A Phosphomimetic Mutation at Threonine-57 Abolishes Transactivation Activity and Alters Nuclear Localization Pattern of Human Pregnane X Receptor

Satyanarayana R. Pondugula, Cynthia Brimer-Cline, Jing Wu, Erin G. Schuetz, Rakesh K. Tyagi, and Taosheng Chen

Departments of Chemical Biology and Therapeutics (S.R.P., J.W., T.C.) and Pharmaceutical Sciences (C.B.-C., E.G.S.), St. Jude Children’s Research Hospital, Memphis, Tennessee; and Special Centre for Molecular Medicine, Jawaharlal Nehru University, New Delhi, India (R.K.T.)

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The pregnane X receptor (PXR) plays crucial roles in multiple physiological processes. However, the signaling mechanisms responsible are not well defined; it is most likely that multiple functions of PXR are modulated by its phosphorylation. Therefore, we sought to determine whether mutation at a highly conserved Thr57 affects human PXR (hPXR) function. Site-directed mutagenesis was performed to generate phosphorylation-deficient (hPXRT57A) and phosphomimetic (hPXRT57D) mutants. Gene reporter, Western blotting, immunocytochemistry, mammalian two-hybrid, and electrophoretic mobility shift assays were used to study cytochrome P450 3A4 (CYP3A4) promoter activation, protein levels, localization, cofactor interaction, and CYP3A4 promoter binding of the hPXR mutants, respectively. hPXRT57D, but not hPXRT57A, lost its transactivation activity. Neither mutation altered hPXR’s protein levels and interaction with steroid receptor coactivator-1. hPXR and hPXRT57A exhibited a homogenous nuclear distribution, whereas hPXRT57D exhibited a distinctive punctate nuclear localization pattern similar to that of hPXR mutants with impaired function that colocalize with silencing mediator of retinoid and thyroid receptors (SMRT), although silencing of SMRT did not rescue the altered function of hPXRT57D. However, hPXRT57D, but not hPXRT57A, impaired hPXR’s ability to bind to the CYP3A4 promoter, consistent with the mutant’s transactivation function. Furthermore, the 70-kDa form of ribosomal protein S6 kinase (p70 S6K) phosphorylated hPXR in vitro and inhibited its transactivation activity, whereas hPXRT57A partially resisted the inhibitory effect of p70 S6K. Our studies identify a functionally significant phosphomimetic mutant (hPXRT57D) and show p70 S6K phosphorylation and regulation of hPXR transactivation to support the notion that phosphorylation plays important roles in regulating hPXR function.

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et al., 2007), cancer cell growth and drug resistance (Chen et al., 2007; Mensah-Osman et al., 2007; Zhou et al., 2008), uterinecontractility and vascular tone (Mitchell et al., 2005; Hagedorn et al., 2007), blood-brainbarrier permeability (Bauer et al., 2006), and neuroprotection (Langmade et al., 2006).Nevertheless, the signaling mechanisms responsible for its diversecellular responses are not well defined, although analogous toother NRs, it is most likely that multiple functions of PXR aremodulated and integrated by its phosphorylation.

Phosphorylation is a dynamic regulatory mechanism that not onlyaffects the function of a protein but also enables both specificity andcross-talk among diverse signaling pathways (Cohen, 2000, 2001).Phosphorylation has been shown to modulate the activity of manyNRs, and it provides an important mechanism for cross-talk betweensignaling pathways (Shao and Lazar, 1999; Weigel and Moore, 2007).All the aspects of NR function are known to be regulated when specific sites are phosphorylated, including expression, stability, subcellular localization, dimerization, ligand and DNA binding, coregulatorinteraction, and transcriptional activity (Ort et al., 1992; Shao andLazar, 1999; Rochette-Egly, 2003; Sun et al., 2007; Weigel andMoore, 2007). Furthermore, it is known that NR phosphorylationplays a crucial role in the development of diseases such as breast,ovarian, and prostate cancers (Rochette-Egly, 2003). In contrast toother NRs, our current understanding on phosphorylation of PXR isextremely poor despite substantial evidence for its considerable roleasa xenobiotic sensor with a promiscuous ligand-binding nature resultingin adverse drug-drug interactions (Kliwer et al., 1998;Lehmann et al., 1998; Harmsen et al., 2007) and as a physiologicalsensor and signal cross-talker in diverse cellular responses. Therefore,an understanding of phosphorylation-dependent events in PXR signalingis crucial for effective drug design and clinical therapeutic strategies.

Similar to steroid receptors, the N-terminal part of the PXR proteincontains a highly conserved DNA binding domain (DBD) (Fig. 1A),and the C-terminal part of the protein contains a ligand bindingdomain (LBD) with an additional ligand-inducible transactivationfunction 2 (AF-2). The DBD and LBD are separated by a hinge region(Fig. 1A). It can be argued that, in contrast to most NRs, PXR does notharbor ligand-independent AF-1 (Fig. 1A), which is the most variableregion in NRs in terms of length and sequence similarities and isbelieved to be regulated by growth factor signaling. NRs are phosphorylated on all the domains at specific sites depending on thecellular context, and such site-specific phosphorylation can lead toeither modulation or termination of the activity of the NRs (Rochetti-Egly, 2003; Weigel and Moore, 2007). Although it is known thatprotein kinase A (PKA) (Ding and Staudder, 2005a), protein kinaseC (PKC) (Ding and Staudder, 2005b), and cyclin-dependent kinase2 (CDK2) (Lin et al., 2008) are involved in the regulation of PXR activity, and that PKA and CDK2 can phosphorylate the PXR in vitro(Ding and Staudder, 2005a; Lin et al., 2008), very little is known aboutthespecific site(s) of phosphorylation. Therefore, we examined one ofthe highly conserved putative phosphorylation sites by performing a phosphomimetic mutation at Thr57 in the DBD and by testing thetranscriptional activity, promoter-binding activity, expression, cofactorinteraction, and localization of the mutant. Furthermore, results fromphosphorylation site prediction and rapamycin-induced PXR modulation (see under Results) led us to test whether the 70-kDaform of ribosomal protein S6 kinase (p70 S6K) regulates thetranscriptional activity of human preganme X receptor (hPXR) throughphosphorylation of Thr57.

In this report, we identify a phosphomimetic mutant of hPXR thatloses its transactivation function and displays a punctate nucleardistribution. Our results indicate that a phosphomimetic mutation atThr57 (hPXR T57S) impairs the ability of hPXR to bind to the promoterof cytochrome P450 3A4 (CYP3A4), an important hPXR target gene.

In addition, we show that p70 S6K phosphorylates hPXR in vitro and inhibits its transactivating function, and that phosphorylation-deficient mutation at Thr57 (hPXR T57S) confers significant resistance to the inhibitory effect of p70 S6K. Taken together, our observations suggest that Thr57 might be regulated by phosphorylation, which in turn regulates thefunction of hPXR. These observations might apply to other NRs dueto the highly conserved nature of this residue.

Materials and Methods

Cell Culture, Plasmids, and Transient Transfections. HepG2 human liver carcinoma and COS7 African green monkey kidney fibroblast cell lines were obtained from the American Type Culture Collection (Manassas, VA). The cells were grown in Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen, Carlsbad, CA) supplemented with 10% fetal bovine serum (FBS; HyClone, Logan, UT), 100 U/ml penicillin, 100 μg/ml streptomycin (Invitrogen), 2 mM l-glutamine (Invitrogen), and 1 mM sodium pyruvate (Invitrogen). The cells were cultured in an incubator with a humidified atmosphere maintained at 5% CO2 and 95% air at 37°C.

To generate the pcDNA3-hPXR, pcDNA3-FLAG-hPXR, and CYP3A4 luciferase reporter (pGL3-CYP3A4-luc) plasmids were generated as described previously (Lin et al., 2008). Site-directed mutagenesis was performed to generate alanine (hPXR T57A) and aspartate (hPXR T57D) mutant hPXR plasmids. Polymerase chain reaction (PCR) was performed using pcDNA3-hPXR as the template and specific mutant primers, with forward primers containing a mutation corresponding to either A or D and a PPUMI restriction enzyme cutting site and a reverse primer containing a BsgI restriction enzyme cutting site. The PCR product was cloned into a TOPO vector (Zero Blunt TOPO PCR Cloning Kit; Invitrogen), followed by cleaving the sequence-verified TOPO-PCR product for the desired mutations with PPUMI (New England Biolabs, Ipswich, MA) and BsgI (New England Biolabs), and ligating the resulting fragment into PPUMI- and BsgI-cleaved pcDNA3-hPXR or pcDNA3-FLAG-hPXR. Only the open reading frame of hPXR, but not pcDNA3 or the FLAG tag, had cleavage sites for PPUMI and BsgI. Sequence-verified pcDNA3-hPXR T57A, pcDNA3-hPXR T57D, pcDNA3-FLAG-hPXR T57A, and pcDNA3-FLAG-hPXR T57D were used for transient transfections. pACT contains the transcriptional activation domain of VP16, a herpes simplex virus protein (Promega, Madison, WI). pBIND contains the DBD of the yeast transcription factor Gal4 (Promega). Full-length wild-type hPXR, hPXR T57A, and hPXR T57D were fused to VP16 in pACT to generate VP16-hPXR, VP16-hPXR T57A, and VP16hPXR T57D, respectively. DNA encoding the NR interaction domain of steroid receptor coactivator (SRC)-1 (amino acids 621–765) (Ohate et al., 1995) was fused to Gal4 in pBIND to generate Gal4-SRC-1. pG5-luc contains the luciferase reporter gene controlled by a promoter containing Gal4 DNA binding sites (Promega). A constitutively active p70 S6K plasmid, Mycz-p70S6K D3E-E389, was a gift from Dr. George Thomas (The Genome Research Institute of The University of Cincinnati, Cincinnati, OH).

Transient transfections were performed using FuGENE 6 (Roche Diagnostics, Indianapolis, IN), unless otherwise mentioned, following the manufacturer’s protocol. Twenty-four hours post-transfection, the cells were grown in DMEM supplemented with 10% charcoal/dextran-treated FBS (HyClone) and above-mentioned antibiotics. They were then treated with 0.1% dimethyl sulfoxide (DMSO; Sigma-Aldrich, St. Louis, MO) or 10 μM compounds such as rifampicin (Sigma-Aldrich), SR12813 (Sigma-Aldrich), and ketocanozole (Sigma-Aldrich) for an additional 24 h (unless otherwise indicated) before performing luciferase assays for transactivation analyses, harvesting of cell lysates for Western blotting analyses for protein expression, and fixing of cells for immunolocalization studies.

For the rifampicin (Sigma-Aldrich) experiments, the final DMSO concentration was 1%. In p70 S6K overexpression experiments, hPXR-transactivating function was analyzed using 1 μM rifampicin with 0.01% final concentration of DMSO. In mammalian two-hybrid assays, 5 μM rifampicin with 0.1% final DMSO concentration was used.
FIG. 1. Thr57 in hPXR is conserved in other NRs. A, top panel showing the schematic comparison of the domain structures of a steroid receptor and hPXR. H, hinge region. Bottom panel shows the schematic representation of the two zinc finger motifs in the DBD of hPXR. The two zinc finger motifs are separated by a linker (from Lys62 to Arg76). The diagram was not drawn as per the scale in terms of number of residues. Note that Thr57 is located within the first zinc finger motif. B, Thr57 in hPXR is highly conserved among other human NRs. Of the 46 human NRs with DBD, 37 receptors (including all the classic steroid receptors) have a conserved Thr57. Expansion of the abbreviated nomenclature for the receptors is as follows: CAR, constitutive androstane receptor; VDR, vitamin D receptor; GR, glucocorticoid receptor; MR, mineralocorticoid receptor; PR, progesterone receptor; AR, androgen receptor; ER, estrogen receptor; TR, thyroid hormone receptor; ROR, retinoid-related orphan receptor; LXR, liver X receptor; FXR, farnesoid X receptor; TR2, testicular receptor 2; TR4, testicular receptor 4; COUP-TF, chicken ovalbumin upstream promoter transcription factor; EAR-2, v-erbA-related; NURR1, NR-related 1; NOR1, neuron-derived orphan receptor 1; Nurr1, orphan-derived orphan receptor; GCNF, germ cell nuclear factor; TLX, human homolog of the Drosophila tailless gene; PNR, photoreceptor cell-specific NR. C, Thr57 in hPXR is conserved in all the vertebrate species with a fully known PXR sequence. Chicken PXR is the only exception and has an S instead of T at that position.

Transfections were performed in six-well culture plates (Corning Inc., Corning, NY). Each well has approximately 10-cm² surface for cell culture growth. Cells in each well were transfected with 2 μg of total plasmid DNA. To test the transactivating activity of wild-type and mutant hPXRs, the cells were transfected with 50 ng of pcDNA3, wild-type or mutant hPXR, and 1000 ng of pGL3-CYP3A4-luc plasmids. For the rapamycin experiments, the cells were transfected with 2 μg of pcDNA3 and wild-type or mutant hPXR plasmids. For the phosphatase inhibitor experiments, the cells were transfected with 50 ng of pcDNA3 or hPXR and 1950 ng of pGL3-CYP3A4-luc plasmids. In p70 S6K overexpression experiments, the cells were transfected with 100 ng of FLAG-pcDNA3, pcDNA3-FLAG-hPXR, or pcDNA3-FLAG-hPXR^{S78A}, 500 ng of constitutively active p70 S6K; and 1000 ng of pGL3-CYP3A4-luc and 100 ng of pGL4-hRluc; FLAG-pcDNA3 was used to make up the total plasmid DNA to 2 μg in each transfection. In mammalian two-hybrid assays, the cells were transfected with 500 ng of pACT (empty vector control for VP16-hPXR), VP16-hPXR, VP16-hPXR^{T57A}, or VP16-hPXR^{T57D}, 500 ng of Gal4-SRC-1; 500 ng of pG5-luc; and 500 ng of pcDNA3 to make up the total plasmid DNA to 2 μg in each transfection.

**Western Blotting Analysis.** HepG2 and COS7 cells were lysed with radioimmunoprecipitation assay buffer (Thermo Fisher Scientific, Waltham, MA) containing protease inhibitor mixture (Thermo Fisher Scientific) and phosphatase inhibitor mixture (Thermo Fisher Scientific). Whole cell lysates containing approximately 25 μg of total protein were solubilized at 95°C for 4 min in Laemmli sample buffer (Bio-Rad, Hercules, CA) containing β-mercaptoethanol and were resolved using electrophoresis on Nupage 4 to 12% bis-Tris gels (Invitrogen) with Nupage MES SDS running buffer (Invitrogen). For immunoblotting, the proteins were transferred from unstained gels to nitrocellulose membranes using the iBlot gel transfer system (Invitrogen) and
iBlot gel transfer stacks (Invitrogen). After blocking with Odyssey blocking buffer (LI-COR Biosciences, Lincoln, NE) for 1 h, membranes were incubated for 1 h with primary antibodies against FLAG-tag (mouse monoclonal anti-FLAG M2 antibody; Sigma-Aldrich) at 1:1000 dilution or silencing mediator of retinoid and thyroid receptor (SMRT) (mouse monoclonal anti-SMRT antibody; GeneTex, Irvine, CA) at 1:500 dilution. All of the steps were conducted at room temperature.

**Immunocytochemistry.** Twenty-four hours post-transfection, approximately 5000 live cells were seeded in each well of a 96-well culture plate (PerkinElmer Life and Analytical Sciences, Waltham, MA) containing either DMSO or rifampicin and grown for an additional 24 h. After aspirating the media, the cells were washed with phosphate-buffered saline (PBS) and fixed in 3.7% formaldehyde for 20 min, followed by four 5-min washes with PBS containing 0.1% Triton X-100 to remove the fixative and to permeabilize the fixed cells. The cells were blocked in Odyssey blocking buffer (LI-COR Biosciences) for 2 h and then incubated with primary antibody in the blocking buffer for 2 h, followed by four 5-min washes with PBS containing 0.1% Tween 20. Cy3-conjugated anti-FLAG M2 mouse monoclonal antibody (Sigma-Aldrich), rabbit polyclonal anti-FLAG pSRX (Saradhi et al., 2005), and mouse monoclonal anti-SMRT antibody (GeneTex) were used at 1:200, 1:500, and 1:100 dilutions, respectively. After incubating with appropriate secondary antibody in the blocking buffer for 1 h, the cells were washed with PBS containing 0.1% Tween 20 for a total of four washes with 5 min/wash. Cy3-conjugated goat anti-mouse IgG (Jackson ImmunoResearch Laboratories Inc., West Grove, PA) and fluorescein isothiocyanate-conjugated donkey anti-rabbit IgG (Jackson ImmunoResearch Laboratories Inc.) were used at a dilution of 1:200. Finally, the cells were incubated with Hoechst dye (AppliChem, Cheshire, CT) for 10 min to stain the nuclear DNA at a final concentration of 0.5 μg/ml in PBS, followed by three PBS washes, before acquiring the fluorescence images using an immunofluorescence microscope (Olympus America, Center Valley, PA). All of the steps were performed at room temperature while gentle shaking of the plate; during cell fixation, there was no shaking of the plate.

**Electrophoretic Mobility Shift Assay.** Double-stranded 32P-labeled oligonucleotides representing the PXR DNA binding sequence within the CYP3A4 promoter (an everted repeat with a 6-base pair spacer; hereafter referred to as CYP3A4-ER6), 5′-GAGTTAAATATGAACTCAAAGGAGGTCAGTG-3′ (bolded bases disrupt the PXR binding half-site) was incubated with in vitro-translated protein was added in each reaction. Oligonucleotide com- petition was performed using a 500-fold molar excess of unlabelled oligonucleotides. To validate the specific binding between FLAG-hPXR and hot mutant CYP3A4-ER6 5′-GATCAATATGCATCAAGAATACAGTTG-3′ (bolded bases disrupt the PXR binding half-site) was incubated with in vitro-translated and -translated protein and binding buffer. Labeled oligonucleotide (600,000 cpm) was used for each reaction together with FLAG-hPXR, FLAG-hPXR TST7, FLAG-hPXR TST7D, and human retinoid X receptor (RXR)-α proteins, which were synthesized in vitro using the TNT rabbit reticulocyte lysate system (Promega) following the manufacturer’s protocol. Equal amount of the in vitro-translated protein was added in each reaction. Oligonucleotide competition was performed using a 500-fold molar excess of unlabelled oligonucleotides. To validate the specific binding between FLAG-hPXR and hot oligonucleotide, 2 μg of either mouse monoclonal anti-FLAG M2 antibody (Sigma-Aldrich) or rabbit polyclonal anti-FLAG pSRX serum (Saradhi et al., 2005) was added to some reactions. To verify the specific binding between FLAG-hPXR and the antibodies, 2 μg of either normal mouse IgG (Santa Cruz Biotechnology, Inc., Santa Cruz, CA) or normal rabbit IgG (Santa Cruz Biotechnology, Inc.) was added to some reactions as negative controls. Binding buffer containing 10 mM Tris, pH 8.0, 40 mM KCl, 0.05% nonidet P-40, 6% glycerol, 1 mM DTT, 0.2 μg of poly(dI-dC), 10 μM zinc chloride, and 2 μl of in vitro-translated protein in a total of 20-μl reaction. Oligonucleotides and synthesized proteins were added to the inside wall of the microcentrifuge tube, mixed via vortexing, and incubated on ice for 30 min. Complexes were resolved using electrophoresis through a nondenaturing 4% polyacrylamide gel and analyzed with a Storm 860 PhosphorImager (GE Healthcare, Little Chalfont, Buckinghamshire, UK).

**Silencing of SMRT Using Small Interfering RNA.** The cells were transfected with Accell nontargeting SMARTpool control small interfering RNA (siRNA) (Thermo Fisher Scientific) or Accell SMARTpool human SMRT siRNA (Thermo Fisher Scientific) using Accell siRNA delivery media (Thermo Fisher Scientific) for 72 h according to the manufacturer’s instructions. Very briefly, when the cell growth reached approximately 75% confluence, growth medium was aspirated and replaced with Accell delivery medium containing a final concentration of 1 μM Accell siRNA. Protein expression of SMRT was analyzed from whole cell lysates collected 72 h after transfection with siRNA. For functional studies, the cells were first transfected with 100 ng of pcdNA3-FLAG-pSRX TST7D, 100 ng of pG4.4-hRluc (Promega), and 3800 ng of pGL3-CYP3A4-luc using Lipofectamine 2000 (Invitrogen) following the manufacturer’s protocol. Five hours after transfection, growth medium was aspirated and replaced with Accell delivery medium containing a final concentration of 1 μM Accell siRNA. Compounds were added 48 h after transfection with siRNA, and luciferase activity was measured 24 h after compound treatments. Renilla luciferase (pGL4-hRluc) was used as a control for transfection efficiency. Firefly luciferase activity followed by Renilla luciferase activity was measured using the Dual-Glo Luciferase Assay System (Promega). Firefly luciferase activity was first normalized with Renilla luciferase activity before normalizing the data for rifampicin with DMSO, as described below for the PXR transactivation assay.

**PXR Transactivation Assay.** The cells were transfected with pcdNA3, hPXR, p70 S6K, pG4.4-hRluc, and CYP3A4-luc plasmids using FuGENE 6 or Lipofectamine. Twenty-four hours after transfection in growth media, approximately 10,000 live cells were plated in each well of a 96-well culture plate (PerkinElmer Life and Analytical Sciences) and grown for an additional 24 h in phenol red-free DMEM (Invitrogen) supplemented with 5% charcoal/dex- tran-treated FBS (HyClone) and other additives, as described in the cell culture section. Forty-eight hours after transfection, luciferase assay was performed to determine luminescence using the Steady Lite luciferase substrate (PerkinElmer Life and Analytical Sciences) and PHERAstar microplate reader (BMG Labtech, Durham, NC). CYP3A4 promoter activity induced by hPXR after incubation with rifampicin, SR12813, or ketoconazole was shown and described as relative luminescence units (RLUs), a normalized ratio between the luminescence observed in the presence of compounds and the DMSO vehicle. In p70 S6K and hPXR cotransfection studies, Renilla luciferase (pG4.4-hRluc) was used as an internal control for transfection efficiency. Firefly luciferase activity followed by Renilla luciferase activity was measured using the Dual-Glo Luciferase Assay System (Promega). Firefly luciferase activity was normalized with Renilla luciferase to determine relative luminescence. Relative luminescence is shown as mean values with S.D. from four to six independent observations. The Student’s t test was used to determine statistical significance of four unpaired samples. Differences were considered significant for p < 0.05 (+), 0.01 < p < 0.05 (++), and 0.001 < p < 0.005 (+++).

For the rapamycin experiments, HepG2 cells were transfected with either pcdNA3 or hPXR and pGL3-CYP3A4-luc. Twenty-four hours after transfection, cells were treated with either DMSO or rapamycin, and firefly luciferase activity was measured 24 h after rapamycin treatment. Luminescence, measured from the samples transfected with hPXR, was normalized with pcdNA3-transfected samples to determine RLU. The Student’s t test was used to determine statistical significance of four unpaired samples by comparing the RLU in the presence of rapamycin to DMSO. Rapamycin treatment did not exert significant cytotoxic effect in HepG2 cells between 10 nM and 10 μM concentrations (data not shown).

**In Vitro p70 S6K Kinase Assay.** The in vitro kinase assays were performed as described previously (Lin et al., 2008). In brief, kinase assays were performed at 30°C for 30 min in 25-μl reactions with approximately 1 μg of substrate protein, 25 ng of p70 S6K (Assay Designs, Ann Arbor, MI), 0.5 μM cold ATP, and 5 μCi of [γ-32P]ATP (PerkinElmer Life and Analytical Sciences; 6000 Ci/mmol). The reactions were loaded to SDS-polyacrylamide gel electrophoresis, and the amount of the substrate samples was visualized by staining with SimplyBlue SafeStain (LCG060; Invitrogen) followed by desic- cation of the stained gel using the LABCONCO gel dryer (Labconco, Kansas City, MO). The dried gel was then subjected to overnight exposure in the Storage Phosphor Screen (GE Healthcare). Phosphoimages were obtained as described from dmd.aspetjournals.org at ASPET Journals on June 23, 2017.
using the Storm scanner (GE Healthcare). Glutathione S-transferase (GST) was expressed and purified using pGex-4T-1 (GE Healthcare) in *Escherichia coli* BL21. Purified hPXR protein was purchased from Origene Technologies, Inc. (Rockville, MD). p70 S6K positive control substrate peptide was purchased from Assay Designs.

**Mammalian Two-Hybrid Assay.** The mammalian two-hybrid system (Promega) consists of VP16-hPXR (wild-type or mutant), Gal4-SRC-1, and a luciferase reporter pG5-luc cotransfected into HepG2 cells. The Gal4 vector (pBIND) also constitutively expresses Renilla luciferase, which was used as an internal transfection control. Dual-Glo Luciferase Assay (Promega) was used to measure luciferase activity, which is an indicator of protein-protein interactions. Expression of hPXR and SRC-1 was confirmed by Western analysis (data not shown). The relative luciferase activity for pG5-luc was determined by normalizing firefly luciferase activity with Renilla luciferase activity. The values represent the means of eight independent experiments, and the bars denote the S.D. The Student’s t test was used to determine statistical significance of unpaired samples. Differences were considered significant for \( p < 0.05 \) (*), 0.01 (**), or 0.001 (***), and nonsignificant (N.S.) for \( p > 0.05 \).

**Results**

**Thr\(^{57} \) in the First Zinc Finger Motif of the DBD of hPXR Is Highly Conserved among NRs.** Sequence alignment of all the human NRs showed that there is a highly conserved threonine (Thr) or serine (Ser) in the DBD one residue N-terminal to a highly conserved cystine (Cys) and five residues C-terminal to a highly conserved histidine (His) (Fig. 1B). Of the 46 human NRs with DBD, all but nine receptors have a Thr or Ser at that position (Fig. 1B). The exceptions—retinoic acid receptor (RAR)-\( \alpha \) and RAR\( \beta \), peroxisome proliferator-activated receptor (PPAR)-\( \alpha \), PPAR\( \beta \), and PPAR\( \gamma \), human homolog of the *Drosophila* tailless gene, photoreceptor cell-specific NR, Rev-erba, and Rev-erb\( \beta \)—have an alanine (Ala) at that position (Fig. 1B). Of the 15 vertebrate species with a fully known PXR NR, Rev-erb\( \alpha \), 0.01 (wild-type) or mutant, Gal4-SRC-1, and an expression vector for hPXR were cotransfected into HepG2 cells. The Gal4 vector (pBIND) also constitutively expresses Renilla luciferase, which was used as an internal transfection control. The mammalian two-hybrid system (Promega) was used to measure luciferase activity, which is an indicator of protein-protein interactions. Expression of hPXR and SRC-1 was confirmed by Western analysis (data not shown). The relative luciferase activity for pG5-luc was determined by normalizing firefly luciferase activity with Renilla luciferase activity. The values represent the means of eight independent experiments, and the bars denote the S.D. The Student’s t test was used to determine statistical significance of unpaired samples. Differences were considered significant for \( p < 0.05 \) (*), 0.01 (**), or 0.001 (***), and nonsignificant (N.S.) for \( p > 0.05 \).

**Phosphomimetic Mutation at Thr\(^{57} \) Abolishes hPXR Transcriptional Activity.** A mutation of a Thr/Ser to a negatively charged aspartic acid (Asp) is often used to mimic phosphorylation. We tested whether such a phosphomimetic mutation at Thr\(^{57} \) affects hPXR transactivation of the *CYP3A4* promoter. The test was performed in HepG2 (Fig. 2A) and COS7 (Fig. 2B) cells transiently transfected with a vector or hPXR plasmid and CYP3A4-luc, in which the expression of luciferase is controlled using an hPXR-regulated *CYP3A4* promoter. When transfected with the vector alone, basal promoter activity was observed after treatment with the hPXR agonists rifampicin (10 \( \mu M \), 24 h) and SR12813 (10 \( \mu M \), 24 h), suggesting activation of the *CYP3A4* promoter by endogenous hPXR (Fig. 2). As expected, wild-type hPXR significantly enhanced the *CYP3A4* promoter activity in the presence of its agonists (Fig. 2). As a negative control, a nonspecific PXR inhibitor, ketocanazole, did not enhance the hPXR activity (Fig. 2). It is interesting to note that a mutation of Thr to Ala (hPXR\(^{T57A} \)), which places a hydrophobic side chain at position 57, did not affect the rifampicin- or SR12813-mediated hPXR transactivation (Fig. 2). In contrast, a mutation to Asp (hPXR\(^{T57D} \)), which mimics phosphorylation, abolished the activation of the *CYP3A4* promoter by hPXR (Fig. 2), suggesting that phosphorylation on Thr\(^{57} \) could confer a tremendous functional impact by negatively regulating hPXR activity. Multiple mechanisms might contribute to the impaired transactivation function of hPXR\(^{T57D} \), including protein stability and expression level, nuclear localization, cofactor interaction, and target promoter binding. Therefore, we sought to elucidate the responsible mechanisms. Fusion of a FLAG-tag to the N-terminal wild-type hPXR (F-hPXR) or mutant hPXR (F-hPXR\(^{T57A} \) or F-hPXR\(^{T57D} \)) did not alter the transactivation activity of the corresponding hPXR (Fig. 2). Therefore, both FLAG-tagged and untagged hPXR constructs were used interchangeably in this study.

**Mutation at Thr\(^{57} \) Does Not Affect hPXR Protein Levels.** It is possible that mutation in a protein could lead to reduced protein stability or expression level. Therefore, we tested whether the remarkable reduced activity of hPXR\(^{T57D} \) was caused by reduced mutant protein levels in HepG2 (Fig. 3A) and COS7 (Fig. 3B) cells. Western blotting results showed that there was no significant change in protein levels of hPXR after mutation to either Ala or Asp at Thr\(^{57} \) (Fig. 3). Furthermore, rifampicin treatment did not alter the levels of mutant protein (Fig. 3), suggesting that the impaired function of the phosphomimetic mutant was not caused by its reduced expression or stability. Equal amounts of actin protein expression were observed in each lane, verifying that equal amounts of total protein were loaded in each lane (Fig. 3).

**Phosphomimetic Mutation at Thr\(^{57} \) Alters hPXR Nuclear Localization Pattern.** hPXR is a transcriptional factor that induces a transcriptional response in the nucleus. Therefore, we asked whether a phosphomimetic mutation alters the localization of hPXR in HepG2 and COS7 cells and thereby contributes to impaired transcriptional activity. Immunocytochemistry experiments revealed that both hPXR\(^{T57A} \) and hPXR\(^{T57D} \), like wild-type, were localized to the nucleus in COS7 cells (Fig. 4). Similar to wild-type hPXR, hPXR\(^{T57D} \) exhibited a homogenous distribution (Fig. 4), whereas hPXR\(^{T57D} \)
nuclear localization pattern of hPXR T57D might contribute to its ability to interact with their coregulators (Rochette-Egly, 2003). In the absence of an agonist, hPXR is associated with transcriptional corepressors such as corepressor 2 (SMRT) (Kliewer et al., 1998; Ding and Staudinger, 2005a,b; Johnson et al., 2006). It is possible that the impaired function of hPXRT57D is caused by its sustained association with SMRT. Therefore, we asked whether the silencing of SMRT can rescue the function of hPXRT57D. This question was answered by knocking down the endogenous SMRT protein (scientific) as described under Materials and Methods, in HepG2 and COS7 cells transiently transfected with hPXRT57D. Western blotting results showed that SMRT siRNA, but not control nonsilencing siRNA, selectively knocked down the protein expression levels of SMRT (Fig. 5B). Equal actin expression indicates that an equal amount of total protein was loaded into each lane (Fig. 5B). However, silencing of SMRT did not significantly rescue the impaired transactivation function of the cells transfected with wild-type or mutant hPXR with the samples transfected with the vector. Likewise, statistical significance was determined from the samples transfected with FLAG-tagged wild-type or mutant hPXR to the samples transfected with the FLAG vector. Furthermore, statistical significance was ascertained similarly by comparing the RLU obtained from the samples transfected with FLAG-tagged plasmids with the samples transfected with corresponding untagged plasmids, and statistics were shown only for one comparison, i.e., between the vector and FLAG vector after rifampicin treatment. No statistical significance was observed between the samples transfected with the FLAG-tagged plasmids and the untagged plasmids (statistics were not shown for all the sample comparisons). Differences were considered significant for p < 0.05 (**), 0.01 (***), or 0.001 (****), and nonsignificant (N.S.) for p > 0.05.

**Fig. 2. Phosphomimetic mutation at Thr57 abolishes hPXR transactivation activity.** HepG2 (A) and COS7 (B) cells were co-transfected with either pcDNA3 or hPXR using CYP3A4-luc. F denotes for FLAG-tag. Twenty-four hours post-transfection, the cells were treated with vehicle DMSO (0.1%) or 10 μM compounds: rifampicin, SR12813, and ketoconazole. Luciferase activity was measured 24 h after compound treatments. CYP3A4 promoter activity induced by hPXR after treatment with the compounds was shown as RLUs. RLUs were determined by normalizing the luminescence observed in the presence of one of the compounds with the luminescence observed in the presence of DMSO. Data are shown as mean values from six independent experiments with bars indicating the S.D. The Student’s t-test was used to determine statistical significance of unpaired samples by comparing the RLU obtained from the samples transfected with wild-type or mutant hPXR with the vector. Likewise, statistical significance was determined from the samples transfected with FLAG-tagged wild-type or mutant hPXR to the samples transfected with the FLAG vector. Furthermore, statistical significance was ascertained similarly by comparing the RLU obtained from the samples transfected with FLAG-tagged plasmids with the samples transfected with corresponding untagged plasmids, and statistics were shown only for one comparison, i.e., between the vector and FLAG vector after rifampicin treatment. No statistical significance was observed between the samples transfected with the FLAG-tagged plasmids and the untagged plasmids (statistics were not shown for all the sample comparisons). Differences were considered significant for p < 0.05 (**), 0.01 (***), or 0.001 (****), and nonsignificant (N.S.) for p > 0.05.

**Silencing of SMRT Cannot Rescue the Impaired Function of hPXRT57D.** Johnson et al. (2006) showed in COS7 cells that transiently transfected PXR deletion mutants with partial or complete loss of function colocalize with SMRT to the nucleus with a punctate pattern similar to that of hPXRT57D (Johnson et al., 2006), suggesting that the altered nuclear localization pattern of hPXRT57D might contribute to its impaired function, and SMRT might be involved.

Phosphorylation of NRs often regulates their ability to interact with their coregulators (Rochette-Egly, 2003). In the absence of an agonist, hPXR is associated with transcriptional corepressors such as nuclear receptor corepressor 1 and nuclear receptor corepressor 2 (SMRT) (Kliewer et al., 1998; Ding and Staudinger, 2005a,b; Johnson et al., 2006). Activation of an NR by its agonist usually causes a release of corepressors and allows the recruitment of coactivators. Activation of hPXR by its agonist rifampicin after binding to the LBD causes an exchange of the coexpressed SMRT with the NR coactivators such as SRC-1 and SRC-3 (Kliewer et al., 1998; Ding and Staudinger, 2005a,b; Johnson et al., 2006). Agonist binding disrupts hPXR-corepressor association and induces the association between hPXR and transcriptional coactivators such as SRC-1 and SRC-3 (Kliewer et al., 1998; Ding and Staudinger, 2005a,b; Johnson et al., 2006).
Phosphomimetic Mutation at Thr57 Impairs hPXR Promoter DNA Binding. hPXR transactivation of the CYP3A4 promoter requires direct binding of hPXR to the CYP3A4 promoter. It is known that the first zinc finger motif in the DBD of hPXR is essential for DNA binding (Staudinger et al., 2001). Thr57 is located in the DBD, whereas PXR interacts with its transcriptional coregulators through its LBD. Although it is a remote possibility that a phosphomimetic mutation at Thr57 in the DBD affects the ability of hPXR to interact with its coregulators, we tested whether the loss of function of the phosphomimetic mutation was caused by a weakened or loss of interaction with its transcriptional coactivator SRC-1. Mammalian two-hybrid assay results showed that rifampicin induced the interaction between hPXR and SRC-1. However, the ability of hPXRThr57D to interact with SRC-1 was similar to that of hPXR and hPXRThr57A (Fig. 6), suggesting that impaired function of the phosphomimetic mutant was not because of its inability to interact with SRC-1. The result also suggests that rifampicin binds to hPXR, hPXRThr57D, and hPXRThr57A with similar affinity. The rifampicin-inducible interaction with SRC-1 was not observed in the absence of hPXR, as indicated by the pACT vector control.

Phosphomimetic Mutation at Thr57 Abolishes PXR Activity. The result also suggests that rifampicin binds to hPXR, hPXRThr57D, and hPXRThr57A with similar affinity. The rifampicin-inducible interaction with SRC-1 was not observed in the absence of hPXR, as indicated by the pACT vector control.

Materials and Methods. pcDNA3-FLAG-PXRT57D, pGL4-hRluc, and pGL3-CYP3A4-luc using Lipo-fectamine. Five hours after transfection with the plasmids, the cells were transfected with 1 μM Accell SMARTpool human SMRT siRNA knocks down the protein expression of SMRT. HepG2 and COS7 cells were transfected with 1 μM Accell nontargeting pool control siRNA (lane 2) or Accell SMARTpool human SMRT siRNA (lane 3) in Accell siRNA delivery media. Whole cell lysates were collected 72 h after transfection and subjected to Western blotting analysis using anti-SMRT and anti-actin antibodies. Protein expression for SMRT without transfection was shown in lane 1. Actin protein expression was shown as a loading control. Data shown are from a representative experiment. C, silencing of SMRT cannot rescue the impaired transactivation function of hPXRThr57D. HepG2 and COS7 cells were transfected with pcDNA3-FLAG-PXRT57D, pGL4-hRluc, and pGL3-CYP3A4-luc using Lipofectamine. Five hours after transfection with the plasmids, the cells were transfected with 1 μM Accell nontargeting SMARTpool control siRNA or Accell SMARTpool SMRT siRNA in Accell siRNA delivery media for 72 h. Forty-eight hours after transfection with the siRNAs, the cells were treated with vehicle DMSO (0.1%) or 10 μM rifampicin. Firefly and Renilla luciferase activities were measured 24 h after rifampicin treatment. Firefly luciferase activity was first normalized with Renilla luciferase activity before normalizing the data for rifampicin with DMSO. Data are shown as mean values from four independent experiments with bars indicating the S.D. The Student’s t test was used to determine statistical significance of unpaired samples by comparing the RLU obtained from the samples transfected with control siRNA or SMRT siRNA with the samples that were not transfected with siRNA. Differences were considered nonsignificant (N.S.) for p > 0.05.
p70 S6K Phosphorylates hPXR in Vivo. The results described in Fig. 8, A through C, suggest that hPXR might be phosphorylated by p70 S6K. In an in vitro kinase assay, we showed that reconstituted complexes of purified p70 S6K directly phosphorylate purified hPXR (Fig. 8D). Purified GST protein was used as a negative substrate control, and a p70 S6K substrate peptide was used as a positive control. Similar amounts of GST and hPXR were used in the kinase assays (Fig. 8D). These results indicate that p70 S6K directly phosphorylates hPXR in vitro and are consistent with the negative regulation of hPXR-transactivating function by p70 S6K.

Discussion

Regulated site-specific phosphorylation provides an important mechanism by which the activity of a protein is regulated (Cohen, 2000, 2001). It is well established for NRs that site-specific phosphorylation occurs on all the domains and plays a vital role to modulate or terminate the activity of NRs (Rochette-Egly, 2003; Weigel and Moore, 2007). It is known that PKA (Ding and Staudinger, 2005a; Lin et al., 2008) are involved in the regulation of PXR activity. It is also known that PKA and CDK2 can phosphorylate the PXR in vitro (Ding and Staudinger, 2005a; Lin et al., 2008). However, very little is known about specific phosphorylation of PXR. Therefore, we examined one of the putative phosphorylation sites by performing a phosphomimetic mutation at Thr<sup>57</sup>—a highly conserved Thr/Ser site in the DBD among the human NRs, including all the classic steroid receptors (Fig. 1B), and among PXR from various species (Fig. 1C)—and by testing the mutant’s transactivation function, promoter-binding activity, expression, cofactor interaction, and localization. This is the first report on the identification and characterization of a PXR phosphomimetic mutant that loses its transactivation function, due to impaired promoter-binding activity, and exhibits a punctate nuclear distribution. In addition, this is the first report to identify hPXR as an in vitro substrate for p70 S6K and to show p70 S6K regulation of hPXR transcriptional activity in HepG2 cells. Our observations suggest that Thr<sup>57</sup> might be regulated by phosphorylation, which in turn regulates hPXR target promoter binding and transactivation function. These observations might apply to other NRs due to the highly conserved nature of this residue.

The DBD is a highly conserved domain among NRs. Phosphorylation of a conserved Thr/Ser in this DBD region has been implicated in regulating NR function. Several studies have revealed that phosphorylation of a conserved Thr/Ser in the DBD can lead to attenuation and/or termination of promoter binding and transcriptional activity (Hsieh et al., 1993; Chen et al., 1999; Delmotte et al., 1999; Rochette-Egly, 2003; Sun et al., 2007). For example, phosphorylation of a highly conserved Ser<sup>78</sup> in the DBD of HNF4α has been shown to...
speculate that the phosphorylation of Thr\textsuperscript{57} in the first zinc finger motif might contribute to the regulation of DNA binding and transactivation activities of the hPXR. Phosphomimetic mutation of this site in other NRs may bring about similar functional effects as a result of the highly conserved nature of this residue.

Wild-type and both hPXR mutants (i.e., hPXR\textsuperscript{T57A} and hPXR\textsuperscript{T57D}) are localized to the nucleus in HepG2 and COS7 cells (Fig. 4), and this observation is in agreement with previously published results (Saradhi et al., 2005; Johnson et al., 2006). Punctate nuclear distribution was observed for the phosphomimetic mutant in approximately 70% of the hPXR\textsuperscript{T57D}-expressing cells. The localization phenotype of both wild-type and mutant hPXR was not affected by treatment with the hPXR agonists rifampicin and SR12813, as well as the nonspecific hPXR inhibitor, ketoconazole (data not shown). Nuclear punctate distribution for some NRs, such as the glucocorticoid receptor (Ogawa et al., 2004) and androgen receptor (Marcelli et al., 2006), has been observed in response to their ligands. It is interesting to note that the function of HNF4\(\alpha\) by altering its intracellular localization and reducing its promoter-binding activity (Sun et al., 2007). Phosphorylation of conserved Thr/Ser residues involved in the recognition of cognate response elements, which results in loss of promoter-binding activity, has also been described for the vitamin D receptor after phosphorylation of Ser\textsuperscript{51} in the DBD by PKC (Hsieh et al., 1993). Inhibition of the transcriptional activity of other NRs also occurs subsequently to phosphorylation of conserved Thr/Ser residues located within the DBD dimerization surface. For example, estrogen receptor \(\alpha\) and RAR\(\alpha\) lose their transcriptional activity after phosphorylation of Ser\textsuperscript{236} or Ser\textsuperscript{157} by PKA (Chen et al., 1999) or PKC (Delmotte et al., 1999), respectively. In agreement with these observations, our results show that the phosphomimetic mutant of hPXR loses its promoter-binding activity and provide a mechanism for the loss of transactivation function.

The fact that the first zinc finger motif in the DBD of hPXR is essential for the DNA binding (Staudinger et al., 2001) leads us to speculate that the phosphorylation of Thr\textsuperscript{57} in the first zinc finger motif might contribute to the regulation of DNA binding and transactivation activities of the hPXR. Phosphomimetic mutation of this site in other NRs may bring about similar functional effects as a result of the highly conserved nature of this residue.

Wild-type and both hPXR mutants (i.e., hPXR\textsuperscript{T57A} and hPXR\textsuperscript{T57D}) are localized to the nucleus in HepG2 and COS7 cells (Fig. 4), and this observation is in agreement with previously published results (Saradhi et al., 2005; Johnson et al., 2006). Punctate nuclear distribution was observed for the phosphomimetic mutant in approximately 70% of the hPXR\textsuperscript{T57D}-expressing cells. The localization phenotype of both wild-type and mutant hPXR was not affected by treatment with the hPXR agonists rifampicin and SR12813, as well as the nonspecific hPXR inhibitor, ketoconazole (data not shown). Nuclear punctate distribution for some NRs, such as the glucocorticoid receptor (Ogawa et al., 2004) and androgen receptor (Marcelli et al., 2006), has been observed in response to their ligands. It is interesting to note
that the nuclear punctate pattern has also been observed for hPXRT57D mutants with impaired function (Johnson et al., 2006), and these mutants have been shown to colocalize with the corepressor SMRT to the nucleus with a punctate pattern. This suggests that mutant hPXRT57D may constitutively associate with SMRT and that altered nuclear localization pattern might contribute to impaired function. Similar results were observed in our study for the phosphomimetic mutant in terms of impaired function and nuclear colocalization with punctate distribution. However, SMRT knockdown did not reverse the functional phenotype of the phosphomimetic mutant, indicating that SMRT is not responsible for the impaired function, although SMRT colocalizes with the phosphomimetic mutant at discrete nuclear foci. At this juncture, we do not have any data to offer a logical speculation of the molecular mechanism(s) responsible for the altered nuclear localization of the phosphomimetic mutant.

The fact that the agonist-mediated transactivation of the endogenous hPXRT57D (i.e., the relative luminescence observed when the cells were transfected with an empty vector) was significantly lower when HepG2 cells were transfected with hPXRT57D (Fig. 2) suggests that hPXRT57D is a properly folded protein and acted as a dominant-negative mutant by competing with one or more of the components involved in the PXR transactivation pathway. The interaction between hPXRT57D and the coactivator SRC-1 is regulated by agonist binding and is critical for hPXRT57D transactivation function. In the mammalian two-hybrid system, hPXRT57D interacts, similarly to hPXRT57A, with SRC-1 in a rifampicin-inducible manner (Fig. 6), showing that the phosphomimetic mutant retains its ability to bind to rifampicin and interact with SRC-1. These data further support the notion that the loss-of-function phosphomimetic mutant is an appropriately folded protein and can act as a dominant-negative protein by competing with wild-type hPXRT57D for interaction with the coactivator SRC-1, which is a crucial protein-protein interaction step in the process of hPXRT57D transactivation pathway. This could partially, if not fully, account for the reduced endogenous hPXRT57D transcriptional activity in HepG2 cells when transfected with hPXRT57D. When a protein is misfolded, it is usually bound to chaperones such as heat shock proteins and subsequently undergoes degradation by proteasomes, and these events occur predominantly in the cytoplasm. The fact that hPXRT57D, when folded, is translocated to the nucleus and stably expressed similarly to the wild-type and nonphosphomimetic mu-
tant hPXR, argues against the idea that hPXR T57S displays the punctate nuclear pattern because of its association with proteasomes or heat shock proteins.

It has been reported that kinases could phosphorylate nonconsensus Thr/Ser sites depending on the cellular context (Lehtinen et al., 2006). Thr57 in hPXR, along with its flanking residues, does not fulfill the known consensus sequence for known kinases. However, our bioinformatics search results predicted that Thr57 is a putative phosphorylation site for S6K and/or protein kinase B (Akt). P38K-Akt pathway, a major signaling pathway activated by insulin and other growth factors, is known to be involved in negatively regulating the transcriptional activity of the PXR by affecting the interaction between PXR and its coactivator forkhead in rhabdomyosarcoma (FKHR) (Kodama et al., 2004). Akt possibly accomplishes this negative regulation by phosphorylating and translocating the nuclear FKHR into the cytoplasm for proteosomal degradation (Tang et al., 1999), consequently minimizing the levels of nuclear FKHR available for interacting with and activating PXR. However, it is not known whether Akt phosphorylates PXR and regulates its activity independently of FKHR.

In this report, we show that p70 S6K, a downstream kinase in the P38K-Akt pathway, phosphorylates and negatively regulates the transcriptional activity of hPXR (Fig. 8). The inhibition of hPXR by p70 S6K was not caused by reduced protein levels of hPXR. In fact, the kinase cotransfection resulted in elevated hPXR protein level (Fig. 8). This particular result might be because of enhanced stability of hPXR by unknown mechanisms. Alamine mutation at Thr57 confers partial but significant resistance to p70 S6K attenuation, hinting that p70 S6K targets phosphorylation and regulation of hPXR, and supports the notion that hPXR may be phosphorylated at multiple sites, resulting in enhanced stability of hPXR.


Address correspondence to: Taosheng Chen, Department of Chemical Biology and Therapeutics, St. Jude Children’s Research Hospital, 262 Danny Thomas Place, Mail Stop 1000, Memphis, TN 38105. E-mail: taosheng.chen@stjude.org