at	SAV1 (D6M6X) Rabbit mAb		Cell Signaling		
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## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: WB, IP, IF-IC	Reactivity: H M	Sensitivity: Endogenous	<b>MW (kDa):</b> 45	Source/Isotype: Rabbit IgG	UniProt ID: #Q9H4B6	Entrez-Gene Id: 60485		
Product Usage		plication				Dilution		
Information	We	stern Blotting				1:1000		
	Imn	nunoprecipitation				1:50		
		nunofluorescence (Ir	mmunocytochen	nistry)		1:400		
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 / Sensiti	vity SAV	SAV1 (D6M6X) Rabbit mAb recognizes endogenous levels of total SAV1 protein.						
Source / Purificatio		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human SAV1 protein.						
Background Background Refere	pair prote activ whic regu cycle epith num sign ences 1. Va 2. OI 3. Ge 4. Ze 5. Ta 6. Le	<ul> <li>Salvador homolog (SAV1), originally named WW45, was first identified as a 45 kDa protein containing a pair of WW domains and a coiled-coil region (1). SAV1 was subsequently shown to function as a scaffold protein, in a protein complex that includes the kinases MST2 and LATS1, and the transcriptional coactivator YAP (2). This protein complex comprises the core components of the Hippo signaling pathway, which regulates important cellular functions, including contact inhibition and apoptosis, that function to regulate tissue growth and organ size (3,4). A genetic screen in <i>Drosophila</i> identified a role for SAV1 in cell cycle regulation and apoptosis (5), while embryonic mice lacking Sav1 displayed hyperplastic growth and epithelial differentiation effects (6). These findings, together with the observation that SAV1 is mutated a number of human cancer cell lines, suggest that SAV1 functions as a tumor suppressor in the Hippo signaling pathway (5, 7).</li> <li>1. Valverde, P. (2000) <i>Biochem Biophys Res Commun</i> 276, 990-8.</li> <li>2. Oka, T. et al. (2008) <i>J Biol Chem</i> 283, 27534-46.</li> <li>3. Guo, C. et al. (2007) <i>Curr Biol</i> 17, 700-5.</li> <li>4. Zeng, Q. and Hong, W. (2008) <i>Cancer Cell</i> 13, 188-92.</li> <li>5. Tapon, N. et al. (2002) <i>Cell</i> 110, 467-78.</li> <li>6. Lee, J.H. et al. (2008) <i>EMBO J</i> 27, 1231-42.</li> </ul>						
	7. D	onninger, H. et al. (2		11 200, 10403-91.				
Species Reactivity	Spec	Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Western Blot Buffe		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.						
<b>Applications Key</b>	WB:	WB: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)						
Cross-Reactivity Ke	<b>X:</b> Xe	<ul> <li>H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster</li> <li>X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse</li> <li>GP: Guinea Pig Rab: rabbit All: all species expected</li> </ul>						
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## SAV1 (D6M6X) Rabbit mAb (#13301) Datasheet Without Images Cell Signaling Technology

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