### **Technical Data Sheet**

# V450 Mouse Anti-Human CD41a

#### **Product Information**

Material Number: 561425

Alternate Name: ITGA2B; Integrin alpha-2b (αIIb); Platelet glycoprotein IIb (GPIIb)

 Size:
 50 tes

 Vol. per Test:
 5 µl

 Clone:
 HIP8

Isotype:Mouse IgG1,  $\kappa$ Reactivity:QC Testing: Human

Workshop: IV P38

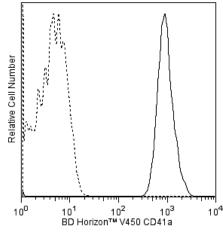
Storage Buffer: Aqueous buffered solution containing protein stabilizer, glycerol and ≤0.09%

sodium azide.

#### Description

The HIP8 monoclonal antibody specifically binds to the  $\alpha$ -chain of CD41. CD41 is also known as Integrin  $\alpha$ IIb or Platelet GPIIb. The calcium-dependent complex of CD41 and CD61 ( $\beta$ 3 integrin or GPIIIa) is normally expressed on platelets and megakaryocytes. The CD41/CD61 complex is the receptor for fibrinogen, fibronectin and von Willebrand factor, and mediates platelet adhesion and aggregation. CD41 (clone HIP8) completely inhibits ADP-, epinephrine-, and collagen-induced platelet activation, and partially inhibits ristocetin- and thrombin-induced platelet activation. This antibody is useful in the morphological and physiological studies of platelets and megakaryocytes.

The antibody is conjugated to BD Horizon<sup>TM</sup> V450, which has been developed for use in multicolor flow cytometry experiments and is available exclusively from BD Biosciences. It is excited by the Violet laser Ex max of 406 nm and has an Em Max at 450 nm. Conjugates with BD Horizon<sup>TM</sup> V450 can be used in place of Pacific Blue<sup>TM</sup> conjugates.



Flow cytometric analysis of CD41a expression on human peripheral blood platelet. Platelets were isolated from fresh whole blood and fixed with 2% formaldehyde. After washing, the fixed platelets were stained with either BD Horizon™ V450 Mouse Anti-Human CD41a antibody (Cat. No. 561425; solid line histogram) or with an BD Horizon™ V450 Mouse IgG1, κ Isotype Control (Cat. No. 560373; dashed line histogram). The fluorescence histograms were derived from events with the forward and side light-scatter characteristics of platelets. Flow cytometry was performed using a BD™ LSR II Flow Cytometer System.

#### **Preparation and Storage**

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

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

The antibody was conjugated with BD Horizon™ V450 under optimum conditions, and unreacted BD Horizon™ V450 was removed.

# **Application Notes**

# Application

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ſ	Flow cytometry	Routinely Tested		

# **Suggested Companion Products**

Catalog Number	Name	Size	Clone
560373	V450 Mouse IgG1, κ Isotype Control	0.1 mg	MOPC-21
554656	Stain Buffer (FBS)	500 ml	(none)

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### **Product Notices**

- This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1 × 10<sup>6</sup> cells in a 100-µl experimental sample (a test).
- 2. An isotype control should be used at the same concentration as the antibody of interest.
- 3. 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.
- For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- BD Horizon<sup>TM</sup> V450 has a maximum absorption of 406 nm and maximum emission of 450 nm. Before staining with this reagent, please
  confirm that your flow cytometer is capable of exciting the fluorochrome and discriminating the resulting fluorescence.
- 7. Pacific Blue<sup>TM</sup> is a trademark of Molecular Probes, Inc., Eugene, OR.

#### References

Knapp W, Dorken B, Rieber EP, et al, ed. Leucocyte Typing IV. New York: Oxford University Press; 1989:1-1208. (Clone-specific)
Schlossman SF, Boumsell L, Gilks W, et al, ed. Leukocyte Typing V: White Cell Differentiation Antigens. New York: Oxford University Press; 1995. (Biology)

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