

## **Product Data Sheet**

## **Anti-OLR1 Antibody**

Catalog #	Source	e Reactivity	Applications	
CQA2124	Rabbit	Н	WB, IH	
Description		Rabbit polyclonal antibody	to OLR1	
Immunogen		Recombinant full length pro	otein of human OLR1	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous lev	els of OLR1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	) - 1/200)	
Gene Symbol		OLR1		
Alternative Names		CLEC8A; LOX1; Oxidized low-density lipoprotein receptor 1; Ox-LDL receptor 1;		
		C-type lectin domain family	8 member A; Lectin-like oxidized LDL receptor 1; LOX-1;	
		Lectin-like oxLDL receptor 1	; hLOX-1; Lectin-type oxidized LDL receptor 1	
Entrez Gene		4973 (Human)		
SwissProt		P78380 (Human)		
Storage/Stabi	lity	Shipped at 4 $^{\circ}$ C. Upon deli	very aliquot and store at -20 $^\circ$ C for one year. Avoid	
		freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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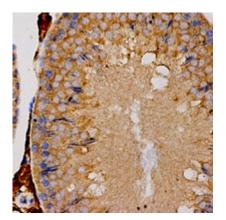
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For research purposes only, not for human use

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Western blot analysis of OLR1 expression in Hela (A), HepG2 (B) whole cell lysates. (Predicted band size: 20; 21; 30 kD; Observed band size: 30 kD)



Immunohistochemical analysis of OLR1 staining in mouse testis formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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