

## **Product datasheet for TP301637M**

#### OriGene Technologies, Inc.

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# Lysosomal acid lipase (LIPA) (NM\_000235) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human lipase A, lysosomal acid, cholesterol esterase (LIPA),

transcript variant 2, 100 µg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC201637 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MKMRFLGLVVCLVLWPLHSEGSGGKLTALDPETNMNVSEIISYWGFPSEEYLVETEDGYILCLNRIPHGR KNHSDKGPKPVVFLQHGLLADSSNWVTNLANSSLGFILADAGFDVWMGNSRGNTWSRKHKTLSVSQDE

 $\mathsf{FW}$ 

AFSYDEMAKYDLPASINFILNKTGQEQVYYVGHSQGTTIGFIAFSQIPELAKRIKMFFALGPVASVAFCT SPMAKLGRLPDHLIKDLFGDKEFLPQSAFLKWLGTHVCTHVILKELCGNLCFLLCGFNERNLNMSRVDVY TTHSPAGTSVQNMLHWSQAVKFQKFQAFDWGSSAKNYFHYNQSYPPTYNVKDMLVPTAVWSGGHDW

LADV

YDVNILLTQITNLVFHESIPEWEHLDFIWGLDAPWRLYNKIINLMRKYQ

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 43 kDa

**Concentration:**  $>0.1 \mu g/\mu L$  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.



#### Lysosomal acid lipase (LIPA) (NM\_000235) Human Recombinant Protein - TP301637M

**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 000226

 Locus ID:
 3988

 UniProt ID:
 P38571

 RefSeq Size:
 2775

Cytogenetics: 10q23.31 RefSeq ORF: 1197

Synonyms: CESD; LAL

**Summary:** This gene encodes lipase A, the lysosomal acid lipase (also known as cholesterol ester

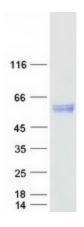
hydrolase). This enzyme functions in the lysosome to catalyze the hydrolysis of cholesteryl esters and triglycerides. Mutations in this gene can result in Wolman disease and cholesteryl ester storage disease. Alternatively spliced transcript variants have been found for this gene.

[provided by RefSeq, Jan 2014]

**Protein Families:** Druggable Genome

Protein Pathways: Lysosome, Steroid biosynthesis

### **Product images:**



Coomassie blue staining of purified LIPA protein (Cat# [TP301637]). The protein was produced from HEK293T cells transfected with LIPA cDNA clone (Cat# [RC201637]) using MegaTran 2.0 (Cat# [TT210002]).