

## **Anti-RAGE Antibody [Clone 16-1]**

Catalog Number	LDG0039YA
Package	100 μg / 200 μg / Customized package

For full product information, images and publications, please visit our website.



## **Overview**

### Description

The immunoglobulin superfamily is a group of transmembrane proteins, contains a variable number of disulfide-stabilized motifs similar to immunoglobulins, associated with the adhesion, binding and recognition processes of cells. Receptor for advanced glycation endproducts (RAGE), a member of the immunoglobulin superfamily, not only binds to advanced glycation endproducts but also interacts with multiple ligands having common motifs. These interactions play important roles in various cellular processes, and associated with many human diseases.

#### **Product Note**

Recommended dilution factor:

ELISA: 1:5000-20000 WB: 1:1000-10000 IFA: 1:200-1000

FACS: Assay dependent

Note: Working dilution for specific application should be determined by the investigator to obtain the best conditions.

# **Specifications** Host Clonality Mouse Monoclonal **Clone Name** Isotype lgG1 clone 16-1 **Immunogen** Reactivity **RAGE** Human

Tainan Headquarter

**Innovation & Research Center** 

**CLD Center** 



**Application** 

ELISA, WB, IFA, FACS

Concentration

1 mg/mL

**Specificity** 

**RAGE** 

Conjugation

Unconjugated

**Buffer** 

Phosphate Buffered Saline containing 0.03% ProClin 300, pH 7.4.

**Form** 

Liquid

## Instruction

### **Shipping**

The product is shipped with polar packs. Upon receipt, store it immediately at -20°C or lower for long term storage.

### Stability & Storage

This product is stable after storage at:

- 2-8°C for 2 weeks under sterile conditions from date of receipt.
- -20°C or -80°C for 12 months under sterile conditions from date of receipt.

Avoid repeated freeze/thaw cycles.

Suggestion: Divide antibody into several vials. Keep only vials for usage at 2-8°C.

### **Image**





Immunofluorescence analysis of Anti-RAGE Antibody [Clone 16-1]
HEK293T cells were transfected with RAGE plasmid and fixed in 4% PFA, permeabilized with PBS containing 0.1% Triton X-100. Cells were stained with mouse anti-RAGE monoclonal antibody (1:200) followed by secondary antibodies (goat anti-Mouse IgG-iFluor 488, 1:200, green) and cell nuclei were stained with Hoechst 33342 (Blue).

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