

AKR1C2 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab)

Catalog # AP12246B

Product Information

Application	WB, IHC-P, E
Primary Accession	P52895
Other Accession	Q95JH7 , Q04828 , NP_995317.1
Reactivity	Human
Predicted	Monkey
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Clone Names	RB32249
Calculated MW	36735
Antigen Region	296-323

Additional Information

Gene ID	1646
Other Names	Aldo-keto reductase family 1 member C2, 1---, 3-alpha-HSD3, Chlordecone reductase homolog HAKRD, Dihydrodiol dehydrogenase 2, DD-2, DD2, Dihydrodiol dehydrogenase/bile acid-binding protein, DD/BABP, Trans-1, 2-dihydrobenzene-1, 2-diol dehydrogenase, Type III 3-alpha-hydroxysteroid dehydrogenase, AKR1C2, DDH2
Target/Specificity	This AKR1C2 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 296-323 amino acids from the C-terminal region of human AKR1C2.
Dilution	WB~~1:2000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.
Format	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	AKR1C2 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	AKR1C2
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Synonyms

DDH2

Function

Cytosolic aldo-keto reductase that catalyzes the NADH and NADPH-dependent reduction of ketosteroids to hydroxysteroids (PubMed:[19218247](#)). Most probably acts as a reductase in vivo since the oxidase activity measured in vitro is inhibited by physiological concentrations of NADPH (PubMed:[14672942](#)). Displays a broad positional specificity acting on positions 3, 17 and 20 of steroids and regulates the metabolism of hormones like estrogens and androgens (PubMed:[10998348](#)). Works in concert with the 5-alpha/5-beta-steroid reductases to convert steroid hormones into the 3-alpha/5-alpha and 3- alpha/5-beta-tetrahydrosteroids. Catalyzes the inactivation of the most potent androgen 5-alpha-dihydrotestosterone (5-alpha-DHT) to 5-alpha-androstane-3-alpha,17-beta-diol (3-alpha-diol) (PubMed:[15929998](#), PubMed:[17034817](#), PubMed:[17442338](#), PubMed:[8573067](#)). Also specifically able to produce 17beta-hydroxy-5alpha-androstan-3-one/5alphaDHT (PubMed:[10998348](#)). May also reduce conjugated steroids such as 5alpha-dihydrotestosterone sulfate (PubMed:[19218247](#)). Displays affinity for bile acids (PubMed:[8486699](#)).

Cellular Location

Cytoplasm, cytosol.

Tissue Location

Expressed in fetal testes. Expressed in fetal and adult adrenal glands.

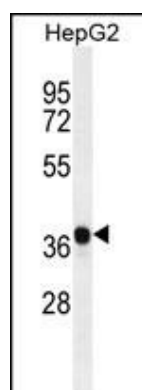
Background

This gene encodes a member of the aldo/keto reductase superfamily, which consists of more than 40 known enzymes and proteins. These enzymes catalyze the conversion of aldehydes and ketones to their corresponding alcohols using NADH and/or NADPH as cofactors. The enzymes display overlapping but distinct substrate specificity. This enzyme binds bile acid with high affinity, and shows minimal 3-alpha-hydroxysteroid dehydrogenase activity. This gene shares high sequence identity with three other gene members and is clustered with those three genes at chromosome 10p15-p14.

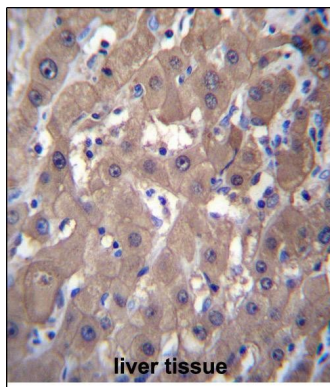
References

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Wang, X., et al. PLoS ONE 5 (8), E11934 (2010) :
Reding, K.W., et al. Am. J. Epidemiol. 170(10):1241-1249(2009)
Cogliati, C., et al. FEBS J. 276(20):6011-6023(2009)
Davies, N.J., et al. Cancer Res. 69(11):4769-4775(2009)

Images



AKR1C2 Antibody (C-term) (Cat. #AP12246b) western blot analysis in HepG2 cell line lysates (35ug/lane). This demonstrates the AKR1C2 antibody detected the AKR1C2 protein (arrow).



AKR1C2 Antibody (C-term) (Cat. #AP12246b) immunohistochemistry analysis in formalin fixed and paraffin embedded human liver tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of AKR1C2 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.

Citations

- [Effects of Curcumin Combined With the 5-alpha Reductase Inhibitor Dutasteride on LNCaP Prostate Cancer Cells](#)
- [Modulation of AKR1C2 by curcumin decreases testosterone production in prostate cancer.](#)

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