

Product datasheet for **TP303625M**

BSCL2 (NM_032667) Human Recombinant Protein

Product data:

Product Type:	Recombinant Proteins
Description:	Recombinant protein of human Bernardinelli-Seip congenital lipodystrophy 2 (seipin) (BSCL2), transcript variant 2, 100 µg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC203625 representing NM_032667 Red =Cloning site Green =Tags(s)

MVNDPPVPALLWAQEVGQVLAGRARRLLLQFGVLFCTILLLLWVSFLYGSFYYSYMPTVSHLSPVHFYY
RTDCDSSTTSLCSFPVANVSLTKGGRDRVLMYGGPYRVTLLELPESPVNQDLGMFLVTISCYTRGGRII
STSSRSVMLHYRSDLLQMLDTLVFSSLLLFGFAEQKQLLEVELYADYRENSYVPTTGAIIEIHSKRIQLY
GAYLRIHAHFTGLRYLLYNFPMTCAFIGVASNFTFLSVIVLFSYMQVWVGGIWPRHRFSLQVNIRKRDNS
RKEVQRRISAHQPGAGPEGQEESTPQSDVTEDGESPEDPSGTEGQLSSEEEKPDQQPLSGEEEELEPEASDG
SGSWEDAALLTEANLPAPAPASASAPVLETLSGSEEPAGGALRQRPTCSSS

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Tag:	C-Myc/DDK
Predicted MW:	44.2 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:	<u>NP_116056</u>



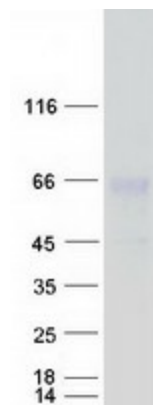
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Locus ID:	26580
UniProt ID:	Q96G97 , A0A024R549
RefSeq Size:	1664
Cytogenetics:	11q12.3
RefSeq ORF:	1200
Synonyms:	GNG3LG; HMN5; HMN5C; PELD; SPG17

Summary: This gene encodes the multi-pass transmembrane protein protein seipin. This protein localizes to the endoplasmic reticulum and may be important for lipid droplet morphology. Mutations in this gene have been associated with congenital generalized lipodystrophy type 2 or Berardinelli-Seip syndrome, a rare autosomal recessive disease characterized by a near absence of adipose tissue and severe insulin resistance. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. Naturally occurring read-through transcription occurs between this locus and the neighboring locus HNRNPUL2 (heterogeneous nuclear ribonucleoprotein U-like 2).[provided by RefSeq, Mar 2011]

Protein Families: Druggable Genome, Transmembrane

Product images:



Coomassie blue staining of purified BSCL2 protein (Cat# [TP303625]). The protein was produced from HEK293T cells transfected with BSCL2 cDNA clone (Cat# [RC203625]) using MegaTran 2.0 (Cat# [TT210002]).