

Product datasheet for **SC337861**

ADNP (NM_001282531) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	ADNP (NM_001282531) Human Untagged Clone
Tag:	Tag Free
Symbol:	ADNP
Synonyms:	ADNP1; HVDAS; MRD28
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
Fully Sequenced ORF:	>NCBI ORF sequence for NM_001282531, the custom clone sequence may differ by one or more nucleotides

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ATGTTCCAACCTCCTGTCAACAATCTTGGCAGTTAAGAAAAGCCCGAAAACCTGTGAAAAAATACTTA
GTGACATTGGGTTGGAATACTGTAAAGAACATATAGAAGATTTTAAACAATTTGAACCTAATGACTTTTA
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ACAAAACCTTTCTGCTGCAGCGCTTGCCATTTTCTCAAATTTCTCTGCCTACAAAAGTCATTTCC
GCAATGTCCATAGTGAAGACTTTGAAAAATAGGATTCTCCTTAATTGCCCTACTGTACCTCAATGCAGA
CAAAAAGACTTTGAAACACACATTAATAATTTTCATGCTCCGAACGCCAGCGCACCAAGTAGCAGCCTC
AGCACTTTCAAAGATAAAAACAAAAATGATGGCCTTAAACCTAAGCAGGCTGACAGTGTAGAGCAAGCTG
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CACAAATGTAGTGGTTCCCCGATCCAAACCCTTGATGCTAATTGCTCCCAAACCTCAAGACAAGAAGAGC
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TGAAGAGATGGGACCTAAAACAGATTCTACTTTGAGTTTTGATTTGACATTGCAGCAGGGTAGTCACACT
AACATCCATCTCCTGGTAACTACATACAATCTGAGGGATGCCCCAGCTGAATCTGTTGCTTACCATGCC
AAAATAATCCTCCAGTTCCTCCAAAGCCACAGCCAAAGGTTCCAGGAAAAGGCAGATATCCCTGTA
TTCACCTCAAGCTGCAGTGCCCTATAAAAAAGATGTTGGGAAAACCCCTTTGCTCTTTGCTTTTCAATC
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ATCCAGTTGAGAAAAAGCTCACCTACAAATGTATCCATTGCCTTGGTGTGTATACCAGCAACATGACCGC
CTCAACTATCACTCTGCATCTAGTTCACTGCAGGGCGTTGAAAAGACCCAAAATGGCCAGGATAAGACA
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CAAAAACGAAACTATTCATTTGACTGAGGAACCAACCAACTAATGCACAATGCATCTGATAGTGGGT
TGACCAAGACGATGTTGTTGAGTGGAAAGACGGTGCTTCTCCATCTGAGAGTGGGCTGGATCCCAACAA
GTGTCAGACTTTGAGGACAATACCTGCGAAATGAAACCAGGAACCTGGTCTGACGAGTCTTCCAAAAGCG
AAGATGCAAGGAGCAGTAAGCCAGCTGCCAAAAAAGGCTACCATGCAAGGTGACAGAGAGCAGTTGAA
ATGGAAGAATAGTTCCTATGGAAAAGTTGAAGGGTTTTGGTCTAAGGACCAGTACAGTGGAAAGAATGCA
TCTGAGAATGATGAGCGTTATCTAACCCCCAGATTGAGTGGCAGAATAGCACAATTGACAGTGGAGATG
GGGAACAGTTTGACAACATGACTGATGGAGTAGCTGAGCCCATGCATGGCAGCTTAGCCGGAGTTAACT
GAGCAGCCAACAGGCCTAA

- Restriction Sites:** SgfI-MluI
- ACCN:** NM_001282531
- Insert Size:** 3309 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).
- 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_001282531.1](#), [NP_001269460.1](#)
- RefSeq Size:** 6242 bp
- RefSeq ORF:** 3309 bp

Locus ID: 23394

UniProt ID: [Q9H2P0](#)

Cytogenetics: 20q13.13

Protein Families: Transcription Factors

Gene Summary: Vasoactive intestinal peptide is a neuroprotective factor that has a stimulatory effect on the growth of some tumor cells and an inhibitory effect on others. This gene encodes a protein that is upregulated by vasoactive intestinal peptide and may be involved in its stimulatory effect on certain tumor cells. The encoded protein contains one homeobox and nine zinc finger domains, suggesting that it functions as a transcription factor. This gene is also upregulated in normal proliferative tissues. Finally, the encoded protein may increase the viability of certain cell types through modulation of p53 activity. Alternatively spliced transcript variants encoding the same protein have been described. [provided by RefSeq, Jul 2008]

Transcript Variant: This variant (3) represents the longest transcript. All five variants encode the same protein. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.