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CUBOLAC POLICLOSTRIDIAL 7/11, SUSPENSION INYECTABLE PARA BOVINO, OVINO Y CAPRINO

Authorised

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

Product identification

Medicine name:
CUBOLAC POLICLOSTRIDIAL 7/11, SUSPENSION INYECTABLE PARA BOVINO, OVINO Y CAPRINO

Active substance:
Clostridium chauvoei, cells and toxin, Inactivated

Clostridium sordellii, toxoid

Clostridium novyi, type B, alpha toxoid

Clostridium septicum, toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type A, alpha toxoid

Clostridium perfringens, type C, beta1 toxoid

Target species:
Cattle
Sheep
Goat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Clostridium chauvoei, cells and toxin, Inactivated
100.00 percentage protection / 1.00 Dose

Clostridium sordellii, toxoid
100.00 percentage protection / 1.00 Dose

Clostridium novyi, type B, alpha toxoid
3.50 Toxicity unit(s) / 1.00 Dose

Clostridium septicum, toxoid
2.50 Toxicity unit(s) / 1.00 Dose

Clostridium perfringens, type D, epsilon toxoid
5.00 Toxicity unit(s) / 1.00 Dose

Clostridium perfringens, type A, alpha toxoid
0.30 Toxicity unit(s) / 1.00 Dose

Clostridium perfringens, type C, beta1 toxoid
10.00 Toxicity unit(s) / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**

- Meat and offal. 0 day
- Milk. 0 day

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

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

Subcutaneous use:

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

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

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

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

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

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Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
CZ Vaccines S.A.U.

Marketing authorisation date:
7/10/1997

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
1187 ESP

Date of authorisation status change:
25/12/2021

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

Documents

Summary of Product Characteristics

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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.