

# Bovilis INtranasal RSP Live nasal spray, lyophilisate and solvent for suspension for cattle

Authorised

- Bovine parainfluenza virus 3, strain INT2-2013, Live
- Bovine respiratory syncytial virus, strain Jencine-2013, Live

## Product identification

### **Medicine name:**

Bovilis INtranasal RSP Live nasal spray, lyophilisate and solvent for suspension for cattle

Bovilis RSP Live Vet. lyofilisat og væske til nesespray, suspensjon til storfe

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### **Active substance:**

Bovine parainfluenza virus 3, strain INT2-2013, Live

Bovine respiratory syncytial virus, strain Jencine-2013, Live

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### **Target species:**

Cattle

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### **Route of administration:**

Nasal use

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## Product details

### **Active substance and strength:**

Bovine parainfluenza virus 3, strain INT2-2013, Live  
63095.70 unit(s) / 1.00 Dose

Bovine respiratory syncytial virus, strain Jencine-2013, Live  
100000.00 unit(s) / 1.00 Dose

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**Pharmaceutical form:**

Nasal spray, suspension

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**Withdrawal period by route of administration:**

**Nasal use:**

• **Cattle**

- Meat and offal. 0 day

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

QI02AD07

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Norway

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**Package description:**

Cardboard box with 50 doses of lyophilisate and cardboard box with 100 ml of solvent

Cardboard box with 25 doses of lyophilisate and cardboard box with 50 ml of solvent

Cardboard box 5 x 5 doses of lyophilisate and 5 x 10 ml of solvent

Cardboard box 5 x 10 doses of lyophilisate and 5 x 20 ml of solvent

Cardboard box with 20 doses of lyophilisate and cardboard box with 40 ml of solvent

Cardboard box with 1 dose of lyophilisate and 2 ml of solvent.

Cardboard box 5 x 1 dose of lyophilisate and 5 x 2 ml of solvent

Cardboard box 5 doses of lyophilisate and 10 ml of solvent.

Cardboard box 10 doses of lyophilisate and 20 ml of solvent.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

This information is not available for this product.

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**Manufacturing sites for batch release:**

INTERVET INTERNATIONAL B.V.

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

18-12322

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**Date of authorisation status change:**

26/06/2019

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0257/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.

### Combined File of all Documents

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