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Deviation Management and CAPA

Workshops on:
- Process Analysis and Failure Investigation
- CAPA Effectiveness & System Performance Check

SPEAKERS:

Dr Martin M. Appel
Cilag AG, Switzerland

Marcus Heinbuch
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Mick Hopper
GxPpro, U.K.

Lea Joos
GMP/GDP Inspectorate, Germany

Dr Bob McDowall
R.D. McDowall Limited, UK

8 - 9 May 2019, Barcelona, Spain

LEARNING OBJECTIVES:

- Rules and Regulations
  - EU
  - FDA
  - What the Inspector is looking for
- Deviations and CAPA
  - Deviations
  - CAPA
  - Classification
  - Failure Investigation
  - Risk Management
  - Root Cause
  - Human Error
- Evaluating and Monitoring
  - Effectiveness of CAPAs
  - KPIs

This education course is recognised for the ECA GMP Certification Programme „Certified QA Manager".
Please find details at www.gmp-certification.eu

Deviation Management and CAPA

8 - 9 May 2019, Barcelona, Spain

Learning Objectives

During this course, you will get to know the principles and discuss all relevant aspects to **implement and/ or work with a Deviation Management and CAPA System.** Furthermore, you will get to know possibilities and tools to **monitor and evaluate your CAPAs**.

Background

Things will go wrong from time to time. In the world of pharmaceuticals, we need to ensure that we have robust processes and procedures in place to deal with such situations. When an unplanned event arises it must be handled accordingly.

FDA's **Quality System Guide, recent Warning Letters** and EU-GMP Chapter 1 clearly emphasise the increasing relevance of a proper deviation management and CAPAs. ICH Q9 on Quality Risk Management and ICH Q10 on Pharmaceutical Quality Systems empower us to handle issues that arise in our daily work on the basis of risk analysis.

In any case a sound failure investigation is the key. Here it is also important to know how to deal with human error based and non-human error based non-conformances. Effective root cause analysis is the key to identifying appropriate CAPAs.

Independent from that, it needs to be pointed out that **CAPA is an excellent Quality Management tool** to continuously improve processes and avoid future failures. All personnel involved in the management of deviations and CAPAs should aim to identify opportunities for further improvement.

Target Audience

This course is designed for all personnel involved in Deviation Management and CAPA activities at their company. It is addressed to persons from Quality Assurance and Control, Manufacturing and R&D.

Programme

International Requirements – Rules and Regulations

- European requirements
- The expectations of the FDA
- GMP and documentation issues
- Harmonisation in sight?

Excerpt from FDA Warning Letter

"...the investigation failed to establish a root cause and your quality unit failed to ensure the implementation of adequate corrective actions to prevent future recurrence."

Deviation Handling

- How to document deviations
- Information and Data Management
- Critical/ major/ minor
- CAPA or not?

CAPA: Principles, System, Implementation and Process Improvements and the use of Risk Management Techniques

- Tools
- Quality Risk Management
- Human Error Overview
- Monitoring & Evaluation Overview

Workshop:

An interactive exercise on scenarios with a focus on using the tools from the presentation

- Human Error based
- Non-human error based

Deviations in the Light of Inspections

- Focus in inspection
- Trends, Product Quality Review and Product Review
- Self-inspection as an important tool

Case Study: how to implement a CAPA System

- How to integrate existing QM Systems (OOS, Complaint Handling, Deviations)
- Examples and lessons learned

Software tools for CAPA management as part of a QMS

- Understanding your paper workflows and processes
- Can you improve the current process using electronic workflows?
- An overview of some of the main software applications for CAPA
- Efficient validation of a CAPA application

CAPA Effectiveness & System Performance Check

As part of the periodic quality review programme, Quality Management should routinely analyse reports of deviations and CAPAs to determine KPIs, trends, recurrence of non-conformances and effectiveness of CAPAs. A summary overview should be reported to the Senior Management team. ICH Q10 identifies this as best practice - but are we doing this as well as we could or should? We will discuss Quality Metrics as well as which are the important ones that will show you have a good Pharmaceutical Quality System.

Workshop on CAPA Effectiveness & System Performance Check

An interactive session with a focus on enhancing the knowledge gained in the presentation

Social Event

In the evening of the first course day, you are cordially invited to a social event. This is an excellent opportunity to share your experiences with colleagues from other companies in a relaxed atmosphere.

Speakers

Dr Martin M. Appel

Cilag AG, Johnson & Johnson, Switzerland

Dr Appel is Director QA for the Global API External Manufacturing and Supplier Quality of Janssen Supply Chain. He has more than 30 years experience in several manager positions in the pharmaceutical industry.

Marcus Heinbuch

B. Braun Melsungen AG, Germany

Marcus Heinbuch is Head of QM Operations in the Quality Management of CoE Pharmaceuticals at B. Braun Melsungen AG.

Michael Hopper

GxPpro, U.K.

Michael (Mick) Hopper set-up GxPpro after leaving Pfizer. Mick has over 30 years and held several Technical, Management and QA roles. He also gained a green belt accreditation and led the implementation of several improvement.

Lea Joos

GMP Inspectorate, Local Authority Munich, Germany

Lea Joos is a Pharmacist working for the local Inspectorate as GMP and GDP Inspector. She also gives lectures on different GMP and GDP topics and published several essays in different professional journals.

Dr Bob McDowall

R.D.McDowall Limited, UK

Bob McDowall is Principal of McDowall Consulting, UK. He has been involved with the validation of computerised systems for over 20 years and is the author of a book on the validation of chromatography data systems. Bob is a member of the Editorial Advisory Boards of several Journals.

Easy Registration

Reservation Form:
CONCEPT HEIDELBERG
P.O. Box 10 17 64
69007 Heidelberg
Germany

Reservation Form:
+ 49 6221 84 44 34

e-mail:
info@concept-heidelberg.de

Internet:
www.gmp-compliance.org

If the bill-to-address deviates from the specifications on the right, please fill out here:

CONCEPT HEIDELBERG
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Fax +49 (0) 62 21/84 44 34

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Reservation Form (Please complete in full)

+ 49 6221 84 44 34

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8 - 9 May 2019, Barcelona, Spain

☐ Mr. ☐ Ms.

Title, first name, surname

Company | Department

Important: Please indicate your company's VAT ID Number | Purchase Order No, if applicable

Street/P.O. Box

City | Zip Code | Country

Phone/Fax | E-Mail (please fill in)

General terms and conditions
If you cannot attend the conference you have two options:
1. We are happy to welcome a substitute colleague at any time.
2. If you have to cancel entirely we must charge the following processing fees: Cancellation
- until 2 weeks prior to the conference 10 %,
- until 1 weeks prior to the conference 50 %
- within 1 week prior to the conference 100 %.

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Important: This is a binding registration and above fees are due in case of cancellation or non-appearance. If you cannot take part, you have to inform us in writing. The cancellation fee will then be calculated according to the point of time at which we receive your message. In case you do not appear at the event without having informed us, you will have to pay the full registration fee, even if you have not made the payment yet. Only after we have received your payment, you are entitled to participate in the conference (receipt of payment will not be confirmed)! (As of January 2012)

German law shall apply. Court of jurisdiction is Heidelberg.

Privacy Policy: By registering for this event, I accept the processing of my Personal Data. Concept Heidelberg will use my data for the processing of this order, for which I hereby declare to agree that my personal data is stored and processed. Concept Heidelberg will only send me information in relation with this order or similar ones. My personal data will not be disclosed to third parties (see also the privacy policy at http://www.gmp-compliance.org/eca_privacy.html). I note that I can ask for the modification, correction or deletion of my data at any time via the contact form on this website.

Date

Wednesday, 08 May 2019, 09.30 – 17.30 h
(Registration and coffee 09.00 – 9.30 h)
Thursday, 09 May 2019, 08.30 – 16.00 h

Venue

Barceló Sants Hotel
Plaça dels Països Catalans, s/n
08014 Barcelona, Spain
Phone +34 (93) 503 53 00
sants@barcelo.com

Fees (per delegate plus VAT)

ECA Members € 1,490
APIC Members € 1,590
Non-ECA Members € 1,690
EU GMP Inspectorates € 845
The course fee is payable in advance after receipt of invoice and includes conference documentation, dinner on the first day, lunch on both days and all refreshments. VAT is reclaimable.

Accommodation

CONCEPT HEIDELBERG has reserved a limited number of rooms in the conference hotel. You will receive a room reservation form when you have registered for the course. Reservation should be made directly with the hotel.
Early reservation is recommended.

Registration

Via the attached reservation form, by e-mail or by fax message. Or you register online at www.gmp-compliance.org.

Conference language

The official conference language will be English.

Organisation and Contact

ECA has entrusted Concept Heidelberg with the organisation of this event.

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