analysis, we selected missense or suspected functional variants with MAF of ≥0.01 (Table S1).

We computed the SLCO1B1 haplotypes from the c.388A>G (p.Asn130Asp, rs2306283), c.463C>A (p.Pro155Thr, rs11045819), c.521T>C, and c.1929A>C (p.Asn643Asp, rs34671512) SNVs with PHASE version 2.1.20,21 Haplotypes were defined according to Pharmacogene Variation Consortium.22,23 By combining variants with similar effect, it is possible to achieve greater statistical power. Therefore, we divided the participants into genotype-predicted OATP1B1 phenotype groups, as described previously24 (Figure S1). As an exploratory analysis, we then classified heterozygous carriers of either CYP3A4 c.664T>C (p. Ser222Pro, rs55785340, and CYP3A4*22) or rs5599367 (CYP3A4*22) as intermediate CYP3A4 metabolizers. One participant was compound heterozygote for these alleles and classified as a poor CYP3A4 metabolizer. CYP3A4*22 was included as it reduces CYP3A4 expression in the liver25 and has been associated with higher plasma concentrations of simvastatin acid and simvastatin lactone.12

**Statistical analysis**

The data were analyzed with the statistical programs JMP Genomics 8.0 (SAS Institute, Cary, NC) and IBM SPSS Statistics for Windows 27.0 (Armonk, NY). Pharmacokinetic variables and body weight were log-transformed before statistical analysis. We investigated the possible effects of demographic covariates (sex and body weight) on the pharmacokinetics of simvastatin lactone and its metabolites using a forward stepwise linear regression analysis. The P value thresholds of 0.05 and 0.10 were used for entry into and removal from the model. Possible effects of genetic variants on pharmacokinetic variables were investigated using stepwise linear regression analysis. Significant demographic covariates, weight for simvastatin acid, and 3′,5′-dihydrodiol simvastatin acid, sex for 3′,5′-dihydrodiol simvastatin lactone, were set as fixed factors. Genomewide meta-analysis was carried out using the inverse variance fixed effect model. A P value of below 5 × 10^{-8} was considered genomewide significant. The candidate gene analysis was performed without correction for multiple testing and with Bonferroni correction for the number of tested genetic variants. For SLCO1B1 and CYP3A4 genotype-predicted phenotype groups, analysis of variance adjusting for significant demographic and appropriate genotype covariates was carried out with pairwise comparisons with the Fisher’s Least Significant Difference method. A P value of below 0.05 was considered statistically significant.

**RESULTS**

In the prospective study, the AUC_{0-∞} of simvastatin lactone varied 20.1-fold, that of simvastatin acid 21.3-fold, and that of 3′,5′-dihydrodiol simvastatin lactone 21.1-fold, whereas the AUC_{0-24hours} of 3′,5′-dihydrodiol simvastatin acid varied 58.5-fold and that of 3′-hydroxy simvastatin acid 39.5-fold. Body weight was a significant covariate for simvastatin acid AUC_{0-∞} and 3′,5′-dihydrodiol simvastatin acid AUC_{0-24hours}, whereas sex was a significant covariate for 3′,5′-dihydrodiol simvastatin lactone AUC_{0-∞}. In the retrospective cohort, the dose-adjusted AUC_{0-∞} of simvastatin lactone varied 14.5-fold and that of simvastatin acid 20.6-fold.

**Genomewide association study**

In the prospective study, we investigated the associations of 261,030 SNVs on AUC_{0-∞} or AUC_{0-24hours} of simvastatin lactone, simvastatin acid, 3′,5′-dihydrodiol simvastatin lactone, 3′,5′-dihydrodiol simvastatin acid, and 3′-hydroxy simvastatin acid. Two SNVs in the SLCO1B1 gene were genomewide significantly associated with increased simvastatin acid AUC_{0-∞}. The strongest association was observed with the SLCO1B1 c.521T>C missense variant, with a 61% (90% confidence interval (CI), 43%, 82%; P = 6.0 × 10^{-10}) larger AUC_{0-∞} per copy of the variant allele (Figure 1). The SLCO1B1 intron variant, rs4363657, was associated with a 45% (90% CI, 31%, 62%; P = 3.3 × 10^{-8}) increase per variant allele. In the prospective study, no other genomewide significant associations were observed.

In the genomewide association study meta-analysis including the retrospective cohort, we investigated the effects of 136,688 SNVs on simvastatin lactone and simvastatin acid AUC_{0-∞}. Only simvastatin acid AUC_{0-∞} showed genomewide significant associations. Five SNVs in the SLCO1B1-SLCO1B3 locus were associated with an increased simvastatin acid AUC_{0-∞} (Table 1, Figure 1). The SLCO1B1 c.521T>C SNV showed the strongest association with a 68% (90% CI, 52%, 86%; P = 1.6 × 10^{-17}) increase in simvastatin acid AUC_{0-∞} per variant allele. The SLCO1B1 intronic variant rs4363657 was associated with a 46% (90% CI, 33%, 60%; P = 1.2 × 10^{-11}) increase per variant allele. In addition, two SLCO1B3 missense variants c.334C>T (p.Ala112Ser, rs4149117) and c.699A>G (p.Ile233Met, rs7311358) were associated with a 43% (90% CI, 29%, 58%; P = 6.9 × 10^{-5}) increase in simvastatin acid AUC_{0-∞} per variant allele. These variants were in a complete linkage (r^2 = 1, P = 9.8 × 10^{-52}) with each other and the intronic SLCO1B3 variant rs1304539 (r^2 = 0.99, P = 3.9 × 10^{-51}). All SNVs showing genomewide significant associations were in
a strong linkage disequilibrium with \( SLCO1B1 \) c.521T>C: \( rs4363657 (r^2 = 0.53, P = 4.2 \times 10^{-28}) \), \( SLCO1B3 \) c.334G>T, and c.699A>G (\( r^2 = 0.35, P = 3.1 \times 10^{-19} \)).

**Candidate gene association study**

Due to the risk of false negative associations with the conservative \( P \) value threshold in the genomewide analysis, we next performed a candidate gene analysis on simvastatin acid AUC\(_{0-\infty}\) among all the 229 participants. In this analysis, we included missense and suspected functional variants with MAF of \( \geq 0.01 \) in genes suggested to be involved in simvastatin pharmacokinetics (Table S1).

In a stepwise linear regression analysis, the strongest association was observed with \( SLCO1B1 \) c.521T>C, with a 57% increase in simvastatin acid AUC\(_{0-\infty}\) per variant allele (90% CI, 43%, 73%; \( P = 1.9 \times 10^{-13} \); Table 2). In addition, two \( SLCO1B1 \) missense variants were associated with decreased simvastatin acid AUC\(_{0-\infty}\): c.463C>A with a 32% (90% CI, 22%, 41%; \( P = 7.2 \times 10^{-6} \)) and c.1929A>C with a 36% (90% CI, 21%, 48%; \( P = 5.3 \times 10^{-4} \)) decrease per variant allele. \( CYP3A4*2 \) allele was associated with a 60% (90% CI, 14%, 125%; \( P = 0.023 \)) increase in simvastatin acid AUC\(_{0-\infty}\) per variant allele.
Table 2 Results of candidate gene analysis on simvastatin acid AUC_{0–∞} in 229 healthy volunteers

| Variable                        | Effecta (%) | 90% CI (%) | P value | Bonferroni P value | r^2 | n (%) | n (%) |
|--------------------------------|-------------|------------|---------|-------------------|-----|-------|-------|
| Weight (per 10% increase)      | −7.7        | −10.5, 4.8 | 2.3 × 10^−6 | 4.8 × 10^−4  | 0.02 |       |       |
| SLC01B1 c.521T>C               | 57.4        | 43.1, 73.2 | 1.9 × 10^{-11} | 4.0 × 10^{-12} | 0.25 | 64 (27.9) | 11 (4.8) |
| SLC01B1 c.463C>A               | −32.4       | −41.3, −22.2 | 7.2 × 10^{-6}  | 1.5 × 10^{-4}  | 0.31 | 37 (16.2) | 1 (0.4) |
| SLC01B1 c.1929A>C              | −35.9       | −47.9, −21.0 | 5.3 × 10^{-4}  | 0.011            | 0.35 | 17 (7.4) | 0 |
| CYP3A4 c.664T>C                | 60.3        | 14.1, 125.3 | 0.023    | 0.48              | 0.36 | 6 (2.6) | 0 |

AUC_{0–∞}, area under the plasma simvastatin acid concentration-time curve from 0 hours to infinity; CI, confidence interval.
aPer variant allele copy or 10% increase in weight.

Figure 2 Area under the plasma simvastatin acid concentration-time curve from 0 hours to infinity (AUC_{0–∞}) values grouped by genotype predicted organic anion transporting polypeptide 1B1 (OATP1B1) and cytochrome P-450 3A4 (CYP3A4) phenotypes. Horizontal lines indicate estimated marginal means, whiskers indicate 90% confidence intervals, and gray dots indicate individual AUC_{0–∞} values. DF, decreased function; HIF, highly increased function; IF, increased function; IM, intermediate metabolizer; NF, normal function; NM, normal metabolizer; PF, poor function; PM, poor metabolizer.

Effects of OATP1B1 and CYP3A4 phenotype classes on simvastatin acid and lactone

To further elucidate the roles of SLC01B1 and CYP3A4 in simvastatin pharmacokinetics, we then grouped the participants as an exploratory post hoc analysis into SLC01B1 and CYP3A4 genotype-predicted phenotype groups (Figure 2). Only one participant was classified as poor CYP3A4 metabolizer and was therefore excluded from these analyses. In addition, the participant had also decreased OATP1B1 function. This participant had the third largest AUC_{0–∞} of simvastatin acid (76.1 ng/mL × hours), and the second largest AUC_{0–∞} of simvastatin lactone (202.6 ng/mL × hours).

When compared with the normal OATP1B1 function group, the AUC_{0–∞} of simvastatin acid was 273% (90% CI, 505%, 2007%; P = 8.1 × 10^{−4}) higher in the poor function group, 40% (90% CI, 57%, 83%; P = 6.1 × 10^{−6}) lower in the highly increased function group, 451% (90% CI, 231%, 817%; P = 9.3 × 10^{−8}) larger than in the increased function group, and 166% (90% CI, 66%, 326%; P = 7.2 × 10^{−4}) larger than in the decreased function group. The simvastatin acid C_{max} was 273% (90% CI, 141%, 477%; P = 3.1 × 10^{−6}) higher in the poor function group and 73% (90% CI, 57%, 83%; P = 6.1 × 10^{−6}) lower in the highly increased function group than in the normal function group.

In the decreased OATP1B1 function group, the C_{max} of simvastatin lactone was 55% (90% CI, 12%, 115%; P = 0.026) higher compared with the normal function group, but there were no other significant associations between OATP1B1 function groups and pharmacokinetic variables of simvastatin lactone (Table 3). Simvastatin acid / simvastatin lactone AUC_{0–∞} ratio was 49% (90% CI, 13%, 71%; P = 0.04) lower in the highly increased, 54% (90% CI, 33%, 68%; P = 8.1 × 10^{−3}) lower in the increased, and 143% (90% CI, 47%, 301%; P = 0.0039) higher in the poor function group than in the normal function group. We also analyzed the associations of OATP1B1 phenotype classes with simvastatin pharmacokinetics without adjusting for CYP3A4 classes, and the associations were similar but stronger (Table S2).

In the CYP3A4 intermediate metabolizer group, simvastatin lactone AUC_{0–∞} was 63% (90% CI, 11%, 140%; P = 0.036)
### Table 3 Pharmacokinetic variables of simvastatin lactone and simvastatin acid in individuals with different genotype-predicted OATP1B1 phenotypes

| OATP1B1 phenotype (n) | Geometric mean$^a$ (90% CI) | Ratio to normal function (90% CI) | P value |
|-----------------------|-----------------------------|----------------------------------|---------|
| **Simvastatin lactone** | | | |
| $C_{\text{max}}$ | | | |
| Highly increased function (3) | 5.8 (3.3, 10.2) | 0.59 (0.33, 1.08) | 0.15 |
| Increased function (38) | 13.5 (9.4, 19.2) | 1.37 (0.91, 2.06) | 0.21 |
| Normal function (113) | 9.9 (8.0, 12.1) | | |
| Decreased function (63) | 15.3 (11.9, 19.7) | 1.55 (1.12, 2.15) | 0.026 |
| Poor function (11) | 15.0 (9.0, 25.1) | 1.52 (0.88, 2.64) | 0.21 |
| **Simvastatin lactone $t_{\frac{1}{2}}$** | | | |
| Highly increased function (3) | 4.1 (2.2, 7.5) | 0.96 (0.50, 1.83) | 0.91 |
| Increased function (38) | 4.1 (2.8, 6.0) | 0.96 (0.61, 1.49) | 0.87 |
| Normal function (113) | 4.2 (3.4, 5.3) | | |
| Decreased function (63) | 5.0 (3.8, 6.6) | 1.18 (0.83, 1.67) | 0.45 |
| Poor function (11) | 3.9 (2.2, 6.7) | 0.91 (0.59, 1.66) | 0.80 |
| **Simvastatin lactone AUC$_{0-\infty}$** | | | |
| Highly increased function (3) | 23.9 (13.2, 43.3) | 0.64 (0.34, 1.21) | 0.25 |
| Increased function (38) | 51.5 (35.4, 75.0) | 1.38 (0.90, 2.13) | 0.22 |
| Normal function (113) | 37.3 (30.0, 46.3) | | |
| Decreased function (63) | 47.2 (36.1, 61.6) | 1.27 (0.90, 1.78) | 0.26 |
| Poor function (11) | 48.0 (27.9, 82.4) | 1.29 (0.72, 2.31) | 0.47 |
| **Simvastatin acid $C_{\text{max}}$** | | | |
| Highly increased function (3) | 0.6 (0.4, 1.0) | 0.27 (0.17, 0.43) | 6.1×10$^{-6}$ |
| Increased function (38) | 1.3 (1.0, 1.8) | 0.56 (0.41, 0.78) | 0.0036 |
| Normal function (113) | 2.4 (2.0, 2.8) | | |
| Decreased function (63) | 3.1 (2.5, 3.8) | 1.30 (1.01, 1.68) | 0.086 |
| Poor function (11) | 8.9 (5.9, 13.3) | 3.73 (2.41, 5.77) | 1.3×10$^{-6}$ |
| **Simvastatin acid $t_{\frac{1}{2}}$** | | | |
| Highly increased function (3) | 3.9 (2.6, 6.0) | 1.14 (0.73, 1.8) | 0.62 |
| Increased function (38) | 4.2 (3.2, 5.5) | 1.23 (0.90, 1.67) | 0.27 |
| Normal function (113) | 3.5 (3.0, 4.0) | | |
| Decreased function (63) | 4.2 (3.4, 5.0) | 1.20 (0.94, 1.54) | 0.21 |
| Poor function (11) | 3.7 (2.5, 5.5) | 1.08 (0.71, 1.63) | 0.76 |
| **Simvastatin acid AUC$_{0-\infty}$** | | | |
| Highly increased function (3) | 6.1 (3.8, 9.6) | 0.33 (0.20, 0.54) | 2.4×10$^{-4}$ |
| Increased function (38) | 12.5 (9.3, 16.7) | 0.68 (0.48, 0.95) | 0.056 |
| Normal function (113) | 18.4 (15.6, 21.8) | | |
| Decreased function (63) | 25.8 (21.0, 31.7) | 1.40 (1.08, 1.83) | 0.036 |
| Poor function (11) | 68.7 (45.1, 104.8) | 3.73 (2.37, 5.88) | 3.1×10$^{-6}$ |
| **Simvastatin acid / lactone AUC$_{0-\infty}$ ratio** | | | |
| Highly increased function (3) | 0.25 (0.15, 0.42) | 0.51 (0.29, 0.87) | 0.040 |
| Increased function (38) | 0.23 (0.17, 0.32) | 0.46 (0.32, 0.67) | 8.1×10$^{-4}$ |
| Normal function (113) | 0.50 (0.41, 0.60) | | |
| Decreased function (63) | 0.56 (0.45, 0.71) | 1.13 (0.84, 1.52) | 0.49 |
| Poor function (11) | 1.21 (0.76, 1.93) | 2.43 (1.47, 4.01) | 0.0039 |

$^a$The data are geometric estimated marginal means adjusted for weight (simvastatin acid AUC$_{0-\infty}$ and $C_{\text{max}}$) and genotype-predicted cytochrome P-450 3A4 (CYP3A4) phenotype.
higher, and simvastatin acid AUC<sub>0–∞</sub> was 87% (90% CI, 39%, 152%; \(P = 6.4 \times 10^{-4}\)) higher than in the normal metabolizer group (Table 4). The intermediate CYP3A4 metabolizers had 54% (90% CI, 7%, 122%; \(P = 0.05\)) higher simvastatin lactone \(C_{\text{max}}\), and 79% (90% CI, 35%, 139%; \(P = 9.0 \times 10^{-4}\)) higher simvastatin acid \(C_{\text{max}}\) than normal metabolizers. CYP3A4 phenotype was not associated with the half-life of either simvastatin acid or simvastatin lactone, or with the simvastatin acid/lactone AUC<sub>0–∞</sub> ratio.

**Effects of OATP1B1 and CYP3A4 phenotype classes on other simvastatin metabolites**

We next investigated the associations of OATP1B1 and CYP3A4 phenotype classes with simvastatin oxidative metabolite pharmacokinetics from the prospective study. As there were only two participants with highly increased OATP1B1 function and only one poor CYP3A4 metabolizer, they were excluded from the analyses. In the poor OATP1B1 function group, 3′,5′-dihydrodiol simvastatin lactone AUC<sub>0–∞</sub> was 80% (90% CI, 29%, 151%; \(P = 0.0040\)) higher, 3′,5′-dihydrodiol simvastatin acid AUC<sub>0–24 hours</sub> was 105% (90% CI, 27%, 229%; \(P = 0.014\)) higher, and 3′-hydroxy simvastatin acid AUC<sub>0–24 hours</sub> was 133% (90% CI, 57%, 246%; \(P = 5.4 \times 10^{-7}\)) higher than in the normal OATP1B1 function group (Figure 3, Table S3).

The 3′,5′-dihydrodiol simvastatin lactone / simvastatin lactone AUC<sub>0–∞</sub> ratio was 43% (90% CI, 10%, 64%; \(P = 0.046\)) lower in the increased function group than in the normal function group, but OATP1B1 phenotype was not associated with differences in metabolite / parent compound AUC ratios of other metabolites. CYP3A4 phenotype was not significantly associated with the AUC of the oxidative simvastatin metabolites. However, intermediate CYP3A4 metabolizers had 49% (90% CI, 26%, 65%; \(P = 0.0034\)) lower 3′,5′-dihydrodiol simvastatin lactone / simvastatin lactone AUC<sub>0–∞</sub> ratio than normal CYP3A4 metabolizers (Table S4).

**DISCUSSION**

In this study, we investigated genetic variation of simvastatin pharmacokinetics in 229 healthy volunteers. Five SNVs in the \(\text{SLCO1B1-SLCO1B3}\) locus associated genomewide significantly with increased simvastatin acid AUC. The strongest association was observed with the \(\text{SLCO1B1 c.521T>C, OATP1B1 no function}\) variant. In addition, a candidate-gene analysis suggested associations of increased function \(\text{SLCO1B1}\) alleles and decreased function \(\text{CYP3A4}\) alleles with simvastatin acid pharmacokinetics. Of note, simvastatin acid AUC was ~10-fold larger in individuals with poor OATP1B1 function than in those with highly increased OATP1B1 function. To our knowledge, this is the first study...
to demonstrate the effects of $SLCO1B1^{*14}$, $SLCO1B1^{*20}$, and $CYP3A4^{*2}$ on simvastatin pharmacokinetics. Taken together, our data show that $SLCO1B1$ and $CYP3A4$ are major determinants of simvastatin acid pharmacokinetics.

The only genomewide significant associations were observed in the $SLCO1B1$-$SLCO1B3$ locus. The $SLCO1B1$ c.521T>C SNV is probably the causative variant underlying these associations, as it showed the strongest association and is in linkage equilibrium with the other genomewide significant SNVs. The c.521T>C SNV is known to cause nearly complete loss-of-function of OATP1B1,\textsuperscript{25–28} and to increase simvastatin acid plasma concentrations and myopathy risk.\textsuperscript{3–6} Mechanistically, the association can be explained by reduced hepatic uptake of simvastatin acid due to impaired OATP1B1 function.\textsuperscript{26} When compared with individuals with normal OATP1B1 function, those with poor OATP1B1 function (c.521T>C homozygotes) showed 273% higher and those with decreased OATP1B1 function (c.521T>C heterozygotes) showed 40% higher simvastatin acid AUC. These data are consistent with previously published data and highlight the important role of $SLCO1B1$ c.521T>C SNV in simvastatin pharmacokinetics.\textsuperscript{6}

In a candidate-gene analysis, the $SLCO1B1$ c.463C>A and c.1929A>C variants, which define the *14 and *20 haplotypes, were associated with a reduced simvastatin acid AUC. Consistent with our results, these haplotypes have been associated with increased OATP1B1 expression in the liver.\textsuperscript{29} Furthermore, they have been associated with reduced OATP1B1 biomarker concentrations\textsuperscript{24,28} and increased methotrexate clearance in humans.\textsuperscript{30} In addition, $SLCO1B1^{*14}$ has been associated with poor response to methotrexate\textsuperscript{31} and enhanced response to

![Figure 3](https://i.imgur.com/3.png)
Although the associations of *14 and *20 were not genomewide significant, the biological mechanism is plausible and the results are consistent with the literature on other OATP1B1 substrates.28,30 Taken together, the data indicate that SLCO1B1*14 and *20 are increased function variants and enhance the hepatic uptake of simvastatin acid thereby reducing its systemic exposure.

In addition, the candidate-gene analysis showed association of CYP3A4*2 with increased simvastatin acid AUC. However, after correction for multiple testing, the association did not remain significant. Studies on the effects of CYP3A4*2 on CYP3A4 activity in vitro are scarce, but one individual with the CYP3A4*2/*2 genotype was reported to have a low 6β-OH-cortisol/cortisol ratio indicating slow CYP3A4 metabolism.33 In vitro studies suggest that the effect of CYP3A4*2 may be substrate-specific as the clearance of some CYP3A4 substrates is reduced whereas the clearance of others, including testosterone,34,36,42 is unaffected or increased.33,44

Another CYP3A4 variant, CYP3A4*22, has associated with increased plasma concentrations of simvastatin lactone and acid, and enhanced cholesterol-lowering efficacy of atorvastatin, lovastatin, and simvastatin.11,12,45 CYP3A4*22 is an intronic variant, which alters RNA splicing and decreases CYP3A4 expression in the liver.46 Based on these data, we classified heterozygous carriers of either CYP3A4*2 or CYP3A4*22 as intermediate CYP3A4 metabolizers. Grouping rare variants with similar effects provides better statistical power, and a similar approach is commonly used with other CYP enzymes, such as CYP2D6. Nevertheless, as the data on these CYP3A4 variants are limited, the results should be interpreted with caution. In our exploratory analyses, the intermediate CYP3A4 metabolizers had 87% higher simvastatin acid AUC and 63% higher simvastatin lactone AUC.

Our analyses of simvastatin metabolite concentrations suggest that CYP3A4 converts simvastatin lactone to 3′,5′-dihydrodiol simvastatin lactone as the CYP3A4 phenotype was significantly associated with the 3′,5′-dihydrodiol simvastatin lactone / simvastatin lactone AUC ratio. In addition, poor OATP1B1 function was associated with an increased exposure to 3′,5′-dihydrodiol simvastatin lactone, 3′,5′-dihydrodiol simvastatin acid, and 3′-hydroxy simvastatin acid. This suggests that these metabolites may be OATP1B1 substrates, but interconversion between the lactone and acid forms of the metabolites could also explain some of the associations. Literature regarding simvastatin metabolites is scarce, and possible contributions of the metabolites to the efficacy or toxicity of simvastatin are not known. In one study in hypercholesterolemic children and adolescents, the SLCO1B1 c.521T>C SNV showed no association with 3′,5′-dihydrodiol simvastatin lactone pharmacokinetics,47 but there seem to be no other pharmacogenetic studies on the oxidative metabolites of simvastatin. Taken together, our data suggest that CYP3A4 contributes to the formation of 3′,5′-dihydrodiol simvastatin lactone, but plays a minor role only in the formation of 3′-hydroxy simvastatin acid or 3′,5′-dihydrodiol simvastatin acid. This knowledge may be useful for interpreting the results of drug–drug interaction studies using simvastatin as an index substrate for OATP1B1 or CYP3A4.48

In this study, we investigated simvastatin single-dose pharmacokinetics only. According to pharmacokinetic theory, the single-dose AUC should be equal to the steady-state dose-interval AUC. Therefore, the differences in simvastatin exposure seen in the present study between individuals with different SLCO1B1 and CYP3A4 genotypes should be similar during continuous treatment. Moreover, the present study was carried out in young healthy volunteers, whereas statin users are typically older and may use other medications concomitantly. Interindividual variability in simvastatin pharmacokinetics may therefore be even larger in patients than what was observed in the present study.

The participants of this study were White Finnish volunteers. The frequencies of SLCO1B1 and CYP3A4 variants differ between populations.49,50 SLCO1B1 poor function variants are more common in the Northern hemisphere, and SLCO1B1*S is prevalent only in Europe (MAF 2%), Middle East (5%), and North Africa (2%).49 SLCO1B1*5 is present in other populations as well: it is more common in Europe (16%) than in East Asia (12%), and Sub-Saharan Africa (2%).49 SLCO1B1*14 is most common in European (16%) and Middle Eastern (15%) populations, rare in sub-Saharan Africans (4%) and not observed in East Asians.49 SLCO1B1*20 has its highest frequency in North Africans (15%), but it is also present in sub-Saharan African (9%) and European (5%) populations, yet rare in East Asia (0.6%).49 CYP3A4*22 is observed only in Europeans (1%), whereas CYP3A4*22 is most common in Europeans (5%), but also observed in South Asians (0.6%).50 Some populations may also have pharmacogenetic variants affecting simvastatin pharmacokinetics that were not present in this study. Nevertheless, the effects of SLCO1B1 and CYP3A4 variants on simvastatin pharmacokinetics should be similar to our results in other populations, but the proportion of variability in simvastatin pharmacokinetics explained by these variants may differ.

The effect of SLCO1B1 c.521T>C on simvastatin-induced myopathy and rhabdomyolysis risk is well-known.2–5 In order to mitigate the risk of myotoxicity, a lower simvastatin dose is advisable for patients with decreased OATP1B1 function and the use of simvastatin is best avoided in individuals with poor OATP1B1 function. The effects of SLCO1B1*14 and *20 on the clinical response to simvastatin are not known. The pharmacological target of statins, HMG-CoA reductase, is expressed in the liver. Increased hepatic uptake caused by SLCO1B1*14 and *20 may increase simvastatin acid concentration at the active site. The higher hepatic/systemic concentration ratio may enhance the efficacy and reduce the risk for systemic adverse effects of simvastatin.

In contrast to decreased OATP1B1 function, reduced CYP3A4 activity should increase both the hepatocyte and systemic exposure to active simvastatin acid. Decreased CYP3A4 activity may therefore enhance the cholesterol-lowering efficacy of simvastatin in addition to an increase in the adverse effect risk. Our data indicate that the effects of decreased activity CYP3A4 variants on simvastatin pharmacokinetics are quite large. Therefore, lowering of simvastatin dose should be considered for intermediate CYP3A4 metabolizers and poor CYP3A4 metabolizers should probably avoid using simvastatin. From a safety perspective, intermediate CYP3A4 metabolizers with decreased OATP1B1 function should probably also avoid the use of simvastatin as the effect may be additive.
In conclusion, this large pharmacokinetic study confirms the effects of SLCO1B1 poor and decreased function genotypes on simvastatin acid. Furthermore, it shows that SLCO1B1*4 and *20 haplotypes lower the systemic exposure to simvastatin acid. Moreover, this study is the first to suggest an association of CYP3A4*2 with increased simvastatin lactone and acid exposure. Simvastatin may be best avoided in individuals with genetically poor OATP1B1 function or CYP3A4 activity and in those with the combination of decreased OATP1B1 function and CYP3A4 activity. Reduced simvastatin doses should be considered for individuals with decreased OATP1B1 function or CYP3A4 activity.

SUPPORTING INFORMATION
Supplementary information accompanies this paper on the Clinical Pharmacology & Therapeutics website (www.cptjournal.com).

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AUTHOR CONTRIBUTIONS
A.M. and M.Ni. wrote the manuscript. S.T., J.T.B., A.T., and M.Ni. designed the research. S.T., M.Ne., M.P.-H., E.K.T., T.L., T.T., J.T.B., A.T., and M.Ni. performed the research. A.M. and M.Ni. analyzed the data.

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CONFLICTS OF INTEREST
The authors declared no competing interests for this work.

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1. Neuvonen, P.J., Niemi, M. & Backman, J.T. Drug interactions with lipid-lowering drugs: mechanisms and clinical relevance. Clin. Pharmacol. Ther. 80, 565–581 (2006).
2. Voora, D. et al. The SLCO1B1*5 genetic variant is associated with statin-induced side effects. J. Am. Coll. Cardiol. 54, 1609–1616 (2009).
3. Wilke, R.A. et al. The clinical pharmacogenomics implementation consortium: CPIC guideline for SLCO1B1 and simvastatin-induced myopathy. Clin. Pharmacol. Ther. 92, 112–117 (2012).
4. Cooper-DeHoff, R.M. et al. The clinical pharmacogenetics implementation consortium (CPIC) guideline for SLCO1B1, ABCB2, and CYPC29 and statin-associated musculoskeletal symptoms. Clin. Pharmacol. Ther. 111, 1007–1021 (2022).
5. SEARCH Collaborative Group et al. SLCO1B1 variants and statin-induced myopathy—a genomewide study. N. Engl. J. Med. 359, 789–799 (2008).
6. Pasanen, M.K., Neuvonen, M., Neuvonen, P.J. & Niemi, M. SLCO1B1 polymorphism markedly affects the pharmacokinetics of simvastatin acid. Pharmacogenet. Genomics 16, 873–879 (2006).
7. Neuvonen, P.J., Kantalainen, T. & Kivistö, K.T. Simvastatin but not pravastatin is very susceptible to interaction with the CYP3A4 inhibitor itraconazole. Clin. Pharmacol. Ther. 63, 332–341 (1998).
8. Prueksaritanont, T. et al. Glucuronidation of statins in animals and humans: a novel mechanism of statin lactonization. Drug Metab. Dispos. 30, 505–512 (2002).
9. Prueksaritanont, T., Ma, B. & Yu, N. The human hepatic metabolism of simvastatin hydroxy acid is mediated primarily by CYP3A, and not CYP2D6. Br. J. Clin. Pharmacol. 56, 120–124 (2003).
10. Filippula, A.M. et al. Comparative hepatic and intestinal metabolism and pharmacodynamics of statins. Drug Metab. Dispos. 49, 658–667 (2021).
11. Wang, D., Guo, Y., Wrighton, S.A., Cooke, G.E. & Sadee, W. Intrinsic polymorphism in CYP3A4 affects hepatic expression and response to statin drugs. Pharmacogenomics J. 11, 274–286 (2011).
12. Kitzmiller, J.P., Luzum, J.A., Baldassarre, D., Krauss, R.M. & Medina, M.W. CYP3A4*2 and CYP3A5*3 are associated with increased levels of plasma simvastatin concentrations in the cholesterol and pharmacogenetics study cohort. Pharmacogenet. Genomics 24, 486–491 (2014).
13. Hochman, J.H. et al. Interactions of human P-glycoprotein with simvastatin, simvastatin acid, and atorvastatin. Pharm. Res. 21, 1686–1691 (2004).
14. Chen, C. et al. Differential interaction of 3-hydroxy-3-methylglutaryl-coa reductase inhibitors with ABCB1, ABCC2, and OATP1B1. Drug Metab. Dispos. 33, 537–546 (2005).
15. Deng, F. et al. Comparative hepatic and intestinal efflux transport of statins. Drug Metab. Dispos. 49, 750–759 (2021).
16. Kesktalo, J.E., Kurkinen, K.J., Neuvoneni, P.J. & Niemi, M. ABCB1 haplotypes differentially affect the pharmacokinetics of the acid and lactone forms of simvastatin and atorvastatin. Clin. Pharmacol. Ther. 84, 457–461 (2008).
17. Becker, M.L., Visser, L.E., van Schaik, R.H.N., Hofman, A., Uitterlinden, A.G. & Stricker, B.H.C. Common genetic variation in the ABCB1 gene is associated with the cholesterol-lowering effect of simvastatin in males. Pharmacogenom. 10, 1743–1751 (2009).
18. Jiang, F. et al. The influences of SLCO1B1 and ABCB1 genotypes on the pharmacokinetics of simvastatin, in relation to CYP3A4 inhibition. Pharmacogenomics 18, 459–469 (2017).
19. Kesktalo, J.E., Pasanen, M.K., Neuvonen, P.J. & Niemi, M. Different effects of the ABCG2 c.421C→A SNP on the pharmacokinetics of fluvastatin, pravastatin and simvastatin. Pharmacogenom. 10, 1617–1624 (2009).
20. Stephens, M., Smith, N.J. & Donnelly, P. A new statistical method for haplotype reconstruction from population data. Am. J. Hum. Genet. 68, 978–989 (2001).
21. Stephens, M. & Donnelly, P. A comparison of bayesian methods for haplotype reconstruction from population genotype data. Am. J. Hum. Genet. 73, 1162–1169 (2003).
22. Gaedigk, A. et al. The Pharmacogene variation (PharmVar) consortium: incorporation of the human cytochrome P450 (CYP) allele nomenclature database. Clin. Pharmacol. Ther. 103, 399–401 (2018).
23. Gaedigk, A. et al. The evolution of PharmVar. Clin. Pharmacol. Ther. 105, 29–32 (2019).
24. Neuvonen, M., Tornio, A., Hirvensalo, P., Backman, J.T. & Niemi, M. Performance of plasma coproporphyrin I and III as OATP1B1 biomarkers in humans. Clin. Pharmacol. Ther. 110, 1622–1632 (2021).
25. Tirona, R.G., Leake, B.F., Merino, G. & Kim, R.B. Polymorphisms in OATP-C: identification of multiple allelic variants associated with altered transport activity among European- and African-Americans. J. Biol. Chem. 276, 35669–35675 (2001).
26. Niemi, M., Pasanen, M.K. & Neuvonen, P.J. Organic anion transporting polypeptide 1B1: a genetically polymorphic transporter of major importance for hepatic drug uptake. Pharmacol. Rev. 63, 157–181 (2011).
27. Hirvensalo, P. et al. Enantiomeric pharmacogenomics of fluvastatin. Clin. Pharmacol. Ther. 106, 668–680 (2019).
28. Neuvonen, M. et al. Identification of glycochenodeoxycholate 3-O-glucuronide and glycochenodeoxycholate 3-O-glucuronide as highly sensitive and specific OATP1B1 biomarkers. Clin. Pharmacol. Ther. 109, 646–657 (2021).
29. Nies, A.T. et al. Genetics is a major determinant of expression of the human hepatic uptake transporter OATP1B1, but not of OATP1B3 and OATP2B1. *Genome Med.* 5, 1 (2013).

30. Ramsey, L.B. et al. Rare versus common variants in pharmacogenetics: SLC01B1 variation and methotrexate disposition. *Genome Res.* 22, 1–8 (2012).

31. Ramsey, L.B. et al. Association of SLC01B1 *14 allele with poor response to methotrexate in juvenile idiopathic arthritis patients. *ACR Open Rheumatol* 1, 58–62 (2019).

32. Couvert, P. et al. Association between a frequent allele of the gene encoding OATP1B1 and enhanced LDL-lowering response to fluvastatin therapy. *Pharmacogenomics* 9, 1217–1227 (2008).

33. Shchepotina, E.G., Vavilin, V.A., Goreva, O.B. & Lyakhovich, V.V. Some mutations of exon-7 in cytochrome P450 gene 3A4 and their effect on 6beta-hydroxylation of cortisol. *Bull. Exp. Biol. Med.* 141, 701–703 (2006).

34. Sata, F. et al. CYP3A4 allelic variants with amino acid substitutions in exons 7 and 12: evidence for an allelic variant with altered catalytic activity. *Clin. Pharmacol. Ther.* 67, 48–56 (2000).

35. Miyazaki, M., Nakamura, K., Fujita, Y., Guengerich, F.P., Horiuchi, R. & Yamamoto, K. Defective activity of recombinant cytochromes P450 3A4 and 3A4 in oxidation of midazolam, nifedipine, and testosterone. *Drug Metab. Dispos.* 36, 2287–2291 (2008).

36. Fang, P. et al. Functional assessment of CYP3A4 allelic variants on lidocaine metabolism in vitro. *Drug Des. Devel. Ther.* 11, 3503–3510 (2017).

37. Xu, R.-A. et al. Functional characterization of 22 CYP3A4 protein variants to metabolize ibritinib in vitro. *Basic Clin. Pharmacol. Toxicol.* 122, 383–387 (2016).

38. Lin, Q.-M. et al. Characterization of genetic variation in CYP3A4 on the metabolism of cabozantinib in vitro. *Chem. Res. Toxicol.* 32, 1583–1590 (2019).

39. Tang, P.-F. et al. Functional measurement of CYP2C9 and CYP3A4 allelic polymorphism on sildenafil metabolism. *Drug Des. Devel. Ther.* 14, 5129–5141 (2020).

40. Chen, B. et al. Effects of 26 recombinant CYP3A4 variants on brexpiprazole metabolism. *Chem. Res. Toxicol.* 33, 172–180 (2020).

41. Cai, Y. et al. Evaluation of recombinant CYP3A4 variants on the metabolism of oxycodone in vitro. *Chem. Res. Toxicol.* 34, 103–109 (2021).

42. Akiyoshi, T. et al. Comparison of the inhibitory profiles ofitraconazole and cimetidine in cytochrome P450 3A4 genetic variants. *Drug Metab. Dispos.* 39, 724–728 (2011).

43. Han, M. et al. Functional assessment of the effects of CYP3A4 variants on acalabrutinib metabolism in vitro. *Chem. Biol. Interact.* 345, 109559 (2021).

44. Han, M. et al. Functional evaluation of vandetanib metabolism by CYP3A4 variants and potential drug interactions in vitro. *Chem. Biol. Interact.* 350, 109700 (2021).

45. Luzum, J.A. et al. Individual and combined associations of genetic variants in CYP3A4, CYP3A5, and SLC01B1 with simvastatin and simvastatin acid plasma concentrations. *J. Cardiovasc. Pharmacol. Pharmacol.* 66, 80–85 (2015).

46. Wang, D. & Sadee, W. CYP3A4 intronic SNP rs35599367 (CYP3A4*22) alters RNA splicing. *Pharmacogenet. Genomics* 26, 40–43 (2016).

47. Ogungbenro, K., Wagner, J.B., Abdel-Rahman, S., Leeder, J.S. & Galetin, A. A population pharmacokinetic model for simvastatin and its metabolites in children and adolescents. *Eur. J. Clin. Pharmacol.* 75, 1227–1235 (2019).

48. Tornio, A., Filppula, A.M., Niemi, M. & Backman, J.T. Clinical studies on drug-drug interactions involving metabolism and transport: methodology, pitfalls, and interpretation. *Clin. Pharmacol. Ther.* 105, 1345–1361 (2019).

49. Pasanen, M.K., Neuvonen, P.J. & Niemi, M. Global analysis of genetic variation in SLCO1B1. *Pharmacogenomics* 9, 19–33 (2008).

50. Zhou, Y., Ingelman-Sundberg, M. & Lauschke, V.M. Worldwide distribution of cytochrome P450 alleles: a meta-analysis of population-scale sequencing projects. *Clin. Pharmacol. Ther.* 102, 688–700 (2017).