

## Anti-Phospho-MAPT-S404 antibody [S5855RM] (STJ11105855)

ST.111105855

## **GENERAL INFORMATION**

Product Type Primary antibodies

**Short Description** 

Applications WB/ELISA Host/Source Rabbit

Reactivity Human/Mouse/Rat

## **PRODUCT PROPERTIES**

Clonality Monoclonal
Clone ID 55855RM

Concentration Lot specific
Conjugation Unconjugated
Purification Affinity purification
Dilution Range WB:1:2000-1:6000

ELISA:Recommended starting concentration is 1 Mu g/mL. Please optimize the concentration based on your specific assay

requirements.

Formulation PBS with 0.05% Proclin300, 0.05% BSA, 50% Glycerol, pH 7.3.

**Isotype** IgG

Storage Instruction Store at-20°C for up to 1 year from the date of receipt, and avoid repeat freeze-thaw cycles.

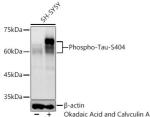
## **TARGET INFORMATION**

Gene ID 4137
Gene Symbol MAPT
Uniprot ID TAU\_HUMAN
Immunogen

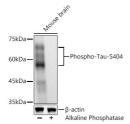
Immunogen Immunogen Region

Specificity A synthetic phosphorylated peptide around S404 of human Tau.

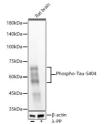
Immunogen Sequence



— + Ukadaic Acid and Calyculin A Western blot analysis of lysates from SH-SYSY cells using A Phospho-Tau-S404 Rabbit monoclonal artibody (STJ1110855) adA 1:500 dilution. SH-SYSY cells were and Calyculin A (100Mg) at 37 A\*C for 30 minutes after serum-starvation overnight. Secondary antibody:HBP Coat Anti-Rabbit 1gG (H+1) (STJ5000856) at 1:10000 dilution. Lysates/proteins: 30Å Mu gper lane. Blocking buffer: 3% norfat dry milk in TBST. Detection:ECL Basic Kit. Exposuretime A 10s.



Western blot analysis of lysates from Mouse brain using AProsphor-Tau-Su4P fabilit monoclonal antibody, (STJ 11105855) at A1:5000 dilution. Mouse brain hysates were treated by Lambda-PP mixed solution (Iul) at 30 A°C for 30 minutes. Secondary antibody-HRP Goa ART-Habbit 19G (H+1) GSTJ5000856) at 1:1000 dilution. Lysates/proteins: 30A Mu g per lane. Blocking buffer: 3% nortiat dry milk in 18ST. Detection.ECL



Western blot analysis of lysates from rat brain, using Phospho-Tau-S404 Rabbit monoclonal antibody (STJ11105855) at 1:5000 dilution. Rat brain cells were treated by Lambda-PP mixed solution (1u) at 30 A°C for 30 mínutes. Secondary antibody HRP Goat Anti-Rabbit IgG (H+I) (STJS000856) at 1:1000 dilution. Lysates/proteins: 25 Mu g per lane. Blocking buffer: 3% confact dry milk in TBST. Detection: ECL Basic Kit.