Covexin 8 suspensão injetável para ovinos e bovinos

Authorised

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

Product identification

Medicine name:
Covexin 8 suspensão injetável para ovinos e bovinos

Active substance:
Clostridium perfringens, type B and C, beta toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium chauvoei, Inactivated

Clostridium novyi, toxoid

Clostridium novyi, toxoid

Clostridium septicum, toxoid

Clostridium tetani, toxoid

Target species:
Cattle

Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium perfringens, type B and C, beta toxoid
10.00 international unit(s) / 5.00 millilitre(s)

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

Clostridium chauvoei, Inactivated
1.00 European Pharmacopoeia Unit(s) / 5.00 millilitre(s)

Clostridium novyi, toxoid
10.00 international unit(s) / 5.00 millilitre(s)

Clostridium novyi, toxoid
3.50 international unit(s) / 5.00 millilitre(s)

Clostridium septicum, toxoid
2.50 international unit(s) / 5.00 millilitre(s)

Clostridium tetani, toxoid
2.50 international unit(s) / 5.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB01

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 8(3) of Directive No 2001/83/EC)

Marketing authorisation holder:
Zoetis Portugal Lda.

Marketing authorisation date:
24/03/1988

Manufacturing sites for batch release:
Gmax Biopharm Belgium

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
168/87

Date of authorisation status change:
1/02/2013

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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