

## Product datasheet for TP508839

### Atl1 (NM\_178628) Mouse Recombinant Protein

#### Product data:

Product Type:	Recombinant Proteins
Description:	Purified recombinant protein of Mouse atlastin GTPase 1 (Atl1), with C-terminal MYC/DDK tag, expressed in HEK293T cells, 20ug
Species:	Mouse
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>MR208839 representing NM_178628 <b>Red</b> =Cloning site <b>Green</b> =Tags(s)

MAKSRDRNSWGGFSEKSSDWSSEEEEPVRKAGPVQVLIVKDDHSFELDEAALNRILLSQAVRDKEVVAV  
SVAGAFRKGKSFLMDFMLRYMYNQESVDWVGDYNEPLTGFSWRGGSERETTGIQIWSEVFLINKLDGKKV  
AVLLMDTQGTFDQSSTLRDSATVFALSTMISSIQVYNLSQNVQEDDLQHLQLFTEYGRAMEETFLKPFQ  
SLIFLVRDWSFPYEFYSGADGGAKFLEKRLKVSGNQHEELQNVKHIHSCFTNISCFLLPHPGLKVATNP  
NFDGKLEIDDEFIKNLKILIPWLLSPERLDIKEINGNKITCRGLLEYFKAYIKIYQGEELPHPKSMMLQA  
TAEANNLAAVATAKDTYNKKMEEVCGGDKPFLAPNDLQSKHLQLKEESVKLFRGVKKMGGEFSSRRYLQQ  
LESEIDELIYQYIKHNSKNIFHAARTPATLFWVIFITYVIAGVTGFIGLDIIASLCNMIMGLTLITLCT  
WAYIRYSGEYRELGAVIDQVAAALWDQGSTNEALYKLYSAAATHRHLCQAFPAPKSEPTQQPEKKKI

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV**

Tag:	C-MYC/DDK
Predicted MW:	63.8 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
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 after receiving vials.
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:	<a href="#">NP_848743</a>



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Locus ID:	73991
UniProt ID:	<a href="#">Q8BH66</a>
RefSeq Size:	2818
Cytogenetics:	12 C2
RefSeq ORF:	1674
Synonyms:	4930435M24Rik; AD-; Adfsp; FS; Fsp1; SP; Spg3; Spg3a
Summary:	This gene encodes a member of the dynamin family of GTPases. The encoded protein interacts with tubule-shaping proteins of the endoplasmic reticulum. Mutations in the homologous human gene can cause hereditary spastic paraplegia. [provided by RefSeq, Feb 2010]