

Anti-IL-32 Antibody

Rabbit polyclonal antibody to IL-32 Catalog # AP59774

Specification

Anti-IL-32 Antibody - Product Information

Application WB, IHC
Primary Accession P24001
Reactivity Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 26676

Anti-IL-32 Antibody - Additional Information

Gene ID 9235

Other Names

NK4; TAIF; Interleukin-32; IL-32; Natural killer cells protein 4; Tumor necrosis factor alpha-inducing factor

Target/Specificity

KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human IL-32. The exact sequence is proprietary.

Dilution

WB~~WB (1/500 - 1/1000), IH (1/100 - 1/200) IHC~~1:100~500

Format

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.

Storage

Store at -20 °C.Stable for 12 months from date of receipt

Anti-IL-32 Antibody - Protein Information

Name IL32

Synonyms NK4, TAIF

Function

Cytokine that may play a role in innate and adaptive immune responses. It induces various cytokines such as TNFA/TNF-alpha and IL8. It activates typical cytokine signal pathways of NF-kappa-B and p38 MAPK.

Cellular Location



Secreted.

Tissue Location

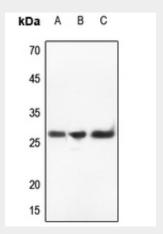
Selectively expressed in lymphocytes. Expression is more prominent in immune cells than in non-immune cells

Anti-IL-32 Antibody - Protocols

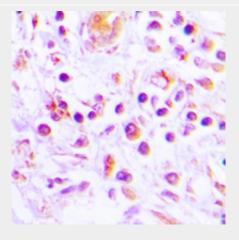
Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Anti-IL-32 Antibody - Images



Western blot analysis of IL-32 expression in mouse liver (A), rat liver (B), rat kidney (C) whole cell lysates.



Immunohistochemical analysis of IL-32 staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with





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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.

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