

Product datasheet for **TP301904L**

CLN6 (NM_017882) Human Recombinant Protein

Product data:

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| Product Type: | Recombinant Proteins |
| Description: | Recombinant protein of human ceroid-lipofuscinosis, neuronal 6, late infantile, variant (CLN6), 1 mg |
| Species: | Human |
| Expression Host: | HEK293T |
| Expression cDNA Clone or AA Sequence: | >RC201904 protein sequence Red =Cloning site Green =Tags(s) |
| | <p>MEATRRRQHLGATGGPGAQLGASFLQARHGVSVADEAARTAPFHLDLWFYFTLQNWVLD FGRPIAMLVFP</p> <p>LEWFPLNKPSVGDYFHMAYNVITPFLLLKLIERSPRTLPRSITYVSIIFIMGASIHVGDVSNHRLLS</p> <p>GYQHHLVRENPPIKNLKPETLIDSFELLYYDEYLGHCMWYIPFFLILFMYFSGCFTASKAESLIPGPA</p> <p>LLLVAAPGLYYWLVTEGQIFILFIFTFAMLALVLHQKRKRLFLDSNGLFLFSSFALTL LLLVALWVAVL</p> <p>WNDPVLRRKKYPGVIVPEPWAFYTLHVSSRH</p> <p>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</p> |
| Tag: | C-Myc/DDK |
| Predicted MW: | 35.7 kDa |
| Concentration: | >0.05 µg/µL as determined by microplate BCA method |
| Purity: | > 80% as determined by SDS-PAGE and Coomassie blue staining |
| Buffer: | 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol |
| Preparation: | Recombinant protein was captured through anti-DDK affinity column followed by conventional chromatography steps. |
| Note: | For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process. |
| Storage: | Store at -80°C. |
| Stability: | Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles. |
| RefSeq: | NP_060352 |



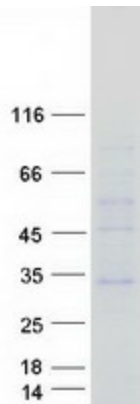
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Locus ID: 54982
UniProt ID: [Q9NWW5](#)
RefSeq Size: 2258
Cytogenetics: 15q23
RefSeq ORF: 933
Synonyms: CLN4A; HsT18960; nclf

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:



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]).