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Stig Pedersen-Bjergaard, Bente Gammelgaard, Trine G. Halvorsen

Voigt's Pharmaceutical Technology Alfred Fahr, 2018-02-21 A textbook which is both comprehensive and comprehensible and that offers easy but scientifically sound reading to both students and professionals Now in its 12th edition in its native German, Voigt's Pharmaceutical Technology is an interdisciplinary textbook covering the fundamental principles of pharmaceutical technology. Available for the first time in English, this edition is produced in full colour throughout, with a concise, clear structure developed after consultation with students, instructors and researchers. This book: Features clear chapter layouts and easily digestible content Presents novel trends, devices and processes Discusses classical and modern manufacturing processes Covers all formulation principles including tablets, ointments, capsules, nanosystems and biopharmaceutics Takes account of legal requirements for both qualitative and quantitative composition Addresses quality assurance considerations Uniquely relates contrasting international pharmacopeia from EU, US and Japan to formulation principles Includes examples and text boxes for quicker data assimilation Written for both students studying pharmacy and industry professionals in the field as well as toxicologists, biochemists, medical lab technicians, Voigt's Pharmaceutical Technology is the essential resource for understanding the various aspects of pharmaceutical technology.

Introduction to Pharmaceutical Analytical Chemistry Stig Pedersen-Bjergaard, Bente Gammelgaard, Trine G. Halvorsen, 2019-04-29 Die umfassend überarbeitete 2. Auflage enthält ein neues Kapitel zur chemischen Analyse von Biopharmazeutika, in dem die Identifizierung, Reinheitsprüfung und die Analyse von Peptiden und proteinbasierten Formulierungen erläutert werden. Die neue Auflage bietet ebenfalls verbesserte farbige Abbildungen und Tabellen, eine gestraffte Kapitelstruktur und überarbeitete Inhalte, die das Fachgebiet klarer und verständlicher präsentieren. - Bietet eine Einführung in die grundlegenden Konzepte der pharmazeutischen analytischen Chemie und Statistik. - Untersucht systematisch pharmazeutische Anwendungen, die in anderen Lehrbüchern zu dem Fachgebiet fehlen. - Untersucht verschiedene Analysetechniken, die in der Regel in Pharmalaboren zur Anwendung kommen. - Präsentiert Fragestellungen aus der Praxis, aktuelle praktische Beispiele und detaillierte Illustrationen. - Die aktualisierten Inhalte entsprechen den aktuellen europäischen und US-amerikanischen Arzneibuchvorschriften und -richtlinien.

The International Pharmacopoeia World Health Organization, 1979 The International Pharmacopoeia contains a collection of recommended methods for analysis and quality specifications for pharmaceutical substances, excipients and

products. Volume five of this publications describes methods and procedures for the quality control of pharmaceutical substances and tablets, tests for dosage forms for suppositories and ophthalmic preparations, and a new section on quality control of anti-malarials. Supplementary information on International Chemical Reference Substances and International Reference Spectra, and on the establishment, maintenance and distribution of chemical reference substances are also included.

Introduction to Pharmaceutical Chemical Analysis Steen Hansen, Steen Honoré Hansen, Stig Pedersen-Bjergaard, Knut Rasmussen, 2011-12-12 This textbook is the first to present a systematic introduction to chemical analysis of pharmaceutical raw materials, finished pharmaceutical products, and of drugs in biological fluids, which are carried out in pharmaceutical laboratories worldwide. In addition, this textbook teaches the fundamentals of all the major analytical techniques used in the pharmaceutical laboratory, and teaches the international pharmacopoeias and guidelines of importance for the field. It is primarily intended for the pharmacy student, to teach the requirements in “analytical chemistry” for the 5 years pharmacy curriculum, but the textbook is also intended for analytical chemists moving into the field of pharmaceutical analysis. Addresses the basic concepts, then establishes the foundations for the common analytical methods that are currently used in the quantitative and qualitative chemical analysis of pharmaceutical drugs Provides an understanding of common analytical techniques used in all areas of pharmaceutical development Suitable for a foundation course in chemical and pharmaceutical sciences Aimed at undergraduate students of degrees in Pharmaceutical Science/Chemistry Analytical Science/Chemistry, Forensic analysis Includes many illustrative examples

European Pharmacopoeia Council of Europe, 2010 The 7th edition of the European Pharmacopoeia was published July 15 2010 and consists of a two-volume main edition. It is complemented by non-cumulative supplements that are to be kept for the duration of the 7th Edition. Two supplements were published in 2010 and three supplements will be published in each 2011 and 2012. It contains information on all types of active substances used to prepare pharmaceutical products: various chemical substances, antibiotics, biological substances, vaccines for human or veterinary use, immunosera, radiopharmaceutical preparations, herbal drugs and homoeopathic preparations. Over 1800 specific and general monographs are included.

European Pharmacopoeia ,2001

Practical Pharmaceutics Paul Le Brun, Sylvie Crauste-Manciet, Irene Krämer, Julian Smith, Herman Woerdenbag, 2023-06-15 Practical Pharmaceutics contains essential knowledge on the preparation, quality control, logistics, dispensing and use of medicines. It features chapters written by experienced pharmacists and scientists working in hospitals, academia and industry throughout Europe, including practical examples as well as information on current GMP and GMP-based guidelines and EU-legislation. In this second edition all chapters have been updated with numerous new as well as

didactically revised illustrations and tables. A completely new chapter about therapeutic proteins and Advanced Therapy Medicinal Products was added. From prescription to production, from usage instructions to procurement and the impact of medicines on the environment, the book provides step-by-step coverage that will help a wide range of readers, students as well as professionals. It offers product knowledge for all pharmacists working directly with patients and it will enable them to make the required medicine available, to store medicines properly, to adapt medicines if necessary and to dispense medicines with the appropriate information for patients as well as caregivers about product care and how to maintain the quality of the product. The basic knowledge presented in the book will also be valuable for industrial pharmacists to remind and focus them on the application of the medicines manufactured. The basic and practical knowledge on the design, preparation and quality management of medicines can directly be applied by the pharmacists whose main duty is production in community and hospital pharmacies and in industry. Undergraduate as well as graduate pharmacy students will find knowledge presented in a coherent way and fully supported with relevant examples. Practical Pharmaceutics has become a reliable and recognised source for the acquisition of pharmaceutical-technological knowledge. The book is used in the curriculum of a number of international universities and schools of Pharmacy.

European Pharmacopoeia Consejo de Europa. Dirección para la Calidad en los Medicamentos,2013

Specialised Pharmaceutical Formulation Geoffrey D Tovey,2022-05-04 Formulation is a key step in the drug design process, where the active drug is combined with other substances that maximise the therapeutic potential, safety, and stability of the final medicinal product. Regulatory and quality demands, in addition to advances in processing technologies, result in growing challenges as well as possibilities for the field. Following on from Pharmaceutical Formulation, which covered traditional dosage forms such as tablets and capsules, this volume expands upon those formulations to cover a more diverse range of less common dosage forms. Novel routes of administration are covered from inhalational, dermal and transdermal formulations to ocular, oral suspensions, vaccines and nanoparticle drug delivery. The methods through which these formulations are processed and manufactured is also covered, providing essential knowledge to ensure quality, efficiency, and acceptable costing. Specialised Pharmaceutical Formulation is an essential, up to date resource for students and researchers working in academia and in the pharmaceutical industry and will equip readers with the ability to effectively and reliably produce products which can be approved, manufactured and made available to administer to patients.

Practical Pharmaceutics Yvonne Bouwman-Boer,V'Iain Fenton-May,Paul Le Brun,2015-08-24 This book contains essential knowledge on the preparation, control, logistics, dispensing and use of medicines. It features chapters written by experienced pharmacists working in hospitals and academia throughout Europe, complete with practical examples as well as information on current EU-legislation. From prescription to production, from usage instructions to procurement and the impact of medicines on the environment, the book provides step-by-step coverage that will help a wide range of readers. It

offers product knowledge for all pharmacists working directly with patients and it will enable them to make the appropriate medicine available, to store medicines properly, to adapt medicines if necessary and to dispense medicines with the appropriate information to inform patients and caregivers about product care and how to maintain their quality. This basic knowledge will also be of help to industrial pharmacists to remind and focus them on the application of the medicines manufactured. The basic and practical knowledge on the design, preparation and quality management of medicines can directly be applied by the pharmacists whose main duty is production in community and hospital pharmacies and industries. Undergraduate as well as graduate pharmacy students will find knowledge and backgrounds in a fully coherent way and fully supported with examples.

British Pharmacopoeia 2005 British Pharmacopoeia Commission,2005 This set comprises of five volumes and a CD-ROM: i) four volumes detailing all current UK pharmacopoeial standards for medicines for human use; ii) a companion volume providing standards for substances, preparations and immunological products used in veterinary medicine; and iii) a fully searchable CD-ROM which contains the contents of these volumes in electronic form together with a user manual, as well as the British Approved Names 2002 and supplements. The Pharmacopoeia is published on the recommendation of the Medicines Commission in accordance with the Medicines Act 1968. This edition is effective from 1 December 2005 and it incorporates the requirements of the 5th edition of the European Pharmacopoeia 2004 and its supplements.

British Pharmacopoeia (Veterinary). ,2008

Establishing quality specifications for medicines, vaccines and in vitro diagnostics World Health Organization,2024-07-10

Pharmacopoea Internationalis ,1994

Modified-Release Drug Delivery Technology Michael J. Rathbone,Jonathan Hadgraft,Michael S. Roberts,2002-11-07 Describing formulation challenges and their solutions in the design, development, and commercialization of modified-release drugs delivery systems, this book contains eighty papers that review recent developments in design and manufacturing techniques. It includes detailed descriptions of extended release drug products for the oral, nasal, ophthalmic, pulmonary, vaginal, dermal and transdermal pathways. With the exception of the final section addressing regulatory issues, each section covers a particular route for drug delivery and opens with an overview of the anatomical, physiological, and pharmaceutical basics of each route before moving on to cover specific technologies.

British Pharmacopoeia 2012 British Pharmacopoeia Commission,2011

European Pharmacopoeia European Pharmacopoeia Commission,1980

British Pharmacopoeia 2001 Great Britain. Medicines Commission,2001 The Pharmacopoeia is published by the Department of Health on behalf of the Medicines Control Agency. The MCA is the body responsible for creating the standards

for medicinal products and the control and licensing of medicines for use in the UK. The 4-volume work, CD-ROM, and web site contain 2,760 monographs for substances and articles used in the practice of medicine, including all the monographs from the updated third edition of the European pharmacopoeia. It provides all the current UK pharmacopoeial standards for medicines for human use.

WHO Drug Information ,2005

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