# **Human MTA2 Knockdown Cell Line (WB-Validated)**



**Catalog #: C62344** 

#### **Aliases**

Metastasis Associated 1 Family Member 2; MTA1L1; Metastasis Associated Gene Family, Member 2; P53 Target Protein In Deacetylase Complex; Metastasis-Associated Protein MTA2; Metastasis-Associated 1-Like 1; MTA1-L1 Protein; MTA1-L1; PID; Metastasis -Associated Gene 1-Like 1; Metastasis-Associated Protein 2

### **Background**

Gene Name: MTA2 NCBI Gene Entry: 9219

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human MTA2 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.

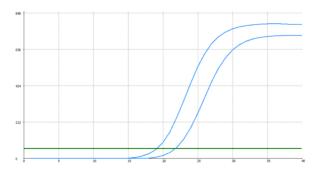
#### **Instructions For Use**

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

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

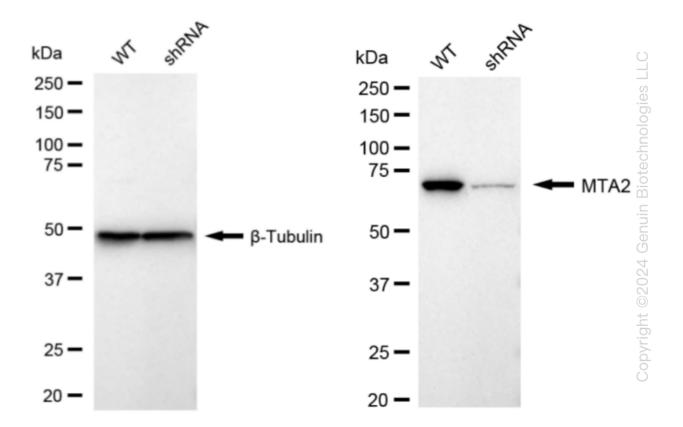
#### Validation Data

## **Human MTA2 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	19.08
Knock-Down	21.56
ΔCt (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	2.48
% mRNA Reduction	<b>↓ 82%</b>
	Spyri

RT-qPCR analysis. HeLa cells were infected with MTA2-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. MTA2 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 (Cat#62344, 1:5,000) against MTA2 and  $\beta$ -Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody (Cat#201, 1:20,000). Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit (Cat#226).

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