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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:
Denmark

Available in:
Denmark

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:
25/10/2022

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:

Danish Medicines Agency

Authorisation number:
67701

Date of authorisation status change:
25/10/2022

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

Combined File of all Documents

Summary of Product Characteristics

English (PDF)
Published on: 27/07/2022
Updated on: 13/03/2026
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