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SODIUM CHLORIDE OSALIA 0.9% SOLUTION FOR INJECTION OR INFUSION FOR CATTLE, HORSES,PIGS, SHEEP, GOATS, DOGS AND CATS

Authorised

- Sodium chloride

Product identification

Medicine name:
SODIUM CHLORIDE OSALIA 0.9% SOLUTION FOR INJECTION OR INFUSION FOR CATTLE, HORSES,PIGS, SHEEP, GOATS, DOGS AND CATS

Active substance:
Sodium chloride

Target species:
Cattle
Pig
Cat
Equid
Horse
Sheep
Goat
Dog

Route of administration:
Intravenous use
Subcutaneous 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**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day

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

Subcutaneous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Goat**
  - Milk. 0 day
  - 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:
Estonia

Package description:
Box of 1 vial of 100 ml
Box of 2 bags of 5000 ml
Box of 1 bag of 5000 ml
Box of 5 bags of 2000 ml
Box of 1 bag of 2000 ml
Box of 10 bags of 1000 ml
Box of 1 bag of 1000 ml
Box of 12 vials of 500 ml
Box of 1 vial of 500 ml
Box of 20 vials of 250 ml
Box of 1 vial of 250 ml
Box of 30 vials of 100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Osalia

Marketing authorisation date:
28/09/2026

Manufacturing sites for batch release:
S.A.L.F. S.p.A. Laboratorio Farmacologico

Responsible authority:
State Agency Of Medicines

Authorisation number:
1273926

Date of authorisation status change:
28/09/2026

Reference member state:
France

Procedure number:
FR/V/0443/001

Concerned member states:
Estonia Latvia Lithuania Romania

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

Documents

Summary of Product Characteristics

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

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.