MINI SUMMIT TOPICS LISTED BY CATEGORIES

NOTE: This is only a guide to help organize and manage topics. Please refer to the brochure for an accurate description of each Mini Summit to facilitate your selections as topics may fall into multiple categories.

AI AND DATA ANALYTICS	MS 2: Enhancing Compliance and Overcoming Challenges under Corporate Integrity Agreements (CIA) through AI and Automation MS 12: Responsible AI at the Cusp of Scale MS 21: AI and Compliance Analytics: Real Case Studies for Managing HCP, HCO and Third-Party Risk MS 43: Detecting Bias in your AI Tools MS 46: Analytics and Monitoring: Understanding the Risks and Underlying Complexity of Today's Copay Program Landscape
COMPLIANCE FRAMEWORK (SEVEN ELEMENTS)	MS 3: Fostering a Speak Up Culture MS 4: Presenting to Senior Executives Effectively MS 9: Compliance Governance and Board of Directors MS 11: Targeted, Tailored, True-to-Life: How DOJ Guidance Should Shape Your Compliance Training MS 15: Balancing Compliance and Legal Roles and Responsibilities MS 16: Risk-Based Compliance Audits: Prioritizing What Matters Most MS 17: ESG Overload – A CCO's Guide to a Balanced Program Implementation MS 18: Compliance Training on the Digital Frontier MS 19: Substance over Form: Practical Approaches to Enterprise Risk Management (ERM) MS 26: Change Management for Compliance Program Transformation MS 29: Advancing Risk Management: A call to action for Integrated Risk Management (IRM) in Pharma MS 30: Attention Please! Elevating E&C Learning & Engagement in the Age of Distraction MS 35: Privacy Top-Ten: Strategies and Tactics to Address Operational Impacts of the Evolving Privacy Landscape MS 42: Best Practices in Investigations MS 51: Negotiating DOJ expectations and business challenges while creating a "Culture of Compliance"
DRUG PRICING/ TRANSPARENCY	MS 28: Government Programs and Compliance Office Oversight MS 31: Key Learnings from CMS Audits MS 41: Intersecting Reforms Affecting Drug Pricing MS 45: US Trends and Updates in HCP Spend Transparency and Drug Pricing Transparency MS 47: Lessons from Spend Transparency and Their Direct Application to Pricing Transparency
HCP ENGAGEMENT	MS 5: Interactions Between Sales and Medical Affairs MS 13: Hot Topics and Compliance Oversight in the Research and Development Area MS 22: Beyond Sales Representatives and MSLs: Considerations for Other FieldBased Roles MS 23: Fair Market Value: Navigating the Ongoing Changes in HCP Compensation Compliance MS 32: Exploring Risks and Enforcement Trends for Buy-and-Bill Drugs MS 39: HCP Expense Auditing and Monitoring MS 44: Evolving Risks in Medical Affairs
PATIENT INTERACTIONS	MS 10: Health Equity Initiatives —Compliance Considerations MS 20: Patient Support Programs and Patient Access Programs—A Look at the Evolution, Risks, and Enforcement Activity MS 25: Compliance Considerations for Rare Disease MS 33: Global Compliance Considerations for Designing Patient Support Programs
REGULATORY ENFORCEMENT UPDATES	MS 1: Federal Criminal and Civil Enforcement: Recent Highlights and Emerging Issues MS 6: Benchmarking and Brainstorming regarding Evolving Regulator Expectations relating to Off-Channel Communication MS 7: OIG New General Compliance Program Guidance MS 8: Update on Medical Device Regulatory and Enforcement Actions MS 14: Recent Developments in Enforcement Actions MS 24: Is Bigger Always Bad?Assessing Developing FTC Antitrust Enforcement Trends in Life Sciences MS 34: Qui Tam Declinations—Analysis and Trends When DOJ Declines to Intervene MS 36: Hot Topics in State Law Compliance MS 37: The Latest and Greatest in Enforcement and Guidelines for Social Media and Influencers MS 38: Compliance Considerations for Genetic Testing MS 48: State Law Updates— 2024 Recap and 2025 MS 50: Market Access— Compliance Considerations When Resolving Disputes
THIRD PARTY RISK MANAGEMENT	MS 27: Service Fees: Hidden Compliance Risks and Downstream Market Access and Government Price Reporting Implications MS 40: The Role of Technology and Process Enhancements for Effective ABAC Third Party Vendor Diligence MS 49: Addressing Nuances in Third-Party Risk Management for Medical Devices