

**HLA-DQA2 & HLA-DQA1 Antibody (C-Term)**  
**Peptide-affinity purified goat antibody**  
**Catalog # AF3351a****Specification**

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**HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Product Information**

Application	WB
Primary Accession	<a href="#">P01906</a>
Other Accession	<a href="#">NP_064440.1</a> , <a href="#">NP_002113.2</a> , <a href="#">3118</a> , <a href="#">3117</a>
Reactivity	Human
Host	Goat
Clonality	Polyclonal
Concentration	0.5 mg/ml
Isotype	IgG
Calculated MW	28033

**HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Additional Information****Gene ID** 3118**Other Names**

HLA class II histocompatibility antigen, DQ alpha 2 chain, DX alpha chain, HLA class II histocompatibility antigen, DQ(6) alpha chain, HLA-DQA1, MHC class II DQA2, HLA-DQA2, HLA-DXA

**Format**

0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

HLA-DQA2 & HLA-DQA1 Antibody (C-Term) is for research use only and not for use in diagnostic or therapeutic procedures.

**HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Protein Information****Name** HLA-DQA2**Synonyms** HLA-DXA**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 a 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 microenvironment has been implicated in the regulation of antigen loading into MHC II molecules, increased acidification produces increased proteolysis and efficient peptide loading.

#### **Cellular Location**

Cell membrane; Single-pass type I membrane protein Endoplasmic reticulum membrane; Single-pass type I membrane protein. Golgi apparatus, trans-Golgi network membrane; Single-pass type I membrane protein Endosome membrane; Single-pass type I membrane protein. Lysosome membrane; Single-pass type I membrane protein. Note=The MHC class II complex transits through a number of intracellular compartments in the endocytic pathway until it reaches the cell membrane for antigen presentation

#### **Tissue Location**

Restricted to skin Langerhans cells, although some expression at low levels may occur at the surface of B lymphoblastoid cells.

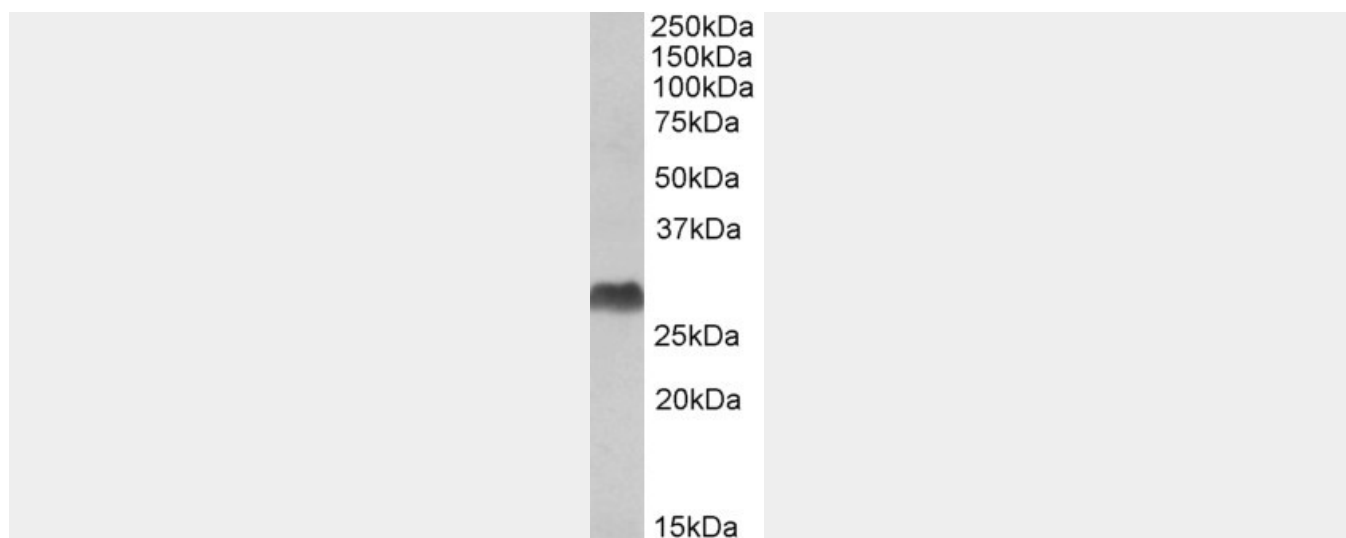
### **HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### **HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Images**





AF3351a (1 µg/ml) staining of Human Bone Marrow lysate (35 µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

#### **HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - Background**

This antibody is expected to recognize: NP\_064440.1 (HLA-DQA2; GeneID: 3118) and also the very similar NP\_002113.2 (HLA-DQA1; GeneID:3117).

#### **HLA-DQA2 & HLA-DQA1 Antibody (C-Term) - References**

New genetic associations detected in a host response study to hepatitis B vaccine. Davila S, Froeling FE, Tan A, Bonnard C, Boland GJ, Snippe H, Hibberd ML, Seielstad M, Genes and immunity 2010 Apr 11 (3): 232-8. PMID: 20237496