

Human RRM1 Knockdown Cell Line (WB-Validated)



Catalog #: C69651

Aliases

Ribonucleotide Reductase Catalytic Subunit M1; Ribonucleoside-Diphosphate Reductase Large Subunit; Ribonucleoside-Diphosphate Reductase Subunit M1; Ribonucleotide Reductase M1 Polypeptide; EC 1.17.4.1; RR1; Ribonucleotide Reductase Large Subunit; Ribonucleotide Reductase, R1 Subunit; RIR1; R1

Background

Gene Name: RRM1

NCBI Gene Entry: [6240](#)

Storage

Store at liquid nitrogen for 1 year.

Kit Components

1. Human RRM1 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

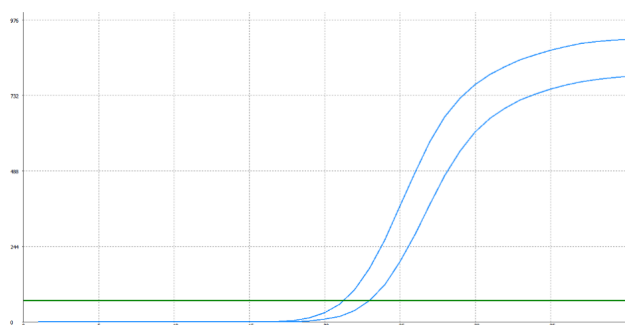
ORDERS

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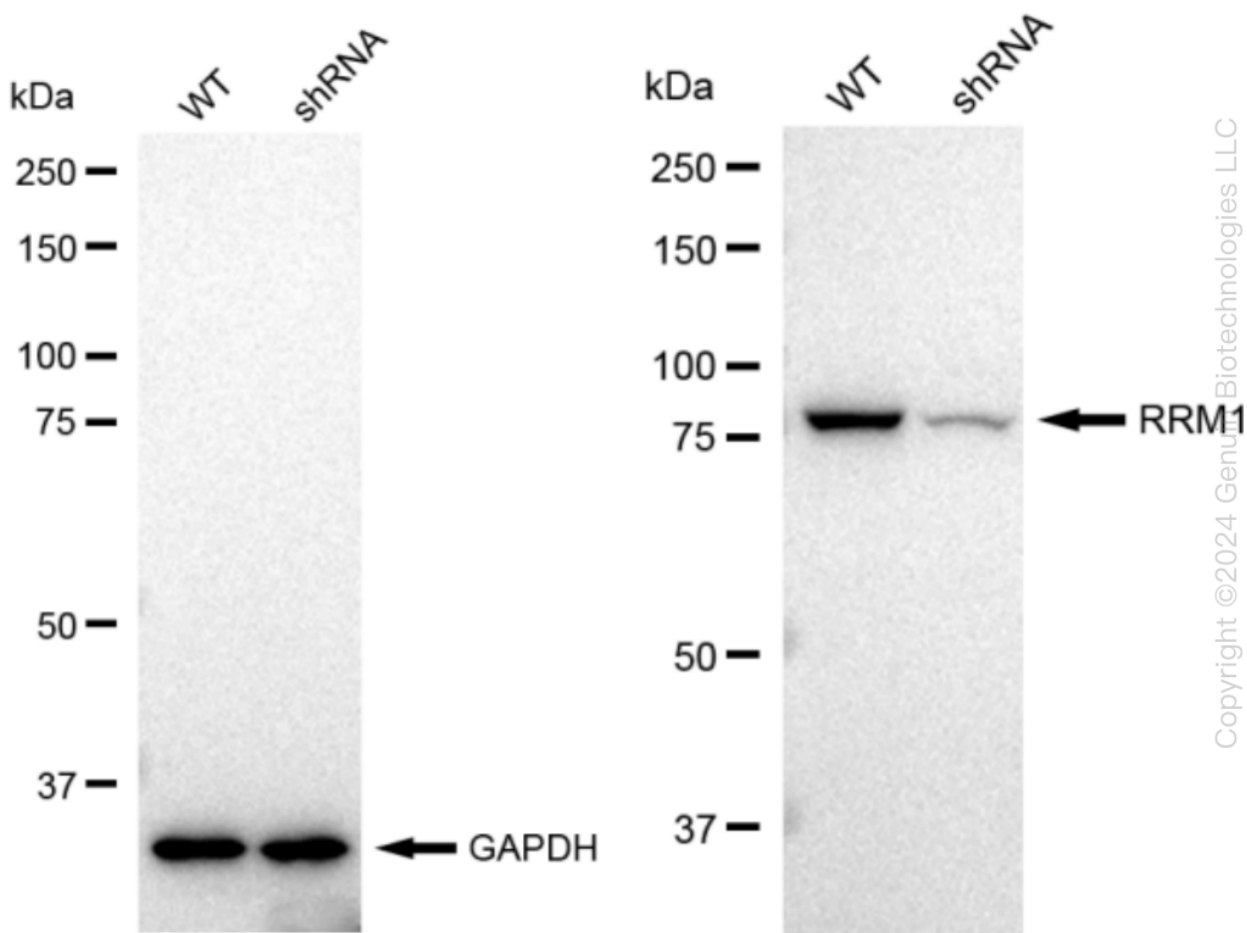
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Genotype	Ct Value
Wild-Type	21.17
Knock-Down	22.59
$\Delta Ct (Ct_{KD} - Ct_{WT})$	1.42
% mRNA Reduction	↓ 63%

RT-qPCR analysis. HeLa cells were infected with RRM1-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. RRM1 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. GAPDH served as a loading control. The blots were incubated with primary antibodies (Cat#69651, 1:5,000) against RRM1 and GAPDH, respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody

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(Cat#201, 1:20,000). Images were developed using FeQ™ ECL Substrate Kit (Cat#226).

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