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Product Information

Anti-phospho-Tau (pSer²⁶²)

Developed in Rabbit, Affinity Isolated Antibody

Product Number **T 7569**

Product Description

Anti-phospho-Tau (pSer²⁶²) is developed in rabbit using a synthetic phosphopeptide derived from the region of human Tau that contains serine 262 as immunogen. The serum is affinity purified using epitope-specific affinity chromatography. The antibody is preadsorbed to remove any reactivity towards a non-phosphorylated Tau.

Anti-phospho-Tau (pSer²⁶²) recognizes human, mouse and rat Tau (pSer²⁶²), 45-68 kDa. It has been used in immunoblotting applications.

Tau is a microtubule-associated phosphoprotein (MAP), localized in neuronal axons. It promotes tubulin polymerization and stabilizes microtubules.¹ The biological activity of Tau is regulated by its degree of phosphorylation.^{1,2} Hyperphosphorylated Tau is the major protein of the paired helical filaments (PHFs), which make up the pathological neurofibrillary tangles of Alzheimer's disease (AD). The PHFs are also found in the lesions of other central nervous system disorders.^{3,4}

Tau phosphorylation involves numerous kinases: glycogen synthase kinase 3 β (GSK-3 β), MARK kinase, MAP kinase, protein kinase A and C, cyclin-dependent kinase 5 (Cdk5), p38 kinase, c-Jun N-terminal kinase, and casein kinase II.^{1,2,5,6,7} Combined Tau protein kinase II (TPKII), which consists of Cdk5 and GSK-3 β , is the most potent phosphorylation agent indirectly involved in the regulation of the phosphorylation state of Tau in neuronal cells.^{6,8} In addition, Tau is phosphorylated *in vitro* by osmotic cellular stress, which activates the stress-activated protein kinases (SAPKs).

To date, a total of 25 abnormal phosphorylation sites have been identified on hyperphosphorylated Tau in AD brain.¹⁰ Normal Tau has approximately eight phosphorylation sites. The abnormal phosphorylation occurs usually on serine and threonine residues. Specifically, TPKII phosphorylates serines 202 and 404.

GSK-3 β transfection phosphorylates serines 199, 202, 235, 396, 404 and 413, and threonines 205 and 231. These sites are among the major abnormal phosphorylation sites of tau.¹¹ Phosphorylation on these sites reduces the ability of given Tau species to promote microtubule self-assembly.^{11,12} Okadaic acid increases phosphorylation at threonine 231 and serines 235, 396 and 404. Phosphorylated serine 422 was found in the biopsies of brains from patients with Down syndrome, amyotrophic lateral sclerosis, corticobasal degeneration, and Pick's disease. It was absent from control group of normal brains.¹³

The opposite process, Tau dephosphorylation, is controlled by different protein phosphatases expressed in neurons. Protein phosphatases PP2A and PP2B efficiently dephosphorylate Tau *in vitro* and restore biological activity in the assembly of microtubules.^{3,10,14}

Recently it was discovered that propyl isomerase (Pin 1) interacts with Tau hyperphosphorylated on threonine 231 and restores the ability of Tau to bind to microtubules.

Reagent

The antibody is supplied at approximately 0.5 mg/ml in 100 μ l of Dulbecco's phosphate buffered saline (without Mg²⁺ and Ca²⁺), pH 7.3, with 50% glycerol, 1.0 mg/ml BSA (IgG, protease free) and 0.05% sodium azide.

Precautions and Disclaimer

Due to the sodium azide content, a material safety data sheet (MSDS) for this product has been sent to the attention of the safety officer of your institution. Consult the MSDS for information regarding hazards and safe handling practices.

Storage/Stability

Store at -20 °C. Due to the presence of 50% glycerol the antibody will remain in solution. For extended storage, centrifuge the vial briefly before opening and prepare working aliquots. To ensure accurate dilutions

mix gently, remove excess solution from pipette tip with clean absorbent paper, pipette slowly. The antibody is stable for at least six months when stored appropriately. Working dilutions should be discarded if not used within 12 hours.

Product Profile

The supplied reagent is sufficient for 10 immunoblots.

A recommended working dilution of 1:1000 is determined by immunoblotting.

Note: In order to obtain best results in different techniques and preparations we recommend determining optimal working concentration by titration test.

Results

Peptide Competition

- NIH 3T3 cell extracts spiked with human recombinant tau left untreated (1) or treated with PKA to become phosphorylated (2-5) were resolved by SDS-PAGE on a 10% polyacrylamide gel and transferred to PVDF.
- Membranes were blocked with a 5% BSA-TBST buffer overnight at 4 °C.
- After blocking, membranes were preincubated with different peptides as follow:

Lane 1,2	no peptide
Lane 3	non phosphorylated peptide corresponding to the immunogen
Lane 4	a generic phosphotyrosine containing peptide
Lane 5	immunogen
- After preincubation membranes were incubated with Tau [pS²⁶²] antibody for two hours at room temperature in a 1% BSA-TBST buffer
- After washing, membranes were incubated with goat F(ab')₂ anti-rabbit IgG alkaline phosphatase and signals were detected using the Pierce SuperSignal[®] method.

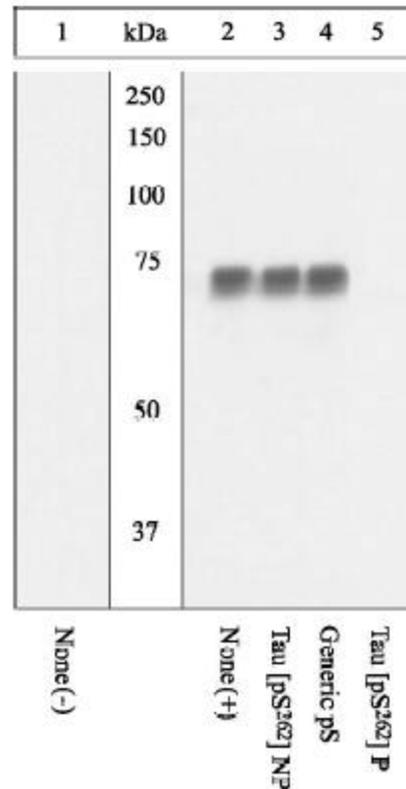


Figure 1 Peptide competition

The data in Figure 1 show that only the peptide corresponding to Tau [pS²⁶²] blocks the antibody signal, thereby demonstrating the specificity of the antibody.

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