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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 46, 48	Source/Isotype: Rabbit	UniProt ID: #Q9Y5Q3	Entrez-Gene Id: 9935	
Product Usage Information	2	Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity/Ser	nsitivity	MAFB Antibody recognizes endogenous levels of total MAFB protein. Based on sequence similarity, this antibody is not predicted to cross-react with MAFA.					
Source / Purifi	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding of human Pro188 protein. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		MAFB belongs to the musculoaponeurotic fibrosarcoma (MAF) family of basic leucine-zipper transcription factors (1). In mouse embryo, MAFB expression is first detected at E10.5 (2,3). Early in development, MAFB drives differentiation of both glucagon-producing α -cells and insulin-producing β -cells in the pancreas, but later plays a more decisive role in the maturation and maintenance of functional α -cells (4,5). Consistent with MAFB playing a critical role in mature α -cells, MAFB is enriched in α -cells within two weeks of birth in the pancreas (6). Glucagon and insulin secretion is tightly regulated, and imbalances in these hormones contribute to metabolic conditions. Therefore, understanding the role of MAFB in α -cell development, maintenance, and physiological function may contribute to developing deeper insights into how these cells contribute to metabolic diseases like diabetes. MAFB also regulates monocyte differentiation, indicating MAFB functions beyond the pancreas (7).					
Background R	eferences	1. Hang, Y. and Stein, R. (2011) <i>Trends Endocrinol Metab</i> 22, 364-73. 2. Nishimura, W. et al. (2006) <i>Dev Biol</i> 293, 526-39. 3. Artner, I. et al. (2006) <i>Diabetes</i> 55, 297-304. 4. Katoh, M.C. et al. (2018) <i>Mol Cell Biol</i> 38, e00504-17. 5. Artner, I. et al. (2007) <i>Proc Natl Acad Sci U S A</i> 104, 3853-8. 6. Artner, I. et al. (2010) <i>Diabetes</i> 59, 2530-9. 7. Wu, X. et al. (2016) <i>J Exp Med</i> 213, 2553-65.					
Species Reacti	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Western Blot E	Buffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications K	ey	W: Western Blotting IP: Immunoprecipitation					
Cross-Reactivi	ty Key	H: Human					
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