# **Human RBBP4 Knockdown Cell Line (WB-Validated)**



**Catalog #: C62522** 

#### **Aliases**

RB Binding Protein 4, Chromatin Remodeling Factor; Retinoblastoma-Binding Protein 4; RbAp48; NURF55; Lin-53; Nucleosome-Remodeling Factor Subunit RBAP48; Chromatin Assembly Factor I P48 Subunit; Chromatin Assembly Factor 1 Subunit C; Retinoblastoma-Binding Protein P48; Histone-Binding Protein RBBP4; CAF-I 48 KDa Subunit; CAF-1 Subunit C; CAF-I P48; RBBP-4; Chromatin Assembly Factor/CAF-1 P48 Subunit; Retinoblastoma Binding Protein 4; MSI1 Protein Homolog; RBAP48

### **Background**

Gene Name: RBBP4 NCBI Gene Entry: 5928

### **Storage**

Store at liquid nitrogen for 1 year.

# **Kit Components**

- 1. Human RBBP4 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.

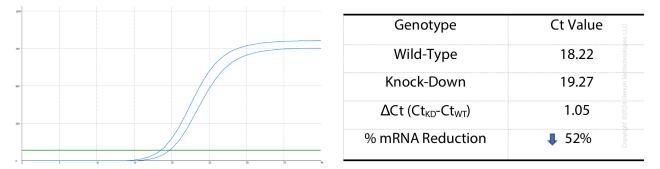
#### **Instructions For Use**

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

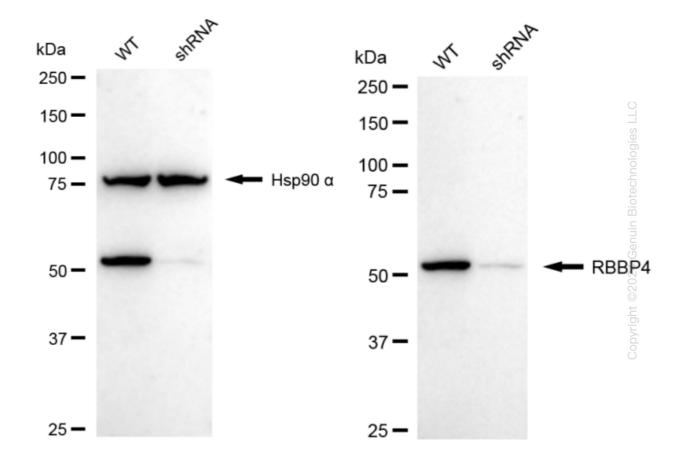
**Note:** This product is for research use only.

#### Validation Data

# **Human RBBP4 Knockdown Cell Line (WB-Validated)**



RT-qPCR analysis. HeLa cells were infected with RBBP4-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. RBBP4 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 RBBP4 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.