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Overview : Legal Aspects of Regulatory Treatment of Banks ...
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This book discusses legal aspects of the regulatory treatment of banks 1 in distress. Banks in distress should be distinguished from banks that are merely weak. "Banks in distress" are defined as

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banks that are not in compliance with prudential banking law.

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Legal Aspects of Regulatory Treatment of Banks in Distress
Release on 2001-04-18 | by T. M. C. Asser This book is based on a comparative study of the law of selected countries with banking legislation that may serve as standard for the treatment of banks in distress.

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1. Compliance. Compliance to new laws and regulation can be a significant expense that may include the cost of changing products, processes and legal structures. The burden of compliance can be greater for small businesses with limited resources. In some cases, compliance can also be a significant expense for large multinational companies that must comply with laws in a number of countries that are perpetually in flux.

What is Regulatory Risk? -

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Legal Aspects Of Regulatory Treatment Of Banks In Distress Get this from a library! Legal aspects of regulatory treatment of banks in distress. [T M C Asser; International Monetary Fund.] -- This book is based on a comparative study of the law of selected countries with banking

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Treatment Of Banks In Distress
Legislation that may serve as a standard for the treatment of banks in distress.

This book analyzes and compares the laws of selected industrial countries that are representative of the different approaches to the treatment of banks in distress. It addresses only those banking and economic policy issues that are required for a proper understanding of the banking law or the legal strategies, procedures, and practices that have evolved in the treatment of banking problems. The book does not cover international aspects of bank insolvency, but rather has a domestic focus, given that bank

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Treatment and supervision are still largely a national endeavor.

“ Patenting Nanomedicines: Legal Aspects, Intellectual Property and Grant Opportunities ” focusses on the fundamental aspects of Patenting Nanomedicines applied in different “ Drug Delivery and Targeting Systems ” . The promoters of new findings in this field of research are numerous and spread worldwide; therefore, managing intellectual property portfolios, and the acquisition and exploitation of new knowledge face several contingency factors. Today, the scientific community is

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discussing issues of economic outcomes in the field of Nanomedicines. Major concerns include questions as to whether the research groups, academics, industry and other stakeholders should work in unison or independently, if innovation or adaptation of new technology should be prioritized, public versus private research funding, and safeguarding versus sharing knowledge. However, despite its increasing importance for humankind, it is a matter of concern as to whether technological development can really be stimulated by patent protection. An intellectual property strategy should aim to develop a qualitative patent portfolio for continuous learning. This book

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addresses questions of ethics, socio-political policies and regulatory aspects of novel Nanomedicine-based products which are currently under development for the diagnosis and treatment of different types of diseases. It is divided in two parts – Part I is composed of the first 3 chapters, which focus on the “ fundamentals ” of legal aspects, emerging threats, advantages and disadvantages of patenting Nanomedicines, whereas Part II collects 12 chapters discussing different types of Nanomedicine-based products, their potential marketing aspects and patent protection. Whenever applied, each chapter offers a list of patents, based on a specific application in drug delivery and targeting. An

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outstanding team of 53 authors have contributed to this book, which will be of interest to professionals from the field of patent examiners, academics, researchers and scientists, students and other practitioners.

For nearly three decades, methadone hydrochloride has been the primary means of treating opiate addiction. Today, about 115,000 people receive such treatment, and thousands more have benefited from it in the past. Even though methadone's effectiveness has been well established, its use remains controversial, a fact reflected by the extensive regulation of its manufacturing, labeling, distribution, and use. The Food

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and Drug Administration regulates the safety and effectiveness of methadone, as it does for all drugs, and the Drug Enforcement Administration regulates it as a controlled substance. However, methadone is also subjected to a unique additional tier of regulation that prescribes how and under what circumstances it may be used to treat opiate addiction. Federal Regulation of Methadone Treatment examines current Department of Health and Human Services standards for narcotic addiction treatment and the regulation of methadone treatment programs pursuant to those standards. The book includes an evaluation of the effect of federal regulations on the provision of methadone treatment services and

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an exploration of options for modifying the regulations to allow optimal clinical practice. The volume also includes an assessment of alternatives to the existing regulations.

Incisive, authoritative and thoughtful, this important and timely collection of papers exploring the unresolved issues left by the recent global financial turmoil, will undoubtedly shape the policy responses to come.

Interdisciplinary in approach and wide-ranging in jurisdictional scope, it draws together influential commentators, practitioners and regulators, to create a new milestone in the search for the fundamentals of a more stable global financial system.? - Eva

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Tomnicka, King's College London, UK ? This book contains a large number of chapters, nearly 30 in all, by acknowledged experts on various aspects of the recent financial crisis. Whichever aspect of this crisis that may interest you, such as bank taxes, deposit insurance, TBTF and how to respond, cross-border issues, and many, many others, you will find chapters that are both authoritative and stimulating in this collection. The editors are to be congratulated not only in their selection of authors but also in the speed with which they have taken them from conference presentation to book chapter.?

- Charles Goodhart, London School of Economics, UK

Managing Risk in the Financial System makes

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Important and timely contributions to our knowledge and understanding of banking law, financial institution restructuring and related considerations, through the production of an innovative, international and interdisciplinary set of contributions which link law and policy issues surrounding systemic risk and crisis management. The recent financial crisis has exposed both the banking industry and financial system safety net players in many countries to a considerable level of distress as well as economic and reputational damage. These circumstances have heightened the need for policymakers to consider remedial measures under a broad umbrella that encompass inter alia prompt corrective actions, early

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closure of distressed entities, deposit insurance, bail-outs, state-aid, bank resolution and restructuring techniques. These essays provide an important contribution to research in this area, at a crucial time in the debate around the future financial industry. Contributors

“ Patenting Nanomedicines: Legal Aspects, Intellectual Property and Grant Opportunities ” focusses on the fundamental aspects of Patenting Nanomedicines applied in different “ Drug Delivery and Targeting Systems ” . The promoters of new findings in this field of research are numerous and spread worldwide; therefore, managing intellectual property portfolios, and the acquisition and

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exploitation of new knowledge face several contingency factors.

Today, the scientific community is discussing issues of economic outcomes in the field of Nanomedicines. Major concerns include questions as to whether the research groups, academics, industry and other stakeholders should work in unison or independently, if innovation or adaptation of new technology should be prioritized, public versus private research funding, and safeguarding versus sharing knowledge. However, despite its increasing importance for humankind, it is a matter of concern as to whether technological development can really be stimulated by patent protection. An intellectual property

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Strategy should aim to develop a qualitative patent portfolio for continuous learning. This book addresses questions of ethics, socio-political policies and regulatory aspects of novel Nanomedicine-based products which are currently under development for the diagnosis and treatment of different types of diseases. It is divided in two parts – Part I is composed of the first 3 chapters, which focus on the “ fundamentals ” of legal aspects, emerging threats, advantages and disadvantages of patenting Nanomedicines, whereas Part II collects 12 chapters discussing different types of Nanomedicine-based products, their potential marketing aspects and patent protection. Whenever applied, each

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chapter offers a list of patents, based on a specific application in drug delivery and targeting. An outstanding team of 53 authors have contributed to this book, which will be of interest to professionals from the field of patent examiners, academics, researchers and scientists, students and other practitioners.

This is an invaluable resource for lawyers, policy makers, and other practitioners with an interest in countries' responses to HIV/AIDS. Legal Aspects of HIV/AIDS: A Guide for Policy and Law Reform covers 65 wide-ranging topics in a concise, accessible format, explaining how laws and regulations can either underpin or undermine public health programs

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and responsible personal behavior. For each topic, the Guide summarizes the key legal or policy issues, provides relevant "practice examples" (citing actual laws and regulations), and offers a selective list of references that may be consulted for more information. Laws relating to many areas of our lives - from intimate physical conduct to international travel - can contribute to stigma, discrimination, and exclusion or, contrariwise, can help remedy these inequities. In order to create a supportive legal framework for responding to HIV/AIDS, it is important that governments effectively address gaps and other problematic aspects in their legislation and regulatory systems. This book, written by a team of

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Leading legal experts, helps them do so.

In recent years, there has been much debate over whether marijuana, an illegal drug, can provide patients with a level of therapeutic relief comparable to existing pharmaceutical treatments. While this idea is hardly new, it is advanced by some proponents as deserving more scientific inquiry. Advocates for the medical use of marijuana contend that there is already sufficient scientific evidence to justify rescheduling marijuana under the Controlled Substances Act, a change that would give it the necessary legal recognition to be used for medicinal purposes. This has already occurred in the

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case of dronabinol, the synthetic form of the main psychoactive ingredient in marijuana, which has been available as an oral prescription drug since 1986 under its brand name Marinol. To address these viewpoints, several comprehensive studies were done in the late 1990s to evaluate medicinal claims made for smoked marijuana and determine whether they are supported by convincing scientific evidence. The medical marijuana debate gained attention at the state level in 1996, when voters in California and Arizona approved ballot initiatives allowing doctors to prescribe the drug for therapeutic uses. In 1998, similar propositions were adopted in Alaska, Nevada, Oregon, and Washington, and reaffirmed in

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Arizona. Voters in Maine adopted a medical marijuana initiative in 1999. In 2000, medical marijuana was approved by voters in Colorado, reconfirmed in Nevada, and passed by the legislature in Hawaii. Federal health officials assert that these initiatives are part of a strategy to soften the nation's drug laws, and that public policy would be better served if science, rather than the ballot box, were used to judge the drug's utility. This book assesses the current issues and examines the controversies regarding the marijuana legalization issue.

Food Law and Policy surveys the elements of modern food law. It broadens the coverage of traditional food and drug law topics

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of safety, marketing, and nutrition, and includes law governing environment, international trade, and other legal aspects of the modern food system. The result is the first casebook that provides a comprehensive treatment of food law as a unique discipline. Key Features: Draws together cases with other regulatory materials such as rulemaking documents and agency requests for proposals for grant funding. Focuses on federal law and includes discussion of innovations in food law happening at the municipal, state and federal level. Covers the latest developments in food law.

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