

Human RAD23B Knockdown Cell Line (WB-Validated)



Catalog #: C63571

Aliases

RAD23B; RAD23 Homolog B, Nucleotide Excision Repair Protein; HHR23B; HR23B; P58; XP-C Repair-Complementing Complex 58 KDa Protein; UV Excision Repair Protein RAD23 Homolog B; XP-C Repair Complementing Complex 58 KDa; XP-C Repair Complementing Protein; RAD23 (S. Cerevisiae) Homolog B; RAD23 Homolog B (S. Cerevisiae); RAD23, Yeast Homolog Of, B

Background

Gene Name: RAD23B
NCBI Gene Entry: [5887](#)

Storage

Store at liquid nitrogen for 1 year.

Kit Components

1. Human RAD23B 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

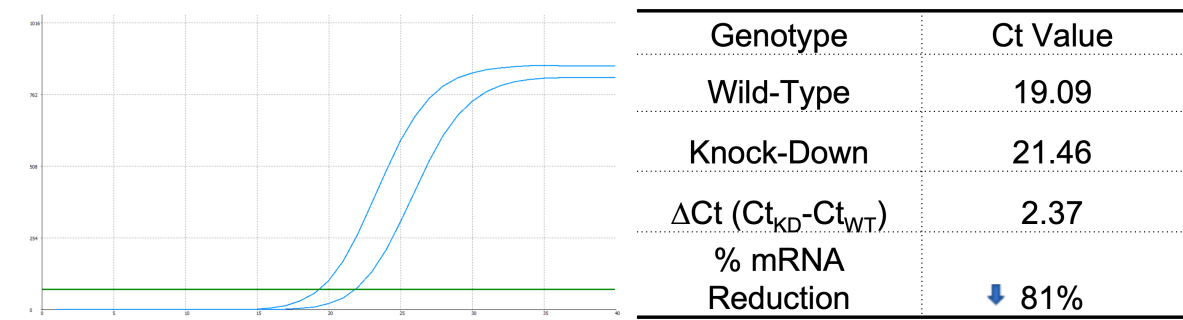
SUPPORT

SUPPORT@GENUINBIOTECH.COM
TEL: +1-540-855-7041

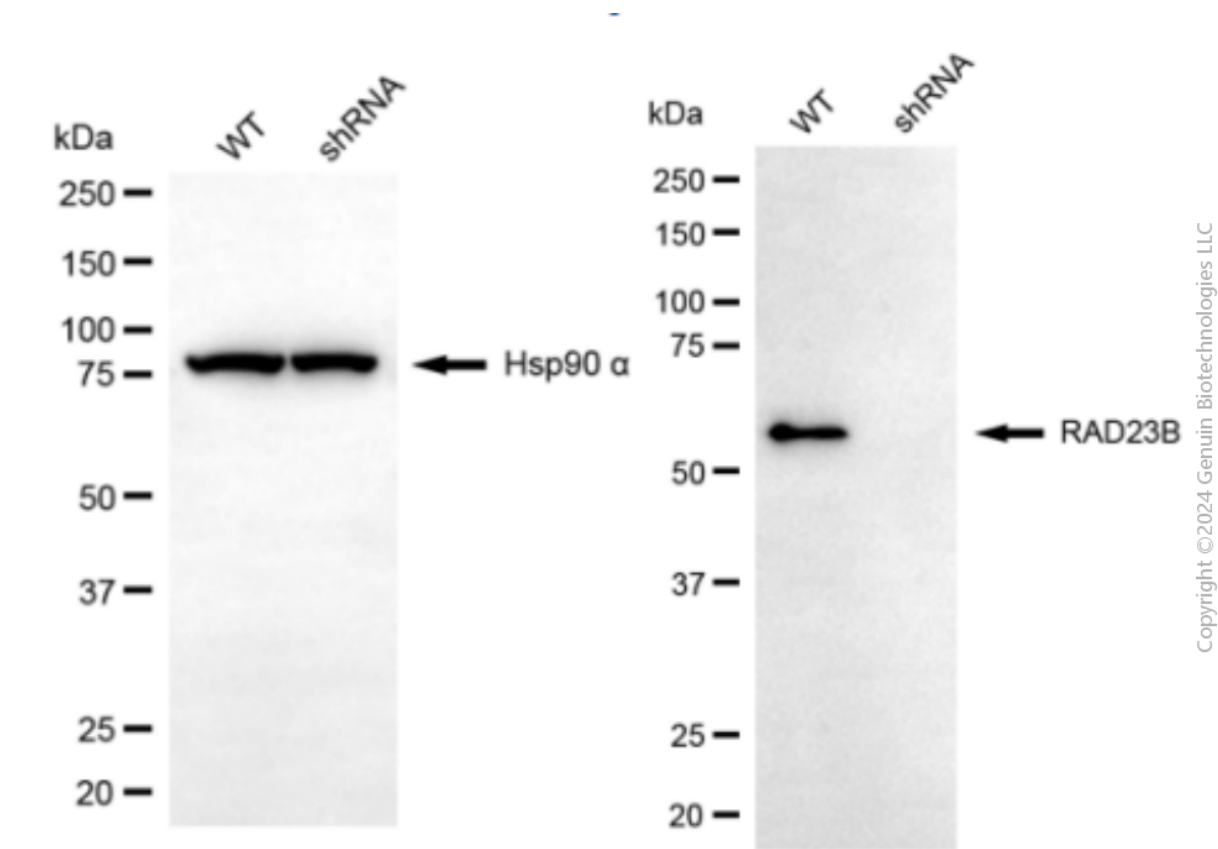
ORDERS

SALES@GENUINBIOTECH.COM
FAX: +1-540-855-7041

WWW.GENUINBIOTECH.COM



RT-qPCR analysis. HT-1080 cells were infected with RAD23B-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: $(1 - 1/2^{\Delta Ct}) \times 100\%$.



Western blotting analysis. RAD23B protein expression in wild-type (WT) and shRNA knockdown (KD) HT-1080 cells was detected using Western blotting. Hsp90 α served as a loading control. The blots were incubated with primary antibodies against RAD23B and Hsp90 α , respectively, followed by incubating with HRP-conjugated goat anti-mouse secondary antibody. Images were developed using FeQ™ ECL Substrate Kit.