

#### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

# Product datasheet for TP710292

# Adracalin (AAAS) (NM\_015665) Human Recombinant Protein

# **Product data:**

Product Type:	Recombinant Proteins
Description:	Purified recombinant protein of Human achalasia, adrenocortical insufficiency, alacrimia (AAAS), transcript variant 1, full length, with C-terminal DDK tag, expressed in sf9, 20ug
Species:	Human
Expression Host:	Sf9
Expression cDNA Clone or AA Sequence:	A DNA sequence from TrueORF clone, RC200154, encoding human full-length AAAS
Tag:	C-DDK
Predicted MW:	59.4 kDa
Concentration:	>0.05 $\mu$ g/ $\mu$ L as determined by microplate BCA method
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Buffer:	50 mM Tris-HCl, 100 mM glycine, pH 8.0, 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.
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 056480</u>
Locus ID:	8086
UniProt ID:	Q9NRG9
RefSeq Size:	1854
Cytogenetics:	12q13.13
RefSeq ORF:	1638
Synonyms:	AAA; AAASb; ADRACALA; ADRACALIN; ALADIN; GL003



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

### **GRIGENE** Adracalin (AAAS) (NM\_015665) Human Recombinant Protein – TP710292

Summary:The protein encoded by this gene is a member of the WD-repeat family of regulatory proteins<br/>and may be involved in normal development of the peripheral and central nervous system.<br/>The encoded protein is part of the nuclear pore complex and is anchored there by NDC1.<br/>Defects in this gene are a cause of achalasia-addisonianism-alacrima syndrome (AAAS), also<br/>called triple-A syndrome or Allgrove syndrome. Two transcript variants encoding different<br/>isoforms have been found for this gene. [provided by RefSeq, Mar 2010]

### **Product images:**

116 — 66 — 45 — 35 — 25 — 18 — 14 —

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US