# **Human STAT6 Knockdown Cell Line (WB-Validated)**



**Catalog #: C69717** 

#### **Aliases**

STAT6; Signal Transducer And Activator Of Transcription 6; IL-4-STAT; D12S1644; Signal Transducer And Activator Of Transcription 6, Interleukin-4 Induced; Transcription Factor IL-4 STAT; STAT, Interleukin4-Induced; EC 2.4.1.227; EC 2.7.7.6; IL-4 Stat; STAT6B; STAT6C

## **Background**

Gene Name: STAT6 NCBI Gene Entry: 6778

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human STAT6 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.

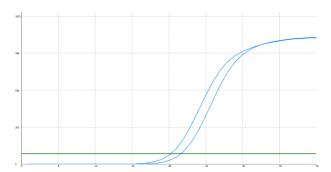
## **Instructions For Use**

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

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

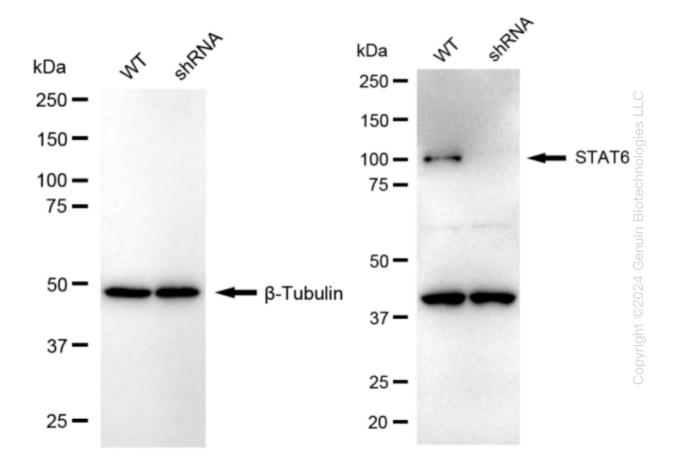
#### Validation Data

## **Human STAT6 Knockdown Cell Line (WB-Validated)**



Genotype	Ct Value
Wild-Type	20.02
Knock-Down	21.35
$\Delta$ Ct (Ct <sub>KD</sub> -Ct <sub>WT</sub> )	1.33
% mRNA Reduction	<b>4</b> 60%

RT-qPCR analysis. HeLa cells were infected with STAT6-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. STAT6 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. β-Tubulin served as a loading control. The blots were incubated with primary antibodies against STAT6 and β-Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit.