## **Human DGCR8 Knockdown Cell Line (WB-Validated)**



**Catalog #: C62316** 

#### **Aliases**

DGCR8; DGCR8 Microprocessor Complex Subunit; DGCRK6; DiGeorge Syndrome Critical Region Gene 8; Microprocessor Complex Subunit DGCR8; DiGeorge Syndrome Critical Region 8; C22orf12; Pasha; Gy1; DGCR8, Microprocessor Complex Subunit; Chromosome 22 Open Reading Frame 12; C22ORF12; PASHA; GY1

### **Background**

Gene Name: DGCR8 NCBI Gene Entry: 54487

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human DGCR8 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.

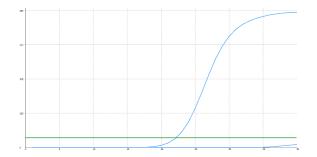
#### **Instructions For Use**

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

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

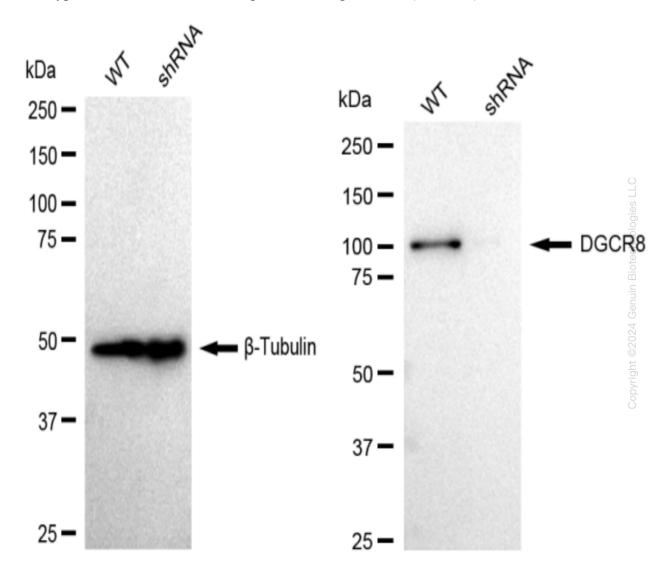
#### Validation Data

## **Human DGCR8 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	22.18
Knock-Down	37.51
$\Delta Ct (Ct_{KD}-Ct_{WT})$	15.33
% mRNA Reduction	<b>4</b> 99.99%

RT-qPCR analysis. HeLa cells were infected with DGCR8-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. DGCR8 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

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blots were incubated with primary antibodies against DGCR8 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.