

#### OriGene Technologies, Inc.

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# Product datasheet for PH321699

### CD105 (ENG) (NM\_000118) Human Mass Spec Standard

### **Product data:**

Product Type:	Mass Spec Standards
Description:	ENG MS Standard C13 and N15-labeled recombinant protein (NP_000109)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC221699
Predicted MW:	67.78 kDa
Protein Sequence:	<pre>&gt;RC221699 representing NM_000118 Red=Cloning site Green=Tags(s)</pre>
	MDRGTLPLAVALLLASCSLSPTSLAETVHCDLQPVGPERGEVTYTTSQVSKGCVAQAPNAILEVHVLFLE FPTGPSQLELTLQASKQNGTWPREVLLVLSVNSSVFLHLQALGIPLHLAYNSSLVTFQEPPGVNTTELPS FPKTQILEWAAERGPITSAAELNDPQSILLRLGQAQGSLSFCMLEASQDMGRTLEWRPRTPALVRGCHLE GVAGHKEAHILRVLPGHSAGPRTVTVKVELSCAPGDLDAVLILQGPPYVSWLIDANHNMQIWTTGEYSFK IFPEKNIRGFKLPDTPQGLLGEARMLNASIVASFVELPLASIVSLHASSCGGRLQTSPAPIQTTPPKDTC SPELLMSLIQTKCADDAMTLVLKKELVAHLKCTITGLTFWDPSCEAEDRGDKFVLRSAYSSCGMQVSASM ISNEAVVNILSSSSPQRKKVHCLNMDSLSFQLGLYLSPHFLQASNTIEPGQQSFVQVRVSPSVSEFLLQL DSCHLDLGPEGGTVELIQGRAAKGNCVSLLSPSPEGDPRFSFLLHFYTVPIPKTGTLSCTVALRPKTGSQ DQEVHRTVFMRLNIISPDLSGCTSKGLVLPAVLGITFGAFLIGALLTAALWYIYSHTREYPRPPQSGP
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 000109</u>
RefSeq Size:	3142
RefSeq ORF:	1884

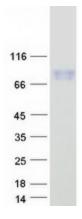


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	CD105 (ENG) (NM_000118) Human Mass Spec Standard – PH321699
Synonyms:	END; HHT1; ORW1
Locus ID:	2022
UniProt ID:	<u>P17813, Q5T9B9</u>
Cytogenetics:	9q34.11
Summary:	This gene encodes a homodimeric transmembrane protein which is a major glycoprotein of the vascular endothelium. This protein is a component of the transforming growth factor beta receptor complex and it binds to the beta1 and beta3 peptides with high affinity. Mutations in this gene cause hereditary hemorrhagic telangiectasia, also known as Osler-Rendu-Weber syndrome 1, an autosomal dominant multisystemic vascular dysplasia. This gene may also be involved in preeclampsia and several types of cancer. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, May 2013]
Protein Families	: Druggable Genome, ES Cell Differentiation/IPS, Transmembrane

## Product images:



Coomassie blue staining of purified ENG protein (Cat# [TP321699]). The protein was produced from HEK293T cells transfected with ENG cDNA clone (Cat# [RC221699]) using MegaTran 2.0 (Cat# [TT210002]).

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