# **Human PCBD1 Knockdown Cell Line (WB-Validated)**



**Catalog #: C62427** 

#### **Aliases**

PCBD1; Pterin-4 Alpha-Carbinolamine DehydRatase 1; PCD; DCOH; PCBD; 6-Pyruvoyl-Tetrahydropterin Synthase/Dimerization Cofactor Of Hepatocyte Nuclear Factor 1 Alpha (TCF1); Pterin-4 Alpha-Carbinolamine DehydRatase/Dimerization Cofactor Of Hepatocyte Nuclear Factor 1 Alpha; Phenylalanine Hydroxylase-Stimulating Protein; 4-Alpha-Hydroxy-Tetrahydropterin DehydRatase; Pterin-4-Alpha-Carbinolamine DehydRatase; Dimerizing Cofactor For HNF1; EC 4.2.1.96; PHS; Pterin-4 Alpha-Carbinolamine DehydRatase/Dimerization Cofactor Of Hepatocyte Nuclear Factor 1 Alpha (TCF1); Pterin-4a-Carbinolamine DehydRatase (Dimerization Cofactor Of Hepatocyte Nuclear Factor 1-Alpha); Dimerization Cofactor Of Hepatocyte Nuclear Factor 1-Alpha; Pterin-4-Alpha Carbinolamine DehydRatase; Pterin Carbinolamine DehydRatase; Dimerization Cofactor Of HNF1; DCoH

## **Background**

Gene Name: PCBD1 NCBI Gene Entry: 5092

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human PCBD1 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.

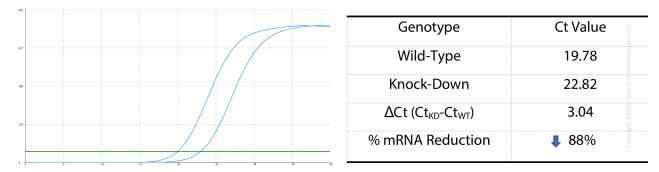
## **Instructions For Use**

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

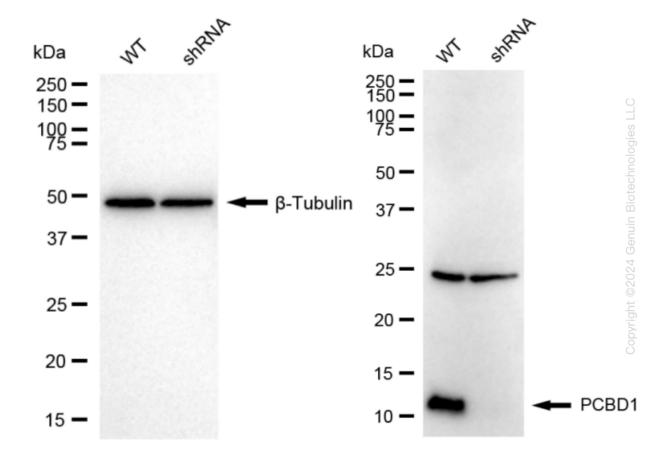
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**Note:** This product is for research use only.

#### **Validation Data**



RT-qPCR analysis. HeLa cells were infected with PCBD1-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%.



Western blotting analysis. PCBD1 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 PCBD1 and  $\beta$ -Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were

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developed using FeQ $^{\text{TM}}$  ECL Substrate Kit.