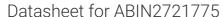
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GDAP1L1 Protein (Myc-DYKDDDDK Tag)



Image



Go to Product page

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Overview	
Quantity:	20 μg
Target:	GDAP1L1
Origin:	Human
Source:	HEK-293 Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This GDAP1L1 protein is labelled with Myc-DYKDDDDK Tag.
Application:	Antibody Production (AbP), Standard (STD)
Product Details	
Characteristics:	 Recombinant human GDAP1L1 protein expressed in HEK293 cells. Produced with end-sequenced ORF clone
Purity:	> 80 % as determined by SDS-PAGE and Coomassie blue staining
Target Details	
Target:	GDAP1L1
Alternative Name:	Gdap1I1 (GDAP1L1 Products)
Background:	The ganglioside GD3 synthase causes cell differentiation with neurite sprouting when transfected into the mouse neuroblastoma cell line Neuro2a. After differentiation, the expression of several genes is upregulated, including one that encodes a protein termed ganglioside-induced differentiation-associated protein 1 (Gdap1). A similar gene was found in humans, and mutations in the human gene are associated with Charcot-Marie-Tooth type 4A
	humans, and mutations in the human gene are associated with Charcot-Marie-Tooth typ

Target Details

Molecular Weight:	41.8 kDa
	variant, have been found for this gene.
	Several transcript variants encoding different isoforms, as well as a noncoding transcript
	disease. The protein encoded by this gene is similar in sequence to the human GDAP1 protein.

Application Details

NP_076939

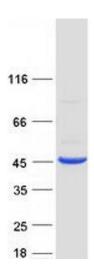
NCBI Accession:

Application Notes:	Recombinant human proteins can be used for:
	Native antigens for optimized antibody production
	Positive controls in ELISA and other antibody assays
Comment:	The tag is located at the C-terminal.
Restrictions:	For Research Use only

Handling

Concentration:	50 μg/mL
Buffer:	25 mM Tris.HCl, pH 7.3, 100 mM glycine, 10 % glycerol.
Storage:	-80 °C
Storage Comment:	Store at -80°C. Thaw on ice, aliquot to individual single-use tubes, and then re-freeze immediately. Only 2-3 freeze thaw cycles are recommended.

Images



Western Blotting

Image 1. Validation with Western Blot