3/14/24, 10:33 AM Revision 5



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

Applications: WB, IHC-P, IF-IC	<b>Reactivity:</b> H M R Hm Mk	Sensitivity: Endogenous	<b>MW (kDa):</b> 80	Source/Isotype: Rabbit IgG	<b>UniProt ID:</b> #O95140	Entrez-Gene Id: 9927	
Product Usage Information	Application			Dilution			
mormation		Western Blotting			1:1000		
		Immunohistochemistry (Paraffin)			1:200 - 1:800		
	Imn	Immunofluorescence (Immunocytochemistry)			1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.					
	For a	For a carrier free (BSA and azide free) version of this product see product #92271.					
Specificity / Sensiti	i <b>vity</b> Mitof	Mitofusin-2 (D2D10) Rabbit mAb recognizes endogenous levels of total mitofusin-2 protein.					
Source / Purificatio		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val573 of human mitofusin-2 protein.					
Background	proc struc mitol mitod deve muta studi seve Rese an in	Mitofusins are mitochondrial transmembrane GTPases that function to regulate mitochondrial fusion, a process that occurs in concert with mitochondrial division and is necessary for the maintenance of structural and genetic mitochondrial integrity (1,2). Two mitofusins have been described in mammals, mitofusin-1 and -2, which share 60% amino acid identity and appear to function coordinately to regulate mitochondrial fusion (3). Mitochondrial fusion is widely recognized as important for normal cell growth and development (4), and may have evolved as a mechanism to offset the deleterious effects of mtDNA mutations (3). Null mutations in either mitofusin-1 and mitofusin-2 in skeletal muscle results in severe mitochondrial dysfunction (3). Research studies have revealed that mutations in mitofusin-2 are linked to Charcot-Marie-Tooth disease, an inherited neurogenerative disease characterized by a progressive loss of muscle tissue and sensory perception (5,6).					
Background Refere	Background References       1. Zhang, Y. and Chan, D.C. (2007) FEBS Lett 581, 2168-73.         2. Chan, D.C. (2006) Annu Rev Cell Dev Biol 22, 79-99.         3. Chen, H. et al. (2010) Cell 141, 280-9.         4. Bereiter-Hahn, J. and Vöth, M. (1994) Microsc Res Tech 27, 198-219.         5. Kijima, K. et al. (2005) Hum Genet 116, 23-7.         6. Züchner, S. et al. (2004) Nat Genet 36, 449-51.						
Species Reactivity	Speci	Species reactivity is determined by testing in at least one approved application (e.g., western				estern blot).	
Western Blot Buffe		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat di milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				% w/v nonfat dry	
Applications Key		WB: Western Blotting IHC-P: Immunohistochemistry (Paraffin) IF-IC: Immunofluorescence (Immunocytochemistry)					
Cross-Reactivity K	X: 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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Limited Uses

Mitofusin-2 (D2D10) Rabbit mAb (#9482) Datasheet Without Images Cell Signaling Technology

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