

# RINGER LATTATO S.A.L.F.

## soluzione per infusione endovenosa per bovini, equini, cani e gatti

Authorised

- Sodium chloride
- Potassium chloride
- Calcium chloride dihydrate
- SODIUM LACTATE 60%

## Product identification

### **Medicine name:**

RINGER LATTATO S.A.L.F. soluzione per infusione endovenosa per bovini, equini, cani e gatti

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### **Active substance:**

Sodium chloride

Potassium chloride

Calcium chloride dihydrate

SODIUM LACTATE 60%

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### **Target species:**

Cattle

Dog

Cat

Horse

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**Route of administration:**

Intraperitoneal use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Sodium chloride

6.00 gram(s) / 1000.00 millilitre(s)

Potassium chloride

0.40 gram(s) / 1000.00 millilitre(s)

Calcium chloride dihydrate

0.27 gram(s) / 1000.00 millilitre(s)

SODIUM LACTATE 60%

5.46 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:**

**Intraperitoneal use:**

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**Cattle**

- Milk. 0 day

- Meat and offal. 0 day

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**Horse**

- Meat and offal. 0 day

**Intravenous use:**

- 

**Cattle**

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

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**Horse**

- Meat and offal. 0 day

**Subcutaneous 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:**

Ministry Of Health

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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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