MAPK1 Recombinant Antibody

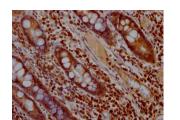
RAC00445

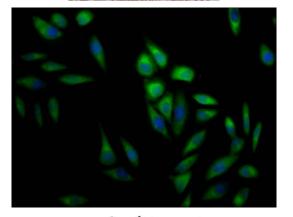


Product Information	
Size:	Protein Background:
50ul	Serine/threonine kinase which acts as an essential component of the MAP kinase signal
Reactivity:	transduction pathway. MAPK1/ERK2 and MAPK3/ERK1 are the 2 MAPKs which play an important role in the MAPK/ERK cascade. They participate also in a signaling cascade initiated by activated KIT and KITLG/SCF. Depending on the cellular context, the MAPK/ERK cascade mediates diverse biological functions such as cell growth, adhesion, survival and differentiation through the regulation of transcription, translation,
Human, Mouse, Rat	
Source:	
Homo sapiens (Human)	cytoskeletal rearrangements. The MAPK/ERK cascade plays also a role in initiation and regulation of meiosis, mitosis, and postmitotic functions in differentiated cells by
lsotype:	phosphorylating a number of transcription factors.
Rabbit IgG	Gene ID:
Applications:	MAPK1
ELISA, WB, IHC, IF	Uniprot
Recommended dilutions:	P28482
WB:1:500-1:5000, IHC:1:50-1:200, IF:1:20- 1:200	Synonyms:
	Mitogen-activated protein kinase 1 (MAP kinase 1) (MAPK 1) (EC 2.7.11.24) (ERT1) (Extracellular signal-regulated kinase 2) (ERK-2) (MAP kinase isoform p42) (p42-MAPK) (Mitogen-activated protein kinase 2) (MAP kinase 2) (MAPK 2), MAPK1, ERK2 PRKM1 PRKM2
	Immunogen:
	A synthesized peptide derived from human ERK2.

Storage:

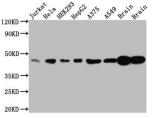
Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.





IHC image of RACO0445 diluted at 1:100 and staining in paraffinembedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

Immunofluorescence staining of Hela Cells with RACO0445 at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).



Western Blot

Positive WB detected in(Jurkat whole cell lysate) Hela whole cell lysate) HEK293 whole cell lysate) HepG2 whole cell lysate) A375 whole cell lysate) A549 whole cell lysate) Rat Brain whole cell lysate) Mouse Brain whole cell lysate) All lanes: ERK2 antibody at 1:1000 Secondary

Goat polyclonal to rabbit IgG at 1:50000 dilution Predicted band size: 42, 37 kDa Observed band size: 42 kDa