

Nucleotide sequence of cDNA covering the complete coding part of the human vimentin gene

B.Honoré, P.Madsen, B.Basse, A.Andersen, E.Walburn, J.E.Celis and H.Leffers

Institute of Medical Biochemistry and Bioregulation Research Centre, Aarhus University, Ole Worms Allé, Build. 170, DK-8000 Aarhus C, Denmark

Submitted October 16, 1990

EMBL accession no. X56134

Seven clones containing all or parts of the coding sequence of the human vimentin gene (1398 bp) were isolated from a λ gt11 human cDNA library constructed by reverse transcription of mRNA purified from SV40 transformed MRC-5 fibroblasts (MRC-5 V2) (1). Together, the clones span a cDNA sequence of 1765 bp. Previous attempts to isolate cDNA for human vimentin have resulted in partial lengths below 1100 bp (2, 3).

A monoclonal antibody, 98C1, recognizing vimentin among other proteins, was used for preliminary screening of 2×10^5 plaques. Two positive clones were obtained. One clone contained a 1.4 kb insert. Both strands were sequenced with the dideoxy chain termination technique (4). This clone was used to re-screen 2×10^5 plaques giving approximately 80 positive clones. Five clones were isolated and end-sequenced. Two of these latter clones extended the preliminary sequence in both directions. Both strands of the extensions were sequenced. The overlapping sequences were identical except for 44 bp on one of the clones considered to be a cloning artefact. The complete cDNA sequence (1765 bp) is shown in Fig. 1. The sequence contained an open reading frame coding for a 466 amino acid protein found to be almost identical with the published sequence for human vimentin based on a combination of partial cDNA sequences and genomic DNA sequences (2, 3). Our sequence, however, shows some mismatches in the coding sequence with the previously published partial sequences. We have thus found Ser-42 and Leu-442 instead of Asp and Phe respectively (2), and Asn-201, Leu-265, Ser-278, Ser-339 and Asn-350 instead of Ser, Ser, Ile, Cys and Lys respectively (3). The 3' untranslated region contains 322 bp from the stop codon to the poly A stretch. In agreement with previous findings (3) we found two putative polyadenylation signals, the first is located 40 bp downstream from the stop codon, the second is located 301 bp from the stop codon.

ACKNOWLEDGEMENTS

We would like to thank O. Sønderskov for reproductions. P. Madsen was supported by a research position from the Medical Faculty, Aarhus University and H. Leffers by a fellowship from the Danish Cancer Foundation. The work was supported by grants from the Danish Biotechnology Programme, the Danish Cancer Society, the EEC Science Programme, the Danish

Medical Research Council, the Danish Rheumatoid Society, NOVO and the Fund for Lægevidenskabens Fremme.

REFERENCES

- Huschtscha, L.I. and Holiday, R. (1983) *J. Cell Sci.* **63**, 77–99.
- Ferrari, S., Battini, R., Kaczmarek, L., Rittling, S., Calabretta, B., de Riel, J.K., Philiponis, V., Wei, J.-F. and Baserga, R. (1986) *Mol. Cell. Biol.* **6**, 3614–3620.
- Perreau, J., Lilienbaum, A., Vasseur, M. and Paulin, D. (1988) *Gene* **62**, 7–16.
- Sanger, F., Nicklen, S. and Coulson, A.R. (1977) *Proc. Natl. Acad. Sci. USA* **74**, 5463–5467.

```

1 CGCGCCACCGCCGCCGCCAGCCATCGCCACCCCTCCGCGAGCCATGTCCACCAGGTCCTG
1 M S T R S V
61 GTCCCTCGTCTCCTACCGCAGGATGTTCCGGCCGCCCGGGCACCAGCGAGCCGGCCGAGTCT
7 S S S S Y R R M F G G P G T A S R P S S
121 CAGCCGGAGCTACGTGACTACGTCACCCCGCCTACAGCCTGGGACAGCGCGCTGCGCCC
27 S R S Y V T T S T R T Y S L G S A L R P
181 CAGCACCAGCCGAGCCTCTACGCTCGTCCCGGGCGGCTGTATGCCACCGCGCTCTC
47 S T S R S L Y A S S P G G V Y A T R S S
241 TGCCGTGGCCTGGGAGCAGCGTGCCTGGGCTGGGCTCTGCAGGACTCGGTGGACTT
67 A V R L R S S V P G V R L L Q D S V D F
301 CTCGTGGCCGACCCATCAACCCAGTTCAGAACCCCGCACCACGAGAAGTGGGA
87 S L A D A I N T E R T N E K V E
361 GTCGAGGAGTGAATGACCGTTCGCCAATCATCGCAAGTTCGGCTCTCTGGAGCA
107 L Q E L N D R F A N Y I D K R L E Q
421 CAGAAATAGATCCTGCTGGCCGAGCTCGAGCAGTCAAGGCCAAGGCAAGTCCGCT
127 Q N K I L L A E L E Q L K G Q G K S R L
481 GGGGACCTCTACGAGGAGGATGCGGGAGCTGCCCGGCGAGTGGACAGCTAACCAA
147 G D L Y E E M R E L R R Q V D Q L T N
541 CGACAAAGCCCGCTCGAGGTGGAGCGGACCACTGGCCGAGGACATCATGCGCCTCCG
167 D K A R V E V E R D N L A E D I M R L R
601 GGAGAAATGCGAGGAGATGCTTCAGAGAGGAGGACCCGAAACACCCCTGCAATCTT
187 E K L Q E E M L Q R E E A E N T L Q S F
661 CAGACAGGATGTTGACAAATCGCTCTGCGCAGCTTGGACCTTGAACGCAAGTGAATC
207 R Q D V D N A S L A R L D L E R K V E S
721 TTTGCAAGAAGATGTCCTTTTGAAGAACTCCACGAAAGGAAATCCAGGAGTGC
227 L Q E E I A F L K K L H E E E I Q E L Q
781 GGCTCAGATTCAGAACAGCATGTCACAAATCGATGGTGTATTTTCCAAGCTGACCTCAC
247 A Q I Q E Q H V Q I D V D V S K P D L T
841 GGCTGCCTCGCTGACGTACGTCAGCAATATGAAAGTGGTGGTCCGAACCTGCAGGA
267 A R L R D V R Q P Y E S V A A K N L Q E
901 GGCAGAAGATGGTACAATCCAGTTTCTGCTCCTCTGAGGCTCCCAACCGGACAA
287 A E E W Y K S K E A D L S E A N R N N
961 TGACGCCCTGCGCAGGCAAGCAGGAGTCCACTGAGTACCGGAGCAGGTGACGTCCT
307 D A L R Q A K Q E S T E Y R Q V Q S L
1021 CACCTGTAAGTGGATGCCCTTAAAGGAACTAGTCCCTGGAGCGCAGATGGCTGA
327 T C E V D A L K G T N E S L E R Q M R E
1081 AATGGAAGAGAACTTTCGGTGAAGTCTGCTATACCAAGACACTATGGCCGCTGCA
347 M E E N F A V E A A N Y Q D T I G R L Q
1141 GGATGAGATTCAGAATATGAGGAGGAAATGGCTCGTCACTTGGTGAATACCAAGCCT
367 D E I Q N M K E E M A R H L R E Y Q D L
1201 GCTCAATGTTAAGTGGCCCTTGACATGAGATGGCCACCTACAGGAGCTGCTGGAAG
387 L N V K M A L D I E I A T Y R K L L E G
1261 CGAGGAGCAGGATTTCTGCTCTTCCAACTTTTCTCCCTGAACCTGAGGAGAAC
407 E E S R I S L P L P N F S S L N L R E T
1321 TAATCTGGATCTCACTCCCTTGGTATACCCACTCAAAAGGACACTCTGTTAATGAC
1381 GGTGAAACTAGAGATGGACAGGTTATACAGAAATCTTCAGCACTCAGCATCAGACTTGA
447 V E T R D G Q V I N E T S Q H D D L E
1441 ATAAATGCACTACTCAGTGCAGCAATATATTACAGCAAGATAAAAAGAAATCC
1501 ATATCTTAAGAACTACTTCAAGTGCCTTTCGAGTTTTTCAGGAGCCAGATGAA
1561 TTTGGAATAGGAATACTTAGTCTTAAACACCCGACCTCTACAAGATTTAGAAAAA
1621 AGTTTACAACATACTACTTACAGAAATCTTGTGCTAGATACTTTTAAAGGTA
1681 TTTGAAATACCAATAAACTGCTTTTTCAGCAGTATCCCAACTTGGTCTT
1741 GCTTCAATAATCTTTGAAAAACTAn

```