For Research Use Only

COL4A5 Polyclonal antibody

Catalog Number:19797-1-AP 16 Publications



	Catalog Number: 19797-1-AP	GenBank Accession Number: NM_000495	Purification Method:
	Size:	GenelD (NCBI):	Antigen affinity purification
	150ul , Concentration: 167 µg/ml by	1287	
	Bradford method using BSA as the standard;	UNIPROT ID: P29400	
	Source: Rabbit	Full Name: collagen, type IV, alpha 5	
	Isotype: IgG	Calculated MW: 161 kDa	
Applications	Tested Applications: ELISA		
	Cited Applications: WB, IHC, IF		
	Species Specificity: human		
	Cited Species: human, mouse, rat		
	COL4A5 belongs to the type IV collagen family. Type IV collagen is the major structural component of glomerular basement membranes (GBM), forming a 'chicken-wire' meshwork together with laminins, proteoglycans and entactin/nidogen. Defects in COL4A5 are the cause of Alport syndrome X-linked (APSX). Deletions covering the N-terminal regions of COL4A5 and COL4A6, which are localized in a head-to-head manner, are the cause of diffuse leiomyomatosis with Alport syndrome (DL-ATS)which also known as esophageal and vulval leiomyomatosis with nephropathy or Alport syndrome and diffuse leiomyomatosis (ATS-DL).		
Background Information	entactin/nidogen. Defects in COL4A5 terminal regions of COL4A5 and COL leiomyomatosis with Alport syndrom	are the cause of Alport syndrom 4A6, which are localized in a hea he (DL-ATS)which also known as e	ther with laminins, proteoglycans and e X-linked (APSX). Deletions covering the N- d-to-head manner, are the cause of diffuse esophageal and vulval leiomyomatosis with
Background Information	entactin/nidogen. Defects in COL4A5 terminal regions of COL4A5 and COL leiomyomatosis with Alport syndrom nephropathy or Alport syndrome and	are the cause of Alport syndrom 4A6, which are localized in a hea he (DL-ATS)which also known as e	other with laminins, proteoglycans and e X-linked (APSX). Deletions covering the N- d-to-head manner, are the cause of diffuse esophageal and vulval leiomyomatosis with).
	entactin/nidogen. Defects in COLAA5 terminal regions of COLAA5 and COL leiomyomatosis with Alport syndrom nephropathy or Alport syndrome and Author Put	are the cause of Alport syndromo 4A6, which are localized in a hea ne (DL-ATS)which also known as e diffuse leiomyomatosis (ATS-DL	ther with laminins, proteoglycans and e X-linked (APSX). Deletions covering the N- d-to-head manner, are the cause of diffuse esophageal and vulval leiomyomatosis with
	entactin/nidogen. Defects in COLAA5 terminal regions of COLAA5 and COL leiomyomatosis with Alport syndrom nephropathy or Alport syndrome and Author Put Jialin Li 360	are the cause of Alport syndrom 4A6, which are localized in a hea he (DL-ATS)which also known as e diffuse leiomyomatosis (ATS-DL omed ID Journal	other with laminins, proteoglycans and a X-linked (APSX). Deletions covering the N- d-to-head manner, are the cause of diffuse esophageal and vulval leiomyomatosis with). Application

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