Lactato-RingerVet Solution for infusion for cattle, horses, sheep, goats, pigs, dogs and cats

Authorised

- Sodium lactate
- Calcium chloride dihydrate
- Sodium chloride
- Potassium chloride

Product identification

Medicine name:
Lactato-RingerVet Solution for infusion for cattle, horses, sheep, goats, pigs, dogs and cats

Active substance:
Sodium lactate
Calcium chloride dihydrate
Sodium chloride
Potassium chloride

Target species:
Cattle
Sheep
Goat
Horse
Pig

Dog
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Sodium lactate
0.31 gram(s) / 100.00 millilitre(s)

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

Sodium chloride
0.60 gram(s) / 100.00 millilitre(s)

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

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**

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

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB05BB01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Romania

Package description:
box containing 2 bags of 5000 ml
box containing 10 bottles of 500 ml
box containing 10 bottles of 1000 ml
box containing 20 bottles of 250 ml

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:
B Braun Vetcare S.A.

Marketing authorisation date:
11/05/2026

Manufacturing sites for batch release:
B BRAUN MEDICAL S.A.
B. Braun Melsungen AG

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
260082

Date of authorisation status change:
11/05/2026

Reference member state:
Spain

Procedure number:
ES/V/0153/001

Concerned member states:
Austria Belgium Denmark France Germany Ireland Italy Netherlands Poland Portugal Romania Sweden United Kingdom (Northern Ireland)

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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