Veyx yl LA 20%, 200 mg/ml süstesuspensioon veistele, sigadele, lammastele, koertele ja kassidele

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Veyx yl LA 20%, 200 mg/ml süstesuspensioon veistele, sigadele, lammastele, koertele ja kassidele

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Pig
Sheep
Dog
Cat

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Amoxicillin trihydrate
229.60 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
- **Pig**
  - Meat and offal. 28 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Estonia

Package description:
Available only in Estonian
Available only in Estonian

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:
Veyx Pharma GmbH

Marketing authorisation date:
6/04/2006

Manufacturing sites for batch release:
Veyx Pharma GmbH

Responsible authority:
State Agency Of Medicines

Authorisation number:
1390

Date of authorisation status change:
6/04/2006

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.