# **Human AKR1C2 Knockdown Cell Line (WB-Validated)**



**Catalog #: C67575** 

#### **Aliases**

AKR1C2; Aldo-Keto Reductase Family 1 Member C2; DD2; HAKRD; MCDR2; BABP; DDH2; DD; Dihydrodiol Dehydrogenase 2; Bile Acid Binding Protein; 3-Alpha Hydroxysteroid Dehydrogenase, Type III; Testicular 17,20-Desmolase Deficiency; Chlordecone Reductase Homolog HAKRD; 3-Alpha-HSD3; DD/BABP; DD-2; TDD; Aldo-Keto Reductase Family 1, Member C2 (Dihydrodiol Dehydrogenase 2; Bile Acid Binding Protein; 3-Alpha Hydroxysteroid Dehydrogenase, Type III); Dihydrodiol Dehydrogenase/Bile Acid-Binding Protein; Trans-1,2-Dihydrobenzene-1,2-Diol Dehydrogenase; Type III 3-Alpha-Hydroxysteroid Dehydrogenase; Type II Dihydrodiol Dehydrogenase; Pseudo-Chlordecone Reductase; Dihydrodiol Dehydrogenase 2; AKR1C-Pseudo; EC 1.1.1.112; EC 1.1.1.209; EC 1.1.1.357; EC 1.1.1.53; EC 1.1.1.62; EC 1.3.1.20; EC 1.-..-; EC 1.1.1; SRXY8; HBAB

## **Background**

Gene Name: AKR1C2 NCBI Gene Entry: 1646

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human AKR1C2 Knockdown Cell Line (Wb-Validated)
- 2. Wild-type cell line

### **Parental Cell Line**

Human cell line supplied by the client

### Validation Methods

RT-qPCR, Western blotting (WB)

## **Shipping**

Shipped on Dry Ice.

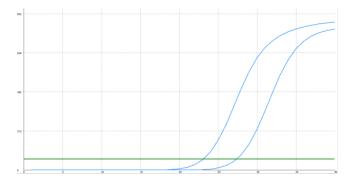
## **Instructions For Use**

This knockdown cell line should be paired with wild-type cell line for use.

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**Note:** This product is for research use only.

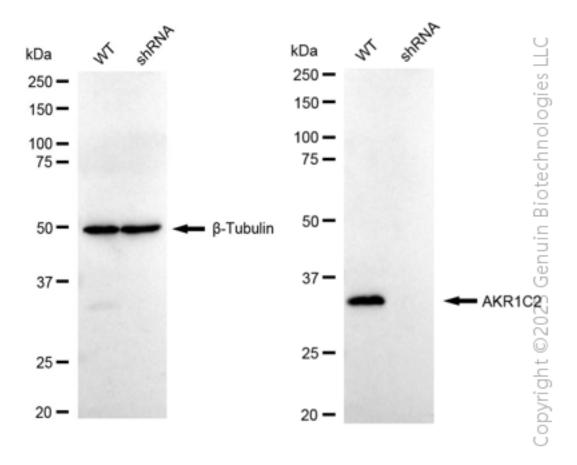
#### **Validation Data**



Genotype	Ct Value
Wild-Type	23.04
Knock-Down	27.21
ΔCt (CtKD-CtWT)	4.17
% mRNA	yright
Reduction	94% <sup>§</sup>

RT-qPCR analysis. HeLa cells were infected with AKR1C2-specific shRNA lentiviral particles, total RNA was extracted from wild-type and knockdown cells, RT-qPCR was performed using gene-specific primers.  $\Delta$ Ct (CtKD-CtWT) was used to calculate mRNA reduction (%) between wild-type and knockdown cells using the following formula:  $(1-1/2\Delta$ Ct) x 100%.

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Western blotting analysis. AKR1C2 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting.  $\beta$ -Tubulin served as a loading control. The blots were incubated with primary antibodies against AKR1C2 and  $\beta$ -Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit.