Consequences of Phenytoin Exposure on Hepatic Cytochrome P450 Expression during Postnatal Liver Maturation in Mice

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Abstract

The induction of cytochrome P450 (P450) enzymes in response to drug treatment is a significant contributing factor to drug-drug interactions, which may reduce therapeutic efficacy and/or cause toxicity. Since most studies on P450 induction are performed in adults, enzyme induction at neonatal, infant, and adolescent ages is not well understood. Previous work defined the postnatal ontogeny of drug-metabolizing P450s in human and mouse livers; however, there are limited data on the ontogeny of the induction potential of each enzyme in response to drug treatment. Induction of P450s at the neonatal age may also cause permanent alterations in P450 expression in adults. The goal of this study was to investigate the short- and long-term effects of phenytoin treatment on mRNA and protein expressions and enzyme activities of CYP2B10, 2C29, 3A11, and 3A16 at different ages during postnatal liver maturation in mice.

Introduction

Drug-drug interactions (DDIs) are a major cause of adverse drug reactions (ADRs) and occur when one medication interferes with the efficacy or toxicity profile of another medication administered concurrently. Pharmacokinetic DDIs occur when the perpetrating drug affects the absorption, distribution, metabolism, or excretion of other victim drugs, shifting plasma concentrations outside of optimal therapeutic windows (Palleria et al., 2013). Plasma concentrations are largely determined by cytochrome P450 (P450)–mediated drug metabolism, which can be inhibited or induced by xenobiotics. Several marketed drugs induce the transcription of P450 genes via the activation of nuclear receptors and are implicated in clinically significant DDIs (Lin, 2006).

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Abbreviations: ADR, adverse drug reaction; AED, antiepileptic drug; BSA, bovine serum albumin; CAR, constitutive androstane receptor; DDI, drug-drug interaction; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; IDA, information-dependent acquisition; MPB, mobile phase B; MS, mass spectrometry; P450, cytochrome P450; PBS, phosphate-buffered saline; RT-PCR, real-time polymerase chain reaction; SWATH-MS, sequential window acquisition of all theoretical mass spectra; TOF, time-of-flight.
2013). Other enzymes, such as CYP3A7, are only expressed in the fetal and early postnatal liver and become undetectable once the liver is fully matured (Lacroix et al., 1997).

Whereas the postnatal ontogeny of P450s in humans has been established, the induction of P450s in response to drug treatment has not been well studied in pediatric populations at different stages of early development. It is largely unknown to what extent P450 activity can be induced in neonates and children compared with adults. Whether the capacity for enzymatic induction varies at different ages during postnatal liver maturation is also not understood. Several studies have additionally suggested neonatal exposures to inducer drugs, particularly the antiepileptic drug (AED) phenobarbital, can cause permanent alterations to expression and activities of P450s in adult rodents (Agrawal and Shapiro, 1996, 2005; Tien et al., 2015, 2017; Piekos et al., 2017). There also appears to be a window of sensitivity during neonatal life that allows the imprinting of lasting changes to P450 expression following treatment (Tien et al., 2015). This effect has not yet been validated with other small molecule drugs.

The goal of this work was to further examine the postnatal liver’s capacity for P450 induction in response to inducer treatment at different ages spanning neonatal, infant, and adolescent stages in mice. This is a continuation of our previous studies that investigated the effects of neonatal phenobarbital treatment on expression of P450s in adult mice. Whereas no animal model is a perfect representation of human drug metabolism, the mouse is a useful in vivo system for studying developmental responses to drug treatment in a relatively human drug metabolism, the mouse is a useful in vivo system for studying developmental responses to drug treatment in a relatively human drug metabolism, the mouse is a useful in vivo system for studying developmental responses to drug treatment in a relatively.

**Materials and Methods**

**Chemicals and Reagents.** Phosphate-buffered saline (PBS), 5,5-diphenylhydantoin salt (phenytoin), trifluoroacetic acid, formic acid, and acetonitrile were purchased from Sigma-Aldrich (St. Louis, MO). Urea and dithiothreitol were obtained from Fisher Scientific Co. (Pittsburgh, PA). Iodoacetamide and ammonium bicarbonate were purchased from Acros Organics (Morris Plains, NJ). Tosyl phenylalanyl chloromethyl ketone (Tosyl phenylalanyl chloromethyl ketone) and ammonium bicarbonate were purchased from Acros Organics (Morris Plains, NJ). Tosyl phenylalanyl chloromethyl ketone–treated trypsin was obtained from Worthington Biochemical Corporation (Freehold, NJ). Standard bovine serum albumin (BSA) was purchased from Thermo Fisher Scientific (Waltham, MA).

**Animals.** C57Bl/6 mice (Jackson Laboratories, Bar Harbor, ME) were housed in accordance with the animal care guidelines outlined by the American Association for Animal Laboratory Sciences and were bred under standard conditions in the Animal Resources Facility at the University of Connecticut. The use of these animals was approved by the University of Connecticut’s Institutional Animal Care and Use Committee. At 12 weeks of age, mice were set up into breeding pairs. To study short-term effects of drug treatment at different ages during postnatal development, mice were administered a single dose of phenytoin (100 mg/kg) or PBS vehicle control intraperitoneally on days 5, 10, 15, 20, 30, or 60 following birth. Twenty-four hours after the treatment, mice were sacrificed and liver samples were collected. To study long-term effects of drug treatment, mice on day 5 after birth were administered either phenytoin (100 mg/kg) or PBS vehicle control intraperitoneally. Following treatment, mice were sacrificed at different ages (day 10, 15, 20, 30, or 60 after birth) and livers were collected.

Livers with gallbladders removed were snap frozen in liquid nitrogen and then stored at −80°C. In both the short- and long-term studies, 4–6 livers were collected at each age for both control and treatment groups. Both male and female mice were included in this study.

**Total RNA Extraction, Reverse Transcription, and Real-Time Polymerase Chain Reaction.** Total RNAs were isolated from whole livers using TRIzol reagent according to the manufacturer’s protocol (Life Technologies, Guilford, CT). RNA concentrations were measured using a NanoDrop spectrophotometer (NanoDrop Technologies, Wilmington, DE) at a wavelength of 260 nm. To obtain cDNA, reverse transcription was performed using an iScript Reverse Transcription Supermix according to the manufacturer’s protocol (Bio-Rad, Hercules, CA). Real-time polymerase chain reaction (RT-PCR) was performed using a CFX-96 thermocycler system (Bio-Rad) with TaqMan gene expression assays for glyceraldehyde-3-phosphate dehydrogenase (GAPDH), CYP3A16, 3A11, 2B10, and 2C29 (Thermo Fisher Scientific). Fold changes of P450 expression in the phenytoin-treated group compared with the PBS control group at each postnatal age were calculated using the 2^(-ΔΔCT) method described by Livak and Schmittgen (2001) with a normalization of P450 expression to GAPDH. A consistent expression level of GAPDH is observed by either RNA-Seq or RT-PCR in mouse liver during postnatal maturation (Supplemental Fig. 1).

**Protein Sample Preparation and Liquid Chromatography–Tandem Mass Spectrometry–Based Quantification.** Whole mouse livers were chopped into small pieces using scissors and homogenized in PBS (pH 7.4) by a tissue grinder equipped with a Teflon pestle on ice. Liver S9 fractions were obtained by centrifuging the homogenate at 4°C at 10,000g for 30 minutes. The supernatant containing the S9 fractions was transferred to a Beckman ultracentrifuge tube and centrifuged at 300,000g for 20 minutes. The liver microsomes were obtained by resuspending the pellet with PBS using the tissue grinder. Protein concentration was determined using a Pierce BCA Protein Assay Kit (Thermo Fisher Scientific). Protein digestion was conducted according to a previously reported Lys-C/Trypsin combinatorial digestion protocol (Glatter et al., 2012) with minor modifications. An aliquot of 80 μg liver microsomal protein was mixed with the internal standard BSA (0.2 μg). After adding a 10-fold volume of precooled acetone, the mixture was incubated at −20°C for 2 hours and centrifuged. The supernatants were removed and the precipitated proteins were washed with 80% ethanol and then air-dried at room temperature. Proteins were resuspended in 4 mM DDT in 8 M urea solution containing 100 mM ammonium bicarbonate. Samples were sonicated and then incubated at 37°C for 45 minutes. After samples were cooled down to room temperature, 20 mM iodoacetamide in 8 M urea/100 mM ammonium bicarbonate was added and incubated for 30 minutes at room temperature for alkylation. Following incubation, the urea concentration was adjusted to 6 M by adding 61.6 μM of 50 mM amonium bicarbonate. Lysyl endopeptidase (Wako Chemicals, Richmond, VA) was added at an enzyme-to-protein ratio of 1:100 for the first step protein digestion for 6 hours at 37°C. Samples were diluted to 1.6 M urea followed by a second digestion with trypsin at an enzyme-to-protein ratio of 1:50 for an overnight incubation at 37°C. Digestion was terminated by adding 1 μl trifluoroacetic acid.

Digested peptides were extracted and purified using Waters Oasis HLB columns (Waters Corporation, Milford, MA) according to the manufacturer’s instructions. Eluted peptides were dried in a Speed Vac SPD1010 (Thermo Scientific, Hudson, NH) and reconstituted in 3% acetonitrile with 0.1% formic acid. The peptide samples were centrifuged and one-half of the supernatant was collected and supplemented with synthetic iRT standard solutions (Biognosys AG, Cambridge, MA) prior to liquid chromatography–tandem mass spectrometry (MS) analysis.

Peptides were analyzed in a TripleTOF 5600+ mass spectrometer (AB Sciex, Framingham, MA) coupled with an Eksigent 2D Plus LC System (Eksigent Technologies, Dublin, CA). Liquid chromatography separation was performed via a trap-elute configuration, which included a trapping column (ChromXP C18-CL, 120 Å, 5 μm, 0.3 mm cartridge; Eksigent Technologies) and an analytical column (ChromXP C18-CL, 120 Å, 150 × 0.3 mm, 5 μm; Eksigent Technologies). The mobile phase consisted of water with 0.1% formic acid (phase A) and acetonitrile with 0.1% formic acid (phase B) (Avantor, Center Valley, PA). A total of 6 μg protein was injected for analysis. Peptides were trapped and cleaned on the trapping column, with phase A delivered at a flow rate of 10 μl/min for 3 minutes, followed by separation on the analytical column with a gradient elution at a flow rate of 5 μl/min. For column equilibration, the gradient time program was set as follows for phase B: 0–68 minutes at 3%–30%; 68–73 minutes at 100% phase B.
Fig. 1. Effects of phenytoin (PHY) treatment on the expression and activity of CYP2B10 at different ages during postnatal maturation. Male and female mice were treated with either vehicle control (PBS) or 100 mg/kg PHY at different ages following birth. The short-term effects of PHY treatment were measured 24 hours after treatment. Changes in gene expression were evaluated at the mRNA level using RT-PCR, and then fold changes were calculated using the $2^{-\Delta\Delta Ct}$ method. Data points represent fold changes in gene expression of individual mice treated with PBS or PHY compared with the average $\Delta Ct$ of mice that received PBS control (A). Ct values from RT-PCR experiments were also compared with the GAPDH Ct value in each individual mouse and the differences are expressed as $2^{(\text{Ct(P450)} - \text{Ct(GAPDH)})}$ (B). The dotted line
30%–40%; 73–75 minutes at 40%–80%; 75–78 minutes at 80%; 78–79 minutes at 80%–3%; and finally 79–90 minutes at 3%. The TripleTOF instrument was operated in positive ion mode with ion spray voltage flowing at 5500 V, ion source gas one at 28 psi, ion source gas two at 16 psi, curtain gas at 25 psi, and source temperature at 280°C.

To generate the reference spectral library for SWATH-MS data analysis, information-dependent acquisition (IDA) was performed for three pooled mouse liver microsome samples with equal protein amounts. The IDA method included a 250 millisecond time-of-flight (TOF)-MS scan from 400 to 1250 Da, followed by a tandem MS scan in high-sensitivity mode from 100 to 1500 Da (50 millisecond accumulation time, 10 ppm mass tolerance, charge state from +2 to +5, rolling collision energy and dynamic accumulation) of the top 30 precursor ions from the TOF-MS scan. The IDA data from the pooled mice liver microsomes were searched by MaxQuant (version 1.5.3.30; Max Planck Institute of Biochemistry, Planegg, Germany). The mouse FASTA file with 16,911 protein entries was downloaded from Uniport on August 29, 2016, and used as the reference sequences for searches with trypsin/P as the protease. Peptide length was set between 7 and 25 residues with up to two missed cleavage sites allowed. Carbamidomethyl was set as a fixed modification. A false discovery rate of 0.01 was used as the cutoff for both peptide and protein identification.

All microsomes samples were analyzed using a SWATH-MS method with the “SWATH (VW 100)” isolation scheme, which is comprised of a 250 millisecond TOF-MS scan from 400 to 1250 Da, followed by MS/MS scans from 100 to 1500 Da performed on all precursors in a cyclic manner. The accumulation time was 50 milliseconds per isolation window, resulting in a total cycle time of 2.8 seconds. The spectral alignment and target data extraction of the SWATH-MS data were performed on Skyline-daily (version 3.7.1.11271; University of Washington, Seattle, WA) with the reference spectral library generated from the aforementioned IDA searches. The M1 and MS/MS filtering were both set as TOF mass analyzer with the resolution power of 30,000 and 15,000, respectively. The retention time prediction was based on the auto-calculate regression implemented in the iRT calculator. Proper peak selection was reviewed manually with the automated assistance of Skyline-daily. The surrogate peptides used for the quantification of CYP2B10, 2C29, 3A11, and 3A16 proteins are listed in Supplemental Table 1. The selections were based on the uniqueness and chromographic performance of these peptides. The peak areas of the top 3–5 fragment ions were summed up and normalized to the internal standard BSA. The BSA-normalized peak area of each peptide was further divided by the average peak area of the 30 samples to calculate the relative abundance of the peptide. The average of relative abundance of all surrogate peptides of a protein was used to determine protein relative abundance.

**Evaluation of恩维克克 Activities.** Liver tissues were homogenized in ice-cold 100 mM PBS (pH 7.4) and S9 fractions were obtained by centrifugation. Protein concentrations were determined using a Pierce BCA Protein Assay Kit (Thermo Fisher Scientific, Rockford, IL). Pentoxysorufin O-dealkylation, paclitaxel 6-hydroxylation, and midazolam 1’-hydroxylation, were used as probes for CYP2B10 expression, in -duct and function of drug-metabolizing P450s throughout postnatal maturation, a mouse model was used. Short-term effects of drug treatment were evaluated at 24 hours following administration of a single dose of phenytoin in mice at different ages. Long-term effects of drug treatment were evaluated by administering a single dose of phenytoin at the age of 5 days after birth and then collecting livers at different time points throughout development, beginning on day 10 after birth. When comparing mouse to human ages (and considering total lifespans), 9 days in the mouse has been compared with 1 year in the human. Additionally, mice are weaned between days 21 and 28 of age, while humans are weaned between 6 and 12 months of age (Dutta and Sengupta, 2016). Based on this, the age on day 5 in mouse would correlate with roughly the first 3–6 months of life in humans when hepatic P450 expression is known to be rapidly changing. The expression of hepatic murine CYP2B10, 2C29, 3A11, and 3A16 enzymes was evaluated as representative members of the CYP2B, 2C, and 3A subfamilies that are known to be involved in the metabolism of xenobiotics and drugs across species (Martignoni et al., 2006). The effect of drug treatment on each enzyme was measured at the mRNA level using RT-PCR, at the protein level using SWATH-MS quantification, and at the enzymatic activity level using in vitro P450 probe substrates and liquid chromatography MS quantification. SWATH-MS is a relatively novel method for measuring protein concentrations, but previous studies have suggested lower limits of peptide quantitation in the 0.1–10 fmol range (Collins et al., 2017). Both male and female mice were evaluated collectively for data presented at ages younger than day 30; however, male and female data were evaluated separately on day 60 after birth. At each time point, mice that received drug treatment were compared against age-matched mice that received a vehicle control.

**Effects of Phenytoin Exposure on CYP2B10 Expression, Induction, and Function throughout Postnatal Maturation.** The fold increase of CYP2B10 mRNA induction 24 hours following phenytoin administration was the greatest of all P450s measured; however, the large value for fold induction may be overstated due to low basal expression of CYP2B10 in the liver compared with other enzymes.

To evaluate the impact of phenytoin exposure on the expression and function of drug-metabolizing P450s throughout postnatal maturation, a mouse model was used. Short-term effects of drug treatment were evaluated at 24 hours following administration of a single dose of phenytoin in mice at different ages. Long-term effects of drug treatment were evaluated by administering a single dose of phenytoin at the age of 5 days after birth and then collecting livers at different time points throughout development, beginning on day 10 after birth. When comparing mouse to human ages (and considering total lifespans), 9 days in the mouse has been compared with 1 year in the human. Additionally, mice are weaned between days 21 and 28 of age, while humans are weaned between 6 and 12 months of age (Dutta and Sengupta, 2016). Based on this, the age on day 5 in mouse would correlate with roughly the first 3–6 months of life in humans when hepatic P450 expression is known to be rapidly changing. The expression of hepatic murine CYP2B10, 2C29, 3A11, and 3A16 enzymes was evaluated as representative members of the CYP2B, 2C, and 3A subfamilies that are known to be involved in the metabolism of xenobiotics and drugs across species (Martignoni et al., 2006). The effect of drug treatment on each enzyme was measured at the mRNA level using RT-PCR, at the protein level using SWATH-MS quantification, and at the enzymatic activity level using in vitro P450 probe substrates and liquid chromatography MS quantification. SWATH-MS is a relatively novel method for measuring protein concentrations, but previous studies have suggested lower limits of peptide quantitation in the 0.1–10 fmol range (Collins et al., 2017). Both male and female mice were evaluated collectively for data presented at ages younger than day 30; however, male and female data were evaluated separately on day 60 after birth. At each time point, mice that received drug treatment were compared against age-matched mice that received a vehicle control.

**Effects of Phenytoin Exposure on CYP2B10 Expression, Induction, and Function throughout Postnatal Maturation.** The fold increase of CYP2B10 mRNA induction 24 hours following phenytoin administration was the greatest of all P450s measured; however, the large value for fold induction may be overstated due to low basal expression of CYP2B10 in the liver compared with other enzymes.

represents GAPDH expression set to 1 (n = 4–6). Protein was quantified using a liquid chromatography–tandem MS method in males only on days 5 and 60 of age of PHY treatment (C) (n = 5). Enzyme activity was confirmed by measuring the amount of pentoxysorufin metabolite produced by S9 fractions isolated from day 5 and 60 male liver samples (n = 5) treated with resorufin (D). The long-term effects of PHY treatment were investigated in a similar fashion. Mice were treated with 100 mg/kg PHY on day 5 after birth, and then livers were collected (n = 3–6) at different later time points for measurements of mRNA (E), and for protein and enzymatic activity at 60 days of age (F and G). The data are presented as mean ± S.E. *P < 0.05; **P < 0.01; ***P < 0.001.
Fig. 2. Effects of phenytoin (PHY) treatment on the expression of CYP2C29 at different ages during postnatal maturation. Male and female mice were treated with either vehicle control (PBS) or 100 mg/kg PHY at different ages following birth. The short-term effects of PHY treatment were measured 24 hours after treatment. Changes in gene expression were evaluated at the mRNA level using RT-PCR, and then fold changes were calculated using the $2^{-\Delta\Delta Ct}$ method. Data points represent fold changes in gene expression of individual mice treated with PBS or PHY compared with the average $\Delta Ct$ of mice that received PBS control (A). Ct values from RT-PCR experiments were also compared with the GAPDH Ct value in each individual mouse and the differences are expressed as $2^{-\Delta Ct(P450)-\Delta Ct(GAPDH)}$ (B). The dotted line represents GAPDH induction of P450s by Phenytoin in Developmental Liver.
Induction of CYP2C29 mRNA expression until the mice reached 60 days of age. The induction throughout postnatal maturation was evident at all ages studied during postnatal development. The greatest fold change between phenytoin-treated mice and control mice occurred in neonates at 5 days of age, presumably because this is when the basal expression of CYP3A11 is the lowest (Fig. 3A). As mice matured following birth, basal expression of CYP3A11 mRNA increased along with increased induced levels of mRNA with age (Fig. 3B). Maximal induction occurred at the adult ages (days 30 and 60), which was also reflected in the expression of CYP3A11 protein (Fig. 3C). Protein quantification revealed that induced levels of CYP3A11 in the neonate at 5 days of age were not statistically different from basal levels of CYP3A11 protein in the adult. This was also confirmed by enzymatic activity analyses with midazolam in liver S9 fractions (Fig. 3D). At 5 days of age, mice treated with phenytoin had similar rates of 1-hydroxymidazolam metabolite formation as adult mice treated with vehicle control, while adult mice treated with phenytoin produced the most metabolite overall.

There also appeared to be no significant long-term effects on CYP3A11 expression or function following neonatal phenytoin exposure. Following administration at day 5, phenytoin seemed to increase the range of CYP3A11 mRNA expression at later ages between mice, such as at 10 days of age where the S.E. in the fold changes of treated mice was larger than in control mice. However, there were no significant differences in the mean expression from control mice, except for a slight increase at 30 days of age (Fig. 3E). This was also confirmed in protein (Fig. 3F) and enzymatic activity (Fig. 3G) quantification. At the adult ages, there were no significant differences in mice that received phenytoin at 5 days of age of CYP3A11 protein expression or midazolam metabolism.

Effect of Phenytoin Exposure on the Postnatal Repression of the Fetal Enzyme CYP3A16. CYP3A16 is considered to be the fetal enzyme of the CYP3A subfamily of enzymes. While it is highly expressed immediately after birth, its expression declines steadily throughout postnatal maturation and is undetectable in the adult liver. Phenytoin treatment also appeared to significantly affect short-term CYP3A16 expression at different ages during postnatal development. Twenty-four hours after phenytoin administration, CYP3A16 mRNA was significantly suppressed in mice at all ages examined (Fig. 4A). The fold change in the decrease of expression was greatest at 20 days of age expression set to 1 (n = 4–6). Protein was quantified 24 hours after PHY treatment using a liquid chromatography–tandem MS method in males only on days 5 and 60 after birth (C) (n = 5). The long-term effects of PHY treatment were investigated in a similar fashion. Mice were treated with 100 mg/kg PHY 5 days after birth, and then livers were collected (n = 3–6) at different later time points for measurements of mRNA (D) and at 60 days of age for measurement of protein concentration (E). The data are presented as mean ± S.E. *P < 0.05; **P < 0.01; ***P < 0.001.
Fig. 3. Effects of phenytoin (PHY) treatment on the expression and activity of CYP3A11 at different ages during postnatal maturation. Male and female mice were treated with either vehicle control (PBS) or 100 mg/kg PHY at different ages following birth. The short-term effects of PHY treatment were measured 24 hours after treatment. Changes in gene expression were evaluated at the mRNA level using RT-PCR, and then fold changes were calculated using the \(2^{-\Delta\Delta Ct}\) method. Data points represent fold changes in gene expression of individual mice treated with PBS or PHY compared with the average \(\Delta Ct\) of mice that received PBS control (A). Ct values from RT-PCR experiments were also compared with the GAPDH Ct value in each individual mouse and the differences are expressed as \(2^{[Ct(P450)-Ct(GAPDH)]}\) (B). The dotted line
and this corresponded to the age in which CYP3A16 mRNA became undetectable in phenytoin-treated mice (Fig. 4B). CYP3A16 mRNA was not detected by RT-PCR in mice at 30 or 60 days of age; therefore, the data are not shown. The decrease in CYP3A16 in response to phenytoin treatment was also confirmed by protein quantification (Fig. 4C). At 5 days of age, CYP3A16 protein was decreased following drug exposure; however, not significantly. CYP3A16 protein was not detected in adults regardless of phenytoin administration.

**Discussion**

The present study reveals both the immediate and persistent effects on the expression and functions of several drug-metabolizing P450 enzymes following exposure to the AED drug phenytoin during murine postnatal development. Short-term induction of CYP2B10, 2C29, and 3A11 after phenytoin administration for 24 hours seems to vary in extent based on age and sex. Younger pups, such as those at 5 and day 10 days of age, had higher fold changes relative to control for P450 induction; however, these values may be overstated due to the low basal expression of P450s in the neonatal period. With increasing age, the values for fold induction decreased compared with control animals. Based on this, it is unclear how induced levels of each enzyme change independent of their basal expression levels. For this reason, we also chose to evaluate mRNA expression of each P450 relative to GAPDH. GAPDH expression was not altered by drug treatment and remained relatively consistent throughout postnatal development. Analyzing gene expression in relation to GAPDH allowed the ontogeny of each P450 enzyme to be observed in basal and induced states at each age. Induction of each enzyme (CYP2B10, 2C29, and 3A11) appeared to be qualitatively proportional to the basal expression level of the enzyme; the higher the mRNA expression of a P450 at a certain age, the greater its induction capacity. On the other hand, CYP3A16 appeared to be repressed to greater extents at increasing ages, which correlated with decreasing basal expression.

For both CYP2B10 and 2C29, day 10 appeared to be the age most sensitive to the inductive effects of phenytoin as evidenced by the greatest fold changes in mRNA expression compared with control animals. Interestingly, this correlates with our previous data that show a surge of CYP2B10 and 2C29 expression beginning at the same age (Peng et al., 2012). The expression patterns of the murine enzymes investigated in this study, namely, CYP2B10, 2C29, 3A11, and 3A16, follow specific ontogenic patterns throughout postnatal maturation, and our data show that induction varies depending on developmental age as well. A better understanding of the mechanisms regulating enzyme induction between the developing and fully mature mouse liver is needed to understand this observation. The expression of P450s in the human fetal, neonatal, and infant liver also follow distinct ontogenic patterns that may be specific to each enzyme. However, there are still marked differences between human and mouse P450-mediated metabolism, especially in terms of enzyme induction and regulation. Other studies have also highlighted the need for pediatric pharmacokinetics to be evaluated in age- and maturation-dependent models, rather than grouping all neonatal and infant guidelines into one pediatric category (Suzuki et al., 1994; Abduljalil et al., 2014). The long-term effects of neonatal phenytoin exposure were also investigated in our study. Following day 5 administration, CYP2B10 was the only P450 that remained overexpressed at day 10. This mechanism will have to be further elucidated since the half-life of phenytoin in adult mice is only 16 hours (Markowitz et al., 2010); however, this might not be true for neonates. At day 60, there appears to be slight, but significant, overexpression of CYP3A11 and CYP2B10 RNA in male mice only. However, the induction was not observed in the protein or enzymatic activity analyses. Interestingly, CYP2C9 protein expression is significantly downregulated at day 60 following neonatal phenytoin exposure, despite no significant changes observed at the mRNA level.

Additional studies will be necessary to further distinguish why neonatal exposure to phenobarbital produces overexpression of P450 mRNA, protein, and function in adulthood, while exposure to phenytoin does not; phenobarbital upregulates the transcription of P450s through its indirect action on the nuclear CAR (Wei et al., 2000). Previous work has shown that direct activation using 1,4-bis[2-(3,5-dichloropyridyloxy)] benzene, a murine-specific CAR ligand, at the neonatal age also produces permanent alterations to P450 expression (Li et al., 2016b). Greater fold changes in RNA expression following xenobiotic treatment are typically observed at the neonatal age. However, our study shows that despite the apparent greater extent of RNA induction at younger ages, protein expression and enzymatic activity following phenytoin treatment is still greater in adults than neonates. The expression of CYP2B10, 2C29, and 3A11 proteins is several magnitudes higher in adults than in neonates following phenytoin treatment, suggesting the adult liver has a greater capacity for enzyme induction following drug treatment than the infant liver. In the case of CYP2B10, protein expression in the neonate following phenytoin exposure exceeds that of adults in basal conditions with vehicle treatment. However, the expression of CYP2C29 protein following phenytoin treatment in the neonate does not reach adult expression levels. These data suggest that induction in response to drug treatment is not equivalent in neonatal and adult mice, even following the same dose, and adult data cannot be extrapolated to younger ages. Only liver samples from day 5 were used for protein and enzymatic activity analyses of early life, thus additional ages will need to be examined in the future.
P450 expression; however, this was not the case. Unlike 1,4-bis[2-(3,5-dichloropyridin-2-yl)]x, both phenobarbital and phenytoin produce CAR activation indirectly. While they both are AEDs, these chemicals have different mechanisms of action for treating seizures. It is plausible that phenobarbital and phenytoin produce indirect CAR activation via separate mechanisms or pathways, and a pathway further upstream of CAR activation is what becomes permanently affected following neonatal phenobarbital treatment that is not impacted by phenytoin treatment.

Phenytoin and phenobarbital are both considered first-line therapy for treating seizures in neonates with phenobarbital appearing to be slightly more efficacious (Painter et al., 1999; Booth and Evans, 2004; Pathak et al., 2013). Both drugs have indications for acute treatment of infant seizures as well as chronic administration for control of epileptic disorders in infants and children. Phenytoin therapy, as with most other AEDs, is initiated with a larger loading dose (18–20 mg/kg for pediatric use), followed by subsequent lower maintenance doses for recurrent seizure control. Appropriate doses are calculated based on the patient’s body weight; however, it is unclear whether other factors are taken into account (Hawcutt et al., 2011). In the current study, we administered mice a single dose of 100 mg/kg phenytoin, which is equivalent to a human dose of 8 mg/kg according to the Food and Drug Administration’s Guidance for Industry to extrapolate human dose concentrations from animal data. This is likely more representative of a maintenance dose of phenytoin; however, this was the maximal dose we could achieve while avoiding excessive toxicity. Perhaps a higher dosing concentration, or multiple-day phenytoin treatment, would have produced the permanent overexpression of P450s that was observed with phenobarbital treatment.

Phenytoin has a narrow therapeutic index with a large propensity to produce ADRs due to patient interindividual variability. Adverse side effects of phenytoin are common and often include rashes, headache, behavioral changes, Steven-Johnson syndrome, cardiovascular collapse, and arhythmicias, especially with rapid intravenous administration, in both children and adults (Bansal et al., 2013; Polat et al., 2015). Many pediatric patients with severe epilepsy are often treated with polytherapy, and interactions between phenytoin and other AEDs and the resulting adverse effects have been documented; for example, concurrent treatment with phenytoin and valproic acid produced phenytoin toxicity in an adolescent epileptic patient despite adhering to pediatric dosing guidelines for phenytoin (Carvalho et al., 2014). A better understanding of the mechanism behind phenytoin-mediated P450 induction via CAR, especially through postnatal development, could help alleviate the high risk of ADRs that commonly accompany phenytoin treatment. Despite the high risk of potentially dangerous side effects, phenytoin might be a better therapeutic option for seizure control in infants since it may not produce the permanent effects on P450 expression and liver function caused by phenobarbital. These data warrant further studies in humans.

**References**

**Authorship Contributions**
**Participated in research design:** Piekos, Zhong, Ma, Zhu.
**Conducted experiments:** Piekos, Chen, Wang, Shi, Yaqoob.
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**Wrote or contributed to the writing of the manuscript:** Piekos, Zhong, Ma, Zhu.

**Fig. 4.** Effects of phenytoin (PHY) treatment on the expression of CYP3A16 during postnatal maturation. Male mouse mice were treated with either vehicle control (PBS) or 100 mg/kg PHY at different ages following birth. Twenty-four hours after treatment, livers were collected for quantification of mRNA by RT-PCR, and then fold changes were calculated using the 2^(-ΔΔCt) method. Data points represent fold changes in gene expression of individual mice treated with PBS or PHY compared with the average ΔCt of mice that received PBS control (A). Ct values from RT-PCR experiments were also compared with the GAPDH Ct value in each individual mouse and the differences are expressed as 2^(-ΔCt) (B). The dotted line represents GAPDH expression set to 1 (n = 4–6). Protein was quantified using liquid chromatography-tandem MS 24 hours following PHY treatment in male neonates (day 5) and male adults (day 60) (C). The data are presented as mean ± S.E. *P < 0.05; **P < 0.01; ***P < 0.001.

**Table 4.** Relative amount of protein with PBS and PHY treatment.


