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Lactolyte Pulver zur Herstellung einer Lösung zum Einnehmen

Not authorised

- Sodium chloride
- SODIUM ACETATE ANHYDROUS
- Sodium propionate
- WHEY
- Potassium chloride
- MAGNESIUM CHLORIDE ANHYDROUS
- Potassium dihydrogen phosphate

Product identification

Medicine name:
Lactolyte Pulver zur Herstellung einer Lösung zum Einnehmen
Lactolyte Poudre pour solution buvable
Lactolyte Poeder voor drank

Active substance:
Sodium chloride
SODIUM ACETATE ANHYDROUS
Sodium propionate
WHEY
Potassium chloride
MAGNESIUM CHLORIDE ANHYDROUS
Potassium dihydrogen phosphate

Target species:
Cattle (calf)

Route of administration:
Oral use

Product details

Active substance and strength:
Sodium chloride
2.92 gram(s) / 90.00 gram(s)

SODIUM ACETATE ANHYDROUS
4.90 gram(s) / 90.00 gram(s)

Sodium propionate
1.92 gram(s) / 90.00 gram(s)

WHEY
76.88 gram(s) / 90.00 gram(s)

Potassium chloride
0.74 gram(s) / 90.00 gram(s)

MAGNESIUM CHLORIDE ANHYDROUS
0.38 gram(s) / 90.00 gram(s)

Potassium dihydrogen phosphate
1.36 gram(s) / 90.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:
Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07CQ01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:
Belgium

Package description:
Lactolyte Powder for oral solution Barrel of 4.5 kg
Lactolyte Powder for oral solution Jar of 900 g
Lactolyte Powder for oral solution Box of 40 sachets of 90g
Lactolyte Powder for oral solution Box of 8 sachets of 90g

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
22/05/1995

Manufacturing sites for batch release:
Virbac

Responsible authority:
Federal Agency For Medicines And Health Products

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

Date of authorisation status change:

30/08/2023

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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