# **Product Name: CD231 Rabbit Polyclonal Antibody**

Catalog #: APRab08293



#### **Summary**

Production Name CD231 Rabbit Polyclonal Antibody

**Description** Rabbit Polyclonal Antibody

Host Rabbit
Application IHC,ELISA

**Reactivity** Human, Mouse, Rat

#### **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

Clonality Polyclonal Form Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw Storage

cycles.

**Buffer** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.

**Purification** Affinity purification

#### **Immunogen**

**Gene Name** TSPAN7 A15 DXS1692E MXS1 TM4SF2

Tetraspanin-7 (Tspan-7;Cell surface glycoprotein A15;Membrane component

Alternative Names chromosome X surface marker 1;T-cell acute lymphoblastic leukemia-associated

antigen 1;TALLA-1;Transmembrane 4 superfamily member 2;CD antigen CD231)

Gene ID 7102.0

**SwissProt ID** P41732.Synthetic peptide from human protein at AA range: 101-150

### **Application**

**Dilution Ratio** IHC 1:50-200, ELISA 1:10000-20000.

**Molecular Weight** 

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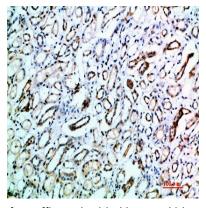


#### **Background**

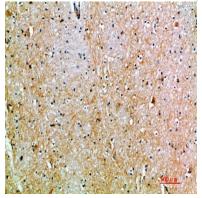
The protein encoded by this gene is a member of the transmembrane 4 superfamily, also known as the tetraspanin family. Most of these members are cell-surface proteins that are characterized by the presence of four hydrophobic domains. The proteins mediate signal transduction events that play a role in the regulation of cell development, activation, growth and motility. This encoded protein is a cell surface glycoprotein and may have a role in the control of neurite outgrowth. It is known to complex with integrins. This gene is associated with X-linked mental retardation and neuropsychiatric diseases such as Huntington's chorea, fragile X syndrome and myotonic dystrophy. [provided by RefSeq, Jul 2008], disease:Defects in TSPAN7 are the cause of mental retardation X-linked type 58 (MRX58) [MIM:300210]. Mental retardation is characterized by significantly sub-average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. Non-syndromic mental retardation patients do not manifest other clinical signs, function:May be involved in cell proliferation and cell motility, similarity:Belongs to the tetraspanin (TM4SF) family, tissue specificity:Not solely expressed in T-cells. Expressed in acute myelocytic leukemia cells of some patients.,

#### Research Area

#### **Image Data**



Immunohistochemical analysis of paraffin-embedded human-kidney, antibody was diluted at 1:200



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Immunohistochemical analysis of paraffin-embedded human-brain, antibody was diluted at 1:200

#### Note

For research use only.

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