Calciject LV Solution for Injection

Not authorised

- Magnesium chloride hexahydrate
- Calcium hydroxide
- Boric acid
- Calcium gluconate

Product identification

Medicine name:
Calciject LV Solution for Injection

Active substance:
Magnesium chloride hexahydrate

Calcium hydroxide

Boric acid

Calcium gluconate

Target species:
Cattle

Route of administration:
Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Magnesium chloride hexahydrate
65.00 milligram(s) / 1.00 millilitre(s)

Calcium hydroxide
13.20 milligram(s) / 1.00 millilitre(s)

Boric acid
73.40 milligram(s) / 1.00 millilitre(s)

Calcium gluconate
388.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:
Ireland

Package description:
Calciject LV is packaged in 100 ml clear grade II glass vial or in 100 ml amber grade II glass vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
22/03/1996

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/044/001

Date of authorisation status change:
31/01/2023

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics