Heptavac P plus Injektionssuspension für Schafe

Authorised

- Clostridium perfringens, type C, beta toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium septicum, toxoid
- Tetanus toxoid adsorbed
- Clostridium novyi, toxoid
- Clostridium chauvoei, cells and toxin, Inactivated
- Mannheimia haemolytica, serotype A1, strain S1006/77, Inactivated
- Mannheimia haemolytica, serotype A2, strain S1126/92, Inactivated
- Mannheimia haemolytica, serotype A6, strain S1084/81, Inactivated
- Mannheimia haemolytica, serotype A7, strain S1078/81, Inactivated
- Mannheimia haemolytica, serotype A9, strain S994/77, Inactivated
- Bibersteinia trehalosi, serotype T4, strain S1085/81, Inactivated
- Bibersteinia trehalosi, serotype T3, strain S1109/84, Inactivated
- Bibersteinia trehalosi, serotype T10, strain S1075/81, Inactivated
- Bibersteinia trehalosi, serotype T15, strain S1105/84, Inactivated

Product identification

Medicine name:
Heptavac P plus Injektionssuspension für Schafe

Active substance:
Clostridium perfringens, type C, beta toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium septicum, toxoid

Tetanus toxoid adsorbed

Clostridium novyi, toxoid

Clostridium chauvoei, cells and toxin, Inactivated

Mannheimia haemolytica, serotype A1, strain S1006/77, Inactivated

Mannheimia haemolytica, serotype A2, strain S1126/92, Inactivated

Mannheimia haemolytica, serotype A6, strain S1084/81, Inactivated

Mannheimia haemolytica, serotype A7, strain S1078/81, Inactivated

Mannheimia haemolytica, serotype A9, strain S994/77, Inactivated

Bibersteinia trehalosi, serotype T4, strain S1085/81, Inactivated

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

Bibersteinia trehalosi, serotype T10, strain S1075/81, Inactivated

Bibersteinia trehalosi, serotype T15, strain S1105/84, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Clostridium perfringens, type C, beta toxoid
10.00 international unit(s) / 1.00 millilitre(s)

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

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

Tetanus toxoid adsorbed
2.50 international unit(s) / 1.00 millilitre(s)

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

Clostridium chauvoei, cells and toxin, Inactivated
0.50 90% protective dose in guinea pig / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A1, strain S1006/77, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A2, strain S1126/92, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A6, strain S1084/81, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A7, strain S1078/81, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A9, strain S994/77, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T4, strain S1085/81, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T3, strain S1109/84, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T10, strain S1075/81, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T15, strain S1105/84, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Milk. 0 day
  - 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:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
29/07/1999

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
73a/97

Date of authorisation status change:
15/07/2009

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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