HEPTAVAC IRP suspensie injectabilă pentru oi

Authorised

- Clostridium perfringens, type C, beta toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium septicum, toxoid
- Clostridium tetani, toxoid
- Clostridium novyi, toxoid
- Clostridium chauvoei, cells and toxin, Inactivated
- Mannheimia haemolytica, serotype A1, Inactivated
- Mannheimia haemolytica, serotype A2, Inactivated
- Mannheimia haemolytica, serotype A6, Inactivated
- Mannheimia haemolytica, serotype A7, Inactivated
- Mannheimia haemolytica, serotype A9, Inactivated
- Bibersteinia trehalosi, serotype T3, Inactivated
- Bibersteinia trehalosi, serotype T4, Inactivated
- Bibersteinia trehalosi, serotype T10, Inactivated
- Bibersteinia trehalosi, serotype T15, Inactivated

Product identification

Medicine name:
HEPTAVAC IRP suspensie injectabilă pentru oi

Active substance:
Clostridium perfringens, type C, beta toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium septicum, toxoid

Clostridium tetani, toxoid

Clostridium novyi, toxoid

Clostridium chauvoei, cells and toxin, Inactivated

Mannheimia haemolytica, serotype A1, Inactivated

Mannheimia haemolytica, serotype A2, Inactivated

Mannheimia haemolytica, serotype A6, Inactivated

Mannheimia haemolytica, serotype A7, Inactivated

Mannheimia haemolytica, serotype A9, Inactivated

Bibersteinia trehalosi, serotype T3, Inactivated

Bibersteinia trehalosi, serotype T4, Inactivated

Bibersteinia trehalosi, serotype T10, Inactivated

Bibersteinia trehalosi, serotype T15, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Clostridium perfringens, type C, beta toxoid
10.00 unit(s) / 1.00 millilitre(s)

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

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

Clostridium tetani, toxoid
2.50 unit(s) / 1.00 millilitre(s)

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

Clostridium chauvoei, cells and toxin, Inactivated
90.00 Relative Percentage Survival / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A1, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A2, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A6, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A7, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Mannheimia haemolytica, serotype A9, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T3, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T4, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T10, Inactivated
8.70 log10 colony forming unit(s) / 1.00 millilitre(s)

Bibersteinia trehalosi, serotype T15, Inactivated
8.70 log10 colony forming unit(s) / 1.00 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:
Romania

Available in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
4/08/2011

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
170027

Date of authorisation status change:
10/01/2024

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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