

MEK1 (D2R10) Rabbit mAb

✓ 100 µl
 (10 western blots)

New 09/13



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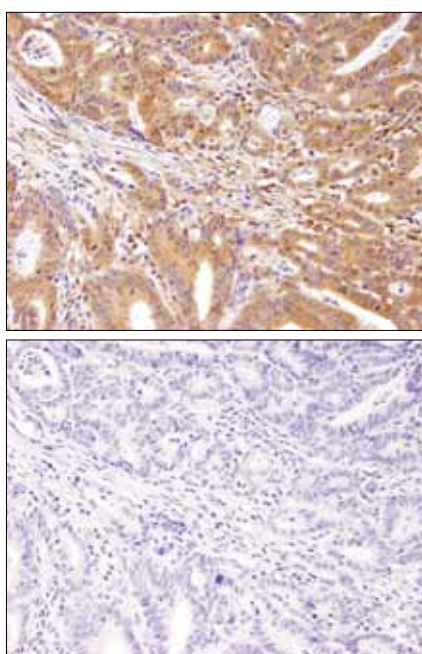
Entrez Gene ID #5604
 UniProt ID #Q02750

Applications W, IHC-P Endogenous	Species Cross-Reactivity* H, M, R, Mk	Molecular Wt. 45 kDa	Isotype Rabbit IgG**
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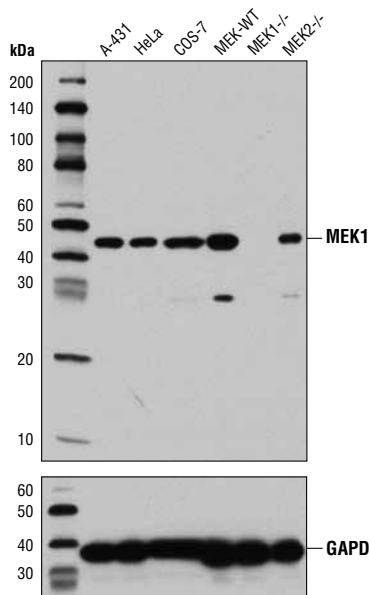
Background: MEK1 and MEK2, also called MAPK or Erk kinases, are dual-specificity protein kinases that function in a mitogen activated protein kinase cascade controlling cell growth and differentiation (1-3). Activation of MEK1 and MEK2 occurs through phosphorylation of two serine residues at positions 217 and 221, located in the activation loop of subdomain VIII, by Raf-like molecules. MEK1/2 is activated by a wide variety of growth factors and cytokines and also by membrane depolarization and calcium influx (1-4). Constitutively active forms of MEK1/2 are sufficient for the transformation of NIH/3T3 cells or the differentiation of PC-12 cells (4). MEK activates p44 and p42 MAP kinase by phosphorylating both threonine and tyrosine residues at sites located within the activation loop of kinase subdomain VIII.

Specificity/Sensitivity: MEK1 (D2R10) Rabbit mAb recognizes endogenous levels of total MEK1 protein. This antibody does not cross-react with MEK2 or other MAPK kinase proteins.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human MEK1 protein.



Immunohistochemical analysis of paraffin-embedded colon adenocarcinoma using MEK1 (D2R10) Rabbit mAb in the presence of control peptide (upper) or antigen specific peptide (lower).



◀ Western blot analysis of extracts from various cell lines using MEK1 (D2R10) Rabbit mAb (upper) or GAPDH (D16H11) XP® Rabbit mAb #5174 (lower). MEK1/2 WT (MEK-WT), MEK1 knockout (MEK1-/-) and MEK2 knockout (MEK2-/-) MEFs were generously provided by Dr. Jean Charron, Centre de recherche du Centre hospitalier de l'Université Laval, Quebec, Canada.

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

***Species cross-reactivity is determined by western blot.**

****Anti-rabbit secondary antibodies must be used to detect this antibody.**

Recommended Antibody Dilutions:

Western blotting	1:1000
Immunohistochemistry (Paraffin)	1:250†
Unmasking buffer:	Citrate
Antibody diluent:	SignalStain® Antibody Diluent #8112
Detection reagent:	SignalStain® Boost (HRP, Rabbit) #8114

† Optimal IHC dilutions determined using SignalStain® Boost IHC Detection Reagent.

For product specific protocols please see the web page for this product at www.cellsignaling.com.

Please visit www.cellsignaling.com for a complete listing of recommended complementary products.

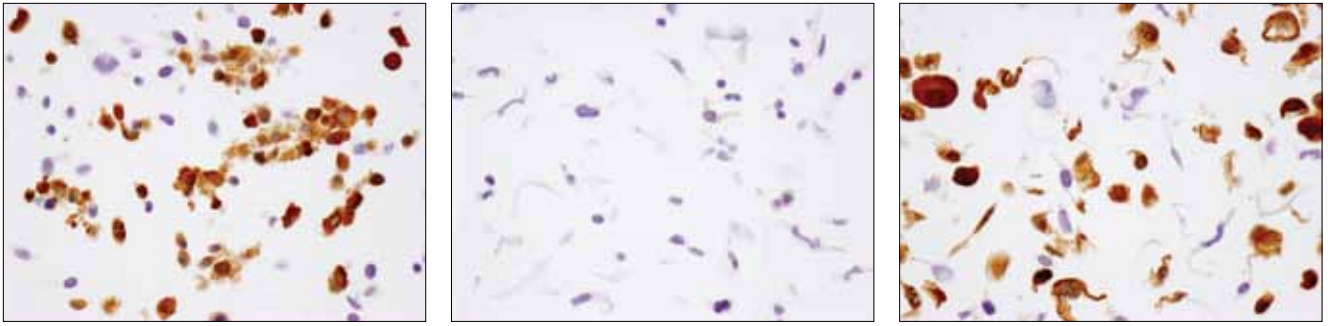
Background References:

- (1) Crews, C.M. et al. (1992) *Science* 258, 478-480.
- (2) Alessi, D.R. et al. (1994) *EMBO J.* 13, 1610-1619.
- (3) Rosen, L.B. et al. (1994) *Neuron* 12, 1207-1221.
- (4) Cowley, S. et al. (1994) *Cell* 77, 841-852.

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

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Applications Key: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide
Species Cross-Reactivity Key: H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine
 Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—horse All—all species expected Species enclosed in parentheses are predicted to react based on 100% homology.



Immunohistochemical analysis of paraffin-embedded cell pellets, wild type MEF (left), MEK1 (-/-) MEF (middle) or MEK2 (-/-) MEF (right), using MEK1 (D2R10) Rabbit mAb. Cells provided by Dr. Jean Charron, Centre de recherche du Centre hospitalier de l'Université Laval, Quebec, Canada.