Vitamine A + D3 + E Drank

Authorised

- Cholecalciferol concentrate (powder form)
- Retinol
- ALPHATOCOPHEROL ACETATE

Product identification

Medicine name:
Vitamine A + D3 + E Drank
Vitamine A + D3 + E Solution buvable
Vitamine A + D3 + E Lösung zum Einnehmen

Active substance:
Cholecalciferol concentrate (powder form)
Retinol
ALPHATOCOPHEROL ACETATE

Target species:
Pig
Cattle
Horse
Sheep
Goat
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:

Cholecalciferol concentrate (powder form)
25000.00 international unit(s) / 1.00 millilitre(s)

Retinol
50000.00 international unit(s) / 1.00 millilitre(s)

ALPHATOCOPHEROL ACETATE
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:

Oral use:

- **Pig**
  - Meat and offal. 28 day
- **Cattle**
  - Meat and offal. 28 day
  - Milk. no withdrawal period 0 days
- **Horse**
  - Meat and offal. 28 day
  - Milk. no withdrawal period 0 days
- **Sheep**
  - Meat and offal. 28 day
  - Milk. no withdrawal period 0 days

- **Goat**
  - Meat and offal. 28 day
  - Milk. no withdrawal period 0 days
- **Chicken**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11BA

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Vitamine A + D3 + E 1 Bottle with 1 l of Oral solution
Vitamine A + D3 + E 1 Bottle with 500 ml of Oral solution
Vitamine A + D3 + E 1 Bottle with 250 ml of Oral solution
Vitamine A + D3 + E 1 Bottle with 100 ml of Oral solution

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
15/04/1976

Manufacturing sites for batch release:
V.M.D.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V102365

Date of authorisation status change:
3/09/2015

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

Documents

Package Leaflet

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

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