

Product Data Sheet

Anti-MKK2 Antibody

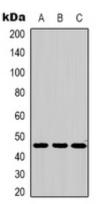
| Catalog # | Source | Reactivity | Applications |
|----------------|----------------|-------------------------------|--|
| CPA9421 | Rabbit | H, M, R | WB, IH |
| Description | Ra | abbit polyclonal antibody t | o MKK2 |
| Immunogen | Re | ecombinant protein corres | ponding to human MKK2. |
| Purification | Tł | he antibody was purified b | y immunogen affinity chromatography. |
| Specificity | Re | ecognizes endogenous leve | els of MKK2 protein. |
| Clonality | Ро | olyclonal | |
| Conjugation | | | |
| Form | Li | quid in 0.42% Potassium p | hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, |
| | ar | nd 0.01% sodium azide. | |
| Dilution | W | /B (1/1000 - 1/2000), IH (1/2 | 00 - 1/500) |
| Gene Symbol | Μ | 1AP2K2 | |
| Alternative Na | ames M | 1EK2; MKK2; PRKMK2; Dua | l specificity mitogen-activated protein kinase kinase 2; |
| | Μ | 1AP kinase kinase 2; MAPK | X 2; ERK activator kinase 2; MAPK/ERK kinase 2; MEK 2 |
| Entrez Gene | 56 | 605 (Human) | |
| SwissProt | P | 36507 (Human) | |
| Storage/Stabi | lity Sł | hipped at 4°C. Upon delive | ry aliquot and store at -20°C for one year. Avoid |
| | fr | eeze/thaw cycles. | |

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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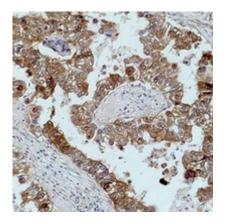




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Western blot analysis of MKK2 expression in Hela (A), NIH3T3 (B), PC12 (C) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 45 kD)



Immunohistochemical analysis of MKK2 staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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