# Technical Data Sheet Purified Rat Anti-Mouse CD62P

Material Number:	553742
Alternate Name:	P-selectin
Size:	0.5 mg
Concentration:	0.5 mg/ml
Clone:	RB40.34
Immunogen:	P-selectin-IgG1 Fusion
Isotype:	Rat (LEW) IgG1, $\lambda$
Reactivity:	QC Testing: Mouse
Storage Buffer:	Aqueous buffered solution containing $\leq 0.09\%$ sodium azide.

## Description

The RB40.34 antibody reacts with mouse P-selectin (CD62P), a 140 kDa protein which is expressed on activated platelets, activated endothelial cells, and megakaryocytes. P-selectin mediates the adhesion of neutrophils and monocytes to activated platelets and endothelial cells, mediates leukocyte rolling, and is involved in the migration of leukocytes into inflamed tissues. CD24 and CD162 (PSGL-1) are ligands of CD62P. mAb RB40.34 can block mouse P-selectin binding to its ligands in vitro and in vivo.

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

Flow cytometry	Routinely Tested	
ELISA	Reported	
Immunoprecipitation	Reported	
Blocking	Reported	
Immunofluorescence	Reported	
Immunohistochemistry-frozen	Reported	

# **Suggested Companion Products**

Catalog Number	Name	Size	Clone
554016	FITC Goat Anti-Rat Igs	0.5 mg	Polyclonal
553993	Purified Rat IgG1 Lambda Isotype Control	0.1 mg	A110-1

# **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.
- 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.
- 4. 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<sup>™</sup> (No Azide/Low Endotoxin) antibody format, if available, for in vitro and in vivo use.

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