

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 (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)
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)

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: 19/01/2026

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Package Leaflet

English (PDF)

Published on: 19/01/2026

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Labelling

English (PDF)

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