CEVAMEC 1 %, injekcinis tirpalas

Not authorised

- Ivermectin

Product identification

Medicine name:
CEVAMEC 1 %, injekcinis tirpalas

Active substance:
Ivermectin

Target species:
Cattle
Sheep
Goat
Pig

Route of administration:
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 49 day
  - Milk. no withdrawal period

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

- **Sheep**
  - Meat and offal. 21 day
  - Milk. no withdrawal period

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

- **Goat**
  - Meat and offal. 28 day
  - Milk. no withdrawal period

Not authorized for use in lactating females, whose milk will be used for human consumption and pregnant females, whose milk will be used for human consumption, 28 days before delivery.

- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:
Surrendered

Authorised in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
23/03/2000

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/00/1080/001-005

Date of authorisation status change:
6/11/2010

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

Documents

RV1080.pdf