Zerofen 100 mg/ml, vet. mixtúra

Authorised

- Fenbendazole

Product identification

Medicine name:
Zerofen 100 mg/ml, vet. mixtúra

Active substance:
Fenbendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Fenbendazole
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 96 hour
- **Sheep**
  - Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC13

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

Authorisation status:
Valid

Authorised in:
Iceland

Available in:
Iceland

Package description:
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
21/09/2001

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/01/004/01

Date of authorisation status change:
11/02/2022

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Package Leaflet and Labelling

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