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

EB11003 - Goat Anti-DGAT1 (aa67-79) Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: DGAT1, diacylglycerol O-acyltransferase 1, ARGP1, DGAT, ACAT related gene product 1, ACAT-related gene product 1, acyl coenzyme A:cholesterol acyltransferase related gene 1, diacylglycerol O-acyltransferase homolog 1, diglyceride acyltransferase

Official Symbol: DGAT1

Accession Number(s): NP_036211.2

Human GeneID(s): [8694](#)

Immunogen

Peptide with sequence RCHRLQDSLFSDD, from the internal region of the protein sequence according to NP_036211.2.

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. 55-58kDa band observed in lysates of cell line NIH3T3, and approx. 58kDa in Mouse Duodenum and Rat Kidney lysates (calculated MW of 56.8kDa according to Mouse NP_034176.1 and 57.1kDa according to Rat NP_445889.2).
Recommended concentration: 0.3-1µg/ml. Primary incubation 1 hour at room temperature.

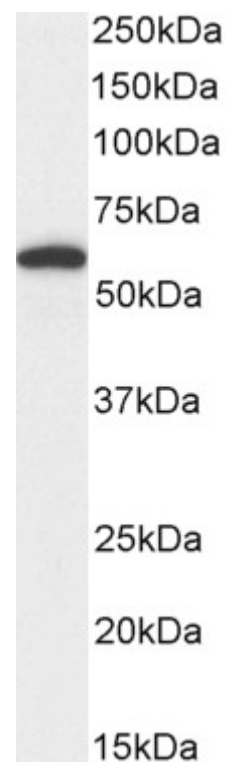
Immunofluorescence: Strong expression of the protein seen in A431 and NIH3T3 cells.
Recommended concentration: 10µg/ml.

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

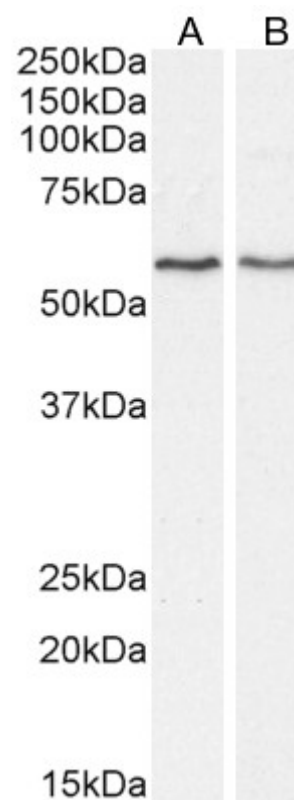
Species Reactivity

Tested: Human, Mouse, Rat

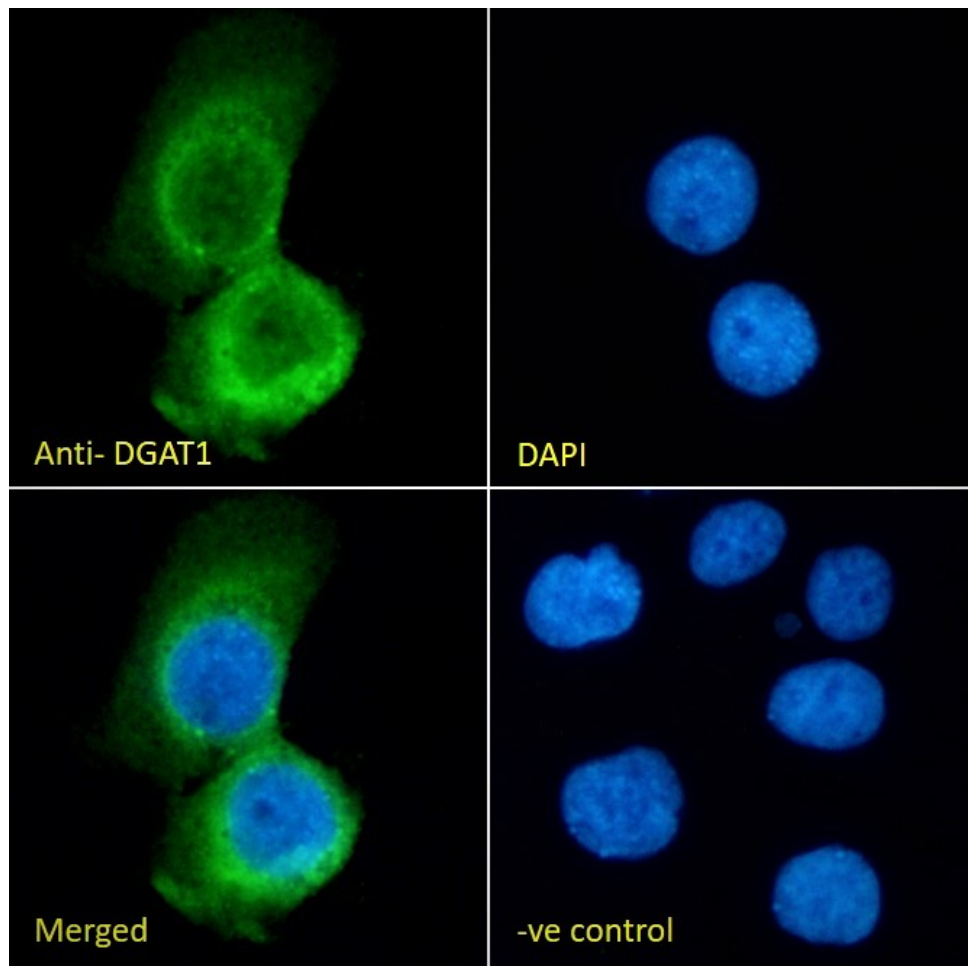
Expected from sequence similarity: Human, Mouse, Rat, Cow



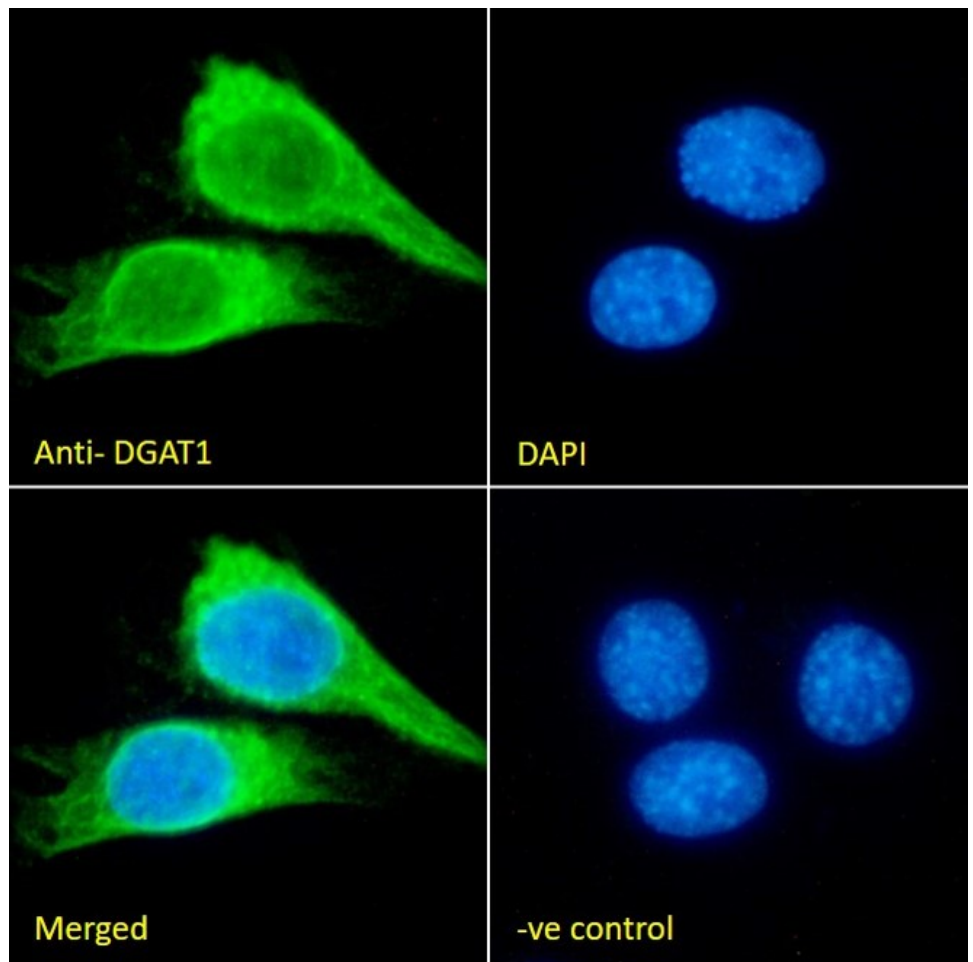
EB11003 (0.5 μ g/ml) staining of NIH3T3 cell lysate (35 μ g protein in RIPA buffer). Detected by chemiluminescence.



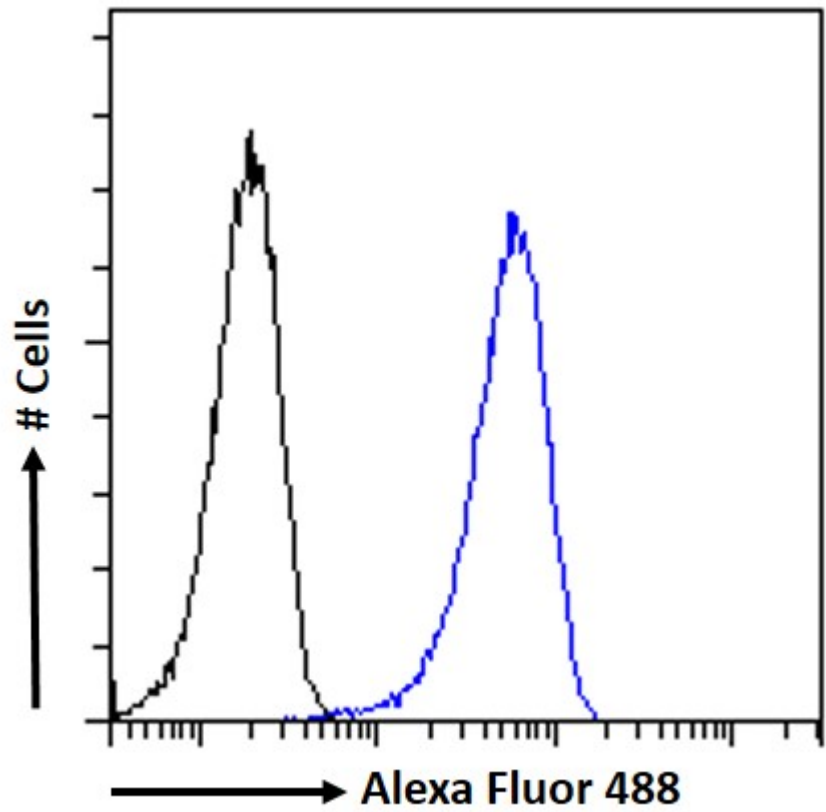
EB11003 (1 μ g/ml) staining of Mouse Duodenum (A) and (0.5 μ g/ml) Rat Kidney (B) lysate (35 μ g protein in RIPA buffer). Detected by chemiluminescence.



EB11003 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 endoplasmic reticulum 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).



EB11003 Immunofluorescence analysis of paraformaldehyde fixed NIH3T3 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



EB11003 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.