Leader in Biomolecular Solutions for Life Science



Catalog No.: A24007 Recombinant



## **Basic Information**

Observed MW 19kDa

Calculated MW 16kDa

**Category** Primary antibody

Applications WB,FC (intra),ELISA

Cross-Reactivity Human

CloneNo number ARC62039

# Background

The protein encoded by this gene is a cytokine that acts as a regulator of a variety of hematopoietic cells. This cytokine stimulates cell proliferation and prevents apoptosis. It functions through the interleukin 9 receptor (IL9R), which activates different signal transducer and activator (STAT) proteins and thus connects this cytokine to various biological processes. The gene encoding this cytokine has been identified as a candidate gene for asthma. Genetic studies on a mouse model of asthma demonstrated that this cytokine is a determining factor in the pathogenesis of bronchial hyperresponsiveness.

### **Recommended Dilutions**

WB	1:1000 - 1:4000
FC (intra)	1:500 - 1:1000
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

## **Immunogen Information**

**Gene ID** 3578 Swiss Prot P15248

#### Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

### Synonyms

P40; HP40; IL-9; IL9

### Contact

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# **Product Information**

**Source** Rabbit **lsotype** lgG Purification Affinity purification

### Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.



Western blot analysis of lysates from 293T(Transfection) and control 293T cells, using IL9 Rabbit mAb (A24007) at 1:1000 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 30s.





1X10^6 293T cells (negative control,left) and 293T(Transfection,right) cells were intracellularly-stained with IL9 Rabbit mAb(A24007,2µg/mL,orange line) or ABflo® 488 Rabbit IgG isotype control (A22069,2µg/mL,blue line), followed by FITC conjugated goat anti-rabbit pAb (1:200 dilution) staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry:1X10^6 293T(Transfection) cells were intracellularly-stained with ABflo® 488 Rabbit IgG isotype control (A22069,2 µg/mL,left) or IL9 Rabbit mAb(A24007,2 µg/mL,right).