

Technical Data Sheet

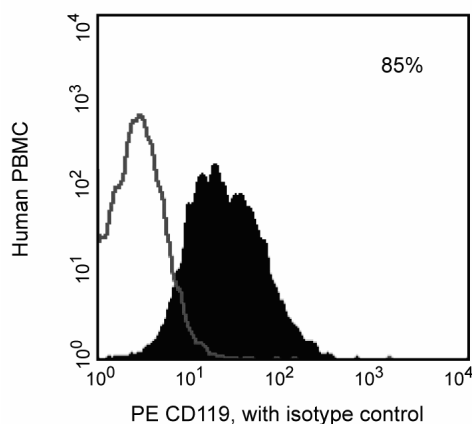
PE Mouse Anti-Human CD119

Product Information

Material Number:	558934
Alternate Name:	IFN gamma Receptor alpha chain
Size:	0.2 mg
Concentration:	0.2 mg/ml
Clone:	GIR-208
Immunogen:	Human IFN- γ R α
Isotype:	Mouse IgG1, κ
Reactivity:	QC Testing: Human
Storage Buffer:	Aqueous buffered solution containing $\leq 0.09\%$ sodium azide.

Description

The GIR-208 antibody recognizes the extracellular region of CD119 which is also known as the alpha chain subunit (80-95 kDa glycoprotein) of the human interferon- γ receptor (IFN- γ R α). The functionally active-form of the human IFN- γ receptor consists of two (or more) subunits, with IFN- γ R α responsible for IFN- γ binding and both the IFN- γ R α and β chains required for the transduction of biologic responses. The IFN- γ receptor α chain (CD119) is expressed on the surface of most human cells (except mature erythrocytes) including monocytes, macrophages, T cells, B cells, NK cells, neutrophils, fibroblasts, epithelial cells, and endothelium. Binding of 125I-labeled GIR-208 antibody to IFN- γ R α + cells is reported to be specifically inhibited in the presence of excess IFN- γ . GIR-208 does not cross react with IFN- γ as tested by ELISA. The ability of this antibody to bind to IFN- γ receptors of species other than human has not been determined. The immunogen used to generate this hybridoma was human IFN- γ R α purified from human placenta. The GIR-208 has been reported to block the binding of 125I-human IFN- γ to IFN- γ R α + cells as well as purified, soluble human IFN- γ R α . GIR-208 is a neutralizing antibody that has been shown to neutralize the anti-viral activity of IFN- γ on WISH cells in a dose-dependent fashion.



Human PBMC were isolated by Lymphoprep (Nycomed) density centrifugation. The cells were stained with R-PE conjugated GIR-208 (0.25 μ g Cat. No. 558934). Staining with the GIR-208 antibody (filled histograms) is compared to staining obtained using R-PE- conjugated immunoglobulin isotype control (MOPC-21, 0.25 μ g) (open histograms). Histograms in Figure 1 are gated on the lymphocyte population defined by its light scattering characteristics.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated with R-PE under optimum conditions, and unconjugated antibody and free PE were removed.

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

BD Biosciences

bdbiosciences.com

United States	Canada	Europe	Japan	Asia Pacific	Latin America/Caribbean
877.232.8995	888.259.0187	32.53.720.550	0120.8555.90	65.6861.0633	55.11.5185.9995

For country-specific contact information, visit bdbiosciences.com/how_to_order/

Conditions: The information disclosed herein is not to be construed as a recommendation to use the above product in violation of any patents. BD Biosciences will not be held responsible for patent infringement or other violations that may occur with the use of our products. Purchase does not include or carry any right to resell or transfer this product either as a stand-alone product or as a component of another product. Any use of this product other than the permitted use without the express written authorization of Becton Dickinson and Company is strictly prohibited.

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Not for resale.

BD, BD Logo and all other trademarks are the property of Becton, Dickinson and Company. ©2008 BD



Application Notes

Application

Flow cytometry	Routinely Tested
----------------	------------------

Neutralization Activity:

In vitro neutralization: The NA/LE format of this antibody (Cat No. 557531) is useful for bioassay.

Recommended Assay Procedure:

Immunofluorescent staining and flow cytometric analysis: The R-PE conjugated GIR-208 antibody (Cat. No. 558934) can be used for the immunofluorescent staining ($\leq 1 \mu\text{g}$ antibody/10[6] cells) and flow cytometric analysis of the levels of membrane IFN- γ R α expressed by human cell lines or human lymphoid cells. An appropriate R-PE conjugated immunoglobulin isotype control is the clone MOPC-21 (Cat. No. 554680).

Since GIR-208 is a neutralizing antibody, it competes with IFN- γ for binding to its receptor. Therefore, the use of the GIR-208 antibody for immunofluorescent staining and flow cytometric analysis in systems where the natural ligand of the receptor is present may give an underestimation of IFN- γ R α chain expression. Based on our testing results, the presence of recombinant human IFN- γ protein at levels above 200 ng/ml is sufficient to completely inhibit the binding of the GIR-208 antibody (at 0.06 μg /10[6] cells). In such cases, it is recommended that the investigator uses a non-neutralizing antibody to detect IFN- γ R α chain expression by flow cytometry, such as the clone GIR-94 (Cat. No. 558937).

WB: The GIR-208 antibody has been reported to be useful for Western blotting. Please note that this application is not routinely tested at BD Biosciences Pharmingen.

Suggested Companion Products

Catalog Number	Name	Size	Clone
554680	PE Mouse IgG1, κ Isotype Control	0.1 mg	MOPC-21
558937	PE Mouse Anti-Human CD119	0.2 mg	GIR-94
557531	Purified Mouse Anti-Human CD119	each	20/EIF-5

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.

References

- Bach E and Shreiber R. Kishimoto T, Kikutani H, von dem Borne A.E.G.K, ed. *White Cell Differentiation*.. New York: Garland Publishing, Inc; 1998:818-821. (Biology)
- Bach EA, Aguet M, Schreiber RD. The IFN gamma receptor: a paradigm for cytokine receptor signaling. *Annu Rev Immunol*. 1997; 15:563-591.(Biology)
- Gumina RJ, Freire-Moar J, DeYoung L, Webb DR, Devens BH. Transduction of the IFN-gamma signal for HLA-DR expression in the promonocytic line THP-1 involves a late-acting PKC activity. *Cell Immunol*. 1991; 138(2):265-279.(Clone-specific: Neutralization)
- Peyman JA, Hammond GL. Localization of IFN-gamma receptor in first trimester placenta to trophoblasts but lack of stimulation of HLA-DRA, -DRB, or invariant chain mRNA expression by IFN-gamma. *J Immunol*. 1992; 149(8):2675-2680.(Biology)
- Sheehan KC, Calderon J, Schreiber RD. Generation and characterization of monoclonal antibodies specific for the human IFN-gamma receptor. *J Immunol*. 1988; 140(12):4231-4237.(Immunogen: Neutralization, Western blot)
- Valente G, Ozmen L, Novelli F. Distribution of interferon-gamma receptor in human tissues. *Eur J Immunol*. 1992; 22(9):2403-2412.(Biology)