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IL-9 Rabbit pAb

Catalog Number: bs-2428R

Target Protein: IL-9

Concentration: 1mg/ml

Form: Liquid

Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Applications: WB (1:500-2000), IHC-P (1:100-500), IHC-F (1:100-500), IF (1:200-800)

Reactivity: Mouse, Rat

Predicted MW: 16 kDa Entrez Gene: 16198

Swiss Prot: P15247

Source: KLH conjugated synthetic peptide derived from mouse IL-9: 21-120/144.

Purification: affinity purified by Protein A

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

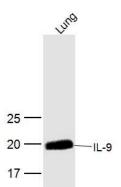
Background: The protein encoded by this gene is a cytokine that actsas a regulator of a variety of

hematopoietic cells. This cytokinestimulates 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 geneencoding this cytokine has been identified as a candidate gene for as thma. Genetic studies on a mouse model of as thma demonstrated that this cytokine is a determining factor in the pathogenesis of bronchial hyperresponsiveness.

[provided by RefSeq, Jul 2008].

VALIDATION IMAGES

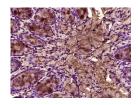


Sample: Lung (Mouse) Lysate at 40 ug Primary: Anti-IL-9 (bs-2428R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 16 kD Observed band size: 16 kD

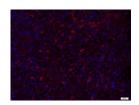


11 -

Tissue/cell: mouse lung tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-IL-9 Polyclonal Antibody, Unconjugated(bs-2428R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Paraformaldehyde-fixed, paraffin embedded (mouse colon tissue); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (IL-9) Polyclonal Antibody, Unconjugated (bs-2428R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse mammary tumor); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (IL-9) Polyclonal Antibody, Unconjugated (bs-2428R) at 1:400 overnight at 4°C, followed by a conjugated Goat Anti-Rabbit IgG antibody (bs-0295G-cy3) for 90 minutes, and DAPI for nuclei staining.

PRODUCT SPECIFIC PUBLICATIONS

[IF=5.6] Yun Sun. et al. Inhibitory effects of catalpol on DNCB-induced atopic dermatitis and IgE-mediated mast cells reaction. INT IMMUNOPHARMACOL. 2024 Jan;126:111274 ELISA, IHC; Mouse . 38041954

[IF=3.7] Yao, Xiuhua, et al. "Neutralization of IL $\,^{-}$ 9 ameliorates symptoms of experimental autoimmune myasthenia gravis in rats by decreasing effector T cells and altering humoral responses." Immunology (2014). Other; = "Rat" $\,^{-}$. 24850614

[IF=1.5] Lin, Yinyao, et al. "Expression of interleukin-9 and its upstream stimulating factors in rats with ischemic stroke." Neurological Sciences (2015): 1-8. IHC; = "Rat". 25652434