MILOXAN, Injekční suspenze

Not authorised

- Clostridium septicum, toxoid
- Clostridium tetani, toxoid
- Clostridium chauvoei, Inactivated
- Clostridium sordellii, toxoid
- Clostridium perfringens, beta toxoid
- Clostridium perfringens, epsilon toxoid
- Clostridium novyi, Inactivated

Product identification

Medicine name:
MILOXAN, Injekční suspenze

Active substance:
Clostridium septicum, toxoid

Clostridium tetani, toxoid

Clostridium chauvoei, Inactivated

Clostridium sordellii, toxoid

Clostridium perfringens, beta toxoid

Clostridium perfringens, epsilon toxoid

Clostridium novyi, Inactivated

Target species:
Goat
Pig
Sheep

Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium septicum, toxoid
2.50 international unit(s) / 1.00 Dose

Clostridium tetani, toxoid
2.50 international unit(s) / 1.00 Dose

Clostridium chauvoei, Inactivated
90.00 percent / 1.00 Dose

Clostridium sordellii, toxoid
90.00 percent / 1.00 Dose

Clostridium perfringens, beta toxoid
10.00 international unit(s) / 1.00 Dose

Clostridium perfringens, epsilon toxoid
5.00 international unit(s) / 1.00 Dose

Clostridium novyi, Inactivated
3.50 international unit(s) / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**

- Meat and offal. 0 day

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB01

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

Authorisation status:
Surrendered

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech

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 France

Marketing authorisation date:
15/12/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/181/98-C

Date of authorisation status change:
18/09/2024

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.

Package Leaflet

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

Labelling

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