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

EB09109 - Goat Anti-GAD1 (isoform GAD67) Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: GAD1, glutamate decarboxylase 1 (brain, 67kDa), FLJ45882, GAD, SCP, OTTHUMP00000041055, glutamate decarboxylase 1, glutamate decarboxylase 1 (brain, 67kD), GAD67, GAD25

Official Symbol: GAD1

Accession Number(s): NP_000808.2

Human GeneID(s): [2571](#)

Non-Human GeneID(s): 14415 (mouse), 24379 (rat)

Important Comments: This antibody is expected to recognize isoform GAD67. There is no cross-reactivity expected with GAD2.

Immunogen

Peptide with sequence C-PDSPQRREKLHK, from the internal region of the protein sequence according to NP_000808.2.

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 70kDa band observed in Human Cerebellum, and in Mouse and Rat Brain lysates (calculated MW of 66.9kDa according to Human NP_000808.2, and 66.6kDa according to Mouse NP_032103.2 and Rat NP_058703.1). Recommended concentration: 0.3-1µg/ml. Primary incubation 1hour at room temperature.

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

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

Species Reactivity

Tested: Human, Mouse, Rat

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

EB09109 (1µg/ml) staining of Human Cerebellum (A) and Mouse Brain (B) and (0.3ug/ml) Rat Brain (C) lysate (35µg protein in RIPA buffer). Detected by chemiluminescence.

EB09109 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 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).

EB09109 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.