# Miloxan suspensão injectável para bovinos, ovinos e caprinos.

Authorised

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

## Product identification

**Medicine name:**
Miloxan suspensão injectável para bovinos, ovinos e caprinos.

---

**Active substance:**
Clostridium chauvoei, toxoid

Clostridium sordellii, toxoid

Clostridium tetani, toxoid

Clostridium novyi, toxoid

Clostridium septicum, toxoid

Clostridium perfringens, type B, epsilon toxoid

Clostridium perfringens, type C, beta toxoid

---

**Target species:**
Cattle
Goat
Sheep

---

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Clostridium chauvoei, toxoid
1.00 90% protective dose in guinea pig / 2.00 millilitre(s)

Clostridium sordellii, toxoid
1.00 90% protective dose in guinea pig / 2.00 millilitre(s)

Clostridium tetani, toxoid
2.50 international unit(s) / 2.00 millilitre(s)

Clostridium novyi, toxoid
3.50 international unit(s) / 2.00 millilitre(s)

Clostridium septicum, toxoid
2.50 international unit(s) / 2.00 millilitre(s)

Clostridium perfringens, type B, epsilon toxoid
5.00 international unit(s) / 2.00 millilitre(s)

Clostridium perfringens, type C, beta toxoid
10.00 international unit(s) / 2.00 millilitre(s)

**Pharmaceutical form:**
Suspension for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Cattle**
- **Goat**
- **Sheep**

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI02AB01

**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

## 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:**
Boehringer Ingelheim Animal Health Portugal Unipessoal Lda.

**Marketing authorisation date:**
21/10/1986

**Manufacturing sites for batch release:**
Boehringer Ingelheim Animal Health France

**Responsible authority:**
DGAV

**Authorisation number:**
424/91 DGV

**Date of authorisation status change:**
1/05/2019

To consult adverse reactions on veterinary medicinal products please go to 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.

---