

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product datasheet for PH319187

Iduronate 2 sulfatase (IDS) (NM_000202) Human Mass Spec Standard

Product data:

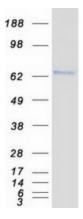
Product Type:	Mass Spec Standards
Description:	IDS MS Standard C13 and N15-labeled recombinant protein (NP_000193)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC219187
Predicted MW:	61.87 kDa
Protein Sequence:	<pre>>RC219187 representing NM_000202 Red=Cloning site Green=Tags(s)</pre>
	MPPPRTGRGLLWLGLVLSSVCVALGSETQANSTTDALNVLLIIVDDLRPSLGCYGDKLVRSPNIDQLASH SLLFQNAFAQQAVCAPSRVSFLTGRRPDTTRLYDFNSYWRVHAGNFSTIPQYFKENGYVTMSVGKVFHPG ISSNHTDDSPYSWSFPPYHPSSEKYENTKTCRGPDGELHANLLCPVDVLDVPEGTLPDKQSTEQAIQLLE KMKTSASPFFLAVGYHKPHIPFRYPKEFQKLYPLENITLAPDPEVPDGLPPVAYNPWMDIRQREDVQALN ISVPYGPIPVDFQRKIRQSYFASVSYLDTQVGRLLSALDDLQLANSTIIAFTSDHGWALGEHGEWAKYSN FDVATHVPLIFYVPGRTASLPEAGEKLFPYLDPFDSASQLMEPGRQSMDLVELVSLFPTLAGLAGLQVPP RCPVPSFHVELCREGKNLLKHFRFRDLEEDPYLPGNPRELIAYSQYPRPSDIPQWNSDKPSLKDIKIMGY SIRTIDYRYTVWVGFNPDEFLANFSDIHAGELYFVDSDPLQDHNMYNDSQGGDLFQLLMP
Tag:	TRTRPLEQKLISEEDLAANDILDYKDDDDKV C-Myc/DDK
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Concentration:	$>0.05 \mu g/\mu L$ as determined by microplate BCA method
Labeling Method:	Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3
Storage:	Store at -80°C. Avoid repeated freeze-thaw cycles.
Stability:	Stable for 3 months from receipt of products under proper storage and handling conditions.
RefSeq:	<u>NP 000193</u>
RefSeq Size:	2504
RefSeq ORF:	1650



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	lduronate 2 sulfatase (IDS) (NM_000202) Human Mass Spec Standard – PH319187
Synonyms:	ID2S; MPS2; SIDS
Locus ID:	3423
UniProt ID:	<u>P22304</u>
Cytogenetics:	Xq28
Summary:	This gene encodes a member of the sulfatase family of proteins. The encoded preproprotein is proteolytically processed to generate two polypeptide chains. This enzyme is involved in the lysosomal degradation of heparan sulfate and dermatan sulfate. Mutations in this gene are associated with the X-linked lysosomal storage disease mucopolysaccharidosis type II, also known as Hunter syndrome. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed. [provided by RefSeq, Jan 2016]
Protein Families:	Druggable Genome
Protein Pathways	: Glycosaminoglycan degradation, Lysosome, Metabolic pathways

Product images:



Coomassie blue staining of purified IDS protein (Cat# [TP319187]). The protein was produced from HEK293T cells transfected with IDS cDNA clone (Cat# [RC219187]) using MegaTran 2.0 (Cat# [TT210002]).

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