Tissue Distribution, Ontogeny, and Chemical Induction of Aldo-Keto Reductases in Mice

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ABSTRACT

Aldo-keto reductases (Akrs) are a conserved group of NADPH-dependent oxido-reductase enzymes. This study provides a comprehensive examination of the tissue distribution of the 16 substrate-metabolizing Akrs in mice, their expression during development, and whether they are altered by chemicals that activate distinct transcriptional pathway. Akrlc6, 1c14, 1c20, and 1c22 are primarily present in liver; Akrla14, 1c18, 1c21, and 7a5 in kidney; Akrld1 in liver and kidney; Akrlb7 in small intestine; Akrlb3 and Akrlc11 in brain; Akrlc8 in testes; Akrlc14 in ovaries; and Akrlc12, 1c13, and 1c19 are expressed in numerous tissues. Liver expression of Akrld1 and Akrlc1 is lowest during prenatal and postnatal development. However, by 20 days of age, liver Akrld1 increases 120-fold, and Akrlc mRNAs increase as much as 5-fold (Akrlc19) to 1000-fold (Akrlc6). Treatment of mice with chemical activators of transcription factors constitutive androgen receptor (CAR), pregnane X receptor (PXR), and the nuclear factor-erythroid-2 (Nrf2) transcription factor alters liver mRNAs of Akrs. Specifically, CAR activation by 1,4-bis-[2-(3,5-dichloropyridyloxy)]benzene (TCPOBOP) increases mRNAs of Akrlb7, Akrlc6, Akrlc19, and Akrld1, whereas PXR activation by 5-pregnenolone-16α-carbonitride (PCN) increases the mRNA of Akrlb7 and suppresses mRNAs of Akrlc13 and Akrlc20. The Nrf2 activator 2-cyano-3,12-dioxooleana-1,9-dien-28-imidazolide (CDDO-Im) induces mRNAs of Akrlc6 and Akrlc19. Moreover, Nrf2-null and Nrf2 overexpressing mice demonstrate that this induction is Nrf2-dependent.

Introduction

Aldo-keto reductases (Akrs) are a conserved group of NADPH-dependent oxido-reductase enzymes with (α/β)3-barrel structural similarities (Jez et al., 1997). Most Akrs are monomeric 34–37 kDa proteins; however, multimeric proteins are found in Akr2, Akr6, and Akr7 family members. Akrs have been identified in multiple species, including eubacteria, fungi, plants, and vertebrates. Akrs metabolize a wide variety of substrates, such as steroid hormones, carbohydrates, and xenobiotics (Seery et al., 1998). The substrate specificity is dependent on three flexible loops (A, B, and C) (Hoog et al., 1994).

Three Akr families have been identified. Mammalian Akrs (Table 1) are found in the Akr1, Akr6, and Akr7 families (Jin and Penning, 2007). The Akr6 family members are involved in inactivation of voltage-gated potassium channels (Xie et al., 2011). The Akr7 family members reduce a metabolite of aflatoxin (B1-dihydropdiol), a toxin produced by the fungus Aspergillus flavus, preventing the aflatoxin metabolite from forming damaging protein-adducts (Guengerich and Johnson, 1999).

Akr1 is the largest family of Akrs. The Akr1c, 1d, and 1e families are essential in the metabolism of hormones, including androgens, estrogens, and progesterone, as well as prostaglandins (Seery et al., 1998; Penning et al., 2000). Akrla4 is involved in the reduction of D-glyceraldehyde to glycerol. The Akr1b family of enzymes (Akr1b3, 1b7, and 1b8) is responsible for the reduction of toxic aldehydes generated during lipid peroxidation and steroidogenesis (Spitz et al., 2007). In addition to detoxification activity, Akr1b3 and Akr1b7 also have prostaglandin F2α synthase activity (Kabututu et al., 2009).

The Akr1c subfamily has been viewed as a drug target because many Akr1c enzymes selectively metabolize steroid hormones as well as xenobiotics. In mice, nine members of the Akr1c family have been identified: Akr1c6, Akr1c12, Akr1c13, Akr1c14, Akr1c18, Akr1c19, Akr1c20, Akr1c21 and Akr1c22 (Deshayeshi et al., 1995; Du et al., 2000; Matsumoto et al., 2006). Akr1c6 is required for the formation of testosterone from 4-androstenedione (Rheault et al., 1999). Akr1c12 and Akr1c13 oxidize acyclic alcohols, aliphatic alcohols, 3α-hydroxysteroids, 17β-hydroxysteroids, and 20α-hydroxysteroids (Endo et al., 2006, 2007). Akr1c18 inactivates progesterone by reducing it to its inactive metabolite, 20α-hydroxyprogesterone (Mao et al., 1997). Akr1c20 and Akr1c21 metabolize multiple steroid hormones (Matsumoto et al., 2006; Dhagat et al., 2008).

Akrs are important for the regulation of many endogenous pathways. Akr1c19 reduces isatin in the gastrointestinal tract. Isatin is a pharmacologically active molecule produced by intestinal bacteria (Isshiki et al., 2005). Akr1l1 reduces anhydro-D-fructose to anhydro-D-glucitol (Sakuma and Kubota, 2008), which is important in the regulation of glycoprotein. Akrs are also important in the synthesis of bile acids from cholesterol (Stolz et al., 1984; Dufort et al., 1996). Akr1c6 is responsible for the 3α-reduction, and Akr1d1 is responsible for the 5β-reduction of...
7,8-dihydroxybenzopyrene. After several auto-oxidation steps, an 7,8-diol to form a ketol. Tautomerization of this product leads to Human Akrs 1B1 and 1B10 catalyze the oxidation of benzopyrene-carbons are byproducts of fossil fuel combustion and tobacco smoke.

2000; Mindnich et al., 2011).

prevents AFB1-dialdehyde from forming lysine adducts (Ellis et al., 1996, 1999).

is cytotoxic because of its ability to form protein adducts. Akr7a (aflatoxin aldehyde reductase) reduces AFB1-dialdehyde to form an alcohol, and this reduction prevents AFB1-dialdehyde from forming lysine adducts (Ellis et al., 1993).

In general, little is known about the regulation of Akrs. Therefore, in the present study, the tissue distribution of mouse Akrs was determined in adult male mice. In addition, an ontogeny study was performed to determine the pattern of Akr expression during liver and kidney development. Lastly, we sought to determine whether chemicals known to induce drug-metabolizing enzymes and transporters will also alter the mRNA expression of various Akr isoforms. The inducers include 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD), which activates the aryl hydrocarbon receptor (AhR); 1,4-bis[3,5-dichloropyridyldioxy]) benzene (TCPOBOP), which activates the constitutive androstane receptor (CAR); 5-pregnenolone-16α-carbonitrile (PCN), which activates the pregnane X receptor (PXR); clofibrate, which activates the peroxisome proliferator-activated receptor (PPARα); and 2-cyano-3,12 dioxygenane-1,9 dien-28-imidazolide (CDDO-Im), which activates nuclear factor-erythroid-2 (Nrf2). Finally, those Akr genes with expression altered by the Nrf2 agonist CDDO-Im, were further examined in genetically altered mice that have either hepatic knockout or overexpression of Nrf2.
Materials and Methods

Chemicals. TCPOBOP and 5-pregnenolone-16α-carbonitrile (PCN) were purchased from Sigma-Aldrich (St. Louis, MO). TCDD was a gift from Dr. Karl Rozman (University of Kansas Medical Center, Kansas City, KS). Clofibrate was purchased from Fisher Scientific (Hampton, NH). (CDDO-Im) was a gift from Reata Pharmaceuticals (Irvine, TX).

Mice. Eight-week-old C57BL/6J mice were purchased from Jackson Laboratory (Bar Harbor, ME) to determine the tissue distribution, ontogeny, and chemical induction of Akr. Nrf2-null mice (Chan et al., 1996) were obtained from Dr. Dr. Masayuki Yamamoto (Tohoku University, Sendai, Japan). Keap1-knockdown (KD) mice (Okada et al., 2008) were supplied by Dr. Dr. Karl Rozman (University of Kansas Medical Center, Kansas City, KS). Nrf2-null and Keap1-KD mice were backcrossed into the C57BL/6 background, and 99% congenicity was confirmed by Jackson Laboratory. The mice were housed on corn cob bedding according to the Association for Assessment and Accreditation of Laboratory Animal Care guidelines. All animals were given ad libitum access to water and standard rodent chow (Harlan Teklad 8604; Harlan Teklad, Madison, WI).

For tissue distribution studies, 8-week-old male (n = 6) and female (n = 6) mice were used. Liver, kidney, lung, stomach, duodenum, jejunum, ileum, colon, brain, testes, ovaries, and heart were collected from mice. The tissues were snap-frozen in liquid nitrogen. The intestine was longitudinally dissected, rinsed in saline, and divided into three equal-length sections (referred to as duodenum, jejunum, and ileum) before being frozen in liquid nitrogen. All tissues were stored at −80°C. For the ontogeny study, liver was collected from male mice (n = 5) at postnatal days −2, 0, 1, 3, 5, 10, 20, and 45. Kidney was collected from male mice (n = 4) at postnatal days −2, 0, 10, 20, and 30. Brain was collected from male mice (n = 4) at postnatal days 0, 10, 20, and 30. For chemical induction of Akr by Cyp inducers, the activators were administered to male mice (n = 5) for 4 days (intraperitoneally) as described previously (Cheng et al., 2005). Tissues were collected 24 hours after the final dose.

RNA Isolation, cDNA Synthesis, and Real-Time Polymerase Chain Reaction. Total RNA was isolated from 50 mg of snap-frozen liver using RNAzol B reagent purchased from Tel-Test Inc. (Friendswood, TX), and quantified using a NanoDrop Spectrophotometer (NanoDrop Technologies, Wilmington, DE) at an UV absorbance of 260 nm. Two micrograms of total RNA was used for cDNA synthesis performed using a high-capacity cDNA synthesis kit from Applied Biosystems (Foster City, CA). Oligonucleotide primer sequences were designed by NCBI Primer-BLAST to be specific to 16 Akr mRNAs (detailed in Table 2). Primers were then aligned to the mouse genome to test for specificity. Real-time polymerase chain reaction (PCR) analysis was performed using a StepOnePlus instrument from Applied Biosystems. The mRNA of each gene was adjusted to the mRNA of RLP13. Relative levels of each gene were evaluated using the ΔΔCt method.

Statistics. Statistical differences between multiple groups were analyzed by a one-way analysis of variance followed by Duncan’s post hoc test. Statistical significance was considered at P < 0.05.

Results

Tissue Distribution of Mouse Akr.s. The mRNAs of sixteen Akr.s were quantified in 12 mouse tissues as shown in Fig. 1. Akr6 members were not quantified because they are only distantly related (15%–30% homology) to other Akr.s, and they function as inactivators of voltage-gated potassium channels (McCormack and McCormack, 1994; Barski et al., 2009). Tissues from six male and six female mice were used to determine the expression of these Akr genes in liver, kidney, lung, stomach, duodenum, jejunum, ileum, colon, brain, testes, ovaries, and heart. No gender differences in expression of any Akr mRNA were detected (unpublished data). Akr1a4 mRNA was highest in kidney; however, there was also expression in liver, lungs, stomach, and small intestine. Akrb3 mRNA was 20 times more prevalent in brain than in other tissues (Fig. 1). Akrb7 mRNA was highest in the small intestine, with similar amounts in duodenum, jejunum, and ileum (Fig. 1). There was also quantifiable Akrb7 mRNA in kidney; however, it was only 1/20 of that in small intestine. Akrb8 mRNA was highest in testes and second highest in stomach (Fig. 1). Akr1c6 mRNA was found to be highest in liver, with levels at least 200-fold higher than those quantified in other tissues (Fig. 1). Akr1c12 and Akr1c13 were expressed in liver, stomach, duodenum, jejunum, ileum, and colon (Fig. 1). Akr1c14 was expressed at levels at least 4-fold higher in liver and ovaries than in other tissues (Fig. 1). Akr1c18 was expressed in kidneys at levels 500-fold higher than those in other tissues (Fig. 1). Akr1c19 mRNA was expressed in liver, kidney, stomach, duodenum, jejunum, ileum, and colon (Fig. 1). Akr1c20 was mainly expressed in liver, at quantities 50-fold higher than seen in other tissues (Fig. 1). Akr1c21 was expressed mainly in kidney. Akr1c22 was expressed highest in liver and stomach, with the mRNA about 20-fold higher than in other tissues (Fig. 1). Akr1d1 was expressed highest in liver and kidney, with the mRNA about 66- and 33-fold higher, respectively, than in other tissues (Fig. 1). Akr1e1 mRNA was highest in brain, with a value 3-fold higher than in other tissues (Fig. 1). Akr1f5 mRNA was 10-fold higher in kidney than in other tissues (Fig. 1).

Ontogeny of Akr.s in Mouse Liver, Kidney, and Brain. A developmental study was performed to determine the pattern of Akr expression during liver, kidney, and brain development. First, the Akr.s

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward</th>
<th>Reverse</th>
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<tr>
<td>Akr1a4</td>
<td>ATAGGCCCTTGCGGCCTTCCT</td>
<td>GACCTGACCTGAGATGACGC</td>
</tr>
<tr>
<td>Akr1b5</td>
<td>GTGACTCTTTAACTGTGCTGGC</td>
<td>TCTCGAGCCTGCTTTCTCTT</td>
</tr>
<tr>
<td>Akr1b7</td>
<td>GCCACAGGAGGCCTGAGCCTT</td>
<td>ACCGCTCCTTCAACAGGAGTGACC</td>
</tr>
<tr>
<td>Akr1b8</td>
<td>CCACTGGTCTGGGACTACAGTACC</td>
<td>TGTGGCAGTAKGACATACCA</td>
</tr>
<tr>
<td>Akr1e6</td>
<td>TCCTCCGAAACTCTGTAAGGGT</td>
<td>TCCCGCTCCTAGGGCAATGG</td>
</tr>
<tr>
<td>Akr1e12</td>
<td>CGACGATCGTCTTCTGCGACCGC</td>
<td>CGACGATCGTCTGCGACCGC</td>
</tr>
<tr>
<td>Akr1c14</td>
<td>TCTGGTGACTTCCGTGTGGTCA</td>
<td>GCTGCTCAAGCCCTCGAGGAG</td>
</tr>
<tr>
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<td>CTTCTGAGTCAACAGGAGTCTAGAG</td>
<td>TGTGGCAGTACGCAAGGGCA</td>
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<tr>
<td>Akr1c19</td>
<td>AGTCAGCTTCCTGCTTGTGAG</td>
<td>GCTGCTCAAGCCCTCAGG</td>
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<tr>
<td>Akr1c20</td>
<td>GAACCTTGCTCGTCAGGCA</td>
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<td>Akr1c21</td>
<td>GCAGCTGTCTGCTGTGAG</td>
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<td>Akr1c22</td>
<td>GGCACCTTTAATCTCCCCT</td>
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<tr>
<td>Akr1d1</td>
<td>TAAGAGTGTGGGTGGC</td>
<td>CGACGATCGTCTGCGACCG</td>
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<td>Akr1e1</td>
<td>AGCCCTGGTTGAGGCTGACGC</td>
<td>GACGCGATCGTCTGCGAGG</td>
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<tr>
<td>Akr7a5</td>
<td>TACTCAGCGAGCCAGTGCAG</td>
<td>CCGTCTCTGCGGCGGCTC</td>
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that were significantly expressed in livers of adult mice were examined, namely Akr1c6, Akr1c14, Akr1c19, Akr1c20, Akr1d1, Akr1e1, and Akr7a5. The mRNAs of these Akrs were quantified at postnatal days 2, 0, 1, 3, 5, 10, 20, and 45.

Akr1c6 mRNA increased between 5 and 10 days of age and reached adult expression by 20 days of age (Fig. 2). Akr1c14 mRNA increased markedly between 10 and 20 days of age (Fig. 2). Akr1c19 and Akr1c20 mRNA also increased between 10 and 20 days of age. Akr1c19 mRNA increased 5-fold, and Akr1c20 increased more than 2-fold during this time interval (Fig. 2). Akr1e1 mRNA also increased between 10 and 20 days of age, which was about a 4-fold increase (Fig. 2). Akr1d1 mRNA was barely detectable before birth (−2), was detectable at day 1 of age, and reached adult levels at 20 days of age (Fig. 2). The mRNA of Akr7a5 did not change markedly in livers from 2 days before birth to 45 days of age.

The Akrs that were significantly expressed in kidney of adult mice were examined to determine how expression changes during development. The mRNAs were quantified at postnatal days 22, 0, 10, 20, and 30. The mRNAs of Akr1a4 and Akr1c19 in kidneys did not change during the first month of life (Fig. 3). Akr1c18 and Akr1c21 mRNAs in kidneys of mice increased 7-fold and 200-fold, respectively, from 2 days before birth to 30 days of age (Fig. 3). Akr1d1 mRNA increased about 3-fold after TCPOBOP administration (Fig. 4).

The Akrs that were expressed in brain of adult mice were also examined to determine whether the mRNA of the Akrs change during the first month of life. The mRNAs of Akr1b7, Akr1c18, Akr1e1, and Akr7a5 were quantified from brains of mice at postnatal days 0, 10, 20, and 30. There were no significant age-relevant changes in the mRNAs observed for any of these Akrs (Supplemental Fig. 1).

Chemical Induction of Mouse Akrs. Five different chemicals known to activate various transcription factors that induce cytochrome P450 enzymes as well as other drug metabolizing enzymes and transporters were used in this study (Aleksunes and Klaassen, 2012). The inducers used were: TCDD for AhR, TCPOBOP for CAR, PCN for PXR, clofibrate for PPARα, and CDDO-Im for Nrf2. Akr1a4, Akr1b3, Akr1b8, Akr1c21, Akr1e1, and Akr7a5 had low amounts of measurable mRNA in liver before and after inducer treatment (Supplemental Fig. 2). Akr1b7 mRNA was very low in liver of control mice but increased about 300-fold after TCPOBOP and 15-fold after PCN administration (Fig. 4).

Akr1c6 mRNA increased about 50% after TCPOBOP and 100% after CDDO-Im administration (Fig. 4). Akr1c12 mRNA was unresponsive to any of the activators. Akr1c13 mRNA decreased more than 50% after PCN and clofibrate treatments (Fig. 4). Akr1c14 was unresponsive to all five activators (Fig. 4). Akr1c18 mRNA increased more than 400% after clofibrate administration. Akr1c19 mRNA increased more than 3-fold after TCPOBOP and more than 5-fold after CDDO-Im administration. Akr1c20 mRNA decreased 80% after TCPOBOP and 75% after treatment with PCN (Fig. 4). Akr1c22 mRNA decreased 80% after TCPOBOP and TCDD administration, and was undetectable after PCN administration. Akr1d1 mRNA more than doubled after TCPOBOP administration (Fig. 4).
Gene Dose-Response of Nrf2 on Akr1c6 and Akr1c14. Because CDDO-Im is a Nrf2 activator and increases the expression of Akr1c6 and Akr1c14, a gene dose–response model was employed to further analyze the role of Nrf2 in the regulation of these genes. Nrf2 is a short-lived protein that is readily ubiquitylated by CUL3-RBX1 and degraded by the 26S proteasome. The ubiquitylation of Nrf2 depends on Kelch-like ECH-associated protein 1 (Keap1), which functions as a substrate adaptor for CUL3-RBX1 (Kobayashi et al., 2004). If Keap1 is not present or inactivated, Nrf2 translocates into the nucleus. The four types of mice used to make the Nrf2 gene dose response were Nrf2-null, wild-type, Keap1-KD, and Keap1-HKO. In this model, Nrf2-null mice possess no Nrf2 activity, and the Keap1-HKO mice possess the highest level of Nrf2, as characterized previously (Wu et al., 2011). Mice that lack Nrf2 had a 50% decrease in Akr1c6 mRNA (Fig. 5). The Keap1-HKO mice had a 5-fold increase in Akr1c19 (Fig. 5). The transcription of a gene whose transcription was not affected by CDDO-Im treatment was also examined (not shown). The elimination of Nrf2 or Keap1 had no effect on the transcription of Akr1c14, which mirrored the results from the chemical treatment study. These results further indicate that Nrf2 is important for the regulation of Akr1c6 and Akr1c19.

**Discussion**

Akr1s are important enzymes for the synthesis of endogenous compounds as well as for the detoxification of xenobiotics. To better
understand the expression of these genes, this study first quantitatively determined the expression of the Akrs in various tissues, although other studies have performed limited tissue distribution studies of a few Akrs. The present study examined more Akrs, more mice, and contained mice of both.

The Akr1b family (Akr1b3, 1b7, and 1b8) of enzymes is responsible for the reduction of toxic aldehydes generated during lipid peroxidation and steroidogenesis (Spite et al., 2007). Akr1b family members (Akr1b3, Akr1b7, and Akr1b8) also possess prostaglandin F2α synthase activity (Kabututu et al., 2009). In addition, Akr1b7 is important for metabolizing 3-keto bile acids to 3β-hydroxy bile acids. Toxic bile acids, such as desoxycorticosterone acetate, are converted to less toxic 3β bile acids, such as 3βDCA, by Akr1b7 (Schmidt et al., 2011). The present data indicates for the first time that Akr1b3 is expressed mainly in brain. Akr1b7 was expressed mainly in intestine, which confirms an earlier published result that was obtained using the RNase protection assay (Lau et al., 1995). In the present study, Akr1b8 was located mainly in testes and stomach. This finding somewhat contradicts a previous report using qualitative reverse transcriptase PCR of Akr1b8 in which expression was observed in equal amounts in most tissues except brain and kidney, which exhibited lower expression (Salabei et al., 2011). The results of the present study may be different because the previous study did not include stomach and was only determined after 28 cycles of PCR.

The Akr1c family members (Akr1c6, 1c12, 1c13, 1c14, 1c18, 1c19, 1c20) are viewed as drug targets because of their roles in steroid

![Fig. 4. Effects of chemical induction on C57BL/6 mouse Akr mRNA expression in liver. The dose of chemical treatment (n = 5) was described under Materials and Methods. Data are presented as mean ± S.E.M. Asterisks (*) indicate statistically significant increase/decrease in mRNA level after treatment compared with the control group by analysis of variance (P < 0.05).](image1)

![Fig. 5. Analysis of mRNA expression of Akr1c6, and Akr1c19 by real-time PCR analysis of the four groups (n = 5 per group). The different genotypes included Nrf2-null, wild-type, Keap-1-KD, and Keap1-HKO. Data are presented as mean ± S.E.M. Asterisks (*) indicate statistically significant differences from wild-type mice (P < 0.05).](image2)
biogenesis and xenobiotic metabolism. Previous reports have demonstrated that many of these Akr1c family members are expressed in livers of mice (Vergnes et al., 2003; Velica et al., 2009). Akr1c6 is almost exclusively expressed in liver (Fig. 1), which corresponds with previously published data (Vergnes et al., 2003; Velica et al., 2009). Akr1c12 and Akr1c13 mRNA are ubiquitously expressed. Akr1c14 expression is highest in liver and in ovaries. This finding was different from a previous report that found the highest expression of Akr1c14 in kidney, and similar expression in liver, kidney, stomach, intestine, colon, lung, and ovary (Velica et al., 2009). Akr1c18 mRNA was found mainly in the kidney and at a lower concentration in brain. This result also differed from the publication of Velica et al., which reported expression only in ovary. These differences could be the result of differences in mouse strain because Velica et al. used CD1 mice, whereas this study used C57BL/6 mice. Another possible cause of the variance is that Velica et al. used pregnant and nonpregnant female mice. Pregnancy might have increased the expression of Akr1c18 mRNA.

The expression of Akr genes was examined at different stages of liver, kidney, and brain development. The liver undergoes marked developmental changes after birth. Liver changes from a hematopoietic organ before birth to an organ necessary for xenobiotic metabolism after birth. The expression of the P450 enzymes and transporters in liver changes over the course of development (Hart et al., 2009; Cui et al., 2012a,b; Lu et al., 2012). An increase in the mRNA of several Akr1c genes in liver was observed after birth (Fig. 2). Akr1d1 mRNA increased over 100-fold between birth and day 45. Akr1c20 mRNA increased over 50-fold, Akr1c6 increased over 10-fold, and Akr1c4 and Akr1c19 increased in liver to a lesser extent between birth and day 45 (Fig. 2).

The expression of several Akr genes in kidney was also quantified in kidney at multiple ages. Similar to some of the Akr genes described previously in liver, two Akr mRNAs increase during development in kidney. Akr1c18 mRNA increased over 7-fold, and Akr1c21 mRNA increased in kidney about 200-fold from birth to 30 days of age (Fig. 3).

Various transcription pathways are known to increase various drug-metabolizing enzymes in liver, therefore we sought to determine whether these pathways also alter the expression of Akr. AHR activation led to the increase of mRNA of Akr1b7. CAR activation led to increased mRNA of Akr1b7, which had been described previously (Liu et al., 2009), and Akr1c6. PXR activation caused an increase of Akr1b7 and a decrease in the mRNAs of Akr1c13, Akr1c20, and Akr1c22 (Fig. 4). PXR activation has been reported previously to activate distinct transcription factor pathways. Drug Metab Dispos XX:1276–1282. Cui JY, Gettena K, Yu H, Li L, Renaud HJ, Lu H, Zheng XB, and Klaassen CD (2012a) RNA-Seq reveals different mRNA abundance of transporters and their alternative transcript isoforms during postnatal liver maturation. Proc Natl Acad Sci USA 109:13934–13948. Cui JY, Gettena K, Yu H, Li L, Renaud HJ, and Klaassen CD (2012b) Ontogeny of novel cytochrome P450 gene isoforms during postnatal liver maturation in mice. Drug Metab Dispos 40:1226–1237.


Ontogeny of mouse Akr genes in brain. Total RNA from mice at each age (n=4) was analyzed by real time PCR. Gene control used was RPL13a. Data are presented as mean ± SEM.
Supplemental Figure 2

Effects of chemical induction on C57BL/6 mouse Akr mRNA expression in liver.

The dose of chemical treatment (n=5) was described under Materials and Methods. Data are presented mean ± SEM. Asterisks (*) indicate statistically significant increase/decrease in mRNA level after treatment compared with the control group by ANOVA (p < 0.05).