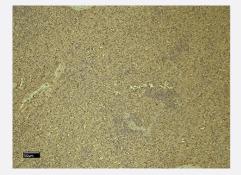
CXCR3 polyclonal antibody

Catalog # PAB7004 Size 100 ug

Applications



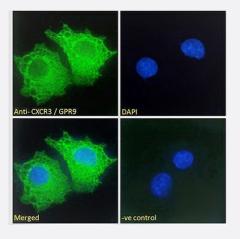
Immunohistochemistry

PAB7004 (8 μ g/mL) staining of paraffin embedded Human Spleen. Heat induced antigen retrieval with citrate buffer pH 6, HRP-staining.

Anti-CXCR3/GPR9 DAPI Merged -ve control

Immunofluorescence

PAB7004 Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10 µg/mL) followed by Alexa Fluor 488 secondary antibody (2 µg/mL), showing membrane staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 µg/mL) followed by Alexa Fluor 488 secondary antibody (2 µg/mL).

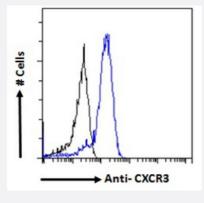


Immunofluorescence

PAB7004 Immunofluorescence analysis of paraformaldehyde fixed HepG2 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10 µg/mL) followed by Alexa Fluor 488 secondary antibody (2 µg/mL), showing membrane staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 µg/mL) followed by Alexa Fluor 488 secondary antibody (2 µg/mL).



Product Information



Flow Cytometry

PAB7004 Flow cytometric analysis of paraformaldehyde fixed HepG2 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10 ug/mL) followed by Alexa Fluor 488 secondary antibody (1 ug/mL). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.

Specification	
Product Description	Goat polyclonal antibody raised against synthetic peptide of CXCR3.
Immunogen	A synthetic peptide corresponding to human CXCR3.
Sequence	C-RRDSSWSETSEA
Host	Goat
Theoretical MW (kDa)	40.7
Reactivity	Human
Form	Liquid
Purification	Antigen affinity purification
Concentration	0.5 mg/mL
Quality Control Testing	Antibody Reactive Against Synthetic Peptide.
Recommend Usage	ELISA (1:64000) Flow Cytometry (10 μg/mL) Immunohistochemistry (6-8 μg/mL) Immunofluorescence (10 μg/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In Tris saline, pH 7.3 (0.5% BSA, 0.02% sodium azide)
Storage Instruction	Store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.



Applications

Immunohistochemistry

PAB7004 (8 µg/mL) staining of paraffin embedded Human Spleen. Heat induced antigen retrieval with citrate buffer pH 6, HRP-staining.

Immunofluorescence

PAB7004 Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10 μ g/mL) followed by Alexa Fluor 488 secondary antibody (2 μ g/mL), showing membrane staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10 μ g/mL) followed by Alexa Fluor 488 secondary antibody (2 μ g/mL).

Immunofluorescence

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- Enzyme-linked Immunoabsorbent Assay
- Flow Cytometry

PAB7004 Flow cytometric analysis of paraformaldehyde fixed HepG2 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10 ug/mL) followed by Alexa Fluor 488 secondary antibody (1 ug/mL). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.

Gene Info — CXCR3

Entrez GenelD	<u>2833</u>
Protein Accession#	<u>NP_001495.1</u>
Gene Name	CXCR3
Gene Alias	CD182, CD183, CKR-L2, CMKAR3, GPR9, IP10-R, Mig-R, MigR
Gene Description	chemokine (C-X-C motif) receptor 3
Omim ID	300574
Gene Ontology	Hyperlink

Abnova	Product Information
Gene Summary	This gene encodes a G protein-coupled receptor with selectivity for three chemokines, termed IP1 0 (interferon-g-inducible 10 kDa protein), Mig (monokine induced by interferon-g) and I-TAC (inter feron-inducible T cell a-chemoattractant). IP10, Mig and I-TAC belong to the structural subfamily of CXC chemokines, in which a single amino acid residue separates the first two of four highly cons erved Cys residues. Binding of chemokines to this protein induces cellular responses that are inv olved in leukocyte traffic, most notably integrin activation, cytoskeletal changes and chemotactic migration. Inhibition by Bordetella pertussis toxin suggests that heterotrimeric G protein of the Gi-subclass couple to this protein. Signal transduction has not been further analyzed but may include the same enzymes that were identified in the signaling cascade induced by other chemokine rece ptors. As a consequence of chemokine-induced cellular desensitization (phosphorylation-depend ent receptor internalization), cellular responses are typically rapid and short in duration. Cellular re sponsiveness is restored after dephosphorylation of intracellular receptors and subsequent recycling to the cell surface. This gene is prominently expressed in in vitro cultured effector/memory T cell Is, and in T cells present in many types of inflamed tissues. In addition, IP10, Mig and I-TAC are c ommonly produced by local cells in inflammatory cells. Therefore, this protein is a target for the de velopment of small molecular weight antagonists, which may be used in the treatment of diverse in flammatory diseases. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq
Other Designations	G protein-coupled receptor 9 IP10 receptor Mig receptor OTTHUMP00000070257 chemokine (C- X-C) receptor 3

Publication Reference

• CXCR3/CXCL10 expression in the synovium of children with juvenile idiopathic arthritis.

Martini G, Zulian F, Calabrese F, Bortoli M, Facco M, Cabrelle A, Valente M, Zacchello F, Agostini C. Arthritis Research & Therapy 2005 Jan; 7(2):R241.

Application: IHC-P, Human, Synovium

Pathway

- <u>Chemokine signaling pathway</u>
- <u>Cytokine-cytokine receptor interaction</u>

Disease

- Asthma
- Bronchiolitis
- <u>Coronary Artery Disease</u>

🗑 Abnova

- Genetic Predisposition to Disease
- Infant
- <u>Respiratory Syncytial Virus Infections</u>