



## Speakers



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# Handling of Foreign Particles in APIs and Excipients

25 /26 January 2022 | Frankfurt/Main, Germany



*Risk analysis, preventive measures and incident management*

## Highlights

- Key preventive measures to minimise foreign particles
- How to deal with technically unavoidable particles in excipients
- Acceptance criteria for particles in APIs
- How to identify the source of insoluble matter
- Analytical control methods for particle detection
- How to minimise the presence of particles – strategies for cleaning and detection
- Foreign particles in excipients and finished product quality and safety

## Objectives

During this course all relevant aspects regarding the control of particles in APIs and excipients will be discussed.

You will learn

- How potential sources of insoluble matter can be identified
- Which acceptance criteria for particles can be applied
- How good practices to minimise the presence of particles in APIs can look like
- What has to be considered regarding control of particles during plant and equipment maintenance and cleaning
- How a particulate contamination profile can be established.

## Background

Visible particles, insoluble particles or matter or foreign particles in Active Pharmaceutical Ingredients (APIs) and pharmaceutical excipients are topics of great interest and of importance to the pharmaceutical industry.

A number of inspectional observations from various Regulatory Authorities related to visible particles in Drug Products and APIs has risen considerable concern. Moreover inappropriate methods of investigation, controls and preventive and corrective actions were all subjects of citations by authorities and observations by API and excipient customers.

Particles have always been present in APIs and excipients but guidance from health authorities (EMA, FDA, others) or Pharmacopoeias (e.g. EP, USP) about particles is very limited. The APIC Guidance on Insoluble Matter and Foreign Particles in APIs and the IPEC Guide on "Technically Unavoidable Particle Profile (TUPP)" are the only best practice documents so far providing guidance for a standard approach towards an appropriate control of foreign particles in APIs and pharmaceutical excipients.

## Target Audience

This course is addressed to employees and senior staff of pharmaceutical companies and manufacturers of APIs and excipients. The course is of particular interest to all those working in Quality Assurance, Quality Control, production and purchasing departments.

## Programme

### Particles and Insoluble Matter in API Manufacturing: Why is it a Topic of Great Interest?

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- Definition of particles
- Types of particles
- Possible reasons for the elevated presence of visible particles
- Hints in guidances on how to deal with visible particles
- Inspectional observations
- Expectations of API manufacturers, API users, API suppliers and supervisory authorities regarding visible particles in APIs

### Foreign Matter in Pharmaceutical Excipients – How to Deal with “Technically Unavoidable Particles” (TUPs)

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- Understanding the nature of contaminants
- Establishing the target profile to support risk assessment
- Establishing the risk profile of unavoidable foreign particles
- Understanding the source and mitigation to minimise the foreign particles

### Incident Management – How to Identify the Source of Insoluble Matter

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- Potential sources of insoluble matter
- Root cause analysis – examples of investigation techniques and aids
- Risk assessment: topics to be considered during the investigation/disposition decision

### Acceptance Criteria for Particles in APIs

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- Types of dosage forms and routes of administration
- Typical limits for particle size seen via a filter test
- Proposal for limits



### Case Studies: Deviations Caused by Foreign Particles

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### How Can Routine Cleaning Procedures Detect or Minimize the Presence of Particles in API Production?

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- Guides and Industry Standards regarding cleaning
- Equipment cleaning
- Production environment cleaning
- Equipment design considerations
- Detection/removal methods of particles
- Preventive measures

## Analytical Control Methods for Particle Detection

- Design of appropriate analytical techniques
- Understanding the operational and investigative analytical methodologies
- Case studies to identify the contaminants

## Foreign Particles in Excipients and Finished Product Quality and Safety

- Contamination Profile of Excipients meets Finished Product Quality Target Product Profile
- Excipient Process Risk Analysis and TUPP/ Particulate Contamination Profiling

## Social Event

In the evening of the first 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 Rajnish Chhabra**  
QAR Solutions, The Netherlands

Dr Chhabra is founder and leader of QAR Solutions B.V. and started his business in September 2018. Before that he worked at DSM Sinochem Pharmaceuticals where Dr Chhabra held a position as Sr. Manager and Director Quality & Regulatory Affairs.



**Dr Martin Melzer**  
gempex GmbH, Germany

Dr Melzer is Principal Consultant at gempex GmbH, Germany. Before that he was consultant for GMP/ GDP aspects, GMP -Inspector in a German Field Inspectorate in Germany, QA/ QC manager at a production site for AP/ and finished products, and head of laboratory for plant medicinal products.



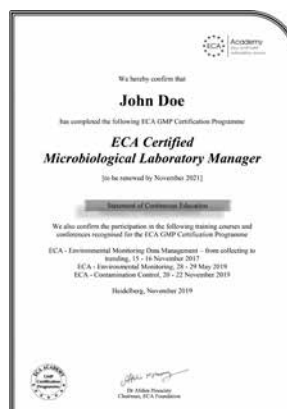
**Peter Mungenast**  
Merck KGaA, Germany

Mr Mungenast studied Biology and Chemistry at the University in Karlsruhe. Then he worked in different functions for Merck KGaA. Since 1996 he is responsible for cleaning validation, training and different projects in the Quality Assurance department.



**Dr Dirk Overrödter**  
Janssen, Schaffhausen, Switzerland

Dr Overrödter joined Cilag AG in 1995 and was employed in various positions in R&D and Compliance. Since May 2014 he is Head of QA Small Molecules (API & Drug Product) at Janssen's site in Schaffhausen, Switzerland.



## Internationally Acknowledged Certificate from ECA Academy

The EU GMP Guide requires: „... All personnel should be aware of the principles of Good Manufacturing Practice that affect them and receive initial and continuing training,...“. This is why you receive an acknowledged participant certificate, which lists the contents of the seminar in detail and with which you document your training.

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Reservation Form (Please complete in full)

## Handling of Foreign Particles in APIs and Excipients, 25 /26 January 2022, Frankfurt/Main, Germany

Title, first name, surname

Department

Company

Important: Please indicate your company's VAT ID Number

Purchase Order Number, if applicable

CONCEPT HEIDELBERG

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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:
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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 <https://www.gmp-compliance.org/privacy-policy>). 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

Tuesday, 25 January 2022, 09.30 – 18.00 h  
(Registration and coffee 09.00 – 09.30 h)

Wednesday, 26 January 2022, 09.00 – 13.00 h

### Venue

Scandic Frankfurt Museumsufer  
Wilhelm-Leuschner-Straße 44

60329 Frankfurt/Main, Germany

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Email [museumsufer@scandichotels.com](mailto:museumsufer@scandichotels.com)

### Fees (per delegate, plus VAT)

ECA Members € 1,590

APIC Members € 1,690

Non-ECA Members € 1,790

EU GMP Inspectorates € 895

The 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/POG when you have registered for the course. Reservation should be made directly with the hotel. Early reservation is recommended. Cancellation of the reserved rooms free of charge is possible up to 2 weeks before arrival.

### Registration

Via the attached reservation form, by e-mail or by fax message.

Or you register online at [www.gmp-compliance.org](http://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.

CONCEPT HEIDELBERG

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