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Daxx (25C12) Rabbit mAb

Applications: W, IF-IC	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 110	Source/Isotype: Rabbit IgG	UniProt ID: #Q9UER7	Entrez-Gene Id: 1616		
Product Usage Information		<b>Application</b> Western Blotting Immunofluorescence	(Immunocytochemi	stry)		<b>Dilution</b> 1:1000 1:25		
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/Sens	itivity	Daxx (25C12) Rabbit mAb detects endogenous levels of total Daxx protein. While Daxx has a calculated MW of 81 kDa, it has been shown to run at an apparent MW of 110 kDa at least in part due to post- translational hyper-phosphorylation (5).						
Species predicte based on 100% s homology	ed to react sequence	Monkey, Bovine, Dog						
Source / Purifica	ation	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to a region surrounding Gln255 of Daxx.						
Background		Daxx is a ubiquitously expressed protein that was originally identified through a yeast two-hybrid screen as an interactor with the cytoplasmic domain of Fas. It was found to enhance Fas-mediated apoptosis and activate the JNK pathway (1). However, additional studies have revealed that Daxx is actually a nuclear protein localizing to promyelocytic leukemia oncogenic domains (PODs) (2,3). Nuclear interactions have since been observed with CENP-C (4), Pax3 (5), DNA methyltransferase I (6) and chromatin-associated proteins, including histone deacetylase II, H2A, H2B, H3, H4 and Dek (5). Roles for Daxx have been suggested in transcriptional repression and cell cycle control. Loss of Daxx in mice leads to embryonic lethality with extensive developmental apoptosis, suggesting a role for Daxx directly or indirectly in suppressing cell death (5). Furthermore, inhibition of Daxx expression using RNAi has confirmed Daxx to be anti-apoptotic and to repress transcriptional activity of targets including NF-kB and E2F-1 (7).						
Background References 1. Yang, X. et al. (1997) Cell 89, 1067-76.   2. Torii, S. et al. (1999) EMBO J 18, 6037-49.   3. Li, H. et al. (2000) Mol Cell Biol 20, 1784-96.   4. Pluta, A.F. et al. (1998) J Cell Sci 111 (Pt 14), 2029-41.   5. Hollenbach, A.D. et al. (1999) EMBO J 18, 3702-11.   6. Michaelson, J.S. et al. (1999) Genes Dev 13, 1918-23.   7. Michaelson, J.S. and Leder, P. (2003) J Cell Sci 116, 345-52.								
Species Peactivi	+\/	Species reactivity is de	atermined by testing	uin at least one annrove	d application (e.g.	western blot)		
Western Blot Bu	ıffer	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 Ke	y	W: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)						
Cross-Reactivity	v Key	H: Human M: Mouse R: Rat						
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