

Epitope Mapping A Practical Approach Practical Approach Series

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In epitope-mapping, a phage-display library is initially scanned with the antibody; following that, the affinity selected peptides need to be mapped onto the antigen structure in order to infer ...

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Epitope Mapping covers all the major methods for the identification and definition of epitopes. The Pepsan assay is used to define B cell epitopes and makes use of synthetic peptides but can only be used if the amino acid sequence is known. It can be adapted for the delineation of both helper T cells and cytotoxic T cells.

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Introduction to epitope mapping Synthetic peptides 1 pepsan assay to define antigenic determinants for antibodies (B cell Epitopes) Delineation of helper T cell epitopes and MHC Combined T and B cell epitopes Cytotoxic T cell eptitopes Peptide mimic libraries Steric competition mapping Oligosaccharide libraries Phage display libraries Site-directed mutagenesis

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There are several methods available for mapping antibody epitopes on target antigens: X-ray co-crystallography and cryogenic electron microscopy (cryo-EM). X-ray co-crystallography has historically been... Array -based oligo-peptide scanning. Also known as overlapping peptide scan or pepsan ...

Epitope mapping - Wikipedia
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An epitope is a structural region of an antigen that is recognized by an antibody and is therefore central to the immune response. Epitope Mapping describes the various methods for their location and characterization.

An epitope is a structural region of an antigen that is recognized by an antibody and is therefore central to the immune response. Epitope Mapping describes the various methods for their location and characterization. This process is an essential part of developing non-pathogenic vaccines.

Peptide antigens and anti-peptide antibodies are widely used in biochemistry and molecular biology for the measurement, location, and purification of specific oligopeptides. More recently the application of these reagents has expanded, for example in the identification and mapping of the binding sites of antibodies and T-cell receptors and in the identification and characterization of proteins which are known only by their primary structure. This volume provides practical guidance to the major techniques used in the exploitation of peptide antigens and anti-peptide antibodies. The chapters give detailed protocols for the prediction of epitopes, peptides synthesis, the preparation of peptide immunogens, immunoaffinity chromatography, immunoassays, and the mapping of epitopes using both synthetic peptides and phage display systems. Peptide Antigens: A Practical Approach covers all practical aspects of this important and growing subject. It is a unique compendium of methods for workers in biochemistry, molecular biology, and immunology who need to use this technology in their research.

This new book is designed to enable researchers to design and undertake all aspects of a phage display project, from designing an experimental strategy and constructing a library to performing selections and analyzing the results.All of the protocols and chapters are extensively cross-referenced, allowing readers to move beyond the specific examples provided in order to customize the procedures for their own protein or selection system of interest. Phage Display is an up-to-date, comprehensive and integrated experimental guide to the technique, which is essential reading for anyone currently using, or wishing to use the technique for basic research and drug discovery.

The latest edition of this highly successful text, covers the major advances in the methods used in cellular and molecular pathology. In recent years, knowledge of the molecular organization of the cell has led to the development of powerful new techniques that bring greater accuracy and objectives to the diagnosis, prognosis and management of many diseases and to the study of pathological states. This book describes the latest molecular techniques available for the analysis of diseases. In particular it includes new techniques using fluorescent dyes, DNA microarrays, protein chemistry, and mass spectrometry. It also incorporates information from the Human Genome Project, and the new disciplines of genomics and proteomics, where relevant to pathology. Color plates are a new feature of this edition, illustrating the advances in fluorescence labeling of cells.

Ebola epidemics have had immediate and lasting impact in Africa and beyond, with its high case fatality and societal disruption. Its rapid spread, coupled with the limited knowledge, serves as a recipe for disaster and panic in the community. Health workers are particularly at risk, paying heavily with their lives. Sharing knowledge from various experts in basic sciences that support vaccine and drug development, as well as improving community surveillance and case management, enriches our understanding of this highly fatal and contagious disease. In a world that is fast becoming a global village, communicable diseases from low-resource setting are gradually becoming a global health threat. This book seeks to discuss emerging advances in the Ebola control.

Enzymology at the Membrane Interface: Intramembrane Proteases, Volume 584, the latest release in the Methods in Enzymology series, covers a subset of enzymes that work in the environment of the biological cell membrane. This field, called interfacial enzymology, involves a special series of experimental approaches for the isolation and study of these enzymes. Covers a subset of enzymes that work in the environment of the biological cell membrane Offers a series of experimental approaches for the isolation and study of enzymes

This third edition volume expands on the previous editions with more detailed research on the characterization of antibody antigen interactions between different users with different requirements. The chapters in this book are divided into four parts: Part One looks at the entire native antigen and covers traditional structural biology techniques such as nuclear magnetic resonance and x-ray crystallography. Part Two talks about protein fragments derived from antigens, and discusses binding regions within antigen sequence using bacterial surface display and ELISA, for example. Part Three describes the use of surface plasmon resonance spectroscopy and biolayer interferometry, and Part Four highlights methods used to identify new antigens and assess antibody cross-reactivity. Written in the highly successful Methods in Molecular Biology series format, chapters include introductions to their respective topics, lists of the necessary materials and reagents, step-by-step, readily reproducible laboratory protocols, and tips on troubleshooting and avoiding known pitfalls. Thorough and cutting-edge, Epitope Mapping Protocols, Third Edition is a valuable resource for anyone interested in furthering their research in this expanding field.

Practical Immunology is a basic text aimed at immunology students and researchers at all levels who need a comprehensive overview of the methodology of immunology. The rapid and startling innovations in immunology over the past two decades have their root in sound experimental practice and it has always been the aim of this book to educate researchers in the design and performance of complex techniques. It will appeal to students of immunology, graduate students embarking on bench science, or specialised immunologists who need to use an immunological technique outside their sphere of expertise. The definitive lab "bench book". A one stop resource. Techniques explained from first principles. Basic forms of apparatus described in detail. Totally revised with new user friendly layout to aid use in the lab. Includes useful hints and tips.

Phage display has become established as a powerful protein engineering method for identifying polypeptides with novel properties, and altering the properties of existing ones. Although the technique is widely used in biological research and drug discovery, it remains technically challenging, and new applications and procedures continue to evolve. Phage Display - A Practical Approach is an up-to-date, comprehensive and integrated experimental guide to the technique, useful for novice and expert alike. The book aims to enable researchers to design and undertake all aspects of a phage display project, from designing an experimental strategy and constructing a library to performing selections and analyzing the results. An introductory chapter provides an overview of phage biology and phage display, including guidelines for planning a successful phage display experiment. Individual chapters provide protocols for constructing libraries using oligonucleotide-directed mutagenesis or DNA recombination, performing binding selections, and analyzing the binding activities of selected phage clones. Separate chapters then cover common applications, including selection of ligands from peptide libraries, generation of phage antibody libraries and isolation and optimization of antibodies, selection of DNA binding proteins, and expression cloning using cDNA display. Further chapters describe alternative selection strategies, such as selection using immune sera, selection based on enzymatic activity or protein stability, and selection in vivo. Protocols and chapters are extensively cross-referenced, allowing readers to move beyond the specific examples given to customize the procedures to their own protein or selection system of interest. Written by experts in the field, Phage Display - A Practical Approach provides a comprehensive guide to the design and execution of phage display projects, for all those using the technique in basic research and drug discovery.