

ADE – vit injekčný roztok

Authorised

- RETINOL ACETATE
- Ergocalciferol
- alfa-Tocopheryl acetate

Product identification

Medicine name:

ADE – vit injekčný roztok

Active substance:

RETINOL ACETATE

Ergocalciferol

alfa-Tocopheryl acetate

Target species:

Cattle

Horse

Pig

Sheep

Goat

Rabbit

Dog

Route of administration:

Intramuscular use

Product details

Active substance and strength:

RETINOL ACETATE

100000.00 international unit(s) / 1.00 millilitre(s)

Ergocalciferol

100000.00 international unit(s) / 1.00 millilitre(s)

alfa-Tocopheryl acetate

30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11CB

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

31/01/2003

Manufacturing sites for batch release:

Bioveta, a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/009/03-S

Date of authorisation status change:

31/01/2003

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

Documents

Combined File of all Documents

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