

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:

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Cattle

- Meat and offal. 60 day

- Milk. 14 day
Pienam: 14 dienas (28 slaukšanas reizes)

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Goat

- Milk. 14 day
Pienam: 14 dienas (28 slaukšanas reizes)

- Meat and offal. 60 day

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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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