

ALPL Rabbit monoclonal antibody

Catalog: MB66312

Host: Rabbit

Reactivity: Human, Rat

Background:

The family of alkaline phosphatases in humans is comprised of four members: intestinal, placental, placental-like (germ cell type), and tissue nonspecific. The tissue nonspecific isozyme is also known as the liver/bone/kidney alkaline phosphatase due to its expression in these tissues. It is also highly expressed in embryonic stem (ES) and embryonic carcinoma (EC) cells and is lost as these cells undergo differentiation. The TRA-2-54 (2J) Mouse mAb specifically detects this isozyme and does not detect the other family members, making it a useful tool for tracking the pluripotency status of ES and EC cells in culture.

Product:

Liquid in 50mM Tris-Glycine (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide and 0.05% BSA.

Molecular Weight:

~ 80 kDa

Swiss-Prot:

P05186

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/1000), IHC (1/50 - 1/100), IF/ICC (1/50 - 1/100), IP (1/10 - 1/50)

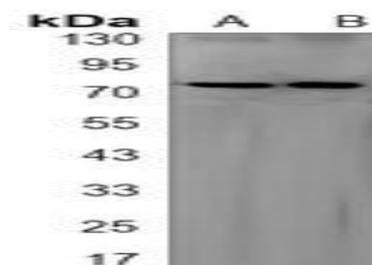
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

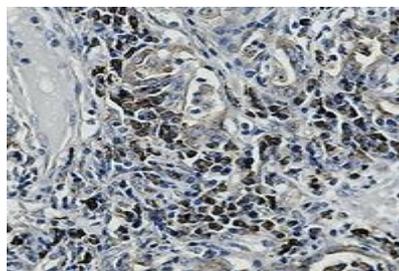
Specificity:

Recognizes endogenous levels of ALPL protein.

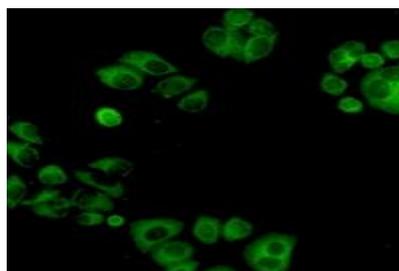
DATA:



Western blot analysis of ALPL expression in HeLa (A), C6 (B) whole cell lysates.



Immunohistochemical analysis of ALPL staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.76). 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 ALPL staining in HeLa 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 humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated sec-

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PRODUCT DATA SHEET

Bioworld Technology, Inc.

secondary antibody (green) in PBS at room temperature in the dark.

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

Note:

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