

**Anti-AHR Antibody**  
**Catalog # ABO11088****Specification**

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**Anti-AHR Antibody - Product Information**

Application	WB
Primary Accession	<a href="#">P35869</a>
Host	Rabbit
Reactivity	Human
Clonality	Polyclonal
Format	Lyophilized

**Description**

Rabbit IgG polyclonal antibody for Aryl hydrocarbon receptor(AHR) detection. Tested with WB in Human.

**Reconstitution**

Add 0.2ml of distilled water will yield a concentration of 500ug/ml.

**Anti-AHR Antibody - Additional Information**

**Gene ID** 196

**Other Names**

Aryl hydrocarbon receptor, Ah receptor, AhR, Class E basic helix-loop-helix protein 76, bHLHe76, AHR, BHLHE76

**Calculated MW**

96147 MW KDa

**Application Details**

Western blot, 0.1-0.5 µg/ml, Human<br>

**Subcellular Localization**

Cytoplasm. Nucleus. Initially cytoplasmic; upon binding with ligand and interaction with a HSP90, it translocates to the nucleus.

**Tissue Specificity**

Expressed in all tissues tested including blood, brain, heart, kidney, liver, lung, pancreas and skeletal muscle. .

**Protein Name**

Aryl hydrocarbon receptor

**Contents**

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na<sub>2</sub>HPO<sub>4</sub>, 0.05mg Thimerosal, 0.05mg NaN<sub>3</sub>.

**Immunogen**

A synthetic peptide corresponding to a sequence at the C-terminus of human AHR (832-848aa HPSEARPFDPDLTSSGFL).

**Purification**

Immunogen affinity purified.

**Cross Reactivity**

No cross reactivity with other proteins

**Storage**

**At -20°C for one year. After r°Constitution, at 4°C for one month. It°Can also be aliquotted and stored frozen at -20°C for a longer time.Avoid repeated freezing and thawing.**

**Sequence Similarities**

Contains 1 bHLH (basic helix-loop-helix) domain.

**Anti-AHR Antibody - Protein Information**

**Name** AHR {ECO:0000303|PubMed:8393992, ECO:0000312|HGNC:HGNC:348}

**Function**

Ligand-activated transcription factor that enables cells to adapt to changing conditions by sensing compounds from the environment, diet, microbiome and cellular metabolism, and which plays important roles in development, immunity and cancer (PubMed:<a href="http://www.uniprot.org/citations/30373764" target="\_blank">30373764</a>, PubMed:<a href="http://www.uniprot.org/citations/23275542" target="\_blank">23275542</a>, PubMed:<a href="http://www.uniprot.org/citations/7961644" target="\_blank">7961644</a>, PubMed:<a href="http://www.uniprot.org/citations/32818467" target="\_blank">32818467</a>). Upon ligand binding, translocates into the nucleus, where it heterodimerizes with ARNT and induces transcription by binding to xenobiotic response elements (XRE) (PubMed:<a href="http://www.uniprot.org/citations/30373764" target="\_blank">30373764</a>, PubMed:<a href="http://www.uniprot.org/citations/23275542" target="\_blank">23275542</a>, PubMed:<a href="http://www.uniprot.org/citations/7961644" target="\_blank">7961644</a>). Regulates a variety of biological processes, including angiogenesis, hematopoiesis, drug and lipid metabolism, cell motility and immune modulation (PubMed:<a href="http://www.uniprot.org/citations/12213388" target="\_blank">12213388</a>). Xenobiotics can act as ligands: upon xenobiotic- binding, activates the expression of multiple phase I and II xenobiotic chemical metabolizing enzyme genes (such as the CYP1A1 gene) (PubMed:<a href="http://www.uniprot.org/citations/7961644" target="\_blank">7961644</a>). Mediates biochemical and toxic effects of halogenated aromatic hydrocarbons (PubMed:<a href="http://www.uniprot.org/citations/7961644" target="\_blank">7961644</a>, PubMed:<a href="http://www.uniprot.org/citations/34521881" target="\_blank">34521881</a>). Next to xenobiotics, natural ligands derived from plants, microbiota, and endogenous metabolism are potent AHR agonists (PubMed:<a href="http://www.uniprot.org/citations/18076143" target="\_blank">18076143</a>). Tryptophan (Trp) derivatives constitute an important class of endogenous AHR ligands (PubMed:<a href="http://www.uniprot.org/citations/32866000" target="\_blank">32866000</a>, PubMed:<a href="http://www.uniprot.org/citations/32818467" target="\_blank">32818467</a>). Acts as a negative regulator of anti-tumor immunity: indoles and kynurenic acid generated by Trp catabolism act as ligand and activate AHR, thereby promoting AHR-driven cancer cell motility and suppressing adaptive immunity (PubMed:<a href="http://www.uniprot.org/citations/32818467" target="\_blank">32818467</a>). Regulates the circadian clock by inhibiting the basal and circadian expression of the core circadian component PER1 (PubMed:<a href="http://www.uniprot.org/citations/28602820" target="\_blank">28602820</a>). Inhibits PER1 by repressing the CLOCK-BMAL1 heterodimer mediated transcriptional activation of PER1 (PubMed:<a href="http://www.uniprot.org/citations/28602820" target="\_blank">28602820</a>). The heterodimer ARNT:AHR binds to core DNA sequence 5'-TGCGTG-3' within the dioxin response

element (DRE) of target gene promoters and activates their transcription (PubMed:<a href="http://www.uniprot.org/citations/28602820" target="\_blank">28602820</a>).

#### Cellular Location

Cytoplasm. Nucleus. Note=Initially cytoplasmic; upon binding with ligand and interaction with a HSP90, it translocates to the nucleus.

#### Tissue Location

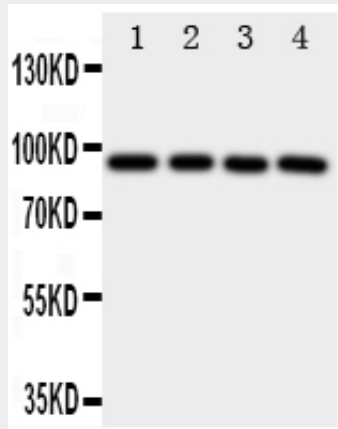
Expressed in all tissues tested including blood, brain, heart, kidney, liver, lung, pancreas and skeletal muscle Expressed in retinal photoreceptors (PubMed:29726989)

### Anti-AHR Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### Anti-AHR Antibody - Images



Anti-AHR antibody, ABO11088, Western blotting All lanes: Anti AHR (ABO11088) at 0.5ug/ml Lane 1: PANC Whole Cell Lysate at 40ug Lane 2: HELA Whole Cell Lysate at 40ug Lane 3: MCF-7 Whole Cell Lysate at 40ug Lane 4: HT1080 Whole Cell Lysate at 40ug Predicted bind size: 96KD Observed bind size: 96KD

### Anti-AHR Antibody - Background

AHR(aryl hydrocarbon receptor), also called bHLHe76, is a member of the family of basic helix-loop-helix transcription factors. Ahr is a cytosolic transcription factor that is normally inactive, bound to several co-chaperones. The AHR gene is mapped on 7p21.1. Estrogenic actions of AHR agonists were detected in wildtype ovariectomized mouse uteri, but were absent in Ahr<sup>-/-</sup> or Er- $\alpha$ <sup>-/-</sup> ovariectomized mice. Complex assembly and ubiquitin ligase activity of CUL4B(AHR) in vitro and in vivo are dependent on the AHR ligand. In the CUL4B(AHR) complex, ligand-activated AHR acts as a substrate-specific adaptor component that targets sex steroid

receptors for degradation.  $\text{Ahr}^{-/-}$  Cd4-positive cells from mice lacking  $\text{Ahr}$  developed Th17 responses but failed to produce IL22 and did not show enhanced Th17 development. Activation of  $\text{Ahr}$  during induction of EAE accelerated disease onset and increased pathology in wildtype mice, but not in  $\text{Ahr}^{-/-}$  mice. The TDO-AHR pathway is active in human brain tumors and is associated with malignant progression and poor survival.  $\text{Ahr}$  activity within ROR- $\gamma$ -t-positive ILC could be induced by dietary ligands such as those contained in vegetables of the family Brassicaceae.