

# SOLUZIONE ELETTROLITICA REIDRATANTE III

Authorised

- Sodium chloride
- Potassium chloride
- Magnesium chloride hexahydrate
- Calcium chloride dihydrate
- Sodium acetate
- Sodium citrate dihydrate

## Product identification

**Medicine name:**

SOLUZIONE ELETTROLITICA REIDRATANTE III

---

**Active substance:**

Sodium chloride

Potassium chloride

Magnesium chloride hexahydrate

Calcium chloride dihydrate

Sodium acetate

Sodium citrate dihydrate

---

**Target species:**

Cattle

Dog

Sheep

Horse (non food-producing)

Cat  
Pig  
Horse

---

**Route of administration:**

Intravenous use

---

## Product details

**Active substance and strength:**

Sodium chloride

5.00 gram(s) / 1000.00 millilitre(s)

Potassium chloride

0.75 gram(s) / 1000.00 millilitre(s)

Magnesium chloride hexahydrate

0.31 gram(s) / 1000.00 millilitre(s)

Calcium chloride dihydrate

0.35 gram(s) / 1000.00 millilitre(s)

Sodium acetate

6.40 gram(s) / 1000.00 millilitre(s)

Sodium citrate dihydrate

0.75 gram(s) / 1000.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for infusion

---

**Withdrawal period by route of administration:**

**Intravenous use:**

•

**Cattle**

- Milk. 0 day

- Meat and offal. 0 day

•

**Sheep**

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

- 

**Pig**

- Meat and offal. 0 day

- 

**Horse**

- Meat and offal. 0 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QB05BB01

---

**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Italy

---

**Package description:**

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Industria Farmaceutica Galenica Senese S.r.l.

---

**Marketing authorisation date:**

29/09/1999

---

**Manufacturing sites for batch release:**

Industria Farmaceutica Galenica Senese S.r.l.

---

**Responsible authority:**

Ministry Of Health

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

29/09/1999

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.