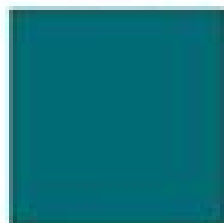
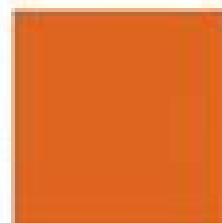
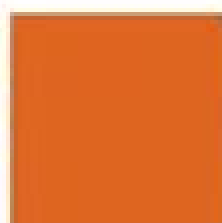
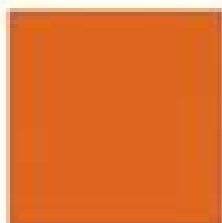
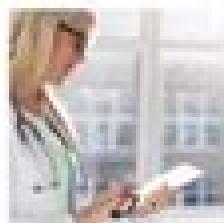
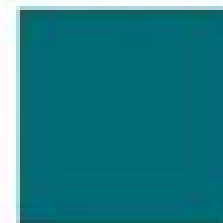


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Author: Julie Bonhomme * Date: June 2016 *



 Presentation



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**Karen M. Facey, Helle Ploug
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The Sedated Society James Davies, 2017-01-23 This edited volume provides an answer to a rising public health concern what drives the over prescription of psychiatric medication epidemic Over 15% of the UK public takes a psychiatric medication on any given day and the numbers are only set to increase Placing this figure alongside the emerging clinical and scientific data revealing their poor outcomes and the harms these medications often cause their commercial success cannot be explained by their therapeutic efficacy Chapters from an interdisciplinary team of global experts in critical psychopharmacology rigorously examine how pharmaceutical sponsorship and marketing diagnostic inflation the manipulation and burying of negative clinical trials lax medication regulation and neoliberal public health policies have all been implicated in ever rising psycho pharmaceutical consumption This volume will ignite a long overdue public debate It will be of interest to professionals in the field of mental health and researchers ranging from sociology of health to medical anthropology and the political economy of health

Guide to EU and UK Pharmaceutical Regulatory Law Pieter Erasmus, Sally Shorthose, 2025-10-30 In the European Union EU its Member States and the United Kingdom UK post Brexit as elsewhere the marketing of pharmaceuticals is subject to an ever more complex web of legislation and regulation resulting from the intense scrutiny necessary to ensure such essential products are not only efficacious but also safe This useful book lays out this system with extraordinary clarity and logic Adopting a Europe wide perspective on the law governing pharmaceuticals expert contributors from the law firm Bird stages and standards for creating a product dossier obtaining a marketing authorisation how and when an abridged marketing authorisation procedure can be used criteria for conditional marketing authorisations generic products and essential similarity paediatric use and the requisite additional trials orphan medicinal products biologicals and biosimilars homeopathic herbal and similar medicines medical devices pandemics epidemics and vaccines pharmacovigilance parallel trade advertising and relevant competition law intellectual property rights and data protection regulation Pharmaceutical lawyers and regulatory advisers both in house and in private practice will welcome this unique book It offers immeasurable value for all who need to understand the process of bringing a medicinal product or medical device to market and the continuing rights and obligations

Institutional Corruption Theory in Pharmaceutical Industry-Medicine Relationships Anna Laskai, 2020-05-06 This book discusses the influence of the pharmaceutical industry on the practice of medicine and the observed and potential pitfalls of such partnerships It argues that the pharmaceutical industry has become indispensable to many of the activities of the medical profession across the pharmaceutical product lifecycle and examines the regulatory ethical professional and institutional difficulties that arise from these interactions With data drawn from over 80 qualitative accounts from medical pharmaceutical regulatory and healthcare professionals this book uses both Hungary and the Netherlands as case studies to demonstrate the potential problem of undue pharmaceutical industry influence within the relationships fostered with the profession of medicine

Chapters systematically describe the lifecycle of a pharmaceutical product from research to distribution demonstrating the interdependency of industry and medicine Arguing that the medical profession should be a buffer between the pharmaceutical industry interests and patient interests the book explores how undue industry influence weakens the ability of the medical profession to do so Using the theory of institutional corruption the book aims to analyze how conflict of interest and the weakening of institutional imperatives is a result of institutional interactions rather than individual actions Appropriate for students and researchers of the pharmaceutical industry corporate corruption and those working in NGOs and policy making this unique volume is an comprehensive look at the complex relationship between medicine and pharmacy

Patient Involvement in Health Technology Assessment Karen M. Facey, Helle Ploug Hansen, Ann N.V. Single, 2017-05-15 This is the first book to offer a comprehensive guide to involving patients in health technology assessment HTA Defining patient involvement as patient participation in the HTA process and research into patient aspects this book includes detailed explanations of approaches to participation and research as well as case studies Patient Involvement in HTA enables researchers postgraduate students HTA professionals and experts in the HTA community to study these complementary ways of taking account of patients knowledge experiences needs and preferences Part I includes chapters discussing the ethical rationale terminology patient based evidence participation and patient input Part II sets out methodology including Qualitative Evidence Synthesis Discrete Choice Experiments Analytical Hierarchy Processes Ethnographic Fieldwork Deliberative Methods Social Media Analysis Patient Reported Outcome Measures patients as collaborative research partners and evaluation Part III contains 15 case studies setting out current activities by HTA bodies on five continents health technology developers and patient organisations Each part includes discussion chapters from leading experts in patient involvement A final chapter reflects on the need to clearly define the goals for patient involvement within the context of the HTA to identify the optimal approach With cohesive contributions from more than 80 authors from a variety of disciplines around the globe it is hoped this book will serve as a catalyst for collaboration to further develop patient involvement to improve HTA If you re not involving patients you re not doing HTA Dr Brian O Rourke President and CEO of CADTH Chair of INAHTA **Democratizing Health** the late Hans Löfgren, Evelyne de Leeuw, Michael Leahy, 2011-01-01 This book examines the important role of consumer activism in health policy in different national contexts In an age of shifting boundaries between state and civil society consumer groups are potentially drivers of democratisation in the health domain The expert contributors explore how their activities bring new dynamics to relations between service providers the medical profession government agencies and other policy actors This book is unique in comprehensively analysing the opportunities and dilemmas of this type of activism including ambiguous partnerships between consumer groups and stakeholders such as the pharmaceutical industry These themes are explored within an internationally comparative framework with case studies from various countries *Health Policy Brief* William W. Lowrance, Organisation for Economic Co-operation and Development.

Directorate for Science, Technology, and Industry, Organisation for Economic Co-operation and Development. Committee for Scientific and Technological Policy, OECD Working Party on Biotechnology, 1999 Europolitics Environment, 2006-12-08

Yearbook of International Organizations 2014-2015, Volumes 1a & 1b (Set) Union of International Associations, 2014-06-16 Volume 1 A and B covers international organizations throughout the world comprising their aims activities and events

Implementierung eines Controllingsystems zur Sicherstellung der Einhaltung der Regelungen des EFPIA-Kodex in der pharmazeutischen Industrie Franziska Protz, 2015-04-24

Jahrelang beeinflusste eine Mitarbeiterin eines Pharmaunternehmens das Verschreibungsverhalten von Vertragsärzten zugunsten des Vertriebes eigener Produkte. Ein Prämiensystem sah vor, dem Arzt 5% des Herstellerabgabepreises zu gewähren und als eine fiktive Zuwendung für wissenschaftliche Beratungsleistung auszuweisen, wenn sich dieser als Gegenleistung dazu entschied, die Produkte des Arzneimittelherstellers bevorzugt zu verordnen. Insgesamt 16 Fällen ergab sich ein Gesamtbetrag von ca. 18000 Euro, der als Vorteilsnahme durch die Ärzte angenommen wurde. Durch eine Gesetzeslücke wurde dieser Sachverhalt zwar unter dem Tatbestand der Bestechung geahndet, der Fall konnte aber für keine der Parteien unter Straftatbestand gestellt werden. Demnach machen sich weder Ärzte bei Annahme von Provisionen strafbar, noch sind Pharmaunternehmen durch die Offerte von Geldern der Bestechung schuldig. Solche und weitere Fälle unlauterer Beeinflussung erwecken in der Gesellschaft kein gutes Gefühl und erzeugen neben Misstrauen und Unsicherheit bezüglich der Unabhängigkeit ärztlicher Entscheidungen und Integrität der Pharmaunternehmen gleichfalls Forderungen nach mehr Transparenz und Offenlegung aller Zahlungsflüsse. Infolgedessen entfachte die Bekämpfung von korruptivem Verhalten im Gesundheitswesen auch in der Politik weitreichende Diskussionen, die den Gesetzgeber in der Pflicht sehen, das Strafgesetz entsprechend zu bearbeiten. Aufgrund der bestehenden Korruptionsrisiken vor allem aber getrieben durch das Urteil im obigen Fall befrachten Pharmaunternehmen nicht nur einen erheblichen Reputationsverlust, sondern auch einen nicht ausreichend rechtlichen Schutz zur Bekämpfung der Machenschaften schwarzer Schafe und zur Unterstützung des eigenen Verhaltenskodex. Um Korruptionsrisiken effektiv vorzubeugen und das Vertrauen der Öffentlichkeit zu verbessern, plädieren sie daher für ein aktives Vorgehen gegen kriminelle Handlungsweisen im Gesundheitswesen und Aufklärung der Gesellschaft. Der EFPIA Kodex (EFPIA Code on Disclosure of Transfer of Value from Pharmaceutical Companies to Healthcare Professionals and Healthcare Organisations) des europäischen Dachverbandes der nationalen Verbände forschender Pharmaunternehmen, kurz EFPIA, ist eine der untergesetzlichen Normen, die seither durch Eigeninitiative in der Ärzteschaft und der Pharmabranche vorangetrieben werden, um integres Verhalten in den Mittelpunkt zu stellen und die Verantwortung dafür zu verdeutlichen.

Reviewing **Efpia Codes Transparency International**: Unlocking the Spellbinding Force of Linguistics

In a fast-paced world fueled by information and interconnectivity, the spellbinding force of linguistics has acquired newfound prominence. Its capacity to evoke emotions, stimulate contemplation, and stimulate metamorphosis is actually astonishing. Within the pages of "**Efpia Codes Transparency International**," an enthralling opus penned by a very acclaimed wordsmith, readers attempt an immersive expedition to unravel the intricate significance of language and its indelible imprint on our lives. Throughout this assessment, we shall delve into the book's central motifs, appraise its distinctive narrative style, and gauge its overarching influence on the minds of its readers.

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