MILOXAN ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Not authorised

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

Product identification

Medicine name:
MILOXAN ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Active substance:
Clostridium chauvoei, toxoid

Clostridium sordellii, toxoid

Clostridium tetani, toxoid

Clostridium novyi, toxoid

Clostridium septicum, toxoid

Clostridium perfringens, type B and C, beta toxoid

Clostridium perfringens, type D, epsilon toxoid

Target species:
Goat
Sheep
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium chauvoei, toxoid
1.00 100% protective dose / 2.00 millilitre(s)

Clostridium sordellii, toxoid
1.00 100% protective dose / 2.00 millilitre(s)

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

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

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

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)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Goat**
  - Not applicable. no withdrawal period
- **Sheep**
  - Not applicable. no withdrawal period
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Cattle

- Not applicable. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR05DA12

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

Authorisation status:
Surrendered

Authorised in:
Greece

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
30/03/1992

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
National Organization For Medicines

Authorisation number:
6739-05-03-1992/31-03-1992/K-0044101

Date of authorisation status change:
29/05/2025

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