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COGLAVAX suspensie injectabilă pentru bovine, oi, capre

Authorised

- Clostridium chauvoei, cells and toxin, Inactivated
- Clostridium tetani, toxoid
- Clostridium novyi, type B, toxoid
- Clostridium septicum, toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium perfringens, type C, beta1 toxoid
- Clostridium perfringens, type A, alpha toxoid

Product identification

Medicine name:
COGLAVAX suspensie injectabilă pentru bovine, oi, capre

Active substance:
Clostridium chauvoei, cells and toxin, Inactivated

Clostridium tetani, toxoid

Clostridium novyi, type B, toxoid

Clostridium septicum, toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type C, beta1 toxoid

Clostridium perfringens, type A, alpha toxoid

Target species:
Sheep

Goat
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium chauvoei, cells and toxin, Inactivated

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

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

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

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

Clostridium perfringens, type C, beta1 toxoid
10.00 international unit(s)/dose / 2.00 millilitre(s)

Clostridium perfringens, type A, alpha toxoid
2.00 international unit(s)/dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB01

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale Romania S.R.L.

Marketing authorisation date:
17/07/2003

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
150319

Date of authorisation status change:
17/11/2025

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

Documents

Combined File of all Documents

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