# **Human MCM7 Knockdown Cell Line (WB-Validated)**



# **Catalog #: C5157**

#### **Aliases**

MCM7; Minichromosome Maintenance Complex Component 7; P1CDC47; MCM2; P85MCM; CDC47; PNAS146; P1.1-MCM3; DNA Replication Licensing Factor MCM7; CDC47 Homolog; Homolog Of S. Cerevisiae Cdc47; MCM7 Minichromosome Maintenance Deficient 7 (S. Cerevisiae); Minichromosome Maintenance Deficient 7; Minichromosome Maintenance Deficient (S. Cerevisiae) 7; EC 3.6.4.12

# **Background**

Gene Name: MCM7 NCBI Gene Entry: 4176

# **Storage**

Store at liquid nitrogen for 1 year.

# **Kit Components**

- 1. Human MCM7 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.

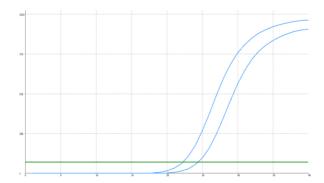
### **Instructions For Use**

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

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

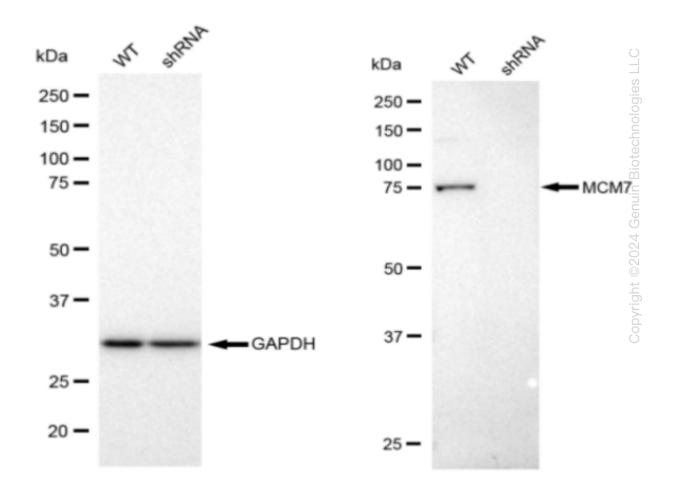
#### Validation Data

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Genotype	Ct Value
Wild-Type	22.20
Knock-Down	24.26
$\Delta Ct (Ct_{KD}-Ct_{WT})$	2.06
% mRNA Reduction	<b>↓ 76%</b>

RT-qPCR analysis. HeLa cells were infected with MCM7-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. MCM7 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. GAPDH served as a loading control. The

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blots were incubated with primary antibodies (Cat#68157, 1:5,000) against MCM7 and GAPDH, 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).