

# Human SNAP29 Knockdown Cell Line (WB-Validated)



**Catalog #: C62640**

## Aliases

SNAP29; Synaptosome Associated Protein 29; SNAP-29; Soluble 29 KDa NSF Attachment Protein; CEDNIK; Cerebral Dysgenesis, Neuropathy, Ichthyosis And KeRatoderma Syndrome; Vesicle-Membrane Fusion Protein SNAP-29; Synaptosomal-Associated Protein, 29kDa; Synaptosomal-Associated Protein 29; Synaptosomal-Associated Protein, 29kD; Synaptosome Associated Protein 29kDa

## Background

Gene Name: SNAP29  
NCBI Gene Entry: [9342](#)

## Storage

Store at liquid nitrogen for 1 year.

## Kit Components

1. Human SNAP29 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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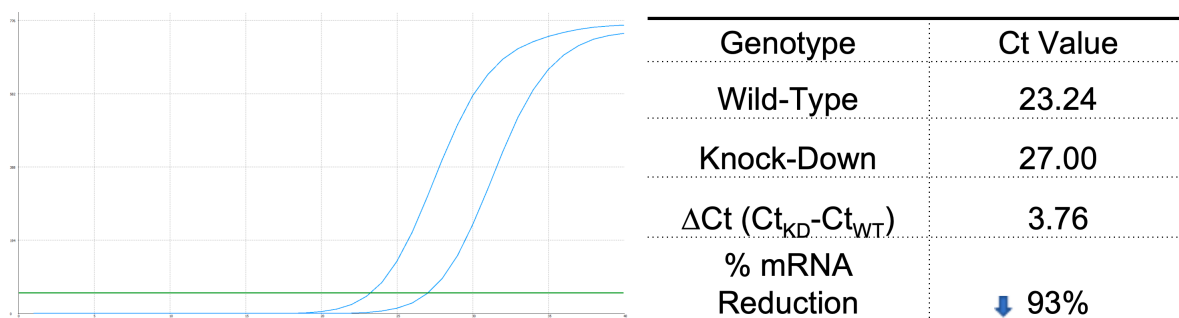
### SUPPORT

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### ORDERS

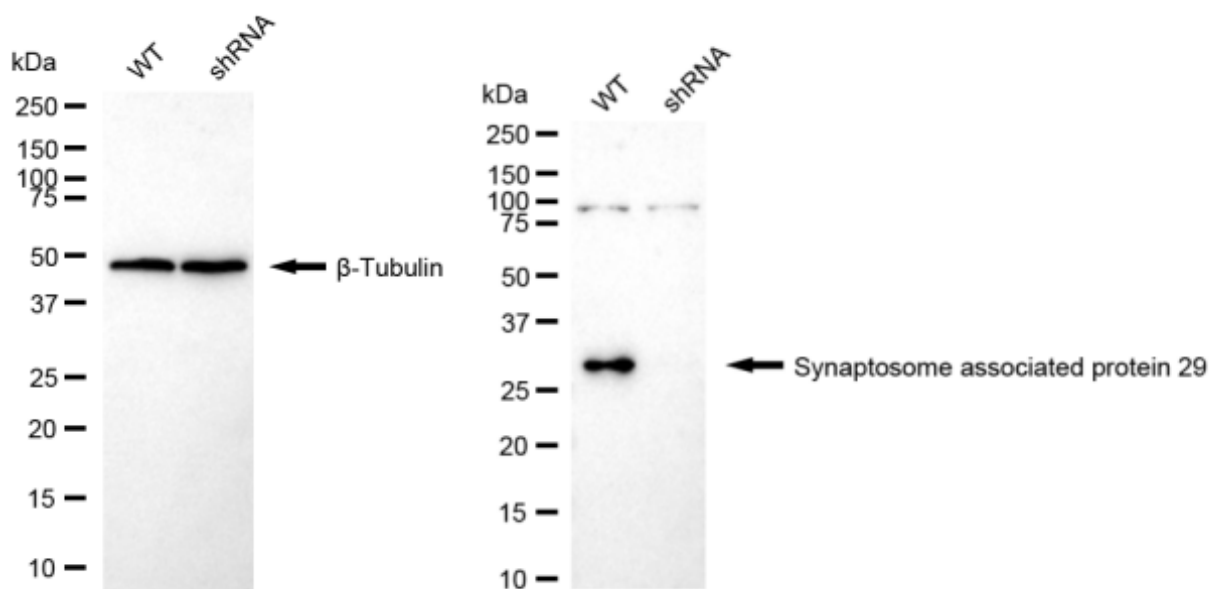
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RT-qPCR analysis. HeLa cells were infected with SNAP29-specific shRNA lentiviral particles, total RNA was extracted from wild-type and knockdown cells, RT-qPCR was performed using gene-specific primers.  $\Delta Ct (Ct_{KD} - Ct_{WT})$  was used to calculate mRNA reduction (%) between wild-type and knockdown cells using the following formula:  $(1 - 1/2^{\Delta Ct}) \times 100\%$ .



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Western blotting analysis. SNAP29 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 SNAP29 and  $\beta$ -Tubulin, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ™ ECL Substrate Kit.