

Nucleotide sequence of a cDNA coding for mouse cyclophilin

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Cyclophilin is an abundant cytosolic protein that is expressed in most tissues and is found in all eukaryotic species examined, including yeast and man (1, 2). The immunosuppressive drug Cyclosporin A has been shown to bind cyclophilin with high affinity (1). The porcine enzyme peptidyl-prolyl cis-trans isomerase, which is inhibited by Cyclosporin A, is identical to the bovine cyclophilin (3, 4). Cyclophilin may represent the intracellular target for Cyclosporin A-mediated immunosuppression. Reported here is the nucleotide sequence of a clone of cyclophilin derived from a cDNA library made from mouse thymus mRNA. The sequence, as determined by primer extension and direct RNA sequencing methods, represents a full-length mRNA except the first nucleotide. The deduced mouse protein sequence shows extremely high identity to all mammalian cyclophilins of known sequence: 96.3%, 95.7%, 95.7%, 97.6%,

and 98.8% amino acid identity with the human (5), bovine (2), porcine (3, 4), rat (6), and hamster (7) sequences, respectively.

REFERENCES

1. Handschumacher, R.E., Harding, M.W., Rice, J., Drugge, R.J. and Speicher, D.W. (1984) *Science* **226**, 544–547.
2. Harding, M.W., Handschumacher, R.E. and Speicher, D.W. (1986) *J. Biol. Chem.* **261**, 8547–8555.
3. Takahashi, N., Hayano, T. and Suzuki, M. (1989) *Nature* **337**, 473–475.
4. Fisher, G., Wittmann-Liebold, B., Lang, K., Kiefhaber, T. and Schmid, F.X. (1989) *Nature* **337**, 476–478.
5. Haendler, B., Hofer-Warbinek, R. and Hofer, E. (1987) *EMBO J.* **6**, 947–950.
6. Danielson, P.E., Forss-Petter, S., Brow, M.A., Calavetta, L., Douglass, J., Milner, R.J. and Sutcliffe, J.G. (1988) *DNA* **7**, 261–267.
7. Bergsma, D.J. and Sylvester, D. (1990) *Nucl. Acids Res.* **18**, 200.

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TTTGCAGACGCCACTGTCGCTTTTCGCCGCTTGCTGCAGCCATGGTCAACCCACCGTGTCTTCGACATC 71
METValAsnProThrValPhePheAspIle 10

ACGCCGATGACGAGCCCTTGGGCCGCGTCTCCTTCGAGCTGTTTGACAGACAAAGTTCCAAAGACAGCAGAA 143
ThrAlaAspAspGluProLeuGlyArgValSerPheGluLeuPheAlaAspLysValProLysThrAlaGlu 34

AACTTTCGAGCTCTGAGCACTGGAGAGAAAGGATTTGGCTATAAGGGTTCCTCCTTTCACAGAATTATTCCA 215
AsnPheArgAlaLeuSerThrGlyGluLysGlyPheGlyTyrLysGlySerSerPheHisArgIleIlePro 58

GGATTCATGTGCCAGGGTGGTGACTTTACAGCCATAATGGCACTGGCGGCAGGTCCATCTACGGAGAGAAA 287
GlyPheMetCysGlnGlyGlyAspPheThrArgHisAsnGlyThrGlyGlyArgSerIleTyrGlyGluLys 82

TTTGAGGATGAGAACTTCATCCTAAAGCATAACAGGTCCTGGCATCTTGTCCATGGCAAATGCTGGACCAAAC 359
PheGluAspGluAsnPheIleLeuLysHisThrGlyProGlyIleLeuSerMetAlaAsnAlaGlyProAsn 106

ACAAACGGTTCAGTTCCTTTATCTGCACTGCCAAGACTGAATGGCTGGATGGCAAGCATGTGGTCTTTGGG 431
ThrAsnGlySerGlnPhePheIleCysThrAlaLysThrGluTrpLeuAspGlyLysHisValValPheGly 130

AAGGTGAAAGAAGGCATGAACATTGTGGAAGCCATGGAGCGTTTTGGGTCCAGGAATGGCAAGACCAGCAAG 503
LysValLysGluGlyMetAsnIleValGluAlaMetGluArgPheGlySerArgAsnGlyLysThrSerLys 154

AAGATCACCATTTCCGACTGTGGACAGCTCTAATTTCTTTGACTTGCGGGCATTTCACCCATCAAACCATT 575
LysIleThrIleSerAspCysGlyGlnLeu*** 164

CCTTCTGTAGCTCAGGAGAGCGTCCCTACCCCATCTGCTCGCAATGTCCTGTAATCTCTGCTCTCACTGAAG 647
TTCTTTGGGTCCATATTTTCTCATTCCCTTCAAGTCTAGCTGGATTGCAAAGTTAAGTTTATGATTATG 719
AATAAAAATAAATAAGAAAAAAAAAAAAAAAAAAAAA 736

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