
Methods in Molecular Biology

Volume 2

Nucleic Acids

Edited by

John M. Walker

Nucleic Acids Vol 2 Methods In Molecular Biology

Nicholas T. Potter



Nucleic Acids Vol 2 Methods In Molecular Biology:

Capillary Electrophoresis of Nucleic Acids Keith R. Mitchelson, Jing Cheng, 2008-02-05 The development of PCR which enables extremely small amounts of DNA to be amplified led to the rapid development of a multiplicity of analytical procedures that permit use of this new resource for the analysis of genetic variation and for the detection of disease causing mutations The advent of capillary electrophoresis CE with its power to separate and analyze very small amounts of DNA has also stimulated researchers to develop analytical procedures for the CE format The advantages of CE in terms of speed and reproducibility of analyses are manifold Furthermore the high sensitivity of detection and the ability to increase sample throughput with parallel analysis has led to the creation of a full range of analysis of DNA molecules from modified DNA adducts and single strand oligonucleotides through PCR amplified DNA fragments and whole chromosomes Capillary Electrophoresis of Nucleic Acids focuses on analytical protocols that can be used for detection and analysis of mutations and modification from precise DNA loci through entire genomes of organisms Important practical considerations for CE such as the choice of separation media electrophoresis conditions and the influence of buffer additives and dyes on DNA mobility are discussed in several key chapters and within particular applications

Capillary Electrophoresis Philippe Schmitt-Kopplin, 2008-02-25 This book presents a selection of current capillary electrophoresis methods used to separate representative types of molecules and particles and in combination with different detection techniques It includes practical details which are hard to find elsewhere The volume is intended for beginners in the field and provides an overview of the technique and a starting point for the exploration of the defined literature on different application topics

Nucleic Acid Probes Robert H. Symons, 1989-03-31 With each chapter written by a distinguished expert in the biochemistry field this comprehensive volume describes the preparation and use of a variety of radioactive and non radioactive probes in situations ranging from research laboratories to routine diagnosis laboratories The enzymatic and chemical techniques for labeling nucleic acid probes with radioisotopes and with non radioactive ligands and haptens are discussed Additionally the associated methods for their detection the use of these probes in the diagnosis of human and microbial pathogens of plant viruses and viroids and of human genetic disorders as well as in the detection of nucleic acids in tissues and cells by in situ hybridization are presented

Peptide Nucleic Acids Peter E. Nielsen, 2008-02-04 Peptide nucleic acids PNAs have now existed for slightly more than ten years with the interest in and applications of this pseudopeptide DNA mimic steadily increasing during the entire period PNAs have rapidly attracted the attention of scientists from a diversity of fields ranging from bio organic and biophysical chemistry to prebiotic evolution and from molecular biology to genetic diagnostics and drug development Many of the applications take advantage of the unique properties of PNA an uncharged pseudopeptide that distinguish this DNA mimic from more traditional DNA analogs Rather than trying to create a comprehensive collection of all published methods and protocols involving PNA many of which have not yet been validated I have decided to concentrate on

select protocols that are either very well established by several groups around the world such as PCR clamping and in situ hybridization or on new methods that may have broader future impact Basic methods for PNA oligomer synthesis and analyses have also been included I am very grateful to those friends and colleagues who have enthusiastically contributed their work discussions and writing and thereby made this book possible Peter E Nielsen v Contents Preface v Contributors ix IINTRODUCTION 1 PNA Technology Peter E Nielsen 3 II CHEMISTRY 2 Solid Phase Synthesis of PNA Oligomers Frederik Beck 29 3 Synthesis of PNA Peptide Conjugates Satish Kumar Awasthi and Peter E Nielsen 43 4 Parallel Synthesis of PNA Peptide Conjugate Libraries Satish Kumar Awasthi and Peter E Nielsen *The ELISA Guidebook* John R.

Crowther, 2008-02-04 John R Crowther provides today's premier practical guide to the understanding and application of ELISA Updating and greatly expanding his widely appreciated earlier publication *ELISA Theory and Practice* 1995 this important work introduces chapters on such major new topics as checkerboard titrations quality control of testing kit production and control novel monoclonal antibodies validation of assays statistical requirements for data examination and epidemiological considerations With its numerous worked examples detailed instructions and extensive illustrations *The ELISA Guidebook* offers a powerful synthesis of all the basic concepts and practical experimental details investigators need to understand develop and apply the new ELISA methodology successfully in day to day basic and clinical research

Protocols for Nucleic Acid Analysis by Nonradioactive Probes Elena Hilario, John F. MacKay, 2008-02-02 *Protocols for Nucleic Acid Analysis by Non radioactive Probes* Second Edition provides a firm background on the basic preparative protocols required for the analysis of nucleic acids by nonradioactive methods Presenting the methodologies using amazing new applications this volume offers guide chapters on nucleic acid extractions preparation of nucleic acid blots and labeling of nucleic acids with nonradioactive haptens New fluorescent techniques such as Real Time PCR and microarrays are also included allowing users to get a nonradioactive protocol implemented in the laboratory with minimum adaptation required and fastest time to results The protocols follow the successful *Methods in Molecular Biology*™ series format each offering step by step laboratory instructions an introduction outlining the principles behind the technique lists of the necessary equipment and reagents and tips on troubleshooting and avoiding known pitfalls **MHC Volume 1** N. Fernandez, G.

Butcher, 1997-08-21 The major histocompatibility complex is the subject of much research in the immunology area There is great interest in the structure of MHC proteins and in their function as antigen presenting molecules Many immunology labs are now faced with the prospect of tackling biochemical and genetic techniques in order to study these molecules and these two Practical Approach volumes provide the ideal introduction MHC 1 Antigen Processing and Presentation is complemented by MHC 2 Genes Cells and Molecules **Neurogenetics** Nicholas T. Potter, 2008-02-02 The rapid identification and characterization of genes of neurological relevance holds great potential for offering insight into the diagnosis management and understanding of the pathophysiologic mechanisms of neurological diseases This volume in the *Methods in Molecular*

BiologyTM series was conceived to highlight many of the contemporary methodological approaches utilized for the characterization of neu logically relevant gene mutations and their protein products Although an emphasis has been placed upon descriptions of methodologies with a defined clinical utility it is hoped that Neurogenetics Methods and Protocols will appeal not only to clinical laboratory diagnosticians but also to clinicians and to biomedical researchers with an interest in advances in disease diagnosis and the functional consequences of neu logically relevant gene mutations To meet this challenge more than 60 authors graciously accepted my invitation to contribute to the 32 chapters of this book Through their collective commitment and diligence what has emerged is a comprehensive and timely treatise that covers many methodological aspects of mutation detection and screening including disc sions on quantitative PCR trinucleotide repeat detection sequence based mutation detection molecular detection of imprinted genes fluorescence in situ hybridization FISH in vitro protein expression systems and studies of protein expression and function I would like to take this opportunity to formally thank my colleagues for their effort and dedication to this work

In Vitro Mutagenesis Protocols Jeff Braman,2008-02-05 Hands on researchers with proven track records describe in stepwise fashion their advanced mutagenesis techniques The contributors focus on improvements to conventional site directed mutagenesis including a chapter on chemical site directed mutagenesis PCR based mutagenesis and the modifications that allow high throughput mutagenesis experiments and mutagenesis based on gene disruption both in vitro and in situ based Additional methods are provided for in vitro gene evolution for gene disruption based on recombination transposon and cassette mutagenesis and for facilitating the introduction of multiple mutations Time tested and highly practical the protocols in In Vitro Mutagenesis Protocols 2nd Edition offer today s molecular biologists reliable and powerful techniques with which to illuminate the proteome

Antibody Phage Display Philippa M. O'Brien,Robert Aitken,2008-02-05 The closing years of the 19th century and the start of the 20th century witnessed the emergence of microbiology and immunology as discrete sci tific disciplines and in the work of Roux and Yersin perhaps the first benefits of their synergy immunotherapy against bacterial infection As we advance into the new millennium microbiology and immunology again offer a c ceptual leap forward as antibody phage display gains increasing acceptance as the definitive technology for monoclonal production and unleashes new portunities in immunotherapy drug discovery and functional genomics In assembling Antibody Phage Display Methods and Protocols we have aimed to produce a resource of real value for scientists who have followed the development of phage display technology over the past decade The founding principles of phage display have always held an elegant simplicity We hope that readers will find similar clarity in the technical guidance offered by the book s contributors In meeting our objectives we have tried to cover the broad scope of the technology and the key areas of library construction scre ing antibody modification and expression Of course the technology cont ues to advance apace but we trust that readers will be able to gage the potential of phage display from our coverage that some of its subtleties will emerge and that our selection of methods will prove

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