

Advances in Clinical and Experimental Medicine

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Advances Clinical Science Experimental Medicine

Jie Xu



Advances Clinical Science Experimental Medicine:

Advances in Clinical Chemistry, 2011-03-10 Volume 53 in the internationally acclaimed *Advances in Clinical Chemistry* contains chapters submitted from leading experts from academia and clinical laboratory science. Authors are from a diverse field of clinical chemistry disciplines and diagnostics ranging from basic biochemical exploration to cutting edge microarray technology. Leading experts from academia and clinical laboratory science. Volume emphasizes novel laboratory advances with application to clinical laboratory diagnostics and practical basic science studies. Clinical and Translational Science David Robertson, Gordon H. Williams, 2016-11-25 *Clinical and Translational Science Principles of Human Research* Second Edition is the most authoritative and timely resource for the broad range of investigators taking on the challenge of clinical and translational science, a field that is devoted to investigating human health and disease interventions and outcomes for the purposes of developing new treatment approaches, devices and modalities to improve health. This updated second edition has been prepared with an international perspective beginning with fundamental principles, experimental design, epidemiology, traditional and new biostatistical approaches and investigative tools. It presents complete instruction and guidance from fundamental principles, approaches and infrastructure especially for human genetics and genomics, human pharmacology, research in special populations, the societal context of human research and the future of human research. The book moves on to discuss legal, social and ethical issues and concludes with a discussion of future prospects, providing readers with a comprehensive view of this rapidly developing area of science. Introduces novel physiological and therapeutic strategies for engaging the fastest growing scientific field in both the private sector and academic medicine. Brings insights from international leaders into the discipline of clinical and translational science. Addresses drug discovery, drug repurposing and development, innovative and improved approaches to go/no-go decisions in drug development and traditional and innovative clinical trial designs. **Antivivisection and Medical Science in Victorian Society** Richard D. French, 2019-03-12 Late nineteenth century England witnessed the emergence of a vociferous and well organized movement against the use of living animals in scientific research, a protest that threatened the existence of experimental medicine. Richard D. French views the Victorian antivivisection movement as a revealing case study in the attitude of modern society toward science. The author draws on popular pamphlets and newspaper accounts to recreate the structure, tactics, ideology and personalities of the early antivivisection movement. He argues that at the heart of the antivivisection movement was public concern over the emergence of science and medicine as leading institutions of Victorian society, a concern he suggests that has its own contemporary counterparts. In addition to providing a social and cultural history of the Victorian antivivisection movement, the book sheds light on many related areas including Victorian political and administrative history, the political sociology of scientific communities, social reform and voluntary associations, the psychoanalysis of human attitudes toward animals and Victorian feminism. Richard D. French is a Science Advisor with the Science Council of Canada. Originally published in 1975.

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Potency Assays for Advanced Stem Cell Therapy Medicinal Products

Jorge S. Burns, 2023-05-31. This volume of the Springer book series *Advances in Experimental Medicine and Biology* covers potency assays, one of the most complex yet fundamental evaluations that critically influence stem cell regenerative medicine. Developing potency assays for cell-based medicinal products comes with numerous challenges due to the highly specialised nature of the application and purpose. This book provides the reader with the knowledge necessary to understand issues governing the successful development of potency assays, highlighting an international outlook of how the various challenges raised are being managed. Stakeholders concerned with potency assay development range from patient and clinician to contract research organisations, small and medium enterprise, regulatory authorities, and even politicians. The value of potency assays is poised to increase given the inevitable watershed as early stage clinical trials addressing safety progress to trials testing efficacy. Contributors from clinical, academic, industrial, and regulatory sectors establish a broad point of view for guidance and timely debate. Potency assays require extensive collaboration across disciplines and sectors, as well as compromise, and the authors aim to constructively address the many key aspects involved. Potency assays provide a quantitative measure of the biological activity of advanced therapy medicinal products (ATMPs) and thus are required for their market authorization. As the pace of ATMP development accelerates, the need to develop specific, accurate, and robust potency assays for each product is also accelerating. The volume *Potency Assays for Stem Cell Advanced Therapy Medicinal Products* presents a broad outlook on the development, quality attributes, and implementation of potency assays for ATMPs. The first few chapters introduce a nuanced historical perspective on the science of potency assay development, describe specific quality attributes of an idealized potency assay, indicate pitfalls associated with developing such assays for ATMPs, and review guidance recommended by regulatory authorities on assay suitability for product approval. Subsequent chapters highlight efforts to develop potency assays for specific ATMPs, including skeletal stem cells, mesenchymal stromal cells, extracellular vesicles, CAR T cells, and discuss emerging technologies/platforms for potency assay design. The volume concludes with a chapter reviewing potency assays used for the release of commercial ATMP products, which amalgamates information contained in previous chapters. Overall, the knowledge contributed from leading authorities in both academia and industry is an ideal resource for technicians, scientists, clinicians, process engineers, and regulators working with ATMPs. Donald G. Phinney, PhD, Professor, Department of Molecular Medicine, Herbert Wertheim UF Scripps Institute for Biomedical Innovation Technology.

List of Journals Indexed in Index Medicus National Library of Medicine (U.S.), 1974. Issues for 1977, 1979.

include also Special List journals being indexed in cooperation with other institutions Citations from these journals appear in other MEDLARS bibliographies and in MEDLINE but not in Index Medicus

The Role of Glycosylation in Health and Disease Gordan Lauc, Irena Trbojević-Akmačić, 2021-09-08 Glycobiology is an emerging field of studying glycans sugars and glycoconjugates that are essentially involved in almost all biological processes from fine tuning glycoprotein function to protein protein interactions signaling immune response host pathogen interactions etc However due to structural complexity of glycans and analytical challenges this exciting field was lagging behind other areas of biology With technological advancements growing number of glycans functions are being discovered and the study of glycans is becoming a cutting edge discipline in basic and clinical research Despite recent developments in glycobiology field many aspects of glycosylation process still remain unknown both in a healthy human organism and in pathological states Human glycome is dynamic and changes with physiological triggers immune challenges and disease Atypical glycosylation is consequently a subject of disease biomarker research and a target for therapeutic interventions On the other hand properties of glycosylated biotherapeutics are predominantly determined by their glycans The Role of Glycosylation in Health and Disease provides a comprehensive overview of types and functions of glycans in a healthy human organism as well as their role in pathophysiology of different diseases and efficiency of glycosylated biotherapeutics Written by the experts in the field this book aims to bring glycobiology field closer to students researchers in life sciences and professionals in biopharmaceutical industry

Notch Signaling in Embryology and Cancer Jörg Reichrath, Sandra Reichrath, 2020-10-08 This thoroughly revised second edition is an up to date overview of the current knowledge of Notch and Notch signaling in embryology and cancer It discusses this topic from Notch's role in the development of the embryo to the Notch signaling pathway's role in the development of a number of cancers including breast cancer malignant melanoma Non melanoma skin cancer intestinal cancer and others In the years since the previous edition there have been numerous developments and insights within this rapidly moving field making this new edition urgently needed This volume also features discussions of current insights on Notch's role in senescence the regulation of Notch signaling by microRNAs Notch's role in the microbiome diet and its influence on Notch signaling and more Taken as a whole with its companion books Notch Signaling and Embryologic Development and Molecular Basis of Notch Signaling this is a definitive discussion of the topic presented by internationally recognized contributors Presented in a coherent and accessible structure this revised and updated second edition is an essential and up to date guide for oncologists embryologists researchers and advanced students

Tumor Microenvironment Alexander Birbaumer, 2020-10-29 This book provides a comprehensive overview of the latest research on the molecular players in the tumor microenvironment including MicroRNAs estrogen Caveolin 1 Nitric Oxide RANK RANKL signaling COX 2 Signaling Renin angiotensin system and more Taken alongside its companion volumes Tumor Microenvironment Molecular Players Part B updates us on what we know about the tumor microenvironment as well as

future directions This book is essential reading for advanced cell biology and cancer biology students as well as researchers seeking an update on research in the tumor microenvironment Virus Entry Inhibitors Shibo Jiang,Lu Lu,2022-04-12 This book focuses on the important discoveries of the small molecule peptide and protein based virus entry inhibitors and discusses advance of the development of different type of virus entry inhibitors as a novel class of antiviral drugs for treatment and prevention of viral infection It includes entry inhibitors of HIV HeV NiV RSV EBOV HCoV HBV HCV HDV HPV flavivirus and influenza virus so on This book aims at providing an updated knowledge on virus entry inhibitors

Regulation of Cancer Immune Checkpoints Jie Xu,2020-03-17 This book systematically reviews the most important findings on cancer immune checkpoints sharing essential insights into this rapidly evolving yet largely unexplored research topic The past decade has seen major advances in cancer immune checkpoint therapy which has demonstrated impressive clinical benefits The family of checkpoints for mediating cancer immune evasion now includes CTLA 4 PD 1 PD L1 CD27 CD70 FGL 1 LAG 3 Siglec 15 VISTA PD 1L VSIG3 CD47 SIRPA APOE LILRB4 TIGIT and many others Despite these strides most patients do not show lasting remission and some cancers have been completely resistant to the therapy The potentially lethal adverse effects of checkpoint blockade represent another major challenge the mechanisms of which remain poorly understood Compared to the cancer signaling pathways such as p53 and Ras mechanistic studies on immune checkpoint pathways are still in their infancy To improve the responses to checkpoint blockade therapy and limit the adverse effects it is essential to understand the molecular regulation of checkpoint molecules in both malignant and healthy cells tissues This book begins with an introduction to immune checkpoint therapy and its challenges and subsequently describes the regulation of checkpoints at different levels In closing it discusses recent therapeutic developments based on mechanistic findings and outlines goals for future translational studies The book offers a valuable resource for researchers in the cancer immunotherapy field helping to form a roadmap for checkpoint regulation and develop safer and more effective immunotherapies

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