

## Product datasheet for PH311422

### Fukutin (FKTN) (NM\_001079802) Human Mass Spec Standard

#### Product data:

Product Type:	Mass Spec Standards
Description:	FKTN MS Standard C13 and N15-labeled recombinant protein (NP_001073270)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC211422
Predicted MW:	53.5 kDa
Protein Sequence:	>RC211422 representing NM_001079802 Red=Cloning site Green=Tags(s)

MSRINKNVVLALLTLTSSAFLLFQLYYYKHYLSTKNGAGLSKSKGSRIGFDSTQWRVAKKFI MLT SNQNV  
PVFLIDPLILEL INKNFEQVKNTSHGSTSQCKFFCVPRDFTAFALQYHLWKNEEGWFRIAENMGFQCLKI  
ESKDPRLDGIDSLSGTEIPLHYICKLATHAIHLVVFHERSGNYLWHGHLRLKEHIDRKFVPPFRKLQFGRY  
PGA FDRPELQQVTVDGLEVLIPKDPMHFVEEVPHSRFIECRYKEARAFFQQYLDNTVEAVAFRKS AKEL  
LQLAAKTLNKLGVPFWLSSTGCLGWYRQCNIIPYSKDVDLGIFIQDYKSDIILAFQDAGLPLKHKFGKVE  
DSLELSFQGDVVKLDVFFYYEETHMWNNGGTQAKTGKFKYLPKFTLCWTEFVDMKVHVP CETLEYIE  
ANYGKTWKIPVKTWDWKRSPPNVQPNGIWP ISEWDEVIQLY

TRTRPLEQKLI SEEDLAANDILDYKDDDDKV

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- <sup>13</sup> C <sub>6</sub> , <sup>15</sup> N <sub>4</sub> ]-L-Arginine and [U- <sup>13</sup> C <sub>6</sub> , <sup>15</sup> N <sub>2</sub> ]-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:	<a href="#">NP_001073270</a>
RefSeq Size:	7456
RefSeq ORF:	1383
Synonyms:	CMD1X; FCMD; LGMD2M; LGMDR13; MDDGA4; MDDGB4; MDDGC4



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Locus ID: 2218

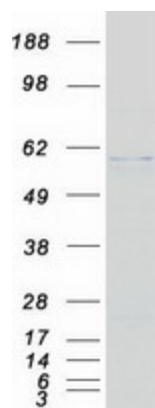
UniProt ID: [O75072](#)

Cytogenetics: 9q31.2

**Summary:** The protein encoded by this gene is a putative transmembrane protein that is localized to the cis-Golgi compartment, where it may be involved in the glycosylation of alpha-dystroglycan in skeletal muscle. The encoded protein is thought to be a glycosyltransferase and could play a role in brain development. Defects in this gene are a cause of Fukuyama-type congenital muscular dystrophy (FCMD), Walker-Warburg syndrome (WWS), limb-girdle muscular dystrophy type 2M (LGMD2M), and dilated cardiomyopathy type 1X (CMD1X). Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Nov 2010]

**Protein Families:** Transmembrane

### Product images:



Coomassie blue staining of purified FKTN protein (Cat# [TP311422]). The protein was produced from HEK293T cells transfected with FKTN cDNA clone (Cat# [RC211422]) using MegaTran 2.0 (Cat# [TT210002]).