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B. Braun Vet Care Hartmann’s Lactated Ringers Solution for infusion for cattle, horse, sheep, goat, pig, dog and cat

Authorised

- Potassium chloride
- Sodium chloride
- Calcium chloride dihydrate
- Sodium (S)-lactate solution

Product identification

Medicine name:
B. Braun Vet Care Hartmann’s Lactated Ringers Solution for infusion for cattle, horse, sheep, goat, pig, dog and cat

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Active substance:
Potassium chloride

Sodium chloride

Calcium chloride dihydrate

Sodium (S)-lactate solution

Target species:
Cattle
Sheep
Goat

Horse
Pig
Dog
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Potassium chloride
0.40 milligram(s) / 1.00 millilitre(s)

Sodium chloride
6.00 milligram(s) / 1.00 millilitre(s)

Calcium chloride dihydrate
0.27 milligram(s) / 1.00 millilitre(s)

Sodium (S)-lactate solution
6.24 milligram(s) / 1.00 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

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Authorisation status:
Valid

Authorised in:
Austria

Available in:
Austria

Package description:
Available only in German
Available only in German
Available only in German
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 Melsungen AG

Marketing authorisation date:
19/05/2010

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00872

Date of authorisation status change:
19/05/2010

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

Package Leaflet

English (PDF)
Published on: 22/10/2025
Updated on: 13/03/2026
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Summary of Product Characteristics

English (PDF)
Published on: 22/10/2025
Updated on: 13/03/2026
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Labelling

English (PDF)
Published on: 22/10/2025
Updated on: 13/03/2026
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Combined File of all Documents

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