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



**Catalog #: C61348** 

#### **Aliases**

AR; Androgen Receptor; NR3C4; Dihydrotestosterone Receptor; HUMARA; SMAX1; DHTR; AIS; Nuclear Receptor Subfamily 3 Group C Member 4; SBMA; Spinal And Bulbar Muscular Atrophy; Testicular Feminization; Kennedy Disease; HYSP1; AR8; TFM; KD

### **Background**

Gene Name: AR

NCBI Gene Entry: 367

### **Storage**

Store at liquid nitrogen for 1 year.

## **Kit Components**

- 1. Human AR 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.

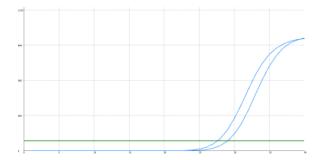
## **Instructions For Use**

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

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

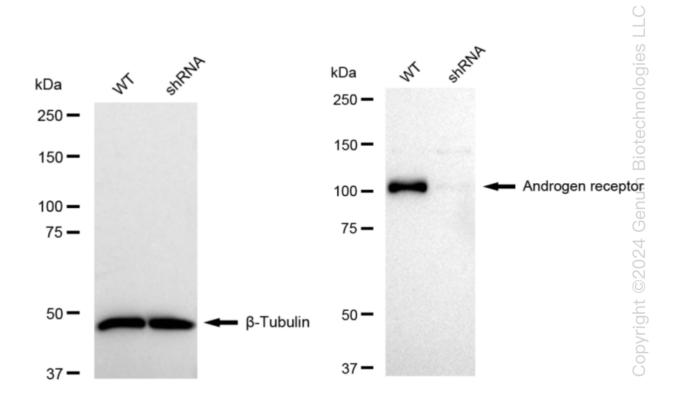
#### Validation Data

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



Genotype	Ct Value
Wild-Type	27.23 gg
Knock-Down	28.66
$\Delta Ct$ ( $Ct_{KD}$ - $Ct_{WT}$ )	1.43
% mRNA Reduction	<b>♣ 63</b> % dipind

RT-qPCR analysis. HeLa cells were infected with AR-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. AR 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 (Cat#61348, 1:5,000) against AR and β-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).