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Research Use Only. Not for diagnostic or therapeutic use.

EB09775 - Goat Anti-SRGAP2 Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: SRGAP2, FNBP2, formin binding protein 2, KIAA0456, SLIT-ROBO

Rho GTPase activating protein 2, srGAP3

Official Symbol: SRGAP2

Accession Number(s): NP_056141.2

Human GeneID(s): 23380

Non-Human GenelD(s): 14270 (mouse), 360840 (rat)

Important Comments: This antibody is expected to recognize isoform a (NP_056141.2).

Immunogen

Peptide with sequence C-KQEDRQTPRSPDST, from the internal region of the protein sequence according to NP_056141.2.

Please note the peptide is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:8000.

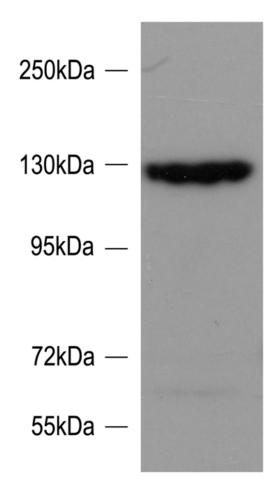
Western blot: Approx 124kDa band observed in lysates of cell line HeLa and in rat primary culture cortical neurons at Day 4 (calculated MW of 121kDa according to human NP_056141.2 but the rat protein sequence is stlll provisional). Recommended concentration: 0.1-0.3µg/ml.

Immunofluorescence: Transfected HEK293 transiently expressing GFP-tagged Human srGAP2 with the primary antibody stained in the red channel (Confocal IF).

Species Reactivity

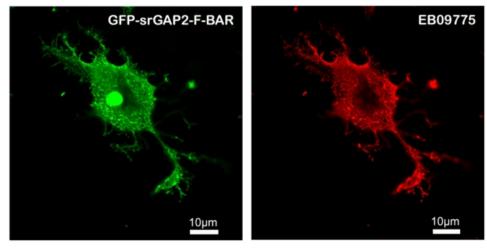
Tested: Human

Expected from sequence similarity: Human, Mouse, Rat, Dog, Cow



IB: EB09775

EB09775 (0.1µg/ml) staining of Rat Cortical Neuron Iysate (35µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence. Data kindly provided by Ms. Ya-Jing Mi and Dr.Wei-Lin Jin, Institute of Neurosciences, Shanghai Jiao Tong University.



HEK293 overexpressing Human srGAP2 and probed with EB09775 at 2.5ug/ml in the right panel. Data kindly provided by Ms. Ya-Jing Mi and Dr.Wei-Lin Jin, Institute of Neurosciences, Shanghai Jiao Tong University.