SHANK2 Antibody

(10 western blots)

New 09/12

For Research Use Only. Not For Use In Diagnostic Procedures.

Applications Species Cross-Reactivity* W, IP M, R, (H, Mk, B) Endogenous	Molecular Wt. 165 kDa	Source Rabbit**
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Background: The SHANK family proteins, also known as proline-rich synapse-associated proteins, consist of SHANK1, SHANK2, and SHANK3. SHANK proteins act as scaffolds at the neuronal post-synaptic density (PSD) (1), where they play a critical role in PSD assembly of excitatory synapses during development (2). While recruitment of SHANK proteins to the synapse is independent of their interaction with Homer (3), proper synaptic targeting of SHANK1 is mediated by interactions between its PDZ domain and PSD proteins (4). At the synapse, SHANK proteins interact with NMDA receptors and metabotropic glutamate receptor complexes (5). Research studies have proposed the involvement of SHANK proteins in autism and neurodegenerative diseases (6)

Specificity/Sensitivity: SHANK2 Antibody recognizes endogenous levels of total SHANK2 protein.

Source/Purification: Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val537 of human SHANK2 protein. Antibodies are purified by protein A and peptide affinity chromatography.





Western blot analysis of extracts from the indicated rat brain synaptic fractions using SHANK2 Antibody. Separation of the different synaptic fractions was confirmed using PSD95 (D27E11) XP[®] Rabbit mAb #3450 and Syntaxin 6 (C34B2) Rabbit mAb #2869. Equal loading of each fraction was assessed using β -Tubulin (9F3) Rabbit mAb #2128. Fractionation of the different synaptic compartments was carried out as described by Phillips, G.R. et al. (2001) Neuron 32, 63-77. PAZ, Pre-synaptic active zone.



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Entrez-Gene ID #22941 Swiss-Prot Acc. #Q9UPX8

Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. 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
Immunoprecipitation		1:50

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

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

Background References:

(1) Grabrucker, A.M. et al. (2011) Trends Cell Biol 21, 594-603.

(2) Boeckers, T.M. et al. (1999) J Neurosci 19, 6506-18.

(3) Boeckers, T.M. et al. (2005) J Neurochem 92, 519-24.

(4) Sala, C. et al. (2001) Neuron 31, 115-30.

(5) Boeckers, T.M. et al. (2002) J Neurochem 81, 903-10.

(6) Grabrucker, A.M. et al. (2011) Trends Cell Biol 21, 594-603.

Western blot analysis of extracts from mouse and rat brain tissue using SHANK2 Antibody.

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.

IF-Immunofluorescence Applications Key: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation 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. melanooaster 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.