ECA Academy
Your GMP/GDP Information Source

European Events

2020/2021

Quality Assurance
Quality Control
Sterile/Aseptic Manufacturing
APIs/Excipients
Computer Validation
Data Integrity
Production/Engineering
Microbiology
Biopharmaceuticals
Validation
GDP
Others

www.gmp-compliance.org

European Compliance Academy
P.O. Box 10 21 68
69011 Heidelberg, Germany
info@gmp-compliance.org

ECA GMP/GDP Certification Programme

One reason for the ECA Academy's excellent reputation is its high-quality Certification Programme. In the past years, thousands of GMP and GDP professionals already relied on the programme to advance their knowledge and to get an additional qualification - and completed the ECA Certification Level.

This comprehensive qualification curriculum comprises 15 programmes, allowing professionals to combine several seminars according to their fields of interest.

New – Attend Online Courses for Certification

Currently many training courses are offered live online. Also some training courses will be offered on demand (recording of live online training courses). These online courses will be recognized for the ECA GMP/GDP Certification Programme as well!

Objectives

Highly qualified personnel is crucial within the field of GMP-compliant manufacturing of APIs and drugs. Although college and university education provide the scientific basis, a continuous advanced training is essential.

And this is where the ECA Academy GMP Certification Programme fills the gap. It offers modular training with an industry-known certification at the end. For most of the Certification Options you will need to attend 3 training courses of a comprehensive number of events. You can:

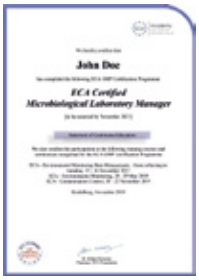
- select courses according to your individual professional demands
- suit the course registration to your company's necessities, i.e. usually there is a period of several months between the courses. If two dates are too close together, you can attend one course in the following year.

Recognition of the Certification Programme

The ECA Academy enjoys an excellent reputation within Europe's pharmaceutical industry. The Certification Programme represents a useful completion of your college and university education. The ECA Certification Programme is the largest programme of its kind in Europe.

Which Training Courses are Recognized for the Certification Programme?

To find out which training courses are recognized for the certification program, please click on the different certification options listed below. There you will find a link to the currently recognized training courses. Please note: On each program flyer for the Training Courses you will find an indication for which certification it is recognized.



How to Obtain the Certificate?

To obtain the certificate, please send an e-mail prior or after the 3rd event you attended to info@gmp-compliance.org. This e-mail should also list the courses you attended in the past. You will then get your certificate during the 3rd course. The Certificate will not only include the title of the Certification Programme, but also lists all ECA courses and conferences you attended. Thus, it will serve as a valid documentation for your continuous GMP and GDP education.

Continuous GMP/GDP Certification – a service offered by the ECA Academy

In order to reflect the development of a continuous advanced education for GMP and GDP professionals from 2014 on the ECA Academy will issue the Certificates of the Certification Programme with a validity date. Every new Certificate (see list of the Certificates below) will be valid for two years. To renew it the applicant has to join one ECA Training Course or Conference within that period. Applicants are quite flexible in selecting training courses or conferences for the renewal. For example, a Validation Training Course or one on OOS Results will also be recognised for the renewal of the ECA Certified QA Manager Certificate, although these courses are not specifically recognized for the QA Manager Certificate (the same applies to all other certificates). This flexibility takes into account that applicants will broaden their knowledge in GMP Compliance.

The renewal process is easy and will be managed by the ECA Academy. If you have obtained one of the Certificates below in 2014 or later you will automatically receive a new version of your certificate which contains the new two years Certification statement every time you will participate in an ECA Training Course or Conference. Please contact us at info@gmp-compliance.org if you have any further question.

We offer the following modules:

- **ECA Certified Biotech Manager**
- **ECA Certified Pharmaceutical Development Manager**
- **ECA Certified Sterile Production Manager**
- **ECA Certified Computer Validation Manager**
- **ECA Certified Microbiological Laboratory Manager**
- **ECA Certified Technical Operations Manager**
- **ECA Certified Regulatory Affairs Manager**
- **ECA Certified API Production Manager**
- **ECA Certified QA Manager**
- **ECA Certified Validation Manager**
- **ECA Certified Quality Control Manager**
- **ECA Certified GMP Auditor**
- **ECA Certified GDP Compliance Manager**
- **ECA Certified Packaging Manager**
- **ECA Data Integrity Manager**

European Conferences & Education Courses

in co-operation with CONCEPT HEIDELBERG

ECA Certified Pharmaceutical Development Manager

- **ICH Q8 / ICH Q11 Training Course**
  Live Online Training on 8/9 October 2020
- **GMP meets Development**
  Live Online Training from 17-19 November 2020
- **Stability by Design**
  28/29 April 2021, Vienna, Austria

ECA Certified Computer Validation Manager

- **Computerised Systems Validation:**
  - Leveraging Suppliers
  - Computerised System Validation Master Class
  Live Online Training on 27 & 28-30 October 2020
- **SAP – Validation and GMP Compliance**
  Live Online Training on 10/11 November 2020
- **IT Infrastructure - Qualification and Operation in a GMP Environment**
  Live Online Training on 12/13 November 2020
- **Computerised Systems Validation: Introduction to Risk Management & The GAMP®5 Approach**
  Live Online Training on 17 and 18-20 November 2020
- **Cloud Computing in a GxP Environment**
  24-26 February 2021, Berlin, Germany
- **Data Integrity**
  Requirements for a GMP-compliant Data Life Cycle
  17 – 19 March 2021, Barcelona, Spain
  With an optional full-day pre-course session Audit Trail Review on 16 March 2021

ECA Certified Data Integrity Manager

- **Lab Data Integrity - Meeting FDA and EU Concerns**
  Live Online Training on 22-24 September 2020
- **Data Integrity and Good Documentation Practice**
  Live Online Training from 24-26 November 2020
  06-08 May 2021, Heidelberg, Germany
- **Data Integrity Audits & Inspections**
  Live Online Training on 24/24 November 2020
- **Data Integrity & optional pre-course session Audit Trail Review**
  Live Online Training on 08 - 11 December 2020
  16 - 19 March 2021, Barcelona, Spain
- **HPLC Data Integrity - Ensuring Control of Chromatographs, Integration and Results**
  02/03 February 2021, Prague, Czech Republic
- **Audit Trail Review for Computerised Systems in Analytical Laboratories**
  03/04 March 2021, Prague, Czech Republic
- **Data Integrity Quality Oversight in the QC Laboratory**
  13/14 April 2021, Hamburg, Germany
  with an optional post-conference Workshop „Audit Trail Review for CDS / Laboratory Systems", 15 April 2021, Hamburg, Germany
- **Lab Data Integrity**
  Part 1: Establishing the Controls for Ensuring Laboratory Data Integrity, 14/15 September 2021, Prague, Czech Republic
  Part 2: Self Inspections and Audits to Confirm Effective Data Integrity Controls, 15/16 September 2021, Prague, Czech Republic
- **Lab Data Integrity Master Class**
  12 – 14 October 2021, Berlin, Germany

ECA Certified Technical Operations Manager

- **Product Transfer**
  Live Online Training on 20/21 October 2020
- **GMPs for Equipment, Utilities and Facilities**
  Live Online Training from 6-8 October 2020
- **Control of Parenterals**
  Container-/Closure-Integrity Testing
  Visual Inspection Systems
  Live Online Conference from 6 - 8 October 2020
- **Pharmaceutical Water**
  Live Online Training on 1/2 December 2020
- **Clean Rooms & HVAC Systems**
  Live Online Training on 15/16 December 2020
- **Spray Drying - with Hands-On Spray Drying Course at the GEA Niro Site**
  02 - 04 March 2021, Copenhagen, Denmark

ECA Certified Regulatory Affairs Manager

- **Drug Master File Procedures in the EU, the US and Japan**
  Live Online Training on 29/30 September 2020
- **API Regulatory Starting Materials**
  Live Online Training on 13/14 October 2020
- **GMP meets Regulatory Affairs**
  Live Online Training on 15/16 October 2020
- **How to provide Process Validation Data in a regulatory submission**
  Live Online Training on 04/05 November 2020
- **How to write the Quality Part of an IMPD**
  Live Online Training on 08/09 December 2020
  06/07 October 2021, Hamburg, Germany

ECA Certified Biotech Manager

- **Quality, Safety and GMP for Advanced Therapy Medicinal Products (ATMPs)**
  Live Online Conference on 10/11 November 2020
- **GMP for Vaccine Manufacturers**
  Live Online Training on 24/25 November 2020
- **Pharmaceutical Biotechnology for Non-Biotechnologists**
  Live Online Training on 15/16 December 2020
- **Annex 2 + Co - GMP Compliance for Biopharmaceuticals**
  16/17 March 2021, Munich, Germany
- **Blood and Plasma – Audits and Inspections**
  03/04 November 2021, Berlin, Germany

ECA Certified GMP Auditor

- **The GMP Auditor**
  Live Online Training on 10-12 November 2020
  16-18 March 2021, Amsterdam, The Netherlands

European Conferences & Education Courses

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ECA Certified Sterile Production Manager

- **Environmental Monitoring**
  Live Online Training on 22/23 September 2020
- **Annex 1 Conference**
  Live Online Confernce on 10/11 November 2020
- **Isolator Technology Workshop – Manufacturing Isolator**
  24/25 November 2020, Basel, Switzerland

ECA Certified Microbiological Laboratory Manager

- **Contamination Control Strategies**
  Live Online Training on 18-20 November 2020
- **Modern Microbiology Laboratory**
  Pharmacopoeial and GMP Compliance
  Live Online Training from 08-10 December 2020,
- **Microbiology for Non-Microbiologists**
  02/03 February 2021, Antwerp, Belgium
- **Virus and TSE Safety made simple**
  02/03 March 2021, Berlin, Germany
- **Bioburden Workshop**
  08/09 June 2021, Berlin, Germany

ECA Certified GDP Compliance Manager

- **The GDP Compliance Manager**
  Live Online Training on 06/07 October 2020
- **The GDP Audit**
  Live Online Training on 28/29 October 2020
- **Ambient Transport and Cold Chain**
  Live Online Training on 04/05 November 2020
- **The Responsible Person for Good Distribution Practices (GDP)**
  24/25 March 2021, Vienna, Austria

ECA Certified API Production Manager

- **ICH Q7 Compliance for APIs Manufactured by Chemical Synthesis**
  30 November – 02 December 2020, Berlin, Germany
- **ICH Q7 Compliance for APIs Manufactured by Cell Culture/Fermentation**
  30 November – 02 December 2020, Berlin, Germany
- **ICH Q7 Auditor Training Course**
  02 – 04 December 2020, Berlin, Germany
- **Handling of Foreign Particles in APIs and Excipients**
  26/27 January 2021, Berlin, Germany

ECA Certified Quality Assurance Manager

- **GMP for Medical Devices**
  Live Online Training on 13/14 October 2020
- **Quality Risk Management - An ICH Q9 Training Course**
  Live Online Training on 14/15 October 2020
- **The GMP Compliance Manager**
  Live Online Training on 4/5 November 2020
- **Complaint Handling and Recall Management**
  Live Online Training on 17/18 November 2020
- **Inspection Management**
  Live Online Training on 17-18 November 2020
- **Pharmaceutical Contracts: GMP and Legal Compliance**
  09/10 February 2021, Amsterdam, The Netherlands
- **Combination Products**
  23/24 February 2021, Heidelberg, Germany
- **GMP and Quality Requirements for Radiopharmaceuticals**
  23/24 March 2021, Berlin, Germany
- **Improve your Quality Reviews**
  25/26 March 2021, Prague, Czech Republic
  With an optional pre-course Session on 24 March:
  Statistical Process Evaluation and Reporting
- **Lean GMP Systems**
  06/07 May 2021, Vienna, Austria
- **Quality Oversight**
  19/20 May 2021, Berlin, Germany
- **Deviation Management and CAPA**
  01/02 June 2021, Prague, Czech Republic
- **Efficient Supplier Qualification**
  23/24 June 2021, Copenhagen, Denmark
  With an optional pre-course Session on 22 June:
  What you need to know about Suppliers in China and India

ECA Certified Validation Manager

- **Process Validation in the light of the revised Annex 15 and FDA Requirements**
  Live Online Training on 6/7 October 2020
  27/28 April 2021, Prague, Czech Republic
  06/07 October 2021, Berlin, Germany
- **Launch Conference – Final version: ECA´s Integrated Qualification and Validation Guide – working with suppliers towards modern qualification and validation**
  27/28 October 2020 in Berlin, Germany
  or broadcasted live to your desk!
- **Understanding Design of Experiments (DoE) in the Pharmaceutical Industry**
  03/04 November 2020, Heidelberg, Germany
  12/13 October 2021, Heidelberg, Germany
- **The Validation Manager in the Pharmaceutical Industry**
  17-19 March 2021, Heidelberg, Germany
- **Ongoing/Continued Process Verification**
  Live Online Training on 17 November 2020
  26/27 May 2021, Heidelberg, Germany
- **Analytical Methods for Cleaning Validation**
  29/30 June 2021, Heidelberg, Germany

ECA Certified Quality Control Manager

- **Ph. Eur., USP and other Pharmacopoeias**
  Dealing with different compendial methods
  Live Online Training on 22/23 September 2020
- **FDA Compliance in Analytical Laboratories**
  Live Online Training on 6/7 October 2020
- **Bioassays and Bioanalytics**
  Live Online Training on 20/21 October 2020

European Conferences & Education Courses

in co-operation with CONCEPT HEIDELBERG

- **Stability Testing for Biological/Biotechnological Drug Substances and Drug Products**
  Live Online Training on 22 October 2020
- **Design Controls for Drug – Device Combination Products**
  Live Online Training on 21/22 October 2020
- **Validation of Analytical Test Procedures & Measurement Uncertainty**
  23-25 November 2020, Berlin, Germany
- **Reduced Sampling / Reduced Testing**
  Live Online Training on 26/27 November 2020
- **The Impurities Workshop**
  01-03 Dezember 2020, Heidelberg, Germany
  Part I - General Strategies for Investigation & Control of Impurities
  Part II - Nitrosamine Impurities
  Part III - Elemental Impurities
- **Practical Statistical Tools for Analytical Laboratories**
  09/10 December 2020, Heidelberg, Germany
- **Leachables & Extractables**
  Testing & Assessment from Packaging to Single Use
  27 - 29 April 2021, Vienna, Austria
- **EU GMP/FDA-Compliant Sampling**
  19 – 21 May 2021, Berlin, Germany

- **PharmaLab Congress 2020 - Live Online Conferences**
  10/11 November 2020
  - Endotoxin and Pyrogen Testing
  - Laboratory Optimization and Automation
  - Rapid Microbiological Methods and Mycoplasma Testing
  - Challenges in Bioanalytical and Analytical Laboratories - From Bioassays to ATMP Analytics
  - Challenges in Bioanalytical and Analytical Laboratories - From Life Cycle Management to Real Time

On behalf of APIC/CEFIC

- **23rd APIC/CEFIC Global GMP & Regulatory API Conference**
  28-30 October 2020 in Amsterdam, The Netherlands or broadcasted live to your desk!
- **ICH Q7 Training Courses**
  ICH Q7 in modern API Manufacturing – what to do and how to do
  30 November - 04 December 2020, Berlin, Germany

On behalf of European Qualified Person Association

- **The Qualified Person Forum 2020**
  26/27 November 2020
  Face-to-face meeting in Berlin and as a live online conference.
- **Qualified Person Education Course - Module A**
  Understand the Implications of becoming a QP
  03/04 March 2021, Munich, Germany
  With an optional pre-course Session:
  "Investigational Medicinal Products (IMP) QP Education Course"
- **Qualified Person Education Course - Module B**
  Mastering the QP Role in daily Practice
  30 June/01 July 2021, Berlin, Germany
  With an optional pre-course Session: "Interpersonal and Soft Skills for the QP" on 29 June 2021

Live Online Trainings/Conferences

- **EP - USP and other Pharmacopoeias: Dealing with different compendial methods**
  Tuesday, 22 September 2020, 09:00 h – 17:00 h CEST
  Wednesday, 23 September 2020, 08:30 h – 16:30 h CEST
- **Lab Data Integrity - Meeting FDA and EU Concerns**
  Tuesday, 22 September 2020, 09.00 h - 17.00 h CEST
  Wednesday, 23 September 2020, 09.00 h - 15.15 h CEST
- **Drug Master File Procedures in the EU, the US and Japan**
  Tuesday, 29 September 2020, 09.00 - 16.45 h CEST
  Wednesday, 30 September 2020, 08.30 - 16.30 h CEST
- **Process Validation**
  Tuesday, 06 October 2020, 09.00 – 17.00 h CEST
  Wednesday, 07 October 2020, 09.00 – 11.30 h CEST
- **Change Control**
  Tuesday, 6 October 2020, 9.00 – 17.15 h CEST
  Wednesday, 7 October 2020, 8.30 – 13.00 h CEST
- **FDA Compliance in Analytical Laboratories**
  Tuesday, 06 October 2020, 09.00 h - approx. 16.35 h CEST
  Wednesday, 07 October 2020, 08.30 h - approx. 16.20 h CEST
- **The GDP Compliance Manager**
  Tuesday, 06 October 2020, 09.00 h – 16.30 h CEST
  Wednesday, 07 October 2020, 09.00 h – 16.30 h CEST
- **Control of Parenterals**
  CCI Testing of Parenterals
  Tuesday, 6 October 2020, 09.00 – 18.00h
  Visual Inspection Systems
  Wednesday, 7 October 2020, 09.00 – 16.45 h
  Thursday, 8 October 2020, 08.30 to approx. 16.30 h
- **GMPs for Equipment, Utilities and Facilities**
  Tuesday, 6 October 2020, 09.00 – 17.00 h CEST
  Wednesday, 7 October 2020, 09.00 – 18.00 h CEST
  Thursday, 8 October 2020, 09.00 – 15.15 h CEST
- **API Regulatory Starting Materials**
  Tuesday, 13 October 2020, 9.00 – 17.30 h CEST
- **Handling OOT Results**
  Tuesday, 13 October 2020, 09.00 – 17.00 h CEST
- **Handling OOS Results**
  Wednesday, 14 October 2020, 9.00– 16.30 h CEST
- **GMP for Medical Devices**
  Tuesday, 13 October 2020, 09.00 – 16.30 h CEST
  Wednesday, 14 October 2020, 08.30 – 17.15 h CEST
- **Analytical Instrument Qualification**
  Thursday, 15 October 2020, 9.00– 17.00 h CEST
- **GMP meets Regulatory Affairs**
  Thursday, 15 October 2020, 09.00 – 17.15 h CEST
  Friday, 16 October 2020, 09.00 – 16.45 h CEST
- **China GMP and Registration of APIs**
  Tuesday, 20 October 2020, 10:00 - 12:30 h CEST
- **GMP for Beginners**
  Tuesday, 20 October 2020, 09.00 h – 17.30 h CEST
  Wednesday, 21 October 2020, 09.00 h – 17.00 h CEST
- **Bioassays and Bioanalytics**
  Tuesday, 20 October 2020, 09.00 – 17.30 h CEST
  Wednesday, 21 October 2020, 09.00 – 17.30 h CEST

- **Product Transfer**
  Tuesday, 20 October, 09.00 to approx. 17.30 h CEST
  Wednesday, 21 October 2020, 09.00 to approx. 17.00 h CEST
- **Stability Testing for Biological/Biotechnological Drug Substances and Drug Products**
  Thursday, 22 October 2020, 08.30 – 17.00 h CEST
- **GMP Auditor Practice**
  Wednesday, 21 October 2020, 9.00h – 17.00 h CEST
  Thursday, 22 October 2020, 8.30h – 17.00 h CEST
- **Data Integrity and Good Documentation Practice**
  Tuesday, 24 November 2020, 9.00 – 17.30 CET
  Wednesday, 25 November 2020, 9.00 – 16.30 CET
  Thursday, 26 November 2020, 9.00 – 15.15 CET
- **GMP for Beginners in Sterile Manufacturing**
  Tuesday, 27 October 2020, 9.30 to 17.30 h CET
  Wednesday, 28 October 2020, 9.00 to 16.15 h CET
- **How to Provide Process Validation Data in a Regulatory Submission**
  Wednesday, 04 November 2020, 9.00 – 17.30 h CET
  Thursday, 05 November 2020, 9.00 – 15.00 h CET
- **Annex 1 Conference - Current requirements for sterile manufacturing**
  Tuesday 10 November 2020, 09.00 – 18.00 h CET
  Wednesday , 11 November 2020, 08.30 – 17.00 h CET
- **Ambient Transport and Cold Chain**
  Wednesday, 04 November 2020, 9.00 h – 17.30 h CET
  Thursday, 05 November 2020, 9.00 h – 16.00 h CET
- **Quality, Safety and GMP for Advanced Therapy Medicinal Products (ATMPs)**
  Tuesday, 10 November 2020, 09.00 – 18.00 h CET
  Wednesday 11 November 2020, 08.30 – 17.30 h CET
- **Serialization - What's on**
  Wednesday, 11 November 2020, 09.00 – 16.45 h CET
  Thursday, 12 November 2020, 08.30 – 16.00 h CET
- **Ongoing/Continued Process Verification**
  Tuesday, 17 November 2020, 09.00 – 16.00 h CET
- **Contamination Control Strategies**
  Thursday, 19 November 2020, 08.30 h – 18.00 h CET
  Friday, 20 November 2020 , 08.30 h – 13.30 h CET
- **Data Integrity Audits & Inspections**
  Tuesday, 24 November 2020, 09:00 – 17:00 h CET
  Wednesday, 25 November 2020, 09:00 – 12.30 h CET
- **GMP for Vaccine Manufacturers**
  Tuesday, 24 November 2020, 09.30 h – 18.00 h CET
  Wednesday, 25 November 2020., 08.30 h – 17.00 h CET
- **GMP for Cannabis - what you need to know**
  Monday, 23 November 2020, 13.30 to approx. 18.00 h CET
  Tuesday, 24 November 2020, 10.30 to approx. 17.45 h CET
  With optional "GACP Post-Conference"
  Wednesday, 25 November 2020, 13.30 to approx. 18.00 h CET
- **Pharmaceutical Water**
  Tuesday, 1 December 2020, 09.00 – 17.00 h CET
  Wednesday, 2 December 2020, 09.00 – 15.30 h CET
- **How to write the Quality Part of an IMPD**
  Tuesday, 08 December 2020, 9.00 – 17.30 h CET
  Wednesday, 09 December 2020, 9.00 – 16.45 h CET
- **Modern Microbiology Laboratory**
  Tuesday, 08 December. 2020, 09:00 – 18.00 h CET
  Wednesday, 09 December 2020, 09.00 – 17.00 h CET
  Thursday, 10 December 2020, 09.00 – 13:30 h CET
- **Data Integrity & optional pre-course session Audit Trail Review**
  Audit Trail Review:
  Tuesday, 8 December 2020: 09.00 h – 17.15 h CET
  Data Integrity:
  Wednesday, 9 December 2020: 09.00 h – 17.30 h CET
  Thursday, 10 December 2020: 09.00 h – 17.30 h CET
  Friday, 11 December 2020: 09.00 h – 13.15 h CET
- **Pharmaceutical Biotechnology for Non-Biotechnologists**
  Tuesday, 15 December 2020, 09.30 h – 17.30 h CET
  Wednesday, 16 December 2020, 08.30 h – 17.00 h CET
- **Clean Rooms & HVAC Systems**
  Tuesday, 15 December 2020, 09.00 – 17.30 h CET
  Wednesday, 16 December 2020, 09.00 – 16.30 h CET

GMP Webinars

- **The risk-based Approach and GDP**
  Monday, 28 September 2020, 14.00 – 15.30 h CEST
- **GDP for APIs**
  Thursday, 01 October 2020, 14.00 – 15.30 h CEST
- **GxP Webinar Series on "Data Integrity"**
  Data Integrity from the view of a GMDP inspector
  Monday, 5 October 2020 10.00 - 12.00 h CEST
- **Data Integrity in the Light of ICH Q7**
  Thursday, 15 October 2020, 14.00 – 15.30 h CEST
- **GDP Partner Selection and Technical/ Quality Agreements**
  Monday, 19 October 2020, 14.00 – 15.30 h CEST
- **Temperature Qualification in Storage**
  Monday, 30 November 2020, 14.00 – 15.30 h CET
- **GMP Update 2020/2021**
  Thursday, 17 December 2020, 15.00 – 16.30 h CET
- **Ongoing/Continued Process Verification (Part 1)**
  Tuesday, 12 January 2021, 14.00 – 15.30 CET
- **Ongoing/Continued Process Verification (Part 2)**
  Wednesday, 13 January 2021, 14.00 -15.30 CET
- **Current Inspection Trends 2021**
  Tuesday, 19 January 2021, 14.00 -15.30 h CET
- **Data Integrity in Qualification and Validation Documentation**
  Thursday, 21 January 2021, 14.00 -15.30 h CET
- **Responsibilities at GDP**
  Wednesday, 27 January 2021, 14.00 -15.30 h CET
- **GDP Update 2021**
  Thursday, 24 February 2021, 14.00 – 15.30 h CET
- **Medical Device Technical Documentation**
  Tuesday, 9 March 2021, 14.00 – 15.30 h CET
- **Key Figures in Analytical Laboratories**
  Tuesday, 16 March 2021, 14.00 – 15.30 h CET
- **Transport Validation / Transport Verification**
  Tuesday, 11 May 2021, 14.00 – 15.30 h CEST