

Product datasheet for PH301904

CLN6 (NM_017882) Human Mass Spec Standard

Product data:

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 Type:	Mass Spec Standards
Description:	CLN6 MS Standard C13 and N15-labeled recombinant protein (NP_060352)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC201904
Predicted MW:	35.9 kDa
Protein Sequence:	<pre>>RC201904 protein sequence Red=Cloning site Green=Tags(s)</pre>
	MEATRRRQHLGATGGPGAQLGASFLQARHGSVSADEAARTAPFHLDLWFYFTLQNWVLDFGRPIAMLVFP LEWFPLNKPSVGDYFHMAYNVITPFLLLKLIERSPRTLPRSITYVSIIIFIMGASIHLVGDSVNHRLLFS GYQHHLSVRENPIIKNLKPETLIDSFELLYYYDEYLGHCMWYIPFFLILFMYFSGCFTASKAESLIPGPA LLLVAPSGLYYWYLVTEGQIFILFIFTFFAMLALVLHQKRKRLFLDSNGLFLFSSFALTLLLVALWVAWL WNDPVLRKKYPGVIYVPEPWAFYTLHVSSRH
	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:	<u>NP 060352</u>
RefSeq Size:	2258
RefSeq ORF:	933
Synonyms:	CLN4A; HsT18960; nclf
Locus ID:	54982



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2024 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

CLN6 (NM_017882) Human Mass Spec Standard – PH301904	
UniProt ID:	<u>Q9NWW5, A0A024R601</u>
Cytogenetics:	15q23
Summary:	This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses (NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive, neurodegenerative disorders affecting children. The genes responsible likely encode proteins involved in the degradation of post-translationally modified proteins in lysosomes. The primary defect in NCL disorders is thought to be associated with lysosomal storage function. [provided by RefSeq, Oct 2008]
Protein Families	Transmembrane

Product images:

116	_	
66	_	
45	_	
35	-	
25	_	
18	_	
14	_	

Coomassie blue staining of purified CLN6 protein (Cat# [TP301904]). The protein was produced from HEK293T cells transfected with CLN6 cDNA clone (Cat# [RC201904]) using MegaTran 2.0 (Cat# [TT210002]).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2024 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US