antibodies -online.com







anti-IL-27 antibody

Images



Publication



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Quantity:	100 μg	
Target:	IL-27 (IL27)	
Reactivity:	Mouse	
Host:	Rat	
Clonality:	Monoclonal	
Conjugate:	This IL-27 antibody is un-conjugated	
Application:	Western Blotting (WB)	
Product Details		
Immunogen:	This Protein A purified monoclonal antibody was produced in rats by repeated immunizations with mature length recombinant mouse p28 protein (produced in E.coli) followed by hybridoma development. Immunogen Type: Recombinant Protein	
Clone:	3H12-F10	
Isotype:	IgG2a kappa	
Specificity:	This product is an IgG fraction antibody purified from monospecific antiserum by a multi-step process which includes delipidation, salt fractionation and ion exchange chromatography followed by extensive dialysis against the buffer stated above. This antibody is specific for mouse and rat p28 protein. Cross-reactivity with IL-27 from other sources has not been determined.	
Characteristics:	Mouse IL-27/p28 Subunit, also known as Interleukin-30, is a member of the IL-12 family of	

Product Details

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	cytokines. When combined with EBI3 (Epstein-Barr virus induced gene 3), the heterodimer formed is IL-27. Mouse p28 is a proinflammatory cytokine inducing immunomodulatory effects Current research is underway to delineate specific biological functions.	
Purification:	purified	
Sterility:	Sterile filtered	
Target Details		
Target:	IL-27 (IL27)	
Alternative Name:	IL-27/p28 (IL27 Products)	
Background:	Mouse IL-27/p28 Subunit, also known as Interleukin-30, is a member of the IL-12 family of cytokines. When combined with EBI3 (Epstein-Barr virus induced gene 3), the heterodimer formed is IL-27. Mouse p28 is a proinflammatory cytokine inducing immunomodulatory effects Current research is underway to delineate specific biological functions. Synonyms: Interleukin-27 subunit alpha, IL-27 subunit alpha, IL-27-A cytokine, IL27-A, p28, IL-30	
Gene ID:	246779	
NCBI Accession:	NP_663611	
UniProt:	Q8K3I6	
Application Details		
Application Notes:	IL-27 is expressed in activated antigen presenting cells including monocytes, endothelial cells, and dendritic cells, for example mouse CD4 splenocytes. This purified antibody has been tested for use in western blotting. Specific conditions for reactivity should be optimized by the end user. Expect a band approximately 26 KDa in size corresponding to the mature mouse p28 protein, a non-glycosylated polypeptide chain consisting of amino acids, by western blotting in appropriate cell lysate or extract.	
Comment:	Gene Name: IL27	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	1.0 mg/mL	

Handling

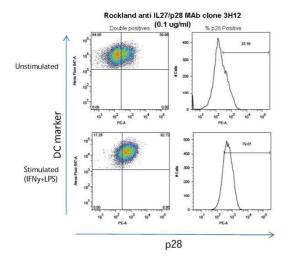
Buffer:	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2	
Preservative:	Sodium azide	
Precaution of Use:	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	4 °C/-20 °C	
Storage Comment:	Store vial at 4 °C prior to restoration. For extended storage aliquot contents and freeze at -20 °C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4 °C as an undiluted liquid. Dilute only prior to immediate use. Expiration date is one (1) year from date of opening.	
Expiry Date:	12 months	

Publications

Product cited in:

Purkey, Woolfrey, Crosby, Stich, Chick, Aoto, DellAcqua: "AKAP150 Palmitoylation Regulates Synaptic Incorporation of Ca2+-Permeable AMPA Receptors to Control LTP." in: **Cell reports**, Vol. 25, Issue 4, pp. 974-987.e4, (2018) (PubMed).

Validation report #104424 for Immunohistochemistry (IHC)



Flow Cytometry

Image 1. Mouse peritoneal macrophages were grown in culture for 24 hours, stimulated with 10ng/mL IFN? and 1ug/mL LPS for 14 hours and incubated for 4 hours with Bredfeldin A. Cells were harvested, washed, aliquoted 1x106 cells per sample, and fixed and permeabilized according to a standard protocol. Samples were stained with biotinylated primary anti-mouse p28 antibody at (0.1 - 10ug/mL primary antibody alongside negative controls of unstimulated cells and isotype controls. Cells were stained with 0.25ug/mL rat anti-mouse CD107b conjugated Alexa Fluor 647 and PHYCOERYTHRIN Conjugated secondary at 1:100 and analyzed by flow cytometry. Stimulated cells showed increase PE staining (horizontal axis) when compared with



unstimulated cells.

Western Blotting

Image 2.