CAMBRIDGE HISTORY OF MEDICINE

# THE PROGRESS OF EXPERIMENT

Science and Therapeutic Reform in the United States, 1900-1990

HARRY M. MARKS



# Progress Of Experiment Science And Therapeutic Reform In The United States 1900 199

**James Colgrove** 

#### **Progress Of Experiment Science And Therapeutic Reform In The United States 1900 199:**

The Progress of Experiment Harry M. Marks,1997 How do we evaluate the safety and benefit of new drugs What tasks do we hold the government responsible for and which ones do we leave to the medical profession Harry Marks explores the origins of our contemporary system of drug regulation and the modern clinical trial He shows that the story of modern drug regulation is synonymous with the history of therapeutic reform Accompanying this history of public policy is a detailed account of changing experimental ideal and practices Marks follows the history of therapeutic experimentation from the collective investigations of the last century to the controlled clinical trial which emerged after 1950 as the paradigm of scientific experimentation The result is the first general history of clinical research in the United States a book which examines therapeutic experiments in a wide range of diseases from syphilis and pneumonia to heart disease and diabetes

**Uncertainty in Pharmacology** Adam LaCaze, Barbara Osimani, 2020-02-20 This volume covers a wide range of topics concerning methodological epistemological and regulatory ethical issues around pharmacology The book focuses in particular on the diverse sources of uncertainty the different kinds of uncertainty that there are and the diverse ways in which these uncertainties are or could be addressed Compared with the more basic sciences such as chemistry or biology pharmacology works across diverse observable levels of reality although the first step in the causal chain leading to the therapeutic outcome takes place at the biochemical level the end effect is a clinically observable result which is influenced not only by biological actions but also psychological and social phenomena Issues of causality and evidence must be treated with these specific aspects in mind In covering these issues the book opens up a common domain of investigation which intersects the deeply intertwined dimensions of pharmacological research pharmaceutical regulation and the related economic environment. The book is a collective endeavour with in depth contributions from experts in pharmacology philosophy of medicine statistics scientific methodology formal and social epistemology working in constant dialogue across disciplinary boundaries Case Studies and the Dissemination of Knowledge Joy Damousi, Birgit Lang, Katie Sutton, 2015-02-11 The case study has proved of enduring interest to all Western societies particularly in relation to questions of subjectivity and the sexed self This volume interrogates how case studies have been used by doctors lawyers psychoanalysts and writers to communicate their findings both within the specialist circles of their academic disciplines and beyond to wider publics At the same time it questions how case studies have been taken up by a range of audiences to refute and dispute academic knowledge As such this book engages with case studies as sites of interdisciplinary negotiation transnational exchange and influence exploring the effects of forces such as war migration and internationalization Case Studies and the Dissemination of Knowledge challenges the limits of disciplinary based research in the humanities The cases examined serve as a means of passage between disciplines genres and publics from law to psychoanalysis and from auto biography to modernist fiction Its chapters scrutinize the case study in order to sharpen understanding of the genre s

dynamic role in the construction and dissemination of knowledge within and across disciplinary temporal and national boundaries In doing so they position the case at the center of cultural and social understandings of the emergence of modern The Development of Scientific Marketing in the Twentieth Century Jean-Paul Gaudillière, 2015-10-06 The global pharmaceutical industry is currently estimated to be worth 1 trillion Contributors chart the rise of scientific marketing within the industry from 1920 1980 This is the first comprehensive study into pharmaceutical marketing demonstrating that many new techniques were actually developed in Europe before being exported to America **Oxford Handbook of Research Ethics** Ana S. Iltis, Douglas MacKay, 2024 The development of new pharmaceutical products and behavioral interventions aimed at improving people s health as well as research that assesses the efficacy and cost effectiveness of public policies such as policies designed to improve children's education or reduce poverty depends on research conducted with human participants It is imperative that research with human subjects is conducted in accordance with sound ethical principles and regulatory requirements Featuring 45 original essays by leading research ethicists The Oxford Handbook of Research Ethics offers a critical overview of the ethics of human subjects research within multiple disciplines and fields including biomedicine public health psychiatry sociology political science and public policy Changing Faces of Childhood Cancer Joanna Baines, 2014-12-18 This book traces the development of British answers to the problem of childhood cancer The establishment of the NHS and better training for paediatricians meant children were given access to experimental chemotherapy sending cure rates soaring Children with cancer were thrust into the spotlight as individuals stories of hope hit the headlines Manufacturing Depression Gary Greenberg, 2010-02-02 Am I depressed or just unhappy In the last two decades antidepressants have become staples of our medicine cabinets doctors now write 120 million prescriptions annually at a cost of more than 10 billion dollars At the same time depression rates have skyrocketed twenty percent of Americans are now expected to suffer from it during their lives Doctors and drug companies claim that this convergence is a public health triumph the recognition and treatment of an under diagnosed illness Gary Greenberg a practicing therapist and longtime depressive raises a more disturbing possibility that the disease has been manufactured to suit and sell the cure Greenberg draws on sources ranging from the Bible to current medical journals to show how the idea that unhappiness is an illness has been packaged and sold by brilliant scientists and shrewd marketing experts and why it has been so successful Part memoir part intellectual history part expos including a vivid chronicle of his participation in a clinical antidepressant trial Manufacturing Depression is an incisive look at an epidemic that has changed the way we have come to think of ourselves The Antibiotic Era Scott H. Podolsky, 2015-01-15 A compelling analysis of nearly seven decades of antibiotic reform framing our current efforts to stave off a post antibiotic era Winner of the CHOICE Outstanding Academic Title of the Choice ACRL In The Antibiotic Era physician historian Scott H Podolsky narrates the far reaching history of antibiotics focusing particularly on reform efforts that attempted to fundamentally change how antibiotics are

developed and prescribed This sweeping chronicle reveals the struggles faced by crusading reformers from the 1940s onward as they advocated for a rational therapeutics at the crowded intersection of bugs and drugs patients and doctors industry and medical academia and government and the media During the post World War II wonder drug revolution antibiotics were viewed as a panacea for mastering infectious disease But from the beginning critics raised concerns about irrational usage and overprescription The first generation of antibiotic reformers focused on regulating the drug industry The reforms they set in motion included the adoption of controlled clinical trials as the ultimate arbiters of therapeutic efficacy the passage of the Kefauver Harris amendments mandating proof of drug efficacy via well controlled studies and the empowering of the Food and Drug Administration to remove inefficacious drugs from the market Despite such victories no entity was empowered to rein in physicians who inappropriately prescribed or overly prescribed approved drugs Now in an era of emerging bugs and receding drugs discussions of antibiotic resistance focus on the need to develop novel antibiotics and the need for more appropriate prescription practices in the face of pharmaceutical marketing pressure from patients and the structural constraints that impede rational delivery of antibiotics worldwide Concerns about the enduring utility of antibiotics indeed about a post antibiotic era are widespread as evidenced by reports from the Centers for Disease Control and Prevention academia and popular media alike Only by understanding the historical forces that have shaped our current situation Podolsky argues can we properly understand and frame our choices moving forward State of Immunity James Colgrove, 2006-10-05 This first comprehensive history of the social and political aspects of vaccination in the United States tells the story of how vaccination became a widely accepted public health measure over the course of the twentieth century One hundred years ago just a handful of vaccines existed and only one for smallpox was widely used Today more than two dozen vaccines are in use fourteen of which are universally recommended for children State of Immunity examines the strategies that health officials have used ranging from advertising and public relations campaigns to laws requiring children to be immunized before they can attend school to gain public acceptance of vaccines Like any medical intervention vaccination carries a small risk of adverse reactions But unlike other procedures it is performed on healthy people most commonly children and has been mandated by law Vaccination thus poses unique ethical political and legal questions James Colgrove considers how individual liberty should be balanced against the need to protect the common welfare how experts should act in the face of incomplete or inconsistent scientific information and how the public should be involved in these decisions A well researched intelligent and balanced look at a timely topic this book explores these issues through a vivid historical narrative that offers new insights into the past present and future of vaccination The Routledge Handbook of the Political Economy of Science David Tyfield, Rebecca Lave, Samuel Randalls, Charles Thorpe, 2017-04-28 The political economy of research and innovation R I is one of the central issues of the early twenty first century Science and innovation are increasingly tasked with driving and reshaping a troubled global economy while also tackling multiple overlapping global

challenges such as climate change or food security global pandemics or energy security But responding to these demands is made more complicated because R I themselves are changing Today new global patterns of R I are transforming the very structures institutions and processes of science and innovation and with it their claims about desirable futures Our understanding of R I needs to change accordingly Responding to this new urgency and uncertainty this handbook presents a pioneering selection of the growing body of literature that has emerged in recent years at the intersection of science and technology studies and political economy The central task for this research has been to expose important but consequential misconceptions about the political economy of R I and to build more insightful approaches This volume therefore explores the complex interrelations between R I both in general and in specific fields and political economies across a number of key dimensions from health to environment and universities to the military The Routledge Handbook of the Political Economy of Science offers a unique collection of texts across a range of issues in this burgeoning and important field from a global selection of top scholars The handbook is essential reading for students interested in the political economy of science technology and innovation It also presents succinct and insightful summaries of the state of the art for more advanced Herbs and Roots Tamara Venit Shelton, 2019-11-26 An innovative deeply researched history of Chinese medicine scholars in America and the surprising interplay between Eastern and Western medical practice Chinese medicine has a long history in the United States with written records dating back to the American colonial period In this intricately crafted history Tamara Venit Shelton chronicles the dynamic systems of knowledge therapies and materia medica crossing between China and the United States from the eighteenth century to the present Chinese medicine she argues has played an important and often unacknowledged role in both facilitating and undermining the consolidation of medical authority among formally trained biomedical scientists in the United States Practitioners of Chinese medicine as racial embodiments of irregular medicine became useful foils for Western physicians struggling to assert their superiority of practice At the same time Chinese doctors often embraced and successfully employed Orientalist stereotypes to sell their services to non Chinese patients skeptical of modern biomedicine What results is a story of racial constructions immigration politics cross cultural medical history and the lived experiences of Asian Americans in American history **Artificial Hearts** Shellev McKellar, 2018 Fighting heart disease with machines and devices Multiple approaches to building artificial hearts technological optimism and political support in the early years Dispute and disappointment heart transplantation and total artificial heart implant cases in the 1960s Technology and risk nuclear powered artificial hearts and medical device regulation Media spotlight the Utah total artificial heart Clinical and commercial rewards ventricular assist devices Securing a place therapeutic clout and second generation VADs Artificial hearts in the 21st century A Historical Guide to the U.S. Government George Thomas Kurian, Joseph P. Harahan, 1998 This volume presents a composite profile of the U S government through a history of the various departments and agencies that constitute it Preface p vii America, History and Life, 1999

Provides historical coverage of the United States and Canada from prehistory to the present Includes information abstracted from over 2 000 journals published worldwide Bulletin of the History of Medicine, 2007 Includes the Transactions of the 15th annual meetings of the American Association of the History of Medicine 1939 The Michigan Historical Review Cancer in the Twentieth Century David Cantor, 2008-05-26 This collection of essays explores efforts to control and prevent cancer in North America and Europe On both sides of the Atlantic control programs emerged in the early twentieth century and most were focused on early detection and treatment Yet those initiatives took very different forms in different countries Experts disagreed on how to persuade the public to go to their doctors what should be the role of public education how cancer services should be delivered who should provide them which forms of therapy were most appropriate to particular cancers and where to draw the line between therapy and prevention Focusing on the United States and Britain this volume examines why these differences emerged how they shaped national programs of control and how control programs in the early twentieth century presaged and set the conditions for the emergence of prevention oriented programs in the 1960s and 1970s Featuring works by leading medical historians on subjects such as the portrayal of cancer in the movies feminist surgeons risk factors for breast cancer and the emergence of clinical trials Cancer in the Twentieth Century will engage historians of medicine and public health as well as health policy analysts medical sociologists and anthropologists and medical researchers and practitioners Komplementäre Methodenlehre der klinischen Forschung Helmut Kiene, 2013-03-07 Die komplement re Methodenlehre bietet die Basis freine wissenschaftliche Fortentwicklung der Medizin ber die Evidence based Medicine hinaus als Cognition based Medicine Dieses Buch pr sentiert was in den gro en blinden Fleck der konventionellen Methoden filt und deshalb inexistent zu sein scheint die Methoden des nicht statistischen Nachweises der therapeutischen Wirksamkeit am individuellen Patienten ebenso das dazugeh rige System klinischer Forschung Die Analyse der individuellen Therapiebeurteilung bildet die Grundlage fr eine Aufwertung des rztlichen Urteils und der rztlichen Erfahrung Die Konsequenzen sind weitreichend und gehen alle an die an der Gestaltung des Gesundheitswesens beteiligt sind rzte Therapeuten Gesundheits konomen Juristen Politiker Methodiker Statistiker

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