

# **Product datasheet for PH311132**

## OriGene Technologies, Inc.

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### GLUD1 (NM\_005271) Human Mass Spec Standard

**Product data:** 

**Product Type:** Mass Spec Standards

**Description:** GLUD1 MS Standard C13 and N15-labeled recombinant protein (NP\_005262)

Species:HumanExpression Host:HEK293

Expression cDNA Clone

RC211132

or AA Sequence:

**Protein Sequence:** 

Predicted MW: 61.4 kDa

>RC211132 representing NM\_005271 Red=Cloning site Green=Tags(s)

MYRYLGEALLLSRAGPAALGSASADSAALLGWARGQPAAAPQPGLALAARRHYSEAVADREDDPNFFKMV EGFFDRGASIVEDKLVEDLRTRESEEQKRNRVRGILRIIKPCNHVLSLSFPIRRDDGSWEVIEGYRAQHS QHRTPCKGGIRYSTDVSVDEVKALASLMTYKCAVVDVPFGGAKAGVKINPKNYTDNELEKITRRFTMELA KKGFIGPGIDVPAPDMSTGEREMSWIADTYASTIGHYDINAHACVTGKPISQGGIHGRISATGRGVFHGI ENFINEASYMSILGMTPGFGDKTFVVQGFGNVGLHSMRYLHRFGAKCIAVGESDGSIWNPDGIDPKELED FKLQHGSILGFPKAKPYEGSILEADCDILIPAASEKQLTKSNAPRVKAKIIAEGANGPTTPEADKIFLER NIMVIPDLYLNAGGVTVSYFEWLKNLNHVSYGRLTFKYERDSNYHLLMSVQESLERKFGKHGGTIPIVPT AEFQDRISGASEKDIVHSGLAYTMERSARQIMRTAMKYNLGLDLRTAAYVNAIEKVFKVYNEAGVTFT

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag: C-Myc/DDK

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Concentration:** >0.05 μg/μL 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:** NP 005262

RefSeq Size: 3051 RefSeq ORF: 1674



#### GLUD1 (NM\_005271) Human Mass Spec Standard - PH311132

Synonyms: GDH; GDH1; GLUD

**Locus ID:** 2746

**UniProt ID:** <u>P00367</u>, <u>E9KL48</u>

Cytogenetics: 10q23.2

**Summary:** This gene encodes glutamate dehydrogenase, which is a mitochondrial matrix enzyme that

catalyzes the oxidative deamination of glutamate to alpha-ketoglutarate and ammonia. This

enzyme has an important role in regulating amino acid-induced insulin secretion. It is

allosterically activated by ADP and inhibited by GTP and ATP. Activating mutations in this gene are a common cause of congenital hyperinsulinism. Alternative splicing of this gene results in multiple transcript variants. The related glutamate dehydrogenase 2 gene on the human X-chromosome originated from this gene via retrotransposition and encodes a soluble form of glutamate dehydrogenase. Related pseudogenes have been identified on chromosomes 10,

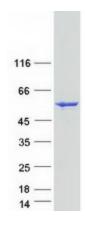
18 and X. [provided by RefSeq, Jan 2016]

**Protein Families:** Druggable Genome

**Protein Pathways:** Alanine, aspartate and glutamate metabolism, Arginine and proline metabolism, D-Glutamine

and D-glutamate metabolism, Metabolic pathways, Nitrogen metabolism

### **Product images:**



Coomassie blue staining of purified GLUD1 protein (Cat# [TP311132]). The protein was produced from HEK293T cells transfected with GLUD1 cDNA clone (Cat# [RC211132]) using

MegaTran 2.0 (Cat# [TT210002]).