IVERMIX, 0,15mg/g, Perorální prášek

Authorised

- Ivermectin

Product identification

Medicine name:
IVERMIX, 0,15mg/g, Perorální prášek

Active substance:
Ivermectin

Target species:
Fallow deer
Roe deer
Mouflon
Wild boar
Red deer

Route of administration:
In-feed use

Product details

Active substance and strength:
Ivermectin
0.15 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
In-feed use:

- **Fallow deer**
  - Meat and offal. 28 day
- **Roe deer**
  - Meat and offal. 28 day
- **Mouflon**
  - Meat and offal. 28 day
- **Wild boar**
  - Meat and offal. 14 day
- **Red deer**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Tekro spol. s r.o.

Marketing authorisation date:
1/11/2011

Manufacturing sites for batch release:
Tekro spol. s r.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/096/11-C

Date of authorisation status change:
8/10/2021

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.