Catalog # CD3-HP2H7



#### Synonym

CD33,SIGLEC3,gp67

#### Source

PE-Labeled Human Siglec-3 Protein, His Tag (CD3-HP2H7) is produced via conjugation of PE to Human Siglec-3 Protein, His Tag with a new generation site-specific technology under Star Staining labeling platform. Human Siglec-3 Protein, His Tag is expressed from human 293 cells (HEK293). It contains AA Asp 18 - His 259 (Accession # <u>AAH28152.1</u>).

Predicted N-terminus: Asp 18

#### **Molecular Characterization**

# Siglec-3(Asp 18 - His 259) AAH28152.1 Poly-his

This protein carries a polyhistidine tag at the C-terminus.

The protein has a calculated MW of 41.5 kDa.

#### Conjugate

PE

Excitation Wavelength: 488 nm / 561 nm

Emission Wavelength: 575 nm

#### Endotoxin

Less than 1.0 EU per  $\mu$ g by the LAL method.

### Purity

>90% as determined by SDS-PAGE.

#### Formulation

Lyophilized from 0.22  $\mu m$  filtered solution in PBS, 0.2% BSA, pH7.4 with trehalose as protectant.

Contact us for customized product form or formulation.

#### Reconstitution

Please see Certificate of Analysis for specific instructions.

For best performance, we strongly recommend you to follow the reconstitution protocol provided in the CoA.

#### Storage

For long term storage, the product should be stored at lyophilized state at -20°C or lower.

Please protect from light and avoid repeated freeze-thaw cycles.

This product is stable after storage at:

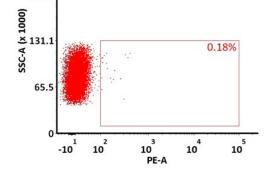
- -20°C to -70°C for 12 months in lyophilized state;
- -70°C for 3 months under sterile conditions after reconstitution.

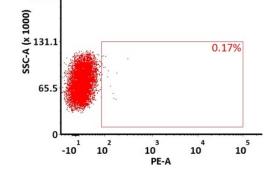
# **Star Staining** fluorescent-labeled products are developed by a new-generation site-specific labeling technology with Star Standard quality at ACROBiosystems

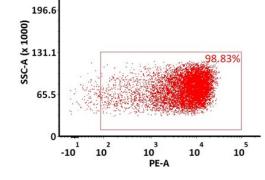
★ Using new-generation site-specific labeling technology ★ High specificity and sensitivity verified by flow cytometry. to maintain natural bioactivity.

 $\star$  No non-specific binding to non-transduced PBMCs.  $\star$  High homogeneity and high batch-to-batch consistency.

Evaluation of CAR expression		
FACS Analysis of Anti-Siglec-3 CAR Expression		
A	В	С
293 cells+PE-Labeled Human Siglec-3 Protein <sup>262.1</sup>	293-CAR cells+Negative control protein 262.1	293-CAR cells+ PE-Labeled Human Siglec-3 Protein
196.6	196.6	









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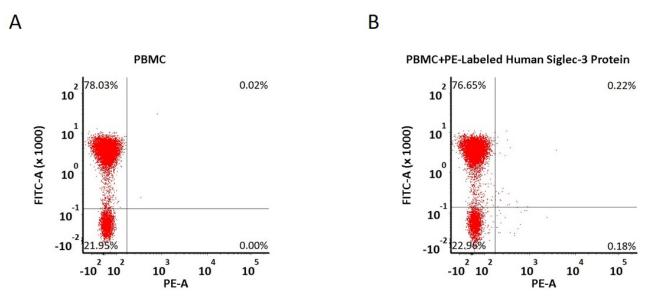
12/11/2024

## PE-Labeled Human Siglec-3 / CD33 Protein, His TagStar Staining



#### Catalog # CD3-HP2H7

5e5 of anti-Siglec-3 CAR-293 cells were stained with 100 μL of 1:50 dilution (2 μL stock solution in 100 μL FACS buffer) of PE-Labeled Human Siglec-3 Protein, His Tag (Cat. No. CD3-HP2H7) and negative control protein respectively (Fig. C and B), and non-transfected 293 cells were used as a control (Fig. A). PE signal was used to evaluate the binding activity (QC tested). FACS Analysis of Non-specific binding to PBMCs



5e5 of PBMCs were stained with PE-Labeled Human Siglec-3 Protein, His Tag (Cat. No. CD3-HP2H7) and anti-CD3 antibody, washed and then analyzed with FACS. FITC signal was used to evaluate the expression of CD3+ T cells in PBMCs, and PE signal was used to evaluate the non-specific binding activity to PBMCs (QC tested).

#### Background

Myeloid cell surface antigen CD33 is also known as SIGLEC3, Siglecs (sialic acid binding Iglike lectins) and GP67, is a single-pass type I membrane protein which belongs to the immunoglobulin superfamily and SIGLEC (sialic acid binding Ig-like lectin) family. Human CD33 / Siglec-3 cD encodes a 364 amino acid (aa) polypeptide with a hydrophobic signal peptide, an N-terminal Ig-like V-type domain, one Ig-like C2-type domains, a transmembrane region and a cytoplasmic tail. CD33 / Siglec-3 usually considered myeloid-specific, but it can also be found on some lymphoid cells. In the immune response, CD33 / Siglec-3 may act as an inhibitory receptor upon ligand induced tyrosine phosphorylation by recruiting cytoplasmic phosphatase(s) via their SH2 domain(s) that block signal transduction through dephosphorylation of signaling molecules. CD33 / Siglec-3 induces apoptosis in acute myeloid leukemia.

#### **Clinical and Translational Updates**



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