

RNA Silencing

Methods and Protocols

Edited by

Gordon G. Carmichael



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Rna Silencing Methods And Protocols Methods In Molecular Biology

Sandra J Rosenthal, David W Wright



Rna Silencing Methods And Protocols Methods In Molecular Biology:

RNA Silencing Gordon Carmichael, 2008-02-04 A collection of readily reproducible methods for the design preparation and use of RNAs for silencing gene expression in cells and organisms The techniques range widely and include methods addressing the biochemical aspects of the silencing machinery RNA silencing in non mammalian organisms and the in vivo delivery of siRNAs and silencing vectors There are also techniques for designing preparing and using RNAs to silence gene expression for fine tuning regulation by targeting specific isoforms of a given gene and for the study and use of microRNAs The protocols follow the successful *Methods in Molecular Biology*TM series format each offering step by step laboratory instructions an introduction outlining the principle behind the technique lists of the necessary equipment and reagents and tips on troubleshooting and avoiding known pitfalls *Phosphodiesterase Methods and Protocols* Claire Lugnier, 2008-02-04 Research leaders in the PDE field describe new concepts and techniques for investigating the role of PDEs in orchestrating normal and pathophysiological responses Presented in step by step detail these readily reproducible methods allow the measurement of cyclic nucleotide variations in living cells as well as their visualization in a spatio temporal manner the localization and characterization of their activities in tissues and living cells and the assessment of targeted PDEs in creating specific tools and drugs *Differential Display Methods and Protocols* Peng Liang, Jonathan Meade, Arthur B. Pardee, 2008-02-04 Since the first edition of this book dedicated to differential display DD technology was published in 1997 we have witnessed an explosive interest in studying differential gene expression The gene hunting euphoria was initially powered by the invention of DD which was gradually overtaken by DNA microarray technology in recent years Then why is there still the need for second edition of this DD book First of all DD still enjoys a substantial lead over DNA microarrays in the ISI citation data see Table 1 despite the hundreds of millions of dollars spent each year on arrays This may come as a surprise to many but to us it implies that many of the DNA microarray studies went unpublished owing to their unfulfilled promises 1 Second unlike DNA microarrays DD is an open ended gene discovery method that does not depend on prior genome sequence information of the organism being studied As such DD is applicable to the study of all living organisms from bacteria fungi insects fish plants to mammals even when their genomes are not sequenced Second DD is more accessible technically and financially to most cost conscious cottage industry academic laboratories So clearly DD still has its unique place in the modern molecular biological toolbox for gene expression analysis **NanoBiotechnology Protocols** Sandra J Rosenthal, David W Wright, 2008-02-04 Hands on experts in nanomaterial synthesis and application describe in detail the key experimental techniques currently employed in novel materials synthesis dynamic cellular imaging and biological assays The author s emphasize diverse strategies to synthesize and functionalize the use of nanoparticles for biological applications Additional chapters focus on the use of biological components peptides antibodies and DNA to synthesize and organize nanoparticles to be used a building block in larger assemblies These new materials make it possible to image

cellular processes for longer durations leading to high throughput cellular based screens for drug discovery drug delivery and diagnostic applications Highlights include overview chapters on quantum dots and DNA nanotechnology and cutting edge techniques in the emerging nanobiotechnology arena **Amyloid Proteins** Einar M. Sigurdsson,2008-02-02 A proven collection of readily reproducible techniques for studying amyloid proteins and their involvement in the etiology pathogenesis diagnosis and therapy of amyloid diseases The contributors provide methods for the preparation of amyloid and its precursors oligomers and protofibrils in vitro assays and analytical techniques for their study and cell culture models and assays for the production of amyloid proteins Additional chapters present readily reproducible techniques for amyloid extraction from tissue its detection in vitro and in vivo as well as nontransgenic methods for developing amyloid mouse models The protocols follow the successful Methods in Molecular Biology™ series format each offering step by step laboratory instructions an introduction outlining the principle behind the technique lists of the necessary equipment and reagents and tips on troubleshooting and avoiding known pitfalls **Receptor Binding Techniques** Anthony P. Davenport,2008-02-02 A comprehensive collection of readily reproducible methods for studying receptors in silico in vitro and in vivo These cutting edge techniques cover mining from curated databases identifying novel receptors by high throughput screening molecular methods to identify mRNA encoding receptors radioligand binding assays and their analysis quantitative autoradiography and imaging receptors by positron emission tomography PET Highlights include phenotypic characterization of receptors in knockout mice imaging receptors using green fluorescent protein and fluorescent resonance energy transfer and quantitative analysis of receptor mRNA by TaqMan PCR These book equips the researcher with techniques for exploring the unprecedented number of new receptor systems now emerging and the so called orphan receptors whose activating ligand has not been identified **Cytochrome P450 Protocols** Ian R. Phillips,2008-02-04 For this second edition of their much praised Cytochrome P450 the editors have collected accounts of the essential core techniques that use the latest methodologies for the investigation of P450s Highlights include protocols for spectral analysis and purification of P450s enzymatic assays of P450s and flavin containing monooxygenases FMOs expression of P450s and FMOs in heterologous systems and the production and use of antipeptide antibodies Additional chapters contain readily reproducible techniques for the transfection of hepatocytes for gene regulation studies P450 reporter gene assays in situ hybridization and analysis of genetic polymorphisms Although the emphasis is on P450s of mammalian origin many of the readily reproducible methods described are suitable for P450s from any source **Chemical Genomics** Edward D. Zanders,2008-02-04 Chemical genomics is an exciting new field that aims to transform biological chemistry into a high throughput industrialized process much in the same way that molecular biology has been transformed by genomics The interaction of small organic molecules with biological systems mostly proteins underpins drug discovery in the pharmaceutical and biotechnology industries and therefore a volume of laboratory protocols that covers the key aspects of chemical genomics

would be of use to biologists and chemists in these organizations Academic scientists have been exploring the functions of proteins using small molecules as probes for many years and therefore would also benefit from sharing ideas and laboratory procedures Whatever the organizational backgrounds of the scientists involved the challenges of extracting the maximum human benefit from genome sequencing projects remains considerable and one where it is increasingly recognized that chemical genomics will play an important part Chemical Genomics Reviews and Protocols is divided into two sections the first being a series of reviews to describe what chemical genomics is about and to set the scene for the protocol chapters The subject is introduced by Paul Caron who explains the various flavors of chemical genomics This is followed by Lutz Weber and Philip Dean who cover the interaction between organic molecules and protein targets from the different perspectives of laboratory experimentation and in silico design The protocols begin with the methods developed in Christopher Lowes laboratory Roque et al *Plant Cell Culture Protocols* Víctor M. Loyola-Vargas, Felipe Vázquez-Flota, 2008-02-04 A comprehensive state of the art collection of the most frequently used techniques for plant cell and tissue culture Readily reproducible and extensively annotated the methods range from general methodologies such as culture induction growth and viability evaluation and contamination control to such highly specialized techniques as chloroplast transformation involving the laborious process of protoplast isolation and culture Most of the protocols are currently used in the research programs of the authors or represent important parts of business projects aimed at the generation of improved plant materials Two new appendices explain the principles for formulating culture media and the composition of the eight most commonly used media formulations and list more than 100 very useful internet sites *DNA Repair Protocols* Daryl S. Henderson, 2008-02-03 The first edition of this book published in 1999 and called DNA Repair Protocols Eukaryotic Systems brought together laboratory based methods for studying DNA damage and repair in diverse eukaryotes namely two kinds of yeast a nematode a fruit fly a toad three different plants and human and murine cells This second edition of DNA Repair Protocols covers mammalian cells only and hence its new subtitle Mammalian Systems There are two reasons for this fresh emphasis both of them pragmatic to cater to the interests of what is now a largely mammalocentric DNA repair field and to expedite editing and production of this volume Although DNA Repair Protocols Mammalian Systems is a smaller book than its predecessor it actually contains a greater variety of methods Fourteen of the book's thirty two chapters are entirely new and areas of redundancy present in the first edition have been eliminated here for example now just two chapters describe assays for nucleotide excision repair NER rather than seven All eighteen returning chapters have been revised many of them extensively In order to maintain a coherent arrangement of topics the four part partitioning seen in the first edition was dispensed with and chapters concerned with ionizing radiation damage and DNA strand breakage and repair were recast to near the front of the book Finally an abstract now heads each chapter

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