

BioBos Respi 2 intranasal, Nasal spray, lyophilisate and solvent for suspension

Authorised

- Bovine respiratory syncytial virus, strain BIO 24/A, Live
- Bovine parainfluenza virus 3, strain BIO 23/A, Live

Product identification

Medicine name:

BioBos Respi 2 intranasal, Nasal spray, lyophilisate and solvent for suspension
BioBos Respi 2 intranasal, nosový sprej, lyofilizát a rozpúšťadlo na suspenziu

Active substance:

Bovine respiratory syncytial virus, strain BIO 24/A, Live
Bovine parainfluenza virus 3, strain BIO 23/A, Live

Target species:

Cattle

Route of administration:

Nasal use

Product details

Active substance and strength:

Bovine respiratory syncytial virus, strain BIO 24/A, Live
7.50 log₁₀ tissue culture infective dose 50 / 1.00 Dose

Bovine parainfluenza virus 3, strain BIO 23/A, Live
6.00 log₁₀ tissue culture infective dose 50 / 1.00 Dose

Pharmaceutical form:

Nasal spray, lyophilisate and solvent for suspension

Withdrawal period by route of administration:

Nasal use:

• **Cattle**

- Milk. 0 hour
 - Meat and offal. 0 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AD07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Glass Vial 5 x 5.0 Dose

Glass Vial 1 x 5.0 Dose

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:

5/05/2017

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/021/MR/17-S

Date of authorisation status change:

5/05/2017

Reference member state:

Czechia

Procedure number:

CZ/V/0136/001

Concerned member states:

Bulgaria Croatia Cyprus Estonia Hungary Latvia Lithuania Poland Slovakia
Slovenia

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

Documents

Combined File of all Documents

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