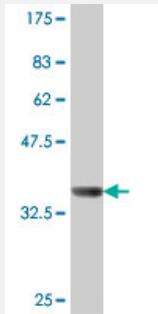


IDS polyclonal antibody (A01)

Catalog # H00003423-A01

Size 50 uL

Applications



Western Blot detection against Immunogen (37.11 KDa) .

Specification

Product Description	Mouse polyclonal antibody raised against a partial recombinant IDS.
Immunogen	IDS (NP_000193, 38 a.a. ~ 137 a.a) partial recombinant protein with GST tag.
Sequence	NVLLIVDDLRLPSLGCYGDKLVRSPNIDQLASHLLFQNAFAQQAVCAPSRVSFLTGRRPDTTRL YDFNSYWRVHAGNFSTIPQYFKENGYVTMSVGKVF
Host	Mouse
Reactivity	Human
Interspecies Antigen Sequence	Mouse (97)
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 (Recombinant protein)

[Protocol Download](#)

- ELISA

Gene Info — IDS

Entrez GeneID [3423](#)

GeneBank Accession# [NM_000202](#)

Protein Accession# [NP_000193](#)

Gene Name IDS

Gene Alias MPS2, SIDS

Gene Description iduronate 2-sulfatase

Omim ID [309900](#)

Gene Ontology [Hyperlink](#)

Gene Summary Iduronate-2-sulfatase is required for the lysosomal degradation of heparan sulfate and dermatan sulfate. Mutations in this X-chromosome gene that result in enzymatic deficiency lead to the sex-linked Mucopolysaccharidosis Type II, also known as Hunter Syndrome. Iduronate-2-sulfatase has a strong sequence homology with human arylsulfatases A, B, and C, and human glucosamine-6-sulfatase. A splice variant of this gene has been described. [provided by RefSeq]

Other Designations Hunter syndrome|OTTHUMP00000024207|OTTHUMP00000024210|alpha-L-iduronate sulfate sulfatase|iduronate-2-sulfatase|idursulfase

Publication Reference

- [In Vitro recapitulation of the site-specific editing \(to Wild-Type\) of mutant IDS mRNA transcripts, and characterization of IDS protein translated from the edited mRNAs.](#)

Lualdi S, Zotto GD, Zegarra-Moran O, Pedemonte N, Corsolini F, Bruschi M, Tomati V, Amico G, Candiano G, Dardis A, Cooper DN, Filocamo M.

Human Mutation 2017 Jul; 38(7):849.

Application: WB, IP, Human, Lymphoblasts from Hunter syndrome patients

Pathway

- [Glycosaminoglycan degradation](#)
- [Lysosome](#)
- [Metabolic pathways](#)

Disease

- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)