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Marc A. LeBeau, Ashraf Mozayani

Countering the Problem of Falsified and Substandard Drugs Institute of Medicine, Board on Global Health, Committee on Understanding the Global Public Health Implications of Substandard, Falsified, and Counterfeit Medical Products, 2013-06-20 The adulteration and fraudulent manufacture of medicines is an old problem, vastly aggravated by modern manufacturing and trade. In the last decade, impotent antimicrobial drugs have compromised the treatment of many deadly diseases in poor countries. More recently, negligent production at a Massachusetts compounding pharmacy sickened hundreds of Americans. While the national drugs regulatory authority (hereafter, the regulatory authority) is responsible for the safety of a country's drug supply, no single country can entirely guarantee this today. The once common use of the term counterfeit to describe any drug that is not what it claims to be is at the heart of the argument. In a narrow, legal sense a counterfeit drug is one that infringes on a registered trademark. The lay meaning is much broader, including any drug made with intentional deceit. Some generic drug companies and civil society groups object to calling bad medicines counterfeit, seeing it as the deliberate conflation of public health and intellectual property concerns. Countering the Problem of Falsified and Substandard Drugs accepts the narrow meaning of counterfeit, and, because the nuances of trademark infringement must be dealt with by courts, case by case, the report does not discuss the problem of counterfeit medicines.

Martindale Sean C. Sweetman, 2006-01-01 This is thirty-fifth edition of Martindale, which provides reliable, and evaluated information on drugs and medicines used throughout the world. It contains encyclopaedic facts about drugs and medicines, with: 5,500 drug monographs; 128,000 preparations; 40,700 reference citations; 10,900 manufacturers. There are synopses of disease treatments which enables identification of medicines, the local equivalent and the manufacturer. It also Includes herbals, diagnostic agents, radiopharmaceuticals, pharmaceutical excipients, toxins, and poisons as well as drugs and medicines. Based on published information and extensively referenced

Drug-Induced Liver Injury , 2019-07-13 Drug-Induced Liver Injury, Volume 85, the newest volume in the Advances in Pharmacology series, presents a variety of chapters from the best authors in the field. Chapters in this new release include Cell death mechanisms in DILI, Mitochondria in DILI, Primary hepatocytes and their cultures for the testing of drug-induced liver injury, MetaHeps an alternate approach to identify IDILI, Autophagy and DILI, Biomarkers and DILI, Regeneration and DILI, Drug-induced liver injury in obesity and nonalcoholic fatty liver disease, Mechanisms of Idiosyncratic Drug-Induced Liver Injury, the Evaluation and Treatment of Acetaminophen Toxicity, and much more. Includes the authority and expertise of leading contributors in pharmacology Presents the latest release in the Advances in Pharmacology series

India and the Patent Wars Murphy Halliburton, 2017-11-15 India and the Patent Wars contributes to an international debate over the costs of medicine and restrictions on access under stringent patent laws showing how activists and drug companies in low-income countries seize agency and exert influence over these processes. Murphy Halliburton contributes to analyses of globalization within the fields of anthropology, sociology, law, and public

health by drawing on interviews and ethnographic work with pharmaceutical producers in India and the United States. India has been at the center of emerging controversies around patent rights related to pharmaceutical production and local medical knowledge. Halliburton shows that Big Pharma is not all-powerful, and that local activists and practitioners of ayurveda, India's largest indigenous medical system, have been able to undermine the aspirations of multinational companies and the WTO. Halliburton traces how key drug prices have gone down, not up, in low-income countries under the new patent regime through partnerships between US- and India-based companies, but warns us to be aware of access to essential medicines in low- and middle-income countries going forward.

Viagra, cialis e levitra Edições Lebooks,2014-03-07 O objetivo deste eBook é mostrar de forma comparativa três dos medicamentos mais utilizados atualmente para a disfunção erétil: o Viagra, Cialis e Levitra, de forma que você possa conhecer o funcionamento de cada um, os cuidados necessários e as recomendações para o seu uso correto. Além dos testes comparativos de eficiência, o ebook fornece orientações e dicas, tais como: uso do viagra, cialis e levitra com álcool, a duração dos efeitos, como maximizar e prolongar os resultados e evitar os efeitos colaterais.

Generic Drug Product Development Isadore Kanfer,Leon Shargel,2016-04-19 Due to a worldwide need for lower cost drug therapy, use of generic and multi-source drug products have been increasing. To meet international patent and trade agreements, the development and sale of these products must conform to national and international laws, and generic products must prove that they are of the same quality and are therapeutica

Rang & Dale's Pharmacology James M. Ritter,Rod J. Flower,Graeme Henderson,Yoon Kong Loke,David MacEwan,Humphrey P. Rang,2018-11-04 World-renowned coverage of today's pharmacology at your fingertips Keeps you up-to-date with new information in this fast-changing field, including significantly revised coverage of CNS drugs, cognitive enhancers, anti-infectives, biologicals/biopharmaceuticals, lifestyle drugs, and more. Includes access to unique features, including more than 100 brand new chapter-specific multiple-choice questions and 6 new cases for immediate self-assessment. Features a color-coded layout for faster navigation and cross-referencing. Clarifies complex concepts with Key Points boxes, Clinical Uses boxes and full-color illustrations throughout.

Private Patents and Public Health Ellen F. M. 't Hoen,2016 Millions of people around the world do not have access to the medicines they need to treat disease or alleviate suffering. Strict patent regimes introduced following the establishment of the World Trade Organization in 1995 interfere with widespread access to medicines by creating monopolies that keep medicines prices well out of reach for many. The AIDS crisis in the late nineties brought access to medicines challenges to the public's attention, when millions of people in developing countries died from an illness for which medicines existed, but were not available or affordable. Faced with an unprecedented health crisis ? 8,000 people dying daily ? the public health community launched an unprecedented global effort that eventually resulted in the large-scale availability of low-priced generic HIV medicines. But now, high prices of new medicines - for example, for cancer, tuberculosis and hepatitis C - are limiting access to

treatment in low-, middle and high-income countries alike. Patent-based monopolies affect almost all medicines developed since 1995 in most countries, and global health policy is now at a critical juncture if the world is to avoid new access to medicines crises. This book discusses lessons learned from the HIV/AIDS crisis, and asks whether actions taken to extend access and save lives are exclusive to HIV or can be applied more broadly to new global access challenges.

Drug-Facilitated Sexual Assault Marc A. LeBeau, Ashraf Mozayani, 2001-08-28 Drug-Facilitated Sexual Assault educates readers on the ways in which drugs are used as weapons in committing sexual assaults and how to successfully investigate these crimes. It looks at the history of these crimes over the years, and includes an in-depth discussion of the drugs and drug classes in use today. It describes the effects of these drugs on the victims, the process for reporting these crimes, details on the type of person who uses drugs to sexually assault an individual, and obstacles to investigating the suspect. The authors show the proper techniques in collecting and analyzing evidence; ways to overcome some of the unique difficulties encountered in these types of investigations; and how to work with other professionals to prosecute these cases successfully. The concluding appendixes are valuable samples of the necessary forms needed to complete these investigations. This book is ideal for anyone involved in investigating these crimes, including forensic scientists, law enforcement officers, lawyers, toxicologists, and medical professionals. * Ideal for everyone involved in the investigation of these crimes, including forensic scientists, police officers, lawyers, toxicologists and medical professionals

Natural Standard Herb & Supplement Reference Natural Standard (Firm), 2005 Provides detailed, evidence-based reviews for 98 herbs and dietary supplements.

EBOOK: Regulating Pharmaceuticals in Europe: Striving for Efficiency, Equity and Quality Elias Mossialos, Monique Mrazek, Tom Walley, 2004-06-16 This thoughtful and comprehensive book represents the best work I have seen on the current situation concerning medication policies in the EU. It is not just that this is a very up-to-date compendium of facts and data across a wide variety of domains that impact on pharmaceutical regulation. The book is also strong on analysis of those facts as well. Jerry Avorn, Harvard Medical School. This book offers a comprehensive examination of approaches to manage pharmaceutical expenditures in Europe. It is a must-read for those who seek to understand and navigate the changing regulatory environment for medicines in the European Union. Bernie O'Brien, McMaster University, Canada. The rising cost of pharmaceutical expenditures in many European countries is of concern to governments required to make effective use of health care budgets. Taking a broad perspective that encompasses institutional, political and supranational aspects of pharmaceutical regulation, this book examines approaches used to manage pharmaceutical expenditure across Europe and what impact these strategies have had on efficiency, quality, equity and cost of pharmaceutical care. Regulating Pharmaceuticals in Europe is an important book for students of health policy, regulation and management, and for health managers and policy makers. The editors: Elias Mossialos is Brian Abel-Smith Professor of Health Policy at the London School of Economics and Political Science and a Research Director of the European Observatory on Health Systems and

Policies. Monique Mrazek is a Health Economist (Europe and Central Asia region) for the World Bank and formerly a Research Officer in Health Economics for the European Observatory on Health Systems and Policies. Tom Walley is Professor of Clinical Pharmacology at the University of Liverpool and Director of the UK National Health Technology Assessment Programme. Contributors: Julia Abelson, Christa Altenstetter, Vittorio Bertele', Christine Bond, Marcel L. Bouvy, Colin Bradley, Steve Chapman, Anna Dixon, Michael Drummond, Pierre Durieux, Edzard Ernst, Armin Fidler, Eric Fortess, Richard Frank, Silvio Garattini, Leigh Hancher, Ebba Holme Hansen, Steve Hudson, Kees de Jonchere, Panos Kanavos, Sjoerd Kooiker, Jean-Marc Leder, Graham Lewis, Donald W. Light, Alistair McGuire, Elias Mossialos, Monique Mrazek, Maria Pia Orru', Govin Permanand, Guenka Petrova, Munir Pirmohamed, Dennis Ross-Degnan, Frans Rutten, Steven Soummerai, David Taylor, Sarah Thomson, Tom Walley.

Drug-Induced Liver Toxicity Minjun Chen,Yvonne Will,2018-03-21 This book provides a comprehensive view of the methodologies used for the study of liver toxicity encountered throughout the whole life cycle of a drug, from drug discovery, to clinical trial, post-marketing, and even clinical practice. Organized into six sections, the first section introduces the mechanisms contributing to drug-induced liver toxicity. The second and third section explore in silico and in vitro approaches used to help mitigate hepatotoxicity liability at the early stages of drug development. The fourth section describes methodologies applied in regulatory processes, including preclinical studies, clinical trials, and post-marketing surveillance. The fifth section discusses clinical hepatotoxicity. Emerging technologies are examined in the final section. As a volume in the Methods in Pharmacology and Toxicology series, chapters include the kind of expert advice that will lead to optimal results. Authoritative and practical, *Drug-Induced Liver Toxicity* serves all those who aim to improve assessment and understanding of hepatotoxic potentials of new medications and marketed drugs. Chapter 30 is open access under a CC BY 4.0 license via link.springer.com.

An Introduction to Critical Discourse Analysis in Education Rebecca Rogers,2011-04-06 Accessible yet theoretically rich, this landmark text introduces key concepts and issues in critical discourse analysis and situates these within the field of educational research. The book invites readers to consider the theories and methods of three major traditions in critical discourse studies – discourse analysis, critical discourse analysis, and multimodal discourse analysis -- through the empirical work of leading scholars in the field. Beyond providing a useful overview, it contextualizes CDA in a wide range of learning environments and identifies how CDA can shed new insights on learning and social change. Detailed analytic procedures are included – to demystify the process of conducting CDA, to invite conversations about issues of trustworthiness of interpretations and their value to educational contexts, and to encourage researchers to build on the scholarship in critical discourse studies. This edition features a new structure; a touchstone chapter in each section by a recognized expert (Gee, Fairclough, Kress); and a stronger international focus on both theories and methods. NEW! Companion Website with Chapter Extensions; Interviews; Bibliographies; and Resources for Teaching Critical Discourse Analysis.

The Rise of Viagra Meika Loe,Associate Professor of Sociology and Women's Studies Meika Loe,2004-08-11 Provides

details on the history of Viagra and the social phenomenon that surrounds it.

Patents and Innovation Trends and Policy Challenges OECD,2004-02-02 Few systematic economic evaluations have been carried out on patent system to better inform policy choices. This report, which covers a range of areas, and highlights some issues that policy makers should address in the near future, including ...

Derecho de marcas y proceso penal Juan José Caselles Fornés,2018-04-09 La falsificación de marcas se ha convertido en un fenómeno de dimensión internacional que alcanza en la actualidad una especial gravedad, afectando no sólo a los titulares de esos derechos de propiedad industrial sino también a la sociedad en su conjunto. La globalización de la economía y, con ello, de la falsificación de marcas, exigen una respuesta también global al fenómeno. Sin embargo, todos los intentos internacionales de armonizar la protección penal de las marcas han terminado fracasando, requiriendo la aportación de nuevos argumentos y sensibilidades. La territorialidad de la ley penal supone un obstáculo a la persecución de estos delitos, resultando necesario un mayor nivel de cooperación policial y judicial en el ámbito internacional. Esta obra cubre tanto los aspectos pre-procesales de la investigación criminal como los del propio proceso penal seguido en los delitos de usurpación de marcas, analizando, entre otras cuestiones, la dimensión económica y la internacionalización de la falsificación de marcas, la delincuencia organizada presente en esta materia, los organismos internacionales con competencia en la persecución de estos delitos en el ámbito nacional e internacional, los intentos internacionales de homogeneizar la protección penal, los instrumentos internacionales en la lucha contra este fenómeno, la tipificación penal de la infracción y el proceso penal seguido por falsificación de marcas, el régimen especial de medidas aduaneras contra las falsificaciones establecido en el Reglamento (UE) núm. 608/2013, incluyendo una referencia a las mercancías en tránsito, así como la necesidad de reforzar los instrumentos procesales en orden a la defensa penal contra la falsificación de marcas.

Integrative Medicine David Rakel,2007-01-01 Drawing on solid scientific evidence as well as extensive first-hand experience, this manual provides the practical information you need to safely and effectively integrate complementary and alternative treatment modalities into your practice. It explains how alternative therapies can help you fight diseases that do not respond readily to traditional treatments... presents integrative treatments for a full range of diseases and conditions, including autism, stroke, chronic fatigue syndrome, and various forms of cancer...explores how to advise patients on health maintenance and wellness...and offers advice on topics such as meditation, diet, and exercises for back pain. 24 new chapters, a new organization, make this landmark reference more useful than ever. Provides dosages and precautions to help you avoid potential complications.Delivers therapy-based pearls to enhance your patient care.Facilitates patient education with helpful handouts. Offers helpful icons that highlight the level and quality of evidence for each specific modality.Includes bonus PDA software that lets you load all of the therapeutic review sections onto your handheld device.Presents a new organization, with numerous section headings and subheadings, for greater ease of reference.Provides additional clinical practice and business considerations for incorporating integrative medicine

into clinical practice.

Antimicrobial Resistance in the Americas, 2001-09 This book makes a valuable contribution to the surveillance of resistance to antibiotics. The text offers noteworthy articles grouped under two major categories: monitoring bacterial resistance to antimicrobial drugs and factors determining the use of antimicrobials. The goal of this work is to increase awareness of the problem to promote surveillance activities and to find the best ways to apply preventive measures so that antibiotics are used judiciously with both humans and animals.

Reproductive Health and Human Rights Rebecca J. Cook, Bernard M. Dickens, Mahmoud F. Fathalla, 2003-04-17 The concept of reproductive health promises to play a crucial role in improving women's health and rights around the world. It was internationally endorsed by a United Nations conference in 1994, but remains controversial because of the challenge it presents to conservative agencies: it challenges policies of suppressing public discussion on human sexuality and regulating its private expressions. *Reproductive Health and Human Rights* is designed to equip healthcare providers and administrators to integrate ethical, legal, and human rights principles in protection and promotion of reproductive health, and to inform lawyers and women's health advocates about aspects of medicine and healthcare systems that affect reproduction. Rebecca Cook, Bernard Dickens, and Mahmoud Fathalla, leading international authorities on reproductive medicine, human rights, medical law, and bioethics, integrate their disciplines to provide an accessible but comprehensive introduction to reproductive and sexual health. They analyse fifteen case-studies of recurrent problems, focusing particularly on resource-poor settings. Approaches to resolution are considered at clinical and health system levels. They also consider kinds of social change that would relieve the underlying conditions of reproductive health dilemmas. Supporting the explanatory chapters and case-studies are extensive resources of epidemiological data, human rights documents, and research materials and websites on reproductive and sexual health. In explaining ethics, law, and human rights to healthcare providers and administrators, and reproductive health to lawyers and women's health advocates, the authors explore and illustrate limitations and dysfunctions of prevailing health systems and their legal regulation, but also propose opportunities for reform. They draw on the values and principles of ethics and human rights recognized in national and international legal systems, to guide healthcare providers and administrators, lawyers, governments, and national and international agencies and legal tribunals. *Reproductive Health and Human Rights* will be an invaluable resource for all those working to improve services and legal protection for women around the world. Updates to this book, and information on translations to French, Spanish, Portuguese, Chinese and Arabic are now available at www.law.utoronto.ca/faculty/cook/ReproductiveHealth.html

Health & Human Development in the New Global Economy World Health Organization, 2000 This publication is the result of a collaborative effort between the Pan American Health Organization and the WHO Collaborating Center for International Health at the University of Texas Medical Branch. The chapters are based on the proceedings from a seminar/workshop held in Galveston Texas in 1998. The objective of this forum was to examine the impact of the economic globalization process and technological change on health trends in the Region of the Americas and even

more specifically to explore how civil society groups can influence the formulation of healthy public policies and monitor equity in terms of both the health situation and the distribution of health care resources. This publication addresses a wide range of issues affecting health in the global economy including various types of economic organization and potential dangers the world faces as a new century begins. It centers on experiences opportunities and risks in the Americas and on the value of health in human development and of international and regional integration processes in the formulation of a new global economy. It includes articles analyzing the impact of economic reform policies on health in general as well as monographs written from the perspective of civil society groups in specific countries. The variety of viewpoints and lessons learned in this publication will be of use to decision-makers health and development experts and emerging civil society groups in Latin America and the Caribbean and can guide them in better addressing the enormous risks facing all sectors of society as a result of globalization by enabling them to take advantage of the tremendous potential this process affords.

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