

Product datasheet

Anti-TCR V beta 5.3 antibody [W112] ab171099

[3 Images](#)

Overview

Product name	Anti-TCR V beta 5.3 antibody [W112]
Description	Mouse monoclonal [W112] to TCR V beta 5.3
Host species	Mouse
Tested applications	Suitable for: Flow Cyt
Species reactivity	Reacts with: Human
Immunogen	Full length protein corresponding to Human TCR V beta 5.3. Native protein Database link: P04435

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As

Properties

Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term. Avoid freeze / thaw cycle.
Storage buffer	Preservative: 0.1% Sodium azide Constituents: 0.5% BSA, 99% PBS
Purity	Protein A purified
Clonality	Monoclonal
Clone number	W112
Isotype	IgG1

Applications

The Abpromise guarantee Our [Abpromise guarantee](#) covers the use of ab171099 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt		Use at an assay dependent concentration. 1- 2 µg/Test ab170190 - Mouse monoclonal IgG1 , is suitable for use as an isotype control with this antibody.

Target

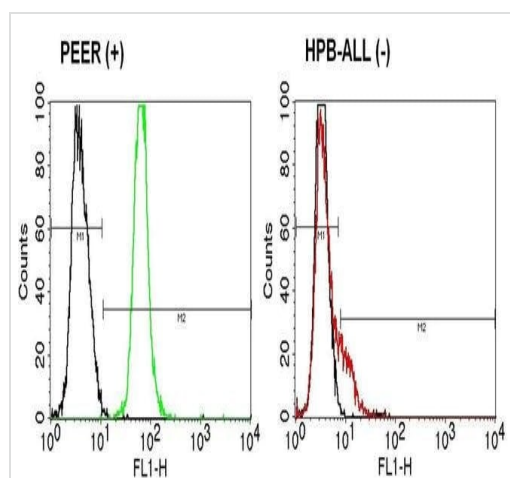
Relevance

T cell receptors (TCR) recognize foreign antigens which have been processed as small peptides and bound to major histocompatibility complex (MHC) molecules at the surface of antigen presenting cells (APC). Each T cell receptor is a dimer consisting of one alpha and one beta chain or one delta and one gamma chain. This region represents the germline organization of the T cell receptor beta locus. The beta locus includes V (variable), J (joining), diversity (D), and C (constant) segments. During T cell development, the beta chain is synthesized by a recombination event at the DNA level joining a D segment with a J segment; a V segment is then joined to the D-J gene. The C segment is later joined by splicing at the RNA level.

Cellular localization

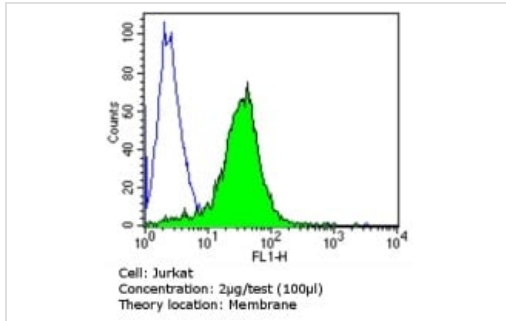
Plasma membrane

Images



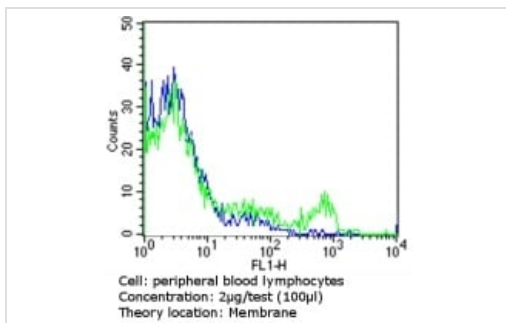
Flow cytometric analysis of TCR V beta 5.3 in positive PEER cells (left) stained with 5 uL ab171099 and FITC labelled secondary. Compared to negative control: HPB-ALL cells (right). A representative 10,000 cells were acquired for each sample.

Flow Cytometry - Anti-TCR V beta 5.3 antibody
[W112] (ab171099)



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Flow cytometry analysis of TCR V beta 5b in Jurkat cells (green) compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of $1-5 \times 10^6$ cells/ml, fixed with 2% paraformaldehyde and washed with PBS. Cells were blocked with a 2% solution of BSA-PBS for 30 min at room temperature and incubated with a TCR V beta 5b monoclonal antibody (ab171099) at a dilution of 2 µg/test for 60 min at room temperature. Cells were then incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated goat anti-mouse IgG (H+L) secondary antibody and re-suspended in PBS for FACS analysis.



Flow Cytometry - Anti-TCR V beta 5.3 antibody
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Flow cytometry analysis of TCR V beta 5b in PBMC cells (green) compared to an isotype control (blue). Human blood was collected, combined with a hydrophilic polysaccharide, centrifuged, transferred to a conical tube and washed with PBS. 50 µl of cell solution was added to each tube at a dilution of 2×10^7 cells/ml, followed by the addition of 50 µl of isotype control and primary antibody (ab171099) at a dilution of 2 µg/test. Cells were incubated for 30 min at 4°C and washed with a cell buffer, followed by incubation with a DyLight 488-conjugated goat anti-mouse IgG (H+L) secondary for 30 min at 4°C in the dark. FACS analysis was performed using 400 µl of cell buffer.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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