

Methods in Molecular Biology™

VOLUME 206

Peptide Research Protocols

Endothelin

Edited by

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Anthony P. Davenport**



HUMANA PRESS

Peptide Research Protocols Endothelin Methods In Molecular Biology

Yijin Wang



Peptide Research Protocols Endothelin Methods In Molecular Biology:

Peptide Research Protocols Janet J. Maguire, Anthony P. Davenport, 2008-02-04 A panel of multidisciplinary experts describes in detail readily reproducible methods to investigate all aspects of the endothelin system from its synthesis and metabolism to its function in health and disease These methods use state of the art molecular techniques to quantify the expression of mRNA for both endothelin receptors and the endothelin converting enzymes They show how peptides precursors receptors and synthetic enzymes can be localized and quantified in plasma culture supernatants tissue homogenate and tissue sections using antibodies Several in vivo protocols illustrate the role of the endothelin peptides in healthy human individuals and describe animal models that can be used to predict the therapeutic potential of cardiovascular drugs that manipulate endothelin synthesis or function *Methods in Molecular Biology: Peptide research protocols: endothelin* John M. Walker, 1984 *The Pharmacology of Functional, Biochemical, and Recombinant Receptor Systems* T. Kenakin, J. A. Angus, 2012-12-06 This the 148th volume of the Handbook of Experimental Pharmacology series focuses on the very core of pharmacology namely receptor theory It is fitting that the originator of receptor pharmacology A J CLARK authored the fourth volume of this series 63 years ago In that volume CLARK further developed his version of receptor theory first described four years earlier in his classic book *The Mode of Action of Drugs* An examination of the topics covered in volume 4 reveals a striking similarity to the topics covered in this present volume pharmacologists today are still as interested in unlocking the secrets of dose response relationships to reveal the biological and chemical basis of drug action as they were over half a century ago Sections in that 1937 volume such as *Curves relating exposure to drugs with biological effects* and *Implications of monomolecular theory* show Clark's keen insight into the essential questions that required answers to move pharmacology forward With the advent of molecular biological cloning of human receptors has come a transformation of receptor pharmacology Thus the expression of human receptors into surrogate host cells helped unlock secrets of receptor mechanisms and stimulus transduction pathways To a large extent this eliminates the leap of faith required to apply receptor activity of drugs tested on animal receptor systems to the human therapeutic arena However a new leap of faith concerning the veracity of the effects found in recombinant systems with respect to natural ones is now required **Peptide Research Protocols** Janet J. Maguire, Anthony P. Davenport, 2002-06-25 A panel of multidisciplinary experts describes in detail readily reproducible methods to investigate all aspects of the endothelin system from its synthesis and metabolism to its function in health and disease These methods use state of the art molecular techniques to quantify the expression of mRNA for both endothelin receptors and the endothelin converting enzymes They show how peptides precursors receptors and synthetic enzymes can be localized and quantified in plasma culture supernatants tissue homogenate and tissue sections using antibodies Several in vivo protocols illustrate the role of the endothelin peptides in healthy human individuals and describe animal models that can be used to predict the therapeutic potential of cardiovascular

drugs that manipulate endothelin synthesis or function **Atomic Force Microscopy** Pier Carlo Braga, Davide Ricci, 2008-02-02 The natural biological medical and related sciences would not be what they are today without the microscope After the introduction of the optical microscope a second breakthrough in morphostructural surface analysis occurred in the 1940s with the development of the scanning electron microscope SEM which instead of light i e photons and glass lenses uses electrons and electromagnetic lenses magnetic coils Optical and scanning or transmission electron microscopes are called far field microscopes because of the long distance between the sample and the point at which the image is obtained in comparison with the wavelengths of the photons or electrons involved In this case the image is a diffraction pattern and its resolution is wavelength limited In 1986 a completely new type of microscopy was proposed which without the use of lenses photons or electrons directly explores the sample surface by means of mechanical scanning thus opening up unexpected possibilities for the morphostructural and mechanical analysis of biological specimens These new scanning probe microscopes are based on the concept of near field microscopy which overcomes the problem of the limited diffraction related resolution inherent in conventional microscopes Located in the immediate vicinity of the sample itself usually within a few nanometers the probe records the intensity rather than the interference signal thus significantly improving resolution Since the most we known microscopes of this type operate using atomic forces they are frequently referred to as atomic force microscopes AFMs **Posttranslational Modification of Proteins** Christoph Kannicht, 2008-02-04 Christoph Kannicht and a panel of highly experienced researchers describe readily reproducible methods for detecting and analyzing the posttranslational modifications of protein particularly with regard to protein function proteome research and the characterization of pharmaceutical proteins **Subject Guide to Books in Print**, 1991 **Forthcoming Books** Rose Arny, 2002 **Biochemical and Organic Compounds for Research and Diagnostic Clinical Reagents** Sigma Chemical Company, 1994 **The British National Bibliography** Arthur James Wells, 2003 *Innovation and Perspectives in Solid Phase Synthesis & Combinatorial Libraries*, 1998 Roger Epton, 1999 Investigative Ophthalmology & Visual Science, 1999 **Abstracts, 24th Annual Meetings, April 2-23, 1995**, 1995 Molecular Biology of the Cell, 2004 Proceedings of the National Academy of Sciences of the United States of America, 2005 *Abstracts, 20th Annual Meetings, April 6-April 25, 1991*, 1991 Doody's Rating Service Daniel J. Doody, 1996 *Current Opinion in Biotechnology*, 1992 *Cancer Research*, 2007-11 **Canadian Journal of Microbiology**, 2002-09

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