Injectio Glucosi Isotonica et Natrii Chlorati Isotonica 1:1 WET Baxter (25 g + 4,5 g)/1000 ml Roztwór do infuzji

Authorised

- Glucose
- Sodium chloride

Product identification

Medicine name:
Injectio Glucosi Isotonica et Natrii Chlorati Isotonica 1:1 WET Baxter (25 g + 4,5 g)/1000 ml Roztwór do infuzji

Active substance:
Glucose
Sodium chloride

Target species:
Goat
Horse
Cat
Pig
Cattle
Dog
Sheep

Route of administration:
Intraperitoneal use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Glucose
25.00 gram(s) / 1000.00 millilitre(s)

Sodium chloride
4.50 gram(s) / 1000.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intraperitoneal use:

- **Goat**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day
- **Sheep**

- Meat and offal. 0 day

Subcutaneous use:

- **Goat**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day

Intravenous use:

- **Goat**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**

- Meat and offal. 0 day

- **Sheep**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB05BB02

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish

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:
Baxter Polska Sp. z o.o.

Marketing authorisation date:
12/08/1996

Manufacturing sites for batch release:

Bieffe Medital S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0275

Date of authorisation status change:
12/08/1996

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

Documents

Summary of Product Characteristics

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Package Leaflet

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.