Prazimex, 46,9mg/ml, Injekční roztok

Authorised

- Praziquantel

Product identification

Medicine name:
Prazimex, 46,9mg/ml, Injekční roztok

Active substance:
Praziquantel

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Praziquantel
46.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 4 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA01

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:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharmagal Bio spol. s r.o.

Marketing authorisation date:
30/11/2020

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

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

Authorisation number:
96/073/20-C

Date of authorisation status change:
30/11/2020

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

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Labelling

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