bs-2611R

[Primary Antibody]

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IL17C Rabbit pAb

DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GenelD: 27189 SWISS: Q9P0M4

Target: IL17C

Immunogen: KLH conjugated synthetic peptide derived from human IL-17C:

101-197/197.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

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 T cell-derived cytokine that

shares the sequence similarity with IL17. This cytokine was reported to stimulate the release of tumor necrosis factor alpha and interleukin 1 beta from a monocytic cell line. The expression of

this cytokine was found to be restricted to activated T cells.

[provided by RefSeq].

Applications: WB (1:500-2000)

IHC-P (1:100-500) **IHC-F** (1:100-500) **IF** (1:100-500)

Reactivity: Human, Rat

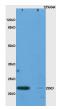
(predicted: Mouse, Rabbit,

Cow, Dog, Horse)

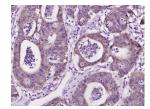
Predicted 20 kDa

Subcellular Secreted

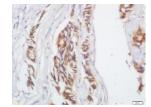
VALIDATION IMAGES



Sample: Lane1: Lung(Rat) Lysate at 30 ug Lane2: Pancreas(Rat) Lysate at 30 ug Primary: Anti-IL-17C (bs-2611R) at 1:200 dilution; Secondary: HRP conjugated Goat Anti-Rabbit IgG(bs-0295G-HRP) at 1: 3000 dilution; Predicted band size: 20kD Observed hand size · 22kD



Paraformaldehyde-fixed, paraffin embedded (human colon carcinoma); 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 (IL17C) Polyclonal Antibody, Unconjugated (bs-2611R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Tissue/cell: Human colon cancer tissue; 4% Paraformaldehyde-fixed and paraffinembedded; 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-IL17C Polyclonal Antibody, Unconjugated(bs-2611R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining

— SELECTED CITATIONS —

- [IF=6.676] Takuya Takeichi. et al. Ceramide analysis in combination with genetic testing may provide a precise diagnosis for self-healing collodion babies. J LIPID RES. 2022 Nov;:100308 IHC; Human. 36332686
- [IF=5.9] Huang et al. IL-17C is required for lethal inflammation during systemic fungal infection. (2016) Cell.Mol.Immunol. 13:474-83 IHC; Mouse. 26166766
- [IF=4.99] Huang, Jinling, et al. "IL-17C/IL-17RE Augments T Cell Function in Autoimmune Hepatitis." The Journal of Immunology (2016): 1600977. IHC ;="Mouse". 27956525
- [IF=2.952] Xiaoyan Pu. et al. Oxidative stress and expression of inflammatory factors in lung tissue of acute mountain sickness rats. Mol Med Rep. 2022 Feb;25(2):1-7 WB; Rat. 34913080

• [IF=preprint] Xiaoying Yang. et al. Serum pharmacochemistry profiling combined with molecular docking analysis to	
reveal the pharmacodynamic material basis of Lonicerae japonicae flos against acute lung injury. SSRN. Western blo; Rat. 10.2139/ssrn.4809218	Эť
,Nat. 10.2159/SSIII.4009210	