

RALY Antibody



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

Product Usage Information Storage		Application				
Storage		Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		RALY Antibody recognizes endogenous levels of total RALY protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro163 of human RALY protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		RALY is a member of the large family of heterogeneous nuclear ribonucleoproteins (hnRNPs). RALY was initially discovered in lethal yellow mice, a condition where heterozygotes display an all yellow coat, obesity, diabetes, and tumors. In this condition, mRNA to the 5' untranslated region of RALY fuses to the agouti transcript (1,2). RALY binds to U-rich elements in coding and non-coding mRNAs undergoing translation (3). RALY has been implicated in certain cancer types, as it can control transcriptional regulation and splicing of E2F1 and PRMT1 mRNAs (4,5).				
Background References		 Michaud, E.J. et al. (1994) Proc Natl Acad Sci U S A 91, 2562-6. Michaud, E.J. et al. (1993) Genes Dev 7, 1203-13. Rossi, A. et al. (2017) Nucleic Acids Res 45, 6775-6792. Bondy-Chorney, E. et al. (2017) Int J Biochem Cell Biol 91, 124-135. Cornella, N. et al. (2017) J Biol Chem 292, 19674-19692. 				
Species Reactiv	rity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western 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 W: Western Blotting

Limited Uses

Cross-Reactivity Key H: Human Mk: Monkey

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