# **Human GLB1 Knockdown Cell Line (WB-Validated)**



# **Catalog #: C62411**

#### **Aliases**

GLB1; Galactosidase Beta 1; ELNR1; EBP; Elastin Receptor 1, 67kDa; Elastin Binding Protein; Acid; Beta-Galactosidase; Beta-Galactosidase; EC 3.2.1.23; Lactase; Elastin Receptor 1 (67kD); Galactosidase, Beta 1; Elastin Receptor 1; MPS4B

## **Background**

Gene Name: GLB1 NCBI Gene Entry: 2720

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human GLB1 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.

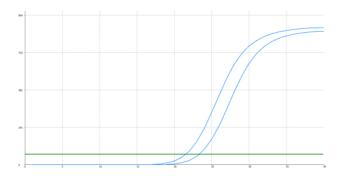
### **Instructions For Use**

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

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

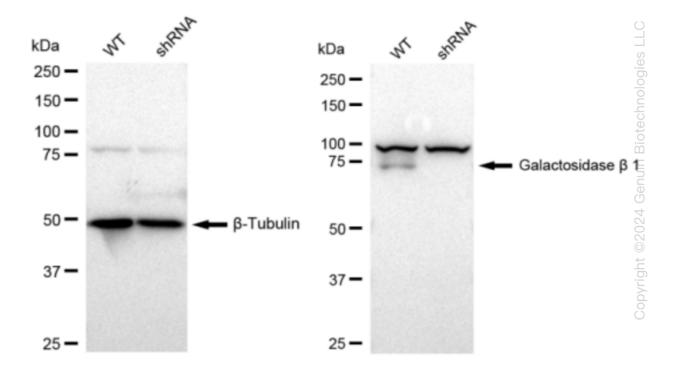
#### **Validation Data**

# **Human GLB1 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	21.39
Knock-Down	23.15
$\Delta Ct (Ct_{KD}-Ct_{WT})$	1.76
% mRNA Reduction	<b>↓</b> 70%

RT-qPCR analysis. HeLa cells were infected with GLB1-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. GLB1 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. β-Tubulin served as a loading control. The blots were incubated with primary antibodies (Cat#62411, 1:5,000) against GLB1 and β-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).