Covexin 10 Suspension for injection for sheep and cattle

Authorised

- Clostridium haemolyticum, toxoid
- Clostridium sordellii, toxoid
- Tetanus toxoid adsorbed
- Clostridium septicum, toxoid
- Clostridium novyi, toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium perfringens, type B and C, beta toxoid
- Clostridium chauvoei, Inactivated
- Clostridium perfringens, type A and C, alpha toxoid

Product identification

Medicine name:
Covexin 10 Suspension for injection for sheep and cattle
COVEXIN 10 suspensie injectabilă pentru ovine și bovine

Active substance:
Clostridium haemolyticum, toxoid

Clostridium sordellii, toxoid

Tetanus toxoid adsorbed

Clostridium septicum, toxoid

Clostridium novyi, toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type B and C, beta toxoid

Clostridium chauvoei, Inactivated

Clostridium perfringens, type A and C, alpha toxoid

Target species:

Cattle

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Clostridium haemolyticum, toxoid

16.50 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium sordellii, toxoid

0.80 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Tetanus toxoid adsorbed

2.50 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium septicum, toxoid

3.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium novyi, toxoid

1.20 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium perfringens, type D, epsilon toxoid

5.10 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium perfringens, type B and C, beta toxoid

12.40 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium chauvoei, Inactivated

90.00 percentage protection / 1.00 millilitre(s)

Clostridium perfringens, type A and C, alpha toxoid

0.90 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB01
QI04AB01

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

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Authorisation status:
Valid

Authorised in:
Romania

Package description:
(ID2) 100 millilitre(s): Box (board) with 1 Bottle (high-density polyethylene) with 100 millilitre(s), closed with Stopfen (rubber) and Cap`` (aluminium)
(ID1) 50 millilitre(s): Box (board) with 1 Bottle (high-density polyethylene) with 50 millilitre(s), closed with Stopfen (rubber) and Cap`` (aluminium)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
19/01/2014

Manufacturing sites for batch release:
Zoetis Belgium
Schering-Plough Limited

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
150422

Date of authorisation status change:
18/02/2024

Reference member state:
Germany

Procedure number:
DE/V/0283/001

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Estonia France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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.

Combined File of all Documents