

## Product datasheet for TP302989

## OriGene Technologies, Inc.

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## UFD1 (NM\_005659) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human ubiquitin fusion degradation 1 like (yeast) (UFD1L), transcript

variant 1, 20 µg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone

or AA Sequence:

>RC202989 protein sequence Red=Cloning site Green=Tags(s)

MFSFNMFDHPIPRVFQNRFSTQYRCFSVSMLAGPNDRSDVEKGGKIIMPPSALDQLSRLNITYPMLFKLT NKNSDRMTHCGVLEFVADEGICYLPHWMMQNLLLEEGGLVQVESVNLQVATYSKFQPQSPDFLDITNPK

Α

VLENALRNFACLTTGDVIAINYNEKIYELRVMETKPDKAVSIIECDMNVDFDAPLGYKEPERQVQHEEST EGEADHSGYAGELGFRAFSGSGNRLDGKKKGVEPSPSPIKPGDIKRGIPNYEFKLGKITFIRNSRPLVKK

VEEDEAGGRFVAFSGEGQSLRKKGRKP

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK
Predicted MW: 34.3 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 005650





**Locus ID:** 7353

 UniProt ID:
 Q92890

 RefSeq Size:
 1783

Cytogenetics: 22q11.21

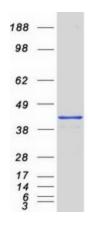
RefSeq ORF: 921
Synonyms: UFD1L

**Summary:** The protein encoded by this gene forms a complex with two other proteins, nuclear protein

localization-4 and valosin-containing protein, and this complex is necessary for the

degradation of ubiquitinated proteins. In addition, this complex controls the disassembly of the mitotic spindle and the formation of a closed nuclear envelope after mitosis. Mutations in this gene have been associated with Catch 22 syndrome as well as cardiac and craniofacial defects. Alternative splicing results in multiple transcript variants encoding different isoforms. A related pseudogene has been identified on chromosome 18. [provided by RefSeq, Jun 2009]

## **Product images:**



Coomassie blue staining of purified UFD1 protein (Cat# TP302989). The protein was produced from HEK293T cells transfected with UFD1 cDNA clone (Cat# [RC202989]) using MegaTran 2.0 (Cat# [TT210002]).