

ERK 1 (G-8): sc-271269



The Power to Question

BACKGROUND

Mitogen-activated protein kinase (MAPK) signaling pathways involve two closely related MAP kinases, known as extracellular-signal-related kinase 1 (ERK 1, p44) and 2 (ERK 2, p42). Growth factors, steroid hormones, G protein-coupled receptor ligands and neurotransmitters can initiate MAPK signaling pathways. Activation of ERK 1 and ERK 2 requires phosphorylation by upstream kinases such as MAP kinase kinase (MEK), MEK kinase and Raf-1. ERK 1 and ERK 2 phosphorylation can occur at specific tyrosine and threonine sites mapping within consensus motifs that include the threonine-glutamate-tyrosine motif. ERK activation leads to dimerization with other ERKs and subsequent localization to the nucleus. Active ERK dimers phosphorylate serine and threonine residues on nuclear proteins and influence a host of responses that include proliferation, differentiation, transcription regulation and development. The human ERK 1 gene maps to chromosome 16p11.2 and encodes a 379 amino acid protein that shares 83% sequence identity to ERK 2.

REFERENCES

- Boulton, T.G., et al. 1991. ERKs: a family of protein-serine/threonine kinases that are activated and tyrosine phosphorylated in response to Insulin and NGF. *Cell* 65: 663-675.
- Crews, C.M., et al. 1992. The primary structure of MEK, a protein kinase that phosphorylates the ERK gene product. *Science* 258: 478-480.

CHROMOSOMAL LOCATION

Genetic locus: MAPK3 (human) mapping to 16p11.2; Mapk3 (mouse) mapping to 7 F3.

SOURCE

ERK 1 (G-8) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 291-335 within subdomain XI of ERK 1 of rat origin.

PRODUCT

Each vial contains 200 µg IgG_{2b} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

ERK 1 (G-8) is available conjugated to agarose (sc-271269 AC), 500 µg/0.25 ml agarose in 1 ml, for IP; to HRP (sc-271269 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-271269 PE), fluorescein (sc-271269 FITC), Alexa Fluor® 488 (sc-271269 AF488), Alexa Fluor® 546 (sc-271269 AF546), Alexa Fluor® 594 (sc-271269 AF594) or Alexa Fluor® 647 (sc-271269 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-271269 AF680) or Alexa Fluor® 790 (sc-271269 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-271269 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

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STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

ERK 1 (G-8) is recommended for detection of ERK 1 p44 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

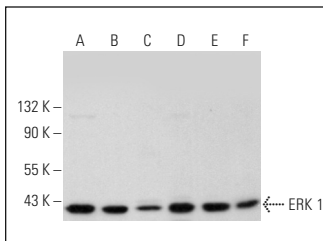
ERK 1 (G-8) is also recommended for detection of ERK 1 p44 in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for ERK 1 siRNA (h): sc-29307, ERK 1 siRNA (m): sc-29308, ERK 1 siRNA (r): sc-156030, ERK 1 shRNA Plasmid (h): sc-29307-SH, ERK 1 shRNA Plasmid (m): sc-29308-SH, ERK 1 shRNA Plasmid (r): sc-156030-SH, ERK 1 shRNA (h) Lentiviral Particles: sc-29307-V, ERK 1 shRNA (m) Lentiviral Particles: sc-29308-V and ERK 1 shRNA (r) Lentiviral Particles: sc-156030-V.

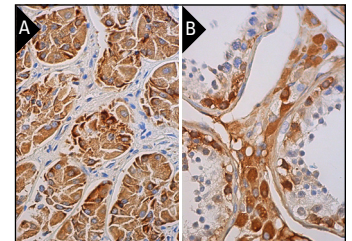
Molecular Weight of ERK 1: 44 kDa.

Positive Controls: MCF7 whole cell lysate: sc-2206, KNRK whole cell lysate: sc-2214 or 3611-RF whole cell lysate: sc-2215.

DATA



ERK 1 (G-8): sc-271269. Western blot analysis of ERK 1 expression in MCF7 (A), THP-1 (B), DU 145 (C), KNRK (D), 3611-RF (E) and HEK293 (F) whole cell lysates.



ERK 1 (G-8): sc-271269. Immunoperoxidase staining of formalin fixed, paraffin-embedded human upper stomach tissue showing cytoplasmic staining of glandular cells (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human testis tissue showing cytoplasmic staining of cells in seminiferous ducts and Leydig cells (B).

SELECT PRODUCT CITATIONS

- Gonzalez-Garcia, J.R., et al. 2014. The dynamics of MAPK inactivation at fertilization in mouse eggs. *J. Cell Sci.* 127: 2749-2760.
- Aurtenetxe, O., et al. 2018. DUSP5 expression associates with poor prognosis in human neuroblastoma. *Exp. Mol. Pathol.* 105: 272-278.
- Najenson, A.C., et al. 2019. The exocrine pancreas is an extracardiac source of atrial natriuretic peptide. *Pflugers Arch.* 471: 915-924.
- Ouyang, H., et al. 2020. p190A RhoGAP induces CDH1 expression and cooperates with E-cadherin to activate LATS kinases and suppress tumor cell growth. *Oncogene* 39: 5570-5587.

RESEARCH USE

For research use only, not for use in diagnostic procedures.