# **Human RPA1 Knockdown Cell Line (WB-Validated)**



**Catalog #: C61865** 

#### **Aliases**

RPA1; Replication Protein A1; REPA1; RPA70; HSSB; RF-A; RP-A; Replication Protein A 70 KDa DNA-Binding Subunit; Single-Stranded DNA-Binding Protein; Replication Factor A Protein 1; Replication Protein A1, 70kDa; RF-A Protein 1; RP-A P70; Replication Protein A1 (70kD); PFBMFT6; MSTP075; MST075

## **Background**

Gene Name: RPA1

NCBI Gene Entry: 6117

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human RPA1 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.

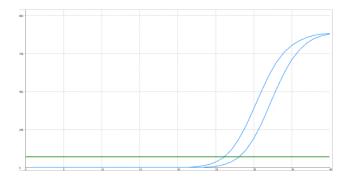
### **Instructions For Use**

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

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

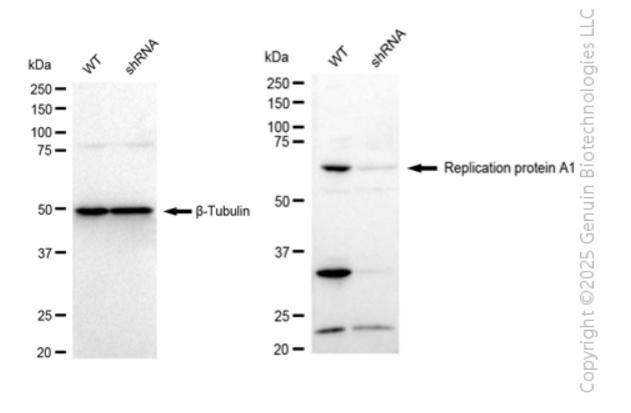
#### Validation Data

# **Human RPA1 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value	_
Wild-Type	25.92	
Knock-Down	27.86	
∆Ct (CtKD-CtWT)	1.94	
% mRNA	yvight	
Reduction	<b>74%</b>	

RT-qPCR analysis. HeLa cells were infected with RPA1-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. RPA1 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 against RPA1 and  $\beta$ -Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit.