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Research Use Only. Not for diagnostic or therapeutic use.

EB05999 - Goat Anti-FOXA1 / HNF3A Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: FOXA1, HNF3A, forkhead box A1, TCF3A, MGC33105, hepatocyte

nuclear factor 3, alpha

Official Symbol: FOXA1

Accession Number(s): NP_004487.2

Human GenelD(s): 3169

Non-Human GenelD(s): 15375 (mouse), 25098 (rat)

Immunogen

 $\label{eq:continuous} \mbox{Peptide with sequence C-GVYSRPVLNTS, from the C Terminus of the protein sequence}$

according to NP_004487.2.

Please note the <u>peptide</u> 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 50kDa band observed in nuclear lysates of cell line HepG2 and approx 50-55kDa in Mouse Liver and in nuclear lysates of cell line MCF7 calculated MW of 49.1kDa according to Human NP_004487.2 and 48.9kDa according to Mouse NP_032285.2). These bands were successfully blocked by incubation with the immunizing peptide. Recommended concentration: 0.1-0.3μg/ml. Primary incubation 1 hour at room temperature.

Immunofluorescence: Strong expression of the protein seen in the nuclei of MCF7 and U2OS cells. Recommended concentration: 10µg/ml.

Flow Cytometry: Flow cytometric analysis of MCF7 cells. Recommended concentration: 10ug/ml.

Species Reactivity

Tested: Human, Mouse

Expected from sequence similarity: Human, Mouse, Rat, Pig, Cow

EB05999 (0.1μg/ml) staining of HepG2 (A) and (0.3ug/ml) MCF7 (B) nuclear cell lysate (35μg protein in RIPA buffer). Detected by chemiluminescence.

EB05999 (0.1μg/ml) staining of Mouse Liver lysate (35μg protein in RIPA buffer). Detected by chemiluminescence.

EB05999 Immunofluorescence analysis of paraformaldehyde fixed MCF7 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing strong nuclear staining. Actin filaments were stained with phalloidin (red) and the nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB05999 Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing strong nuclear staining. Actin filaments were stained with phalloidin (red) and the nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB05999 Flow cytometric analysis of paraformaldehyde fixed MCF7 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.