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Geepenil® vet.24 g, powder and solvent for solution for injection

Authorised

- Benzylpenicillin sodium

Product identification

Medicine name:
Geepenil® vet.24 g, powder and solvent for solution for injection

Active substance:
Benzylpenicillin sodium

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Benzylpenicillin sodium
24.00 gram(s) / 1.00 Vial

Pharmaceutical form:
Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 10 day
  - Milk. 2 day
- **Pig**
  - Meat and offal. 10 day

Intravenous use:

- **Horse**
  - Meat and offal. 10 day
- **Cattle**
  - Meat and offal. 10 day
  - Milk. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE01

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish

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:
Orion Corporation

Marketing authorisation date:
15/05/1998

Manufacturing sites for batch release:
Orion Corporation

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
13563

Date of authorisation status change:
15/05/1998

Reference member state:
Sweden

Procedure number:
SE/V/0118/001

Concerned member states:
Iceland

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

Documents

Package Leaflet

English (PDF)
Published on: 16/11/2026
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Summary of Product Characteristics

English (PDF)
Published on: 16/11/2026
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Labelling

English (PDF)
Published on: 14/10/2025
Updated on: 13/03/2026
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