Discovery and Validation of Pyridoxic Acid and Homovanillic Acid as Novel Endogenous Plasma Biomarkers of Organic Anion Transporter (OAT) 1 and OAT3 in Cynomolgus Monkeys

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ABSTRACT

Perturbation of organic anion transporter (OAT) 1- and OAT3-mediated transport can alter the exposure, efficacy, and safety of drugs. Although there have been reports of the endogenous biomarkers for OAT1/3, none of these have all of the characteristics required for a clinical useful biomarker. Cynomolgus monkeys were treated with intravenous probenecid (PROB) at a dose of 40 mg/kg in this study. As expected, PROB increased the area under the plasma concentration-time curve (AUC) of coadministered furosemide, a known substrate of OAT1 and OAT3, by 4.1-fold, consistent with the values reported in humans (3.1- to 3.7-fold). Of the 233 plasma metabolites analyzed using a liquid chromatography–tandem mass spectrometry (LC–MS/MS)–based metabolomics method, 29 metabolites, including pyridoxic acid (PDA) and homovanillic acid (HVA), were significantly increased after either 1 or 3 hours in plasma from the monkeys pretreated with PROB compared with the treated animals. The plasma of animals was then subjected to targeted LC–MS/MS analysis, which confirmed that the PDA and HVA AUCs increased by approximately 2- to 3-fold by PROB pretreatments. PROB also increased the plasma concentrations of hexadecanedioic acid (HDA) and tetradecanedioic acid (TDA), although the increases were not statistically significant. Moreover, transporter profiling assessed using stable cell lines constitutively expressing transporters demonstrated that PDA and HVA are substrates for human OAT1, OAT3, OAT2 (HVA), and OAT4 (PDA), but not OCT2, MATE1, MATE2K, OATP1B1, OATP1B3, and sodium taurocholate co-transporting polypeptide. Collectively, these findings suggest that PDA and HVA might serve as blood-based endogenous probes of cynomolgus monkey OAT1 and OAT3, and investigation of PDA and HVA as circulating endogenous biomarkers of human OAT1 and OAT3 function is warranted.

Introduction

The organic anion transporters (OATs) OAT1 (SLC22A6) and OAT3 (SLC22A8) function as influx transporters that are mainly expressed on the basolateral membrane of renal proximal tubular cells (RPTCs) and mediate cellular uptake of substrates from blood into the cells (Motohashi et al., 2002). OAT1 and OAT3 can serve as the loci of drug-drug interactions (DDIs), and such DDIs can lead to undesired changes in systemic and local exposures of victim drugs and toxins (Morrissey et al., 2013; Nigam et al., 2015). For example, furosemide (FSM) is taken up from the blood in RPTCs via OAT1 and OAT3 (Hasanmejad et al., 2004), and probenecid (PROB), an inhibitor of OAT1 and OAT3, causes significant alterations in the pharmacokinetic parameters of FSM in humans through the increased area under the plasma drug concentration-time curve (AUC), decreased total plasma clearance (CL\textsubscript{TOT}) and renal clearance (CLR), and decreased fraction excreted in the urine (f\textsubscript{R}) (Chennavasin et al., 1979; Smith et al., 1980; Vree et al., 1995). Moreover, a few uremic toxins recently have been demonstrated to be inhibitors of OAT1 and OAT3, and have the potential to inhibit renal secretion clearance of drug substrates of OAT1 and OAT3 in patients with chronic kidney disease (Hsueh et al., 2016). On the other hand, concomitant use of PROB is recommended by the U.S. Food and Drug Administration for reducing the pronounced nephrotoxicity of cidofovir by inhibiting OAT1 and OAT3 to reduce the exposure of cidofovir in RPTCs to an extent that results in an acceptable benefit-risk balance (http://www.accessdata.fda.gov/drugsatfda_docs/label/1999/020638s003lbl.pdf).

ABBREVIATIONS: ACN, acetonitrile; AUC, area under the plasma concentration-time curve; BSA, bovine serum albumin; CL\textsubscript{NPR}, nonrenal plasma clearance; CL\textsubscript{R}, renal clearance; CL\textsubscript{TOT}, total plasma clearance; DDI, drug-drug interaction; Enal-d5, enalapril maleate-d5; FSM, furosemide; HBSS, Hanks’ balanced salt solution; HDA, hexadecanedioic acid; HEK, human embryonic kidney; HILIC, hydrophilic interaction liquid chromatography; HVA, homovanillic acid; LC, liquid chromatography; LC–MS/MS, liquid chromatography–tandem mass spectrometry; MS, mass spectrometry; NTCP, sodium taurocholate co-transporting polypeptide; OAT, organic anion transporter; PBS, phosphate-buffered saline; PDA, pyridoxic acid; PROB, probenecid; QC, quality control; RP, reverse phase; RPTC, renal proximal tubular cell; TDA, tetradecanedioic acid; V\textsubscript{dss}, volume of distribution at steady-state.
Animal models have often been used to assess preclinical drug-OAT interactions. For example, the clinical famotidine-PROB interaction was reproduced in cynomolgus monkeys recently since PROB caused an approximate 90% reduction in the tubular secretion clearance of famotidine in monkeys (4.58 ± 1.25 vs. 0.38 ± 0.36 ml/min per kilogram) (Tahara et al., 2006). Similarly, PROB significantly decreased the renal tubular secretion clearance of famotidine from 2.80 ± 0.36 to 0.37 ± 0.07 ml/min per kilogram in humans. Furthermore, the protective effect of PROB treatment on the nephrotoxicity of cidofovir in humans was able to be recapitulated in monkeys (Lacy et al., 1998). These results suggest that monkey is the more suitable animal model to predict the clinical DDIs involving OAT1 and OAT3.

Recently, endogenous biomarkers have been envisioned as simple, fast, and cost-effective tools to monitor transporter activity in preclinical and clinical settings to facilitate development of a drug candidate (Ito et al., 2012; Müller et al., 2015; Lai et al., 2016; Yee et al., 2016; Bergagnini-Kolev et al., 2017). Therefore, the identification of a sensitive endogenous biomarker for OAT1 and OAT3 would be of great value. In this regard, Imamura et al. (2014) reported that 6α-hydroxyhydrocortisol could be an endogenous probe for OAT3 inhibition evaluation since PROB significantly changed the AUC and CLA of 6α-hydroxy cortisol in healthy subjects. However, 6α-hydroxy cortisol is formed from cortisol by hepatic CYP3A4, and many drugs known to affect this drug-metabolizing enzyme can also change exposure of 6α-hydroxy cortisol (Peng et al., 2011). Very recently, Tsuura et al. (2016) reported that taurine and glycochenodeoxycholate sulfate were endogenous biomarkers of OAT1 and OAT3, respectively. However, the plasma levels of taurine and glycochenodeoxycholate sulfate were not changed significantly by PROB treatment compared with control even though the renal secretion and CLA were significantly decreased (Tsuura et al., 2016). Reduced activities of blood-facing OAT1 and OAT3 by PROB are supposed to increase plasma concentration of a sensitive and selective endogenous probe, mimicking systemic alterations of probe drugs such as FSM. To our knowledge, all published endogenous biomarkers of OAT1 and OAT3 could not recapitulate plasma drug concentration-time profiles in the presence of OAT1 inhibitors. Several features including specificity, sensitivity, predictability, reproducibility, acute response, and accessibility have been considered in the identification and validation of ideal endogenous biomarkers for drug transporters (Chu et al., 2017; Mariappan et al., 2017; Rodrigues et al., 2017). Given the aforementioned reasons, there is a need for novel plasma biomarkers of renal OATs.

In the present studies we provide direct experimental evidence that several organic anionic compounds, including pyridoxic acid (PDA), homovanillic acid (HVA), hexadecanedioic acid (HDA), and tetradecanedioic acid (TDA) are potential endogenous biomarkers of OAT1 and OAT3 in cynomolgus monkeys. Specifically, an untargeted metabolomics analysis was applied to plasma samples to screen endogenous compounds that were associated with OAT1 and OAT3 inhibition in monkeys. Follow-up quantitative liquid chromatography–tandem mass spectrometry (LC-MS/MS) analysis further characterized the time–plasma concentration profiles of selected endogenous compounds (i.e., PDA, HVA, had, and TDA) after PROB administration in cynomolgus monkeys. Moreover, transporter profiling assessed using human embryonic kidney (HEK) 293 cells stably transfected with major human renal and hepatic drug transporters demonstrated that PDA and HVA are OAT1 and OAT3 substrates and thus potential novel plasma endogenous biomarkers of OAT1 and OAT3 inhibition.

Materials and Methods

Materials. [3H]Penciclovir (1.3 Ci/mmol) and [14C]metformin (98 mCi/mmol) were purchased from Moravek Biochemicals, Inc. (Brea, CA), and [3H]para-aminohippuric acid (4.5 Ci/mmol), [3H]estrone-3-sulfate (44.0 Ci/mmol), [3H]estradiol-17b-D-glucuronide (34.3 Ci/mmol), and [3H]holecystokinin octapeptide (97.5 mCi/mmol) were purchased from PerkinElmer Life and Analytical Sciences (Waltham, MA). Nonradiolabeled FSB, PROB, and taurocholic acid, and the corresponding stable isotope-labeled internal standards were purchased from Toronto Research Chemicals Inc. (North York, Ontario, Canada). 4-PDA (≥98%) was purchased from Sigma-Aldrich (St. Louis, MO), and HVA (≥98%) was purchased from Acros Organics (Geel, Belgium). Hexadecanedioic acid-d28, tetradecanedioic acid-d24, and palmitoyl maleate-d5 (Enal-d5) were obtained from CDN Isotopes (Pointe-Claire, Quebec, Canada). Other nonradiolabeled compounds were purchased from either Sigma-Aldrich or Toronto Research Chemicals Inc. and were of analytical grade. Cell culture media and reagents were purchased from Invitrogen (Carlsbad, CA) or Mediatech, Inc., A Corning Subsidiary (Manassas, VA).

Cynomolgus Monkeys Pharmacokinetic FSB-PROB Interaction Study Protocol. To identify and verify the endogenous biomarkers of OAT1 and OAT3, we conducted a series of experiments: cynomolgus monkey FSB-PROB interaction, metabolomics, targeted LC tandem MS (LC-MS/MS), and transporter profiling experiments. The experimental workflow is shown in Fig. 1.

A single-dose, three-period crossover intravenous pharmacokinetic DD1 study was carried out at the Bristol-Myers Squibb Company, and three male cynomolgus monkeys with body weights ranging from 5.3 to 6.0 kg during the study periods were included in this study. The experiment was performed in accordance with the National Institutes of Health guidelines (https://grants.nih.gov/grants/dawr/guide-for-the-care-and-use-of-laboratory-animals.pdf) and approved by the Bristol-Myers Squibb Animal Care and Use Committee. The animals were housed in a temperature- and humidity-controlled room with a 12-hour light/dark cycle.

In the first period, 40 mg/kg PROB dissolved in a sodium hydroxide solution (0.1 N) and titrated to neutral pH with hydrochloric acid was intravenously infused via a femoral vein catheter over 5 minutes (5 ml/kg). Venous blood samples (2 ml) were collected before and 0.08, 0.17, 0.25, 0.5, 0.75, 1, 2, 3, 5, 7, and 24 hours after administration in K2-EDTA-containing tubes. In the second period, after a washout period of 7 days, 2 mg/kg FSB dissolved in saline was intravenously infused via a femoral vein over 5 minutes (5 ml/kg), and blood samples were collected before and 0.08, 0.17, 0.25, 0.5, 0.75, 1, 2, 3, 5, 7, and 24 hours after administration. In the third period, 8 days after administrations of the previous dose of FSB, a FSB saline solution (2 mg/kg) was given by femoral vein infusion over 5 minutes to each monkey 30 minutes after PROB administration (40 mg/kg, i.v. infusion for 5 minutes). Blood was sampled before and 0.08, 0.17, 0.25, 0.5, 0.75, 1, 2, 3, 5, 7, and 24 hours after start of the FSB infusion. Blood samples were spun for 5 minutes at 13,000 rpm within 1 hour to obtain plasma. Plasma samples were stored at −80°C until LC-MS/MS and metabolomics analyses were conducted.

LC-MS/MS Analysis of Furosemide and Probencid. The LC-MS/MS analysis was performed on a SCIEX Triple Quad API-4000 system (AB SCIEX, Framingham, MA) coupled with a Shimadzu LD-30AD ultra-performance LC system (Shimadzu, Columbia, MD). The chromatographic separation was performed on an Agilent Zorbax RRHD SB-C8 column (2.0 × 100 mm, 1.7 μm) from Agilent Technologies (Santa Clara, CA) using mobile phases of 0.1% formic acid in water and 0.1% formic acid in acetonitrile (ACN). The flow rate was 0.7 ml/min and total run time was 3.7 minutes. The LC column was maintained at 60°C. The analytes were monitored using selected reaction monitoring in negative ionization mode with the optimized nebulizing and desolvation gases. The source temperature was set at 400°C, and the declustering potential and collision energy were optimized. FSB, furosemide-d5, and PROB were detected at the selected reaction monitoring transitions of m/z 329.1 → 285.0, 334.1 → 291.0, and 283.9 → 239.9, respectively.

Before the analysis, urine samples were diluted 100-fold into blank plasma and treated as previously described for plasma. The plasma and diluted urine samples were then extracted using protein precipitation. Specifically, the 100 μl plasma and diluted urine samples were mixed with 100 μl of ACN containing 400 nM of furosemide-d5 (internal standard), followed by vortex mixing on a mixer for 5 minutes at room temperature. The mixed solutions were then filtered using a MultiScreen hydrophilic filtration plate (Billerica, Millipore, MA) by
centrifugation at 4000g for 10 minutes, and the filtered solution was then injected (5 μl) onto LC-MS/MS. The assay was qualified over the analytical range of 1–5000 nM using a linear 1/x2 weighed regression.

**Plasma Metabolomic Profiling.** Frozen plasma samples were thawed and 50 μl aliquots were subjected to protein precipitation by the addition of 600 μl of methanol containing 0.1% formic acid and stable-isotopic internal standards. The samples were mixed with a vortex mixer and subjected to centrifugation for 10 minutes at 2700g (4000 rpm). An aliquot (50 μl) of the resulting supernatants was transferred to a 96-well plate for further hydrophilic interaction LC (HILIC) LC-MS analysis and an additional 100 μl was transferred to a separate 96-well plate for reverse phase (RP) LC-MS analysis. Both plates were dried to determine whether the selected probes were selective substrates for human OAT1 and OAT3.

**Schematic overview of experimental workflow for identification and validation of endogenous biomarkers of OAT1 and OAT3.** Plasma samples were collected from monkey FSP-PROB interaction study, and metabolites were quantified by LC-MS/MS. Raw data were extracted, and analyzed by various tools to identify associations between drug treatments and concentrations, determine significant correlations, and integrate results with transporter knowledge. Three selected potential endogenous biomarkers were further validated by LC-MS/MS with deuterated internal standards to confirm metabolomics observations. Transporter uptake studies were performed to determine whether the selected probes were selective substrates for human OAT1 and OAT3.

**Fig. 1.** Schematic overview of experimental workflow for identification and validation of endogenous biomarkers of OAT1 and OAT3. Plasma samples were collected from monkey FSP-PROB interaction study, and metabolites were quantified by LC-MS/MS. Raw data were extracted, and analyzed by various tools to identify associations between drug treatments and concentrations, determine significant correlations, and integrate results with transporter knowledge. Three selected potential endogenous biomarkers were further validated by LC-MS/MS with deuterated internal standards to confirm metabolomics observations. Transporter uptake studies were performed to determine whether the selected probes were selective substrates for human OAT1 and OAT3.
substrate concentration at which the uptake rate is half of the \( V_{\text{max}} \) value has been reported in rat Oat2-mediated transport for HVA (Mori et al., 2003), which is significantly greater than the selected testing concentrations (274 ± 100 vs. 5 \( \mu \)M). No corresponding data were available for PDA. Kinetic transport experiments were conducted under linear-uptake conditions or for shortest incubation duration with acceptable analytical sensitivity (Supplemental Fig. 1).

Cells were grown to confluence in 24-well poly-D-lysine-coated plates 2–3 days after seeding at a cell density of 500,000 cells per well (BD Biosciences, San Jose, CA). All experiments were conducted at 37 °C using a working solution containing Hank’s balanced salt solution (HBSS) supplemented with 10 mM HEPES (pH 7.4) for OAT1-, OAT2-, OAT3-, OCT4-, OCT2-, OATP1B1, OATP1B3, and NTCP-HEK cells, and pH 8.4 for MATE1- and MATE2K-HEK cells, respectively, with a probe substrate, PDA, or HVA. The probe substrates used were \([\text{H}]\)-paraxanthine (OAT1), \([\text{H}]\)-penciclovir (OAT2), \([\text{H}]\)-estrone-3-sulfate (OAT3 and OAT4), \([\text{H}]\)-nicotinamide (OCT2, MATE1, and MATE2K), \([\text{H}]\)-estradiol-17β-D-glucuronide (OATP1B1), \([\text{H}]\)-cholic acid octapeptide (OATP1B3), and taurocholic acid (NTCP). Compounds were dissolved in dimethyl sulfoxide and diluted in HBSS (maximum 0.2% dimethyl sulfoxide). Plating medium was removed, and new monolayers were rinsed twice with prewarmed HBSS. Incubations were started by the addition of 100 \( \mu \)l of substrate prewarmed at 37 °C. After incubation for desired time at 37 °C, the cell monolayers were rinsed three times with 500 \( \mu \)l of ice-cold HBSS. The cells were then lysed with 300 \( \mu \)l buffer (0.1% Triton X-100 or methanol), and compound concentrations in the cell lysates were measured by either liquid scintillation counting (Tri-Carb 2910 TR Liquid Scintillation Analyzer; PerkinElmer Life and Analytical Sciences) or LC-MS/MS as described subsequently.

LC-MS/MS Analysis of PDA and HVA. Stock PDA solution (390 \( \mu \)g/ml, 2.1 \( \mu \)mol) was prepared by dissolving PDA in 0.5% NH₄OH solution. Stock HVA (1.8 mg/ml, 9.5 mM) solution was prepared in water. Stock solutions were stored in the dark at 4 °C. The highest calibration standard (5000 ng/ml) was prepared by diluting appropriate aliquots of PDA and HVA stock solutions to a final volume of 2 ml with 1% bovine serum albumin (BSA) in phosphate-buffered saline (PBS) (pH 7.4). Additional standard solutions were obtained by serial dilution from the 5000 ng/ml standard with 1% BSA in PBS (pH 7.4) to final concentrations of 2500, 1000, 500, 250, 100, 50, 25, 10, and 5 ng/ml. Quality control (QC) samples were also prepared with 1% BSA in PBS (pH 7.4) at three concentration levels: 7.5, 75, and 750 ng/ml. The internal standard stock solution of Enal-d5 was prepared at 1 mg/ml in methanol. The internal standard working solution containing 1000 ng/ml of Enal-d5 was prepared by dilution of internal standard stock solution with methanol and stored at 4°C.

Calibration curves for PDA and HVA were fitted by the 1/x weighted least-squares quadratic. The calibration curves for PDA and HVA ranged from 5 to 5000 ng/ml. All coefficients of determination (\( R^2 \)) of the calibration lines were >0.98. The mean accuracy (percentage of true value) of individual calibrators was ±15%. The lower limit of quantification, defined as the lowest concentration that could be determined with precision and accuracy of ±20%, was 5 ng/ml for PDA and HVA.

Sample stability was also determined for QC samples (\( n = 5 \)) that were extracted and stored in the instrument autosampler under refrigerated conditions (5°C). Samples were stable for 24 hours. Average recovery for QC samples (\( n = 5 \)) prepared with 1% BSA in PBS (pH 7.4) was 101% and 102% for PDA and HVA, respectively.

Urine and plasma samples were diluted by factors of 1:1, 1:2, 1:10 or 1:100 with 1% BSA in PBS (pH 7.4) to ensure the levels were within the range of the calibration curve. Aliquots (50 \( \mu \)l) of diluted animal samples (urine and plasma), calibrator standards, or QCs were transferred to a 2 ml 96-well plate and mixed for 1 minute at 2000 rpm, and then centrifuged at 24,000 g for 5 minutes, 10 minutes at 4000 g. The supernatant (170 \( \mu \)l) was transferred to a 500 \( \mu \)l 96-well plate and dried under heated nitrogen (45°C). Samples were then reconstituted with 50 \( \mu \)l of water/methanol (98.2) containing 0.1% formic acid, vortex mixed for 1 minute at 2000 rpm, followed by 10 minutes of centrifugation at 4000 g at 4°C before MS analysis. To avoid PDA and HVA degradation, all samples were protected from direct light exposure during sample preparation and analysis.

LC-MS/MS analyses were carried out on a Waters Acquity UPLC system, consisting of a Acquity binary solvent manager and Acquity sample manager with an electrospray ionization source. The analytes (10 \( \mu \)l) were separated on an Acquity UPLC BEH130 C18 (2.1 × 100 mm; 1.7 \( \mu \)m particle size) column and eluted by a gradient program as follows: held 2% B for 0.5 minutes, 2% B–20% B for 2.5 minutes, 20% B–98% B for 1 minute, 98% B–2% B for 0.01 minute, and then held 1 minute for equilibration. The column was heated at 45°C, and the flow rate was 500 \( \mu \)l/min. The mobile phase consisted of an aqueous phase (A: 0.1% formic acid in water) and an organic phase (B: 0.1% formic acid in ACN).

The electrospray ionization source was operated in negative ion mode and its main working parameters were set as follows: ion spray voltage, −4.5 kV; ion source temperature, 550°C; declustering potential, −40 V for PDA and HVA and −150 V for Enal-d5; collision energy, −35 V; entrance potential, −10 V; and collision cell exit potential, −10 V. Multiple reactions monitoring measurements of PDA and HVA analytes were performed using individually optimized cone voltage and collision energy. The multiple reactions monitoring precursor/product ion transitions were as follows: m/z 182 > 138.0 for PDA, 181.0 > 122.0 for HVA, and 380.3 > 114.2 for the internal standard, Enal-d5. The dwell time established for each transition was 50 milliseconds. All peak integration and data processing methods were performed using SCIEX Analyst 1.6.2 (Applied Biosystems/MDS SCIEX).

Pharmacokinetic, Transport, and Statistical Analyses. The AUC from zero to 24 hours (AUC_{0–24 h}) was calculated using the mixed trapezoidal rule. The AUC from zero to infinity includes AUC_{0–24 h} and one extrapolated to infinity from the last measured concentration. The volume of distribution at steady-state (V_{ss}) was determined by the noncompartmental method.

\[ V_{\text{dss}} = \frac{Dose \cdot (AUMC)}{(AUC)^2} \]  
(1)

where AUMC is the area under the curve of the first moment of the concentration-time curve. The CL_{TOT} was calculated from the following equation:

\[ \text{CL}_{\text{TOT}} = \frac{Dose}{AUC} \]  
(2)

The pharmacokinetics parameters including AUC_{0–24 h}, AUC_{ss}, and CL_{TOT} for FSM, PROB, PDA, and HVA, following single intravenous administration of PROB and FSM with or without coadministration of PROB were analyzed with a mixed trapezoidal model using the Kinetica program (Thermo Electron, Philadelphia, PA). The CL_{R} was estimated from the following equation:

\[ \text{CL}_{R} = \frac{X_{\text{Enal-d5}-24\ h}}{AUC_{\text{Enal-d5}-24\ h}} \]  
(3)

where \( X_{\text{Enal-d5}-24\ h} \) is the cumulative amount of unchanged FSM excreted in urine over 24 hours. The renal extraction ratio (ER_{R}) of FSM was calculated by the following equation:

\[ \text{ER}_{R} = \frac{\text{CL}_{R}}{f_{sc} \cdot \text{GFR}} \]  
(4)

where \( f_{sc} \) is the fraction of unbound FSM in human plasma reported (i.e., 0.041) (Rane et al., 1978), and GFR is the glomerular filtration rate in cynomolgus monkeys (i.e., 10.4 ml/min) (Davies and Morris, 1993). The fraction of FSM excreted unchanged in the urine (\( f_{sc} \)) was calculated by dividing \( X_{\text{Enal-d5}-24\ h} \) by the dose. The nonrenal plasma clearance (CL_{NR}) was estimated as the difference between the total plasma and renal clearances. The fraction excreted by nonrenal routes (\( f_{NR} \)) was calculated by dividing the nonrenal clearance by the total plasma clearance. Paired Student’s t test was performed to compare pharmacokinetic parameters between groups using GraphPad Prism version 7 (GraphPad Software Inc., San Diego, CA). A P value of less than 0.05 was considered to be statistically significant (*\( P < 0.05 \), **\( P < 0.01 \), and ***\( P < 0.001 \)). To compare pharmacokinetic parameters between groups, the data are also reported as the geometric mean ratio with a two-sided 90% confidence interval.

Transport data represent the results from a single study run in triplicate and a minimum of two experiments on different days. The results are reported as mean ± S.D. (\( n = 3 \)). To estimate the transport kinetics parameters of PDA and HVA into transporter-expressing HEK 293 cells, the transporter-mediated uptake was calculated by subtracting the uptake in MOCK-HEK cells from that in transporter-expressing HEK 293 cells. The following equation was used to estimate the parameters:

\[ V = \frac{V_{\text{max}} \times [S]}{K_m + [S]} \]  
(5)
where \( V \) is the rate of uptake measured at the given concentration; \( V_{\text{max}} \) is the maximal rate of uptake; \( K_m \) represents the Michaelis-Menten constant at which the transport rate is half its maximal value; and \( [S]_i \) is the substrate concentration.

Statistical differences between cell lines or treatments were determined by an unpaired two-tailed Student \( t \) test (GraphPad Prism version 7; GraphPad Software Inc.). A \( P \) value of less than 0.05 was considered to be statistically significant (\(* P < 0.05\), \(** P < 0.01\), and *** \( P < 0.001\)).

**Results**

**Effects of PROB on Pharmacokinetics of FSM in Cynomolgus Monkeys.** The mean FSM plasma concentration-time profiles after intravenous administration of 2 mg/kg of FSM in the absence and presence of PROB (40 mg/kg, i.v.) are illustrated in Fig. 2A. The plasma concentrations of FSM were higher in the presence than in the absence of PROB at all time points and in all animals. This statement was supported by the data shown in Table 1.

![Graph showing the effects of PROB on FSM concentrations](image1)

**Table 1**

<table>
<thead>
<tr>
<th>Analyte/Parameter</th>
<th>PROB Alone</th>
<th>FSM Alone</th>
<th>FSM with PROB</th>
<th>Ratio (90% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>( \text{AUC}_{0-24} ) (( \mu \text{M} \times \text{h} ))</td>
<td>3436 ± 655</td>
<td>3386 ± 860</td>
<td>3386 ± 860</td>
<td>0.98 (0.58–1.6)</td>
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<tr>
<td>( C_{24} ) (( \mu \text{M} ))</td>
<td>7.3 ± 5.7</td>
<td>8.9 ± 6.5</td>
<td>1.1 (0.37–3.2)</td>
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<tr>
<td>( CL_{\text{GFR}} ) (ml/min per kilogram)</td>
<td>0.69 ± 0.12</td>
<td>0.71 ± 0.18</td>
<td>1.0 (0.60–1.7)</td>
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<tr>
<td>( V_{dSS} ) (l/kg)</td>
<td>0.20 ± 0.01</td>
<td>0.21 ± 0.02</td>
<td>1.0 (0.85–1.3)</td>
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<tr>
<td>( T_{1/2} ) (h)</td>
<td>3.7 ± 0.8</td>
<td>3.7 ± 0.9</td>
<td>1.0 (0.87–1.1)</td>
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**CI**, confidence interval; \( ER_{\text{R}} \), renal extraction ratio; NA, not applicable; \( T_{1/2} \), terminal elimination half-life.

**TABLE 1**

Pharmocokinetic parameters of FSM and PROB in cynomolgus monkeys (\( n = 3 \)) following intravenous administration of PROB alone (40 mg/kg), FSM alone (2 mg/kg), and PROB with FSM.

Pharmocokinetic parameters were determined as described in Materials and Methods. Data are shown as mean ± S.D. (\( n = 3 \)). PROB was intravenously dosed 30 minutes prior to FSM administration in the coadministration treatment group.

Fig. 2. (A) Plasma concentration-time curves of FSM administered intravenously alone (2 mg/kg; open circles) or with PROB (40 mg/kg, i.v.; closed circles). (B) Plasma concentration time curves of PROB administered intravenously alone (40 mg/kg; open triangles) or with FSM (2 mg/kg, i.v.; closed triangles). (C) Effects of PROB on urinary exertion rate of FSM administered intravenously alone (2 mg/kg; open bar) or with PROB (40 mg/kg, i.v.; closed bar). Data are shown as mean ± S.D. (\( n = 3 \)). **\( P < 0.01\), significantly different from urinary excretion rate in the absence of PROB.
Consistently, the CLTOT of FSM was reduced significantly by PROB pretreatment (8.8 ± 0.6 vs. 2.1 ± 0.3 ml/min per kilogram; P < 0.001). However, the administration of PROB did not cause a significant difference in Vdss and terminal elimination half-life of FSM although the mean values were decreased by PROB pretreatment (0.33 ± 0.11 vs. 0.21 ± 0.03 l/kg and 5.9 ± 1.4 vs. 4.7 ± 0.8 hour, respectively; P > 0.05) (Table 1).

The mean urinary excretion rates of FSM in cynomolgus monkeys are shown in Fig. 2C. A 2.1- to 2.4-fold reduction of FSM urinary excretion in the presence of PROB was observed (P < 0.01). The CLR and extraction ratio of FSM were reduced significantly with PROB pretreatment (3.7 ± 0.6 vs. 0.44 ± 0.12 ml/min per kilogram and 49.5 ± 8.8 vs. 5.9 ± 1.4, respectively; P < 0.01). Furthermore, the CLNR of FSM in the presence of PROB was also reduced markedly (5.0 ± 0.6 vs. 1.7 ± 0.2 ml/min per kilogram; P < 0.01). Therefore, the reduced CLTOT by PROB pretreatment was not solely due to either decreased CLIR or CLNR.

Figure 2B illustrates the PROB concentration in plasma after intravenous PROB alone and concurrently with FSM. The PROB plasma concentrations 24 hours after PROB administration (C24 h) were not significantly different (7.3 ± 5.7 and 8.9 ± 6.5 μM, P > 0.05) (Table 1). Comparing other PROB pharmacokinetic parameters in the absence or presence of FSM indicates that the mean PROB AUC0–24 h, C24 h, CLTOT, and terminal elimination half-life values were almost identical (Table 1).

### Identification of Potential Plasma Endogenous Biomarkers of OAT1 and OAT3 by Metabolomics

To identify potential endogenous plasma probes of OAT1 and OAT3, LC-MS-based metabolomics was used to determine the alterations in plasma concentrations of endogenous compounds between treatments in cynomolgus monkeys. A total of 233 metabolites of known structural identity by matching accurate mass and retention time values with an in-house database of endogenous metabolites were measured, and the concentration values were used to determine statistical significance using the paired Student’s t-test (Supplemental Table 1). Of those metabolites monitored, 29 endogenous molecules were identified to be present at concentrations at least 3-fold higher at 1 and/or 3 hours in PROB pretreatment groups (administered alone or with FSM) (Table 2), suggesting that the changes of endogenous metabolites in the plasma were associated with inhibition of monkey OAT1 and OAT3, with associated reduction of OAT-mediated renal clearance. However, only some metabolites were able to return to baseline at 24 hours, which were included as a criterion for transporter biomarker candidacy.

Based on the structure similarities, the identified metabolites can be categorized into three subset groups. The first subset of metabolites are long-chain dicarboxylic acids and derivatives (Table 2), which include TDA [HOOC(CH2)12COOH] and HDA [HOOC(CH2)10COOH]. The second subset consists of small acids that are produced by gut microflora.
The metabolites include indole-3 acetic acid, cresol sulfate, phenyl sulfate, phenylacetic acid, indoxyl sulfate, and phenylacetyl glycine. The third are amino acids and derivatives, which include hydroxy isovaleric acid, tyrosine, and aspartic acid. The metabolites whose plasma concentrations were significantly increased by PROB pretreatments include PDA, HVA, glucuronic acid, pantothenic acid, xanthurenic acid, and kynurenic acid (Table 2). The administration of PROB caused increases in plasma PDA and HVA concentrations at 1 and 3 hours (approximately 3- to 6-fold), and the concentrations returned to baseline at 24 hours (Table 2). PDA and HVA represent novel types of endogenous biomarkers not known previously to interact with monkey and human OAT1 and OAT3. Therefore, we conducted follow-up experiments to determine the time course of change in plasma PDA and HVA concentrations and transporter-expressing cell uptake to comprehend the sensitivity and specificity of the probes as described subsequently.

Time-Dependent Effects on Plasma PDA, HVA, HDA, and TDA Levels of PROB Pretreatment. Administration of 40 mg/kg PROB intravenously caused increases in plasma PDA and HVA concentrations in monkeys. The plasma PDA concentration increased by approximately 3-fold over the first 24 hours (113.5 ± 23.3 and 120.0 ± 17.9 vs. 40.5 ± 2.2 μM·h, P < 0.05), and gradually declined to the basal level (1.5 ± 0.3 μM) at 24 hours after PROB pretreatment (Fig. 3A; Table 3). The PDA concentrations were greater at any time points after PROB pretreatment compared with those after FSM administration. Similarly, the increase in plasma HVA concentration was persistent. The plasma HVA concentration peaked at 3 to 4 hours, returning to the baseline (85 ± 32 nM) within 24 hours after PROB pretreatment with approximately 3-fold increase in AUC (6.5 ± 1.0 and 8.7 ± 1.9 vs. 4.1 ± 0.7 μM·h, P < 0.05) (Fig. 3B; Table 3). As shown in Table 3, PROB pretreatment significantly decreased CLR of PDA compared with FSM treatment (P < 0.05). The pretreatment also decreased the CLR value of HVA, although the reduction was not statistically significant.

We also detected the effects of PROB on plasma concentrations of TDA and HDA since they are reported as substrates of OAT1 and OAT3. PROB pretreatment increased plasma concentrations of TDA and other long-chain dicarboxylic acids in monkeys (Table 2). Figure 3, C and D show that PROB pretreatments increase plasma concentrations, although the increases were not statistically significant (Table 3).

Measurement of PDA and HVA Uptake in Stable Cell Lines Constitutively Expressing Renal Transporter. To determine whether PDA and HVA are substrates for major renal drug transporters, the cellular uptake of the molecules was measured at a single concentration (1 and 5 μM for PDA and HVA, respectively) in human OAT1-, OAT2-, OAT3-, OAT4-, OCT2-, MATE1-, MATE2K-, OATP1B1, OATP1B3-, and NTCP-HEK cells after 5-minute incubation. The cellular uptake of PDA and HVA into the HEK cells stably transfected with the cyromolgus monkey renal OATs was not evaluated because in vitro models are not available in the United States (Tahara et al., 2005). The uptake of PDA in OAT1- and OAT3-HEK cells was approximately 55- and 52-fold higher than that in the control cells (1757 ± 703 vs. 30.6 ± 6.7 and 1548 ± 40.0 vs. 29.8 ± 5.9 pmol/mg after 5 minutes, respectively) (Fig. 4A). In addition, the PDA uptake in OAT1- and OAT3-HEK cells is significantly inhibited by PROB, a potent OAT1 and OAT3 inhibitor (Fig. 4A). Furthermore, PDA is a substrate for human OAT4 because the uptake of PDA in OAT4-HEK cells is significantly greater than that in Mock-HEK cells (1.8-fold; P < 0.01) and the uptake is reduced by 1 mM PROB (Fig. 5A). On the other hand, there was no significant OCT2-, OCT2-, MATE1-, and MATE2K-mediated uptake of PDA compared with the control (P > 0.05) (Figs. 4A and 5A). Similarly, the uptake of HVA in OAT1- and OAT3-expressing cells was approximately 45- and 6-fold higher than those in the control cells (8.33 ± 0.40 vs. 0.18 ± 0.02 and 1.99 ± 0.09 vs. 0.18 ± 0.02 pmol/mg after 5 minutes, respectively; P < 0.001) (Fig. 4B). Additionally, there was also significant OCT2-mediated uptake of HVA compared with the control (5-fold; P < 0.001). However, there was no significant OCT2-, MATE1-, and MATE2K-mediated uptake of HVA compared with the control (P > 0.05). Moreover, both PDA and HVA are not substrates for human hepatic transporters OATP1B1, OATP1B3, and NTCP because there are no significant differences in the uptake between the transporter-overexpressing cells and mock cells (P > 0.05) (Supplemental Fig. 2).

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**Figure 3.** Effect of PROB on plasma concentration-time curves of PDA (A), HVA (B), TDA (C), and HDA (D) in three cynomolgus monkeys after intravenous administration of PROB alone (40 mg/kg; open squares), FSM alone (2 mg/kg, close triangles), and PROB with FSM (open circles).
The affinities of OAT1- and OAT3-mediated transport for PDA and HVA were further determined over a range of PDA and HVA concentrations (0.2–500 μM) in OAT1- and OAT3-HEK cells after 2.5-minute incubation. The incubation time was set by the linear-uptake condition and the lower limit of bioanalytical methods (Supplemental Fig. 1). As shown in Fig. 6, the apparent K_m values were 33.0 ± 6.1, 124 ± 15.3 μM for OAT1- and OAT3-mediated uptake of PDA, respectively, while the apparent K_m values of OAT1-, OAT-2, and OAT3-mediated uptake of HVA were 108 ± 6.2, 15.3 ± 1.6, and 438 ± 63 μM, respectively.

**Discussion**

OAT transporters are mainly expressed in RPTCs and mediate active renal secretion of their substrates. Inhibition of OAT transporter function could lead to decreased renal elimination and increased plasma exposure of xenobiotics and endogenous metabolites that are transported by these transporters. The general approach of using cynomolgus monkey as a model for study of human transporter function has become routine and takes advantage of the similarity and function of drug transporters between humans and monkeys (Tahara et al., 2006; Shen et al., 2013, 2016b). It is recognized that such approaches can identify endogenous biomarker candidates that are selective for a specific transporter (Chu et al., 2015; Shen et al., 2016a; Thakare et al., 2017). To demonstrate that cynomolgus monkey permits quantitative prediction of human OAT-mediated DDIs, we measured the effects of the clinically relevant OAT inhibitor PROB on the pharmacokinetics of the OAT1 and OAT3 probe substrate FSM in monkeys, and compared these results with the changes observed in human subjects. As shown in Fig. 2, the intravenous

**Fig. 4.** Profiling of the transport of PDA and HVA by major drug transporters expressed at the basolateral membrane of RPTC. Uptake in the HEK cells stably transfected with the control vector (Mock-HEK), OAT1, OAT2, OAT3, or OCT2 was measured after 5-minute incubation at 37°C with PDA (1 μM) (A), HVA (5 μM) (B), and radiolabeled probe substrates [3H]pyridoxal-5-phosphate (OAT1), 1 μM [3H]estrone-3-sulfate (OAT3), and 2 μM [3H]metformin (OAT2) (C). Incubations were conducted in the absence and presence of 1 mM PROB (OAT1 and OAT3), 100 μM indomethacin (OAT2), or 100 μM pyrimethamine (OCT2) to evaluate the effects of these inhibitors on PDA and HVA uptake. Each value represents the mean ± S.D. (n = 3). *P < 0.05; **P < 0.01; ***P < 0.001 statistically significantly different from uptake in Mock-HEK cells, and #P < 0.05; ##P < 0.01; ###P < 0.001 statistically significantly different from uptake in the absence of an inhibitor.

<table>
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<tr>
<th>Analyte/Parameter</th>
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<th>FSM Alone</th>
<th>FSM with PROB</th>
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<td><strong>Ratio (90% CI)</strong></td>
<td><strong>Value</strong></td>
<td><strong>Ratio (90% CI)</strong></td>
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<td>3.1 (2.2–4.3)</td>
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<td>Cl_{0-24 h} (ml/min per kilogram)</td>
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<td>AUC_{0-24 h} (μM-h)</td>
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<td>1.6 (1.1–2.3)</td>
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<td>2.0 (1.1–3.9)</td>
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CI, confidence interval.

*Ratio of pharmacokinetics parameter of PROB treatment to FSM treatment.

Cage malfunction prevented urine collection from one monkey in the coadministration treatment group (n = 2).

**P < 0.05 and ***P < 0.01, when the parameter was compared with that administered with FSM alone.
administration of PROB generated a plasma total $C_{24\text{h}}$ of 7.3 ± 5.7 μM (Fig. 2; Table 1). Based on the IC$_{50}$ values for both monkey OAT1 and OAT3 (<10 μM) (Tahara et al., 2005), sustained inhibition of OAT1 and OAT3 transport function is anticipated. In agreement with the in vitro inhibition, a pronounced increase in AUC of FSM, a known substrate of OAT1 and OAT3, resulting from PROB pretreatment was evident (Fig. 2; Table 1). At the relevant dose (i.e., 40 mg/kg, i.v.), PROB decreased the FSM CL$_R$ value by 8.3-fold in cynomolgus monkeys. The decreases in the CL$_R$ values of coadministered FSM observed in the clinic were 3.6- to 5.1-fold for PROB (Chennavasin et al., 1979; Smith et al., 1980). Furthermore, PROB decreased the FSM CLR value by 8.3-fold in cynomolgus monkeys pretreated with PROB alone and with FSM (Fig. 2; Table 1). Furthermore, PROB decreased FSM CL$_{NR}$ in monkeys and humans by 2.9- and 1.5-fold, respectively, suggesting inhibition of FSM metabolism in both species. In the current study, we demonstrated that intravenous pretreatment of cynomolgus monkeys with PROB (40 mg/kg) increased in the AUC of FSM by 4.1-fold (Figs. 2 and 3; Tables 1 and 3). Furthermore, we demonstrated that PDA and HVA are substrates for human OAT1, OAT3, OAT2 (HVA), and OAT4 (PDA) but not OCT2, MATE1, MATE2K, OATP1B1, OATP1B3, and NTCP (Figs. 4 and 5; Supplemental Fig. 2). However, we did not study the transport of PDA and HVA by cynomolgus monkey OAT1 and OAT3 because in vitro cell models are not available in the United States. The species-dependent differences in OAT1- and OAT3-mediated transport of PDA and HVA cannot be excluded, although monkey OAT1 and OAT3 exhibit similar transport kinetics (i.e., $K_m$ and $V_{max}$ values) to human orthologs (Tahara et al., 2005). Collectively, the results from the present study demonstrate that circulating PDA and HVA can be potentially used as endogenous biomarkers of OAT1 and OAT3 inhibition.

A few years ago, an investigation was conducted examining the changes of plasma and urine metabolites with murine Oat1 (Slc22a6) deficiency. Using untargeted metabolomics analysis, the researchers identified several physiologically important metabolites, including PDA, that were increased in the plasma from Oat1 knockout mice compared with wild-type animals, suggesting they are substrates for mouse Oat1 (Wikoff et al., 2011). In agreement, PROB increased the plasma HVA levels in rhesus monkey (Bacopoulos et al., 1978). PDA has been shown to undergo active renal secretion in humans, and elevated plasma PDA was observed in patients with renal insufficiency (Coburn et al., 2002). It is worth noting that the present study shows a considerable variation in the urinary excretion and CLR of PDA and HVA in cynomolgus monkeys and humans (Tahara et al., 2005), but not OCT2, MATE1, MATE2K, OATP1B1, OATP1B3, and NTCP (Fig. 5). Profiling of the transport of PDA and HVA by major drug transporters expressed at the apical membrane of RPTC. Uptake in the HEK cells stably transfected with the control vector (Mock-HEK), OAT4, MATE1, or MATE2K was assessed after 5-minute incubation at 37°C with PDA (1 μM) (A), HVA (5 μM) (B), and [3H]estrone-3-sulfate (1 μM) and [14C]metformin (2 μM) (C). Incubations were conducted in the absence and presence of 1 nM PROB (OAT4) or 100 μM pyrimethamine (MATE1 and MATE2K) to evaluate the effects of these inhibitors on PDA and HVA uptake. Each value represents the mean ± S.D. (n = 3). ***P < 0.001 statistically significantly different from uptake in Mock-HEK cells, and $^a$ $^b$ $^c$ S$^s$ $^d$ $^e$ $^f$ P < 0.001 statistically significantly different from uptake in the absence of an inhibitor.
In conclusion, the use of monkey as a transporter-mediated DDI model along with metabolomics has been demonstrated to be a useful approach for transporter biomarker identification. Our investigations showed that PDA and HVA are novel cynomolgus monkey plasma OAT biomarkers, it cannot be assumed that the results translate directly to human subjects. In addition, we cannot exclude that the formation of the biomarkers studied may be affected by PROB. Moreover, PDA is the dead-end catabolite of the B6 vitamins including pyridoxine, pyridoxamine, and pyridoxal (Merrill and Henderson, 1990). Although the plasma PDA baseline level exhibits acceptable inter-individual variability in monkeys and humans (1.6 ± 0.4 μM and 43 ± 17 nM) (Coburn et al., 2002), the use of vitamin B6 supplements causes elevation of PDA levels far above the normal physiologic level (Zempleni, 1995; Coburn et al., 2002). Therefore, the use of vitamin B6 supplements should be excluded in a clinical study with the goal of assessing a change in PDA plasma level.

Emerging metabolomics and genome-wide association data revealed that HDA and TDA, two fatty acid dicarboxylates, were potential endogenous biomarkers of human OATP1B1 (Yee et al., 2016). Although HDA and TDA are substrates of OATP1B1, OAT1 and OAT3 were also involved in the disposition of HDA and TDA (Yee et al., 2016). Consistently, we have observed that administration of rifampin, a selective inhibitor of hepatic OATPs over renal OATs, increased the AUCs of HDA and TDA by approximately 2-fold in human subjects (Shen et al., 2017). Although these results indicated that HDA and TDA are endogenous biomarkers of OATP1B1, the in vitro transport profiling suggested the involvement of OAT1 and OAT3 in the disposition of HDA and TDA (Yee et al., 2016). Metabolomics analysis in this study demonstrated the significant elevations in plasma levels of at least 20 dicarboxylic acids and fatty acids in the monkeys pretreated with PROB (Supplemental Table 1; Table 2). A similar trend was more evident for TDA with significant increases of AUC in the LC-MS/MS data set (Fig. 3C; Table 3). A comparable degree of increase has been observed with HDA (Fig. 3D; Table 3). As a result, HDA and TDA may serve as dual hepatic OATP and renal OAT biomarkers although further studies are needed for the relevance of clinical interactions. The concentration-time profiles for TDA and HDA appear different in the PROB-alone and with FSM groups although the concentrations in both groups are greater than those in that of FSM alone (Fig. 3, C and D). This is likely due to the additional inhibitory effect of FSM toward OAT1- and OAT3-mediated transport of TDA and HDA (Hasannejad et al., 2004; Nieskens et al., 2016).

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Fig. 6. Concentration-dependent uptake of PDA (A) and HVA (B) by human OAT1 and OAT3. Cells were incubated with PDA or HVA (0.23–500 μM) for 2 minutes (linear range). The net OAT1- (open triangle) or OAT3-mediated uptake (open circle) was calculated by subtracting that in mock cells from that in OAT1- or OAT3-HEK cells. The curves represent the best fit of the Michaelis-Menten equation. Each value represents the mean ± S.D. (n = 3).
in this monkey study on the extent of these endogenous compounds after pretreatment with PROB along with the results from stably transporter-overexpressing cell lines that suggest PDA and HVA are candidate biomarkers of OAT1 and OAT3 for use in clinical settings.

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Participated in research design: Shen, Nelson, Oliveira, Lai, Humphreys.
Conducted experiments: Shen, Oliveira, Zhang, Mcanney, Gu, Chen, Su.
Contributed new reagents or analytic tools: Shen, Nelson, Oliveira.
Performed data analysis: Shen, Nelson, Oliveira, Zhang, Shijpкова, Reily, Gan, Marathe, Humphreys.
Wrote or contributed to the writing of the manuscript: Shen, Nelson, Oliveira, Lai, Marathe, Humphreys.

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