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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.

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## **What is Regulatory Risk? - Simplicable**

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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 regulation and supervision are still largely a national endeavor.

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The Medical-Legal Aspects of Acute Care Medicine: A Resource for Clinicians, Administrators, and Risk Managers is a comprehensive resource intended to provide a state-of-the-art overview of complex ethical, regulatory, and legal issues of importance to clinical healthcare professionals in the area of acute care medicine; including, for example, physicians, advanced practice providers, nurses, pharmacists, social workers, and care managers. In addition, this book also covers key legal and regulatory issues relevant to non-clinicians, such as hospital and practice administrators; department heads, educators, and risk managers. This text reviews traditional and emerging areas of ethical and legal controversies in healthcare such as

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resuscitation; mass-casualty event response and triage; patient autonomy and shared decision-making; medical research and teaching; ethical and legal issues in the care of the mental health patient; and, medical record documentation and confidentiality. Furthermore, this volume includes chapters dedicated to critically important topics, such as team leadership, the team model of clinical care, drug and device regulation, professional negligence, clinical education, the law of corporations, telemedicine and e-health, medical errors and the culture of safety, regulatory compliance, the regulation of clinical laboratories, the law of insurance, and a practical overview of claims management and billing. Authored by experts in the field, *The Medical-Legal Aspects of Acute Care Medicine: A*

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## Resource for Clinicians,

Administrators, and Risk Managers is a valuable resource for all clinical and non-clinical healthcare professionals.

When the end of life makes its inevitable appearance, people should be able to expect reliable, humane, and effective caregiving. Yet too many dying people suffer unnecessarily. While an "overtreated" dying is feared, untreated pain or emotional abandonment are equally frightening. Approaching Death reflects a wide-ranging effort to understand what we know about care at the end of life, what we have yet to learn, and what we know but do not adequately apply. It seeks to build understanding of what constitutes good care for the dying and

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**Offers recommendations to decisionmakers that address specific barriers to achieving good care. This volume offers a profile of when, where, and how Americans die. It examines the dimensions of caring at the end of life: Determining diagnosis and prognosis and communicating these to patient and family. Establishing clinical and personal goals. Matching physical, psychological, spiritual, and practical care strategies to the patient's values and circumstances. Approaching Death considers the dying experience in hospitals, nursing homes, and other settings and the role of interdisciplinary teams and managed care. It offers perspectives on quality measurement and improvement, the role of practice guidelines, cost concerns, and legal issues such as assisted suicide. The book proposes**



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how health professionals can become better prepared to care well for those who are dying and to understand that these are not patients for whom "nothing can be done."

While subordinated debt instruments are widely used in the financial sector, their legal aspects are far from clear, particularly if the instrument is to qualify as a quasi-capital instrument for regulatory capital purposes. Since regulatory capital treatment is one of the key objectives of the instrument, it is very important to understand the distinctive features of the instruments that distinguish this instrument. This article discusses the development of case law on subordinated instruments.

For nearly three decades, methadone hydrochloride has been the primary

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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 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

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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 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.

A wide variety of legal issues surround caring for older individuals. Health and human service practitioners need to plan, provide and evaluate geriatric care, while also understanding public policies. Legal knowledge is an essential part of caring for the elderly. Students and professionals must be

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able to deliver appropriate care while also being aware of any legal, ethical and political issues that may arise. Legal Aspects of Elder Care provides a clear overview of geriatric policies and laws, enabling the reader to use informed decision-making with older clients.

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

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change that would give it the necessary legal recognition to be used for medicinal purposes. This has already occurred in the 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,

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Nevada, Oregon, and Washington, and reaffirmed in 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.

“Patenting Nanomedicines: Legal Aspects, Intellectual Property and Grant Opportunities” focusses on the fundamental aspects of Patenting

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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 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

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**Durable Diseases** 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 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



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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 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.

As the volume of transactions in European financial markets continues to grow, the use of financial collateral, be it in the form of cash, shares, bonds or credit claims, has become a critical tool in supporting and managing risk in financial transactions. This book is the first of its kind to offer a systematic examination of the whole law relating

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to financial collateral. It does so in two parts. First, it explains the law created by the Financial Collateral Arrangements (No 2) Regulations 2003, the Directive it implemented and related legislation. Second, it examines how financial collateral is used in practice in a range of different markets. It will be an essential reference point for all legal practitioners operating in financial markets. Key features:

- Analytical rigour combined with insight into how financial collateral works in practice, covering both English and Scots law
- Valuable discussion of control and possession tests, right of use, remedy of appropriation, close-out netting and impact of 'bail-in'
- Explains use of financial collateral in the derivatives market, clearing houses, direct and indirect securities holding systems and

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