

## 6889

## **BRM Antibody**



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

Applications: W	<b>Reactivity:</b> H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 200	<b>Source/Isotype:</b> Rabbit	UniProt ID: #P51531	Entrez-Gene Id: 6595
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		BRM Antibody recognizes endogenous levels of total BRM protein. This antibody does not cross-react with BRG1 protein.				
Species predicted to react based on 100% sequence homology		Dog				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly264 of human BRM protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		ATP-dependent chromatin remodeling complexes play an essential role in the regulation of various nuclear processes, such as gene expression, DNA replication, and repair (1,2). The SWI/SNF chromatin remodeling complex consists of more than 10 subunits with a single molecule of the ATPase catalytic subunit BRM or BRG1, but not both. The activities of these two subunits drive the disruption of histone-DNA contacts that lead to changes in accessibility of crucial regulatory elements within chromatin (2-5). The BRM/BRG1 containing SWI/SNF complexes are recruited to target promoters by transcription factors, such as nuclear receptors, p53, RB, and BRCA1 to regulate gene activation, cell growth, the cell cycle, and differentiation processes (1,6-9). BRM and BRG1 are also considered to be tumor suppressors and their expression levels are severely reduced in several cancer cell lines (10-13).				
Background References		1. Ho, L. and Crabtree, G.R. (2010) <i>Nature</i> 463, 474-84. 2. Becker, P.B. and Hörz, W. (2002) <i>Annu Rev Biochem</i> 71, 247-73. 3. Eberharter, A. and Becker, P.B. (2004) <i>J Cell Sci</i> 117, 3707-11. 4. Bowman, G.D. (2010) <i>Curr Opin Struct Biol</i> 20, 73-81. 5. Gangaraju, V.K. and Bartholomew, B. (2007) <i>Mutat Res</i> 618, 3-17. 6. Lessard, J.A. and Crabtree, G.R. (2010) <i>Annu Rev Cell Dev Biol</i> 26, 503-32. 7. Morettini, S. et al. (2008) <i>Front Biosci</i> 13, 5522-32. 8. Wolf, I.M. et al. (2008) <i>J Cell Biochem</i> 104, 1580-6. 9. Simone, C. (2006) <i>J Cell Physiol</i> 207, 309-14. 10. Yamamichi, N. et al. (2005) <i>Oncogene</i> 24, 5471-81. 11. Reisman, D.N. et al. (2002) <i>Oncogene</i> 21, 1196-207. 12. Shen, H. et al. (2008) <i>Cancer Res</i> 68, 10154-62. 13. Weissman, B. and Knudsen, K.E. (2009) <i>Cancer Res</i> 69, 8223-30.				
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

**Cross-Reactivity Key** 

H: Human Mk: Monkey

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