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Datasheet for ABIN6263820 anti-OLR1 antibody (N-Term)

3 Images



Overview

| Quantity: | 100 µL |
|----------------------|---|
| Target: | OLR1 |
| Binding Specificity: | N-Term |
| Reactivity: | Human, Mouse, Rat |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This OLR1 antibody is un-conjugated |
| Application: | Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF) |

Product Details

| Immunogen: | A synthesized peptide derived from human OLR1, corresponding to a region within N-terminal amino acids. |
|---------------|--|
| Isotype: | lgG |
| Specificity: | OLR1 Antibody detects endogenous levels of total OLR1. |
| Purification: | The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific). |

Target Details

| Target: | OLR1 |
|-------------------|----------------------|
| Alternative Name: | OLR1 (OLR1 Products) |

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Target Details

| Background: | Description: Receptor that mediates the recognition, internalization and degradation of |
|-------------------|--|
| | oxidatively modified low density lipoprotein (oxLDL) by vascular endothelial cells. OxLDL is a |
| | marker of atherosclerosis that induces vascular endothelial cell activation and dysfunction, |
| | resulting in pro-inflammatory responses, pro-oxidative conditions and apoptosis. Its association |
| | with oxLDL induces the activation of NF-kappa-B through an increased production of |
| | intracellular reactive oxygen and a variety of pro-atherogenic cellular responses including a |
| | reduction of nitric oxide (NO) release, monocyte adhesion and apoptosis. In addition to binding |
| | oxLDL, it acts as a receptor for the HSP70 protein involved in antigen cross-presentation to |
| | naive T-cells in dendritic cells, thereby participating in cell-mediated antigen cross-presentation. |
| | Also involved in inflammatory process, by acting as a leukocyte-adhesion molecule at the |
| | vascular interface in endotoxin-induced inflammation. Also acts as a receptor for advanced |
| | glycation end (AGE) products, activated platelets, monocytes, apoptotic cells and both Gram- |
| | negative and Gram-positive bacteria. |
| | Gene: OLR1 |
| Molecular Weight: | 31kDa |
| Gene ID: | 4973 |

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|----------|--------|
| UniProt: | P78380 |

Application Details

| Application Notes: | WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000 |
|--------------------|---|
| Restrictions: | For Research Use only |

Handling

| Format: | Liquid |
|--------------------|--|
| Concentration: | 1 mg/mL |
| Buffer: | Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol. |
| Preservative: | Sodium azide |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| Storage: | -20 °C |

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Handling

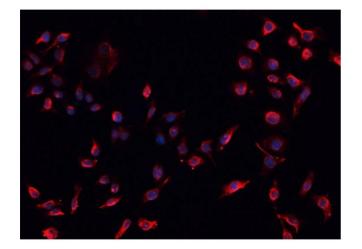
Storage Comment:

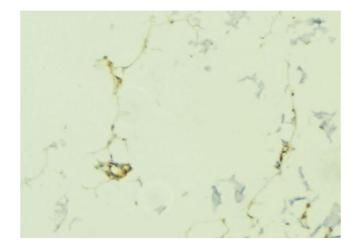
Store at -20 °C. Stable for 12 months from date of receipt.

Expiry Date:

12 months

Images



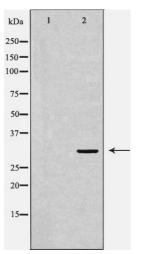


Immunofluorescence (fixed cells)

Image 1. ABIN6276785 staining HepG2? cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25jãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37jãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod

Immunohistochemistry

Image 2. ABIN6276785 at 1/100 staining Human lung tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22_jãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary



Western Blotting

Image 3. Western blot analysis of Jurkat whole cell lysates, using OLR1 Antibody. The lane on the left is treated with the antigen-specific peptide.

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