Oramec Drench vet. 0,8 mg/ml Mixtúra, lausn Handa sauðfé og geitum.

Authorised

- Ivermectin

Product identification

Medicine name:
Oramec Drench vet. 0,8 mg/ml Mixtúra, lausn Handa sauðfé og geitum.

Active substance:
Ivermectin

Target species:
Sheep
Goat

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
0.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:
Oral use:

- **Sheep**
  - Meat and offal. 6 day

Ekki má meðhöndla dýr sem gefa af sér mjólk til manneldis.

- **Goat**
  - Meat and offal. 8 day

Ekki má meðhöndla dýr sem gefa af sér mjólk til manneldis.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
1/04/1992

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France
Merck Sharp & Dohme B.V.

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
910066

Date of authorisation status change:
18/01/2022

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.

Labelling

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