File downloaded on 2026-11-24
Source URL: https://medicines.health.europa.eu/veterinary/en/600000092649

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:
Czechia

Available in:

Czechia

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:
28/06/2022

Manufacturing sites for batch release:
Serumwerk Bernburg AG

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/024/22-C

Date of authorisation status change:
28/06/2022

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

Summary of Product Characteristics

English (PDF)
Published on: 11/05/2026
Download

Package Leaflet

English (PDF)
Published on: 11/05/2026
Download

Labelling

English (PDF)

Published on: 11/05/2026

Download

eu-puar-czv0175001-mr-nacl_bernburg-en.pdf