RESEARCH COMMUNICATION

Identification of a novel mechanism of regulation of Ah (dioxin) receptor function

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Ah receptor (AhR) is a ligand-activated transcription factor that mediates pleiotropic effects of environmental pollutants such as 2,3,7,8-tetrachlorodibenzo-*p*-dioxin on host animals. In addition to induction of drug-metabolizing enzymes, the liganded AhR complex was found to activate gene expression of a factor designated AhR repressor (AhRR), which inhibits AhR function by competing with AhR for dimerizing with Arnt and binding to the XRE sequence. Thus, AhR and AhRR form a regulatory circuit in the xenobiotic signal transduction pathway and provide a novel mechanism of regulation of AhR function that may determine tissue-specific sensitivity to environmental pollutants.

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AhR (arylhydrocarbon receptor, or dioxin receptor) has been known to mediate pleiotropic biological effects of various environmental contaminants, mainly polycyclic aromatic hydrocarbons usually represented by 2,3,7,8-tetrachlorodibenzo-*p*-dioxin (TCDD). These biological effects include teratogenesis, tumor promotion, thymic atrophy, epithelial hyperplasia, hepatotoxicity, and induction of drug-metabolizing enzymes (Poland and Knutson 1982; Swanson and Bradfield 1993; Hankinson 1995; Sogawa and Fujii-Kuriyama 1997). AhR is usually present in cytoplasm in association with Hsp90 (Perdew 1988; Pongratz et al. 1992). Subsequently, upon high-affinity binding of inducing chemicals, the liganded AhR translocates to nuclei, where it switches the partner from Hsp90 to Arnt (AhR nuclear translocator) and binds the cognate enhancer sequence, XRE, upstream of the target genes for CYP1A1, GST, and others to activate their expressions (Fujisawa-Sehara et al. 1987; Telakowski-Hopkins et al. 1988). Recently, involvement of AhR in TCDD-induced teratogenesis, such as cleft palate and hydronephrosis in fetal development and cytotoxicity in adult animals, has been demonstrated by using AhR knockout mice (Fernandez-Salguero et al. 1996; Mimura et al. 1997).

Structurally, AhR and Arnt belong to a superfamily of bHLH transcription factors that include MyoD and Myc

(Murre et al. 1989). A characteristic common domain of AhR and Arnt, designated PAS [conserved sequence among Per (Jackson et al. 1986), Arnt/AhR (Hoffman et al. 1991), and Sim (Nambu et al. 1991)], which abuts on the carboxyl terminus of the bHLH, defines a growing family of bHLH-PAS factors among the bHLH superfamily.

These transcription factors with bHLH motif form homo- and/or heterodimers with themselves or other members of the same family, to bind the cognate binding site upstream or downstream of their target genes, resulting in activation of gene expression (Murre et al. 1989). Because many of the bHLH transcription factors are involved in physiologically and developmentally important functions such as cell proliferation and differentiation, their transcription activities are negatively regulated by competitive heterodimer formation with inhibitory bHLH proteins. Two Myc-related proteins, Mad (Ayer et al. 1993) and Mxi1 (Zervos et al. 1993), dimerize with Max, and these heterodimers bind the same core sequence (CACGTG) as Myc/Max heterodimer. Thereby, Mad and Mxi1 interfere with Myc function either by sequestering Max or by direct competition for the DNA target sequence. During differentiation of certain myeloid cell lines in vitro, relative changes in the intracellular concentration of Myc and Mad (or Mxi1) rapidly modulate the expression of a set of genes responsive to these transcription factors (Ayer and Eisenman 1993; Larsson et al. 1994). In another group of bHLH transcription factors such as MyoD and E12/E47, inhibitory proteins, Ids, which lack a basic region adjacent to the HLH, are able to dimerize with a member of bHLH proteins including MyoD and E12/E47, resulting in inhibition of their transcription activation activity via sequestration into dimers that cannot bind DNA (Christy et al. 1991; Neuhold and Wold 1993). It has been reported that Id inhibits muscle differentiation by associating with E12 and prevents it from forming the active MyoD/E12 heterodimer (Benezra et al. 1990; Jen et al. 1992). During terminal differentiation, the Id levels decrease, suggesting that Id can act as an inhibitor of differentiation.

Although two kinds of suppressive forms of bHLH transcription factors change in concentration in association with cell proliferation and differentiation, the mechanism of their gene expression remains unknown.

During investigation of AhR and Arnt of a third group of bHLH transcription factors, we isolated cDNA clones that encode a polypeptide with high similarity to the sequence of the bHLH/PAS of AhR. This polypeptide was found to repress the transcription activity of AhR by competing with AhR in forming a heterodimer with Arnt and binding with the XRE sequence, and is thus designated AhRR or AhR repressor. Furthermore, the expression of AhRR is induced by the AhR/Arnt heterodimer through binding to the enhancer sequence XRE, upstream of the *AhRR* gene; thus the AhR function is regulated by the feedback inhibition of AhRR. A similar

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mechanism has been suggested recently: the circadian rhythmic regulation of the mammalian clock system, consisting of the same bHLH-PAS factors, that is, mClock, BMAL1, and mPer1 (Gekakis et al. 1998).

Results and Discussion

During the screening of a mouse genomic library with *AhR* cDNA used as a hybridization probe, we isolated a genomic clone that has a sequence with high similarity to a part of *AhR* cDNA encoding the bHLH region (Ema et al. 1992). Subsequently, we isolated cDNA clones from a mouse small intestine cDNA library with the genomic DNA fragment showing high similarity to the *AhR* used as probe. The longest insert of the isolated cDNA clones was estimated to be 4.5 kb, and the determined sequence contains a long ORF of 2103 nucleotides, encoding a polypeptide of 701 amino acids. By comparison with other bHLH-PAS proteins, the encoded sequence shows the highest degree of sequence similarity to *AhR* in the sequence of the bHLH and PAS-A regions (Fig. 1). However, the sequence carboxy-terminal to PAS-A is quite variable. Notably, the PAS-B sequence, which functions as a ligand binding site and an interaction interface with Hsp90 in *AhR*, is missing in the deduced sequence and the sequence of the carboxy-terminal half, which corresponds to the transactivation domain for *AhR* (Sogawa et al. 1995), differs greatly from that of *AhR* (we designated the isolated factor *AhRR* for the reasons described below).

Close similarity between *AhRR* and *AhR* in the bHLH and PAS-A regions led us to investigate, by immunoprecipitation assay, whether *AhRR* interacts with Arnt. When incubated with Arnt, *AhRR* was coimmunoprecipitated efficiently by an anti-Arnt antibody in a manner independent of the presence of a ligand, 3-methylcholanthrene (3MC) (Fig. 2B, lanes 3,4), whereas interaction between *AhR* and Arnt was significantly enhanced (twofold) by 3MC (Fig. 2B, lanes 8,9) as reported previously (Hirose et al. 1996). The degree of enhanced interaction between in vitro-synthesized *AhR* and Arnt by

Figure 1. Predicted mouse *AhRR* primary structure and sequence comparison with that of *AhR*. (A) Predicted amino acid sequence of m*AhRR*, in single-letter symbols. bHLH (26–82 amino acids) and PAS-A regions (116–167) are underlined. (B) Domain structures of m*AhRR* and m*AhR*. Percentages of identity between m*AhRR* and m*AhR* are indicated.

Figure 2. Subcellular localization of *AhRR*, heterodimer formation of *AhRR* with Arnt, and its binding to XRE sequence. (A) Subcellular localization of GFP-*AhRR* fusion protein. To construct pCMX-GFP-*AhRR*, an *EcoRI*-*SalI* fragment of *AhRR* was ligated with the *SalI*-*BamHI* site of pCMX-GFP-hGR (Ogawa et al. 1995). GFP (left) or GFP-*AhRR* fusion protein (right) was expressed in COS7 cells and visualized as described (Ogawa et al. 1995). (B) Coimmunoprecipitation of *AhRR* and *AhR* with an anti-Arnt antibody in the presence or absence of 3MC. Open and closed arrowheads indicate coprecipitated *AhRR* and *AhR*, respectively. (C) Interaction between *AhRR* or *AhR* and Arnt revealed by the mammalian two-hybrid method. A fusion gene encoding GAL4-DBD and Arnt-bHLH-PAS as bait and those encoding *AhR* or *AhRR* bHLH-PAS and VP16 AD were constructed. Various prey/bait combinations were cotransfected into 293 cells with pG3E-Luc; interaction of *AhR* or *AhRR* with Arnt was assessed by measuring the expressed luciferase activity. 3MC (1 μM) was used as inducer. (D) In vitro interaction of *AhRR*/Arnt heterodimer with XRE sequence. (Lanes 6,10) Non-labeled XRE (250-fold) was used as competitor. Open and closed arrowheads indicate *AhRR*-Arnt-XRE and *AhR*-Arnt-XRE complexes, respectively.

3MC was variable, but the enhancement was reproducibly observed. This interaction between *AhRR* or *AhR* and Arnt was confirmed by the mammalian two-hybrid system (Dang et al. 1991). A fusion gene encoding the VP16 activation domain (AD) and the bHLH-PAS region of *AhRR* was transfected into 293T cells together with a fusion gene encoding GAL4-DBD and the bHLH-PAS region of Arnt, and the pG3E-Luc reporter gene. As a control experiment, a fusion gene of *AhR* bHLH-PAS and VP16 AD was used as prey. Whereas the *AhR* fusion gene enhanced the reporter gene expression in response to 3MC (Fig. 2C, lane 3), the *AhRR* fusion gene activated the luciferase expression in a manner independent of the

Figure 4. Promoter activity of *AhRR* gene in response to 3MC. (A) Structure of the *mAhRR* gene promoter. (X1 and X2) Relative location of two XRE sequences in the regulatory region upstream of the *AhRR* gene. The *AhRR* gene is a TATA box-less gene; GC boxes are indicated by shaded ovals. (B) Activation of the mouse *AhRR* gene promoter in response to 3MC. pGL3 or pAhRR-Prom-Luc was cotransfected into HeLa cells and various combinations of AhR and Arnt expression vectors, and the transfected cells were cultured for 44 hr in the presence or absence of 3MC. The promoter activity of the *AhRR* gene was assessed by measuring expressed luciferase activities. (C) Inducible expression of *AhRR* mRNA in mouse various tissues in response to 3MC. Male mice (C57BL/6J) were injected intraperitoneally with 3MC in sesame oil (80 mg/kg body weight). Total RNAs from various tissues were subjected to RT-PCR analysis.

of the gene carries two and three copies of the XRE and GC box, respectively, as shown in Figure 4A, which suggests that *AhRR* gene expression is regulated by the AhR/Arnt heterodimer in response to 3MC.

When the 5'-flanking sequence of the *AhRR* gene was fused to the luciferase gene and transfected into HeLa cells along with the expression plasmids of AhR and Arnt, expression of luciferase activity was markedly activated by the administration of 3MC (Fig. 4B, lane 4), indicating that the XRE in the *AhRR* gene is functional as inducible enhancer. These experiments established *AhRR* as a novel target gene of AhR in addition to genes encoding drug metabolizing enzymes. Against this background we investigated inducible expression of *AhRR* mRNA in various tissues of mice in response to 3MC by RT-PCR. In untreated animals, essentially no expression of the mRNA was detected, in liver, heart, lung or other tissues. Upon treatment with 3MC, however, *AhRR* mRNA levels were induced in these tissues, although the mode of expression was different from tissue to tissue. As shown in Figure 4C, heart and lung showed most abundant *AhRR* mRNA expression, whereas liver, thymus, kidney, and intestine expressed relatively small amounts of mRNA. Interestingly, tissues such as liver and thymus, which are known to be most susceptible to the toxic effects of dioxin (Poland and Knutson 1982; Fernandez-Salguero et al. 1996) appear to express *AhRR* mRNA poorly in response to inducers. Expression of the

AhRR gene and susceptibility of various tissues to dioxin are now under detailed investigation.

It has long been known that superinduction of TCDD-induced *CYP1A1* mRNA was generated in cultured cells such as Hepa-1c1c7 by treatment with inhibitors of protein synthesis such as cycloheximide (Israel et al. 1985), that the AhR function is down-regulated after a short interval of treatment with the inducers, and this down-regulation is blocked by inhibitors of protein synthesis (Lusska et al. 1992). The interesting phenomena of superinduction and block of down-regulation caused by protein synthesis inhibitors have been postulated to be due to inhibited synthesis of a short-lived repressor of AhR function, the entity of which remains elusive. The present data suggest that this repressor is most likely AhRR. Recently, a similar regulatory circuit has been suggested with a mammalian circadian system carried out by the same bHLH-PAS proteins as AhR, such as Clock, BMAL1 and mPer, although their transcriptional activities have not been demonstrated clearly. An elaborate regulatory circuit of AhR function may support the fact that it is involved in cell cycle regulation, as suggested previously (Ma and Whitlock 1996; Weiss et al. 1996).

In conclusion, we have identified *AhRR* as a target gene of the AhR, providing a novel mechanism of feedback inhibition of the receptor function in that a transcription factor induces directly the expression of its repressor gene through binding to the cognate regulatory sequence of the gene. As shown in Figure 5, this regulatory circuit involves activation of AhR by xenobiotics to induce expression of the *AhRR* gene and many others through binding to the XRE as a heterodimeric complex with Arnt, and induced levels of AhRR inhibit in a tissue-specific manner the AhR function by competing with AhR for Arnt and XRE binding activity.

Figure 5. Mechanism of negative feedback regulation of AhR function by AhRR. Ligand-activated AhR/Arnt heterodimer transactivates the expression of target genes (including *AhRR* gene) driven by the XRE sequence. Expressed *AhRR* mRNA is translated into AhRR protein, and the resulting AhRR inhibits AhR function by competing with AhR for heterodimerizing with Arnt and binding with the XRE sequence. Consequently, inhibition of translation of *AhRR* mRNA by inhibitors of protein synthesis results in superinduction of mRNAs by TCDD or 3MC.

Materials and methods

Isolation of mouse *AhRR* cDNA

A mouse genomic library was screened with a *mAhR* cDNA to obtain a genomic DNA clone. The genomic fragment encoding the bHLH region was used as a probe to screen a mouse cDNA library under stringent hybridization conditions to isolate *AhRR* cDNA. The cDNA sequence has been registered in DDBJ (accession no. AB015140).

Immunoprecipitation assay

AhRR and *AhR* proteins were synthesized *in vitro* with a TnT-coupled transcription-translation kit (Promega) in the presence of [³⁵S]methionine (Amersham). Arnt protein was synthesized with nonlabeled methionine. Labeled *AhRR* (5 μ l) or *AhR* (5 μ l) was mixed with nonlabeled Arnt (5 μ l) and incubated for 2 hr at 30°C in the absence or presence of 3MC (1 μ M). The reaction mixtures were immunoprecipitated with anti-Arnt or non-immune sera and then adsorbed on protein A-Sepharose. Immunoprecipitates were eluted by adding 2 $\times$ SDS sample buffer (20 μ l), boiled for 5 min, and subjected to SDS-PAGE. The radioactivity was visualized with the Image Analyzer BAS 1000 (Fuji film).

Plasmid construction

For construction of pBOS*AhR*, we first produced a *mAhR* cDNA fragment that contains the Kozak sequence (Kozak 1987) by PCR, using pBSK-*mAhR* as a PCR template and the following pair of primers: 5'-GTCGAAGCTTCCGCCACCATTGGCCAGCAGCGCGCC-3' (sense) and 5'-AGTCCGACGAATAGGTTTC-3' (antisense). The PCR product was inserted into the blunt-ended *Xba*I site of pBluescript vector to generate an *Xba*I site at the 3' end of cDNA [pBSK-*AhR*(K)]. pBOS*AhR* was produced by inserting the blunt-ended *Hind*III-*Xba*I fragment of pBSK-*AhR*(K) into the blunt-ended *Xba*I site of pEFBOS vector (Mizushima and Nagata 1990). To obtain pBOS-Arnt, the mouse Arnt cDNA fragment was excised by *Eco*RI-*Bam*HI digestion of pBSK-*mArnt* (Numayama-Tsuruta et al. 1997), blunt-ended by Klenow and was then ligated with the blunt-ended *Xba*I site of pEFBOS. The blunt-ended *Sma*I-*Hind*III fragment (2.4 kb) of *AhRR* cDNA was inserted into the blunt-ended *Xba*I site of pEFBOS to produce pBOS-*AhRR*. To construct VP16 fusion constructs, we first generated VP16 cDNA by PCR with pSR α -GAL4-DBD-VP16 used as a template, and the resulting fragment was inserted into pBSK-NLS, which also contained the *Hind*III-*Bam*HI fragment of pENL (Mimura et al. 1997) in pBluescript vector to obtain pNV. The blunt-ended *Hind*III-*Eco*RV fragment of pBSK-*AhR*(K) and the *Eco*RI-*Sma*I fragment of *AhRR* cDNA were inserted into the *Eco*RV and *Eco*RI-*Eco*RV sites of pNV to produce pNV-*AhR* Δ C and pNV-*AhRR* Δ C, respectively. Excision of *Hind*III-*Sal*I or *Not*I-*Sal*I from these pNV constructs and subsequent ligation of these fragments (blunt-ended) with the *Sma*I site of pEFBOS(Stop) (containing an *Xba*I nonsense linker at the 5' end of the polyadenylation signal) gave pBOS-VP16-*AhR* Δ C and pBOS-VP16-*AhRR* Δ C, respectively. For construction of pBOS-GAL-DBD-Arnt Δ C, pBOS-GAL-DBD-Arnt, and pBOS-GAL-DBD-Arnt Δ bHLH-PAS, we first produced pGBT-Arnt Δ C, pGBT-Arnt, and pGBT-Arnt Δ bHLH-PAS by inserting blunt-ended *Nco*I-*Pvu*II, *Nco*I-*Bam*HI, and *Pvu*II-*Bam*HI fragments of pBSK-Arnt into the blunt-ended *Eco*RI site of the pGBT9 vector, respectively. pBOS-GAL-DBD-Arnt Δ C, pBOS-GAL-DBD-Arnt, and pBOS-GAL-DBD-Arnt Δ bHLH-PAS were constructed by excising the *Hind*III-*Sal*I fragments from pGBT-Arnt Δ C, pGBT-Arnt, and pGBT-Arnt Δ bHLH-PAS and, subsequently, subcloning these fragments into the blunt-ended *Xba*I site of pEFBOS vector, respectively.

pG3E-Luc was produced by inserting three copies of the GAL4 binding site and Elb TATA sequence excised from pG5EC vector (Sogawa et al. 1995) into the *Sma*I site of pGL3 vector (Clontech). pX4TK-Luc was constructed by subcloning four copies of synthesized XRE1 (Kubota et al. 1991) and the TK promoter of pBLCAT2 (*Bam*HI-*Xho*I fragment) into the *Xho*I site of pGL3. p*AhRR*-Prom-Luc was produced by subcloning the blunt-ended *Hind*III-*Sac*II fragment in the promoter region of the *AhRR* genomic clone into the *Sma*I site of pGL3.

GMSA

A double-stranded oligonucleotide XRE probe (Matsushita et al. 1993) was end-labeled with [γ -³²P]ATP (Amersham). *In vitro*-translated *AhRR* or *AhR* (5 μ l) was mixed with *in vitro*-translated Arnt (5 μ l) or reticulocyte lysate without mRNA and incubated for 2 hr at 30°C in the absence or presence of 3MC (1 μ M). The reaction mixtures were diluted by adding

10 ml of 2 $\times$ binding buffer [200 mM HEPES-KOH (pH 7.9), 1 M KCl, 2 mM EDTA, 60 mM MgCl₂, 20 mM DTT, 0.2 mg/ml salmon sperm DNA]. After 15 min incubation at 25°C, the labeled XRE probe (2 $\times$ 10⁴ cpm) was added and incubated at 25°C for another 15 min. Protein-DNA complexes were resolved by 4.5% PAGE and subjected to autoradiography.

RT-PCR

Total RNAs (3 μ g) from various tissues were used for cDNA synthesis (20 μ l), and an aliquot (2 μ l) of synthesized cDNA was amplified in a total volume of 20 ml containing 150 mM dNTP, 0.2 units of *Taq* polymerase, and 0.12 μ g of each primer and [α -³²P]dCTP. Samples were amplified by repeated cycles of 94°C for 45 sec, 60°C for 45 sec, and 72°C for 1 min. Amplifications of 30 and 28 cycles were applied for *AhRR*, *AhR*, *CYP1A1*, and β -actin, respectively. PCR products were separated on 4% polyacrylamide gels and subjected to autoradiography. PCR primers for amplification of *AhR*, *CYP1A1*, and β -actin cDNA were described previously (Mimura et al. 1997), and those for *AhRR* were 5'-GGCTTAC-CATGGGCGCTGAG-3' (sense) and 5'-CCACCAGAGCGAAGCCATTGAG-3' (antisense). Mice were treated in accordance with institutional guidelines.

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References

- Ayer, D.E. and R.N. Eisenman. 1993. A switch from Myc:Max to Mad:Max heterocomplexes accompanies monocyte/macrophage differentiation. *Genes & Dev.* 7: 2110-2119.
- Ayer, D.E., L. Kretzner, and R.N. Eisenman. 1993. Mad: A heterodimeric partner for Max that antagonizes Myc transcriptional activity. *Cell* 72: 211-222.
- Benezra, R., R.L. Davis, D. Lockshon, D.L. Turner, and H. Weintraub. 1990. The protein Id: A negative regulator of helix-loop-helix DNA binding proteins. *Cell* 61: 49-59.
- Christy, B.A., L.K. Sanders, L.F. Lau, N.G. Copeland, N.A. Jenkins, and D. Nathans. 1991. An Id-related helix-loop-helix protein encoded by a growth factor-inducible gene. *Proc. Natl. Acad. Sci.* 88: 1815-1819.
- Dang, C.V., J. Barrett, M. Villa-Garcia, L.M. Resar, G.J. Kato, and E.R. Fearon. 1991. Intracellular leucine zipper interactions suggest c-Myc hetero-oligomerization. *Mol. Cell. Biol.* 11: 954-962.
- Ema, M., K. Sogawa, N. Watanabe, Y. Chujoh, N. Matsushita, O. Gotoh, Y. Funae, and Y. Fujii-Kuriyama. 1992. cDNA cloning and structure of mouse putative Ah receptor. *Biochem. Biophys. Res. Commun.* 184: 246-253.
- Fernandez-Salguero, P.M., D.M. Hilbert, S. Rudikoff, J.M. Ward, and F.J. Gonzalez. 1996. Aryl-hydrocarbon receptor-deficient mice are resistant to 2,3,7,8-tetrachlorodibenzo-p-dioxin-induced toxicity. *Toxicol. Appl. Pharmacol.* 140: 173-179.
- Fujisawa-Sehara, A., K. Sogawa, M. Yamane, and Y. Fujii-Kuriyama. 1987. Characterization of xenobiotic responsive elements upstream from the drug-metabolizing cytochrome P-450c gene: A similarity to glucocorticoid regulatory elements. *Nucleic Acids Res.* 15: 4179-4191.
- Gekakis, N., D. Staknis, H.B. Nguyen, F.C. Davis, L.D. Wilsbacher, D.P. King, J.S. Takahashi, and C.J. Weitz. 1998. Role of the CLOCK protein in the mammalian circadian mechanism. *Science* 280: 1564-1569.
- Hankinson, O. 1995. The aryl hydrocarbon receptor complex. *Annu. Rev. Pharmacol. Toxicol.* 35: 307-340.
- Hirose, K., M. Morita, M. Ema, J. Mimura, H. Hamada, H. Fujii, Y. Saijo,

- O. Gotoh, K. Sogawa, and Y. Fujii-Kuriyama. 1996. cDNA cloning and tissue-specific expression of a novel basic helix-loop-helix/PAS factor (Arnt2) with close sequence similarity to the aryl hydrocarbon receptor nuclear translocator (Arnt). *Mol. Cell. Biol.* **16**: 1706–1713.
- Hoffman, E.C., H. Reyes, F.F. Chu, F. Sander, L.H. Conley, B.A. Brooks, and O. Hankinson. 1991. Cloning of a factor required for activity of the Ah (dioxin) receptor. *Science* **252**: 954–958.
- Israel, D.I., M.G. Estolano, D.R. Galeazzi, and J. Whitlock, Jr. 1985. Superinduction of cytochrome P1-450 gene transcription by inhibition of protein synthesis in wild type and variant mouse hepatoma cells. *J. Biol. Chem.* **260**: 5648–5653.
- Jackson, F.R., T.A. Bargiello, S.H. Yun, and M.W. Young. 1986. Product of per locus of *Drosophila* shares homology with proteoglycans. *Nature* **320**: 185–188.
- Jen, Y., H. Weintraub, and R. Ben Ezra. 1992. Overexpression of Id protein inhibits the muscle differentiation program in vivo association of Id with E2A proteins. *Genes & Dev.* **6**: 1466–1479.
- Kozak, M. 1987. At least six nucleotides preceding the AUG initiator codon enhance translation in mammalian cells. *J. Mol. Biol.* **196**: 947–950.
- Kubota, M., K. Sogawa, Y. Kaizu, T. Sawaya, J. Watanabe, K. Kawajiri, O. Gotoh, and Y. Fujii-Kuriyama. 1991. Xenobiotic responsive element in the 5'-upstream region of the human P-450c gene. *J. Biochem.* **110**: 232–236.
- Larsson, L.G., M. Pettersson, F. Oberg, K. Nilsson, and B. Lüscher. 1994. Expression of mad, mx1, max and c-myc during induced differentiation of hematopoietic cells: opposite regulation of mad and c-myc. *Oncogene* **9**: 1247–1252.
- Lusska, A., L. Wu, and J. Whitlock, Jr. 1992. Superinduction of CYP1A1 transcription by cycloheximide. Role of the DNA binding site for the liganded Ah receptor. *J. Biol. Chem.* **267**: 15146–15151.
- Ma, Q. and J.P. Whitlock, Jr. 1996. The aromatic hydrocarbon receptor modulates the Hepa 1c1c7 cell cycle and differentiated state independently of dioxin. *Mol. Cell. Biol.* **16**: 2144–2150.
- Matsushita, N., K. Sogawa, M. Ema, A. Yoshida, and Y. Fujii-Kuriyama. 1993. A factor binding to the xenobiotic responsive element (XRE) of P-4501A1 gene consists of at least two helix-loop-helix proteins, Ah receptor and Arnt. *J. Biol. Chem.* **268**: 21002–21006.
- Mimura, J., K. Yamashita, K. Nakamura, M. Morita, T.N. Takagi, K. Nakao, M. Ema, K. Sogawa, M. Yasuda, M. Katsuki, and Y. Fujii-Kuriyama. 1997. Loss of teratogenic response to 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD) in mice lacking the Ah (dioxin) receptor. *Genes Cells* **2**: 645–654.
- Mizushima, S. and S. Nagata. 1990. pEF-BOS, a powerful mammalian expression vector. *Nucleic Acids Res.* **18**: 5322.
- Murre, C., P.S. McCaw, and D. Baltimore. 1989. A new DNA binding and dimerization motif in immunoglobulin enhancer binding, daughterless, MyoD, and myc proteins. *Cell* **56**: 777–783.
- Nambu, J.R., J.O. Lewis, K. Wharton, Jr., and S.T. Crews. 1991. The *Drosophila* single-minded gene encodes a helix-loop-helix protein that acts as a master regulator of CNS midline development. *Cell* **67**: 1157–1167.
- Neuhold, L.A. and B. Wold. 1993. HLH forced dimers: Tethering MyoD to E47 generates a dominant positive myogenic factor insulated from negative regulation by Id. *Cell* **74**: 1033–1042.
- Numayama-Tsuruta, K., A. Kobayashi, K. Sogawa, and Y. Fujii-Kuriyama. 1997. A point mutation responsible for defective function of the aryl-hydrocarbon-receptor nuclear translocator in mutant Hepa-1c1c7 cells. *Eur. J. Biochem.* **246**: 486–495.
- Ogawa, H., S. Inouye, F.I. Tsuji, K. Yasuda, and K. Umesono. 1995. Localization, trafficking, and temperature-dependence of the Aequorea green fluorescent protein in cultured vertebrate cells. *Proc. Natl. Acad. Sci.* **92**: 11899–11903.
- Perdew, G.H. 1988. Association of the Ah receptor with the 90-kDa heat shock protein. *J. Biol. Chem.* **263**: 13802–13805.
- Poland, A. and J.C. Knutson. 1982. 2,3,7,8-tetrachlorodibenzo-p-dioxin and related halogenated aromatic hydrocarbons: examination of the mechanism of toxicity. *Annu. Rev. Pharmacol. Toxicol.* **22**: 517–554.
- Pongratz, I., G.G. Mason, and L. Poellinger. 1992. Dual roles of the 90-kDa heat shock protein hsp90 in modulating functional activities of the dioxin receptor. Evidence that the dioxin receptor functionally belongs to a subclass of nuclear receptors which require hsp90 both for ligand binding activity and repression of intrinsic DNA binding activity. *J. Biol. Chem.* **267**: 13728–13734.
- Sogawa, K., K. Iwabuchi, H. Abe, and Y. Fujii-Kuriyama. 1995. Transcriptional activation domains of the Ah receptor and Ah receptor nuclear translocator. *J. Cancer Res. Clin. Oncol.* **121**: 612–620.
- Sogawa, K. and Y. Fujii-Kuriyama. 1997. Ah receptor, a novel ligand-activated transcription factor. *J. Biochem.* **122**: 1075–1079.
- Swanson, H.I. and C.A. Bradfield. 1993. The AH-receptor: Genetics, structure and function. *Pharmacogenetics* **3**: 213–230.
- Telakowski-Hopkins, C.A., R.G. King, and C.B. Pickett. 1988. Glutathione S-transferase Ya subunit gene: Identification of regulatory elements required for basal level and inducible expression. *Proc. Natl. Acad. Sci.* **85**: 1000–1004.
- Weiss, C., S.K. Kolluri, F. Kiefer, and M. Göttlicher. 1996. Complementation of Ah receptor deficiency in hepatoma cells: negative feedback regulation and cell cycle control by the Ah receptor. *Exp. Cell Res.* **226**: 154–163.
- Zervos, A.S., J. Gyuris, and R. Brent. 1993. Mx1, a protein that specifically interacts with Max to bind Myc-Max recognition sites. *Cell* **72**: 223–232.

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