

Anti-CPI17 Antibody

Catalog # AP53927

#### Specification

# Anti-CPI17 Antibody - Product Information

Application Primary Accession Reactivity Host Clonality Calculated MW WB, IF, IHC <u>O96A00</u> Human, Mouse, Rat Rabbit Polyclonal 16693

# **Anti-CPI17 Antibody - Additional Information**

Gene ID 94274

**Other Names** CPI17; PPP1INL; Protein phosphatase 1 regulatory subunit 14A; 17 kDa PKC-potentiated inhibitory protein of PP1; Protein kinase C-potentiated inhibitor protein of 17 kDa; CPI-17

Target/Specificity Recognizes endogenous levels of CPI17 protein.

**Dilution** WB~~1/500 - 1/1000 IF~~1/50 - 1/200 IHC~~1:100~500

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

# Anti-CPI17 Antibody - Protein Information

Name PPP1R14A

Synonyms CPI17, PPP1INL

Function

Inhibitor of PPP1CA. Has over 1000-fold higher inhibitory activity when phosphorylated, creating a molecular switch for regulating the phosphorylation status of PPP1CA substrates and smooth muscle contraction.

Cellular Location Cytoplasm.



Tissue Location

Isoform 1 is detected in aorta and testis. Isoform 2 is detected in aorta.

#### Anti-CPI17 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- <u>Western Blot</u>
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

#### **Anti-CPI17 Antibody - Images**



Western blot analysis of CPI17 expression in A2780 (A), HEK293T (B), EC9706 (C), mouse brain (D), rat brain (E) whole cell lysates.



Immunohistochemical analysis of CPI17 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.





Immunofluorescent analysis of CPI17 staining in Jurkat cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

# Anti-CPI17 Antibody - Background

Rabbit polyclonal antibody to CPI17