

# PharmaFORUM Webcast International

Global Drug Safety & Global Regulatory Affairs

5-years  
ANNIVERSARY!  
One extra free  
webcast as a  
thank you.

## THE UPCOMING WEBCASTS AT A GLANCE

- India - Regulatory affairs & pharmacovigilance challenges
- China: Regulatory affairs & CMC requirements
- Food supplements & OTC
- Iran & Iraq - Marketing authorisation
- QPPV worldwide oversight

## YOUR BENEFITS

- One live webcast with international experts every two months
- Consolidated information in a short period of time at your work place
- Opportunity to interact directly with the speaker

## Concept

Do you work in international regulatory affairs or in pharmacovigilance? We would like to invite you to join us every two months for our live webcasts, where international regulatory affairs and vigilance experts will inform you of the latest news and trends in global marketing authorisation and drug safety within and beyond the ICH region.

You will meet our experts in a virtual conference room. Each meeting will take place as a 1.5-2 hour live webcast, presenting the latest news with supporting presentation slides (also for your personal download). Your questions will be coordinated by the chairperson and forwarded directly to the speaker.

---

## Additional benefits

Are you unable to attend one of the webcasts? No problem! After each live meeting, you will be able to retrieve the recorded webcast from our e-learning centre. This allows you to view each webcast at any time and as often as you like. An optional multiple choice test finalizes each webcast, giving you the possibility to receive a personal certificate.

Get a first impression at [www.forum-institut.com/ pharma-forum-webcast-international](http://www.forum-institut.com/pharma-forum-webcast-international)

## Programme: Webcast International 2022

---

### Your expert



#### Indu Nambiar

Boehringer Ingelheim India  
Pvt. Ltd., Mumbai, India  
Head-PV



#### Pooja Maingi

PharmaLex India Pvt. Ltd.,  
Noida, India,  
Head of Regulatory Affairs  
India



#### Yingying Liu

CSL Behring, Bern,  
Switzerland  
Associate Director  
Regulatory Affairs

### Date, time and programme

27 January 2022, 11:00 CET

#### Pharmacovigilance in India

- PV framework in India
- Similarities and differences compared to Europe
- Recent developments

27 January 2022, 14:00 CET

#### Regulatory Affairs in India

- Focus on procedures and procedure management

14 March 2022, 14:00 CET

#### Regulatory Update China

## Programme Webcast International 2022

---



**Dr Karola  
Krell Zbinden**  
FoodLex, Bern, Switzerland  
Lawyer

23 May 2022, 14:00 CET

### **Food supplements/over the counter drugs – export outside the EU area**

- Strategic aspects to consider upfront
- Typical pitfalls and questions to be answered
- Examples from specific countries



**Dr Makram Nehme**  
PAREXEL International Ltd.,  
Jdeide - Bouchrieh, Metn,  
Lebanon  
Regulatory Independent  
Consultant

6 July 2022, 14:00 CET

### **Marketing Authorisation of Drugs in Iraq**

- National marketing authorisation
- Procedure and recent changes

### **Marketing Authorisation of Drugs in Iran**

- Legal complexity due to sanctions against Iran

### **Dr Sabine Jeck-Thole**

Boehringer Ingelheim  
Pharma GmbH & Co. KG,  
Ingelheim, Germany  
EU QPPV & Head Regional PV

6 September 2022, 14:00 CET

### **A company's QPPV Network**

- EU QPPV: Worldwide oversight
- The post Brexit scenario
- QPPVs in other regions (EAEU, Arabian countries, etc.)
- Collaboration, exchange and overall responsibilities



**Dr Christina Juli**  
Boehringer Ingelheim  
Pharma GmbH & Co. KG,  
Biberach an der Riss,  
Germany  
Biopharma CMC Project  
Mgmt & Tech RA

23 November 2022, 14:00 CET

### **CMC requirements in China**

- Specific Chinese requirements to be considered in drug development, for submission and post-approval
- Specific CMC requirements – Challenges for global industry

# PharmaFORUM Webcast International

## HOW TO REGISTER

service@forum-institut.de      Tel +49 6221 500-500  
www.forum-institut.com      Fax +49 6221 500-555  
www.forum-institut.com/pharma-forum-webcast-international

## REGISTRATION

Yes, I want to join the

- PharmaFORUM Webcast International  
(you will receive a confirmation email  
with your login details)
- Yes, I agree that FORUM Institut may inform me  
about events and relevant expert content by:  
 email; and/or  telephone.  
I may withdraw my consent at any time.

---

Name

---

E-Mail (required for your login details)

---

Position

---

Company

---

Street address

---

Postal Code/City/Country

---

Tel. No.

---

Date, Signature

### Fee:

Membership of the PharmaFORUM Webcast International is available for one year. **The annual membership fee of € 900** (plus German VAT) for six webcasts is due upon registration.

Membership is automatically extended by one year, unless written notice has been submitted no later than six weeks before the end of the membership. A 12-month membership may be started at any time.

**If you are interested in a group account, please contact us.**

### Benefits:

The webcast will be recorded and made available to participants in their online archive.

Six live webcasts per year

- Recorded presentations since 2017 available at our e-learning centre
- Documentations for your personal download
- Multiple choice test and personal certificate after each webcast

## CANCELLATION POLICY

Our general terms and conditions apply (as of 01.01.2016) and are available upon request. We can send them to you anytime or you can find them online at [www.forum-institut.com/t&c](http://www.forum-institut.com/t&c)

## YOUR CONTACT



**Dr Henriette Wolf-Klein**  
Head of Department  
Pharma & Healthcare  
Phone: +49 6221 500-680  
[h.wolf-klein@forum-institut.de](mailto:h.wolf-klein@forum-institut.de)