Structural and Functional Similarity of Amphibian Constitutive Androstane Receptor with Mammalian Pregnan X Receptor

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Abstract

The nuclear receptors and xenosensors constitutive androstane receptor (CAR, NR1I3) and pregnane X receptor (PXR, NR1I2) induce the expression of xenobiotic metabolizing enzymes and transporters, which also affects various endobiotics. While human and mouse CAR feature a high basal activity and low induction upon ligand exposure, we recently identified two constitutive androstane receptors in Xenopus laevis (xlCARa and a) that possess PXR-like characteristics such as low basal activity and activation in response to structurally diverse compounds. Using a set of complementary computational and biochemical approaches we provide evidence for xlCARa being the structural and functional counterpart of mammalian PXR. A three-dimensional model of the xlCARa ligand-binding domain (LBD) reveals a human PXR-like L-shaped ligand binding pocket with a larger volume than the binding pockets in human and murine CAR. The shape and amino acid composition of the ligand-binding pocket of xlCAR suggests PXR-like binding of chemically diverse ligands which was confirmed by biochemical methods. Similarly to PXR, xlCARa possesses a flexible helix 11'. Modest increase in the recruitment of coactivator PGC-1α may contribute to the enhanced basal activity of three gain-of-function xlCARa mutants humanizing key LBD amino acid residues. xlCARa and PXR appear to constitute an example of convergent evolution.

Introduction

Ligand-modulated transcription factors of the nuclear receptor (NR) superfamily are involved in regulation of various physiological processes such as development and homeostasis and also play a prominent role in metabolism and disposition of exogenous compounds from the body [1], [2]. In mammals the latter process is mainly governed by pregnane X receptor (PXR, NR1I2) and constitutive androstane receptor (CAR, NR1I3). CAR and PXR arose in early vertebrates from a common ancestral gene (Figs. 2 and 4 in [3]) and they both recognize chemically diverse substances including pesticides, plasticizers as well as prescription drugs [4]. Target genes of CAR and PXR comprise two overlapping panels of phase I-III metabolizing enzymes and transporters. Represented by cytochrome P450 (CYP) enzymes CYP2B6 and CYP3A4, the most relevant human drug metabolizing CYPs are under transcriptional control of CAR and PXR [5].

In contrast to most other NRs, including PXR, the major splice variant of human CAR (CAR SV1) reveals a pronounced ligand-independent activity in vitro accompanied by a limited activation potential upon agonist binding [6],[7]. In contrast, the recently identified Xenopus laevis CARα (xlCARα) possesses a low basal activity in vitro and a large induction potential by agonists [3]. The ligand spectrum (including natural products, prescription drugs and endogenous substances) was found to be significantly broader compared to the human ortholog. Further investigations revealed that the constitutive CAR activity typical for mammals emerged first with fully terrestrial land vertebrates [3].

By means of experimental and theoretical studies, the structural basis for the constitutive activity of human and mouse CAR has been addressed in several previous studies. Using homology modeling approaches and X-ray crystallography studies, 3-dimensional structures of the CAR ligand-binding domain (LBD) have been generated and used for identification of single amino acids relevant for constitutive activity with subsequent experimental validation by means of site-directed mutagenesis [8–12]. In many cases the standard procedure – replacing the amino acid of interest by alanine – resulted in complete loss of basal activity and ligand responsiveness, thus demonstrating the limiting informative value of this approach. Recently, we identified two amino acids within the xlCARα ligand-binding pocket (LBP) whose mutation towards the human receptor resulted in a substantially increased
basal activity [3]. The gain in receptor activity was accompanied by a reduced ligand induction potential. The mutated receptor phenotypically resembled human CAR, providing a model system for further investigating the structural basis of its constitutive activity.

In addition, our recent study provided first evidence for xlCAR involvement in physiological processes different from those described previously for mammalian CAR as suggested by a much wider xlCAR organ expression. Besides liver, the receptor is strongly expressed in lung, skin, stomach, kidney and ovary, with lower expression levels in brain and heart [3]. This is in clear contrast to human CAR expression that – besides liver and intestine – is only expressed at low levels in other tissues [4]. Therefore, xlCAR inducers may be capable to affect more organs in amphibians, which may contribute to the extraordinary sensitivity of amphibians to environmental pollutants compared to mammals [3]. A more detailed investigation of the structural and functional properties of amphibian constitutive androstane receptor may support future toxicological and ecological risk assessments. Moreover, as shown by the previously identified gain-of-function mutations, the receptor may represent a model system for further investigating the structural determinants of mammalian CAR basal activity.

In this study, we intended to characterize the xlCAR ligand-binding domain from a structural and functional point of view by investigating the overall LBD structure, its interaction possibilities with ligands and the heterodimerization partner RXRα (retinoid X receptor α) as well as two coactivators, using a combination of in silico and in vitro approaches. As no x-ray crystal structure is available for the xlCAR LBD, we utilized a homology modeling approach, similar to previous studies on human CAR [11–13], by which the three-dimensional structure of the xlCAR ligand-binding domain was constructed based on a sequentially related x-ray crystal structure. The model guided the selection of amino acids to be mutated in in vitro assays, validating the outcome of the modeling process. Putative binding modes of previously identified agonists were determined using a molecular docking approach. This procedure was validated and complemented by various in vitro assays using receptor mutants by which the effects of ligand binding on receptor activation and heterodimerization with RXRα as well as on recruitment of coactivators SRC-1 and PGC-1α were investigated.

Materials and Methods

Homology model generation
In order to generate a homology model of the xlCAR ligand-binding domain, a BLAST search querying the Protein Data Bank (PDB) was performed for identification of sequentially related protein structures. Subsequently, x-ray crystal structures of proteins with highest sequence similarity (human CAR and PXR) were downloaded from the PDB [14]. Similar to human VDR and PXR, xlCAR contains a H1–H3 insert [3]. According to secondary structure predictions using PSIPRED (http://bioinf.cs.ucl.ac.uk/psipred/, see also Figure S1), the xlCAR H1–H3 insert lacks any secondary structural elements. Thus, PXR x-ray crystal structures are not suited as template structures as these contain a helix and two β-strands within the H1–H3 which are associated with further structural changes in helices 6 and 7 [15].

The identification of a suitable template structure from the set of available CAR x-ray crystal structures was guided by stereochemical parameters as determined with PROCHECK and the folding reliability as assessed with ProSa 2003 [16], [17]. Finally, PDB code 1xv9 was chosen as main modeling template [9]. Including results from secondary structure prediction for xlCARα using PSIPRED, two modifications were introduced using MOE (Chemical Computing Group, Montreal, Canada): as helix 3 was predicted to be shorter in xlCARα than in human CAR, amino acids 155–157 were deleted from the original template structure. In return, PSIPRED predicted helix 10/11 to possess a C-terminal extension. Therefore, coordinates for residues 410–416 of human PXR (PDB code 1nrl), that contains an extended helix 10/11, were transferred into the template structure upon superpositioning of PXR with CAR. As protein modeling tools generally fail to reliably predict structures of loops longer than 10–12 residues, amino acids 199–230 of the insert were not considered for the homology modeling procedure. The homology model of xlCARα was constructed employing MOE thereby using the sequence alignment shown in Figure S1. Initially, ten structures were generated which were energy minimized using the Amber force field with Born solvation as implemented in MOE. Coordinates of the final protein model are available on request from the authors. Shapes of the ligand-binding pockets of xlCARα, hsCAR and hsPXR and determination of their volumes were calculated using MOLCAD within SYBYL-X.1.2 (Tripos Int., St Louis, MO, USA) after addition of all hydrogen atoms.

Molecular Dynamics Simulations
Human CAR x-ray crystal structure (PDB code 1xvp, chain D) was used for the simulations. The Thr232 mutation was generated using SCWRL3 [18]. All simulations were performed using GROMACS version 3.3.3 [19]. The setup of the simulation system as well as the equilibration and production procedures were performed as described previously [20].

Molecular docking
Three-dimensional structures of known xlCAR ligands artemisinin, fenofibrate and pregnanedione were generated within SYBYL-X1.2 and subsequently energetically minimized. Molecular docking was performed using GOLD version 5.1 (Cambridge Crystallographic Data Centre). For each ligand, 10 independent docking runs were performed. The early termination option was switched off. Of the scoring options provided with GOLD, ChemScore was selected.

Plasmids and mutagenesis
The following firefly luciferase reporter gene plasmids have been described previously: human CYP3A4 enhancer/promoter plasmid pGL3-CYP3A4(7830/A7208–364) [21], hereinafter called CYP3A4 enhancer/promoter; human MDR1 enhancer DR4 (I) element-dimer/Tk promoter plasmid [22], hereinafter called DR4-Tk promoter; Gal4-dependent reporter gene construct pGL3-G5 [23]. The Renilla luciferase reporter gene plasmid pRL-EF1α was described previously [24]. The following eukaryotic expression plasmids have been published elsewhere: encoding full-length human CAR1 [25], xlCARα [3] and human PXR [22]; encoding fusion proteins of the GAL4 DNA-binding domain (DBD) and receptor interaction domain (RID) of human coactivator SRC-1 (amino acids 583–783) [23], or RID of human PGC-1α (amino acids 38–215) [3], or ligand binding domain (LBD) of human RXRα (amino acids 226–462) [23]; encoding fusion proteins of the VP16 activation domain (AD) and LBD of human CAR1 (amino acids 105–348) [23].

Base pair mutations and deletions were introduced into the respective expression plasmids using appropriate oligonucleotide primers (Table S1) and the QuikChange Site-Directed Mutagen-
esis Kit (Agilent, Santa Clara, CA), according to manufacturer’s instructions. All mutated constructs were verified by sequencing.

Cell culture, transient transfections and reporter gene assays

Human colon adenocarcinoma LS174T cells (ATCC, Manassas, VA) and human hepatoma HepG2 cells (ATCC) were maintained as previously described [23]. Cells were plated into 96-well plates at a density of 2.5 × 10^4 cells/well (LS174T) or 24-well plates at 1.5 × 10^5 cells/well (HepG2) 24 h before transfection, which was performed using GeneJuice (Merck Millipore, Darmstadt, Germany) or Effectene (Qagen, Hilden, Germany) reagents for LS174T or HepG2 cells, respectively, according to manufacturer’s instructions. The combinations of co-transfected firefly luciferase reporter gene and expression plasmids are indicated in the respective Figure legends. Firefly luciferase activities, determined in cell lysates as described [26], were normalized with respect to Renilla luciferase activities of co-transfected pRL-EF1α. At least three independent experiments, each done in triplicates, were performed using a minimum of two different plasmid DNA preparations of the respective nuclear receptor expression plasmids.

For mammalian two-hybrid assays, COS7 cells were plated into 96-well plates at a density of 1,500 cells per well. Cells were transfected using Gene Juice and 36.7 ng of pGL3-G5, 3.3 ng of the respective GAL4-DBD-fusion protein expression plasmid, together with 26.7 ng of respective expression plasmids encoding fusion proteins of VP16-AD and the wild type or mutant xlCAR LBDs and 7.7 ng of pRL-EF1α. Protein expression of the VP16-AD/CAR-LBD fusion proteins were determined by Western blot using anti-VP16 (14–5) antibody (Santa Cruz).

Recombinant protein expression

Recombinant human SRC-1 protein (amino acids 583–783, comprising the receptor interaction domain), fused at the amino-terminus to GST, was expressed in bacteria as described previously [27]. In vitro transcription/translation (TXT T7 quick coupled transcription/translation system, Promega, Madison, WI, USA) was used to synthesize [35S]-methionine labelled full-length X. laevis and human CAR proteins in 25 μl reactions, containing 0.5 μg of the respective expression plasmids and 10 μCi [35S]-methionine (specific activity 1175 Ci/mmol, radioactive concentration 10 μCi/μl, MP Biomedicals, Santa Ana, CA, USA). Equal protein expression of mutant receptors (Fig. S2) was confirmed by protein gel electrophoresis of an aliquot. Gels were stained with Coomassie, dried and exposed to BAS-IP MS 2325 imaging plates (Fuji, Kanagawa, Japan), which were quantified using the CR35 Bio radioluminography laser scanner and AIDA software (Raytest, Straubenhardt, Germany).

Coactivator-dependent receptor ligand assay (CARLA)

CARLA was performed essentially as described before [27]. Briefly, 1 ml reactions were set up in NETN (100 mM NaCl, 1 mM EDTA, 1 mM DTT, 20 mM Tris-Cl pH 8.0, 0.5% (v/v) Nonidet P40), with 0.5% (w/v) skimmed milk powder, using 3–5 μg of GST-tagged SRC-1 RID protein, bound to 25 μl bead volume of glutathione-sepharose 4B beads (GE Healthcare, Freiburg, Germany), 2 μl of respective [35S]-labelled CAR protein, and the respective chemicals or 1% solvent DMSO only. After incubation over night at 4°C with constant rotation, beads were washed three times in 1 ml NETN buffer, supplemented with the respective chemicals. Bound GST/SRC-1 fusion protein/CAR complexes were extracted from the beads by boiling in SDS-sample buffer and separated on 10% SDS-polyacrylamide gels, which were subsequently stained with Coomassie, dried and exposed to BAS-IP MS 2325 imaging plates. CAR protein bound to SRC-1 was detected by reading the image plates with CR35 Bio radioluminography laser scanner and quantified by densitometric scanning of the image, using AIDA software. Coomassie staining of the protein gels demonstrated the use of equal amounts of GST/SRC-1 fusion protein in each reaction. Respective control experiments, which had been set up with GST protein only, demonstrated negligible binding of CAR proteins to the GST moiety of the GST/SRC-1 fusion protein.

Statistical analysis

Multiple comparisons were performed by one-way or two-way analysis of variance (ANOVA) with Dunnett’s multiple comparisons test, as indicated. Pairwise comparisons were done by paired t-test and comparisons to a hypothetical mean by one sample t-test. In the latter case, the resulting P-values were corrected for multiple testing using the method of Bonferroni. All calculations were performed with GraphPad Prism version 6.03 (GraphPad Software, La Jolla, CA). Statistical significance was defined as P<0.05.

Results

Homology model of xlCARα

In order to investigate the 3-dimensional structure of X. laevis constitutive androstane receptor xlCARα, we constructed a homology model of its ligand-binding domain (residues 145–425). Utilizing sequence comparisons with related nuclear receptors and secondary structure predictions, a chimeric template structure for the homology modeling process was constructed from coordinates of x-ray crystal structures solved for human CAR and PXR.

As no suitable template structure was available for modeling the 62 residues long and, according to results from secondary structure prediction tools, unstructured H1–H3 insert, this part of the LBD was largely excluded (amino acids 199–230) from the modeling process. In order to justify this approach we experimentally assessed this region for any involvement in receptor activity or ligand-binding as seen in x-ray crystal structures of human PXR [15]. At first, a deletion mutant excluding amino acids 186–229 of xlCARα was generated. For further comparison with PXR, which possesses a similar H1–3 insert contributing to LBP-formation, the corresponding region (residues 182–233) was removed. While the PXR deletion mutant resulted in a loss-of-function, the truncated xlCARα revealed no change in basal activity and responsiveness to ligands (Fig. 1), suggesting that the H1–H3 insert is not involved in LBP formation.

The structural and functional outcome of the resulting protein model was evaluated using PROCHECK and ProSa. The Ramachandran plot revealed 97.8% of all amino acids adopting most favorable or additionally allowed ϕ/ψ backbone torsion angles. Residues with less favorable geometries were found to be largely located in the remaining parts of the H1–H3 insert. The ProSa Z-score of -10.28 for the 247 amino acids comprising xlCARα LBD indicated a reliable overall fold and was better than all scores determined for human CAR x-ray crystal structures (varying from -9.70 to -10.05) which possess an almost identical number of residues. Protein regions with unfavorable geometries were mainly restricted to the truncated H1–H3 insert (data not shown).

Figure 2A shows the three-dimensional structure of the resulting LBD homology model. Due to the small extension of helix 10/11 at its C terminus, the xlCARα model suggests the absence of a
helical segment in between helix 10/11 and the activation helix (H12) found in x-ray crystal structures of murine and human CAR. Instead, the loop shows an extended conformation.

The L-shaped ligand-binding pocket is framed by 33 amino acids located on helices 3, 5–7, 10 and 12 as well as β-strands 1–3 (Table 1). Additionally, few amino acids of the undeleted parts of the H1–H3 insert were found to line the LBP. However, as a major part of the insert was excluded from the modeling procedure, any contribution of those residues to LBP formation remains speculative and thus they are not listed in Table 1. The ligand-binding pocket captures a volume of 915 Å³ which is larger compared to human and murine CAR (750–810 Å³), but smaller than in human PXR (1320–1420 Å³). The shape of the pocket less resembles the ellipsoid shape of human or mouse CAR and is more alike the T-shaped form observed in human PXR x-ray crystal structures (Fig. 2B&C).

Similar to human CAR, the LBP is almost completely separated from the activation helix by a barrier formed by amino acids Phe234, Ser238 and Ile242 (all on H3), Met272 (on H5) as well as Tyr399 and Ile403 (both on H10/11). A hydrogen bond is shared between Ser238 and Tyr399 side chains, connecting helices 3 and 10/11. A similar interaction is seen in mammalian CAR, involving Asn165 (H3) and Tyr326 (H10/11) of the human receptor.

In hsCAR, a hydrogen bond connects Tyr224 on the β3 strand and His160 (H3). This interaction cannot be established in xlCAR as the corresponding amino acids (Phe297 and His233) do not allow hydrogen bond formation between their side chains. Instead, the model suggests a hydrogen bond between His295 and His233.

As indicated by the lipophilic potential mapped on the surfaces of the ligand-binding pockets in Figure 2, the xlCAR LBP amino
Acid composition is less hydrophobic compared to human CAR and provides more possibilities for sharing hydrogen bonds with ligands. A third of the amino acids constituting the ligand-binding pocket are capable of serving as hydrogen bond donor or acceptor via their side chain and/or backbone atoms (see Table 1).

**Basal activity**

Based on the homology model structure described above as well as on sequence comparisons with human CAR, we tested the relevance of selected amino acids within the LBD for basal receptor activity by mutating the respective amino acids to their corresponding residues in human CAR (Fig. 3). The selection focused on amino acids lining and surrounding the ligand-binding pocket of which some have been already identified as important for human CAR basal activity. Mutations towards the human ortholog not only provides the opportunity to investigate differences between human and *Xenopus* CAR but it is also beneficial compared to the conventional alanine mutation approach as the small alanine side chain may result in unwanted and non-predictable rearrangements of neighbor amino acids within the LBP [12].

Previously, we have shown a marked increase in basal activity upon mutation of Met272 into valine and of Met316 into phenylalanine, as well as an additive effect of both replacements [3], which was confirmed in the present study. We additionally report here the 8.6-fold increase in basal activity upon mutating Thr305, located on helix 6, into the corresponding human CAR amino acid valine (Val232) (Fig. 3A). The already low basal activity was further decreased 3-fold upon mutation of amino acid Val315 into isoleucine. Not unexpectedly, mutation of Tyr311 into the closely related phenylalanine did not reveal a significant change in basal activity. Also the mutation of His295, predicted to form a hydrogen bond with His233 on helix 3 and located more than 15 Å away from the LBD/H12 interface, did not influence basal activity.

In order to evaluate the molecular basis of the significant increase in basal activity of the three gain-of-function mutants, as well as of the double mutant Met272Val/Met316Phe, we

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**Table 1.** Amino acids constituting the xlCAR ligand-binding pocket.

<table>
<thead>
<tr>
<th>Location</th>
<th>Amino Acid</th>
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<tbody>
<tr>
<td>H1′</td>
<td>Phe171</td>
</tr>
<tr>
<td>H1–H3 Loop</td>
<td>Leu231</td>
</tr>
<tr>
<td>H3</td>
<td>His233, Phe234, Leu237, Ser238, Met241, Ile242</td>
</tr>
<tr>
<td>H5</td>
<td>Met272, Ala275, His276, Phe279</td>
</tr>
<tr>
<td>H5–S1 Loop</td>
<td>Tyr283</td>
</tr>
<tr>
<td>S1–S2 Loop</td>
<td>Asn288</td>
</tr>
<tr>
<td>S2</td>
<td>Phe290, Cys292</td>
</tr>
<tr>
<td>S3</td>
<td>His295, Phe297, Ser298</td>
</tr>
<tr>
<td>H6</td>
<td>Ile299, Asp301, Gly302, Ile304, Thr305</td>
</tr>
<tr>
<td>H6–H7 Loop</td>
<td>Phe307</td>
</tr>
<tr>
<td>H7</td>
<td>Tyr311, Leu312, Val315, Met316, Gln319</td>
</tr>
<tr>
<td>H10</td>
<td>Tyr399</td>
</tr>
<tr>
<td>H12</td>
<td>Met418</td>
</tr>
</tbody>
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Underlined residues provide hydrogen bond acceptor or donors interaction capabilities for ligands via backbone and/or side chain atoms.

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investigated xlCAR heterodimerization with RXRα, as well as recruitment of coactivator proteins SRC-1 and PGC-1α using mammalian two-hybrid assays. Neither the deactivating mutant Val315Ile, nor the non-effective His295Leu revealed any differences in RXRα heterodimerization to wild type receptor (Fig. 3B). Also, neither the two previously identified gain-of-function mutations Met272Val and Met316Phe nor the double mutant, revealed an altered interaction with RXRα. However, a substantial increase in RXRα interaction was observed for the Thr305Val mutant. Interestingly, also the non-effective Tyr311Phe mutation also showed improved heterodimerization (Fig. 3B). Subsequently, interaction studies with coactivators SRC-1 and PGC-1α were performed. In contrast to human CAR1, which showed a strong constitutive interaction with these coactivators [3], wild type xlCARα did not interact constitutively (Fig. 3C&D). A slight but statistically significant constitutive interaction with SRC-1 was only observed for mutant Thr305Val (Fig. 3C). All gain-of-function mutants revealed modest but statistically significant interaction with PGC-1α, with the exception of Met316Phe, where the difference to empty pVP16-AD did not reach statistical significance (P = 0.12) (Fig. 3D). With the exception of Met272Val, the enhanced interaction of these mutants with heterodimerization partner RXRα and coactivators SRC-1 and PGC-1α in the mammalian two-hybrid assays was most likely not an artifact of increased protein expression levels of the respective VP16-AD/ xlCARα-LBD fusion proteins, as all mutants, with the exception of Met272Val, showed protein expression equal to or even lower (Met316Phe) than wild type (Fig. S3). In contrast, the expression of the fusion protein of Met272Val appeared to be increased, which may explain its enhanced interaction with PGC-1α.

Figure 3. Basal activity of xlCARα mutants and their constitutive interaction with RXRα and coactivators. (A) LS174T cells were co-transfected with the DR4-TK promoter reporter gene plasmid and expression plasmids encoding the indicated xlCARα mutants. Data are presented as % of normalized activity of wild type xlCARα, which was set as 100% and is indicated as a red line. Bars show means ± SEM (n=8 independent experiments). Statistically significant differences to wild type xlCARα, as determined by one-sample t-tests (corrected for multiple testing by the method of Bonferroni) are indicated by asterisks. (B–D) Mammalian two-hybrid assays were used to analyze the interaction of xlCARα mutants with RXRα (B), SRC-1 (C) and PGC-1α (D). To this end COS7 cells were co-transfected with the GAL4-dependent pGL3SG reporter, and expression plasmids encoding fusions of VP16-AD and the respective LBD of the indicated xlCARα mutants together with expression plasmids encoding fusions of GAL4-DBD/RXRα-LBD (B), or GAL4-DBD/SRC1-RID (C), or GAL4-DBD/PGC-1α-RID (D). For (B) bars show means ± SEM (n = 6 independent experiments), with activity of cells transfected with empty vector pVP16-AD set as 1. Statistically significant differences to the activity of wild type xlCARα, as identified by one-way ANOVA with Dunnett’s multiple comparisons test are indicated by asterisks. For (C) and (D) bars show means ± SEM (n = 9 independent experiments), with activity of cells transfected with the combination of both empty expression plasmids pVP16-AD and pM set as 1. Statistically significant differences to the activity of pVP16-AD and the respective GAL4-DBD/coactivator RID, as identified by two-way ANOVA with Dunnett’s multiple comparisons test, are indicated by asterisks. *, P<0.05; **, P<0.01; ***, P<0.001, no asterisks indicate no differences. Red vertical lines in C and D indicate the activity of the empty vector pVP16-AD.
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The significantly enhanced interaction of the Thr305Val mutant with both RXRα and PGC-1α prompted us to further investigate this mutation on a molecular level of detail. Thus, we intended to perform comparative molecular dynamics simulations of the wild type and Thr305Val mutant xlCARα protein model. However, due to the largely missing H1–H3 insert, interactions between the truncated H1–H3 region and the Thr305-carrying helix 6 may lead to artificial results. Therefore, we applied a reverse approach in which a human-to-frog mutation (Val232Thr) was introduced into the human CAR x-ray crystal structure (PDB code 1xvp). Analysis of the molecular dynamics trajectories revealed significant differences compared to simulations performed for the wild type receptor published recently [19]. Helices in close proximity of Thr232 were found to be less stable throughout the simulations compared to the wild type receptor. While in one simulation a complete unfolding of helix 6 was observed, a second and third simulation revealed larger rearrangements within the C-terminal region of the LBD (Fig. S4).

Evaluation of the H10/11–H12 region

X-ray crystal structures of human and mouse CAR possess short helices H11’ (also termed helix X) in between helices 10/11 and 12 which has been suggested to represent an important determinant of the high basal activity of mammalian CAR [9]. In contrast, available structures of human PXR show a loop at the corresponding location and our protein model predicts the same for xlCARα. We assessed the importance of length differences of the H10/11–H12 region in human and Xenopus CAR by deleting Asp408 and Ile409 from xlCARα. The shortening was accompanied by a further reduction in basal activity (Fig. 4). Conversely, the extension of the H10/11–H12 region of human CAR by inserting aspartate and isoleucine after Gln334 of human CAR reduced the basal activity by more than 50% (Fig. 4). Next, we aimed to clarify the existence of a helical structure in the H10/11–H12 region. Therefore, we mutated Asn413 in the constitutively active xlCARα double-mutant Met316Phe/Met272Val into the helix-breaking residue proline, in order to destroy any potentially existing helical fold in this region. As a proof of concept, we also mutated H11’ amino acid Ala338 of hsCAR into proline. While the constitutive activity of human CAR dropped significantly, that of the xlCARα double mutant did not change (Fig. 4).

Identification of ligand binding modes

In order to investigate the binding modes of structurally diverse xICARα agonists artemisinin, fenofibrate and 5β-pregnanediol-3,20-dione (in the following denoted as pregnanedione), three-dimensional structures of the compounds were docked into the ligand-binding pocket of the xICARα model using the docking program GOLD in combination with the ChemScore scoring function. In order to take ligand efficiency into account, docking scores were corrected as described previously [27]. For pregnanedione only a single binding mode was obtained while for both artemisinin and fenofibrate two slightly differing binding orientations within the LBP were found (data not shown). Due to the barrier between LBP and activation helix, formed by amino acids from helices H3, H5 and H10/11, none of the ligands directly interacted with residues of the activation helix.

Docking poses for artemisinin revealed lowest docking scores of all three ligands (Table 2). The compound is placed in the upper part of the LBP and contacts 10 amino acids (Table 3). Besides extensive van der Waals (vdW) contacts with amino acids of the ligand-binding pocket, the compound shares a hydrogen bond with the Tyr311 side chain (Fig. 5A). An alternative docking pose indicates a hydrogen bond shared with Gln319. Similar to human CAR, both binding modes indicate partial occupation of the S1 sub-pocket, which has been identified as an important area for potent receptor activation [19].

Docking poses for fenofibrate revealed significantly higher docking scores compared to artemisinin (Table 2). The compound interacts with 18 amino acids (Table 3) and occupies a large part of the LBP (Fig. 5B), including the S1 sub-pocket. Most docking poses were found to share a hydrogen bond with His295. Although the compound possesses further hydrogen bond acceptors, no additional H-bond formation with the receptor was observed.

Molecular docking calculations suggest the steroid pregnanedione to bind with highest affinity (Table 2). The ligand contacts 12 amino acids in the LBP via vDW interactions and in addition shares hydrogen bonds with amino acids His233, His295 and Tyr311 (Fig. 5C). The steroid scaffold stacks with the phenyl ring of Phe279 and the compound completely occupies the S1 sub-pocket.

Ligand binding affinity and ligand-dependent receptor activation

In order to validate the predicted binding mode exemplarily for the ligand pregnanedione, we performed CARLA assays in which the induction of the interaction of in vitro translated CAR proteins with bacterially expressed SRC-1 RID was used as a measure of ligand binding (Fig. 6A&B). Amino acids shown here to contact pregnanedione (Table 3) and to differ from their counterparts in human CAR were mutated towards the situation in the human receptor. Mutation of each individual amino acid, predicted to contact the ligand, reduced the affinity of in vitro ligand binding of the respective mutant receptor, shown by a shift of the dose response curves to the right (Fig. 6B). Especially Phe279Leu, His295Leu, Phe297Tyr and Tyr311Phe mutants demonstrated nearly complete loss of binding. The reduced ligand binding affinity of mutants Leu237Ile (hsCAR: Ile164), Phe297Leu (hsCAR: Leu206), Phe297Tyr (hsCAR: Tyr224) and Val315Ile (hsCAR: Leu242), in which the corresponding hsCAR amino acid residues have been shown to make hydrophobic contacts with pregnanedione [9], suggests that xICARα binds pregnanedione with higher affinity than hsCAR. CARLA performed with human CAR confirmed this assumption as no significant induction of the interaction with SRC-1 was observed (Fig. 6C). Furthermore, the mutants did not show any significant change in basal interaction with SRC-1 in the absence of ligand, as compared to wild type xICARα (Fig. 6D), thereby confirming the data obtained with the SRC-1 mammalian two hybrid interaction assay (Fig. 3C). In contrast, human CAR demonstrated a significantly higher level of basal interaction with SRC-1 than xICARα (Fig. 6D).

Next we analyzed whether the impaired ligand binding, which was observed in the SRC-1 CARLA assay, further translates into lower ligand-dependent induction of the respective receptor transactivation activity in reporter gene assays. Figure 7A shows that the dose response curves of fold induction by pregnanedione of six mutants, all contacting pregnanedione by vDW interactions, were shifted to the right, compared to wild type xICARα, even if three of them (Phe279Leu, Phe297Tyr, Val315Ile) caught up to wild type at the highest dose of 30 µM. In contrast, the dose response curves of His295Leu and Tyr311Phe mutants, both interacting with pregnanedione via hydrogen bonds, did not show clear dose-dependent differences as compared to wild type (Fig. 7B).
confirmed experimentally. Conversely, the importance of helix 11 activity of xlCAR human CAR suggest the salt bridge to be instable [20].

In order to expand our knowledge on these unusual CAR properties in X. laevis, we investigated the structural characteristics of the xlCAR\(x\) ligand-binding domain by constructing a homology model based on sequentially related human CAR and PXR x-ray crystal structures. The selection of our approach is well-founded in the findings of a previous study describing a model of the human CAR LBD utilizing crystallographic data for human VDR and PXR as modeling templates that has been shown to be in very good agreement with published x-ray crystal structures, in particular within the ligand-binding pocket [13].

On the basis of x-ray crystal structures, several structural features have been suggested as important for human CAR basal activity [9]. Besides LBP amino acids directly contacting helix 12, an additional helix (helix 11') in between helices 10/11 and 12 as well as a salt bridge formed by Lys195 and the C terminus of helix 12 were considered to contribute to the pronounced basal activity. While our model suggests the presence of LBP residues interacting with helix 12, it lacks a helix 11' like structure, a finding that was confirmed experimentally. Conversely, the importance of helix 11' for human CAR activity has been verified.

Although a salt bridge between Lys268 and the C terminus was observed in the protein model. Amino acid side chains in close proximity to Thr305 are either aromatic or aliphatic (Leu231, Phe297, Phe307, Ile406), thus the non-polar side chain of the mutant may provide better vdW interaction possibilities with closely neighbored amino acids on helices H5 (Met272Val), H6 (Thr305Val) and H7 (Met316Phe), whose mutation towards hsCAR revealed a substantial gain-of-function that reached or even exceeded basal activity levels measured for the human receptor.

In contrast to these mutations, humanizing Thr305 to the more hydrophobic amino acid valine revealed both reinforced RXX\(x\) heterodimerization and enhanced PGC-1\(x\) recruitment. Molecular dynamics simulations of a reversed mutation inserted into the hsCAR LBD suggested a partially instable LBD fold upon introduction of a polar amino acid. According to the protein model, Thr305 is located at the very C-terminal end of helix 6 with its polar side chain buried within the protein interior. Mutation into the corresponding human amino acid (Val232) may disrupt hydrogen bond formation with Asp301 on helix 6 which is observed in the protein model. Amino acid side chains in close proximity to Thr305 are either aromatic or aliphatic (Leu231, Phe234, Phe297, Phe307, Ile406), thus the non-polar side chain of the mutant may provide better vdW interaction possibilities with closely neighbored amino acids on helices 3, 6 and 10/11 and thereby stabilize the overall receptor fold.

Tyr311 is predicted to be located at the N terminus of helix 7 with its side chain pointing into the ligand-binding pocket, thus providing an anchor point for bound ligands via its hydroxyl group. As the surrounding amino acid side chains are either of aromatic or aliphatic nature, mutation into phenylalanine may ameliorate vdW contacts with closely neighbored residues and RXX\(x\) and coactivators SRC-1 and PGC-1\(x\) upon introduction of humanizing mutations was investigated. Including the previously reported results, we now have identified altogether three amino acids located on helices H5 [Met272Val], H6 [Thr305Val] and H7 (Met316Phe), whose mutation towards hsCAR revealed a substantial gain-of-function that reached or even exceeded basal activity levels measured for the human receptor.

Previously, based on luciferase activity data and the protein model, the structural basis for the increased constitutive activity had been suggested to rely on improved interactions with the activation helix and therefore better coactivator binding [Met272Val] or enhanced heterodimerization with RXX\(x\) (Met316Phe). While the two-hybrid data did not confirm increased interaction with RXX\(x\) of these mutants, they suggest significant basal interaction with the coactivator PGC-1\(x\), which was not observed for the wild type. However, the effects appeared to be small and in the case of Met272Val may result from an increased expression of the respective VP16-AD fusion protein. Therefore, the molecular basis for the significantly increased basal activity of Met272Val and Met316Phe remains unknown.

In contrast to these mutations, humanizing Thr305 to the more hydrophobic amino acid valine revealed both reinforced RXX\(x\) heterodimerization and enhanced PGC-1\(x\) recruitment. Molecular dynamics simulations of a reversed mutation inserted into the hsCAR LBD suggested a partially instable LBD fold upon introduction of a polar amino acid. According to the protein model, Thr305 is located at the very C-terminal end of helix 6 with its polar side chain buried within the protein interior. Mutation into the corresponding human amino acid (Val232) may disrupt hydrogen bond formation with Asp301 on helix 6 which is observed in the protein model. Amino acid side chains in close proximity to Thr305 are either aromatic or aliphatic (Leu231, Phe234, Phe297, Phe307, Ile406), thus the non-polar side chain of the mutant may provide better vdW interaction possibilities with closely neighbored amino acids on helices 3, 6 and 10/11 and thereby stabilize the overall receptor fold.
thereby stabilize the local fold of the receptor which includes the heterodimerization motif (helix 10/11), thus providing an explanation for the experimentally observed improvement in RXRα binding. The unaltered basal activity is probably due to the unchanged PGC-1α coactivator binding. Mutation of Val315 towards isoleucine may sterically slightly reorient the neighbored amino acid Tyr311. Using MD simulations, a mutation of the corresponding human CAR amino acid Phe238 to alanine has shown to result in reorientation of the closely neighbored Tyr326 (xlCARα: Tyr399) away from helix 12 into the LBP that may impair stabilization of the activation helix [12]. Therefore, loss of interactions between Tyr311 and Tyr399 may result in decreased basal activity. Of all mutants tested, Val315Ile revealed least interactions with RXRα and coactivator proteins. Although not statistically significant, the observed trend may explain the loss in basal activity.

Pregnanedione, one of the most potent known xlCARα agonists [3], is predicted to be tightly anchored within the LBP via several hydrogen bonds, thus providing a convincing explanation for its extraordinary induction potential. The binding mode suggests the compound to act as a cross-linking agent of several secondary structural elements within the LBD, namely helices 3 and 11 as well as the β-strand, thereby stabilizing the overall LBD fold which in turn facilitates heterodimerization with RXRα and coactivator binding. While pregnanedione is a highly potent xlCARα agonist, the compound revealed a modest 2-fold activation in human receptor [29]. Based on an x-ray crystal structure of human CAR a hydrogen bond between pregnanedione and His203 (His276 in xlCARα) has been reported [9]. However, molecular dynamics simulations did not reveal the presence of such an interaction [13]. Instead, the ligand was found to contact the receptor exclusively by vdW interactions and thus is expected to bind with significantly lower affinity to hsCAR than to xlCARα.

Both fenofibrate and artemisinin are less potent agonists, a result that was confirmed by molecular docking calculations. The docking poses suggest that both ligands share only a single hydrogen bond with the receptor and thus bind with lower affinity compared to pregnanedione. While the docked fenofibrate almost completely occupies the LBP and thereby forms many more vdW interactions with amino acids of the ligand-binding pocket than artemisinin, the better activation potential of fenofibrate can be explained by a higher receptor affinity.

Docking predictions were exemplarily confirmed for the pregnanedione by determining ligand binding affinities of receptor mutants in vitro using the SRC-1 CARLA assay. Mutation of any single amino acid predicted to contact pregnanedione resulted in markedly impaired ligand binding. Mutations of the two H-bonding amino acids His295 and Tyr311 were among the four most affected in the SRC-1 CARLA assay. In contrast, mutation of these two amino acid residues did not clearly affect ligand-dependent transactivation potential of the respective receptor mutants, which may indicate that they may still be able to recruit coactivators other than SRC-1. Furthermore, this demonstrates the limitations of indirect methods, such as CARLA, for the analysis of ligand binding affinity, which seems to be influenced by the coactivator used. The mutation of any other amino acid,
except His295 and Tyr311, contacting pregnanedione did also impair the ligand-dependent transactivation potential of the respective receptors, thereby further confirming molecular docking predictions as well as CARLA results.

His295 and Tyr311 each interact with a single ligand oxygen atom located at both ends of the steroid scaffold, thereby the mutation into the corresponding amino acid in human CAR may less impact ligand binding compared to mutations such as Phe297 with the phenyl side chains extensively contacting A, B and C ring of the compound. Moreover, a new hydrogen bond with His276, located in close proximity of Tyr311 might be formed. While in wild type receptor His276 interacts with pregnanedione only via vdW contacts, a mutated receptor may result in small rearrangements within the LBP, thereby allowing hydrogen bonding and neutralizing the loss of binding affinity resulting from removal of a hydrogen bonding partner. Docking pregnanedione into receptors carrying either the His295Leu or Tyr311Phe mutation revealed the ligand to be placed closer to His276 so that only a small imidazole reorientation is required to form a strong hydrogen bond with the ligand (data not shown).

Compared with all known human CAR splice variants, xICARα is most similar to splice variant 2 (SV2, UniProt identifier Q14994-8), which contains a five amino acid insert (sequence: APYLT) after Pro270, located on the H8–H9 loop [23]. In contrast to the SV1 isoform (UniProt identifier Q14994-2), often considered as the wild type splice variant, this isoform possesses a low basal activity and substantial responsiveness to CAR ligands [30]. Although hepatic mRNA levels of both isoforms have been shown to reach similar levels [31], [32], research has concentrated mainly on the SV1 isoform. The significant reduction of the xICARα ligand-dependent activation upon introduction of gain-of-function mutation suggests that the high activity of human CAR-SV1 masks the inducing effects of binding agonists which can be exposed by introducing just a single amino acid of the APYLT motif [33].

Considering also the structural information derived from our homology model as well as the accompanying assays, X. laevis CARα clearly more resembles the mammalian pregnane X receptor rather than mammalian CAR. Compared to human CAR (SV1 and SV2 isoforms), the ligand-binding pocket appears to be substantially larger and provides significantly more possibilities for establishing hydrogen bonds with bound ligands. Shaped similar to the hsPXR LBP, the xICARα LBP may also harbor larger and chemically more diverse ligands as human CAR. Previously, we have shown that SR12813 (MW 505 Da), initially identified as human PXR inducer [34], is a potent xICARα agonist [3]. By contrast, the compound does not induce human CAR activity [3, 35]. As predicted by the protein model and confirmed by mutational studies, the X. laevis constitutive androstane receptor does not possess a helical secondary structural element in between helices 10/11 and 12 as found in human CAR x-ray crystal structures of splice variant 1 (H119) but rather is expected to possess a PXR-like loop conformation. Moreover, our results experimentally confirm the important role of H119 in hsCAR basal activity.

Similar to PXR and also the related vitamin D receptor (VDR, NR1I1), xICARα possesses a long H1–H3 insert. However, this part of the LBD is not involved in the LBP formation as shown for PXR and therefore is not expected to affect the fold of helices 6 and 7 as observed in all PXR x-ray crystal structures [36].

### Table 3. Protein-ligand interactions between xICARα and docked agonists.

<table>
<thead>
<tr>
<th>Location</th>
<th>Artemisinin</th>
<th>Fenofibrate</th>
<th>Pregnanedione</th>
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<tr>
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<td>Gln177</td>
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<tr>
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For each agonist docked into the ligand-binding pocket of xICARα the contacting amino acids (distance criterion of 4.0 Å) and their location on secondary structural elements of the ligand-binding domain is given. Amino acids sharing hydrogen bonds with docked ligands are underlined.

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Figure 6. Constitutive interaction and ligand-dependent induction of the interaction of xlCARα mutants with coactivator SRC-1. CARLA assays were performed using bacterially expressed GST-SRC-1 RID protein and in vitro translated 35S-labeled xlCARα wild type or mutant proteins or human CAR1 protein (hsCAR1), respectively, in the presence of the indicated concentrations of pregnanedione or solvent only (DMSO). (A) The results of representative experiments with xlCARα wild type and mutant proteins are shown. (B,C) Dose response analysis of ligand-dependent induction of SRC-1 interaction of xlCARα mutants (B) and human CAR1 (C), with graphs showing means ± SD (n = 3 independent experiments) of fold induction by respective concentrations of pregnanedione. Activity in the presence of DMSO only was designated as 1. In the upper part of (C), a representative experiment with human CAR1 is shown. Data were analyzed by two-way ANOVA with Dunnett’s multiple comparisons test (B) or one sample t-test (C). Significant differences to wild type xlCARα are shown by asterisks. The boxed asterisks indicate significant difference for all mutants at the respective dose. (D) Columns show means ± SD (n = 3 independent experiments) of the constitutive interaction with SRC-1 in the absence of ligand. Data were analyzed by one-way ANOVA with Dunnett’s multiple comparisons test. Significant difference to wild type xlCARα is shown by asterisks. *** P<0.001.

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deletion of a homologous region from hsVDR and xlVDR had also no effect on the selectivity for bile acids of these receptors [37]. Therefore, the functional role of the H1–H3 insert in xlCAR3 still needs to be elucidated.

Taken together, our study provides the first in-depth view into the structural and functional characteristics of a non-mammalian xenosensor that suggests its structural and functional similarity to the mammalian pregnane X receptor using a combination of computational and biochemical methods. In addition, our approach proved to be a procedure for investigating the contribution of single amino acids to basal activity of human CAR. Due to the ligand promiscuity of the receptor and to its wide organ distribution, exogenous xlCAR agonists might be problematic as environmental pollution increases. Therefore, our study may represent a preliminary step for the development of an in silico assessment of chemicals for unwanted effects in amphibians.

Supporting Information

Figure S1 Sequence alignment of xlCARα template structure generated using CLUSTALW. Secondary structure prediction for xlCARα (H, helix; E, strand; C, coil) was performed using PSIPRED (http://bioinf.cs.ucl.ac.uk/psipred/). Predictions for single amino acids with a confidence level higher than 5 are highlighted in bold.

(TIF)

Figure S2 Equal expression of in vitro synthesized CAR proteins. Wild type and mutant xlCARα, and hsCAR1 proteins were labeled with 35S-methionine by in vitro transcription/translation using the respective expression plasmids. Aliquots of the reactions were analyzed by protein gel electrophoresis.

(TIF)

Figure S3 Expression levels of CAR-VP16 fusion proteins in COS 7 cells. Expression plasmids encoding fusion proteins of VP16-AD with wild type or mutant xlCARα or hsCAR LBD used in the mammalian two-hybrid assays were cotransfected with a Renilla luciferase reporter gene plasmid. Protein expression was analyzed in Western blots with anti-VP16 antibody and normalized for transfection efficiency via Renilla luciferase activity. The upper panel shows the respective quantification of VP16-AD/CAR-LBD fusion protein expression as mean ± SEM of 2 independent experiments. The two respective Western Blots are shown in a) and b).

(TIF)

Figure S4 Conformational rearrangements within the ligand-binding domain of hsCAR upon the introduction of a Val232Thr mutation. Structures of wildtype (mauve) and mutant (yellow) receptor emerging after 50 ns of 3 different unconstrained molecular dynamics simulations are superimposed on the hsCAR x-ray crystal structure (PDB code 1xvp, colored in green). For the sake of clarity the structures are shown in ribbon representation and large parts of the LBD are excluded. The side chain of Val232 of the x-ray crystal structure is shown explicitly (green).

(TIF)

Table S1 Primers used for generation of point mutations and deletions within the hsCAR, xlCARα and hsPXR ligand-binding domains.

(DOCX)

Author Contributions

Conceived and designed the experiments: MM CN OB LW BW. Performed the experiments: MM CN OB UG BW. Analyzed the data: MM CN OB UG HH BW. Contributed reagents/materials/analysis tools: OB LW BW. Wrote the paper: OB HH LW BW.

References