Fendov 1250 avec Systamex, dispositif intraruminal à libération séquentielle pour bovins

Authorised

- Oxfendazole

Product identification

Medicine name:
Fendov 1250 avec Systamex, dispositif intraruminal à libération séquentielle pour bovins

Active substance:
Oxfendazole

Target species:
Cattle

Route of administration:
Oral use

Product details

Active substance and strength:
Oxfendazole
1250.00 milligram(s) / 1.00 Dose

Pharmaceutical form:
Pulsatile-release intraruminal device

Withdrawal period by route of administration:

Oral use:

- **Cattle**

  - Meat and offal. 7 month

  - Milk. no withdrawal period

Do not use in animals producing milk for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Luxembourg

Package description:
Fendov 1250 avec Systamex 12 Container Pulsatile-release intraruminal device
Fendov 1250 avec Systamex 24 Container Pulsatile-release intraruminal device

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
14/07/1989

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Le Ministère De La Santé Division De La Pharmacie Et Des Médicaments

Authorisation number:
V 087/91/11/0335

Date of authorisation status change:
28/08/2007

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.
