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Research Use Only. Not for diagnostic or therapeutic use.

EB07113 - Goat Anti-TIM3 / HAVCR2 Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: HAVCR2, TIM3, hepatitis A virus cellular receptor 2, FLJ14428, KIM-3, TIMD3, Tim-3, T cell immunoglobulin mucin 3, T cell immunoglobulin mucin-3, kidney

injury molecule-3

Official Symbol: HAVCR2

Accession Number(s): NP_116171.3

Human GeneID(s): 84868

Immunogen

Peptide with sequence C-KWYSHSKEKIQN, from the internal region of the protein sequence according to NP_116171.3.

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 35kDa band observed in lysates of cell lines A549 and HepG2 and in preliminary testing of Jurkat and U937 cell lysate (calculated MW of 33.4kDa according to NP_116171.3). Recommended concentration:0.1-1μg/ml. Primary incubation 1 hour at room temperature.

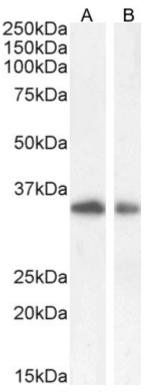
Immunofluorescence: Strong expression of the protein seen in the nucleus and plasma membrane of HepG2 cells. Recommended concentration: 10µg/ml.

Flow Cytometry: Flow cytometric analysis of HepG2 cells. Recommended concentration: 10ug/ml.

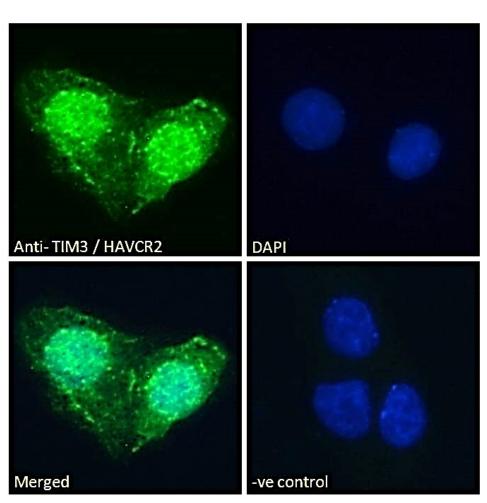
Species Reactivity

Tested: Human

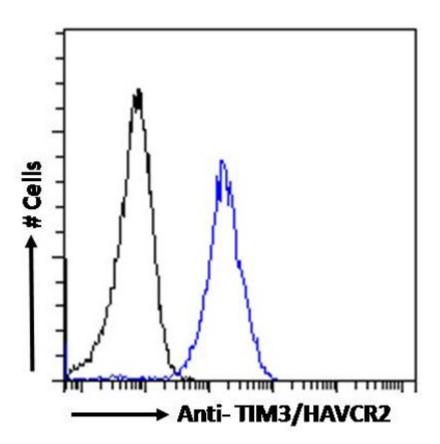
Expected from sequence similarity: Human, Dog



EB07113 (0.1μg/ml) staining of A549 (A) and (0.5ug/ml) HepG2 (B) cell lysate (35μg protein in RIPA buffer). Detected by chemiluminescence.



EB07113 Immunofluorescence analysis of paraformaldehyde fixed HepG2 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing



EB07113 Flow cytometric analysis of paraformaldehyde fixed HepG2 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.