

# Speakers



Peter Kralinger Carrymed Pharma & Transport, Austria



Kane Edgeworth Biomap, UK

# COVID-19 Vaccines and Temperature-Sensitive Pharmaceuticals

- Transport and Vehicle Qualification



Live Online Training on 04 March 2021, 10:00 – 13:00 h



# Highlights

- Temperature-Controlled Transports of Medicinal Products
  - Insulated packaging for small and large quantities
  - Road transport and air freight of cold chain products (2 8 °C)
  - Transport and storage at ultra-frozen range (-70 °C)
- Vehicle Qualification
  - Qualification of temperature-controlled vehicles (IQ/OQ/PQ)
  - Regulatory landscape
  - Matrix approach for larger fleets
  - Vehicle mapping case study
- **Questions and Answers Session**

# Objectives

This Live Online Training aims to give participants a comprehensive yet compact overview of expectations concerning the transportation of products requiring special conditions. The focus will be on the requirements for the transport of **COVID-19 vaccines** and other **temperature-sensitive products**.

General aspects of road transport and air fright at cold chain (2 – 8 °C) and at ultra-frozen (-70 °C) conditions will be discussed. Furthermore, vehicle qualification and the effective mapping of vehicles will be covered. A Q&A session ensures interaction and that all questions are answered.

# 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.

The distribution of temperature-sensitive pharmaceuticals is extremely challenging. The EU GDP-Guidelines (Guidelines of 5 November 2013 on Good Distribution Practice of medicinal products for human use) require that if temperature-controlled vehicles are used, the temperature monitoring equipment used during transport should be maintained and calibrated at regular intervals. Temperature mapping under representative conditions should be carried out, taking into account seasonal variations. For temperature-sensitive products, qualified equipment (e.g. thermal packaging, temperature-controlled containers or temperature controlled vehicles) should be used to ensure correct transport conditions.

With the approvals of various **COVID-19 vaccines** in many countries, the question arises how to safely transport the serum from manufacturing facilities to storage and distribution centres, and then onwards to local healthcare facilities and vaccination centres. However, this question does not only arise for COVID-19 vaccines, but also for **all temperature-sensitive products in general.** 

The specific packaging, transport, handling and storage requirements of the various COVID-19 vaccines are partly still open. The transport routes may also depend on the future production sites. In any case, transport companies must take special safety precautions for the transport of COVID-19 vaccines. The serum is not only very valuable, but also particularly sensitive. Damages can quickly lead to a total loss, as in case of doubt, the entire load must be destroyed. In addition, there are risks such as interruption of the cold chain. Therefore, the vaccine may only be distributed with controlled packaging solutions and in special vehicles that are appropriately qualified and whose temperature is monitored permanently.

# **Target Audience**

This Live Online Training was developed for managers, executives, Responsible Persons (RP's), technical staff and other employees from companies involved in the distribution and supply of COVID-19 vaccines and temperature-sensitive pharmaceutical products.

It will be of interest in particular for personnel from the following departments:

- Quality Assurance
- Validation
- Engineering
- Logistics
- Cold Chain
- Regulatory Compliance

# Moderator

Dr Markus Funk

# Programme

Welcome and Introduction

(Dr Markus Funk)



Guidelines of 5 November 2013 on Good Distribution Practice of medicinal products for human use (2013/C 343/01)

### Chapter 2.4. (Training)

[...] Personnel dealing with any products which require more stringent handling conditions should receive specific training. Examples of such products include [...] temperature-sensitive products.

A record of all training should be kept, and the effectiveness of training should be periodically assessed and documented.

# Temperature-Controlled Transports of Medicinal Products

(Peter Kralinger)

- Insulated packaging for small and large quantities
- Road transport and air freight of cold chain products (2 8 °C)
- Transport and storage at ultra-frozen range (-70 °C)

### Vehicle Qualification

### (Kane Edgeworth)

- Qualification of temperature-controlled vehicles (IQ/OQ/PQ)
- Regulatory landscape
- Matrix approach for larger fleets
- Vehicle mapping case study



### Questions and Answers Session

(Peter Kralinger and Kane Edgeworth)

Participants are invited to ask questions



Guidelines of 5 November 2013 on Good Distribution Practice of medicinal products for human use (2013/C 343/01)

### **Chapter 9.4. (Products requiring special conditions)**

In relation to deliveries containing medicinal products requiring special conditions such as narcotics or psychotropic substances, the wholesale distributor should maintain a safe and secure supply chain for these products in accordance with requirements laid down by the Member States concerned. There should be additional control systems in place for delivery of these products. There should be a protocol to address the occurrence of any theft.

[...]

For temperature-sensitive products, qualified equipment (e.g. thermal packaging, temperature-controlled containers or temperature-controlled vehicles) should be used to ensure correct transport conditions are maintained between the manufacturer, wholesale distributor and customer.

If temperature-controlled vehicles are used, the temperature monitoring equipment used during transport should be maintained and calibrated at regular intervals. Temperature mapping under representative conditions should be carried out and should take into account seasonal variations.

If requested, customers should be provided with information to demonstrate that products have complied with the temperature storage conditions.

[...]

The process for delivery of sensitive products and control of seasonal temperature variations should be described in a written procedure.



Peter Kralinger

Carrymed Pharma & Transport GmbH, Austria Peter Kralinger is Managing Director of Carrymed, the first licensed pharma company providing inter-

national transport of temperature-sensitive pharmaceuticals. Before that he was in charge of the global transportation activities for all manufacturing sites in Europe of a large manufacturer of the pharmaceutical industry.



Kane Edgeworth Biomap Ltd, UK

Kane Edgeworth is Director at Biomap, providing validation & temperature monitoring solutions for the

Life Sciences industry. Before that, he was UK Operations Director at one of the world's largest data logger manufacturers.

# Agenda

10:00 – 10:10 h Welcome and Introduction

10:10 – 11:10 h **Presentation 1** 

11:10 – 11:20 h Break

11:20 – 12:20 h **Presentation 2** 

12:20 - 12:30 h Break

12:30 – 13:00 h Questions and Answers Session



# Reservation Form (Please complete in full)

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

COVID-19 Vaccines and Temperature-Sensitive Pharmaceuticals – Transport and Vehicle Qualification Purchase Order Number, if applicable Live Online Training on 04 March 2021, 10:00 – 13:00 h Company Important: Please indicate your company's VAT ID Numbe Title, first name, surname Department City

E-mail (Please fill in)

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# Date of the Live Online Training Thursday, 04 March 2021, 10:00 - 13:00 h All times mentioned are CET.

# Technical Requirements

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# Fees (per delegate, plus VAT)

ECA Members € 490 European GDP Association Members € 490 APIC Members € 540 Non-ECA Members € 590 EU GMP Inspectorates € 490

The conference fee is payable in advance after receipt of invoice.

# Registration

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

### Presentations/Certificate

The presentations will be made available to you prior to the Live Online Training as PDF files. After the event, you will automatically receive your certificate of participation.

# Conference language

The official conference language will be English.

# Ordering a Recording

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## Organisation and Contact

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

### **CONCEPT HEIDELBERG**

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### For questions regarding content:

Dr Markus Funk (Director Operations) at +49(0) 62 21/84 44 40, or per e-mail at funk@concept-heidelberg.de

### For questions regarding organisation please contact:

Mr Rouwen Schopka (Organisation Manager) at +49(0) 62 21/84 44 13, or per e-mail at schopka@concept-heidelberg.de

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