27th Pharmaceutical and Medical Device Ethics and Compliance Congress

October 13 – 16, 2026

Washington Hilton • Washington, DC

Sponsored by the Pharmaceutical Compliance Forum (PCF)

GHC Life Sciences
Global Health Care, LLC

A Hybrid Event — Onsite or Virtual Online Event Live and Archived

All Times are EDT · Agenda Current as of September 25, 2026 • Subject to Change and Regular Updates

C-SUITE CONGRESS VIDEO WELCOME:

Albert Bourla, DVM, PhD, *Chairman and Chief Executive Officer, Pfizer; Former Chair, Pharmaceutical Research and Manufacturers of America (PhRMA); Former President, International Federation of Pharmaceutical Manufacturers and Associations (IFPMA)*

GOVERNMENT KEYNOTE SPEAKERS:

Jacob Foster, JD, *Acting Chief, Health Care Fraud Unit, Criminal Division, Fraud Section, US DOJ*

Brenna Jenny, JD, MPH, *Deputy Assistant Attorney General, Commercial Litigation Branch, Civil Division, US DOJ*

Mary E. Riordan, JD, *Senior Counsel to the Inspector General, HHS Inspector General*

Veronika Peleshchuk Fradlin, MPP, *Director, Open Payments Data Group (OPDG), Office of Enterprise Data and Analytics (OEDA), US HHS*

Twyla Mosey, PharmD, RPh, *Director, Office of Prescription Drug Promotion, US FDA*

PCF BOARD:

Donna White, CCEP, *Vice President and Compliance Officer, Chiesi Farmaceutici S.p.A., Americas Region (Board Chair)*

Jill Dailey, JD, *Vice President & Chief Compliance Officer, Oncology, Menarini Stemline*

Daryl Kreml, JD, *Vice President, Head of Ethics & Compliance, Global Oncology, Takeda*

Christie Camelio, *Senior Vice President and Chief Compliance Officer, Insmed*

Anisa Dhalla, *Vice President, Global Head, Ethics and Business Integrity, UCB*

Melissa T. Lozner, JD, *Senior Vice President, Chief Compliance Officer, Regeneron*

MEDICAL DEVICE EXECUTIVE STEERING COMMITTEE:

Sujata Dayal, JD, LLM, *Founder & Senior Advisor, Arziyan LLC; Independent Board Director, Emergent BioSolutions*

Mona Peterson Rosow, JD, MPH, *Former Vice President, Chief Risk, Ethics & Compliance Officer, Mozarc Medical*

Daniel Spicehandler, JD, *Vice President Compliance, MSNT and Global Programs, Stryker*

JJ Kuhn, JD, *Chief Ethics & Compliance Officer, Danaher*

Tara Shewchuk, JD, LLM, *Senior Vice President, Chief Privacy, Integrity & Compliance Officer, Medtronic*

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- **Afternoon Half-day Specialty Focused Sessions**

Day 2: Wednesday, October 14, 2026

- **Networking Breakfast in the Exhibit Hall**
- **Morning Concurrent Mini Summits**
- **Networking Luncheon in the Exhibit Hall**
- **Afternoon Opening Plenary Session**
- **Networking Reception**

Day 3: Thursday, October 15, 2026

- **Networking Breakfast in the Exhibit Hall**
- **Morning PCF CCO Roundtable (Invitation-only, Breakfast Included)**
- **Morning Concurrent Mini Summits**
- **Networking Luncheon in the Exhibit Hall**
- **Afternoon Closing Plenary Session**

Day 4: Friday, October 16, 2026

- **Networking Breakfast**
- **Morning PCF Industry-only Think Tank**
- **Noon: Congress Adjournment**

IMPORTANT THINGS TO REMEMBER AS YOU PREPARE TO ATTEND THE CONGRESS

- For the first time, the Congress is offering 5 half-day Focused Sessions on Hot Topics on Tuesday afternoon, October 13.
- Onsite attendees may register onsite as early as 10 am ET on Tuesday, October 13.
- All onsite attendees will have the ability to scan QR codes on badges to exchange contact information.
- For the first time, all Congress attendees (both onsite and online) will be able to text each other via the Congress App.
- All onsite attendees will also have access to the Congress. Virtual Video/Audio Archive for at least 9 months following the Congress. This will allow attendees to watch sessions that they missed at the Congress.

ABOUT THE CONGRESS SPONSOR, THE PHARMACEUTICAL COMPLIANCE FORUM

Since 1999, PCF has been dedicated to the advancement of the pharmaceutical compliance profession. Our meetings provide a forum for members to share knowledge, explore innovative solutions to complex compliance challenges and network with peers to build long lasting career relationships.

The PCF's Vision is to be the leading forum in the pharmaceutical and biotech industry for promoting excellence in the compliance profession and advance effective and sustainable risk-based compliance programs through solutions-oriented collaboration and innovative best practice sharing.

PCF is proud of its Board, a team of six senior compliance executives who dedicate and volunteer their time to serve, each appointed by the membership for three-year terms. Our leadership team is responsible for setting the organization's strategic direction, overseeing business affairs, and ensuring we deliver meaningful value to our members.

For more information regarding the PCF, please contact PCF Administrator Debra Scanlon at deb_admin@pcf-org.com.

PCF Administrator Debra Scanlon (Deb)

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AGENDA AT A GLANCE

DAY 1: TUESDAY, OCTOBER 13, 2026

(Included in standard Congress registration)

11:00 am — **Registration Opens**

1:00 pm – 5:45 pm — **IN-DEPTH, HALF-DAY, AFTERNOON FOCUSED SESSIONS** *(Included in your Congress registration)*

Focused Session 1: **Whistleblower Complaints: From Hotline to Resolution: Managing Internal Investigations, Whistleblower Risk & Escalation Decisions**

1:00 pm • Chair's Welcome and Introductions & DOJ/ FCA Enforcement Trends: Lessons from Whistleblower Cases
1:45 pm • Qui Tam Counsel Roundtable
2:30 pm • With Data and AI, Whistleblowers Set Off an FCA Tidal Wave
3:45 pm • Hotline Isn't Enough: Building a Smart Investigations Ecosystem
4:30 pm • Anatomy of an Internal Investigation
5:15 pm • Managing Whistleblower Risk and Internal Escalation
5:45 pm **Adjournment**

Focused Session 2: **Pharmaceutical and Medical Device Advanced Privacy and Security/Cybersecurity Updates**

1:00 pm • Chair's Welcome, Overview and Introductions
1:15 pm • The Basics of Healthcare Privacy for Compliance Professionals and an Overview of Privacy Issues Unique to Pharmaceutical and Medical Device Enterprises
1:45 pm • IAPP Presents: Privacy, AI Compliance, and Beyond for the Future of Pharma and Medical Devices
2:30 pm • The International Pharmaceutical and Medical Device Privacy Consortium Presents Advanced Strategies for Addressing Privacy and Cybersecurity Issues for Life Sciences Companies
3:15 pm • Companies from the Cybersecurity Working Group, Health Sector Coordinating Council
4:15 pm • Cybersecurity Best Practices for Pharma and Device
5:00 pm • Session Faculty Q&A
5:45 pm **Adjournment**

Focused Session 3: **Managing Medical — Commercial Interactions: Strategic Collaboration, Evolving Roles & Risk-Based Governance**

1:00 pm • Co-Chair's Introductions & Session Overview
1:15 pm • Strategic Planning Together, Tactical Execution Separately: Practical Case Studies in Medical–Commercial Collaboration
2:00 pm • Beyond Sales Representatives and MSLs: Understanding Emerging Field-Based Roles and Governance Challenge
3:15 pm • Protecting Medical Independence While Enabling Strategic Cross-Functional Collaboration
4:00 pm • Medical–Commercial Boundaries: Congresses, Sponsorships & Promotional Risk
4:45 pm • Have We Become Too Restrictive? Rethinking Medical–Commercial Governance for the Next Era
5:30 pm **Adjournment**

Focused Session 4: **AI in Action: Real-World Compliance Case Studies from the Front Lines of AI Use in Life Science Compliance Departments**

1:00 pm • AI Basics, Life Science Industry Ethics and Compliance Applications
1:30 pm • A Brief Overview of the Epsilon & qordata Annual AI Use Compliance Survey
2:00 pm • Facilitated Case Study Discussion: From Transcript to Insights: How Compliance Teams Are Using AI to Transform Investigations
2:30 pm • Facilitated Case Study Discussion: AI Governing AI: Leveraging Artificial Intelligence to Develop Effective Governance Policies
3:15 pm • Facilitated Case Study Discussion: AI-Assisted Monitoring and Detection of Potential Policy Violations
3:45 pm • Facilitated Case Study Discussion: Agentic AI Use for Continuous Risk Assessment (vs Annual)
4:15 pm • The Unspoken Risk of AI: Moral Outsourcing — and Why "Add Human Judgment" Isn't Enough
5:00 pm • The Health Care Compliance Association Presents: Advanced Strategies in Using AI to Monitor and Improve Compliance: Lessons from Other Healthcare Sectors
5:45 pm **Adjournment**

Focused Session 5: **GP 101: A Primer on the Fundamentals of Government Pricing for Compliance Officers & The Evolving Pricing and Contracting Landscape: What Compliance Officers Need to Know**

1:00 pm • Chair's Introductions & Session Overview
1:15 pm • GP 101: A Primer on the Fundamentals of Government Pricing for Compliance Officers
4:00 pm • The Evolving Pricing and Contracting Landscape: What Compliance Officers Need to Know
5:30 pm **Adjournment**

AGENDA AT A GLANCE

DAY 2: WEDNESDAY, OCTOBER 14, 2026

7:00 am Registration Opens & Networking Breakfast in the Exhibit Hall

8:00 am – 8:50 am MORNING MINI SUMMITS: GROUP 1

MS1: Compliance Independence in the Real World: Governance Models That Work	MS2: The Agentic Compliance Function of the Future	MS3: Evolution of Patient Finding Activities	MS4: First Product Launch: Policy Development Strategies and Techniques	MS5: Speaker Programs in 2026: Practical Legal and Compliance Considerations for Executing from Beginning to End Amid Ongoing Scrutiny

9:00 am – 9:50 am MORNING MINI SUMMITS: GROUP 2

MS6: From Pronouncements to Practice: Understanding DOJ Enforcement Priorities	MS7: Boring is Risky! Want Stronger Ethics and Compliance Engagement? Train & Communicate Colorfully	MS8: Best Practices for Data Analytics	MS9: On the Front Lines: Navigating Field Force & Sales Compliance Risk	MS10: Communicating the Story on Integrity: What Compliance Leaders Can Learn from Health Stakeholders	MS11: Designing HCP Engagement with Transparency Built In

9:50 am – 10:10 am NETWORKING BREAK (in the Exhibit Hall)

10:10 am – 11:00 am MORNING MINI SUMMITS: GROUP 3

MS12: Compliance and M&A: Why Early Engagement Matters	MS13: The State of Medical Device Compliance: Annual Industry Roundtable	MS14: The State We're In: Legal & Enforcement Trends Reshaping Life Science Companies' Compliance	MS15: External Funding Risks and Mitigation: Latest Trends in Exhibit & Display and Sponsorship Requests	MS16: Beyond the Questionnaire: Building a Risk-Based Third-Party Management Program That Actually Works	MS17: Navigating Today's Enforcement Landscape with Culture, Data, and Ethics

11:10 am – 12:00 pm MORNING MINI SUMMITS: GROUP 4

MS18: Risk Advantage: How Resilient Organizations are Capturing Opportunity through Uncertainty	MS 19: View from the Top: Practical Advice for Accelerating Your Compliance Career	MS 20: Direct-to-Consumer (DTC) Models: Emerging Compliance Risks and Governance Considerations	MS 21: Fair Market Value: Practical Strategies and Targeted Insights for Building Defensible FMV Frameworks	MS 22: Navigating the Evolving US Privacy Landscape	MS 23: Agentic Risk Management: Transforming Compliance Across Key Business Processes

12:00 pm – 1:15 pm NETWORKING LUNCHEON — OR — OPTIONAL LUNCHEON DISCUSSION

ENJOY THE CONGRESS NETWORKING LUNCHEON OR ATTEND THIS OPTIONAL SMALL AND MID-SIZED LIFE SCIENCE COMPANIES LUNCHEON

12:10 pm – 1:10 pm Compliance Risks and Challenges in Small and Emerging Life Sciences Companies

Grab your box lunch and join a luncheon discussion designed for emerging and smaller life sciences companies that have limited to no existing compliance program elements. The focus is on helping compliance leaders determine where to begin and how to build a program in a practical, risk-based way.

The discussion will address questions such as:

- Which risks should be prioritized first?
- When should key compliance program elements and controls be implemented?
- How do you build a roadmap that aligns with the company's stage of growth?
- How do you maximize limited resources while establishing an effective compliance program

1:15 pm – 5:45 pm PHARMA/DEVICE CONGRESS OPENING PLENARY SESSION

1:15 pm	Welcome and Introduction: PCF Board
1:30 pm	C-Suite Congress Video Welcome
1:40 pm	Keynote Annual OIG Update
2:30 pm	Keynote US DOJ Criminal Division Update Fireside Chat
3:00 pm	Tech & Human-in-the-Loop Compliance (Why Human Judgment Matters More Than Ever)
3:30 pm	Networking Break in the Exhibit Hall
4:00 pm	The Next Generation of Compliance: Leveraging Data to Stay Ahead of Risk
4:30 pm	Integrated Risk Transformation in Pharma/Medical Device: From Compliance Burden to Strategic Advantage
5:00 pm	Annual Chief Compliance Officers Fireside Chat

5:45 pm OPENING PLENARY SESSION ADJOURNMENT AND NETWORKING RECEPTION

AGENDA AT A GLANCE

DAY 3: THURSDAY, OCTOBER 15, 2026

7:00 am — **Registration Opens & Networking Breakfast in the Exhibit Hall**

PCF CHIEF COMPLIANCE OFFICER ROUNDTABLE *(PCF-Sponsored, Exclusive Invitation-Only, Closed-Door Session)*

- **7:00 am** Breakfast Buffet
- **8:00 am** CCO Roundtable Welcome and Introductions
- **8:15 am** Future of the Compliance Profession in an AI Driven Environment
- **9:00 am** Executive Breakout by Company Size — *Leveraging AI to Support Compliance Programs*
- **9:40 am** Networking Break
- **10:00 am** Executive Breakout by Company Size — *Building AI Ready Teams*
- **10:40 am** Board Oversight of Compliance: What Audit Firms Observe and What Regulators Require
- **11:30 am** Open Forum: Opportunities for Benchmarking and Peer-driven Q&A. Bring your questions!
- **11:55 am** Wrap-up
- **12:00 pm** Adjournment

8:00 am – 8:50 am — MORNING MINI SUMMITS: GROUP 5

- MS24: Identifying and Managing Risk Associated with AI
- MS25: Transparency Reporting; Update on the Centers for Medicare and Medicaid Services Open Payments Program
- MS26: From Speak-Up to Proof: Turning DOJ and OIG Expectations into Defensible, AI-Enabled Compliance Operations
- MS27: Manufacturer–Specialty Pharmacy Relationships: Managing Compliance Risk in Specialty and Rare-Disease Commercial Models
- MS28: The Future of Compliance Monitoring: AI Agents, Risk Intelligence and Practical Adoption

9:00 am – 9:50 am — MORNING MINI SUMMITS: GROUP 6

- MS29: Building the Compliance Operating Model for What's Next
- MS30: Leveraging AI for More Efficient and Effective Investigations
- MS31: Compliance Considerations, Enforcements and Guidelines on Social Media Activities
- MS32: Building a Compliance Roadmap: Practical Strategies for Scaling Compliance Programs
- MS33: Fair Market Value in Flux: Navigating Complex Fee Arrangements, HCP & Patient Engagement, and Influencer Compensation Risks
- MS34: Future Proofing the Compliance Function: Operating Models for an AI-Enabled World

9:50 am – 10:10 am — NETWORKING BREAK IN THE EXHIBIT HALL

10:10 am – 11:00 am — MORNING MINI SUMMITS GROUP 7

- MS35: Unstructured Data and Advanced Monitoring Strategies
- MS36: Leading Practices for Managing Investigations Panel
- MS37: From Makary to Overton: Reflections and Predictions on the FDA's Priorities
- MS38: Compliance Considerations for Rare Disease Products
- MS39: From Oversight to Partnership: Strengthening Collaboration Between Compliance and the Business
- MS40: Digital Ecosystem Considerations for Life Science Companies

11:10 am – 12:00 pm — MORNING MINI SUMMITS GROUP 8

- MS41: Enabled Technology can Transform Transactional Testing and Other Components of Your Monitoring Program
- MS42: Emerging Commercial & Market Access Risk in Life Sciences: Navigating Innovation, Compliance, and Evolving Access Models
- MS43: Effectively Leveraging Post-Investigation Mitigation to Create a Narrative of Continuous Improvement
- MS44: Data Analytics & Compliance Monitoring in an AI-Powered World
- MS45: Can Learning Data Reduce Risk? Using Measurement to Drive Defensible, Risk-Based Compliance Training
- MS46: AI in Clinical Practice: Compliance, Governance & FDA Oversight

12:00 PM – 1:10 PM — NETWORKING LUNCHEON WITH DESSERT AND COFFEE IN THE EXHIBIT HALL

12:10 pm – 1:00 pm — LUNCHEON MINI SUMMITS GROUP 9

- MS47: Direct-to-Consumer (DTC) Models in the Pharmaceutical Space
- MS48: ROI of AI: Separating Value from Hype
- MS49: Lean, Credible, Defensible: Compliance Programs That Work Without Big Budgets
- MS50: A Look Under the Hood: Exploring PBM and Payer Arrangement Risks and Emerging Reporting Requirements
- MS51: Collaborating to Create Lasting Solutions During and After Monitoring or Investigations
- MS52: Annual FDA Office of Prescription Drug Promotion Update

1:10 pm – 5:15 pm — CLOSING PLENARY SESSION

- **1:10 pm** Welcome and Introduction: PCF Board
- **1:15 pm** Keynote Annual DOJ Civil Division Update Fireside Chat
- **1:45 pm** Investigations, Enforcement and Prosecutions Roundtable
- **2:15 pm** Updates from AdvaMed, BIO and PhRMA
- **2:45 pm** Market Access in 2026 and Beyond: A Shifting Compliance Landscape
- **3:15 pm** Networking Break in the Exhibit Hall
- **3:45 pm** Burgeoning Compliance Risks
- **4:15 pm** AI and the Evolving Compliance Function
- **4:45 pm** Managing Risk in Unsettled Times

5:15 pm — **CLOSING PLENARY SESSION ADJOURNMENT**

AGENDA AT A GLANCE

DAY 4: FRIDAY, OCTOBER 16, 2026

7:00 am **Breakfast Buffet**

8:00 am – 12:00 pm **PCF INDUSTRY-ONLY COMPLIANCE BEST PRACTICES THINK TANK**

(Hosted by PCF: Industry-only Session for Pharmaceutical and Medical Device Company Ethics and Compliance Professionals and In-house Counsel only)

7:00 am Breakfast Buffet
8:00 am Welcome and Introductions
8:10 am Inside the New DOJ Initiative: The Rising Importance of Data Miners
8:45 am Wins & Losses in the Early Days of AI – Audience Discussion Session
9:15 am Benchmarking and Q&A Forum
10:00 am Networking Break
10:20 am Customer-Centric Complexity: Meeting Customer Needs While Balancing Field Pressures
11:00 am Breakouts by Company Size
11:30 am Last Minute Questions – Wrap Up
12:00 pm Congress Adjournment

12:00 pm **CONGRESS ADJOURNMENT**

DETAILED AGENDA

TUESDAY, OCTOBER 13, 2026

PHARMACEUTICAL AND MEDICAL DEVICE ETHICS AND COMPLIANCE CONGRESS DAY 1 (Included in standard Congress registration.)

11:00 am Registration Opens /Lunch on Your Own

FEATURING FIVE CONCURRENT SPECIAL FOCUSED SESSIONS

Focused Session 1: Whistleblower Complaints: From Hotline to Resolution: Managing Internal Investigations, Whistleblower Risk & Escalation Decisions

1:00 pm **Chair's Welcome and Introductions & DOJ/FCA Enforcement Trends: Lessons from Whistleblower Cases**

John E. Kelly, JD, *Partner and Chair, Healthcare Industry Practice, Barnes & Thornburg; Former Assistant Chief for Health Care Fraud, Criminal Division, Fraud Section, US Department of Justice; Washington, DC (Chair)*

1:15 pm **A Fireside Chat with Nicholas C. Perros, JD, Senior Trial Counsel, Civil Division, Fraud Section, US DOJ**

Nicholas C. Perros, JD, *Senior Trial Counsel, Civil Division, Fraud Section, US Department of Justice; Washington, DC*

Interviewed by:
John E. Kelly, JD, *Partner and Chair, Healthcare Industry Practice, Barnes & Thornburg; Former Assistant Chief for Health Care Fraud, Criminal Division, Fraud Section, US Department of Justice; Washington, DC (Chair)*

1:45 pm **Qui Tam Counsel Roundtable**

Mary A. Inman, JD, *Partner, Whistleblower Partners; Founding Member & Board Chair, Psst.org; Member of Advisory Board, Whistleblowing Canada Research Society; San Francisco, CA*

Linda Severin, JD, *Managing Member, Whistleblower Law Collaborative; Former Assistant US Attorney, Southern District of New York, US Department of Justice; Boston, MA*

Gregg Shapiro, JD, *Founder, Gregg Shapiro Law; Former Assistant US Attorney and Chief, Affirmative Civil Enforcement Unit, US Attorney's Office, District of Massachusetts, US Department of Justice; Boston, MA*

Sarah M. Hall, JD, *Partner, Member of the Firm, Epstein Becker Green; Former Trial Attorney, Fraud Section, Criminal Division, US Department of Justice; Washington, DC (Moderator)*

2:30 pm **With Data and AI, Whistleblowers Set Off an FCA Tidal Wave**

Avia M. Dunn, JD, MBA, *Partner, Life Sciences, Skadden; Washington, DC*

Stewart Kameen, JD, *Partner, Davis Wright Tremaine; Former Senior Counsel, Industry Guidance Branch, Office of Counsel to the Inspector General, US Department of Health and Human Services, Washington, DC*

3:15 pm **Networking Break**

3:45 pm **Hotline Isn't Enough: Building a Smart Investigations Ecosystem**

Michael Clarke, JD, CCEP, NACD.DC®, *Founding Principal, Jaeger-Plymouth Advisors; Retired Vice President, Deputy General Counsel & Global Chief Compliance Officer, ConvaTec; Former Vice President, Corporate Compliance, Indivior Inc.; Former Medical Device Executive Committee; Bridgewater, NJ*

Mona Peterson Rosow, JD, MPH, *Former Vice President, Chief Risk, Ethics & Compliance Officer, Mozarc Medical; Former Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Steering Committee)*

Saraswatha Lalitha Putcha, MA, *Chief Operations Officer, Cresen Solutions; Philadelphia, PA (Moderator)*

4:30 pm **Anatomy of an Internal Investigation**

David J. Derusha, JD, *Assistant General Counsel, Vertex Pharmaceuticals; Former Assistant US Attorney, US Attorney's Offices, District of Massachusetts, US Department of Justice; Boston, MA*

Naveen Ganesh, MA, JD, *Vice President, Global Head of Investigations, Ethics & Compliance, Takeda; Former Head of Investigations & Monitoring, North America & Global Specialty Care, Sanofi; Former Compliance Assurance & Investigations, Lead, Verily; Boston, MA*

Sarah L. Whipple, JD, *Principal, WhipEdge Solutions, Member, Rare Disease Compliance Consortium (RDCC); Former Vice President, Global Chief Compliance Officer, Apellis; Former Vice President, Legal & Chief Compliance Officer, Akebia; Needham, MA*

Amanda P. Masselam Strachan, JD, *Partner, WilmerHale; Former Chief, Criminal Division, US Attorney's Office for the District of Massachusetts; Former Assistant US Attorney, US Attorneys' Offices, US Department of Justice; Boston, MA (Moderator)*

5:15 pm **Managing Whistleblower Risk and Internal Escalation**

Jennifer Beaudoin, CFE, CPA, *Executive Director, Head of Global Compliance Program Strategy, Governance and Transformation, Vertex Pharmaceuticals; Boston, MA*

Warren T. Allen II, MPP, JD, *Principal, Miles & Stockbridge; Professorial Lecturer in Law (White Collar Crime), The George Washington University Law School; Washington, DC (Co-Moderator)*

David Amendola, *Director, Life Sciences, Resolution Economics; Former Global Compliance Risk, Evaluation & Management, ConvaTec; New York, NY (Co-Moderator)*

5:45 pm **Adjournment**

Focused Session 2: Pharmaceutical and Medical Device Advanced Privacy and Security/ Cybersecurity Updates

1:00 pm **Chair's Welcome, Overview and Introductions**

Mary Devlin Capizzi, JD, MBA, *Partner, Faegre Drinker; General Counsel, International Pharmaceutical and Medical Device Privacy Consortium; Washington, DC (Chair)*

1:15 pm **The Basics of Healthcare Privacy for Compliance Professionals and an Overview of Privacy Issues Unique to Pharmaceutical and Medical Device Enterprises**

Adam Greene, JD, MPH, *Partner and Co-chair, Health Information & HIPAA Practice, Davis Wright Tremaine; Former Senior Health Information Technology and Privacy Specialist, Office for Civil Rights, US Department of Health & Human Services, Washington, DC*

1:45 pm **IAPP Presents: Privacy, AI Compliance, and Beyond for the Future of Pharma and Medical Devices**

Melissa Bianchi, JD, *Partner and Leader, Firm's Digital Health Initiative, Hogan Lovells Cadwalader; Washington, DC*

Patrice Ettinger, JD, CIPP/US, *Vice President, Chief Privacy Officer, Regeneron; Former Chief Privacy Officer, Pfizer; Former Chief Privacy Officer, Avon; Past Chair, International Association of Privacy Professionals; New York, NY*

Gabe Maldoff, JD, *Partner, Data, Privacy, and Cybersecurity practice, Goodwin Procter; Adjunct Professor, Maine Law School, Washington, DC*

Cobun Zweifel-Keegan, JD, CIPP/US, CIPM, *Managing Director, IAPP; Former Deputy Director, Privacy Initiatives, Better Business Bureau; Washington DC (Moderator)*

2:30 pm **The International Pharmaceutical and Medical Device Privacy Consortium Presents Advanced Strategies for Addressing Privacy and Cybersecurity Issues for Life Sciences Companies**

Kimberly J. Gold, JD, *Chief Privacy Officer, Senior Associate General Counsel and Executive Director, Privacy Law, Genentech; Former Privacy Secondee, Pfizer; San Francisco, CA*

Tahvia Jenkins, JD, CIPP/US, *US Privacy Officer & Counsel, Sanofi; Former Assistant Counsel, Health Affairs, Privacy, and Data Protection, University of California, Cambridge, MA*

Danielle Manner, JD, *Chief Privacy Officer and Senior Associate General Counsel, Gilead; Former Senior Privacy Counsel, Pfizer; Chicago, IL*

Mary Devlin Capizzi, JD, MBA, *Partner, Faegre Drinker; General Counsel, International Pharmaceutical and Medical Device Privacy Consortium; Washington, DC (Moderator)*

3:15 pm **Companies from the Cybersecurity Working Group, Health Sector Coordinating Council**

Gregory T. Garcia, *Executive Director, Cybersecurity Working Group, Health Sector Coordinating Council; Former Assistant Secretary for Cybersecurity, US Department of Homeland Security; Washington, DC*

3:45 pm **Networking Break**

4:15 pm **Cybersecurity Best Practices for Pharma and Medical Device**

David J. Scott, MS, *Managing Director, Forensic & Integrity Services, EY; Former Assistant Director, Counterterrorism, and Head, Operations Branch, Cyber Division, Director, National Cyber Investigative Joint Task Force, Federal Bureau of Investigation; Leesburg, VA*

5:00 pm **Session Faculty Q&A**

Jamie E. Darch, JD, *Partner, HIPAA and State Data Privacy & Cybersecurity Law, Ropes & Gray; Chicago, IL (Moderator)*

5:45 pm **Adjournment**

Focused Session 3: Managing Medical-Commercial Interactions: Strategic Collaboration, Evolving Roles & Risk-Based Governance

1:00 pm **Co-Chair's Introductions & Session Overview**

Peter Agnoletto, CPA, *Compliance Officer, General Medicines and Consumer Health Care, Sanofi; Former Senior Director, Chief Compliance Officer & Chief Audit Executive & ERM Leader, Par Pharmaceutical Companies; Bridgewater, NJ (Co-Chair)*

Jillian A. Gillespie, MBA, *Senior Director, Compliance, Medical & Medical Affairs, Pfizer; Former Director, Compliance Business Partner, R&D, Medical Affairs, HEOR; Chicago, IL (Co-Chair)*

1:15 pm **Strategic Planning Together, Tactical Execution Separately: Practical Case Studies in Medical–Commercial Collaboration**

Jillian A. Gillespie, MBA, *Senior Director, Compliance, Medical & Medical Affairs, Pfizer; Former Director, Compliance Business Partner, R&D, Medical Affairs, HEOR; Chicago, IL*

Samantha Brabant Philis, *Director, E&C Policy, Training, and Business Operations, Jazz Pharmaceuticals; Former North America Compliance Manager, Cushman & Wakefield; Chicago, IL*

Samantha Sutherland, *Director, Life Sciences Consulting Practice, Baker Tilly; Jersey City, NJ (Moderator)*

2:00 pm **Beyond Sales Representatives and MSLs: Understanding Emerging Field-Based Roles and Governance Challenges**

Sri Burra, MBA, *Director, Ethics and Compliance, Organon US; Former Ethics & Compliance Lead, North America, Haleon; Former Transformation Lead, Ethics & Compliance Workstream GSK; Malvern, PA*

Amy Wilson, MSJ, *Vice President, Chief Compliance Officer, Esperion; Former Head of U.S. Compliance, MorphoSys; Former Director, Enterprise Compliance, Alexion Pharmaceuticals; Boston, MA*

Matt Chandler, JD, *Director, Epsilon Life Sciences; Richmond, VA (Moderator)*

Session continued next page

Focused Session 3 *continued*

2:45 pm **Networking Break**

3:15 pm **Protecting Medical Independence While Enabling Strategic Cross-Functional Collaboration**

Peter Lee, JD, LLM, *Chief Compliance Officer, Kura Oncology; Former Chief Compliance Officer, Mirati Therapeutics; Former Chief Compliance Officer, Heron Therapeutics; San Diego, CA*

Stefanie A. Doebler, JD, *Partner and Co-Chair, Health Care Practice Group, Covington & Burling; Washington, DC (Co-Moderator)*

Sarah A. Franklin, JD, *Partner and Vice-Chair, Life Sciences Investigations Practice, Covington & Burling; Former Attorney, US Federal Trade Commission; Washington, DC (Co-Moderator)*

4:00 pm **Medical–Commercial Boundaries: Congresses, Sponsorships & Promotional Risk**

Maureen Cawley, JD, *Senior Director, Corporate Counsel, Incyte; Former Global Agreements Attorney, Merck; Philadelphia, PA*

Akosua Tuffuor, JD, *Corporate Counsel, US General Medicines, Sanofi; Morristown, NJ/Washington, DC*

Achsah Warren, MSJ, *Vice President, Head of Compliance, Sumitomo Pharma America; Former Vice President Compliance, Urovant Sciences; Former Senior Compliance Manager, Allergan, Irvine, CA*

Mahnu V. Davar, MA, JD, *Partner and Co-chair, Life Sciences & Healthcare Regulatory, Arnold & Porter; Washington, DC (Moderator)*

4:45 pm **Have We Become Too Restrictive? Rethinking Medical–Commercial Governance for the Next Era**

Giulia Del Cane, MBA; *Ethics & Business Integrity Strategic Advisor, Global and US Specialty Care, Sanofi; Boston, MA*

Peter Agnoletto, CPA, *Compliance Officer, General Medicines and Consumer Health Care, Sanofi; Former Senior Director, Chief Compliance Officer & Chief Audit Executive & ERM Leader, Par Pharmaceutical Companies, Bridgewater, NJ (Co-Chair)*

Jillian A. Gillespie, MBA, *Senior Director, Compliance, Medical & Medical Affairs, Pfizer; Former Director, Compliance Business Partner, R&D, Medical Affairs, HEOR; Chicago, IL (Co-Chair)*

5:30 pm **Adjournment**

Focused Session 4: AI in Action: Real-World Compliance Case Studies from the Front Lines of AI Use in Life Science Compliance Departments

1:00 pm **Chair's Welcome and Introductions and AI Basics, Life Science Industry Ethics and Compliance Applications**

John Poulin, *Chief Technology Officer and Partner, HELIO; Boston, MA (Chair)*

1:30 pm **A Brief Overview of the Epsilon & qordata AI Use Compliance Survey**

Casey J. Horton, CFE, *Managing Director, Epsilon Life Sciences; Former Director, Compliance Operations, AbbVie; Chicago, IL*

Mohammad Ovais, *Founder & Board Member, qordata; Princeton, NJ*

2:00 pm **Facilitated Case Study Discussion: From Transcript to Insights: How Compliance Teams Are Using AI to Transform Investigations**

Briana Cabrera, MS, CFE, *Senior Director, Global Privacy & Compliance, Investigations, Medtronic, Miami, FL*

Alexis Shaw, *Vice President, Head of Healthcare Compliance, Paratek Pharmaceuticals; Former Compliance Officer, Endo Pharmaceuticals; Former Global Compliance Investigations Manager, Boston Scientific; King of Prussia, PA*

Minna Bak, MBA, *Managing Director, HELIO; Boston, MA (Moderator)*

2:30 pm **Facilitated Case Study Discussion: AI Governing AI: Leveraging Artificial Intelligence to Develop Effective Governance Policies**

Marla Izbicky, JD, *Senior Director, Senior Corporate Counsel, Harmony Biosciences; Chicago, IL*

Neeraj Gupta, MBA, *President, Chief Technology Officer, Cresen Solutions; Chester Springs, PA (Moderator)*

3:00 pm **Networking Break**

3:15 pm **Facilitated Case Study Discussion: AI-Assisted Monitoring and Detection of Potential Policy Violations**

Ron Strauss, *Vice President Compliance, MiMedx; Former Sales Leader, Osteo Specialty, Eli Lilly and Company; Marietta, GA*

John Poulin, *Chief Technology Officer and Partner, HELIO; Boston, MA (Moderator)*

3:45 pm **Facilitated Case Study Discussion: Agentic AI Use for Continuous Risk Assessment (vs Annual)**

Matthew Stasiunas, PMP, CCEP, LPEC, *RAMP & Project Management Lead, Indivior; Former Senior Specialist, Integrity and Compliance, Kaléo; Richmond, VA*

Michael L. Shaw, JD, *Global Head of Compliance, Privacy and Risk, ZS; Former Vice President and Compliance Officer, GSK; Former Senior Counsel, Office of Inspector General, US Department of Health and Human Services; Princeton, NJ (Moderator)*

4:15 pm **The Unspoken Risk of AI: Moral Outsourcing — and Why "Add Human Judgment" Isn't Enough**

Benjamin Correa, JD, *Partner, Food, Drug & Medical Device, Sidley Austin; Washington, DC*

Stacy E. Yeung, JD, *Vice President, General Counsel and Corporate Secretary, SELLAS Life Sciences Group; Former Assistant General Counsel, Arena Pharmaceuticals; Former Deputy General Counsel, DBV Technologies; New York, NY*

L. Stephan Vincze, JD, LLM, MBA, *President and Chief Executive Officer, TRESTLE Compliance; Former Senior Vice President and Chief Compliance Officer, Warner Chilcot; Former Vice President, Ethics and Compliance Officer, TAP Pharmaceutical; Boston, MA (Moderator)*

Session continued next page

Focused Session 4 *continued*

5:00 pm **The Health Care Compliance Association Presents: Advanced Strategies in Using AI to Monitor and Improve Compliance: Lessons from Other Healthcare Sectors**

Joan Podleski, CCEP, CHPC, CHRC, CHC, *Vice President & Chief Privacy Officer, Children's Health System of Texas; Former Director, Institutional Ethics & Compliance Program, Duke University; Dallas, TX*

R. Brett Short, CHC, CHPC, CHRC, *Chief Compliance Officer and Chief Privacy Officer, University of Kentucky; Board & Immediate Past Chair, Health Care Compliance Association; Lexington, KY*

Betsy Wade, MPH, CHC, CHPC, CNA, *President, eitic healthcare compliance solutions; Faculty, Health Care Compliance Association (HCCA) Basic Compliance Academy and Privacy Academy, HCCA; Former Chief Compliance & Ethics Officer, Signature Healthcare; Louisville, KY*

5:45 pm **Adjournment**

Focused Session 5: GP 101: A Primer on the Fundamentals of Government Pricing for Compliance Officers & The Evolving Pricing and Contracting Landscape: What Compliance Officers Need to Know

1:00 pm **Chair's Introductions & Session Overview**

Chris Cobourn, MS, *Managing Director Government Programs Practice Lead, HELIO; Placida, FL (Chair)*

1:15 pm **GP 101: A Primer on the Fundamentals of Government Pricing for Compliance Officers**

Tracy Mumpower, *Director Managed Services, Life Science Revenue Management, Government Pricing & Commercial Contacting, HELIO; Former Manager, Commercial Contract Administration, King Pharmaceuticals; Bristol, VA*

John Shakow, JD, *Pharmaceutical Government Price Reporting Expert; Partner, FDA & Life Sciences Practice, King & Spalding; Washington, DC*

Chris Cobourn, MS, *Managing Director Government Programs Practice Lead, HELIO; Placida, FL (Moderator)*

3:30 pm **Networking Break**

4:00 pm **The Evolving Pricing and Contracting Landscape: What Compliance Officers Need to Know**

Kristie Champlin Gurley, JD, *Partner, Covington & Burling; Former Counsel, Office of the General Counsel, CMS Division, US Department of Health and Human Services; Washington, DC*

Kristin M. Hicks, JD, *Partner, Life Sciences & Healthcare Regulatory, Arnold & Porter; Washington, DC*

Tracy Mumpower, *Director Managed Services, Life Science Revenue Management, Government Pricing & Commercial Contacting, HELIO; Former Manager, Commercial Contract Administration, King Pharmaceuticals; Bristol, VA*

Stephanie Trunk, JD, *Partner, Life Science Practice Group, ArentFox Schiff; Former Legal Counsel, Catalyst Health Solutions; Washington, DC*

Chris Cobourn, MS, *Managing Director Government Programs Practice Lead, HELIO; Placida, FL (Moderator)*

5:30 pm **Adjournment**

WEDNESDAY, OCTOBER 14, 2026

PHARMACEUTICAL AND MEDICAL DEVICE ETHICS AND COMPLIANCE CONGRESS DAY 2

7:00 am **Registration Opens & Networking Breakfast in the Exhibit Hall**

MORNING MINI SUMMITS: GROUP 1 (8:00 am - 8:50 am)

Mini Summit 1: Compliance Independence in The Real World: Governance Models That Work

What does compliance independence look like in day-to-day decisions? This discussion will explore governance models that give Compliance a meaningful voice while supporting effective collaboration with Legal and business teams. Panelists will consider reporting relationships, decision-making authority, escalation paths, and practical ways to address disagreements when priorities compete.

8:00 am **Introductions, Discussions and Q&A**

Kathy Johnson, JD, *Vice President, Commercial and Development Legal, Vertex Pharmaceuticals; Former US Legal Counsel Consultant, Takeda; Former Senior Director, Legal Counsel, Shire; Boston, MA*

Stefanie A. Doebler, JD, *Partner and Co-Chair, Health Care Practice Group, Covington & Burling; Washington, DC (Co-Moderator)*

Sarah A. Franklin, JD, *Partner and Vice-Chair, Life Sciences Investigations Practice, Covington & Burling; Former Attorney, US Federal Trade Commission; Washington, DC (Co-Moderator)*

Mini Summit 2: The Agentic Compliance Function of the Future

As life sciences organizations adopt AI, automation, and agentic technologies across the enterprise, compliance must evolve beyond traditional oversight models. This session will explore how an AI-enabled operating model can transform compliance from a reactive control function into a proactive, scalable capability that strengthens trust, accountability, and responsible innovation.

Attendees will gain insights into:

- Embedding "Trust by Design" into AI-enabled processes and digital initiatives
- Translating regulatory expectations into practical guardrails and governance structures
- Using agentic monitoring and AI-native surveillance for continuous oversight and anomaly detection
- Designing scalable human-in-the-loop oversight for AI-assisted and autonomous activities
- Integrating risk intelligence across compliance, operational, cyber, privacy, and strategic risk domains

The session will also address implementation considerations, organizational design, governance challenges, and emerging practices. Participants will leave with actionable perspectives on using agentic technologies and AI-enabled oversight to make compliance a more strategic enabler of innovation, trust, and enterprise performance.

8:00 am **Introductions, Discussions and Q&A**

Jennifer Beaudoin, CFE, CPA, *Executive Director, Head of Global Compliance Program Strategy, Governance and Transformation, Vertex Pharmaceuticals; Boston, MA*

Emily Cunningham, *Associate Director, Compliance, Ultragenyx; New York, NY*

Lauren H. Thal, *Vice President, Compliance, Tolmar; Former Compliance Officer, Cerapedics; Former Senior Director of Compliance, Beta Bionics; Chicago, IL*

Emily (Jaffe) Singh, *Partner, Global Spend Transparency Practice Leader, PwC; Chicago, IL (Moderator)*

Mini Summit 3: Evolution of Patient Finding Activities

As therapies become more targeted and bespoke, particularly in the treatment of rare diseases, the accurate identification of appropriate patients has become a growing priority for manufacturers. Sponsored genetic testing programs are among the more widely adopted approaches to support diagnosis and patient identification, but additional patient finding strategies are emerging rapidly. This panel will explore new and evolving patient identification activities and examine the legal, compliance, and privacy considerations they may raise.

8:00 am **Introductions, Discussions and Q&A**

Betania Glorio, *Former Senior Vice President, Chief Compliance Officer, Avidity Biosciences; Former Chief Compliance Officer and Head of Group Compliance and Data Privacy, EMD Serono; Former Global Head of Compliance, Healthcare, Merck KGaA Boston, MA*

Jesse Siegel, JD, *Head of Business Ethics, Americas, Ipsen; Former Senior Counsel, Neurology & Immunology, EMD Serono; Cambridge, MA*

Sara K. Frank, JD, *Principal, Choate Hall & Stewart; Boston, MA (Moderator)*

Mini Summit 4: First Product Launch: Policy Development Strategies and Techniques

A forward-looking discussion on how geopolitical developments are influencing enforcement priorities and how compliance programs should be structured today to withstand future DOJ scrutiny (e.g., 2028–2032 outlook). The session would also explore the balance between conduct and cooperation, and the importance of documentation and strategic planning.

8:00 am **Introductions, Discussions and Q&A**

Michael Morgan, MBA, CCEP, *Global Compliance Officer, Telix Pharmaceuticals (US); Member, Business Ethics Indiana; Former Ethics and Compliance Officer-North America, Corteva; Indianapolis, IN*

Sandra Park, JD, *Senior Vice President, Legal & Chief Compliance Officer, Vera Therapeutics; Former General Counsel & Chief Compliance Officer, IDEOlogy Health; Former Vice President, Commercial Legal, Kite, a Gilead Company; Brisbane, CA*

Hannah Putnam, MHA, CHC, CHRC, *Head of Ethics & Compliance Operations, Oncology, Takeda; Former Senior Director, Ethics & Compliance, Veloxis; Former Senior Corporate Ethics & Compliance Officer, Fresenius; Cambridge, MA*

Yogesh Bahl, MBA, *Partner and Practice Leader, Resolution Life Sciences and Healthcare; Chief Financial Officer and Head of Investor Relations, IACTA Pharmaceuticals; New York, NY (Moderator)*

Mini Summit 5: Speaker Programs in 2026: Practical Legal and Compliance Considerations for Executing from Beginning to End Amid Ongoing Scrutiny

1. What patterns can be observed and lessons learned from continued government scrutiny, enforcement activity, and evolving expectations? What are some key legal and compliance themes?
2. How can compliance, legal, and regulatory teams work with their colleagues to implement compliance standards and strategies?
3. What types of challenges can arise and how can they be addressed?

8:00 am **Introductions, Discussions and Q&A**

Sharon R. Delshad, JD, *Senior Director, Legal-Compliance Counsel, Travere Therapeutics; San Diego, CA*

Tracey Hannon, JD, *Director, Compliance Business Partner, Alexion Pharmaceuticals; Former Compliance Counsel, Ionis Pharmaceuticals; Boston, MA*

Lauren Ann Pond, JD, *Assistant General Counsel, BridgeBio; Former Commercial Counsel, Biogen; Former Counsel, Takeda; Palo Alto, CA*

Noah C. Goldstein, JD, *Counsel, Pharma, Biotech & Medical Device Legal, Regulatory, & Compliance, Porzio Bromberg & Newman; Westborough, MA (Moderator)*

8:50 am **Transition Break**

MORNING MINI SUMMITS: GROUP 2 (9:00 am - 9:50 am)

Mini Summit 6: From Pronouncements to Practice: Understanding DOJ Enforcement Priorities

Join this session for a critical discussion of how DOJ's enforcement priorities are taking shape in practice. We'll share insights on enforcement trends and draw from recent matters (from the sources of these cases to the activities that were scrutinized and how the results aligned with DOJ policy) for an engaging discussion for in-house compliance and legal professionals around the practical considerations and implications of these latest developments for compliance programs.

9:00 am **Introductions, Discussions and Q&A**

Christina O. Hud, JD, *Senior Counsel, Global Investigations, Pfizer; Former Health Care Fraud Acting Unit Chief and Senior Trial Counsel at US Department of Justice, US Attorney's Office for the District of New Jersey; New York, NY*

Michelle Dineen Jerrett, JD, *Regional Head of Investigations & Monitoring, North America, Sanofi; Former Legal Director, Endoscopy, Boston Scientific; Former Assistant United States Attorney, US Attorney's Office, District of Massachusetts; Cambridge, MA*

Emily Treanor, JD, *Head of Global Investigations & Enterprise Risk Management, Alnylam Pharmaceuticals; Washington, DC*

Jane H. Yoon, JD, *Partner, Litigation Department, Paul Hastings; Former Assistant United States Attorney, Health Care and Government Fraud Unit, US Attorney's Office for the District of New Jersey; New York, NY (Moderator)*

Mini Summit 7: Boring is Risky! Want Stronger Ethics and Compliance Engagement? Train & Communicate Colorfully

Boring is your program's worst enemy. You have so many important things to share, but it all falls apart if employees are tuning you out. People do not speak up to ask questions or report concerns when they're bored, annoyed or afraid.

In this session we'll share creative, practical strategies and success stories and we'll show you real examples inspired by great TV that employees want to binge like Netflix. Tune in to learn how to:

- Change E&C's vibe to create a positive identity and brand
- Get attention and increase E&C awareness all year
- Improve learning engagement while taking up less time
- Generate influence & create E&C influencers

9:00 am **Introductions, Discussions and Q&A**

Regina Gore Cavaliere, JD, *Chief Ethics and Compliance Officer, Vice President Regulatory and Compliance, Henry Schein; Melville, NY*

Ronnie H. Feldman, *Founder, Chief Executive Officer and Creative Director, Learnings & Entertainment: Comedians Who Improve Compliance; Chicago, IL*

Mini Summit 8: Best Practices for Data Analytics

Recent enforcement actions have turned manufacturers' own speaker logs, meal records, call notes, and prescribing trends into evidence, putting a pointed question to compliance teams: can they get to the same data, and act on what it shows in time to matter? This panel moves beyond the dashboard. We'll examine how teams set analytics strategy, close the loop from signal to remediation, and govern the use of AI in monitoring and review, including where detection creates its own obligation to act. Panelists will speak to building a credible analytics function at any scale, from a one-person program to a global monitoring team.

9:00 am **Introductions, Discussions and Q&A**

Bobby Carnathan, *Senior Manager, Ethics and Compliance Business Partner, Novo Nordisk; Plainsboro, NJ*

Christian Malias, MBA, *Associate Director, Compliance Program Development, Insmed; Former Associate Director Compliance Business Partner, Sage Therapeutics; Former Global Corporate Compliance Manager, Adaptimmune; Pittsburgh, PA*

Erica Powers, CHC, *Founder & Chief Executive Officer, Ethical Paths; Former Vice President, Chief Compliance Officer, Sage Therapeutics; Former Director Corporate Compliance, Vertex Pharmaceuticals; Boston, MA (Moderator)*

Mini Summit 9: On the Front Lines: Navigating Field Force and Sales Compliance Risk

- Compliance risks associated with speaker programs, meals, and HCP interactions
- Sampling controls, documentation risks, and recent enforcement focus areas
- State law developments impacting field activities like disclosures and licensing
- Sales incentive compensation and unintended behavioral risk

9:00 am **Introductions, Discussions and Q&A**

Ron Strauss, *Vice President Compliance, MiMedx; Former Sales Leader, Osteo Specialty, Eli Lilly and Company; Marietta, GA*

Sarah L. Whipple, JD, *Principal, WhipEdge Solutions; Member, Rare Disease Compliance Consortium (RDCC); Former Vice President, Global Chief Compliance Officer, Apellis; Former Vice President, Legal & Chief Compliance Officer, Akebia; Needham, MA*

Brian Van Hoy, RPh, *Vice President of Compliance, G&M Health; Former Director, Compliance & Ethics, Eli Lilly and Company; Somerset, NJ (Moderator)*

Mini Summit 10: Communicating the Story on Integrity: What Compliance Leaders Can Learn from Health Stakeholders

Compliance leaders are asked not only to build and sustain effective integrity programs with employees, but to inform business decisions that impact external stakeholder trust. They are also increasingly asked to engage directly with external stakeholders to share and improve their company's practices, from trade associations to collective action initiatives. This session will provide practical insights to help you and your team better communicate the story of integrity when it impacts an external audience, drawing on diverse stakeholder perspectives and experiences from partnerships such as the U.S. Consensus Framework for Ethical Collaboration.

9:00 am **Introductions, Discussions and Q&A**

Paul M. Johnson, JD, *Associate Vice President, Deputy Chief Compliance Officer, Global Compliance, Amgen; Former Senior Counsel, Office of Counsel to the Inspector General, US Department of Health and Human Services; Thousand Oaks, CA*

Karen Mancera-Cuevas, DrPH, MS, MPH, *Senior Director, Research Advancement & Equity, National Health Council; Former Deputy Director, Health Promotion, Illinois Department of Public Health, Washington, DC*

Julie Ritchie Wagner, JD, *Assistant General Counsel, PhRMA; Former Senior Counsel, Office of Counsel to the Inspector General, US Department of Health and Human Services; Washington, DC*

Andrew Blasi, MBA, *Founder & Chief Executive Officer, Ethicist International; Founding Member, Anti-Corruption Leaders Hub, OECD; Faculty, International Anti-Corruption Academy; Washington, DC (Moderator)*

Mini Summit 11: Designing HCP Engagement with Transparency Built In

As regulatory scrutiny around HCP engagement increases, organizations must ensure transparency is embedded into their processes. This session explores how to design systems that capture required data and leverage AI.

9:00 am **Introductions, Discussions and Q&A**

Kaoutar Afif, *Manager, Ethics & Compliance, Cardinal Health; Dublin, OH*

Peggy Kruk, HMCC, *Compliance Manager, MiMedX; Former Director, Compliance, Program Services, Vaniam Group; Dallas, GA*

Jacob Wolf, MBA, *Chief Commercial Officer, Cresen Solutions; Orlando, FL (Moderator)*

9:50 am **Networking Break in the Exhibit Hall**

MORNING MINI SUMMITS: GROUP 3 (10:10 am – 11:00 am)

Mini Summit 12: Compliance and M&A: Why Early Engagement Matters

Compliance issues can affect deal value, integration, and post-close risk—yet compliance is often brought in too late. This session explores how early engagement helps teams identify risks sooner, make better decisions, and avoid costly surprises throughout the deal lifecycle.

10:10 am **Introductions, Discussions and Q&A**

Eliza L. Andonova, JD, *Partner, Global Regulatory, Hogan Lovells Cadwalader; Washington, DC*

Eric Nichols, JD, *Associate Vice President, Ethics & Compliance, R&D and Business Development, Eli Lilly and Company; Indianapolis, IN*

Megan Roberts, CPA, MS, *Head of Integrity & Compliance, Par Health; Former Senior Director, Integrity & Compliance; St. Louis, MO*

Heather Young, JD, *Senior Director, Compliance Officer, Global Business Partner, Olympus Corporation of the Americas; Former Assistant District Attorney, Philadelphia District Attorney's Office, Philadelphia, PA; Center Valley, PA*

Russell Rose, JD, *Managing Director, Deloitte Consulting; Dallas, TX (Moderator)*

Mini Summit 13: The State of Medical Device Compliance: Annual Industry Roundtable

Medical device compliance is evolving quickly, shaped by changing regulations, heightened enforcement attention, new technologies, and increasingly complex commercial and quality-system risks. This panel brings together experienced compliance leaders and industry experts for a candid discussion of the issues defining today's environment. This discussion will explore:

Enforcement trends: Analyze recent actions involving medical device manufacturers, the underlying compliance breakdowns, and the areas drawing increased regulatory attention.

Compliance strategies: Learn practical approaches for responding to evolving requirements and building durable, risk-based compliance frameworks aligned with industry expectations.

What's next: Hear expert perspectives on upcoming regulatory and enforcement trends so your organization can anticipate change, identify exposure early, and prepare for future compliance demands.

10:10 am **Introductions, Discussions and Q&A**

Kate Godfrey, JD, CCEP, *Senior Vice President, Global Chief Compliance Officer, Karl Storz North America; El Segundo, CA*

JJ Kuhn, JD, *Chief Ethics & Compliance Officer, Danaher; Former Senior Vice President and Chief Ethics & Compliance Officer, Solventum; Former Vice President Chief Counsel, Global Investigations, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Committee)*

Diane Seol, JD, *Of Counsel, Choate Hall & Stewart; Former Assistant United States Attorney, Affirmative Civil Enforcement Unit, US Attorney's Office, District of Massachusetts; Boston, MA*

Casey J. Horton, CFE, *Managing Director, Epsilon Life Sciences; Former Director, Compliance Operations, AbbVie; Chicago, IL (Moderator)*

Mini Summit 14: The State We're In: Legal & Enforcement Trends Reshaping Pharmaceutical Compliance

- Recent and upcoming state law developments affecting pharmaceutical manufacturers
- Trends in proposed legislation, such as A.I., disclosures, advertising, and more
- Commonly overlooked requirements creating hidden compliance exposure
- Review of recent OIG enforcement actions and Advisory Opinions impacting life sciences organization
- Key themes and patterns emerging from enforcement activity across the industry

10:10 am **Introductions, Discussions and Q&A**

Brian Van Hoy, RPh, *Vice President of Compliance, G&M Health; Former Director, Compliance & Ethics, Eli Lilly and Company; Somerset, NJ*

Olivia Krzeminski, JD, *Director of Compliance & Legal Research, G&M Health; Somerset, NJ (Moderator)*

Mini Summit 15: External Funding Risks and Mitigation: Latest Trends in Exhibit & Display and Sponsorship Requests

Exhibit, display, and sponsorship requests can offer legitimate opportunities for engagement while presenting evolving compliance, legal, and reputational risks. This session will examine current trends and practical approaches for evaluating these requests consistently and appropriately.

Panelists will discuss key considerations, including establishing a legitimate business purpose, assessing value and benefits, applying fair market value principles, maintaining appropriate separation from commercial influence, and documenting funding decisions. The session will also explore common red flags, the role of cross-functional review, and strategies for developing clear, risk-based approval standards. Attendees will gain practical insights to help strengthen oversight while supporting appropriate external engagement.

10:10 am **Introductions, Discussions and Q&A**

Asha Green, JD, MPA, CCEP, CHC, *Director, US Compliance, Ferring; Former Global Head, I&C Risk Assessment and Mitigation Planning and Senior Business Advisor, Indivior; Parsippany, NJ*

Alison J. Page, JD, *Assistant General Counsel, Oncology and US Market Access & Patient Services Compliance Lead, Pfizer; New York, NY*

Samantha Barrett Badlam, JD, *Partner, Litigation & Enforcement Group, Ropes & Gray; Washington, DC (Moderator)*

Mini Summit 16: Beyond the Questionnaire: Building a Risk-Based Third-Party Management Program That Actually Works

Third party risk management too often calcifies into a check-the-box exercise with a questionnaire at onboarding, then silence until renewal. Meanwhile, real risk emerges dynamically: ownership changes, sanctions designations, sub-distributors operating three tiers beyond visibility. This panel examines how leading life science companies are rebuilding their third-party risk management around how risk actually behaves. Panelists will share practical approaches to meaningful risk-tiering, the shift from point-in-time diligence to continuous monitoring, and where AI-driven screening ends and human intelligence must begin.

10:10 am **Introductions, Discussions and Q&A**

Ariadna Quesada, MSJ, *Vice President of Compliance - Global Compliance & Privacy lead, Zoll Medical; Former Compliance Director Europe & MEATI, Hillrom; Former Ethics and Compliance Officer, The Netherlands, AbbVie; Madrid, Spain*

Maria Villanueva, MBA, CFE, *Director, Ethics and Compliance, BioTissue; Former VP, Compliance Officer and Head of HR, Ra Medical Systems, Manufacturer of DABRA excimer laser system; Former Director, Compliance, Zimmer Biomet; Miami, FL*

Suki Wiltman, *Vice President, Global Ethics & Compliance, Ipsen; Former Business Partner, Governance, Ethics & Compliance, GSK; Former Senior Corporate Compliance Manager, Allergan Canada; Toronto, Canada*

David Fisher, MS, *President, TDI Solutions; Former Vice President & General Manager, Cyber Innovations, Battelle; Washington, DC (Moderator)*

Mini Summit 17: Navigating Today's Enforcement Landscape with Culture, Data, and Ethics

Participants will explore:

- Are traditional assumptions about enforcement risk still valid?
- When does self-disclosure make strategic sense, and when might it not?
- How can organizations measure whether compliance programs are actually working?
- What data boards and executives are demanding to make better risk decisions?
- How can companies thrive in an environment where certainty is decreasing and scrutiny remains high?

10:10 am **Introductions, Discussions and Q&A**

Hui Chen, JD, *Co-Founder, Culture. Data. Ethics; Former Compliance Counsel Expert (Consultant), US Department of Justice; Former Assistant General Counsel, Pfizer; Honolulu, HI*

Zachary Coseglia, JD, *Co-Founder, Culture. Data. Ethics; Co-Founder & Former Managing Principal & Head of Innovation R&G Insights Lab, Ropes & Gray; Former Vice President & Assistant General Counsel, Global Compliance Monitoring, Analytics and Digital, Pfizer; Boston, MA*

11:00 am **Transition Break**

MORNING MINI SUMMITS: GROUP 4 (11:10 am – 12:00 pm)

Mini Summit 18: Risk Advantage: How Resilient Organizations are Capturing Opportunity through Uncertainty

Recent enforcement actions have turned manufacturers' own speaker logs, meal records, call notes, and prescribing trends into evidence, putting a pointed question to compliance teams: can they get to the same data, and act on what it shows in time to matter? This panel moves beyond the dashboard. We'll examine how teams set analytics strategy, close the loop from signal to remediation, and govern the use of AI in monitoring and review, including where detection creates its own obligation to act. Panelists will speak to building a credible analytics function at any scale, from a one-person program to a global monitoring team.

11:10 am **Introductions, Discussions and Q&A**

Anisa Dhalla, *Vice President, Chief Ethics and Compliance Officer; Global Head, Ethics and Compliance, UCB; Atlanta, GA (PCF Board Member)*

Kate Godfrey, JD, CCEP, *Senior Vice President, Global Chief Compliance Officer, Chief Compliance Officer KSNA, Karl Storz North America; El Segundo, CA*

Russell Schaefer, MBA, *Managing Director, BCG; Washington, DC (Moderator)*

Mini Summit 19: View from the Top: Practical Advice for Accelerating Your Compliance Career

What distinguishes a strong compliance professional from a future compliance leader? In this interactive session, accomplished compliance executives will share candid, practical advice on the skills, experiences, and behaviors that can accelerate career growth in today's rapidly evolving environment. Attendees will learn how to increase their visibility, build credibility with business leaders, move from execution to strategy, and identify the next experience needed to advance their careers.

11:10 am **Introductions, Discussions and Q&A**

Terra Buckley, JD, *Senior Vice President, Chief Compliance & Ethics Officer, Bristol Myers Squibb; Former Chief Compliance Officer, LivaNova; Former Executive Director, Head, Business Advisory Center of Excellence, US Healthcare Compliance, Celgene; Summit, NJ*

Christie Camelio, *Senior Vice President and Chief Compliance Officer, Insmed; Former Vice President & Deputy Global CCO, Celgene; Florham Park, NJ (PCF Board)*

Daryl Kreml, JD, *Head, Ethics & Compliance, Global Oncology, Takeda; Former Senior Vice President, Chief Enterprise, Risk & Compliance Officer, Sage; Former US Compliance & Global Program Management, Biogen; Boston Scientific; Boston, MA (PCF Board Member)*

Daniel O'Connor, *Senior Vice President, NXLevel Compliance; Lambertville, NJ (Moderator)*

Mini Summit 20: Direct-to-Consumer (DTC) Models: Emerging Compliance Risks and Governance Considerations

As pharmaceutical and medical device companies continue to explore direct-to-consumer models, including telehealth partnerships, online prescribing platforms, patient support services, and direct distribution channels, compliance professionals are facing new and evolving risk areas. The Panel will explore these evolving models, including:

- The growing use of AI in consumer engagement models
- Current trends in patient access and patient support tools and programs
- Best practices in partnerships with healthcare providers

The Panel will also address the compliance risks posed by these models and how to best address them, including:

- Anti-Kickback Statute and False Claims Act considerations
- Patient inducement concerns
- Marketing and promotional oversight
- Data privacy and patient information considerations
- Lessons learned from recent enforcement actions and regulatory scrutiny

11:10 am **Introductions, Discussions and Q&A**

Amy Bentsen, JD, *Head of Investigations for Americas, Ethics & Compliance, Takeda; Cambridge, MA*

Michele Guerino, MS, *Executive Director, Commercial Compliance Office, Human Health Ethics and Compliance, Merck; Former Senior Regional Business Director, Cubist Pharmaceuticals; Hopewell, NJ*

Edward "Ted" B. Diskant, JD, *Partner, McDermott Will & Schulte; Former Assistant US Attorney, Southern District of New York, US Department of Justice; New York, NY (Moderator)*

Mini Summit 21: Fair Market Value: Practical Strategies and Targeted Insights for Building Defensible FMV Frameworks

Building and deploying a consistent fair market value (FMV) program remains a cornerstone of compliance programs across the life sciences industry. Designed for professionals at all experience levels, this discussion will cover both foundational concepts and new challenges, including how to assess FMV for non-traditional stakeholders, ensure appropriate valuation for travel and related expenses, navigate some common pitfalls of operating in global markets, leverage data and new technology while maintaining defensibility, and more! Our expert panel of esteemed compliance professionals will share practical insights, lessons learned, and strategies to help organizations build adaptable, risk-based FMV frameworks that keep pace with innovation across the industry.

11:10 am **Introductions, Discussions and Q&A**

Yelena Ferreira, JD, *Head of Compliance and Assistant General Counsel, Pharmacosmos Therapeutics; Former Senior Director, Associate General Counsel, Pacira BioSciences; Morristown, NJ*

Abe Kassis, MBA, PhD, *Global Head Compliance and Legal Operations, Bausch + Lomb; Former Director of Corporate Compliance (Ethics & Compliance); Former Global Compliance, Director of Compliance, Takeda; New York, NY*

Eric Bolesh, *Chief Executive Officer, Cutting Edge Information; Research Triangle Park, NC (Moderator)*

Mini Summit 22: Navigating the Evolving US Privacy Landscape

As state privacy laws continue to multiply, understanding the common principles and requirements across these laws has never been more important. In this interactive session, our panelists will provide their insights and concerns regarding risks and compliance demands. Through real-world scenarios, attendees will learn how leading organizations prioritize compliance efforts and navigate the continuously evolving privacy landscape.

11:10 am **Introductions, Discussions and Q&A**

Simonne Brousseau, JD, *Associate, Faegre Drinker; Washington, DC*

Tom Hiney, CIPP/US, CIPM, JD, *Head, Digital Ethics & Compliance, US Business Unit, Takeda; Former Privacy Counsel, Blueprint Medicines; Boston, MA*

Courtney Lyles, *Manager, Privacy and Data Protection, Dovetail Consulting Group, Atlanta, GA*

Erin Wagenberg, CIPP/US, CIPP/E, CIPT, FIP, *Director, Dovetail Consulting Group; Former Deputy Privacy Officer, Deloitte Global; Atlanta, GA (Moderator)*

Mini Summit 23: Agentic Risk Management: Transforming Compliance Across Key Business Processes

Agentic AI is redefining the compliance operating model. Compliance teams are managing more signals than ever, often without a clear way to act on them. By the time a violation is confirmed, the moment to prevent it has usually already passed. This panel makes the case for a different model: AI embedded directly into the workflow, flagging and guiding risk in real time, before it becomes a problem. Topics include:

- The hidden risk of over-engineered controls, which create friction that drives workarounds you can't see
- Catching speaker program risk in real time instead of weeks after the event closes
- Why reviewing a monitoring/audit sample can look clean while most of the program goes unseen
- As field tools move toward open, free-text entry, how compliance oversight must evolve to keep pace
- What audit-ready documentation looks like when an AI agent takes the action instead of a person

Panelists will share a practical framework for identifying which compliance processes are ready for agentic transformation, along with what should be in place before that shift begins.

11:10 am **Introductions, Discussions and Q&A**

Kevin Espinoza, *Chief Integrity & Compliance Officer, Indivior; Former Chief Integrity & Compliance Officer, Kaléo; Former Senior Vice President, Corporate Compliance, Boston Scientific; Richmond, VA*

Jeanille Gatta, MA, *Director, Compliance Business Partner, AstraZeneca, North America; Philadelphia, PA*

Stephanie Santana, MPH, PMP, *Associate Principal, ZS; New York, NY (Co-Moderator)*

Michael L. Shaw, JD, *Global Head of Compliance, Privacy and Risk, ZS; Former Vice President and Compliance Officer, GSK; Former Senior Counsel, Office of Inspector General, US Department of Health and Human Services; Princeton, NJ (Co-Moderator)*

12:00 pm **Luncheon Activities**

LUNCHEON ACTIVITIES

Choose between two lunch activities:

12:00 pm – 1:15 pm **Networking Lunch in the Exhibit Hall**
(Pick up your box lunch in the exhibit hall.)

—— OR ——

12:10 pm – 1:10 pm **Optional Luncheon Discussion:**

Compliance Risks and Challenges in Small and Emerging Life Sciences Companies

Grab your box lunch in the exhibit hall and join an interactive discussion designed for emerging and smaller life sciences companies with limited or no existing compliance program infrastructure. This practical, risk-based session will help compliance leaders determine where to begin, which risks and program elements to prioritize, and how to build an effective compliance program as the organization grows.

The discussion will address questions such as:

- Which risks should be prioritized first?
- When should key compliance program elements and control be implemented?
- How do you build a roadmap that aligns with the company's stage of growth?
- How do you maximize limited resources while establishing an effective compliance program?

Discussion Facilitators:

Greg Demske, JD, *Partner, Goodwin Procter; Former Chief Counsel, Office of Inspector General, US Department of Health and Human Services; Washington, DC*

Jeffrey Kawalek, MBA, *Chief Compliance Officer, Zambon US; Former Deputy Chief Compliance Officer US, Jazz Pharmaceuticals; New York, NY (Former PCF Board Chair)*

PHARMACEUTICAL AND MEDICAL DEVICE CONGRESS OPENING PLENARY SESSION

1:15 pm **Welcome and Introduction: PCF Board**

Donna White, CCEP, *Vice President, Compliance Officer, America Region, Chiesi USA; Cary, NC (PCF Board Chair)*

1:30 pm **C-Suite Congress Video Welcome**

Albert Bourla, DVM, PhD, *Chairman and Chief Executive Officer, Pfizer; Co-Chair, Partnership for New York City; Former Chair, Pharmaceutical Research and Manufacturers of America (PhRMA); Former President, International Federation of Pharmaceutical Manufacturers and Associations (IFPMA); New York, NY*

1:40 pm **Keynote Annual OIG Update**

Mary E. Riordan, JD, *Senior Counsel, Office of Counsel to the Inspector General, Office of Inspector General, Department of Health and Human Services; Washington, DC*

2:30 pm **Keynote US DOJ Criminal Division Update Fireside Chat**

Jacob Foster, JD, *Acting Chief, Health Care Fraud (HCF) Unit, Criminal Division, Fraud Section, US Department of Justice (DOJ); Former Trial Advocacy Project Deputy District Attorney, Los Angeles County District Attorney's Office; Washington, DC*

Gary F. Giampetruzzi, JD, *Partner & Global Chair Life Sciences, Paul Hastings; Former Vice President and Acting General Counsel, Head of Government Investigations, Pfizer; New York, NY (Moderator)*

3:00 pm **Tech & Human-in-the-Loop Compliance (Why Human Judgment Matters More Than Ever)**

Jason DiPietro, *Associate Director, Compliance Innovation & Analytics, Lunbeck; Deerfield, IL*

Rawan Fusisi, *Head, Transparency and Third Party Compliance, Indivior; Former Transparency and Analytics Manager, Stryker; Washington, DC*

Rajiv Mehrotra, MS, *Senior Vice President, Data Science, HELIO; New York, NY*

Marci Juneau, MBA, *A Founding Partner, HELIO, Atlanta, GA (Moderator)*

3:30 pm **Networking Break in the Exhibit Hall**

Mona Peterson Rosow, JD, MPH, *Former Vice President, Chief Risk, Ethics & Compliance Officer, Mozarc Medical; Former Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Steering Committee)*

4:00 pm **The Next Generation of Compliance: Leveraging Data to Stay Ahead of Risk**

Lindsey LaMoria Gavin, *Senior Director, Deputy Compliance Officer, Lundbeck; Former Compliance Director, Meetings & Events International; Deerfield, IL*

Al Park, *Senior Compliance Officer, Global Monitoring, Auditing and Third-Party Risk Management, Global Ethics and Compliance, Jazz Pharmaceuticals; Former Global Program Effectiveness & Insights Leader, Worldwide Compliance & Business Ethics, Amgen; Phoenix, AZ*

Robert Melillo, JD, *Co-founder and Managing Director, G&M Health; Cocoa Beach, FL (Moderator)*

4:30 pm **Integrated Risk Transformation in Pharma/Medical Device: From Compliance Burden to Strategic Advantage**

Michael Reyes, *Director, Risk & Regulatory Consulting, PwC; Former Analyst, Compliance Analytics, Edwards Lifesciences; Dallas, TX*

Brian Riewerts, MPA, *Partner, Pharmaceutical and Life Sciences Risk and Regulatory Leader, PwC; Washington, DC (Moderator)*

5:00 pm **Annual Chief Compliance Officers Fireside Chat**

Terra Buckley, JD, *Chief Compliance & Ethics Officer, Bristol Myers Squibb; Former Chief Compliance Officer, LivaNova; Former Executive Director, Head, Business Advisory Center of Excellence, US Healthcare Compliance, Celgene; Summit, NJ*

Christine Gordon, JD, *Vice President, Risk, Assurance, & Compliance and Chief Compliance Officer, RAC Business Partner, Global SIS Division, Olympus Corporation of the Americas; Former Assistant District Attorney, Philadelphia District Attorney's Office; Bethlehem, PA*

Melissa T. Lozner, JD, *Senior Vice President, Chief Compliance Officer, Regeneron; Former Chief Compliance & Ethics Officer and Special Counsel, Rafael Holdings; Former Vice President, Head of Ethics, Risk & Compliance, Novartis; Tarrytown, NY (PCF Board Member)*

Paul J. Silver, *Principal, Life Sciences Regulatory, Compliance, and Legal Leader, Deloitte Consulting, Atlanta, GA (Moderator)*

5:45 pm **ADJOURNMENT**

Melissa T. Lozner, JD, *Senior Vice President, Chief Compliance Officer, Regeneron; Former Chief Compliance & Ethics Officer and Special Counsel, Rafael Holdings; Former Vice President, Head of Ethics, Risk & Compliance, Novartis; Tarrytown, NY (PCF Board Member)*

5:45 pm **NETWORKING RECEPTION In the Exhibit Hall**

PHARMACEUTICAL AND MEDICAL DEVICE ETHICS AND COMPLIANCE CONGRESS DAY 3

7:00 am **Registration Opens: Breakfast Buffet in the Exhibit Hall**

CHIEF COMPLIANCE OFFICER ROUNDTABLE

PCF-Sponsored, Exclusive Invitation-Only, Closed-Door session

7:00 am **Breakfast will be served in the CCO Roundtable Meeting room**

8:00 am **CCO Roundtable Welcome and Introductions**

Jill Dailey, JD, *Vice President, Chief Compliance Officer, Oncology, Menarini Stemline; Former Vice President & Chief Compliance Officer, Incyte; Former Assistant General Counsel, Asia Pacific Compliance Lead, Pfizer; New York, NY (PCF Board Member)*

Antitrust Admonition
Kristin Graham Koehler, JD, *Managing Partner, DC Office & Executive Committee Member, Sidley Austin; Washington, DC*

8:15 am **Future of the Compliance Profession in an AI Driven Environment**

AI is reshaping how compliance functions operate, how teams are structured, and what skills tomorrow's leaders must bring to the table. This panel brings together organizations of varying sizes to explore how AI is changing expectations, budgets, staffing models, and the competencies required for the next generation of compliance leadership.

The conversation will focus on practical realities: what companies are doing, where pressures are emerging, and how to prepare talent for a future where AI is embedded in every workflow.

Kevin Espinoza, *Chief Integrity & Compliance Officer, Indivior; Former Chief Integrity & Compliance Officer, Kaléo; Former Senior Vice President, Corporate Compliance, Boston Scientific; Richmond, VA*

Jim Gibney, *Vice President, Global Compliance Operations, Corporate Compliance, Regeneron; Former Director, Worldwide Programs and US Investigations, Pfizer; Ridgewood, NJ*

Rebekah Latchis, JD, *Vice President, Risk Intelligence & Investigations, Office of Privacy, Integrity & Compliance, Medtronic; Former Corporate Responsibility Director, Bon Secours; Washington, DC*

Natasha Trifun, JD, *Head of Enterprise Compliance & Risk and Privacy, AstraZeneca; Former Head of Ethics and Compliance for EUCAN/ International, R&D, and Operations for Alexion AstraZeneca Rare Disease; Former Corporate Counsel, Brazil Compliance Officer, Pfizer; Boston, MA*

Joy Dowdle, JD, *Partner, Litigation Department, Paul Hastings and Co-Chair of the Houston Office; Houston, TX (Moderator)*

9:00 am **Executive Breakout by Company Size - *Leveraging AI to Support Compliance Programs***

Participants will dig into practical, real-world AI implementation, sharing how AI is actually being used inside their organizations—what's working, what's stalled, and where the biggest opportunities and risks are beginning to emerge.

Daniel Spicehandler, JD, *Vice President Compliance, MSNT and Global Programs, Stryker; New York, NY (Pharma Congress Medical Device Executive Steering Committee)*

9:40 am **Networking Break**

10:00 am **Executive Breakout by Company Size - *Building AI Ready Teams***

Participants will share how they are upskilling their team to effectively leverage AI. Sharing how they're training current staff, recruiting future compliance talent, and building AI-ready skill sets across their programs.

Mona Peterson Rosow, JD, MPH, *Former Vice President, Chief Risk, Ethics & Compliance Officer, Mozarc Medical; Former Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Steering Committee)*

10:40 am **Board Oversight of Compliance: What Audit Firms Observe and What Regulators Require**

As regulatory expectations intensify and AI accelerates business risk, Boards are reexamining what "good" compliance looks like—and whether their organizations are structured to deliver it. Yet many Boards still misunderstand the purpose, scope, and value of Compliance, creating gaps in oversight and weakening independence.

This session offers an inside look at what audit firms are seeing in board governance and cultural attitudes toward compliance. We will also explore the government's perspective on why robust compliance programs matter, and the oversight behaviors regulators expect Boards to demonstrate in the boardroom.

Katherine Buckley, MBA, *Partner, Pharmaceutical and Life Sciences Risk & Regulatory Consulting Solutions, PwC; Philadelphia, PA*

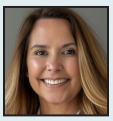
Saul B. Helman, MD, MBA, *President and Managing Director, Epsilon Life Sciences; Chicago, IL*

Tony Jordan, *Partner, EY; Former Accounting Branch Chief - Boston District Office, US Securities and Exchange Commission; Boston, MA*

Kenneth A. Polite, Jr, JD, *Partner & Global Co-leader, White Collar Defense, Sidley Austin; Former Assistant Attorney General, Criminal Division, US Department of Justice; Washington, DC*

Christie Camelio, *Senior Vice President and Chief Compliance Officer, Insmed; Former Vice President & Deputy Global CCO, Celgene; Florham Park, NJ (PCF Board) (Moderator)*

11:30 am **Open Forum: Opportunities for Benchmarking and Peer-driven Q&A. Bring your questions!**



Jill Dailey, JD, *Vice President, Chief Compliance Officer, Oncology, Menarini Stemline; Former Vice President & Chief Compliance Officer, Incyte; Former Assistant General Counsel, Asia Pacific Compliance Lead, Pfizer; New York, NY (PCF Board Member)*

Mona Peterson Rosow, JD, MPH, *Former Vice President, Chief Risk, Ethics & Compliance Officer, Mozarc Medical; Former Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Steering Committee)*

Daniel Spicehandler, JD, *Vice President Compliance, MSNT and Global Programs, Stryker; New York, NY (Pharma Congress Medical Device Executive Steering Committee)*

11:55 am **Wrap-up**

Jill Dailey, JD, *Vice President, Chief Compliance Officer, Oncology, Menarini Stemline; Former Vice President & Chief Compliance Officer, Incyte; Former Assistant General Counsel, Asia Pacific Compliance Lead, Pfizer; New York, NY (PCF Board Member)*

Noon **Chief Compliance Officer Roundtable Adjournment**

MORNING MINI SUMMITS: GROUP 5 (8:00 am - 8:50 am)

Mini Summit 24: Identifying and Managing Risk Associated with AI

As AI use expands across life sciences organizations, compliance teams must identify and manage risks without unnecessarily slowing responsible innovation. This session will explore practical approaches for:

- Evaluating and prioritizing AI use cases based on risk
- Establishing appropriate governance, accountability, and human oversight
- Addressing privacy, security, bias, accuracy, and transparency concerns
- Assessing third-party AI tools and vendors
- Monitoring AI systems as their use and capabilities evolve

Panelists will share perspectives on building a risk-based framework that supports innovation while maintaining effective compliance oversight.

8:00 am **Introductions, Discussions and Q&A**

Amy Bentsen, JD, *Head of Investigations for Americas, Ethics & Compliance, Takeda; Cambridge, MA*

Jerel Smith, MSJ, *Chief Compliance Officer-Director Integrity & Compliance, Upsher-Smith; Former Director, Compliance Operations, Karyopharm Therapeutics; Former, Director, Biopharma Compliance, Pfizer; New York, NY*

Mohammad Ovais, *Founder & Board Member, qordata; Princeton, NJ (Moderator)*

Mini Summit 25: Transparency Reporting; Update on the Centers for Medicare and Medicaid Services Open Payments Program

This session will examine recent developments in the CMS Open Payments Program and their implications for reporting entities, including:

- Trump Administration priorities and the program's realignment within CMS
- Persistent reporting challenges, including covered recipient matching
- CMS audits, data analytics, and enforcement activity—and how to respond to educational letters and investigations
- Documentation, internal auditing, error correction, and common preventable mistakes
- Frequently misinterpreted requirements, including ownership and investment interests

Panelists will share practical guidance for improving reporting accuracy, addressing identified errors, and strengthening transparency compliance.

8:00 am **Introductions, Discussions and Q&A**

Veronika Peleshchuk Fradlin, MPP, *Director, Open Payments Data Group (OPDG), Office of Enterprise Data and Analytics (OEDA), Centers for Medicare & Medicaid Services, US Department of Health and Human Services; Washington, DC*

Lisa M. Re, JD, *Partner, Life Sciences/Healthcare Regulatory Practice Group, Arnold & Porter; Former Assistant Inspector General for Legal Affairs, Office of Inspector General, US Department of Health and Human Services; Washington, DC (Moderator)*

Mini Summit 26: From Speak-Up to Proof: Turning DOJ and OIG Expectations into Defensible, AI-Enabled Compliance Operations

Regulators no longer view an effective compliance program as a set of policies on paper. They expect organizations to demonstrate, with evidence, that risks are being detected, escalated, investigated, and remediated in a consistent, measurable way. Recent DOJ guidance emphasizes the role of data, analytics, AI, and strong speak-up programs in supporting continuous risk detection.

The session will focus on practical, regulator-aligned approaches for moving from reactive case handling to proactive, evidence-based compliance management. Attendees will see how data, analytics, and AI can strengthen continuous risk detection, surface meaningful trends, and create the audit-ready records needed to demonstrate program effectiveness over time.

8:00 am **Introductions, Discussions and Q&A**

Allison Benkert, MBA, *Senior Director, Compliance Policies, Training & Investigations, Jazz Pharmaceuticals; Former Senior Director, Global Investigations, Pfizer; Former Head of Global Compliance Investigations, Seagen; Doylestown, PA*

Melissa (Missy) Kaschak, CPA, CFE, CISA, *GRC Operations, Olympus Corporations of the Americas; Center Valley, PA*

Aaron Lewis, MBA, *Director, Legal and Compliance Operations, Cordis; Former Director, Compliance Insights and Technologies, Cardinal Health; Miami Lakes, FL*

Brook Mishler, RN, MSN, *Global Director of Compliance Solutions, Life Sciences, Case IQ; Former Compliance Officer for the Americas, Cordis; Ontario, Canada (Co-Moderator)*

Shannon Walker, *Executive Vice President, Thought Leadership & Strategy, Case IQ; Founder and President, WhistleBlower Security; Ontario, Canada (Co-Moderator)*

Mini Summit 27: Manufacturer–Specialty Pharmacy Relationships: Managing Compliance Risk in Specialty and Rare Disease Commercial Models

Relationships between manufacturers and specialty pharmacies are foundational to modern specialty and rare disease commercial models, supporting patient access, adherence, and complex therapy management. At the same time, these arrangements operate in a legally sensitive space, raising recurring fraud and abuse questions around financial support, data sharing, service arrangements, and the degree of manufacturer involvement in pharmacy operations.

As manufacturers increasingly rely on specialty pharmacies to execute patient support and distribution strategies, compliance and legal teams are frequently asked to help define the appropriate level of collaboration with specialty pharmacies. This panel will examine the core compliance considerations that arise in manufacturer–specialty pharmacy relationships, focusing on practical risk identification, program design, and oversight approaches that reflect how these models function in the real world. Panelists will share perspectives on common friction points, recurring questions from internal stakeholders, and strategies for supporting patient access while maintaining appropriate compliance guardrails.

8:00 am **Introductions, Discussions and Q&A**

Elizabeth Conti, JD, *Senior Legal Counsel, Nestlé Health Science; Arlington, VA*

Rakan F. Ghubej, JD, *Senior Director, Senior Corporate Counsel & Assistant Corporate Secretary, Kyowa Kirin; New York, NY*

Aaron P. Maskery, CPA, CIA, *Senior Director, Compliance Operations, Mighty Therapeutics; Former Senior Director, Compliance, Karuna Therapeutics; Former Director, Compliance Auditing & Monitoring, Idorsia; Wilmington, DE*

Eliza L. Andonova, JD, *Partner, Global Regulatory, Hogan Lovells Cadwalader; Washington, DC (Moderator)*

Mini Summit 28: The Future of Compliance Monitoring: AI Agents, Risk Intelligence and Practical Adoption

With AI capabilities maturing rapidly, compliance leaders are evaluating how these technologies can move beyond experimentation and create measurable value within monitoring programs. Join us for a discussion on where AI is delivering value today within compliance monitoring programs, including a live demonstration of AI-enabled capabilities and practical examples. We will explore the evolving role of agentic AI relative to traditional analytics approaches, key cost considerations, governance implications, and lessons learned from organizations developing and deploying AI-enabled monitoring programs.

Session continued next page

8:00 am **Introductions, Discussions and Q&A**

Kristiana Bankova, MS, *Senior Manager, Forensics, EY; New York, NY*

Rachael Hu, MS, MPP CIPM, CISP, CIPP/E, CIPP/US, *Data Privacy Senior Compliance Program Manager, Medtronic; Santa Clara, CA*

Libby M. Rosenberry, *Senior Manager, Forensic & Integrity Services, EY; Boston, MA*

8:50 am **Transition Break**

MORNING MINI SUMMITS: GROUP 6 (9:00 am - 9:50 am)

Mini Summit 29: Building the Compliance Operating Model for What's Next

As AI, automation, and advanced technologies reshape work, compliance can no longer rely on traditional operating models. This session explores how leading organizations are redesigning governance, talent, and oversight to manage new risks, enable innovation, and build a more connected, data-driven compliance function.

9:00 am **Introductions, Discussions and Q&A**

Carly Greenberg, JD, *Vice President, Risk Assessment, Monitoring, Governance & Legal Operations, Pfizer, New York, NY*

Monica Schroeter, CPA, CISA, *VP, Head of Compliance, F2G; Former Head of Compliance Operations and Risk, LivaNova; Former Head of Compliance & Privacy, SK Life Sciences; Goshen, NY*

Clarissa Crain, *Managing Director, Life Sciences Regulatory, Compliance, and Legal Leader, Deloitte Consulting; Phoenixville, PA (Moderator)*

Mini Summit 30: Leveraging AI for More Efficient and Effective Investigations

Legal, compliance, and investigations professionals are increasingly leveraging AI throughout the lifecycle of investigations. Not only do these tools drive cost and time efficiencies, but they also can help investigators get to the truth faster and with less business disruption. But at the same, deploying this technology can create risks if not done with appropriate care and planning. During this session we will:

• Offer practical and creative AI use cases (beyond the basics) throughout the investigation lifecycle, from intake to interviews to investigative output;

• Dig into legal issues presented by the use of AI in investigations, including how to avoid foot faults with respect to the attorney client privilege and attorney work product;

• Provide "watch-outs" or common mis-steps to avoid when leveraging AI tools in this context; and

• Consider how to leverage AI to not only respond to individual allegations, but also to use broader investigations data to identify trends

9:00 am **Introductions, Discussions and Q&A**

Mike D'Alessandro, CFE, CHC, CPA, *Senior Manager, Forensics & Integrity Services, EY; Former Chief Ethics and Compliance Officer, ahma Rx; Iselin, NJ*

Susana Franco, MBA, *Director, Field Compliance and Investigations, Lundbeck; Former Director of Compliance Operations, Orthofix; Dallas, TX*

Tracey Tran, *Senior Manager, Forensic and Integrity Services, EY; New York, NY*

Erin Brown Jones, MA, JD, *Partner, Latham & Watkins; Washington, DC (Moderator)*

Mini Summit 31: Compliance Considerations, Enforcements and Guidelines on Social Media Activities

As companies use social media to engage with patients, healthcare professionals and the public, legal and compliance teams must navigate evolving promotional rules and enforcement expectations. This discussion will explore practical considerations for reviewing social media content, managing employee and third-party activity, and responding when a post raises regulatory concerns.

9:00 am **Introductions, Discussions and Q&A**

Rosemary McKenna, JD, *Executive Director, Regulatory Legal, Organon; Former Director, Regulatory-Legal, Merck; Philadelphia, PA*

Joseph Philipose, JD, *Former Vice President, Chief Ethics and Compliance Officer, Emergent BioSolutions; Former Vice President, U.S. and Enterprise Compliance, Alexion Pharmaceuticals; Former U.S. Compliance Officer, Salix Pharmaceuticals; Boston, MA*

Nikki Reeves, JD, *Partner, FDA & Life Sciences Practice Group, Co-Chair, Government Matters & Regulatory Practice Group, King & Spalding; Washington, DC (Moderator)*

Mini Summit 32: Building a Compliance Roadmap: Practical Strategies for Scaling Compliance Programs

Building a compliance program is not a one-size-fits-all exercise. For life sciences companies navigating clinical/pre-commercial, commercialization, and growth stages, leaders must determine which risks require immediate attention, what elements are needed to address them, and how to build a roadmap that aligns compliance priorities with business objectives.

In this panel discussion, experienced compliance leaders from companies at different maturity stages will share practical approaches to developing and implementing compliance roadmaps that appropriately address the risk at your organization, including:

• Identifying and prioritizing risks across different stages of organizational growth

• Determining what and when program components are needed and how they should evolve over time

• Leveraging existing resources, AI, and cross-functional partnerships to enhance compliance effectiveness

• Balancing competing priorities and resource constraints while maintaining a risk-based approach

• Real-world examples and lessons learned from building and maturing compliance programs

9:00 am **Introductions, Discussions and Q&A**

Geeta Kaveti, MHA, JD, *Chief Compliance Officer & US General Counsel, Nyxoah; Former Vice President, Chief Compliance & Privacy Officer, Nevro; Former Division Counsel - Legal Regulatory & Compliance, Abbott; Vernon Hills, IL*

Fabiana Morette-Lattea, LLM, LLB, *Global Compliance Program Lead, Telix Pharmaceuticals; Former Associate Director, Compliance, Merck; Former Compliance Officer Europe and North America, Roche; Fishers, IN*

Meghan Naas, MA, *Senior Director, Head of Compliance, Innoviva Specialty Therapeutics; Former Head of Corporate Compliance, Adaptimmune; Former Senior Director, Global Operations and US Compliance Officer, Zai Lab; Waltham, MA*

Eric Ortman, MBA, *Senior Director, Legal Operations & Healthcare Compliance, 4D Molecular Therapeutics; Former Senior Director, Legal Operations, BeiGene; Former Director, Legal Operations, Ultragenyx Pharmaceutical; Walnut Creek, CA*

Adam T. Oakley, *Senior Director, Potomac River Partners; Washington, DC (Moderator)*

Mini Summit 33: Fair Market Value in Flux: Navigating Complex Fee Arrangements, HCP & Patient Engagement, and Influencer Compensation Risks

Explore the evolving fair market value (FMV) landscape for healthcare professionals (HCPs), patients, and social media influencers with a focus on the unique market, regulatory, and compliance considerations that influence engagement strategies.

Key Details

- Examine the complexities of establishing FMV for HCPs and patients across markets with varying regulatory requirements, healthcare infrastructure, and economic conditions.
- Discuss challenges created by HCP scarcity, supply-and-demand dynamics, and market-specific restrictions that can affect engagement and compensation.
- Explore FMV considerations for social media influencers/digital opinion leaders, including the lack of standardized compensation benchmarks and platform variability.
- Review emerging compliance risks, including transparency and disclosure expectations, as well as heightened regulatory scrutiny over promotional versus educational content.

9:00 am **Introductions, Discussions and Q&A**

Sarah diFrancesca, JD, *Head of US Compliance – Immunology/Dermatology, Incyte; Wilmington, DE*

Erin B. Pulice, JD, *Litigation & Investigations Counsel, Medtronic; Minneapolis, MN*

Ravneet Talwar, MBA, *Senior Manager, Life Sciences & Healthcare Consulting Group, Paul Hastings; New York, NY*

Laura A. Skinner, MBA, *Managing Director, Life Sciences & Healthcare Consulting Group, Paul Hastings; Managing Director, Rare Disease Compliance Consortium; New York, NY (Moderator)*

Mini Summit 34: Future Proofing the Compliance Function: Operating Models for an AI-Enabled World

As organizations navigate AI adoption, increasing regulatory expectations, and evolving business models, compliance leaders are reevaluating how their functions are structured and staffed. This session could explore emerging operating models, changing skill requirements, the impact of AI on compliance roles, and how organizations can position compliance teams for long-term success.

9:00 am **Introductions, Discussions and Q&A**

Craig B. Bleifer, JD, *Partner, McGuire Woods; Former Corporate Vice President, General Counsel North America; Novo Nordisk; Former Senior Vice President, General Counsel & Secretary, Daiichi Sankyo; New York, NY*

Brett Barlag, MBA, *Senior Managing Director, Healthcare Risk Management and Advisory, FTI Consulting; New York, NY (Moderator)*

9:50 am **Networking Break in the Exhibit Hall**

MORNING MINI SUMMITS GROUP 7 (10:10 am – 11:00 am)

Mini Summit 35: Unstructured Data and Advanced Monitoring Strategies

As communications expand across email, mobile devices, and social media, organizations need more sophisticated strategies for monitoring unstructured data and identifying emerging risks. This session will examine:

- Updating policies and procedures
- Using AI in monitoring programs and establishing appropriate governance
- Identifying common violations and areas for improvement
- Monitoring email, mobile devices, and social media
- Using monitoring results to identify training opportunities

Panelists will share practical approaches for strengthening monitoring programs and translating findings into meaningful compliance improvements.

10:10 am **Introductions, Discussions and Q&A**

Ashley Blackwood, *Manager, Compliance, MiMedx; Marietta, GA*

Christina Woods, MHA, *Partner, Pharmaceutical and Life Sciences Risk & Regulatory Consulting, PwC; New York, NY*

Jeff Hyre, *Vice President Sales, Healthcare and Life Sciences, Global Relay; Denver, CO (Moderator)*

Mini Summit 36: Leading Practices for Managing Investigations Panel

When a report drops, the first 72 hours can define the outcome—this panel will show you how leading organizations respond with speed, rigor, and control. Hear practical, real-world approaches to critical steps in the investigation process, such as preserving evidence and privilege, aligning quickly with counsel and governance teams, reporting out to the organizations, and using various tools and approaches to get to the facts faster. You'll leave with actionable ideas that you can tailor to enhance your investigation practices.

10:10 am **Introductions, Discussions and Q&A**

John S. Rah, JD, *Counsel and Head, Life Sciences Compliance Practice, Potter & Murdock; Potomac, MD*

Olga Zinavenka, CPA, MBA, *Head of Compliance Operations, Keenova; Former Executive Director, Head of Compliance Operations, Endo; Former Compliance Officer – Europe, Middle East and Africa; Paoli, PA*

David Amendola, *Director, Life Sciences, Resolution Economics; Former Global Compliance Risk, Evaluation & Management, ConvaTec; New York, NY (Moderator)*

Mini Summit 37: From Makary to Overton: Reflections and Predictions on the FDA's Priorities

With a new FDA Commissioner announced in August and "micro" trends at the FDA continuing to shift every two to three months, this presentation tracks recent and rapidly evolving developments from peptides to tariffs. The discussion will evaluate where the agency stands today and what to watch considering the risks and opportunities likely to shape the life sciences industry in the coming months.

10:10 am **Introductions, Discussions and Q&A**

Dominick P. DiSabatino, JD, *Partner, Sheppard Mullin Richter & Hampton; Washington, DC*

Scott S. Liebman, JD, *Partner, Chair of FDA Regulatory & Compliance, Co-Chair of Life Sciences Practice, Sheppard Mullin Richter & Hampton; New York, NY*

Mini Summit 38: Compliance Considerations for Rare Disease Products

Rare disease products present distinct compliance questions, from engaging with small patient communities to supporting education about conditions that may be difficult to diagnose. This discussion will explore practical approaches to patient and HCP engagement, interactions with advocacy organizations, and applying compliance principles when the patient population is small and relationships are especially close.

10:10 am **Introductions, Discussions and Q&A**

Cheryl Lee, MBA, *Head of Compliance, Cycle Pharmaceuticals; Former Vice President, ECG Strategy and Execution; Forner Vice President, Head of Compliance Worldwide Markets; New York, NY*

Daniel S. Murdock, MBA, JD, *Associate General Counsel, Sobi - Swedish Orphan Biovitrum AB (publ); Former Associate General Counsel - Life Sciences, McKinsey & Co; Former Assistant General Counsel, Indivior; Washington, DC*

Craig B. Bleifer, JD, *Partner, McGuire Woods; Former Corporate Vice President, General Counsel North America; Novo Nordisk; Former Senior Vice President, General Counsel & Secretary, Daiichi Sankyo; New York, NY (Moderator)*

Mini Summit 39: From Oversight to Partnership: Strengthening Collaboration Between Compliance and the Business

Compliance and business functions share a common goal: enabling organizations to achieve their objectives responsibly and ethically. Yet too often, these functions operate in silos, creating friction, inefficiencies, and missed opportunities for collaboration. This panel discussion will explore how organizations can move from a traditional oversight model to a partnership approach that strengthens both business performance and risk management. Leaders from Compliance, Commercial Excellence, and Business Operations will share practical strategies for building trust, aligning objectives, leveraging data and analytics, and creating governance models that support informed decision-making. Attendees will leave with actionable insights for fostering stronger cross-functional partnerships and positioning Compliance as a strategic business partner that enables growth while maintaining appropriate oversight.

10:10 am **Introductions, Discussions and Q&A**

Jessica Constant, MS, *Associate Director, Compliance & Business Controls – US Oncology, AstraZeneca; Former Global Compliance Business Partner, QIAGEN; Washington, DC*

Edmund Greenidge JD, LLM, CCEP, *Senior Director, Compliance, Summit Therapeutics; Former R&D Ethics and Compliance Lead, Takeda; Former Vice; President, Corporate Compliance, Aclaris Therapeutics; Menlo Park, CA*

Angela Schwartz, JD, *Senior Principal, Global Commercial Compliance Consulting Leader, IQVIA; Former Special Assistant District Attorney, Montgomery County District Attorney's Office; Philadelphia, PA (Moderator)*

Mini Summit 40: Digital Ecosystem Considerations for Life Science Companies

Pharmaceutical companies have increasingly developed complex digital ecosystems that underly their patient support programs, provider outreach, digital marketing, data analytics, and AI initiatives. This session will review regulatory and practical considerations related to overseeing and safeguarding these digital ecosystems. During this session we will speak specifically regarding state and federal regulatory updates, heightened IT security expectations, and litigation and enforcement trends.

10:10 am **Introductions, Discussions and Q&A**

Siyka Bozadjieva-Galiova, JD, *Global Data and Privacy Senior Advisor, Medtronic; Minneapolis, MN*

Christine Moundas, JD, MPH, *Healthcare and Data Partner, Co-Head, Digital Health Initiative, Ropes & Gray; Former Analyst, Office of Inspector General, US Department of Health and Human Services; New York, NY (Moderator)*

11:00 am **Transition Break**

MORNING MINI SUMMITS GROUP 8 (11:10 am – 12:00 pm)

Mini Summit 41: Enabled Technology can Transform Transactional Testing and Other Components of Your Monitoring Program

Disparate systems, unstructured and inconsistent data and documentation, lack of resources, geographic variability….all historic and real challenges to your compliance program, and specifically your monitoring program. Once anomalies and risk are identified, further assessment and testing are required. Historically, this is a manual process, conducted by valuable human resources and team members with limited bandwidth that need to focus on higher risk and impact matters. The use of data and technology is an expectation of our industry's regulators and leadership is continuously asking compliance teams to "do more with less." Compliance professionals need to start considering how automation and AI can be leveraged to enhance their programs. This session will discuss these issues and highlight real-world examples of how compliance and risk assurance functions are addressing these challenges. In this session, we will discuss:

- Current challenges including resources, organizational structure, and geo market variability
- Regulators' expectations around leveraging data, AI, and automation
- What are the core functions and/or components of your compliance program where automation and AI can play a significant role?
- How can technology modernize your transactional testing program?

11:10 am **Introductions, Discussions and Q&A**

Devin Bates, *Global Director, Compliance Operations, Life Sciences, Danaher Corporation; Former Director of Compliance, Walgreens; Chicago, IL*

Terri Segura, *Global Chief Compliance Officer, LivaNova*

Angela Rodin, MBA, *Principal, Global Ethics, KPMG; Former Vice President, Global Head of Investigations and Monitoring, GSK; Washington, DC (Co-Moderator)*

Jordan Seiferas, MS, *Principal, Life Sciences Data Modernization, KPMG US; Columbus, OH (Co-Moderator)*

Mini Summit 42: Emerging Commercial & Market Access Risk in Life Sciences: Navigating Innovation, Compliance, and Evolving Access Models

Review emerging compliance risks, including transparency and disclosure expectations, as well as heightened regulatory scrutiny over promotional versus educational content.

11:10 am **Introductions, Discussions and Q&A**

Sara Simon, JD, *Director of Compliance, IBSA USA; Former Director, US Health Care Compliance Certification Program, Center for Health & Pharmaceutical Law & Policy, Seton Hall University School of Law, Parsippany, NJ*

Anthony S. Greco, MBA, *Managing Director, Pharmaceutical & Life Sciences Advisory Services, PwC; Philadelphia, PA (Moderator)*

Mini Summit 43: Effectively Leveraging Post-Investigation Mitigation to Create a Narrative of Continuous Improvement

In many investigations the focus is often on intentional or negligent conduct of individuals. In the long run; however, the ability to draw lessons learned for your compliance program and to make programmatic improvements in areas like policies, controls, training, and communications provides more value. This panel will explore best practices in managing and tracking programmatic mitigation, when to keep outside counsel involved, when to turn to other SMEs, and ultimately how to incorporate this work into a narrative of continuous improvement. The 2024 DOJ Guidance on Compliance programs expects programs to focus on continuous improvement as a measure of effectiveness. Currently, a lot of focus is around the front end, the use of AI and analytics to identify instances of non-compliance or programmatic gaps, but equal attention is required on the back end, the pull through of mitigation efforts, to be able to tell a convincing story of actionable evaluation of effectiveness.

11:10 am **Introductions, Discussions and Q&A**

Aurélie Ercoli, LLM, PhD, *Partner, Life Sciences Practice Group, DLA Piper; Washington, DC*

Perri Pomper, JD, *Vice President, Compliance - Orthopaedics, Strategy & Education, Stryker; Former Vice President, Compliance & Legal Affairs, Clinical Genomics; Former Director, Ethics and Compliance Strategy & Education, Novo Nordisk; Mahwah, NJ*

Emily Treanor, JD, *Head of Global Investigations & Enterprise Risk Management, Alnylam Pharmaceuticals; Washington, DC*

Jeffrey Scott, JD, *Of Counsel, Life Sciences Practice Group, DLA Piper; Former Lead Compliance Counsel, Digital, Reporting, and Analytics, Pfizer; Philadelphia, PA (Moderator)*

Mini Summit 44: Data Analytics & Compliance Monitoring in an AI-Powered World

Structured and unstructured data is ubiquitous throughout our organizations, as is the opportunity – and the expectation – for compliance officers to actively engage with company data while using it to assess risk, monitor trends, and identify areas of potential concern.

With so many functions, so much data, so many possibilities, and so little time – where do you even begin? And what does "good" look like? How might the data we are collecting and analyzing be used to advance compliance programs and prevent or reduce revenue leakage, and how might the insights that data brings create concerns in its own right, such as concerns regarding violations of FDA or DEA regulations, or implications of the FCA and AKS? What tools work well, and how can they be used most effectively? What are the implications of this harnessing and monitoring data-driven information in investigations and litigation?

This session will discuss a range of data analytics and monitoring tools and techniques ranging from the simple to the complex, and will examine case studies to illuminate the life cycle of structured and unstructured data capture, analysis and ongoing monitoring – in addition to insights into how this same data can be analyzed in reactive scenarios.

11:10 am **Introductions, Discussions and Q&A**

Ignatius Grande, JD, *Of Counsel, Winston Taylor; New York, NY*

Omar S. Richardson, MBA, *Executive Director, Compliance, Shionogi; Former Compliance Officer, Grünenthal Group; New York, NY*

Roberta Rima, MS, *Vice President, Global Compliance Audit, Monitoring & Analytics, Zimmer Biomet; Former Senior Director, Head of Third Party & Corporate Compliance Audit, Gilead; San Francisco, CA*

Stephanie M. Lewko, CFE, *Managing Director, Healthcare Disputes & Economics, Ankura; Washington, DC (Moderator)*

Mini Summit 45: Can Learning Data Reduce Risk? Using Measurement to Drive Defensible, Risk-Based Compliance Training

Regulators and auditors are no longer satisfied with completion rates and pass scores — they want documented evidence that training actually changed behavior and reduced risk. This session focuses specifically on compliance training programs: how to use assessment data, role-based segmentation, and adaptive learning pathways to build defensible proof of competency — not just activity. Attendees will leave with practical strategies for connecting training outcomes to behavioral indicators, right-sizing learning investments by risk level, and producing the audit-ready documentation that holds up under DOJ and OIG scrutiny.

11:10 am **Introductions, Discussions and Q&A**

Adriana Davies, JSM, *Vice President Compliance and Privacy US and China, Smith+Nephew; Former Senior Program Director. Compliance Officer Surgical Innovations, Renal Care Solutions and Robotics, Medtronic; Former Regional Director Americas Ethics and Compliance & Group Corporate Programs, Terex Corporation; Jacksonville, FL*

A.D. Detrick, CPTM, *President, Founder, MetriVerse Analytics; Columbus, OH*

Jaya Saigal, *Senior Director, C&E Policy, Education, Communications, Culture of Integrity and Ombuds, Bristol Myers Squibb; Princeton, NJ*

Lily Zimmerman, *Learning Solution Architect, Tier1 Performance; Covington, KY (Moderator)*

Mini Summit 46: AI in Clinical Practice: Compliance, Governance & FDA Oversight

Regulation of AI in Clinical Practice, Including Clinical Decision Support Tools, Documentation, FDA Clearance Requirements, and Provider Credentialing Considerations

11:10 am **Introductions, Discussions and Q&A**

Julie K. Taitsman, MD, JD, *Managing Director, BDO, US; Former Chief Medical Officer, Office of Inspector General (OIG), US Department of Health & Human Services; Former Special Counsel for Health & Science, United States Senate Committee on Finance; Washington, DC*

Georgia Ravitz, JD, *Partner, FDA & Healthcare Regulatory, Orrick, Herrington & Sutcliffe; Washington, DC; (Moderator)*

12:00 pm **Networking Luncheon**

NETWORKING LUNCHEON

12:00 pm **Networking Luncheon with Dessert and Coffee in the Exhibit Hall**

LUNCHEON MINI SUMMITS GROUP 9 (12:10 pm – 1:00 pm)

Pickup your box lunch in the Exhibit Hall and take it to the Luncheon Mini Summit room of your choice.

Mini Summit 47: Direct-to-Consumer (DTC) Models in the Pharmaceutical Space

- Discuss different DTC program models
- Provide an overview of recent OIG guidance and key laws and regulations impacting DTC programs
- Discuss compliance risk considerations related to execution of DTC programs with pharmacies, telehealth providers and other partners

12:10 pm **Introductions, Discussions and Q&A**

Kala Shankle, JD, *Vice President, Regulatory Affairs, Healthcare Distribution Alliance; President Elect, American Society for Pharmacy Law; Washington, DC*

Alison J. Fethke, JD, *Member of the Firm, Epstein Becker Green; Former Division Counsel, Legal Regulatory & Compliance, AbbVie; Chicago, IL (Moderator)*

Mini Summit 48: ROI of AI: Separating Value from Hype

AI has rapidly become a strategic priority across organizations, but the path from adoption to measurable value remains uneven. Many teams are piloting tools, investing in platforms, and exploring use cases, yet fewer can clearly articulate where AI is delivering sustained ROI.

This discussion will focus on separating high-value AI applications from hype-driven initiatives. Participants will share how their organizations are measuring success, what metrics they are using, which use cases are producing tangible impact, and where AI has fallen short of expectations.

- Difference definitions of ROI for AI
- Measuring Value in quantitative and qualitative terms
- Creating value while avoiding AI overload and fatigue
- Continued success with and without AI

12:10 pm **Introductions, Discussions and Q&A**

Michelle Dineen Jerrett, JD, *Regional Head of Investigations & Monitoring, North America, Sanofi; Former Legal Director, Endoscopy, Boston Scientific; Former Assistant United States Attorney, US Attorney's Office, District of Massachusetts; Cambridge, MA*

Stephanie Macholtz, JD, MBA, *Adjunct Faculty, Suffolk University Law School; Former Vice President, Healthcare Compliance, Organogenesis; Former Senior Director Global Compliance and Deputy General Counsel, Clinical Affairs, Apellis Pharmaceuticals; Cambridge, MA*

Omar S. Richardson, MBA, *Executive Director, Compliance, Shionogi; Former Compliance Officer, Grünenthal Group; New York, NY*

Katie Jacyna, *Chief Product Officer, HELIO; Denver, CO (Moderator)*

Mini Summit 49: Lean, Credible, Defensible: Compliance Programs That Work Without Big Budgets

Building an effective compliance program does not always require a large team or budget. This session will explore practical, low-cost strategies for creating a risk-based compliance program that is credible, sustainable, and responsive to regulatory expectations. Panelists will discuss how organizations can prioritize limited resources, focus on high-impact activities, and demonstrate that their compliance programs are working effectively.

12:10 pm **Introductions, Discussions and Q&A**

Loren Becher, MPH, *Former Senior Director, Quality, Ethics, Compliance & Risk Enterprise Programs, Emergent BioSolutions; Former Director, Compliance Operations & Planning, AstraZeneca; Former Director, Chief of Staff & Enterprise Programs, Alexion Pharmaceuticals; Rye Brook, NY*

Julian Gabbard, JD, *Associate General Counsel, Meso Scale Diagnostics; Former Head of Legal and Compliance, Corporate Counsel, Kolon TissueGene; Former Senior Director, Arcellx; Washington, DC*

Elizabeth R. Weiss, JD, *Vice President, Chief Compliance & Privacy Officer, Invivyd; Former Head of Compliance for Americas & EMEA, Lupin; Former Senior Director, US Risk and Compliance, LEO Pharma; Former Assistant General Counsel, Compliance Lead, Global Health & Patient Access, Pfizer; Livingston, NJ*

Robert A. Wells, JD, *Health Care Regulatory Shareholder, Chair, Health Law and Government Solutions, Baker, Donelson, Bearman, Caldwell & Berkowitz; Baltimore, MD (Moderator)*

Mini Summit 50: A Look Under the Hood: Exploring PBM and Payer Arrangement Risks and Emerging Reporting Requirements

PBM and payer arrangements face increasing scrutiny as federal and state authorities focus on drug pricing, rebates, and transparency. This session will examine:

- Federal and state scrutiny of PBMs and their financial relationships with manufacturers
- Manufacturer, PBM, and prescription drug plan arrangements and the flow of funds
- Auditing and monitoring strategies for channel partner arrangements
- The intersection of these arrangements with drug pricing reforms
- Emerging PBM reporting requirements involving formularies, pricing, and manufacturer arrangements
- Implications for Market Access interactions and formulary negotiations

Panelists will share practical considerations for evaluating these complex arrangements as part of a broader compliance risk-management strategy.

8:00 am **Introductions, Discussions and Q&A**

Daryl Berke, JD, *Assistant General Counsel, Specialty US Market Access Lead, Sanofi; Washington DC*

Craig B. Bleifer, JD, *Partner, McGuire Woods; Former Corporate Vice President, General Counsel North America; Novo Nordisk; Former Senior Vice President, General Counsel & Secretary, Daiichi Sankyo; New York, NY*

Andrew Bonasera, MSF, *Senior Managing Director, FTI Consulting; Washington, DC*

Katherine C. Norris, MPA, *Senior Managing Director, Life Sciences Compliance, Disputes & Economics, Ankura; Washington, DC (Moderator)*

Mini Summit 51: Collaborating to Create Lasting Solutions During and After Monitoring or Investigations

When monitoring or investigatory teams come back with findings, frequently there is a wish to manage the response narrowly rather than taking the more challenging step of identifying and addressing the root cause. The presentation will discuss strategies and practicalities of collaborating across compliance and business teams during and after monitoring and investigations. The presentation will cover collaboration within compliance, collaboration with the business, and managing both mundane and higher risk findings—all with the goal of creating lasting results. It will include audience interaction. Presenters have served in both business partnering and monitoring/investigatory roles and have managed internal and external investigations.

12:10 pm **Introductions, Discussions and Q&A**

Katherine "Kate" Dyson, JD, *Head of US Compliance, Alexion Pharmaceuticals; Former Head of Assurance, Alexion, AstraZeneca RDU; Former Senior Director, Neurology Compliance Business Partner; Alexion Pharmaceuticals; Boston, MA*

Naveen Ganesh, MA, JD, *Vice President, Global Head of Investigations, Ethics & Compliance, Takeda; Former Head of Investigations & Monitoring, North America & Global Specialty Care, Sanofi; Former Compliance Assurance & Investigations (Lead), Verily;; Boston, MA*

Kathryn Harris, JD, *Head of US Compliance, Galderma; Former Senior Director, Global Compliance, Emergent BioSolutions; Former Compliance Business Partner (Market Access, Patient Services, and Strategic Accounts); Alexion Pharmaceuticals; Boston, MA*

Audrey DeGuarde, MSJ, *VVice President, General Manager, Commercial Compliance Solutions, Medispend; Morristown, NJ (Moderator)*

Mini Summit 52: Annual FDA Office of Prescription Drug Promotion Update

This session will explore the evolving compliance landscape for drug advertising and promotion from the perspective of Twyla Mosey, the Director of the Office of Prescription Drug Promotion ("OPDP") in the Center for Drug Evaluation and Research at FDA. The discussion will examine enforcement trends, including the surge in warning and untitled letters since the administration's "crackdown" on deceptive drug advertising announced in September 2025, as well as OPDP's current priorities and areas of focus. The discussion will provide practical guidance for companies on responding to OPDP enforcement letters, engaging constructively with OPDP, and strengthening compliance processes to mitigate advertising and promotion risk.

12:10 pm **Introductions, Discussions and Q&A**

Twyla Mosey, PharmD, RPh, *Director, Office of Prescription Drug Promotion, US Food and Drug Administration, Washington, DC*

Joshua (Josh) Oyster, JD, *Partner, Life Sciences Regulatory & Compliance Practice, Ropes & Gray; Former Counsel, Humana; Washington, DC (Moderator)*

1:00 pm **Transition Break**

CLOSING PLENARY SESSION

1:10 pm **Welcome and Introduction**

Tara Shewchuk, JD, LLM, *Senior Vice President, Chief Privacy, Integrity & Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Committee)*

1:15 pm **Keynote Annual DOJ Civil Division Update Fireside Chat**

Brenna Jenny, JD, MPH, *Deputy Assistant Attorney General, Commercial Litigation Branch, Civil Division, US Department of Justice; Former Principal Deputy General Counsel & Chief Legal Officer, Centers for Medicare & Medicaid Services, US Department of Health and Human Services; Washington, DC*

Interviewed by:
Scott A. Memmott, JD, *Partner, Morgan, Lewis & Bockius LLP; Former Special Assistant US Attorney, Criminal Division, US Attorney's Office, Eastern District of Virginia and Former Trial Attorney, US Department of Justice, Civil Division, US Department of Justice; Washington, DC*

1:45 pm

Investigations, Enforcement and Prosecutions Roundtable

Jennifer L. Bragg, JD, *Partner, Latham & Watkins; Former Associate Chief Counsel for Enforcement, Office of Chief Counsel, US Food and Drug Administration; Washington, DC*

Greg Demske, JD, *Partner, Goodwin Procter; Former Chief Counsel, Office of Inspector General, US Department of Health and Human Services; Washington, DC*

Preeya Noronha Pinto, JD, *Partner, DLA Piper; Former Acting General Counsel, US Department of Health and Human Services; Washington, DC*

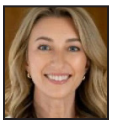
Kip F. Ebel, MBA, *Principal, Forensic & Integrity Services, EY; New York, NY (Moderator)*

2:15 pm

Updates from AdvaMed, BIO and PhRMA

John Delacourt, JD, *Deputy General Counsel, Biotechnology Innovation Organization; Former Chief Antitrust Counsel, Office of Policy Planning, US Federal Trade Commission; Washington, DC*

Ida Nassar, JD, *Vice President, Assistant General Counsel, Ethics & Compliance, AdvaMed; Former Senior Attorney, Office of Chief Counsel, US Drug Enforcement Administration; Washington, DC*



Julie Ritchie Wagner, JD, *Assistant General Counsel, PhRMA; Former Senior Counsel, Office of Counsel to the Inspector General, US Department of Health and Human Services; Washington, DC*

Mark C. Scallon, MHA, *Principal, Risk Advisory, Life Sciences Consulting, Compliance and Ethics Market Leader, Baker Tilly; Richmond, VA (Moderator)*

2:45 pm

Market Access in 2026 and Beyond: A Shifting Compliance Landscape

Jill Dailey, JD, *Vice President & Chief Compliance Officer, Oncology, Menarini Stemline; Former Vice President and Chief Compliance Officer, Incyte; Former Assistant General Counsel & Asia Pacific Compliance Lead, Pfizer; New York, NY (PCF Board Member)*

Christian Scognamillo, JD, *Associate General Counsel, US Market Access & Pricing, Abbvie; Former Senior Corporate Counsel, Market Access & Commercial Transactions, Exelixis; Former Senior Director, Legal, Access & Transactions, Alnylam, San Francisco, CA*

Meena Datta, JD, *Partner and Global Co-leader, Healthcare Practice, Sidley Austin; Chicago, IL (Moderator)*

3:15 pm

Networking Break in the Exhibit Hall

JJ Kuhn, JD, *Chief Ethics & Compliance Officer, Danaher; Former Senior Vice President and Chief Ethics & Compliance Officer, Solventum; Former Vice President Chief Counsel, Global Investigations, Medtronic; Minneapolis, MN(Pharma Congress Medical Device Executive Committee)*

3:45 pm

Burgeoning Compliance Risks

Brian C. Barry, JD, *Vice President, Chief Compliance Officer, Vertex Pharmaceuticals; Former Chief Compliance Officer & Compliance Counsel, EMD Serono; Boston, MA*

Jennifer Dubas, JD, *Chief Ethics and Compliance Officer, Jazz Pharmaceuticals; Former Senior Vice President, Chief Compliance Officer, Cencora; Former Senior Vice President, Chief Compliance Officer, AmerisourceBergen; Ambler, PA*

Shannon Kelley, JD, *Executive Vice President, Chief Legal Officer, Madrigal; Former Vice President, Head of Compliance; North America and Global Specialty Care GBU, Sanofi; Former Assistant US Attorney & Deputy Chief of Litigation, US Attorney's Office District of Massachusetts; Boston, MA*

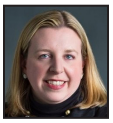
John S. Rah, JD, *Counsel and Head, Life Sciences Compliance Practice, Potter & Murdock; Potomac, MD*

Yogesh Bahl, MBA, *Partner and Practice Leader, Resolution Life Sciences and Healthcare; Chief Financial Officer and Head of Investor Relations, IACTA Pharmaceuticals; New York, NY (Moderator)*

4:15 pm

AI and the Evolving Compliance Function

Peter Brensilver, JD, MPH, *Senior Vice President, Chief Global Commercial Compliance & Risk Counsel, Pfizer; New York, NY*



Jennifer Dubas, JD, *Chief Ethics and Compliance Officer, Jazz Pharmaceuticals; Former Senior Vice President, Chief Compliance Officer, Cencora; Former Senior Vice President, Chief Compliance Officer, AmerisourceBergen; Ambler, PA*

Rebekah Latchis, JD, *Vice President, Risk Intelligence & Investigations, Office of Privacy, Integrity & Compliance, Medtronic; Former Corporate Responsibility Director, Bon Secours; Washington, DC*

Neeraj Gupta, MBA, *President, Chief Technology Officer, Cresen Solutions; Chester Springs, PA (Moderator)*

4:45 pm

Managing Risk in Unsettled Times

Deborah A. Curtis, JD, *Partner & Co-Chair, White Collar Defense & Investigations Practice, Arnold & Porter; Former Deputy General Counsel, Central Intelligence Agency; Former Deputy Chief, Counterespionage Section, National Security Division, US Department of Justice; Former Deputy Chief, National Security Section, District of Columbia US Attorneys' Office; Washington, DC*

Ambassador Barbara A. Leaf, JD, *Senior International Policy Advisor, Arnold & Porter; Former Special Assistant to the President and Senior Director for the Middle East and North Africa; Former Assistant Secretary of State for Near Eastern Affairs, and Former US Ambassador to the United Arab Emirates, US State Department; Washington, DC*

Forrest Deegan, JD, *Principal Consultant, Spark Compliance Consulting, a Diligent brand; Co-Chair, Corporate Compliance & Ethics Subcommittee, Business Law Section, American Bar Association; Former Chief Ethics and Compliance Officer, Abercrombie & Fitch and Victoria's Secret; Columbus, OH (Moderator)*

5:15 pm

ADJOURNMENT

Daryl Kreml, JD, *Head, Ethics & Compliance, Global Oncology, Takeda; Former Senior Vice President, Chief Enterprise, Risk & Compliance Officer, Sage; Former US Compliance & Global Program Management, Biogen; Boston Scientific; Boston, MA (PCF Board Member)*

PHARMACEUTICAL AND MEDICAL DEVICE ETHICS AND COMPLIANCE CONGRESS DAY 4

(Industry-only Session for Pharmaceutical and Medical Device Company Ethics and Compliance Professionals and In-house Counsel only. Hosted by PCF)

PCF INDUSTRY-ONLY COMPLIANCE BEST PRACTICES THINK TANK

8:00 am **Welcome and Introductions**

Daryl Kreml, JD, *Head, Ethics & Compliance, Global Oncology, Takeda; Former Senior Vice President, Chief Enterprise, Risk & Compliance Officer, Sage; Former US Compliance & Global Program Management, Biogen; Boston Scientific; Boston, MA (PCF Board Member)*

Antitrust Admonition

Brian A. Bohnenkamp, MHA, JD, *Partner, FDA and Life Sciences, King & Spalding; Washington, DC*

8:10 am **Inside the New DOJ Initiative: The Rising Importance of Data Miners**

The DOJ has added a powerful new tool to its enforcement toolbox: Data Miners. Under its new initiative, the Department is partnering with select data-miner relators who can sift massive public datasets to uncover statistical anomalies, hidden patterns, and potential False Claims Act violations. We'll break down what this initiative means for compliance teams, how data-driven allegations may originate and evolve, and the practical steps organizations should take now to strengthen monitoring, documentation, and response readiness.

This session features a panel moderated by Mark Jensen, a 30-year defense-side veteran of pharmaceutical investigations and FCA litigation, with insights from economist Andrée-Anne Fournier from the Analysis Group, who brings extensive expertise in data analysis and has paired with Mark to develop data-driven defenses in life sciences matters for over a decade, and with further perspective from Corey Amundson, a former Acting U.S. Attorney, Section Chief in DOJ's Criminal Division, head of DOJ's internal affairs, and longtime Assistant U.S. Attorney specializing in healthcare fraud and other white-collar crimes. In these roles, Corey led multiple crisis-response efforts and developed proactive enforcement initiatives aligned with DOJ priorities—including one of the nation's first healthcare fraud strike forces, the Criminal Division Corporate Whistleblower Awards Pilot Program, and a financial-fraud task force built around data-mining analytics.

Featuring:
Corey R. Amundson, JD, *Partner, Special Matters & Government Investigations Practice Group, King & Spalding; Former Head, Public Integrity Section, US Department of Justice; Former Acting US Attorney, Middle District of Louisiana; Washington, DC*

Andrée-Anne Fournier, MA, *Managing Principal, Analysis Group; Brookline, MA*

Mark A. Jensen, JD, *Co-Chair & Partner, Special Matters & Government Investigations, King & Spalding; Washington, DC*

8:45 am **Wins & Losses in the Early Days of AI – Audience Discussion Session**

AI has entered compliance workflows with remarkable speed—and with uneven results. Some teams are already seeing meaningful efficiency gains, while others are wrestling with adoption challenges, accuracy concerns, governance gaps, or cultural resistance.

We'll discuss early wins, early failures, and the real-world lessons learned from attendees implementing AI in compliance programs. This is a candid, practical discussion about what's working, what isn't, and how organizations are navigating the realities of building AI-enabled compliance functions. Come prepared to share!

Anisa Dhalla, *Vice President, Chief Ethics and Compliance Officer; Global Head, Ethics and Compliance, UCB; Atlanta, GA (PCF Board Member)*

Daryl Kreml, JD, *Head, Ethics & Compliance, Global Oncology, Takeda; Former Senior Vice President, Chief Enterprise, Risk & Compliance Officer, Sage; Former US Compliance & Global Program Management, Biogen; Boston Scientific; Boston, MA (PCF Board Member)*

Tara Shewchuk, JD, LLM, *Senior Vice President, Chief Privacy, Integrity & Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Committee)*

9:15 am **Benchmarking and Q&A Forum**

Opportunity for polling, asking questions and benchmark against peers for valuable insights-Come prepared with questions!

Anisa Dhalla, *Vice President, Chief Ethics and Compliance Officer; Global Head, Ethics and Compliance, UCB; Atlanta, GA (PCF Board Member)*

Daryl Kreml, JD, *Head, Ethics & Compliance, Global Oncology, Takeda; Former Senior Vice President, Chief Enterprise, Risk & Compliance Officer, Sage; Former US Compliance & Global Program Management, Biogen; Boston Scientific; Boston, MA (PCF Board Member)*

Tara Shewchuk, JD, LLM, *Senior Vice President, Chief Privacy, Integrity & Compliance Officer, Medtronic; Minneapolis, MN (Pharma Congress Medical Device Executive Committee)*

10:00 am **Networking Break**

10:20 am **Customer-Centric Complexity: Meeting Customer Needs While Balancing Field Pressures**

Customer expectations and field realities are shifting faster than traditional governance models can absorb. Field teams are navigating mounting office restrictions, expanding reimbursement responsibilities, role overlapping, emerging roles, and an escalating need for data, all within the constraints of firewalls and privacy requirements. Panelists will discuss how organizations can modernize governance to keep pace, enabling coordinated, compliant, and customer-centric engagement amid accelerating change. Drawing on diverse real-world experiences, panelists will share practical perspectives, including frontline field insights, on navigating these shifts and balancing customer needs with organizational realities.

Michelle Murphy, *Executive Director, Compliance, Regeneron; New York, NY*

Denis Jacob, *Ethics & Compliance Americas Regional Leader, Solventum; Former Ethics & Compliance Business Partner, Orthopedics, Henry Schein; Former Head of Compliance Audit and Investigations, GE; New York, NY*

George Kuri, *Former Area Business Leader, Novartis Oncology; Former, District Sales Leader, Uro-Oncology/GI, Ferring Pharmaceuticals; Atlanta, GA*

Laura E. Heyduk, *Vice President, Head of Global Compliance, KalVista Pharmaceuticals; Former Head of US Compliance, Biogen; Former Compliance Director, Healthcare Compliance, Insulet Corporation; Cambridge, MA (Moderator)*

11:00 am **Breakout by Company Size: "Cross Functional Account Planning & Guardrails"**

Topics Include: Cross-Functional Account Planning – roles, permissible topics, frequency and guardrails

11:30 am **Last Minute Questions & Benchmarking**

Anisa Dhalla, *Vice President, Chief Ethics and Compliance Officer; Global Head, Ethics and Compliance, UCB; Atlanta, GA (PCF Board Member)*

Noon **CONGRESS ADJOURNMENT**