Dewo 100mg/ml šķīdums injekcijām liellopiem, aitām, kazām, cūkām, suņiem, kaķiem

Not authorised

- Levamisole hydrochloride

Product identification

Medicine name:
Dewo 100mg/ml šķīdums injekcijām liellopiem, aitām, kazām, cūkām, suņiem, kaķiem

Active substance:
Levamisole hydrochloride

Target species:
Cat
Cattle
Sheep
Goat
Pig
Dog

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Levamisole hydrochloride
117.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Intramuscular use:

- **Cattle**

- Meat and offal. 8 day

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Nelietot dzīvniekiem, no kuriem iegūst pienu, ko paredzēts izmantot cilvēku uzrturā.

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AE01

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

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Authorisation status:
Surrendered

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Bremer Pharma GmbH

Marketing authorisation date:
25/10/1995

Manufacturing sites for batch release:
Bremer Pharma GmbH

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/95/0155

Date of authorisation status change:
29/06/2022

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

Documents

Summary of Product Characteristics

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

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.