Multivitamin injection (15 000 j.m. + 1000 j.m. + 20 mg + 10 mg + 5 mg + 3 mg + 35 mg + 25 mg + 0,05 mg) /ml Roztwór do wstrzykiwań

Not authorised

- Retinol
- Colecalciferol
- ALPHA-TOCOPHEROL
- Thiamine
- Pyridoxine hydrochloride
- Riboflavin
- Dexpanthenol
- Nicotinamide
- Cyanocobalamin

Product identification

Medicine name:
Multivitamin injection (15 000 j.m. + 1000 j.m. + 20 mg + 10 mg + 5 mg + 3 mg + 35 mg + 25 mg + 0,05 mg) /ml Roztwór do wstrzykiwań

Active substance:
Retinol

Colecalciferol

ALPHA-TOCOPHEROL

Thiamine

Pyridoxine hydrochloride

Riboflavin

Dexpanthenol

Nicotinamide

Cyanocobalamin

Target species:

Cattle
Sheep
Goat
Horse
Cat
Dog
Pig

Route of administration:

Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Retinol

Colecalciferol

ALPHA-TOCOPHEROL

Thiamine

Pyridoxine hydrochloride

Riboflavin

Dexpanthenol

Nicotinamide

Cyanocobalamin
50.00 microgram(s) / 1.00 microgram(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Subcutaneous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
-

Sheep

- Meat and offal. 0 day

•

Goat

- Meat and offal. 0 day
- Milk. 0 day

•

Horse

- Meat and offal. 0 day
- Milk. 0 day

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11EX

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

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Authorisation status:
Surrendered

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Scanvet Poland Sp. z o.o.

Marketing authorisation date:
28/05/1998

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0735

Date of authorisation status change:
21/07/2025

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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