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PORTER RUSH

Formulation tools for Pharmaceutical Development Lippincott Williams & Wilkins
Developing Solid Oral Dosage Forms: Pharmaceutical Theory and Practice, Second Edition illustrates how to develop high-quality, safe, and effective pharmaceutical products by discussing the latest techniques, tools, and scientific advances in preformulation investigation, formulation, process design, characterization, scale-up, and production operations. This book covers the essential principles of physical pharmacy, biopharmaceutics, and industrial pharmacy, and their application to the research and development process of oral dosage forms. Chapters have been added, combined, deleted, and completely revised as necessary to produce a comprehensive, well-organized, valuable reference for industry professionals and academics engaged in all aspects of the development process. New and important topics include spray drying, amorphous solid dispersion using hot-melt extrusion, modeling and simulation, bioequivalence of complex modified-released dosage forms, biowaivers, and much more. Written and edited by an international team of leading experts with experience and knowledge across industry, academia, and regulatory settings Includes new chapters covering the pharmaceutical applications of surface phenomenon, predictive biopharmaceutics and pharmacokinetics, the development of formulations for drug discovery support, and much more Presents new case studies throughout, and a section completely devoted to regulatory aspects, including global product regulation and international perspectives
The Art and Science of Dermal Formulation Development Disha Publications
A needed resource for pharmaceutical scientists and cosmetic chemists, Essential Chemistry for Formulators of Semisolid and Liquid Dosages provides insight into the basic chemistry of mixing different phases and test methods for the stability study of nonsolid formulations. The book covers foundational surface/colloid chemistry, which forms the necessary background for making emulsions, suspensions, solutions, and nano drug delivery systems, and the chemistry of mixing, which is critical for further formulation of drug delivery systems into semisolid (gels, creams, lotions, and ointments) or liquid final dosages. Expanding on these foundational principles, this useful guide explores stability testing methods, such as particle size, rheological/viscosity, microscopy, and chemical, and closes with a valuable discussion of regulatory issues. Essential Chemistry for Formulators of Semisolid and Liquid Dosages offers scientists and students the foundation and practical guidance to make and analyze semisolid and liquid formulations. Unique coverage of the underlying chemistry that makes possible stable dosages Quality content written by experienced experts from the drug development industry Valuable information for academic and industrial scientists developing topical and liquid dosage formulations for pharmaceutical as well as skin care and cosmetic products
UGC NET Political Science (Paper I & II) 2021 | 10 Full-length Mock Test (New Exam Pattern) CRC Press
How to Develop Robust Solid Oral Dosage Forms from Conception to Post-Approval uses a practical and hands-on approach to cover the development process of solid oral dosage forms in one single source. The book details all of the necessary steps from formulation through the post-approval phase and contains industry case studies, real world advice, and troubleshooting tips. By merging the latest scientific information with practical instructions, this book provides pharmaceutical scientists in formulation research and development with a concrete look at the key aspects in the development of solid oral dosage forms. Focuses on important topics, such as robustness, bioavailability, formulation design, continuous processing, stability tests, modified release dosage forms, international guidelines, process scale-up, and much more Part of the Expertise in Pharmaceutical Process Technology series edited by Michael Levin Discusses common, real-world problems and offers both theoretical and practical solutions to these everyday issues
Formulation Development of Niacin Extended Release Tablets CRC Press
The development of paediatric medicines can be challenging since this is a different patient population with specific needs. A medicine designed for use in paediatric patients must consider the following aspects: patient population variability; the need for dose flexibility; route of administration; patient compliance; excipient tolerability. For example, the toxicity of excipients may differ in children compared to adults and children have different taste preferences. Globally, about 75% of drugs do not carry regulatory approval for use in children; worldwide, many medications prescribed for the treatment of paediatric diseases are used off-label, and less than 20% of package inserts have sufficient information for treating children. This book provides an update on both state-of-the-art methodology and operational challenges in paediatric formulation design and development. It aims at re-evaluating what is needed for more progress in the design and development of age-appropriate treatments for paediatric diseases, focusing on: formulation development; drug delivery design; efficacy, safety, and tolerability of drugs and excipients.
The Science of Dosage Form Design Woodhead Publishing
The Future of Pharmaceutical Product Development and Research examines the latest developments in the pharmaceutical sciences, also highlighting key developments, research and future opportunities. Written by experts in the field, this volume in the Advances in Pharmaceutical Product Development and Research series deepens our understanding of the product development phase of drug discovery and drug development. Each chapter covers fundamental principles, advanced methodologies and technologies employed by pharmaceutical scientists, researchers and the pharmaceutical industry. The book focuses on excipients, radiopharmaceuticals, and how manufacturing should be conducted in an environment that follows Good Manufacturing Practice (GMP) guidelines. Researchers and students will find this book to be a comprehensive resource for those working in, and studying, pharmaceuticals, cosmetics, biotechnology, foods and related industries. Provides an overview of practical information for clinical trials Outlines how to ensure an environment that follows Good Manufacturing Practice (GMP) Examines recent developments and suggests future directions for drug production methods and techniques
From Conception to Post-Approval MDPI
A real-world guide to the production and manufacturing of biopharmaceuticals While much has been written about the science of biopharmaceuticals, there is a need for practical, up-to-date information on key issues at all stages of developing and manufacturing commercially viable biopharmaceutical drug products. This book helps fill the gap in the field, examining all areas of biopharmaceuticals manufacturing, from development and formulation to production and packaging. Written by a group of experts from industry and academia, the book focuses on real-world methods for maintaining product integrity throughout the commercialization process, clearly explaining the fundamentals and essential pathways for all development stages. Coverage includes: Research and early development phase–appropriate approaches for ensuring product stability Development of commercially viable formulations for liquid and lyophilized dosage forms Optimal storage, packaging, and shipping methods Case studies relating to therapeutic monoclonal antibodies, recombinant proteins, and plasma fractions Useful analysis of successful and failed products Formulation and Process Development Strategies for Manufacturing Biopharma-ceuticals is an essential resource for scientists and engineers in the pharmaceutical and biotech industries, for government and regulatory agencies, and for anyone with an interest in the latest developments in the field.
USDA Forest Service Research Paper INT. John Wiley & Sons
The aim of present study is to design and develop a solid oral extended release dosage form (tablet) of Niacin to deliver controlled release of drug at desired site at specific time comparable to the innovator product with better stability, high production feasibility, and excellent patient compatibility. The objective of the study is to evaluate the release pattern of the drug from fabricated extended release tablets and compare with marketed sample of the same drug Niaspan 1000mg ER tablet over a period of 24 hours. To carry out the stability for the optimized formulations.
A Study of the Role of the Federal Government in the First Five Years of the Social Security Act, 1935-1940 John Wiley & Sons
Engineering Drug Delivery Systems is an essential resource on a variety of biomaterials engineering approaches for creating drug delivery systems that have market and therapeutic potential. The book comprehensively discusses recent advances in the fields of biomaterials and biomedical sciences in relation to drug delivery. Chapters provide a detailed introduction to various engineering approaches in designing drug delivery systems, delve into the engineering of body functions, cover the selection, design and evaluation of biomaterials, and discuss the engineering of colloids as drug carriers. The book's final chapters address the engineering of implantable drug delivery systems and advances in drug delivery technology. This book is an invaluable resource for drug delivery, materials scientists and bioengineers within the pharmaceutical industry. Examines the properties and synthesis of biomaterials for successful drug delivery Discusses the important connection between drug delivery and tissue engineering Includes techniques and approaches applicable to a wide range of users Reviews innovative technologies in drug delivery systems such as 3-D printed devices for drug delivery
Design and Development Academic Press
A teacher is a person who not only teaches but also guides his/her student in building a successful career. The future of a nation lies upon the level of knowledge the people in the country are having. Thus, the responsibility of a teacher goes far beyond what we think of it at an individual level. We have seen people are interested in making their career in many other professions but teaching as a profession is not the first choice in most cases. Nevertheless, teaching is one of the most interesting professions as it involves a continuous learning exercise and at the same time making others learned by delivering the knowledge one is having. The teachers assess their students but at first, they also get assessed under UGC NET conducted by the National Testing Agency. The National Eligibility Test (NET), also known as UGC NET or NTA-UGC-NET, is the test for determining the eligibility for the post of Assistant Professor and/or Junior Research Fellowship (JRF) award in Indian universities and colleges. UGC NET is considered as one of the toughest exams in India, with success ratio of merely 6%. Previously, the passing ratio was around 3% - 4%. Assistant Professors in private colleges may or may not be NET qualified but NET qualification is mandatory for universities & government colleges.
UGC NET English 2021 | 10 Full-length Mock Test (Paper I & II) | With Latest Exam Pattern Elsevier
Drug Delivery Aspects reviews additional features of drug delivery systems, along with the standard formulation development, like preclinical testing, conversion into solid dosage forms, roles of excipients and polymers used on stability and sterile processing. There is a focus on formulation engineering and related large scale (GMP) manufacturing, regulatory, and functional aspects of drug delivery systems. A detailed discussion on biologics and vaccines gives insights to readers on new developments in this direction. The series Expectations and Realities of Multifunctional Drug Delivery Systems examines the fabrication, optimization, biological aspects, regulatory and clinical success of wide range of drug delivery carriers. This series reviews multifunctionality and applications of drug delivery systems, industrial trends, regulatory challenges and in vivo success stories. Throughout the volumes discussions on diverse aspects of drug delivery carriers, such as clinical, engineering, and regulatory, facilitate insight sharing across expertise area and form a link for collaborations between industry-academic scientists and clinical researchers. Expectations and Realities of Multifunctional Drug Delivery Systems connects formulation scientists, regulatory experts, engineers, clinical experts and regulatory stake holders. The wide scope of the book ensures it as a valuable reference resource for researchers in both academia and the pharmaceutical industry who want to learn more about drug delivery systems. Encompasses engineering and large-scale manufacturing of nanocarriers Considers preclinical, regulatory and ethical guidelines on nanoparticles Contains in-depth discussions on delivery of biologics, vaccines and sterilisation Industrial view on solid dispersions, milling techniques
Oral Controlled Release Formulation Design and Drug Delivery Woodhead Publishing
A teacher is a person who not only teaches but also guides his/her student in building a successful career. The future of a nation lies upon the level of knowledge the people in the country are having. Thus, the responsibility of a teacher goes far beyond what we think of it at an individual level. We have seen people are interested in making their career in many other professions but teaching as a profession is not the first choice in most cases. Nevertheless, teaching is one of the most interesting professions as it involves a continuous learning exercise and at the same time making others learned by delivering the knowledge one is having. The teachers assess their students but at first, they also get assessed under UGC NET conducted by the National Testing Agency. The National Eligibility Test (NET), also known as UGC NET or NTA-UGC-NET, is the test for determining the eligibility for the post of Assistant Professor and/or Junior Research Fellowship (JRF) award in Indian universities and colleges. UGC NET is considered as one of the toughest exams in India, with success ratio of merely 6%. Previously, the passing ratio was around 3% - 4%. Assistant Professors in private colleges may or may not be NET qualified but NET qualification is mandatory for universities &

government colleges.
Paediatric Formulation BoD – Books on Demand
Paediatric FormulationDesign and DevelopmentMDPI
How to Think in Medicine CRC Press
Today, the pressure on healthcare costs and resources is increasing, and especially for biopharmaceuticals that require parenteral administration, the inherent complex and invasive dosing procedure adds to the demand for efficient medical management. In light of the COVID-19 pandemic the value of drug delivery technologies in enabling a flexible care setting is broadly recognized. In such a setting, patients and their caregivers can choose the place of drug administration based on individual preferences and capabilities. This includes not only dosing in the clinic but also supervised at-home dosing and self-administration for eligible patients. Formulation and Device Lifecycle Management of Biotherapeutics: A Guidance for Researchers and Drug Developers covers the various aspects of improving drug delivery of biological medicines with the ultimate goal to reduce dosing complexity associated with parenteral administration and, thus, enhance patient experience and drug administration-related healthcare capacity. The target audience are multidisciplinary researchers and drug developers in the pharmaceutical industry, biotech companies, and academia involved in formulation and device development. This includes pharmacology and medical experts in charge of generating nonclinical and clinical data to support approval of novel dosing regimens, and drug delivery scientists and engineers responsible for technical particulars of product optimizations. Moreover, professionals in market access and commercial functions are expected to benefit from the discussions about the impact of patient and healthcare provider needs and country-specific reimbursement models on realizing a truly convenient and cost and resource efficient drug delivery solution. Summarizes formulation and device lifecycle management activities that enable customer-centric and sustainable drug delivery for biotherapeutics Describes the pharmacokinetic-based clinical development pathway for subcutaneous dosing alternatives to established intravenous formulations for monoclonal antibodies Details established clinical development pathways supporting the approval of automated subcutaneous injection devices and proposes novel concepts Discusses how to realize home- and self-administration of biotherapeutics in cancer care Highlights aspects of multidisciplinary formulation and device lifecycle management that can be leveraged across different disease areas and introduces a decision architecture on when and how drug developers should embark into related development activities
Practical Guide to Methods and Applications CRC Press
Pharmaceutical Preformulation and Formulation: A Practical Guide from Candidate Drug Selection to Commercial Dosage Form reflects the mounting pressure on pharmaceutical companies to accelerate the new drug development and launch process, as well as the shift from developing small molecules to the growth of biopharmaceuticals. The book meets the need for advanced information for drug preformulation and formulation and addresses the current trends in the continually evolving pharmaceutical industry. Topics include: Candidate drug selection Drug discovery and development Preformulation predictions and drug selections Product design to commercial dosage form Biopharmaceutical support in formulation Development The book is ideal for practitioners working in the pharmaceutical arena—including R&D scientists, technicians, and managers—as well as for undergraduate and postgraduate courses in industrial pharmacy and pharmaceutical technology.
UGC NET History 2021 | 10 Full-length Mock Test (Paper I & II) | With Latest Exam Pattern Academic Press
This comprehensive book encompasses various facets of sterile product development. Key concepts relevant to the successful development of sterile products are illustrated through case studies and are covered under three sections in this book: • Formulation approaches that discuss a variety of dosage forms including protein therapeutics, lipid-based controlled delivery systems, PEGylated biotherapeutics, nasal dosage form, and vaccines • Process, container closure and delivery considerations including freeze-thaw process challenges, best practices for technology transfer to enable commercial product development, innovations and advancement in aseptic fill-finish operations, approaches to manufacturing lyophilized parenteral products, pen / auto-injector delivery devices, and associated container closure integrity testing hurdles for sterile product closures • Regulatory and quality aspects in the areas of particulate matter and appearance evaluation, sterile filtration, admixture compatibility considerations, sterilization process considerations, microbial contamination investigations and validation of rapid microbiological methods, and dry and moist heat sterilizers This book is a useful resource to scientists and researchers in both industry and academia, and it gives process and product development engineers insight into current industry practices and evolving regulatory expectations for sterile product development.
Pharmaceutical Dosage Forms - Tablets LAP Lambert Academic Publishing
This book describes the theories, applications, and challenges for different oral controlled release formulations. This book differs from most in its focus on oral controlled release formulation design and process development. It also covers the related areas like preformulation, biopharmaceutics, in vitro-in vivo correlations (IVIVC), quality by design (QbD), and regulatory issues.
Formulation Development and Release Kinetics Study John Wiley & Sons
A comprehensive textbook covering the design of dosage forms and all aspects of drug delivery systems. 'Pharmaceutics' in its broadest sense is the 'art of the apothecary' or, in simple terms, pharmaceutical preparations. It remains a diverse subject in the pharmacy curriculum, encompassing design of drugs, their manufacture, and the elimination of micro-organisms from the products. This books encompasses all those areas and pays particular attention to the design of dosage forms and their manufacture.
UGC NET Sociology 2021 | 10 Full-length Mock Test (New Exam Pattern) LAP Lambert Academic Publishing
The Art and Science of Dermal Formulation Development is a comprehensive guide to the theory and practice of transdermal and topical formulation development, covering preclinical studies, evaluation, and regulatory approval. It enables the reader to understand the opportunities and challenges in developing products and how risks can be mitigated. Over the last 25 years, expertise in this area has declined whilst drug delivery systems for other administration routes have developed significantly. The advantages offered by transdermal and topical drug delivery remain compelling for sectors including the pharmaceutical industry, personal care, and cosmetics. This text addresses the dearth of expertise and discusses how skin can be a route of delivery and the processes in formulation development, but how such an application is very different to that used for oral, IV, and other administration routes. Key Features: Presents a practical guide for both industry and academia Focuses on and draws together the fundamental principles behind transdermal and topical drug delivery Illustrates the practicalities of formulation design using key case studies Gives an understanding of the skin as a route of delivery and how formulation development for such application differs from that for other administration routes
Design and Development at Early Stage Academic Press
There are unique challenges in the formulation, manufacture, analytical chemistry, and regulatory requirements of low-dose drugs. This book provides an overview of this specialized field and combines formulation, analytical, and regulatory aspects of low-dose development into a single reference book. It describes analytical methodologies like dissolution testing, solid state NMR, Raman microscopy, and LC-MS and presents manufacturing techniques such as granulation, compaction, and compression. Complete with case studies and a discussion of regulatory requirements, this is a core reference for pharmaceutical scientists, regulators, and graduate students.
A Study of the Formulation and Development of a Pre-kindergarten Program in a Public School EduGorilla Community Pvt. Ltd.
The ultimate goal of drug product development is to design a system that maximizes the therapeutic potential of the drug substance and facilitates its access to patients. Pharmaceutical Dosage Forms: Tablets, Third Edition is a comprehensive resource of the design, formulation, manufacture, and evaluation of the tablet dosage form, an