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\$174 • Members ISBN 9781632820679 eISBN 9781632820693

### \$204 • Non-members ISBN 9781632820686

eISBN 9781632820709

Pub. #28190, softbound 325 pages, © 2015

# The ACO Handbook: A Guide to **Accountable Care Organizations**

### Second Edition

Peter A. Pavarini, Charlene L. McGinty and Michael F. Schaff, executive editors Mark L. Mattioli, coordinating editor

Gregory D. Anderson, Peggy L. Barlett, Clifford E. Barnes, Troy A. Barsky, Thomas E. Bartrum, Elisabeth Belmont, Julian D. "Bo" Bobbit Jr., Timothy J. Cahill, Michelle E. Calloway, Robert James Cimasi, Sarah S. Fallows, Robert A. Gerberry, Jenny E. Gladieux, David W. Grauer, Lauren N. Haley, Rick L. Hindman, Robert G. Homchick, Travis F. Jackson, John M. Kirsner, Teresa Koenig, David E. Kopans, Marilyn Lamar, Kelly A. Leahy, Alyson M. Leone, Kim Harvey Looney, Daniel C. Lyons, Grace D. Mack, Mary C. Malone, Charlene L. McGinty, David W. McMillan, Rodney A. Myer, Thomas E. Miller, Kimberly A. Mobley, M. Daria Niewenhous, Peter A. Pavarini, David R. Pearl, Craig Pederson, Glenn P. Prives, Martie Ross, Michael F. Schaff, Thomas N. Shorter, Toby G. Singer, Donald B. Stuart, Sarah E. Swank, Elizabeth E.H. Trende, Danielle L. Trostorff, Claire M. Turcotte, John R. Washlick, Stephen M. Weiner, Keith Wright Ill, Todd A. Zigran, authors

The authors and editors contributing to this *Handbook* have strived to produce the most comprehensive guide to ACOs available today. It is a critical work whether you're advising a hospital or health system in establishing an ACO model, or counseling organizations already operating within the delivery structure.

The book analyzes various components of an optimized delivery system, and examines issues ranging from the financial aspects of ACOs to the unique nature of academic medical center, pediatric, and commercial ACOs. Coverage includes:

- New chapters examining compliance plans, non-Medicare commercial ACOs, best practices, and payer perspectives
- > A review of integration models used by hospitals and physicians prior to enactment of the ACA
- > Examinations of cost containment in the "Medical Home"
- > Waivers by the federal government of aspects of the Stark Law, the Anti-Kickback Statute, and certain provisions of the civil monetary penalty law (Gainsharing CMP)
- > The implications for various organizations in their pursuit of integration and coordination of services, whether as an ACO or otherwise
- > In-depth discussions of why understanding health information technology, antitrust, financing, and risk-sharing issues are critical to the operation and success of ACOs

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# > Watch for coming new titles/new editions

for Accountable Care Organizations

Ancillary Providers in Health Care: A Primer, First Edition

Data Breach Notification Laws: Fifty State Survey, Second Edition

Diagnostic Imaging Centers: Legal and Regulatory Issues, Third Edition

Health Care Entity Bylaws and Related Documents, Fourth Edition

Health Plans Contracting Handbook: A Guide for Payers and Providers,

Seventh Edition

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\$104 • Members ISBN 9781632817549 eISBN 9781632828637

\$134 • Non-members ISBN 9781632817556 eISBN 9781632828644

Pub. #30230, softbound, 122 pages, © 2015

# **Health Plan Disputes and Litigation Practice Guide**

### First Edition

Joseph Scott Schoeffel and Julie A. Simer

The passage of the Affordable Care Act has led to numerous changes to the delivery of health care, including new marketplaces to purchase insurance to cover health care expenses. As more people become insured, their health plan decisions become even more important to them and to the health plan industry in general.

Experienced practitioners Joseph Scott Schoeffel and Julie A. Simer provide thorough treatment of those areas likely to give rise to disputes. The book will be a welcome addition to the libraries of both new practitioners in this area and veteran practitioners alike.

#### Table of Contents

- Ch. 1 The Role of Health Plans in the United States
- Ch. 2 Who is the Client?
- Ch. 3 Is This a Managed Care Issue?
- Ch. 4 Is the Health Plan Sponsored by an Employer?
- Ch. 5 What are Some of the Relevant Documents?
- Ch. 6 What Other Causes of Actions May be Filed Against a Health Plan?
- Ch. 7 What are Common Reasons for Denial of Benefits?
- Ch. 8 What Defenses are Available to Defeat an FRISA Claim?
- Ch. 9 Was Provider In-Network?
- Ch. 10 Does Dispute Involve a Medicare or Medicaid Health Plan?
- Ch. 11 What Causes of Action May Health Plan Assert?
- Ch. 12 When May Arbitration be Required?
- Ch. 13 Has Market Competition Been Affected?
- Ch. 14 Is Member's Health Information at Risk?
- Ch. 15 Does Dispute Implicate Affordable Care Act?
- Ch. 16 Conclusion
- Appendix A Commonly Used Health Care Terms
- Appendix B Case Summaries
- Appendix C Managed Care Litigation Decision Tree

# > RECENTLY PUBLISHED



# **Corporate Practice of Medicine: Fifty State Survey**

### First Edition with CD-ROM

Stuart Silverman, Chair, AHLA Corporate Practice of Medicine Project

Anthony H. Choe, Terri A. DeSio, Alyson M. Leone, Glenn P. Prives, Daniel Z. Sternthal, and Rose J. Willis, team leaders; with numerous authors and editorial board members

The corporate practice of medicine doctrine (CPOM) has a long history as an effort by the American Medical Association to distinguish physicians from non-physicians offering services and cures for various afflictions. The doctrine seeks to prohibit a non-physician from interfering with a physician's professional judgment by prohibiting corporations not owned or controlled by physicians from employing physicians to practice medicine and charge for those professional services.

This valuable resource is for anyone needing to determine how a particular state addresses the corporate practice of medicine doctrine. The doctrine is most often invoked in contract disputes, such as enforcement of noncompetition agreements and the right to receive reimbursement from third parties. Some courts have cited the doctrine in refusing to enforce an insurance carrier's reimbursement to a medical corporation operating in violation of a state's CPOM.

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# \$209 • Non-members

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\$204 • Non-members ISBN 9781632806161 eISBN 9781632816436

Pub. #27060, softbound with CD-ROM, 570 pages, © 2014

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# **Health Care Compliance** Legal Issues Manual

### Fourth Edition with CD-ROM

Harry R. Silver and Cynthia F. Wisner, editors

Amy Bailey, Douglas A. Blair, Nancy Bonifant, Elizabeth Carder-Thompson, Thomas S. Crane, Gerald "Jud" E. DeLoss, Andrew Dick, Gerald M. Griffith, Jacob J. Harper, Jessica M. Herron, Ann T. Hollenbeck, Kenneth Hooper, Gabriel L. Imperato, Richard Korman, Ronald H. Levine, Melissa L. Markey, Robert A. Pelaia, Vanessa A. Reynolds, Albert W. Shay, Harry R. Silver, E. John Steren, Cynthia F. Wisner, Howard J. Young, authors

The latest edition of Health Care Compliance Legal Issues Manual continues to act as the authoritative source for every stakeholder in the health care arena. With contributions from more than 20 esteemed authors and editors writing in their respective areas of expertise, the latest edition of this acclaimed Manual covers regulatory and legal developments, as well as practical guidance for complying with requirements.

Thorough coverage includes:

- What constitutes a compliance program
- > How to conduct an internal investigation
- Audit basics
- > What to consider prior to deciding on repayments and disclosures
- Substantive overviews of the False Claims Act
- > The Stark and Anti-Kickback Laws
- HIPAA privacy and security
- > Issues in life sciences entities
- Tax compliance
- > A searchable CD-ROM containing the full text of the Manual with links to selected cases, statutes, and regulations

(See also Pharmaceutical and Medical Device Compliance Manual, First Edition, p. 24.)



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\$204 • Non-members ISBN 9781630435042 eISBN 9781630435271

Pub. #28293, softbound with CD-ROM, 518 pages, © 2014

# Post-Acute Care Handbook: Regulatory, Risk, and Compliance Issues

### First Edition with CD-ROM

Ari J. Markenson, Editor in Chief

Jeannie A. Adams, Heather O. Berchem, Dena M. Castricone, Phillip J. Chapman, Michael H. Cook, Sheryl Tatar Dacso, Robin Dale, Janet K. Feldkamp, James P. Holloway, Alan C. Horowitz, Matthew E. Jassak, John S. Linehan, Michael F. McGahan, James F. Miles, Julie B. Mitchell, Randi S. Nathanson, Daniel J. O'Brien, Michael A. Okaty, Emily M. Park, Paula G. Sanders, Howard L. Sollins, Daniel Z. Sternthal, Steven M. Swirsky, Sanford V. Teplitzky, authors Caroline Berdzik, Joseph A. Donchess, Alan C. Horowitz, James F. Miles, Barbara L. Miltenberger,

Michelle Petersen, Kelly A. Priegnitz, Christopher C. Puri, Lawrence W. Vernaglia, editors

This invaluable guide provides insight into what's ahead for professionals in the field and the evolving compliance and regulatory issues, including:

- > The evolution of the nursing home survey process
- > Residents' rights and facility practices
- > The importance of nursing facility agreements
- Compliance issues, including federal and state reimbursement requirements
- > Facilities' ability to challenge enforcement remedies imposed against them
- > Fraud and abuse issues that affect the industry
- Development of new systems and relationships that respond to incentives under the Affordable Care Act

#### Learn more about:

- > What to consider when building, buying, or selling a post-acute care facility
- > Financing for construction or acquisition of nursing homes
- > Continuing care
- > Assisted-living facilities

# > RECENTLY **PUBLISHED**



### \$129 • Members ISBN 9781632806109 eISBN 9781632811837

#### \$154 • Non-members ISBN 9781632806116 eISBN 9781632811844

Pub. #27010, softbound with CD-ROM, 172 pages, © 2014

# The Stark Law: Comprehensive Analysis + **Practical Guide**

### Fifth Edition

Charles B. Oppenheim

This completely updated, revised, and significantly expanded new edition supersedes and replaces the prior editions. More than a summary of the law, this publication addresses the legal effect of final Stark regulations, after the completion of the formal rule-making process. Written by Charles B. Oppenheim, a leading expert in the interpretation and application of the Stark Law, this publication provides an in-depth critical analysis, and contains a wealth of pointers and practical advice, as well as a look at what future direction the Stark regimen might take. Coverage includes:

- > The legal effect of the regulations and the regulatory process
- > Discussion of several key themes that emerge in the regulations, including the tempering of the previous trend of broadening exceptions for compensation arrangements
- > Key definitions and interpretive changes
- > Impact on the physician practice world including issues relating to physician recruitment, specialty hospitals, gain-sharing arrangements, and Stark-compliant physician joint ventures
- > Analysis of group practices and the exceptions applicable to them, as well as the recent guidance from CMS concerning the period of disallowance for referrals and claims, and analysis addressing the final rule



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ISBN 9780769857305

Pub. #26742, softbound, 3 volumes, with CD-ROM, 6,000 pages, © 2013

# AHLA's Federal Healthcare Laws & Regulations

2013 – 2014 Edition with CD-ROM and 2014 – 2015 Supplement

William W. Horton, editor; W. Scott Hardy, Alan M. Kirschenbaum, Arthur N. Lerner, Patricia A. Marcus, John A. Meyers, Daniel F. Murphy, Paul W. Shaw, Donald B. Stuart, and Judith A. Waltz, editorial advisory board

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 health care practitioner. It includes:

- The Affordable Care Act and the many new regulations that accompany it
- > E-prescribing
- > Enhanced enrollment requirements
- > The HIT certification program
- > Health insurance reform requirements for individual and group markets
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Title 42 The Public Health and Welfare

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Title 26 Internal Revenue

Title 29 Labor

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Selected Sections from the Code of Federal Regulations

Title 42 Public Health, continued

Title 45 Public Welfare

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

### Second Edition with 2015 Supplement and CD-ROM

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



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Pub. #26900, softbound with CD-ROM, 280 pages, © 2014 

# **Ambulatory Surgery Centers: Legal and Regulatory Issues**

Fifth Edition with CD-ROM

Scott Becker, Megan Michelle Bushee, LauraLee Lawley, Melissa Szabad, Anna Timmerman, Barton Walker, and Amber Walsh

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.



\$174 • Members
ISBN 9780769865560
eISBN 9780327185956

\$219 • Non-members ISBN 9780769865577 eISBN 9780327185963

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

# Antitrust and Healthcare: A Comprehensive Guide

### First Edition with CD-ROM

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

If you represent clients in the health care 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 health care industry—including providers and third-party payers—and the specific issues they confront.

(See also Healthcare Antitrust FAQ Handbook, p. 18.)



\$124 • Members ISBN 9780327175216 eISBN 9780769860954

\$159 • Non-members ISBN 9780327175223 eISBN 9780769860961

Pub. #27755, softbound with CD-ROM, 200 pages, © 2013

# Clinical Research Practice Guide

### Second Edition with CD-ROM

R. Harold McCard, coordinating editor

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

With this important book, those who represent health care 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 quidances
- > 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.)



# > SECOND EDITION COMING SOON

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\$139 • Non-members ISBN 9781422495292 eISBN 9780327174950

Pub. #28156, softbound, with supplement, 145 pages, © 2011

# 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.



# \$164 • Members ISBN 9781422491157 eISBN 9780327175230

# \$194 • Non-members ISBN 9781422491164 eISBN 9780327175247

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

# Deciphering Codes: Fraud & Abuse for Coders and Coding Insight 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 health care coding and the law. This valuable manual discusses the nuances of coding, why proper coding is essential for health care facilities and professionals, and the legal implications of improper coding.

It covers the various sources for codes, provides an overview of the applicable fraud and abuse statutes and regulations, and then discusses ten specific health care 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 health care 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.



\$169 • Members ISBN 9780769861166 eISBN 9780327175254

### \$209 • Non-members ISBN 9780769861173 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, Sheila Hagg-Rickert, 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 Norris, Gisele Norris, Richard S. Porter, Yvonne K. Puig, Steven M. Puiszis, Emily Rhinehart, Fay A. Rozovsky, Joshua Rozovsky, Mary S. Schaefer, and Kathryn K. Wire

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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# False Claims Act & The Healthcare Industry: Counseling & Litigation

# Second Edition with 2014 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 2013 it recovered nearly \$3.8 billion, of which \$2.6 billion was from health-related FCA cases.

With the stakes this high, professionals involved in this area will benefit from this comprehensive work. Coverage in the 2014 Supplement includes:

- > Actions alleging a violation of the Anti-Kickback statute and Stark Law
- > How defense 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
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Robert A. Griffith and Paul W. Shaw

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

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



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# **Fundamentals of Health Law**

# Sixth Edition with CD-ROM and 2015 Medicaid Supplement

Barry D. Alexander, Bernadette M. Broccolo, Anthea R. Daniels, Sandra M. DiVarco, Anjali N.C. Downs, Geoff A. Drucker, Catherine A. Hurley, Cynthia B. Hutto, Raymond J. Lindholm, Carol Colborn Loepere, Vicki L. Lung, David E. Matyas, Thomas Wm. Mayo, John J. Miles, Eli A. Poliakoff, Jouya Rastegar, Ross E. Sallade, Michael F. Schaff, Susan O. Scheutzow, Daniel J. Schwartz, Nancy A. Sheliga, Kerrin B. Slattery, and Craig H. Smith

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- > 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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Jeremy M. Alexander, Lauren Battaglia, Brian A. Bohnenkamp, Nancy E. Bonifant, Eve M. Brunts, Elizabeth Carder-Thompson, Carl H. Coleman, Susan A. Edwards, Paige A. Fillingame, David C. Gibbons, Daniel G. Gottlieb, Eric C. Greig, Simone Handler-Hutchinson, Clinton D. Hermes, Stuart S. Kurlander, Robert F. Leibenluft, Vicki L. Lung, Melissa L. Markey, Juliet M. McBride, Leigh L. Oliver, Jordan K. Paradise, Heather H. Pierce, Jennifer L. Pike, Preeya Noronha Pinto, Thomas J. Quinlan, Kelly N. "Nikki" Reeves, Corey W. Roush, Jason W. Sapsin, Richard B. Smith, Judith L. Toffenetti, Susan L. Walker, and Constance A. Wilkinson, authors

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- > Fraud and Abuse
- > Federal Agencies
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### Healthcare Antitrust FAQ Handbook

### First Edition with CD-ROM

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 health care organizations. Whether you are in-house counsel with a health care organization, a health care attorney not generally involved with antitrust issues, or even one more familiar with how issues impact health care organizations, you will benefit from this handy quide.

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 health care, such as:

- Antitrust implications in the peer-review and medical-staff arena
- > Physician and other health care provider networks
- > Sharing health care 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 Care 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, First Edition, p. 11.)



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(See also Peer Review Hearing Guidebook, Second Edition, p. 23.)



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Patricia D. King, editor

Jeffry Adest, Lara Jean Ancona, Ann M. Bittinger, Damon G. Carpenter, Alisa L. Chestler, Lisa L. Dahm, William P. Dillon, Gregory Ewing, Valita Marie Fredland, Jill M. Girardeau, Jennifer J. Hennessy, Caroline Kubovy, Morris A. Landau, Wendy C. Maneval, Valerie Breslin Montague, John Murdoch, Cheryl Camin Murray, Stephen Page, Jennifer L. Rathburn, Lisa Pierce Reisz, Stephen D. Rose, Amita A. Sanghvi, Sarah E. Swank, Leslie M. Tector, authors

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(See also The Law of Health Information Technology, First Edition, p. 21.)

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# Institutional Review Boards: A Primer

### Second Edition with CD-ROM

Teresa A. Williams, Richard G. Korman, Janet M. Lis, Melinda G. Murray, Kate Bowen, Amy L. Bradshaw, Tara R. Cowell, Amy K. Dow, Kimberly H. Gillespie, Jennifer R. Henderson, Marta J. Hoffman, Veronica A. Marsich, Tamara J. O'Black, Kay M. Perry, Carol A. Poindexter, Stacey A. Ries, Martha C. Romney, Jennifer Sharp, Sarah Shulman Swank, Alexandra Trinkoff, Leah A. Voigt, and Catherine M. With

The expanded second edition of Institutional Review Boards: A Primer outlines the regulatory requirements and legal challenges associated with the evolving area of human subject research, and presents a thorough discussion of the role that institutional review boards serve in this area. This useful guide not only explains the difference between the various regulations that govern human subject research, but it also details the steps to take to set up an institutional review board that can adequately perform its role.

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(See also Clinical Research Practice Guide, Second Edition, p. 13.)



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Gary L. Kaplan

This publication provides a road map to health information technology (HIT) and its challenges. The ever-evolving world of HIT requires a new way of planning for subjects as disparate as delivery of health care, retention of records for standard business reasons, and preparation for litigation. Successful HIT projects necessitate coordination between four critical constituencies: medical, business, technical, and legal. This practice guide delineates many of the considerations that health systems and other providers face today, and offers detailed guidance on complying with laws and regulations concerning privacy, security, copyright infringement, as well as marketing and advertising rules.

Highlights include:

- Guidance on how to avoid the pitfalls of negotiating for new technology systems
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(See also HIPAA/HITECH Resource Guide, First Edition, p. 20.)



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# Legal Issues in Healthcare Fraud and Abuse: Navigating the Uncertainties

Fourth Edition with 2015 Supplement

David E. Matyas, Carrie Valiant, Jason Eric Christ, and Anjali N.C. Downs

The fight against fraud in the health care industry will increase in intensity and sophistication in the 21st century. This bestselling road map describes the legal theories governments use to identify and eliminate fraud in health care.

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- A thorough review of governmental enforcement entities including the Department of Justice, the Department of Health and Human Services, as well as other federal agencies, state governments, and private payers
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[See also False Claims Act & The Healthcare Industry: Counseling & Litigation, Second Edition, p. 15.]



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Gregory G. Gosfield

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### **Medicare Law**

### Third Edition with CD-ROM

Thomas W. Coons, Editor in Chief

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

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- Medicare reimbursement rules, which are tied much closer to the quality of the services delivered
- > The addition of significant Medicare reporting requirements
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- New payment approaches, such as Accountable Care Organizations, that are being or have been introduced

Comprehensive coverage also includes in-depth examinations of:

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This publication provides a thorough understanding of Medicare, whether you are just beginning the journey or are continuing to learn about this complex and important aspect of health care law.



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# **Peer Review Guidebook**

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Daniel Mulholland, Lauren M. Massucci, and Charles J. Chulack, editors

Peer Review Guidebook is a comprehensive resource for understanding the peer review process and disciplinary proceedings involving physicians. The fourth edition contains information about new Joint Commission standards on Focused Professional Practice Evaluation and Ongoing Professional Practice Evaluation, a discussion of the Patient Safety and Quality Improvement Act, as well as the latest from the National Practitioner Data Bank. Also included is a searchable CD-ROM containing the full text of the publication with links to selected cases, statutes, and regulations.

(See also Peer Review Hearing Guidebook, Second Edition, below.)



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# **Peer Review Hearing Guidebook**

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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:

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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 health care. This 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 health care 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
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- > 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, Fourth Edition, p. 6.)



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Gerald M. Griffith

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- Virtual Equity Models
- > Other Physician Recruitment and Retention Incentives
- Lawyer & Compliance Officer Liability
- > Preparing for Audit and Investigation
- > Practice Acquisitions

(See also Representing Physicians Handbook, Third Edition, p. 26.)



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# Representing Physicians Handbook

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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

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 health care 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.

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- Non-Physician Practitioners
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- Physician Recruitment Agreements
- Concierge Medicine

(See also Physician Recruitment and Compensation Arrangements Practice Guide, Third Edition, p. 25.)

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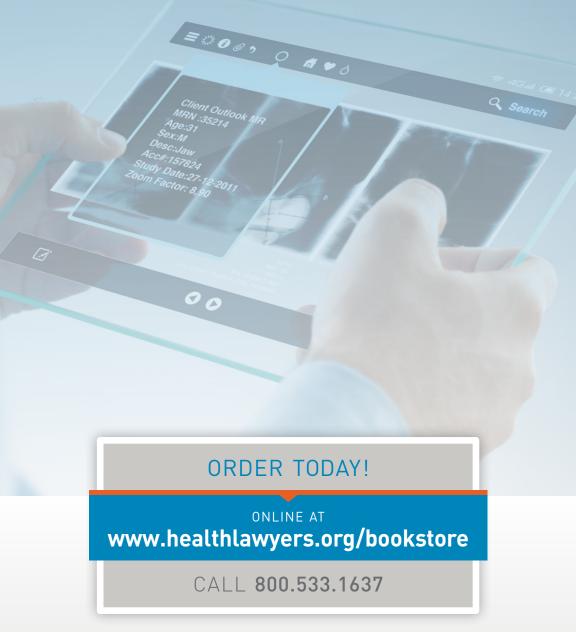
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