# **Human CLP1 Knockdown Cell Line (WB-Validated)**



## **Catalog #: C61566**

#### **Aliases**

CLP1; Cleavage Factor Polyribonucleotide Kinase Subunit 1; HEAB; Cleavage And Polyadenylation Factor I Subunit 1; Polyribonucleotide 5'-Hydroxyl-Kinase Clp1; Pre-MRNA Cleavage Complex II Protein Clp1; Polyadenylation Factor Clp1; Polynucleotide Kinase Clp1; EC 2.7.1.78; HClp1; CLP1, Cleavage And Polyadenylation Factor I Subunit, Homolog (S. Cerevisiae); Polyribonucleotide 5'-Hydroxyl-Kinase; Homolog Of Yeast CFIA Subunit Clp1p; ATP/GTP-Binding Protein; ATP/GTPbinding Protein; HCLP1

# **Background**

Gene Name: CLP1

NCBI Gene Entry: 10978

## **Storage**

Store at liquid nitrogen for 1 year.

# **Kit Components**

- 1. Human CLP1 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.

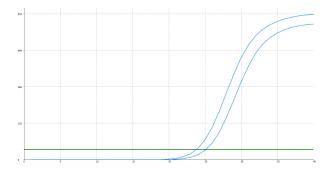
### **Instructions For Use**

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

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

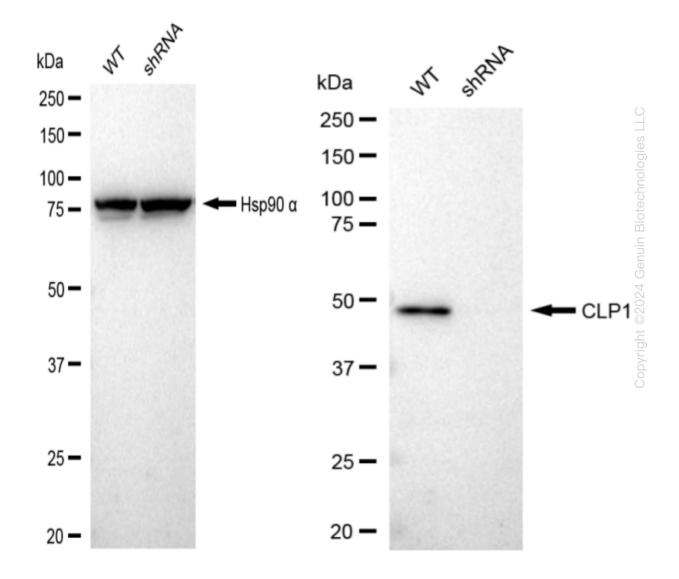
#### Validation Data

# **Human CLP1 Knockdown Cell Line (WB-Validated)**



| Genotype                      | Ct Value |  |
|-------------------------------|----------|--|
| Wild-Type                     | 23.83    |  |
| Knock-Down                    | 25.03    |  |
| $\Delta Ct (Ct_{KD}-Ct_{WT})$ | 1.2      |  |
| % mRNA Reduction              | J 57%    |  |

RT-qPCR analysis. HeLa cells were infected with CLP1-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%.



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# **Human CLP1 Knockdown Cell Line (WB-Validated)**

Western blotting analysis. CLP1 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 (Cat#61566, 1:5,000) against CLP1 and Hsp90  $\alpha$ , 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).