Bicormicina L.A. suspensija injekcijām liellopiem, zirgiem, kazām, cūkām, suņiem un kaķiem

Authorised

- Benzylpenicillin procaine
- Benzathine benzylpenicillin
- Dihydrostreptomycin sulfate
- Dexamethasone sodium phosphate
- Dexamethasone isonicotinate

Product identification

Medicine name:
Bicormicina L.A. suspensija injekcijām liellopiem, zirgiem, kazām, cūkām, suņiem un kaķiem

Active substance:
Benzylpenicillin procaine
Benzathine benzylpenicillin
Dihydrostreptomycin sulfate
Dexamethasone sodium phosphate
Dexamethasone isonicotinate

Target species:
Cat
Cattle
Horse (non food-producing)

Goat
Pig
Dog

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Benzylpenicillin procaine
125000.00 international unit(s) / 1.00 millilitre(s)

Benzathine benzylpenicillin
125000.00 international unit(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate
250.00 milligram(s) / 1.00 millilitre(s)

Dexamethasone sodium phosphate
0.20 milligram(s) / 1.00 millilitre(s)

Dexamethasone isonicotinate
0.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 60 day
  - Milk. 14 day Pienam: 14 dienas (28 slaukšanas reizes)
- **Goat**
  - Milk. 14 day Pienam: 14 dienas (28 slaukšanas reizes)

- Meat and offal. 60 day

- **Pig**
  - Meat and offal. 60 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RV01

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

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
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:
Fatro S.p.A.

Marketing authorisation date:
27/09/2002

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/02/1495

Date of authorisation status change:
29/09/2002

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

Documents

Combined File of all Documents

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