

**Product Name: CD151 Rabbit Polyclonal Antibody**  
**Catalog #: APRab08219**



## Summary

<b>Production Name</b>	CD151 Rabbit Polyclonal Antibody
<b>Description</b>	Rabbit Polyclonal Antibody
<b>Host</b>	Rabbit
<b>Application</b>	IHC,ELISA
<b>Reactivity</b>	Human,Mouse,Rat

## Performance

<b>Conjugation</b>	Unconjugated
<b>Modification</b>	Unmodified
<b>Isotype</b>	IgG
<b>Clonality</b>	Polyclonal
<b>Form</b>	Liquid
<b>Storage</b>	Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.
<b>Buffer</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.
<b>Purification</b>	Affinity purification

## Immunogen

<b>Gene Name</b>	CD151 TSPAN24
<b>Alternative Names</b>	CD151 antigen (GP27;Membrane glycoprotein SFA-1;Platelet-endothelial tetraspan antigen 3;PETA-3;Tetraspanin-24;Tspan-24;CD antigen CD151)
<b>Gene ID</b>	977.0
<b>SwissProt ID</b>	P48509.Synthetic peptide from human protein at AA range: 91-140

## Application

<b>Dilution Ratio</b>	IHC 1:50-200 ELISA 1:10000-20000
<b>Molecular Weight</b>	

## Background

The protein encoded by this gene is a member of the transmembrane 4 superfamily, also known as the tetraspanin family.

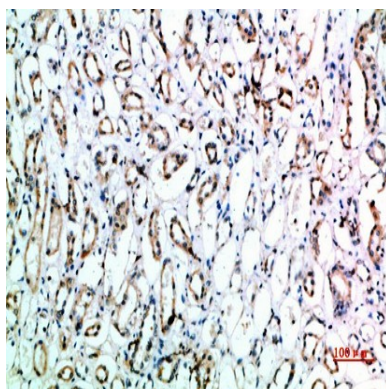
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Most of these members are cell-surface proteins that are characterized by the presence of four hydrophobic domains. The proteins mediate signal transduction events that play a role in the regulation of cell development, activation, growth and motility. This encoded protein is a cell surface glycoprotein that is known to complex with integrins and other transmembrane 4 superfamily proteins. It is involved in cellular processes including cell adhesion and may regulate integrin trafficking and/or function. This protein enhances cell motility, invasion and metastasis of cancer cells. Multiple alternatively spliced transcript variants that encode the same protein have been described for this gene. [provided by RefSeq, Jul 2008],disease:Defects in CD151 are the cause of nephropathy with pretibial epidermolysis bullosa and deafness (NPEBD) [MIM:609057]. NPEBD is characterized by the association of hereditary nephritis, epidermolysis bullosa, deafness, and beta-thalassemia minor.,function:Essential for the proper assembly of the glomerular and tubular basement membranes in kidney.,induction:By HTLV-1.,online information:Blood group antigen gene mutation database,polymorphism:CD151 defines the MER2=RAPH1 antigen of the RAPH blood group system. 92% of Caucasians are MER2-positive and 8% are apparently MER2-negative.,similarity:Belongs to the tetraspanin (TM4SF) family.,subunit:Interacts with integrins alpha3beta1, alpha5beta1, alpha3beta1 and alpha6beta4, with CD9 and CD181.,tissue specificity:Expressed in a variety of tissues including vascular endothelium and epidermis. Expressed on erythroid cells, with a higher level of expression in erythroid precursors than on mature erythrocytes.,

## Research Area

## Image Data



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

## Note

For research use only.