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



**Catalog #: C62452** 

#### **Aliases**

PPIA; Peptidylprolyl Isomerase A; CYPA; Peptidyl-Prolyl Cis-Trans Isomerase A; Cyclosporin A-Binding Protein; Cyclophilin A; Rotamase A; EC 5.2.1.8; PPIase A; Epididymis Secretory Sperm Binding Protein Li 69p; Peptidylprolyl Isomerase A (Cyclophilin A); T Cell Cyclophilin; HEL-S-69p; CYPH

## **Background**

Gene Name: PPIA

NCBI Gene Entry: 5478

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human PPIA 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.

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

#### Validation Data

Ct Value

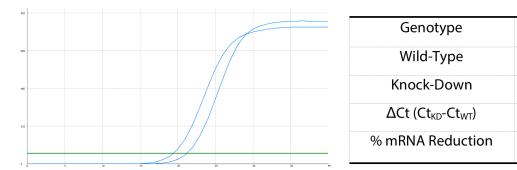
19.26

21.13

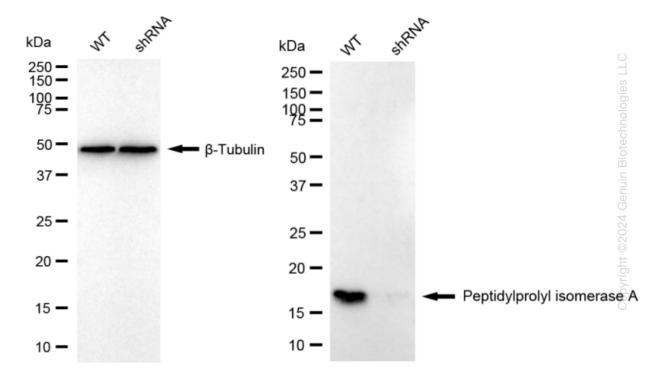
1.87

**J** 73%

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RT-qPCR analysis. HeLa cells were infected with PPIA-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. PPIA 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 PPIA 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.