

## DESCRIPTION

|                           |   |
|---------------------------|---|
| <b>Species Reactivity</b> | Human   |
| <b>Specificity</b>        | Detects human MFG-E8 in direct ELISAs and Western blots.  |
| <b>Source</b>             | Monoclonal Mouse IgG <sub>2A</sub> Clone # 278918   |
| <b>Purification</b>       | Protein A or G purified from hybridoma culture supernatant  |
| <b>Immunogen</b>          | Mouse myeloma cell line NS0-derived recombinant human MFG-E8<br>Leu24-Cys387<br>Accession # Q08431      |
| <b>Conjugate</b>          | Alexa Fluor 488<br>Excitation Wavelength: 488 nm<br>Emission Wavelength: 515-545 nm                     |
| <b>Formulation</b>        | 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.

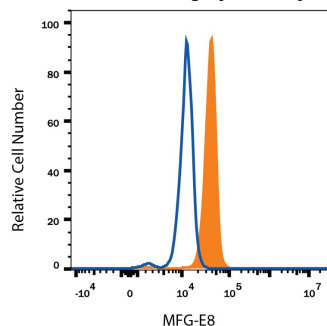
## APPLICATIONS

**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.

**Intracellular Staining by Flow Cytometry** Titration recommended for optimal concentration with starting range of 0.1-1 µg/1 million cells. Sample used for this experiment was Human immature dendritic cells and TF-1 cells fixed with FC012 and permeabilized with FoxP3 Perm.

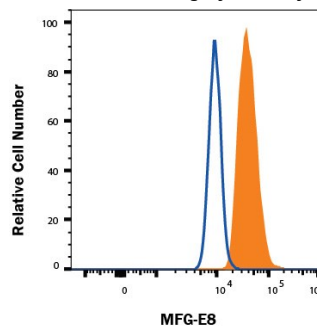
## DATA

### Intracellular Staining by Flow Cytometry



**Detection of MFG-E8 in Human Dendritic Cells by Flow Cytometry** Human immature dendritic cells were stained with Mouse Anti-Human MFG-E8 Alexa Fluor® 488-conjugated Monoclonal Antibody (Catalog # IC27671G, filled histogram) or isotype control antibody (Catalog # IC003G, open histogram). To facilitate intracellular staining, cells were fixed with Flow Cytometry Fixation Buffer (Catalog # FC004) and permeabilized with Flow Cytometry Permeabilization/Wash Buffer I (Catalog # FC005). View our protocol for [Staining Intracellular Molecules](#).

### Intracellular Staining by Flow Cytometry



**Detection of MFG-E8 in TF-1 cells by Flow Cytometry.** TF-1 cells were stained with Mouse Anti-Human MFG-E8 Alexa Fluor® 488-conjugated Monoclonal Antibody (Catalog # IC27671G, filled histogram) or isotype control antibody (Catalog # IC003G, open histogram). To facilitate intracellular staining, cells were fixed with FC012 and permeabilized with FoxP3 Perm. View our protocol for [Staining Intracellular Molecules](#).

## 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.

## BACKGROUND

Milk Fat Globulin Protein E8 (MFG-E8), also known as Lactadherin, MP47, breast epithelial antigen BA46, and SED1, is a 66-75 kDa pleiotropic secreted glycoprotein that promotes mammary gland morphogenesis, angiogenesis, and tumor progression. MFG-E8 also plays an important role in tissue homeostasis and the prevention of inflammation (1). Human MFG-E8 contains one N-terminal EGF-like domain and two C-terminal F5/8-type discoidin-like domains (2). It shares 63% and 61% aa sequence identity with comparable regions of mouse and rat MFG-E8, respectively. Shorter isoforms of human MFG-E8 may have N-terminal deletions (beginning near the end of the first discoidin-like domain), internal deletions (lacking either the EGF-like domain or the central region of the second discoidin-like domain), or C-terminal deletions (truncated within the second discoidin-like domain) (3). A 50 aa internal proteolytic fragment of human MFG-E8 (known as Medin) is a major component of aortic medial amyloid deposits (4). MFG-E8 is released into the milk in complex with lipid-containing milk fat globules. It is also found in multiple other cell types including endothelial cells and smooth muscle cells of the vasculature, immature dendritic cells, at the acrosomal cap of testicular and epididymal sperm, and in epithelial cells of the endometrium (1). MFG-E8 binds to the Integrins  $\alpha V\beta 3$  and  $\alpha V\beta 5$  and potentiates the angiogenic action of VEGF through VEGF R2 (5, 6). It reduces inflammation and tissue damage in a variety of settings. MFG-E8 functions as a bridge between phosphatidylserine on apoptotic cells and Integrin  $\alpha V\beta 3$  on phagocytes, leading to the clearance of apoptotic debris (7). It mediates the engulfment of apoptotic bodies in atherosclerotic plaques and prion-infected brain (8, 9) and of apoptotic B cells during germinal center reactions (10, 11). MFG-E8 also promotes the removal of excess Collagen in fibrotic lungs and the regeneration of damaged intestinal epithelia (12, 13). Its tissue-protective role impairs anti-tumor immunity and chemotherapy-induced apoptosis (14). MFG-E8 in the breastmilk blocks rotavirus infection in nursing babies (15).

## References:

1. Raymond, A. *et al.* (2009) J. Cell. Biochem. **106**:957.
2. Couto, J.R. *et al.* (1996) DNA Cell Biol. **15**:281.
3. Yamaguchi, H. *et al.* (2010) Eur. J. Immunol. **40**:1778.
4. Haggqvist, B. *et al.* (1999) Proc. Natl. Acad. Sci. USA **96**:8669.
5. Silvestre, J.-S. *et al.* (2005) Nat. Med. **11**:499.
6. Borges, E. *et al.* (2000) J. Biol. Chem. **275**:39867.
7. Hanayama, R. *et al.* (2002) Nature **417**:182.
8. Ait-Oufella, H. *et al.* (2007) Circulation **115**:2168.
9. Kranich, J. *et al.* (2010) J. Exp. Med. **207**:2271.
10. Hanayama, R. *et al.* (2004) Science **304**:1147.
11. Kranich, J. *et al.* (2010) J. Exp. Med. **205**:1293.
12. Atabai, K. *et al.* (2009) J. Clin. Invest. **119**:3713.
13. Bu, H.-F. *et al.* (2007) J. Clin. Invest. **117**:3673.
14. Jinushi, M. *et al.* (2009) J. Exp. Med. **206**:1317.
15. Kvistgaard, A.S. *et al.* (2004) J. Dairy Sci. **87**:4088.

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