Pen & Strep Suspension for Injection

Authorised

- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate

Product identification

Medicine name:
Pen & Strep Suspension for Injection

Active substance:
Benzylpenicillin procaine
Dihydrostreptomycin sulfate

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Benzylpenicillin procaine
200.00 milligram(s) / 1.00 millilitre(s)

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 48 hour
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 48 hour
- **Pig**
  - Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RA01

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:

Multidose 50 ml uncoloured Type II (Ph. Eur.) plastic (PET) vial sealed with a bromobutyl rubber stopper and aluminium cap, containing a white to off-white sterile suspension.
Multidose 100 ml uncoloured Type II (Ph. Eur.) plastic (PET) vial sealed with a bromobutyl rubber stopper and aluminium cap, containing a white to off-white sterile suspension.
250 ml plastic (PET) vials sealed with a bromobutyl rubber stopper andaluminium cap, containing a white to off-white sterile suspension.
Multidose 100 ml uncoloured Type II (Ph. Eur.) glass vial sealed with a bromobutyl rubber stopper and aluminium cap, containing a white to off-white sterile suspension.
Multidose 50 ml uncoloured Type II (Ph. Eur.) glass vial sealed with a bromobutyl rubber stopper and aluminium cap, containing a white to off-white sterile suspension.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
1/10/1987

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/009/001

Date of authorisation status change:

1/10/1987

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

Documents

Summary of Product Characteristics