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ELETTROLITICA REIDRATANTE III ACME, soluzione perfusionale per cavalli, bovini, cani e gatti

Authorised

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

Product identification

Medicine name:
ELETTROLITICA REIDRATANTE III ACME, soluzione perfusionale per cavalli, bovini, cani e gatti

Active substance:
Sodium chloride

Potassium chloride

Magnesium chloride hexahydrate

Calcium chloride dihydrate

Sodium acetate trihydrate

Sodium citrate dihydrate

Target species:
Cattle

Dog
Horse
Cat

Route of administration:
Intraperitoneal use
Transdermal use
Intravenous use

Product details

Active substance and strength:
Sodium chloride
0.50 gram(s) / 100.00 millilitre(s)

Potassium chloride
0.08 gram(s) / 100.00 millilitre(s)

Magnesium chloride hexahydrate
0.03 gram(s) / 100.00 millilitre(s)

Calcium chloride dihydrate
0.04 gram(s) / 100.00 millilitre(s)

Sodium acetate trihydrate
0.64 gram(s) / 100.00 millilitre(s)

Sodium citrate dihydrate
0.08 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intraperitoneal use:

- **Cattle**
  - Meat and offal, milk. 0 day
- **Horse**

- Meat and offal, milk. 0 day

Transdermal use:

- **Cattle**
  - Meat and offal, milk. 0 day
- **Horse**
  - Meat and offal, milk. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal, milk. 0 day
- **Horse**
  - Meat and offal, milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB05BB01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian

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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Acme S.r.l.

Marketing authorisation date:
29/11/1995

Manufacturing sites for batch release:
Fresenius Kabi Italia S.r.l. In Breve Fki S.r.l.

Responsible authority:
Ministry Of Health

Authorisation number:
100330

Date of authorisation status change:
10/02/2017

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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