DIA LEARNING

Mastering GDPR Compliance in Clinical Trials

Virtual Live Training Course

16 April 2026 | 09:00-13:30 CEST

Overview

This virtual live training course covers the essential concepts of EU and UK data protection laws that sponsors need to consider when running clinical trials.

Understanding and complying with data protection laws, such as the General Data Protection Regulation (GDPR), is crucial for clinical trial sponsors to ensure legal compliance, protect participant rights, maintain data security, and uphold ethical standards in clinical research. It is also often a prerequisite for obtaining regulatory approval to conduct the clinical trial.

This training course will equip participants with the knowledge and tools to navigate GDPR requirements confidently and ensure clinical trials meet the (sometimes) onerous GDPR requirements. Case studies will be used throughout to provide context.

Participants benefit from the hands-on experience of instructors who are experts in this area.

Learning Objectives

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

- Appreciate the national-level nuances when it comes to identifying the legal bases for processing and meeting the transparency requirements under the GDPR and learn how best to reflect these in the ICF/PIS to ensure (as far as possible) a streamlined approach to a multi-jurisdiction study
- Understand the implications for the potential differing roles of a site i.e., in different countries and how this is in turn reflected in the CTA and ICF/PIS
- Recognize key considerations and necessary contractual provisions when engaging CT vendors
- Be able to identify cross-border data flows, and the mechanisms to address including, the nuances in approach in clinical trials
- Understand what constitutes a personal data breach and the potential consequences for this under the GDPR
- Identify key actions for GDPR-compliant processing of study personnel data
- Learn how to meet data treatment and accountability requirements in clinical studies including, how to conduct a DPIA and the potential expectations of sites/ECs in relation to the same

Who Will Attend

This course is designed for professionals involved in:

- Commercial and Non-commercial sponsors of clinical trials
- CROs
- Consultants

Faculty

Francesca Blythe
Partner: Healthcare, Privacy and Cybersecurity
Sidley Austin, United Kingdom

Eleanor Dodding
Senior Managing Associate
Sidley Austin, United Kingdom

PharmaTrain
MASTERING MEDICINES DEVELOPMENT
CENTRE OF EXCELLENCE

16 April 2026

09:00 WELCOME AND INTRODUCTIONS

09:15 SESSION 1

- GDPR overview
- Pseudonymised and anonymised personal data
- Key considerations from a clinical trials perspective:
    - Lawful processing of study subject data
    - Transparency – study subjects and study personnel

10:15 SESSION 2

- Role of the site
- Engaging vendors
- Cross border transfers

11:15 BREAK

11:45 SESSION 3

- Safety data reporting activities
- Data treatment principles and accountability
- Security and personal data breaches

13:15 Q&A

13:30 END OF THE TRAINING COURSE

Group Discounts

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

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.

**Terms and Conditions apply. Please contact DIA EMEA office for more information.*

Customized Professional Development for Your Team

Get a customized training for your department (or even across different departments!) and benefit from increased:

- Knowledge of a topic of your choice
- Flexibility & Convenience
- Cost Effectiveness

Or explore eLearning to allow self-paced learning.

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

Technical Requirements

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

Continuing Education

The Swiss Association of Pharmaceutical Professionals (SwAPP) and the Swiss Society for Pharmaceutical Medicine (SGPM) have accredited this training course with 3.5 credits.

REGISTRATION FORM

GDPR Compliance in CTs | Virtual Live Training Course | # 26532
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DIA LEARNING

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 19 Mar 2026	MEMBER valid from 20 Mar 2026	NON-MEMBER
INDUSTRY/ REPRESENTATIVE	€ 450.00 ❑	€ 500.00 ❑	€ 760.00 ❑
ACADEMIA/CHARITABLE/GOVERNMENT/NON-PROFIT (FULL-TIME)	NA	€ 250.00 ❑	€ 510.00 ❑
A special discount is available for organisations which are listed in the EMA SME register. Number of discounted seats is limited.			

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 CE(S)T. 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
DIA respects the privacy of all of its members and customers. To view our privacy policy, click https://www.diaglobal.org/about-us/privacy-policy.

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 METHOD

DIA accepts only Credit Card as a payment method.

Payments by VISA, Mastercard or AMEX are accepted. Other types of credit card are not accepted.

You will receive a payment link in the coming days to complete the payment.

Please complete payment within 7 days of receipt of the payment link.

Payments will be net of all charges and bank charges will be borne by the payer.

If you have not received your confirmation within five working days, please contact basel@diaglobal.org.

By signing below, I confirm that I read and agree with DIA's Terms and Conditions of booking.

These are available from the office or online by clicking:

http://www.diaglobal.org/EUterms

Date	Signature