

# 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

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**Target species:**

Cattle

Sheep

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**Route of administration:**

Subcutaneous use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Milk. 0 day
- Meat and offal. 0 day

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**Sheep**

- Meat and offal. 0 day
- Milk. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AB01

QI04AB01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Lithuania

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**Available in:**

Lithuania

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Zoetis Belgium

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**Marketing authorisation date:**

29/06/2010

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**Manufacturing sites for batch release:**

Zoetis Belgium

Schering-Plough Limited

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/10/1859/002-003

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**Date of authorisation status change:**

4/11/2023

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0283/001

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**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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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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