

## HLA-DOα rabbit pAb

## Cat No.:ES2536

For research use only

## Overview

Product Name	HLA-DOα rabbit pAb
Host species	Rabbit
Applications	WB;IHC;IF;ELISA
Species Cross-Reactivity	Human
<b>Recommended dilutions</b>	Western Blot: 1/500 - 1/2000.
	Immunohistochemistry: 1/100 - 1/300. ELISA:
	1/40000. Not yet tested in other applications.
Immunogen	The antiserum was produced against synthesized
	peptide derived from human HLA-DOA. AA
	range:71-120
Specificity	HLA-DOα Polyclonal Antibody detects endogenous
	levels of HLA-DOα protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and
	0.02% sodium azide.
Storage	Store at -20°C. Avoid repeated freeze-thaw cycles.
Protein Name	HLA class II histocompatibility antigen DO alpha
	chain
Gene Name	HLA-DOA
Cellular localization	Endosome membrane; Single-pass type I membrane
	protein. Lysosome membrane; Single-pass type I
	membrane protein. Complexes with HLA-DM
	molecule during intracellular transport and in
	endosomal/lysosomal compartments.
	Heterotetramerization is necessary to ex
Purification	The antibody was affinity-purified from rabbit
	antiserum by affinity-chromatography using
	epitope-specific immunogen.
Clonality	Polyclonal
Concentration	1 mg/ml
Observed band	34kD
Human Gene ID	3111
Human Swiss-Prot Number	P06340
Alternative Names	HLA-DOA; HLA-DNA; HLA-DZA; HLA class II



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Background

COLO205

(kD)

117-85-

48-

34-

26-

19-

histocompatibility antigen; DO alpha chain; MHC DN-alpha; MHC DZ alpha; MHC class II antigen DOA HLA-DOA belongs to the HLA class II alpha chain paralogues. HLA-DOA forms a heterodimer with HLA-DOB. The heterodimer, HLA-DO, is found in lysosomes in B cells and regulates HLA-DM-mediated peptide loading on MHC class II molecules. In comparison with classical HLA class II molecules, this gene exhibits very little sequence variation, especially at the protein level. [provided by RefSeq, Jul 2008],

Western Blot analysis of various cells using HLA-DO $\alpha$  Polyclonal Antibody



Immunohistochemical analysis of paraffin-embedded Human breast cancer. Antibody was diluted at 1:100(4° overnight). High-pressure and temperature Tris-EDTA,pH8.0 was used for antigen retrieval. Negetive contrl (right) obtaned from antibody was pre-absor



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Western blot analysis of lysates from COLO cells, using HLA-DOA Antibody. The lane on the right is blocked with the synthesized peptide.



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