COGLAVAX suspension for injection for cattle, sheep, goats

Authorised

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

Product identification

Medicine name:
COGLAVAX suspension for injection for cattle, sheep, goats

Active substance:
Clostridium chauvoei, cells and toxin, Inactivated

Clostridium tetani, toxoid

Clostridium septicum, toxoid

Clostridium novyi, type B, alpha toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type C, beta toxoid

Clostridium perfringens, type A, alpha toxoid

Target species:
Cattle

Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium chauvoei, cells and toxin, Inactivated
90.00 80% Protective Dose / 1.00 millilitre(s)

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

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

Clostridium novyi, type B, alpha toxoid
3.50 international unit(s) / 1.00 millilitre(s)

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

Clostridium perfringens, type C, beta toxoid
10.00 international unit(s) / 1.00 millilitre(s)

Clostridium perfringens, type A, alpha toxoid
2.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB01

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

Authorisation status:
Valid

Authorised in:

Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Animal Health Bulgaria EOOD

Marketing authorisation date:
28/06/2006

Manufacturing sites for batch release:
Ceva-Phylaxia Veterinary Biologicals Co. Ltd

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1505

Date of authorisation status change:
28/06/2006

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.