

OriGene Technologies, Inc.

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Product datasheet for PH307050

METTL27 (NM_152559) Human Mass Spec Standard

Product data:

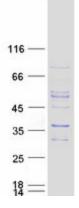
Product Type:	Mass Spec Standards
Description:	WBSCR27 MS Standard C13 and N15-labeled recombinant protein (NP_689772)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC207050
Predicted MW:	26.7 kDa
Protein Sequence:	>RC207050 protein sequence <mark>Red=</mark> Cloning site Green=Tags(s)
	MAQEEGGSLPEVRARVRAAHGIPDLAQKLHFYDRWAPDYDQDVATLLYRAPRLAVDCLTQALPGPPHSAL ILDVACGTGLVAAELRAPGFLQLHGVDGSPGMLEQARAPGLYQRLSLCTLGQEPLPSPEGTFDAVLIVGA LSDGQVPCNAIPELHVTKPGGLVCLTTRTNWSNLQYKEALEATLDRLEQAGMWEGLVAWPVDRLWTAGSW LPPSWWWYPASLPRMASSPALSTCTESGRRPRLRK
	TRTRPLEQKLISEEDLAANDILDYKDDDDKV
Tag:	C-Myc/DDK
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Labeling Method:	Labeled with [U- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-L-Lysine
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3
Storage:	Store at -80°C. Avoid repeated freeze-thaw cycles.
Stability:	Stable for 3 months from receipt of products under proper storage and handling conditions.
RefSeq:	<u>NP 689772</u>
RefSeq Size:	958
RefSeq ORF:	735
Synonyms:	WBSCR27
Locus ID:	155368
UniProt ID:	<u>Q8N6F8</u>



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	METTL27 (NM_152559) Human Mass Spec Standard – PH307050
Cytogenetics:	7q11.23
Summary:	This gene encodes a protein belonging to ubiE/COQ5 methyltransferase family. The gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.22-q11.23. [provided by RefSeq, Jul 2008]

Product images:



Coomassie blue staining of purified METTL27 protein (Cat# [TP307050]). The protein was produced from HEK293T cells transfected with METTL27 cDNA clone (Cat# [RC207050]) using MegaTran 2.0 (Cat# [TT210002]).

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