The Risks Of Prescription Drugs A Columbia Ssrc Privatization Of Risk

Getting the books **The Risks Of Prescription Drugs A Columbia Ssrc Privatization Of Risk** now is not type of challenging means. You could not deserted going afterward book hoard or library or borrowing from your associates to admittance them. This is an totally simple means to specifically acquire guide by on-line. This online pronouncement The Risks Of Prescription Drugs A Columbia Ssrc Privatization Of Risk can be one of the options to accompany you later having new time.

It will not waste your time. give a positive response me, the e-book will no question freshen you extra event to read. Just invest tiny get older to contact this on-line pronouncement **The Risks Of Prescription Drugs A Columbia Ssrc Privatization Of Risk** as well as evaluation them wherever you are now.

Sexology for the Wise Omar Zaid 2022-07-29 This essay collection applies wide-ranging optics to myths of LGBT normality. The author compares and contrasts biological, metaphysical, psychological, moral, and social dynamics that define and delimit normal heterosexual duality with elements of the gender confused. He does this in terms that illustrate spiritual and physical absolutes that are denied yet manipulated by postmodern nihilists who serve the occult governance that institutionalizes evil. The heterosexual dyad is rigorously defended as cardinal, essential, existential, naturally hegemonic, and not the least bit ambiguous. Zaid's comprehensive acumen is both frightening and captivating. His race through the Holocene irremediably shakes and changes the reader's world view via this careful amalgamation of Religion, Theology, Scripture, History, Science, Geo-Politics, Human Nature, Magick, Philosophy, and Occult Mystery Systems. Sexology For The Wise is an intense dot-connecting narrative that crosses all bounds of taboo to reveal much we do not wish to acknowledge.

Benefits, Risks and Costs of Prescription Drugs in Ontario : a Scientific Basis to Evaluate Policy Options : a Submission of the Inter-University Working Group to the Lowy Commission of Inquiry Inter-University Working Group on Prescription Drug Use (Ont.) 1988

Syringe Exchange Programs and the Opioid Epidemic Joaquin Jay Gonzalez III 2022-01-28 Syringe exchange programs and safe injection services are outside-the-box interventions increasingly being used by governments, nonprofits and citizens to address dire issues percolating in tandem with America's burgeoning opioid epidemic. People who inject drugs (PWID)--almost a million Americans annually--commonly use painkillers such as heroin and fentanyl, as well as methamphetamine, benzodiazepines, barbiturates and cocaine. Yet the users themselves are often obscured or marginalized by the bigger picture. This collection of essays covers policies and practices aimed at preventing both opioid-related deaths and related infections of hepatitis and HIV.

Issues in Clinical Psychology, Psychiatry, and Counseling: 2011 Edition 2012-01-09 Issues in Clinical Psychology, Psychiatry, and Counseling: 2011 Edition is a ScholarlyEditions™ eBook that delivers timely, authoritative, and comprehensive information about Clinical Psychology, Psychiatry, and Counseling. The editors have built Issues in Clinical Psychology, Psychiatry, and Counseling: 2011 Edition on the vast information databases of ScholarlyNews.™ You can expect the information about Clinical Psychology, Psychiatry, and Counseling in this eBook to be deeper than what you can access anywhere else, as well as consistently reliable, authoritative, informed, and relevant. The content of Issues in Clinical Psychology, Psychiatry, and Counseling: 2011 Edition has been produced by the world's leading scientists, engineers, analysts, research institutions, and companies. All of the content is from peer-reviewed sources, and all of it is written, assembled, and edited by the editors at ScholarlyEditions™ and available exclusively from us. You now have a source you can cite with authority, confidence, and credibility. More information is available at http://www.ScholarlyEditions.com/.

Safety and soundness standards in the mail order prescription industry United States. Congress. Senate. Committee on Governmental Affairs. Subcommittee on Government Efficiency, Federalism, and the District of Columbia 1987

Communicating Therapeutic Risks Louis A. Morris 2012-12-06 I guess everyone has a cousin Ernest. He is the fellow of whom your mother asks . . . "Why can't you be more like your cousin Ernest?" Cousin Ernest went to the high school for genius children and got all A's, even in French. As the years went by, I lost contact with Cousin Ernest. Then last year, at a family gathering, I met him again. Sure enough, he had gone to Harvard and become a doctor, a radiologist. We began discussing his practice and he mentioned that he performs some fairly risky diagnostic tests. While legally he was compelled to tell patients about the risks they were undertaking, he said that risk disclosure was a useless exercise. "No one has ever refused to undergo the procedure," he said. It was difficult to argue with his observation that no patient ever refused to undergo his tests. I understood that the lack of refusals did not necessarily mean that risk disclosure was a useless exercise, but his underlying argument was quite compelling.

How to Raise a Drug-Free Kid Joseph A. Califano 2014-09-09 The highly acclaimed comprehensive guide to getting your child through the formative pre-teen, teen, and college years drug-free—now completely revised and updated. Nearly every child will be offered drugs or alcohol before graduating high school, and excessive drinking is common at most colleges. But the good news is that a child who gets to age twenty-one without smoking, using illegal drugs, or abusing alcohol or prescription drugs is virtually certain never to do so. Drawing on more than two decades of research at The National Center on Addiction and Substance Abuse at Columbia University (CASAColumbia), founder Joseph A. Califano, Jr., presents a clear, common-sense guide to helping kids stay drug-free. All parents dream of a healthy, productive, and fulfilling future for their children; Califano shows which specific actions work and what parents can do to teach, protect, and empower their children to have the greatest chance of making that future come true. Teenagers who learn about the risks of drugs from their parents are twice as likely never to try them, and this book provides the tools parents need to prepare their children for those crucial decision-making moments. In this revised and updated edition, Califano tackles some of the newest obstacles standing between our kids and a drug-free life—from social media sites and cell phone apps to the explosion in prescription and over-the-counter drug abuse and the increased dangers and addictive power of marijuana. He reveals what teens can't or won't tell their parents about their thoughts on drugs and alcohol, and combines the latest research with his discussions with thousands of parents and teens about the challenges that widespread access to drugs and alcohol present, and how parents can instill in their teens the will and skills to choose not to use. Califano's insightful and lively guide is as readable as it is informative.

Effects of Prescription Drugs During Pregnancy United States. Congress. House. Committee on Science and Technology. Subcommittee on Investigations and Oversight 1982

Health at Risk Jacob S. Hacker 2008 A collection of essays dealing with the health care system.

Under the Rug 1998

The Pill Book (14th Edition) Harold M. Silverman 2011-07-20 THE CONSUMER'S GUIDE TO PILLS—COMPLETELY REVISED 14th EDITION FOR 2010 WITH MORE THAN 20 IMPORTANT NEW DRUGS AND DOZENS OF NEW BRAND NAMES For more than three decades, millions of consumers have trusted The Pill Book to provide official, FDA-approved information on more than 1,800 of the most commonly prescribed drugs in the United States with guidelines from leading pharmacists. Each drug is profiled in a concise, readable, easy-to-understand entry, making The Pill Book the perfect reference when you have questions about the medications your doctor prescribes. Inside you'll discover • generic and brand-name listings that can help you save money • What each drug is for, and how it works • usual dosages, and what to do if a dose is skipped • side effects and possible adverse reactions, highlighted for quick reference • interactions with other drugs and food • overdose and addiction potential • alcohol-free and sugar-free medications • the most popular self-injected medications and their safe handling • information for seniors, pregnant and breast-feeding women, children, and others with special needs • cautions and warnings, and when to call your doctor • 32 pages of actual-size color photographs of prescription pills* No home should be without this book! *Not all ereading devices will show the images in color and at the exact size.

Unnatural Selection Mark Roeder 2014-10-14 Unnatural Selection is the first book to examine the rise of the "technocentric being"—or geek—who personifies a distinct new phase in human evolution. People considered geeks often have behavioral or genetic traits that were previously considered detrimental. But the new environment of the Anthropocene period—the Age of Man—has created a kind of digital greenhouse that actually favors their traits, enabling many non-neurotypical people to bloom. They resonate with the technological Zeitgeist in a way that turns their weaknesses into strengths. Think of Mark Zuckerberg versus the towering, Olympics-bound Winklevoss twins in the movie Social Network. Roeder suggests that the rise of the geek is not so much the product of Darwinian "natural selection" as of man-made—or unnatural—selection. He explains why geeks have become so phenomenally successful in such a short time and why the process will further accelerate, driven by breakthroughs in genetic engineering, neuropharmacology, and artificial intelligence. His book offers a fascinating synthesis of the latest trends in these fields and predicts a twenty-first century "cognitive arms race" in which new technology will enable everyone to become more intelligent and "geek-like."

Departments of Labor, Health and Human Services, Education, and Related Agencies Appropriations for 2008: Dept. of Education FY 2008 budget justifications United States. Congress. House. Committee on Appropriations. Subcommittee on the Departments of Labor, Health and Human Services, Education, and Related Agencies 2007

The Pill Book Harold M. Silverman 2002 Based on information from the Food and Drug Administration, an updated consumer's guide offers up-to-date profiles of more than 1,500 of the most commonly prescribed drugs in America, including generic and brand names, usual dosages, and side effects, as well as color photographs. Original

Benefits, Risks and Costs of Prescription Drugs in Ontario 1989*

Pharmaceutical and Medical Device Safety Sonia Macleod 2019-02-21 This book examines how regulatory and liability mechanisms have impacted upon product safety decisions in the pharmaceutical and medical devices sectors in Europe, the USA and beyond since the 1950s. Thirty-five case studies illustrate the interplay between the regulatory regimes and litigation. Observations from medical practice have been the overwhelming means of identifying post-marketing safety issues. Drug and device safety decisions have increasingly been taken by public regulators and companies within the framework of the comprehensive regulatory structure that has developed since the 1960s. In general, product liability cases have not identified or defined safety issues, and function merely as compensation mechanisms. This is unsurprising as the thresholds for these two systems differ considerably; regulatory action can be triggered by the possibility that a product might be harmful, whereas establishing liability in litigation requires proving that the product was actually harmful. As litigation normally post-dates regulatory implementation, the 'private enforcement' of public law has generally not occurred in these sectors. This has profound implications for the design of sectoral regulatory and liability regimes, including associated features such as extended liability law, class actions and contingency fees. This book forms a major contribution to the academic debate on the comparative utility of regulatory and liability systems, on public versus private enforcement, and on mechanisms of behaviour control.

H.R. 4489, the FEHBP Prescription Drug Integrity, Transparency, and Cost Savings Act United States. Congress. House. Committee on Oversight and Government Reform. Subcommittee on Federal Workforce, Postal Service, and the District of Columbia 2010

Code of Federal Regulations 2013 Special edition of the Federal Register, containing a codification of documents of general applicability and future effect ... with ancillaries.

Public Health Reports 1978

2017 CFR Annual Print Title 42 Public Health Parts 414 to 429 Office of The Federal Register 2017-07-01

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

Social, Political and Cultural Dimensions of Health Kevin Dew 2016-05-09 This book comprehensively explores social, political and cultural dimensions of health in contemporary society. It addresses many issues and pertinent questions, including the following: Are we over diagnosed and over medicated? How can patients participate in their own care? Do pharmaceutical companies coerce us into medication regimes? What drives inequalities in health outcomes? What is the experience of health care for indigenous communities? Why do different countries have such different health care systems? How do we respond to life-changing conditions? Can we achieve a 'good death'? How do new genetics shape our identities? Is public health a force of liberation or disempowerment? The book incorporates the range of levels of influence on health, covering individual patient experiences, the health professions, multinational corporations, the state, global organisations as well as examining trends in social organisation, cultural expression and technological developments. It volume provides an accessible, yet in-depth, overview and discussion of the sociology of health. The chapters include an illustrative case study and further readings relating to the topic.

HSMHA Health Reports United States. Health Services and Mental Health Administration 1978

Silent Cells Anthony Ryan Hatch 2019-04-30 A critical investigation into the use of psychotropic drugs to pacify and control inmates and other captives in the vast U.S. prison, military, and welfare systems For at least four decades, U.S. prisons and jails have aggressively turned to psychotropic drugs—antidepressants, antipsychotics, sedatives, and tranquilizers—to silence inmates, whether or not they have been diagnosed with mental illnesses. In Silent Cells, Anthony Ryan Hatch demonstrates that the pervasive use of psychotropic drugs has not only defined and enabled mass incarceration but has also become central to other forms of captivity, including foster homes, military and immigrant detention centers, and nursing homes. Silent Cells shows how, in shockingly large numbers, federal, state, and local governments and government-authorized private agencies pacify people with drugs, uncovering patterns of institutional violence that threaten basic human and civil rights. Drawing on publicly available records, Hatch unearths the coercive ways that psychotropics serve to manufacture compliance and docility, practices hidden behind layers of state secrecy, medical complicity, and corporate profiteering. Psychotropics, Hatch shows, are integral to "technocorrectional" policies devised to minimize public costs and increase the private profitability of mass captivity while guaranteeing public safety and national security. This broad indictment of psychotropics is therefore animated by a radical counterfactual question: would incarceration on the scale practiced in the United States even be possible without psychotropics?

Departments of Labor, Health and Human Services, Education, and Related Agencies Appropriations for 2008 United States. Congress. House. Committee on Appropriations. Subcommittee on the Departments of Labor, Health and Human Services, Education, and Related Agencies 2007

Prescription Drugs United States. General Accounting Office 1994

Effect Modification by Socioeconomic Conditions on the Effects of Prescription Opioid Supply on Drug Poisoning Deaths in the United States David S. Fink 2020 As federal and state policies continue to target the rising rates of fatal drug poisonings, these findings show that area-level socioeconomic conditions may represent an important target for policy intervention during the current drug poisoning crisis and a critical piece of information necessary for predicting any future drug-related crises.

Drug Safety Nigel S. B. Rawson 2016-11-08 With "Big Pharma" garnering an increasing number of negative headlines due to reports of adverse drug reactions and a surge in prescription drug addiction and overdose deaths, many people are increasingly skeptical about the safety of modern pharmaceutics and the moral integrity of the pharmaceutical industry. This book was written to provide a balanced perspective on drug safety risks. No therapeutic prescription drug is entirely risk-free. Before receiving marketing approval, new drugs go through arduous and expensive testing processes that can take up to a decade and cost over two billion dollars. While not perfect, the process is far from a "Wild West" environment where big pharmaceutical companies ride roughshod over government regulators. However, author and pharmacoepidemiologist Nigel Rawson argues, the antipathy that is common between governments, pharmaceutical industry and academic experts in Canada needs to change to an environment of collaboration and partnership to enhance our ability to respond in a timely fashion to future pharmaceutical crises. While directed mainly at students in the health sciences and pharmaceutical professionals, this book will be of interest to anyone, including lay people and policy makers, who would like to know more about the evolution of the prescription drug evaluation and risk assessment process. Although the book focuses primarily on Canada, it makes comparisons with the United States and Europe, and several of the author's recommendations for how to improve the prescription drug evaluation process are applicable worldwide....

Law and the Regulation of Medicines Emily Jackson 2012-03-01 The principal purpose of this book is to tell the story of a medicine's journey through the regulatory system in the UK, from defining what counts as a medicine, through clinical trials, licensing, pharmacovigilance, marketing and funding. The question of global access to medicines is addressed because of its political importance, and because it offers a particularly stark illustration of the consequences of classifying medicines as a private rather than a public good. Two further specific challenges to the future of medicine's regulation are examined separately: first, pharmacogenetics, or the genetic targeting of medicines to subgroups of patients, and second, the possibility of using medicines to enhance well-being or performance, rather than treat disease. Throughout, the emphasis is on the role of regulation in shaping and influencing the operation of the medicines industry, an issue that is of central importance to the promotion of public health and the fair and equitable distribution of healthcare resources.

Pharmaceutical R & D 1993-01-31 Analyzes the costs, risks, and economic rewards of pharmaceutical R&D and the impact of public policy on both costs and returns. Examines the rapid increase in pharmaceutical R&D that began in the 1980s in the light of trends in science, technology, drug discovery, and health insurance coverage; Government regulation; product liability; market competition; Federal tax policy; and Federal support of prescription drug research. 12 appendices, including a glossary of terms.

Hidden Addictions Marilyn Freimuth 2009-03-19 Media portrayals and diagnostic criteria convey an image of an addicted person as someone whose deficient coping skills and severely compromised functioning are readily apparent. Yet addictions remain some of the most frequently missed diagnoses in health and mental health care settings. This occurs, in large part, because most people with addictions do not fit the stereotype. In the context of psychotherapy, the typical patient with an addiction will present depression, anxiety, marital problems or a general sense that life is not working. This book addresses how addictions can be recognized more often and accurately assessed in the context of psychotherapy. Along with learning about the standard assessment instruments, the reader is introduced to methods for asking the appropriate questions and listening to the clinical dialogue for signs of a undisclosed addiction. This book provides a great deal of knowledge about addictions and their assessment in a way that is relevant to clinical practice.

The Pill Book Guide to Safe Drug Use Harold M. Silverman 1989 Describes the drugs most frequently prescribed for the elderly, discussing the often subtle warning signs that signal dangerous over medication for common anti-arthritis, anti-depressant, and other drugs

The Risks of Prescription Drugs Donald Light 2010 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 or risk borne by individuals in the world of pharmaceuticals. "Although many are aware that pharmaceutical industry lobbyists influence policy decisions, few know the full consequences. This book is enlightening." Jill Quadagno, author of The Transformation of Old Age Security "Clear, concise, and unflinching, this book provides consumers with tools for self-defense and concerned citizens with a road map for rebalancing American medicine." John Abramson, author of Overdosed America: The Broken Promise of American Medicine "This book introduces important debates on pharmaceutical promotion and marketing, needed drug evaluation and regulation, professional conflicts of interest, and increased medicalization of behavior. It explores important trends and policy questions that all engaged citizens should consider." David Mechanic, author of The Truth About Health Care: Why Reform is not Working in America

Improving Drug Safety — A Joint Responsibility Rolf Dinkel 2013-03-07 As the focus on pharmaceuticals has broadened from concern for their cost and effectiveness to their real and potential risks and benefits, a critical question has been raised: whose responsibility is it to improve drug safety? In April 1990, this question became the theme for a conference at Wolfsberg, Switzerland, near the shores of Lake Constance. Called an "international dialogue conference" by its organizers, the meeting brought together leaders from the pharmaceutical industry, regulatory authorities, academia, medicine, consumer organizations and the media. Opening addresses were given by representatives of the Council for International Organizations of Medical Sciences (CIOMS), the International Federation of Pharmaceutical Manufacturers Associations (IFPMA), the Swiss International Pharmaceutical Agency, and the RAD-AR Consortium. This book documents the papers presented and discussions held at this conference, which took the topic of risks and benefits of drug therapy one step further to responsibility. It includes a rich menu of issues for those who care about the evaluation of drug therapy, the ethics behind it, the expectations of the patient, and the role of traditional and nontraditional drug safety communica tions. The ideas expressed here come from different parts of the world but relate to common drug safety problems, observations, and scientific assessments; they provide insights into innovative approaches, cautious changes, and desired actions. The papers in this volume are broadly divided into conceptual perspectives (ethics, how the knowledge about drug risks and benefits is generated and appraised, the expectations in drug safety) and operational perspectives (communication, discussion, and action).

Good Pharma Donald W. Light 2015-06-30 Drawing on key concepts in sociology and management, this history describes a remarkable institute that has elevated medical research and worked out solutions to the troubling practices of commercial pharmaceutical research. Good Pharma is the answer to Goldacre's Bad Pharma: ethical research without commercial distortions.

Overcoming Prescription Drug Addiction Rod Colvin 2008-06-01 This newly revised third edition delves into the most widely abused narcotic in the U.S.—prescription drugs. The book offers help to those suffering from this type of addiction as well as their families. The topics discussed include dynamics of addiction and the newest treatment options, who is at risk for addiction, why more teens are abusing prescription drugs, the symptoms of withdrawal, and methods of intervention for family members. Personal stories from addicts who describe their journeys into recovery are also included.

Shared Responsibility, Shared Risk Jacob Hacker 2012-01-19 How can the American social welfare system be repaired so that workers and families receive adequate protection and, if necessary, provision from the ravages of the market? This book addresses this fundamental problem and analyses how the 'privatization of risk' has increased hardships for American families and increased inequality. It also proposes a series of solutions that would distribute the burdens of risks more broadly and expand the social safety net.

Treating Heroin Addiction in Norway Aleksandra Bartoszko 2021-07-06 Focusing on the world of Norwegian Opioid Substitution Treatment (OST) in the aftermath of significant reforms, this book casts a critical light on the intersections between medicine and law, and the ideologies infusing the notions of "individual choice" and "patient involvement" in the field of addiction globally. With ethnographic attention to the encounters between patients, clinicians, and bureaucrats, the volume shows that OST sustains the realities it is meant to address. The chapters follow one particular patient through complex clinical and legal battles as they fight to achieve a better quality of life. The study provides ethnographic insight that captures the individual, experiential aspects of addiction treatment, and how these experiences find a register within different domains of treatment and policy, including the familial, social, legal, and clinical. Offering a rare view of addiction treatment in a Scandinavian welfare state, this book will be of interest to scholars of medical and legal anthropology and sociology, and others with an interest in drug policy and addiction treatment.

Women Under the Influence 2006 This comprehensive and accessible book documents the physical and emotional effects of substance abuse in girls and women, explores the role of the advertising and entertainment industries in popularizing various substances of abuse, and discusses the way America responds to this enormous health problem. Covering a broad range of substances—nicotine, alcohol, prescription and illicit drugs—the book addresses the unique reasons that girls and women smoke and abuse alcohol and drugs. It provides the most current information about the use of prescription and club drugs, key warning signs of addiction, and options for prevention and treatment. The book includes historical anecdotes and testimonies from recovering women. Incorporating more than a decade of extensive research, Women under the Influence will help women, health care professionals, educators, and policy makers understand the scope of substance abuse in girls and women, the urgency of responding to the problem, the key points of intervention, and potential roads to recovery.

Foute farma Ben Goldacre 2013-10-23 Medicijnfabrikanten voeren structureel slechte experimenten uit op hun medicijnen, verhullen nadelige uitkomsten en verdraaien goede uitkomsten. Schadelijke praktijken, die ieder jaar levens van patiënten kosten. Arts en schrijver Ben Goldacre licht de farmaceutische industrie door, een branche waar miljoenen in omgaan en waar wij allemaal mee te maken hebben.