e at -20C	TRIB2 (D8P2X) Rabbit mAb		Cell Signaling		
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For Research Use Only	Not for Use	in Diagnostic I	Procedures
FOI INESCAICH USE OIN	. NOLIDI 030	in Diagnostic i	Toccuures.

Applications: WB, IP	Reactivity: H M	Sensitivity: Endogenous	<b>MW (kDa):</b> 42	Source/Isotype: Rabbit IgG	<b>UniProt ID:</b> #Q92519	Entrez-Gene Id: 28951	
Product Usage Information	١	Application Western Blotting mmunoprecipitation			<b>Dilution</b> 1:1000 1:200		
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		TRIB2 (D8P2X) Rabbit mAb recognizes endogenous levels of total TRIB2 protein. This antibody does not cross-react with other TRIBBLES family proteins.					
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant protein corresponding to full- length human TRIB2 protein.					
Background	pi a R re cc (T e) le (5 u u w (7	resence of a variant pr C-terminal COP1 site Drosophila genetic so esearch studies in Dro egulating turnover of th ontains a single <i>tribble</i> TRIB1-3), which exhibi xample, TRIB1 and TF oucine zipper transcript 5,6). TRIB2 is overexpin ndergoing proliferation hile retroviral-mediate	otein kinase mot that binds ubiqui creens for genes <i>bsophila</i> suggest e cell cycle prote s gene, the geno t both distinct an RIB2, but not TRI tion factor C/EBF ressed in a subse arrest (7), and p d overexpression ectively suggest t	mily of serine-threonine if (lacking a canonical A tin ligase. The <i>tribbles</i> g that regulate cell divisio ed that TRIBBLES funct sin String/cdc25. In cont ones of mice and humar d overlapping patterns o B3, were reported to pro Pa, a function that appea et of human AML patient positively regulated by th of TRIB2 in mice was s hat TRIB2 functions as a	TP binding site), a ME gene was first identified n, gastrulation, and oo ions to coordinate cell rast to the <i>Drosophila</i> as encode three known of expression and funct prote degradation of the urs to be conserved fro a samples, downregula e NOTCH signaling pa- shown to induce transp	K1 binding site, and d and characterized ogenesis (1-3). division by genome, which n TRIBBLES proteins tions (4). For he basic region- m flies to humans ted in leukemic cells athway in T cells (8), olantable leukemia	
Background Refer	2. 3. 4. 5. 6. 7. 8.	Grosshans, J. and W Seher, T.C. and Lepti Mata, J. et al. (2000) Dobens, L.L. and Bou Dedhia, P.H. et al. (200) Keeshan, K. et al. (2000) Keeshan, K. et al. (201 Hannon, M.M. et al. (201	n, M. (2000) <i>Cur</i> . <i>Cell</i> 101, 511-22 Jyain, S. (2012) <i>I</i> 010) <i>Blood</i> 116, 2 Mol Cell 6, 23-3 006) <i>Cancer Cell</i> 2012) <i>Br J Haerr</i>	r Biol 10, 623-9. 2. Dev Dyn 241, 1239-48. 1321-8. 0. 10, 401-11. natol 158, 626-34.			
Species Reactivity		ecies reactivity is dete	ermined by testing	g in at least one approve	ed application (e.g., we	estern blot).	
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications Key		B: Western Blotting IF	: Immunoprecip	itation			
Cross-Reactivity K	X:	<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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