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Schillerstr. 5

AP53172PU-N Polyclonal Antibody to PATL1 (C-term) - Aff - Purified

Alternate names: OK/KNS-cl.5, PAT1-like protein 1, Protein PAT1 homolog 1

Quantity: 0.4 ml

 Concentration:
 lot specific

 Uniprot ID:
 086TB9

 NCBI:
 NP_689929

 GeneID:
 219988

 Host / Isotype:
 Rabbit / Ig

Immunogen: KLH conjugated synthetic peptide between 496~526 amino acids from the C-terminal

region of human PATL1

Format: State: Liquid purified lg fraction

Purification: Protein A column, followed by peptide affinity purification **Buffer System:** PBS containing 0.09% (W/V) Sodium Azide as preservative

Applications: ELISA: 1/1000.

Western Blot: 1/100-1/500. Flow Cytometry: 1/10-1/50.

Immunohistochemistry on Paraffin Sections: 1/50-1/100.

Other applications not tested. Optimal dilutions are dependent on conditions and

should be determined by the user.

Specificity: This antibody recognizes Human PATL1 (C-term).

Add. Information: Molecular Weight: 86850 Da

Storage: Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing. Shelf life: one year from despatch.

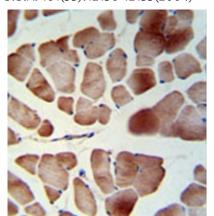
General Readings: Scheller, N., et al. Biochim. Biophys. Acta 1773(12):1786-1792(2007)

Beausoleil, S.A., et al. Proc. Natl. Acad. Sci. U.S.A. 101(33):12130-12135(2004)

Pictures: Immunohistochemistry analysis in

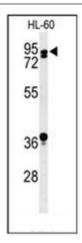
formalin fixed and paraffin embedded skeletal muscle reacted with PATL1 Antibody (C-term) Cat.-No AP53172PU-N, which was peroxidase conjugated to the secondary antibody and followed by DAB staining. This data demonstrates the use

of the PATL1 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.





Western blot analysis of PATL1 Antibody (C-term) Cat.-No AP53172PU-N in HL-60 cell line lysates (35ug/lane). PATL1 (arrow) was detected using the purified Pab.



Flow cytometric analysis of HL-60 cells using PATL1 Antibody (C-term) Cat.-No AP53172PU-N (right histogram) compared to a negative control cell (left histogram). FITC-conjugated goat-antirabbit secondary antibodies were used for the analysis.

