

#### **UK Office**

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Research Use Only. Not for diagnostic or therapeutic use.

### EB11754 - Goat Anti-ICAM1 (aa313-327) Antibody

Size: 100µg specific antibody in 200µl

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#### **Target Protein**

**Principal Names:** BB2, CD54, cell surface glycoprotein P3.58, human rhinovirus receptor, ICAM1, ICAM-1, intercellular adhesion molecule 1, intercellular adhesion molecule 1 (CD54), human rhinovirus receptor, major group rhinovirus receptor, P3.58 **Official Symbol:** ICAM1

Accession Number(s): NP\_000192.2

#### Human GeneID(s): 3383

**Important Comments:** The immunizing peptide represents part of the extracellular domain, and it does not overlap any known glycosite..

#### Immunogen

Peptide with sequence C-PNVILTKPEVSEGTE, from the internal region of the protein sequence according to NP\_000192.2.

Please note the peptide is available for sale.

#### **Purification and Storage**

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

#### **Applications Tested**

Peptide ELISA: antibody detection limit dilution 1:128000.

**Western blot:** Approx 100kDa band observed in Human Lung lysates (calculated MW of 57.8kDa according to Human NP\_000192.2). This molecular weight is routinely observed by other sources. Recommended concentration: 0.1-0.3µg/ml. Primary incubation 1 hour at room temperature.

**Immunofluorescence:** Strong expression of the protein seen in the cytoplasm of A431 cells Recommended concentration: 10µg/ml.

**Flow Cytometry:** Flow cytometric analysis of Jurkat cells and Human peripheral blood monocytes. Recommended concentration: 10ug/ml.

#### **Species Reactivity**

Tested: Human Expected from sequence similarity: Human

## EB11754 (0.3µg/ml) staining of Human Lung lysate (35µg protein in RIPA buffer). Detected by chemiluminescence.

EB11754 Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB11754 Flow cytometric analysis of paraformaldehyde fixed Jurkat cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.

EB11754 Flow cytometric analysis of paraformaldehyde fixed human peripheral blood monocytes (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.