EudraVigilance Data Analysis System (EVDAS): Practical approach on use for Signal Management in the EU
Virtual Live Training Course

24-25 June 2021 09:00-13:00 CEST

OVERVIEW

This virtual live training course will teach concepts, access policy, and use of the EudraVigilance Data Analysis System (EVDAS) for signal detection. Experienced trainers using EVDAS on a regular basis will share practical advice on how to download and interpret the data and use it for signal detection.

The course is based on the latest guidelines on EU Good Pharmacovigilance Practices (GVP): Module IX – Signal management, Commission Implementing Regulation (EU) No. 520/2012.

Time has been set aside for practical exercises, questions, and discussions.

LEARNING OBJECTIVES

After the completion of this virtual live training course, participants will be able to:

- Understand regulatory requirements for Signal management in Europe (GVP IX and Addendum I)
- Describe principles of screening EudraVigilance for adverse reactions and the EudraVigilance data access policy
- Identify levels and methods of access to Individual Case Safety Reports (ICSRs) data and understand the terminology of EVDAS
- Learn to use active substance grouping reports, electronic reaction monitoring reports (eRMRs)
- Analyse eRMRs with various reference periods and line listings and document your assessments
- Identify potential signals and know how to escalate them to the National Competent Authorities (NCAs) and the European Medicines Agency (EMA)

KEY TOPICS

- GVP IX
- Principles of access in EudraVigilance
- EVDAS
- eRMR analysis and documentation
- Line listing analysis
- Literature search including non-clinical safety findings
- How to escalate signals, e.g. stand-alone notification, ESI

WHO WILL ATTEND

This virtual live training course is aimed at professionals who work in:

- Pharmacovigilance (including QPPVs)
- Drug safety and patient safety risk management
- Information Technology
- Regulatory Affairs
- Pharmacovigilance consultancies
- Quality and Compliance
- Legal

Course level: Intermediate, for professionals with 2-3 years of experience in Pharmacovigilance, or related functions who are working in Pharmacovigilance in the area of signal management.

FACULTY

Calin A. Lungu
CEO
Drug Development Consulting Services
Luxembourg

Vojtech Kvita
Pharmacovigilance Consultant
Czech Republic

DAY 1

09:00 WELCOME AND INTRODUCTION

09:30 SESSION 1

GVP IX

10:15 SESSION 2

ADDENDUM I TO GVP IX AND SCREENING FOR ADVERSE
REACTIONS IN EUDRAVIGILANCE

11:00   BREAK

11:15 SESSION 3

PRINCIPLES OF ACCESS IN EUDRAVIGILANCE

12:00 SESSION 4

EVDAS PILOT

12:45 QUESTIONS AND ANSWERS

13:00 END OF DAY 1

DAY 2

09:00 SESSION 5

EVDAS

09:45 SESSION 6

PRACTICAL EXERCISES
- eRMR analysis and documentation
- Line listing analysis
- Communication of potential duplicates to the EMA

11:00   BREAK

11:15   SESSION 6 CONTINUED

PRACTICAL EXERCISES
- Other sources of information
  - CMDh list of safety concerns
  - Referrals page on the EMA website
  - List of signals discussed by the PRAC since 2012
  - Literature search including non-clinical safety findings
- How to escalate signals, e.g. stand-alone notification, ESI

12:45 QUESTIONS AND ANSWERS

13:00 END OF THE TRAINING COURSE

Group Discounts

**Register 3 individuals from the same company 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 basel@diaglobal.org.

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
- Flexibility
- Convenience
- Cost Effectiveness

For more information please contact Basel@DIAglobal.org

About DIA

DIA is the global connector in the life sciences product development process. Our association of more than 18,000 members builds productive relationships by bringing together regulators, innovators, and influencers to exchange knowledge and collaborate in an impartial setting. DIA's network creates unparalleled opportunities for exchange of knowledge and has the inter-disciplinary experience to prepare for future developments.

The dedicated efforts of DIA staff, members and speakers enable DIA to provide a comprehensive catalogue of conferences, workshops, training courses, scientific publications and educational materials. DIA is a global community representing thousands of stakeholders working together to bring safe and effective products to patients.

DIA is an independent, non-profit organisation has its Global Center in Washington, DC, USA with the European office in Basel, Switzerland, and additional regional offices in Horsham, Pennsylvania, USA; Tokyo, Japan; Mumbai, India; and Beijing, China

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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 6 credits.

SwAPP | Swiss Association of Pharmaceutical Professionals

Technical Requirements

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

Operating Systems
- Windows: XP 32-bit (SP3), 2003, Vista 32-bit/64-bit, Windows 7 32-bit/64-bit
- Mac OS X: 10.5, 10.6, 10.7
- Linux: 32-bit Ubuntu 10.x,11.x 32-bit Fedora 15/16, 32-bit Red Hat 5/6, 32-bit OpenSuSE 11.4

Minimum System Requirements
- Windows: Processor – Requires Sun Java 5 or higher, Recommend ActiveX be enabled for Internet Explorer
- Mac OS X: Processor – JavaScript and cookies enabled, Requires Apple Java 5 or higher, No support for Remote Access
- Linux: Processor – JavaScript and cookies enabled, Requires Apple Java 5 or higher, No support for Remote Access

Browsers
- Windows: Internet Explorer 6, 7, 8, 9, (Win7 Only), Firefox latest (32-bit), Chrome latest
- Mac OS X: Safari 4-Mar; Firefox 2/3/3.5
- Linux: Mozilla 1.7, Firefox 2/3/3.5

Internet Connection Speed
- Windows: Intel or AMD processor (1GHz or faster), At least 512 MB RAM (at least 2 GB RAM for Vista)
- Mac OS X: Intel processor, At least 512 MB RAM
- Linux: At least 512 MB RAM

Display

800x600 pixel resolution or greater (1024x768 pixels recommended).

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 29 April 2021	MEMBER valid from 30 April 2021	NON-MEMBER
INDUSTRY/ REPRESENTATIVE	€ 720.00 ❑	€ 800.00 ❑	€ 985.00 ❑
ACADEMIA/CHARITABLE/GOVERNMENT/NON-PROFIT (FULL-TIME)	NA	€ 400.00 ❑	€ 585.00 ❑
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 DIAglobal.org/Membership.

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 DIAglobal.org. 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 CET. **Tel.** :+41 61 225 51 51

Email: Basel@DIAglobal.org Mail: DIA, Küchengasse 16, 4051 Basel, Switzerland
Web: www.DIAglobal.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 non-member 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 https://www.diaglobal.org/general/photography-policy.

Privacy Policy
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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

First Name

Job Title

Company

Address

Postal Code

City

Country

Telephone Number

Attendee email required for course material access

PAYMENT METHODS

Credit cards: Payments by VISA, Mastercard or AMEX can be made by completing the details below. Please note that other types of credit card cannot be accepted.

❑ **Please charge my** ❑ VISA ❑ MC ❑ AMEX

Card N°

Exp. Date ☐☐ / ☐☐

Cardholder's Name

❑ **Bank transfers:** When DIA completes your registration, an email will be sent to the address on the registration form with instructions on how to complete the bank transfer. Payments in EURO should be addressed to "Account Holder: DIA." Please include your name, company, Course ID #21559 as well as the invoice number to ensure correct allocation of your payment.

Payments must be net of all charges and bank charges must be borne by the payer. **If you have not received your confirmation within five working days, please contact DIA.**

By signing below, I confirm that I agree with DIA's Terms and Conditions of booking. These are available from the office or on http://www.diaglobal.org/EUTerms

Date	Signature