CYP26C1 Is a Hydroxylase of Multiple Active Retinoids and Interacts with Cellular Retinoic Acid Binding Proteins

Guo Zhong, David Ortiz, Alex Zelter, Abhinav Nath, and Nina Isoherranen

Departments of Pharmaceutics (G.Z., N.I.) and Medicinal Chemistry (D.O., A.N.), School of Pharmacy, and Department of Biochemistry, School of Medicine (A.Z.), University of Washington, Seattle, Washington

Received October 31, 2017; accepted February 22, 2018

ABSTRACT

The clearance of retinoic acid (RA) and its metabolites is believed to be regulated by the CYP26 enzymes, but the specific roles of CYP26A1, CYP26B1, and CYP26C1 in clearing active vitamin A metabolites have not been defined. The goal of this study was to establish the substrate specificity of CYP26C1, and determine whether CYP26C1 interacts with cellular retinoid acid binding proteins (CRABPs). CYP26C1 was found to effectively metabolize all-trans retinoic acid (atRA), 9-cis-retinoic acid (9-cis-RA), 13-cis-retinoic acid, and 4-oxo-atRA with the highest intrinsic clearance toward 9-cis-RA. In comparison with CYP26A1 and CYP26B1, CYP26C1 resulted in a different metabolite profile for retinoids, suggesting differences in the active-site structure of CYP26C1 compared with other CYP26s. Homology modeling of CYP26C1 suggested that this is attributable to the distinct binding orientation of retinoids within the CYP26C1 active site. In comparison with other CYP26 family members, CYP26C1 was up to 10-fold more efficient in clearing 4-oxo-atRA (intrinsic clearance 153 μl/min/pmol) than CYP26A1 and CYP26B1, suggesting that CYP26C1 may be important in clearing this active retinoid. In support of this, CRABPs delivered 4-oxo-atRA and atRA for metabolism by CYP26C1. Despite the tight binding of 4-oxo-atRA and atRA with CRABPs, the apparent Michaelis-Menten constant in biological matrix (Km) value of these substrates with CYP26C1 was not increased when the substrates were bound with CRABPs, in contrast to what is predicted by free drug hypothesis. Together, these findings suggest that CYP26C1 is a 4-oxo-atRA hydroxylase and may be important in regulating the concentrations of this active retinoid in human tissues.

Introduction

Vitamin A (retinol; ROL), through its active metabolite retinoic acid (RA), plays a critical role in regulation of gene transcription and cell proliferation, differentiation, and apoptosis (Noy, 2010; Tang and Gudas, 2011). Retinol is believed to be devoid of biologic activity and is metabolized by alcohol and aldehyde dehydrogenases to retinoic acid (RA) (Fig. 1; Napoli, 2012; Kedishvili, 2016). The endogenous vitamin A metabolites detected in human serum include all-trans (atRA), 9-cis-, 13-cis-, and 9,13-di-cis-RA, 4-oxo-13-cis-RA, and 4-oxo-atRA (Figs. 1 and 2; Arnold et al., 2012). Of these retinoids, atRA activates retinoid acid receptors (RARs) and is believed to be the most important and biologically active endogenous RA isomer (Allenby et al., 1994; Chambon, 1996; Stevison et al., 2015). However, 13-cis-RA, 9-cis-RA, and 4-oxo-atRA can also activate RARs (Idres et al., 2002; Topletz et al., 2015). Owing to this pharmacological activity and their favorable in vivo pharmacokinetics, 13-cis-RA and 9-cis-RA are used to treat acne, high risk neuroblastoma (Veal et al., 2007), and chronic hand eczema (Schmitt-Hoffmann et al., 2011). Likewise, both 13-cis-RA and 9-cis-RA are teratogenic, demonstrating classic retinoid effects (Willhite et al., 1986; Kraft and Juchau, 1993). 9-cis-RA is also a ligand of retinoid X receptors (RXRs) with significantly higher binding affinity than atRA and 13-cis-RA (Aström et al., 1990; Heyman et al., 1992; Allenby et al., 1993), but the in vivo significance of 9-cis-RA is unclear. 4-oxo-atRA exhibits higher affinity to RARα and similar affinity to RARβ as does atRA (Pijnappel et al., 1993; Idres et al., 2002; Topletz et al., 2015), consistent with its teratogenicity (Herrmann, 1995) and ability to modulate positional specification in early embryo (Pijnappel et al., 1993). However, the role of the different RA isomers and 4-oxo-atRA in human physiology is unclear.

Biologic effects of RA isomers and metabolites are dependent on the RAR binding affinity and the cellular concentrations of the retinoids. Therefore, strict regulation of the physiologic concentrations of endogenous retinoids via tissue-specific expression of retinoid-synthesizing and -metabolizing enzymes and retinoid binding proteins (Fig. 1) is critical. Two enzymes of the CYP26 family, CYP26A1 and CYP26B1, have been identified as key enzymes responsible for clearing atRA and controlling atRA concentrations (Fig. 1). CYP26A1 appears to be the liver atRA hydroxylase

This research was supported by grants from National Institutes of Health [Grants R01 GM111772 and R01 GM081569].

This article has supplemental material available at molpharm.aspetjournals.org.

ABBREVIATIONS: ACN, acetonitrile; ANOVA, analysis of variance; atRA, all-trans retinoic acid; CRABP, cellular retinoic acid binding protein; EPI, enhanced product ion; FA, formic acid; HPLC, high-performance liquid chromatography; IPTG, isopropyl β-D-1-thiogalactopyranoside; IS, internal standard; Km, apparent Michaelis-Menten constant in biological matrix; LC-MS/MS, liquid chromatography–tandem mass spectrometry; LC-UV, liquid chromatography UV; MD, molecular dynamics; MRM, multiple reaction monitoring; P450, cytochrome P450; RAR, retinoic acid receptor; ROL, retinol.
eliminating biologically active retinoids, whereas CYP26B1 appears to be responsible for RA clearance in extrahepatic tissues (Thatcher et al., 2010; Topletz et al., 2015). CYP26A1 and CYP26B1 have also been shown to metabolize 4-oxo-RA and 9-cis-RA, but the catalytic efficiency is much lower than that toward RA (Thatcher et al., 2011; Topletz et al., 2012; Diaz et al., 2016), suggesting that other enzymes may be responsible for the elimination of 4-oxo-RA and RA isomers in humans. The sequence similarity of human CYP26C1 is only 43% with CYP26A1 and 51% with CYP26B1 (Taimi et al., 2004), suggesting structural differences between these cytochrome P450s (P450s). Hence, CYP26C1 may have different substrate specificity than CYP26A1 and CYP26B1. Indeed, in CYP26C1-transfected COS-1 cells, both 9-cis-RA and atRA were identified as potential substrates of CYP26C1 (Taimi et al., 2004). However, the role of CYP26C1 in retinoid clearance or in formation of active metabolites such as 4-oxo-atRA has not been defined, and biochemical characterization of CYP26C1 in retinoid metabolism is lacking.

A distinct characteristic of biologically important enzymes contributing to retinoid metabolism is that they interact with and obtain their substrates from cellular retinoid binding proteins (CRABPs), which appear to deliver atRA either to nuclear RARs to regulate gene transcription or to CYP26A1 and CYP26B1 for catabolism (Napoli, 2017). Hence, whether metabolic enzymes interact with CRABPs can define the biologic importance of the enzyme and the substrates. For example, CRABP1-bound atRA was metabolized in rat testes microsomes at a rate similar to free atRA, and CRABP1-binding of 4-oxo-atRA abolished 4-oxo-atRA metabolism, suggesting substrate-specific delivery to metabolism (Fiorella and Napoli, 1991, 1994). Recently, atRA was shown to be channeled to recombinant CYP26B1 by CRABPs (Nelson et al., 2016), but other substrates that bind to CRABPs were not studied, and the specificity of CRABP interactions with CYP26 enzymes was not established. To better understand the role of CYP26 enzymes in retinoid metabolism, the CRABP-specific deliveries of substrates to metabolism by P450s need to be defined. The goal of this study was to establish the ligand specificity of CYP26C1 and characterize the CRABP-CYP26C1 interactions to delineate the biologic significance and the role of CYP26C1 in regulating retinoid homeostasis.

Materials and Methods

Chemicals and Reagents. atRA, NADPH, and ketoconazole were purchased from MilliporeSigma (St. Louis, MO). Talarozole was purchased from MedChem Express (Princeton, NJ). 4(S)-OH-atRA and 4(R)-OH-atRA were synthesized as described previously (Shimshoni et al., 2012). All other retinoids were from Toronto Research Chemicals (North York, ON, Canada). All solvents were high-performance liquid chromatography (HPLC) or Optima grade and were purchased from Thermo Fisher Scientific (Waltham, MA).

Cloning, Expression, and Characterization of Recombinant CYP26C1. Human CYP26C1 cDNA was purchased from OriGene Technologies (Rockville, MD) (SKU: SC307567). Primers were designed to amplify the CYP26C1 coding sequence minus the stop
experiments repeated three times. NADPH. Incubations were conducted in duplicate or triplicate and all UV-Vis detection. All product formation was confirmed to be NADPH-an internal standard (IS) for all the incubations except those using experiment below. 4-Oxo-
reductase concentration used for incubations are specified for each 5 minutes potassium phosphate buffer in a total volume of 1 ml. After another to incorporate into the membranes at room temperature for 5 minutes The rat reductase was added into CYP26C1 microsomes and allowed incubations and retinoid extractions were performed following pub-
lar antibodies, respectively, to detect the His-tagged CYP26C1. Valencia, CA) and IRDye 680RD anti-mouse antibody (1: 10000; Carlsbad, CA) supplemented with 2.5% fetal bovine serum was used during protein expression, media was supplemented with 80°C. The microsomal fractions of insect cells were prepared by ultracentrifugation on the basis of published methods (Topletz et al., 2012). The content of active P450 was measured by a CO-difference spectrum (Omura and Sato, 1964), using an Amino DW-2 dual-beam spectrophotometer (Olis Inc, Bogart, GA). Supersomes containing human P450 oxidoreductase and b5 prepared from insect cells (Corning Inc., Corning, NY) were used as negative control for CO-difference spectra. Protein concentra-
tion of the CYP26C1 microsomes was measured using BCA protein assay kit according to manufacturer's recommendations (Thermo Fisher Scientific). The CYP26C1 microsomes were used in all catalytic experiments. Rat P450 reductase was expressed in Escherichia coli and purified as described previously (Woods et al., 2011). Qualitative Western blotting was performed on the basis of a published method for CYP26A1 with minor modifications (Lutz et al., 2009). CYP26C1 microsomes containing 2 μg total protein were loaded on the gel. Mouse anti-6His antibody (1:2000; Qiagen, Valencia, CA) and IRDye 680RD anti-mouse antibody (1: 10000; LI-COR Biosciences, Lincoln, NE) were used as primary and secondary antibodies, respectively, to detect the His-tagged CYP26C1.

**General Protocol for Incubations.** Unless otherwise described, incubations and retinoid extractions were performed following published procedures used for analysis of CYP26A1 and CYP26B1 activity (Thatcher et al., 2010; Topletz et al., 2012). Similar incubation conditions and reductase to P450 ratios were used for consistency. The rat reductase was added into CYP26C1 microsomes and allowed to incorporate into the membranes at room temperature for 5 minutes (Thatcher et al., 2010; Topletz et al., 2012), before membranes were added to the incubation mixture containing substrate and 100 mM potassium phosphate buffer in a total volume of 1 ml. After another 5 minutes' preincubation at 37°C with the substrate, the reaction was initiated with the addition of 1 mM NADPH. Incubations were quenched and extracted with 3 ml of ethyl acetate, as previously described (Topletz et al., 2012). The ethyl acetate layer was collected, evaporated to dryness under N₂ flow, and reconstituted with 100 μl of methanol. Incubation time, micromolar CYP26C1 content and rat reductase concentration used for incubations are specified for each experiment below. 4-Oxo-arRa-d₅ (2 μl of 5 μM solution) was added as an internal standard (IS) for all the incubations except those using UV-Vis detection. All product formation was confirmed to be NADPH-dependent by comparison to identical incubations in the absence of NADPH. Incubations were conducted in duplicate or triplicate and all experiments repeated three times.

**Evaluation of Potential CYP26C1 Substrates.** To identify substrates of CYP26C1, individual retinoids at 1 μM concentration were incubated with 18 nM microsomal CYP26C1 and 36 nM rat reductase for 30 minutes. Tested retinoids included RA isomers (aRA, 13-cis-RA, and 9-cis-RA), 4-oxo-RA isomers (4-oxo-aRA, 4-oxo-13-cis-RA, and 4-oxo-9-cis-RA), 4-0H-RA isomers (4-0H-aRA, 4-0H-13-cis-RA, and 4-0H-9-cis-RA), ROL isomers (aROL, 13-cis-ROL, and 9-cis-ROL), and 3-dehydro-aROL. For incubations with RA isomers, 4-oxo-RA isomers, or 4-0H-RA isomers, samples were extracted and analyzed by liquid chromatography (LC-UV) as described previously (Thatcher et al., 2010) using an Agilent 1200 series HPLC system (Agilent Technologies, Santa Clara, CA) equipped with a Zorbax C18 column (3.5 μm, 2.1 x 100 mm; Agilent Technologies) and by using water instead of 50 mM ammonium acetate (pH 4.5) in the mobile phase. The column temperature was maintained at room temperature. Analytes were monitored at UV wavelength of 360 nm. For incubations with ROL isomers and 3-dehydro-aROL, hexanes (4 ml) were used to quench the reaction and extract analytes. After the hexane layer was collected and evaporated, samples were reconsti-
tuted with 100 μl acetonitrile (ACN). Retinyl acetate (5 μl of 5 μM solution) was added to the samples as IS. Samples were analyzed by LC-UV using the Agilent 1200 series HPLC system coupled with an Ascentis RP-amide column (2.7 μm, 15 cm × 2.1 mm; MilliporeSigma). Column temperature was maintained at 40°C. The flow rate was 0.5 ml/min. The gradient elution started with initial condition of 40% aqueous with 0.1% formic acid (FA) and 60% acetonitrile with 0.1% FA, which was maintained for 2 minutes, followed by a gradient to 100% ACN with 0.1% FA over 11 minutes, and held at that for 8 minutes before being returned to the initial conditions before next injection. Analytes were monitored at UV wavelength of 325 nm.

**Steroselective Formation of 4-0H-RA Enantiomers from RA Isomers by CYP26C1.** The stereoselectivity in the formation of 4-0H-RA by CYP26C1 was determined for individual RA isomers by incubating aRA, 13-cis-RA, or 9-cis-RA at 0.75 μM with 15 nM CYP26C1 and 30 nM reductase for 15 minutes. For comparison, CYP26A1 was incubated under identical conditions with the same substrates. Samples were extracted and analyzed by LC-UV using an Agilent 1200 series HPLC system coupled with an Chiralcel OD-RH column (2.1 mm; Chiral Technologies Inc., West Chester, PA) as described previously to detect 4(S)- and 4(R)-OH-RA formation (Fig. 2; Shimshoni et al., 2012). The peak area ratio of 4(S)- and 4(R)-OH-RA was calculated and used as an indicator of the stereoselectivity of 4-0H-RA formation.

**Identification of Metabolites Formed by CYP26C1.** To iden-
tify metabolites formed by CYP26C1, the incubations described above for substrate identification were further analyzed by liquid chromatography–tandem mass spectrometry (LC-MS/MS) using an Sciex API5500 Q/LIT mass spectrometer (Sciex, Concord, On,
Canada) equipped with Agilent 1290 Infinity UHPLC (Agilent Technologies). All analytes were monitored with electrospray ionization under negative ion mode. To evaluate the identity of metabolites of RA isomers, samples were first separated using a Zorbax C18 column (3.5 μm, 2.1 x 100 mm) and a mobile phase flow rate of 0.35 ml/min, with a linear gradient from 90% aqueous with 0.1% FA and 10% ACN to 5% aqueous with 0.1% FA and 95% ACN over 10 minutes. The column temperature was maintained at 40°C. The multiple-reaction monitoring (MRM) transitions monitored included m/z 315 > 253 (4-OH- and 18-OH-
RA), m/z 315 > 241 (16-OH-RA), and m/z 313 > 269 (4-oxo-RA), as described previously (Topletz et al., 2012). All MS parameters set for MRM scans are listed in Supplemental Table 1. Together with monitoring MRM transitions, parallel information of [M-H] ions, including m/z 315 and m/z 313 Da, were acquired via enhanced product ion (EPI) scan. The MS/MS spectra were collected from m/z 50–330 Da. EPI scan rate was set at 10,000 Da/s, linear ion trap fill time at 100 milliseconds, Q₁ at low resolution and Q₃ entry barrier at 8 V. Q₃ trap was turned on. Other mass spectrometer detection settings were the same as those used in MRM scan, except that declustering potential (DP) of –80 V and collision energy (CE) of –35 V with a spread of ±10 V for both [M-H] ions was used. For further separation of the metabolites, the MS/MS spectra were also collected after chiral separation of the products using the Chiralcel OD-RH column and chromatography as described above. Mass spectrometry settings for MRM and EPI scans were as described above. To determine the identity of 4-oxo-arRa metabolites, a Zorbax C18 column was used for analyte separation, and the incubations were analyzed by LC-MS/MS. The flow rate was 0.2 ml/min, with a linear gradient from 90% aqueous with 0.1% FA and 10% ACN to 10% aqueous with 0.1% FA and 90% ACN over 15 minutes, held for 8 minutes before being returned to initial conditions. The MRM transitions monitored included m/z 329 > 255 (4-oxo-16-OH-ORa)
and m/z 329 > 267 Da (4-oxo-18-OH-αRA), as described previously (Topletz et al., 2012). Parameters set for MRM scans are provided in Supplemental Table 1. The EPI scan with the spectrum range of 50–350 Da was acquired for the [M-H] ion of m/z 329 Da. EPI settings were similar to those described for metabolites of RA isomers except that the scan rate was 1000 Da/5 s, and Q1 was set at unit resolution.

**Enzyme Kinetics of RA Isomers and 4-oxo-αtRA with CYP26C1.** To determine enzyme kinetic parameters for RA isomers, varying concentrations of αtRA (10–250 nM), 13-cis-RA (5–250 nM), or 9-cis-RA (3–100 nM) were incubated with 2 nM CYP26C1 and 4 nM reductase for 30 seconds. Standard curves of RA isomers, 4-OH-RA isomers, and 4-oxo-RA isomers were constructed for quantification. Samples were extracted and analyzed by LC-MS/MS using Sciex QTAPR 4500 mass spectrometer (Sciex) coupled with Shimadzu UFLC XR DGU-20A5 (Shimadzu Corporation, Kyoto, Japan) and a Zorbax C18 column, as described above for metabolite identification. 4-OH-, 18-OH- and 16-OH-RA isomers, 4-oxo-RA isomers, 4-oxo-αtRA-d₃, and RA isomers were monitored by MRM, and the MRM transitions and MS parameters used are listed in Supplemental Table 2. On the basis of the peak area ratio of metabolites to the IS, the formation of major metabolites of RA isomers was quantified using Analyst software (Sciex). Owing to the sequential metabolism of 4-OH-RA to 4-oxo-RA, to quantify the 4-OH-RA formation from αtRA and 13-cis-RA, the formation of 4-OH-RA and 4-oxo-RA were summed, as previously described for CYP26A1 and CYP26B1 (Topletz et al., 2015) (Supplemental Fig. 1A). For 9-cis-RA, only the formation of 4-oxo-RA was determined because the formation of 4-oxo-9-cis-RA was quantitatively insignificant. The velocity of metabolite formation was plotted as a function of RA concentration, and substrate concentration was corrected for depletion by calculating the average of the initial added concentration and final measured concentration, as described previously (Lutz et al., 2009). Enzyme kinetic parameters, including kcat and apparent Michaelis-Menten constant in biological matrix (Km), were obtained from GraphPad Prism 5.0 (GraphPad Software, La Jolla, CA) by fitting the Michaelis-Menten equation to the data. Intrinsic clearance (Clint) was calculated as kcat/Km.

To obtain enzyme kinetic parameters for 4-oxo-αtRA, both product formation and substrate depletion of 4-oxo-αtRA by CYP26C1 were measured. For product formation, varying concentrations of 4-oxo-αtRA (3–100 nM) were incubated with 2 nM CYP26C1 and 4 nM reductase for 2 minutes. Samples were extracted and analyzed by LC-MS/MS using Sciex 4500 QTRAP mass spectrometer and the Zorbax C18 column with the same LC conditions as described above. Monitored analytes included 4-oxo-16-OH-RA, 4-oxo-αtRA, and 4-oxo-αtRA-d₃. MRM transitions and MS parameters are listed in Supplemental Table 2. Owing to the tight binding of 4-oxo-αtRA to CYP26C1, enzyme kinetic parameters, including kcat and km, were obtained from GraphPad Prism 5.0 by fitting the Morrison equation to the data, as described previously (Shimshoni et al., 2012), and assuming a CYP26C1 concentration of 2 nM in the incubations. Because of the lack of synthetic 4-oxo-16-OH-αtRA standard, the product formation was estimated as the ratio of peak area of 4-oxo-16-OH-αtRA to IS to allow km determination. For substrate depletion, experiments were conducted following published procedures (Shimshoni et al., 2012) with varying concentrations of 4-oxo-αtRA (5–150 nM) incubated with 2 nM CYP26C1 and 4 nM reductase in a total volume of 3 ml. At 0.5, 1, 2, and 4 minutes, 0.5-ml aliquots were taken and quenched with 3 ml of ethyl acetate. 4-oxo-αtRA-d₃ was added as an IS and 4-oxo-αtRA quantified as described for product formation assays. km and Clint were obtained using GraphPad Prism 5.0, as described previously (Shimshoni et al., 2012).

Differences in the kinetic parameters between substrates were evaluated using Graphpad Prism 5.0. The differences for each kinetic parameter (km, kcat, and Clint) were tested via analysis of variance (ANOVA), including all different substrates in the analysis, and Tukey’s test was used as a post-hoc test to identify differences between substrates. Owing to the multiple ANOVA comparisons done (comparison of km, kcat, and Clint values), a Bonferroni adjustment for the P value was done (0.05/3) and a P value less than 0.01 was considered significant.

**Expression and Purification of CRABPs.** CRABP expression vectors were a gift from Dr. Noa Noy (Case Western Reserve University). His-tagged CRABP-I and CRABP-II vectors were used to transform E. coli. Transformed cells were plated into LB agar plates containing 30 µg/ml of kanamycin. Overnight cultures were prepared from freshly streaked plates. LB media (500 ml) with kanamycin added was inoculated with overnight cultures and incubated at 37°C with shaking until optical density at the wavelength of 600 nm reached 0.6. IPTG was added at a final 1 mM concentration and the culture was incubated at 37°C for another 2 hours. Induced cells were collected by centrifugation at 6000g for 20 minutes. Cell pellets were stored at −80°C until protein purification. For purification of CRABPs, frozen cell pellets were thawed on ice, resuspended with 20 ml of lysis buffer (1 mg/ml of lysosome, 500 mM NaCl, 20 mM Tris, 5 mM imidazole, pH 7.4), and incubated at room temperature for 20 minutes with gentle shaking. Resuspended mixture was placed on ice and sonicated for 30 seconds twice with a 30-second interval followed by centrifugation at 12,000g for 30 minutes. Supernatant was applied to a HisTrap HP column (GE Healthcare Bio-Sciences, Marlborough, MA) using Biologic DualFlow chromatography system (Bio-Rad Laboratories, Hercules, CA) following manufacturer’s instructions. Loaded samples were first washed with 5× column volume of 20 mM Tris-HCl, 500 mM NaCl, 20 mM imidazole (pH 7.4), and then CRABPs were eluted with 5× column volume of 20 mM Tris-HCl, 500 mM NaCl, 300 mM imidazole (pH 7.4). Fractions containing CRABPs were concentrated and exchanged into HEDK buffer (50 mM Hepes, 0.1 mM EDTA, 0.5 mM DTT, 100 mM KCl, pH 8.0) using Amicon 10-kD molecular-weight cutoff filters (MilliporeSigma, Billerica, MA). Purified CRABPs were stored at −20°C in HEDK buffer with 50% glycerol.

**Effect of CRABPs on Retinoid Metabolism by CYP26C1.** To investigate how CRABPs affect αtRA, 9-cis-RA, and 4-oxo-αtRA metabolism by CYP26C1, three experiments were conducted as previously described with minor modifications (Nelson et al., 2016). For all the CRABP-related incubations, microsomal CYP26C1 (2 nM) and rat reductase (4 nM) were first preincubated with 1 mM NADPH in a total volume of 1 ml at 37°C for 3 minutes. The reaction was initiated by adding free substrate (αtRA, 4-oxo-αtRA, or 9-cis-RA) or substrates premixed with CRABP-I or CRABP-II in 1:1 concentration ratio. One-to-one binding of substrates (αtRA or 4-oxo-αtRA) with CRABPs was confirmed with fluorescence titration and via isolation of the holo-CRABP from CRABP-ligand mixture using the method adapted from Fiorella et al. (Fiorella and Napoli, 1991; Fiorella et al., 1993) in which the ligand is bound with CRABP and then holo-CRABP is separated from any free ligand using a mini-desalting column (Thermo Fisher Scientific). After the incubation time specified for each experiment, the reaction was quenched with ACN (1 ml) followed by the addition of ethyl acetate (3 ml) and 4-oxo-αtRA-d₃ (2 µl of 5 µM solution), as previously described (Nelson et al., 2016). The rest of the procedures were as described above and metabolite formation was measured as described above using LC-MS/MS. To characterize the effect of CRABPs on CYP26C1 activity, first, CYP26C1 was incubated with 100 nM free substrate or 100 nM CRABP-ligand mixture (1:1) for 2 minutes. The formation rates of 4-OH-αtRA, 4-OH-9-cis-RA, and 4-oxo-16-OH-αtRA were compared in the absence and presence of CRABPs. Second, the kinetics of 4-OH-αtRA and 4-oxo-16-OH-αtRA formation as metabolites of αtRA and 4-oxo-αtRA were characterized, respectively. For this, varying concentrations of CRABP-I and CRABP-II-bound αtRA or 4-oxo-αtRA (3–200 nM; 1:1 CRABP-ligand ratio) were incubated with CYP26C1 for 1 (αtRA) or 2 minutes (4-oxo-αtRA) and extracted as described above. The km and kcat of αtRA and 4-oxo-αtRA bound to CRABP were obtained by fitting the Michaelis-Menten equation and the Morrison equation to the data, respectively. Third, the effect of excess apo-CRABPs on CYP26C1 activity was evaluated as described previously for CRABPs and CYP26B1 (Nelson et al., 2016). Individual substrate at 100 nM was premixed with 0–400 nM CRABP-I or CRABP-II and incubated with CYP26C1 in the
presence of NADPH for 2 minutes. Metabolite formation rates were measured and compared with the control, which contained no CRABP in the incubation. Differences in kinetic values were compared together with all substrates by ANOVA followed by Tukey’s test as a post-hoc test as described above for enzyme kinetic analysis.

**Ligand-Induced Binding Spectra and Determination of IC$_{50}$ Values of Talarozole and Ketoconazole with CYP26C1.** To explore the active-site characteristics of CYP26C1 and test if talarozole and ketoconazole bind to the active site of CYP26C1 and interact with the heme, CYP26C1 microsomes at 0.7 μM in 100 mM potassium phosphate buffer (pH 7.4) were added into the sample and reference cuvettes, with 0.45 ml in each. Microsomal CYP26C1 was titrated with ketoconazole (100, 149, and 198 μM) or talarozole (20, 79, and 157 μM) in methanol in the sample cuvette, and an equal volume of methanol was added into the reference cuvette. The binding spectra were obtained using an Aminco DW2 dual-beam spectrophotometer (Olis Inc.). The spectra were normalized to 415 or 420 nm for comparison.

To determine if talarozole and ketoconazole are also inhibitors of CYP26C1 and to obtain IC$_{50}$ values, talarozole (0.01–25 μM dissolved in dimethyl sulfoxide) and ketoconazole (25–200 μM dissolved in methanol) were incubated with 2 nM CYP26C1, 4 nM reductase, and 50 nM arRa for 2 minutes. Because the $K_a$ values for 9-cis-RA were much lower than arRa with CYP26C1, instead of using 9-cis-RA as a substrate in the inhibition assays, as previously described (Thatcher et al., 2011), assays were performed using arRa as a substrate at 50 nM, a concentration similar to the $K_a$ value of arRa. Control incubation contained a volume of dimethyl sulfoxide or methanol equal to those used with talarozole or ketoconazole. Samples were analyzed by LC-MS/MS as described above for enzyme kinetic experiments. The peak area ratio of 4-OH-Ra to IS was used as a measure of CYP26C1 activity. The percent remaining activity compared with control was calculated and plotted as a function of inhibitor concentration. The IC$_{50}$ values were determined by nonlinear regression using GraphPad Prism 5.0, as described previously (Thatcher et al., 2011).

**Generation of a CYP26C1 Homology Model and Docking of Retinoids.** To explore the ligand binding interactions of CYP26C1, a CYP26C1 homology model was created with I-TASSER (Zhang Lab, University of Michigan, Ann Arbor, MI; Roy et al., 2010), using the sequence for human CYP26C1 (NCBI sequence number NP_899230.2). The following structures were used as templates: the retinoic acid-binding cyano bacterial CYP120A1 (PDB code 2VE3; Kühnel et al., 2008); the lanosterol- (PDB code 4LXJ), itraconazole- (PDB code 5EQB; Monk et al., 2014), and fluconazole- (PDB code 4WMB; Sagatova et al., 2015) bound structures of a CYP51 from Saccharomyces cerevisiae; and the thioperamide-bound human CYP46A1 (PDB code 3MDM; Mast et al., 2010). The heme group was positioned in I-TASSER models according to its position in structural analogs. To model a structure in solution capable of binding CYP26 substrates, molecular dynamics (MD) simulations were performed with GROMACS version 4.6.5 (Van Der Spoel et al., 2005). The topology files were modified to reflect the cysteine thiolate linkage to the heme iron, as previously described (Autenrieth et al., 2004; Oda et al., 2005). Atomic interactions were defined with the GROMOS 54a7 force field. The model was placed in a dodecahedron periodic box of diameter 3.75 nm and then energy was minimized by the steepest-descent and conjugate-gradient methods. The system was solvated with ~89,000 simple point-charge water molecules, and Na$^+$ and Cl$^-$ ions were added to neutralize the system with an effective 0.15 M ionic strength. Molecular dynamics simulations reached equilibrium in ~7 nanoseconds and were allowed to proceed for 40 nanoseconds. Root-mean-square deviation and active-site volume during the MD simulation are shown in Supplementary Fig. 2. To select a conformation suitable for docking studies, the active-site volume throughout the MD simulation was measured with Pocke tVolume M Easurer 2.0 (POVME; Durrant et al., 2014). Candidate models were selected from different time points representing conformations with maximum, minimum, and intermediate active-site volumes. Each candidate was then subjected to in silico ligand docking screens in Autodock 4.0 (Osterberg et al., 2002). The heme charges were manually edited to match the high-spin ferric state, as described previously (Shahrkoh et al., 2012). The following ligands were obtained from PubChem and docked to each candidate model: all-trans-RA (compound identifier CID 444795), 9-cis-RA (CID 449171), 13-cis-RA (CID 5282379), and 4-oxo-arRa (CID 6437063). Side-chains from the following residues were selected to be flexible during the docking simulations for all substrates: Thr-117, Phe-295, Phe-299, and Ile-497. Additionally, Leu-131 was made flexible for the docking of 4-oxo-arRa, and Thr-300 was made flexible for the docking of 9-cis-RA and 4-oxo-arRa. Initial docking screens were performed with 10 simulations per ligand and 2 × 10$^5$ energy evaluations per docking simulation. The model derived from the 8-nanosecond time point has an active-site volume that reaches a local maximum (Supplemental Fig. 2B) and permits the binding of each ligand with high affinity. This model was chosen as the most suitable candidate. The final CYP26C1 homology model was subjected to more extensive docking studies of 200 simulations per ligand with the same parameters as above. The quality of the model before and after refinement was validated by Ramachandran plot analysis (RAMPAGE server; Lovell et al., 2003) and Verify3D score (Eisenberg et al., 1997). Parameters of homology model quality assessment and the secondary structure assignments are listed in Supplemental Tables 3 and 4.

**Results**

**Expression and Characterization of CYP26C1.** Microsomal CYP26C1 prepared from baculovirus-infected Sf9 insect cells showed a typical CO-difference spectrum with a defined peak at 448 nm, which was not observed in the negative control (Fig. 3). The P450 content, calculated from the CO-difference spectrum and from the microsomal protein content, was 84.3 pmol/mg of protein (2.2 μM) in the membrane preparation. The Western blot developed with anti-6His antibody indicated expression of a protein with an approximate size of 50–60 kD, corresponding to the predicted molecular weight of CYP26C1, 57 kD (Fig. 3), demonstrating CYP26C1 expression in Sf9 insect cell microsomes.

**Identification and Characterization of CYP26C1 Substrates.** Among all tested retinoids, all three RA isomers, 4-0H-RA isomers, and 4-oxo-arRa were substrates of CYP26C1, showing significant metabolite formation in the incubations with CYP26C1 on the basis of UV absorbance (Figs. 4–6). In contrast, no significant metabolite formation...
was detected in incubations with retinol isomers, 3-dehydro-atROL (Supplemental Fig. 3), 4-oxo-13-cis-RA (Fig. 6B), or 4-oxo-9-cis-RA (Fig. 6C). With all RA isomers (atRA, 13-cis-RA, or 9-cis-RA), the 4-hydroxylation pathway was the primary metabolic pathway resulting in formation of 4-OH-RA isomers, and subsequent minor formation of 4-oxo-RA and other hydroxylation products (M1–M8). The primary hydroxylation products at the C16 and C18 positions were observed from atRA, and C16 hydroxylation was observed from 13-cis-RA and 9-cis-RA (Fig. 4). The identity of these minor metabolites was confirmed by LC-MS/MS (Fig. 4, G–I). Notably, the polar metabolites (M1–M8) observed in the incubations with atRA, 13-cis-RA and 9-cis-RA (Fig. 4) corresponded exactly to those observed as products from the incubations of the corresponding 4-OH-RA isomers after incubation with CYP26C1 (Fig. 5), suggesting sequential metabolism of 4-OH-RA isomers formed from RA by CYP26C1. For 4-oxo-RA isomers, CYP26C1 metabolized 4-oxo-atRA to a single metabolite identified as 4-oxo-16-OH-atRA by LC-MS/MS (Fig. 6, A and D), but metabolism of 4-oxo-13cisRA and 4-oxo-9-cis-RA was negligible (Fig. 6). The identity of the metabolites formed by CYP26C1 was confirmed via collecting MS/MS spectra (Supplemental Figs. 4–6). The fragmentation patterns of metabolites were consistent with those described in previous studies with other CYP26s (Thatcher et al., 2011; Topletz et al., 2012).

To further define the metabolite pattern for CYP26C1, the stereospecificity of the formation of 4-OH-RA (Fig. 2) from atRA, 13-cis-RA, or 9-cis-RA by CYP26C1 was evaluated using chiral chromatography (Fig. 7). In the incubation with atRA, 4(R)-OH-atRA was the main enantiomer produced by CYP26C1, with a 4S/4R ratio of 0.2 (Fig. 7G). In comparison, similar amounts of 4(S)- and 4(R)-OH-13-cis-RA were formed by CYP26C1 with a 4S/4R ratio of 1.3 (Fig. 7H). From 9-cis-RA only 4(S)-OH-9-cis-RA was detected, demonstrating distinct stereospecificity in 4-OH-RA formation (Fig. 7I). The stereo-specificity of 4-OH-RA formation was clearly different for CYP26A1 in comparison with CYP26C1. CYP26A1 produced mainly 4(S)-OH-RA from all three RA isomers. The 4S/4R

Fig. 4. Characterization of the metabolism of RA isomers by CYP26C1. From left to right, CYP26C1 was incubated with atRA, 13-cis-RA, and 9-cis-RA, respectively. (A–C) LC-UV chromatograms of synthetic standards. Middle panels are UV (D–F) and LC-MS/MS chromatograms (G–I) of metabolite formation by CYP26C1. Indication of 16-OH-RA formation in LC-UV chromatograms (D–F) was based on MRM chromatograms (G–I) and EPI scan (Supplemental Figs. 4–6) despite the lack of 16-OH-RA standards. (L–N) Representative Michaelis-Menten plots from three repeated experiments for each substrate for the formation of the primary metabolites. To quantify the 4-OH-RA formation from atRA (L) and 13-cis-RA (M), the formation of 4-OH-RA and 4-oxo-RA were summed (Supplemental Fig. 1A). For 9-cis-RA (N), only the formation of 4-OH-9-cis-RA was quantified because the formation of 4-oxo-9-cis-RA was quantitatively insignificant. The data points shown are the mean values of duplicate measurements with error bars as range. Each kinetic experiment was repeated three times to obtain kinetic value estimates (Table 1).
ratios were 5.5 and 7.6 with atRA and 13-cis-RA as CYP26A1 substrates, respectively (Fig. 7, D and E). With 9-cis-RA as the substrate, only the peak of (4S)-OH-9-cis-RA was observed with CYP26A1 (Fig. 7F).

In addition to the separation of the 4-OH-RA enantiomers, the separation of 16-OH-13-cis-RA from 4-OH-13-cis-RA and 16-OH-9-cis-RA from 4-OH-9-cis-RA was much better when chiral chromatography (Supplemental Figs. 4–6) was used, rather than standard reversed phase separation (Fig. 4, E and F), allowing confirmation of the formation of these metabolites and collection of good quality MS/MS spectra of the metabolites of 13-cis-RA and 9-cis-RA (Supplemental Figs. 5 and 6).

The metabolites identified included the hydroxylations at C4 and C16 positions and sequential formation of the 4-oxo metabolites.

Enzyme Kinetic Parameters of RA Isomers and 4-oxo-atRA as Substrates of CYP26C1. The $k_{cat}$ values with CYP26C1 for all RA isomers were similar, whereas $K_m$ values of atRA and 13-cis-RA were 5.1- and 1.6-fold higher than those of 9-cis-RA, respectively (Table 1). The $Cl_{int}$ value for 9-cis-RA was 4.2- and 2.4-fold higher than that for atRA and 13-cis-RA, respectively. This suggests that CYP26C1 prefers 9-cis-RA as a substrate over atRA and 13-cis-RA (Fig. 4, L–N; Table 1). The relatively robust formation of the 16-OH metabolite from 9-cis-RA allowed the determination of the $K_m$ value toward this metabolite formation (5.9 ± 0.3 nM; mean ± S.D., $n = 3$; Supplemental Fig. 1B), as well as for 4-OH-9-cis-RA formation, and the $K_m$ values for the two metabolites were similar (Table 1). In the case of 4-oxo-atRA as a substrate of CYP26C1, similar $K_m$ values were obtained by measuring product formation and substrate depletion ($P < 0.01$; Fig. 6, E and F; Table 1). These $K_m$ values were similar to those of 9-cis-RA but significantly lower than those of atRA and 13-cis-RA ($P < 0.01$; Table1), suggesting that 4-oxo-atRA has higher affinity to CYP26C1 than atRA and 13-cis-RA, and CYP26C1 may prefer 4-oxo-atRA as a substrate. Because of

Fig. 5. Metabolism of 4-OH-RA isomers by CYP26C1. LC-UV chromatograms of 4-OH-atRA (A), 4-OH-13-cis-RA (B), and 4-OH-9-cis-RA (C) metabolism by CYP26C1 are shown in the presence and absence of NADPH. Sequential metabolites are labeled as M1–M8 corresponding to the numbering in Fig. 4, D–F.

Fig. 6. Metabolism of 4-oxo-RA isomers. Top panels depict LC-UV chromatograms of 4-oxo-atRA (A), 4-oxo-13-cis-RA (B), and 4-oxo-9-cis-RA (C) incubated with CYP26C1 in the presence and absence of NADPH. (D) shows the MRM chromatogram of 4-oxo-atRA incubated with CYP26C1 and MS/MS spectra of 4-oxo-16-OH-atRA. (E) Enzyme kinetics of 4-oxo-16-OH-atRA formation; Morrison equation was fit to the data owing to tight binding of 4-oxo-atRA to CYP26C1; each data point is shown as mean ± S.D., $n = 3$. (F) Representative depletion kinetic curve with each data point representing the $k_{dep}$ value obtained from the depletion experiment (inset).
the lack of synthetic 4-oxo-16-OH-atRA standard, the \( k_{\text{cat}} \) for product formation was not measured but the efficiency of 4-oxo-atRA metabolism was determined via substrate depletion (Fig. 6F). Of all the substrates, 9-cis-RRA had the highest Cl\(_{\text{int}}\) by CYP26C1, which was significantly greater (3.3-fold, \( P < 0.01; \) Table 1) than that of 4-oxo-atRA, whereas the Cl\(_{\text{int}}\) of atRA, 13-cis-RRA, and 4-oxo-atRA were not significantly different from each other (\( P > 0.01; \) Table 1).

**Effect of CRABPs on Metabolism by CYP26C1.** Since retinoids are extensively bound to CRABPs in cellular environment, the delivery of CRABP-bound substrates to metabolic enzymes has often been used as an indicator of the

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**TABLE 1**

| Enzyme kinetic parameters of RA isomers, 4-oxo-atRA, CRABP-bound atRA, and CRABP-bound 4-oxo-atRA as substrates with CYP26C1
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<td>( h_{\text{cat}} )</td>
<td>( K_m )</td>
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<tr>
<td></td>
<td>min(^{-1})</td>
<td>nM</td>
</tr>
<tr>
<td>atRA</td>
<td>5.9 ± 1.9</td>
<td>50.4 ± 6.2</td>
</tr>
<tr>
<td>13-cis-RRA</td>
<td>7.7 ± 2.9</td>
<td>38.2 ± 4.9</td>
</tr>
<tr>
<td>9-cis-RRA</td>
<td>4.7 ± 1.2</td>
<td>9.7 ± 2.6(^c)</td>
</tr>
<tr>
<td>4-oxo-atRA(^a)</td>
<td>15.7 ± 4.3(^d)</td>
<td>153 ± 20</td>
</tr>
<tr>
<td>4-oxo-atRA(^b)</td>
<td>10.2 ± 2.2(^d)</td>
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<tr>
<td>CRABP-I-atRA</td>
<td>2.1 ± 0.2</td>
<td>50.8 ± 8.0(^e)</td>
</tr>
<tr>
<td>CRABP-II-atRA</td>
<td>2.1 ± 0.8</td>
<td>34.1 ± 7.1(^r)</td>
</tr>
<tr>
<td>CRABP-I-4oxo-atRA</td>
<td>3.5 ± 0.5(^d)</td>
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<tr>
<td>CRABP-II-4oxo-atRA</td>
<td>5.4 ± 1.5(^d)</td>
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\(^a\)Kinetic parameters were obtained by measuring substrate depletion.

\(^b\)\( K_m \) value was obtained by measuring product formation.

\(^c\)\( P < 0.01, \) significantly different from the \( K_m \) value of atRA.

\(^d\)\( P < 0.01, \) significantly different from the \( K_m \) value of 13-cis-RRA.

\(^e\)\( P < 0.01, \) significantly different from the \( K_m \) value of 9-cis-RRA.

\(^f\)\( P < 0.01, \) the Cl\(_{\text{int}}\) value of 9-cis-RRA is significantly different from the other Cl\(_{\text{int}}\) values.
biologic importance of a metabolic enzyme in retinoid homeostasis. In addition, the delivery of atRA for metabolism by P450s has been shown to be enzyme-dependent (Nelson et al., 2016). For example, binding of atRA to CRABP-I and CRABP-II abolished the metabolism of atRA by CYP3A4 and CYP2C8 (Nelson et al., 2016), as predicted by the free drug hypothesis, i.e., that only drug free in solution and not protein-bound is available for metabolism. However, CRABP-I and CRABP-II delivered atRA for metabolism by CYP26B1 (Nelson et al., 2016). On the other hand, atRA binding to albumin resulted in reduced metabolism of atRA by CYP26B1, as predicted by free drug hypothesis and by albumin acting as a binding sink of atRA (Nelson et al., 2016). To investigate whether CYP26C1 takes any of its substrates from CRABP-I or CRABP-II, and interacts with the binding proteins, CYP26C1 was incubated with CRABP-bound atRA, 4-oxo-atRA, and 9-cis-RAL (CRABP-ligand ratio of 1:1; Fig. 8A). In the case of atRA, compared with the no CRABP incubations, 4-OH-atRA formation was decreased by 60%–70% in the presence of CRABP-I or CRABP-II (Fig. 8A). However, no change of 4-OH-9-cis-RAL formation by CYP26C1 was observed in the presence of CRABPs when 9-cis-RAL was the substrate (Fig. 8A). With 4-oxo-atRA as a substrate, 4-oxo-atRA metabolism by CYP26C1 was about 75% and 45% lower in the presence of CRABP-I and CRABP-II, respectively, in comparison with the incubations with 4-oxo-atRA as a substrate in the absence of the CRABPs (Fig. 8A).

To further investigate the interactions between CRABPs and CYP26C1 with different CYP26C1 substrates, the kinetics of metabolite formation were characterized using CRABP-bound substrates. In kinetic studies with CYP26C1 (Fig. 8B; Table 1), the $k_{cat}$ of atRA was 64% lower in the presence of

![Fig. 8. Effects of CRABPs on metabolism catalyzed by CYP26C1. (A) Bar graph of metabolite formation by CYP26C1 in the absence or presence of CRABPs; *** $P < 0.001$, analyzed by two-way ANOVA with Bonferroni post-test. For 9-cis-RAL, CRABP-ligand mixtures were prepared by premixing CRABPs with 9-cis-RAL in 1:1 concentration ratio but one-to-one binding was not confirmed as described for atRA and 4-oxo-atRA in the Materials and Methods section owing to the low binding affinity of 9-cis-RAL to CRABPs. Representative enzyme kinetic plots of product formation as a function of substrate concentration are shown for atRA (B) and 4-oxo-atRA (C) in the presence and absence of CRABP-I and CRABP-II. Data were fit to Michaelis-Menten equation for atRA and to Morrison equation for 4-oxo-atRA owing to tight binding. (D and E) Effects of excess apo-CRABP-I (D) and apo-CRABP-II (E) on product formation from atRA, 4-oxo-atRA, and 9-cis-RAL (100 nM for each substrate).]
CRABP-I and CRABP-II compared with free aTRA, but this difference was not statistically significant. The $K_m$ value of CRABP-I-αTRA and CRABP-II-αTRA remained unchanged compared with that of free αTRA ($P > 0.01$; Table 1). As αTRA binds to CRABPs tightly, virtually no unbound αTRA is expected to be present in the incubations, including CRABPs. Therefore, this data suggests that CRABPs deliver αTRA to CYP26C1. Although there was a trend toward decreased intrinsic clearance of αTRA in the presence of CRABPs, the intrinsic clearances were not significantly different ($P > 0.01$; Table 1) in the presence and absence of CRABPs. CRABP-I and CRABP-II affected 4-oxo-αTRA metabolism in a manner similar to αTRA (Fig. 8C; Table 1). The $K_m$ values for 4-oxo-αTRA obtained in the presence of CRABP-I and CRABP-II were similar to those obtained in the absence of CRABPs ($P > 0.01$; Table 1). The estimated maximum peak area ratio of 4-oxo-16OH-αTRA to the IS obtained in the presence of CRABP-I and CRABP-II (Fig. 8C), which is an indicator of the $k_{cat}$, was reduced about 80% and 60%, respectively, compared with that obtained in the absence of CRABPs, suggesting a similar decrease in the intrinsic clearance of 4-oxo-16OH-αTRA formation. However, owing to the lack of reference standard for this metabolite, the true $k_{cat}$ values could not be quantified.

To study the effect of excess apo-CRABPs on product formation by CYP26C1, a fixed concentration of substrate was mixed with varying concentrations of CRABP-I and CRABP-II, with the ratio of CRABP to substrate ranging from 0 to 4 (Fig. 8, D and E). In the case of αTRA and 4-oxo-αTRA, CYP26C1 activity decreased as the ratio of CRABP to substrate increased, reaching a plateau after the ratio exceeded 1. This suggests that apo-CRABPs do not compete with holo-CRABPs to interact with CYP26C1, but CRABPs may decrease metabolism by CYP26C1 via allosteric modulation or via noncompetitive inhibition. Surprisingly, neither CRABP-I nor CRABP-II had any effect on 9-cis-RA metabolism and there was no significant change in 4-OH-9-cis-RA formation from 9-cis-RA by CYP26C1 as the concentration of CRABP-I or CRABP-II increased from 0 to 400 nM. This further suggests that CRABP-P450 interactions are substrate-dependent.

**Binding Spectra and IC$_{50}$ Values of Talarozole and Ketoconazole.** The substrate specificity and stereoselectivity of the product formation with CYP26C1 suggests that CYP26C1 has different ligand interactions than do other members of the CYP26 family. To explore the ligand specificity of CYP26C1, the binding of the two azole inhibitors of CYP26A1 and CYP26B1, ketoconazole, and talarozole, were tested with CYP26C1. The coordination of the triazole or imidazole nitrogen with the heme iron causes a high-spin–to–low-spin shift of the iron, resulting in a type 2 binding spectrum. The binding of ketoconazole or talarozole to microsomal CYP26C1 induced a type 2 binding spectrum with some atypical characteristics, with the minimum absorbance observed between 410 and 420 nm and the maximum absorbance between 420 and 430 nm (Fig. 9, A and B). This suggests that theazole nitrogen in these two inhibitors can coordinate with the CYP26C1 heme. The binding was inhibitor concentration–dependent and showed increased spectral intensity with increasing inhibitor concentration. In agreement with the binding spectra, both ketoconazole

![Fig. 9. Characterization of the inhibition of CYP26C1 by imidazole- and triazole-containing inhibitors. Binding spectra of ketoconazole (A) and talarozole (B) with CYP26C1, and the inhibitor concentration–dependent decrease in CYP26C1 activity with ketoconazole (C) and talarozole (D). Inhibitor concentrations are indicated in binding spectra. Data points in (C) and (D) are shown as mean ± S.D., n = 4. For fitting of the IC$_{50}$ values, the y-intercept was constrained as less than 125% and the minimum activity remaining as greater than 0.](https://doi.org/10.1093/molpharm/081.2.498)
and talarozole inhibited CYP26C1 activity in an inhibitor concentration–dependent manner (Fig. 9, C and D). However, the IC\textsubscript{50} values, 124 \mu M (26–594; 95% confidence intervals) with ketoconazole and 3.8 \mu M (1.8–8.1; 95% CI) with talarozole, were much higher than the IC\textsubscript{50} values observed with CYP26A1 (<10 nM for talarozole and 0.55 \mu M for ketoconazole) and CYP26B1 (<10 nM for talarozole and 0.59 \mu M for ketoconazole) (Thatcher et al., 2011; Diaz et al., 2016; Foti et al., 2016a).

**Homology Model of CYP26C1 and Molecular Docking of Retinoic Acid Isomers.** The experimental data of substrate selectivity, metabolic specificity, ligand binding, and CRABP interactions suggest that CYP26C1 has functional differences compared with CYP26A1 and CYP26B1, and that substrates have different orientations in the CYP26C1 active site compared with other CYP26 enzymes. To investigate the basis of substrate recognition by CYP26C1, a homology model of CYP26C1 was developed and refined by energy minimization and MD simulations (Supplemental Fig. 2; Supplemental Tables 3 and 4). RA isomers (\(\alpha\)-RA, 9-cis-RA, 13-cis-RA) and 4-oxo-\(\alpha\)-RA were then docked in silico to the CYP26C1 active site. On the basis of the docking simulations, \(\alpha\)-RA, 9-cis-RA, and 13-cis-RA all positioned in a similar fashion within the CYP26C1 active site, with the \(\beta\)-ionone ring closest to the heme iron. 4-oxo-\(\alpha\)-RA was rotated about 90° relative to RA isomers in its preferred orientation within the active site (Fig. 11). The docking simulations could appropriately predict the sites of oxidation of all four CYP26C1 substrates. According to the distance between the heme and carbons at positions 4, 16, and 18 on the \(\beta\)-ionone ring, the CYP26C1 model predicted that \(\alpha\)-RA, 9-cis-RA, and 13-cis-RA are primarily oxidized at carbon 4, and 4-oxo-\(\alpha\)-RA is mainly oxidized on carbon 16 (Table 2). At the prochiral carbon 4 position of \(\alpha\)-RA, the hydrogen that would account for the formation of 4(R)-OH-\(\alpha\)-RA oriented toward the heme iron with a closer distance (3.9 Å) compared with the hydrogen that, when abstracted, would lead to the formation of 4(S)-OH-\(\alpha\)-RA (5.0 Å; Fig. 11A). Thus, the model predicted that CYP26C1 preferentially forms 4(R)- over 4(S)-OH-\(\alpha\)-RA from \(\alpha\)-RA. For 13-cis-RA, pro-S and pro-R hydrogen atoms at C4 position were 3.7 and 3.8 Å away from the heme iron, respectively, leading to the prediction of racemic 4-oxo-13-cis-RA formation by CYP26C1 (Fig. 11B). Because the \(\beta\)-ionone rings of \(\alpha\)-RA and 9-cis-RA are flipped 180° relative to each other, the formation of 4(S)-OH-9-cis-RA was more favorable than the formation of 4(R)-OH-9-cis-RA (Fig. 11C). The distance between the heme iron and the hydrogen atom at the pro-S and pro-R position of 9-cis-RA were 3.1 and 4.6 Å, respectively.

Compared with the crystal structure of \(\alpha\)-RA-bound CYP120A1 (Kühnel et al., 2008), the CYP26C1 model revealed that a C-terminal loop (amino acid L484-L504) occludes the binding pocket (Fig. 10). The amino acids proposed to interact with RA isomers in the active site are shown in Fig. 12 and Supplemental Table 5. Leu-221 and Phe-222 are predicted to interact with the carboxylic acid tail of \(\alpha\)-RA, 9-cis-RA, and 4-oxo-\(\alpha\)-RA, whereas Ser-120 is predicted to interact with that of 13-cis-RA. Notably, on the basis of the homology model, CYP26C1 active site lacks the positively charged amino acids such as Arg within the active site to interact with the carboxylic acid moiety in \(\alpha\)-RA.

**Discussion**

To date, most studies related to the CYP26 family have focused on CYP26A1 and CYP26B1, which are known to be important RA hydroxylases (Thatcher and Isoherranen, 2009; Toplzeit et al., 2012). The importance of CYP26C1 in retinoid clearance has been questioned, as Cyp26C1\textsuperscript{-/-} mice do not show any of the phenotypic malformations typical for knock-out models of retinoid metabolism (Uehara et al., 2007). However, the tissue-specific importance of the individual CYP26 enzymes in retinoid clearance has not been defined, and depends on both the expression levels of the CYP26 enzymes and their catalytic activity toward the specific substrates. As the sequence homology of CYP26C1 with CYP26A1 and CYP26B1 is low (43%–51%), we hypothesized that CYP26C1 differs functionally from CYP26A1 and CYP26B1. Our data demonstrates that CYP26C1 has different ligand (inhibitor and substrate) specificity than CYP26A1 and CYP26B1, illustrated most strongly by the differences in the IC\textsubscript{50} values for talarozole and ketoconazole with CYP26C1 enzymes. Although talarozole and ketoconazole binding to CYP26C1 active site induced a type II binding spectrum with some atypical characteristics similar to CYP26A1 (Thatcher and Isoherranen, 2009), talarozole and ketoconazole IC\textsubscript{50} values were more than 100-fold higher with CYP26C1 than with CYP26A1 and CYP26B1 (Thatcher et al., 2011; Diaz et al., 2016). Comparisons of IC\textsubscript{50} values between enzymes can be confounded by different substrates and by different substrate concentration/K\textsubscript{m} ratios used, and the inhibitory potency of competitive inhibitors can be underestimated if substrate is used at concentrations approaching its K\textsubscript{m}. Even still, the IC\textsubscript{50} data obtained demonstrate clear differences in ligand interactions within the CYP26 family, and CYP26C1 displays a clear lack of susceptibility to these inhibitors. The different ligand interactions were also evident with retinoids. Whereas \(\alpha\)-RA has a lower K\textsubscript{m} with CYP26A1 and CYP26B1 than 9-cis-RA, with CYP26C1 9-cis-RA exhibited the lowest K\textsubscript{m} and highest intrinsic clearance. In addition, the K\textsubscript{m} of 9-cis-RA with CYP26C1 (K\textsubscript{m} = 9.7 nM) was much lower than...
with CYP26A1 ($K_m = 134$ nM; Thatcher et al., 2011) and CYP26B1 ($K_m = 555$ nM; Diaz et al., 2016). This is in agreement with a previous study in which radiolabeled substrate competition assays showed that 9-cis-RA was a better ligand for CYP26C1 than for the other CYP26s (Taimi et al., 2004; Helvig et al., 2011). Interestingly, 9-cis-RA metabolism by CYP26C1 was unaffected by the CRABPs (Fig. 8), possibly because the binding affinity of 9-cis-RA to CRABPs is much weaker than that of 9tRA and 4-oxo-9tRA (Fiorella et al., 1993) resulting in higher free fraction of 9-cis-RA. These findings also indicate a high clearance of 9-cis-RA in vivo, a process unprotected by CRABPs. This may explain the lack of detection of 9-cis-RA in common tissue samples. In human serum 9-cis-RA was detected at a very low level (0.08–0.1 nM) only in a subset of individuals (Arnold et al., 2012). Likewise, in mice 9-cis-RA was only detected in the pancreas (Kane et al., 2010; Kane, 2012). The results shown here suggest that the detection of 9-cis-RA may be dependent on CYP26C1 expression, and further studies are needed to clarify the expression pattern of CYP26C1 in humans, and the role of CYP26C1 in 9-cis-RA clearance. CYP26C1 mRNA has been detected in human fetal liver and brain and in adult human brain, lung, liver, spleen, and testis (Xi and Yang, 2008), suggesting that CYP26C1 is expressed in these retinoid target tissues.

All the data presented here suggest that CYP26C1 plays a unique role in the clearance of the active 9tRA metabolite 4-oxo-9tRA as a 4-oxo-9tRA hydroxylase, and that 4-oxo-9tRA may be an important active retinoid in humans. However, the microsomal enzyme(s) responsible for 4-oxo-9tRA formation is still unknown. As with CYP26A1 and CYP26B1 (Lutz et al., 2009; Topletz et al., 2012), CYP26C1 forms 4-oxo-9tRA from 4-OH-9tRA but with minimal efficiency, supporting the prior report that CYP26 enzymes are important in the elimination of bioactive retinoids, not in their formation (Lutz et al., 2009; Topletz et al., 2012). Previously, 4-oxo-9tRA has been shown to be a substrate of both CYP26A1 ($K_m = 63$ nM; $Cl_{int} = 91$ μl/min per picomol) and CYP26B1 ($K_m = 29$ nM; $Cl_{int} = 15$ μl/min per picomol) (Topletz et al., 2012). The $K_m$ values of 4-oxo-9tRA with CYP26A1 and CYP26B1 were 4- and 1.8-fold higher than the $K_m$ of 4-oxo-9tRA with CYP26C1, respectively, and the $Cl_{int}$ of 4-oxo-9tRA was 1.7- and 10-fold higher with CYP26C1 than with CYP26A1 and CYP26B1, respectively. Most notably, CYP26C1 appeared to take CRABP-bound 4-oxo-9tRA as a substrate, similar to its acceptance of CRABP-bound 9tRA as a substrate. This suggests that CRABPs play a role in modulating 4-oxo-9tRA clearance in the cellular environment. In contrast to our findings, in rat testes microsomes CRABP binding eliminated 4-oxo-9tRA metabolism completely (Fiorella and Napoli, 1994), suggesting that the substrate delivery is P450-specific. This is consistent with the general observation that retinoid exchange between cellular binding proteins and metabolic enzymes is unique to specific binding protein–enzyme pairings (Napoli, 2017). Substrate delivery to CYP26C1 by CRABPs together with the relative expression

![Fig. 11. Comparison of the predicted binding orientations of RA isomers with CYP26C1. The predicted binding orientations of 9tRA ((A), magenta), 13-cis-RA ((B), orange), 9-cis-RA ((C), light green), and 4-oxo-9tRA ((D), cyan) with respect to the heme are shown. Dashed lines compare the distances between the heme iron and the hydrogens at C4 position (A–C) or at C16 position (D).](image-url)
levels of CYP26 enzymes, probably determines the physiologic importance of CYP26C1.

The data shown here reveal important information about the P450-CRABP interactions. Owing to the high binding affinity of 4-oxo-ara and ara with CRABPs ($K_d < 20$ nM) (Fiorella and Napoli, 1991; Fiorella et al., 1993), 4-oxo-ara and ara are expected to be bound with CRABPs in the incubations and in cells with virtually no free retinoids in solution. This lack of free ara or 4-oxo-ara in the holo-CRABP–containing incubations was confirmed via back-purification of the holo-CRABPs prior to incubations. If CRABPs acted as a sink of retinoids and did not deliver the retinoids for metabolism (i.e., according to free drug hypothesis), the $K_m$ for retinoid metabolism should have been increased in these incubations, as was observed in the presence of albumin (Nelson et al., 2016). In contrast, the $K_m$ values of ara and 4-oxo-ara bound to CRABP were similar to those of free ara and 4-oxo-ara, respectively, with a trend toward lower $k_{cat}$ with CRABP-bound substrates. These findings are similar to decreased $k_{cat}$ or $V_{max}$ and $K_m$ for CRABP-bound ara reported in rat testis microsomes and recombinant CYP26B1 (Fiorella and Napoli, 1991; Nelson et al., 2016). The data with CYP26C1 supports the notion that CRABPs deliver active retinoids for metabolism via substrate channeling, and strongly suggest that CRABPs deliver both ara and 4-oxo-ara to CYP26C1 for metabolism. In contrast, the lack of effect of CRABPs on 9-cis-ara metabolism suggests that CRABPs do not modulate 9-cis-ara clearance. The lack of effect of CRABPs on 9-cis-ara metabolism can be explained by the low binding affinity of 9-cis-ara with CRABPs and the low $K_m$ of 9-cis-ara with CYP26C1.

The differences in substrate specificity, inhibition profiles, and stereospecificity of metabolite formation by CYP26C1, all suggest that the retinoids have different binding orientations within the CYP26C1 active site compared with CYP26A1 and CYP26B1. Specifically, CYP26C1 formed 4-OH-9-cis-ara as the major metabolite of 9-cis-ara and the formation of 16-OH-9-cis-ara appeared very minor, whereas CYP26A1 made these metabolites at equal levels (Thatcher et al., 2011). Likewise, CYP26C1 formed predominantly the 4(R)-OH-ara as the major metabolite of 9-cis-ara and the formation of 4(S)-OH-ara appeared very minor, whereas CYP26A1 made these metabolites at equal levels (Thatcher et al., 2011). Likewise, CYP26C1 formed predominantly the 4(R)-OH-ara, whereas CYP26A1 formed the 4(S)-OH-ara, suggesting different orientation of ara within the CYP26A1 and CYP26C1 active sites. To explore the binding pocket of CYP26C1, a homology model of CYP26C1 was built on the basis of a complement of P450 crystal structures. Previously, CYP26A1 and CYP26B1 homology models were constructed mainly on the basis of the CYP120A1 crystal structure (Karlsson et al., 2008; Kühnel et al., 2008; Shimshoni et al., 2012; Foti et al., 2016b). The
predicted orientation of αrTA in the active site in these homology models (Shimshoni et al., 2012; Foti et al., 2016b) was similar to that observed in the crystal structure of CYP120A1 (Kuhnel et al., 2008), with the carboxylic acid group of αrTA interacting with Arg64, Arg86, and Arg90 of CYP26A1 or Trp65, Arg76, Tyr372, and Arg373 of CYP26B1. However, the CYP26C1 homology model revealed that a hairpin loop located at the C terminus protrudes ~5 Å deeper into the active site than the homologous loop in CYP120A1, which forces αrTA and other docked substrates to orient differently, with carboxylic acid interacting with Leu221 and Phe222 (αrTA) and 9-cis-RA or Ser120 (13-cis-RA). This orientation was predicted to be different in the steric specificity of the C4-hydroxylation of αrTA. The overall active-site volume of CYP26C1 was predicted to be approximately 200–250 Å³, much smaller than the predicted active-site volume of CYP26A1 (918 Å³) and CYP26B1 (977 Å³) (Foti et al., 2016b). This indicates that the CYP26C1 active site may offer fewer potential substrate binding orientations than the other two CYP26 enzymes, an observation consistent with the much less diverse metabolite profile observed with CYP26C1 than with CYP26A1 and CYP26B1 (Thatcher et al., 2011; Topletz et al., 2012). The small active-site volume may also explain the high IC₅₀ of talazopar and ketoconazole with CYP26C1. Bearing in mind the inherent limitations of homology modeling, the distinct structural properties of the CYP26C1 active site predicted by the structural model are consistent with the experimental findings presented here, and successfully captured the regio- and steric specificity of the enzyme, providing insights into the CYP26C1 active site.

Acknowledgments

The authors thank Dr. Jay Kirkwood for his skilful assistance on LC-MS/MS experiments and Brian Buttrick for his help in cloning the CYP26C1.

Authorship Contributions

Participated in research design: Zhong, Ortiz, Nath, Isiherranen.
Conducted experiments: Zhong, Ortiz.
Contributed new reagents or analytic tools: Zelter.
Wrote or contributed to the writing of the manuscript: Zhong, Ortiz, Zelter, Nath, Isiherranen.

References


**Address correspondence to:** Dr. Nina Isoherranen, Department of Pharmaceutics, School of Pharmacy, University of Washington, Health Science Building, Room H-272M, Box 357610, Seattle, WA 98195-7610. E-mail: n2@uw.edu