# **Human STING1 Knockdown Cell Line (WB-Validated)**



**Catalog #: C1236** 

#### **Aliases**

STING1; Stimulator Of Interferon Response CGAMP Interactor 1; STING; ERIS; MITA; Transmembrane; Protein 173; TMEM173; NET23; MPYS; Endoplasmic Reticulum Interferon Stimulator; Stimulator Of Interferon Genes Protein; Endoplasmic Reticulum IFN Stimulator; FLJ38577; HSTING; HMITA; N-Terminal Methionine-Proline-Tyrosine-Serine Plasma Membrane Tetraspanner; Mitochondrial Mediator Of IRF3 Activation; Stimulator Of Interferon Protein; Stimulator Of Interferon Gene; Mediator Of IRF3 Activation; STING-Beta; Sting 1; SAVI

## **Background**

Gene Name: STING1 NCBI Gene Entry: 340061

### **Storage**

Store at liquid nitrogen for 1 year.

### **Kit Components**

- 1. Human STING1 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.

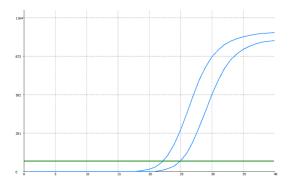
#### **Instructions For Use**

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

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

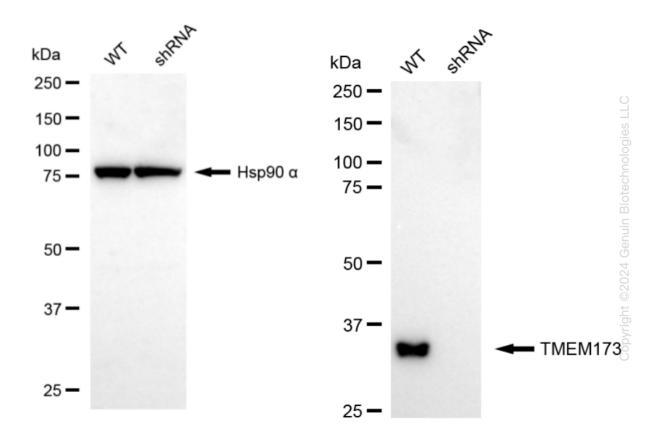
#### Validation Data

# **Human STING1 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	22.08
Knock-Down	24.71
ΔCt (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	2.63
% mRNA Reduction	<b>♣</b> 84%

RT-qPCR analysis. HeLa cells were infected with STING1-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. STING1 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. Hsp90  $\alpha$  served as a loading control. The blots were incubated with primary antibodies against STING1 and Hsp90  $\alpha$ , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit.