

Product datasheet for **SC328848**

Adractalin (AAAS) (NM_001173466) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	Adractalin (AAAS) (NM_001173466) Human Untagged Clone
Tag:	Tag Free
Symbol:	Adractalin
Synonyms:	AAA; AAASb; ADRACALA; ADRACALIN; ALADIN; GL003
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)



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Fully Sequenced ORF: >SC328848 representing NM_001173466.
 Blue=Insert sequence Red=Cloning site Green=Tag(s)

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GCTCGTTTAGTGAACCGTCAGAATTTTGTAAACGACTCACTATAGGGCGGCCGGGAATTCGTGACTG
GATCCGGTACCGAGGAGATCTGCCGCCCGCATCGCC
ATGTGCTCTCTGGGGTTGTTCCCTCCTCACCGCCTCGGGTCAAGTCACCTATATGAGCACAATAAC
GAGCTGGTGACGGCAGTAGCTATGAGAGCCCGCCCCGACTTCCGGGGCCAGTGGATCAATCTTCTC
GTCTACAACGACAAAGGATCCCTAAAGACCCCTGGAAGGCTGGACCATGGCACAAGAAGTGCCTTC
ATCCATCACCGGGAGCAAGTGTGAAGAGATGCATCAACATTTGGCGTGATGTGGCCCTTTTTGGGGTG
CTAAATGAAATTGCAAACCTCAGAAGAAGAGGTGTTTGTAGTGGTGAAGACGGCATCCGGCTGGGCCCTG
GCACTCTGTGATGGCCCTTCCCTCCATGGTCCCTGTTCCCCATCTGTCTCTCAGGAGCGAAGAT
CTGATCGTGAATTTGCCAAGTCACAAATTGCACCATAGTCCCCTCCGAAGCACCAGGCTGCAGCGA
AATGTGGCGTCTCTGGCCTGGAAGCCCTTAGTGCCTCTGTCTTGGCTGTGGCCTGCCAGAGCTGCATT
CTTATCTGGACCCTGGACCCTACCTCCTTGTCTACCCGACCCTCTCTGGCTGTGCCAAGTGTGTCT
CACCTGGGCATACACCTGTTACCAGCTTGGCCTGGGCCCCAGTGGGGGGCGGCTGCTCTCAGCTTCA
CCCGTGGATGCTGCTATCCGGGTATGGGATGTCTCAACAGAGACCTGTGTCCCCCTTCCCTGGTTCCGA
GGAGGTGGGGTGACCAACCTGCTCTGGTCCCCAGACGGCAGCAAAATCCTGGCTACCCTCCTTACGCT
GTCTTTGAGTCTGGGAGGCCAGATGTGGACTTGTGAGAGGTGGCCTACTCTATCAGGGCGCTGTGAG
ACTGGCTGCTGGAGCCCAGATGGCAGCCGACTGCTGTTCACTGTATTGGGAGAGCCACTGATTTACTCC
CTGTCTTTCCAGAACGTTGTGGTGAGGGAAAGGGTGCCTTGGAGGTGCAAAGTACAGCAACGATTGTG
GCAGATCTGTCTGAGACAACAATACAGACACCAGATGGTGAAGAGAGGCTTGGGGGAGAGGCTACTCC
ATGGTCTGGGACCCAGTGGGGAACGCTGGCTGTGCTTATGAAAGGAAAGCCAAGGGTACAGGATGGT
AAACCAGTCATCCTCCTTTTTCGCACTCGAAACAGCCCTGTGTTGAGCTCCTTCCCTGTGGCATTATC
CAGGGGGAGCCAGGAGCCAGCCAGCTCATCACTTTCCATCCTTCAACAAAGGGGCCCTGCTC
AGTGTGGGCTGGTCCACAGGCCGAATTGCCACATCCCGCTGTACTTTGTCAATGCCAGTTTCCACGT
TTAGCCAGTGCTTGGGCGGGCCAGGAACCCCTGCTGGGGTGGAGGCTCTATTATGACCTGCC
CTCTTTACTGAGACATCCCAACCTCTGCCCTTGGGACCCTCTCCAGGGCCACCACCTGTTCTGCC
CACTCCCCACATTCCCACCTTAA
ACGGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTAAACGGCCGGC
  
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Restriction Sites: SgfI-MluI

ACCN: NM_001173466

Insert Size: 1542 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method:

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: [NM_001173466.1](#)

RefSeq Size: 1755 bp

RefSeq ORF: 1542 bp

Locus ID: 8086

UniProt ID: [Q9NRG9](#)

Cytogenetics: 12q13.13

MW: 55.8 kDa

Gene Summary: The protein encoded by this gene is a member of the WD-repeat family of regulatory proteins and may be involved in normal development of the peripheral and central nervous system. The encoded protein is part of the nuclear pore complex and is anchored there by NDC1. Defects in this gene are a cause of achalasia-addisonianism-alacrima syndrome (AAAS), also called triple-A syndrome or Allgrove syndrome. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Mar 2010]
Transcript Variant: This variant (2) lacks an alternate in-frame exon compared to variant 1. The resulting isoform (2) has the same N- and C-termini but is shorter compared to isoform 1.