

A Regulatory Update on Animal Feed and Feed Additives in the EU, USA and China - To book online go to: management-forum.co.uk/2091

Date and venue

7-8 December 2017

Ref: 9912

The Rembrandt Hotel
11 Thurloe Place
London SW7 2RS
Tel: +44 (0)20 7589 8100

Registration and refreshments - 09.00

Day one	9.30 - 17.00
Drinks reception	17.00 - 18.00
Day two	9.00 - 16.30

Three ways to book

 management-forum.co.uk

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Fees and payment

EARLY BOOKING DISCOUNT

Book BEFORE 23 October 2017

£1199.00 + VAT = £1438.80 • €1679.00 + VAT = €2014.80

FULL PRICE

Book BEFORE 23 October 2017

£1399.00 + VAT = £1678.80 • €1959.00 + VAT = €2350.80

Multiple booking discount for 2nd or subsequent delegates - 15%

£1189.15 + VAT = £1426.98 • €1665.15 + VAT = €1998.18

Payment options

1. Invoice which can be paid by bank transfer or credit / debit card
2. Online through our secure website when registering

In-house training

Elements of this course can be run in-house and be tailored to your specific needs. Our experts come to you, saving you time and money. For more information contact Customer Services on +44 (0)20 7749 4730 or email inhouse@management-forum.co.uk

The small print

FEE: The fee includes all meals and refreshments for the duration of the course and a complete set of course materials. If you have any particular requirements please advise customer services when booking.

HOW TO REGISTER AND PAY: A VAT invoice and booking confirmation will be sent within 7 days, please contact us if you have not heard anything after that time. Payment can be made by credit/debit card, by bank transfer (for bank account details please see payment details section on our website). VAT no GB 341232109. Any questions please contact Customer Services on +44 (0)20 7749 4730. **ALL PAYMENTS MUST BE RECEIVED IN ADVANCE OF THE EVENT.**

MULTIPLE BOOKING DISCOUNTS: This discount may not be used in conjunction with any other offer

CANCELLATIONS AND TRANSFER: Once we have received your booking the place(s) are confirmed.

Delegate	Up to 28 days before course	27 to 14 days before course	13 to 0 days before course
Cancellation	10% admin fee	100% admin fee	100% admin fee
Transfers	Free	10% admin fee	100% admin fee
Substitution	Free	Free	Free

A maximum of one transfer is allowed. After the transfer no cancellation can be accepted and the full invoiced fee will be charged. Transfers are subject to payment of the difference on higher value courses. All cancellations must be received in written form.

For event cancellation policy and T&Cs see website

Annual Conference

A Regulatory Update on Animal Feed and Feed Additives in the EU, USA and China

7-8 December 2017 London



Programme to include:

- "REFIT" of EU Regulation (EC) N° 1831/2003 on feed additives
- The changing face of EFSA - engaging stakeholders, status of new EFSA guidance
- Re-evaluation of EU feed additives – challenges remaining after seven years
- EU ten-year renewals - a smaller hurdle for approved feed additives?
- EU procedures and timelines in relation to feed additive applications
- EFSA application handling and scientific evaluation
- Permitted claims and borderlines between feeds, veterinary medicines and feed additives - differences between the EU, USA and China
- US FDA approaches to the regulation of nutritional products for animals
- The legal framework for feed additives in China
- How will BREXIT affect the EU with respect to feed and feed additives?

To be chaired by:

Dr Elinor McCartney, Director, Pen & Tec Consulting Group, Spain

With an international panel of experts:

Dr Francisco Javier Piquer, Head of Animal Nutrition, General Directorate Agricultural Products & Markets, Spanish Ministry of Agriculture, Fisheries, Food & the Environment, (MAPAMA), Spain (invited)

Ruud Bremmers, Regal BV, The Netherlands

Dr Hannah Lester, Pen & Tec Consulting, Spain

Dr Marta Ponghellini, Veterinary & International Affairs, Unit G1, Animal Nutrition, DG Health and Consumers, EU Commission (invited)

Dr Manuela Tiramani, Head of Feed Unit, European Food Safety Authority (EFSA) (invited)

Raymond Ng, Food & Feed Regulatory & BD Manager, ChemLinked, REACH24 Consulting, China (invited)

Dr George Burdock, President, Burdock Consulting. USA

Maximise your networking by attending the drinks reception

Introduction

Meeting regulatory requirements for feed and feed additives in the EU and other key markets such as China and the USA are major challenges for businesses in the field of animal nutrition. This conference will review EU legislation, examine procedures and data requirements, and discuss to what extent EFSA-compliant data can be used to achieve approvals in China and the USA (or vice versa).

Where the industry is today

The EU has transformed its food legislation in the last two decades, creating EFSA (European Food Safety Authority) in 2002 and adopting a harmonised approach to food safety, "from farm to fork". The 2003 feed additive regulation introduced a central ("one door-one key") approval system for feed additives, involving the EU Commission, the EURL (European Union Reference Laboratory), EFSA and the Standing Committee on the Food Chain and Animal Health, with delegates from 28 Member States ('Comitology').

Regulation (EC) N° 1831/2003 re-categorised feed additives and extended the scope to include amino acids, silage agents and urea. New classes of feed additives were added, e.g. mycotoxin inactivators, feed hygiene condition enhancers. The EU completed its ban on antibiotic growth promoters in January 2006 and although coccidiostats remain as feed additives, maintaining approvals presents considerable challenges for FBOs (Feed Business Operators). Re-evaluation of around 500 feed additives started in 2010 and the EU has prohibited feed additives for which no re-evaluation dossier was submitted, or which fail EFSA's scrutiny. A new feed regulation, the feed material register and the catalogue of feed materials have all improved transparency in feed labelling, while allowing some physiological and functional claims.

Depending on intended use, the US FDA may regulate a product added to animal feed as either a drug or a feed ingredient. In either case, the Food Drug and Cosmetic Act requires the sponsor to obtain FDA approval or GRAS recognition prior to marketing. In the past, FDA has followed a policy of enforcement discretion to allow marketing of unapproved products if evaluated by the AAFCO feed ingredient definition process and listed in the AAFCO Official Publication. The FDA now encourages sponsors to use the food additive petition procedure for new products.

The Chinese Ministry of Agriculture introduced new guidance and legislation on feed additives, adopting some aspects of EU/EFSA, other aspects of USA FDA, and some uniquely Chinese approaches in the area of animal nutrition.

For these reasons many companies manufacturing or marketing feed additives wish to address, as far as possible, the regulatory requirements of EU, Chinese and US authorities in a single project. Informal workshops will enable delegates to work together to solve regulatory problems.

Who should attend

- Regulatory
- Business development
- Feed business operators
- Strategic marketing managers
- Registration managers
- Product managers
- R&D scientists
- Project managers
- Senior managers seeking a "snapshot" of current regulatory trends and challenges in feeds and feed additives in Europe, China and the USA.

To be Chaired by

Dr Elinor McCartney, Director, Pen & Tec Consulting Group, Spain

Expert faculty

Dr Francisco Javier Piquer, Head of Animal Nutrition, General Directorate Agricultural Products & Markets, Spanish Ministry of Agriculture, Fisheries, Food & the Environment, (MAPAMA), Spain (invited)

Ruud Bremmers, Regal BV, The Netherlands

Dr Hannah Lester, Pen & Tec Consulting, Spain

Dr Marta Ponghellini, Veterinary & International Affairs, Unit G1, Animal Nutrition, DG Health and Consumers, EU Commission (invited)

Dr Manuela Tiramani, Head of Feed Unit, European Food Safety Authority (invited)

Raymond Ng, Food & Feed Regulatory & BD Manager, ChemLinked, REACH24 Consulting, China (invited)

Dr George Burdock, President, Burdock Consulting, USA

Elements of this programme can be run within your organisation as an in-house programme. To find out more please contact Customer Services on +44 (0)20 7749 4730 or email inhouse@management-forum.co.uk

Day one

7 December 2017

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|---------|---|
| 09.30 ► | Welcome and introduction
Dr Elinor McCartney , Pen & Tec Consulting |
| 09.45 ► | Implementation of Regulation (EC) N° 1831/2003 on Additives for Use in Animal Nutrition - An update
<ul style="list-style-type: none"> • Re-evaluation of existing feed additives: where are we? • Recent developments / ongoing discussions Dr Francisco Javier Piquer , MAPAMA (invited) |
| 11.00 ► | Discussion |
| 11.15 ► | Refreshments |
| 11.45 ► | The Role of the EU and Comitology
<ul style="list-style-type: none"> • Scope of EU feed additive legislation - in or out of scope? • Differences between feeds, feed additives and veterinary products • Feed and feed additive claims - best practices • Post-EFSA opinion - Comitology • Special case of supplementary dossiers Dr Marta Ponghellini , EU Commission (invited) |
| 12.45 ► | Discussion |
| 13.00 ► | Lunch |
| 14.00 ► | Assessment of feed additives and application handling
<ul style="list-style-type: none"> • EFSA Feed Unit structure • Application handling - administrative and technical • The roles of the EFSA FEED Unit, the FEEDAP Panel and its Working Groups, Plenary and Working Group meetings • When and how to consult EFSA • Work programme for the next three years (2015-2018): priorities and perspectives Dr Manuela Tiramani , European Food Safety Authority (invited) |
| 15.00 ► | Discussion |
| 15.10 ► | Refreshments |
| 15.30 ► | Workshop one: Strategic options for animal nutrition products in the EU
Led by Ruud Bremmers , Regal BV and Dr Hannah Lester , Pen & Tec Consulting |
| 16.30 ► | Workshop feedback and discussion |
| 17.00 ► | End of day one |

17.00 - 18.00 Networking drinks reception for delegates and speakers

Day two

8 December 2017

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| 09.00 ► | Review of day one - Focus on EU/EFSA - BREXIT update
Dr Elinor McCartney , Pen & Tec Consulting |
| 09.15 ► | Study design and statistics - What EFSA wants
<ul style="list-style-type: none"> • Basic EFSA-compliant study designs and key end-points • Recent changes in statistics • Frequently asked questions from EFSA Dr Hannah Lester , Pen & Tec Consulting |
| 10.15 ► | Refreshments |
| 10.30 ► | The US FDA approach to the regulation of feed ingredients
<ul style="list-style-type: none"> • Legal and regulatory framework • Authorisation pathways for feed ingredients • Recent changes • Key differences from EU Dr George Burdock , Burdock Consulting |
| 11.30 ► | Discussion |
| 11.45 ► | The Chinese approach to feed additives
<ul style="list-style-type: none"> • Legal and regulatory framework • Recent changes Raymond Ng , REACH24 Consulting (invited) |
| 12.45 ► | Discussion |
| 13.00 ► | Lunch |
| 14.00 ► | Workshop two: Study design and statistics
Led by Dr Hannah Lester , Pen & Tec Consulting and Ruud Bremmers , Regal BV |
| 14.45 ► | Discussion, questions and answers |
| 15.00 ► | Refreshments |
| 15.15 ► | Workshop three: Managing dossier projects - best practices
Led by Ruud Bremmers , Regal BV & Dr Hannah Lester , Pen & Tec Consulting |
| 16.00 ► | Workshop feedback and discussion |
| 16.15 ► | Discussion, questions and answers |
| 16.30 ► | Close of forum |

A certificate of attendance for professional development will be available to each participant who completes the course.