COGLAVAX

Authorised

- Clostridium perfringens, type A, alpha toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium perfringens, type C, beta toxoid
- Clostridium tetani, toxoid
- Clostridium septicum, toxoid
- Clostridium novyi, type B, toxoid
- Clostridium chauvoei, strain Hung 89, Inactivated

Product identification

Medicine name:
COGLAVAX

Active substance:
Clostridium perfringens, type A, alpha toxoid
Clostridium perfringens, type D, epsilon toxoid
Clostridium perfringens, type C, beta toxoid
Clostridium tetani, toxoid
Clostridium septicum, toxoid
Clostridium novyi, type B, toxoid
Clostridium chauvoei, strain Hung 89, Inactivated

Target species:
Cattle
Rabbit (for reproduction)
Rabbit
Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium perfringens, type A, alpha toxoid
4.00 international unit(s) / 1.00 unit(s)

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

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

Clostridium tetani, toxoid
5.00 international unit(s) / 1.00 unit(s)

Clostridium septicum, toxoid
5.00 international unit(s) / 1.00 unit(s)

Clostridium novyi, type B, toxoid
7.00 international unit(s) / 1.00 unit(s)

Clostridium chauvoei, strain Hung 89, Inactivated
1.00 100% protective dose / 1.00 unit(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - All relevant tissues. 0 day
- **Rabbit (for reproduction)**
  - All relevant tissues. 0 day
- **Rabbit**
  - All relevant tissues. 0 day
- **Sheep**
  - All relevant tissues. 0 day

- **Goat**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB01
QI03AB
QI04AB01
QI08AB

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
4/10/1982

Manufacturing sites for batch release:
Cz Veterinaria S.A.

Responsible authority:
National Veterinary Medicines Agency

Authorisation number:
FR/V/5726681 3/1982

Date of authorisation status change:
4/10/2012

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 and Labelling

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