

# Rabbit anti Phosphothreorine(pThr) Polyclonal Antibody Alternative Name(s): nan

Order Information

- Description: Phosphothreonine(pThr)
- Catalogue: 620-670
- Lot: See label
- Size: 100ug/200ul
- Host: Rabbit
- Clone: nan
- Application: IHC(P), WB
- Reactivity: all species

# ANTIGEN PREPARATION

A chemically linked phosphothreonine.

#### BACKGROUND

Protein phosphorylation is involved in cell signaling pathways. These cascades are mediated by three types of kinases: serine, threonine and tyrosine kinases which phosphorylate serine, threonine and tyrosine amino acid side chains. These three amino acids are phosphorylated by its specific kinases. These processes are regulated by kinases and phosphatases.

## PURIFICATION

The Rabbit IgG is purified by Epitope Affinity Purification

# FORMULATION

This affinity purified antibody is supplied in sterile Phosphatebuffered saline (pH7.2) containing antibody stabilizer

## SPECIFICITY

This antibody recognizes phosphothreonine (pThr) only. It does not cross react with phosphoserine or phosphotyrosine

#### STORAGE

The antibodies are stable for 24 months from date of receipt when stored at -200C to -700C. The antibodies can be stored at 20C-80C for three month without detectable loss of activity. Avoid repeated freezing-thawing cycles.

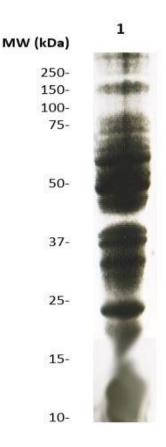
## **APPLICATIONS/SUGGESTED WORKING DILUTIONS\***

- Western Blot: 0.1-1 µg/ml
- ELISA: 0.01-0.1 µg/ml
- Immunoprecipitation: 2-5 µg/ml
- IHC: 2-10 µg/ml
- Flow cytometry: Not tested
- Molecular Weight: nan
- Positive Control: Kidney Tissue
- Cellular Location: Cell Membrane

\*Optimal dilutions should be determined by researchers for the specific applications.

# FOR RESEARCH USE ONLY.





Western Blot: The cell lysate derived from EGFstimulated A431was resolved onto 12% SDS-PAGE and immunoblotted by Rabbit anti Phosphothreonine (Cat#620-670) at 1:500. A panel of phosphorylated proteins was observed.

#### REFERENCES

Trinidad JC, Thalhammer A, Specht CG, Lynn AJ, Baker PR, Schoepfer R, Burlingame AL. Quantitative analysis of synaptic phosphorylation and protein expressio. Mol. Cell Proteomics 7 (4): 684–96, 2008