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**Research Use Only. Not for
diagnostic or therapeutic use.**

EB06868 - Goat Anti-SDHB Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: SDHB, IP, SDH, SDH1, SDHIP, succinate dehydrogenase complex, subunit B, iron sulfur (Ip), OTTHUMP00000044624, iron-sulfur subunit, FLJ92337, SDH2, CWS2, PGL4

Official Symbol: SDHB

Accession Number(s): NP_002991.2

Human GeneID(s): [6390](#)

Immunogen

Peptide with sequence C-ATYKEKKASV, from the C Terminus of the protein sequence according to NP_002991.2.

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 1:8000.

Western blot: Approx 28-30kDa band observed in lysates of cell lines HEK293, Jurkat and HeLa (calculated MW of 31.6kDa according to NP_002991.2). Recommended concentration: 0.1-0.3µg/ml. Primary incubation 1 hour at room temperature.

IHC: Paraffin embedded Human Kidney. Recommended concentration: 6-8µg/ml.

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

Flow Cytometry: Flow cytometric analysis of A431 cells. Recommended concentration: 10ug/ml.

Species Reactivity

Tested: Human

Expected from sequence similarity: Human

EB06868 (0.1µg/ml) staining of HEK293 (A), Jurkat (B) and HeLa (C) cell lysate (35µg protein in RIPA buffer).
Detected by chemiluminescence.

EB06868 (6µg/ml) staining of paraffin embedded Human Kidney. Heat induced antigen retrieval with citrate
buffer pH 6, HRP-staining.

EB06868 Negative Control showing staining of paraffin embedded Human Kidney, with no primary antibody.

EB06868 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 plasma
membrane and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG
(10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB06868 Flow cytometric analysis of paraformaldehyde fixed A431 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.