# **Human VPS35 Knockdown Cell Line (WB-Validated)**



**Catalog #: C61728** 

#### **Aliases**

VPS35; VPS35 Retromer Complex Component; MEM3; PARK17; Vacuolar Protein Sorting-Associated Protein 35; Maternal-Embryonic 3; FLJ10752; HVPS35; Vacuolar Protein Sorting 35 Homolog (S. Cerevisiae); Vacuolar Protein Sorting 35 Homolog; VPS35, Retromer Complex Component; Vesicle Protein Sorting 35

## **Background**

Gene Name: VPS35

NCBI Gene Entry: 55737

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human VPS35 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.

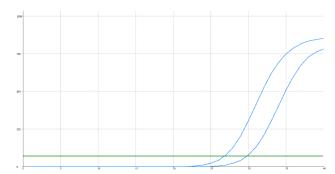
## **Instructions For Use**

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

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

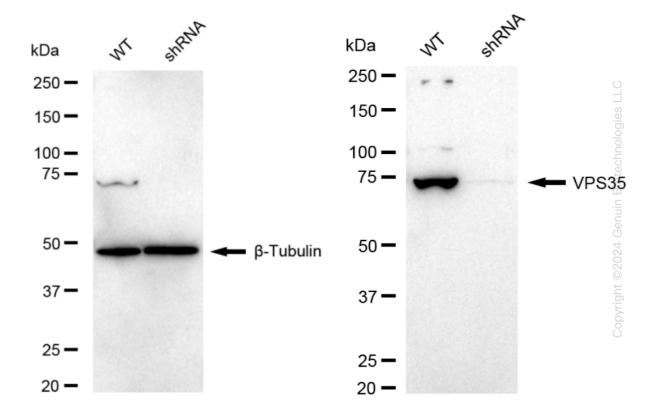
#### Validation Data

# **Human VPS35 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	26.53
Knock-Down	29.38
ΔCt (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	2.85
% mRNA Reduction	<b>↓</b> 86%

RT-qPCR analysis. HeLa cells were infected with VPS35-specific shRNA lentiviral particles, total RNA was extracted from wild-type and knockdown cells, RT-qPCR was performed using genespecific 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. VPS35 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 VPS35 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.