Veyxyl LA 20% 200 mg/ml suspensija injekcijām liellopiem, cūkām, aitām, suņiem un kaķiem

Authorised

- Amoxicillin

Product identification

Medicine name:
Veyxyl LA 20% 200 mg/ml suspensija injekcijām liellopiem, cūkām, aitām, suņiem un kaķiem

Active substance:
Amoxicillin

Target species:
Cattle
Dog
Cat
Sheep
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Amoxicillin
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 28 day

Nav reģistrēts lietošanai aitām, kuru pienu paredzēts izmantot cilvēku uzturā.

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Veyx Pharma GmbH

Marketing authorisation date:
26/10/1999

Manufacturing sites for batch release:
Veyx Pharma GmbH

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/99/1035

Date of authorisation status change:
26/10/1999

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.