Combivit (35 mg + 0,5 mg + 7 mg + 23 mg + 70 mg)/ml Roztwór do wstrzykiwań

Not authorised

- Ascorbic acid
- Thiamine
- Pyridoxine hydrochloride
- Riboflavin
- Nicotinamide

Product identification

Medicine name:
Combivit (35 mg + 0,5 mg + 7 mg + 23 mg + 70 mg)/ml Roztwór do wstrzykiwań

Active substance:
Ascorbic acid
Thiamine
Pyridoxine hydrochloride
Riboflavin
Nicotinamide

Target species:
Cat
Dog
Horse
Goat

Pig
Cattle
Sheep

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Ascorbic acid

Thiamine

Pyridoxine hydrochloride

Riboflavin

Nicotinamide

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

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

- Meat and offal. 0 day

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

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

Subcutaneous use:

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

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

- **Pig**
  - Meat and offal. 0 day

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

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

Intravenous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11BA

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:
0736

Date of authorisation status change:
10/07/2025

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

Documents

Labelling

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.

Summary of Product Characteristics

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