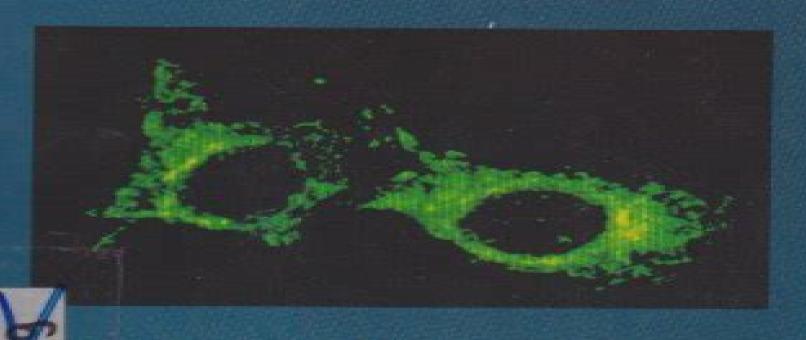
METHODS IN NONRADIOACTIVE DETECTION

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Gary C. Howard

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Danny L. Wiedbrauk, Daniel H. Farkas

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but vital details not always explicit in the methods sections of journal papers In recent years most biological laboratories have been invaded by computers and a wealth of new DNA technology and this will be reflected in many of the titles appearing in the series The books will be of value to advanced researchers and graduate students seeking to learn and apply new techniques and will be useful to teachers of advanced undergraduate courses involving practical or project work Labelled biomolecules are an essential tool in life science research and non radioactive labels are becoming increasingly important due to their convenience of measurement greater safety and lack of disposal problems compared to radioactive labels This book provides practical information background theory and protocols to allow a beginner to label many types of biomolecules including proteins peptides nucleic acids and small molecules This book is essential for biochemists molecular biologists and cell biologists wanting to use non radioactively labelled molecules Aimed at researchers without specific expertise in chemistry the book includes A review of the main signal systems and labels available indicating their strengths and weaknesses Discussion of the most useful strategies for labelling the various biomolecules 32 protocols covering common labelling needs Descriptions of the factors governing protocol design enabling protocols to be modified for different applications Sources of information including references data and suppliers **PCR Mutation Detection Protocols** Bimal D. M. Theophilus, Ralph Rapley, 2008-02-02 1Bimal D Theophilus and Ralph Rapley provide biological and clinical investigators with a comprehensive collection of new recent and updated PCR based screening methods suitable for detecting the presence of both known and novel mutations The methods cover point mutations e g ASO PCR SSCP DGGE chemical cleavage deletions multiplex PCR FISH blotting non sense mutations PTT and more The new and exciting techniques of DNA array analysis along with such recently developed experimental methods as conformation sensitive gel electrophoresis are also included Each chapter explains the basic theory behind the technique and provides valuable notes essential for its successful execution Molecular Methods for Virus Detection Danny L. Wiedbrauk, Daniel H. Farkas, 1995-02-08 Molecular diagnostic procedures have been described in a number of recent books and articles However these publications have not focused on virus detection nor have they provided practical protocols for the newer molecular methods Written by the inventors or principal developers of these technologies Molecular Methods for Virus Detection provides both reviews of individual methods and instructions for detecting virus nucleic acid sequences in clinical specimens Each procedure includes quality assurance protocols that are often ignored by other methodology books Molecular Methods for Virus Detection provides clinically relevant procedures for many of the newer diagnostic methodologies Provides state of the art PCR methods for amplification quantitation in situ hybridization and multiplex reactions Goes beyond PCR with protocols for 3SR NASBA LCR SDA and LAT Covers important virus detection methods such as in situ hybridization Southern dot and slot blots branched chain signal amplification and chemiluminescence Includes quality control information crucial in research and clinical laboratories Most chapters are written by the inventors and principal developers of the methodologies

Includes color plates 77 figures and 18 tables Nonisotopic DNA Probe Techniques Larry J. Kricka, 2012-12-02 Recently many nonisotopic methods of probing specific DNA sequences have been developed as replacements for radioactive labels such as 32phosphorous and 125iodine This book brings all of these new methods together in one convenient easily accessible source It enables researchers to select the nonisotopic method best suited to their application and to use it to maximum advantage by following the straightforward instructions provided This book contains chapters on colorimetric bioluminescent chemiluminescent fluorescent and time resolved fluorescent detection methods Each chapter has been written by the inventor or developer of a particular nonisotopic method and thus provides an expert account of the method Each chapterpresents useful background information and detailed step by step easy to follow experimental procedures for labeling and detection Gives extensive practical informationCovers major types of nonisotopic labels and proceduresPresents background information for each methodProvides strategies and detailed experimental procedures for labeling and detecting DNA sequences by Fluorescence Chemiluminescence Bioluminescence Colorimetry **Technologies for Detection of DNA Damage and Mutations** G.P. Pfeifer, 2013-11-11 Man made carcinogens natural genotoxic agents in the environment as well as ionizing and ultraviolet radiation can damage DNA and are a constant threat to genome integrity Throughout the evolution of life complex DNA repair systems have developed in all living organisms to cope with this damage Unrepaired DNA lesions can promote genetic alterations mutations that may be linked to an altered phenotype and if growth controlling genes are involved these mutations can lead to cell transformation and the development of malignant tumors Proto oncogenes and tumor suppressor genes may be critical targets for DNA damaging agents In a number of animal model systems correlations between exposure to a carcinogen tumor develop ment and genetic changes in tumor DNA have been established To understand mutagenesis processes in more detail at the molecular level we need to know the type and frequency of DNA adducts within cells their distribution along genes and specific DNA sequences as well as the rates at which they are repaired We also need to know what types of mutations are produced and which gene positions are most prone to mutagenesis This book provides a collection of techniques that are useful in mutagenesis research The book is divided into three parts In Part I methods for DNA damage and repair analysis are provided

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