

Natural Killer Cell Protocols

Cellular and Molecular Methods
Second Edition

Edited by

Kerry S. Campbell

 Humana Press

Natural Killer Cell Protocols Cellular And Molecular Methods

Ambikaipakan Balasubramaniam



Natural Killer Cell Protocols Cellular And Molecular Methods:

Natural Killer Cell Protocols Kerry S. Campbell, Marco Colonna, 2008-02-03 In *Natural Killer Cell Protocols Cellular and Molecular Methods* Kerry S Campbell and Marco Colonna have assembled a comprehensive collection of readily reproducible methods designed to study natural killer NK cells from the broadest variety of viewpoints These include not only classic techniques but also new approaches to standard methods newly evolved techniques that have become valuable for specific applications and unique models for manipulating and studying NK cells Among the advanced methods covered are those for in vitro transendothelial migration in vivo detection of cells migrating into tumors immunofluorescence staining of intracellular cytokines and in vitro NK cell development Valuable techniques for specific applications include vaccinia virus protein expression soluble KIR Fc fusions for HLA class I binding assays calcium mobilization in cell conjugates and identification of heterodimeric receptor complexes using cDNA library expression cloning No less important are accounts of such classic methods as hybrid resistance ADCC viral defense target cell cytotoxicity assays cloning and culturing tumor immunotherapy and generation of HLA class I transfected target cells *Natural Killer Cell Protocols Cellular and Molecular Methods* offers immunologists cancer researchers virologists and cell biologists today's most comprehensive collection of both established and cutting edge techniques methods that will contribute significantly to advancing our understanding of this fascinating and critically important class of cells

Extracellular Matrix Protocols Charles Streuli, Michael Grant, 2008-02-05 It is now widely accepted that much of the dynamic function of cells and tissues is regulated from outside the cell by the extracellular matrix In addition to its conventional role in providing a scaffold for building tissues the extracellular matrix acts as a directional highway for cellular movement and provides instructional information for promoting survival proliferation and differentiation Indeed the extracellular matrix is beginning to take a starring role in the choreography of cell and tissue function The diverse roles of the extracellular matrix are reflected in its highly complicated structure consisting of an ever increasing number of components Yet the mechanisms of extracellular matrix assembly and how they influence cell behavior are only just beginning to be understood In order to solve these problems new methodologies are of necessity being developed Many of these technologies are highly sophisticated and are currently available only in a handful of laboratories However we believe that they can readily be transported and established by other researchers Thus the purpose of *Extracellular Matrix Protocols* is to present some of these complicated techniques in a style that is relatively easy to reproduce

Neuropeptide Y Protocols Ambikaipakan Balasubramaniam, 2008-02-05 The observation that neuropeptide Y NPY is the most abundant peptide present in the mammalian nervous system and the finding that it elicits the most powerful orexigenic signal have led to active investigations of the properties of the NPY family of hormones including peptide YY PYY and pancreatic polypeptide PP Nearly two decades of research have led to the identification of several NPY receptor subtypes and the development of useful receptor selective ligands Moreover these

investigations have implicated NPY in the pathophysiology of a number of diseases including feeding disorders seizures memory loss anxiety depression and heart failure Vigorous efforts are therefore continuing not only to understand the biochemical aspects of NPY actions but also toward developing NPY based treatments for a variety of disorders To facilitate these efforts it was decided to produce the first handbook on NPY research techniques as part of the Methods in Molecular Biology Series In compiling Neuropeptide Y Protocols I have gathered contributions on techniques considered critical for the advancement of the NPY field from experts in various disciplines Each chapter starts with a brief introduction with Materials and Methods sections following The latter sections are presented in an easy to follow step by step format The last section of the chapter Notes highlights pitfalls and the maneuvers employed to overcome them This information not usually disseminated in standard research publications may prove extremely useful for investigators employing these techniques in NPY research Eicosanoid Protocols Elias A Lianos,2008-02-03 **NMDA Receptor Protocols** Min Li,2008-02-03 Min Li and a panel of hands on experimentalists detail state of the art molecular techniques for studying NMDA ligand gated ion channels and developing assays for nontherapeutic lead selection The topics range from cDNA cloning to in vitro and in vivo investigation of the channel complex in the mammalian brain Additional topics include the biochemical analysis of the channel protein and the construction of various heterologous systems for both basic research and high throughput screens HTS for pharmaceutical chemicals Although the focus is on NMDA receptors the methods are applicable to other ligand gated ion channels and with some modification may be extended to related membrane signaling receptors NMDA Receptor Protocols offers today's scientists powerful methods for basic research on NMDA receptor structure and function as well as enormous opportunities for clinical investigation toward the development of novel bioactive compounds

Immunocytochemical Methods and Protocols Lorette C. Javois,2008-02-02 Lorette Javois timely new 2nd edition revises and updates her widely acclaimed collection of step by step immunocytochemical methods one that is now used in many biological and biomedical research programs The methods are designed for researchers and clinicians who wish to visualize molecules in plant or animal embryos tissue sections cells or organelles In addition to cutting edge protocols for purifying and preparing antibodies light microscopic analysis confocal microscopy FACS and electron microscopy this revised edition contains many new methods for applying immunocytochemical techniques in the clinical laboratory and in combination with in situ hybridization Chromatin Protocols Peter B. Becker,2008-02-03 More than 40 years after the discovery of the nucleosome as the fundamental unit of chromatin the multifaceted problem of how variations in chromatin structure affect the activity of the eukaryotic genome has not been solved However during the past few years research on chromatin structure and function has gained considerable momentum and impressive progress has been made at the level of concept development as well as filling in crucial detail The structure of the nucleosome has been visualized at unprecedented resolution Powerful multisubunit enzymes have been identified that alter histone DNA interactions in ways that expose

regulatory sequences to factors initiating and regulating such nuclear processes as transcription. Though the importance of posttranslational modifications of histones, notably their acetylation, has long been known, the finding that a number of bona fide regulators increase transcription by acetylating nucleosomes has lent new support to the old idea that the process of gene regulation is intimately related to the nature of the chromatin environment. A wealth of nonhistone proteins contribute to a continuum of structures with distinct biochemical properties and varying degrees of DNA condensation. Perhaps the most important conclusion from a large number of studies is a fresh appreciation of the dynamic nature of chromatin structure, the built-in flexibility providing the basis for regulation.

Calpain Methods and Protocols John S. Elce, 2008-02-05 The purpose of *Calpain Methods and Protocols* is quite straightforward: it is to present the actual experimental methods used in many different laboratories for the study of calpain. It will provide the vital experimental detail and the discussion of possible pitfalls for which the standard journals no longer provide space. This will make it as easy as possible for investigators interested in calpain to adopt established methods without repeating old mistakes and to adapt and apply these methods in novel approaches to the many outstanding calpain questions. These questions range from purely biochemical problems of protein structure and enzyme regulation at the molecular level through large areas of cell biology to applied and clinical aspects of calpain function in human disease. Within this panoply of topics, a wide range of investigators will find many fascinating and as yet unanswered questions about calpain. *Calpain Methods and Protocols* will provide instant access to many essential techniques while saving them the time and effort involved in developing a new method. In addition to questions relating to the normal physiological roles of the calpains, there is considerable evidence that inappropriate calpain activity may have pathological effects in many tissues, for example following ischemia. This provides a major stimulus for the development of specific calpain inhibitors for therapeutic purposes and for the development of methods to evaluate such inhibitors.

Molecular Methods in Developmental Biology Matt Guille, 2008-02-03 The process whereby a single cell, the fertilized egg, develops into an adult has fascinated for centuries. Great progress in understanding that process has never been made in the last two decades when the techniques of molecular biology have become available to developmental biologists. By applying these techniques, the exact nature of many of the interactions responsible for forming the body pattern are now being revealed in detail. Such studies are a large and ever-expanding part of most life science groups. It is at newcomers to this field that this book is primarily aimed. A number of different plants and animals serve as common model organisms for developmental studies. In *Molecular Methods in Developmental Biology*, *Xenopus* and zebrafish, a range of the molecular methods applicable to two of these organisms are described; these are the South African clawed frog *Xenopus laevis* and the zebrafish *Brachydanio rerio*. The embryos of both of these species develop rapidly and externally, making them particularly suited to investigations of early vertebrate development. However, both *Xenopus* and zebrafish have their own advantages and disadvantages. *Xenopus* has large, robust embryos that can be manipulated surgically with ease, but their pseudotetraploidy

and long generation time make them unsuitable candidates for genetics This disadvantage may soon be overcome by using the diploid *Xenopus tropicalis* and early experiments are already underway The transparent embryos of zebrafish render them well suited for in situ hybridization and immunohistochemistry and good for observing mutations in genetic screens

Chemokine Protocols Amanda E.I. Proudfoot, Timothy N.C. Wells, Christine Power, 2008-02-05 The chemokines family of small proteins are involved in numerous biological processes ranging from hematopoiesis angiogenesis and basal leukocyte trafficking to the extravasation and tissue infiltration of leukocytes in response to inflammatory agents tissue damage and bacterial or viral infection Chemokines exert their effects through a family of seven G protein coupled transmembrane receptors Worldwide interest in the chemokine field surged dramatically early in 1996 with the finding that certain chemokine receptors were the elusive coreceptors required along with CD4 for HIV infection Today though over 40 human chemokines have been described the number of chemokine receptors lags behind only 17 human chemokine receptors have been identified so far What has emerged over the years is that most chemokine receptors bind several distinct ligands and indeed the majority of chemokines are able to bind to multiple chemokine receptors explaining to some extent the apparent disparity in the numbers of chemokines and receptors Yet in spite of the apparent redundancy in chemokine chemokine receptor interactions it is clear that in vivo spatial temporal and indeed cell and tissue specific expression of both chemokines and their receptors are important factors in determining the precise nature of cellular infiltrates in physiological and pathological processes

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