Ovilis Enzovax Lyophilisate and solvent for suspension for injection for sheep

Authorised

- Chlamydia abortus, strain 1B (thermosensitive), Live

Product identification

Medicine name:
Ovilis Enzovax Lyophilisate and solvent for suspension for injection for sheep

Active substance:
Chlamydia abortus, strain 1B (thermosensitive), Live

Target species:
Sheep

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Chlamydia abortus, strain 1B (thermosensitive), Live
100000.00 Inclusion forming unit(s) / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 7 day

Intramuscular use:

- **Sheep**
  - Meat and offal. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AE01

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID3) 10 Dose; 20 millilitre(s): Box (board) with 1 Bottle with 10 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle with 20 millilitre(s)
(ID7) 100 Dose; 200 millilitre(s): Box (board) with 1 Bottle with 100 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle with 200 millilitre(s)
(ID6) 50 Dose; 100 millilitre(s): Box (board) with 1 Bottle with 50 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle with 100 millilitre(s)
(ID4) 20 Dose; 40 millilitre(s): Box (board) with 1 Bottle with 20 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle with 40 millilitre(s)

(ID11) 100 Dose; 200 millilitre(s): Box (board) with 1 Bottle with 100 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle (polyethylene terephthalate) with 200 millilitre(s)
(ID10) 50 Dose; 100 millilitre(s): Box (board) with 1 Bottle with 50 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle (polyethylene terephthalate) with 100 millilitre(s)
(ID8) 10 Dose; 20 millilitre(s): Box (board) with 1 Bottle with 10 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle (polyethylene terephthalate) with 20 millilitre(s)
(ID9) 20 Dose; 40 millilitre(s): Box (board) with 1 Bottle with 20 Dose, closed with (Gummi) and (Aluminium) and 1 Bottle (polyethylene terephthalate) with 40 millilitre(s)

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:
22/03/2000

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
163a/95

Date of authorisation status change:

12/03/2010

Reference member state:
Germany

Procedure number:
DE/V/0246/001

Concerned member states:
Czechia Portugal Slovakia

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

Documents

Combined File of all Documents

English (PDF)
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