

# Human PON2 Knockdown Cell Line (WB-Validated)



**Catalog #: C62478**

## Aliases

Paraoxonase 2; Serum Paraoxonase/Arylesterase 2; Serum Aryldialkylphosphatase 2; Aromatic Esterase 2; Paraoxonase Nirs; Arylesterase 2; A-Esterase 2; PON 2; EC 3.1.1.81; EC 3.1.1.2

## Background

Gene Name: PON2

NCBI Gene Entry: [5445](#)

## Storage

Store at liquid nitrogen for 1 year.

## Kit Components

1. Human PON2 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

---

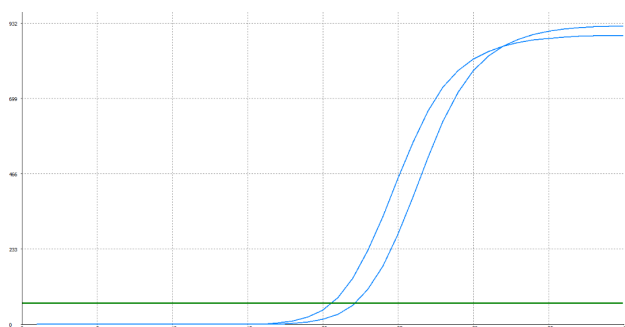
### SUPPORT

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

### ORDERS

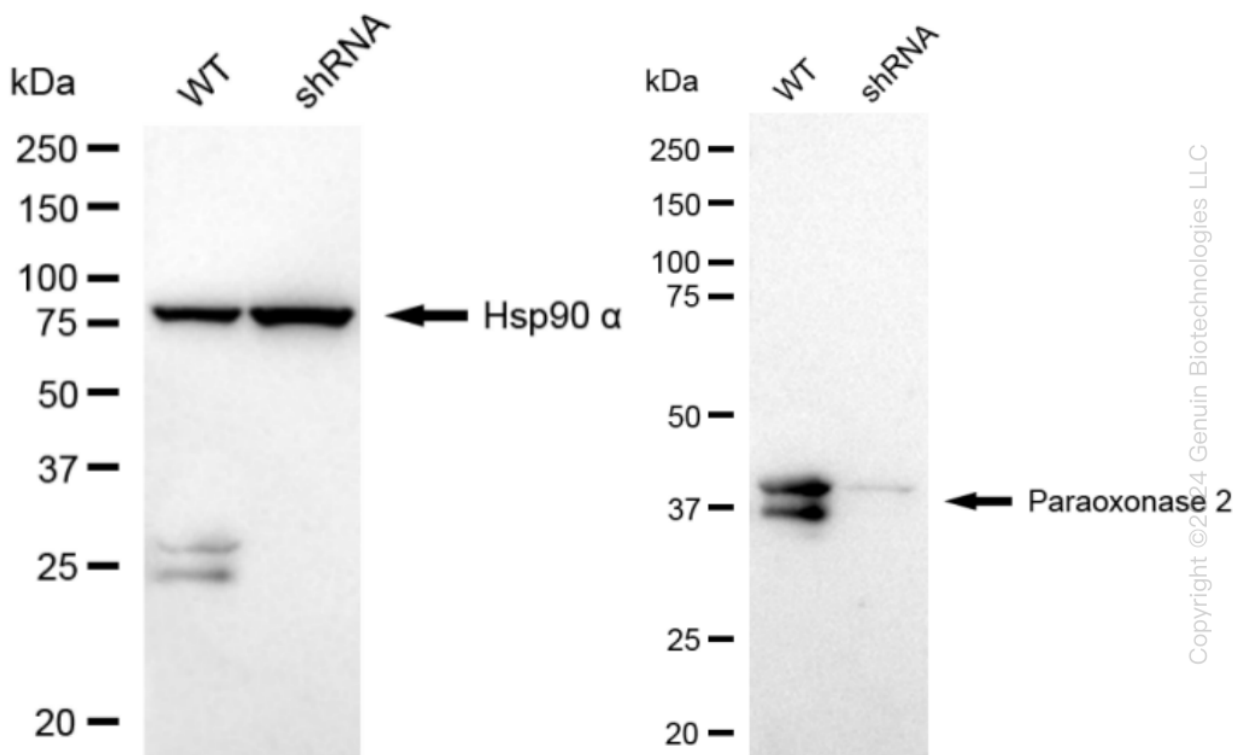
SALES@GENUINBIOTECH.COM  
FAX: +1-540-855-7041

[WWW.GENUINBIOTECH.COM](http://WWW.GENUINBIOTECH.COM)



Genotype	Ct Value
Wild-Type	20.50
Knock-Down	22.13
$\Delta Ct (Ct_{KD} - Ct_{WT})$	1.63
% mRNA Reduction	↓ 68%

RT-qPCR analysis. HeLa cells were infected with PON2-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. PON2 protein expression in wild-type (WT) and shRNA knockdown (KD) HeLa cells was detected using Western blotting. Hsp90  $\alpha$  served as a loading control. The blots were incubated with primary antibodies against PON2 and Hsp90  $\alpha$ , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ™ ECL Substrate Kit.