Vitol-Ject Forte, oplossing voor injectie

Not authorised

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

Product identification

Medicine name:
Vitol-Ject Forte, oplossing voor injectie

Active substance:
Ascorbic acid

Nicotinamide

Pyridoxine hydrochloride

DL-ALPHA TOCOPHEROL ACETATE

Dexpanthenol

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

Cyanocobalamin

Colecalciferol

Retinyl propionate

Target species:
Cattle
Sheep
Cattle (calf)
Horse
Pig
Dog
Cat

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Ascorbic acid
2.00 milligram(s) / 1.00 millilitre(s)

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

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

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

Dexpanthenol
3.00 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)

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

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

Retinyl propionate
50000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. no withdrawal period 0 days
  - Meat and offal. no withdrawal period 0 days
- **Sheep**
  - Milk. no withdrawal period 0 days
  - Meat and offal. no withdrawal period 0 days
- **Cattle (calf)**
  - Meat and offal. no withdrawal period 0 days
- **Horse**
  - Meat and offal. no withdrawal period 0 days
- **Pig**
  - Meat and offal. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11BA

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
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:
Dopharma Research B.V.

Marketing authorisation date:
16/01/1992

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 4142

Date of authorisation status change:
6/12/2022

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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