Anti-Human CD2 (GA110; Ms.IgG1) Technical Data Sheet



Specificity	CD2	Clone	GA110		
Hybridoma	Mice myeloma cells x Mice spleen cells				
Isotype	IgG ₁	Host	Mouse		
Source and Purification	The antibody is purified from cellular supernatant or ascitic fluid via affinity chromatography (protein A/G); after fluorochrome conjugation, the antibody is purified by means of size exclusion chromatography.				
Storage Buffer	1 ml of PBS pH 7.4 containing 0.5% BSA and 0.1% NaN ₃ .				
Intended use	The antibody is intended for identification and in vitro enumeration of CD2+ cells, according to customer's protocol. Relevant protocols are available upon request, including troubleshooting.				
Main clinical applications	Characterization of T precursor leukaemia and peripheral T-cell lymphomas. For Research Use Only - Not for use in diagnostic procedures				

Stability and storage

- Store at 2-8 °C. Do not freeze!
- Do not expose the reagent to direct light during storage or incubation with cells. In these conditions the product is stable until the expiration date stated on the vial label. Do not use after the expiration date.
- Use a fresh micropipette tip to take the reagent from the vial to preserve its performance characteristics and to avoid contaminations, which can cause erroneous results. Do not use the reagent if it discolours, or if precipitate forms.
- It is recommended to centrifuge before use.
- The pellet formation after centrifugation is a normal event which does not modify the product performances.

Performance characteristics

Specificity

CD2 is a surface antigen of the human T-lymphocyte lineage that is expressed on all peripheral blood T cells. It is one of the earliest T-cell markers, being present on more than 95% of thymocytes; it is also found on some natural killer cells but not on B lymphocytes. Monoclonal antibodies directed against CD2 inhibit the formation of rosettes with sheep erythrocytes, indicating that CD2 is the erythrocyte receptor or is closely associated with it.CD2 functions as an adhesion, between T cells and antigen-presenting cells or NK cells and target cells, and costimulatory molecule.

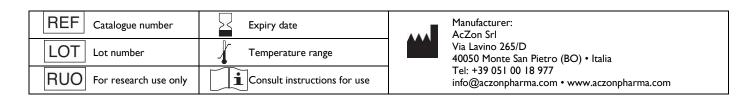
Sensitivity

The antibody sensitivity is defined by the positive CD2 population resolution from the negative CD2 population, obtained by analysing several antibody concentrations on PBMC from healthy donors.

Reproducibility and repeatability

The antibody meets the specifications defined by the Human Leukocyte Differentiation Workshop.

(Bernard AR, Boumsell L, Dausset J, et al. eds. Leucocyte Typing: Human Leucocyte Differentiation Antigens Detected by Monoclonal antibodies. Berlin: *Springer-Verlag*, 1984).



To determine the repeatability of staining with each reagent, samples were stained with different lots of reagents using several samples.

Limits

When analysing samples, it should be considered that the use of monoclonal antibodies in patient treatment can interfere with recognition of target antigens by this reagent. Using pathological specimens (e.g., leukaemia or lymphomas), it is possible to obtain more information with combined reagents rather than single reagents.

Since reagents can be used in different combinations, laboratories need to become familiar with the properties of each antibody in conjunction with other markers in normal and abnormal samples.

Instructions and precautions

The reagent contains sodium azide, a toxic and dangerous compound, and should be handled by trained staff only.

H302 - Harmful if swallowed.

EUH032 - Contact with acids liberates very toxic gas.

P102 - Keep out of reach of children.

P270 – Do not eat, drink or smoke when using this product.

P280 - Wear protective gloves/protective clothing/eye protection/face protection/hearing protection/...

P301+P310 – IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician...

The reagent, the biological specimens and materials coming in contact with them are considered biohazards and handled as if capable of transmitting infections. Dispose in accordance with federal, state and local regulations.

Available packages

Form	Quantity	Code	F/P ratio	Tested Application
Purified concentrated	100 µg/1mL	CD2AMS100H	n.d.	Flow cytometry
Biotin concentrated	100 µg/1mL	CD2FMS100H	4-8	-
FITC ready to use	100 tests/1mL	CD2BMS100H	3-9	Flow cytometry
R-PE ready to use	100 tests/1mL	CD2CMS100H	0.5-1.5	Flow cytometry
APC ready to use	100 tests/1mL	CD2EMS100H	0.5-1.5	Flow cytometry
PerCP ready to use	100 tests/1mL	CD2GMS100H	1-2	Flow cytometry
TDR5 ready to use	100 tests/1mL	CD2DMS100H	2-4	Flow cytometry
TDP55 ready to use	100 tests/1mL	CD2HMS100H	2-4	Flow cytometry
TDR7 ready to use	100 tests/1mL	CD2IMS100H	0.5-1.5	Flow cytometry
TDA7 ready to use	100 tests/1mL	CD2JMS100H	1-2	Flow cytometry

I. For ready to use formats,10 µl are sufficient to label 0.1-1x106 leukocytes (or PBMCs) in 100 µl.

II. See label for lot-specific concentration values.

- TDR5 is a R-PE-Cy5 substitute dye.
- TDP55 is a PerCP-Cv5.5 substitute dve.
- TDR7 is a R-PE-Cy7 substitute dye.
- TDA7 is an APC-Cy7 substitute dye.

Only for professional use • MSDS and protocols available on request

References

Espagnolle N, Depoil D, Zaru R, Demeur C, Champagne E, Guiraud M, Valitutti S, *Int Immunol. 2007*, 19:239-48 Crawford K, Stark A, Kitchens B, Sternheim K, Pantazopoulos V, Triantafellow E, Wang Z, Vasir B, Larsen CE, Gabuzda D, Reinherz E, Alper CA, *Blood 2003*, 102(5):1745-52

Beyers AD, Barclay AN, Law DA, He Q, Williams AF, Immunol Rev. 1989, 111:59-77 Ocklind G, *J Immunol Methods* 1988, 112(2):169-72