Miloxan vakcina A.U.V.

Authorised

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

Product identification

Medicine name:
Miloxan vakcina A.U.V.

Active substance:
Clostridium sordellii, toxoid

Clostridium tetani, toxoid

Clostridium septicum, toxoid

Clostridium novyi, toxoid

Clostridium chauvoei, Inactivated

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type B and C, beta toxoid

Target species:
Cattle
Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium sordellii, toxoid
1.00 90% protective dose in guinea pig / 1.00 unit(s)/dose

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

Clostridium septicum, toxoid
2.50 international unit(s) / 1.00 unit(s)/dose

Clostridium novyi, toxoid
3.50 international unit(s) / 1.00 unit(s)/dose

Clostridium chauvoei, Inactivated
1.00 90% protective dose in guinea pig / 1.00 unit(s)/dose

Clostridium perfringens, type D, epsilon toxoid
5.00 international unit(s) / 1.00 unit(s)/dose

Clostridium perfringens, type B and C, beta toxoid
10.00 international unit(s) / 1.00 unit(s)/dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
-

Goat

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB01

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

Authorisation status:
Valid

Authorised in:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
1/07/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
1/07/1998

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