

Workshop

16 February 2012

Radisson Blu Royal Hotel
Brussels, Belgium

Intertek

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CANTOX
HEALTH SCIENCES INTERNATIONAL

Regulation of Animal Feed Ingredients in the United States





Event Summary

Intertek Cantox is pleased to invite you to this workshop. Highlights include:

- **Comparison of current approval processes in the US, including AAFCO ingredient definition, food additive petition, and GRAS notification**
- **Aligning regulatory strategies in the EU and US to achieve expedient routes to market in both jurisdictions**
- **Similarities and differences between US and EU data requirements**

Who should participate?

If you are part of an animal feed company interested in understanding the routes to obtaining approval for new feed ingredients in the US, this workshop is for you!

Why should you participate?

Regulatory processes in the US are changing, and it is critical to understand the options open to any company wishing to gain acceptance to market a new feed ingredient. This workshop will provide a comparison of the current approval processes, with up-to-date information on the regulatory options available to an animal feed company. Particular focus will be given to the US Food and Drug Administration (FDA) Center for Veterinary Medicine (CVM) Generally Recognized as Safe (GRAS) Notification Pilot Program, which was recently introduced. The role of the Association of American Feed Control Officials (AAFCO) in the regulation of feed ingredients also will be covered. Understanding the importance of aligning feed strategies globally, this workshop will consider the design of studies in feed or target animals, which meet both US and EU requirements.

Speakers

Dr. David A. Dzanis, DVM, DACVN
CEO, Regulatory Discretion, Inc.

Dr. Robert Waltz
President
AAFCO

Dr. Ashley Roberts
Senior Vice President
Intertek Cantox

Dr. Elizabeth Lewis
Scientific & Regulatory Consultant
Intertek Cantox

Rajinder Kaur
Scientific & Regulatory Consultant
Intertek Cantox

Tentative Agenda (8:30-4:30) Thursday, 16 February 2012

- **Registration; Welcome & Introduction**
- **Understanding US Regulations for Substances Added to Feed**
Comparative analysis of the various approaches for obtaining regulatory clearance in the US
- **Role of AAFCO in the Regulation of Animal Feed**
Outline of the role AAFCO plays in the sale and distribution of animal feed in the US, including its function in developing feed ingredient definitions
- **Overview of the GRAS Process**
Review of the regulatory elements of the GRAS determination process
- **Data Requirements for a GRAS Dossier**
Discussion of the technical, safety and efficacy data requirements to support a successful GRAS determination and CVM notification
- **Case Studies**
Regulatory considerations for specific types of feed ingredients or their uses, such as GMOs, direct-fed microbials, enzymes, and nutraceuticals
- **Making the Transition from Food to Feed Ingredients**
Points to consider when extending the use of your food ingredient to animal feed
- **Developing Strategies to Meet Both US and EU Regulatory Requirements**
Evaluation of the similarities and differences in the data requirement and regulatory processes to gaining acceptance for feed ingredients in the EU and US
- **Workshop Q & A**



Being a leader in the preparation of successful GRAS notifications for human food substances, Intertek Cantox is well-positioned to assist clients with their GRAS determination for animal feed ingredients. In addition, Intertek Cantox has a wealth of expertise in developing regulatory strategies and preparing submissions for feed ingredients in the EU and US.

Whether you are working to market your new ingredient or are in the process of research and development of new ingredients, this workshop will guide you through the regulatory opportunities and changes in the US. The similarities and differences between the US and EU regulatory processes for substances added to feed will be highlighted allowing you to develop a global strategy.





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Regulation of Animal Feed Ingredients in the United States

Brussels, Belgium, 16 February 2012

Title: Dr. Mr. Mrs. Ms. Other:.....

Name:..... Surname:.....

Company:.....

Address:.....

City/Zip/Country:.....

Tel:..... Fax:.....

Email:.....

Payment

By Bank Wire Transfer (details will be provided upon registration)

By Credit Card: Mastercard Visa

Number:..... Expiry:...../..... CVC

Name on Card:

Date:.....

Signature:.....

Return by Fax to: +1 905 542-1011 or email to hcormack@cantox.com.

General information

Participation Fee

€ 495 by Bank Wire Transfer or \$675.00 USD by Credit Card

Cancellation Policy

You may cancel 10 business days before the seminar date and receive a full refund less the cancellation fee of € 50. If you cancel less than 10 business days before the seminar date, no refund will be given, however, participation substitutions will be accepted.

Hotel Reservation

Delegates are responsible for organizing their own travel and accommodation. A limited number of rooms have been reserved for conference delegates wishing to stay at the Radisson Blu at a special rate. Wherever possible accommodation should be reserved as soon as possible as rooms cannot be guaranteed and rates are subject to change.

To book your accommodation, please contact the hotel directly, quoting "USFeed" when booking.

Conference Venue

Radisson Blu Royal Hotel
Rue du Fossé-aux-Loups 47
Wolvengracht
1000 Brussels, Belgium

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Workshop Coordinator

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