

# ELETTROLITICA REIDRATANTE III S.A.L.F. Soluzione isotonica per infusione endovenosa per bovini, equini, cani e gatti

Authorised

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

## Product identification

### **Medicine name:**

ELETTROLITICA REIDRATANTE III S.A.L.F. Soluzione isotonica per infusione endovenosa per bovini, equini, cani e gatti

### **Active substance:**

Sodium acetate

Calcium chloride dihydrate

Magnesium chloride hexahydrate

Potassium chloride

Sodium chloride

Sodium citrate dihydrate

**Target species:**

Cattle

Dog

Cat

Horse

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Sodium acetate

6.40 gram(s) / 1000.00 millilitre(s)

Calcium chloride dihydrate

0.35 gram(s) / 1000.00 millilitre(s)

Magnesium chloride hexahydrate

0.31 gram(s) / 1000.00 millilitre(s)

Potassium chloride

0.75 gram(s) / 1000.00 millilitre(s)

Sodium chloride

5.00 gram(s) / 1000.00 millilitre(s)

Sodium citrate dihydrate

0.75 gram(s) / 1000.00 millilitre(s)

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**Pharmaceutical form:**

Solution for infusion

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**Withdrawal period by route of administration:****Intravenous use:**

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**Cattle**

- Milk. 0 day

- Meat and offal. 0 day

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## Horse

- Meat and offal. 0 day

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

QB05BB01

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### Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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### Authorisation status:

Valid

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### Authorised in:

Italy

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### Available in:

Italy

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### 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](#)

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## Additional information

### Entitlement type:

Marketing Authorisation

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### Legal basis of product authorisation:

Bibliographic application (Article 22 of Regulation (EU) 2019/6)

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### Marketing authorisation holder:

S.A.L.F. S.p.A. Laboratorio Farmacologico

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**Marketing authorisation date:**

15/11/2005

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**Manufacturing sites for batch release:**

S.A.L.F. S.p.A. Laboratorio Farmacologico

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**Responsible authority:**

MdS

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

15/11/2010

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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

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