

# AKR1C2 Antibody

Purified Rabbit Polyclonal Antibody (Pab)  
Catalog # AP53284

## Product Information

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<b>Application</b>	WB
<b>Primary Accession</b>	<a href="#">P52895</a>
<b>Reactivity</b>	Human
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Calculated MW</b>	36735

## Additional Information

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<b>Gene ID</b>	1646
<b>Other Names</b>	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, 1.3.1.20, Type III 3-alpha-hydroxysteroid dehydrogenase, 1.1.1.357, AKR1C2, DDH2
<b>Target/Specificity</b>	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human AKR1C2. The exact sequence is proprietary.
<b>Dilution</b>	WB~~ 1:1000
<b>Format</b>	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.09% (W/V) sodium azide and 50% glycerol
<b>Storage</b>	Store at -20 °C.Stable for 12 months from date of receipt

## Protein Information

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<b>Name</b>	AKR1C2
<b>Synonyms</b>	DDH2
<b>Function</b>	Cytosolic aldo-keto reductase that catalyzes the NADH and NADPH-dependent reduction of ketosteroids to hydroxysteroids (PubMed: <a href="#">19218247</a> ). Most probably acts as a reductase in vivo since the oxidase activity measured in vitro is inhibited by physiological concentrations of NADPH (PubMed: <a href="#">14672942</a> ). 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: <a href="#">10998348</a> ). 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.

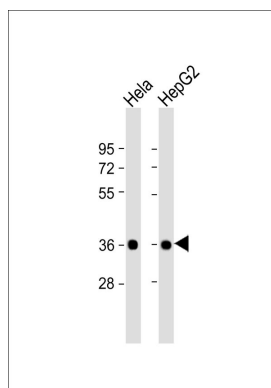
## Background

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). Has a high bile-binding ability.

## References

- Qin K.-N.,et al.J. Steroid Biochem. Mol. Biol. 46:673-679(1993).  
Ciaccio P.J.,et al.Biochim. Biophys. Acta 1186:129-132(1994).  
Qin K.-N.,et al.Gene 149:357-361(1994).  
Dufort I.,et al.Biochem. Biophys. Res. Commun. 228:474-479(1996).  
Shiraishi H.,et al.Biochem. J. 334:399-405(1998).

## Images



All lanes : Anti-AKR1C2 Antibody at 1:1000 dilution Lane 1: HeLa whole cell lysate Lane 2: HepG2 whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 37 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.