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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

Route of administration:
Intravenous use

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)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Milk. 0 day
  - 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

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

Marketing authorisation holder:
S.A.L.F. S.p.A. Laboratorio Farmacologico

Marketing authorisation date:
15/11/2005

Manufacturing sites for batch release:
S.A.L.F. S.p.A. Laboratorio Farmacologico

Responsible authority:
MdS

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/11/2010

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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