Procipen 300 mg/ml suspension for injection for cattle, sheep and pigs

Authorised

- Benzylpenicillin procaine

Product identification

Medicine name:
Procipen 300 mg/ml suspension for injection for cattle, sheep and pigs
Procipen 300 mg/ml Suspensie voor injectie
Procipen 300 mg/ml Suspension injectable
Procipen 300 mg/ml Injektionssuspension

Active substance:
Benzylpenicillin procaine

Target species:
Cattle
Sheep
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. 10 day — For treatment duration 3 days.
  - Meat and offal. 12 day — For treatment duration of 4-7 days.
  - Milk. 108 hour
- **Sheep**
  - Meat and offal. 4 day — For treatment duration of 3 days.
  - Meat and offal. 6 day — For treatment duration of 4-7 days.
- **Pig**
  - Meat and offal. 7 day — For treatment duration of 3 days.
  - Meat and offal. 9 day — For treatment duration of 4-7 days.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE09

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

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Authorisation status:
Valid

Authorised in:
Belgium

Package description:
The product is packaged in clear 100 ml Type I and 250 ml clear Type III glass vials sealed with red or grey, siliconised bromobutyl rubber stoppers and aluminium overseal, containing a sterile aqueous suspension. Carton box with 48 boxes containing 1 vial of 250 mlThe vials are colourless.
The product is packaged in clear 100 ml Type I glass vials sealed with red or grey, siliconised bromobutyl rubber stoppers and aluminium overseal, containing a sterile aqueous suspension. Carton box with 48 boxes containing 1 vial of 100 mlThe vials are colourless.
The product is packaged in clear 100 ml Type I and 250 ml clear Type III glass vials sealed with red or grey, siliconised bromobutyl rubber stoppers and aluminium overseal, containing a sterile aqueous suspension. 12 shrink-wrapped boxes containing 1 vial of 250 mlThe vials are colourless.
The product is packaged in clear 100 ml Type I glass vials sealed with red or grey, siliconised bromobutyl rubber stoppers and aluminium overseal, containing a sterile aqueous suspension. 12 shrink-wrapped boxes containing 1 vial of 100 mlThe vials are colourless.
The product is packaged in clear 100 ml Type I and 250 ml clear Type III glass vial sealed with red or grey, siliconised bromobutyl rubber stopper and aluminium overseal, containing a sterile aqueous suspension. Carton box with 1 vial of 250 mlThe vial is colourless.
The product is packaged in clear 100 ml Type I glass vial sealed with red or grey, siliconised bromobutyl rubber stopper and aluminium overseal, containing a sterile aqueous suspension. Carton box with 1 vial of 100 mlThe vial is colourless.

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:
Bimeda Animal Health Limited

Marketing authorisation date:
22/06/2021

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
22/06/2021

Reference member state:
Ireland

Procedure number:
IE/V/0416/001

Concerned member states:
Austria Belgium Denmark Estonia Finland France Germany Italy Latvia Lithuania Netherlands Norway Poland Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 16/01/2026
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Labelling

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