

### Speakers



Dr Michael Aigner University Erlangen



Cheryl Chia Lotus Phoenix Consulting



Dr Martin Egger Pharmaserv Logistics



Prof René Geyeregger St. Anna Children's Cancer Research Institute



Dr Christian Grote-Westrick B. Braun Avitum AG



Jochen Heinzel F. Hoffmann-La Roche



Dr Afshin Hosseiny European GDP Association



Saddam Huq GlaxoSmithKline



Saskia Killer Oncotec Pharma Produktion



Sue Mann Sue Mann Consultancy



Axel Radke Trust Expertenservice



Dr Laura Ribeiro OCP Portugal



Dr Torsten Schmidt-Bader moveproTEC



Emil Schwan Swedish Medical Products Agency



**Maria Sekamwa** Surgipharm



Gundula Warnecke Oncotec Pharma Produktion



# GDP Forum

A GDP Conference designed by the European GDP Association

24/25 June 2020 | Barcelona, Spain



## Highlights

- KPIs and Benchmarking
- Brexit Implications
- Risk Management
- Failure Investigation
- Challenges in local Markets
- Sea Freight
- Customer Master Data
- Insourcing of Logistics
- Distribution of ATMPs
- With five parallel Workshops

Every participant will get a free copy of the ECA/PQG GDP Guidance Document!

## Objectives

The aim of this conference is to inform about recent experiences made in the implementation of the EU-GDP Guidelines. Challenges and possible solutions will be discussed and examples will demonstrate how the requirements can be put into practice.

## Background

It is of key importance that medicinal products are not only made to a high quality in accordance with Good Manufacturing Practice (GMP), but that the quality and integrity of these products are maintained through the entire supply chain to the patient. This is where Good Distribution Practice (GDP) comes into play.

In the last few years the world of GDP has changed in scope and complexity and continues to evolve to meet new challenges with the continued aim of safeguarding public health. The Falsified Medicines Directive (2011/62/EU) was transposed into the laws and regulations of the EU Member States alongside the introduction of revised EU guidelines on Good Distribution Practice. Many other countries followed adopting these Guidelines or implementing very similar approaches.

## **Target Audience**

All personnel involved in pharmaceutical storage, transportation, cold chain and distribution activities and the control of those activities.

### Moderator

Dr Afshin Hosseiny

### About the GDP Association:

The European GDP Association aims to support Pharmaceutical Industry, Authorities and Logistic Providers with regard to the implementation of Good Distribution Practice.

It represents all stakeholders e.g. from Pharmaceutical Industry, Authorities and Logistic Providers and supports all members and stakeholders by providing them information and support in the implementation of GDP.

The Association is a not for profit organisation under the umbrella of the ECA Foundation. Membership is free to all individuals involved in Good Distribution Practice (currently more than 2.000 members).

www.good-distribution-practice-group.org

## Programme

### Performance beyond Compliance

- Benchmarks with sense
- When is a system or process robust?
- Easily understandable key figures
- Case studies

### **Brexit Implications**

- Import/ export issues
- Drug shortage challenges
- Falsification
- Serialisation
- Co-operation of the agencies

### GDP Challenges in Africa

- Background
- Evolution of GDP over the years
- Problems involved
- Steps taken to improve

#### Customer Master Data

- Definitions
- Parties involved
- Responsibilities and RP oversight
- Managing Customer Master Data
- Lifecycle Management

## 3 parallel Workshops:

- 1. Risk Management in GDP Warehousing
  - Challenges and possible solutions
  - Examples
- 2. When things go wrong what can happen in the real World
  - Experiences made in examining problems in the supply chain – and what to learn from them.
- 3. How to deal with suspected and confirmed falsified Medicines
  - What is a suspected falsified medicine
  - How to identify falsified medicine in your workplace
  - How to approach the investigation of a suspected pack
  - What should the end user do with the pack who should manage communication with the end users

You will be able to attend TWO of these workshops. Please choose the ones you like to attend when you register.

#### Case Studies:

The Roche Quality Governance Model for Distribution to Direct Markets

- Selection, implementation, operation and decommissioning of partners
- Key Performance Indicators (KPIs)
- Compliance score card

### The Switch from Air-freight to Sea-Freight

- Challenges and Benefits
- Examples

From external to internal Solutions - Insourcing of Warehousing and Logistics

- Implementation of own storage areas for minimum supply at the site
- Container and transport vehicle purchasing (refrigerated, warm, hazardous)

## 2 parallel Workshops:

- 1. Master Data and Serialisation
  - Master Data for Serialisation
  - Alerts Handling
  - Exercises

### 2. Distribution of ATMPs

- Challenges when dealing with ATMP
- What information is needed to help service providers
- Examples

You will be able to attend ONE of these workshops. Please choose the one you like to attend when you register.

### Question and Answers Session

You will be able to send in GDP related questions before the conference. Additionally, questions can be posted on question cards during the conference.

Answers will be given by the expert speakers in a dedicated session and published in the members are of the GDP Association's website (www.good-distribution-practice-group.org).

## Speakers

Dr Michael Aigner, University Hospital Erlangen, Germany Michael Aigner is Head of Production and Qualified Person for cellular ATMP/ATIMPs.

Cheryl Chia, Lotus Phoenix Consulting, Netherlands Cheryl Chia is Consultant for GMP and GDP compliance in the pharmaceutical supply chain.

Dr Martin Egger, Pharmaserv, Germany

Dr Martin Egger is Managing Director of Pharmaserv and responsible for the brand Pharmaserv Logistics.

Prof René Geyeregger, St. Anna Children's Cancer Research Institute, Austria

Prof René Geyeregger is associated Professor and principal investigator at the Clinical Cell Biology and FACS Core Unit.

Dr Christian Grote-Westrick, B. Braun Avitum AG, Germany Dr Grote-Westrick is Head of Quality Assurance and lead auditor with responsibilities in GMP / GDP compliance.

Jochen Heinzel, F. Hoffmann-La Roche, Switzerland Jochen Heinzel is Global Quality Manager Supply Chain & Affiliate Quality EEMEA Region.

Afshin Hosseiny, Ph.D., Tabriz Consulting Ltd., U.K. Dr Afshin Hosseiny is Managing Director of Tabriz Consulting Ltd. and Chairman of ECA's GDP Association.

### Saddam Huq, GlaxoSmithKline, U.K.

Saddam Huq is Senior Manager Quality for Distribution & Cold Chain Management Vaccines, Quality Assurance Shared Services.

Saskia Killer, Oncotec Pharma Produktion, Germany
Saskia Killer is head of production and responsible for the aseptic processing of cytotoxic drugs. She is a Pharmaceutical Engi-

tic processing of cytotoxic drugs. She is a Pharmaceutical Engineer and at Oncotec since 2014.

### Sue Mann, Sue Mann Consultancy, U.K.

Sue Mann is GMP/GDP consultant with more than 35 years experience in the Pharmaceutical Industry, mainly in Quality Assurance, Clinical Trials supply and production support.

### Axel Radke, Trust Expertenservice, Germany

Axel Radke is Responsible Person and technical expert for investigations of issues in the storage and transport of pharmaceuticals.

### Dr Laura Ribeiro, OCP Portugal

Laura Ribeiro is Director Quality and Regulatory Affairs, managing a team of Responsible Persons.

Dr Torsten Schmidt-Bader, moveproTEC Compliance & Innovation Advisory, Germany

Dr Torsten Schmidt-Bader is Managing Director at moveproTEC and a GMP/GDP lead auditor and compliance advisor.

Emil Schwan, Medical Products Agency, Sweden Emil Schwan is Pharmaceutical Inspector at the Drug Inspectorate of the MPA and a member of the PIC/S Working Group on GDP.

### Maria Sekamwa, Surgipharm, Uganda

Maria Sekamwa is Assistant Warehouse Manager and Qualified Person for Pharmacovigilance.

Gundula Warnecke, Oncotec Pharma Produktion, Germany Gundula Warnecke is a project manager for Oncotec and was responsible for the installation of a GDP-compliant warehouse at Oncotec.



Every participant will get a free copy of the GDP Guidance on interpretation and implementation, a joint publication of the ECA Foundation and the Pharmaceutical Quality Group of the Chartered Quality Institute.

Date

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