

Product datasheet for AP51489PU-N

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EYA4 (C-term) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: FC, IHC, WB

Recommended Dilution: ELISA: 1/1000.

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

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

Reactivity: Human
Host: Rabbit

Isotype: lg

Clonality: Polyclonal

Immunogen: KLH conjugated synthetic peptide from the C-terminal region of human EYA4

Specificity: This antibody recognizes Human EYA4 (C-term).

Formulation: PBS containing 0.09% (W/V) Sodium Azide as preservative

State: Aff - Purified

State: Liquid purified Ig fraction

Concentration: lot specific

Purification: Protein A column, followed by peptide affinity purification

Conjugation: Unconjugated

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

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: EYA transcriptional coactivator and phosphatase 4

Database Link: Entrez Gene 2070 Human

<u>095677</u>

Background: EYA4 encodes a member of the eyes absent (EYA) family of proteins. The encoded protein

may act as a transcriptional activator through its protein phosphatase activity, and it may be important for eye development, and for continued function of the mature organ of Corti.

Synonyms: Eyes absent homolog 4

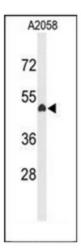




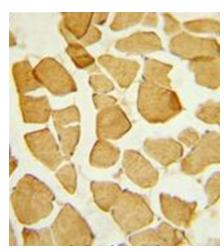
Note: Molecular Weight: 69505 Da

Protein Families: Druggable Genome, Phosphatase, Transcription Factors

Product images:



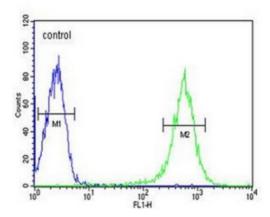
Western blot analysis of EYA4 Antibody (C-term) in A2058 cell line lysates (35ug/lane). EYA4 (arrow) was detected using the purified Pab.



Formalin-fixed and paraffin-embedded human skeletal muscle reacted with EYA4 Antibody (Cterm), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining.



HL-60



Flow cytometry analysis of HL-60 cells using EYA4 Antibody (C-term) (bottom histogram) compared to a negative control cell (top histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.