Technical Data Sheet

BV711 Mouse Anti-Human CD14

Product Information

Material Number: 56337.

Alternate Name: LPS receptor; LPS-R; Myeloid cell-specific leucine-rich glycoprotein

 Size:
 25 tes

 Vol. per Test:
 5 μl

Clone: $M\phi P9$ (also known as $M\phi P-9$)

Immunogen:Human MonocytesIsotype:Mouse (BALB/c) IgG2b, κ Reactivity:QC Testing: Human

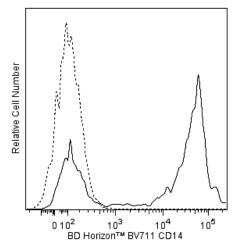
Workshop: I M35

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

Description

The $M\phi P9$ monoclonal antibody specifically binds to human CD14, a 53-55 kDa glycosylphosphatidylinositol (GPI)-anchored and single chain glycoprotein expressed at high levels on monocytes. Additionally, this CD14-specific antibody reacts with interfollicular macrophages, reticular dendritic cells and some Langerhans cells. CD14 has been identified as a high affinity cell-surface receptor for complexes of lipopolysaccharide (LPS) and serum LPS-binding protein, LPB. The purified antibody is suitable for staining acetone-fixed, frozen tissue sections.

The antibody was conjugated to BD HorizonTM BV711 which is part of the BD HorizonTM Brilliant VioletTM family of dyes. This dye is a tandem fluorochrome of BD HorizonTM BV421 with an Ex Max of 405-nm and an acceptor dye with an Em Max at 711-nm. BD HorizonTM BV711 can be excited by the violet laser and detected in a filter used to detect CyTM5.5 / Alexa Fluor® 700-like dyes (eg, 712/20-nm filter). Due to the excitation and emission characteristics of the acceptor dye, there may be moderate spillover into the Alexa Fluor® 700 and PerCP-CyTM5.5 detectors. However, the spillover can be corrected through compensation as with any other dye combination.



Flow cytometric analysis of CD14 expression on human peripheral blood monocytes. Human whole blood was stained with either BD Horizon™ BV711 Mouse Anti-Human CD14 antibody (Cat. No. 563372/563373; solid line histogram) or BD Horizon™ BV711 Mouse IgG2b, k Isotype Control (Cat. No. 563125; dashed line histogram). The erythrocytes were lysed with BD Pharm Lyse™ Lysing Buffer (Cat. No. 555899). The fluorescence histograms were derived from events with the forward and side light-scatter characteristics of viable monocytes. 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 HorizonTM BV711 under optimum conditions, and unconjugated antibody and free BD HorizonTM BV711 were removed.

Application Notes

Application

Flow cytometry Routinely Tested

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Suggested Companion Products

Catalog Number	Name	Size	Clone
554656	Stain Buffer (FBS)	500 ml	(none)
563125	BV711 Mouse IgG2b, κ Isotype Control	50 μg	27-35
563372	BV711 Mouse Anti-Human CD14	100 tests	ΜφΡ9
555899	Lysing Buffer	100 ml	(none)
349202	FACS Lysing Solution		(none)

Product Notices

- This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use $1 \times 10^{\circ}6$ cells in a 100- μ l experimental
- 2. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- An isotype control should be used at the same concentration as the antibody of interest.
- 4. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- Alexa Fluor® is a registered trademark of Molecular Probes, Inc., Eugene, OR. 5
- 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.
- 7. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- 8 Cy is a trademark of Amersham Biosciences Limited.
- Brilliant VioletTM 711 is a trademark of Sirigen. 9.

References

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Dimitriu-Bona A, Burmester GR, Waters SJ, Winchester RJ. Human mononuclear phagocyte differentiation antigens. I. Patterns of antigenic expression on the surface of human monocytes and macrophages defined by monoclonal antibodies. J Immunol. 1983; 130(1):145-152. (Immunogen: Flow cytometry, Immunofluorescence)

Goyert SM, Ferrero E. Biochemical analysis of myeloid antigens and cDNA expression of gp55 (CD14). In: McMichael AJ, ed. Leucocyte Typing III: White Cell Differentiation Antigens. New York, NY: Oxford University Press;; 1987:613-619. (Biology)

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