LODEVIL

Authorised

- Glucose
- Magnesium chloride
- Potassium chloride
- Sodium chloride
- Sodium hydrogen carbonate
- Sodium acetate

Product identification

Medicine name:
LODEVIL

Active substance:
Glucose

Magnesium chloride

Potassium chloride

Sodium chloride

Sodium hydrogen carbonate

Sodium acetate

Target species:
Cattle (calf)

Route of administration:
Intravenous use

Product details

Active substance and strength:

Glucose
10.80 milligram(s) / 1.00 millilitre(s)

Magnesium chloride
0.20 milligram(s) / 1.00 millilitre(s)

Potassium chloride
0.45 milligram(s) / 1.00 millilitre(s)

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

Sodium hydrogen carbonate
4.20 milligram(s) / 1.00 millilitre(s)

Sodium acetate
4.08 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

- **Cattle (calf)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Package description:
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
22/01/1982

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3830006 4/1982

Date of authorisation status change:
22/01/2012

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.
