Technical Data Sheet

Purified Rat Anti-Mouse CD49e

Product Information

Material Number: 553319 Integrin a5 chain Alternate Name: 0.5 mg Size 0.5 mg/mlConcentration: 5H10-27 (MFR5) Clone:

Mouse mast cell line MC/9 Immunogen: Rat (LEW) IgG2a, ĸ Isotype: QC Testing: Mouse Reactivity:

Aqueous buffered solution containing ≤0.09% sodium azide. Storage Buffer:

Description

The 5H10-27 (MFR5) antibody reacts with the α5 chain of the integrin α5β1 fibronectin receptor (CD49e/CD29, VLA-5) on thymocytes, activated T lymphocytes, mast cells, and a variety of mouse cell lines, but not splenocytes. Soluble 5H10-27 (MFR5) antibody has been reported to inhibit VLA-5-mediated functions in vitro. In addition, immobilized mAb 5H10-27 (MFR5) has been demonstrated to costimulate the proliferative response of CD8+ T cells to plate-bound anti-CD3e antibody.

This antibody is routinely tested by flow cytometric analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at 4°C.

Application Notes

Application

Application		
Flow cytometry	Routinely Tested	
Blocking	Reported	
(Co)-stimulation	Reported	
Immunofluorescence	Reported	
Immunohistochemistry-frozen	Reported	
Immunohistochemistry-paraffin	Not Recommended	

Suggested Companion Products

Catalog Number	Name	Size	Clone	
553927	Purified Rat IgG2a, κ Isotype Control	0.5 mg	R35-95	
554016	FITC Goat Anti-Rat Ig	0.5 mg	Polyclonal	

Product Notices

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 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.
- Sodium azide is a reversible inhibitor of oxidative metabolism; therefore, antibody preparations containing this preservative agent must not be used in cell cultures nor injected into animals. Sodium azide may be removed by washing stained cells or plate-bound antibody or dialyzing soluble antibody in sodium azide-free buffer. Since endotoxin may also affect the results of functional studies, we recommend the NA/LE (No Azide/Low Endotoxin) antibody format, if available, for in vitro and in vivo use.

References

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