Streptocillin vet. injeksjonsvæske, suspensjon

Authorised

- Dihydrostreptomycin
- Benzylpenicillin procaine

Product identification

Medicine name:
Streptocillin vet. injeksjonsvæske, suspensjon

Active substance:
Dihydrostreptomycin
Benzylpenicillin procaine

Target species:
Cattle
Pig
Sheep
Goat
Horse
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use
Intrauterine use
Intraperitoneal use

Intrasynovial use

Product details

Active substance and strength:
Dihydrostreptomycin
250.00 milligram(s) / 1.00 millilitre(s)

Benzylpenicillin procaine
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat. 60 day
  - Milk. 3 day
- **Pig**
  - Meat. 60 day
- **Sheep**
  - Meat. 60 day
  - Milk. 3 day
- **Goat**
  - Meat. 60 day
  - Milk. 3 day
- **Horse**

- Meat. 60 day

Subcutaneous use:

- **Cattle**
  - Meat. 60 day
  - Milk. 3 day
- **Pig**
  - Meat. 60 day
- **Sheep**
  - Meat. 60 day
  - Milk. 3 day
- **Goat**
  - Meat. 60 day
  - Milk. 3 day
- **Horse**
  - Meat. 60 day

Intrauterine use:

- **Cattle**
  - Meat. 6 day
  - Milk. 3 day
- **Pig**
  - Meat. 6 day
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Sheep

- Meat. 6 day
- Milk. 3 day

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Goat

- Meat. 6 day
- Milk. 3 day

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Horse

- Meat. 6 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:
Norway

Available in:
Norway

Package description:
Available only in Norwegian
Available only in Norwegian
Available only in Norwegian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Denmark A/S

Marketing authorisation date:
19/11/1952

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
3600

Date of authorisation status change:
29/04/2007

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

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