Belavit AD3E, Solution for Injection for Horses, Cattle, Pigs, and Dogs

Not authorised

- DL-ALPHA TOCOPHEROL ACETATE
- COLECALCIFEROL CONCENTRATE (OILY FORM)
- Retinol palmitate

Product identification

Medicine name:
Vitamin AD3E pro injectione, solution for injection for horses, cattle, pigs, and dogs
Belavit AD3E, Solution for Injection for Horses, Cattle, Pigs, and Dogs

Active substance:
DL-ALPHA TOCOPHEROL ACETATE
COLECALCIFEROL CONCENTRATE (OILY FORM)
Retinol palmitate

Target species:
Cattle
Dog
Horse
Pig

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

DL-ALPHA TOCOPHEROL ACETATE
50.00 milligram(s) / 1.00 millilitre(s)

COLECALCIFEROL CONCENTRATE (OILY FORM)
100.00 milligram(s) / 1.00 millilitre(s)

Retinol palmitate
176.46 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Milk. 120 hour
  - Meat and offal. 259 day
- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Intramuscular use:

- **Cattle**
  - Milk. 120 hour

- Meat and offal. 259 day

- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

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

---

Authorisation status:
Expired

Authorised in:
United Kingdom (Northern Ireland)

Package description:
(ID3): 1 unspecified outer container with 12 Bottle (Glass) with 100 millilitre(s) (1200 millilitre(s))
(ID2): 1 unspecified outer container with 6 Bottle (Glass) with 100 millilitre(s) (600 millilitre(s))
(ID1): 1 unspecified outer container with 1 Bottle (Glass) with 100 millilitre(s) (100 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
12/07/2019

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 41816/4004

Date of authorisation status change:
15/05/2023

Reference member state:
Germany

Procedure number:
DE/V/0313/001

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

Documents

Combined File of all Documents