Revision 1	
Phospho-MEK1/2 (Ser217/221) (166F8) Rabbit mAb	Cell Signaling TECHNOLOGY®
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MW (kDa): 45	Source/Isotype Rabbit IgG	: UniProt ID: #P36507, #Q02750	Entrez-Gene Id: 5605, 5604
Storage			HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than at –20°C. Do not aliquot the antibody.
Specificity/Sens	itivity	Phospho-MEK1/2 (Ser217/2 phosphorylated at serine 2	221) (166F8) Rabbit mAb detects endogenous levels of MEK1/2 only when 17/221.
Source / Purifica	ation		duced by immunizing animals with a synthetic phosphopeptide surrounding Ser217/221 of human MEK1/2.
Background		mitogen activated protein I MEK1 and MEK2 occurs thr located in the activation loc variety of growth factors ar Constitutively active forms differentiation of PC-12 cell	d MAPK or Erk kinases, are dual-specificity protein kinases that function in kinase cascade controlling cell growth and differentiation (1-3). Activation c ough phosphorylation of two serine residues at positions 217 and 221, p of subdomain VIII, by Raf-like molecules. MEK1/2 is activated by a wide nd cytokines and also by membrane depolarization and calcium influx (1-4) of MEK1/2 are sufficient for the transformation of NIH/3T3 cells or the s (4). MEK activates p44 and p42 MAP kinase by phosphorylating both dues at sites located within the activation loop of kinase subdomain VIII.
Background Rei	ferences	1. Crews, C.M. et al. (1992) . 2. Alessi, D.R. et al. (1994) <i>E</i> 3. Rosen, L.B. et al. (1994) <i>A</i> 4. Cowley, S. et al. (1994) <i>Ce</i>	MBO J. 13, 1610-19. Jeuron 12, 1207-21.
Species Reactiv	ity	Species reactivity is determ	ined by testing in at least one approved application (e.g., western blot).
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