

Glucobel 40 g/100 ml

Authorised

Infusionslösung für Pferde, Rinder,
Schafe, Ziegen, Schweine, Hunde
und Katzen

- Glucose monohydrate

Product identification

Medicine name:

Glucobel 40 g/100 ml Infusionslösung für Pferde, Rinder, Schafe, Ziegen, Schweine, Hunde und Katzen

Active substance:

Glucose monohydrate

Target species:

Cattle

Dog

Goat

Sheep

Horse

Cat

Pig

Route of administration:

Intravenous use

Product details

Active substance and strength:

Glucose monohydrate

440.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

-

Cattle

- Meat and offal. 0 day
- Milk. 0 hour

-

Goat

- Meat and offal. 0 day
- Milk. 0 hour

-

Sheep

- Meat and offal. 0 day
- Milk. 0 hour

-

Horse

- Meat and offal. 0 day
- Milk. 0 hour

-

Pig

- Meat and offal. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BA03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

1 x 500 ml polypropylene bottle with bromobutyl stopper and aluminium cap

1 x 750 ml polypropylene bottle with bromobutyl stopper and aluminium cap

12 x 500 ml polypropylene bottle with bromobutyl stopper and aluminium cap

12 x 750 ml polypropylene bottle with bromobutyl stopper and aluminium cap

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

21/12/2022

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Authorisation number:

63340

Date of authorisation status change:

21/12/2022

Reference member state:

Austria

Procedure number:

AT/V/0028/001

Concerned member states:

Denmark Finland Germany Ireland Netherlands Norway Portugal Romania
Sweden

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 27/07/2022

Updated on: 13/03/2026

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Package Leaflet

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Package Leaflet and Labelling

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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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