

**Marco Mamone Capria** 

Science, Politics And The Pharmaceutical Industry John Abraham, John Abraham University of Reading., 2023-05-31 Drug disasters from Thalidomide to Opren and other less dramatic cases of drug injury raise questions about whether the testing and control of medicines provides satisfactory protection for the public In this revealing study John Abrahan develops a theoretically challenging realist approach in order to probe deeply into the work of scientists in the pharmaceutical industry and governmental drug regulatory authorities on both sides of the Atlantic Through the examination of contemporary controversial case studies he exposes how the commercial interest of drug manufacturers are consistently given the benefit of the scientific doubts about medicine safety and effectiveness over and above the best interests of patients A highly original combination of philosophical rigour historical sensitivity and empirical depth enables the black box of industrial and government science to be opened up to critical scrutiny much more than in previous social scientific study All major aspects of drug testing and regulation are considered including pre clinical animal tests clinical trials and postmarketing surveillance of adverse drug reactions The author argues that drug regulators are too dependent on pharmaceutical industry resources and expertise and too divorced from public accountability. The problem of corporate bias is particularly severe in the UK where regulatory decisions about medicine safety are shrouded in greater secrecy than in the US Since the purpose of drug regulation should be to maximize the safety and effectiveness of medicines for patients the public needs and deserves policies to counteract corporate bias in drug testing and evaluation John Abraham's realist analysis provides a robust basis for policy interventions at the institutional and legislative levels He proposes that corporate bias could be reduced by more extensive freedom of information greater autonomy of government scientists from pharmaceutical industry the development of independent drug testing by the regulatory authority increased patient representation on regulatory committees and more frequent and thorough oversight of regulatory performance by the legislature This book should be of interest to anyone who cares about how medicines should be controlled in modern society It should prove particularly rewarding for students and researchers in the sociology of science and technology science and medicines policy medical sociologists the medical and pharmaceutical professions and consumer organizations Politics, and the Pharmaceutical Industry John Abraham, Drug disasters from Thalidomide to Opren and other less dramatic cases of drug injury raise questions about whether the testing and control of medicines provides satisfactory protection for the public In this revealing study John Abrahan develops a theoretically challenging realist approach in order to probe deeply into the work of scientists in the pharmaceutical industry and governmental drug regulatory authorities on both sides of the Atlantic Through the examination of contemporary controversial case studies he exposes how the commercial interest of drug manufacturers are consistently given the benefit of the scientific doubts about medicine safety and effectiveness over and above the best interests of patients A highly original combination of philosophical rigour historical sensitivity and empirical

depth enables the black box of industrial and government science to be opened up to critical scrutiny much more than in previous social scientific study All major aspects of drug testing and regulation are considered including pre clinical animal tests clinical trials and postmarketing surveillance of adverse drug reactions. The author argues that drug regulators are too dependent on pharmaceutical industry resources and expertise and too divorced from public accountability. The problem of corporate bias is particularly severe in the UK where regulatory decisions about medicine safety are shrouded in greater secrecy than in the US Since the purpose of drug regulation should be to maximize the safety and effectiveness of medicines for patients the public needs and deserves policies to counteract corporate bias in drug testing and evaluation John Abraham s realist analysis provides a robust basis for policy interventions at the institutional and legislative levels. He proposes that corporate bias could be reduced by more extensive freedom of information greater autonomy of government scientists from pharmaceutical industry the development of independent drug testing by the regulatory authority increased patient representation on regulatory committees and more frequent and thorough oversight of regulatory performance by the legislature. This book should be of interest to anyone who cares about how medicines should be controlled in modern society. It should prove particularly rewarding for students and researchers in the sociology of science and technology science and medicines policy medical sociologists the medical and pharmaceutical professions and consumer organizations

Cranston's Consumers and the Law Colin Scott, Julia Black, 2000-08 The third edition of this text is designed to bring the reader up to date with developments in consumer law up to 1999 It includes material on utilities and financial services The ^AOxford Handbook of Expertise and Democratic Politics ,2023-05-05 In The Oxford Handbook of regulation Expertise and Democratic Politics Gil Eyal and Thomas Medvetz have brought together a broad group of scholars who have engaged substantively and theoretically with debates regarding the nature of expertise and the social roles of experts to examines these areas within sociology and allied disciplines The analyses take an historical and relational approach to the topic and are motivated by the sense that growing mistrust in experts represents a danger to democratic politics today Bringing together investigations from social scientists philosophers and legal scholars into the political dimensions of expertise this Handbook connects interdisciplinary work done in science and technology studies with the more classic concerns topics and concepts of sociologists of professions and intellectuals Regulation of the Pharmaceutical Industry J. Abraham, H. Lawton Smith, 2003-06-10 How are pharmaceutical technologies developed and controlled in our societies To what extent should the availability of these technologies be determined by scientific experts a democratic state the interests of final users or ethical principles This unique collection brings together the work of social scientists ethicists lawyers and policy analysts on regulation ethics and innovation in the pharmaceutical industry Regulatory systems and their implications for public health in North America Europe and developing countries are discussed including case studies of norplant interferon and anti fertility vaccines The Politics of Pharmaceutical Policy Reform Elize Massard da Fonseca, 2014-11-17

Brazil is renowned worldwide for its remarkable reforms in pharmaceutical regulation which have enhanced access to essential medicines while lowering drug costs This book innovates by analysing the generic drug reform in Brazil demonstrating that pharmaceutical regulation is only partially influenced by non state actors Little is known about the institutional antecedents and policy process that channeled this regulatory reform This is particularly intriguing because a regulatory shift in the pharmaceutical sector requires the participation of a number of stakeholders and interest groups in the policy process Fonseca examines the generic drug reform s causes and consequences No study has approached the generic drug regulation in Brazil from this perspective The Politics of Pharmaceutical Policy Reform A Study of Generic Drug Regulation in Brazil explores the following The politics of pharmaceutical regulation in Brazil over the last 25 years The political negotiations to approve the Generic Drug Act which involved a hard to reach agreement between the pharmaceutical industry national and multinational the Ministry of Health and Congress The controversial decisions to regulate packaging and pharmaceutical equivalence The surprising success of Brazilian pharmaceutical firms which became market champions in a sector largely dominated by multinational firms Comparative lessons from the Brazilian case for the political construction of regulatory standards to regulate generic drugs and its effects on global health This book will interest political scientists and health policy scholars concerned with the political conflicts in the pharmaceutical sector It argues against well established approaches to regulatory capture such as control of the regulatory process by interest groups and policy diffusion It can be used as evidence for graduate courses in public policy health policy and political science Because Brazil is one of the largest markets for pharmaceuticals in the world business leaders and consultancy firms would also be Ways of Regulating Drugs in the 19th and 20th Centuries V. Hess, 2012-12-03 This collection takes the interested perspective that the historiography of science technology and medicine needs a broader approach toward regulation The authors explore the distinct social worlds involved in regulation the forms of evidence and expertise mobilized and means of intervention chosen to tame drugs in factories consulting rooms and courts Science and the Citizen Marco Mamone Capria, 2013-06-07 This collection of essays comes from the international project Science and Democracy It offers an examination of several controversial issues within and about science of wide ranging social relevance A partial list runs as follows the role of scientific technology in shaping our life the influence of corporations on contemporary medicine grass roots activism and new technologies environmental constraints on economical growth the HIV AIDS controversy the Wakefield trial and the MMR vaccine autism link the organ transplant ideology and business the debate on the terrorist attacks in USA of September 11 2001 the role of whistleblowers in science etc Contributions by J Barretto Bastos Filho H Bauer M Brown M C Danhoni Neves F Fabbri P Ghisellini S Lang A Liversidge C Lor M Mamone Capria R Maruotti D Mastrangelo S Maurano M Mazzucco D Rasnick S Siminovic S Ulgiati M Walker The Risks of Prescription Drugs Donald W. Light, 2010-10-14 Few people realize that prescription drugs have become a leading cause of death disease and

disability Adverse reactions to widely used drugs such as psychotropics and birth control pills as well as biologicals result in FDA warnings against adverse reactions The Risks of Prescription Drugs describes how most drugs approved by the FDA are under tested for adverse drug reactions yet offer few new benefits Drugs cause more than 2 2 million hospitalizations and 110 000 hospital based deaths a year Serious drug reactions at home or in nursing homes would significantly raise the total Women older people and people with disabilities are least used in clinical trials and most affected Health policy experts Donald Light Howard Brody Peter Conrad Allan Horwitz and Cheryl Stults describe how current regulations reward drug companies to expand clinical risks and create new diseases so millions of patients are exposed to unnecessary risks especially women and the elderly They reward developing marginally better drugs rather than discovering breakthrough life saving drugs The Risks of Prescription Drugs tackles critical questions about the pharmaceutical industry and the privatization of risk To what extent does the FDA protect the public from serious side effects and disasters What is the effect of giving the private sector and markets a greater role and reducing public oversight This volume considers whether current rules and incentives put patients health at greater risk the effect of the expansion of disease categories the industry s justification of high U S prices and the underlying shifts in the burden of risk borne by individuals in the world of pharmaceuticals Chapters cover risks of statins for high cholesterol SSRI drugs for depression and anxiety and hormone replacement therapy for menopause A final chapter outlines six changes to make drugs safer and more effective Suitable for courses on health and aging gender disability and minority studies this book identifies the Risk Proliferation Syndrome that maximizes the number of people exposed to these risks Additional Columbia SSRC books on the privatization of risk and its implications for Americans Bailouts Public Money Private ProfitEdited by Robert E Wright Disaster and the Politics of InterventionEdited by Andrew Lakoff Health at Risk America's Ailing Health System and How to Heal ItEdited by Jacob S Hacker Laid Off Laid Low Political and Economic Consequences of Employment InsecurityEdited by Katherine S Newman Pensions Social Security and the Privatization of RiskEdited by Mitchell A Orenstein The Therapeutic Nightmare John Abraham, Julie Sheppard, 2014-03-18 How do drugs get to the market What controls are there and what procedures for monitoring their effects And how adequate are the regulators in protecting public health when new drugs have serious side effects The Therapeutic Nightmare tells the story of the sleeping pill Halcion a story which is far from over First marketed in the 1970s Halcion has been taken by millions of patients around the world For many years it has been associated with serious adverse effects such as amnesia hallucinations aggression and in extreme cases homicide Thirteen years after its first release it was banned by the British government It remains on sale in the United States and many other countries This book explains why patients have come to be exposed to Halcion's risks and examines the corporate interests of the manufacturers the professional interests of the scientists and medical researchers and the interests of patients in safe and effective medication It reveals how these contending forces shape the regulatory decision making process about drug safety As the number of new drugs and health products grows a major challenge facing regulators and the medical profession is how to put the interests of public health decisively and consistently above the commercial interests of the drugs industry while becoming more accountable to patient and consumer organizations

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