

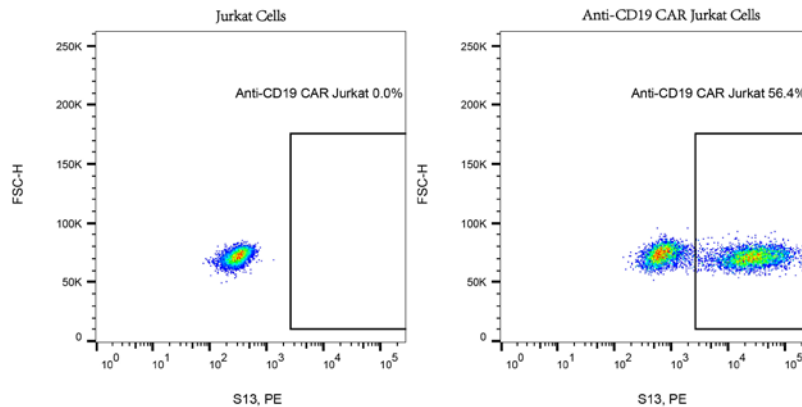
Technical Data Sheet

Rabbit Anti-G4S Linker Monoclonal Antibody, PE

| Product Information | |
|---------------------------|---|
| Product No. | 212306 |
| Size | 100 tests |
| Recommended Vol. per Test | 1 μ L |
| Antibody Types | Monoclonal |
| Antibody Format | Whole IgG |
| Clone | S13 |
| Immunogen | Synthetic 15-mer (G4S) polypeptide |
| Conjugate | PE |
| Excitation/Emission Max | 488/578nm, 561/578nm |
| Host Species | Rabbit |
| Storage Buffer | Aqueous buffered solution containing protein stabilizer and $\leq 0.05\%$ ProClin 300 |
| Storage conditions | 2-8°C, store in dark |

Description

This antibody reacts with cell membrane-expressed CARs of varying specificity containing a G4S linker within the scFv of the extracellular domain. The poly-Glycine-Serine (G4S) linker is a kind of synthetic peptide linker sequence that is flexible and unstructured. It is frequently used to link the variable heavy (VH) domain and the variable light (VL) domain of single-chain variable fragments (scFvs) and chimeric antigen receptors (CARs) that have an extracellular domain scFv for recognizing target antigens. The linker is made up of a core pentapeptide sequence, Gly-Gly-Gly-Gly-Ser, that is repeated and usually seen as either a 15-mer (G4S) or 20-mer (G4S) in scFv-based CARs and scFv fragments.



Flow cytometric analysis of a mixed population containing live wild-type Jurkat cells and Jurkat cells engineered to express an scFv-based Anti-CD19 CAR containing a G4S linker. Jurkat cells were transduced with lentivirus encoding anti-CD19 CAR containing a G4S linker and cultured for 7 days. 2×10^5 cells were stained for the expression of anti-CD19 CAR with Rabbit Anti-G4S Linker Monoclonal Antibody, PE (Product No. 212306, right panel). Non-transduced Jurkat cells were used as a control for gating of CAR expression (left panel).

Preparation & Storage

- Store undiluted at 2-8°C.
- Avoid prolonged exposure to light.
- The monoclonal antibody was purified by Protein A.
- The antibody was conjugated with PE under optimum conditions, and unincorporated dye was removed.

Application Notes

Application

| | |
|----------------|------------------|
| Flow cytometry | Routinely Tested |
|----------------|------------------|

Recommended Antibodies to Include in the Detection Process

| Product name | Product No. |
|--------------------------|---------------|
| Anti-human CD45 Antibody | 602148 |
| Anti-human CD14 Antibody | 602240 |
| Anti-human CD8 Antibody | 602044 |
| Anti-human CD3 Antibody | 603943/604043 |
| Anti-human CD4 Antibody | 604344 |

FACS Protocol

(Optional) For Whole Blood Sample

- Pipette 1 μ L Rabbit Anti-G4S Linker Monoclonal Antibody, PE into the bottom of the tube.
- Add dead cell staining solution and additional fluorochrome-conjugated antibodies into the bottom of the tube.
- Pipette 100 μ L of well-mixed, anticoagulated whole blood into the bottom of the tube. Mix gently and thoroughly.
Note Avoid smearing sample down the side of the tube. If the sample remains on the side of the tube, it will not be stained with the reagents.
- Incubate for 25 minutes in the dark at room temperature (18-25°C).
- Pipette Red Blood Cell Lysis Solution to the tube. Mix gently and thoroughly. Incubate for 15 minutes in the dark at room temperature (18-25°C).
- Add 500 μ L FACS buffer to the tube. Mix well and centrifuge at 300g for 5 minutes at room temperature (18-25°C). Aspirate supernatant completely.
- Repeat step 6 twice.
- Add a suitable amount of FACS buffer to resuspend cell and analysis by flow cytometry.

(Optional) For Cell Sample

- Harvest the cells and wash the cells twice by FACS buffer.
- Count the cells number and the viability.
- Resuspend the cell suspension to a concentration up to 1×10^6 nucleated cells per 100 μ L of buffer.
- Add 1 μ L Rabbit Anti-G4S Linker Monoclonal Antibody, PE, dead cell staining solution and additional fluorochrome. Mix gently and thoroughly.
- Incubate for 25 minutes in the dark at room temperature (18-25°C).
- Add 500 μ L FACS buffer to the tube. Mix well and centrifuge at 300g for 5 minutes at room temperature (18-25°C). Aspirate supernatant completely.
- Repeat step 6 twice.
- Add a suitable amount of FACS buffer to resuspend cell and analysis by flow cytometry.

Product Notices

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Caution: Antibody solutions containing ProClin 300 should be handled with care. Do not take internally and avoid all contact with the skin, mucosa and eyes.

Intellectual Product Notices

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