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



**Catalog #: C62009** 

#### **Aliases**

GLA; Galactosidase Alpha; GALA; Galactosylgalactosylglucosylceramidase GLA; Alpha-D-Galactosidase A; Alpha-Galactosidase A; EC 3.2.1.22; Melibiase; Alpha-D-Galactoside Galactohydrolase 1; Alpha-D-Galactoside Galactohydrolase; Agalsidase Alfa; Alpha-Gal A; Agalsidase; EC 3.2.1

### **Background**

Gene Name: GLA

NCBI Gene Entry: 2717

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human GLA 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.

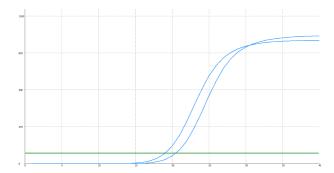
#### **Instructions For Use**

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

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

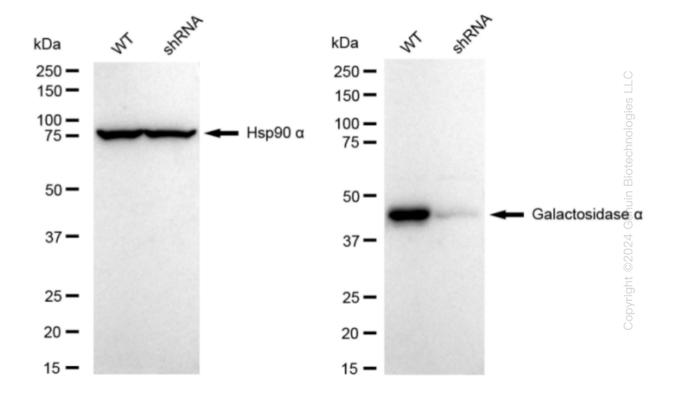
#### Validation Data

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



Genotype	Ct Value
Wild-Type	18.75
Knock-Down	20.26
ΔCt (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	1.51
% mRNA Reduction	<b>4</b> 65%

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