Product Datasheet

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Anti-TAU Rabbit pAb

WL03184

For Research Use Only. Not For Use In Diagnostic Procedures

Product Information

Product name Anti-TAU Rabbit pAb

Source Rabbit

Species reactivity Human, Mouse, Rat

Tested applications Western blot 1:1000-1:2000

Immunohistochemistry 1:100-1:400 Immunofluorescence 1:100-1:400

Molecular Wt. 50-80 kDa

Cytoplasm. Cell membrane. Cytoplasm. Cell projection.

Pack size 50/100/200/500/1000μl

Storage Store at -20°C. Avoid freeze/thaw cycles.

Storage buffer Supplied in 20 mM phosphate (pH 7.5), 150 mM NaCl, 100 µg/ml

BSA, 50% glycerol and less than 0.02% sodium azide

General Information

Background Tau is a heterogeneous microtubule-associated protein that promotes

and stabilizes microtubule assembly, especially in axons.

Phosphorylation decreases the ability of tau to bind to microtubules.

Neurofibrillary tangles are a major hallmark of Alzheimer's disease;
these tangles are bundles of paired helical filaments composed of
hyperphosphorylated tau. Tau is found to be the major component of the
paired helical filaments (PHFs) found in the brains of patients with
Alzheimer disease (AD). Tau is hyperphosphorylated in PHFs, and specific

phosphorylation sites have been implicated in the loss of Taus association with the membrane cortex during AD disease state.

Immunogen Polyclonal antibody is produced by immunizing animals with a synthetic

peptide of TAU.

Purification Polyclonal antibody was purified by Protein A affinity chromatography.

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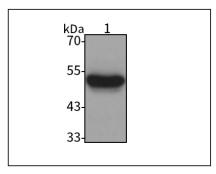


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Product Images



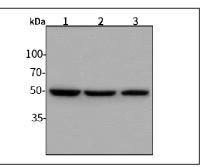
Western blot-Anti-TAU pAb

Lane 1: Human SH-SY5Y cell lysate 30µg Separation gel: 10% polyacrylamide Electrophoresis: 100V, 4°C, 3h Transmembrane: 100V, 4°C, 1h

 $\textbf{Blocking:}~5\%~\text{w/v}~\text{nonfat}~\text{dry}~\text{milk},~1\times\text{TBST},~\text{at}~\text{RT}~\text{with}~\text{gentle}~\text{shaking}$

Primary antibody: 1:1000 in blocking buffer, 4°C, overnight Secondary antibody (WLA023a):1:5000-1:10000, 45min

Detection: ECL, 30s-2min



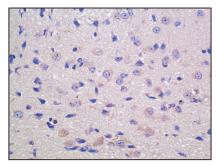
Western blot-Anti-TAU pAb

Lane 1: Mouse cerebellum tissue lysate 30µg Lane 2: Mouse ridge tissue lysate 30µg Lane 3: Rat brain tissue lysate 30µg Separation gel: 10% polyacrylamide Electrophoresis: 100V, 4°C, 3h Transmembrane: 100V, 4°C, 1h

 $\textbf{Blocking:} \, 5\% \, \text{w/v} \, \text{nonfat} \, \text{dry milk,} \, 1 \times \text{TBST, at RT with gentle shaking}$

Primary antibody: 1:1000 in blocking buffer, 4°C, overnight Secondary antibody (WLA023a):1:5000-1:10000, 45min

Detection: ECL, 30s-2min



Immunohistochemistry-Anti-TAU pAb

Sample: Mouse brain tissue

Antigen retrieval: pH 9.0 Tris-EDTA buffer Primary antibody: 1:200, 4°C, overnight Secondary antibody-Biotin: 1:150, 37°C, 1h Streptavidin-HRP: 1:200, 37°C, 30min

Color Developing: DAB

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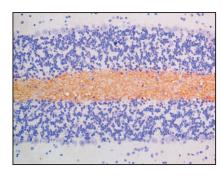
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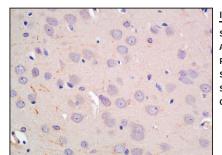
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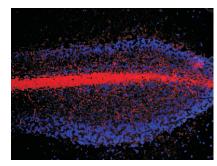
Immunohistochemistry-Anti-TAU pAb

Sample: Mouse cerebellun tissue Antigen retrieval: pH 9.0 Tris-EDTA buffer Primary antibody: 1:200, 4°C, overnight Secondary antibody-Biotin: 1:150, 37°C, 1h Streptavidin-HRP: 1:200, 37°C, 30min Color Developing: DAB



Immunohistochemistry-Anti-TAU pAb

Sample: Rat brain tissue Antigen retrieval: pH 9.0 Tris-EDTA buffer Primary antibody: 1:200, 4°C, overnight Secondary antibody-Biotin: 1:150, 37°C, 1h Streptavidin-HRP: 1:200, 37°C, 30min Color Developing: DAB



Immunofluorescence-Anti-TAU pAb

Sample: Rat cerebellun tissue Primary antibody: 1:200, 4°C, overnight

Secondary antibody-CY3: 1:200, at room temperature, 1h