PUBLICATIONS **CATALOG**

2013 | 2014

AHLA American Health Lawyers Association

AHLA

Leading health law to excellence through education, information, and dialogue, the **American Health Lawyers Association** (AHLA) is the nation's largest nonpartisan, 501(c)(3) educational organization devoted to legal issues in the healthcare field. AHLA provides resources to address the issues facing its active members who practice in law firms, government, in-house settings, and academia and who represent the entire spectrum of the health industry: physicians, hospitals and health systems, health maintenance organizations, health insurers, managed care companies, nursing facilities, home care providers, and consumers.

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AHLA's Guide to Healthcare Legal Forms, Agreements, and Policies

Second Edition with CD-ROM

$369 • **Members**
ISBN 9780769860749

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ISBN 9780769861005

Pub. #27635, loose-leaf with CD-ROM, 1,900 pages, © 2012

Collected from expert health law attorneys and members of AHLA, this resource is designed to meet the needs of health law attorneys who must regularly create such documents for their clients. It is conveniently organized by topic in a loose-leaf format for easy updating.

The Second Edition of this best-selling forms set includes more than 300 new forms, contracts, agreements, checklists, and other legal documents. For even further utility you'll also have access to form completion tips, pointers, and other references. Here is a sampling of some of the areas for which you'll find forms and practice aids for completion:

- Accountable Care Organizations
- Alternative Dispute Resolution
- Business Transactions
- Clinical Trials and Research
- Conflicts of Interest
- Corporate Compliance Programs
- Facility Operations
- Fraud and Abuse
- Governance
- Health Information Exchanges
- Health Information Technology
- Health Plan Contracting
- Internal Investigations
- Labor and Employment
- Legal Services
- Long Term Care
- Medical Group Practices
- Patient Care Policies
- Patient Safety and Adverse Outcomes
- Physician Employment
- Reimbursement
- Risk Management

The *Guide* comes complete with a companion CD-ROM which contains all the documents in an electronic .rtf format.

Antitrust and Healthcare: A Comprehensive Guide

First Edition

Christine L. White, Saralisa C. Brau, and David Marx Jr., authors and editors; David A. Argue, Martin Bienstock, Robert S. Canterman, David Narrow, Joshua H. Soven, and Shoshana Speiser, contributing authors

$174 • **Members**
ISBN 9780769865560
eISBN 9780327185956

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ISBN 9780769865577
eISBN 9780327185963

Pub. #28252, hardbound with CD-ROM, 350 pages, © 2013

If you represent clients in the healthcare arena, or are a professional with business interests in this area, *Antitrust and Healthcare: A Comprehensive Guide* is a must-have resource for informing your advice, understanding the inherent risks, opportunities and alternative strategies for effective transactions, and evaluating the antitrust issues associated with marketplace conduct. The accessible *Guide* covers:

- Substantive antitrust law
- Important case-law developments
- Formal and informal guidance issued by federal and state enforcement agencies

Consult *Antitrust and Healthcare* for guidance on any of the business activities your clients or organization are likely to be involved with:

- Mergers, acquisitions, and other transactions
- Joint ventures, provider networks, and other collaborative arrangements—including clinically integrated networks and ACOs
- Pre-merger notification and transaction planning
- Exclusive contracting
- Medical staff membership, clinical privileges, and peer review activities
- Trade association and group purchasing activities
- Dominant firm conduct

Additionally *Antitrust and Healthcare* provides invaluable "practice pointers" to help minimize antitrust risk and more successfully plan and execute business and litigation strategies. The expert authors draw on their significant government enforcement and private sector counseling and litigation experience to provide practical insights for:

- Pre-merger notification and transaction planning
- Developing antitrust compliance and "sensitivity training" programs
- Identifying conduct and language that could create antitrust "red flags"
- The creation, distribution, and use of emails, electronic documents, and other materials
- Antitrust safety zones, defenses, and immunities

Antitrust and Healthcare explains the application of antitrust principles to the different segments of the healthcare industry—including providers and third party payors—and the specific issues they confront.

(See also *Healthcare Antitrust FAQ Handbook*, p. 5.)

$169 • **Members**
ISBN 9780769861166
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eISBN 9780327175261

Pub. #27997, softbound with CD-ROM, 800+ pages, © 2013

Enterprise Risk Management Handbook for Healthcare Entities

Second Edition with CD-ROM

Roberta L. Carroll, Editor in Chief; Peggy Nakamura and Rachel V. Rose, Editors; Jessica K. Bae, Ellen Barron, Ellen L. Barton, India K. Brim, Patchin C. Curtis, John R. Evancho, Mark Faccenda, Amanda J. Flanagan, Phyllis F. Granade, Steven O. Grubbs, Daniel G. Hale, Peter J. Hoffman, Mark A. Kadzielski, Christopher N. Kanagawa, Jee-Young Kim, Marilyn Lamar, Eileen Lampe, R.Jeffrey Layne, Dana B. Mehlman, Elizabeth M. Mills, Jennifer C. Monroe, Joshua Moore, Cheryl Camin Murray, Deborah Martin Norcross, Amy B. Norris, Gisele A. Norris, Richard S. Porter, Yvonne K. Puig, Steven M. Puiszis, Emily Rhinehart, Sheila Hagg-Rickert, Fay A. Rozovsky, Joshua I. Rozovsky, Mary S. Schaefer, and Kathryn K. Wire, authors

The Second Edition of this popular *Handbook* addresses the need for and implementation of a comprehensive risk management process that encompasses the entire enterprise and crosses departmental barriers. Coverage begins with an overview of enterprise risk management (ERM) and its evolution, and goes on to address the structuring of an ERM system, as well as risk financing methods.

The authors delineate how to manage risk in a variety of settings, including:

- Contract management
- Claims management
- Environmental compliance
- Human research
- Peer review and credentialing
- Due diligence in business transactions
- Consent to treatment
- And numerous others

The publication also includes insight on the impact that electronic health record (EHR) systems, combined with the advent of e-discovery rules, will have on traditional documentation issues.

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Healthcare Antitrust FAQ Handbook

First Edition

Mark L. Mattioli, Alexander M. McIntyre, Jr., David M. Narrow, Stephen P. Murphy, Patricia M. Wagner, and Hillary A. Webber

Here is an important resource for anyone needing to understand how antitrust issues impact healthcare organizations. Whether you are in-house counsel with a healthcare organization, a healthcare attorney not generally involved with antitrust issues, or even one more familiar with how issues impact healthcare organizations, you will benefit from this handy guide.

The book employs a practical question-and-answer approach for understanding the antitrust implications for structuring deals in this sector.

The subjects covered include not only the traditional antitrust questions but other topics specific to healthcare, such as:

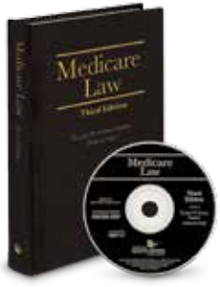
- Antitrust implications in the peer review and medical staff arena
- Physician and other healthcare provider networks
- Sharing healthcare price information

Questions include:

- When is it permissible to talk to a competitor about merging and what information can be shared?
- How do federal antitrust agencies determine when to challenge mergers?
- How can you take advantage of the immunity protection of the Health Quality Improvement Act of 1986 (HCQIA) in credentialing matters?
- When are exclusive contracts with primary payers permissible?
- What type of provider network arrangements can avoid per se condemnation?
- What authority do state attorneys general have to investigate federal antitrust violations?

The questions and answers are bolstered by extensive footnotes for further expanding research into each area of coverage.

(See also *Antitrust and Healthcare: A Comprehensive Guide*, p. 3.)



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eISBN 9780327175155

Pub. #26965, hardbound with CD-ROM, 525 pages, © 2012

Medicare Law

Third Edition

Thomas W. Coons, Editor in Chief; Rodney A. Johnson, Emily Jane Cook, John R. Hellow, David W. Hilgers, Elizabeth T. Thomas, Matthew Philip Utech, Kathleen Ann Peterson, Anne W. Hance, Gary Scott Davis, Adam J. Rogers, Donald H. Romano, Robert L. Roth, William T. Mathias, and James F. Flynn, authors

Since the last edition of *Medicare Law*, the rules that govern Medicare have changed substantially, with many changes being driven by the Affordable Care Act. Here are a few of the areas of coverage that have been updated in the latest edition:

- New or modified payment systems
- Medicare reimbursement rules, which are tied much closer to the quality of the services delivered
- The addition of significant Medicare reporting requirements
- Modified billing including new time limits on the filing of claims
- Strengthening of the fraud and abuse rules
- New payment approaches, such as Accountable Care Organizations, that are being or have been introduced

Comprehensive coverage also includes in-depth examinations of:

- Payment for hospital services, including PPS and PPS-exempt hospitals
- The DRG system
- The urban/rural distinctions
- Outlier payments
- The physician fee schedule
- Cost reimbursement
- Prohibited acts under the Anti-Kickback and Stark Laws
- Administrative and judicial appeals processes
- The new incentive payment programs

This is the road map to a thorough understanding of Medicare, whether you are just beginning the journey or are continuing to learn about this complex and important aspect of healthcare law.

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ISBN 9780769849577
eISBN 9780327175346

Pub. #28198, softbound with CD-ROM, 350 pages, © 2012

Pharmaceutical and Medical Device Compliance Manual

First Edition

Kathleen M. Boozang and Simone Handler-Hutchinson, Editors; Bret A. Campbell, Sujata Dayal, Michael Andre Donnella, Katie Rose Fink, Brett R. Friedman, Gary F. Giampetruzzi, Christopher R. Hall, Patrick M. Hromisin, Elizabeth H. Kim, Daniel A. Kracov, Bruce A. Levy, Ann E. Lewis, Benjamin S. Martin, Joseph W. Metro, Lewis Morris, Kiaema R. Reid, Linda Pissott Reig, Margaret Renner, Mary Riordan, Lynn Shapiro Snyder, Brian Tretick, Robert E. Wanerman, and Mara E. Zazzali-Hogan, authors

Since 1996 the federal government has strengthened its efforts to detect and prevent fraud and abuse in healthcare. This recently published *Manual* is your key to effectively protecting companies from investigation and prosecution.

The publication is a joint project of the Center for Health & Pharmaceutical Law & Policy at Seton Hall University School of Law, the Food and Drug Law Institute, and the American Health Lawyers Association.

The *Manual* offers an in-depth overview of the federal and state enforcement agencies that are responsible for investigating and resolving violations of the law by healthcare entities, with an emphasis on pharmaceutical and medical device companies. Coverage includes:

- The federal Anti-Kickback Statute and the False Claims Act
- Promotion of off-label uses for drugs and reimbursement for prescription drugs and medical devices
- Distinctions between manufacturers' lawful dissemination of scientific information, and the unlawful promotion of off-label usage
- The Foreign Corrupt Practices Act
- Implications of extra-territorial reach outside the United States
- Prescription drug price regulations

This *Manual* is a must-have for anyone involved in building a compliance program: health and life sciences attorneys, compliance officers, and other professionals in the industry.

(See also ***Healthcare Compliance Legal Issues Manual, Third Edition***, p. 18.)

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ISBN 9780769859507
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Pub. #26965, softbound with CD-ROM, 800 pages, © 2012

Representing Physicians Handbook

Third Edition

Michael F. Schaff, Task Force Chair; Glenn P. Prives, Coordinating Editor; Bradford E. Adatto, Steven R. Antico, Todd R. Bartos, Philip F. Berg, Brian C. Betner, Ann M. Bittinger, Andrew E. Blustein, Mark A. Bonanno, Steven J. Chananie, Nora A. Colangelo, Kathleen L. DeBruhl, Richard E. Gardner III, Nicholas J. Giampetro, Peter A. Greenbaum, Rick L. Hindmand, David J. Hyman, Bernard E. Jacques, Kimberly Kempton-Serra, Tara Kepler, Mark S. Kopson, Alyson M. Leone, David T. Lewis, Kim Harvey Looney, Rolf E. Lowe, Theresamarie Mantese, Jay A. Martus, Jeremy N. Miller, Gregory M. Nowakowski, Mark W. Peters, Cynthia Y. Reisz, Todd A. Rodriguez, David L. Rogers, Stephanie A. Roth, Joseph Rugg, Gary S. Sastow, James W. Saxton, Andrew Stathopulos, Jennifer Pearson Taylor, Rhonda Teitelbaum, Mark R. Thompson, and Sidney S. Welch, authors and/or editors

The world of physician practice continues to evolve, as do the complexities facing both physicians and the attorneys who represent them. With the advent of healthcare reform, the increased consolidation of medical practices, and ownership by physicians of ancillary services, a greater understanding of the issues impacting solo and small practices is more and more vital.

The latest edition of this popular handbook thoroughly addresses:

- The need to understand the Stark regulations
- Compliance issues
- Business and reimbursement
- The growing effort by state and federal government agencies to attack fraud and abuse
- And many other areas

Summary Table of Contents

- Regulatory Issues Affecting Physicians
- Telemedicine
- Employment Agreements
- Life Cycle of Association: The Buy-In and the Buy-Out
- Tax Principles Concerning Buy-Outs and Related Post-Withdrawal Compensation Issues
- Practice Breakups and Physician Departures
- Physician/Hospital Relationships
- Malpractice Insurance and Risk Management
- Physician Joint Ventures
- The Sale and Purchase of a Medical Practice
- Hospital-Based Physician Representation
- Use of Non-Competition Covenants in Physician Employment Relationships
- Compliance, Compliance Plan, and Process for the Physician Practice
- Non-Physician Practitioners
- Physician Ancillary Services
- Physician Recruitment Agreements
- Concierge Medicine

(See also *Physician Recruitment and Compensation Arrangements Practice Guide, Third Edition*, p. 22.)

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Clinical Research Practice Guide

Second Edition

R. Harold McCard, Coordinating Editor; Melissa L. Markey, David S. Weinstock, and Jamie K. Wolszon, Monica R. Chmielewski, M. Leeann Habte, Jonathan M. Holda, E. Scott Johnson, Christopher F. Lonegro, Aaron J. Rabinowitz, Sarah E. Swank, Lawrence W. Vernaglia, and Torrey K. Young, authors

With this important book, those who represent healthcare entities will gain insight into the growing interrelationship between health law and life sciences, as well as the growing risks and compliance issues facing clients involved in this complex area. The Second Edition of the popular *Practice Guide* is extensively enhanced and updated with coverage of:

- Changes to HIPAA that affect research and IRBs
- New Food and Drug Administration guidances and finalization of draft guidances
- Changes to patent law due to the America Invents Act
- HITECH and enhanced coverage of the Security Rule
- New requirements for group health plans and health insurers to cover routine patient costs in an approved clinical trial

Comprehensive coverage in the *Clinical Research Practice Guide* is both analytical and practical, with thorough treatment of:

- The development of human subject protections
- The federal approval process for pharmaceuticals, medical devices, and biologics
- The responsibility for regulatory oversight and investigations, with a discussion of which agency in the U.S. Department of Health and Human Services has responsibility for ensuring compliance
- Intellectual property considerations, including an extensive discussion of the basics
- Information management, including the sources for human subject protection: the Federal Common Rule, FDA Guidance, and HIPAA
- Insurance payments for clinical trial services, whether through private insurance or federal reimbursement

(See also *Institutional Review Boards: A Primer, Second Edition*, p. 20.)

Fraud & Abuse Investigations Handbook for the Healthcare Industry

COMING FALL 2013

Robert A. Griffith and Paul W. Shaw

This *Handbook* is designed to provide healthcare administrators, executives, medical practice directors and managers as well as attorneys, with a broad overview of healthcare fraud investigations.

The authors' discussion is supplemented with a comprehensive set of sample government documents, including subpoenas and search warrants, as well as helpful letters and memoranda generated by experienced healthcare fraud and abuse defense attorneys. Understanding the powers, procedures, and remedies available to investigative and law enforcement agencies is critical for healthcare industry executives, managers, and attorneys who find themselves the focus of such investigations.

(See also *Legal Issues in Healthcare Fraud and Abuse, Fourth Edition*, p. 21.)

COMING FALL 2013

HIPAA/HITECH Handbook

First Edition

The final regulations governing HIPAA and HITECH are out, and the compliance date is approaching quickly. This is the ideal guide for understanding how the laws have evolved and for advising your clients. Coverage includes:

- The original HIPAA regulations
- HITECH Act Privacy provisions and rulemaking
- Modifications to HIPAA Privacy, Security, and Enforcement Rules
- Includes valuable practice tools such as sample Business Associate Agreements, sample privacy practice notices, authorizations, policies, and training materials

This *Handbook* is the all-in-one resource that you need to keep up with HIPAA/HITECH compliance.

COMING FALL 2013

Peer Review Hearing Guidebook

Second Edition with CD-ROM

S. Allan Adelman and Ann O'Connell, Co-Editors and authors

Peer Review Hearing Guidebook focuses on the peer review hearing process, including steps that should be taken long before a medical staff hearing is contemplated. This is a critical resource that will help you be sure that all necessary procedures are in place to facilitate an effective and fair hearing.

The practical new *Guidebook* includes:

- Coverage of all of the legal issues involved in peer review hearings
- Practical steps for improving the peer review and hearing processes
- A companion CD-ROM that provides sample forms, checklists, bylaws, and fair hearing provisions—customizable for individual clients
- A survey of statutes, regulations, and cases from all 50 states relating to the right to and conduct of peer review hearings

(See also *Peer Review Guidebook, Fourth Edition*, p. 22.)

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eISBN 9780327174981

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ISBN 9780769846507
eISBN 9780327174998

Pub. #28190, softbound with CD-ROM, 325 pages, © 2012

The ACO Handbook, A Guide to Accountable Care Organizations

First Edition with CD-ROM

Peter A. Pavarini, Charlene L. McGinty, Michael F. Schaff, Editors; Thomas E. Bartrum, Elisabeth Belmont, Rudolf M. Blumentritt, Mark W. Browne, Timothy J. Cahill, Lauren N. Haley, Robert G. Homchick, Travis F. Jackson, Marilyn Lamar, Alyson M. Leone, Nathan L. Lutz, Barbara L. Miltenberger, Kimberly A. Mobley, Jan E. Murray, David R. Pearl, Glenn P. Prives, Toby G. Singer, Harvey M. Tettlebaum, Danielle L. Trostorff, John R. Washlick, authors

It is the importance of accountable care organizations (ACOs), as much as the uncertainty about their future, that makes *The ACO Handbook: A Guide to Accountable Care Organizations* necessary for those grappling with the changes brought about by healthcare reform. The contributors to this *Handbook* are among the best professionals in America today who are seriously considering what it will take to succeed under the new healthcare environment. Their astute observations about the legal issues, both novel and familiar, likely to be encountered by those contemplating ACO development will be useful to the reader no matter what role these organizations ultimately play in the reformed U.S. healthcare system.

The authors begin with a discussion of the importance of this new model, review the integration models used by hospitals and physicians prior to enactment of the PPACA, and continue with examinations of issues such as cost containment as envisioned in the "Medical Home," as well as waivers by the federal government of aspects of the Stark Law, the Anti-Kickback Statute, and certain provisions of the Civil Monetary Penalties Law.

AHLA's Federal Healthcare Laws & Regulations

2011 – 2012 Edition with CD-ROM and 2013 Supplement

William W. Horton, Editor; W. Scott Hardy, Alan M. Kirschenbaum, Arthur N. Lerner, Patricia A. Marcus, R. Harold McCard, John A. Meyers, Paul W. Shaw, Donald B. Stewart, and Judith A. Waltz, Editorial Advisory Board

New edition Fall 2013

$324 • **Members**
ISBN 9781422496558

$374 • **Non-members**
ISBN 9781422496565

Pub. #26742, softbound, 3 volumes with current supplement, 6,000 pages, © 2011

Federal Healthcare Laws & Regulations from AHLA is a comprehensive three-volume compilation that incorporates the most significant and timely federal statutes and regulations for the healthcare practitioner. It includes:

- The Affordable Care Act and the many new regulations that accompany it
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- Enhanced enrollment requirements
- The HIT certification program
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- The Stark Law
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- Key regulations governing CMS and the Office of Inspector General
- Rules and regulations on many other critical aspects of your healthcare law practice

A fully searchable CD-ROM allows you to research more efficiently and copy and paste the text of statutes and regulations easily into your documents.

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- **Title 21** Food and Drugs
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- **Title 29** Labor
- **Title 31** Money and Finance
- **Title 35** Patents
- **Title 38** Veterans' Benefits
- **Title 42** The Public Health and Welfare

Volume II

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- **Title 16** Commercial Practices
- **Title 20** Employees' Benefits
- **Title 21** Food and Drugs
- **Title 24** Housing and Urban Development
- **Title 26** Internal Revenue
- **Title 29** Labor
- **Title 42** Public Health (through Part 412)

Volume III

Selected Sections from the Code of Federal Regulations

- **Title 42** Public Health (Part 413 – 1008)
- **Title 45** Public Welfare
- **Index**

CD-ROM

The companion CD-ROM contains the complete contents of the 3 volumes in a fully searchable Folio format.

Ambulatory Surgery Centers: Legal and Regulatory Issues

Fourth Edition with 2011 Supplement and CD-ROM

Scott Becker, Sarah Abraham Chacko, Ronald E. Lundeen, Jr., Elissa Koch Moore, Melissa Szabad, Gretchen Heinze Townshend, and Amber McGraw Walsh

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eISBN 9780327174011

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ISBN 9781422461846
eISBN 9780327174134

Pub. #26900, softbound with 2011 supplement and CD-ROM, 280 pages, © 2009

This publication addresses the unique nature of ASCs, emphasizing their physical and organizational separation from other providers, regardless of whether they are owned by a hospital, a physician practice, or other entity. This revised and updated edition focuses on current issues for ASCs, and offers practical and useful guidance for those involved in giving legal advice to them. Coverage includes: key trends and tensions facing ASCs, such as the points of conflict that often arise between ASCs, hospitals, and physicians; Medicare and Medicaid fraud and abuse concerns, tax exemption and other tax-related issues, and the safe harbors for ASCs; relevant Stark issues, as well as the impact of state self-referral laws; and antitrust considerations for ASCs that face often intense competition.

Corporate Governance Implications of Nonprofit Executive Compensation

First Edition with CD-ROM

Douglas K. Anning, John B. Beard, Ralph DeJong, Susan G. Duffy, Stuart Harvey, Marci Rose Levine, Karen B. McAfee, Susan Schalla, Thomas C. Schroeder, and Michael J. Stewart

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ISBN 9781422457795

Pub. #26837, softbound, 110 pages, © 2009

This publication discusses the need for a proper governance process for the approval and implementation of executive compensation programs in nonprofit healthcare organizations. It also focuses on the federal tax concepts and guidance applicable to executive compensation paid by tax-exempt healthcare providers; recent Congressional, IRS and state-level inquiries, compliance initiatives and audits; the use of comparability data and expert opinion in building a credible governance process; governance structure and policies; and reporting and disclosure of executive compensation.

Data Breach Notification Laws: A Fifty State Survey

First Edition with 2013 Supplement

Jonathan M. Joseph

The risk of a data breach has increased tremendously in recent years. While the U.S. Congress has enacted breach notification requirements in a number of Acts, data breach concerns are not limited to federal law. This is due to actual breaches of state information systems, and has led state legislatures to enact an array of data breach notification laws.

This publication is a guide to existing state laws, and includes selected statutes for a number of states to illustrate not only the approach that larger states have taken, but also to highlight some of the more unusual approaches that some states have followed.

Highlights include:

- A fifty state survey of data breach notification statutes
- The text for the relevant act in CA, NY, IL, TX, FL, MA, MD, MI, NH, NJ, NC, WV, and WI
- Sample data breach notification to consumers, reporting forms, sample letter to a state Attorney General, and breach notification requirements for the Commonwealth of Virginia

The 2013 Supplement brings the information in this publication up to date with changes enacted in numerous states and also includes additional sample notice letters and forms.

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ISBN 9781422495285
eISBN 9780327174929

$139 • **Non-members**
ISBN 9781422495292
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Pub. #28156, softbound with 2013 supplement, 150 pages, © 2011

Deciphering Codes: Fraud & Abuse for Coders and Coding Issues for Healthcare Lawyers

First Edition with CD-ROM

Rhonda Buckholtz and Robert A. Pelaia, Project Leaders; Brad Ericson and Renee Dustman, Editors; Katherine Abel, Nicole Benjamin, Richard J. Brooderson, Rhonda Buckholtz, Gregory A. Chaires, Julie E. Chicoine, Shelly Cronin, Kerin Draak, Stephanie G. Ellis, Brad Ericson, Amy E. Fouts, Michael A. Gardner, Nate Gilmer, Deborah Grider, JoAnn M. Guerrero, Jillian Harrington, Raemarie Jimenez, Jodi B. Laurence, Terry Leone, Kelly Loya, Theresamarie Mantese, Jonnie Massey, Robert E. Mazer, Leslie Murphy, Gregory M. Nowakowski, Christopher A. Parrella, Abby Pendleton, Liston E. Radney III, David L. Rogers, Donna SanGiovanni, Harry R. Silver, Kevin Solinsky, and Jennifer L. Weaver, authors

This unique publication is designed to help coders and attorneys understand and navigate the complex intersection of healthcare coding and the law. This valuable manual discusses the nuances of coding, why proper coding is essential for healthcare facilities and professionals, and the legal implications of improper coding.

It covers the various sources for codes, provides an overview of the applicable fraud & abuse statutes and regulations, and then discusses ten specific healthcare settings where coding is a critical element. The importance of a detailed compliance plan for coding is discussed throughout the book, as are cases brought by federal and state governments against healthcare providers for improper coding. Also included is a searchable CD-ROM containing the full text of the manual with links to selected cases, statutes, and regulations.

$164 • **Members**
ISBN 9781422491157

$194 • **Non-members**
ISBN 9781422491164

Pub. #28007, loose-leaf with CD-ROM, 450 pages, © 2010

False Claims Act & The Healthcare Industry: Counseling & Litigation

Second Edition with 2013 Cumulative Supplement

Robert S. Salcido

Significant events have occurred regarding the government's enforcement and administration of the False Claims Act (FCA) in the last few years.

In 2012, the Department of Justice announced that since the FCA was substantially amended in 1986, it had recovered more than $30 billion under the FCA, and in 2012 it recovered nearly $5 billion, of which $3 billion was from health-related FCA cases.

With the stakes this high, professionals involved in this area will benefit from this comprehensive two volume work. Coverage in the 2013 supplement includes:

- Actions alleging a violation of the Anti-Kickback statute and Stark Law
- Important defenses the courts have applied to dismiss false claim actions
- Dismissing FCA actions under the FCA public disclosure jurisdictional bar
- Whether the plaintiff can establish that the defendant retaliated against the plaintiff for investigating fraudulent conduct
- Whether the tolling provisions of the FCA apply when the United States declines to intervene in a qui tam action

(See also *Legal Issues in Healthcare Fraud and Abuse, Fourth Edition*, p. 21.)

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ISBN 9781422443163
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Fifty State Survey of Certificate of Need and Licensure: Nursing Homes, Assisted Living, Home Health, and Hospice

First Edition with CD-ROM

Ari J. Markenson and Carrie K. Rabbitt, Editors

This publication is designed to provide attorneys and managers in the long term care, sub-acute and post-acute care industries with a detailed overview of the legal and regulatory landscape for obtaining certificate of need and/or licensure for nursing homes, assisted living facilities, home, health, and hospice services in each of the fifty states and the District of Columbia.

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ISBN 9781422457740

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Fundamentals of Health Law

Fifth Edition with CD-ROM

Barry D. Alexander, Bernadette M. Broccolo, Anthea R. Daniels, Sandra M. DiVarco, Cynthia B. Hutto, Carol Colborn Loepere, David E. Matyas, Thomas Wm. Mayo, John J. Miles, Eli A. Poliakoff, Ross E. Sallade, Michael F. Schaff, Susan O. Scheutzow, Daniel J. Schwartz, and Kerrin B. Slattery

This edition covers the basic issues of health law practice, from patient to facility issues, from permits and regulation issues to compliance and investigation issues, and includes issues raised by new laws, regulations, and guidelines promulgated since the Fourth Edition, including the Affordable Care Act (ACA).

This publication covers fundamental legal principles and issues to assist:

- New practitioners or experienced attorneys entering their first years of health-law practice
- Professors of health law searching for a comprehensive text for their students
- Users of any law library looking for answers on the health law resource shelf

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The Fundamentals of Life Sciences Law: Drugs, Devices and Biotech

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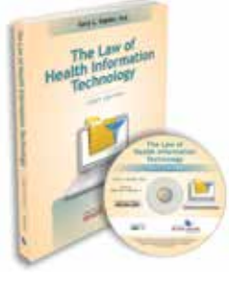

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Rebecca F. Cady, Editor; Anne M. Murphy, Susan Wood O'Leary, Mary Anne Hilliard, Ronni P. Solomon, Maria T. Currier, Reetu Dua, Jose I. Fernandez, authors

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