Penstrep-ject, suspensie voor injectie

Authorised

- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate

Product identification

Medicine name:
Penstrep-ject, suspensie voor injectie

Active substance:
Benzylpenicillin procaine
Dihydrostreptomycin sulfate

Target species:
Cattle
Pig
Pig (piglet)
Cattle (calf)
Sheep
Sheep (lamb)
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Benzylpenicillin procaine
200000.00 international unit(s) / 1.00 millilitre(s)

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

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 5 day
  - Meat and offal. 56 day
- **Pig**
  - Meat and offal. 49 day
- **Pig (piglet)**
  - Meat and offal. 49 day
- **Cattle (calf)**
  - Meat and offal. 56 day
- **Sheep**
  - Meat and offal. 56 day
- **Sheep (lamb)**

- Meat and offal. 56 day

Subcutaneous use:

- **Cattle**
  - Milk. 5 day
  - Meat and offal. 56 day
- **Pig**
  - Meat and offal. 49 day
- **Pig (piglet)**
  - Meat and offal. 49 day
- **Cattle (calf)**
  - Meat and offal. 56 day
- **Sheep**
  - Meat and offal. 56 day
- **Sheep (lamb)**
  - Meat and offal. 56 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RA01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

---

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
30/08/2007

Manufacturing sites for batch release:
Dopharma International B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 100679

Date of authorisation status change:
16/07/2019

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.