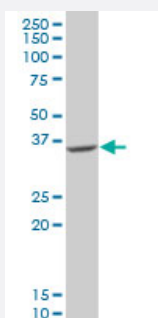


GDAP1 polyclonal antibody (A01)

Catalog # H00054332-A01

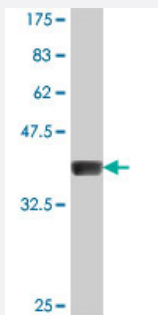
Size 50 uL

Applications



Western Blot (Cell lysate)

GDAP1 polyclonal antibody (A01), Lot # 060613JCS1 Western Blot analysis of GDAP1 expression in HepG2 (Cat # L019V1).



Western Blot detection against Immunogen (37.11 kDa) .

Specification

Product Description	Mouse polyclonal antibody raised against a partial recombinant GDAP1.
Immunogen	GDAP1 (NP_061845, 158 a.a. ~ 257 a.a) partial recombinant protein with GST tag.
Sequence	TRIRSQIGNTESELKKLAEENPDLQEAYIAKQKRLKSKLLDHDNVKYLKKILDELEKVLQVETELQ RRNEETPEEGQQPWLCGESFTLADVSLAVTLHR
Host	Mouse
Reactivity	Human
Interspecies Antigen Sequence	Mouse (96); Rat (95)

Quality Control Testing

Antibody Reactive Against Recombinant Protein.
Western Blot detection against Immunogen (37.11 KDa) .

Storage Buffer

50 % glycerol

Storage Instruction

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Cell lysate)

GDAP1 polyclonal antibody (A01), Lot # 060613JCS1 Western Blot analysis of GDAP1 expression in HepG2 (Cat # L019V1).

[Protocol Download](#)

- Western Blot (Recombinant protein)

[Protocol Download](#)

- ELISA

Gene Info — GDAP1

Entrez GeneID

[54332](#)

GeneBank Accession#

[NM_018972](#)

Protein Accession#

[NP_061845](#)

Gene Name

GDAP1

Gene Alias

-

Gene Description

ganglioside-induced differentiation-associated protein 1

Omim ID

[214400](#) [606598](#) [607706](#) [607831](#)

Gene Ontology

[Hyperlink](#)

Gene Summary

This gene encodes a member of the ganglioside-induced differentiation-associated protein family , which may play a role in a signal transduction pathway during neuronal development. Mutations in this gene have been associated with various forms of Charcot-Marie-Tooth Disease and neuropathy. Two transcript variants encoding different isoforms have been identified for this gene. [provided by RefSeq]

Other Designations

-

Publication Reference

- [Effective therapeutic strategies in a pre-clinical mouse model of Charcot-Marie-tooth disease.](#)

Cristina Nuevo-Tapióles, Fulvio Santacatterina, Brenda Sánchez-Garrido, Cristina Núñez Arenas, Adrián Robledo-Bérgamo, Paula Martínez-Valero, Lara Cantarero, Beatriz Pardo, Janet Hoenicka, Michael P Murphy, Jorgina Satrustegui, Francesc Palau, José M Cuezva.

Human Molecular Genetics 2021 Nov; 30(24):2441.

Application: WB-Tr, Human, SH-SY5Y cells

- [AMPK activation negatively regulates GDAP1, which influences metabolic processes and circadian gene expression in skeletal muscle.](#)

Lassiter DG, Sjögren RJO, Gabriel BM, Krook A, Zierath JR.

Molecular Metabolism 2018 Oct; 16:12.

Application: WB-Ce, Human, Skeletal muscle cells

- [CMT-linked loss-of-function mutations in GDAP1 impair store-operated Ca²⁺ entry-stimulated respiration.](#)

González-Sánchez P, Pla-Martin D, Martínez-Valero P, Rueda CB, Calpena E, Del Arco A, Palau F, Satrustegui J.

Scientific Reports 2017 Feb; 7:42993.

Application: WB, Human, HEK 293T cells

- [Silencing of the Charcot-Marie-Tooth disease-associated gene GDAP1 induces abnormal mitochondrial distribution and affects Ca²⁺ homeostasis by reducing store-operated Ca²⁺ entry.](#)

Pla-Martin D, Rueda CB, Estela A, Sanchez-Piris M, Gonzalez-Sanchez P, Traba J, de la Fuente S, Scorrano L, Renau-Piqueras J, Alvarez J, Satrustegui J, Palau F.

Neurobiology of Disease 2013 Jul; 55:140.

Application: WB-Ti, Mouse, Brain

- [Charcot-Marie-Tooth-related Gene GDAP1 Complements Cell Cycle Delay at G2/M Phase in Saccharomyces cerevisiae fis1 Gene-defective Cells.](#)

Estela A, Pla-Martin D, Sanchez-Piris M, Sesaki H, Palau F.

The Journal of Biological Chemistry 2011 Oct; 286(42):36777.

Application: WB-Tr, Yeast, Yeast cells

- [Mitochondrial complex I deficiency in GDAP1-related autosomal dominant Charcot-Marie-Tooth disease \(CMT2K\).](#)

Cassereau J, Chevrollier A, Gueguen N, Malinge MC, Letournel F, Nicolas G, Richard L, Ferre M, Verny C, Dubas F, Procaccio V, Amati-Bonneau P, Bonneau D, Reynier P.

Neurogenetics 2008 Dec; 10(2):145.

Application: WB, Human, Human skin fibroblasts

Disease

- [Kidney Failure](#)