

## SMARCAL1 (D3P5I) Rabbit mAb



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

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 105	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q9NZC9	Entrez-Gene Id: 50485
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50	
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		SMARCAL1 (D3P5I) Rabbit mAb recognizes endogenous levels of total SMARCAL1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala933 of human SMARCAL1 protein.				
Background		SMARCAL1 was first identified as a ubiquitously expressed member of the SNF2 family with homology to the <i>E. coli</i> protein HepA (1). Mutations in the gene encoding SMARCAL1 were subsequently shown to be the cause of Schimke immuno-osseous dysplasia (SIOD), an autosomal recessive disorder characterized by phenotypes in multiple systems, including spondyloepiphyseal dysplasia, renal dysfunction, immunodeficiency, and impaired neurological function (2). Researchers have also associated SMARCAL1 deficiency with predisposition to non-Hodgkin's lymphoma (3). The array of phenotypes associated with SMARCAL1 is likely due to its role as an annealing helicase in the DNA damage response. During DNA replication stress, SMARCAL1 is phosphorylated by DNA repair kinases (ATM, ATR, DNA-PK) (4). SMARCAL1 deficiency sensitizes cells to replication stress agents, and appears to increase the frequency of replication fork breakdown (4,5). SMARCAL1 is also required for efficient DNA double strand break repair via the nonhomologous end joining (NHEJ) DNA repair pathway (6). Researchers have suggested that inhibitors targeting SMARCAL1 may be effective in sensitizing cancer cells to chemotherapeutic agents (reviewed in 7).				
Background References		<ol> <li>Coleman, M.A. et al. (2000) Genomics 65, 274-82.</li> <li>Boerkoel, C.F. et al. (2002) Nat Genet 30, 215-20.</li> <li>Baradaran-Heravi, A. et al. (2012) Am J Med Genet A 158A, 2204-13.</li> <li>Bansbach, C.E. et al. (2009) Genes Dev 23, 2405-14.</li> <li>Silverberg, M.J. et al. (2009) AIDS 23, 2337-45.</li> <li>Keka, I.S. et al. (2015) Nucleic Acids Res 43, 6359-72.</li> <li>Zhang, L. et al. (2012) Biochem Biophys Res Commun 427, 232-5.</li> </ol>				

**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.

W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key

**Applications Key** 

**H:** Human

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