

PRODUCT DATA SHEET

Bioworld Technology,Inc.

TRAP220 (phospho-T1457) polyclonal antibody

Catalog: BS67078 Host: Rabbit Reactivity: Human, Mouse, Rat

BackGround:

Component of the Mediator complex, a coactivator involved in the regulated transcription of nearly all RNA polymerase II-dependent genes. Mediator functions as a bridge to convey information from gene-specific regulatory proteins to the basal RNA polymerase II transcription machinery. Mediator is recruited to promoters by direct interactions with regulatory proteins and serves as a scaffold for the assembly of a functional preinitiation complex with RNA polymerase II and the general transcription factors (PubMed:10406464, PubMed:11867769, PubMed:12037571, PubMed:12218053, Med:12556447, PubMed:14636573, PubMed:15340084, PubMed:15471764. PubMed:15989967. Pub-Med:16574658, PubMed:9653119).

Product:

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Molecular Weight:

~ 170 kDa

Swiss-Prot:

Q15648

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB (1/500 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500), ChIP (Use at an assay dependent concentration)

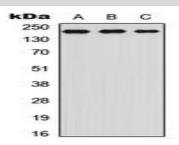
Storage&Stability:

Store at $4\,\mathrm{C}$ short term. Aliquot and store at $-20\,\mathrm{C}$ long term. Avoid freeze-thaw cycles.

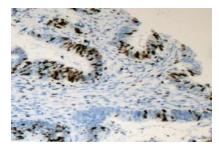
Specificity:

Recognizes endogenous levels of TRAP220 (pT1457) protein.

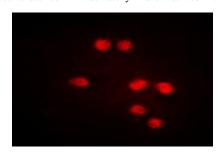
DATA:



Western blot analysis of TRAP220 (pT1457) expression in Jurkat (A), HeLa (B), NIH3T3 (C) whole cell lysates.



Immunohistochemical analysis of TRAP220 (pT1457) staining in human colon cancer 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 TRAP220 (pT1457) 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.

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Cells were washed with PBST and incubated with a DyLight

Note:

594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

For research use only, not for use in diagnostic procedure.

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