WEBCAST SERIES

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PharmaFORUM Webcast International

Global Drug Safety & Global Regulatory Affairs

THE UPCOMING WEBCASTS AT A GLANCE

- Marketing authorisation in Australia
- Pharmacovigilance in China
- Therapeutic Products Act (TPA) in Switzerland
- Global eCTD Lifecycle Management
- Regulatory Affairs in EAEU
- Post authorisation safety related changes – the process from signal to artwork

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 2020

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Your Programme	Date and Time	Your Expert
Marketing Authorisation in Australia	22 January 2020 (09:00 CET)	Greg Plunkett
PV in China • Legal background • Similarities to, and differences from, EMA • Role of affiliates, external consultants and agents • Practical examples and future trends	9 March 2020 (14:00 CET)	Dr Tanja Peters
Therapeutic Products Act (TPA) in Switzerland • Change of the "Heilmittelgesetz", focus regulatory affairs and pharmacovigilance	28 May 2020 (14:00 CEST)	Sylvia Schüpbach
Global eCTD Lifecycle Management • Change management and Variation management • Further product development and the challenges in eCTD • National eCTD peculiarities in a global environment	21 July 2020 (14:00 CEST)	Remco Munnik
Regulatory Affairs in EAEU • Update Customs Union • MRP, DCP • Cessation of national procedures?	22 Sept. 2020 (14:00 CEST)	Anna Kramar
Post authorisation safety related changes – the process from signal to artwork • Worldwide changes	17 Nov. 2020 (14:00 CET)	Susanne Weber

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HOW TO REGISTER

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REGISTRATION

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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
- Recorded presentations since 2017 available at our e-learning centre
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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

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