Multivitamin Injection

Not authorised

- Dexpanthenol
- Nicotinamide
- Retinol palmitate
- Pyridoxine
- Colecalciferol
- RIBOFLAVIN SODIUM PHOSPHATE
- Thiamine hydrochloride
- TOCOPHERYL ACETATE
- VITAMIN B12

Product identification

Medicine name:
Multivitamin Injection

Active substance:
Dexpanthenol

Nicotinamide

Retinol palmitate

Pyridoxine

Colecalciferol

RIBOFLAVIN SODIUM PHOSPHATE

Thiamine hydrochloride

TOCOPHERYL ACETATE

VITAMIN B12

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Dexpanthenol
25.00 milligram(s) / 1.00 millilitre(s)

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

Retinol palmitate
15000.00 international unit(s) / 1.00 millilitre(s)

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

Colecalciferol
25.00 microgram(s) / 1.00 millilitre(s)

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

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

TOCOPHERYL ACETATE
21.95 milligram(s) / 1.00 millilitre(s)

VITAMIN B12
25.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 28 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

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

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Authorisation status:

Surrendered

Authorised in:
Ireland

Package description:
100 ml type II glass (amber) containers, sealed with nitryl rubber bungs and aluminium caps.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Chem-Pharm

Marketing authorisation date:
1/10/2001

Manufacturing sites for batch release:
Norbrook Laboratories Limited

Responsible authority:
HPRA

Authorisation number:
VPA10823/017/001

Date of authorisation status change:
22/12/2021

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

Documents

Summary of Product Characteristics