



CD231 rabbit pAb

Cat#: orb771573 (Manual)

For research use only. Not intended for diagnostic use.

Product Name CD231 rabbit pAb

Host species Rabbit

Applications IHC;IF;ELISA

Species Cross-Reactivity Human; Mouse; Rat

Recommended dilutions IHC-p 1:50-200, ELISA 1:10000-20000

Immunogen Synthetic peptide from human protein at AA range: 101-150

The antibody detects endogenous CD231 **Specificity**

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium

azide..

Store at -20°C. Avoid repeated freeze-thaw cycles. **Storage**

Protein Name 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

TSPAN7 A15 DXS1692E MXS1 TM4SF2 Gene Name

Cellular localization Membrane; Multi-pass membrane protein.

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.





Clonality Polyclonal

Concentration 1 mg/ml

Observed band

Human Gene ID 7102

Human Swiss-Prot Number P41732

Alternative Names Tetraspanin-7 (Tspan-7; Cell surface glycoprotein A15; Membrane

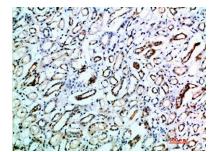
component chromosome X surface marker 1;T-cell acute lymphoblastic leukemia-associated antigen 1;TALLA-1;Transmembrane 4 superfamily

member 2;CD antigen CD231)

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],



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







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