abcam

Product datasheet

Anti-IL-6 antibody [EPR21710] ab229381

Recombinant RabMAb

20 References 4 Images

Overview

Product name Anti-IL-6 antibody [EPR21710]

Description Rabbit monoclonal [EPR21710] to IL-6

Host species Rabbit

Tested applications Suitable for: WB, IP Species reactivity Reacts with: Mouse

Recombinant fragment. This information is proprietary to Abcam and/or its suppliers. **Immunogen**

Positive control WB: RAW 264.7 treated with LPS and BFA, whole cell lysate. IP: RAW 264.7 treated with LPS

and BFA, whole cell lysate.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: PBS, 40% Glycerol, 0.05% BSA

Purity Protein A purified

Clonality Monoclonal Clone number EPR21710

Isotype ΙgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab229381 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes	
WB		1/1000. Detects a band of approximately 24 kDa (predicted molecular weight: 24 kDa).	
IP		1/30.	

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Function

Cytokine with a wide variety of biological functions. It is a potent inducer of the acute phase response. Plays an essential role in the final differentiation of B-cells into Ig-secreting cells Involved in lymphocyte and monocyte differentiation. It induces myeloma and plasmacytoma growth and induces nerve cells differentiation Acts on B-cells, T-cells, hepatocytes, hematopoeitic progenitor cells and cells of the CNS. Also acts as a myokine. It is discharged into the bloodstream after muscle contraction and acts to increase the breakdown of fats and to improve insulin resistance.

Involvement in disease

Genetic variations in IL6 are associated with susceptibility to rheumatoid arthritis systemic juvenile (RASJ) [MIM:604302]. An inflammatory articular disorder with systemic-onset beginning before the age of 16. It represents a subgroup of juvenile arthritis associated with severe extraarticular features and occasionally fatal complications. During active phases of the disorder, patients display a typical daily spiking fever, an evanescent macular rash, lymphadenopathy, hepatosplenomegaly, serositis, myalgia and arthritis.

 ${\tt Note=A\ IL6\ promoter\ polymorphism\ is\ associated\ with\ a\ lifetime\ risk\ of\ development\ of\ Kaposiana and the promoter polymorphism\ is\ associated\ with\ a\ lifetime\ risk\ of\ development\ of\ Kaposiana and the promoter\ polymorphism\ is\ associated\ with\ a\ lifetime\ risk\ of\ development\ of\ Kaposiana\ polymorphism\ po$

sarcoma in HIV-infected men.

Belongs to the IL-6 superfamily.

Sequence similarities

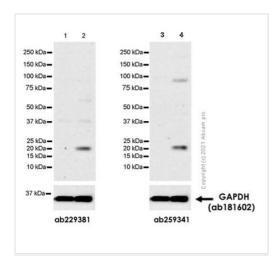
Post-translational modifications

N- and O-glycosylated.

Cellular localization

Secreted.

Images



Western blot - Anti-IL-6 antibody [EPR21710] (ab229381)

Lanes 1-2: Anti-IL-6 antibody [EPR21710] (ab229381) at 1/1000 dilution

Lanes 3-4: Anti-IL-6 antibody [EPR23819-11] (**ab259341**) at 1/1000 dilution

Lanes 1 & 3: Untreated RAW264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage) whole cell lysate

Lanes 2 & 4: RAW264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage) treated with 100ng/ml

lipopolysaccharide (LPS) for 6 hours, then added 300 ng/ml BFA for the last 3 hours, whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

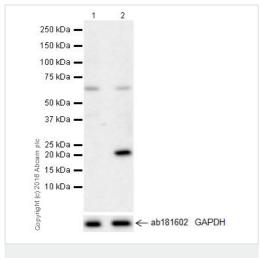
All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 24 kDa **Observed band size:** 24 kDa

Exposure time: 180 seconds

Western blot analysis with ab229381 (left) and control product <u>ab259341</u> (right).

Blocking and diluting buffer: 5% NFDM/TBST



Western blot - Anti-IL-6 antibody [EPR21710] (ab229381)

All lanes : Anti-IL-6 antibody [EPR21710] (ab229381) at 1/1000 dilution

Lane 1: Untreated RAW 264.7 (mouse macrophage cell line transformed with Abelson murine leukemia virus), whole cell lysate

Lane 2: RAW 264.7 (mouse macrophage cell line transformed with Abelson murine leukemia virus) treated with 100 ng/ml

Lipopolysaccharides (LPS) for 6 hours, then added 300 ng/ml BFA for the last 3 hours, whole cell lysate, whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$ at 1/100000 dilution

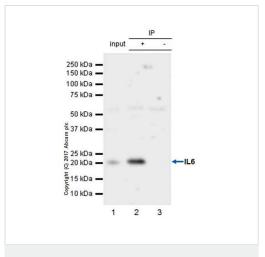
Developed using the ECL technique.

Predicted band size: 24 kDa **Observed band size:** 24 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

We detect a non-specific band at approximately 60 KDa.



Immunoprecipitation - Anti-IL-6 antibody [EPR21710] (ab229381)

IL6 was immunoprecipitated from 0.35 mg of RAW 264.7 (mouse macrophage cell line transformed with Abelson murine leukemia virus) treated with 100 ng/ml Lipopolysaccharide (LPS) for 6 hours, then added 300 ng/ml BFA the last 3 hours, whole cell lysate with ab229381 at 1/30 dilution. Western blot was performed from the immunoprecipitate using ab229381 at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/5000 dilution.

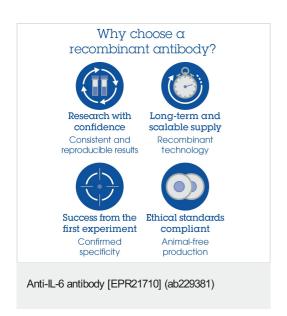
Lane 1: RAW 264.7 treated with 100 ng/ml Lipopolysaccharide (LPS) for 6 hours, then added 300 ng/ml BFA the last 3 hours, whole cell lysate 10 μ g (Input).

Lane 2: ab229381 IP in RAW 264.7 treated with 100 ng/ml Lipopolysaccharide (LPS) for 6 hours, then added 300 ng/ml BFA the last 3 hours, whole cell lysate.

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab229381 in RAW 264.7 treated with 100 ng/ml Lipopolysaccharide (LPS) for 6 hours, then added 300 ng/ml BFA the last 3 hours, whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 2 minutes.



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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