Multivit inj. oplossing voor injectie

Authorised

- Retinol palmitate
- Colecalciferol
- Cyanocobalamin
- Dexpanthenol
- DL-ALPHA TOCOPHEROL ACETATE
- RIBOFLAVIN SODIUM PHOSPHATE
- Thiamine hydrochloride
- Nicotinamide
- Ascorbic acid
- Pyridoxine hydrochloride

Product identification

Medicine name:
Multivit inj. oplossing voor injectie

Active substance:
Retinol palmitate

Colecalciferol

Cyanocobalamin

Dexpanthenol

DL-ALPHA TOCOPHEROL ACETATE

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

Nicotinamide

Ascorbic acid

Pyridoxine hydrochloride

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

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Retinol palmitate
50000.00 international unit(s) / 1.00 millilitre(s)

Colecalciferol
25000.00 international unit(s) / 1.00 millilitre(s)

Cyanocobalamin
30.00 microgram(s) / 1.00 millilitre(s)

Dexpanthenol
3.00 milligram(s) / 1.00 millilitre(s)

DL-ALPHA TOCOPHEROL ACETATE
4.00 milligram(s) / 1.00 millilitre(s)

RIBOFLAVIN SODIUM PHOSPHATE
2.00 milligram(s) / 1.00 millilitre(s)

Thiamine hydrochloride
2.50 milligram(s) / 1.00 millilitre(s)

Nicotinamide
12.50 milligram(s) / 1.00 millilitre(s)

Ascorbic acid
2.00 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride
1.25 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. no withdrawal period nul dagen
  - Meat and offal. no withdrawal period nul dagen
- **Cattle (calf)**
  - Meat and offal. no withdrawal period nul dagen
- **Sheep (lamb)**
  - Meat and offal. no withdrawal period nul dagen
- **Horse**
  - Meat and offal. no withdrawal period nul dagen
- **Pig (piglet)**
  - Meat and offal. no withdrawal period nul dagen
- **Horse (foal)**
  - Meat and offal. no withdrawal period nul dagen

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11BA

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:
Feramed B.V.

Marketing authorisation date:
23/07/1993

Manufacturing sites for batch release:
Feramed B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 8010

Date of authorisation status change:
11/04/2011

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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