## **Human ANXA10 Knockdown Cell Line (WB-Validated)**



**Catalog #: C61578** 

#### **Aliases**

ANXA10; Annexin A10; ANX14; Annexin-10; Annexin 14; Annexin-14

### **Background**

Gene Name: ANXA10 NCBI Gene Entry: 11199

## **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human ANXA10 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.

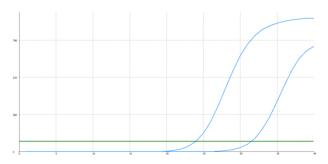
#### **Instructions For Use**

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

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

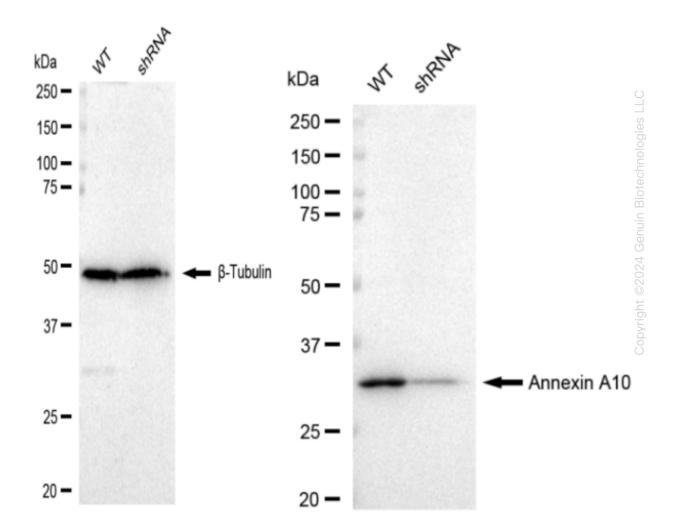
#### Validation Data

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| Genotype                      | Ct Value       |
|-------------------------------|----------------|
| Wild-Type                     | 23.81          |
| Knock-Down                    | 30.98 Genuin B |
| $\Delta Ct (Ct_{KD}-Ct_{WT})$ | 7.17 °         |
| % mRNA Reduction              | <b>J</b> 99%   |

RT-qPCR analysis. HeLa cells were infected with ANXA10-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. ANXA10 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

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blots were incubated with primary antibodies (Cat#61578, 1:5,000) against ANXA10 and  $\beta$ -Tubulin , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody (Cat#201, 1:20,000). Images were developed using FeQ<sup>TM</sup> ECL Substrate Kit (Cat#226).