LABORATORY EXERCISE #3. Detection of genetically modified food products by gel electrophoresis

Introduction

PCR products of differing lengths can be separated from each other based on size by an analytical method called gel electrophoresis. In this method charged macromolecules (such as DNA, RNA or proteins) migrate through a porous gel typically made from agarose (polysaccharide polymer derived from sea weed) or polyacrylamide when an electrical current is applied. The DNA sample is loaded into a well within the gel and placed near the anode (negative charge). As DNA has a very high negative charge due to the presence of the ionized phosphate groups on its backbone, it will migrate away from the negative anode toward the positively charged cathode as electrophoresis takes place. During the migration, smaller DNA fragments will pass through the gel pores more easily and thus migrate through the gel faster. Larger DNA molecules take longer to get through the pores and will migrate slower. The separated DNA "bands" or fragments can be visualized within the gel by staining with **ethidium bromide**. This dye intercalates (inserts) itself into the DNA and will fluoresces under ultraviolet light. In other words, DNA fragments will appear as fluorescent bands in the gel when exposed to UV light (Fig 1).

Concept check: If you compared two DNA fragments, one of 200 base pairs (bp) and the other of 1000 bp, which fragment will appear lower on the gel/have migrated faster?

Q: During gel electrophoresis, DNA is separated based on what two properties? A: Size and charge

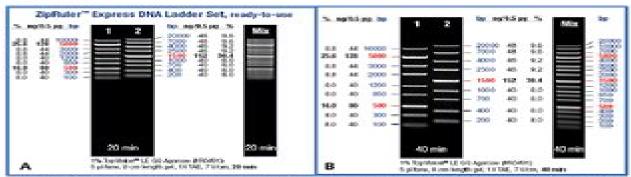


Figure 1. DNA fragments of known size are separated on an agarose gel for 20 min (A) or 40 min (B). By comparing the results, you can see how at 20 min, the DNA fragments are not well separated but by 40 min, the individually sized fragments are well separated and the size is easily determined. Figure adapted from Biogen.

Gel Electrophoresis Lab Manual Answers

Marilyn S. Arsham, Margaret J. Barch, Helen J. Lawce

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Laboratory Manual For Genetic Engineering VENNISON, S. JOHN, 2009-01-01 This systematically designed laboratory manual elucidates a number of techniques which help the students carry out various experiments in the field of genetic engineering The book explains the methods for the isolation of DNA and RNA as well as electrophoresis techniques for DNA RNA and proteins It discusses DNA manipulation by restriction digestion and construction of recombinant DNA by ligation Besides the book focuses on various methodologies for DNA transformation and molecular hybridization While discussing all these techniques the book puts emphasis on important techniques such as DNA isolation from Gram positive bacteria including Bacillus sp the slot lysis electrophoresis technique which is useful in DNA profile analysis of both Gram negative and positive bacteria plasmid transduction in Bacillus sp and the conjugal transfer of plasmid DNA in cyanobacteria Bacillus and Agrobacterium tumefaciens This book is intended for the undergraduate and postgraduate students of biotechnology for their laboratory courses in genetic engineering Besides it will be useful for the students specializing in genetic engineering molecular biology and molecular microbiology KEY FEATURES Includes about 60 different experiments Contains several figures to reinforce the understanding of the techniques discussed Gives useful information about preparation of stock solutions DNA protein conversions restriction enzymes and their recognition sequences and so on in Laboratory Manual for Biotechnology and Laboratory Science Lisa A. Seidman, Mary Ellen Kraus, Diana **Appendices** Lietzke Brandner, Jeanette Mowery, 2022-12-23 Provides the basic laboratory skills and knowledge to pursue a career in biotechnology Written by four biotechnology instructors with over 20 years of teaching experience it incorporates instruction exercises and laboratory activities that the authors have been using and perfecting for years These exercises and activities help students understand the fundamentals of working in a biotechnology laboratory Building skills through an organized and systematic presentation of materials procedures and tasks the manual explores overarching themes that relate to all biotechnology workplaces including forensic clinical quality control environmental and other testing laboratories Features Provides clear instructions and step by step exercises to make learning the material easier for students There are Lab Notes for Instructors in the Support Material see tab below Emphasizes fundamental laboratory skills that prepare students for the industry Builds students skills through an organized and systematic presentation of materials procedures and tasks Updates reflect recent innovations and regulatory requirements to ensure students stay up to date Supplies skills suitable for careers in forensic clinical quality control environmental and other testing laboratories Lab Manual Biology Class 12 Rajesh Kumar. Lab Manual BIOCHEMISTRY LABORATORY MANUAL PALLAB BASU,2016-01-01 The AGT Cytogenetics Laboratory Manual Marilyn S. Arsham, Margaret J. Barch, Helen J. Lawce, 2017-04-24 Cytogenetics is the study of chromosome morphology structure pathology function and behavior The field has evolved to embrace molecular cytogenetic changes now termed cytogenomics Cytogeneticists utilize an assortment of procedures to investigate the full complement of

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