

Electrovit poeder voor toediening via drinkwater voor kalveren en biggen

Not
authorised

- Nicotinamide
- Thiamine hydrochloride
- Pyridoxine hydrochloride
- Calcium pantothenate
- Potassium chloride
- Sodium chloride
- Calcium lactate
- Lysine hydrochloride
- Albumin tannate
- Ascorbic acid
- Sodium hydrogen carbonate
- Methionine
- Colecalciferol
- RETINOL ACETATE

Product identification

Medicine name:

Electrovit poeder voor toediening via drinkwater voor kalveren en biggen

Active substance:

Nicotinamide

Thiamine hydrochloride

Pyridoxine hydrochloride
Calcium pantothenate
Potassium chloride
Sodium chloride
Calcium lactate
Lysine hydrochloride
Albumin tannate
Ascorbic acid
Sodium hydrogen carbonate
Methionine
Colecalciferol
RETINOL ACETATE

Target species:

Cattle (calf)
Pig (piglet)

Route of administration:

In drinking water use

Product details

Active substance and strength:

Nicotinamide
4.00 milligram(s) / 1.00 gram(s)
Thiamine hydrochloride
1.80 milligram(s) / 1.00 gram(s)
Pyridoxine hydrochloride
0.80 milligram(s) / 1.00 gram(s)
Calcium pantothenate
0.80 milligram(s) / 1.00 gram(s)
Potassium chloride
0.04 gram(s) / 1.00 gram(s)
Sodium chloride

0.10 gram(s) / 1.00 gram(s)

Calcium lactate

0.01 gram(s) / 1.00 gram(s)

Lysine hydrochloride

0.01 gram(s) / 1.00 gram(s)

Albumin tannate

0.01 gram(s) / 1.00 gram(s)

Ascorbic acid

0.02 gram(s) / 1.00 gram(s)

Sodium hydrogen carbonate

0.10 gram(s) / 1.00 gram(s)

Methionine

0.01 gram(s) / 1.00 gram(s)

Colecalciferol

250.00 international unit(s) / 1.00 gram(s)

RETINOL ACETATE

500.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

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Cattle (calf)

- Meat and offal. no withdrawal period Zero days

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Pig (piglet)

- Meat and offal. no withdrawal period Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11AB

Legal status of supply:

Medicinal product not subject to medical prescription

Authorisation status:

Surrendered

Authorised in:

Netherlands

Package description:

Available only in [Dutch](#)

Available only in [Dutch](#)

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:

Feramed B.V.

Marketing authorisation date:

21/06/1993

Manufacturing sites for batch release:

Feramed B.V.

Responsible authority:

MEB

Authorisation number:

REG NL 7652

Date of authorisation status change:

17/01/2011

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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