abcam

Product datasheet

PerCP Anti-HLA-DR antibody [MEM-12] ab239298

1 Image

Overview

Product name	PerCP Anti-HLA-DR antibody [MEM-12]	
Description	PerCP Mouse monoclonal [MEM-12] to HLA-DR	
Host species	Mouse	
Conjugation	PerCP. Ex: 482nm, Em: 675nm	
Specificity	ab239298 recognizes a common epitope on human HLA-DR which is dependent on the association of alpha and beta chains.	
Tested applications	Suitable for: Flow Cyt	
Species reactivity	Reacts with: Human	
Immunogen	Tissue, cells or virus corresponding to Human HLA-DR. (Thymocyte membrane).	
Positive control	Flow Cyt: Human peripheral blood cells.	
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. Store In the Dark.	
Storage buffer	pH: 7.4 Preservative: 0.0975% Sodium azide Constituent: PBS	
Purity	Size exclusion	
Purification notes	Purified antibody is conjugated with Peridinin-chlorophyll-protein complex (PerCP) under optimum conditions. The conjugate is purified by size-exclusion chromatography and adjusted for direct use. No reconstitution is necessary.	
Clonality	Monoclonal	

Clone number	MEM-12
lsotype	lgG1

Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab239298 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 10µl for 10^6 cells. (or 100 µl of whole blood)

Target

Function

Binds peptides derived from antigens that access the endocytic route of antigen presenting cells (APC) and presents them on the cell surface for recognition by the CD4 T-cells. The peptide binding cleft accommodates peptides of 10-30 residues. The peptides presented by MHC class II molecules are generated mostly by degradation of proteins that access the endocytic route, where they are processed by lysosomal proteases and other hydrolases. Exogenous antigens that have been endocytosed by the APC are thus readily available for presentation via MHC II molecules, and for this reason this antigen presentation pathway is usually referred to as exogenous. As membrane proteins on their way to degradation in lysosomes as part of their normal turn-over are also contained in the endosomal/lysosomal compartments, exogenous antigens must compete with those derived from endogenous components. Autophagy is also a source of endogenous peptides, autophagosomes constitutively fuse with MHC class II loading compartments. In addition to APCs, other cells of the gastrointestinal tract, such as epithelial cells, express MHC class II molecules and CD74 and act as APCs, which is an unusual trait of the GI tract. To produce a MHC class II molecule that presents an antigen, three MHC class II molecules (heterodimers of an alpha and a beta chain) associate with a CD74 trimer in the ER to form an heterononamer. Soon after the entry of this complex into the endosomal/lysosomal system where antigen processing occurs, CD74 undergoes a sequential degradation by various proteases, including CTSS and CTSL, leaving a small fragment termed CLIP (class-II-associated invariant chain peptide). The removal of CLIP is facilitated by HLA-DM via direct binding to the alpha-beta-CLIP complex so that CLIP is released. HLA-DM stabilizes MHC class II molecules until primary high affinity antigenic peptides are bound. The MHC II molecule bound to a peptide is then transported to the cell membrane surface. In B-cells, the interaction between HLA-DM and MHC class II molecules is regulated by HLA-DO. Primary dendritic cells (DCs) also to express HLA-DO. Lysosomal miroenvironment has been implicated in the regulation of antigen loading into MHC II molecules, increased acidification produces increased proteolysis and efficient peptide loading.

Sequence similarities

Post-translational modifications

Cellular localization

Cell membrane. Endoplasmic reticulum membrane. Golgi apparatus > trans-Golgi network membrane. Endosome membrane. Lysosome membrane. Late endosome membrane. The MHC class II complex transits through a number of intracellular compartments in the endocytic pathway

Ubiquitinated by MARCH1 or MARCH8 at Lys-244 leading to down-regulation of MHC class II.

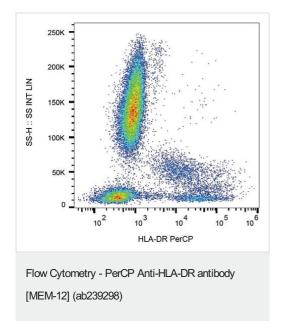
When associated with ubiquitination of the beta subunit of HLA-DR: HLA-DRB4 'Lys-254', the

Belongs to the MHC class II family.

Contains 1 lg-like C1-type (immunoglobulin-like) domain.

down-regulation of MHC class II may be highly effective.

Images



Flow cytometric analysis of human peripheral blood cells labeling HLA-DR with ab239298. Surface staining. Gated on leukocytes.

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