

Product datasheet for TP750166

TBL1 (TBL1X) (NM_005647) Human Recombinant Protein

Product data:

Product Type:	Recombinant Proteins
Description:	Purified recombinant protein of Human transducin (beta)-like 1X-linked (TBL1X), transcript variant 1, full length, Tag free, expressed in E.coli, 50ug
Species:	Human
Expression Host:	E. coli
Expression cDNA Clone or AA Sequence:	A DNA sequence encoding the full length of human TBL1.
Tag:	Tag Free
Predicted MW:	62.3 kDa
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Buffer:	50 mM Tris-HCl, pH 8.0, 500 mM NaCl, 10% glycerol
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:	<u>NP 005638</u>
Locus ID:	6907
UniProt ID:	<u>O60907, A0A024RBV9</u>
RefSeq Size:	5886
Cytogenetics:	Хр22.31-р22.2
RefSeq ORF:	1731
Synonyms:	CHNG8; EBI; SMAP55; TBL1



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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

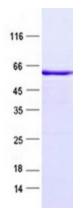
GRIGENE TBL1 (TBL1X) (NM_005647) Human Recombinant Protein – TP750166

Summary: The protein encoded by this gene has sequence similarity with members of the WD40 repeatcontaining protein family. The WD40 group is a large family of proteins, which appear to have a regulatory function. It is believed that the WD40 repeats mediate protein-protein interactions and members of the family are involved in signal transduction, RNA processing, gene regulation, vesicular trafficking, cytoskeletal assembly and may play a role in the control of cytotypic differentiation. This encoded protein is found as a subunit in corepressor SMRT (silencing mediator for retinoid and thyroid receptors) complex along with histone deacetylase 3 protein. This gene is located adjacent to the ocular albinism gene and it is thought to be involved in the pathogenesis of the ocular albinism with late-onset sensorineural deafness phenotype. Four transcript variants encoding two different isoforms have been found for this gene. This gene is highly similar to the Y chromosome TBL1Y gene. [provided by RefSeq, Nov 2008]

Protein Families: Transcription Factors

Protein Pathways: Wnt signaling pathway

Product images:



Purified recombinant protein TBL1 was analyzed by SDS-PAGE gel and Coomossie Blue Staining.

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