

CD62E/E-Selectin Rabbit mAb

Catalog No.: A23836 **Recombinant**

Basic Information

Observed MW

110-120kDa

Calculated MW

66kDa

Category

Primary antibody

Applications

ELISA, WB, FC

Cross-Reactivity

Human

CloneNo number

ARC61987

Background

The protein encoded by this gene is found in cytokine-stimulated endothelial cells and is thought to be responsible for the accumulation of blood leukocytes at sites of inflammation by mediating the adhesion of cells to the vascular lining. It exhibits structural features such as the presence of lectin- and EGF-like domains followed by short consensus repeat (SCR) domains that contain 6 conserved cysteine residues. These proteins are part of the selectin family of cell adhesion molecules. Adhesion molecules participate in the interaction between leukocytes and the endothelium and appear to be involved in the pathogenesis of atherosclerosis.

Recommended Dilutions

WB 1:500 - 1:1000

FC 1:100 - 1:500

Immunogen Information

Gene ID

6401

Swiss Prot

P16581

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 22-230 of human CD62E/E-Selectin (NP_000441.2).

Synonyms

SELE; CD62E; ELAM; ELAM1; ESEL; LECAM2; E-selectin; CD62E/E-Selectin

Contact

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Product Information

Source

Rabbit

Isotype

IgG

Purification

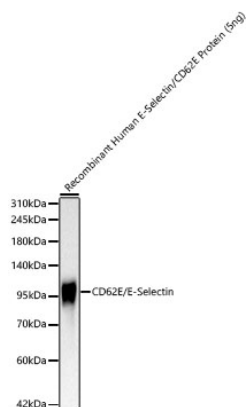
Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300, 0.05% BSA, 50% glycerol, pH7.3.

Validation Data



Western blot analysis of recombinant Human E-Selectin/CD62E Protein, using CD62E/E-Selectin Rabbit mAb (A23836) at 1:1000 dilution.

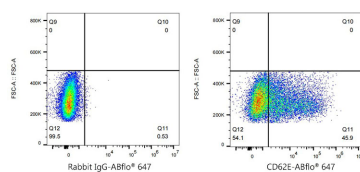
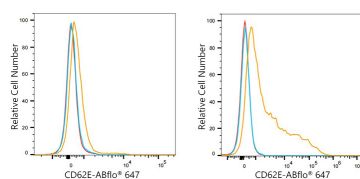
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 5ng per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 30s.



Flow cytometry: 1×10^6 HUVEC cells (negative control, Left) and HUVEC cells (treated with $\text{TNF-}\alpha$, Right) were surface-stained with CD62E/E-Selectin Rabbit mAb (A23836, $2 \mu\text{g/mL}$, orange line) or ABflo® 647 Rabbit IgG isotype control (A22070, $2 \mu\text{g/mL}$, blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb (1:200 dilution) staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry: 1×10^6 HUVEC cells (treated with $\text{TNF-}\alpha$, Right) were surface-stained with ABflo® 647 Rabbit IgG isotype control (A22070, $2 \mu\text{g/mL}$, left) or CD62E/E-Selectin Rabbit mAb (A23836, $2 \mu\text{g/mL}$, right).