

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, cells and toxin, Inactivated
- Clostridium perfringens, type A and C, alpha toxoid

Product identification

Medicine name:

Covexin 10 Suspension for injection for sheep and cattle

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, cells and toxin, 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, cells and toxin, 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

Authorisation status:

Valid

Authorised in:

Latvia

Available in:

Latvia

Package description:

(ID2) 100 millilitre(s): Box (cardboard) with 1 Bottle (high-density polyethylene) with 100 millilitre(s), closed with Stopfen (rubber) and Cap`` (aluminium)

(ID2) 100 millilitre(s): Box (cardboard) with 1 Bottle (high-density polyethylene) with 100 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:

9/07/2010

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/10/0020

Date of authorisation status change:

11/07/2010

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)

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www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

Published on: 16/01/2026

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