

Phospho-GSK3 beta (Ser25) Ab

Cat.#: AF7314 Concn.: 1mg/ml Mol.Wt.: 46kDa Size: 100ul.200ul Source: Rabbit Clonality: Polyclonal

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

Reactivity: Human, Mouse, Rat

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

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-GSK3 beta (Ser25) Ab detects endogenous levels of

GSK3 beta only when phosphorylated at Ser25.

A synthesized peptide derived from human GSK3 beta Immunogen:

around the phosphorylation site of Ser25.

Uniprot: P49841

Subcellular Location: Cytoplasm. Nucleus. Cell membrane. The phosphorylated

> form shows localization to cytoplasm and cell membrane. The MEMO1-RHOA-DIAPH1 signaling pathway controls localization of the phosophorylated form to the cell

membrane.

Tissue Specificity: Expressed in testis, thymus, prostate and ovary and weakly

> expressed in lung, brain and kidney. Colocalizes with EIF2AK2/PKR and TAU in the Alzheimer disease (AD) brain.

Similarity: Belongs to the protein kinase superfamily. CMGC Ser/Thr

protein kinase family. GSK-3 subfamily.

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.



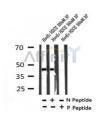
Western blot analysis of extracts from HuvEc H2O2 100uM 30', using Phospho-GSK3 beta (Ser25) Ab. Lane1 was treated with phospho-blocking peptide, Lane2 was treated with non-

phospho-blocking peptide.



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Western blot analysis of Phospho-GSK3 beta (Ser25) in lysates of HuvEc H2O2 100uM 30', using Phospho-GSK3 beta (Ser25) Ab(AF7314).



AF7314 at 1/100 staining mouse brain tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue 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.



AF7314 at 1/100 staining human brain tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue 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 , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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