COMBI-kel 40 suspensija injekcijām zirgiem, liellopiem, cūkām, aitām, kazām, suņiem, kaķiem

Authorised

- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate

Product identification

Medicine name:
COMBI-kel 40 suspensija injekcijām zirgiem, liellopiem, cūkām, aitām, kazām, suņiem, kaķiem

Active substance:
Benzylpenicillin procaine
Dihydrostreptomycin sulfate

Target species:
Cattle
Goat
Sheep
Horse
Pig
Dog
Cat

Route of administration:
Intraperitoneal use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Benzylpenicillin procaine
200000.00 international unit(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate
200000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intraperitoneal use:

- **Cattle**
  - Meat and offal. 30 day
  - Milk. 4 day
- **Goat**
  - Milk. 4 day
  - Meat and offal. 30 day
- **Sheep**
  - Milk. 4 day
  - Meat and offal. 30 day
- **Horse**

- Milk. no withdrawal period

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

Intramuscular use:

- **Cattle**
  - Meat and offal. 30 day
  - Milk. 4 day
- **Sheep**
  - Meat and offal. 30 day
  - Milk. 4 day
- **Goat**
  - Meat and offal. 30 day
  - Milk. 4 day
- **Horse**
  - Meat and offal. no withdrawal period

Nav reģistrēts lietošanai zirgiem, kuru gaļu paredzēts izmantot cilvēku uzturā.

- **Pig**
  - Meat and offal. 30 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RA01

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

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Authorisation status:
Valid

Authorised in:
Latvia

Package description:
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:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
1/08/2003

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/98/0716

Date of authorisation status change:
3/08/2003

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.