

#### **International Office**

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Research Use Only. Not for diagnostic or therapeutic use.

## EB09588 - Goat Anti-LIGHT / CD258 Antibody

Size: 100µg specific antibody in 200µl



## **Target Protein**

**Principal Names:** TNFSF14, tumor necrosis factor (ligand) superfamily, member 14, CD258, HVEML, LIGHT, LTg, TR2, delta transmembrane LIGHT, herpesvirus entry mediator A, ligand for herpesvirus entry mediator, tumor necrosis factor ligand superfamily, member 14, tumor necrosis factor receptor-like 2, tumor necrosis factor superfamily member LIGHT

Official Symbol: TNFSF14

Accession Number(s): NP\_003798.2; NP\_742011.1

Human GenelD(s): 8740

Important Comments: This antibody is expected to recognize both reported isoforms

(NP\_003798.2; NP\_742011.1).

## **Immunogen**

Peptide with sequence C-HEVNPAAHLTGANS, from the internal region of the protein sequence according to NP\_003798.2; NP\_742011.1.

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 1:64000.

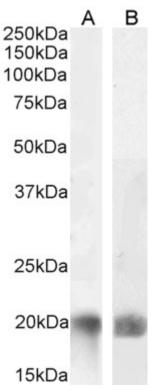
**Western blot:** Approx 20-22kDa band observed in Human Cerebellum and Frontal Cortex Brain lysates (calculated MW of 22.4kDa according to NP\_742011.1) Recommended concentration: 0.03-0.1µg/ml. Primary incubation 1 hour at room temperature.

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

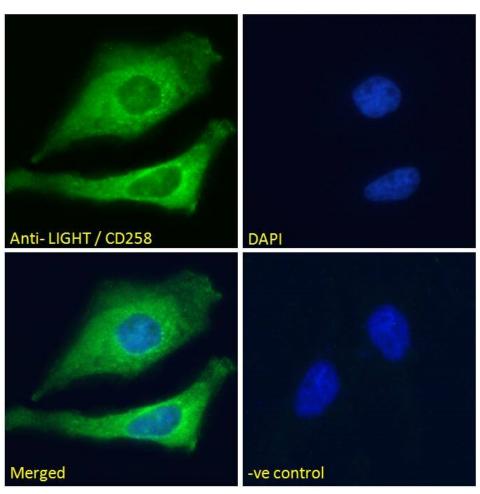
# **Species Reactivity**

Tested: Human

Expected from sequence similarity: Human



EB09588 (0.03μg/ml) staining of Human Cerebellum (A) and Frontal Cortex (B) lysate (35μg protein in RIPA buffer). Detected by chemiluminescence.



EB09588 Immunofluorescence analysis of paraformaldehyde fixed HeLa cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic

and membrane staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).