#88652

## DJ-1 (D29E5) XP<sup>®</sup> Rabbit mAb (HRP Conjugate)



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Applications: W	<b>Reactivity:</b> H M R Hm Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 22	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q99497	<b>Entrez-Gene Id:</b> 11315		
Product Usage Information		<b>Application</b> Western Blotting	Dilution 1:1000					
Storage		Supplied in 136 mM NaCl, 2.6 mM KCl, 12 mM sodium phosphate (pH 7.4) dibasic, 2 mg/ml BSA, and 50% glycerol. Store at –20°C. Do not aliquot the antibody.						
Specificity/Sensitivity		DJ-1 (D29E5) XP $^{ extsf{@}}$ Rabbit mAb (HRP Conjugate) recognizes endogenous levels of total DJ-1 protein.						
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Lys148 of human DJ-1 protein.						
Description		This Cell Signaling Technology antibody is conjugated to the carbohydrate groups of horseradish peroxidase (HRP) via its amine groups. The HRP conjugated antibody is expected to exhibit the same species cross-reactivity as the unconjugated DJ-1 (D29E5) XP <sup>®</sup> Rabbit mAb #5933.						
Background		Parkinson's disease (PD) is characterized by the presence of Lewy bodies (intracellular inclusions) and by the loss of dopaminergic neurons. Research studies have shown that mutations in α-synuclein, Parkin, and DJ-1 are linked to PD (1). α-synuclein is a major component of the aggregates found in Lewy bodies. Parkin is involved in protein degradation through the ubiquitin-proteasome pathway, and investigators have shown that mutations in Parkin cause early onset of PD (1). Loss-of-function mutations in DJ-1 cause early onset of PD, but DJ-1 is associated with multiple functions: it cooperates with Ras to increase cell transformation, it positively regulates transcription of the androgen receptor, and it may function as an indicator of oxidative stress (2-5). Dopamine D2 receptor-mediated functions are greatly impaired in DJ-1 (-/-) mice, resulting in reduced long-term depression (6).						
Background R	eferences	1. Borrelli, E. (2005) <i>Neuron</i> 45, 479-81. 2. Bonifati, V. et al. (2003) <i>Science</i> 299, 256-9. 3. Nagakubo, D. et al. (1997) <i>Biochem. Biophys. Res. Commun.</i> 231, 509-13. 4. Takahashi, K. et al. (2001) <i>J. Biol. Chem.</i> 276, 37556-63. 5. Mitsumoto, A. and Nakagawa, Y. (2001) <i>Free Radic. Res.</i> 35, 885-93. 6. Goldberg, M.S. et al. (2005) <i>Neuron</i> 45, 489-96.						
Species Reacti	vity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Western Blot I	Buffer	er 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.				ר 5% w/v BSA, 1X		
Applications <b>K</b>	ley	W: Western Blotting						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey						
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