

Anti-CD9 Rabbit Monoclonal Antibody

Catalog Number: M01202

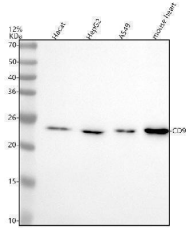
Overview

Product Name	Anti-CD9 Rabbit Monoclonal Antibody
Reactive Species	Human, Mouse, Rat
Description	Boster Bio Anti-CD9 Rabbit Monoclonal Antibody catalog # M01202. Tested in WB, IHC, ICC/IF, IP, Flow Cytometry applications. This antibody reacts with Human, Mouse, Rat.
Application	Flow Cytometry, IP, IF, IHC, ICC, WB
Clonality	Monoclonal CED-3
Formulation	Rabbit IgG in stabilizing components, phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. *This antibody is supplied in a stabilized formulation. Compatibility with conjugation reactions depends on the chemistry of the conjugation method used. For conjugation methods that are not compatible with the stabilizing components present in this formulation, a carrier-free antibody format is required.
Storage Instructions	Store at -20°C for one year. For short term storage and frequent use, store at 4°C for up to one month. Avoid repeated freeze-thaw cycles.
Host	Rabbit
Uniprot ID	P21926

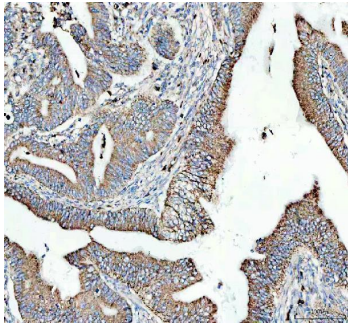
Technical Details

Immunogen	A synthesized peptide derived from human CD9
Isotype	Rabbit IgG
Form	Liquid
Concentration	0.5mg/ml
Purification	Affinity-chromatography
Suggested Dilutions	WB 1:500-2000 IHC 1:50-200 ICC/IF 1:50-200 IP 1:20 FC 1:20

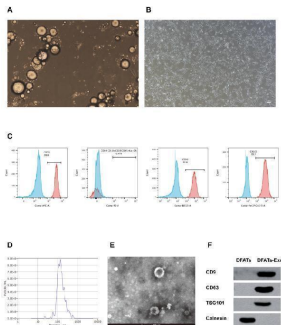
Anti-CD9 Rabbit Monoclonal Antibody (M01202) Images



Western blot analysis of CD9 using anti-CD9 antibody (M01202). Electrophoresis was performed on a 12% SDS-PAGE gel at 80V (Stacking gel) / 120V (Resolving gel) for 2 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: human Hacat whole cell lysates, Lane 2: human HepG2 whole cell lysates, Lane 3: human A549 whole cell lysates, Lane 4: mouse heart tissue lysates. After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-CD9 antigen affinity purified monoclonal antibody (M01202) at 1:1000 overnight at 4°C, then washed with TBS-0.1% Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an ECL Plus Western Blotting Substrate (Catalog # AR1196-200) with Tanon 5200 system. A specific band was detected for CD9 at approximately 23 kDa. The expected band size for CD9 is at 25 kDa.

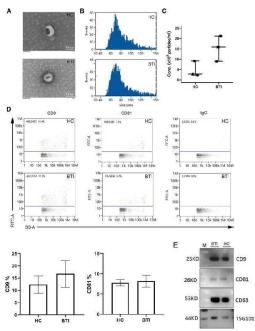


IHC analysis of CD9 using anti-CD9 antibody (M01202). CD9 was detected in a paraffin-embedded section of human colon cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1:100 rabbit anti-CD9 Antibody (M01202) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

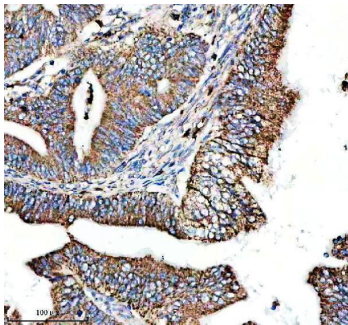


Illustrates the characterization of DFATs and exosomes. DFATs exhibited a spindle-shaped morphology (A-B). Flow cytometry analysis showed that DFATs were positive for markers CD90, CD105, and CD73, and negative for CD34, CD11b, CD19, CD45, and HLA-DR (C). NTA showed that the average diameter of the exosomes was 136.3 nm (D). TEM showed that DFATs-Exos exhibited a typical bilayer membrane structure (E). Western blot analysis revealed that DFATs exosomes were enriched in markers such as CD63, CD9, and TSG101, while lacking the marker Calnexin (F). Full-length blots are presented in Supplementary Digital Material 1 Index in PubMed under a CC BY license. PMID: 40022232

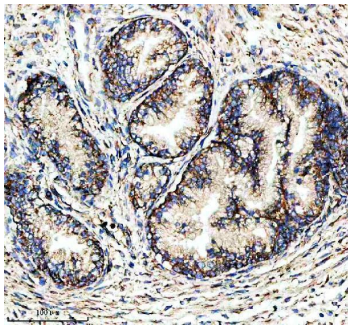
Characterization of EVs (extracellular vesicles) from HC (healthy control) and BTI (biliary tract infection) samples. (A) Representative images of EVs, which were captured by TEM (Transmission Electron Microscope), from HC or BTI



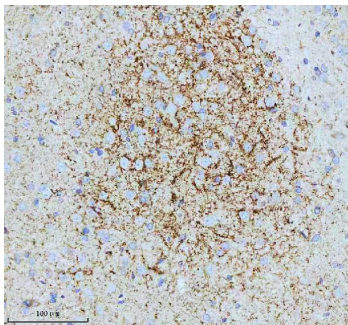
samples (bar = 100 nm). (B , C , D) Plot showing the sizes (B) or concentrations (Conc.; C) and cell surface expression of CD9/CD81/IgG (D) of EVs which were measured by the NFCM (Nano-Flow Cytometry Measurement). (E) CD9, CD81, CD63 and TSG101 were detectable by Western blot (WB) analysis. Index in PubMed under a CC BY license. PMID: 38459197



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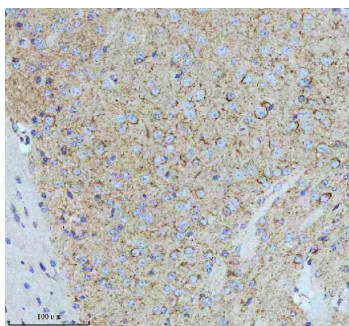


IHC analysis of CD9 using anti-CD9 antibody (M01202). CD9 was detected in a paraffin-embedded section of human prostate cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with a dilution of 1:100 rabbit anti-CD9 Antibody (M01202) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

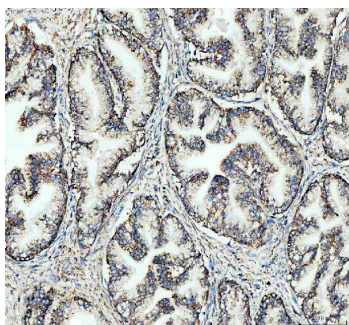


IHC analysis of CD9 using anti-CD9 antibody (M01202). CD9 was detected in a paraffin-embedded section of rat brain tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with a dilution of 1:100 rabbit anti-CD9 Antibody (M01202) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

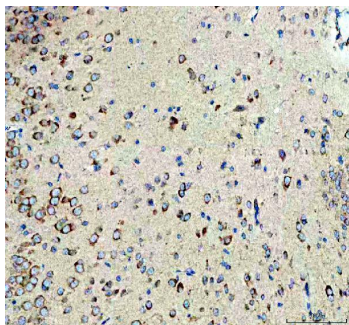
IHC analysis of CD9 using anti-CD9 antibody (M01202). CD9 was detected in a paraffin-embedded section of mouse brain tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue



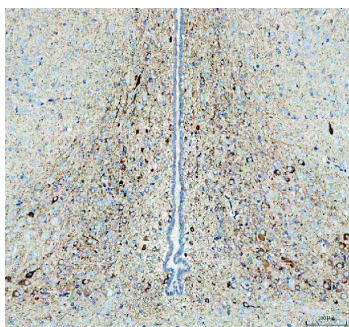
section was then incubated with a dilution of 1:100 rabbit anti-CD9 Antibody (M01202) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.



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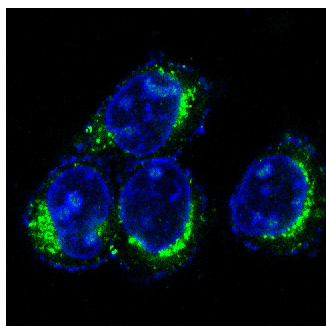


IHC analysis of CD9 using anti-CD9 antibody (M01202). CD9 was detected in a paraffin-embedded section of mouse brain tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1:100 rabbit anti-CD9 Antibody (M01202) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.



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Immunofluorescent analysis of Hela cells, using CD9 Antibody .



2 Publications Citing This Product

1. PubMed ID: , The bone marrow microenvironment at single-cell resolution
2. PubMed ID: 32059163, Wang D,Hao C,Zhang L,Zhang J,Liu S,Li Y,Qu Y,Zhao Y,Huang R,Wei J,Yao W.Exosomal miR-125a-5p derived from silica-exposed macrophages induces fibroblast transdifferentiation.Ecotoxicol Environ Saf.2020 Apr 1;192:110253.doi:10.1016/j.ecoenv.2020.110253.Epub

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Anti-CD9 Rabbit Monoclonal Antibody

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