

Artificial Intelligence in Pharmacovigilance

Virtual Live Training Course

10-12 September 2024 | 15:00-19:30 CEST



Overview

A practical orientation course for the next level of pharmacovigilance technology as it is being implemented across the industry. Over the last 7 years, major advances in artificial intelligence (AI) technology have reached the pharmacovigilance (PV) industry and are changing how we operate with ever-increasing speed. A new skillset for pharmacovigilance professionals is required.

The course is designed to give all senior pharmacovigilance professionals sufficient orientation to make an informed decision and use the new technology in a smart and compliant way.

The course is given by pharmacovigilance and IT veterans who are fearless in enabling sustainable innovation and have hands-on experience with AI tools in pharmacovigilance. A regulatory point of view is represented in panel discussions.

Learning Objectives

At the conclusion of this virtual live training course, participants will be able to:

- Understand how main AI methodologies work - machine learning, natural language processing, classifications, large language models, other ML
- Clearly explain pros and cons of AI models that are already being used in PV, including OpenAI ChatGPT (3.5 and 4.0), Google Gemini, Anthropic Claude 2, MS Copilot etc.
- Identify and manage risks to your data security, data privacy, bias and ethical concerns
- Discuss the management perspective and challenges - people, skills, governance, audits, and compliance
- Review what today's PV IT platforms deliver in terms of implementations, AI, and automation, including major systems such as Oracle Argus, IQVIA IVP, RxLogix, LifeSphere by ArisGlobal etc.
- Brainstorm what would be the perfect PV IT intelligent system of the future
- Roadmap for successful AI implementation in your PV system

Who Will Attend

Pharmacovigilance and IT professionals with 2+ years of PV experience who wish to gain a practical overview of all AI technologies coming to the pharmacovigilance industry.

Faculty

Jan Petracek

Director
Institute of Pharmacovigilance
EU

Robert Scheiner

CIO
iVigee
USA

Phil Tregunno

Deputy Director - Patient Safety Monitoring
MHRA
United Kingdom

Key Topics

- PV journey from manual process through RPA to Generative AI
- Foundations of AI in PV: Supervised vs Unsupervised, NLP, LLMs...
- Ensuring compliance while using AI
- Today's PV interfaces and systems using AI
- PV AI barriers and challenges
- Developing regulatory environment for AI in PV
- Near future PV IT developments

DAY 1

15:00 WELCOME AND INTRODUCTION

15:30 SESSION 1

PV JOURNEY FROM MANUAL THROUGH ROBOTIC PROCESS AUTOMATION (RPA) TO AI

Jan Petracek

- Business experience from 7-year-long development and implementation of intelligent automation in pharmacovigilance
- What is changing in pharmacovigilance and why, the main trends and competencies needed for the near future
- AI initiatives from major regulatory authorities and CIOMS

17:00 BREAK

17:30 SESSION 2

AI AND NATURAL LANGUAGE LEARNING (NLP) IN PV

Robert Scheiner

- Main principles of AI technology, especially in the PV context
- Main use cases where AI is used within PV, where it augments human workflows and vice versa - where humans may complement what AI is primarily (or will be soon) doing

19:00 DISCUSSION AND Q&A

19:30 END OF DAY 1

DAY 2

15:00 SESSION 3

COMPLIANCE AND MANAGEMENT PERSPECTIVE ON IMPLEMENTING INTELLIGENT AUTOMATION

Jan Petracek

- Business case and management views - how senior management and CEOs view investments in innovative technology
- Prepare your departments, teams, hire the right talent, and create your roadmaps
- What are some of the myths on either side - management and business and how might we bridge them
- Exercise a clever approach to validation and quality management requirements to stay compliant while using the most modern technology
- Defence of more intelligent data management practices in front of auditors and inspectors

16:30 BREAK

17:00 SESSION 4

TODAY'S PV INTERFACES AND SYSTEMS

Robert Scheiner

- Current PV IT platforms and patterns
- What to expect from FDA, EMA, and MHRA in terms of intelligent automation
- Ongoing Prove of Concepts (PoCs) and development AI labs at major and smaller pharma, regulators, and service organisations

18:30 DISCUSSION AND Q&A

19:30 END OF DAY 2

DAY 3

15:00 SESSION 5

NEAR FUTURE PV IT DEVELOPMENTS

Robert Scheiner

- Future of PV IT platforms
- Technology architectural patterns supporting high scalability, throughput, and flexibility
- Wise and informed decisions about the future use of technology like NLP, RPA, and advanced ChatGPT-like technologies in their daily work
- Distinguish buzz from reality when assessing options
- How to go from labs to future production and build a real ML Ops
- How to build in explainability from early on and avoid compliance issues later

16:30 BREAK

17:00 SESSION 6

DEVELOPING REGULATORY ENVIRONMENT FOR AI IN PV

Jan Petracek, Robert Scheiner and Phil Tregunno

- Case studies
- View of leading regulators
- 2023 as a breakthrough year and what will happen in 2024+
- Discussion and Q&A

19:30 END OF THE VIRTUAL LIVE TRAINING COURSE



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To take advantage of this offer, please print the registration form for EACH of the four registrants from your company. Include the names of all four group registrants on each of the forms and return them together via email to basel@diaglobal.org.

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To test your system compatibility, please click on the link: <https://diaglobal.zoom.us/test>

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Continuing Education

The Swiss Association of Pharmaceutical Professionals (SwAPP) and the Swiss Society for Pharmaceutical Medicine (SGPM) have accredited this training course with 10 credits.



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REGISTRATION FORM

AI in PV | Virtual Live Training Course | # 24559
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REGISTRATION FEES

Registration fee includes full admission to virtual course, electronic access to training course materials. **Please note that the full amount must be received by DIA by commencement of the course to get the electronic access to the material.** Please check:

FEES	MEMBER EARLY-BIRD valid until 16 Jul 2024	MEMBER valid from 17 Jul 2024	NON-MEMBER
INDUSTRY/ REPRESENTATIVE	€ 1'215.00 <input type="checkbox"/>	€ 1'350.00 <input type="checkbox"/>	€ 1'610.00 <input type="checkbox"/>
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A special discount for SMEs on the standard fee is available for a limited number of places. To prove your status as an SME, a confirmation of the European Medicines Agency is necessary. Please contact DIA for more information.

All registration fees are subject to VAT if applicable.

Please enter your company's VAT number: _____

If DIA cannot verify your membership upon receipt of registration form, you will be charged the non-member fee.

Payment is due 30 days after registration and must be paid in full by commencement of the course.

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Email: Basel@DIAGlobal.org **Mail:** DIA, KÜchengasse 16, 4051 Basel, Switzerland

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ATTENDEE DETAILS

Please complete in block capital letters or attach the attendee's business card here.

Prof Dr Ms Mr

Last Name

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Attendee email required for course material access

TERMS AND CONDITIONS

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- Industry (Member/Non-member) € 200.00
- Academia/Charitable/Government/Non-profit (Full-time) (Member/Non-member) € 100.00

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Date Signature