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New Models of Inclusive Innovation for Development
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The Second Coming of Steve Jobs
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Process Simulation and Data Modeling in Solid Oral Drug Development and Manufacture
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International Encyclopedia of Abbreviations and Acronyms in Science and Technology Yearbook
Continuous Manufacturing of Pharmaceuticals
Redesigning Education
Data Acquisition and Conversion Handbook
Handbook of Business Ethics
Modern Techniques for Food Authentication
Encyclopedia of Pharmaceutical Science and Technology, Fourth Edition, Six Volume Set (Print)
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HBR's 10 Must Reads on Strategic Marketing (with featured article ÒMarketing Myopia,Ó by Theodore Levitt)
Waking the Dead
Pharmaceutical Blending and Mixing
The Contribution of Mixed Legal Systems to European Private Law
A Deal with the Devil
ACCA P6 Advanced Taxation FA2016
Chemical Tradename Dictionary
The Reflexology Handbook
Human Rights Law in Europe
Cry Wolf
New Trends in Pharmacokinetics
40 Years of Chemometrics
The Six Sigma Way: How to Maximize the Impact of Your Change and Improvement Efforts, Second edition
Save Money: Good Food - Family Feasts for a Fiver
Pharmaceutical Process Development
International Political Economy

WILLIS ESTHER

New Models of Inclusive Innovation for Development Hachette UK
Clinical & internal medicine.
It Ain't Half Racist, Mum Piatkus Books
Pharmaceutical process research and development is an exacting, multidisciplinary effort but a somewhat neglected discipline in the chemical curriculum. This book presents an overview of the many facets of process development and how recent advances in synthetic organic chemistry, process technology and chemical engineering have impacted on the manufacture of pharmaceuticals. In 15 concise chapters the book covers such diverse subjects as route selection and economics, the interface with medicinal chemistry, the impact of green chemistry, safety, the crucial role of physical organic measurements in gaining a deeper understanding of chemical behaviour, the role of the analyst, new tools and innovations in reactor design, purification and separation, solid state chemistry and its role in formulation. The book ends with an assessment of future trends and challenges. The book provides a valuable overview of: both early and late stage chemical development, how safe and scaleable synthetic routes are designed, selected and developed, the importance of the chemical engineering, analytical and manufacturing interfaces, the key enabling technologies, including catalysis and biocatalysis, the importance of the green chemical perspective and solid form issues. The book, written and edited by experts in the field, is a contemporary, holistic treatise, with a logical sequence for process development and mini-case histories within the chapters to bring alive different aspects of the process. It is completely pharmaceutical themed, encompassing all essential aspects, from route and reagent selection to manufacture of the active compound. The book is aimed at both graduates and postgraduates interested in a career in the pharmaceutical industry. It informs them about the breadth of the work carried out in chemical research and development departments, and gives them a feel for the challenges involved in the job. The book is also of value to academics who often understand the drug discovery arena, but have far less appreciation of the drug development area, and are thus unable to advise their students about the relative merits of careers in chemical development versus discovery.

OUP Oxford
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NEW from the bestselling HBR’s 10 Must Reads series. Stop pushing products—and start cultivating relationships with the right customers. If you read nothing else on marketing that delivers competitive advantage, read these 10 articles. We’ve combed through hundreds of articles in the Harvard Business Review archive and selected the most important ones to help you reinvent your marketing by putting it—and your customers—at the center of your business. Leading experts such as Ted Levitt and Clayton Christensen provide the insights and advice you need to: • Figure out what business you’re really in • Create products that perform the jobs people need to get done • Get a bird’s-eye view of your brand’s strengths and weaknesses • Tap a market that’s larger than China and India combined • Deliver superior value to your B2B customers • End the war between sales and marketing Looking for more Must Read articles from Harvard Business Review? Check out these titles in the popular series: HBR’s 10 Must Reads: The Essentials HBR’s 10 Must Reads on Communication HBR’s 10 Must Reads on Collaboration HBR’s 10 Must Reads on Innovation HBR’s 10 Must Reads on Leadership HBR’s 10 Must Reads on Making Smart Decisions HBR’s 10 Must Reads on Managing Yourself HBR’s 10 Must Reads on Teams
Army Aviation Digest Harvard Business Review Press
A comprehensive look at existing technologies and processes for continuous manufacturing of pharmaceuticals As rising costs outpace new drug development, the pharmaceutical industry has come under intense pressure to improve the efficiency of its manufacturing processes. Continuous process manufacturing provides a proven solution. Among its many benefits are: minimized waste, energy consumption, and raw material use; the accelerated introduction of new drugs; the use of smaller production facilities with lower building and capital costs; the ability to monitor drug quality on a continuous basis; and enhanced process reliability and flexibility. Continuous Manufacturing of Pharmaceuticals prepares professionals to take advantage of that exciting new approach to improving drug manufacturing efficiency. This book covers key aspects of the continuous manufacturing of pharmaceuticals. The first part provides an overview of key chemical engineering principles and the current regulatory environment. The second covers existing technologies for manufacturing both small-molecule-based products and protein/peptide products. The following section is devoted to process analytical tools for continuously operating manufacturing environments. The final two sections treat the integration of several individual parts of processing into fully operating continuous process systems and summarize state-of-art approaches for innovative new manufacturing principles. Brings together the essential know-how for anyone working in drug manufacturing, as well as chemical, food, and pharmaceutical scientists working on continuous processing Covers chemical engineering principles, regulatory aspects, primary and secondary manufacturing, process analytical technology and quality-by-design Contains contributions from researchers in leading pharmaceutical companies, the FDA, and academic institutions Offers an extremely well-informed look at the most promising future approaches to continuous manufacturing of innovative pharmaceutical products Timely, comprehensive, and authoritative, Continuous Manufacturing of Pharmaceuticals is an important professional resource for researchers in industry and academe working in the fields of pharmaceuticals development and manufacturing.
The Second Coming of Steve Jobs CRC Press
This book brings together a number of essays on the contribution that the so-called mixed legal systems can make to the emergence of a European private law. The contributions discuss different aspects of the law of Scotland, South Africa and Louisiana, as well as some general methodological aspects of mixing legal systems, all in their relationship with the development from a new ius commune for Europe. This book contains contributions from Robin Evans-Jones, Gerhard Lubbe, Johann Neethling, Anthony Ogus, Vernon Palmer, Alan Watson and Jan Smits.
Continuous Pharmaceutical Processing Christina Tetreault
This thorough volume aims to provide pharmaceutical engineers with an introduction to the current state of the art in modeling and simulation of pharmaceutical processes and to summarize a number of practical applications of relevant methodologies in drug product development. Chapters include explorations of simulation and modeling methodologies, data collection and analysis, development of novel sensing techniques, development and integration of individual unit models, optimization approaches for data-based models, design space evaluation techniques, informatics-based methodologies, and emerging topics in pharmaceutical process development. As a part of the Methods in Pharmacology and Toxicology series, the chapters contain the kind of detail and implementation advice that will make the transition into the laboratory as smooth as possible. Authoritative and cutting edge, Process Simulation and Data Modeling in Solid Oral Drug Development and Manufacture seeks to promote research into process systems methodologies and their application in pharmaceutical product and process development, which will undoubtedly become an increasingly important area in the future.
Process Simulation and Data Modeling in Solid Oral Drug Development and Manufacture Routledge
Inequality and innovation are both rising issues on the international development agenda. Their intersection is inclusive innovation; defined as the inclusion within some aspect of innovation of groups who are currently marginalised. This is a topic of increasing interest and activity. Large firms have been working to deliver innovative goods and services for base-of-the-pyramid consumers: the c.3 billion who live on less than US$2 per day. Within poor communities, an influx of new technology, finance and capabilities has spurred more localised innovation. A variety of different models have been identified by which this activity is organised and implemented, such as inclusive innovation clusters, grassroots innovation, frugal innovation, innovation platforms, and inclusive user-producer interactions. This book explores the operation, conceptualisation and impact of these models, and analyses the nature of inclusive innovation practice and research. It will be of interest to researchers, policy-makers, strategists and other practitioners associated with these new forms of innovation. This book was originally published as a special issue of Innovation and Development.
McMeel and Virgo on Financial Advice and Financial Products Chemical Tradename Dictionary
Are you complying with health and safety regulations in the

workplace? Making mistakes in many areas of health and safety can be both incredibly dangerous and hugely costly. So what can you do to avoid hazards and expensive, time-consuming legal battles? That's where Health & Safety at Work For Dummies comes in. Cutting through the clutter, it provides you with the practical, must-know information you need to ensure your workplace is a suitably safe environment that complies with government health and safety rules and regulations. Did you know that in 2014, 1.2 million working people suffered from work-related illnesses, 2,535 mesothelioma deaths occurred due to past asbestos exposure and 133 workers were killed on the job? The list goes on – and the statistics are staggering. Health & Safety at Work For Dummies shows you how to keep your employees safe from becoming another statistic in this frightening data. Arming you with critical information needed to adhere to health and safety regulations, it offers expert guidance on managing and implementing health and safety in your business, controlling workplace risks, going the extra mile in following orders and much more. Offers an easy-to-follow overview for getting started with health and safety Provides tips and advice for planning your health and safety management Includes guidance on monitoring and reviewing your health and safety systems Clearly demonstrates how to organize and motivate your workforce to comply with rules and regulations You can't afford to run a business that doesn't provide a safe work environment. Be smart, safe and proactive with the help of this essential guide.

Health and Safety at Work For Dummies Springer Science & Business Media

This book provides analysis and critique of the dual protection of human rights in Europe by assessing the developing legal relationship between the Court of Justice of the European Union (CJEU) and the European Court of Human Rights (ECtHR). The book offers a comprehensive consideration of the institutional framework, adjudicatory approaches, and the protection of material rights within the law of the European Union and the European Convention on Human Rights (ECHR). It particularly explores the involvement and participation of stakeholders in the functioning of the EU and the ECtHR, and asks how well the new legal model of ‘the EU under the ECtHR’ compares to current EU law, the ECHR and general international law. Including contributions from leading scholars in the field, each chapter sets out specific case-studies that illustrate the tensions and synergies emergent from the EU-ECHR relationship. In so doing, the book highlights the overlap and dialectic between Europe’s two primary international courts. The book will be of great interest to students and researchers of European Law and Human Rights.

The Economics of Innocent Fraud BPP Learning Media

The scream heard by no one is the deadliest. In the rural parishes of Louisiana's French Triangle, young women are disappearing one by one, only to turn up on the banks of the bayou, strangled and cast aside where they are sure to be found. But there is one trophy the killer prizes above all others, one woman who must be silenced forever.... Attorney Laurel Chandler did not come back to Bayou Breaux to seek justice. That once-burning obsession had destroyed her credibility, her career, her marriage—and nearly her sanity. But when a ruthless predator strikes too close to home, she's lured into a perverse game from which there may be no escape. Once before, Laurel's cries against a monstrous evil went unanswered. Who will listen now?

Scottish Legal System Penguin Books, Limited (UK)

Chemical Tradename DictionaryJohn Wiley & Sons

Introduction to Design of Experiments with JMP Examples Edinburgh University Press

The last decade or so has witnessed tremendous progress in methodology in the field of drug development in general and pharmacokinetics in particular. Clinical pharmacokinetics is using new tools for probing into the "black box" once being ac cessible only partly through experimental techniques and, mostly through mathemati cal and computer means. Development of computerized scanning, positron emission tomography (PET), stereoselectivity and other techniques are now enabling investi gators to have better pictures of the systems they are studying. Mathematical models through computer simulation and statistical estimation, mostly due to easy access be cause of inexpensive yet powerful personal computers, are enabling us to investigate ultrastructures and their functional connectivity in more detail. As a consequence, new hypotheses are being formed and tested in various related fields. In clinical pharmacokinetics, mostly due to mathematical modeling, more accurate interspecies scaling of pharmacokinetic parameters and dosimetry can be done now-a-days. The concept of "a human is a bigger rat" does not necessarily fly as a consequence. Pharmacokinetic concepts are becoming powerful tools in meaningful carcinogenic and toxic risk extrapolation of different chemicals in humans. New dose delivery designs are being formulated using pharmacokinetic techniques for different pharmaceutical compounds. Investigations continue in the academia, research institutions, pharmaceutical, biotechnological, and agricultural industries in developmental and physiological aspects of different chemicals for the benefit of mankind. The idea of a school on "New Trends in Pharmacokinetics", from which the present pUblication was made possible, took shape over almost a year.

Data Acquisition Handbook Lynne Rienner Pub

Hyacinth always assumed dying would simplify her life. But when her new boss, Archangel Michael, sends her on her first official mission—to retrieve a powerful rock from a collector in Germany—things go downhill fast. For one thing, the Dead keep popping up, expecting her to guide them to the Afterlife. For another, her part-demon nephew Geordi's powers are starting to leak out, at age seven. What if Michael finds out about him? Worse, what if Satan does? Then there's her love life-after-death. Rooming with a dead French cop no one else can see is complicated enough. But when Jason, Geordi's lying Dioguardi Demon cousin, resurrects himself—so to speak—all Hell breaks loose. Literally. Can Hyacinth get Michael's rock back before Satan steals its powers and breaks free of his prison? Or will her single-minded pursuit put those she loves—and the rest of the world—in the path of Satan's fury?

Interpreting Diffuse Reflectance and Transmittance John Wiley & Sons

A man can only have one great love, and mine was the sea. Until I pulled a half-drowned heiress out of the water. Until I kept her as mine. Until I hurt her to prove I didn’t love her. Now we’re surrounded. Dangerous pirates want to use her for their own purposes. I won’t give her up that easy. I have a lifetime of treasure to protect. But with the ship going down, there’s only one thing I want to save. Her. A Deal with the Devil is book two in the Devil trilogy by USA Today bestselling author Amelia Wilde. The Devil and the Deep Blue Sea must be read first.

International Encyclopedia of Abbreviations and Acronyms in Science and Technology Yearbook Routledge

From the acclaimed Vanity Fair and GQ journalist–an unprecedented, in-depth portrait of the man whose return to Apple precipitated one of the biggest turnarounds in business history. With a new epilogue on Apple’s future survival in today’s roller-coaster economy, here is the revealing biography that blew away the critics and stirred controversy within industry and media circles around the country.

Continuous Manufacturing of Pharmaceuticals Nicholson

Whether youOCOre studying Law in Scotland or looking to convert to Scots law, this invaluable guide will quickly equip you with all the basics of the Scottish legal system. Fully updated for the third edition, it is the ideal textbook for busy law students and revising for those all-important exams. Summary sections of Essentials Facts and Essential Cases will help you to identify, understand and remember the key elements of the subject.&quote;

Redesigning Education Currency

As we hurtle through the second decade of the 21st century, the pressure for radical change to mainstream education is becoming ever more urgent, and images for what that future might look like are emerging and coalescing. But there is a problem. There is no version of this complex, exciting new world of learning that can arrive fully operational and ready to open for business on Monday morning. Redesigning Education captures the journeys of cities and countries around the world as they travel from the education systems they have to the education systems they want and need- and highlights the tools and processes they are using along the way. The Global Education Leaders' Program, or GELP, is a partnership of teams of education system leaders and world-class organizations collaborating to transform education. Its members include some of the highest performing countries in the world and those where providing education for all is an extraordinary challenge. As a community, these teams have developed, refined and shared a compelling and-critically-actionable vision for the future of education: Education 3.0. Redesigning Education is about what it takes to transform education systems when the context in which they've flourished fundamentally alters. While a fully operational 21st century education system does not yet exist, GELP teams are identifying a clear set of models and practices that together form a "roadmap" to the future, backed up by the practical lessons drawn from their experiences. More than simply a compelling case for change, Redesigning Education offers real insights grown from stories of system transformation around the globe.

Data Acquisition and Conversion Handbook Humana Press

Written in four parts, this book provides a dedicated and in-depth reference for blending within the pharmaceutical manufacturing industry. It links the science of blending with regulatory requirements associated with pharmaceutical manufacture. The contributors are a combination of leading academic and industrial experts, who provide an informed and industrially relevant perspective of the topic. This is an essential book for the pharmaceutical manufacturing industry, and related academic researchers in pharmaceutical science and chemical and mechanical engineering.

Handbook of Business Ethics Amelia Wilde

Pharmaceutical science deals with the whole spectrum of drug development from start to finish. There are many different facets to the pharmaceutical industry, from initial research to the finished product, including the equipment used, trials performed, and regulations that must be followed. Presenting an overview of all of these different aspects, the Encyclopedia of Pharmaceutical Science and Technology, Fourth Edition is a must-have reference guide for all laboratories and libraries in the pharmaceutical field. Bringing together leaders from every specialty related to pharmaceutical science and technology, this is the single-source reference at the forefront of pharmaceutical R&D. The strength of this work is not only its breadth but also the caliber of contributing writers, all experts in their field, writing on all aspects of pharmaceutical science and technology. The fourth edition offers 29 new chapters ranging from biomarkers, computational chemistry, and contamination control to high-throughput screening, orally disintegrating tablets, and quality by design. The encyclopedia details best practices of equipment used, methods for manufacturing, options for packaging, and routes for drug delivery. The volumes also provide a thorough understanding of the choices behind each method. In addition, the regulations, safety aspects, patent guidance, and methods of analysis are presented. Key Areas Covered: Analytics Biomarkers Dosage forms Drug delivery Formulation Informatics Manufacturing Packaging Processing Regulatory affairs Systems validation This is an authoritative reference source for those practicing in any area of pharmaceutical science and technology, enabling the pharmaceutical specialist and novice alike to keep abreast of developments in this constantly evolving and highly competitive field. * Online version coming soon. Contact us to inquire about subscription options and print/online combination packages. US: (Tel) 1.888.318.2367 / (E-mail) e-reference@taylorandfrancis.com International: (Tel) +44 (0) 20 7017 6062 / (E-mail) online.sales@tandf.co.uk