

Recombinant Human SLAMF2 (C-Fc-6His)

Catalog No: CB90

Description Recombinant Human SLAM Family Member 2 is produced by our Mammalian expression system and the

target gene encoding Gln27-Ser220 is expressed with a Fc, 6His tag at the C-terminus.

Source Human Cells

Alternative name CD48 antigen; B-lymphocyte activation marker BLAST-1; BCM1 surface antigen; Leukocyte

antigen MEM-102; TCT.1; CD48; BCM1; BLAST1

Accession No. P09326

Predicted Molecular Weight

50.3kDa

AP Molecular Weight

71kDa, reducing conditions.

Formulation Lyophilized from a 0.2 µm filtered solution of PBS, pH 7.4.

Reconstitution

Always centrifuge tubes before opening. Do not mix by vortex or pipetting.

It is not recommended to reconstitute to a concentration less than 100µg/ml.

Dissolve the lyophilized protein in distilled water.

Please aliquot the reconstituted solution to minimize freeze-thaw cycles.

Quality Control Purity: Greater than 95% as determined by reducing SDS-PAGE.

Endotoxin: Less than 0.1 ng/μg (1 IEU/μg) as determined by LAL test.

Shipping The product is shipped at ambient temperature.

Upon receipt, store it immediately at the temperature listed below.

Storage Lyophilized protein should be stored at < -20°C, though stable at room temperature for 3 weeks.

Reconstituted protein solution can be stored at 4-7°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.

Background

CD48 antigen, also known as B-lymphocyte activation marker BLAST-1, BCM1 surface antigen, Leukocyte antigen MEM-102, TCT.1, CD48, BCM1, and BLAST1, CD48 contains one Ig-like C2-type domain and one Ig-like V-type domain, but does not have a transmembrane domain, however, but is held at the cell surface by a GPI anchor via a C-terminal domain which maybe cleaved to yield a soluble form of the receptor. CD48 may facilitate interaction between activated lymphocytes and be involved in regulating T-cell activation. CD48 plays a vital role as an environmental sensor for regulating progenitor cell numbers and inhibiting tumor development. It is suggested that the anti-CD48 mAb has the potential to become an effective therapeutic mAb against multiple myeloma.

SDS-Page

