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Biopect, poeder voor gebruik in drinkwater/melk voor kalveren, lammeren en biggen

Authorised

- Sodium hydrogen carbonate
- Ascorbic acid
- Potassium dihydrogen phosphate
- Sodium chloride
- Citric acid
- Glucose monohydrate
- Magnesium chloride
- Potassium chloride
- Magnesium oxide

Product identification

Medicine name:
Biopect, poeder voor gebruik in drinkwater/melk voor kalveren, lammeren en biggen

Active substance:
Sodium hydrogen carbonate

Ascorbic acid

Potassium dihydrogen phosphate

Sodium chloride

Citric acid

Glucose monohydrate

Magnesium chloride

Potassium chloride

Magnesium oxide

Target species:
Cattle (calf)
Sheep (lamb)
Pig (piglet)

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Sodium hydrogen carbonate
36.80 milligram(s) / 1.00 gram(s)

Ascorbic acid
4.40 milligram(s) / 1.00 gram(s)

Potassium dihydrogen phosphate
5.30 milligram(s) / 1.00 gram(s)

Sodium chloride
38.50 milligram(s) / 1.00 gram(s)

Citric acid
13.10 milligram(s) / 1.00 gram(s)

Glucose monohydrate
301.00 milligram(s) / 1.00 gram(s)

Magnesium chloride
0.90 milligram(s) / 1.00 gram(s)

Potassium chloride
12.30 milligram(s) / 1.00 gram(s)

Magnesium oxide
1.80 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water/milk

Withdrawal period by route of administration:
In drinking water/milk use:

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

- **Sheep (lamb)**
  - Meat and offal. 0 day

- **Pig (piglet)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07CQ02

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vee-Service Drunen B.V.

Marketing authorisation date:
21/11/1991

Manufacturing sites for batch release:
Biofiber-Damino A/S

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 5010

Date of authorisation status change:
22/12/2015

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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