

Goat Anti-Farnesoid X receptor / FXR Antibody
Peptide-affinity purified goat antibody
Catalog # AF1401a

Specification

Goat Anti-Farnesoid X receptor / FXR Antibody - Product Information

Application	WB
Primary Accession	O96RI1
Other Accession	NP_005114 , 9971
Reactivity	Human
Predicted	Dog
Host	Goat
Clonality	Polyclonal
Concentration	100ug/200ul
Isotype	IgG
Calculated MW	55914

Goat Anti-Farnesoid X receptor / FXR Antibody - Additional Information

Gene ID 9971

Other Names

Bile acid receptor, Farnesoid X-activated receptor, Farnesol receptor HRR-1, Nuclear receptor subfamily 1 group H member 4, Retinoid X receptor-interacting protein 14, RXR-interacting protein 14, NR1H4, BAR, FXR, HRR1, RIP14

Format

0.5 mg IgG/ml in Tris saline (20mM Tris pH7.3, 150mM NaCl), 0.02% sodium azide, with 0.5% bovine serum albumin

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

Goat Anti-Farnesoid X receptor / FXR Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Goat Anti-Farnesoid X receptor / FXR Antibody - Protein Information

Name NR1H4

Synonyms BAR, FXR, HRR1, RIP14

Function

Ligand-activated transcription factor. Receptor for bile acids (BAs) such as chenodeoxycholic acid (CDCA), lithocholic acid, deoxycholic acid (DCA) and allocholic acid (ACA). Plays a essential role in

BA homeostasis through the regulation of genes involved in BA synthesis, conjugation and enterohepatic circulation. Also regulates lipid and glucose homeostasis and is involved innate immune response (PubMed: [10334992](http://www.uniprot.org/citations/10334992) [10334993](http://www.uniprot.org/citations/10334993) [21383957](http://www.uniprot.org/citations/21383957) [22820415](http://www.uniprot.org/citations/22820415)). The FXR-RXR heterodimer binds predominantly to farnesoid X receptor response elements (FXREs) containing two inverted repeats of the consensus sequence 5'-AGGTCA-3' in which the monomers are spaced by 1 nucleotide (IR-1) but also to tandem repeat DR1 sites with lower affinity, and can be activated by either FXR or RXR-specific ligands. It is proposed that monomeric nuclear receptors such as NR5A2/LRH-1 bound to coregulatory nuclear responsive element (NRE) halfsites located in close proximity to FXREs modulate transcriptional activity (By similarity). In the liver activates transcription of the corepressor NR0B2 thereby indirectly inhibiting CYP7A1 and CYP8B1 (involved in BA synthesis) implicating at least in part histone demethylase KDM1A resulting in epigenomic repression, and SLC10A1/NTCP (involved in hepatic uptake of conjugated BAs). Activates transcription of the repressor MAFG (involved in regulation of BA synthesis) (By similarity). Activates transcription of SLC27A5/BACS and BAAT (involved in BA conjugation), ABCB11/BSEP (involved in bile salt export) by directly recruiting histone methyltransferase CARM1, and ABCC2/MRP2 (involved in secretion of conjugated BAs) and ABCB4 (involved in secretion of phosphatidylcholine in the small intestine) (PubMed: [12754200](http://www.uniprot.org/citations/12754200) [15471871](http://www.uniprot.org/citations/15471871) [17895379](http://www.uniprot.org/citations/17895379)). Activates transcription of SLC27A5/BACS and BAAT (involved in BA conjugation), ABCB11/BSEP (involved in bile salt export) by directly recruiting histone methyltransferase CARM1, and ABCC2/MRP2 (involved in secretion of conjugated BAs) and ABCB4 (involved in secretion of phosphatidylcholine in the small intestine) (PubMed: [10514450](http://www.uniprot.org/citations/10514450) [15239098](http://www.uniprot.org/citations/15239098) [16269519](http://www.uniprot.org/citations/16269519)). In the intestine activates FGF19 expression and secretion leading to hepatic CYP7A1 repression (PubMed: [12815072](http://www.uniprot.org/citations/12815072) [19085950](http://www.uniprot.org/citations/19085950)). The function also involves the coordinated induction of hepatic KLB/beta-klotho expression (By similarity). Regulates transcription of liver UGT2B4 and SULT2A1 involved in BA detoxification; binding to the UGT2B4 promoter seems to imply a monomeric transactivation independent of RXRA (PubMed: [12806625](http://www.uniprot.org/citations/12806625) [16946559](http://www.uniprot.org/citations/16946559)). Modulates lipid homeostasis by activating liver NR0B2/SHP-mediated repression of SREBF1 (involved in de novo lipogenesis), expression of PLTP (involved in HDL formation), SCARB1 (involved in HDL hepatic uptake), APOE, APOC1, APOC4, PPARA (involved in beta-oxidation of fatty acids), VLDLR and SDC1 (involved in the hepatic uptake of LDL and IDL remnants), and inhibiting expression of MTP (involved in VLDL assembly (PubMed: [12554753](http://www.uniprot.org/citations/12554753) [12660231](http://www.uniprot.org/citations/12660231) [15337761](http://www.uniprot.org/citations/15337761)). Increases expression of APOC2 (promoting lipoprotein lipase activity implicated in triglyceride clearance) (PubMed: [11579204](http://www.uniprot.org/citations/11579204)). Transrepresses APOA1 involving a monomeric competition with NR2A1 for binding to a DR1 element (PubMed: [11927623](http://www.uniprot.org/citations/11927623) [21804189](http://www.uniprot.org/citations/21804189)). Also reduces triglyceride clearance by inhibiting expression of ANGPTL3 and APOC3 (both involved in inhibition of lipoprotein lipase) (PubMed: [12891557](http://www.uniprot.org/citations/12891557)). Involved in glucose homeostasis by modulating hepatic gluconeogenesis through activation of NR0B2/SHP-mediated repression of respective genes. Modulates glycogen synthesis (inducing phosphorylation of glycogen synthase kinase-3) (By similarity). Modulates glucose-stimulated insulin secretion and

is involved in insulin resistance (PubMed:20447400). Involved in intestinal innate immunity. Plays a role in protecting the distal small intestine against bacterial overgrowth and preservation of the epithelial barrier (By similarity). Down-regulates inflammatory cytokine expression in several types of immune cells including macrophages and mononuclear cells (PubMed:21242261). Mediates trans- repression of TLR4-induced cytokine expression; the function seems to require its sumoylation and prevents N-CoR nuclear receptor corepressor clearance from target genes such as IL1B and NOS2 (PubMed:19864602). Involved in the TLR9-mediated protective mechanism in intestinal inflammation. Plays an anti-inflammatory role in liver inflammation; proposed to inhibit pro-inflammatory (but not antiapoptotic) NF-kappa-B signaling) (By similarity).

Cellular Location

Nucleus. [Isoform 2]: Nucleus [Isoform 4]: Nucleus

Tissue Location

Liver and hepatocyte-related cells express mainly FXRalpha1-type isoforms with isoform 3 and isoform 4 in approximately equal proportions. In intestine and kidney mainly FXRalpha2-type isoforms are expressed with isoform 1 and isoform 2 in approximately equal proportions. Expressed in pancreatic beta cells and macrophages

Goat Anti-Farnesoid X receptor / FXR 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](#)

Goat Anti-Farnesoid X receptor / FXR Antibody - Images



AF1401a (1 µg/ml) staining of Human Lung lysate (35 µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

Goat Anti-Farnesoid X receptor / FXR Antibody - References

The nuclear receptor FXR is expressed in pancreatic beta-cells and protects human islets from lipotoxicity. Popescu IR, et al. FEBS Lett, 2010 Jul 2. PMID 20447400.

Bile acids and their nuclear receptor FXR: Relevance for hepatobiliary and gastrointestinal disease. Gadaleta RM, et al. Biochim Biophys Acta, 2010 Jul. PMID 20399894.

Genetic risk factors for hepatopulmonary syndrome in patients with advanced liver disease. Roberts KE, et al. Gastroenterology, 2010 Jul. PMID 20346360.

Regulation of the human bile acid UDP-glucuronosyltransferase 1A3 by the farnesoid X receptor and bile acids. Erichsen TJ, et al. J Hepatol, 2010 Apr. PMID 20189675.

Farnesoid X receptor, through the binding with steroidogenic factor 1-responsive element, inhibits aromatase expression in tumor Leydig cells. Catalano S, et al. J Biol Chem, 2010 Feb 19. PMID 20026603.