

## CD231 Polyclonal Antibody

|                              |  |
|------------------------------|--|
| <b>Catalog No :</b>          | YT5946   |
| <b>Reactivity :</b>          | Human;Mouse;Rat  |
| <b>Applications :</b>        | IHC;IF;ELISA   |
| <b>Target :</b>              | CD231  |
| <b>Fields :</b>              | >>Transcriptional misregulation in cancer  |
| <b>Gene Name :</b>           | TSPAN7 A15 DXS1692E MXS1 TM4SF2  |
| <b>Protein Name :</b>        | Tetraspanin-7 (Tspan-7) (Cell surface glycoprotein A15) (Membrane component chromosome X surface marker 1) (T-cell acute lymphoblastic leukemia-associated antigen 1) (TALLA-1) (Transmembrane 4 superfa |
| <b>Human Gene Id :</b>       | 7102   |
| <b>Human Swiss Prot No :</b> | P41732   |
| <b>Mouse Gene Id :</b>       | 21912  |
| <b>Mouse Swiss Prot No :</b> | Q62283   |
| <b>Immunogen :</b>           | Synthetic peptide from human protein at AA range: 101-150  |
| <b>Specificity :</b>         | The antibody detects endogenous CD231  |
| <b>Formulation :</b>         | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.  |
| <b>Source :</b>              | Polyclonal, Rabbit,IgG   |
| <b>Dilution :</b>            | IHC 1:50-200, ELISA 1:10000-20000. IF 1:50-200   |
| <b>Purification :</b>        | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.  |
| <b>Concentration :</b>       | 1 mg/ml  |

**Storage Stability :** -15°C to -25°C/1 year (Do not lower than -25°C)

**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],

**Function :** 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 adaptive 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.,

**Subcellular Location :** Membrane; Multi-pass membrane protein.

**Expression :** Not solely expressed in T-cells. Expressed in acute myelocytic leukemia cells of some patients.

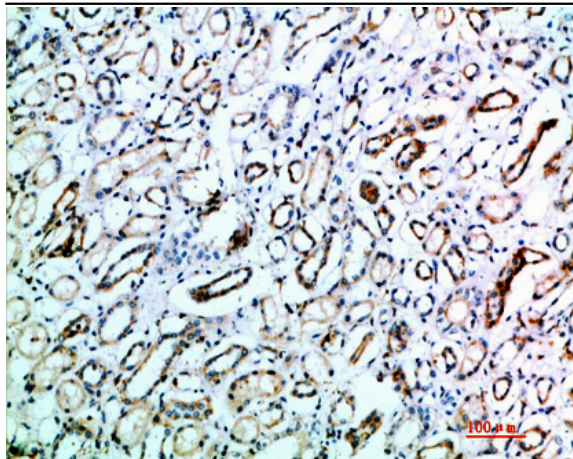
**Sort :** 3482

**No4 :** 1

**Host :** Rabbit

**Modifications :** Unmodified

## Products Images



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