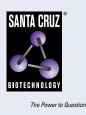
## SANTA CRUZ BIOTECHNOLOGY, INC.

# HLA-DR/DP (HL-40): sc-51617



## BACKGROUND

Major histocompatibility complex (MHC) class II molecules destined for presentation to CD4+ helper T cells is determined by two key events. These events include the dissociation of class II-associated invariant chain peptides (CLIP) from an antigen binding groove in MHC II- $\alpha/\beta$  dimers through the activity of MHC molecules HLA-DM and -DO, and subsequent peptide antigen binding. Accumulating in endosomal/lysosomal compartments and on the surface of B cells, HLA-DM, -DO molecules regulate the dissociation of CLIP and the subsequent binding of exogenous peptides to HLA class II molecules (HLA-DR, -DQ and -DP) by sustaining a conformation that favors peptide exchange. RFLP analysis of HLA-DM genes from rheumatoid arthritis (RA) patients suggests that certain polymorphisms are genetic factors for RA susceptibility. HLA-B belongs to the HLA class I heavy chain paralogs. Class I molecules play a central role in the immune system by presenting peptides derived from the endoplasmic reticulum lumen. HLA-B and -C can form heterodimers consisting of a membrane-anchored heavy chain and a light chain (β-2-Microglobulin). Polymorphisms yield hundreds of HLA-B and -C alleles.

## REFERENCES

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- Momburg, F., et al. 1987. B cell lymphomas of high-grade malignancy frequently lack HLA-DR, -DP and -DQ antigens and associated invariant chain. Int. J. Cancer 40: 598-603.
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- 4. Siegmund, T., et al. 1999. HLA-DMA and HLA-DMB alleles in German patients with type 1 diabetes mellitus. Tissue Antigens 54: 291-294.
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### SOURCE

HLA-DR/DP (HL-40) is a mouse monoclonal antibody raised against Burkitt's lymphoma cell line Raji of human origin.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

## PRODUCT

Each vial contains 100  $\mu g~lg G_{2a}$  in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

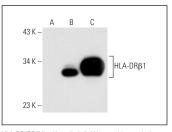
### **APPLICATIONS**

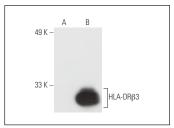
HLA-DR/DP (HL-40) is recommended for detection of a common epitope on the  $\beta$  chain of HLA-DR and HLA-DP of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)] and flow cytometry (1  $\mu$ g per 1 x 10<sup>6</sup> cells).

Molecular Weight of HLA-DR/DP: 30/29 kDa.

Positive Controls: HuT 78 whole cell lysate: sc-2208, BJAB whole cell lysate: sc-2207 or HLA-DR $\beta$ 3 (h): 293T Lysate: sc-110914.

### DATA





HLA-DR/DP (HL-40): sc-51617. Western blot analysis of HLA-DR/1 expression in non-transfected 2931: sc-117752 (**A**), human HLA-DR/1 transfected 2937: sc-115102 (**B**) and BJAB (**C**) whole cell lysates. HLA-DR/DP (HL-40): sc-51617. Western blot analysis of HLA-DR/3D expression in non-transfected: sc-117752 (A) and human HLA-DR/3 transfected: sc-110914 (B) 293T whole cell lysates.

#### SELECT PRODUCT CITATIONS

- Neerincx, A., et al. 2014. The N-terminal domain of NLRC5 confers transcriptional activity for MHC class I and II gene expression. J. Immunol. 193: 3090-3100.
- Burbelo, P.D., et al. 2015. Lack of evidence for molecular mimicry in HIVinfected subjects. PLoS ONE 10: e0127662.
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- Anczurowski, M., et al. 2018. Mechanisms underlying the lack of endogenous processing and CLIP-mediated binding of the invariant chain by HLA-DP<sup>84Gly</sup>. Sci. Rep. 8: 4804.



See **HLA-DR (520B): sc-69673** for HLA-DR antibody conjugates, including AC, HRP, FITC, PE, and Alexa Fluor<sup>®</sup> 488, 546, 594, 647, 680 and 790.