RESPIBOV

Not authorised

- Bovine herpesvirus 1, strain Colorado, Inactivated
- Bovine viral diarrhoea virus, strain Singer, Inactivated
- Bovine parainfluenza virus 3, strain SF-4, Inactivated
- Mannheimia haemolytica, strain Salamanca, Inactivated
- Pasteurella multocida, strain Dario, Inactivated

Product identification

Medicine name:
RESPIBOV

Active substance:
Bovine herpesvirus 1, strain Colorado, Inactivated
Bovine viral diarrhoea virus, strain Singer, Inactivated
Bovine parainfluenza virus 3, strain SF-4, Inactivated
Mannheimia haemolytica, strain Salamanca, Inactivated
Pasteurella multocida, strain Dario, Inactivated

Target species:
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Bovine herpesvirus 1, strain Colorado, Inactivated
8.00 serum neutralising unit(s) / 1.00 Dose

Bovine viral diarrhoea virus, strain Singer, Inactivated
32.00 serum neutralising unit(s) / 1.00 Dose

Bovine parainfluenza virus 3, strain SF-4, Inactivated
64.00 serum neutralising unit(s) / 1.00 Dose

Mannheimia haemolytica, strain Salamanca, Inactivated
80.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pasteurella multocida, strain Dario, Inactivated
80.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Laboratorios Syva S.A.

Marketing authorisation date:
10/08/1994

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
2862 ESP

Date of authorisation status change:
17/02/2023

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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