

## Human adenovirus C, complete genome

NCBI Reference Sequence: NC\_001405.1

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LOCUS NC\_001405 35937 bp DNA linear VRL 13-AUG-2018

DEFINITION Human adenovirus C, complete genome.

ACCESSION NC\_001405

VERSION NC\_001405.1

DBLINK BioProject: [PRJNA485481](#)

KEYWORDS RefSeq.

SOURCE Human mastadenovirus C

ORGANISM [Human mastadenovirus C](#)

Viruses; Varidnaviria; Bamfordvirae; Preplasmiviricota;  
Tectiliviricetes; Rowavirales; Adenoviridae; Mastadenovirus.

REFERENCE 1 (sites)

AUTHORS Roberts,R.J., Akusjaervi,G., Alestroem,P., Gelinas,R.E.,  
Gingeras,T.R., Sciaky,D. and Pettersson,U.

TITLE A consensus sequence for the adenovirus-2 genome

JOURNAL (in) Doerfler,W. (Ed.);

ADENOVIRUS DNA: 1-51;

Martinus Nijhoff Publishing, Boston (1986)

REFERENCE 2 (bases 1 to 35937)

CONSRM NCBI Genome Project

TITLE Direct Submission

JOURNAL Submitted (30-SEP-2004) National Center for Biotechnology  
Information, NIH, Bethesda, MD 20894, USA

COMMENT PROVISIONAL [REFSEQ](#): This record is based on preliminary  
annotation

provided by Dr Andrew Davison (MRC Virology Unit, Glasgow, UK) and  
Dr Balazs Harrach (Veterinary Medical Research Institute,  
Budapest,

Hungary). The reference sequence was derived from [J01917](#).

CURATION:

Proteins: A standard nomenclature has been applied so that  
orthologs have the same name. Functional information may have been  
propagated from other adenovirus species.

Protein-coding regions: The initiation codon for each CDS is  
assigned with as much confidence as possible. CDSs presumably  
inherited from the last common ancestor of all adenoviruses are  
indicated as genus-common, and all other CDSs as genus-specific.  
CDSs that have paralogues in one or more genomes are listed as  
members of named families. The evolutionary history of E3 and E4

is

complex, frequently involving generation or loss of paralogues in

whole or in part, accompanied by rapid sequence divergence. In E3, this is evident in a set of tandemly arranged CDSs encoding transmembrane proteins. All of these are listed as members of the MP family, regardless of whether sequence conservation is detectable.

**Transcripts:** In mastadenoviruses, transcription arises from the early genes (E1A, E1B, E2A, E2B, E3 and E4), the intermediate genes (IX and IVa2) and the late genes (L, U and E2A-L). Certain transcriptional features are supported by experimental data for some of these genes in a few adenoviruses (particularly human adenovirus C). All transcriptional features not marked as experimentally supported are predictions made on the basis of conservation and relevance to CDS expression. The degree of confidence in each prediction depends on the feature under consideration. In primate adenoviruses, predictions for E3, E4 and the 5'-regions of E2A and E2B are generally the least certain. In non-primate mastadenoviruses, this trend is accentuated to the degree that predictions for introns in E3 and E4 are not provided, and those for the 5'-regions of E2A, E2B and IVa2 and for the intron separating the protein-coding exons of 33K are particularly tentative. In human adenovirus C, rightward-oriented transcripts from the late genes (L1-L5) are polyadenylated close downstream from polyadenylation signals at five locations (the 3'-ends of the pIIIa, pX, protease, control protein E3 12.5K and fiber CDSs), and those from the E3 genes (E3A and E3B) at two locations (the 3'-

ends

of the membrane protein E3 CR1-beta and control protein E3 14.7K CDSs). Only the polyadenylation signals for L3, E3B and L5 are conserved among all mastadenoviruses, whereas those for L1 and L2 are conserved in most mastadenoviruses and those for L4 and E3A in a few. In addition, potential additional polyadenylation signals exist at the 3'-ends of the penton base, pVII, V, pVI and fiber-2 CDSs in some mastadenoviruses, particularly those with non-primate hosts; the corresponding genes are listed as L1A, L1B, L1C, L2A

and

L5A, respectively. The gene corresponding to E3B is termed E3 in genomes lacking the E3A polyadenylation signal. In some instances where a TATA signal is not convincingly evident (particularly for E2A/E2B), a tentative assignment is made and marked as nominal.

**ACCURACY:** No potential sequence errors are noted.

**COMPLETENESS:** full length.

FEATURES	Location/Qualifiers
source	1..35937 /organism="Human mastadenovirus C"

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repeat_region 1..102
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              PKLVPAILRRPTSPVSRECNSSTDSCDSGSPNTPPEIHPVVPLCPKIPVAVRVGRRQ  
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other human mastadenoviruses, but their conservation may
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polymerase encoded on the opposing strand; protein
expression has been detected in infected cells; a mutant
has been analysed functionally; genus-specific"
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RRVPP
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 gene 10866..11023  
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EQT V NQEINFQKSFNNHVRTL VAREEVAIGLMHLWDFVSALEQNPNSKPLMAQLFLIV
QH SRDNEAFRDALLNIVEPEGRWLLDLINILQSIVVQERSLSLADKVAAINYSMLS LG
KFYARKIYHTPYVPI DKEVKIEGFYMRMALKVLTLSDDLGVYRNERIHKAVSVSRRE
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CDS 18001..18753  
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CDS 18838..21744

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FDSSVSWPGNDRLLTPNEFEIKRSVDGEGYNVAQCNMTKDWFLVQMLANYNIGYQGFY
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AYPANVPYPLIGKTAVDSITQKKFLCDRTLWRIPFSSNFMSMGALTDLQGNLLYANSA
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CDS 21778..22392

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 prim\_transcript complement(22420..27092)  
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 CDS complement(22490..24079)  
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 SEAESESTVINPLSLPIVSAWEKGMEEAARALMDKYHVDNDLKANFKLLPDQVEALAAV  
 CKTWLNEEHRGLQLTFTSNKTFVTMMGRFLQAYLQSF AEVTKYHHEPTGCALWLHRCA  
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CDS 24108..26525  
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 FAVPEVLATYHIFQNKIPLSCRANRSRADKQLALRQGAVIDIASLDEVPKIFEGL  
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 GILPATCCALPSDFVPIKYRECPLWGHCYLLQLANYLAYHSDIMEDVSGDGLLECH  
 CRCNLCTPHRSLVCNSQLLSESIIGTFELQGPSPDEKSAAPGLKLTPLWTSAYLRK  
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intron complement(24792..27024)  
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CDS join(26239..26551,26754..27127)  
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CDS      26239..26826
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gene 27580..29804  
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intron 27632..30437  
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intron 27632..30046  
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/note="precedes membrane protein E3 RID-beta CDS"

intron 27632..29764  
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/note="precedes membrane protein E3 RID-alpha CDS"

intron 27632..29465  
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genus-specific; MP family"
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CDS      28812..29291
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/old_locus_tag="HAdVCgp28"
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6.7K; probable relic; apoptosis; genus-specific; MP
family"
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CDS      29468..29773
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probable relic; cell egress; genus-specific; MP family"
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regulatory      29769..29774
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