abcam

Product datasheet

Native Human Collagen IV protein ab7536

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製品の詳細

製品名 Native Human Collagen IV protein

 発現系
 Native

 アクセッション番号
 P02462

タンパク質長 Full length protein

Animal free No

由来 Native

生物種Human予測される分子量161 kDa

配列の追加情報 Prepared from human placenta and is chromatographically and immunologically pure.

特性

Our Abpromise guarantee covers the use of ab7536 in the following tested applications.

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

アプリケーション SDS-PAGE

Western blot

Immunoprecipitation

ELISA

Blocking

製品の状態 Liquid

備考 This product is free from other collagens, human serum proteins and non-collagen extracellular

matrix proteins. This product reacts with anti-Collagen Type IV. Reaction with anti-Collagen I, II, III,

V or VI is negligible (typically less than 1% cross reactivity was detected by ELISA).

前処理および保存

保存方法および安定性 Shipped at 4°C. Upon delivery aliquot. Store at +4°C. Store undiluted.

Preservative: 0.01% Sodium azide Constituent: 3% Acetic acid buffer

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関連情報

機能

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. Arresten, comprising the C-terminal NC1 domain, inhibits angiogenesis and tumor formation. The C-terminal half is found to possess the anti-angiogenic activity. Specifically inhibits endothelial cell proliferation, migration and tube formation. Inhibits expression of hypoxia-inducible factor 1alpha and ERK1/2 and p38 MAPK activation. Ligand for alpha1/beta1 integrin.

組織特異性

関連疾患

Highly expressed in placenta.

Defects in COL4A1 are a cause of brain small vessel disease with hemorrhage (BSVDH) [MIM:607595]. Brain small vessel diseases underlie 20 to 30 percent of ischemic strokes and a larger proportion of intracerebral hemorrhages. Inheritance is autosomal dominant.

Defects in COL4A1 are the cause of hereditary angiopathy with nephropathy aneurysms and muscle cramps (HANAC) [MIM:611773]. The clinical renal manifestations include hematuria and bilateral large cysts. Histologic analysis revealed complex basement membrane defects in kidney and skin. The systemic angiopathy appears to affect both small vessels and large arteries. Defects in COL4A1 are a cause of porencephaly familial (PCEPH) [MIM:175780]. Porencephaly is a term used for any cavitation or cerebrospinal fluid-filled cyst in the brain. Porencephaly type 1 is usually unilateral and results from focal destructive lesions such as fetal vascular occlusion or birth trauma. Type 2, or schizencephalic porencephaly, is usually symmetric and represents a primary defect or arrest in the development of the cerebral ventricles.

配列類似性

ドメイン

翻訳後修飾

Belongs to the type IV collagen family.

Contains 1 collagen IV NC1 (C-terminal non-collagenous) domain.

Alpha chains of type IV collagen have a non-collagenous domain (NC1) at their C-terminus, frequent interruptions of the G-X-Y repeats in the long central triple-helical domain (which may

cause flexibility in the triple helix), and a short N-terminal triple-helical 7S domain.

Lysines at the third position of the tripeptide repeating unit (G-X-Y) are hydroxylated in all cases

and bind carbohydrates.

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

of the chains.

Type IV collagens contain numerous cysteine residues which are involved in inter- and intramolecular disulfide bonding. 12 of these, located in the NC1 domain, are conserved in all

known type IV collagens.

The trimeric structure of the NC1 domains is stabilized by covalent bonds between Lys and Met

residues.

Proteolytic processing produces the C-terminal NC1 peptide, arresten.

細胞内局在

Secreted > extracellular space > extracellular matrix > basement membrane.

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