

# Multivac-9

Authorised

- Clostridium perfringens, type A, alpha toxoid
- Clostridium perfringens, type C, beta toxoid
- Clostridium perfringens, type B, epsilon toxoid
- Clostridium novyi, toxoid
- Clostridium septicum, toxoid
- Clostridium tetani, toxoid
- Clostridium sordellii, toxoid
- Clostridium chauvoei, Inactivated

## Product identification

### **Medicine name:**

Multivac-9

---

### **Active substance:**

Clostridium perfringens, type A, alpha toxoid

Clostridium perfringens, type C, beta toxoid

Clostridium perfringens, type B, epsilon toxoid

Clostridium novyi, toxoid

Clostridium septicum, toxoid

Clostridium tetani, toxoid

Clostridium sordellii, toxoid

Clostridium chauvoei, Inactivated

---

### **Target species:**

Cattle

Goat

Sheep

Pig

---

**Route of administration:**

Subcutaneous use

---

## Product details

**Active substance and strength:**

Clostridium perfringens, type A, alpha toxoid

1.00 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium perfringens, type C, beta toxoid

10.00 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium perfringens, type B, epsilon toxoid

5.00 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium novyi, toxoid

3.50 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium septicum, toxoid

2.50 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium tetani, toxoid

2.50 international unit(s)/millilitre / 2.00 millilitre(s)

Clostridium sordellii, toxoid

100.00 percentage protection / 2.00 millilitre(s)

Clostridium chauvoei, Inactivated

90.00 percentage protection / 2.00 millilitre(s)

---

**Pharmaceutical form:**

Suspension for injection

---

**Withdrawal period by route of administration:**

**Subcutaneous use:**

• **Cattle**

- Meat and offal. 0 day

• **Goat**

- Meat and offal. 0 day

• **Sheep**

- Meat and offal. 0 day

• **Pig**

- Meat and offal. 0 day

---

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

QI04AB01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Portugal

---

**Available in:**

Portugal

---

**Package description:**

Available only in Portuguese

Available only in Portuguese

Available only in Portuguese

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Ceva Saude Animal Produtos Farmaceuticos E Immunologicos Lda.

---

**Marketing authorisation date:**

This information is not available for this product.

---

**Manufacturing sites for batch release:**

Ceva-Phylaxia Veterinary Biologicals Co. Ltd.  
Cz Veterinaria S.A.

---

**Responsible authority:**

Directorate General For Food And Veterinary

---

**Authorisation number:**

208/87 DGV

---

**Date of authorisation status change:**

1/03/2022

---

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

This document does not exist in this language (English). You can find it in another language below.

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000098280>