FLAD1 (G-4): sc-376819



The Power to Question

BACKGROUND

FLAD1 (FAD1 flavin adenine dinucleotide synthetase), also known as FAD1, FADS, PP591 or molybdenum cofactor biosynthesis protein-like, is a 587 amino acid protein where its N-terminus belongs to the moaB/mog family and its C-terminus belongs to the PAPS reductase family. Existing as five alternatively spliced isoforms, FLAD1 localizes to the cytoplasm and utilizes magnesium as a cofactor. FLAD1 is a key enzyme in the metabolic pathway that converts riboflavin into the redox cofactor flavin adenine dinucleotide (FAD). It is suggested that the molybdenum cofactor biosynthesis protein-like region of FLAD1 may not be functional. FLAD1 is encoded by a gene located on human chromosome 1, which spans 260 million base pairs, contains over 3,000 genes and comprises nearly 8% of the human genome. Aberrations in chromosome 1 are found in a variety of cancers, including head and neck cancer, malignant melanoma and multiple myeloma.

REFERENCES

- Wu, M., et al. 1995. Cloning and characterization of FAD1, the structural gene for flavin adenine dinucleotide synthetase of *Saccharomyces* cerevisiae. Mol. Cell. Biol. 15: 264-271.
- Barile, M., et al. 2000. The riboflavin/FAD cycle in rat liver mitochondria.
 Eur. J. Biochem. 267: 4888-4900.
- Brizio, C., et al. 2006. Over-expression in *Escherichia coli* and characterization of two recombinant isoforms of human FAD synthetase. Biochem. Biophys. Res. Commun. 344: 1008-1016.

CHROMOSOMAL LOCATION

Genetic locus: FLAD1 (human) mapping to 1q21.3; Flad1 (mouse) mapping to 3 F1.

SOURCE

FLAD1 (G-4) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 157-195 within an internal region of FLAD1 of human origin.

PRODUCT

Each vial contains 200 $\mu g \; lg G_1$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

FLAD1 (G-4) is available conjugated to agarose (sc-376819 AC), 500 μ g/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-376819 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-376819 PE), fluorescein (sc-376819 FITC), Alexa Fluor® 488 (sc-376819 AF488), Alexa Fluor® 546 (sc-376819 AF546), Alexa Fluor® 594 (sc-376819 AF594) or Alexa Fluor® 647 (sc-376819 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-376819 AF680) or Alexa Fluor® 790 (sc-376819 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-376819 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

APPLICATIONS

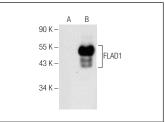
FLAD1 (G-4) is recommended for detection of FLAD1 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

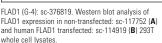
Suitable for use as control antibody for FLAD1 siRNA (h): sc-88309, FLAD1 siRNA (m): sc-145197, FLAD1 shRNA Plasmid (h): sc-88309-SH, FLAD1 shRNA Plasmid (m): sc-145197-SH, FLAD1 shRNA (h) Lentiviral Particles: sc-88309-V and FLAD1 shRNA (m) Lentiviral Particles: sc-145197-V.

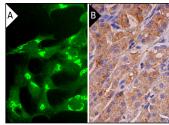
Molecular Weight of FLAD1: 63 kDa.

Positive Controls: FLAD1 (h): 293T Lysate: sc-114919, K-562 whole cell lysate: sc-2203 or MIA PaCa-2 cell lysate: sc-2285.

DATA







FLAD1 (G-4): sc-376819. Immunofluorescence staining of formalin-fixed Hep G2 cells showing cytoplasmic localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human upper stomach tissue showing cytoplasmic staining of glandular cells (B).

SELECT PRODUCT CITATIONS

- 1. Marceau, C.D., et al. 2016. Genetic dissection of Flaviviridae host factors through genome-scale CRISPR screens. Nature 535: 159-163.
- Genc, A.M., et al. 2020. Elimination of a retinal riboflavin binding protein exacerbates degeneration in a model of cone-rod dystrophy. Invest. Ophthalmol. Vis. Sci. 61: 17.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.