For Research Use Only

fibrillin 2-Specific Polyclonal antibody

Catalog Number:20252-1-AP 3 Publications

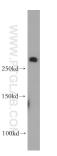


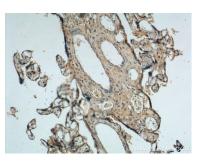
Basic Information	Catalog Number: 20252-1-AP	GenBank Accession Number NM_001999	r: Purification Method: Antigen affinity purification
	Size:	GenelD (NCBI):	Recommended Dilutions:
	150ul , Concentration: 350 ug/ml by	2201	WB 1:500-1:2000
	Nanodrop and 253 ug/ml by Bradford	UNIPROT ID:	IHC 1:50-1:500
	method using BSA as the standard; Source:	P35556	
	Rabbit	Full Name: fibrillin 2	
	Isotype:	Calculated MW:	
	lgG	315 kDa	
		Observed MW: 290 kDa, 160 kDa	
Applications	Tested Applications:	Positive Controls:	
	WB, IHC, ELISA Cited Applications:		: human brain tissue, HepG2 cells, LO2 cells, mouse g tissue
	WB, IHC		: human placenta tissue,
	ipecies Specificity:		
	human, mouse, rat Cited Species:		
	human, mouse		
	Note-IHC: suggested antigen ra TE buffer pH 9.0; (*) Alternativ retrieval may be performed w buffer pH 6.0	vely, antigen	
	FBN2 belongs to the fibrillin family. Fibrillins are structural components of 10-12 nm extracellular calcium-binding microfibrils which occur either in association with elastin or in elastin-free bundles. Fibrillin-2-containing microfibrils regulate the early process of elastic fiber assembly. Defects in FBN2 are the cause of congenital contractural arachnodactyly (CCA) which also known as Beals syndrome or distal arthrogryposis type 9 (DA9). This antibody is specific to FBN2. It recognizes both two isoforms ~300 kDa and 160 kDa.		
Background Information	microfibrils which occur either in asso microfibrils regulate the early proces contractural arachnodactyly (CCA) wh	ociation with elastin or in el s of elastic fiber assembly. I nich also known as Beals syr	astin-free bundles. Fibrillin-2-containing Defects in FBN2 are the cause of congenital ndrome or distal arthrogryposis type 9 (DA9). This
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	microfibrils which occur either in asso microfibrils regulate the early process contractural arachnodactyly (CCA) wh antibody is specific to FBN2. It recogn Author Put Sreenivasulu Chintala 346 Pan Long 394	ociation with elastin or in el s of elastic fiber assembly. I nich also known as Beals syn izes both two isoforms ~300 omed ID Journal 593597 Head Neck	astin-free bundles. Fibrillin-2-containing Defects in FBN2 are the cause of congenital ndrome or distal arthrogryposis type 9 (DA9). This IkDa and 160 kDa. Application WB rmacol WB,IHC
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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

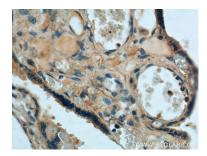
This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

Selected Validation Data





human brain tissue were subjected to SDS PAGE followed by western blot with 20252-1-AP (fibrillin 2-Specific antibody) at dilution of 1:500 incubated at room temperature for 1.5 hours. Immunohistochemical analysis of paraffinembedded human placenta tissue slide using 20252-1-AP (fibrillin 2-Specific antibody at dilution of 1:100 (under 10x lens).



Immunohistochemical analysis of paraffinembedded human placenta tissue slide using 20252-1-AP (fibrillin 2-Specific antibody at dilution of 1:100 (under 40x lens).