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NaCl Bernburg, 0,9%, Solution for injection/infusion

Authorised

- Sodium chloride

Product identification

Medicine name:
NaCl Bernburg, 0,9%, Solution for injection/infusion

Active substance:
Sodium chloride

Target species:
Cattle
Horse
Sheep
Goat
Pig
Dog
Cat
Rabbit
Guinea pig

Route of administration:
Intravenous use
Subcutaneous use
Intraperitoneal use

Product details

Active substance and strength:
Sodium chloride
9.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection/infusion

Withdrawal period by route of administration:
Intravenous use:

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

Rabbit

- Meat and offal. 0 day

Subcutaneous use:

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

Intraperitoneal use:

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

- Milk. 0 hour

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

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

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

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB05BB01

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:

Germany

Package description:
Plastic (PE) Bottle 10 x 1000.0 millilitre(s)
Plastic (PE) Bottle 1 x 500.0 millilitre(s)
Plastic (PE) Bottle 10 x 500.0 millilitre(s)
Plastic (PE) Bottle 20 x 500.0 millilitre(s)
Plastic (PP) Sachet 1 x 500.0 millilitre(s)
Plastic (PP) Sachet 10 x 500.0 millilitre(s)
Plastic (PP) Sachet 20 x 500.0 millilitre(s)
Sachet 1 x 5000.0 millilitre(s)
Sachet 2 x 5000.0 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Serumwerk Bernburg AG

Marketing authorisation date:
14/06/2022

Manufacturing sites for batch release:
Serumwerk Bernburg AG

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7005382.00.00

Date of authorisation status change:
14/06/2022

Reference member state:
Czechia

Procedure number:
CZ/V/0175/001

Concerned member states:
Germany

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

Labelling

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