

PRODUCT INFORMATION

Clone ID DM9 CS1 **Target**

Synonyms SLAM7 (19A; CD319; CRACC; CS1)

Host Species Rabbit

Description Anti-CS1 antibody(DM9); Rabbit mAb

Delivery In Stock **Uniprot ID** Q9NQ25 IgG type Rabbit IgG Clonality Monoclonal Reactivity Human

Applications ELISA; Flow Cyt Recommended Flow Cyt 1:100

Dilutions

Storage & Shipping

Purified from cell culture supernatant by affinity **Purification** chromatography

Lyophilized from sterile PBS, pH 7.4. Normally 5 % Formulation & 8% trehalose is added as protectants before Reconstitution

lyophilization. Please see Certificate of Analysis for specific instructions of reconstitution.

Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not intended for use within a month, aliquot and store at -80°C (Avoid repeated freezing and thawing).

Lyophilized proteins are shipped at ambient temperature.

Self-ligand receptor of the signaling lymphocytic activation molecule (SLAM) family. SLAM

receptors triggered by homo- or heterotypic cell-cell interactions are modulating the activation and differentiation of a wide variety of immune cells and thus are involved in the regulation and interconnection of both innate and adaptive immune response. Activities are controlled by presence or absence of small cytoplasmic adapter proteins; SH2D1A:SAP and:or SH2D1B:EAT-2. Isoform 1 mediates NK cell activation through a SH2D1A-independent extracellular signal-

regulated ERK-mediated pathway (PubMed:11698418). Positively regulates NK cell Background

functions by a mechanism dependent on

phosphorylated SH2D1B. Downstream signaling implicates PLCG1; PLCG2 and PI3K (PubMed:16339536). In addition to heterotypic NK cells-target cells interactions also homotypic interactions between NK cells may contribute to activation. However; in the absence of SH2D1B; inhibits NK cell function. Acts also inhibitory in Tcells (By similarity). May play a role in lymphocyte adhesion (PubMed:11802771). In LPS-activated

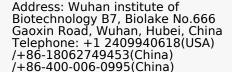
monocytes negatively regulates production of proinflámmatory cytókines (PubMed:23695528).

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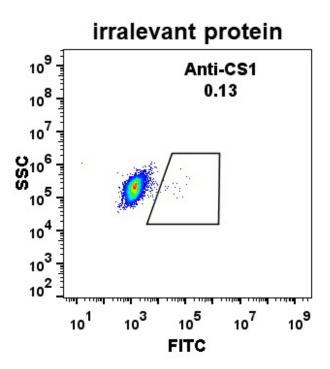
Usage Research use only

Conjugate Unconjugated









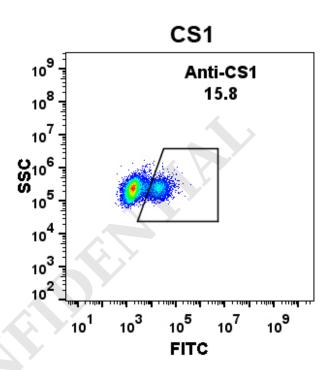


Figure 1. Expi 293 cell line transfected with irrelevant protein (left) and human CS1 (right) were surface stained with Rabbit anti-CS1monoclonal antibody $1\mu g/ml$ (clone: DM9) followed by Alexa 488-conjugated anti-rabbit IgG secondary antibody.

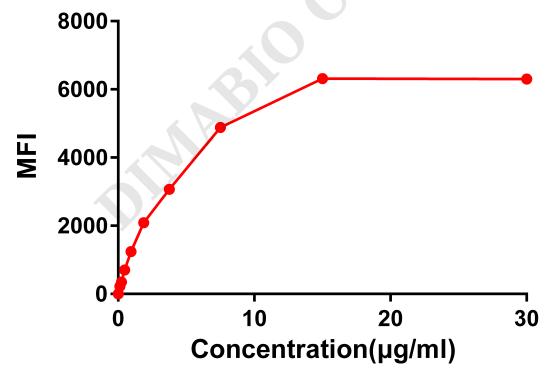


Figure 2. Flow cytometry data of serially titrated Rabbit anti-CS1 monoclonal antibody (clone: DM9) on Raji cells. The Y-axis represents the mean fluorescence intensity (MFI) while the X-axis represents the concentration of IgG used.

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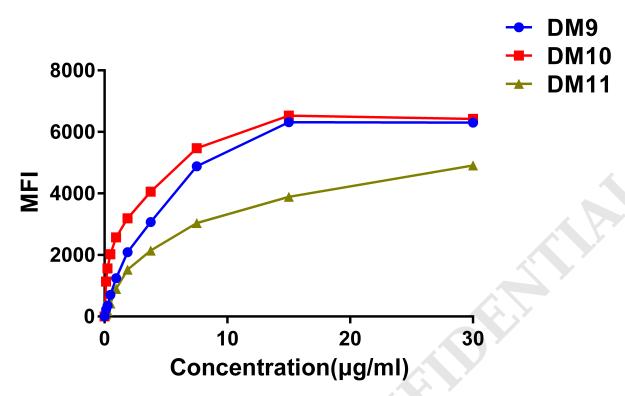


Figure 3. Affinity ranking of different Rabbit anti-CS1 mAb clones by titration of different concentration onto Raji cells. The Y-axis represents the mean fluorescence intensity (MFI) while the X-axis represents the concentration of IgG used.

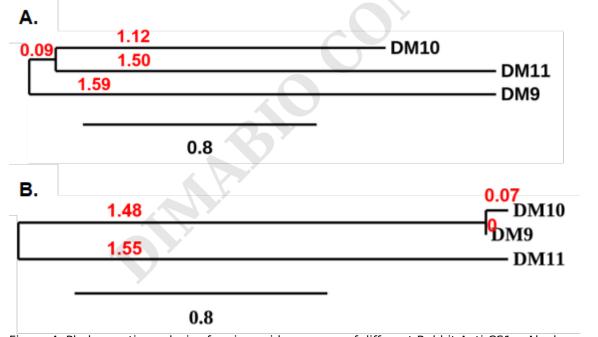


Figure 4. Phylogenetic analysis of amino acid sequence of different Rabbit Anti-CS1 mAb clones. A) Heavy chain and B) Light cháin.

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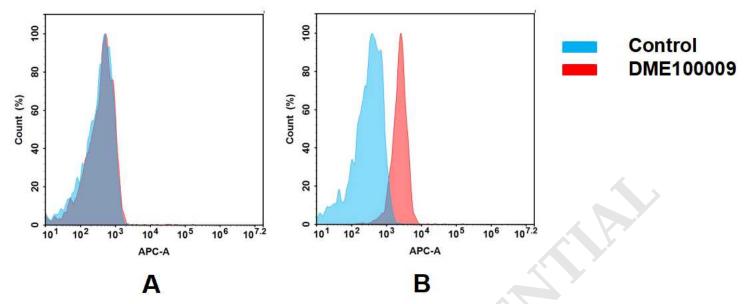


Figure 5. Flow cytometry analysis of antigen binding of rabbit anti-human CS1 mAb(DME100009).

(A) DME100009 does not bind to 293T cells that do not express CS1. (B) A clear peak shift of DME100009 was seen compared to the control when incubated with CS1-expressing MM.1S cells, indicating strong binding of DME100009 to CS1. Antibodies were incubated at 2 μ g/mL.

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