# **Human MRPS26 Knockdown Cell Line (WB-Validated)**



## **Catalog #: C65321**

#### **Aliases**

MRPS26; Mitochondrial Ribosomal Protein S26; MRP-S13; MRP-S26; RPMS13; DJ534B8.3; C20orf193; MS26; 28S Ribosomal Protein S13, Mitochondrial; 28S Ribosomal Protein S26, Mitochondrial; Small Ribosomal Subunit Protein MS26; S13mt; S26mt; Serologically Defined Breast Cancer Antigen NY-BR-87; Mitochondrial Small Ribosomal Subunit Protein MS26; Chromosome 20 Open Reading Frame 193; NY-BR-87; MRPS13; GI008

## **Background**

Gene Name: MRPS26 NCBI Gene Entry: 64949

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human MRPS26 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.

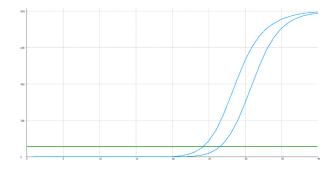
#### **Instructions For Use**

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

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

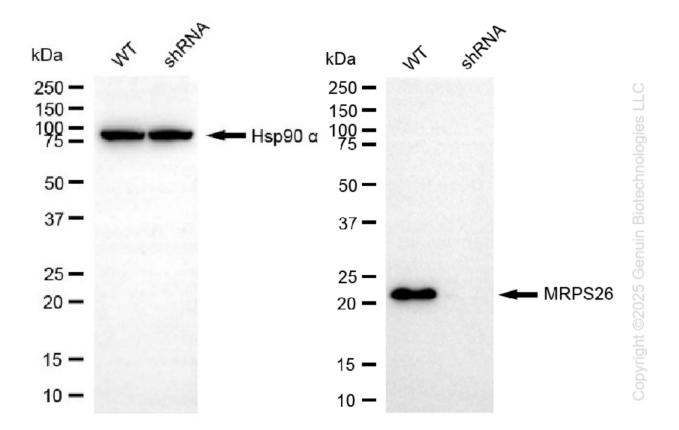
#### Validation Data

## **Human MRPS26 Knockdown Cell Line (WB-Validated)**



| Genotype                                  | Ct Value     |
|---|--------------|
| Wild-Type                                 | 24.21        |
| Knock-Down                                | 26.49        |
| ∆Ct (Ct <sub>KD</sub> -Ct <sub>WT</sub> ) | 2.28         |
| % mRNA<br>Reduction                       | <b>↓</b> 79% |

RT-qPCR analysis. HeLa cells were infected with MRPS26-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.MRPS26 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 MRPS26 and Hsp90  $\alpha$ , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using NaQ<sup>TM</sup> ECL Substrate Kit.