RISPOVAL 4 BVD RS PI3 IBR LYOPHILISAT ET SUSPENSION POUR SUSPENSION INJECTABLE POUR BOVINS

Authorised

- Bovine parainfluenza virus 3, strain RLB103, Live
- Bovine viral diarrhoea virus 1, strain 5960, Inactivated
- Bovine herpesvirus 1, strain Cooper, Inactivated
- Bovine viral diarrhoea virus 1, strain 6309, Inactivated
- Bovine respiratory syncytial virus, strain 375, Live

Product identification

Medicine name:
RISPOVAL 4 BVD RS PI3 IBR LYOPHILISAT ET SUSPENSION POUR SUSPENSION INJECTABLE POUR BOVINS

Active substance:
Bovine parainfluenza virus 3, strain RLB103, Live

Bovine viral diarrhoea virus 1, strain 5960, Inactivated

Bovine herpesvirus 1, strain Cooper, Inactivated

Bovine viral diarrhoea virus 1, strain 6309, Inactivated

Bovine respiratory syncytial virus, strain 375, Live

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bovine parainfluenza virus 3, strain RLB103, Live
5.00 log10 50% cell culture infectious dose / 1.00 Dose

Bovine viral diarrhoea virus 1, strain 5960, Inactivated
0.75 unit(s) / 1.00 Dose

Bovine herpesvirus 1, strain Cooper, Inactivated
0.75 unit(s) / 1.00 Dose

Bovine viral diarrhoea virus 1, strain 6309, Inactivated
0.75 unit(s) / 1.00 Dose

Bovine respiratory syncytial virus, strain 375, Live
5.00 log10 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AH

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

Authorisation status:
Valid

Authorised in:

France

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis France

Marketing authorisation date:
12/10/2000

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0567671 0/2000

Date of authorisation status change:
27/04/2015

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.