

# Mouse Monoclonal Antibody to

# MAPK2/erk2 (internal sequence)

## clone 12A4

0239-100/MAPK2-12A4 Order No.:

100 Size (µg) 0239S Lot No.:



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Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope	Immunogen
lgG1	human	WB	42 kDa	A431	aa 200 - 250	peptide conjugated to hemocyanin

#### **Background and Specificity:**

Extracellular signal/mitogen activated protein kinases (erk/MAPK) are a group of proline-directed serine/threonine kinases that are activated by dual phosphorylation of conserved threonine and tyrosine residues within a characteristic T X Y peptide motif. The mitogen-activated kinases erk1 (MAPK1) and erk2 (MAPK2) acquire full enzymatic activity upon phosphorylation of both threonine and tyrosine residues within the sequence motif T E Y.

Mab MAPK2-12A4 recognizes MAP kinase 2 (erk2) at 42 kDa in Western-blot applications.

The antibody was purified from serum-free cell culture **Purification:** 

supernatant by subsequent ultrafiltration and size exclusion

chromatography.

lyophilized from 1 ml PBS / 0.09 % Na-azide / PEG and Formulation:

Sucrose.

Reconstitute with 1 ml H<sub>2</sub>O (15 min, RT). Reconstitution:

For long-term storage, freeze lyophilizate upon arrival (-20°C). Stability:

Upon reconstitution, aliquote and freeze in liquid nitrogen; reconstituted antibody can be stored frozen at -80°C up to 1 year. Thaw aliquots at 37°C. Thawed aliquots may be stored at 4°C up to

3 months.

Avoid repeated freeze / thaw cycles.

**Positive Control:** #0831: Cell lysate from untreated A431 cells

Immunoblotting: 0.5 µg/ml for HRPO/ECL detection

> Recommended blocking buffer: Casein/Tween 20 based blocking and blot incubation buffer, e.g. nanoTools product

#3031-500/CPPT or #3031-3000/CPPT.

Immunoprecipitation: ND ND Immunocytochemistry: ND **ELISA:** 

> All products are supplied for research and investigational use only. Not for use in humans or laboratory animals.

## **Related Products**

mab to MEK1 (pS218/222) mab to MEK2 (pS222/226) #0174-100/MEK1/2-7E10 mab to MEK1 (N-terminus) #0186-100/MEK1.10B1

mab to MEK1/2 #0150-100/MEK1/2-9G3

mab to MEK2 (N-terminus) #0148-100/MEK2-8È8

mab to MKK3 (N-terminus) #0166-100/MKK3-5

mab to MKK5 (N-terminus) #0224-100/MKK5-14B5

mab to MKK7 (N-terminus) #0189-100/MKK7-10

mab to MAPK 1/2 (pT-E-pY)

#0012-100/MAPK-12D4

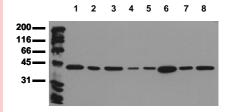
mab to MAPK 2 (C-terminus)

mab to MAPK 2 (N-terminus)

mab to MAPK7/erk5 #0223-100/MAPk7/erk5-12F2 mab to Fos (pS374) #0118-100/Fos-3 mab to Fos (N-terminus)

#0122-100/Fos-8B mab to C-Raf (pS621) #0102-100/C-Raf-6B mab to C-Raf

#0120-100/C-Raf-PBB-1



## **Detection of endogenous MAPK2**

Whole cell lysates of serum starved tumor cells (20.000 cells per lane) were applied to SDS-PAGE and transferred to a PVDF membrane. The immunoblot wasprobed with mab MAPK2-12A4 (0.5 µg/ ml) for 1h at RT and developed by ECL (exp. time: 30 sec).

lane 1: A431; lane 2: A549; lane 3: SKOV3; lane 4: OVCAR5; lane 5: HaCaT; lane 6: PC3; lane 7: HeLa; lane 8: HepG2