# **Human CCNE2 Knockdown Cell Line (WB-Validated)**



**Catalog #: C61601** 

### **Aliases**

CCNE2; Cyclin E2; CYCE2; G1/S-Specific Cyclin-E2

# **Background**

Gene Name: CCNE2 NCBI Gene Entry: 9134

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human CCNE2 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.

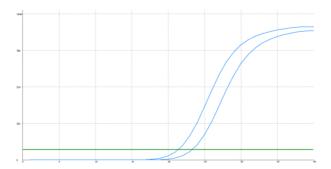
## **Instructions For Use**

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

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

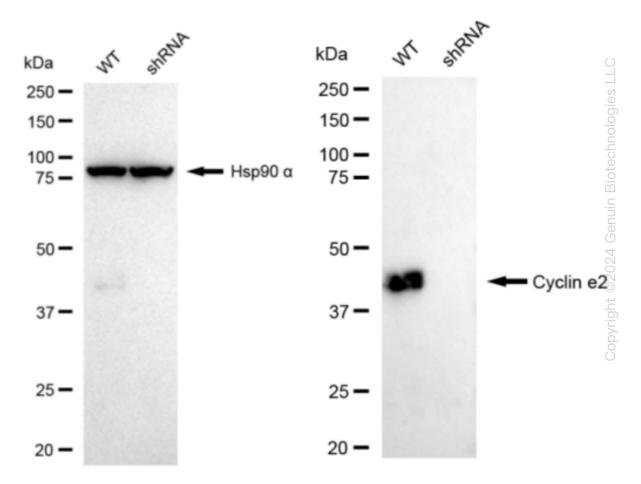
#### **Validation Data**

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Genotype	Ct Value
Wild-Type	21.24
Knock-Down	23.13
$\Delta$ Ct (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	1.89
% mRNA Reduction	<b>♣ 73%</b>

RT-qPCR analysis. HeLa cells were infected with CCNE2-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. CCNE2 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#61601, 1:5,000) against CCNE2 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).