

The nucleotide sequence of a human smooth muscle alpha-actin (aortic type) cDNA

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To isolate the human smooth muscle (SM) alpha-actin (aortic type) cDNA, we constructed the cDNA library from the artery of umbilical cord. By using the human SM alpha-actin specific probe containing 3' untranslated (UT) region (1), we got a clone (pHSM α A-2) which is 1345 bp long with 5' and 3' UT regions of 47 bp and 149bp, respectively. The amino acid sequence deduced from the nucleotide sequence is composed of 377 amino acid residues containing an initial Met-Cys dipeptide, which is removed post-translationally. The remainder of the amino acid sequence matched those of bovine, rat and chicken SM alpha-actins (2,3,4). In 5' UT region, the 23 bp region immediately preceding the initiation codon is identical with the corresponding sequence of the human SM alpha-actin gene (5), but the initial 24bp region did not match with the corresponding genomic DNA. From this result, the human SM alpha-actin gene has at least one more non-coding exons in 5' upstream region. In 5' UT region, there are 83% sequence conservation between human and mouse genes (6), and 76% between human and rat genes (3). Although the human SM alpha-actin 3' UT region did not have a 32bp long G/C rich segment that is present in the rat SM alpha-actin 3' UT region (3), there are 80% sequence conservation between human and rat genes, 75% between human and mouse genes, and 60% between human and chicken genes (3,6,5).

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CGAGCCCGCAACGACCTGTCCAGGAATCCTGTGAAGCAGCTCCAGCTATGTGTGAAGAAGAGGACAGCACTCCCTGGTGTGTGACAATGGCTCTGGCCCT 100
MetCysGluGluGluAspSerThrAlaLeuValCysAspAsnGlySerGlyLe
CTGTAAAGCCGGCTTTGCTGGGGACGATGCTCCCAAGGGCTGTTTTCCCATCCATTGTGGGACGTCCAGACATCAGGGGGTGTGTGGGAATGGGACAA 200
uCysLysAlaGlyPheAlaGlyAspAlaProArgAlaValPheProSerIleValGlyArgProArgHisGlnGlyValMetValGlyMetGlyGln
AAAGACAGCTACCTGGGTGACGAAGCAGACAGCAAAGAGGAATCCTGACCCCTGAAGTACCCGATAGAATCGGCATCACCAACTGGGACGACATGG 300
LysAspSerTyrValGlyAspGluAlaGlnSerLysArgGlyIleLeuThrLeuLysTyrProIleGluHisGlyIleIleThrAsnTrpAspAspMetG
AAAAGATCTGGCACCCTCTTCTACAATGAGCTTCGTGTGGCCCTGAAGAGCATCCCACTGCTCAGGAGGCCACCCCTGAACCCCAAGGCCAACCC 400
luLysIleTrpHisHisSerPheTyrAsnGluLeuArgValAlaProGluGluHisProThrLeuLeuThrGluAlaProLeuAsnProLysAlaAsnAr
GGAGAAAATGACTCAAAATATGTTTGAAGCTTTCAATGTCCAGCCATGATGTGGCTATCCAGGCGGTGCTGTCTCTATGGCTCTGGACGCCAACCT 500
gGluLysMetThrGlnIleMetPheGluThrPheAsnValProAlaMetTyrValAlaIleGlnAlaValLeuSerLeuTyrAlaSerGlyArgThrThr
GGCATCTGTGGACTCTGGAGATGGTGTCAACCCACATGCTCCCATCTATGAGGGCTATGCCTTGCCCATGCCATCATGGCTGTGGATCTGGCTGGCC 600
GlyIleValLeuAspSerGlyAspGlyValThrHisAsnValProIleTyrGluGlyTyrAlaLeuProHisAlaIleMetArgLeuAspLeuAlaGlyA
GAGATCTCACTGACTACCTCATGAAGATCCTGACTGAGCGTGGCTATTCTCTGTTACTACTGCTGAGCCTGAGATGTGTCGGGACATCAAGGAGAACT 700
rAspLeuThrAspTyrLeuMetLysIleLeuThrGluArgGlyTyrSerPheValThrThrAlaGluArgGluIleValArgAspIleLysGluLysLe
GTGTTATGTAGCTCTGGACTTTGAAAATGAGATGGCCATGCCGCATCCTCATCCTCCCTGAGAAGAGTTACGAGTTGCCTGATGGGCAAGTGATCACC 800
uCysTyrValAlaLeuAspPheGluAsnGluMetAlaThrAlaAlaSerSerSerSerLeuGluLysSerTyrGluLeuProAspGlyGlnValIleThr
ATCGGAAATGAACGTTTCGCTGCCAGAGACCCCTGTCAGCCATCCTCATCGGGATGGAGTCTGCTGGCATCCATGAAACCCACTACAACAGCATCA 900
IleGlyAsnGluArgPheArgCysProGluThrLeuPheGlnProSerPheIleGlyMetGluSerAlaGlyIleHisGluThrThrTyrAsnSerIleM
TGAAGTGTGATATTGACATCAGGAAGGACCTCTATGCTAACAATGCTCTATCAGGGGGCACCCTATGTACCTGGCATTCGGACCGCAATGCAGAAGGA 1000
eLysCysAspIleAspIleAspLysAspLeuTyrAlaAsnAsnValLeuSerGlyGlyThrThrMetTyrProGlyIleAlaAspArgMetGlnLysGl
GATCAGGCCCTAGCACCCAGCACCATGAAGATCAAGATCATTGCCCTCCGGAGCCGCAAAATCTCTGTCTGGATCGGTGCTCCATCTGGCTCTCTG 1100
uIleThrAlaLeuAlaProSerThrMetLysIleLysIleIleAlaProProGluArgLysTyrSerValTrpIleGlyLysSerIleLeuAlaSerLeu
TCCACCTCCAGCAGATGGATCAGAAACAGGAATACGATGAAGCCGGCCCTCCATTGTCCACCCGCAATGTCTCTAAACACTTTCCTGCTCCTCT 1200
SerThrPheGlnGlnMetTrpIleSerLysGlnGluTyrAspGluAlaGlyProSerIleValHisArgLysCysPheEnd
CTGTCTAGCACACAAGCTGTGAATGCTGTTGGAATTATGCCTTCAGTCTCTTTCCAAATCATTCTAGCCAAAGCTCTGACTCGTTACCTATGTGTTT 1300
TTTAATAAATCTGAAATAGGCTACTGGTAAAAAATAAAAAAAAAA 1345

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