RINGER LATTATO ACME, soluzione perfusionale per cavalli, bovini, cani e gatti

Authorised

- Sodium chloride
- Potassium chloride
- Sodium hydroxide
- Calcium chloride dihydrate
- Lactic acid

Product identification

Medicine name:
RINGER LATTATO ACME, soluzione perfusionale per cavalli, bovini, cani e gatti

Active substance:
Sodium chloride
Potassium chloride
Sodium hydroxide
Calcium chloride dihydrate
Lactic acid

Target species:
Cattle
Dog
Horse
Cat

Route of administration:

Intraperitoneal use
Transdermal use
Intravenous use

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)

Sodium hydroxide
1.17 gram(s) / 1000.00 millilitre(s)

Calcium chloride dihydrate
0.27 gram(s) / 1000.00 millilitre(s)

Lactic acid
2.60 gram(s) / 1000.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intraperitoneal use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cat**

Transdermal use:

- **Cattle**
  - Meat and offal. 0 day

- Milk. 0 hour

- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cat**

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cat**

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:
FKI S.r.l.

Responsible authority:
Ministry Of Health

Authorisation number:
100351

Date of authorisation status change:
29/11/1995

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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