## **TCP1 Antibody**



## PACO55802

Reactivity:

## **Product Information**

Size: Protein Background:

50ug Molecular chaperone; assists the folding of proteins upon ATP hydrolysis. As part of the

BBS/CCT complex may play a role in the assembly of BBSome, a complex involved in ciliogenesis regulating transports vesicles to the cilia. Known to play a role, in vitro, in

ciliogenesis regulating transports vesicies to the cilia. Known to play a role, in vitro, ir

Human the folding of actin and tubulin.

Source: Gene ID:

Rabbit TCP1

Isotype: Uniprot

IgG P17987

Applications: Synonyms:

ELISA, IHC, IF T-complex protein 1 subunit alpha (TCP-1-alpha) (CCT-alpha), TCP1, CCT1 CCTA

Recommended dilutions: Immunogen:

ELISA:1:2000-1:10000, IHC:1:500-1:1000, Recombinant Human T-complex protein 1 subunit & Alpha; protein (406-506AA).

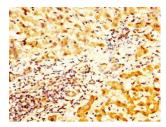
Stora

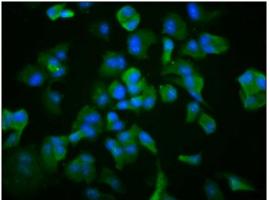
IF:1:200-1:500

Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, pH 7.4

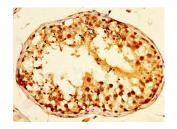
## **Product Images**





IHC image of PACO55802 diluted at 1:800 and staining in paraffinembedded human liver cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunofluorescence staining of MCF-7 cells with PACO55802 at 1:266, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



IHC image of PACO55802 diluted at 1:800 and staining in paraffinembedded human testis tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.