# **Human CYFIP1 Knockdown Cell Line (WB-Validated)**



## **Catalog #: C1921**

#### **Aliases**

CYFIP1; Cytoplasmic FMR1 Interacting Protein 1; P140SRA-1; KIAA; Cytoplasmic FMRP Interacting Protein 1; Cytoplasmic FMR1-Interacting Protein 1; Specifically Rac1-Associated Protein 1; Selective Hybridizing Clone; P140sra-1; SRA-1; Sra-1; SRA1

### **Background**

Gene Name: CYFIP1 NCBI Gene Entry: 23191

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human CYFIP1 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.

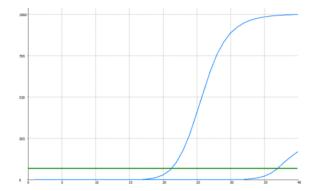
### **Instructions For Use**

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 CYFIP1 Knockdown Cell Line (WB-Validated)**



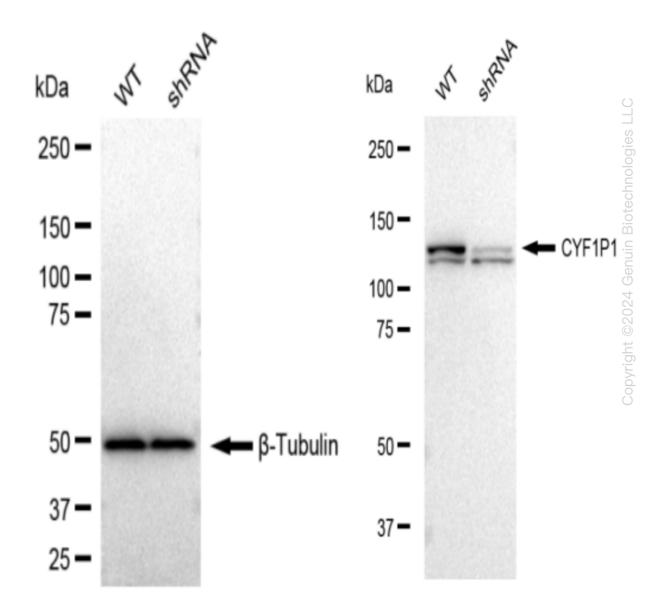
Genotype	Ct Value
Wild-Type	<b>21.26</b> $_{_{\mathrm{2}}}$
Knock-Down	34.00
$\Delta Ct (Ct_{KD}-Ct_{WT})$	12.74
% mRNA Reduction	<b>4</b> 99.99%
70 mm 47 meddetion	<b>33.3370</b> 0 4505

**Figure 1.** RT-qPCR analysis. HeLa cells were infected with *CYFIP1*-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<sub>KD</sub>-Ct<sub>WT</sub>) was used to calculate mRNA reduction (%) between wild-type and knockdown cells using the following formula: (1-1/2<sup> $\Delta$ Ct</sup>) x 100%.

RT-qPCR analysis. HeLa cells were infected with CYFIP1-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%.

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Western blotting analysis. CYFIP1 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 (Cat#61923, 1:5,000) against CYFIP1 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).