

X-ray structure of the orphan nuclear receptor ROR β ligand-binding domain in the active conformation

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The retinoic acid-related orphan receptor β (ROR β) exhibits a highly restricted neuronal-specific expression pattern in brain, retina and pineal gland. So far, neither a natural ROR β target gene nor a functional ligand have been identified, and the physiological role of the receptor is not well understood. We present the crystal structure of the ligand-binding domain (LBD) of ROR β containing a bound stearate ligand and complexed with a coactivator peptide. In the crystal, the monomeric LBD adopts the canonical agonist-bound form. The fatty acid ligand–coactivator peptide combined action stabilizes the transcriptionally active conformation. The large ligand-binding pocket is strictly hydrophobic on the AF-2 side and more polar on the β -sheet side where the carboxylate group of the ligand binds. Site-directed mutagenesis experiments validate the significance of the present structure. Homology modeling of the other isotypes will help to design isotype-selective agonists and antagonists that can be used to characterize the physiological functions of RORs. In addition, our crystallization strategy can be extended to other orphan nuclear receptors, providing a powerful tool to delineate their functions.

Keywords: crystal structure/nuclear receptor/orphan/ROR/RZR

Introduction

Nuclear receptors (NRs) form a superfamily of sequence-specific transcription factors that regulate diverse biological processes including cell growth and differentiation, development, homeostasis and various organ functions in the adult by stimulating or repressing target gene expression (Gronemeyer and Laudet, 1995; Mangelsdorf *et al.*, 1995). NRs all share a common modular structure composed of several domains denoted A–F. Receptor dimerization, ligand binding, repression in the absence of

ligand and ligand-dependent transactivation are mediated by the C-terminal region of NRs, termed the ligand-binding domain (LBD), by generating the proper interaction surfaces for multiple partners, including corepressors, coactivators and mediators. The surface to which coactivators and mediators bind is assembled upon ligand binding and comprises H3, H4 and H12; it corresponds to the AF-2, the ligand-dependent transactivation function (reviewed in Renaud and Moras, 2000). In fact, ligand binding appears to trigger a switch in the LBD from a corepressor-binding to a coactivator-binding conformation (Glass and Rosenfeld, 2000; Renaud *et al.*, 2000).

In addition to the ligand-dependent receptors, a vast number of structurally related gene products are described for which no ligands have yet been identified and, therefore, they are referred to as orphan nuclear receptors (Willy and Mangelsdorf, 1998; Giguère, 1999). The retinoic acid-related orphan receptor β [ROR β ; NR1F2 (Nuclear Receptors Nomenclature Committee, 1999)], also called retinoid Z receptor β (RZR β), is an orphan member of family 1, which contains receptors such as RAR or TR. So far, three ROR isotypes, α , β and γ , have been described. Both ROR α (NR1F1) and ROR γ (NR1F3) are expressed in various tissues (Hirose *et al.*, 1994; Matysiak-Scholze and Nehls, 1997; Koibuchi and Chin, 1998) and seem to be involved in cerebellum development, immune responses (Delerive *et al.*, 2001), lymph node organogenesis and apoptosis during thymopoiesis (Kurebayashi *et al.*, 2000), bone metabolism (Meyer *et al.*, 2000) and adipocyte differentiation (Kurebayashi and Hirose, 1998). In contrast, ROR β is expressed exclusively in areas of the central nervous system (CNS) that are involved in the processing of sensory information, including spinal cord, thalamus and cerebellar cortices, and also the three principal anatomical components of the mammalian timing system, the suprachiasmatic nuclei, the retina and the pineal gland (André *et al.*, 1998). Therefore, it seems that this orphan NR regulates genes whose products play important roles in the context of sensory input integration as well as in the context of the biological clock. ROR β knockout mice exhibit a behavioral phenotype with similarities to a phenotype described some 40 years ago for a spontaneous mouse mutation called *vacillans* (Sirlin, 1956). These mice display a duck-like gait, transient male incapability to sexually reproduce and a severely disorganized retina that suffers from post-natal degeneration. Biochemical analyses indicated that ROR β can bind as a monomer to hormone response elements formed by the extended half-site sequence motif AnnTAGGTCA and activate reporter genes containing multiple copies of this half-site motif (Greiner *et al.*, 1996). However, in spite of the simplicity of the extended half-site sequence motif, no natural target gene regulated by ROR β could be identified up to now. In addition, ROR β

is classified as an orphan receptor, and the lack of a putative ligand has complicated the identification of physiologically relevant targets further. To understand better the role of ROR β in physiology, i.e. regulation of neuronal gene expression, the identification of specific ligands is of utmost importance. To gain insight into the geometry of the potential ligand-binding pocket (LBP) of ROR β and the nature of putative ligands, we concentrated on elucidating the crystal structure of this NR.

Here we present the 1.9 Å crystal structure of a complex between the LBD of the rat ROR β , a fortuitous ligand (stearate) and a peptide from the NR-interacting domain of the coactivator SRC-1 (Onate *et al.*, 1995), a member of the p160 coactivator family (Torchia *et al.*, 1998). The atomic level description of this orphan NR in the active conformation, stabilized by the combined action of the coactivator peptide and the pseudo-ligand, provides an accurate image of the LBP. This information greatly spurs the ability and the rationale for the design of isotype-specific agonists and antagonists that could be used to characterize and modulate the physiological functions of ROR β .

Results

ROR β LBD structure determination

Initially, the ROR β LBD (residues 201–459) was overproduced in *Escherichia coli*. However, the protein stability was strongly affected by oxidation problems, probably due to the presence of nine cysteines. Homology modeling of the RAR γ LBD, the closest homolog with known structure (28% identity, 56% similarity), bound to all-*trans* retinoic acid (ATRA) (Renaud *et al.*, 1995) shows that five cysteines are buried and four are solvent-exposed. Noticeably, the C-terminus contains two cysteines at positions 454 and 458 (after the AF2-AD) that are not conserved in ROR α and γ (Figure 1), and thus are probably involved in ROR β -specific regulation but not in the proper folding and ligand-dependent activity of the LBD. As these two cysteines are probably the most accessible ones and thus the most sensitive to oxidation, we chose to remove them by truncation. The minimal LBD (residues 201–452), overproduced in *E.coli* and purified to homogeneity, proved to be stable towards oxidation. Moreover, it is still transcriptionally active when fused to the GAL4 DNA-binding domain (data not shown).

During the purification process, the presence of a fortuitous ligand was found, which was shown by mass spectrometry to be stearic acid (N.Potier, personal communication). However, crystallization trials in the presence of an excess of stearic acid were unsuccessful. Crystals could only be obtained in the presence of a peptide from the coactivator SRC1 containing the LXXLL motif that was shown to be necessary and sufficient to bind NR LBDs (Heery *et al.*, 1997; Torchia *et al.*, 1997). Crystals of the ROR β LBD–SRC1 peptide complex grew in 1 week from a polyethylene glycol (PEG) 6000 solution without added stearic acid. The structure was solved by molecular replacement using the RAR γ holo-LBD (Renaud *et al.*, 1995) as a search model (see Materials and methods for details). The final R_{cryst} and R_{free} were 22.4% and 24.9%, respectively (see Table I). The

Table I. Data collection and refinement statistics

Data collection	
Resolution (Å)	30.0–1.85 (1.92–1.85)
Unique reflections	28 901
Completeness	99.9% (99.8%)
R_{sym}^a	3.6% (23.1%)
Multiplicity	4.2
Refinement	
Resolution (Å)	30.0–1.9
Reflections used	23 854
Completeness	100%
R_{cryst}^b	22.4%
R_{free}^c	24.9%
R.m.s.d. on bond length (Å)	0.008
R.m.s.d. on bond angles (°)	1.282
Non-hydrogen atoms	
Protein	1977
Peptide	91
Ligand	20
Water molecules	137
Average <i>B</i> factor for non-hydrogen atoms (Å ²)	
Protein	29.1
Peptide	37.1
Ligand	47.9
Water molecules	39.8

In the data collection, the last shell values are presented in parentheses.

^a $R_{\text{sym}}(I) = \sum_{\text{hkl}} \sum_i |I_{\text{hkl},i} - \langle I_{\text{hkl}} \rangle| / \sum_{\text{hkl}} \sum_i I_{\text{hkl},i}$, where $\langle I_{\text{hkl}} \rangle$ is the average intensity of the multiple $I_{\text{hkl},i}$ observations for symmetry-related reflections.

^b $R_{\text{cryst}} = \sum_{\text{hkl}} |F_{\text{obs}} - F_{\text{calc}}| / \sum_{\text{hkl}} |F_{\text{obs}}|$.

^c $R_{\text{free}} = \sum_{\text{hkl} \in T} |F_{\text{obs}} - F_{\text{calc}}| / \sum_{\text{hkl} \in T} |F_{\text{obs}}|$, where the T set (10% of reflections) is omitted in the refinement.

Ramachandran plot showed only one outlier, a residue located in loop 9–10, which is partially disordered according to the $2F_c - F_o$ electron density map.

Overall structure of the ROR β LBD

The ROR β LBD (Figure 2) presents the canonical fold for the NRs (Wurtz *et al.*, 1996) with two additional α -helices, H2' and H11'. It is in the agonist-bound state, H12 joining the H3–H4 region to form the proper interaction surface (the complete AF-2) for the coactivator (reviewed in Renaud and Moras, 2000). An additional H2' helix is also found between H2 and H3 in peroxisome proliferator-activated receptors (PPARs; Nolte *et al.*, 1998). H2 helices in PPAR and ROR are roughly superposed, but H2' helices are almost perpendicular, pointing directly toward the N-terminus of H3 in the case of ROR and toward the solvent parallel to the β -sheet in the case of PPAR. H11' is unique to ROR β among known LBD structures; it roughly superposes with the middle part of loop 11–12 of RAR γ . PROCHECK analysis (Laskowski *et al.*, 1993) indicates that the dimerization helix H10 is kinked at Ala411–Lys412, which should affect the homo- or heterodimerization of ROR β . Noticeably, there is a one amino acid deletion at the corresponding position in ROR γ (Figure 1).

The RAR γ and ROR β LBDs were superposed using the LSQ options from the program O (Jones *et al.*, 1991) (Figure 3A). The root mean square deviation (r.m.s.d.) was 1.2 Å on 148 C α s using a cut-off of 2.5 Å; the aligned regions comprise H1 (second half), H3 (second half)–H5,

	201	211	221	231	241	251
dmHR3	PEFISHADGD	INDVLIKTLA	EAHANT.NTK	LEAVHDMFRK	QPDVSRILYY	KNLQGEELWL
hRORg	TPEAPYASLT	EIEHLVQSVK	KSYRETCQLR	LEDLLRQRSN	IFSREEVTPGY	QRKSMWEMWE
mRORg	APEVPYASLT	DIEYLVQNVK	KSFRETCQLR	LEDLLRQRTN	LFSREEVTSY	QRKSMWEMWE
hRORa	GETSPVTVMA	ELEHLAQNIS	KSHLETCQYL	REELQQITWQ	TFLQEEIENY	QNKQREVWQ
mRORa	GETSPVTVMA	ELEHLAQNIS	KSHLETCQYL	REELQQITWQ	TFLQEEIENY	QNKQREVWQ
hRORb	GQLAPGITMT	EIDRIAQNII	KSHLETCQYT	MEELHQLAWQ	THTYEEIKAY	QSKSREALWQ
rRORb	GQLAPGITMS	EIDRIAQNII	KSHLETCQYT	MEELHQLAWQ	THTYEEIKAY	QSKSREALWQ
		H1		H2		H2'
hRARg	...DSYELSP	OLEELITKVS	KAHOETFP	PSLCQLGK.	.YTTNSSADH
						RVQLDLGLWD
	261	271	281	291	301	311
dmHR3	DCAEKITQMI	QNIIEFAKLI	PGFMRLSQDD	QILLLLKTGS	ELATVRMSRL	LDLSQNAVLV
hRORg	RCAHITTEAI	QYVVEFAKRL	SGFMELCQND	QIVLLKAGA	EVVLRMCRA	YNADNRTVVF
mRORg	RCAHITTEAI	QYVVEFAKRL	SGFMELCQND	QIVLLKAGA	EVVLRMCRA	YNADNRTVVF
hRORa	LCAITTEAI	QYVVEFAKRI	DGFMELCQND	QIVLLKAGSL	EVVLRMCRA	FDSQNNITVVF
mRORa	LCAITTEAI	QYVVEFAKRI	DGFMELCQND	QIVLLKAGSL	EVVLRMCRA	FDSQNNITVVF
hRORb	QCAIQITTHAI	QYVVEFAKRI	TGFMELCQND	QIVLLKAGSL	EVVLRMCRA	FNPLNNTVLF
rRORb	QCAIQITTHAI	QYVVEFAKRI	TGFMELCQND	QIVLLKAGSL	EVVLRMCRA	FNPLNNTVLF
		* H3		H4	* H5** *	S1 *
hRARg	KFSELA	TKIKIVEFAKRL	PGFTGLSIAD	QITLLKAA	CL	DILMLRICTR
		* * *				** ** *
	321	330	338	348	358	368
dmHR3	.GDVMLPQEA	FYTSDSSEMR	LVSRFQTA	K SIAELKLTET	ELALYQSLVL	LWPERNGVRG
hRORg	.BGKYGMEL	FRALGC..SE	LSSPDFSH	SLSALHFS	ED EIALY TALVL	INAHRPLQ
mRORg	.BGKYGMEL	FRALGC..SE	LSSPDFSH	SLSALHFS	ED EIALY TALVL	INAHRPLQ
hRORa	.DGKYASPDV	FKSLGC..ED	TSFPEFGK	SLCSMHLT	ED EIALFSAPVL	MSADRSWLQ
mRORa	.DGKYASPDV	FKSLGC..ED	TSFPEFGK	SLCSMHLT	ED EIALFSAPVL	MSADRSWLQ
hRORb	.BGKYGMOM	FKALGS..DD	LVNEAPDFAK	NLC SLQLTEE	EIALFSSAVL	ISPDRAWLIE
rRORb	.BGKYGMOM	FKALGS..DD	LVNEAPDFAK	NLC SLQLTEE	EIALFSSAVL	ISPDRAWLIE
	S2	H6*	* H7		H8	
hRARg	SDGLTLNR	TQ MHNAGFGP..	LTDLVFAFAG	QLLPLEMDDT	ETGLLSAICL	ICGDRMDLEE
	*	→	*			*
	378	388	398	404	413	423
dmHR3	NTEIQRLFNL	SMNAIRQELE	TNHAPLKGDV	T.VLDTLLNN	IPNFRD	SIL HMESLSKFKL
hRORg	KRKVEQLQYN	LLEAFHHHLC	KTHR....Q	S.ILAKLPPK	.GKLRSCSQ	HVERLQIFQH
mRORg	KRKVEQLQYN	LLEAFHHHLC	KTHR....Q	G.LLAKLPPK	.GKLRSCSQ	HVEKLIQIFQH
hRORa	KVKIEKLQOK	IQLALQHVLO	KNHR....ED	G.ILTKLICK	VSTLRACGR	HTEKLMAFKA
mRORa	KVKIEKLQOK	IQLALQHVLO	KNHR....ED	G.ILTKLICK	VSTLRACGR	HTEKLMAFKA
hRORb	PRKVQKLQEK	IYFALQHVIO	KNHL....DD	E.TLAKLIAK	IPTITAVCNL	HGEKLVQFKQ
rRORb	PRKVQKLQEK	IYFALQHVIO	KNHL....DD	E.TLAKLIAK	IPTITAVCNL	HGEKLVQFKQ
		H9		H10		H11
hRARg	PEKVDKLOEP	LLEALRLYAR	RRRP....SQ	PYMFPRMLMK	ITDLRGISTK	GAERAITLKM
						* *
	433	443	453			
dmHR3	QHPNVV...F	PALYKELFSI	DSQQDLT...			
hRORg	LHPIVVQAAF	PPLYKELFST	ETESVPGCPS			
mRORg	LHPIVVQAAF	PPLYKELFST	DVESPEGLSK			
hRORa	IYPDIVRLHF	PPLYKELFST	EFEPAMQIDG			
mRORa	IYPDIVRLHF	PPLYKELFST	EFEPAMQIDG			
hRORb	SHPEIVNTLF	PPLYKELFNP	DCATACK...			
rRORb	SHPEIVNTLF	PPLYKELFNP	DCATACK...			
		H11'	H12			
hRARg	EIPG....PM	PPLIREMLEN	PEMFEDSSQ			

Fig. 1. Sequence alignment of ROR LBDs with hRAR γ LBD. The LBD sequences of hRAR γ (Krust *et al.*, 1989), hROR γ (Hirose *et al.*, 1994), mROR γ , hROR α (Giguère *et al.*, 1994), mROR α (Carlberg *et al.*, 1994), hROR β (Carlberg *et al.*, 1994), rROR β (Carlberg *et al.*, 1994) and the *Drosophila* homolog DHR3 (Koelle *et al.*, 1992) were aligned using the pileup option of the Genetics Computer Group (GCG, 1994). The secondary structure elements of the rROR β LBD (present study) and the hRAR γ LBD (Renaud *et al.*, 1995) are underlined with a plain line (α -helices) or with an arrow (β -strands). The residues involved in the LBP are shown in red; those of rROR β in close contact with stearate and those of hRAR γ in close contact with retinoate are indicated by an asterisk (4 Å cut-off). The residues whose mutation affects the shape of the LBP are colored in yellow (β -specific or β -like), green (α -specific), pink (γ -specific) or blue (DHR3-specific). The amino acid numbering is given for rROR β .

H7–H9, H10–H11 (first half) and H12. In ROR β , the LBP is shifted laterally toward H7 (Figure 3B). This is due to a different location of H6, which is shifted outwards and tilted downwards, loop 6–7 now lying at the LBD surface; on the other side of the pocket, H3 and s1, the first strand of the β -sheet, are shifted inwards. Secondly, the LBP is larger in the vertical direction, extending toward both the top and the bottom. At the top, the LBP is limited by H5. In RAR γ , M272 from this helix points towards the center of the pocket. In ROR β , the side chain of the corresponding residue L304 lies aside. At the bottom, the pocket in RAR γ

is limited by F230 (H3), F288 (s1) and F304 (loop 6–7). In ROR β , (i) the corresponding C262 in H3 points downwards rather than upwards as F230 in RAR γ ; (ii) due to the rotation of the plane of the β -turn with strand s1 being more inside the LBP, F320 in ROR β lies ~5 Å lower than the corresponding F288 in RAR γ ; and, finally, (iii) loop 6–7 in ROR β is located at the LBD surface and does not contribute to the floor of the pocket, which is now limited on this side by F330 from H6, ~4 Å lower than F304 from loop 6–7 in RAR γ . Globally, the volume of the LBP is much larger in ROR β (766 Å³) than in RAR γ (429 Å³).

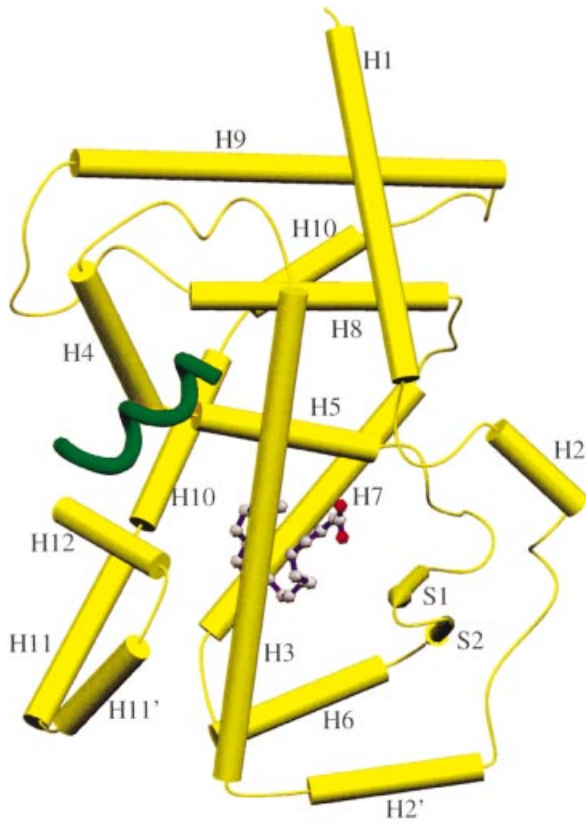


Fig. 2. Schematic representation of the rROR β LBD in complex with stearate (ball-and-stick) and a SRC-1 peptide (ribbon representation). The kink in H10 has been emphasized by breaking H10 into two segments.

A transcriptionally active LBD conformation

The LBD is in the canonical, transcriptionally active conformation, and the coactivator peptide binds the AF-2 surface as described previously for other NR-coactivator peptide complexes (Darimont *et al.*, 1998; Nolte *et al.*, 1998; Shiau *et al.*, 1998). The LXXLL motif-containing peptide used for crystallization corresponds to the second NR-box from the p160 coactivator SRC-1 (residues 686–700) (Onate *et al.*, 1995; Heery *et al.*, 1997; Torchia *et al.*, 1997). The following residues of the peptide are seen in the crystal structure: HKILHRLQEQ. The LXXLL motif forms the hydrophobic face of an amphipathic α -helix interacting with a hydrophobic cleft on the LBD surface. In particular, the side chains of L313 and L314 make van der Waals contacts with V274 (H3), and I292 and L295 (H4) from the ROR β LBD.

The side chain carboxylate of the conserved E448 (H12) known to be important for transactivation (see mutant studies below) forms hydrogen bonds with the backbone amide nitrogens of I309 and L310 at the N-terminus of the peptide helix. At the other end, the main chain oxygen atoms of L313 and L314 are hydrogen-bonded to K278 (H3) from the ROR β LBD. These ‘capping interactions’ are similar to those already described for other NRs (Darimont *et al.*, 1998; Nolte *et al.*, 1998; Shiau *et al.*, 1998).

The ligand-binding pocket

Compared with the human ortholog, the rat ROR β LBD sequence differs at only three amino acid positions: S210T

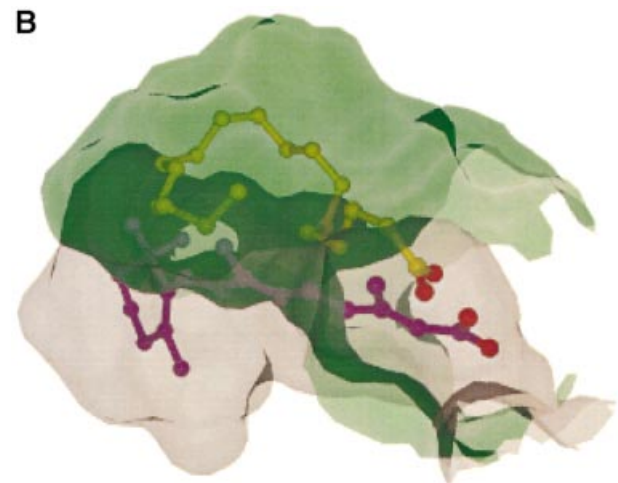
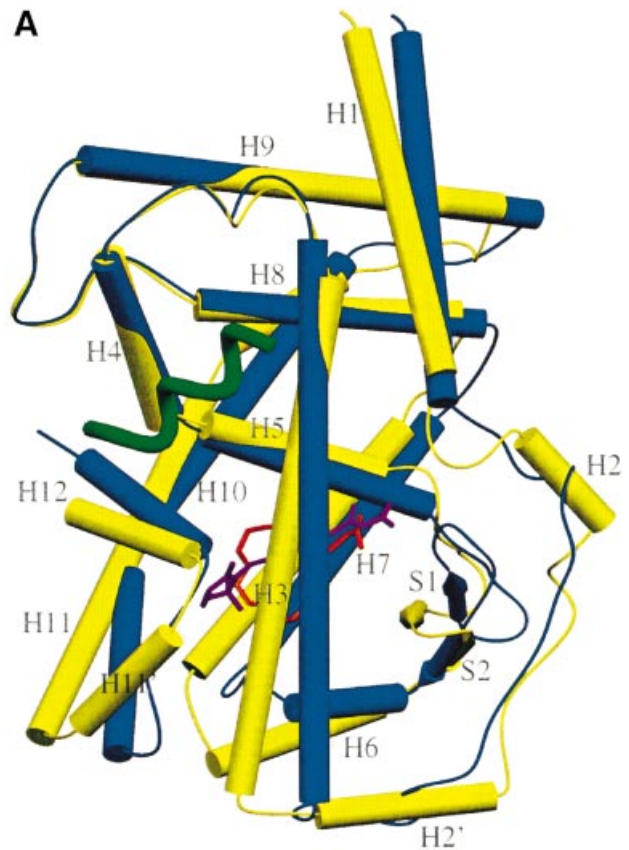


Fig. 3. (A) Backbone superposition of the structures of the rROR β LBD (yellow) in complex with stearate (orange) and the SRC1 peptide (green) and of the hRAR γ LBD (blue) in complex with retinoate (purple). The superposition was done using the LSQ option of O (Jones *et al.*, 1991). The r.m.s.d. was 1.2 Å for 148 matched C α s. (B) Superposition of the probe-occupied cavities of rROR β and hRAR γ (view from the top) calculated by MSMS with a probe radius of 1.4 Å. This figure was prepared with DINO (Philippsen, 1999).

(beginning of H1), L376I (loop H7–H8) and D436E (H11'). A456T and V457A are located at the C-terminus after H12 and are not present in our construct. None of these residues are part of the LBP; therefore, the present analysis can be extrapolated to the human receptor.

The probe-occupied volume of the LBP (766 Å³) is larger than that of the vitamin D receptor (VDR) (660 Å³;

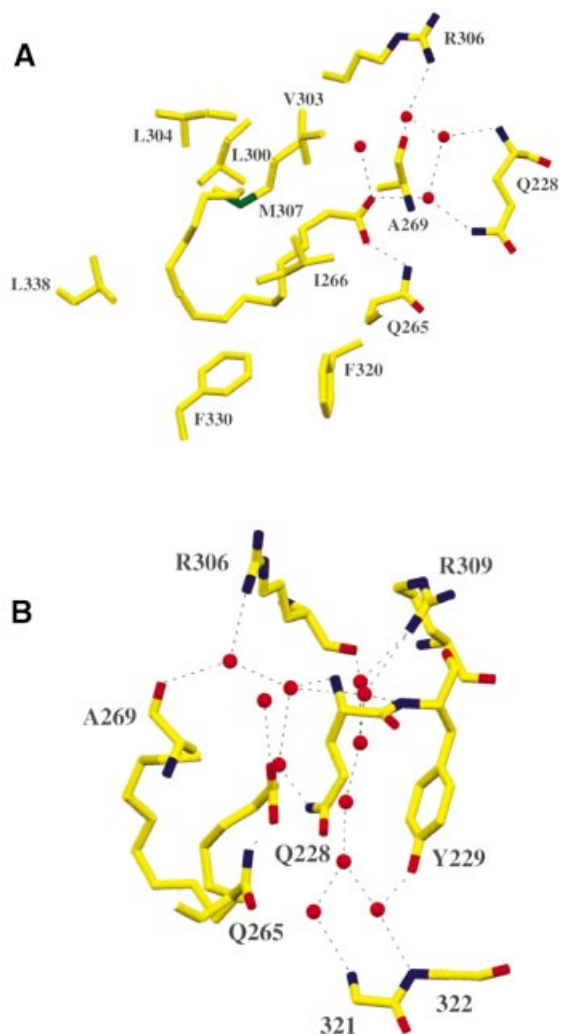


Fig. 4. (A) Detailed view of the LBP showing the stearate ligand and the residues in close van der Waals contact (4.0 Å cut-off) or interacting through hydrogen bonds. (B) Another view, emphasizing the second layer of water molecules forming a channel.

Rochel *et al.*, 2000) and ranks third among the LBDs of known structure, after PPAR (1400 Å³; Nolte *et al.*, 1998) and PXR (1150 Å³; Watkins *et al.*, 2001). The LBP is essentially hydrophobic on the AF-2 side (H5 N-terminus, H6, H7, H10, H12) and more polar on the H3 side (L1–2, H3, H5 C-terminus). Stearic acid was co-purified fortuitously from *E.coli* and co-crystallized with the heterologously expressed RORβ LBD. The hydrophobic side of the pocket is partially filled up with the aliphatic chain of stearic acid, while the polar side is occupied by the carboxylate group of the fatty acid and 11 ordered water molecules (Figure 4). Upon binding, stearate adopts a U-shaped conformation similar to that observed for oleate in the mutant RXRαF318A LBD (Bourguet *et al.*, 2000) and also for several fatty acids in fatty acid-binding proteins (Young *et al.*, 1994; Thompson *et al.*, 1997; Balendiran *et al.*, 2000). One oxygen atom of the carboxylate group forms a hydrogen bond with Q265 NE2. This residue differs in RORα and β. The other oxygen atom of the carboxylate group is hydrogen-bonded through water molecules to other conserved residues among RORα and

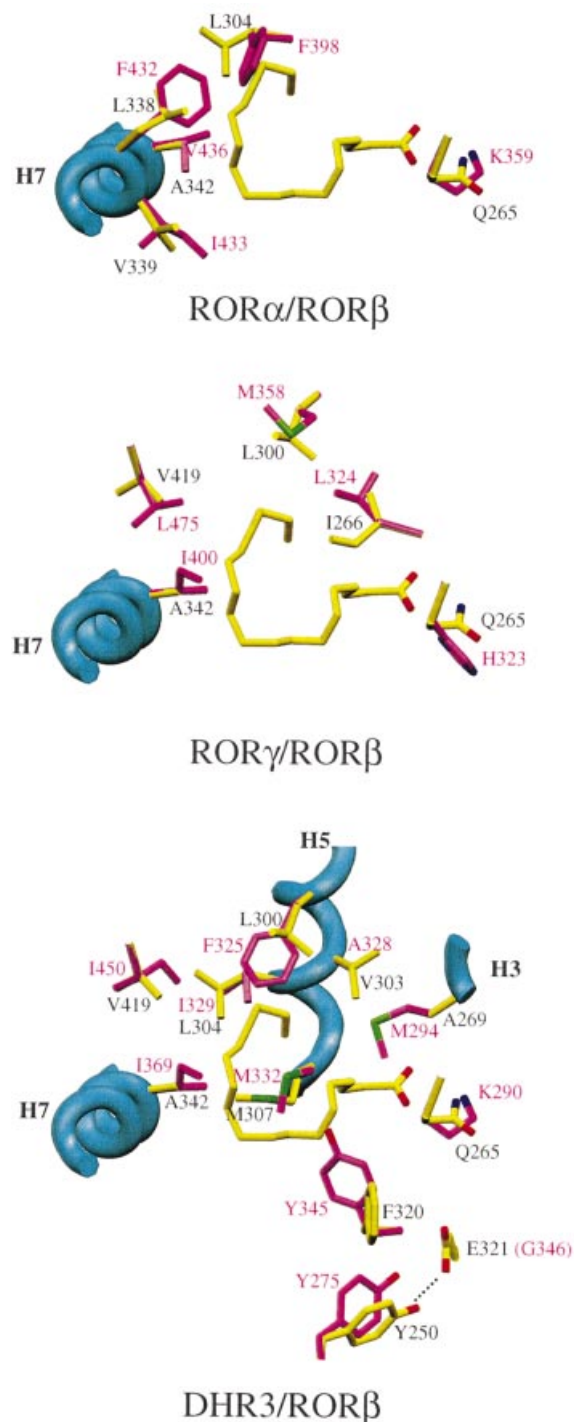


Fig. 5. Homology modeling of RORα, RORγ and DHR3 LBDs, showing the non-conserved residues affecting the pocket's shape. Figures 2, 3A, 4 and 5 were prepared with SETOR (Evans, 1993).

β of the LBP, namely Q228 and R306, and to the carbonyl group of V303 (Figure 4). The higher average *B*-value for the ligand (48 Å² compared with 29 Å² for the protein) suggests that the fatty acid thus probably adopts multiple low-energy conformations inside the pocket, in good agreement with the few distant van der Waals contacts between the aliphatic chain and the pocket. Indeed, the electron density map is less well defined in the middle portion of the chain.

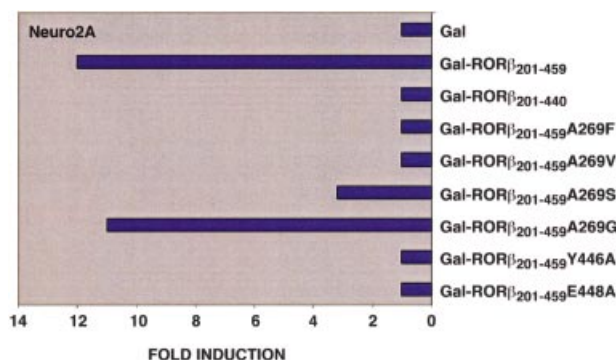


Fig. 6. Histogram showing the results of the transactivation assays.

Isotype variability

The amino acid sequence conservation among ROR LBDs is not very high, ROR β being 61% identical (74% similar) to ROR α and 49% identical (67% similar) to ROR γ ; it is also 33% identical (49% similar) to DHR3 (NR1F4), the *Drosophila* homolog of RORs. Moreover, only 17 out of the 32 residues whose side chain contributes to the LBP are strictly conserved in the three isotypes (Figure 1), raising the possibility of different ligands.

Homology models for the hROR α , hROR γ and DHR3 LBDs were built using the present rROR β structure as a template. The hROR α LBP is smaller on the hydrophobic side (toward H7) due to a cluster of four substitutions for bulkier side chains: hROR α F398/rROR β L304, F432/L338, I433/V339 and V436/A342 (Figure 5A). Accordingly, the volume of the pocket is 568 Å³, compared with 766 Å³ for rROR β . This suggests that the natural ligand for the α and β isotypes could be different. The hROR γ LBP is more similar in size (705 Å³) but exhibits a slightly different shape due to four conservative substitutions—hROR γ L324/rROR β I266 and M358/L300—which slightly enlarge the cavity on one side, and I400/A342 and L475/V419 that decrease the size of the cavity on another side (Figure 5B). Strikingly, the homology model of the DHR3 LBD displays a much smaller pocket (221 Å³) due to: (i) a few substitutions for bulkier side chains: DHR3 M294/rROR β A269, F325/L300, I329/L304 and I369/A342, which makes M332 protrude more into the pocket than the corresponding M307, and I450/V419; and (ii) a reorientation of Y275 and Y345 (Figure 5C).

In vivo mutational analysis of the LBD of ROR β

Prior to the determination of the crystal structure, we chose to model the ROR β LBD from the hRAR γ LBD structure in the agonist-bound form even though no natural or synthetic agonist was known. Homology modeling allowed us to emphasize some important residues in the activation helix H12 and in the putative LBP that were subjected to mutational analysis. The crystal structure now validates this choice.

To test whether ROR β might be a ligand-dependent NR and thus support the existence of an endogenous ligand, a first series of site-directed mutagenesis experiments was designed. A269 was chosen because its side chain points into the pocket. We replaced it by bulkier residues in order

to hinder the binding of a putative ligand. Increasing the size of the side chain decreased transactivation by Gal-ROR $\beta_{201-459}$ in Neuro2A cells (Figure 6) and other cell lines (HT22 and NIH 3T3). All mutants were expressed to similar levels in the tested cells (data not shown). In the A269S mutant, a partial transcriptional activity of ROR β is still observed, whereas both mutants with bulkier side chains, A269V and A269F, failed to transactivate the reporter gene at all. Additional reporter gene assays showed that the transcriptional activity of Gal-ROR $\beta_{201-459}$ A269G in various cell lines does not differ from that of Gal-ROR $\beta_{201-459}$. Altogether, these data suggest that transactivation by ROR β is ligand dependent and that the binding of the endogenous ligand is hampered by bulky side chains at position 269. Indeed, according to the crystal structure, mutation of A269 to a phenylalanine residue prevents the binding of stearate but should not affect the correct folding of the LBD.

Deletion of the AF2-AD domain in both the full-length receptor and the fusion protein Gal-ROR $\beta_{201-440}$ resulted in a complete loss of transcriptional activation by ROR β (Figure 6). Previous mutational analyses also demonstrated that the activation domains AF2-AD of ROR β and RAR γ are functionally interchangeable (Greiner *et al.* 1996). To test the functional importance of the ROR β AF2-AD, two residues located in the activation helix H12 were specifically mutated. Both mutants Gal-ROR $\beta_{201-459}$ E448A and Gal-ROR $\beta_{201-459}$ Y446A are transcriptionally inactive on a (GALp)₃-TK luciferase reporter in Neuro2A (Figure 6) and all other cell lines analyzed (data not shown). In the crystal structure, the side chain carboxylate of E448 is hydrogen-bonded to the coactivator peptide residues L310 and I309. The maintenance of this interaction is thus important for transactivation, most probably by acting in the recruitment of the coactivator. Y446 OH makes a hydrogen bond with H423 NE2 from H11. Moreover, the Y446 side chain is at the core of a highly conserved hydrophobic cluster comprising W259, A263 and I266 (H3), L427(H11), F442 (L11'-12) and F450 (H12). The Y446A mutation directly affects this cluster, probably causing a local collapse perturbing the coactivator-binding surface.

Discussion

The strategy developed in the present study in order to crystallize ROR β LBD without prior knowledge of a natural or synthetic ligand is applicable to other orphan NR LBDs. In our study, three major problems were overcome: (i) oxidation due to the presence of accessible cysteines not involved in the LBP or other critical contacts (in principle, exposed cysteines should be mutated unless located at potentially critical functional sites such as the LBP or its vicinity or interfaces; Gangloff *et al.*, 2001); (ii) stabilization of the LBD fold by a pseudo-ligand coming from the expression host or from a parallel search; and (iii) use of a coactivator peptide to stabilize the agonist-bound, transcriptionally active conformation.

LBD stabilization through the presence of a fortuitous ligand

Several structures of NR LBDs in complex with natural or synthetic ligands have been reported. In most cases (RAR,

TR, ER and VDR), the natural ligand was known and was co-crystallized with the LBD (Renaud *et al.*, 1995; Wagner *et al.*, 1995; Brzozowski *et al.*, 1997; Rochel *et al.*, 2000). In the case of PPARs, the natural ligands are not known with certainty, though it has been proposed that PPARs act as lipid sensors since they bind to and are activated by many low-affinity fatty acids and fatty acid derivatives; indeed, the PPAR β LBD was co-crystallized with eicosapentaenoic acid (Xu *et al.*, 1999). Besides, many PPAR isotype-specific ligands have been synthesized (Willson, 2000). Apo-LBDs are rather unstable in the absence of cofactors; indeed, up to now, crystals of the apo-LBD could only be obtained in the case of RXR α (Bourguet *et al.*, 1995), PPAR β (Xu *et al.*, 1999), PPAR γ (Nolte *et al.*, 1998) and PXR (Watkins *et al.*, 2001).

Two independent structural observations pointed to the unexpected ability of lipids to stabilize mainly hydrophobic LBPs. The first was the discovery of the presence of oleate in RXR α within the heterodimer RXR α -RAR α (Bourguet *et al.*, 2000). More recently, the crystal structure of an insect USP LBD was reported (Billas *et al.*, 2001), showing a very large, fortuitous ligand that co-purifies with the protein, a phospholipid of mass 745. In the present study, we observe a similar situation, where the pseudo-ligand stearate is co-purified from the expression host.

Use of a coactivator peptide as a tool for crystallization

The idea of masking the hydrophobic cleft between H3 and H4 in order to make the LBD more soluble and more amenable to crystallization originated from the observation that in all reported LBD crystal structures, the cleft is always occupied by an amphiphilic helix or molecule: (i) H12 of the same molecule in the known antagonist-bound complexes; (ii) H12 of a neighboring LBD in RXR and PPAR apo-LBDs; (iii) a detergent molecule in the case of RAR holo-LBD (Klaholz and Moras, 2000); or (iv) a coactivator peptide when added to the agonist-bound LBD complex (Darimont *et al.*, 1998; Nolte *et al.*, 1998; Shiau *et al.*, 1998). In fact, the addition of a high-affinity, cognate coactivator peptide removes a major constraint on crystallization. Another reason for using the coactivator peptide is the low affinity of stearate and its lack of agonistic capability in transactivation assays (data not shown), suggesting that this fortuitous ligand alone is not able to stabilize the active conformation. This reasoning is consistent with the equilibrium model where a coactivator can stabilize the active conformation of an NR LBD by shifting the equilibrium between the ligand-bound and unbound forms (Gangloff *et al.*, 2001; Steinmetz *et al.*, 2001). In addition, the present structure, showing that an SRC1 peptide can bind the ROR β LBD coactivator-binding site in the classical conformation, confirms the assumption of multiple possible combinations between NRs and coactivators. Any minimal LXXLL-containing helix should bind to the hydrophobic coactivator-binding site, the specific pairwise interaction being modulated by the flanking regions. In summary, the present structure argues strongly in favor of a classical coactivation mechanism for ROR β .

ROR β probably functions as a monomer

ROR α and ROR β have been proposed to function as both monomers and homodimers (Carlberg *et al.*, 1994; Giguère *et al.*, 1994). However, according to *in vitro* and *in vivo* data, ROR β is unable to form homodimers (Greiner *et al.*, 1996). ROR β binds to monomeric response elements formed by the extended half-site sequence motif AnnTAGGTCA but cannot transactivate reporter genes containing only a single copy of this motif (Greiner *et al.*, 1996). On the other hand, transactivation of ROR β -dependent reporter genes is only achieved with direct repeat binding sites (DR6–DR9) or by two binding sites oriented as inverted palindromes (P0). However, even for the transcriptionally active response elements, no cooperative binding is detected, which indicates that ROR β occupies both sites independently. Furthermore, in contrast to most members of family 1, ROR β does not heterodimerize with RXR (Greiner *et al.*, 1996). In all NR LBD homo- and heterodimer structures, the dimerization interface is topologically conserved, and for the residues in contact no significant conformational change upon dimerization is observed. A homodimer built upon this assumption generates important clashes at the level of H10: a steric one between the 2-fold related I410 and a repulsive interaction involving E386 from one LBD and E404 from the other. These contacts would be sufficient to destabilize a canonical homodimer and explain the biochemical observations.

Is stearate close to the human physiological ligand?

Over the past 10 years, it has been shown that fatty acids can act as signaling molecules in regulating gene expression (Duplus *et al.*, 2000). The mechanism by which fatty acids modulate gene transcription still remains largely unknown. The CNS contains large amounts of polyunsaturated fatty acids such as arachidonic acid (20:4 *n*-6) and docosahexaenoic acid (DHA) (22:6 *n*-3). The total synthesis of these two polyunsaturated fatty acids is not possible in mammals so that their precursors, linoleic acid (18:2 *n*-6) and α -linolenic acid (18:3 *n*-3), must be provided in the diet (Contreras *et al.*, 2000). DHA has been shown to be a ligand for RXR in mouse brain (de Urquiza *et al.*, 2000). However, two facts argue against stearate being close to the real ligand: (i) stearate does not activate ROR β in a cell reporter assay; and (ii) the low percentage of pocket occupancy by stearate (33%) and its partially disordered conformation. Thus, stearate most probably acts as a LBD stabilizer by filling the pocket, but it is unable on its own to generate sufficient interactions with the pocket to stabilize the active conformation, which is achieved by addition of the coactivator peptide. This conformational heterogeneity is reflected by the fact that LBD crystals could not be obtained in the absence of the peptide.

The low sequence conservation among the residues mapping the LBP and the large volume of the cavity (766 Å³) could suggest that the three isotypes do not share the same physiological ligand. Indeed, homology models display important differences at the LBP level, especially between ROR α and ROR β/γ , the ROR α pocket being 25% smaller. ROR β and ROR γ LBPs have similar volumes but their shape is slightly different.

Among the four polar residues interacting directly or indirectly with the stearate carboxylate group, only Gln265 varies. This amino acid is replaced by a lysine residue in ROR α and by a histidine residue in ROR γ . Both residues could bind the carboxylate group of stearate with slightly different geometries. Thus, two possibilities may exist: (i) the natural ligand is the same for the three isotypes and residue 265 is not involved in the anchoring of the ligand, explaining its lack of conservation; or (ii) different ligands exist and the difference of side chain at position 265 is crucial in the ligand specificity for the three isotypes. However, one cannot exclude the possibility of a common ligand accommodated with a subtle rearrangement of the side chains at position 265 to maintain the interaction, although this would affect the binding affinity.

Conclusion

Escherichia coli allows the production of large amounts of heterologous proteins but also contains biomolecules that can be trapped by the heterologously produced proteins. In the present study, stearate was found in the LBP of the orphan NR ROR β LBD. Both the presence of the fortuitous ligand and the addition of a peptide containing the LXXLL motif stabilize the active holo-LBD conformation, the low affinity of stearate for ROR β being compensated by the high-affinity binding of the coactivator peptide, which prevents alternative conformations. Despite the fact that stearate does not function as an ROR β agonist in transactivation assays, a longer fatty acid or related lipids endogenously present in neuronal cells may bind with higher affinity to the ROR β LBP. Such a ligand might explain the neuronal-specific activity of ROR β . The high concentration of fatty acids in neuronal tissues and their role in brain development support this idea. The question remains open of whether there is a unique ligand or a family of compounds that bind to the ROR β LBP and regulate transcription. Nevertheless, the structure of the ROR β LBD provides a detailed picture of its LBP. This knowledge will greatly enhance the ability to design agonist and antagonist molecules that can be used to characterize in detail the physiological functions of RORs. Such synthetic compounds will be powerful tools to investigate ROR signaling pathways without prior knowledge of the natural ligands. Taken together, we have developed an efficient strategy to crystallize the ROR β LBD. This strategy can be extended to orphan NRs in general and will thus provide a very useful tool to delineate novel functions of orphan NRs.

Materials and methods

Expression, purification and crystallization

The rat ROR β LBD (residues 201–459) was overproduced as a histidine-tagged protein in *E. coli* BL21(DE3) by using pET-15b vector (Novagen). Due to aggregation problems, rROR β was recloned as a truncated version (residues 201–452). Two non-conserved cysteines are deleted while maintaining the integrity of H12 (Figure 1). The purified monomer was concentrated to 6 mg/ml. A native gel reveals that the protein of interest migrates as a single species. The cells were grown in LBM at 37°C to an OD 0.6 and induced with 0.8 mM isopropyl- β -D-thiogalactopyranoside (IPTG). The incubation was continued at 16°C overnight. Cells from 1 l of culture were resuspended in 50 ml of 20 mM Tris-HCl pH 8.5, 100 mM NaCl, 10% glycerol, 2 mM CHAPS and 2 mM β -mercaptoethanol (buffer A), and sonicated. The lysate was centrifuged at 50 000 r.p.m. for 2 h at

4°C, and the supernatant was loaded on a 2 ml cobalt affinity column. The column was washed with 10 ml of buffer A and the protein was eluted with a gradient of 0–1 M imidazole in buffer A. Subsequent gel filtration was performed on a Superdex S-200 HiLoad 16/60 from Pharmacia using as elution buffer 20 mM Tris-HCl pH 8.5, 100 mM NaCl, 2 mM CHAPS and 5 mM dithiothreitol (DTT) (at 1 ml/min). The protein eluted at 87 ml and was estimated to be >95% pure and homogeneously by SDS-PAGE. Co-crystallization with a 3 M excess of SRC-1 NR-interacting peptide (686-RHKILHRLQLQEGSPS-700) was carried out with the hanging drop vapor diffusion method (2 μ l of LBD-peptide complex solution + 2 μ l of reservoir solution against 500 μ l of reservoir solution). A proprietary NR LBD screening kit (D.Zeyer, S.Duclaud, D.Moras and J.P.Renaud, unpublished results) allowed us to find preliminary crystallization conditions. In the refined conditions, crystals grow within 1 week at 22°C to a size of $\sim 110 \times 60 \times 30 \mu\text{m}$ with a reservoir of 100 mM Tris-HCl pH 8.0, 15% PEG 6000. Crystallization trials without the SRC-1 peptide were unsuccessful, even in the presence of an excess of stearic acid.

Data collection, structure determination and refinement

Crystals were cryoprotected by equilibration in 15% PEG 6000 at pH 8.0 containing 15% glycerol and then flash-frozen in liquid ethane at liquid nitrogen temperature. X-ray diffraction data were collected at liquid nitrogen temperature from a single frozen crystal at the ID14-3 beamline at the ESRF Grenoble, France. Crystals diffracted X-rays to a resolution limit of 1.9 Å. All data were integrated and scaled using DENZO and SCALEPACK (Otwinowski and Minor, 1997) (Table I). The space group is P212121, with unit cell parameters $a = 52.302 \text{ \AA}$, $b = 58.490 \text{ \AA}$, $c = 106.036 \text{ \AA}$, $\alpha = \beta = \gamma = 90^\circ$. There is one monomer per asymmetric unit and a solvent content of 52%. The estimated B -factor by Wilson plot is 29 Å². The structure was solved by molecular replacement using the program AMoRe (Navaza, 1994) and the RAR γ holo-LBD [Protein Data Bank (PDB) accession code 2lbd] as search model. The top solution had a correlation coefficient of 27.8% and an R -factor of 52.7% after AMoRe rigid-body refinement. Automated model building using ARP/wARP (Perrakis *et al.*, 1999) yielded three chains (243 residues, connectivity index 0.98). The partial model was subjected to alternating rounds of manual building using O (Jones *et al.*, 1991) and refinement using CNS (Brünger *et al.*, 1998). The final model, refined at 1.9 Å ($R_{\text{cryst}} = 22.4\%$, $R_{\text{free}} = 24.9\%$), comprises 244 residues (208–451), the ligand, 10 residues of the SRC1 peptide (687–696) and 137 water molecules. According to PROCHECK (Laskowski *et al.*, 1993), 99.2% of peptide ϕ/ψ angle pairs lie in allowed regions of the Ramachandran plot, 0.4% in generously allowed regions and 0.4% in disallowed regions. This last percentage corresponds to one residue (D403) from loop 9–10 for which the electron density is not well defined. The probe-occupied volume of the cavity was calculated with VOIDOO (Kleywegt, 1994) using a probe radius of 1.4 Å, and the volume of stearate with GRASP (Nicholls, 1993). In the ROR β LBD homodimer modeling study, the buried surface was also calculated with GRASP. The atomic coordinates have been deposited with the PDB (accession code: 1K4W; Berman *et al.*, 2000).

Wild-type and mutant expression vectors and reporter plasmids

The luciferase reporter plasmid (GALp)₃-TKLuc containing three copies of the GAL4-binding site upstream of a thymidine kinase (TK) promoter and the expression vectors for pCMXGal-ROR β _{201–459} and pCMXGal-ROR β _{201–440} were described previously (Greiner *et al.*, 2000). The prokaryotic expression vector pRSETB-ROR β _{201–459} was generated by subcloning ROR β _{201–459} from pCMXGal-ROR β _{201–459} into pRSETB purchased from Invitrogen (Carlsbad, CA). Point mutations in the ROR LBD were generated by using an overlapping PCR approach. PCR products were cloned in-frame into pCMXGal at *EcoRI* and *Sall* restriction sites (Umesono *et al.*, 1991). Detailed descriptions on the cloning of all expression vectors for Gal-ROR β fusion proteins described in the manuscript are available on request. All generated plasmids were verified by sequencing.

Cell culture and transfection assays

HT22 and NIH 3T3 cells were cultured in Dulbecco's modified Eagle's medium (DMEM). Neuro2A cells were cultured in Earl's modified Eagle's medium (EMEM). Both media were supplemented with 10% fetal calf serum, penicillin, streptomycin and glutamine. Transient transfection assays were carried out in 12-well plates (4×10^4 cells per well) using the standard calcium phosphate co-precipitation technique (Pfitzner *et al.*, 1995) or DOTAP lipofection (Roche Molecular Biochemicals) according to the manufacturer's protocol. Luciferase

activity was assayed as recommended by the manufacturer (Promega) in a Luminometer ML 3000 (Dynatech). Relative light units were normalized according to Pfitzner *et al.* (1995). All experiments were repeated at least five times.

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