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

Documents

Combined File of all Documents

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