Webcast series

FORUM

PharmaFORUM Webcast International

Global Drug Safety & Global Regulatory Affairs

The upcoming webcasts at a glance

- Market Access in Europe
- IDMP & SPOR worldwide
- Regulatory Affairs and Pharmacovigilance in Mexico
- Pharmacovigilance in Russia
- Clinical Trials in China
- Marketing Authorisation in Turkey

First impressions at
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Your benefits

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

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PharmaFORUM Webcast International

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 be held as a 1.5-2 hour live webcast, presenting the latest news with supporting presentation slides (also for your personal download). Your practical questions will be addressed directly via the chat function coordinated by the meeting chairperson.

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.

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Your experts in 2019

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Your Programme	Date and Time	Your Expert
Market Access in Europe ▌HTA assessment in UK, France, Germany and how they influence other countries' methodology and decisions ▌Early access schemes & compassionate use – how health systems evolve to accept earlier access while managing risk	23th January 2019 (14:00 CET)	Christian Hill
IDMP & SPOR ▌Status of SPOR implementation ▌Master Data Management and how to link/ align SPOR internally ▌Implementation of SPOR/IDMP and the impact on regulatory processes ▌GAP analyses & data collection with regard to IDMP in pharmaceutical companies	21st March 2019 (14:00 CET)	Remco Munnik
RA and PV in Mexico	6th May 2019 (12:00 CET)	Anita Patel
Pharmacovigilance in Russia ▌Legal background and national authorities ▌Local PSMF ▌EAEU QPPV ▌PV inspections ▌Future trends	9th July 2019 (14:00 CET)	Anna Kramar
Clinical trials in China	September 2019 (14:00 CET - Date will be shortly announced)	Yingying Liu
Marketing authorisation in Turkey ▌How to become a marketing authorisation holder in Turkey ▌Marketing authorisation application & procedure ▌Localization, Turkish GMP inspections for foreign manufacturers and further topics	20th November 2019 (14:00 CET)	Seda Kadioglu

PharmaFORUM Webcast International

Join our webcast series!

Would you like to register?

Via Fax: +49 6221 500-618 or

online: www.forum-institut.com/pharma-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.

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E-Mail (required for your login details)

Position

Company

Street address

Postal Code/City/Country

Tel. No.

Date, Signature

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How to register

Conference-No. 19 11 214

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:
- Six live webcasts per year
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