

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:

Iceland

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:

7/06/2012

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Icelandic Medicines Agency

Authorisation number:

IS/2/12/004/01

Date of authorisation status change:

7/06/2012

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

Summary of Product Characteristics

English (PDF)

Published on: 28/07/2025

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Package Leaflet

English (PDF)

Published on: 28/07/2025

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Labelling

English (PDF)

Published on: 14/10/2025

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