

## ARG82699 Mouse IDS / Iduronate 2 Sulfatase ELISA Kit

Package: 96 wells Store at: 4°C

## Component

Cat. No.	Component Name	Package	Temp
ARG82699-001	Antibody-coated microplate	8 X 12 strips	4°C. Unused strips should be sealed tightly in the air-tight pouch.
ARG82699-002	Standard	2 X 10 ng/vial	4°C
ARG82699-003	Standard/Sample diluent	30 ml (Ready to use)	4°C
ARG82699-004	Antibody conjugate concentrate (100X)	1 vial (100 μl)	4°C
ARG82699-005	Antibody diluent buffer	12 ml (Ready to use)	4°C
ARG82699-006	HRP-Streptavidin concentrate (100X)	1 vial (100 μl)	4°C
ARG82699-007	HRP-Streptavidin diluent buffer	12 ml (Ready to use)	4°C
ARG82699-008	25X Wash buffer	20 ml	4°C
ARG82699-009	TMB substrate	10 ml (Ready to use)	4°C (Protect from light)
ARG82699-010	STOP solution	10 ml (Ready to use)	4°C
ARG82699-011	Plate sealer	4 strips	Room temperature

### Summary

Product Description	ARG82699 Mouse IDS / Iduronate 2 Sulfatase ELISA Kit is an Enzyme Immunoassay kit for the quantification of Mouse IDS / Iduronate 2 Sulfatase in serum, plasma (EDTA, heparin) and cell culture supernatants.	
Tested Reactivity	Ms	
Tested Application	ELISA	
Target Name	IDS / Iduronate 2 Sulfatase	
Conjugation	HRP	
Conjugation Note	Substrate: TMB and read at 450 nm.	
Sensitivity	78 pg/ml	
Sample Type	Serum, plasma (EDTA, heparin) and cell culture supernatants.	
Standard Range	156 - 10000 pg/ml	
Sample Volume	100 μΙ	
Precision	Intra-Assay CV: 5.4%	

~ 5 hours

#### **Application Instructions**

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/ (JJU)	

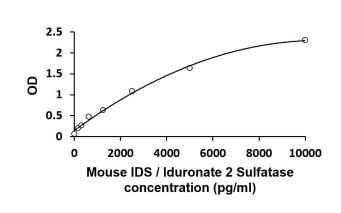
#### **Properties**

Form	96 well
Storage instruction	Store the kit at 2-8°C. Keep microplate wells sealed in a dry bag with desiccants. Do not expose test reagents to heat, sun or strong light during storage and usage. Please refer to the product user manual for detail temperatures of the components.
Note	For laboratory research only, not for drug, diagnostic or other use.

### **Bioinformation**

Gene Symbol	IDS
Gene Full Name	iduronate 2-sulfatase
Background	The protein encoded by this gene belongs to the sulfatase family, is localized to the lysosome, and is involved in 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. Alternatively spliced transcript variants have been described for this gene. [provided by RefSeq, Aug 2013]
Function	Required for the lysosomal degradation of heparan sulfate and dermatan sulfate. [UniProt]
Highlight	Related products: <u>Iduronate 2 Sulfatase antibodies;</u> <u>Iduronate 2 Sulfatase ELISA Kits;</u> New ELISA data calculation tool: <u>Simplify the ELISA analysis by GainData</u>
РТМ	The conversion to 3-oxoalanine (also known as C-formylglycine, FGly), of a serine or cysteine residue in prokaryotes and of a cysteine residue in eukaryotes, is critical for catalytic activity. [UniProt]
Cellular Localization	Lysosome. [UniProt]

#### Images



# ARG82699 Mouse IDS / Iduronate 2 Sulfatase ELISA Kit standard curve image

ARG82699 Mouse IDS / Iduronate 2 Sulfatase ELISA Kit results of a typical standard run with optical density reading at 450 nm.