DIA CALL FOR ABSTRACTS

SUBMISSION DEADLINE: AUGUST 8

Global Pharmacovigilence and Risk Management Strategies Conference

JANUARY 27-29, 2025 | BALTIMORE, MD

CALL FOR ABSTRACTS

ABSTRACT SUBMISSION DETAILS AND GUIDELINES
SUBMISSION DEADLINE: AUGUST 8

Are you a professional involved in pharmacovigilance and risk management? If so, DIA wants to hear from YOU as we prepare for the DIA *Global Pharmacovigilance and Risk Management Strategies Conference*. This conference is a neutral event developed by regulators and industry experts discussing the updates, opportunities, and challenges alongside fresh problem-solving strategies that matter most to safety professionals.

To ensure that we have the most comprehensive and cutting-edge program, we are seeking abstract submissions from professionals like yourself, who are pushing the boundaries in their respective fields. We encourage you to submit abstracts that reflect the latest trends, innovations, and best practices in pharmacovigilance and risk management. We will be accepting the following formats:

- **Presentations:** 15-20 minute presentations to be bundled with other presentations to create a session
- **Sessions:** 60-75 minute total session
- **Short Courses:** Three-hour interactive workshop delivered in a small group format (require a separate fee from attendees)

The Program Committee is seeking abstracts on the following topics (keep in mind, business use cases and lessons learned are encouraged in all topic areas). Please note that topics in addition to those listed below, that you feel are relevant, may be submitted for evaluation and possible selection.

Interest Areas:

- Aggregate Safety Assessment Plan (ASAP)
- ChaptGPT and its effect on Safety Professionals
- Clinical Trial Safety Evaluations
- Emerging Trends in Pharmacovigilance
- Epidemiology and Pharmacovigilance
- FDA's standard tables and figures, and FMQs
- Future of the Pharmacovigilance Professional
- Good Pharmacovigilance Practices (GVP)
- Global Pharmacovigilance Updates from Regulatory Authorities
- GVP Inspections (In-Person and Virtual)
- ICH Safety Related Updates
- Implementation Strategies for Additional Risk Minimization Measures
- Literature Surveillance
- Patient Centricity in Pharmacovigilance
- Pharmacovigilance Perspectives from Startups and Small Pharmaceutical Companies
- Post-Approval Safety Studies
- Reference Safety Information (RSI) Use in Clinical Trials
- Risk Communication
- Risk Evaluation and Mitigation Strategy (REMS)
- Risk Management Plan (RMP)
- Risk Management Strategies
- Safety and Pharmacovigilance for Advanced Therapies
- Safety Governance Models
- Safety for Special Populations (e.g., children, pregnant women, etc.)
- Safety for Biosimilars and Generics
- Safety for Combination Products, Diagnostics, and Medical Devices
- Signal Detection
- Strategies to Effectively Adopt Innovation
- Use of Artificial Intelligence (AI), Machine Learning (ML), Automation, and Technology in Pharmacovigilance

Abstract Submission Deadline: Thursday, August 8

Notification: Week of September 16

Final PowerPoint Presentations Due: January 6, 2025

Please submit all abstracts online at: DIAglobal.org/Abstracts

Questions: Contact Damisha White, Project Manager at Damisha.White@DIAglobal.org

Submit Your Abstract at **DIAglobal.org/Abstracts**

CALL FOR PRESENTATION ABSTRACTS

SUBMISSION DEADLINE: AUGUST 8

GENERAL SUBMISSION REQUIREMENTS

- All submissions must be submitted online
- For complete submission requirements and to submit your abstract go to **DIAglobal.org/Abstracts**
- DIA requires speaker presentations to be non-commercialized, objective,and fair balanced. Company logos are not permitted to be included in slide presentations, per ACCME Standards for Commercial Support. In addition, speaker clothing/backgrounds may not carry logos or other company specific emblems. In this way, DIA activities will be educational, rather than commercial and promotional. Please view full policy concerning promotion of productsand services from the podium.

SUBMISSION TIPS

- Ideal submissions will contain practical content and shared experiences
- Theoretical topics and content is acceptable, however, it should be supported with proof of concepts and use cases
- Diverse topics and sessions are welcomed and encouraged within the scope of the forum
- Please select the interest area that best fits with your proposal. If your topic is relevant to more than one interest area, please indicate that in your abstract summary.
- Abstracts should be written using clear language and descriptions to provide enough clarity for the selection committee to review and understand

REQUIRED DOCUMENTATION FOR ALL ABSTRACTS

- **Participant Disclosure Information:** All abstract authors must disclose any relevant financial relationships with any commercial interest associated with this activity that exist or have existed within the past 12 months, as well as any discussion of unlabeled or unapproved drugs or devices.
- All authors must agree to the DIA Speaker Authorization for Use of Presentation Materials in order for the abstract to be a part of the Program. Accepted abstracts will be available on DIA's website for attendee download.

SUBMISSION GUIDELINES

Submitting a PRESENTATION ABSTRACT (All abstracts must be submitted online)

15-20-minute presentation, bundled with other presentations to create a session. Abstract author is considered the presenter (co-presenters are not permitted) and will be responsible for:

- Adhering to the program development guidelines and timelines
- Working with chair and other presenters in creating a balanced program offering
- Preparing and delivering a PowerPoint presentation

Submitting a SESSION ABSTRACT (All abstracts must be submitted online)

60-75-minute total session. Abstract author will be responsible for:

- Adhering to the program development guidelines and timelines
- Recruiting speakers and ensuring good representation/diversity in their selection (no more than one speaker per company per session). Maximum of 3 speakers per session. Please list all prospective speaker names and organizations in submission.
- Working with the Session Chair to communicate with speakers regarding their role in the session

Submitting a SHORT COURSE ABSTRACT (All abstracts must be submitted online)

Three-hour, interactive presentation delivered in small group format. Abstract author is considered the Short Course Lead Instructor and will be responsible for:

- Adhering to the program development guidelines and timelines
- Recruiting co-instructors and ensuring good representation/diversity in their selection
- Communicating with co-instructors regarding their role in the short course and reviewing presentation materials (note: PowerPoint presentations are required from each instructor)
- Managing the short course, including the facilitation of audience questions and interactions

SUBMISSION GUIDELINES

The following information will be requested at the time of submission. ***DIAglobal.org/Abstracts***

To streamline your submission process and avoid possible delays, DIA strongly encourages you to submit your abstract as early as possible. **Do not wait until the last day.**

Prepare your abstract in advance of accessing the DIA website. Abstract information should be copied and pasted from a prepared document as plain text. **All of the below fields are required.**

Author Information

Abstract Information

Interest Area: Choose from the drop down

Keywords: Provide one or more keywords to highlight your abstract.

Examples of keywords: Personalized Medicine, Health Technology Assessment, etc. (100 characters)

Level of Difficulty: Beginner, Intermediate, or Advanced

Learning Objectives: Provide 2-3 learning objectives that clearly explain what participants should be able to do after attending this event. For a list of suggested verbs to create these objectives, click here. (400 Characters)

Overview: *Please provide 2-3 sentences summarizing your abstract. This summary will be used as the overview description in the DIA program for marketing purposes (250 Characters including spaces)

Abstract Details: *Please provide complete details about your abstract. Information such as scientific, technical, process issues, design/methods, results/outcomes, case studies, statistics, key findings, etc., that would support your proposal should be included here. This information will be used by the Program Committee to learn more about the purpose of your abstract. Is there an interactive component to your topic? If so, please indicate in the abstract details how you would be able to include an interactive learning experience for attendees.
(2000 Characters including spaces)

Submit Your Abstract at **DIAglobal.org/Abstracts**