abcam

Product datasheet

Anti-Collagen VI antibody [EPR7888(N)] ab172606

Recombinant RabMAb

2 References 2 Images

Overview

Product name Anti-Collagen VI antibody [EPR7888(N)]

Description Rabbit monoclonal [EPR7888(N)] to Collagen VI

Host species Rabbit

Suitable for: WB **Tested applications**

Unsuitable for: ICC/IF,IHC-P or IP

Reacts with: Human Species reactivity

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide within Human Collagen VI aa 200-300. The exact sequence is proprietary.

Database link: P12110

Positive control Human skin, placenta and fetal heart lysates.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb** patents.

Properties

Form

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

Purity Tissue culture supernatant

Clonality Monoclonal

Clone number EPR7888(N)

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab172606 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/1000 - 1/10000. Detects a band of approximately 140 kDa (predicted molecular weight: 109 kDa).

Application notes Is unsuitable for ICC/IF,IHC-P or IP.

Target

Function Collagen VI acts as a cell-binding protein.

Involvement in disease Defects in COL6A1 are a cause of Bethlem myopathy (BM) [MIM:158810]. BM is a rare

autosomal dominant proximal myopathy characterized by early childhood onset (complete penetrance by the age of 5) and joint contractures most frequently affecting the elbows and

ankles.

Defects in COL6A1 are a cause of Ullrich congenital muscular dystrophy (UCMD) [MIM:254090];

also known as Ullrich scleroatonic muscular dystrophy. UCMD is an autosomal recessive

congenital myopathy characterized by muscle weakness and multiple joint contractures, generally

Prolines at the third position of the tripeptide repeating unit (G-X-Y) are hydroxylated in some or all

noted at birth or early infancy. The clinical course is more severe than in Bethlem myopathy.

Sequence similarities Belongs to the type VI collagen family.

Contains 3 VWFA domains.

Post-translational

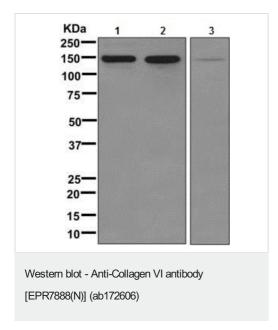
modifications

of the chains.

Secreted > extracellular space > extracellular matrix.

Cellular localization

Images



All lanes : Anti-Collagen VI antibody [EPR7888(N)] (ab172606) at 1/1000 dilution

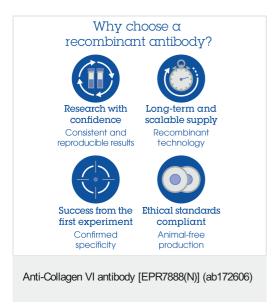
Lane 1: Human skin lysate

Lane 2: Human placenta lysate

Lane 3: Human fetal heart lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 109 kDa



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