Pendistrep, 200 mg/mL, 250 mg/mL, suspenzija za injekciju, za goveda, ovce, svinje i konje koji se ne koriste za hranu

Authorised

- Dihydrostreptomycin sulfate
- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
Pendistrep, 200 mg/mL, 250 mg/mL, suspenzija za injekciju, za goveda, ovce, svinje i konje koji se ne koriste za hranu

Active substance:
Dihydrostreptomycin sulfate
Benzylpenicillin (procaine) monohydrate

Target species:
Cattle
Sheep
Pig
Horse (non food-producing)

Route of administration:
Intramuscular use

Product details

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

Benzylpenicillin (procaine) monohydrate
200.00 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. 6 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 6 day
- **Pig**
  - Meat and offal. 28 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:
Croatia

Available in:
Croatia

Package description:
Available only in Croatian
Available only in Croatian

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:
Laboratorios Syva S.A.

Marketing authorisation date:
14/03/2021

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/21-01/204

Date of authorisation status change:
5/04/2024

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

Documents

Summary of Product Characteristics

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