

Basic Information

Product Name	Anti-TCP1 Antibody (Clone#2E7)	
Gene Name	TCP1	
Source	Mouse	
Clonality	Monoclonal	
Isotype	IgG1	
Species Reactivity	human	
Tested Application	WB, IHC, ICC/IF, FCM	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence at the C-terminus of human TCP1 alpha, different from the related mouse sequence by one amino acid, and from the related rat sequence by two amino acids.	
Concentration	200ug/ml	
Purification	protein G purified.	
Observed MW	60 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 Immunocytochemistry/Immunofluorescence (ICC/IF): 1:50-400 Flow Cytometry (Fixed): 1:50-200 (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied.

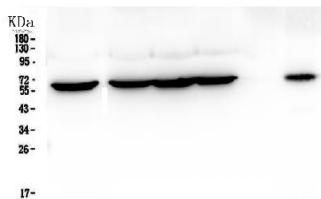
Background Information

T-complex protein 1 subunit alpha is a protein that in humans is encoded by the TCP1 gene. The protein encoded by this gene is a molecular chaperone that is a member of the chaperonin containing TCP1 complex (CCT), also known as the TCP1 ring complex (TRiC). This complex consists of two identical stacked rings, each containing eight different proteins. Unfolded polypeptides enter the central cavity of the complex and are folded in an ATP-dependent manner. The complex folds various proteins, including actin and tubulin. Alternate transcriptional splice variants of this gene, encoding different isoforms, have been characterized. In addition, three pseudogenes that appear to be derived from this gene have been found.

Reference

Anti-TCP1 Antibody (Clone#2E7)被引用在1文献中。

Selected Validation Data



① human Hela ② human MCF-7 ③ human COLO-320
④ human HepG2 ⑤ Human A431 ⑥ human HT1080

Western blot analysis of anti-TCP1 antibody (M02389). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Hela whole cell lysates,

Lane 2: human MCF-7 whole cell lysates,

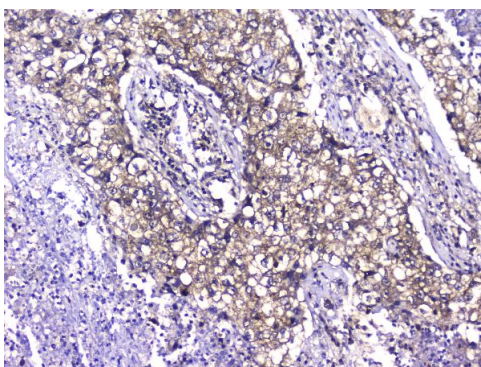
Lane 3: human COLO-320 whole cell lysates,

Lane 4: human HepG2 whole cell lysates,

Lane 5: human A431 whole cell lysates,

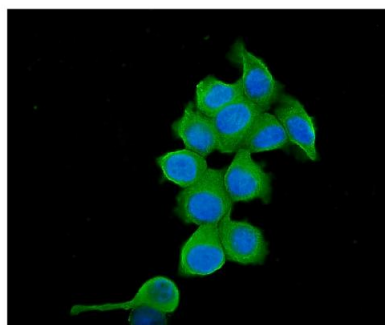
Lane 6: human HT1080 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with mouse anti-TCP1 antigen affinity purified monoclonal antibody (M02389) and probed with a goat anti-mouse IgG-HRP secondary antibody (Catalog # BA1050). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for TCP1 at approximately 60 kDa. The expected band size for TCP1 is at 60 kDa.



IHC analysis of TCP1 using anti-TCP1 antibody (M02389).

TCP1 was detected in a paraffin-embedded section of human lung cancer tissue. The tissue section was incubated with mouse anti-TCP1 Antibody (M02389) at a dilution of 1:200 and developed using HRP Conjugated mouse IgG Super Vision Assay Kit (Catalog # SV0001) with DAB (Catalog # AR1027) as the chromogen.



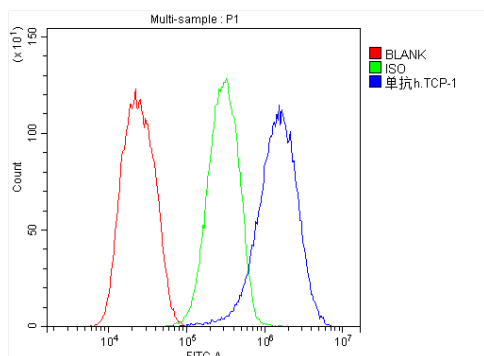
IF analysis of TCP1 using anti-TCP1 antibody (M02389).

TCP1 was detected in an immunocytochemical section of MCF-7 cells.

Dylight488-conjugated Anti-mouse IgG Secondary Antibody

(green)(Catalog#BA1126) was used as secondary antibody. The section

was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of HepG2 cells using anti-TCP1 antibody (M02389).

Overlay histogram showing HepG2 cells stained with A04887-1 (Blue line).

The cells were blocked with 10% normal goat serum. And then incubated

with mouse anti-TCP1 Antibody ((M02389, 1:100). DyLight®488

conjugated goat anti-mouse IgG (BA1126, 1:100) was used as secondary

antibody. Isotype control antibody (Green line) was mouse IgG (Catalog # BA1046) (1:100) used under the same conditions. Unlabelled sample (Red

line) was also used as a control.