HEPTAVAC P PLUS SUSPENSION INYECTABLE PARA OVINO

Authorised

- Clostridium septicum, toxoid
- Clostridium novyi, toxoid
- Clostridium chauvoei, Inactivated
- Mannheimia haemolytica, Inactivated
- Bibersteinia trehalosi, serotype T3, strain S1109/84, Inactivated
- Clostridium perfringens, type C, beta1 toxoid
- Clostridium tetani, toxoid
- Clostridium perfringens, type D, epsilon toxoid

Product identification

Medicine name:
HEPTAVAC P PLUS SUSPENSION INYECTABLE PARA OVINO

Active substance:
Clostridium septicum, toxoid

Clostridium novyi, toxoid

Clostridium chauvoei, Inactivated

Mannheimia haemolytica, Inactivated

Bibersteinia trehalosi, serotype T3, strain S1109/84, Inactivated

Clostridium perfringens, type C, beta1 toxoid

Clostridium tetani, toxoid

Clostridium perfringens, type D, epsilon toxoid

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium septicum, toxoid
2.50 international unit(s) / 2.00 millilitre(s)

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

Clostridium chauvoei, Inactivated
1.00 Protective Dose / 2.00 millilitre(s)

Mannheimia haemolytica, Inactivated
9.00 relative potency / 2.00 millilitre(s)

Bibersteinia trehalosi, serotype T3, strain S1109/84, Inactivated
9.00 relative potency / 2.00 millilitre(s)

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

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

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB05

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

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:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
6/07/2016

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:

3425 ESP

Date of authorisation status change:
7/07/2016

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.