

Product datasheet for AR51843PU-S

TOM1L2 (1-507, His-tag) Human Protein

Product data:

Product Type: Recombinant Proteins

Description: TOM1L2 (1-507, His-tag) human protein, 50 μg

Species: Human **Expression Host:** E. coli

Expression cDNA Clone

MGSSHHHHHH SSGLVPRGSH MGSEFMEFLL GNPFSTPVGQ CLEKATDGSL QSEDWTLNME or AA Sequence: ICDIINETEE GPKDAIRALK KRLNGNRNYR EVMLALTVLE TCVKNCGHRF HILVANRDFI DSVLVKIISP

> KNNPPTIVQD KVLALIQAWA DAFRSSPDLT GVVHIYEELK RKGVEFPMAD LDALSPIHTP QRSVPEVDPA ATMPRSQSQQ RTSAGSYSSP PPAPYSAPQA PALSVTGPIT ANSEQIARLR SELDVVRGNT KVMSEMLTEM VPGQEDSSDL ELLQELNRTC RAMQQRIVEL ISRVSNEEVT EELLHVNDDL NNVFLRYERF ERYRSGRSVQ NASNGVLNEV TEDNLIDLGP GSPAVVSPMV GNTAPPSSLS SQLAGLDLGT ESVSGTLSSL QQCNPRDGFD MFAQTRGNSL AEQRKTVTYE DPQAVGGLAS ALDNRKQSSE GIPVAQPSVM DDIEVWLRTD LKGDDLEEGV TSEEFDKFLE

ERAKAAEMVP DLPSPPMEAP APASNPSGRK KPERSEDALF AL

Tag: His-tag Predicted MW: 58.2 kDa Concentration: lot specific

>80% by SDS - PAGE **Purity:**

Buffer: Presentation State: Purified

State: Liquid purified protein

Buffer System: Phosphate buffered saline (pH 7.4) containing 10% glycerol, 1 mM DTT

Preparation: Liquid purified protein

Storage: Store undiluted at 2-8°C for one week or (in aliquots) at -20°C to -80°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

RefSeq: NP 001028723

Locus ID: 146691

UniProt ID: Q6ZVM7, Q6ZVM7-2

Cytogenetics: 17p11.2



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

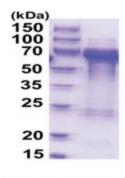
Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



Summary:

This gene belongs to a small gene family whose members have an N-terminal VHS domain followed by a GAT domain; domains which typically participate in vesicular trafficking. The canonical protein encoded by this gene also has a C-terminal clathrin binding motif. This protein has been shown to interact with Tollip, clathrin and ubiquitin and is thought to play a role in endosomal sorting. This gene resides in the 3.7 Mb deletion of chromosome region 17p11.2 that is associated with Smith-Magenis syndrome. Alternative splicing results in multiple transcript variants encoding distinct proteins. [provided by RefSeq, Apr 2017]

Product images:



15% SDS-PAGE (3ug)