

Product Data Sheet

Anti-CD8 Antibody-AF647 labled

| Catalog # | Source | e Reactivity | Applications | | | |
|--------------------|---|--|--|--|--|--|
| CFN8554 | Mouse | e H | IF, FC | | | |
| Description | Description Mouse monoclonal antibody AF647 labled to CD8 | | | | | |
| Immunogen | | Native purified human CD8 | | | | |
| Purification | | The antibody was purified by affinity chromatography. | | | | |
| Specificity | | Recognizes human CD8 | | | | |
| Clonality | | Monoclonal (clone: Riv11) | | | | |
| Conjugation | | AF647 | | | | |
| Form | | Mouse IgG1. Liquid in PBS, pH 7.3, 0.2% BSA, and 0.02% sodium azide. | | | | |
| Dilution | | 10 μl / assay | | | | |
| Gene Symbol | | CD8A; CD8B | | | | |
| Alternative Names | | MAL; T-cell surface glycoprotein CD8 alpha chain; T-lymphocyte differentiation | | | | |
| | | antigen T8/Leu-2; CD antig | en CD8a; CD8B1; T-cell surface glycoprotein CD8 beta | | | |
| | | chain; CD antigen CD8b | | | | |
| Entrez Gene | | 925, 926 (Human) | | | | |
| SwissProt | | P01732, P10966 (Human) | | | | |
| Directions for Use | | 1. Take 100 μl peripheral blood anticoagulated by EDTA and add to the bottom of 5 | | | | |
| | | ml tube. | | | | |
| | | 2. Add 10 μ l labeled antibo | dy to the bottom of flow tube mixing with the whole | | | |
| | | blood, incubate for 20 min | utes at room temperature away from light. | | | |
| | | 3. Add 2 ml RBC lysis buffe | r, incubate for 10 minutes away from light after mixing, | | | |
| | | dissolve red blood cells. | | | | |
| | | 4. Sample tube is set to 10 | 00 rpm centrifugation for 5 minutes, discard the | | | |

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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supernatant.

5. Add 2 ml PBS wash buffer to resuspend the cells, then 1000 rpm centrifugation for 5 minutes, discard the supernatant.

6. Add 0.5 ml PBS wash buffer to resuspend the cells and detect by flow cytometry (sample should be determined on the day on the machine and can also be added fixation overnight at 4 °C then measured).

Storage/Stability Shipped and store at 4°C for one year. Do not freeze.

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