Human MACROH2A1 Knockdown Cell Line (WB-Validated)



Catalog #: C61119

Aliases

MACROH2A1; MacroH2A.1 Histone; H2AFY; Medulloblastoma Antigen MU-MB-50.205; H2A Histone Family Member Y; Core Histone Macro-H2A.1; Histone MacroH2A1; Histone H2A.Y; MacroH2A1.2; H2A/Y; MH2A1; H2A Histone Family, Member Y; Histone MacroH2A1.1; Histone MacroH2A1.2; MACROH2A1.1; MACROH2A1.2; H2AF12M; H2A.Y

Background

Gene Name: MACROH2A1 NCBI Gene Entry: 9555

Storage

Store at liquid nitrogen for 1 year.

Kit Components

- 1. Human MACROH2A1 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.

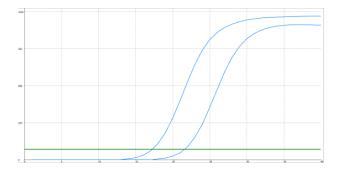
Instructions For Use

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

Note: This product is for research use only.

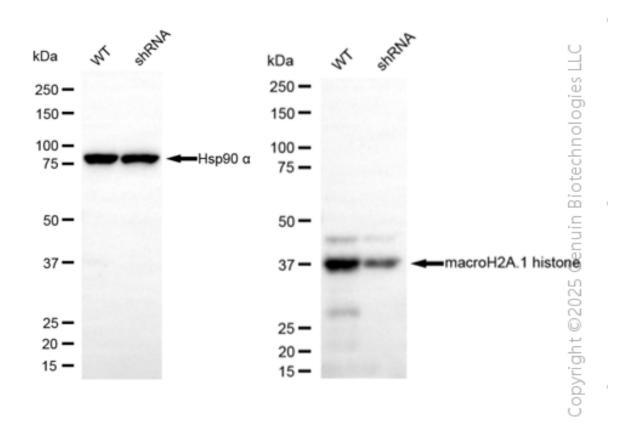
Validation Data

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Genotype	Ct Value	— [
Wild-Type	17.20	,
Knock-Down	21.43	
ΔCt (CtKD-CtWT)	4.23	
% mRNA	in in	2
Reduction	95%)

RT-qPCR analysis. HeLa cells were infected with MACROH2A1-specific shRNA lentiviral particles, total RNA was extracted from wild-type and knockdown cells, RT-qPCR was performed using gene-specific primers. Δ 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. MACROH2A1 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. Hsp90 α served as a loading control. The blots were incubated with primary antibodies against MACROH2A1 and Hsp90 α ,

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respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using NaQ^{TM} ECL Substrate Kit.