

Product datasheet for AP52188PU-N

IL17F (N-term) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: FC, WB

Recommended Dilution: ELISA: 1/1000.

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

Reactivity: Human
Host: Rabbit
Isotype: Ig

Clonality: Polyclonal

Immunogen: KLH conjugated synthetic peptide between 16-46 amino acids from the N-terminal region of

human IL17F.

Specificity: This antibody recognizes Human Interleukin-17F / IL17F (N-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: interleukin 17F

Database Link: Entrez Gene 112744 Human

Q96PD4



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Background: The protein encoded by this gene is a cytokine that shares sequence similarity with IL17. This

cytokine is expressed by activated T cells, and has been shown to stimulate the production of several other cytokines, including IL6, IL8, and CSF2/GM_CSF. This cytokine is also found to inhibit the angiogenesis of endothelial cells and induce endothelial cells to produce IL2,

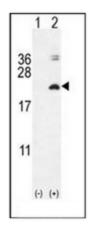
TGFB1/TGFB, and monocyte chemoattractant protein-1. [provided by RefSeq].

Synonyms: IL24, Cytokine ML-1, Interleukin-24, IL-24

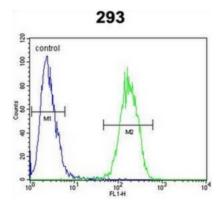
Note: Molecular Weight: 18045 Da

Protein Families: Druggable Genome, Secreted Protein

Product images:



Western blot analysis of IL17F (arrow) using Interleukin-17F / IL17F Antibody (N-term). 293 cell lysates (2 ug/lane) either nontransfected (Lane 1) or transiently transfected with the IL17F gene (Lane 2).



Flow cytometric analysis of 293 cells using Interleukin-17F / IL17F Antibody (N-term) (right histogram) compared to a negative control cell (left histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis