

Human CD83 PE-conjugated Antibody

Monoclonal Mouse IgG₁ Clone # HB15e

Catalog Number: FAB1774P

100 TESTS

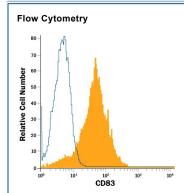
DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human CD83 in flow cytometry.		
Source	Monoclonal Mouse IgG ₁ Clone # HB15e		
Purification	Protein A or G purified from hybridoma culture supernatant		
Immunogen	COS-7 African green monkey SV40 transformed kidney fibroblast-like cells transfected with human CD83		
Conjugate	Phycoerythrin Excitation Wavelength: 488 nm Emission Wavelength: 565-605 nm		
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.		
	*Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.		

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Flow Cytometry	10 μL/10 ⁶ cells	See Below

DATA



Detection of CD83 in CD14+ Monocytederived Dendritic Cells by Flow Cytometry. CD14+ monocyte-derived dendritic cells were stained with Mouse Anti-Human CD83 PE-conjugated Monoclonal Antibody (Catalog # FAB1774P, filled histogram) or isotype control antibody (Catalog # IC002P, open histogram). View our protocol for Staining Membraneassociated Proteins.

PREPARATION AND STORAGE

Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage

Protect from light. Do not freeze.

• 12 months from date of receipt, 2 to 8 °C as supplied.



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BACKGROUND

Human CD83 is a 40-50 kDa member of the Siglec (or Sialic-acid-binding Immunoglobulin-like Lectin) family of transmembrane proteins (1, 2, 3). CD83 is synthesized as a type I transmembrane glycoprotein that contains a 125 amino acid (aa) extracellular region, a 22 aa transmembrane segment, and 39 aa cytoplasmic domain. It contains one V type Ig-like domain in the extracellular region with no inhibitory cytoplasmic motif(s). Although *in vitro* studies suggest CD83 may form membrane-bound covalent homodimers, *in vivo* this does not appear to be the case (1, 4). In the extracellular region, mouse and human CD83 are 66% aa identical (1, 2, 4, 5). Relative to human, mouse CD83 is 11 aa shorter in its extracellular domain and is expressed as a 30-35 kDa protein (1, 4, 5). Human CD83 is active in the mouse system (4). One alternate splice form has been reported. This leads to a small monomeric soluble form of 74 aa that includes aa 20-52 and aa 164-205 (6, 7). In human, proteolytic cleavage and solubilization of CD83 has also been suggested, and this could lead to dimeric circulating CD83 (4, 6). CD83 is a primary marker for dendritic cells (3, 6, 8). It is also found on B cells (6, 9), neutrophils (10), monocytes and macrophages (11). Except for dendritic cells, CD83 expression is often transient. CD83 binds to sialic acids on target cells (12). Membrane CD83 appears to promote T cell proliferation, particularly of CD8* cytotoxic T cells (13, 14). Soluble CD83, however, appears to be immunosuppressive and blocks T cell activation (15, 16). On monocytes, CD83 is suggested to drive monocytes into a fibrocyte phenotype (13). A lack of membrane-expressed CD83 leads to an unusual IL-4/IL-10 producing CD4* T cell phenotype (17).

References:

- 1. Zhou, L-J. et al. (1992) J. Immunol. 149:735.
- 2. Kozlow, E.J. et al. (1993) Blood 81:454.
- 3. Fujimoto, Y and T.F. Tedder (2006) J. Med. Dent. Sci. 53:85.
- Lechmann, M. et al. (2005) Biochem. Biophys. Res. Commun. 329:132.
- 5. Berchtold, S. et. al. (1999) FEBS Lett. 461:211.
- 6. Hock, B.D. et al. (2001) Int. Immunol. 13:959.
- 7. Dudziak, D. et al. (2005) J. Immunol. 174:6672.
- 8. Velten, F.W. et al. (2007) Mol. Immunol. 44:1544.
- 9. Cramer, S.O. et al. (2000) Int. Immunol. 12:1347.
- 10. Yamashiro, S. et al. (2000) Blood 96:3958.
- 11. Cao, W. et al. (2005) Biochem. J. 385:85.
- 12. Scholler, N. et al. (2001) J. Immunol. 166:3865.
- 13. Scholler, N. et al. (2002) J. Immunol. 168:2599.
- 14. Hirano, N. et al. (2006) Blood 107:1528.
- 15. Kotzor, N. et al. (2004) Immunobiology 209:129.
- 16. Zinser, E. et al. (2006) Immunobiology 211:449.
- 17. Garcia-Martinez, L.F. et al. (2004) J. Immunol. 173:2995.

