

# PHARMACOVIGILANCE FOR THE VETERINARY INDUSTRY

27 & 28 October 2009, Conf. No. V10-8209



## APPLICATION TO REGISTER

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## YOU MAY REGISTER BY:-

 +44 (0) 1483 730008

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 E-mail: registrations@management-forum.co.uk

If you have NOT received confirmation seven days after registering, please contact Registration Department.

## REGISTRATION INFORMATION

### Dates

27 October 2009 Start: 09.30 – Finish: 17.00

28 October 2009 Start: 09.00 – Finish: 16.30

### Registration & Coffee

27 October 2009 09.00

### Venue

The Rembrandt Hotel, 11 Thurloe Place, London SW7.

### Directions

Opposite V&A Museum.

Nearest Underground station: South Kensington.

Map available on Website under Hotels and Venues.

### Accommodation

A limited number of bedrooms have been reserved at The Rembrandt Hotel, 11 Thurloe Place, London SW7, at a special rate of £133.92 (Superior), £152.18 (Executive) both including English breakfast.

Excluding VAT – subject to availability.

A special rate for Friday, Saturday and Sunday of

£120.87 (Superior) including English breakfast

excluding VAT – subject to availability when booked as additional nights.

Hotel Tel: +44(0)20 7589 8100.

Hotel Fax: +44(0)20 7225 3476.

Email: reservations\_rembbrandt@sarova.co.uk

**All bookings should be made directly with the hotel or online at [www.sarova.com/rembrandt](http://www.sarova.com/rembrandt), quoting promo code 'manforum'.**

### Fee

**£1,150 + VAT.** The fee includes course documentation as well as mid-session refreshments and lunch. Invoice and confirmation will be forwarded to you.

### Conference No. V10-8209

### Discounted Rates

Available on application for personnel from non-profit making organisations and registered charities.

**Group discount available on request.**

### Cancellation Policy:

Over 14 days prior to the Seminar: Cancellation fee of £75. 7/14 days prior to the Seminar: 50% of the fee. Fewer than 7 days or if no notification received:

Registrant liable to pay FULL seminar fee.

**NB: Cancellations must be received in writing by**

**lesley@management-forum.co.uk.**

In the event of circumstances beyond its control,

Management Forum reserves the right to alter the

programme, the speakers, the date or the venue.

# PHARMACOVIGILANCE FOR THE VETERINARY INDUSTRY



**Gain a Practical Understanding of the Requirements to Ensure Your Company's Pharmacovigilance Strategy is Compliant**

## Benefits in attending:

- Understand the EU Pharmacovigilance Requirements
- Know How to Handle Animal SARs
- Learn How to Handle Human SARs
- Discover the Wider Scope of Pharmacovigilance
- Take Part in Pharmacovigilance Case Studies
- Hear About Electronic Communication in Pharmacovigilance
- Discover Principles of Pharmacovigilance and Data Resources
- Understand the Requirements of Signal Detection and Analysis
- Be Prepared for Pharmacovigilance Inspections
- Gain Practical Experience with a Pharmacovigilance Workshop on PSURs
- Discuss Risk Management and Know How to Deal with an Alert

With:

**Declan O'Rourke** Director, Ortec Consultancy, UK

Book on-line at [www.management-forum.co.uk](http://www.management-forum.co.uk) or telephone +44 (0) 1483 730071

**27 & 28 October 2009**  
**The Rembrandt Hotel, London**

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**Website: [www.management-forum.co.uk](http://www.management-forum.co.uk)**



## BENEFITS IN ATTENDING

**'Ensure your company's pharmacovigilance strategy is compliant.'**

As truly pan-European requirements for product development and registration have become a reality, there is an accompanying need for a consistent, homogenous and harmonised approach for the monitoring of adverse drug reactions and the sharing of safety information across the EU. The various EU Pharmacovigilance guidelines intend to achieve this. This seminar will provide an interpretation of the regulations and detailed guidance on how SARs should be handled, the requirements of PSURs and Pharmacovigilance Inspections. This will be reinforced by interactive case-based discussions and workshops. In addition, an overview of some of the future issues in respect of global harmonisation and electronic communication will be considered. **If you require any further information regarding this event please contact Andrea James on:**

**andrea.james@management-forum.co.uk**

There will be an option to discuss individual cases for review by the group. Please bring case details with you.

## WHO SHOULD ATTEND

Personnel in the veterinary industry from the following departments: **Research and Development, Veterinary Services, Adverse Reaction Monitoring, Regulatory Affairs and Registration, Drug Safety Officers** and all those involved and interested in the daily practice of pharmacovigilance within the EU will find this seminar beneficial.

**Reserve your place on-line at [www.management-forum.co.uk](http://www.management-forum.co.uk) or telephone +44 (0) 1483 730071. If you require further information contact [andrea.james@management-forum.co.uk](mailto:andrea.james@management-forum.co.uk)**

## COURSE LEADER

**Declan O'Rourke** Director, Ortec Consultancy. He has over 20 years experience in industry where he held technical, marketing, product development, clinical development, production and pharmacovigilance roles. He is a veterinary surgeon, holds a Diploma in Marketing, a Master of Business Administration and was awarded a Fellowship of the Royal College of Veterinary Surgeons in 1990. He now directs Ortec Consultancy specialising in EU pharmacovigilance and currently represents IFAH-EU in the VICH Working Group on pharmacovigilance.

## DOCUMENTATION

Delegates will receive a course material folder containing comprehensive documentation provided by the speakers, which will be a valuable source of reference for the future.

## ATTENDANCE LIMITED – EARLY REGISTRATION RECOMMENDED

This limitation, a unique feature of all MANAGEMENT FORUM seminars, will give participants the opportunity for a thorough discussion of the complex issues to be covered by the programme.

**A Certificate of Attendance for Professional Development will be given to each participant who completes the course.**

## PROGRAMME

### Day One

27 October 2009

- ▶ **Welcome and Introduction**
- ▶ **An Overview of EU Pharmacovigilance**
  - What are the requirements?
- ▶ **Pharmacovigilance (PhV) in the Revised EU Legislation**
- ▶ **How to Handle Animal SARs**
- ▶ **Handling Human SARs**
- ▶ **Understanding the Wider Scope of Pharmacovigilance**
- ▶ **Discussion followed by Lunch**
- ▶ **Pharmacovigilance Case Studies**
- ▶ **Electronic Communication in Pharmacovigilance (including VEDDRA)**
- ▶ **Discussion and Close of Day One**

### Day Two

28 October 2009

- ▶ **Principles of Pharmacovigilance and Data Resources**
- ▶ **Signal Detection and Analysis: What is Required?**
  - Company
  - Authority
- ▶ **Pharmacovigilance Inspections: Are you Ready?**
- ▶ **Discussion followed by Lunch**
- ▶ **Practical Workshop on PSURs**
  - Format
  - What is required?
  - Analysis of data
  - Incidence calculation
- ▶ **Risk Management**
  - Dealing with an alert
- ▶ **Discussion and Close of Seminar**