# Bovalto Respi 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:**
Bovalto Respi Intranasal, Nasal spray, lyophilisate and solvent for suspension
Bovalto Respi 2 Nässpray, frystorkat pulver och vätska till suspension

**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 50% tissue culture infectious dose / 1.00 Dose

Bovine parainfluenza virus 3, strain BIO 23/A, Live
6.00 log10 50% tissue culture infectious dose / 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

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI02AD07

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

**Authorisation status:**
Valid

**Authorised in:**
Sweden

**Available in:**
Sweden

**Package description:**
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

## 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:**
Boehringer Ingelheim Animal Health Denmark A/S

**Marketing authorisation date:**
9/03/2018

**Manufacturing sites for batch release:**
Bioveta a.s.

**Responsible authority:**
Swedish Medical Products Agency

**Authorisation number:**
56912

**Date of authorisation status change:**
9/03/2018

**Reference member state:**
Czechia

**Procedure number:**
CZ/V/0141/001

**Concerned member states:**
Austria Belgium Denmark France Germany Greece Ireland Italy Luxembourg Malta Netherlands Norway Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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

## Documents

Package Leaflet

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

Summary of Product Characteristics

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.