For Research Use Only

CISD2-Specific Monoclonal antibody

Catalog Number:66082-1-lg Featured Product 4 Publications

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Basic Information	Catalog Number: 66082-1-lg	GenBank Accession N NM_001008388	umber:	Purification Method: Protein A purification		
	Size:	GenelD (NCBI): ug/ml by 493856		CloneNo.: 3D7A3 Recommended Dilutions: WB 1:2000-1:16000 IHC 1:20-1:200 IF/ICC 1:50-1:500		
	150ul , Concentration: 1653 ug/ml by					
	Bradford method using BSA as the standard;					
	Source:					
	Mouse					
	lsotype: IgG2b					
Applications	Tested Applications:		Positive Controls:			
	WB, IHC, IF/ICC, ELISA Cited Applications:	Cited Applications: ROS1728 cells, RAW 2 WB, CoIP IHC : human kidney tis Species Specificity: IF/ICC : MCF-7 cells		fetal human brain tissue, MCF-7 cells, HeLa cel 1728 cells, RAW 264.7 cells, zebrafish tissue : human kidney tissue, human testis tissue		
	Species Specificity: human, zebrafish, rat, mouse			-7 cells,		
	Cited Species: human, mouse					
	Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0					
	CISD2 gene encodes a 15 kDa CDGSH iron-sulfur domain-containing protein 2, which is also named Miner1 or NAF- this protein was reported on endoplasmic reticulum membrane or mitochondrion outer membrane. Defects in CISD are the cause of Wolfram syndrome type 2 (WFS2), a rare disorder characterized by juvenile-onset insulin- dependent diabetes mellitus with optic atrophy. CISD2 regulates autophagy program by interacting BCL2, contributing to antagonize BECN1-mediated cellular autophagy at the endoplasmic reticulum. This monoclonal antibody is specific to CISD2 and does not cross-react with CISD1.					
Background Informatio	are the cause of Wolfram syndrome ty dependent diabetes mellitus with opt contributing to antagonize BECN1-me	smic reticulum membr. /pe 2 (WFS2), a rare dis tic atrophy. CISD2 regu ediated cellular autoph	ane or mitocho sorder characte Ilates autophag nagy at the end	ndrion outer membrar rized by juvenile-onse gy program by interact	ne. Defects in CISI et insulin- ting BCL2,	
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Background Information	Author Public Mailis Liiv 3902 Huiwen Xu 3900	smic reticulum membri rpe 2 (WFS2), a rare dis- tic atrophy. CISD2 regu- ediated cellular autoph s not cross-react with C med ID Journa 34309 Nat Co 03419 Comm 73330 Mol No er shipment. % glycerol pH 7.3.	ane or mitocho sorder characte ilates autophag agy at the end CISD1. al ommun nun Biol	ndrion outer membrar rized by juvenile-onse gy program by interact	ne. Defects in CISI et insulin- ting BCL2, his monoclonal Application WB,CoIP	

For technical support and original validation data for this product please contact: T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free E: proteintech@ptglab.com in USA), or 1(312) 455-8498 (outside USA) W: ptglab.com

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Selected Validation Data



WB result of CISD2 antibody (66082-1-1g, 1:3000) with si-Control and si-CISD2 transfected HeLa cells.



Immunohistochemical analysis of paraffinembedded human kidney using 66082-1-lg(CISD2-Specific antibody) at dilution of 1:50 (under 10x lens). $150 \text{ kDa} \rightarrow$ $100 \text{ kDa} \rightarrow$ $70 \text{ kDa} \rightarrow$ $50 \text{ kDa} \rightarrow$ $30 \text{ kDa} \rightarrow$ $20 \text{ kDa} \rightarrow$ $15 \text{ kDa} \rightarrow$

fetal human brain tissue were subjected to SDS PAGE followed by western blot with 66082-1-lg (CISD2-Specific Antibody) at dilution of 1:8000 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (-20°C Ethanol) fixed MCF-7 cells using 66082-1-Ig(CISD2-Specific antibody) at dilution of 1:100 and Alexa Fluor 488conjugated Goat Anti-Mouse IgG(H+L).