

Anti-IL-23 Antibody

Catalog Number: A01097-2

About IL23A

Like interleukin-27 (IL-27), IL-23 is a recently discovered member of the IL-6/IL-12 family of proinflammatory and immunoregulatory cytokines. It exists as a heterodimer composed of the IL-12p40 subunit and a novel p19 subunit. IL-23 is secreted by activated dendritic cells, macrophages, and monocytes. Its biological activities include enhancing the proliferation of memory T cells and the production of IFN-gamma, IL-12, and TNF-alpha from activated T cells, and can stimulate macrophages to produce TNF-alpha and nitric oxide. It has also been shown to possess potent anti-tumor and anti-metastatic activity in mouse models of cancer, suggesting a potential role for IL-23 in therapeutic treatment of cancer.

Overview

| Product Name | Anti-IL-23 Antibody |
|----------------------|---|
| Reactive Species | Human, Mouse |
| Description | Boster Bio Anti-IL-23 Antibody (Catalog # A01097-2). Tested in ELISA, WB, IHC-P, IF applications. This antibody reacts with Human, Mouse. |
| Application | ELISA, IF, IHC-P, WB |
| Clonality | Polyclonal |
| Formulation | IL-23 Antibody is supplied in PBS containing 0.02% sodium azide. |
| Storage Instructions | IL-23 antibody can be stored at 4°C up to one year. Antibodies should not be exposed to prolonged high temperatures. |
| Host | Rabbit |
| Uniprot ID | Q9NPF7 |

Technical Details

| Immunogen | Anti-IL-23 antibody was raised against a peptide corresponding to 17 amino acids near the amino terminus of human IL-23. The immunogen is located within amino acids 40 - 90 of IL-23. |
|----------------------------|--|
| Predicted Reactive Species | Guinea Pig, Horse, Pig, Rat |
| Cross Reactivity | At least two isoforms of SCARB1 are known to exist; this antibody can detect both isoforms. |
| Isotype | IgG |
| Form | Liquid |
| Concentration | 1 mg/mL |
| Purification | IL-23 Antibody is affinity chromatography purified via peptide column. |



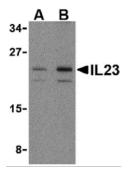
BOSTER BIOLOGICAL TECHNOLOGY 3942 B Valley Ave, Pleasanton, CA 94566

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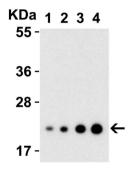
| Suggested Dilutions | Dilute the sample so that the expected range of concentrations fall within the detection range of this kit. If the expected range of concentration is unknown, a pilot test should be conducted to decide the |
|---------------------|--|
| | optimal dilution ratio for your samples. Some PubMed article(s) citing the expression level of this target are as follows: |
| | Boster Bio's internal QC testing used: |
| | WB: 0.25-2 ug/mL; IF: 20 ug/mL, IHC: 2 ug/mL. Antibody validated: Western Blot in human and mouse samples; Immunofluorescence and |
| | Immunohistochemistry in mouse samples. All other applications and species not yet tested. Optimal dilutions for each application should be determined by the researcher. |



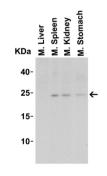
Anti-IL-23 Antibody (A01097-2) Images



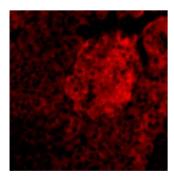
Western Blot Validation in Mouse Pancreas Tissue Lysate Loading: 15 ug of lysates per lane. Antibodies: IL-23 A01097-2 (A: 1 ug/mL, B: 2 ug/mL), 1h incubation at RT in 5% NFDM/TBST. Secondary: Goat anti-rabbit IgG HRP conjugate at 1:10000 dilution.



Western Blot Validation with Recombinant Protein Loading: 30 ng of human IL-23 recombinant protein per lane. Antibodies: IL-23 A01097-2 (1: 0.25 ug/mL, 2: 0.5 ug/mL, 3: 1 ug/mL and 4: 2ug/mL), 1h incubation at RT in 5% NFDM/TBST. Secondary: Goat anti-rabbit IgG HRP conjugate at 1:10000 dilution.



Western Blot Validation in Mouse Tissue Lysate Loading: 15 ug of lysates per lane. Antibodies: IL-23 A01097-2 (1 ug/mL), 1h incubation at RT in 5% NFDM/TBST. Secondary: Goat anti-rabbit IgG HRP conjugate at 1:10000 dilution.



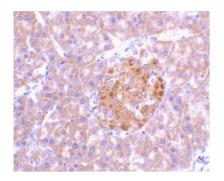
Immunofluorescence Validation of IL-23 in Mouse Pancreas Tissue

Immunofluorescent analysis of 4% paraformaldehyde-fixed mouse pancreas tissue labeling IL-23 with A01097-2 at 20 $\,$ ug/mL

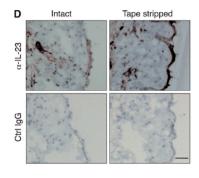
Immunohistochemistry Validation of IL-23 in Mouse Pancreas Tissue

Immunohistochemical analysis of paraffin-embedded mouse pancreas tissue using anti-IL-23 antibody (A01097-2) at 2 ug/ml. Tissue was fixed with formaldehyde and blocked with 10% serum for 1 h at RT; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were





incubated with primary antibody overnight at 4° C. A goat anti-rabbit IgG H&L (HRP) at 1/250 was used as secondary. Counter stained with Hematoxylin.



Induced Expression Validation of IL-23 in Mouse Keratinocytes of Epidermis (Yoon et al.

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