# IRS2 Rabbit mAb

Catalog No.: A25714 Recombinant



# **Basic Information**

#### **Observed MW**

170kDa

### **Calculated MW**

137kDa

### Category

Primary antibody

### **Applications**

ELISA,WB,IF/ICC

#### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC65786

# **Background**

This gene encodes the insulin receptor substrate 2, a cytoplasmic signaling molecule that mediates effects of insulin, insulin-like growth factor 1, and other cytokines by acting as a molecular adaptor between diverse receptor tyrosine kinases and downstream effectors. The product of this gene is phosphorylated by the insulin receptor tyrosine kinase upon receptor stimulation, as well as by an interleukin 4 receptor-associated kinase in response to IL4 treatment.

# **Recommended Dilutions**

**WB** 1:1000 - 1:5000

**IF/ICC** 1:50 - 1:200

# **Immunogen Information**

**Gene ID**Swiss Prot
8660
Q9Y4H2

#### **Immunogen**

A synthetic peptide corresponding to a sequence within amino acids 1239-1338 of human IRS2 (NP $\_003740.2$ ).

## **Synonyms**

IRS-2; Irs2; IRS 2;

## **Contact**

<b>a</b>		400-999-6126
$\bowtie$		cn.market@abclonal.com.cn
$\overline{a}$	ı	www.ahclonal.com.cn

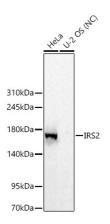
# **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

### **Storage**

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.



Western blot analysis of various lysates using IRS2 Rabbit mAb (A25714)at 1:3000 dilution incubated overnight at 4°C.

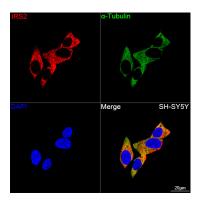
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L)(AS014) at 1:10000 dilution.

Lysates/proteins: 25  $\mu g$  per lane.

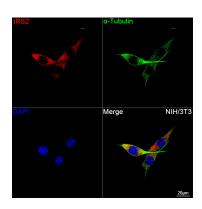
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020) Negative control (NC): U-2 OS

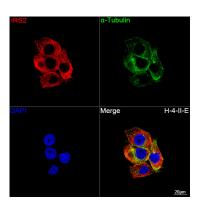
Exposure time: 5 s.



Confocal immunofluorescence analysis of SH-SY5Y cells using IRS2 Rabbit mAb (A25714, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with Alpha-tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal immunofluorescence analysis of NIH/3T3 cells using IRS2 Rabbit mAb (A25714, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with Alpha-tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal immunofluorescence analysis of H-4-II-E cells using IRS2 Rabbit mAb (A25714, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with Alpha-tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.