

ANTXR2 Polyclonal Antibody

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP54361

Product Information

Application WB, IHC-P, IHC-F, IF, ICC, E

Primary Accession P58335

Reactivity Rat, Pig, Dog, Bovine

Host Rabbit
Clonality Polyclonal
Calculated MW 53666
Physical State Liquid

Immunogen KLH conjugated synthetic peptide derived from human ANTXR2

Epitope Specificity 101-200/489

Isotype IgG

Purity affinity purified by Protein A

Buffer 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

SUBCELLULAR LOCATION Secreted; Cell membrane. Expressed at the cell surface and Endoplasmic

reticulum membrane. Expressed predominantly within the endoplasmic

reticulum and not at the plasma membrane.

SIMILARITY Belongs to the ATR family.Contains 1 VWFA domain.

SUBUNIT Binds laminin, and possibly also collagen type IV. Binds to the protective

antigen (PA) of Bacillus anthracis in a divalent cation-dependent manner, with the following preference: calcium > manganese > magnesium > zinc. Binding

of PA leads to heptamerization of the receptor-PA complex.

DISEASEDefects in ANTXR2 are the cause of infantile systemic hyalinosis (ISH). This

autosomal recessive syndrome is similar to JHF, but has an earlier onset and a more severe course. Symptoms appear at birth or within the first months of life, with painful, swollen joint contractures, osteopenia, osteoporosis and livid red hyperpigmentation over bony prominences. Patients develop multiple subcutaneous skin tumors and gingival hypertrophy. Hyaline deposits in multiple organs, recurrent infections and intractable diarrhea often lead to death within the first 2 years of life. Surviving children may

suffer from severely reduced mobility due to joint contractures.

Important Note This product as supplied is intended for research use only, not for use in

human, therapeutic or diagnostic applications.

Background Descriptions This gene encodes a receptor for anthrax toxin. The protein binds to collagen

IV and laminin, suggesting that it may be involved in extracellular matrix adhesion. Mutations in this gene cause juvenile hyaline fibromatosis and infantile systemic hyalinosis. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Mar 2009].

Additional Information

Gene ID 118429

Other Names Anthrax toxin receptor 2, Capillary morphogenesis gene 2 protein, CMG-2,

ANTXR2, CMG2

Target/Specificity Expressed in prostate, thymus, ovary, testis, pancreas, colon, heart, kidney,

lung, liver, peripheral blood leukocytes, placenta, skeletal muscle, small

intestine and spleen.

Dilution WB=1:500-2000,IHC-P=1:100-500,IHC-F=1:100-500,ICC=1:100-500,IF=1:100-50

0,ELISA=1:5000-10000

Format 0.01M TBS(pH7.4) with 1% BSA, 0.09% (W/V) sodium azide and 50% Glyce

Storage Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When

reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody

is stable for at least two weeks at 2-4 °C.

Protein Information

Name ANTXR2 (HGNC:21732)

Function Necessary for cellular interactions with laminin and the extracellular matrix.

Cellular Location [Isoform 1]: Cell membrane; Single-pass type I membrane protein.

Note=Expressed at the cell surface [Isoform 3]: Secreted.

Tissue Location Expressed in prostate, thymus, ovary, testis, pancreas, colon, heart, kidney,

lung, liver, peripheral blood leukocytes, placenta, skeletal muscle, small

intestine and spleen

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.