Bovilis Bovipast RSP suspension for injection for cattle

Authorised

- Bovine respiratory syncytial virus, strain EV 908, Inactivated
- Bovine parainfluenza virus 3, strain SF-4, Inactivated
- Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated

Product identification

Medicine name:
Bovilis Bovipast RSP suspension for injection for cattle
Bovilis Bovipast RSP injektionsvæske, suspension

Active substance:
Bovine respiratory syncytial virus, strain EV 908, Inactivated
Bovine parainfluenza virus 3, strain SF-4, Inactivated
Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Bovine respiratory syncytial virus, strain EV 908, Inactivated
281838.00 unit(s) / 5.00 millilitre(s)

Bovine parainfluenza virus 3, strain SF-4, Inactivated
70794.60 unit(s) / 5.00 millilitre(s)

Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated
100000.00 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

50 ml bottles of type I glass (10 doses), Ph. Eur., closed with injection stoppers type I rubber, Ph. Eur., sealed with an aluminiumcrimp cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
24/11/2005

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
38180

Date of authorisation status change:
24/11/2005

Reference member state:
Ireland

Procedure number:
IE/V/0537/001

Concerned member states:
Belgium Denmark Finland Greece Italy Luxembourg Norway Poland Portugal Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 25/09/2024
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Combined File of all Documents