For Research Use Only

Collagen Type VII Polyclonal antibody

Catalog Number:19799-1-AP

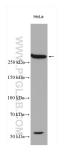


Basic Information	Catalog Number: 19799-1-AP	GenBank Accession Number: NM_000094	Purification Method: Antigen affinity purification
	Size: 150ul , Concentration: 400 ug/ml by Nanodrop and 300 ug/ml by Bradford method using BSA as the standard; Source: Rabbit Isotype: IgG	GenelD (NCBI): 1294	Recommended Dilutions: WB 1:500-1:1000
		UNIPROT ID: Q02388	
		Full Name: collagen, type VII, alpha 1	
		Calculated MW: 295 kDa	
		Observed MW: 290 kDa	
Applications	Tested Applications:	Positive Controls: WB : HeLa cells,	
	WB, ELISA Species Specificity: human		
Background Information	COL7A1, also named as LC collagen, is a stratified squamous epithelial basement membrane protein that forms anchoring fibrils which may contribute to epithelial basement membrane organization and adherence by interacting with extracellular matrix (ECM) proteins such as type IV collagen. Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica (DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica Pasini type (P-DEB) which also known as albopapuloid dominant dystrophic epidermolysis bullosa. Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica Hallopeau-Siemens type (HS-DEB). Defects in COL7A1 are the cause of transient bullous dermolysis of the newborn (TBDN). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica pretibial type (PR-DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica Bart type (B-DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica distrophica in COL7A1 are the cause of epidermolysis bullosa dystrophica bart type (B-DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica bart type (B-DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica bart type (B-DEB). Defects in COL7A1 are the cause of epidermolysis bullosa dystrophica is sollosa dystrophica with subcorneal cleavage (EBDSC) which also known as epidermolysis bullosa simplex superficialis (EBSS). This antibody is specific to COL7A1.		
Storage	Storage: Store at -20°C. Stable for one year after shipment. Storage Buffer: PBS with 0.02% sodium azide and 50% glycerol pH 7.3.		
	Aliquoting is unnecessary for -20°C s		

For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll freeE: proteintech@ptglab.comin 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



HeLa cells were subjected to SDS PAGE followed by western blot with 19799-1-AP (Collagen Type VII antibody) at dilution of 1:600 incubated at room temperature for 1.5 hours.