

# Artificial Intelligence in Pharmacovigilance

**Virtual Live Training Course** 

29-31 January 2024 13:00-17:30 CET

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# **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.

- 50:50 mix of technology vs business contents. Technology topics require no prior technical knowledge; all are explained clearly.
- Each day mixes business and technology topics.

# **LEARNING OBJECTIVES**

- Understand how main Al 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 Bard, 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, Al, 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

# **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

# 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 UK



# DAY 1

# 13:00 WELCOME AND INTRODUCTION

13:30 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
- Al initiatives from major regulatory authorities and CIOMS

# 15:00 BREAK

15:30 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

17:00 DISCUSSION AND Q&A

17:30 END OF DAY 1

### DAY 2

# 13:00 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

# 14:30 BREAK

15:00 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 Al labs at major and smaller pharma, regulators, and service organisations

16:30 DISCUSSION AND Q&A

17:30 END OF DAY 2

# 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.



Unless otherwise disclosed, DIA acknowledges that the statements made by speakers are their own opinion and not necessarily that of the organisation they represent, or that of the DIA. Speakers and agenda are subject to change without notice. Recording during DIA sessions is strictly prohibited without prior written consent from DIA.

### DAY 3

# 13:00 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

# 14:30 BREAK

# 15:00 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

# 17:30 END OF THE VIRTUAL LIVE TRAINING COURSE

# **Group Discounts**

Register 3 individuals from the same company for the same course and receive complimentary registration for a 4th!

- All 4 individuals must register and prepay at the same time no exceptions
- DIA will apply the value of the lowest applicable fee to this complimentary registration; it does NOT include fees for optional events or DIA membership
- You may substitute group participants of the same membership status at any time; however, administrative fees may be incurred.

Group registration is not available online and does not apply to the already discounted fees for government or charitable nonprofit/academia.

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 <a href="mailto:basel@diaglobal.org">basel@diaglobal.org</a>.

# **Technical Requirements**

In preparation for the course, please create a free account (or use existing account) for OpenAI GPT and Google Bard!

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

For further information on system requirements, please visit the website:

https://www.diaglobal.org/General/System-Requirements

# | Plan Your Team's Professional Development

Why not take advantage and train your whole department (or even across different departments!) and benefit from increased:

- Focus
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For more information please contact Basel@diaglobal.org.

# **REGISTRATION FORM Virtual Live Training Course**

Artificial Intelligence in Pharmacovigilance # 24537 29-31 January 2024 13:00-17:30 CET



# **REGISTRATION FEES**

Registration fee includes full admission to virtual course, and electronic access to 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 4 Dec 2023	MEMBER valid from 5 Dec 2023	NON- MEMBER
INDUSTRY/ REPRESENTATIVE	€ 1'215.00 🗖	€ 1'350.00 🗖	€ 1'610.00 🗖
ACADEMIA/CHARITABLE/GOVERNMENT/NON-PROFIT (FULL-TIME)	NA	€ 675.00 🗖	€ 935.00 🗖

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.

### DIA MEMBERSHIP

All nonmember fees include a one year DIA membership, at no additional cost. Explore membership benefits at <u>DIAglobal.org/Membership</u>.

DIA membership will renew automatically at the end of the complimentary membership term, at the then current membership rates. You may cancel automatic membership renewal at any time by accessing your account online at <a href="DIAglobal.org">DIAglobal.org</a>. If you would like to decline complimentary membership, please indicate your preference below.

☐ I would like to decline a one year complimentary DIA membership.

The DIA Contact Centre Team will be pleased to assist you with your registration from Monday to Friday between 09:00 and 17:00 CE(S)T. Tel.:+41 61 225 51 51

Email: Basel@DlAglobal.org Mail: DIA, Küchengasse 16, 4051 Basel, Switzerland Web: www.DlAglobal.org

# **TERMS AND CONDITIONS**

### **Cancellation Policy**

All cancellations must be made in writing and be received at the DIA office four weeks prior to the event start date. Cancellations are subject to an administrative fee:

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

If you do not cancel four weeks prior to the event start date and do not attend, you will be responsible for the full registration fee.

DIA reserves the right to alter the venue and dates if necessary. If an event is cancelled or postponed, DIA is not responsible for airfare, hotel or other costs incurred by registered attendees. Registered attendees are responsible for cancelling their own hotel and travel reservations.

# Transfer Policy

You may transfer your registration to a colleague prior to the start of the event but membership is not transferable. Substitute attendees will be responsible for the nonmember fee, if applicable. Please notify the DIA office of any such substitutions as soon as possible.

# **Event Stream and Recording**

If you attend a DIA event, we make video and audio recordings of events (both face-to-face and online) that may include your participation in the event, including your image, questions and comments. To view our full photography and video recording policy, click <a href="https://www.diaglobal.org/general/photography-policy">https://www.diaglobal.org/general/photography-policy</a>.

# **Privacy Policy**

DIA respects the privacy of all of its members and customers. To view our privacy policy, click <a href="https://www.diaglobal.org/about-us/privacy-policy">https://www.diaglobal.org/about-us/privacy-policy</a>. You agree that your personal data will be transferred to DIA in the US.

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