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Physiologically-Based Pharmacokinetic Modeling Involving Nonlinear Plasma and Tissue

Binding: Application to Prednisolone and Prednisone in Rats

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a) **Running Title:** PBPK of Prednisolone and Prednisone in Rats

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d) **List of non-standard abbreviations:** CBG, corticosteroid binding globulin; CS, corticosteroid; GR, glucocorticoid receptor; 11 β -HSD, 11beta-hydroxysteroid dehydrogenase; P-gp, P-glycoprotein; PBPK, physiologically-based pharmacokinetic; PNL, prednisolone; PN, prednisone.

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ABSTRACT

The pharmacokinetics (PK) of prednisolone (PNL) exhibit nonlinearity related to plasma protein binding, tissue binding, metabolic interconversion with prednisone (PN), and renal elimination. Blood and 11 tissues were collected from male Wistar rats after steady-state (SS) infusion and after subcutaneous (SC) boluses of 50 mg/kg of PNL. Concentrations of PNL and PN were measured by LC-MS/MS. Plasma and tissue profiles were described using a complex PBPK model. Concentrations of PN and PNL were in rapid equilibrium in plasma and tissues. The tissue partition coefficients (K_p) of PNL calculated from most SC-dosed tissue and plasma areas were similar to steady-state (SS) infusion and *in silico* values. The blood-to-plasma ratio of PNL was 0.71 with similar RBC and unbound plasma concentrations. Plasma protein binding (60 to 90%) was related to corticosteroid-binding globulin (CBG) saturation. Tissue distribution was nonlinear. The equilibrium dissociation constant (K_d) of PNL shared by all tissues was 3.01 ng/mL with the highest binding in muscle, followed by liver, heart, intestine, bone, and the lowest in skin, spleen, fat, kidney, lung and brain. Fat and bone distribution assumed access only to interstitial space. Brain PNL concentrations ($K_p = 0.05$) were low owing to presumed P-glycoprotein-mediated efflux. Clearances of CBG-free PNL were 1789 from liver and 191.2 mL/hr from kidney. The PN/PNL ratio was nonlinear for plasma, spleen, heart, intestine, bone, fat, and linear for the remaining tissues. Our PBPK model with multiple complexities well described the PK profiles of PNL and PN in blood, plasma and diverse tissues.

SIGNIFICANCE STATEMENT

As steroids such as prednisolone and prednisone have similar and complex PK properties in various species, receptors in most tissues, and multiple therapeutic and adverse actions, this PBPK model may provide greater insights into the pharmacodynamic complexities of corticosteroids. The complex properties of these compounds required innovative PBPK modeling approaches that may be instructive for other therapeutic agents.

Introduction

Corticosteroids (CS) have long been frequently and intensively used in clinical practice for various indications such as inflammatory (e.g. rheumatoid arthritis and asthma) and autoimmune diseases (e.g. solid organ transplant rejection and systemic lupus erythematosus) owing to their potent anti-inflammatory and immunosuppressive properties (Czock et al., 2005; Rhen and Cidlowski, 2005; Bergmann et al., 2012; Kadmiel and Cidlowski, 2013). As synthetic glucocorticoids (GC), CS act primarily through genomic mechanisms by binding to cytosolic glucocorticoid receptors (GR) in tissues and causing inhibition of transcriptional factors such as nuclear factor- κ B and activator protein-1, thus suppressing the activation of genes encoding various proinflammatory mediators that promote inflammation and tissue damage (Coutinho and Chapman, 2011; Oakley and Cidlowski, 2011). As a result of the ubiquitous expression of GR across the body, the CS show multiple organ sites of activity (both desired and undesired), of which the commonly occurred dose- and time-dependent side effects including growth retardation, metabolic disturbances, and osteoporosis (Saag et al., 1994) often compromise the overall therapeutic effects and limit the sustained use of CS.

Prednisolone (PNL) and prednisone (PN) are among the top widely prescribed oral CS (Overman et al., 2013). PNL is the biologically active substance, while PN is both the prodrug and inactive metabolite of PNL, with their interconversion mediated by 11 β -hydroxysteroid dehydrogenases (11 β -HSD) (Raza et al., 2010). The pharmacokinetics (PK) of PNL is extremely complex with dose-dependency exhibited in both animals (Frey et al., 1980; Rocci and Jusko, 1981; Boudinot and Jusko, 1986) and humans (Rose et al., 1981). PNL binds to corticosteroid-binding globulin (CBG) with high affinity and low capacity, while it binds to albumin with low affinity and high capacity (Rocci et al., 1980). In humans, the dose-dependent increase in plasma

clearance (CL) and volume of distribution (Vd) of PNL at doses over 20 mg is attributed to the saturable binding to CBG (Rose et al., 1981; Frey and Frey, 1990; Barth et al., 1992). A decreasing trend of CL and Vd with dose occurred in humans at very high doses of PNL (Derendorf et al., 1985), which may be explained by saturable elimination mechanisms (e.g. saturable conversion from PNL to PN and saturable renal elimination) (Frey et al., 1980; Legler et al., 1982). The PK of PNL in rats, when calculated based on total or free plasma concentrations, exhibits nonlinear distribution and elimination, with decreased CL and volume of distribution at steady-state (V_{ss}) and increased free fraction with increasing doses (Boudinot et al., 1986; Boudinot and Jusko, 1986; Huang and Jusko, 1990). Capacity-limited binding to CBG alone cannot account for the decrease in CL and V_{ss} as these are opposite to predictions that these parameters would increase with increased availability of free drug; thus saturable elimination and tissue distribution might be involved. The tissue distribution of PNL was assessed in rabbits, where the SS tissue/unbound plasma concentration ratios were linear for all the examined tissues except the liver (Khalafallah and Jusko, 1984b); consistent results were obtained using rabbit tissue slices under *in vitro* conditions (Khalafallah and Jusko, 1984a).

Considering their clinical importance and the fact that CS exert their pharmacologic actions as a direct consequence of their presence in numerous target tissues (Melby, 1977), it is important to understand the whole-body disposition characteristics of PNL. This requires the direct examination of drug concentration in tissues. Physiologically-based pharmacokinetic (PBPK) modeling, which is based on physiological tissue sizes and blood/plasma flow rates, allows characterization of tissue distribution in a diverse array of situations and also offers a way to extrapolate tissue concentrations from preclinical species to humans (Gerlowski and Jain, 1983; Sager et al., 2015).

This work characterizes the diverse complexities in disposition of PNL and PN in rats using a combination of experimental, whole-body PBPK modeling, and in silico approaches.

Materials and Methods

Reagents and Chemicals

Prednisolone hemisuccinate sodium salt, prednisolone (purity \geq 99%), prednisone (purity \geq 98%), liquid chromatography-mass spectrometry (LC/MS)-grade acetonitrile, methanol and high performance liquid chromatography (HPLC)-grade formic acid and phosphoric acid were purchased from Sigma-Aldrich (St Louis, MO). Prednisolone-d8 (internal standard [IS] for prednisolone, purity \geq 97.71%) and prednisone-d8 (IS for prednisone, purity \geq 98%) were obtained from Toronto Research Chemicals (Toronto, ON, Canada). Milli-Q water (Millipore Corporation, Bedford, MA) was used.

Animals

Twenty-nine healthy male Wistar rats aged 5 to 6 weeks (weighing 225-249 g) were purchased from Envigo (Chicago, IL) and housed individually in the University Laboratory Animal Facility under controlled conditions with free access to water and food. All rats were acclimated for 1 week before the experiments. The research protocol adhered to the *Guide for the Care and Use of Laboratory Animals* (National Research Council, 2011) and were approved by the University at Buffalo Institutional Animal Care and Use Committee.

Experimental design

The prednisolone dosing solution was prepared freshly as the hemisuccinate sodium salt in sterile saline and filtered through 0.22 μ m filters before use. The drug was administered subcutaneously (SC) in a volume of 1 ml/kg.

PNL was first infused SC to two rats through Alzet osmotic pumps at a concentration of 336 mg/mL at a rate of 8 μ L/h for 7 hr in order to achieve steady-state (SS). Animals were then sacrificed by exsanguination. Whole blood was drained from the abdominal aorta into

ethylenediaminetetraacetic acid (EDTA)-containing syringes. The 0.5 mL of whole blood was aliquoted and the remaining blood centrifuged ($2000 \times g$, 15 min) immediately to obtain plasma samples. Various tissues (heart, liver, spleen, lung, kidney, muscle (gastrocnemus), skin, fat, brain, bone (tibia), intestine) were collected, rinsed in ice-cold PBS and gently blotted, weighed, and immediately frozen in liquid nitrogen.

A second group of rats were given a SC bolus dose of 50 mg/kg prednisolone and sacrificed by exsanguination at 0.25, 0.5, 1, 2, 4, 6, 8, 12 and 24 hr (n=3 per time point). The following procedures for tissue collection were the same as those described in the infusion study. To assess the unbound concentrations of PNL and PN in plasma, an aliquot of plasma samples was subject to ultrafiltration using Centrifree micropartition devices (Millipore Corporation, Bedford, MA) with a 30 kDa molecular mass cutoff filter. Aliquots of plasma samples (1 mL) were added into the ultrafiltration devices and incubated for 30 min at 37°C. Then the samples were centrifuged at $2000 \times g$ for 20 min (37°C) to obtain ultrafiltrates for measurement of free PNL plasma concentrations. Tissues were powdered under liquid nitrogen before drug extraction. All samples were stored at -80°C before further analysis.

Drug analysis

A liquid chromatography-tandem mass spectrometry (LC-MS/MS) method was established to simultaneously determine the concentrations of PNL and PN in each of the collected samples. The LC-MS/MS system consisted of Shimadzu (Kyoto, Japan) modules including a binary pump, a degasser, an autosampler, and column oven, and an Applied Biosystems (Foster City, CA) PE/SCIEX API3000 mass spectrometer equipped with a turbo ion spray interface. Sample separations were achieved on a HydroBond AQ C8 reverse phase column (2.1×150 mm, particle size 3 µm; MAC-MOD Analytical, Inc. Chadds Ford, PA). The mobile phase consisted of eluent

A [water/methanol (95:5, v/v) containing 0.1% formic acid] and eluent B [methanol containing 0.1% formic acid] and was pumped at a flow rate of 0.2 mL/min with a gradient elution. The gradient profile was: 0-0.5 min, 30% B; a linear increase to 70% B from 0.5 to 6.5 min; a linear increase to 95% B over 0.1 min; 95% B for 5.4 min; a linear decrease to 30% B over 0.1 min; 30% B for 3.9 min and stop at 16.00 min. The mass spectrometer was operated in the positive ion mode for the detection of ion transitions at m/z 361.2/343.2 (prednisolone), 359.3/341.3 (prednisone), 369.4/351.3 (prednisolone-d8) and 367.3/349.5 (prednisone-d8). The LC-MS/MS system was controlled by Analyst software version 1.4 (Applied Biosystems SCIEX) for data acquisition and analysis.

Linearity was achieved over a concentration range of 1-2000 ng/mL for all samples. The coefficients of variation (CV%) for intra- and inter-day accuracies and precisions were within $\pm 15\%$. The lower limit of quantification (LLOQ) was 1 ng/mL for both PNL and PN. The recovery of the sample pretreatment method approaches 100%. The PNL and PN in various matrices, during freeze-thaw cycles and long-term storage at 4°C or a lower temperature are known to be stable (Difrancesco et al., 2007; Methlie et al., 2013; Yao et al., 2020).

Preparation of plasma and ultrafiltrate samples

Plasma and ultrafiltrate samples (100 μ L) were spiked with 10 μ L of IS working solutions (300 ng/mL) and 100 μ L of 4% phosphoric acid. The mixture was vortexed for 30 s and centrifuged at $13000 \times g$ for 10 min (4°C). A 190 μ L aliquot of the supernatant was then subject to solid-phase extraction (SPE) using Oasis HLB 1 cc (30 mg) cartridges (Waters Corporation, Milford, MA). Prior to extraction, each cartridge was conditioned using 1 mL of methanol and twice followed by 1 mL of water using a Vac Elut SPS24 solid phase extraction manifold (Varian, Palo Alto, CA). The cartridges were washed twice with 500 μ L of water containing 5%

(v/v) methanol and eluted thrice with 500 μ L of methanol. The eluates were dried under nitrogen for 30 min and reconstituted with 100 μ L of methanol/water (30:70 v/v). The mixture was vortexed for 30 s and centrifuged ($13000 \times g$, 10 min, 4°C). Finally, 10 μ L of the supernatant was injected into the LC-MS/MS for analysis.

Preparation of whole blood and tissue samples

The PNL and PN concentrations in whole blood and tissue samples were pretreated using a slightly modified method. Briefly, whole blood samples were directly homogenized at 4°C using a PRO-200 BIO-GEN homogenizer (ProScientific, Oxford, CT), and tissue samples were homogenized in PBS at a 3-fold dilution using a bullet blender (Next Advance, Inc., Troy, NY) at 4°C. Whole blood and tissue homogenates (100 μ L) were spiked with 10 μ L of IS working solutions (300 ng/mL) and 980 μ L of methanol. The mixture was vortexed for 30 s and centrifuged at $14000 \times g$ for 20 min (4°C). A 900 μ L aliquot of the supernatant was then evaporated to dryness. The residue was reconstituted with methanol (50 μ L) and vortexed twice for 30 s, and then diluted with water (450 μ L) and vortexed for another 30 s. A 450 μ L aliquot of the mixture was acidified with 4% phosphoric acid (450 μ L) and centrifuged at $13000 \times g$ for 10 min (4°C). A 850 μ L aliquot of the supernatant was subject to SPE using pre-conditioned Oasis PRiME HLB 1 cc (30 mg) cartridges (Waters Corporation, Milford, MA). The remaining SPE extraction procedures were similar to those described for the plasma samples except that the analytes were eluted 2x with 500 μ L of acetonitrile/methanol (90:10, v/v). The eluates were dried under nitrogen for 30 min and reconstituted with 150 μ L of methanol/water (30:70, v/v). The mixture was vortexed for 30 s and centrifuged ($13000 \times g$, 10 min, 4°C). Finally, 10 μ L of the supernatant was injected into the LC-MS/MS for analysis.

Blood-to plasma ratio (R_B)

The blood-to-plasma concentration ratio (R_B) was obtained using the measured blood (C_{Bl}) and plasma (C_p) concentrations of PNL from the SC bolus and the SC SS infusion studies in male rats:

$$R_B = \frac{C_{Bl}}{C_p} \quad (1)$$

The concentration of PNL in red blood cells (RBC) was calculated according to:

$$C_{RBC} = \frac{[C_{Bl} - C_p \cdot (1 - Hct)]}{Hct} \quad (2)$$

where C_{RBC} is the concentration of PNL in RBC and Hct is the hematocrit of male Wistar rats (0.45) (Kampfmann et al., 2012).

Correction of tissue concentrations

The concentrations of PNL and PN in all measured tissues were corrected for residual blood using:

$$C_T = \frac{C_{T(meas)} \cdot V_{T(meas)} - C_{Bl} \cdot V_{T(meas)} \cdot F_{vv}}{V_T} \quad (3)$$

$$V_T = V_{T(meas)} - V_{T(meas)} \cdot F_{vv} \quad (4)$$

where C_T and $C_{T(meas)}$ are the residual blood-corrected and measured tissue concentrations; $V_{T(meas)}$ is the measured or estimated tissue volume; V_T is the tissue vascular-corrected volume; and F_{vv} is the fractional vascular volume of blood trapped in tissues, which was obtained from the literature concentrations (Bernareggi and Rowland, 1991). The C_T and V_T values were used for further model analysis.

Tissue-to-plasma partition coefficients (K_p)

The values of K_p were calculated and compared using several methods:

1. Calculation from the SS plasma ($C_{P,SS}$) and tissue ($C_{T,SS}$) concentrations in the SC infusion study. For non-elimination organs:

$$K_p = \frac{C_{T,SS}}{C_{P,SS}} \quad (5)$$

For elimination organs (liver and kidney):

$$K_{p,Liver} = \frac{C_{liver,ss}}{C_{P,SS}} \cdot \left(\frac{Q_{liver} + CL_{int}}{Q_{liver}} \right) \quad (6)$$

$$K_{p,Kidney} = \frac{C_{kidney,ss}}{C_{P,SS}} \cdot \left(\frac{Q_{kidney} + CL_k}{Q_{kidney}} \right) \quad (7)$$

where Q_{liver} and Q_{kidney} are the liver and kidney blood flows; CL_{int} and CL_k are the hepatic intrinsic and renal (metabolic) clearances.

2. Apparent K_p values were calculated from the PBPK model-predicted data (detailed description will follow):

$$K_p = \frac{AUC_{T,PBPK}}{AUC_{P,PBPK}} \quad (8)$$

where $AUC_{T,PBPK}$ and $AUC_{P,PBPK}$ are the area under the curve (AUC) of the PBPK model-predicted tissue and plasma curves. The same correction method for liver and kidney K_p as described in Eq. (6)-(7) were applied here based on Eq. (8).

3. In silico methods implemented by the GastroPlusTM PBPK Simulator (version 9.6.2, Simulations Plus Inc., Lancaster, CA) were also used for K_p prediction. Published methods (Poulin and Theil, 2002a; Poulin and Theil, 2002b), (Berezhkovskiy, 2004), (Rodgers and Rowland, 2006) for neutral compounds were used and are listed as Methods 1-3.

Plasma protein binding of PNL

The concentration-dependent plasma protein binding of PNL in various species owing to saturable binding to CBG has been well investigated previously (Sandberg et al., 1957; Rocci et

al., 1980; Rocci and Jusko, 1981; Boudinot and Jusko, 1984; Boudinot and Jusko, 1986). The binding equation is:

$$C_b = \frac{N_C K_C P_C \cdot C_f}{(1 + K_C \cdot C_f)} + N_A K_A P_A \cdot C_f \quad (9)$$

where C_b and C_f are the bound and free plasma PNL concentrations; N is the number of binding sites; K is the association constant; P is the plasma protein concentration; and subscripts C and A refer to CBG and albumin.

The total (C_p) plasma concentrations can be described as:

$$C_p = \frac{N_C K_C P_C \cdot C_f}{(1 + K_C \cdot C_f)} + N_A K_A P_A \cdot C_f + C_f \quad (10)$$

Rearrangement of Eq. (10) gives:

$$C_f = \frac{-(N_A K_A P_A + N_C K_C P_C - K_C C_p + 1) + \sqrt{(N_A K_A P_A + N_C K_C P_C - K_C C_p + 1)^2 + 4(K_C + N_A K_A P_A K_C) C_p}}{2(K_C + N_A K_A P_A K_C)} \quad (11)$$

The CBG-free PNL plasma concentrations (C_{fCBG}) were determined as the sum of albumin-bound ($N_A K_A P_A \cdot C_f$) and free (C_f) PNL plasma concentrations at various total and free drug concentrations:

$$C_{fCBG} = N_A K_A P_A \cdot C_f + C_f \quad (12)$$

Combining Eq. (11) and (12), the CBG-free plasma concentration can be calculated from total plasma concentration as:

$$C_{fCBG} = (N_A K_A P_A + 1) \cdot \frac{-(N_A K_A P_A + N_C K_C P_C - K_C C_p + 1) + \sqrt{(N_A K_A P_A + N_C K_C P_C - K_C C_p + 1)^2 + 4(K_C + N_A K_A P_A K_C) C_p}}{2(K_C + N_A K_A P_A K_C)} \quad (13)$$

Combining Eq. (10) and (12), C_p can be computed from C_{fCBG} by:

$$C_p = C_{fCBG} \cdot \left(\frac{N_C K_C P_C}{\left(1 + \frac{K_C \cdot C_{fCBG}}{N_A K_A P_A + 1}\right) \cdot (N_A K_A P_A + 1)} + 1 \right) \quad (14)$$

Tissue binding of PNL

The nonlinear tissue binding of PNL was expressed by:

$$C_{bT} = \frac{B_{max} \cdot C_{uT}}{K_d + C_{uT}} \quad (15)$$

where C_{bT} and C_{uT} are the bound and free concentrations of PNL in tissue; B_{max} is the binding capacity of the tissue (ng/mL); and K_d is the equilibrium dissociation constant. Therefore, total tissue concentrations (C_T) can be described by:

$$C_T = C_{uT} + \frac{B_{max} \cdot C_{uT}}{K_d + C_{uT}} \quad (16)$$

Assuming only CBG-free drug has access to tissues, and tissue free concentration (C_{uT}) is the same as the CBG-free drug concentration in tissue venous plasma ($C_{fCBG,TV}$), then the tissue-to-CBG free plasma concentration ratio (K_{pu}) is given by:

$$K_{pu} = \frac{C_T}{C_{fCBG,TV}} = \frac{C_T}{C_{uT}} \quad (17)$$

Combining Eq. (16)-(17) yields:

$$K_{pu} = \frac{-(C_T - K_d - B_{max}) + \sqrt{(C_T - K_d - B_{max})^2 + 4K_d C_T}}{2K_d} \quad (18)$$

PBPK Model Development and Modeling Strategy

The proposed PBPK model of prednisolone (Figure 1) consists of 2 plasma/blood compartments (arterial and venous plasma) and 12 tissue compartments (liver, kidney, lung, heart, spleen, intestine, muscle, fat, bone, skin, brain, and remainder). Tissues accounting for about 90% of total body weight were evaluated. Unmeasured tissues were lumped as a Remainder compartment to attain whole body mass-balance. The model was based upon the following observations and assumptions: (1) tissue concentrations are considered to be in instant equilibrium with the CBG-free drug in venous plasma, which is assumed to be in equilibrium with the tissue unbound concentration; (2) circulatory transport occurs by plasma/blood flow; (3) brain has two sub-compartments (vascular and extravascular) with a presumed P-glycoprotein

(P-gp)-mediated efflux process; the distribution of PNL into fat and bone is limited to the interstitial space; (4) only CBG-free PNL is available for tissue distribution and elimination; (5) the elimination of PNL occurs from liver and kidney; (6) tissue binding of PNL is saturable; and (7) interconversion of PNL and PN was in continual rapid equilibrium. Physiological parameters including tissue weights and blood flow rates to different organs were fixed to literature values in the PBPK model (Table 1) (Shah and Betts, 2012).

The model differential equations are listed below, with all initial conditions set to 0 except for the dosing depot in Eq. (33).

Rates of change of drug concentration in tissues. Except for lung, liver, kidney, brain, fat, bone, arterial, and venous plasma, the mass-balance in each tissue was expressed as:

$$V_{T,i} \cdot \frac{dC_{T,i}}{dt} = Q_i \cdot (C_{fCBG,art} - C_{fCBG,TV,i}) \quad (19)$$

where

$$C_{fCBG,TV,i} = \frac{C_{T,i}}{K_{pu,i}} \quad (20)$$

where $V_{T,i}$, Q_i , $C_{T,i}$ and $K_{pu,i}$ are volume, blood flow, PNL concentration and unbound partition coefficient for tissue i ; $C_{fCBG,TV,i}$ is the CBG-free venous plasma concentration leaving tissue i ; $C_{fCBG,art}$ is the CBG-free PNL concentration in arterial plasma, which can be calculated from the total arterial plasma concentration (C_{art}) using Eq. (13).

Rates of change of drug concentrations in the venous and arterial plasma. The total PNL concentration in venous plasma leaving tissue i ($C_{TV,i}$) and entering the venous and arterial plasma pools can be computed according to Eq. (14) and (20) as:

$$C_{TV,i} = \frac{C_{T,i}}{K_{pu,i}} \cdot \left(\frac{N_C K_C P_C}{\left(1 + \frac{K_C \cdot C_{T,i}}{K_{pu,i} (N_A K_A P_A + 1)} \right)} + 1 \right) \quad (21)$$

Therefore, the rates of change of PNL concentration in venous and arterial plasma pools were:

$$V_{ven} \cdot \frac{dC_{ven}}{dt} = Input + (Q_{liver} \cdot C_{TV,liver} + Q_{kidney} \cdot C_{TV,kidney} + Q_{heart} \cdot C_{TV,heart} + Q_{muscle} \cdot C_{TV,muscle} + Q_{fat} \cdot C_{TV,fat} + Q_{bone} \cdot C_{TV,bone} + Q_{skin} \cdot C_{TV,skin} + Q_{remainder} \cdot C_{TV,remainder}) + Q_{brain} \cdot C_{brain,cap} - Q_{lung} \cdot C_{ven} \quad (22)$$

$$V_{art} \cdot \frac{dC_{art}}{dt} = Q_{lung} \cdot (C_{TV,lung} - C_{art}) \quad (23)$$

Rate of change of drug concentrations in the lung:

$$V_{lung} \cdot \frac{dC_{lung}}{dt} = Q_{lung} \cdot (C_{fCBG,ven} - \frac{C_{lung}}{K_{pu,lung}}) \quad (24)$$

where the CBG-free venous plasma concentration of PNL ($C_{fCBG,ven}$) was computed from the total venous plasma concentration (C_{ven}) according to Eq. (13); and Q_{lung} is the lung plasma flow corresponding to the cardiac plasma flow.

Rate of change of drug concentrations in the liver:

$$V_{liver} \cdot \frac{dC_{liver}}{dt} = Q_{liver,art} \cdot C_{fCBG,art} + Q_{spleen} \cdot \frac{C_{spleen}}{K_{pu,spleen}} + Q_{intestine} \cdot \frac{C_{intestine}}{K_{pu,intestine}} - Q_{liver} \cdot \frac{C_{liver}}{K_{pu,liver}} - CL_{int} \cdot \frac{C_{liver}}{K_{pu,liver}} \quad (25)$$

where $Q_{liver} = Q_{liver,art} + Q_{intestine} + Q_{spleen}$.

Rate of change of drug concentrations in the kidney:

$$V_{kidney} \cdot \frac{dC_{kidney}}{dt} = Q_{kidney} \cdot (C_{fCBG,art} - \frac{C_{kidney}}{K_{pu,kidney}}) - CL_k \cdot \frac{C_{kidney}}{K_{pu,kidney}} \quad (26)$$

Rates of change of drug concentration in bone and fat:

$$V_{bone_ISF} \cdot \frac{dC_{bone}}{dt} = Q_{bone} \cdot (C_{fCBG,art} - \frac{C_{bone}}{K_{pu,bone}}) \quad (27)$$

$$V_{fat_ISF} \cdot \frac{dC_{fat}}{dt} = Q_{fat} \cdot (C_{fCBG,art} - \frac{C_{fat}}{K_{pu,fat}}) \quad (28)$$

where V_{bone_ISF} and V_{fat_ISF} are the interstitial volumes of bone and fat.

The measured PNL concentrations in bone ($C_{bone(meas)}$) and fat ($C_{fat(meas)}$) were modeled as:

$$C_{bone(meas)} = \frac{C_{bone} \cdot V_{bone_ISF}}{V_{bone}} \quad (29)$$

$$C_{fat(meas)} = \frac{C_{fat} \cdot V_{fat_ISF}}{V_{fat}} \quad (30)$$

Rates of change of drug concentration in brain capillary and brain:

$$V_{brain\ cap} \cdot \frac{dC_{brain\ cap}}{dt} = Q_{brain} \cdot (C_{art} - C_{brain\ cap}) - CL_{up} \cdot \left(C_{fCBG,brain\ cap} - \frac{C_{brain}}{K_{pu,brain}} \right) + CL_{eff} \cdot \frac{C_{brain}}{K_{pu,brain}} \quad (31)$$

$$V_{brain} \cdot \frac{dC_{brain}}{dt} = CL_{up} \cdot \left(C_{fCBG,brain\ cap} - \frac{C_{brain}}{K_{pu,brain}} \right) - CL_{eff} \cdot \frac{C_{brain}}{K_{pu,brain}} \quad (32)$$

where CL_{up} and CL_{eff} are the bidirectional uptake and unidirectional efflux clearance of PNL in brain; $C_{fCBG,brain\ cap}$ is the CBG-free PNL concentration in brain capillary that can be calculated from total brain capillary concentration ($C_{brain\ cap}$) using Eq. (13).

The SC injection was treated as a bolus into a dosing depot (A_{depot}) as described by:

$$\frac{dA_{depot}}{dt} = -ka \cdot A_{depot}, \quad A_{depot,0} = Dose \cdot F \quad (33)$$

where A_{depot} is the amount of drug to be absorbed into the venous plasma; ka is the first-order absorption rate constant; $A_{depot,0}$ is the initial condition of A_{depot} ; F is the bioavailability of SC dose calculated to be 0.65 from literature-reported iv data in rats (Huang and Jusko, 1990).

Therefore, the Input function in Eq. (22) is:

$$Input = ka \cdot A_{depot} \quad (34)$$

Given the rapid appearance of PN in plasma and tissues upon PNL administration and often parallel profiles, the interconversion between PNL and PN was assumed to be in continual rapid equilibrium. The concentration of PN in tissue i ($C_{PN,i}$) including lung, kidney, muscle,

skin, liver and brain was modeled as a linear fraction of PNL concentration in the corresponding tissue ($C_{PNL,i}$):

$$C_{PN,i} = f_{m,i} \cdot C_{PNL,i} \quad (35)$$

where $f_{m,i}$ is the metabolite/drug ratio of PN/PNL for tissue i .

The PN concentration in arterial plasma, spleen, heart, intestine, bone and fat was fitted to a Michaelis-Menten type equation:

$$C_{PN,i} = \frac{PN_{max,i} \cdot C_{PNL,i}}{PNL_{50,i} + C_{PNL,i}} \quad (36)$$

where $PN_{max,i}$ is the maximal PN concentration achievable and $PNL_{50,i}$ is the PNL concentration achieving 50% $PN_{max,i}$.

The measured PNL concentrations in arterial blood were described as:

$$C_{PNL,bl} = R_B \cdot C_{PNL,p} \quad (37)$$

where $C_{PNL,bl}$ and $C_{PNL,p}$ are the PNL concentration in arterial blood and plasma.

The relationship between systemic (CL_s) and intrinsic clearances (CL_{int}) of CBG-free PNL is:

$$CL_s = \frac{Q_{liver} \cdot CL_{int}}{Q_{liver} + CL_{int}} \quad (38)$$

A similar calculation related renal systemic clearance to kidney blood flow (Q_{kidney}) and renal intrinsic clearance (CL_k).

Model fitting

The blood-to-plasma concentration ratio (R_B) and the metabolite/drug ratio-related parameters (f_m , PN_{max} and PNL_{50}) were estimated by Orthogonal Regression (Package ‘onls’) in R (version 3.4.3) according to Eq. (1) and Eq. (35)-(36), and fixed in the PBPK model. Naïve-pooled plasma and tissue time-concentration data from all rats were analyzed jointly using the established PBPK model. The plasma protein binding parameters ($N_C K_C P_C$, $N_A K_A P_A$ and K_C) were

fixed to literature-reported values (Boudinot and Jusko, 1986), with the estimation of the SC first-order absorption rate constant (ka), hepatic intrinsic (CL_{int}) and renal (CL_k) clearance, tissue binding parameters (K_d and B_{max}), the bidirectional uptake (CL_{up}) and unidirectional efflux (CL_{eff}) clearance of PNL in brain, and metabolite/drug ratio for liver (f_{m_liver}). The whole-body PBPK model fittings involved nonlinear regression using the maximum likelihood algorithm in ADAPT 5 (Biomedical Simulations Resource, Los Angeles, CA) (D'Argenio et al., 2009). The variance model used was:

$$V_i = (\sigma_1 + \sigma_2 \cdot Y_i)^2 \quad (39)$$

where V_i represents the variance of the i th data point; Y_i is the i th model-predicted concentration; σ_1 and σ_2 are variance model parameters. Model selection was based on the goodness-of-fit criteria, which included the Akaike Information Criterion (AIC), visual inspection of the fitted profiles, and CV% of the parameter estimates.

The Adapt code for the PBPK model is provided in the Supplemental Materials.

Results

Distribution of PNL between blood fractions

Figure 2A shows the relationship between the blood-to-plasma concentration ratio (R_B) vs. plasma concentrations of PNL from both the SC bolus and SC infusion studies. The combined average R_B value calculated using Eq. (1) was 0.703 ± 0.10 (n=18) which was very close to the estimated R_B value by Orthogonal Regression (0.71, Table 3). The calculated PNL concentrations in RBC according to Eq. (2) closely matches the measured free drug in plasma (Figure 2B), indicating limited entry of PNL into RBC.

Plasma protein binding of PNL

The plasma protein binding of PNL *in vivo* was assessed by measuring the unbound PNL concentration in plasma samples collected at each time point after SC bolus dosing using ultrafiltration. The protein binding of PNL in rats was concentration-dependent, with a lower bound fraction (60%) at higher total plasma concentrations and 90% bound at low concentrations (Figure 3A). The protein binding parameters of PNL in rats were similar across different studies (Rocci et al., 1980; Boudinot and Jusko, 1984; Boudinot and Jusko, 1986) and thus were applied to relate the bound and unbound PNL plasma concentrations using Eq. (9). The model simulations were in very good agreement with the observed data (Figure 3B). Therefore, these protein binding parameter values (Table 3) were fixed in the subsequent PBPK model to calculate CBG-free drug in plasma. This assumes that tissue uptake of PNL is not restricted to free drug, but includes rapid dissociation from albumin. The PNL is bound to CBG with affinity that is several orders of magnitude higher than that of albumin. There is evidence that albumin-bound drug does not retard steroid diffusion into tissues owing to the rapid dissociation from albumin (Pardridge, 1981). Our previous study showed that administration of exogenous CBG

significantly decreased the apparent CL and V_{ss} of PNL. The PK of methylprednisolone (MPL), which does not bind to CBG, did not change in rats (Ko et al., 1995). Therefore, we assumed that only CBG-free drug has access to tissues and is subject to elimination in the current PBPK model. In contrast, the protein binding of PN was essentially linear with an average bound fraction of 55% (Figure 3A), which is consistent with previous observations in rabbit and human plasma (Ferry and Wagner, 1987).

Application of whole-body PBPK model

The tissue-to-unbound plasma concentration ratios (K_{pu}) decreased in all tissues with CBG-free plasma concentrations obtained using Eq. (13) (calculations not shown), indicating nonlinear tissue distribution. Therefore, nonlinear K_{pu} functions for each tissue were incorporated into the PBPK model in the form of Eq. (18).

The whole-body PBPK model was applied to jointly fit the concentration-time data of PNL and PN from plasma and all tissues after SC bolus of 50 mg/kg PNL. The observed and model-predicted PK profiles are shown in Figure 4. The observed PNL concentrations in arterial blood were fitted using Eq. (37) (Figure 5). In general, this model described all profiles reasonably well with slight overestimation for muscle around the maximum concentration. All model parameters were estimated precisely with reasonable CV% values (Table 3). The absorption of PNL after SC dosing was relatively fast, with an estimated ka value of 2.59 hr^{-1} producing immediate high concentrations. The concentrations of PNL in tissues increased rapidly, indicating that PNL distributes quickly within blood and between blood and tissues upon dosing. The PNL PK profiles in most tissues exhibited similar patterns with a dominant early decline phase and a slow terminal phase. The model-estimated binding equilibrium dissociation constant (K_d) of PNL shared by all tissues was 3.01 ng/mL with highest binding observed in muscle ($B_{max_1} = 690.5$

ng/mL), followed by liver, heart, intestine, and bone ($Bmax_2=188.5$ ng/mL), with the lowest in kidney, lung, spleen, skin, fat and brain ($Bmax_3= 50.8$ ng/mL). Clearances of CBG-free PNL were 1789 mL/hr from liver (CL_{int}) and 191.2 mL/hr from kidney (CL_k). Low brain PNL concentrations ($K_{p,Brain}= 0.05$, Table 4) in contrast to its expected high permeability ($\log P=1.62$) were well captured by incorporating a presumed unidirectional active efflux process ($CL_{eff}=4.58$ mL/hr) based upon the bidirectional uptake process ($CL_{up}=0.13$ mL/hr). The much higher CL_{eff} value as compared to that of CL_{up} indicates that the limited brain distribution of PNL is strongly affected by the active efflux process. The PK profiles of PNL in fat and bone were reasonably characterized by assuming only the interstitial space was available for distribution.

Plasma and tissue PN concentrations at each time point after dosing were well characterized simply as a fraction of PNL concentration reflecting their rapid equilibrium (Figure 4). The metabolite/drug ratio-related parameters (f_m , PN_{max} and PNL_{50}) were fixed in the PBPK model to the estimated values from the individual organ data (Table 2). Orthogonal regression of the PN versus PNL relationships listed in Table 2 resulted in reasonable fittings, but with inexplicable high RSE% values. Using ordinary least-squares regression in Adapt yielded CV% values that were usually less than 30%, but less reliable fittings. Within the assessed concentration range, the conversion from PNL to PN, defined by the PN/PNL ratio, exhibited saturable characteristics in plasma, heart, intestine, spleen, bone and fat, with the lowest PNL_{50} (PNL concentration at half-maximal saturation) in fat and bone and the highest in intestine. While linear PN/PNL ratios were found for lung, kidney, muscle, skin, liver, brain and liver, the linear ratio gave the highest f_m for kidney (0.076, Table 2) and the lowest for liver (0.01, Table 3).

The PNL K_p values obtained using different methods are listed in Table 4. For methods based on *in vivo* data, the K_p for liver and kidney were corrected for tissue elimination. The uncorrected liver and kidney K_p values are also listed. The time- and concentration-averaged K_p values calculated from the PBPK model-fitted AUC values were generally around 1, and similar to those obtained from the SS infusion study and the *in silico* methods for most tissues except for brain and liver. The $K_{p,Brain}$ obtained by *in silico* methods were more than 10-fold higher than those obtained from the *in vivo* data. The corrected $K_{p,Liver}$ (3.67) obtained from the SS infusion study was higher than those calculated by the other methods.

Discussion

The PBPK model incorporating nonlinear plasma and tissue binding well characterized the disposition characteristics of PNL and its major metabolite PN in plasma and 11 tissues of rats following SS infusion and a 50 mg/kg SC bolus dose of PNL. The blood partitioning of PNL ($R_B=0.703/0.71$) is comparable to that of dexamethasone (DEX) ($R_B=0.664$) (Figure 2A) (Song et al., 2020). The RBC uptake of PNL was limited to the unbound drug in plasma (Figure 2B), as also found in humans (Araki et al., 1966) and rabbits (Khalafallah and Jusko, 1984b). Initial modeling was attempted based on blood PNL concentrations, but fittings were much better as presented. The model thus assumes that CBG-free drug is available for distribution from both plasma and RBC.

Moderate binding to cellular constituents may account for tissue distribution of PNL, a neutral molecule. The K_{pu} declined with higher free plasma concentrations and the slow terminal decline of plasma and tissue concentrations required nonlinear tissue binding for best fittings. Preliminary model fittings without considering nonlinear tissue binding resulted in more rapid decline of the terminal phases in all tissues to below the LLOQ by 4 hr after dosing. The rabbit exhibited linear distribution of PNL into tissues except liver; but these studies were performed over a small range of free plasma concentrations (66-1213 ng/mL) (Khalafallah and Jusko, 1984b).

The moderate lipid solubility ($\log P=1.62$) and molecular weight (360.4) allow for rapid equilibrium of PNL between blood and tissues, as indicated by the rapid increase in PNL concentrations in tissues (Figures 4 and 5). Similar PNL profiles across tissues allowed use of one universal tissue binding equilibrium dissociation constant (K_d) and three groups of binding capacities (B_{max}). The GR, which are ubiquitously expressed in tissues and most abundant in the

liver (Ballard et al., 1974), may contribute to the tissue binding of PNL. We have shown that the *in vitro* K_d value for the PNL-GR interaction in liver is about 61 nM (Boudinot et al., 1986). Also, 11 β -HSD, which interconverts biologically active GC (cortisol and PNL) with their inactive counter-parts (cortisone and PN), may also contribute to tissue binding. There are two distinct 11 β -HSD enzymes, of which 11 β -HSD1 is a bidirectional enzyme but primarily an activator of GC, while 11 β -HSD2 unidirectionally inactivates GC (Raza et al., 2010). The affinity constant (K_d) of the weaker binding cortisol to human 11 β -HSD2 is 25~55 nM (Chapman et al., 2013). The PBPK model-estimated K_d of PNL (3.01 ng/mL= 8.4 nM) is close to this and the GR K_d value. Further, using a highly purified rat plasma membrane free of CBG and GR, the presence of a GC-responsive site mediating high affinity active uptake of PNL was demonstrated in rat liver (Lackner et al., 1998). Thus, there are several possible explanations for the nonlinear tissue distribution of PNL.

Several tissues including brain, fat and bone exhibited relatively little drug uptake consistent with our finding for DEX (Song et al., 2020). PNL is a substrate for P-gp, which is highly expressed at the blood-brain barrier and hampers brain exposure of steroids (Karszen et al., 2002; Yates et al., 2003). Despite the moderate lipid solubility of PNL, the *in vivo* $K_{p,Brain}$ values (0.05/0.03) were much lower than *in silico* values and for other tissues. This difference was accounted for in the PBPK model by incorporating P-gp-mediated active efflux in brain. Although this phenomenon may reflect lower binding in the brain, our preliminary modeling without the P-gp-mediated efflux process showed an approximate 20-fold overestimation of brain exposure even though the B_{max} value was estimated to be low. This was also seen for DEX in rats ($K_{p,Brain}$ =0.06) (Song et al., 2020), another P-gp substrate (Ueda et al., 1992; Schinkel et al., 1995).

The concentrations of PNL in fat, where drug uptake usually depends on lipid solubility, were low ($K_{p,Fat} = 0.17$), as also found for DEX ($K_{p,Fat} = 0.15$) (Song et al., 2020). These results are in line with findings that the SS fat K_p of PNL in rabbits was 0.13 (Khalafallah and Jusko, 1984b). This explains why volumes of distribution of PNL only increased modestly in obese men (Milsap et al., 1984) and rats (Nichols et al., 1989). Further, CBG is expressed in rat white adipose tissue (WAT) and the CBG layer surrounding WAT cells near the plasma membrane may act as a protective barrier limiting the access of GC to adipocytes (del Mar Grasa et al., 2001).

The relatively low concentrations of PNL in bone ($K_{p,Bone} = 0.22$) is consistent with observations for DEX ($K_{p,Bone} = 0.23$) (Song et al., 2020). Why bone has such low values is not clear. Nevertheless, knowledge of factors responsible for steroid access to bone may be important for understanding their deleterious effects (Lukert and Raisz, 1990). The limited distribution of PNL into fat and bone was well described by assuming only interstitial space is accessible.

Hepatic and renal metabolism and renal excretion are responsible for PNL clearance (Caspi and Pechet, 1958; Vermeulen and Caspi, 1958; Glenn, 1959; Bailey and West, 1967; Rocci et al., 1981). The model-estimated CBG-free hepatic intrinsic clearance was high (1789 mL/h), accounting for rapid conversion of PNL to PN. Using Eq. (38) and the hepatic blood flow (1424 mL/h), this converts to a systemic clearance of 793 ml/h. For kidney, the CLs is 149 ml/h. The overall systemic clearance is thus 942 ml/h, comparable to the value of 1129 mL/h for CBG-free PNL clearance in rats (Boudinot et al., 1986; Boudinot and Jusko, 1986). These previous studies showed similar plasma concentration profiles of PNL and PN after a 50 mg/kg IV dose as ours. The liver thus accounts for 90% of PNL disposition and the kidney 10%. However, these clearance values do not account for PNL and PN interconversion as recycling augments the PNL

AUC and thus underestimate the true clearance (Ebling and Jusko, 1986). There is evidence of an enterohepatic circulation (EHC) of GC which might partially explain their appreciable liver clearance. The intestinal absorption and biliary excretion of DEX in rats was reduced by dicyclomine, a cholinergic blocking agent, and consequently resulted in a decrease in AUC (Ogiso et al., 1985), indicating the involvement of EHC in DEX disposition. The possibility for EHC in rats might also apply to PNL (Mueller and Potter, 1981), although total radioactivity was measured in their study which would include PNL metabolites.

As shown in Figure 4, the appearance of PN in each tissue is very rapid upon PNL administration with similar times to reach maximum concentrations observed for both compounds. Modeling PN concentration simply as a fraction of PNL described the PN data reasonably well. This supports the assumption that the interconversion between PNL and PN is in continual rapid equilibrium. The interconversion of PNL and PN is mediated by 11β -HSDs, of which 11β -HSD1 is widely distributed with high expression in the liver and also in adipose tissue, muscle, bone, synovium, gonadal tissue, decidua, ocular tissues and within the CNS. The 11β -HSD2 is largely restricted to the mineralocorticoid-target tissues and the placenta (Raza et al., 2010). There has been evidence showing that the different tissue ratios of PN/PNL are mainly due to the tissue-specific expression and activity of 11β -HSDs (Escher et al., 1994). The conversion from PNL to PN is mediated by 11β -HSD2 and some tissues exhibit 11β -HSD2 saturation. The nonlinear PN/PNL ratio in rat plasma is consistent with observations in dogs (Frey et al., 1980) and humans (Legler et al., 1982). The 11β -HSD2 has tissue-specific distribution with highest enzyme activity in kidney (Monder, 1991; Yau et al., 1991), which may explain the appreciable and non-saturable production of PN in kidney. Our liver concentrations of PN were very low. This could be explained by the high 11β -HSD1 expression in the liver, and

also agrees with findings in rats (Escher et al., 1994) that the PN/PNL ratio in liver was far lower than in other tissues. Liver perfusions were carried out in rabbits where the extraction ratio of PNL is 50%, but no measurable PN was formed (Hale and Benet, 1991). Redistribution of PN between various tissues cannot be ruled out owing to the likely high permeability of this compound, which partly necessitated our metabolite modeling approach.

The K_p predictions using three *in silico* methods were similar and generally comparable to those obtained from *in vivo* data except for liver and brain (Table 4). The difference between liver K_p before (1.63) and after (3.67) metabolic correction can be attributed to substantial hepatic extraction of PNL and moderate partitioning. Similar findings of moderate liver partitioning were found for DEX ($K_p= 5.06$) (Song et al., 2020) and MPL ($K_p= 13.5$) (Ayyar et al., 2019). The extensive accumulation of GC within liver may be largely attributed to the abundant expression of GR (Ballard et al., 1974). Also, the GC importer-mediated active uptake of PNL, DEX and MPL into highly purified rat liver plasma membranes free of CBG and GR (Lackner et al., 1998) may be another factor contributing to their high K_p values. However, the PBPK model-derived K_p (1.31) and the *in silico* K_p (0.48-0.77) for liver were 2.8- to 10-fold smaller than obtained with the SS infusion. The plasma concentrations of PNL after the 50 mg/kg bolus SC dose (Figure 4) are substantially higher than from (1032.5 ng/mL) the infusion study. These nonlinear K_p differences agree with observations in rabbits that the tissue distribution of PNL is saturable in liver (Khalafallah and Jusko, 1984b). The *in silico* methods assume that tissue distribution is largely determined by neutral and phospholipids drug partitioning. They do not consider the roles of transport proteins, complex binding of drugs in liver, and active efflux in brain.

Limitations of the current study are that only one bolus dose level of PNL was used and the SC bioavailability (F) was based on the IV data from another study (Huang and Jusko, 1990). However, the 50 mg/kg dose of SC PNL yielded a wide range of plasma and tissue concentrations spanning several orders of magnitude. The calculated F appeared reliable in that the same rat strain (male Wistar rats) with comparable body weights were used and the IV AUC values are consistent across studies (Boudinot et al., 1986; Boudinot and Jusko, 1986; Huang and Jusko, 1990). The metabolic interconversion rates between PNL and PN were observed to be in rapid equilibrium. It would be necessary to directly give PN to assess its clearance parameters (Ebling and Jusko, 1986). Through incorporating nonlinear plasma and tissue binding, the current PBPK model was able to reasonably capture the PK profiles of PNL and PN in all tissues. Although the late low concentrations of PNL in muscle were over-predicted, these values are complicated by being close to the LLOQ of 1 ng/mL, account for less than 0.1% of early concentrations, and are the latest values in individual animals after SC dosing and subject to diverse sources of variation. The similarity of PNL and PN profiles across tissues supports the general behavior of all tissues as modeled.

The metabolic and pharmacokinetic properties of PNL and PN are generally similar in rat and man, making this PBPK model applicable to man and perhaps other species. In particular, as described for man in the Introduction, this includes nonlinear binding to CBG in plasma, reversible metabolism with PNL dominating over PN in a similar ratio in plasma, metabolism by liver and kidney, involvement of CYP3A in liver (Pichard et al., 1992; Penzak et al., 2005) and 11β -HSDs in various tissues, a similar apparent V_{ss} /Body Weight in rat (1.31 L/kg) (Huang and Jusko, 1990) and man (1.04 L/kg) (Czock et al., 2005) implying similar tissue binding. There is also similar binding to glucocorticoid receptors in both species. Differences would include

metabolic rates and possible EHC only in the rat. As corticosteroids generally have similar PK properties in various species, receptors in most tissues, and multiple therapeutic and adverse actions, this PBPK model may provide greater insights into their pharmacodynamic complexities.

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Authorship Contributions

Participated in research design: Li, DuBois, Jusko.

Conducted experiments: Li, DuBois.

Performed data analysis: Li, Jusko.

Wrote or contributed to the writing of the manuscript: Li, DuBois, Almon, Jusko.

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Footnote

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Figure Legends

Figure 1. Whole-body PBPK model scheme for prednisolone and prednisone disposition in rats.

Parameters and symbols are defined in the text and tables. Lines with arrows indicate blood flows and drug transport. Each box represents one tissue compartment as indicated by the label.

Figure 2. Blood partitioning of prednisolone. (A) blood-to-plasma concentration ratio (R_B) vs. plasma concentration of PNL from the SC bolus (closed squares) and SC infusion (open squares) studies; (B) Calculated RBC (closed triangles) and measured free plasma (open triangles) prednisolone concentrations in rats after a single 50 mg/kg SC dose of PNL. The dashed line indicates the mean value of all points.

Figure 3. Plasma protein binding of prednisolone and prednisone. (A) Bound fractions versus total concentrations of prednisolone and prednisone; (B) Relationship of bound versus unbound concentrations of prednisolone. Red open circles are PNL and black closed circles are PN data. The curve depicts model simulations using Eq. (9) with protein binding parameters values listed in Table 2.

Figure 4. The prednisolone and prednisone concentration-time profiles in plasma and tissues after 50 mg/kg SC bolus dose of PNL in rats. Red open circles are PNL and black closed circles are PN data. Curves depict PBPK model fittings of PNL (solid line) and PN (dashed line).

Figure 5. The prednisolone concentration-time profiles in blood after 50 mg/kg SC bolus dose of PNL in rats. Red open circles are PNL data. Curves depict PBPK model fittings.

Table 1. Physiological parameters of tissues in 294-g male rats.

Tissue	Volume (V , mL)	Blood Flow (Q , mL/h)
Lung	0.96 ^a	5548.6 ^b
Brain	1.41 ^a	123.1 ^b
Brain Capillary	0.02 ^a	
Heart	0.87 ^a	285 ^b
Intestine	8.14 ^b	1047.7 ^b
Spleen	0.43 ^a	336.8 ^b
Kidney	1.81 ^a	687.1 ^b
Muscle	127.52 ^b	1743.1 ^b
Liver	9.94 ^a	1424.4 ^c
Liver Artery	NA	39.8 ^b
Skin	52.17 ^b	376.2 ^b
Bone	21.61 ^b	115.7 ^b
Bone ISF	4.02 ^b	
Fat	34.44 ^b	422.8 ^b
Fat ISF	5.85 ^b	
Arterial Blood	5.76 ^b	5548.6 ^b
Venous Blood	11.52 ^b	5548.6 ^b
Remainder	17.08 ^c	371.2 ^c

^aTissue vascular-corrected volume (V_T) calculated based on experiment value using Eq. (4).

^b V_T calculated based on the value from (Shah and Betts, 2012).

‘Calculated value assuming 1.) 1 ng/mL tissue density, volume for remainder compartment = body weight – summation of volume for all measured tissues; 2.) Blood flow for remainder compartment = cardiac output - summation of blood flow for all measured tissues; 3.) Liver blood flow = summation of blood flow of liver artery, spleen and intestine).

NA: not applicable.

Table 2. PN/PNL ratios as determined by Orthogonal Regression in each tissue^a.

Tissue	Parameters (RSE%)		
	PN_{max} (ng/mL)	PNL_{50} (ng/mL)	f_m
Fat	166 (41.9)	283 (202)	
Bone	108 (28.4)	314 (173)	
Spleen	230 (48.2)	794 (252)	
Heart	338 (63.3)	2575 (251)	
Intestine	589 (62.8)	3174 (231)	
Plasma	249 (47.2)	1276 (224)	
Kidney			0.076 (39.5)
Lung			0.016 (7.7)
Brain			0.059 (26.7)
Skin			0.064 (17.2)
Muscle			0.060 (28.3)

^a f_m : linear PN/PNL ratio estimated according to Eq. (35); PN_{max} and PNL_{50} : the maximal PN concentration achievable and the PNL concentration at which half of the maximal PN concentration is achieved estimated based on Eq. (36).

Table 3. PBPK model parameter estimates after a single SC dose of 50-mg/kg PNL in rats.

Parameter (Unit)	Definition	Estimate (CV%)
$N_C P_C$ (ng/mL)	Binding capacity of CBG	297.8 (Fixed)
K_C (mL/ng)	Association constant of CBG	0.0046 (Fixed)
$N_A P_A$ (ng/mL)	Binding capacity of albumin	164342 (Fixed)
K_A (mL/ng)	Association constant of albumin	1.02×10^{-5} (Fixed)
B_{max_1} (ng/mL)	Binding capacity of muscle	690.5 (12.5)
B_{max_2} (ng/mL)	Binding capacity of liver, heart, intestine, bone	188.5 (11.6)
B_{max_3} (ng/mL)	Binding capacity of kidney, lung, skin, spleen, fat, brain	50.77 (10.8)
K_d (ng/mL)	Dissociation constant for PNL-tissue interaction	3.01 (23.8)
ka (hr^{-1})	SC first-order absorption rate constant	2.59 (5.1)
CL_{int} (mL/hr)	CBG-free hepatic intrinsic clearance	1789 (16.3)
CL_k (mL/hr)	CBG-free renal clearance	191.2 (40.3)
CL_{up} (mL/hr)	Brain bidirectional uptake clearance	0.13 (22.6)
CL_{eff} (mL/hr)	Brain unidirectional efflux clearance	4.58 (25.9)
f_{m_liver}	Metabolite/drug ratio of PN/PNL for liver	0.01 (30.1)
R_B^a	Blood-to-plasma concentration ratio	0.71 (Fixed)

^a R_B was estimated by Orthogonal Regression according to Eq. (1).

Table 4. Summary of prednisolone tissue partition coefficients by several methods.

Tissue	PBPK	Infusion ^a	Method 1	Method 2	Method 3
$K_{p,Lung}$	1.00	1.21±0.47	0.86	0.7	0.53
$K_{p,Brain}$	0.05	0.03±0.003	1.24	0.89	0.63
$K_{p,Kidney}$	1.02	0.22±0.05	0.79	0.66	0.47
$K_{p,Kidney}$ (uncorrected)	0.79	0.17±0.04			
$K_{p,Heart}$	1.17	0.92±0.19	0.74	0.64	0.47
$K_{p,Intestine}$	1.17	1.37±0.78	NA ^b	NA ^b	NA ^b
$K_{p,Spleen}$	1.01	0.47±0.16	0.67	0.64	0.46
$K_{p,Muscle}$	1.88	0.32±0.02	0.67	0.6	0.36
$K_{p,Liver}$	1.31	3.67±0.19	0.77	0.63	0.48
$K_{p,Liver}$ (uncorrected)	0.58	1.63±0.08			
$K_{p,Skin}$	1.01	0.57±0.30	0.81	0.63	0.46
$K_{p,Bone}$	0.22	0.29±0.13	NA ^c	NA ^c	NA ^c
$K_{p,Fat}$	0.17	0.14±0.05	0.51	0.4	0.42
$K_{p,Remainder}$	1.00 (Fixed)				

^aCalculated based on SS infusion study with n = 2 (SD provided).

^bTissues associated with GI tract were lumped together.

^cDifferent bone marrows were listed separately in software.

Figure 1.

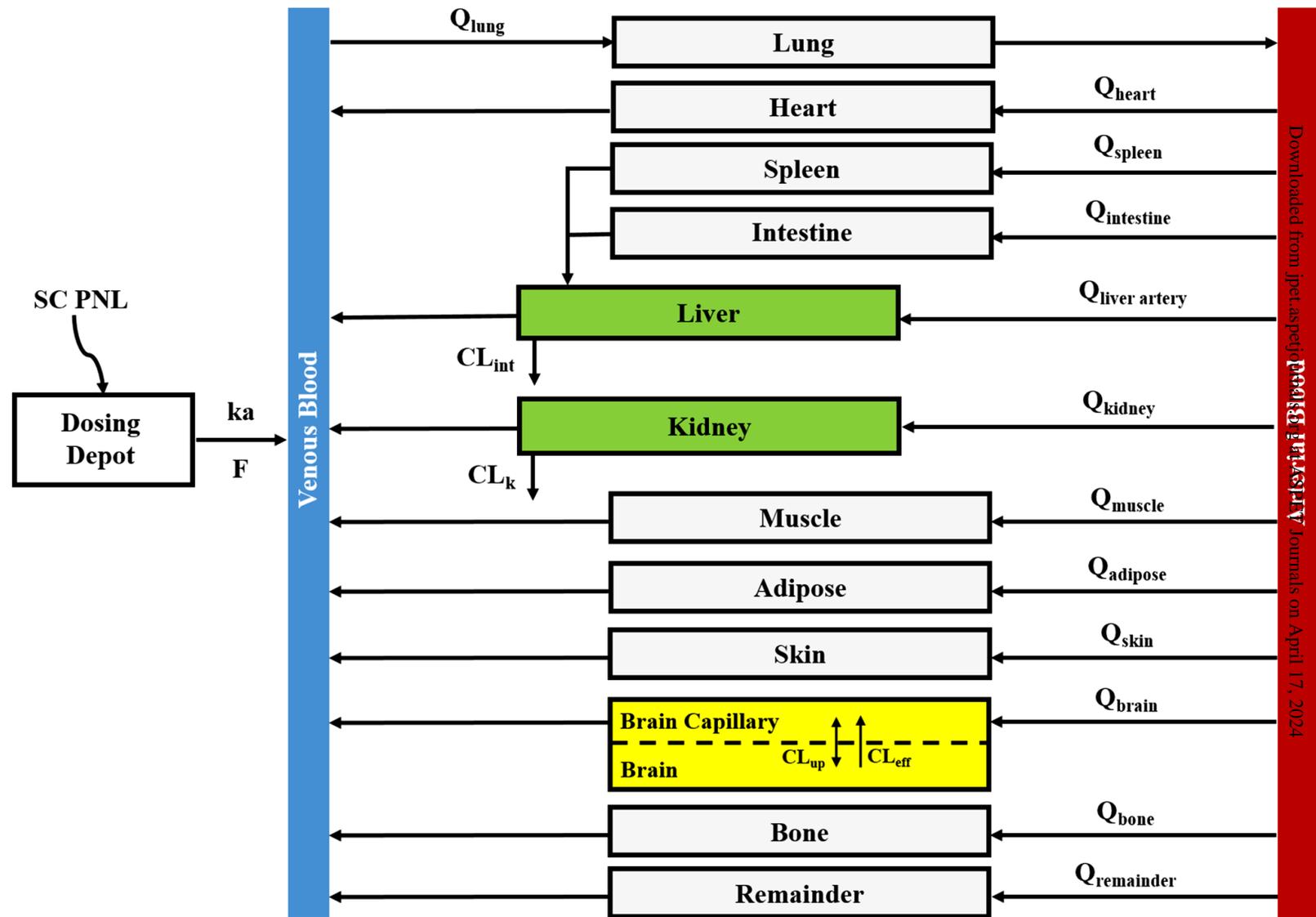
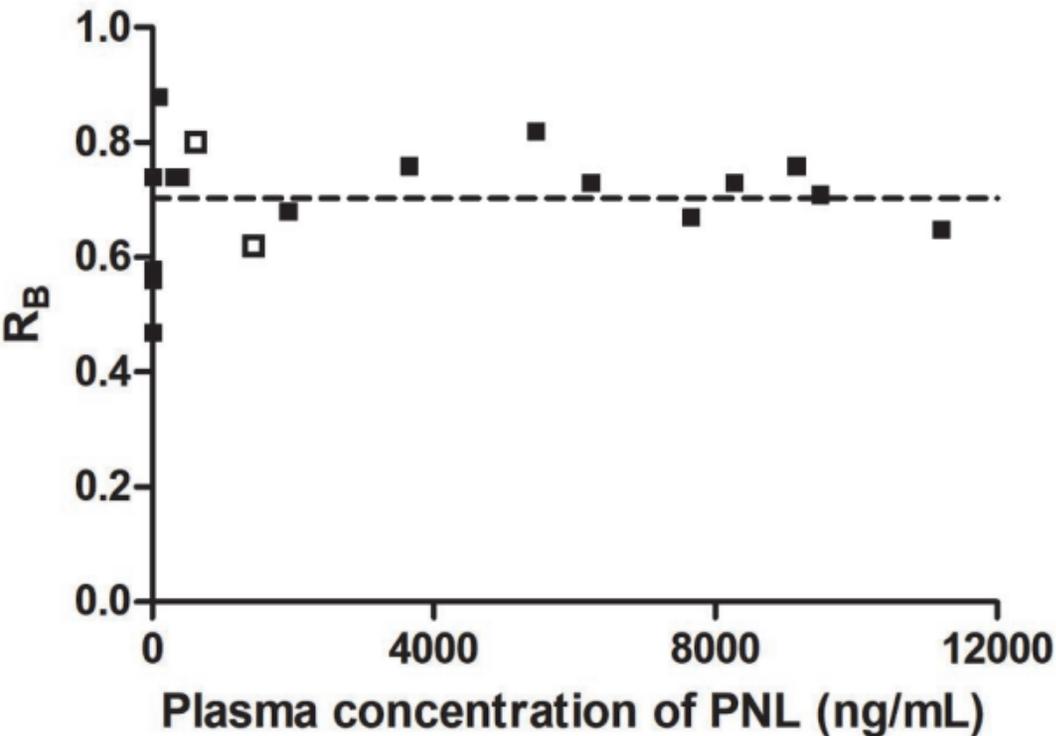


Figure 2.

A



B

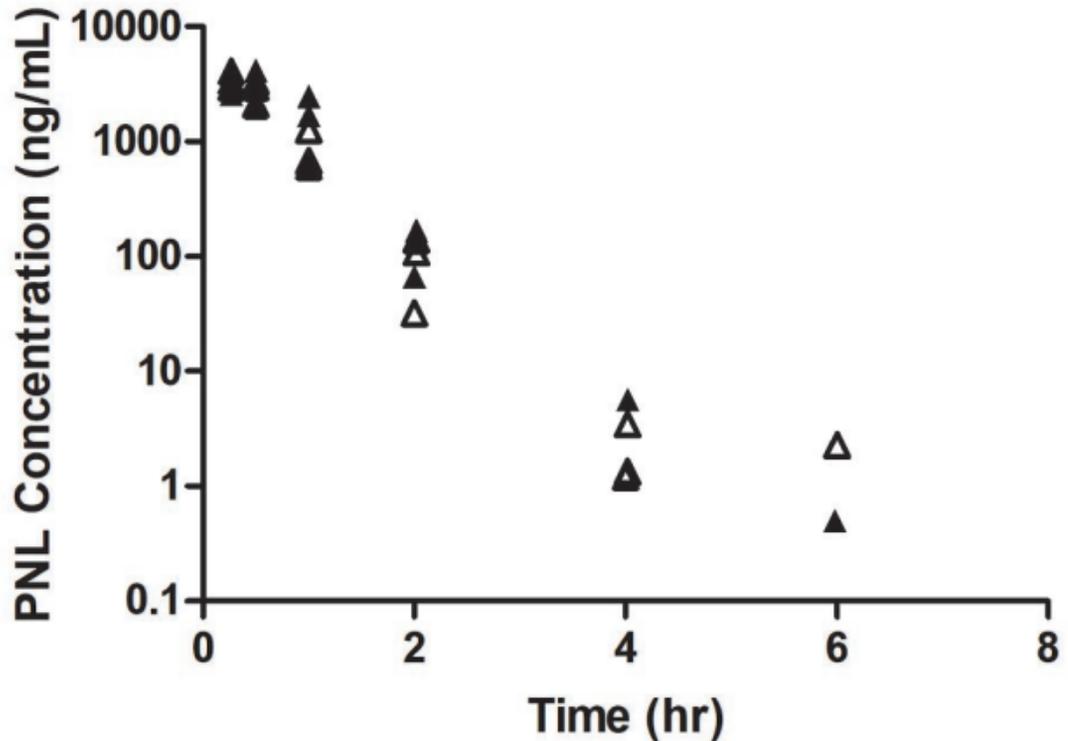


Figure 3.

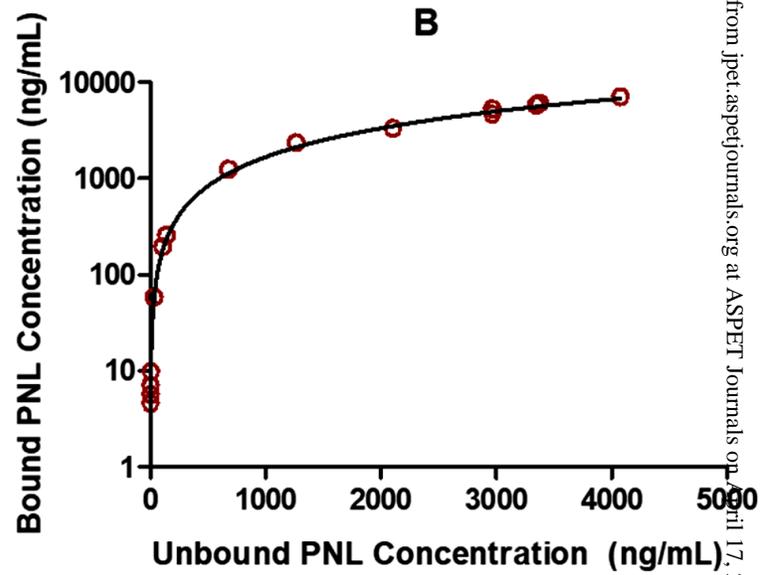
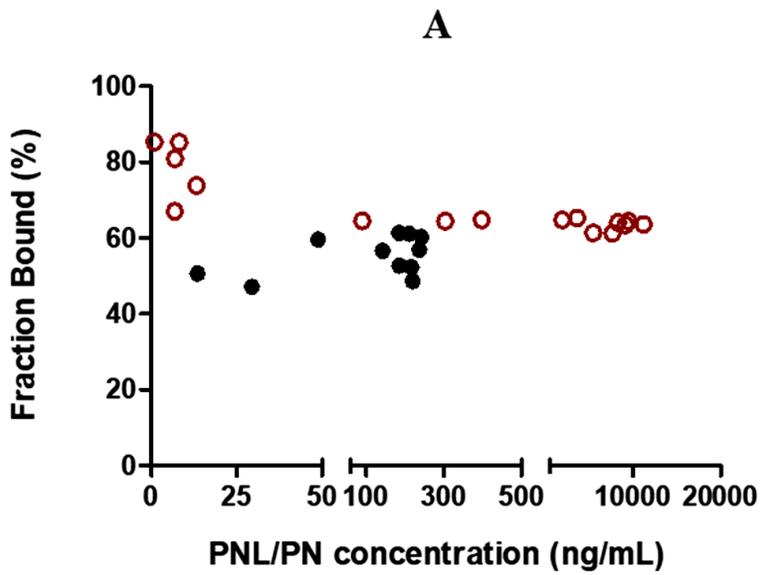
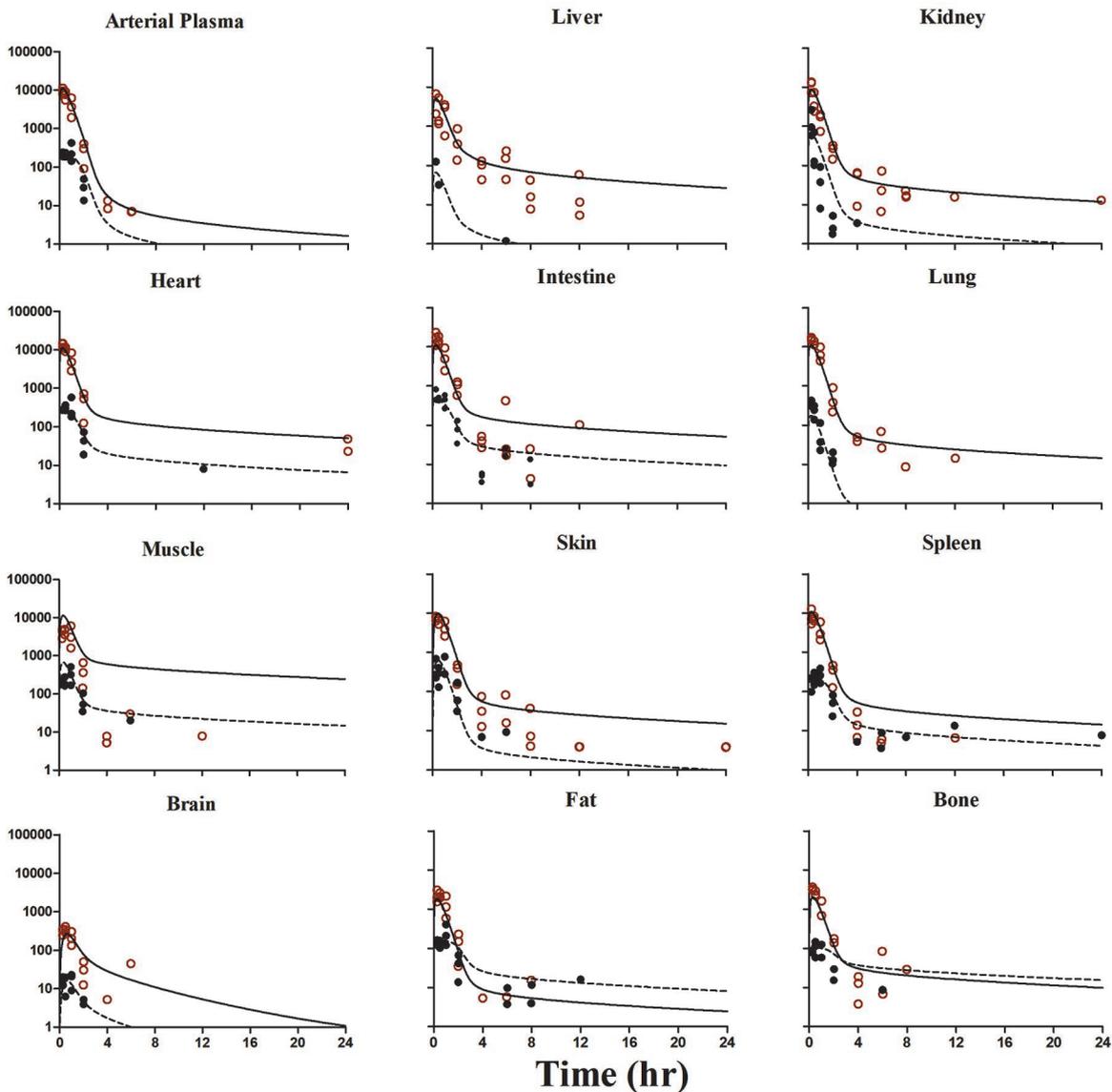


Figure 4.

PNL/PN Concentration (ng/mL)



Time (hr)

Figure 5.

Arterial Blood

