

SMARCD3/BAF60C (D6F1S) Rabbit mAb



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Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 60	Source/Isotype: Rabbit IgG	UniProt ID: #Q6STE5	Entrez-Gene Id: 6604
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		SMARCD3/BAF60A (D6F1S) Rabbit mAb recognizes endogenous levels of total SMARCD3/BAF60C protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human SMARCD3/BAF60C protein.				
Background		ATP-dependent chromatin remodeling complexes play an essential role in the regulation of nuclear processes such as transcription and DNA replication and repair (1,2). The SWI/SNF chromatin remodeling complex consists of more than 10 subunits and contains a single molecule of either BRM or BRG1 as the ATPase catalytic subunit. The activity of the ATPase subunit disrupts histone-DNA contacts and changes the accessibility of crucial regulatory elements to the chromatin. The additional core and accessory subunits play a scaffolding role to maintain stability and provide surfaces for interaction with various transcription factors and chromatin (2-5). The interactions between SWI/SNF subunits and transcription factors, such as nuclear receptors, p53, Rb, BRCA1, and MyoD, facilitate recruitment of the complex to target genes for regulation of gene activation, cell growth, cell cycle, and differentiation processes (1,6-9). SMARCD3/BAF60C is essential for the development of the skeletal and cardiac muscle (10,11). Knockdown of SMARCD3/BAF60C using RNAi in mouse embryos leads to defects in heart development as well as abnormalities in the development of the cardiac and skeletal muscle, mimicking the cardiac defects observed in congenital heart disease (12). In addition, SMARCD3/BAF60C is part of the lipoBAF complex that remodels the chromaitn structure to activate lipogenic genes, promoting lipogenesis in response to feeding and insulin (13).				
Background References		 Ho, L. and Crabtree, G.R. (2010) <i>Nature</i> 463, 474-84. Becker, P.B. and Hörz, W. (2002) <i>Annu Rev Biochem</i> 71, 247-73. Eberharter, A. and Becker, P.B. (2004) <i>J Cell Sci</i> 117, 3707-11. Bowman, G.D. (2010) <i>Curr Opin Struct Biol</i> 20, 73-81. Gangaraju, V.K. and Bartholomew, B. (2007) <i>Mutat Res</i> 618, 3-17. Lessard, J.A. and Crabtree, G.R. (2010) <i>Annu Rev Cell Dev Biol</i> 26, 503-32. Morettini, S. et al. (2008) <i>Front Biosci</i> 13, 5522-32. Wolf, I.M. et al. (2008) <i>J Cell Biochem</i> 104, 1580-6. Simone, C. (2006) <i>J Cell Physiol</i> 207, 309-14. Puri, P.L. and Mercola, M. (2012) <i>Genes Dev</i> 26, 2673-83. Vallaster, M. et al. (2012) <i>Acta Biochim Biophys Sin (Shanghai)</i> 44, 92-102. Lickert, H. et al. (2004) <i>Nature</i> 432, 107-12. Wang, Y. et al. (2013) <i>Mol Cell</i> 49, 283-97. 				

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot 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 Key W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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