

The nucleotide sequence of chicken smooth muscle myosin light chain two

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Chicken smooth muscle regulatory, phosphorylatable myosin light chain (MLC_2) was isolated from a chicken gizzard cDNA library constructed in lambda gt10. The 1075 base pair clone (including poly A tail) was obtained by screening the library with a rat aorta smooth muscle MLC_2 cDNA obtained from Dr. Mark Taubman(1). Prior to obtaining this cDNA, two sequences had been published from amino acid analysis of chicken smooth muscle MLC_2 tryptic peptides which differed at the amino terminus(2,3). The coding region (deduced from the cDNA sequence presented here, initiation and termination codons boxed) predicts a 171 amino acid MLC_2 , which is identical to that obtained by Pearson et al.(2) from analysis of the purified chicken gizzard protein over the first seventeen amino acids containing the phosphorylation site. The remainder of the predicted protein is identical to that obtained by Maita et al.(3). Although the DNA sequence for the MLC_2 coding region shares 82% homology with the rat cDNA(1), the proteins encoded by the DNAs are 94% homologous. Since most of the DNA changes are at third base positions (and the untranslated regions are divergent) there is a strong evolutionary conservation pressure to maintain the smooth muscle MLC_2 protein sequence.

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100 GCTGTCACCT TCCCCCAAC GCACTGAGGC CGGATCCCCC CCCCAGGTAA CCACCCAGC TGCAACATG TCCAGCAAAC GTGCCAAAGC CAAGACCACCC
200 AAGAACGGCC CGCAGCGC GCACCTCAAT GTCTTCGCTA TGTTGACCA GTCACAGATC CAGGAGTTCA AGGAAGCTTT CAACATGATC GACCAGAACCC
300 GTGACGGGGT CATTGACAAG GAGGATCTGC ATGACATGCT GGCTTCCATG GGGAAAAAACCC CACCGACGA GTACCTGGAG GGGATGATGA GTGAGGCCACCC
400 GGGGCCATC AACTTCACCA TGTTCTCAC CATGTTGGG GAGAAGCTGA ATGGCACCGA CCCGGAGGAT GTAATCCGCA ATGCCTTGC CTGCTTGCAC
500 GAGGAGGCGT CAGGGTCTAT TCACGAGGAC CATCTGCCTG AACTGCTGAC CACCATGGGA GACAGGTTCA CTGACGAGGA GGTGGACGAG ATGTAACCGGG
600 AGGCCCCAT CGACAAGAAG GGCAACTTCA ACTATGTGGA GTTCACCCGC ATCCCTGAAGC ACAGGAGCTAA GGACAAGGAC GATTAGCT GAGAGCCGCC
700 CCCCTGCCTT CCCGCACGTC CCACCCACGC TCCCGCACGT GCCACCCACG CCCCTGCCCC TCCTGCCTCA GCCCCACCAT AGGCTCCCCC TGCCGGGACCA
800 GCCCCGTCCT TGCTGGTGC TGCCATCGCG TGCTCTCCCA TTGCCCCGGG ATCCCAGCCA GGCTCCCCC CACGACGCC GCTGGCTCAC
900 CCTGCTCAGC AGAGGGTCCA GACAGGGGCA GGCAGCCGGG GAGTGCTGGC CGGGGGAGTT CCCGCTGGCT CAAAGCAGTG AACCTCTCC AGAGGACCTA
1000 ACACAGAAGG AAGCTCTCCC TTCTCCACA CTCCCTTCT TAGGAAAGAA AGAAAAGAAC TTTGTCTTC CCCTCTTGG CTGTTTATG GCTTTAGAGC
CTGTGATCTA CGGGATTCAAG AAAGCTGAAG CAGCCTATAA AGTCTGATGG GTGTGACAAA AAAAAAAA AAAAAA

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REFERENCES

1. Taubman, M. B., Grant, J. W. and Nadal Ginard, B. (1987) J. Cell. Biol 104:1505-1513.
2. Pearson, R. B., Jakes, R., John, M., Kendrick-Jones, J. and Kemp, B. E. (1984) FEBS Lett. 168:108-112.
3. Maita,T., Chen, J. I. and Matsuda, G. (1981) Eur. J. Biochem 117:417-424.