BioBos Respi 3, Suspension for injection

Authorised

- Bovine respiratory syncytial virus, strain BIO-24, Inactivated
- Bovine parainfluenza virus 3, strain BIO-23, Inactivated
- Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

Product identification

Medicine name:
BioBos Respi 3, Suspension for injection

Active substance:
Bovine respiratory syncytial virus, strain BIO-24, Inactivated

Bovine parainfluenza virus 3, strain BIO-23, Inactivated

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Bovine respiratory syncytial virus, strain BIO-24, Inactivated
1.00 relative potency / 1.00 Dose

Bovine parainfluenza virus 3, strain BIO-23, Inactivated
1.00 relative potency / 1.00 Dose

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 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:
Lithuania

Package description:
Glass Vial 10 x 5.0 Dose krabička plast s jamkami
Glass Vial 1 x 5.0 Dose krabička
Glass Vial 1 x 25.0 Dose krabička
Glass Vial 1 x 50.0 Dose krabička
Glass Vial 10 x 5.0 Dose kartonáž pro hromadné balení
Plastic Vial 10 x 5.0 Dose krabička plast s jamkami
Plastic Vial 1 x 5.0 Dose krabička

Plastic Vial 1 x 25.0 Dose krabička
Plastic Vial 1 x 50.0 Dose krabička
Plastic Vial 10 x 5.0 Dose kartonáž pro hromadné balení

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:
Bioveta a.s.

Marketing authorisation date:
1/07/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/18/2478/001-004

Date of authorisation status change:
24/07/2025

Reference member state:
Czechia

Procedure number:
CZ/V/0143/001

Concerned member states:
Estonia Latvia Lithuania

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

Labelling