

Bovigrip RSP plus

Injektionssuspension für Rinder

Authorised

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

Product identification

Medicine name:

Bovigrip RSP plus Injektionssuspension für Rinder

Active substance:

Bovine respiratory syncytial virus, strain EV 908, Inactivated

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

Bovine parainfluenza virus 3, strain SF-4, 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 enzyme-linked immunosorbent assay unit/dose / 5.00 millilitre(s)
Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated
100000.00 enzyme-linked immunosorbent assay unit/dose / 5.00 millilitre(s)
Bovine parainfluenza virus 3, strain SF-4, Inactivated
70794.60 enzyme-linked immunosorbent assay unit/dose / 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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL04

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

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:

4/07/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

PEI.V.02900.01.1

Date of authorisation status change:

2/03/2009

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

Documents

Package Leaflet

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Combined File of all Documents

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