Rispoval 3 BRSV Pi3 BVD Lyophilisate and suspension for suspension for injection for cattle

Authorised

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

Product identification

Medicine name:
RISPOVAL 3 BRSV PI3 BVD LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR CATTLE
Rispoval 3 BRSV Pi3 BVD Lyophilisate and suspension for suspension for injection for cattle

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

Bovine viral diarrhoea virus 1, strain 5960, 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
100000.00 cell culture infective dose 50 / 1.00 unit(s)

Bovine viral diarrhoea virus 1, strain 5960, Inactivated
3.00 log2 serum neutralising unit(s) / 1.00 unit(s)

Bovine viral diarrhoea virus 1, strain 6309, Inactivated
3.00 log2 serum neutralising unit(s) / 1.00 unit(s)

Bovine respiratory syncytial virus, strain 375, Live
100000.00 cell culture infective dose 50 / 1.00 unit(s)

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 not subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Available only in French

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Belgium S.A.

Marketing authorisation date:
9/12/2013

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/060/001

Date of authorisation status change:
9/12/2013

Reference member state:
France

Procedure number:
FR/V/0146/001

Concerned member states:
Belgium Bulgaria Czechia Estonia Germany Hungary Ireland Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

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