File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000106687

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

Authorised

- 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:

Swedish Medical Products Agency

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
Download

Package Leaflet

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

Package Leaflet and Labelling

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

at-puar-atv0028001-mr-gluecoebel-en.pdf