CHLORURE DE SODIUM OSALIA 0,9 % SOLUTION INJECTABLE OU POUR PERFUSION POUR BOVINS, CHEVAUX, PORCS, OVINS, CAPRINS, CHIENS ET CHATS

Authorised

- Sodium chloride

Product identification

Medicine name:
CHLORURE DE SODIUM OSALIA 0,9 % SOLUTION INJECTABLE OU POUR PERFUSION POUR BOVINS, CHEVAUX, PORCS, OVINS, CAPRINS, CHIENS ET CHATS

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**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

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

Subcutaneous use:

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

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
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Available only in French
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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:
17/01/2019

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/4058207 8/2019

Date of authorisation status change:
17/01/2019

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 and Labelling

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

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