PROBENCIL 300 mg/ml suspension for injection for cattle and pigs

Authorised

- Benzylpenicillin procaine

Product identification

Medicine name:
PROBENCIL 300 mg/ml suspension for injection for cattle and pigs
Probencil 300 mg/ml Zawiesina do wstrzykiwań

Active substance:
Benzylpenicillin procaine

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 6 day — For treatment duration of 3-5 days
  - Meat and offal. 8 day — For a treatment duration of 6-7 days
  - Milk. 96 hour
- **Pig**
  - Meat and offal. 6 day — For a treament duration of 3-5 days
  - Meat and offal. 8 day — For a treatment duration of 6-7 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE09

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

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
250 ml colourless polyethylene terephthalate (PET) bottle with type I bromobutyl rubber stopper and flip-off cap.Pack size: Carton box with 1 bottle of 250 ml.

100 ml colourless polyethylene terephthalate (PET) bottle with type I bromobutyl rubber stopper and flip-off cap.Pack size: Carton box with 1 vial of 100 ml.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Mevet S.A.

Marketing authorisation date:
16/05/2019

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2873

Date of authorisation status change:
16/05/2019

Reference member state:
Ireland

Procedure number:
IE/V/0398/001

Concerned member states:
Poland Spain

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 19/01/2025
Download

Package Leaflet

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

Labelling

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

Combined File of all Documents

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