

Vitamin AD3E

Not authorised

- Retinol palmitate
- TOCOPHERYL ACETATE
- Colecalciferol

Product identification

Medicine name:

Vitamin AD3E

Active substance:

Retinol palmitate

TOCOPHERYL ACETATE

Colecalciferol

Target species:

Cattle (calf)

Cattle

Horse

Pig

Pig (weaned piglet)

Pig (suckling piglet)

Dog

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Retinol palmitate

176.47 milligram(s) / 1.00 millilitre(s)

TOCOPHERYL ACETATE

50.00 milligram(s) / 1.00 millilitre(s)

Colecalciferol

2.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle (calf)

- Meat and offal. 259 day

-

Cattle

- Meat and offal. 259 day

- Milk. 5 day

-

Horse

- Meat and offal. 259 day

- Milk. 0 day

-

Pig

- Meat and offal. 194 day

-

Pig (weaned piglet)

- Meat and offal. 194 day

-

Pig (suckling piglet)

- Meat and offal. 194 day

Intramuscular use:

-

Cattle (calf)

- Meat and offal. 20 day

-

Cattle

- Meat and offal. 50 day

- Milk. 0 day

-

Horse

- Meat and offal. 50 day

- Milk. 0 day

-

Pig

- Meat and offal. 20 day

-

Pig (suckling piglet)

- Meat and offal. 10 day

-

Pig (weaned piglet)

- Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11JA

Legal status of supply:

This information is not available for this product.

Authorisation status:

Surrendered

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

5/10/2004

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

150306

Date of authorisation status change:

23/04/2025

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.