Miloxan - Injektionssuspension für Rinder, Schafe und Ziegen

Authorised

- Clostridium perfringens, type B and C, beta toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium septicum, toxoid
- Clostridium novyi, toxoid
- CLOSTRIDIUM TETANI
- Clostridium chauvoei, Inactivated
- Clostridium sordellii, toxoid

Product identification

Medicine name:
Miloxan - Injektionssuspension für Rinder, Schafe und Ziegen

Active substance:
Clostridium perfringens, type B and C, beta toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium septicum, toxoid

Clostridium novyi, toxoid

CLOSTRIDIUM TETANI

Clostridium chauvoei, Inactivated

Clostridium sordellii, toxoid

Target species:
Cattle
Goat
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium perfringens, type B and C, beta toxoid
10.00 international unit(s) / 2.00 millilitre(s)

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

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

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

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

Clostridium chauvoei, Inactivated
90.00 percentage protection / 2.00 millilitre(s)

Clostridium sordellii, toxoid
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
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB01
QI03AB
QI04AB01

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
14/02/1994

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-20126

Date of authorisation status change:
14/02/1994

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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

Summary of Product Characteristics

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.
