

CONTINUING LEGAL EDUCATION CREDIT SELF REPORTING ATTENDANCE AND EVALUATION FORM — TWENTY-FOURTH PHARMACEUTICAL AND MEDICAL DEVICE ETHICS AND COMPLIANCE CONGRESS

The Pharma/Device Congress is an approved Distance Learning Provider for PA MCLE. As such, the Congress will submit and pay for claimed CLE credits to PA MCLE for attorneys barred in Pennsylvania. For attorneys barred in other states, the Summit will post a record of their attendance at the Pharma Congress in the PA MCLE databases and will issue an Pharma Congress Certificate of Attendance that the attorney seeking CLE credits may use to self-report their participation to states. The Pharma Congress does not guarantee that states will accept the Certificate of Attendance.

As a pre-requisite to the submission of claimed CLE credits to PA MCLE for attorneys barred in Pennsylvania and the issuance of a Pharma Congress Certificate of Attendance, a Pharma Congress attorney attendee must:

- Pay a fee of \$100 to the Pharma Congress @ www.pharmacongress.com, and
- Fully complete and execute this CONTINUING LEGAL EDUCATION CREDIT SELF REPORTING ATTENDANCE AND EVALUATION FORM FOR THE PHARMA CONGRESS. Payment must be made and the completed and executed form submitted via email to cindyr Ramirezgh@outlook.com no later than **November 30, 2023**.

PERSONAL CONTACT INFORMATION

COMPLETE THE FOLLOWING:

NAME	ADDRESS
SIGNATURE OF REGISTRANT - REQUIRED	CITY/STATE/ZIP
JOB TITLE	TELEPHONE
STATE WHERE YOU PRACTICE	E-MAIL
	PA BAR NUMBER IF FROM PA
	NUMBER OF HOURS YOU PARTICIPATED

SELF REPORTING OF ATTENDANCE AT THE TWENTY-FOURTH PHARMA CONGRESS

Please mark those Pharma/Device Congress sessions below that you attended.

DAY I: WEDNESDAY, OCTOBER 25, 2023

SPECIAL INVITATION ONLY SESSION: CHIEF COMPLIANCE OFFICER ROUNDTABLE

What is the Role of Compliance in the Era of ESG?	1 hour
The Rising Stakes of Data Governance and How Companies are Responding	1 hour
A Fireside Chat on Whistleblowers with Kirsten Mayer, JD and Gregg Shapiro, JD	1 hour
Open Forum: Q&A	.5 hour

MINI SUMMITS GROUP I

MINI-SUMMIT 1: Compliance Primer and How to Make the Most of Your Time at the PCF Congress	.83 hour
MINI-SUMMIT 2 : Data Analytic Strategies for Compliance	.83 hour
MINI-SUMMIT 3: Insights from Medical Device Corporate Integrity Agreements	.83 hour
MINI-SUMMIT 4: Update on Federal Government Pharmaceutical Price Negotiations	.83 hour
MINI-SUMMIT 5: The Latest in Social Media Enforcements	.83 hour

MINI SUMMITS GROUP II

MINI-SUMMIT 6: Government Enforcement Actions Piggy-Backing on Product Liability Litigation	.83 hour
MINI-SUMMIT 7 : HCP Engagement: Business Needs Assessment, Contracting, and Activities Management	.83 hour
MINI-SUMMIT 8: Preparing for Government Intervention in Drug Pricing	.83 hour
MINI-SUMMIT 9: Alternative Funding Vendors: The Future and Options for Responding to New Challenges to Patient Support Programs	.83 hour
MINI-SUMMIT 10: R&D and Clinical Trials Compliance Update	.83 hour

MINI SUMMITS GROUP III

MINI-SUMMIT 11: Proving the Value of Corporate Compliance	.83 hour
MINI-SUMMIT 12 : Internal Investigations: Best Practices to Address Compliance Concerns and Reduce Risk	.83 hour
MINI-SUMMIT 13: Insights and Actionable Deliverables in Response to the DOJ Self-Disclosure Policy	.83 hour
MINI-SUMMIT 14: Compliance Considerations for Market Access Initiatives	.83 hour
MINI-SUMMIT 15: Data Analytics to Monitor Payments to HCPs, HCOs, Distributors, and Other Third Parties	.83 hour

MINI SUMMITS GROUP IV

MINI-SUMMIT 16: All Things Patients	.83 hour
MINI-SUMMIT 17 : Evolving Risks in Medical Affairs	.83 hour
MINI-SUMMIT 18: Contemporary Trends in Fair Market Value	.83 hour
MINI-SUMMIT 19: Risk & Ethics Perspective of Artificial Intelligence	.83 hour
MINI-SUMMIT 20: Drug Pricing — Considerations around Price Controls, Negotiation, and Access	.83 hour

LUNCHEON MINI SUMMITS GROUP V

MINI-SUMMIT 21: The Role of Fair Market Value in Pharma and Medical Device Compliance Programs	.83 hour
MINI-SUMMIT 22 : Responsibly Harnessing the Power of AI	.83 hour
MINI-SUMMIT 23: Is the “No Patient Left Behind” Approach to Patient Support Programs Viable?	.83 hour
MINI-SUMMIT 24: Assessing Compliance Priorities: Considerations for Conducting an Effective and Collaborative Compliance Risk Assessment	.83 hour

OPENING PLENARY SESSION

Keynote Fireside Chat with Geoff S. Martha, Chairman and Chief Executive Officer, Medtronic	.5 hour
Keynote: OIG Update	.75 hour
US DOJ Keynote	.5 hour
Strategic, Behavioral Compliance, and Economic Considerations Regarding DOJ’s Pilot Program	.5 hour
Prosecution and Enforcement Actions Update	.75 hour
Annual Chief Compliance Officer Fireside Chat	1 hour

DAY II: THURSDAY, OCTOBER 26, 2023

BREAKFAST MINI SUMMITS GROUP VI

MINI-SUMMIT 25: Sanctions — Compliance Monitoring Approaches and Tools	.83 hour
MINI-SUMMIT 26 : A Fireside Chat with Jim Sheehan, JD	.83 hour
MINI-SUMMIT 27: Key Learnings from CMS Audits of Open Payments	.83 hour
MINI-SUMMIT 28: New Levels of Transparency and Governance Across the Device and Pharma Ecosystems	.83 hour

MINI SUMMITS GROUP VII

MINI-SUMMIT 29: Hiring and Developing Compliance Leaders	.83 hour
MINI-SUMMIT 30 : Privacy & Data Protection: Changes in Laws and the Impact on Our Industry	.83 hour
MINI-SUMMIT 31: Annual FCPA Update	.83 hour
MINI-SUMMIT 32: Developing an Effective AI Governance Model: What you Need to Know	.83 hour
MINI-SUMMIT 33: Navigating M&A of FDA-Regulated Companies	.83 hour

MINI-SUMMIT 34: Enterprise Communications Monitoring and the Recent DOJ Guidance	.83 hour
MINI-SUMMIT 35: DEI Training: Why It Matters and How to Do It?	.83 hour

MINI SUMMITS GROUP VIII

MINI-SUMMIT 36: Outsourced Compliance Programs: How to Make them Work?	.83 hour
MINI-SUMMIT 37 : Fostering a Speak Up Culture at Your Organization	.83 hour
MINI-SUMMIT 38: AI — Art of the Possible	.83 hour
MINI-SUMMIT 39: Third Party Risk Management: Onboarding Diligence, Oversight and Exercising Audit Rights	.83 hour
MINI-SUMMIT 40: Global Aspects and Challenges of FCPA/Compliance Investigations	.83 hour
MINI-SUMMIT 41: Compliance Considerations for Small and Emerging Companies	.83 hour
MINI-SUMMIT 42: Legal Risks Around Digital Health Technology	.83 hour

MINI SUMMITS GROUP IX

MINI-SUMMIT 43: Global Ethics and Compliance: Evolution of Legislation	.83 hour
MINI-SUMMIT 44: DOJ Agreements, Dual Reporting and the Role of the Compliance Officer	.83 hour
MINI-SUMMIT 45: Compliance in Life Science: Unleashing AI's Potential	.83 hour
MINI-SUMMIT 46: Engaging Your Board Effectively in an Era of Heightened Scrutiny and Enforcement Risk	.83 hour
MINI-SUMMIT 47: Pixels and Privacy: Navigating the Litigation and Enforcement Landscape of Website and Mobile App Privacy	.83 hour
MINI-SUMMIT 48 : The Modern Investigation: Current Best Practices for Investigation	.83 hour
MINI-SUMMIT 49: Using Behavioral Compliance to Improve Your Compliance Program — Practical Implementation Suggestions	.83 hour

MINI SUMMITS GROUP X

MINI-SUMMIT 50: Global Ethics and Compliance: ESG Governance	.83 hour
MINI-SUMMIT 51: Navigating Compliance Challenges: Real-World Case Studies and Solutions	.83 hour
MINI-SUMMIT 52 : Compliance Considerations for Rare Disease	.83 hour
MINI-SUMMIT 53: What Do Health Equity Initiatives Mean for Compliance?	.83 hour
MINI-SUMMIT 54: Fireside Chat: Reflections on the Role of the False Claims Act Liability over the past 50 Years into a Major Force in the Regulation of the Pharmaceutical and Medical Device Industries	.83 hour
MINI-SUMMIT 55: Evolving Board of Directors and Compliance Committees Oversight Obligations	.83 hour
MINI-SUMMIT 56: Comedy & Compliance — Shifting Culture & Building Trust with Entertainment	.83 hour

LUNCHEON MINI SUMMITS GROUP XI

MINI-SUMMIT 57: Recent Developments in DOJ and FTC Enforcement Actions	.83 hour
MINI-SUMMIT 58: Building Fearlessness into Organizations	.83 hour
MINI-SUMMIT 59: Compliance Monitoring — Operational Insights and Lessons Learned	.83 hour
MINI-SUMMIT 60: Old Concepts New Risks: Trends in Medical Education Support Compliance	.83 hour

CLOSING PLENARY SESSION

Keynote Fireside Chat with Richard Simkin, Chief Commercial Officer, Indivior	.5 hour
The Future Chief Compliance Officer	.75 hour
FDA Keynote	.5 hour
Updates from AdvaMed and PhRMA	.5 hour
Strategic Resource Management for Optimized Compliance	.5 hour
What's Next in Pharma: Confronting a Challenging Macroeconomic Environment with Innovation	.5 hour

DAY III: FRIDAY, OCTOBER 27, 2023

INDUSTRY-ONLY COMPLIANCE BEST PRACTICES THINK TANK

AI, ChatGPT, and Machine Learning — What Every
Compliance Professional Needs to Know!

1 hour

Key Takeaway Table Discussions	.25 hour
Best Practice Sharing: Latest DOJ Guidances: How Is Your Company Preparing?	.5 hour
How to be a Wildly Effective Compliance Officer?	1 hour
Key Takeaway Table Discussions	.25 hour
Open Forum — Q&A and Best Practice Sharing	.5 hour

WEDNESDAY, NOVEMBER 15, 2023

VIRTUAL GLOBAL PHARMA & MEDICAL DEVICE ETHICS & COMPLIANCE DAY

Keynote Address: Challenges in Developing
a Truly Global Ethics and Compliance Program

.5 hour

EU, CEE and MEA Ethics and Compliance Developments Update Roundtable	1 hour
Asia Pac Ethics and Compliance Developments Update Roundtable	1 hour
LatAm Ethics and Compliance Developments Update Roundtable	1 hour
A Fireside Chat on Navigating the Cultural Challenges in Global Compliance Programs with Hui Chen, JD	.5 hour

EVALUATION FORM FOR THE PHARMA CONGRESS

You must also complete the following Pharma/Device Congress evaluation form:

Failed to Meet Expectations	Needs Improvement	Met Expectations	Exceeded Expectations	Excellent
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Overall Quality

Powerpoints

Speakers

Ease of Use

EXECUTION OF THE CONTINUING LEGAL EDUCATION CREDIT SELF REPORTING ATTENDANCE AND EVALUATION FORM FOR THE PHARMA CONGRESS

By executing this self-reporting form, the attorney hereby warrants that the information provided herein is complete, true and correct.

Executed by:

Date:

Payment must be made and the completed and executed form submitted via email to cindyramirezghc@outlook.com no later than **November 30, 2023**.