Unique requirements for retinoid-dependent transcriptional activation by the orphan receptor LXR

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LXR is an orphan nuclear receptor that confers retinoid responsiveness to the retinoid X receptor (RXR) by its interaction on a specific response element called an LXRE. To understand the mechanism of this response, three characteristics were identified that are crucial to activation of the RXR–LXR complex. First, the orientation of the RXR–LXR heterodimer on DNA indicates that as the ligand-binding partner, RXR occupies the 5' half-site of the response element. Next, the sequence specificity of the LXRE was determined in order to identify residues required for retinoid activation of the heterodimer. Remarkably, subtle changes in the nucleotide sequence of the LXRE half-sites that do not substantially alter DNA binding of the RXR–LXR heterodimer have a significant effect on the ability of the complex to be activated by ligand. Finally, we characterized the contributions of the activation domains of each receptor to the trans-activation potential of the RXR–LXR heterodimer. Surprisingly, our results show that only the activation domain of LXR is required for retinoid activation. Taken together, these results demonstrate the existence of a unique form of communication between heterodimer partners in which the activation potential of one receptor (LXR) is enabled by ligand binding to its partner (RXR). Furthermore, we conclude that RXR ligand activation potential is not dictated solely by its position on DNA, but is influenced by other factors such as the receptor partner and sequence of the response element.

[Key Words: RXR; LXR; retinoid receptors; heterodimers; trans-activation domain; nuclear receptors]

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Members of the nuclear hormone receptor superfamily, including the receptors for retinoic acid, vitamin D, thyroid hormone, and steroid hormones, are transcription factors that influence a variety of cellular processes in response to ligand binding (for review, see Mangelsdorf et al. 1995). These receptors act by binding to specific cis-acting elements in the promoters of target genes, thereby repressing or activating gene expression. A subgroup of receptors within the superfamily, including the retinoic acid receptor (RAR), thyroid hormone receptor (TR), vitamin D receptor (VDR), peroxisome proliferator activated receptor (PPAR), and many orphan receptors, bind to specific DNA sequences known as hormone response elements as heterodimers with the retinoid X receptor (RXR) (for review, see Mangelsdorf and Evans 1995). The hormone response elements generally consist of two direct repeat half-sites of a consensus hexameric sequence (AGGTCA). Response element specificity resides in subtle differences in the sequence of the half-sites as well as the number of nucleotides in the spacer region between the two half-sites. Receptors that bind DNA as heterodimers with RXR exhibit a specific polarity on direct-repeat response elements such that the silent, non-ligand-binding partner (RXR) occupies the 5' half-site whereas the ligand-binding receptor (i.e., RAR, TR, or VDR) occupies the 3' half-site of the response element (Kurokawa et al. 1993; Perlmann et al. 1993; Schrädler et al. 1995). These data have led to the suggestion that the position of the receptor on DNA (i.e., binding to the 3' half-site) enables the receptor to be activated by ligand.

Nuclear receptors share a common structure consisting of several modular domains. Along with the highly conserved DNA-binding domain and a carboxy-terminal domain that specifies both ligand binding and receptor dimerization, these receptors also contain regions required for transcriptional activation. The amino-terminal region of most of these receptors contains a transcriptional activation domain referred to as AF-1 that contributes to constitutive activation by the receptor (Hollenberg and Evans 1988; Nagpal et al. 1992). A second transcriptional activation domain termed the AF-2 is located in the carboxyl terminus, but unlike the AF-1 domain, the AF-2 is ligand-dependent and conserved among members of the nuclear hormone receptor super-
family (Danielian et al. 1992). Mutations within this AF-2 region have been shown to abolish ligand-activated transcription in several receptors including the glucocorticoid receptor, estrogen receptor, RAR, TR, and RXR (Hollenberg and Evans 1988; Webster et al. 1988; Nagpal et al. 1993; Saatcioglu et al. 1993; Barettoni et al. 1994; Durand et al. 1994; Tone et al. 1994; Zhang et al. 1994; Baniahmad et al. 1995; Leng et al. 1995). Transcriptional activation by these receptors is thought to involve association with a class of proteins referred to as coactivators that enhance the transcriptional response by interacting with the basal transcriptional machinery (McKnight 1996). Recently, several putative coactivators have been identified that associate with nuclear receptors in a ligand-dependent manner (Halachmi et al. 1994; Cavailles et al. 1995; Lee et al. 1995; Le Douarin et al. 1995; Ohate et al. 1995; vom Baur et al. 1996). Along with transcriptional activation, TR and RAR recently have been shown to interact with corepressors that mediate repression of target genes in the absence of ligand binding (Chen and Evans 1995; Hörlein et al. 1995).

LXR is an orphan receptor that heterodimerizes with RXR and binds to a specific response element called the LXRE (Willy et al. 1995). The unique association between RXR and LXR on this element allows the complex to be activated by the RXR ligand 9-cis retinoic acid [9cRA]. To understand the mechanism by which LXR enables RXR ligand activation, we examined several parameters that are crucial to nuclear receptor heterodimer action. Taken together, our results define the requisite contributions made by RXR, LXR, and the LXRE to the transcriptionally active heterodimer. We find that in contrast to other RXR heterodimers, ligand activation occurs while the ligand-binding receptor (RXR) occupies the 5' half-site of the response element. In addition, we find that ligand binding by one receptor (RXR) confers activation through the AF-2 domain of the other receptor (LXR). These data suggest the existence of a novel signaling mechanism between heterodimeric partners.

Results

RXR–LXR heterodimer polarity

Previous studies have shown that RXR is the ligand-binding partner in the retinoid-activated RXR–LXR heterodimer (Willy et al. 1995). Therefore, it was of interest to determine whether the polarity of the RXR–LXR heterodimer is different from that of other heterodimers in which it is known that the ligand-binding partner occupies the 3' half-site of the response element and RXR, which does not bind ligand, occupies the 5' half-site (Kurokawa et al. 1993; Perlmann et al. 1993). To deduce the polarity of the RXR–LXR heterodimer on the LXRE sequence, we used site-directed mutagenesis to alter the amino acids in the DNA-binding domain F box of LXR to those of the glucocorticoid receptor [GR] (Perlmann et al. 1993), thereby changing LXR's half-site binding specificity to AGAACCA. The mutant receptor, referred to as LXRpg, was used in electrophoretic mobility shift assays (EMSA) and transient transfections to ascertain the polarity of binding of the RXR–LXR heterodimer using two mutant LXRE sequences [Fig. 1A]. The two mutant sequences, designated G4L and L4G, have their 5' or 3' half-site nucleotides changed to those that specify GR binding (Strähle et al. 1987). As shown in Figure 1B, none of the receptors alone could bind to either element; however, in the presence of RXR, LXRpg bound the L4G element in which the GR-specific half-site is in the 3' position [lane 5]. In contrast, no binding was observed with the G4L element using any of the receptor combinations.

To confirm that the in vitro binding of RXR–LXRpg to the L4G response element was functional, LXRpg was tested for ligand-dependent transcriptional activation in

Figure 1. Polarity of the RXR–LXR heterodimer on DNA. (A) Sense-strand sequences of double-stranded oligonucleotides used to demonstrate RXR–LXR polarity. The glucocorticoid receptor-specific half-site is outlined. (B) EMSA analysis using in vitro synthesized receptor proteins in the combinations shown above the gel with one of the three 32P-labeled oligonucleotide probes shown in A. (C) Transient transfections in CV-1 cells using LXR, RXR, or LXRpg and luciferase reporters containing three copies of either L4G or G4L response elements shown in A. In transfections containing two receptors, equivalent amounts of both receptor plasmids were added to the transfections. (Open bars) Etoh; (shaded bars) methoprene acid [MA]; [RLU] relative light units.
transient transfections of CV-1 cells using a luciferase reporter containing three copies of either the L4G or G4L response element upstream of a minimal promoter (TK-L4G-LUC and TK-G4L-LUC, respectively). The RXR-selective ligand, methylretinene acid (Harmon et al. 1995), which has been shown previously to activate the RXR–LXR heterodimer [Willy et al. 1995], was used to activate transcription of the reporter in this and subsequent experiments. Consistent with the in vitro binding results, LXRα-dependent transcription of the reporter gene occurred with the L4G, but not the G4L reporter (Fig. 1C). Furthermore, when exogenous RXR was added to the transient transfections along with LXRα, an increase in both basal and ligand-activated transcription was observed [Fig. 1C], which is characteristic of the LXR response on the LXRE in the presence of exogenous RXR. Taken together these results demonstrate that the RXR–LXR heterodimer binds to the LXRE with RXR occupying the 5’ half-site and LXR the 3’ half-site. This is the same polarity observed with other RXR heterodimers including RAR, TR, and VDR [Kurokawa et al. 1993; Perlmann et al. 1993; Schröder et al. 1995]. However, in contrast to our results with LXR, in other RXR heterodimers it has been shown that ligand must first bind to the 3’ partner to initiate trans-activation [Forman et al. 1995; Roy et al. 1995]. The fact that RXR directs ligand-dependent transactivation while maintaining its 5’ position within the LXR heterodimer is clear evidence that the orientation of the receptor on DNA alone does not dictate a receptor’s ability to be activated by ligand.

**Identification of critical nucleotides in the LXRE sequence that specify DNA binding and retinoid-induced transcriptional activation**

Previous studies have shown that the sequence of the LXRE is essential for retinoid activation of the RXR–LXR heterodimer [Willy et al. 1995]. The LXRE sequence has been defined as a degenerate DR4-type element containing nucleotide substitutions in both half-sites that deviate from the canonical AGGTCA motif [Willy et al. 1995]. To determine which nucleotides are essential and/or optimal for retinoid induced transcriptional activation, numerous substitutions were made in the LXRE sequence. Because we have shown previously that the RXR–LXR heterodimer can not trans-activate all DNA sequences that it binds to with high affinity, our strategy for analyzing LXRE sequences was based on activation rather than DNA binding. Thus, a single copy of each mutated response element was cloned into a luciferase reporter and the ability to confer retinoid activation by LXR was tested in transient transfections of CV-1 cells. Sequences and activation data from these experiments are summarized in Figure 2.

Initially, the 5’ half-site was mutated to make it more closely resemble the consensus AGGTCA sequence. The most notable deviation of the 5’ half-site from the consensus sequence is that the highly conserved C residue normally found in position 5 of the consensus half-site is a T in the LXRE sequence. When this T was changed to C, activation by the element was similar to that of the original LXRE (Fig. 2, sequence 1). However, when this mutation was coupled with a T to G conversion at position 3 in the 5’ half-site, generating a GGTTCA sequence, there was a twofold increase in activation on the element [Fig. 2, sequence 2]. Interestingly, this is the best retinoid-inducible LXRE we have tested to date. Altering either the 5’ (sequence 3) or 3’ (sequence 4) half-site alone to give the consensus AGGTCA sequence also resulted in activation above that of the wild-type LXRE. Surprisingly, however, changing both half-sites to the consensus DR-4 resulted in relatively poor activation [Fig. 2, sequence 5]. As we have noted previously [Willy et al. 1995], the weak activation on the consensus DR-4 is not attributable to a notable difference in the DNA-binding affinity of the RXR–LXR heterodimer.

Besides nucleotide substitutions in the half-sites, the spacer and flanking nucleotides were also changed to determine their effect on binding and activation. As can be seen in Figure 2, alterations in the nucleotides flanking the half-sites resulted in lower levels of activation for some response elements [sequences 6–9, 12, and 13], and complete loss of activation for others [sequences 10 and 11]. Changing the sequence of the spacer resulted in either an increase or decrease in activation [sequences 14

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**Trans-activation by RXR–LXR heterodimers**

| LXRE MUTANT | Activation |
|-------------|------------|
|            |            |
| 5’ ccag GGTTTA aata AGGTCA tcac | 123456 |
| 3’            | 123456 |
| 1. ccag GGTCTA aata AGGTCA tcag | 5.5 |
| 2. ccag GGTCTA aata AGGTCA tcag | 11.0 |
| 3. ccac AGGTCTA aata AGGTCA tcag | 7.1 |
| 4. ccag GGTCTA aata AGGTCA tcag | 6.9 |
| 5. ccag AGGTCTA aata AGGTCA tcag | 2.8 |
| 6. cccag GGTCTA aata AGGTCA tcag | 3.6 |
| 7. tggc GGTCTA aata AGGTCA tcag | 4.7 |
| 8. cttc GGTCTA aata AGGTCA tcag | 4.8 |
| 9. cccg GGTCTA aata AGGTCA tcag | 2.5 |
| 10. ccag GGTCTA aata AGGTCA tcag | N/A |
| 11. ccag GGTCTA aata AGGTCA tcga | N/A |
| 12. ccag GGTCTA aata AGGTCA agag | 3.4 |
| 13. ccag GGTCTA aata AGGTCA tcct | 3.1 |
| 14. ccag GGTCTA aata AGGTCA tcag | 7.6 |
| 15. ccag GGTCTA aata AGGTCA tcag | 2.8 |
| 16. ccag GGTCTA aata AGGTCA tcac | N/A |
| 17. ccag GGTCTA aata AGGTCA tcca | N/A |

**Figure 2.** Multiple sequences can function as LXREs. Shown are the sense-strand sequences of the wild-type and mutant LXRE double-stranded oligonucleotides that were cloned as single copies into the TK-LUC reporter. Nucleotides altered from the original LXRE sequence are underlined. Each response element cloned into TK-LUC was tested for LXR activation in transient transfections in CV-1 cells. Fold activation by meth­retinene acid is given next to each response element sequence. All data were generated from the same experiment performed in triplicate. Nucleotides in the half-sites are numbered and shown in bold. [N/A] Not active.
and 15), whereas changing the number of nucleotides in the spacer to a DR-3 or DR-5 created an inactive response element (sequences 16 and 17). EMSA experiments have demonstrated a decreased affinity by RXR-LXR for some, but not all of these sequences (Willy et al. 1995; data not shown). For example, sequence 11, which is inactive, binds the RXR-LXR heterodimer with high affinity [Willy et al. 1995]. These results indicate that the inability to activate is not simply a result of a lower affinity for the element, but involves other critical interactions that are altered when RXR-LXR heterodimers bind to certain sequences.

Although no strict consensus sequence can be defined, these data clearly demonstrate that the LXRE is not the only sequence that can confer retinoid responsiveness to the RXR-LXR heterodimer, and, in fact, several nucleotide substitutions in the 5' half-site of the LXRE gave enhanced activation by the heterodimer. Our data also suggest that RXR-LXR binding to some DNA sequences does not necessarily equate with activation in response to retinoids.

The LXR AF-2 domain is necessary for RXR-LXR activation by retinoids on an LXRE

The carboxy-terminal region of most nuclear receptors contains a conserved transcriptional activation function (AF-2) that is required for ligand-dependent activation [Fig. 3A] [Danielian et al. 1992]. For all characterized RXR heterodimers [i.e., RXR-RAR and RXR-TR], the AF-2 domain of the ligand-binding partner is absolutely required for ligand-dependent activation. Interestingly, in most of these heterodimer complexes, the AF-2 domain of the non-ligand-binding partner [i.e., RXR] is not required and can be mutated or deleted without complete loss of function [Durand et al. 1994; Zhang et al. 1994]. To determine whether the AF-2 domain of the ligand-binding partner [RXR] contributes to the ligand-dependent transcriptional activation of the RXR-LXR heterodimer, the RXR AF-2 domain was mutated and tested in transient transfections with LXR for retinoid-dependent transactivation. Site-directed mutagenesis was used to generate two RXR AF-2 mutants, RXR-L451A, in which leucine 451 was replaced with an alanine, and a double mutant, RXR-L455A/E456A, in which both leucine 455 and glutamic acid 456 were replaced with alanines. Based on previous studies, these RXR mutants should retain both ligand-binding and dimerization abilities [Schulman et al. 1995; data not shown]. However, consistent with previous work [Zhang et al. 1994; Leng et al. 1995; Schulman et al. 1995], RXR AF-2 mutants are not capable of ligand-activated transcription of a reporter containing an RXR response element (TK–CRBPII–LUC) [Fig. 3B]. Surprisingly, when the RXR AF-2 mutants were tested with LXR on the LXRE response element, retinoid-dependent induction of the reporter gene was still observed [Fig. 3B]. The overall level of activation for wild-type RXR was slightly higher than for the AF-2 mutants, although the fold activation remained similar, suggesting that the RXR AF-2 domain has more of an effect on the basal than the ligand-inducible transcriptional response. These results indicate that the RXR AF-2 domain contributes to, but is not required for, retinoid induction of the RXR–LXR heterodimer.

The above results imply that the AF-2 domain of LXR may contribute to the ligand-dependent transcriptional activation of the heterodimer on an LXRE. To address this possibility, the carboxy-terminal portion of LXR corresponding to the region in and around the putative AF-2 domain was characterized by alanine scanning mutagenesis [Fig. 4A]. Five consecutive amino acids of LXR were systematically changed to alanine residues and the resulting LXR mutants were tested for activation by the RXR-selective ligand methoprene acid. Retinoid inducibility did not vary dramatically when any of the last three amino acids that are outside of the putative AF-2 region were mutated to alanines [Fig. 4A]. However, mutations to the tryptophan at position 443 and the aspartic...
acid at 444 indicate that both are required for full transcriptional activation [Fig. 4A]. Whereas the LXR-W443A mutant exhibited no detectable activation under any conditions tested, the LXR-D444A mutant was able to confer a low level of activation that increased slightly when RXR was also overexpressed [Fig. 4B]. Overexpressed RXR could not rescue the LXR-W443A mutant under any conditions, indicating a crucial role for this amino acid in transcriptional activation. Consistent with this finding, when tryptophan 443 was changed to a leucine, the resulting mutant was also inactive under all conditions tested [data not shown]. Because the mutated amino acids in LXR that affect activation are contained within the region that corresponds to the conserved activation domain in RXR and other receptors [Danielian et al. 1992], we designate this region as the LXR AF-2 domain [see Fig. 4A]. As was the case with wild-type LXR [Fig. 3B], a similar response was seen with the LXR mutants regardless of whether RXR or the RXR AF-2 mutant were also expressed [Fig. 4B]. In vitro transcription/translation and EMSA analysis of LXR AF-2 mutants showed that all were made in equivalent amounts to wild-type LXR and all exhibited comparable DNA binding as compared with wild-type LXR [data not shown]. Transient transfection experiments with wild-type LXR showed a decrease in reporter activation by methoprene acid when increasing amounts of LXR-W443A were added, demonstrating that the LXR-W443A protein was expressed in transfected cells. Western blot analysis of LXR and LXR AF-2 mutants expressed in transfected cells also indicated that wild-type and mutant proteins were expressed at comparable levels [data not shown].

Although we have shown previously that methoprene acid is strictly an RXR-selective ligand [Harmon et al. 1995], the possibility existed that the observed effects on the RXR-LXR heterodimer were in some way peculiar to methoprene acid. To rule out this possibility, dose response experiments [Fig. 5A–F] were performed with methoprene acid and other known RXR ligands, including the natural ligand 9-cis retinoic acid and the potent synthetic ligand LG69 [Boehm et al. 1994]. Using both RXR and LXR AF-2 mutants, these experiments recapitulated completely the results seen with methoprene acid alone [Fig. 4], confirming that the AF-2 domain of LXR, but not RXR, is required for activation of the heterodimer. Significantly, the dose responses for these ligands on the wild-type RXR–LXR heterodimer are identical to those observed on the wild-type RXR homodimer [Boehm et al. 1994; Harmon et al. 1995; Willy et al. 1995], implying that in both cases RXR alone binds ligand.

To further demonstrate that the LXR AF-2 domain is required for activation of the RXR–LXR heterodimer, trans-activation assays were performed with DNA-binding orientation-independent heterodimers. For these experiments, GAL4-chimeric receptors were used with a luciferase reporter containing four copies of the GAL4 DNA-binding site [TK-MH100X4–LUC]. A chimeric receptor containing the amino terminus and DNA-binding domain of GAL4 fused to the RXR ligand-binding do-

Figure 4. Retinoid-induced activation of the RXR–LXR heterodimer requires the LXR AF-2 domain. (A, left) Amino acid sequence of the wild-type LXR and LXR AF-2 domain mutants. Amino acids that were mutated to alanines are underlined. (Right) Transient transfections of CV-1 cells with the LXR AF-2 mutant receptors shown using the TK-LXRE3–LUC reporter. (B) Transient transfections of CV-1 cells using LXR and the LXR AF-2 mutant receptors in combination with RXR or the RXR AF-2 mutant, RXR-L455A/E456A. In transfections containing both LXR and RXR constructs, one-tenth as much RXR or its mutant plasmid was used. (A, B) Bars as in Fig. 1.
Figure 5. The LXR AF-2 domain is required for RXR–LXR trans-activation by several different RXR ligands. CV-1 cells were transiently transfected with a TK–LXRE3–LUC reporter and the CMX receptor expression plasmids indicated in A–F. Cells were then exposed to increasing concentrations of the indicated ligands and analyzed for luciferase activity. In transfections containing both LXR and RXR, one-tenth as much RXR or RXR mutant plasmid was used. RXR–ΔAF2 represents CMX–RXR–L455A/E456A. (RLU) Relative light units.

Discussion

In this paper we have defined several properties of the RXR–LXR heterodimer that distinguish it from other RXR heterodimer signaling pathways. One conserved feature of RXR heterodimers with RAR, TR, and VDR that sets them apart from the RXR–LXR heterodimer is the inability of these other complexes to be activated from the basal state by RXR ligands. In the case of at least one of these complexes, the RXR–RAR heterodimer, Glass and coworkers have shown that the inability of RXR to bind ligand is a result of an allosteric block imposed by RAR (Kurokawa et al. 1994). Thus, activation of these heterodimers is accomplished by ligand binding to the receptor partner of RXR, which exclusively occupies the 3' half-site of the direct repeat response element (Kurokawa et al. 1993; Perlmann et al. 1993; Schrader et al. 1995). In the one known example (RAR–RXR) where this polarity is switched and the partner of RXR occupies the 5' half-site, the heterodimer is no longer ligand-activated (Kurokawa et al. 1994). One interpretation of these results is that the ligand inducibility of a receptor is determined in part by its orientation on DNA. In this work, we have shown that the polarity of the RXR–LXR heterodimer, although similar to that of other RXR heterodimers, is unique in that the
Figure 6. Polarity-independent trans-activation of the RXR–LXR heterodimer requires a functional LXR AF-2 domain. CV-1 cells were transiently transfected with a luciferase reporter containing GAL4 DNA-binding sites and the indicated CMX–receptor expression plasmids. Each transfection contained 25 ng of CMX–GAL4-chimeric receptor alone or with 25, 50, or 100 ng of CMX–LXR, CMX–LXR-W443A, or CMX–TRβ expression plasmid. The LXR, LXR-W443A, and TR controls (first three lanes) contained 100 ng of receptor expression plasmid and 25 ng of CMX–GAL4DI5D. GAL4-RXR-AAF2 represents CMX–GAL4–RXR-LXB–L455A/E456A. The methoprene acid response is expressed as the fold activation of triplicate assays ±S.E.

ligand-binding partner (RXR) occupies the 5' position of the response element, whereas the partner that contributes to DNA-binding specificity (LXR) occupies the 3' half-site. Similar results have also been found with the orphan receptors NGFI-B and NURR1, which can bind specific response elements as monomers or as heterodimers with RXR, provided that NGFI-B occupies the 3' half-site of the response element (Perlmann and Janson 1995). As RXR heterodimers, these orphan receptors can also direct retinoid-dependent transcriptional activation. The ability of the 3' partner to enable RXR to bind and be activated by ligand appears to be a unique feature of these receptors and further strengthens the notion that one function of orphan receptors such as LXR and NGFI-B is to increase the complexity of 9cRA signaling through RXR (Leblanc and Stunnenberg 1995). From these observations, we conclude that RXR ligand-activation potential is dictated not solely by its position on DNA, but rather by other factors such as the receptor partner and sequence of the response element. In further agreement with this conclusion, we note that RXR–LXR heterodimers bind with the same fixed polarity to LXREs (which are retinoid-responsive) and canonical DR4 elements (which are not retinoid-responsive). In addition, RXR–TR heterodimers, which are not responsive to RXR ligands, bind with the same fixed polarity to all DR4 elements, including LXRE sequences (Perlmann et al. 1993; P. Willy, unpubl.). Finally, polarity-independent experiments using GAL4–RXR–LXR heterodimers and their AF-2 mutants (Fig. 6) further emphasize the importance of the partner receptor in mediating the RXR ligand response.

This work has also identified critical nucleotides in the LXRE sequence that contribute to activation by the heterodimer. From this analysis it is clear that the LXRE sequence is not the only element capable of mediating the RXR–LXR response. Indeed, mutagenesis has defined a sequence that as a single copy gives a twofold greater activation than the original LXRE sequence (Fig. 2, sequence 2). Nevertheless, it has not been possible to generate a consensus LXRE sequence, because several individual changes that increase activity do not act cooperatively (or even additively) when combined. For example, changing either half-site to the canonical AGGTCA sequence results in an increase in retinoid activation. However, altering both half-sites to AGGTCA lowers activity dramatically. Consistent with other receptor response elements (Umesono et al. 1991; Mader et al. 1993; Katz and Koenig 1994), in addition to the half-site sequences, both the spacing and flanking nucleotides of the LXRE are important for binding and activation. The LXRE has an absolute spacing requirement of four nucleotides (Fig. 2), the sequence of which can modulate, but is not critical, for activation. The nucleotides immediately upstream and downstream of the core element are also important for activation but are less critical for in vitro DNA binding.

In addition to the contributions made by the LXR ligand- and DNA-binding domains as well as the response element, a unique mechanism of ligand-dependent transcriptional activation through the RXR–LXR heterodimer has been identified. Our results demonstrate that the functional AF-2 domain of LXR is critical for retinoid activation of the RXR–LXR heterodimer. Surprisingly, the AF-2 domain of RXR, the ligand-binding receptor in this heterodimer, is not required for activation and its apparent contribution is minimal. These results differ from previous observations with RXR–TR and RXR–RAR heterodimers in several ways. In these heterodimers the AF-2 domain of the ligand-binding receptor is required for ligand-dependent transcriptional activation (Saatioglu et al. 1993; Barettoni et al. 1994; Durand et al. 1994; Tone et al. 1994; Banaihmad et al. 1995). With these other receptors it has therefore been suggested that ligand binding to the receptor induces a conformational change (Renaud et al. 1995) that dissociates a corepressor (Chen and Evans 1995; Holstein et al. 1995) and attracts a coactivator (Onate et al. 1995), thereby permitting positive interactions with components of the transcriptional machinery (Fig. 7A). Together with the ability of at least one of these receptors (i.e., RAR) to allosterically block RXR ligand binding, these observations likely explain why RXR cannot respond to retinoids in the context of these heterodimeric interactions (Kurokawa et al. 1995). Our results suggest a different mechanism of action for the RXR–LXR heterodimer (Fig. 7B), namely that the entire heterodimer complex, not simply the ligand-binding partner, must undergo a conformational change that allows transcrip-
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Figure 7. Models for two mechanisms of ligand activation of RXR heterodimers. (A) In many RXR heterodimers, RXR cannot be activated by 9cRA. In the basal state, the 3′ receptor partner (e.g., TR) binds a corepressor [CoRep.] that suppresses promoter activity. Ligand binding to the 3′ partner induces a conformational change in the receptor that simultaneously releases the corepressor and alters the activation domain (AF-2) for interaction with coactivator [CoAct.] proteins. (B) In the RXR–LXR heterodimer, basal transcription is not repressed and RXR ligand binding is enabled while RXR maintains its position on the 5′ half-site. The ligand-mediated conformational change in RXR induces a change in LXR, which can then interact with putative coactivators to affect transcription. In both A and B ligand-induced trans-activation requires the AF-2 of the 3′ receptor partner, regardless of which partner binds ligand.

Materials and methods

Plasmids

Constructions of wild-type receptors inserted into the expression vector pCMX have been detailed elsewhere (Umesceno et al. 1991; Willy et al. 1995). CMX–LXRpg and RXR AF-2 mutants were generated by site-directed mutagenesis using the Morph Mutagenesis Kit (5 Prime → 3 Prime, Inc., Boulder, CO). CMX–LXR–AF-2 alanine scanning mutants were generated as follows: mutant oligonucleotides encoding the last five amino acids of LXR with single alanine substitutions were ligated into CMX–hLXRα that had been digested with BglII and Nhel to remove nucleotides encoding the last five amino acids and 3′-untranslated sequence. Luciferase reporter plasmids were constructed by ligation of the appropriate double-stranded oligonucleotides into the HindIII site of the TK–LUC vector. TK–LXRE3–LUC (Willy et al. 1995), TK–CRBPII–LUC (Mangelsdorf et al. 1991), TK–MH100X4–LUC, and CMX–GAL4–hRXRa have been described. The sequences for L4G and G4L reporters are shown in Figure 1 and contain HindIII ends for cloning. TK–L4G–LUC contains three copies of the response element, which are all in a 5′ → 3′ orientation with respect to the transcription start site. TK–G4L–LUC also contains three copies of the response element, with the first two copies in the 5′ → 3′ direction and the last in the opposite orientation. LXRE mutants were cloned into TK–LUC as a single copy in the 5′ → 3′ orientation with respect to the transcription start site. All plasmid constructs and mutations were verified by sequenc-

EMSAs

Receptor proteins were generated using coupled in vitro transcription/translation as per manufacturer’s instructions (Promega). Sequences of double-stranded oligonucleotides were as shown in the figures and text and were synthesized with HindIII overhangs. Binding reactions were performed in a total volume of 20 μl consisting of 75 mM KCl, 20 mM HEPES at pH 7.4, 2 mM DTT, 7.5% glycerol, 0.1% NP-40, 2 μg of poly(dI–C) [Phar macia], 60 pmoles of a nonspecific single-stranded oligonucleotide [for removal of nonspecific binding in the lysates], and 2 μl of each receptor lysate or unprogrammed (control) lysates to give a total of 4 μl of lysate per reaction. Binding reactions were incubated for ~90 min on ice followed by the addition of 40 fmoles of 32P-labeled oligonucleotide probe [labeled by end-filling] and incubated further for 30 min at room temperature. Samples were analyzed on 5% polyacrylamide gels run in 0.5x TBE buffer at 275 V for 1.5 hr.

Cell culture and cotransfection assays

CV-1 cells were maintained at 37°C, 7% CO2 in Dulbecco’s modified Eagle medium (DMEM) containing 5% calf bovine serum (CBS). Transfections were performed in 48-well plates in

Nucleotides in the response element dictate whether or not binding is coupled to activation by the heterodimer. Ligand binding by RXR then confers a change in LXR, allowing for transcriptional activation that is mediated through the LXR AF-2 domain. This finding suggests the intriguing possibility that this novel mechanism of activation through the AF-2 domain of the non-ligand-binding partner may be used by other receptor heterodimers that work through RXR signaling.
media containing 5% dextran-charcoal stripped CBS by the calcium phosphate coprecipitation technique as described previously (Willy et al. 1995). Eight hours after transfections ligands were delivered to cells at 10-fold dilutions [0.1% (vol/vol) of solvent in medium]. Retinoids were manipulated under cold light and stored dark in ethanol or methanol under nitrogen gas at −80°C. Cells were harvested 36 hr after addition of ligand and analyzed for luciferase and β-galactosidase activity using a Dye natech ML3000 luminometer and MR5000 spectrophotometer, respectively. For most experiments, cotransfection of DNA into mammalian cells was accomplished with 50 ng reporter plasmid, 50 ng of CMX-βgal, 25 ng of each receptor expression plasmid, and pGEM carrier to give 375 ng of DNA/well (see figure legends for details on individual experiments). Transfections with mutant LXRE reporters [Fig. 2] contained 65 ng of receptor, 25 ng of reporter, 50 ng of CMX-βgal, and pGEM carrier to give 375 ng of DNA/well. For controls, CMX vector was used in place of the receptor plasmid. Ligand concentrations were 10−4 M methoprene acid and 10−6 M thyroid hormone. All transfection data points were normalized using an internal P-galactosidase marker (Mangelsdorf et al. 1990) and represent the mean of triplicate assays ± S.E.

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