PharmaLab 2015

Analytics, Bioanalytics and Microbiology
– Congress & Exhibition –

Swissôtel Düsseldorf/Neuss – 10/11 November 2015

www.pharmalab-congress.com

Put together your own programme:
- over 50 Lectures
- almost 40 Speakers

The Conferences

10 November 2015

- ECA – Laboratory Informatics – Update 2015
- ECA – Endotoxin and Pyrogen Testing (Day 1)
- ECA – Rapid Microbiological Methods

11 November 2015

- ECA – cGMP Compliance Trends in Analytical Quality Control
- ECA – Validation Approach of Bioassays Using Statistical Methods
- ECA – Endotoxin and Pyrogen Testing (Day 2)
- ECA – Adventitious Agents - Impurities and Contaminants

Exhibitor Information in the back

Media Partner:
European Biotechnology
Life Sciences and Industry Magazine

CONCEPT HEIDELBERG
Pharmaceutical Quality
Training. Conferences. Services.

The Congress Objective

On 10 and 11 November 2015 the PharmaLab Congress will take place in Düsseldorf/ Neuss for the third time. This Congress addressing staff and executives in all lab areas of the pharmaceutical industry will be comprised of six international and four German language conferences plus the parallel exhibition. It will provide you with current developments of laboratory methods, materials as well as the current status of the regulatory requirements of the Pharmacopoeias.

Furthermore, experts from authorities, industrial quality control and contract laboratories will share their experience with the use and the qualification of analytical systems as well with the validation of analytical methods and microbiological tests.

Use this unique opportunity to get an update on the state of the art in pharmaceutical laboratories and to discuss current developments with speakers and colleagues.

PharmaLab 2015 Overview	
Conferences	One day ticket 690,- EUR
10 November 2015	
ECA – Laboratory Informatics – Update 2015	
ECA – Endotoxin and Pyrogen Testing (Day 1)	
ECA – Rapid Microbiological Methods	
11 November 2015	
ECA – cGMP Compliance Trends in Analytical Quality Control	
ECA – Validation Approach of Bioassays Using Statistical Methods	
ECA – Endotoxin and Pyrogen Testing (Day 2)	
ECA – Adventitious Agents - Impurities and Contaminants	
Exhibition (10 and 11 November 2015)	

Subject Areas: Analytics | Bioanalytics | Microbiology

Background

Laboratory work is an important part of pharmaceutical research, development and quality control. During inspections the responsible authorities significantly increased their emphasis on the quality management and performance of laboratories and their quality standards. This scrutiny of the regulators require laboratories to establish GLP and GMP appropriate systems and methods, and in particular:

- General GLP or cGMP understanding and particularly relating to compliance with written procedures
- Validation, performance and transfer of analytical procedures and microbial tests
- Equipment qualification and calibration
- Computer validation (including the interpretation of EU GMP Annex 11 and 21 CFR Part 11 and the actual requirements for lab data integrity)
- Operator training

Especially for pharmaceutical products and active substances from biological origin, classic analytical and testing methods don't fit. New developed methods e.g. MAT for pyrogen testing, rapid methods for sterility testing or necessary bioassays require a permanent knowledge update and advanced training of laboratory personnel and of the involved staff.

Target Audience

This conference will be of interest to

- Laboratory Managers, Supervisors and Analysts in pharmaceutical quality control departments
- Laboratory Staff of Research and Development
- Responsible Authorities
- Suppliers on Laboratories
- Associates of Contract Laboratories

The fees

A one day ticket/two days ticket will enable you to visit the congress either only on day 1 or only on day 2 or on both days. Charges for the one day tickets are € 690,- plus VAT, for the two days ticket € 1.380,- plus VAT (due to the special fees for the congress, ECA membership discounts are not applicable). They include lunch and beverages during the conferences and in breaks as well as the social event on the evening of the first congress day. Charges are payable after receipt of invoice.

Particularities of PharmaLab 2015:

- The registration allows you to access the 6 conferences with close to 50 lectures. In a word, you can create your individual conference programme.
- Move any time to any conference room. Thanks to one day tickets, you can attend only the first or the second day - but also both days of PharmaLab.
- You will receive a USB stick including all the conference lectures of the Congress.
- Learn about the latest products and services relating to analytics, bioanalytics and microbiology at the exhibition.
- Take advantage of PharmaLab – and particularly of the Social Event on the evening of the first day – for an information exchange with delegates, speakers and exhibitors

The Social Event

On the evening of the first congress day, on 10 November 2015, all congress delegates and speakers are invited to a „Get together" in the Congress Center. Take advantage of this opportunity for an information exchange and enjoy the laid-back atmosphere and the entertainment programme.

The Location

Swissôtel Congress Centrum Düsseldorf/Neuss
Rheinallee 1
41460 Neuss
Tel.: +49 (0) 2131 77 - 00
Fax: +49 (0) 2131 77 - 1367
Emailus@swissotel-duesseldorf.de

The Organiser

CONCEPT HEIDELBERG – On behalf of the ECA Academy
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www.gmp-navigator.com

ECA ACADEMY

The Contacts

For questions regarding content:
Endotoxin and Pyrogen Testing / Rapid Microbiological Methods / Validation Approach of Bioassays Using Statistical Methods / Adventitious Agents – Impurities and Contaminants:
Axel H. Schroeder (Operations Director), Phone +49 (0) 6221 84 44 10,
E-Mail: schroeder@concept-heidelberg.de.

Laboratory Informatics – Update 2015 / cGMP Compliance Trends in Analytical Quality Control:
Dr Günter Brendelberger (Operations Director), Phone +49 (0) 6221 84 44 40,
E-Mail: brendelberger@concept-heidelberg.de.

For questions regarding reservation, hotel, organisation, exhibition etc.:
Detlef Benesch, (Organisation), Phone +49 (0) 6221 84 44 45,
E-Mail: benesch@concept-heidelberg.de.

The Media Partner

European Biotechnology
Life Sciences and Industry Magazine

European Biotechnology Magazine reports about the latest political, economic and technical developments in the life sciences sector in all 28 EU countries plus Switzerland and Norway. To find out more please visit www.eurobiotechnews.eu.

Speakers

Arjan Bannink	**Waters, The Netherlands** Sales Development Manager, Laboratory Informatics, responsible for the adoption of the Laboratory Management Software Solutions from Waters in QC departments within Europe.
Ulla Bondegaard	**Novo Nordisk AS, Bagsværd, Denmark** Responsible for maintaining cross-organisational (and cross-country) laboratory processes.
Peter J. Boogaard	**Industrial Lab Automation, Moordrecht, The Netherlands** Founder of Industrial Lab Automation.
Emmanuelle Charton, Ph. D.	**European Directorate for the Quality of Medicines and HealthCare (EDQM)** Head of Division B and Deputy Head of the European Pharmacopoeia Department.
Peter Cornelis	**Toxikon Europe NV, Leuven, Belgium** Department Supervisor Microbiology & In Vitro Toxicology.
Dr Markus Dathe	**F. Hoffmann-La Roche AG, Basel, Switzerland** GMP and CSV coordinator in the chemical development and supply since 2011.
Dr Anja Fritsch	**Confarma France Sarl, Molecular Biology** Chief Scientific Officer.
Stefan Gärtner	**Labor L+S AG, Bad Bocklet, Germany** Head Special Department Testing of Sterile Products.
Prof Frank Oliver Glöckner	**Max Planck Institut and Jacobs University, Bremen, Germany** Head of Microbial Genomics and Bioinformatics Research Group.
Michael Goetter	**Lonza Walkersville, Inc., Wayne, USA** General Manager of Informatics.
Dr Fatma Gökşin Bahar	**Arven Pharmaceuticals, Turkey** Biotechnology Quality Control Specialist.
Dr Rajesh Gupta	**Biologics Quality & Regulatory Consultants, LLC, North Potomac, USA** Co-Owner and Principal Consultant; formerly Deputy Division Director and Lab Chief, Div of Product Quality (Div of Biological Standards & QC) FDA.
Elena Gustchina	**Lonza Walkersville, Inc., Walkersville, USA** Scientist, Enzyme and Protein Chemistry, Assay and Process Development.
Prof Edwin van den Heuvel	**University of Technology, Eindhoven, The Netherlands** Professor at the TU/e department of Mathematics and Computer Science where he will be closely involved with the development of the Data Science Center Eindhoven (DSC/e).
Cornelia Horoiu	**UCB Pharma SA, Brussels, Belgium** Quality Control, currently in a global position for management of the corporate Critical Materials.
Patricia Hughes, Ph.D.	**U.S. Food and Drug Administration (FDA), Silver Spring, USA** Branch Chief (Actg), Division of Microbiology Assessment, OPF/ OPQ/ CDER.
Peter Huonker	**Zimmer GmbH, Winterthur, Switzerland** Manager Micobiological Services.
Dr Pieta IJzerman-Boon	**MSD B.V., Oss, The Netherlands** Associate Director Quantitative Sciences/Center for Mathematical Sciences-Europe
Liselotte Kamper	**Novo Nordisk A/S, Bagsværd, Denmark** Author of Novo Nordisk internal procedure for handling of OOT results in stability studies.
Jan-Oliver Karo	**Paul-Ehrlich-Institut, German Agency for Vaccines and Biomedicines, Langen, Germany** Scientist Section 1/3 Microbial Safety.
Dr Marc Kelly	**MiCRA-Biodiagnostics, Institute of Technology Tallaght, Dublin, Ireland** Senior Scientist on Development of Process Sensors for Bacterial Contamination.
Dr Manuela Leitner	**AGES – Austrian Agency for Health and Food Safety** Quality Assessor for Biopharmaceuticals and Plasma Master File.
Jack Levin, M.D.	**University of California School of Medicine**
Heiko Linde	**Agilent Technologies, Waldbronn, Germany** Senior Sales Product Specialist responsible for Lab Informatics.
Frans A. Maris	**MSD B.V., Oss, The Netherlands** Head of the Quality Control Analytical Chemistry laboratories for finished products.

Robert J. Mello, Ph.D.	**Mello PharmAssociates, LLC, USA** Former Senior Microbiology Reviewer, New Drug Microbiology Staff, FDA.
Anna Mills	**Rapid Micro Biosystems, Inc., Bedford, USA** Senior Field Application Specialist.
Dr Jelena Novaković Jovanović	**Galenika A.D., Belgrad-Zemun, Serbia** Deputy Head of Microbiology in QC Sterile and Non Sterile Products.
Matthew Paquette	**Pfizer Biotech, USA** Quality Control Scientist II in Microbiology.
Dr France Audrey Peltier	**Merck Millipore, Germany** Product Manager Mycoplasma Media.
Dr Kent Persson	**Octapharma AB, Stockholm, Sweden** Project Manager, PCR Department.
Johannes Reich	**University Regensburg, Germany** PhD Student with focus on the aggregation and interaction of Lipopolysaccharides as well as the related activities in limulus based detection systems.
Prof Dr Renate Rosengarten	**Mycoplasma Biosafety Services GmbH, Wien, Austria** Managing Director, COO, CSO, BioTech Center.
Henrik Salling	**Novo Nordisk A/S, Gentofte, Denmark** Development Scientist, Biopharm Downstream Development & Virology.
Jan Jaap Schot	**MSD B.V., Oss, The Netherlands** Specialist Microbiology for Manufacturing & Quality / Center of Expertise Microbiology.
Dr Ingo Spreitzer	**Paul-Ehrlich-Institut, German Agency for Vaccines and Biomedicines, Langen, Germany** Deputy Section of Microbial Safety.
Dr Ferdinand Steierhoffer	**Boehringer Ingelheim Pharma GmbH & Co. KG, Biberach, Germany** Heading a dissolution lab within the Analytical Development Department.
Dr Masakazu Tsuchiya	**Charles River Laboratories, Charlston, USA** Senior Research Scientist in Endotoxin and Microbial Detection.
Dr Astrid Visser	**Sanquin Plasma Products B.V., Amsterdam, The Netherlands** Business Development Manager, Project Leader MAT Testing.
Helena Windsor	**Mycoplasma Experience, UK** Head and founder of Mycoplasma Experience Ltd. ; Testing Services, Contract Media, Contract R & D and consultancy services.
Dr Friedrich von Wintzingerode	**Roche Diagnostics GmbH, Penzberg, Germany** Senior Manager QC Microbiology. Lead of Endotoxin Expert Group Roche/Genentech.
Prof. Ulrich Zähringer	**Research Center Borstel, Germany**

ECA – Laboratory Informatics – Update 2015

Analytics

Similarities, Differences and Potential of Different Applications like LIMS, ELN, SDMS, PLM, ERP, etc.
➲ Peter J. Boogaard, Industrial Lab Automation

Regulatory Requirements Update (EU and US)
➲ Peter J. Boogaard, Industrial Lab Automation

Implementation and Validation of Lab Standard Systems - a risk-based Approach
➲ Dr Markus Dathe, F. Hoffmann-La Roche

Lab Systems - Going Paperless
➲ Dr Markus Dathe, F. Hoffmann-La Roche

Hot Topic: Lab Data Integrity
➲ Dr Markus Dathe, F. Hoffmann-La Roche

Trending, Data Integrity and Regulatory Guidance - Why QC Needs Paperless Informatics Tools Now
➲ Michael Goetter, Lonza Bioscience

Challenge Lab Data Integrity - Water's Approach for QA/QC
➲ Arjan Bannink, Waters

From Instruments to Decisions, Unifying and Integrating Laboratory Informatics
➲ Heiko Linde, Agilent Technologies

ECA – Endotoxin and Pyrogen Testing (Day 1)

Microbiology

The Limulus Amebocyte Lysate (LAL) Test for Bacterial Endotoxins Discovery, Development and Applications
➲ Jack Levin, M.D., University of California School of Medicine

Current Developments in Endotoxin / Pyrogen testing in the European and other Pharmacopeias
➲ Dr Ingo Spreitzer, Paul-Ehrlich-Institut, German Agency for Vaccines and Biomedicines

Kinetic Bacterial Endotoxin Assay Challenges for Biologics
➲ Dr Fatma Gökşin Bahar, Arven Pharmaceuticals

Increasing LAL Testing Efficiency with Endosafe® Nexus™ Robotic Endotoxin Testing System
➲ Matthew Paquette, Pfizer Biotech

An Improved Monocyte Activation Test Using Cryopreserved Pooled Human Mononuclear Cells
➲ Dr Astrid Visser, Sanquin Plasma Products

Challenges on Performing LAL in Oil Products
➲ Dr Jelena Novaković Jovanović, Galenika

Pyrogen and Endotoxin Analysis: Concepts and Considerations As We Move Forward
➲ Robert Mello, Ph.D., Mello PharmAssociates

LPS Aggregation Changes in Low Endotoxin Recovery –Seeing is believing
➲ Dr Masakazu Tsuchiya, Charles River Laboratories

Bild: HHAC Labor Dr. Heusler

ECA – Rapid Microbiological Methods

Mikrobiologie

SILVA & ARB: high quality ribosomal RNA gene databases and services
➲ Prof Frank Oliver Glöckner, Max Planck Institut & Jacobs University Bremen

Revision of European Pharmacopoeia Chapter 5.1.6
➲ Dr Emmanuelle Charton, EDQM

MICROPRINT BIOCARD: Imprinted Polymer Technology for the Rapid Detection of Microorganisms
➲ Dr Marc Kelly, MiCRA-Biodiagnostics, Institute of Technology Tallaght

Validation of a Sterility Test
➲ Anna Mills, Rapid Micro Biosystems

Rapid Enumeration with MU-scan: Risk or Improvement
➲ Jaap Schot/Dr Pieta Ijzerman-Boon, MSD

Modern Microbiological Safety Concepts – A Regulator´s View on Cell-based Products
➲ Jan-Oliver Karo, Paul-Ehrlich-Institut, German Agency for Vaccines and Biomedicines

Approaches for Validation of Rapid Sterility Testing Methods
➲ Dr Rajesh Gupta, Biologics Quality & Regulatory Consultants

Identification with MALDI-TOF
➲ Peter Huonker, Zimmer

The Conferences
11 November 2015

ECA – cGMP Compliance Trends in analytical Quality Control

Analytics

New GMP Requirements for the Analytical Lab: EU GMP Chapter 5 & 6 and the New FDA Guidance July 2015
➲ Ulla Bondegaard, Novo Nordisk

Regulations on Elemental (Metal) Impurities: Update on the status of the USP, FDA and ICH Guidelines – Path forward at Merck & Co.
➲ Frans Maris, MSD

New USP Chapter <1029> Good Documentation Guidelines - Impact on the Analytical Laboratories
➲ Ulla Bondegaard, Novo Nordisk

Chemical Reference Standards for Quality Control
➲ Cornelia Horoiu, UCB Pharma

Expiry Dates for Reagents, Solvents, Solutions, ...
➲ Cornelia Horoiu, UCB Pharma

Interface Method Development - Validation (Focus: Dissolution Testing)
➲ Dr Ferdinand Steierhoffer, Boehringer Ingelheim

Case Study: Handling of OOT Results
➲ Liselotte Kamper, Novo Nordisk

Automation in QC Labs / Efficient Documentation
➲ Dr Ferdinand Steierhoffer, Boehringer Ingelheim

ECA – Validation Approach of Bioassays Using Statistical Methods

Bioanalytics

Introduction
- Guidelines / Type of bioassays / Basic Statistics

Bioactivity (USP <111>, EP5.3)
- Calculation / Combination / Test set-up

Statistics for validation (USP<1032>, <1033>, <1034>)
- Accuracy / Precision / Sensitivity & Specificity / Linearity & Range / Limit of detection & quantitation / Robustness

➲ Dr. Pieta Ijzerman-Boon und Prof Edwin van den Heuvel

ECA – Endotoxin and Pyrogen Testing (Day 2)

Microbiology

Key Note: Everything You Always Wanted to Know About Endotoxin, But Were Afraid to Ask
➲ Prof. Ulrich Zähringer, Forschungszentrum Borstel

FDAs Current Thinking on LER
➲ Dr Patricia Hughes, CDER, FDA

Case Study: Overcoming Endotoxin Masking in a Drug Product
➲ Johannes Reich, University Regensburg

Endotoxin Masking – Origin, Natural Occuring Endotoxins and Demasking
➲ Peter Cornelis, Toxikon Europe

Development of a LAL-based method to overcome LER in a Biologics product
➲ Dr Friedrich v. Wintzingerode, Roche Diagnostics

Recombinant Factor C : Sustainable Alternative for Endotoxin Detection
➲ Elena Gustchina, Lonza

Reduction of Test-Interferences by Using a Recombinant Limulus Factor c ELISA
➲ Stefan Gärtner, Labor L+S

MAT testing with Cell Lines
➲ Dr. Anja Fritsch, Confarma France

ECA – Adventitious Agents – Impurities and Contaminants

Microbiology

Viral safety in biologicals – The regulatory perspective
➲ Dr Manuela Leitner, AGES – Austrian Agency for Health & Food Safety

Challenges in Testing for Adventitious Agents during Manufacture of Biological Products
➲ Dr Rajesh Gupta, Biologics Quality & Regulatory Consultants

Mycoplasma – Standards and Validation
➲ Prof Dr Renate Rosengarten, Mycoplasma Biosafety Services

Dive into traditional Mycoplasma culture method
➲ Dr France Audrey Peltier, Merck Millipore

PCR - Complementing Culture Expertise - The introduction of PCR into a culture based laboratory
➲ Helena Windsor, Mycoplasma Experience

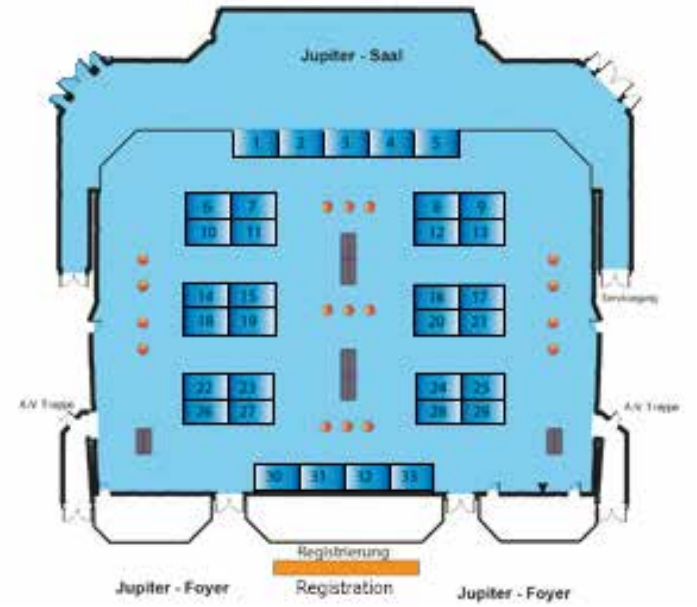
Selecting a rapid mycoplasma assay supporting recombinant production
➲ Dr Kent Persson, Octapharma

Experiences with in-house qPCR assay for Mycoplasma detection
➲ Henrik Salling, Novo Nordisk

The Exhibition

Is your company specialised in products and services for pharma laboratories?

As an exhibitor in the PharmaLab exhibition you can take advantage of the unique opportunity to directly address users and decision makers in the areas analytics, bioanalytics, from microbiological laboratories, Quality Assurance and Quality Control. In addition to high-level discussions during the Congress you can also get in touch with Congress delegates with speakers during the Social Event.



The **charges per stand are 3.980,- Euro** plus VAT. The following services are included:

- 2 one day tickets per 690,- Euro[1]
- Reduced one day tickets for inviting your customers
- Participation for the person mentioned on the form below is free of charge
- Lunch and refreshments during the conferences
- Participation in the Social Event
- Maximum size of the stand: app. 3 x 2 m
- 1 table, 2 chairs and power
- On-site support

If you want to be part of this industry meeting you should register your stand soon.

Materials for your Marketing

As an exhibitor you can take advantage of various marketing materials for promoting your presence. These include

- online exhibition banner – for your website and as signature in your e-mails.
- exhibition stickers – for your business mail
- an ad in the GMP Journal (subject to extra charges) – get directly in touch with your target group

You will find more detailed information on these materials on the Congress website at www.pharmalab-congress.com.

Sponsoring

Moreover, take advantage of the sponsoring opportunities to make Congress delegates aware of your company. In addition to sponsoring coffe breaks or lunches on the 1st and 2nd Congress day there are plenty of other sponsor possibilities. You will find the details on the website.

The Contacts

Do you have any questions with regard to the exhibition? Then please contact:
Detlef Benesch (Organisation Head), Phone +49 (0) 6221 84 44 45,
E-Mail: benesch@concept-heidelberg.de.

[1] *One day tickets will be mailed. Guests will need to register on the PharmaLab website at www.pharmalab-congress.com. Please note that one day tickets are not for exhibitor staff.*

Registration for the Exhibition – PharmaLab 2015

Registration for a stand at the PharmaLab 2015 on 10/11 November 2015 in Düsseldorf/Neuss.
For the registration of your stand you can also alternatively use the online registration form, which you will find on the website at **www.pharmalab-congress.com**. The charges for a stand are 3.980,- Euro plus VAT.
(Please note that exhibitors will be responsible for all charges for building and taking down of stand as well as for all materials related to the presentation.)

☐ **I want to register a stand with the stand number below.**
(Please note that for cancellation after 31 July 2015 the full registration fee of 3.980,- Euro will be charged. In addition, the General Terms and Conditions for Fairs/Exhibitions as available on the PharmaLab website do apply.)

The exhibitor plan on the website at www. pharmalab-congress.com is updated every day. Please take a look at this plan to see what spaces are still open and to pick your stand number which you then fill in here:

Preferred Stand Number: _____ or alternatively _____

Registration / Reservation – Company Information / Invoice Address:

Company	
Contact	
Department	
Phone / Fax	
E-Mail	

Contact on site – this person is also free to attend all conferences (registration as delegate included):

First & Last Name	
Department	
Street, ZIP & City	
Phone / E-Mail	
Invoice Address	
Participation in Social Event on 10 November 2015: Yes ☐ No ☐	

Additional Stand Personnel:
For additional stand personnel a flat rate of € 300, - will be charged per person. Please register additional personnel together with your registration as exhibitor. The participation of conferences is not included.

	Stand Personnel – Person 1:	Stand Personnel – Person 2:
Company		
First & Last Name		
Street, ZIP & City		
Phone / E-Mail		
Invoice Address		
Participation in Social Event on 10 November 2015:	Yes ☐ No ☐	Yes ☐ No ☐

<u>Conference Selection for Congress Delegate (not for Stand Personnel):</u>
PharmaLab 2015 delegates are free to attend the conferences they are interested in. To set up the conference rooms, though, we would appreciate it if you let us know what conference you are specifically interested in – **please mark your choice per day below**.

10 November	☐ ECA – Laboratory Informatics – Update 2015	11 November	☐ ECA – cGMP Compliance in Analytical Quality Control
	☐ ECA – Endotoxin and Pyrogen Testing (Day 1)		☐ ECA – Validation Approach of Bioassays
	☐ ECA – Rapid Microbiological Methods		☐ ECA – Endotoxin and Pyrogen Testing (Day 2)
			☐ ECA – Adventitious Agents – Impurities & Contaminants

<u>Room Reservation</u>:
Direct room reservation by reservation form! Reservations/bookings cannot be made through Concept Heidelberg. Receipt with confirmation/invoice.
CONCEPT HEIDELBERG has reserved a limited number of rooms in the Swissôtel Düsseldorf/Neuss. You can make your room reservation directly with the reservation form you will receive together with the registration confirmation. We recommend to register early.

Court of jurisdiction is Heidelberg, German law is applicable. In addition, the General Terms and Conditions for Fairs/Exhibitions as available on the PharmaLab website at www.pharmalab-congress.com do apply.

City and Date _____ Signature _____

Please complete the form and return to CONCEPT HEIDELBERG, Fax +49 (0) 6221 84 44 34.

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(06221) 84 44 34

E-Mail:
info@concept-heidelberg.de

Internet:
www.pharmalab-congress.de
www.pharmalab-congress.com

Registration Options PharmaLab 2015

☐ Attending the PharmaLab Conferences – One Day Ticket for € 690,-

☐ Attending the PharmaLab Conferences – Two Days Ticket for € 1.380,-

With a one day ticket/two days ticket you can attend any conference offered that day/both days. It includes participation in any conference on that day/on both days and the visit of the exhibition. In addition, it comprises lunch and beverages during the conferences and in breaks (on one or both days) as well as the social event on the evening of the first congress day. Please mark if you would like to attend the Social Event.

To be able to prepare the conference rooms, we would appreciate it if you marked the conference you are interested in. Please also mark the day you plan on attending the Congress. **Please mark only one conference per day.**

☐ I would like to attend on **day 1 (10 November 2015)** and I'm primarily interested in the conference:
- ☐ ECA – Laboratory Informatics – Update 2015
- ☐ ECA – Endotoxin and Pyrogen Testing (Day 1)
- ☐ ECA – Rapid Microbiological Methods

☐ I would also like to take part in the Social Event on the evening of 10 November.

☐ I would like to attend on **day 2 (11 November 2015)** and I'm primarily interested in the conference:
- ☐ ECA – cGMP Compliance Trends in Analytical QC
- ☐ ECA – Validation Approach of Bioassays Using Statistical Methods
- ☐ ECA – Endotoxin and Pyrogen Testing (Day 2)
- ☐ ECA – Adventitious Agents - Impurities and Contaminants

PLEASE NOTE:
- There will be no reservations via Concept Heidelberg. Please book your **hotel room directly with the reservation form** which you will receive together with your confirmation/invoice! Charges are payable after receipt of the invoice.
- There will **not be any print-outs** at the Congress. Instead you will receive all presentations prior to the Congress as Downloads. All Congress delegates will also receive the presentations on a USB stick at the registration center.

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

Reservation Form (Please complete in full)

☐ Mr ☐ Ms ☐ Dr

First name, Surname

Company

Department

Important: Please indicate your company's VAT ID Number P.O. Number (if applicable)

Street/P.O. Box

City Zip Code

Country

Phone/Fax

E-Mail (please fill in)

CONCEPT HEIDELBERG
P.O. Box 101764
Fax +49 (0) 62 21/84 44 34
D-69007 Heidelberg
GERMANY

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 %.

CONCEPT HEIDELBERG reserves the right to change the materials, instructors, or speakers without notice or to cancel an event. If the event must be cancelled, registrants will be notified as soon as possible and will receive a full refund of fees paid.

CONCEPT HEIDELBERGwill not be responsible for discount airfare penalties or other costs incurred due to a cancellation. Terms of payment: Payable without deductions within 10 days after receipt of invoice. 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)!

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.

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