

## Phospho-NMDAR2B (Ser1284) Ab

Cat.#: AF2371 Concn.: 1mg/ml Mol.Wt.: 190kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000, IHC 1:50-1:200

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via seguential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-NMDAR2B (Ser1284) Ab detects endogenous levels

of NMDAR2B.

Immunogen: A synthesized peptide derived from human NMDAR2B

around the phosphorylation site of Ser1284.

Uniprot: Q13224

Subcellular Location: Cell membrane. Cell junction > synapse > postsynaptic cell

membrane.

Tissue Specificity: Primarily found in the fronto-parieto-temporal cortex and

hippocampus pyramidal cells, lower expression in the basal

ganglia.

Similarity: A hydrophobic region that gives rise to the prediction of a

transmembrane span does not cross the membrane, but is part of a discontinuously helical region that dips into the membrane and is probably part of the pore and of the selectivity filter.Belongs to the glutamate-gated ion channel

(TC 1.A.10.1) family. NR2B/GRIN2B subfamily. [View

classification]

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Store at -20

°C.Stable for 12 months from date of receipt.



Western blot analysis of extracts from rat brain tissue, using Phospho-NMDAR2B (Ser1284) Ab. Lane1 was treated with phospho-blocking peptide, Lane2 was treated with non-phospho-blocking peptide.



## Affinity Biosciences website: www.affbiotech.com order: order@affbiotech.com



Western blot analysis of extracts from rat brain tissue, using Phospho-NMDAR2B (Ser1284) Ab.



AF2371 at 1/100 staining Human thyroid cancer tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1% TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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