

Technical Data Sheet

Purified Mouse Anti-EBP50

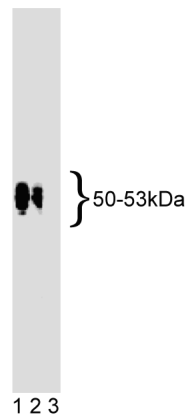
Product Information

Material Number:	611161
Size:	150 µg
Concentration:	250 µg/ml
Clone:	6/EBP50
Immunogen:	Human EBP50 aa. 128-249
Isotype:	Mouse IgG1
Reactivity:	QC Testing: Human
Target MW:	50-53 kDa
Storage Buffer:	Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium azide.

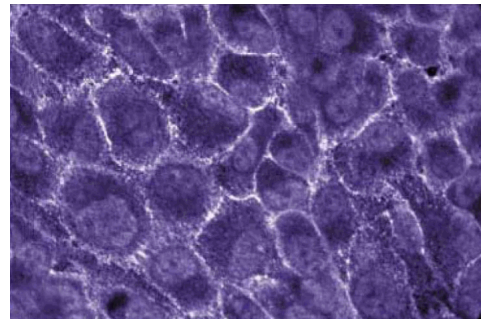
Description

Integral plasma membrane proteins are stabilized by linkages to the cortical actin cytoskeleton, which structurally supports the membrane and contributes to processes such as endocytosis, exocytosis, and transmembrane signaling. The ERM (ezrin-radixin-moesin) family of proteins provides these structural linkages. The ERM proteins contain a 300-residue N-terminal domain, a 170-residue α -helical region, and a C-terminal 100-residue domain that contains F-actin binding sites. The N-terminal domain interacts with the cytoplasmic domain of CD44, the regulatory subunit of PKA, and the PDZ domain containing ERM-binding phosphoprotein (EBP-50). In polarized epithelial cells, EBP-50 links ezrin and the cytoplasmic regions of transmembrane proteins such as the cystic fibrosis transmembrane conductance regulator (CFTR) and the β 2-adrenergic receptor. EBP50 contains two PDZ domains followed by a C-terminal tail. It colocalizes with ezrin in apical microvilli and is also thought to interact with the Na⁺-H⁺ exchanger NHE3 to confer cAMP-mediated inhibition of Na⁺-H⁺ exchange. Thus, EBP-50 mediates membrane attachment to the cytoskeleton and may also function in the regulation of ion exchange.

This antibody is routinely tested in western blot analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.



Western blot analysis of EBP50 on human endothelial cell lysate. Lane 1: 1:250, lane 2: 1:500, lane 3: 1:1000 dilution of anti-EBP50.



Immunofluorescent staining of Human Endothelial cells with anti-EBP50.

Preparation and Storage

Store undiluted at -20° C.

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

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Application Notes

Application

Western blot	Routinely Tested
Immunofluorescence	Tested During Development

Suggested Companion Products

Catalog Number	Name	Size	Clone
611450	Human Endothelial Cell Lysate	500 µg	(none)
554002	HRP Goat Anti-Mouse Ig	1.0 ml	(none)
554001	FITC Goat Anti-Mouse Ig	0.5 mg	Polyclonal

Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to www.bdbiosciences.com/pharming/en/protocols for technical protocols.
3. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
4. 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.

References

Bretscher A, Reczek D, Berryman M. Ezrin: a protein requiring conformational activation to link microfilaments to the plasma membrane in the assembly of cell surface structures. *J Cell Sci.* 1997; 110(Pt 24):3011-3018.(Biology)

Glynn PA, Darling KE, Picot J, Evans TJ. Epithelial inducible nitric-oxide synthase is an apical EBP50-binding protein that directs vectorial nitric oxide output. *J Biol Chem.* 2002; 277(36):33132-33138.(Clone-specific: Immunofluorescence, Western blot)

Naren AP, Cobb B, Li C. A macromolecular complex of beta 2 adrenergic receptor, CFTR, and ezrin/radixin/moesin-binding phosphoprotein 50 is regulated by PKA. *Proc Natl Acad Sci U S A.* 2003; 100(1):342-346.(Clone-specific: ELISA, Western blot)

Reczek D, Berryman M, Bretscher A. Identification of EBP50: A PDZ-containing phosphoprotein that associates with members of the ezrin-radixin-moesin family. *J Cell Biol.* 1997; 139(1):169-179.(Biology)

Reczek D, Bretscher A. The carboxyl-terminal region of EBP50 binds to a site in the amino-terminal domain of ezrin that is masked in the dormant molecule. *J Biol Chem.* 1998; 273(29):18452-18458.(Biology)