



TRAINING

« FEED REGULATION IN THE EU: WHAT ARE THE REGULATORY CONSTRAINTS AND OPPORTUNITIES? »

The aim of this training course is to introduce or validate the knowledge of any fed business operator wishing to learn more about the regulations governing animal feed in Europe. You will find all the information you need to better understand the constraints and opportunities involved in marketing a product that complies with the regulations.



MONDAY 08 & TUESDAY 09 DECEMBER 2025

- Validate your regulatory knowledge on feed.
- Understand the constraints.
- Study and work on one or more concrete cases.
- Q&A - Exchange and debate about the subject.

Public

- Anyone involved in the process of developing and/or marketing animal feed (CEO, Project Managers, R&D, Regulatory or Marketing departments).

Prerequisites

- No specific prerequisites.

Level

No level required

Terms

Remote Training (Webinar)

Accessibility

If a specific adaptation is needed, particularly in relation to a disability, please do not hesitate to let us know.



REGISTRATION FORM TO BE RETURNED TO
CONTACT@PHARMANAGER-DEVELOPMENT.COM

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Two Afternoons : 08 & 09 DECEMBER 2025 (2.00pm- 5.30pm)

Overview of regulatory texts

- Introduction
 - The main regulatory reference texts
 - General texts
 - Specific texts
- Complementary feeds (Animal) vs Dietary supplements (Humans): 2 different regulations!
- Feed Compliance
 - Undesirable substances
 - Pesticides
 - Official controls

=> Quizzes, exercises: validate your knowledge

Placing on the market of animal feed

- Definitions
- The different types of feeds (complementary feed, complete feed, compound feed, etc.)
- Case studies
- Registration

Raw materials and additives

- How can they be differentiated?
- Special case of plant extracts
- Registers and catalogue
- New applications for authorisation: what are the obligations?

=> Quizzes, exercises: validate your knowledge

Labelling rules

- General mandatory information
- Specific mandatory information
- Voluntary labelling

=> Quizzes, exercises: validate your knowledge



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TRAINERS



Marie Coulange

Regulatory and Scientific
Affairs Officer

5 years of experience

MEANS AND MODALITIES

The training will be conducted via the Teams video conferencing tool (a link will be sent to participants before the session for access). The training material (PowerPoint) will be presented through a combination of lectures and discussions with the trainers. Educational documentation will be provided in electronic format. A knowledge assessment must be completed at the end of the training. A certificate of attendance will be issued.



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REGISTRATION FORM

« FEED REGULATION IN THE EU: WHAT ARE THE REGULATORY CONSTRAINTS AND OPPORTUNITIES? »

Company name:

Name of signatory:

Intracommunity VAT number:

Company Training Manager:

(email required - Otherwise: Person signing the training agreement)

PARTICIPANTS AND CHOICE OF SESSIONS

Name :
First name :
Function :
E-mail :

Name :
First name :
Function :
E-mail :

Name :
First name :
Function :
E-mail :

If a particular adaptation of the training is necessary, linked in particular to a particular disability situation, do not hesitate to let us know.

Before registering, make sure you have read the training prerequisites and validate that the objectives are in line with expectations. Do not hesitate to let us know your expectations before the training if necessary.

Individual expectations and objectives:

Total : € tax ex.
Name and address of the company to be invoiced:
Date :
Signature :

Pharmanager Development reserves the right to cancel the training in the event of insufficient registrations. In the event of cancellation, Pharmanager Development will notify participants at least 5 working days before the scheduled training date.

IMPORTANT: Our training courses are eligible for support by an OPCO. However, the invoice will be established and due by the participating company, in the amount of the training.



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