

#### www.immunoway.com.cn

# COL5A2 Rabbit pAb

CatalogNo: YT5533

### Key Features

Host SpeciesReactivity• Rabbit• Human,MouseMWIsotype

• 145kD (Observed)

Isotype • IgG Applications • WB,IHC,IF,ELISA

#### **Recommended Dilution Ratios**

WB 1:500-1:2000 IHC: 1:100-1:300 ELISA 1:20000 IF 1:50-200

### **Storage**

Storage\*-15°C to -25°C/1 year(Do not lower than -25°C)FormulationLiquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

# **Basic Information**

Clonality Polyclonal

# Immunogen Information

Immunogen	The antiserum was produced against synthesized peptide derived from the N-terminal
	region of human COL5A2. AA range:1-50

**Specificity** COL5A2 Polyclonal Antibody detects endogenous levels of COL5A2 protein.

# Target Information

Gene	name	COL5A2
Gene	name	COL5A2

#### **Protein Name** Collagen alpha-2(V) chain

Organism	Gene ID	UniProt ID
Human	<u>1290;</u>	<u>P05997;</u>
Mouse	<u>12832;</u>	<u>Q3U962;</u>

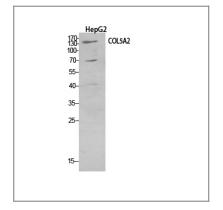
Cellular Secreted, extracellular space, extracellular matrix .

Localization

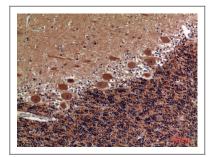
Tissue specificity Bone, Brain, Chondrosarcoma, Placenta, Skin,

**Function** Disease:Defects in COL5A2 are a cause of Ehlers-Danlos syndrome type 1 (EDS1) [MIM:130000]; also known as Ehlers-Danlos syndrome gravis or severe classic type Ehlers-Danlos syndrome. EDS is a connective tissue disorder characterized by hyperextensible skin, atrophic cutaneous scars due to tissue fragility and joint hyperlaxity. EDS1 is the severe form of classic Ehlers-Danlos syndrome.,Disease:Defects in COL5A2 are a cause of Ehlers-Danlos syndrome type 2 (EDS2) [MIM:130010]; also known as Ehlers-Danlos syndrome mitis or mild classic type Ehlers Danlos syndrome., Disease: Genetic variation in COL5A2 is associated with spontaneous cervical artery dissections (sCAD), sCAD are an important cause of stroke among young and middle-aged patients. Ultrastructural abnormalities are observed in skin biopsies of most patients with sCAD. Major findings included enlarged and irregular collagen fibrils and pronounced elastic fibers fragmentation., Function: Type V collagen is a member of group I collagen (fibrillar forming collagen). It is a minor connective tissue component of nearly ubiguitous distribution. Type V collagen binds to DNA, heparan sulfate, thrombospondin, heparin, and insulin. Type V collagen is a key determinant in the assembly of tissue-specific matrices., PTM: Prolines at the third position of the tripeptide repeating unit (G-X-Y) are hydroxylated in some or all of the chains., similarity: Belongs to the fibrillar collagen family., similarity: Contains 1 VWFC domain., subunit: Trimers of two alpha 1(V) and one alpha 2(V) chains in most tissues and trimers of one alpha 1(V), one alpha 2(V), and one alpha 3(V) chains in placenta.,

### Validation Data



Western Blot analysis of HepG2 cells using COL5A2 Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000



Immunohistochemical analysis of paraffin-embedded human-brain, antibody was diluted at 1:100

# **Contact information**

Orders:	order.cn@immunoway.com
Support:	support.cn@immunoway.com
Telephone:	400-8787-807(China)
Website:	http://www.immunoway.com.cn
Address:	2200 Ringwood Ave San Jose, CA 95131 USA



Please scan the QR code to access additional product information: **COL5A2 Rabbit pAb** 

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

Antibody | ELISA Kits | Protein | Reagents