

Product datasheet for LY300236

JNK1 (MAPK8) Human Knockdown Lysate

Product data:

Product Type: Knockdown Lysates

Description: WB-validated MAPK8 Knockdown HeLa Cell Lysate

Species: Human **Expression Host:** HeLa

Tag: Tag Free

Synonyms: MAPK8; Mitogen-Activated Protein Kinase 8; SAPK1; INK1; PRKM8; INK; Stress-Activated

> Protein Kinase 1c; C-Jun N-Terminal Kinase 1; JUN N-Terminal Kinase; MAP Kinase 8; EC 2.7.11.24; JNK-46; SAPK1c; Stress-Activated Protein Kinase JNK1; Stress-Activated Protein

Kinase 1; JNK21B1/2; EC 2.7.11; JNK1A2; SAPK1C; MAPK 8

Predicted MW: 48 kDa

Components: 1 vial of 100 ug WT HeLa cell lysate

1 vial of 100 ug MAPK8 KD HeLa cell lysate

Storage: Store at -20 °C for two years.

Concentration: Lot-specific

Buffer: IntactProtein Cell-Tissue Lysis buffer

Locus ID: 5599 **UniProt ID:** P45983

Protein Families: Druggable Genome, ES Cell Differentiation/IPS, Protein Kinase

Protein Pathways: Adipocytokine signaling pathway, Colorectal cancer, Epithelial cell signaling in Helicobacter

> pylori infection, ErbB signaling pathway, Fc epsilon RI signaling pathway, Focal adhesion, GnRH signaling pathway, Insulin signaling pathway, MAPK signaling pathway, Neurotrophin signaling pathway, NOD-like receptor signaling pathway, Pancreatic cancer, Pathways in cancer, Progesterone-mediated oocyte maturation, RIG-I-like receptor signaling pathway, Toll-

like receptor signaling pathway, Type II diabetes mellitus, Wnt signaling pathway



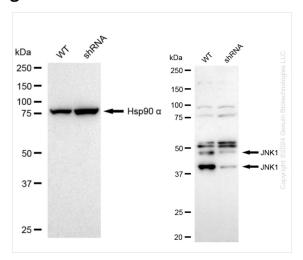
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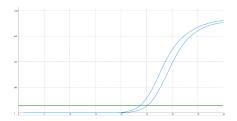
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Product images:



Western blotting analysis. MAPK8 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 MAPK8 and Hsp90 α , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ $^{\rm IM}$ ECL Substrate Kit.



	
Genotype	Ct Value
Wild-Type	23.53
Knock-Down	24.91
Δ Ct (Ct _{KD} -Ct _{WT})	1.38
% mRNA Reduction	4 62%

RT-qPCR analysis. HeLa cells were infected with MAPK8-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 Δ Ct) x 100%.