# **Product Datasheet** HLA DRB1 Antibody (LN-3 + HLA-DRB/1067) - Azide and BSA Free NBP2-47672-0.1mg

Unit Size: 0.1 mg

Store at -20 to -80C. Avoid freeze-thaw cycles.

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Updated 10/23/2024 v.20.1

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## NBP2-47672-0.1mg

HLA DRB1 Antibody (LN-3 + HLA-DRB/1067) - Azide and BSA Free

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Product Information		
Unit Size	0.1 mg	
Concentration	1.0 mg/ml	
Storage	Store at -20 to -80C. Avoid freeze-thaw cycles.	
Clonality	Monoclonal	
Clone	LN-3 + HLA-DRB/1067	
Preservative	No Preservative	
Isotype	IgG2b Kappa	
Purity	Protein A or G purified	
Buffer	10 mM PBS	
Product Description		
Description	<ul> <li>1.0 mg/ml of antibody purified from Bioreactor Concentrate by Protein A/G.</li> <li>Prepared in 10mM PBS WITHOUT BSA &amp; azide. Also available at 200 ug/ml WITH BSA &amp; azide (NBP2-45316).</li> <li>Antibody with azide - store at 2 to 8C. Antibody without azide - store at -20 to -80C.</li> </ul>	
Host	Mouse	
Gene ID	3123	
Gene Symbol	HLA-DRB1	
Species	Human, Monkey	
Specificity/Sensitivity	This monoclonal antibody reacts with the beta-chain of HLA-DRB1 antigen, a member of MHC class II molecules. It does not cross react with HLA-DP and HLA-DQ. HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36kDa alpha (heavy) chain and a 28kDa beta (light) chain. It is expressed on B-cells, activated T-cells, monocytes/macrophages, dendritic cells and other non-professional APCs. In conjunction with the CD3/TCR complex and CD4 molecules, HLA-DR is critical for efficient peptide presentation to CD4+ T cells. It is an excellent histiocytic marker in paraffin sections producing intense cytoplasmic staining. True histiocytic neoplasms are similarly positive. HLA-DR antigens also occur on a variety of epithelial cells and their corresponding neoplastic counterparts. Loss of HLA-DR expression is related to tumor microenvironment and predicts adverse outcome in diffuse large B-cell lymphoma.	
Immunogen	Activated human peripheral blood mononuclear cells (LN-3 & HLA-DRB/1067)	
Product Application Details		
Applications	Western Blot, Flow Cytometry, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin, CyTOF-ready, Immunofluorescence	
Recommended Dilutions	Western Blot 0.5 - 1.0 ug/ml, Flow Cytometry 0.5 - 1 ug/million cells in 0.1 ml, Immunohistochemistry, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry-Paraffin 0.25 - 0.5 ug/ml, Immunofluorescence 0.5 - 1.0 ug/ml, CyTOF-ready	

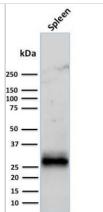


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pplication Notes	Immunohistochemistry (Formalin-fixed) Staining of formalin-fixed tissues require with 1mM EDTA, pH 9.0, for 45 min at 9 minutes. Optimal dilution for a specific applicatio	es heating tissue sections in 10mM Tris 95C followed by cooling at RT for 20
Images		
Western Blot: HLA DRB1 Antiboo BSA Free [NBP2-47672] - Weste using HLA DRB1 Antibody (LN-3		kDa kano 250
+ HLA-DRB/1067) - Azide and BS	Raji cells labeling HLA-DR with HLA 3/1067) conjugated with CF594	
	HLA DRB1 Antibody (LN-3 + HLA- e [NBP2-47672] - Human Tonsil stained ody (LN-3 + HLA-DRB/1067).	
and BSA Free [NBP2-47672] - Fl	oody (LN-3 + HLA-DRB/1067) - Azide ow Cytometric Analysis of Human Raji LN-3 + HLA-DRB/1067). conjugated (Red).	Normalized To Model Mo

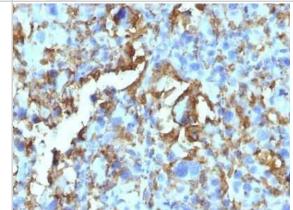
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Western Blot: HLA DRB1 Antibody (LN-3 + HLA-DRB/1067) - Azide and BSA Free [NBP2-47672] - Western Blot Analysis of Human Spleen tissue lysate using HLA DRB1 Antibody (LN-3 + HLA-DRB/1067).



Immunohistochemistry-Paraffin: HLA DRB1 Antibody (LN-3 + HLA-DRB/1067) - Azide and BSA Free [NBP2-47672] - Human Histiocytoma stained with HLA-DR Monoclonal Antibody (LN-3 + HLA-DRB/1067).







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#### Products Related to NBP2-47672-0.1mg

HAF007	Goat anti-Mouse IgG Secondary Antibody [HRP]
NB720-B	Rabbit anti-Mouse IgG (H+L) Secondary Antibody [Biotin]
NBP1-43317-0.5mg	Mouse IgG2b Kappa Light Chain Isotype Control (MG2b)
NBP2-51599-0.1mg	Recombinant Human, HLA DRB1 His Protein

#### Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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