## **Human CASP3 Knockdown Cell Line (WB-Validated)**



**Catalog #: C68673** 

#### **Aliases**

CASP3; Caspase 3; Apoptain; CPP32; CPP32B; Caspase 3, Apoptosis-Related Cysteine Peptidase; Caspase 3, Apoptosis-Related Cysteine Protease; SREBP Cleavage Activity 1; Cysteine Protease CPP32; Protein Yama; EC 3.4.22.56; Caspase-3; CASP-3; CPP-32; SCA-1; Yama; PARP Cleavage Protease; Procaspase3; EC 3.4.22

### **Background**

Gene Name: CASP3 NCBI Gene Entry: 836

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human CASP3 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.

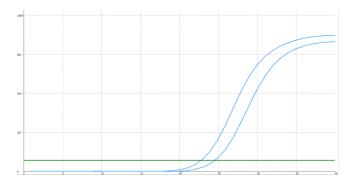
#### **Instructions For Use**

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

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

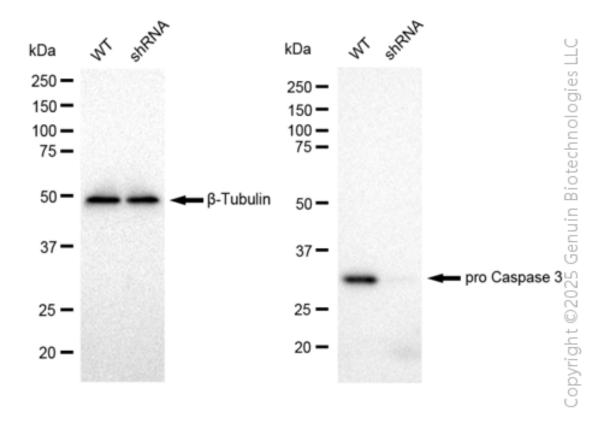
#### Validation Data

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Genotype	Ct Value	
Wild-Type	22.50	
Knock-Down	<b>24.20</b>	
∆Ct (CtKD-CtWT)	1.70	
% mRNA	yright (	•
Reduction	69% ੈ	

RT-qPCR analysis. HeLa cells were infected with CASP3-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. CASP3 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 CASP3 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.