

ERK1 Monoclonal Antibody (ERK-6B11)

Catalog Number13-8600

Product data sheet

Details		Species Reactivity	
Size	100 µg	Species reactivity	Dog, Human, Mouse, Rat
Host/Isotope	Mouse / IgG1, kappa	Published species	Tag, Pig, Rat, Human, Mouse, Not Applicable
Class	Monoclonal		
Type	Antibody	Tested Applications	Dilution *
Clone	ERK-6B11	ELISA (ELISA)	Assay-dependent
Immunogen	Synthetic Peptide corresponding to a section of the C-terminus of rat ERK1.	Immunoprecipitation (IP)	Assay-dependent
Conjugate	Unconjugated	Western Blot (WB)	1:1,000
Form	Liquid	Immunocytochemistry (ICC/IF)	1:250
Concentration	0.5 mg/mL	Published Applications	
Purification	Affinity chromatography	Western Blot (WB)	See 8 publications below
Storage buffer	PBS	Miscellaneous PubMed (Misc)	See 1 publications below
Contains	0.1% sodium azide	Immunoprecipitation (IP)	See 1 publications below
Storage Conditions	-20°C	* Suggested working dilutions are given as a guide only. It is recommended that the user titrate the product for use in their own experiment using appropriate negative and positive controls.	

Background/Target Information

MAPK3 (ERK1) is a serine/threonine kinase that is an important activator of p90, RSK, MSK, ELK1, and Stat3, and is involved in the MAPK pathway. ERK1 is widely expressed and involved in the regulation of meiosis, mitosis, and post-mitotic functions in differentiated cells. Many different stimuli, including growth factors, cytokines, virus infection, ligands for heterotrimeric guanine nucleotide-binding protein (G protein)-coupled receptors and transforming agents, activate the ERK1 and ERK2 pathways. Functionally, ERK1 is involved with a signaling cascade that regulates various cellular processes such as proliferation, differentiation, and cell cycle progression in response to a variety of extracellular signals. ERK1 is activated by upstream kinases, resulting in its translocation to the nucleus where it phosphorylates nuclear targets. Alternatively spliced transcript variants encoding different protein isoforms of ERK1 have been described.

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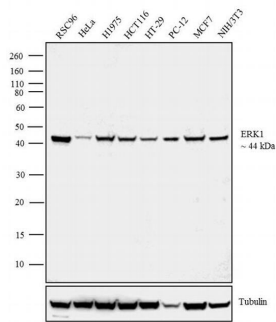
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Product Images For ERK1 Monoclonal Antibody (ERK-6B11)

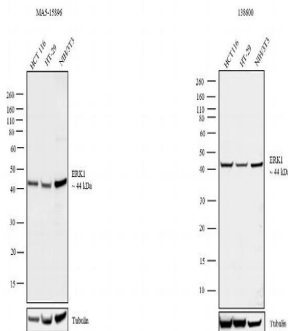
ERK1 Antibody (13-8600) in WB

Western blot analysis was performed on whole cell extracts (30 µg lysate) of RSC96 (Lane 1), HeLa (Lane 2), H1975 (Lane 3), HCT116 (Lane 4), HT-29 (Lane 5), PC12 (Lane 6), MCF7 (Lane 7) and NIH/3T3 (Lane 8). The blot was probed with Anti-ERK1 Mouse Monoclonal Antibody (Product # 13-8600, 1:1000 dilution) and detected by chemiluminescence using Goat anti-Mouse IgG (H+L) Superclonal™ Secondary Antibody, HRP conjugate (Product # A28177, 0.25 µg/mL, 1:4000 dilution). A 44 kDa band corresponding to ERK1 was observed across the cell lines tested. Known quantity of protein samples were electrophoresed using Novex® NuPAGE® 4-12 % Bis-Tris gel (Product # NP0321BOX), XCell SureLock™ Electrophoresis System (Product # EI0002) and Novex® Sharp Pre-Stained Protein Standard (Product # LC5800). Resolved proteins were then transferred onto a nitrocellulose membrane using the wet transfer system. The membrane was probed with the relevant primary and secondary antibody following blocking with 5 % skimmed milk. Chemiluminescent detection was performed using Pierce™ ECL Western Blotting Substrate (Product # 32106).



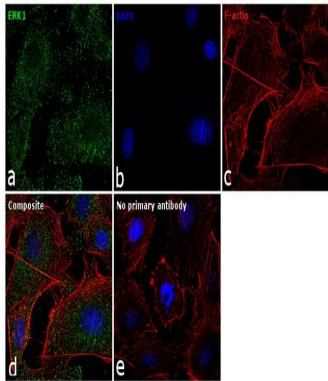
ERK1 Antibody (13-8600)

Antibody specificity was demonstrated by showing that antibodies raised against the same target protein perform similarly. Western blot of ERK1 using Anti-ERK1 Monoclonal Antibody (Product # 13-8600), tested in parallel with ERK1 Monoclonal Antibody (Product # MA5-15896), shows similar expression of ERK1 in the cell lines tested. {IAV}



ERK1 Antibody (13-8600) in ICC/IF

Immunofluorescence analysis of ERK1 was performed using 70% confluent A549 cells. The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 10 minutes, and blocked with 1% BSA for 1 hour at room temperature. The cells were labeled with ERK1 (ERK-6B11) Mouse Monoclonal Antibody (Product # 13-8600) at 1:250 dilution in 0.1% BSA and incubated overnight at 4 degree Celsius and then labeled with Goat anti-Mouse IgG (H+L) Superclonal™ Secondary Antibody, Alexa Fluor® 488 conjugate (Product # A28175) at a dilution of 1:2000 for 45 minutes at room temperature (Panel a: green). Nuclei (Panel b: blue) were stained with SlowFade® Gold Antifade Mountant with DAPI (Product # S36938). F-actin (Panel c: red) was stained with Rhodamine Phalloidin (Product # R415, 1:300). Panel d represents the merged image showing cytoplasmic localization. Panel e represents control cells with no primary antibody to assess background. The images were captured at 60X magnification.



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PubMed References For ERK1 Monoclonal Antibody (ERK-6B11)

8 Western Blot References

Species / Dilution	Summary
	13-8600 was used in Western Blotting to suggest that ethanolic and acetone extracts from Physalis peruviana have potential to overcome oxidative damage induced by neurotoxic compounds.
Human / Not Cited	Frontiers in chemistry (2020; 6:) "Extracts of <i>Physalis peruviana</i> Protect Astrocytic Cells Under Oxidative Stress With Rotenone." Author(s):Areiza-Mazo N,Robles J,Zamudio-Rodriguez JA,Giraldez L,Echeverria V,Barrera-Bailon B,Aliev G,Sahebkar A, Ashraf GM,Barreto GE PubMed Article URL: http://dx.doi.org/10.3389/fchem.2018.00276
Human / Not Cited	Molecular cancer research : MCR (2003; 1: 801) "Distinct mechanisms mediate the initial and sustained phases of cell migration in epidermal growth factor receptor-overexpressing cells." Author(s):Kruger JS,Reddy KB PubMed Article URL: http://www.ncbi.nlm.nih.gov/pubmed/14517342
Pig / Not Cited	The Journal of biological chemistry (2005; 280: 31208) "Roscovitine targets, protein kinases and pyridoxal kinase." Author(s):Bach S,Knockaert M,Reinhardt J,Lozach O,Schmitt S,Baratte B,Koken M,Coburn SP,Tang L,Jiang T,Liang DC, Galons H,Dierick JF,Pinna LA,Meggio F,Totzke F,Schächtele C,Lerman AS,Carnero A,Wan Y,Gray N,Meijer L PubMed Article URL: http://dx.doi.org/10.1074/jbc.M500806200
Rat / Not Cited	13-8600 was used in Western Blot to explore the potential relationships among curcumin, circRNAs, and nasopharyngeal carcinoma (NPC).
Human / Not Cited	Journal of Cancer (2020; 11: 2360) "Curcumin Enhances Radiosensitization of Nasopharyngeal Carcinoma via Mediating Regulation of Tumor Stem-like Cells by a CircRNA Network." Author(s):Zhu D,Shao M,Yang J,Fang M,Liu S,Lou D,Gao R,Liu Y,Li A,Lv Y,Mo Z,Fan Q PubMed Article URL: http://dx.doi.org/10.7150/jca.39511
Not Applicable / 1:3000	13-8600 was used in western blot to study molecular pathways required for fear learning
	The Journal of neuroscience : the official journal of the Society for Neuroscience (2009; 29: 10131) "TrkB modulates fear learning and amygdalar synaptic plasticity by specific docking sites." Author(s):Musumeci G,Sciarretta C,Rodríguez-Moreno A,Al Banchaabouchi M,Negrete-Díaz V,Costanzi M,Berno V, Egorov AV,von Bohlen Und Halbach O,Cestari V,Delgado-García JM,Minichiello L PubMed Article URL: http://dx.doi.org/10.1523/JNEUROSCI.1707-09.2009
Not Applicable / Not Cited	13-8600 was used in western blot to demonstrate that phosphatase of activated cells 1 phosphatase is a direct transcription target of E2F-1 and mediates apoptosis
	Oncogene (2007; 26: 6526) "PAC1 is a direct transcription target of E2F-1 in apoptotic signaling." Author(s):Wu J,Jin YJ,Calaf GM,Huang WL,Yin Y PubMed Article URL: http://dx.doi.org/10.1038/sj.onc.1210484
Tag / Not Cited	The Journal of biological chemistry (2003; 278: 52116) "Regulation of SPIN90 phosphorylation and interaction with Nck by ERK and cell adhesion." Author(s):Lim CS,Kim SH,Jung JG,Kim JK,Song WK PubMed Article URL: http://dx.doi.org/10.1074/jbc.M310974200
Human / Not Cited	13-8600 was used in western blot to test if an increase in fat cell size modulates signaling pathways by changing the relationships between the cell and the extracellular matrix.
Rat / Not Cited	International journal of obesity and related metabolic disorders : journal of the International Association for the Study of Obesity (2003; 27: 1178) "Adipocyte functions are modulated by cell size change: potential involvement of an integrin/ERK signalling pathway." Author(s):Farnier C,Krief S,Blache M,Diot-Dupuy F,Mory G,Ferre P,Bazin R PubMed Article URL: http://dx.doi.org/10.1038/sj.ijo.0802399

1 Miscellaneous PubMed References

Species / Dilution	Summary
	13-8600 was used in western blot to generate and characterize mice conditionally deleted for Ngf or Trka in the central nervous system.
Mouse / 1:3000	The Journal of neuroscience : the official journal of the Society for Neuroscience (2012; 32: 14885) "Loss of NGF-TrkA signaling from the CNS is not sufficient to induce cognitive impairments in young adult or intermediate-aged mice." Author(s):Müller M,Triaca V,Besusso D,Costanzi M,Horn JM,Koudelka J,Geibel M,Cestari V,Minichiello L PubMed Article URL: http://dx.doi.org/10.1523/JNEUROSCI.2849-12.2012

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1 Immunoprecipitation References

Species / Dilution	Summary
Human / Not Cited	The Journal of biological chemistry (2003; 278: 31547) "Alpha 7 nicotinic acetylcholine receptors mediate beta-amyloid peptide-induced tau protein phosphorylation." Author(s):Wang HY,Li W,Benedetti NJ,Lee DH PubMed Article URL: http://dx.doi.org/10.1074/jbc.M212532200

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