

# Bovalto 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:**

Bovalto Respi 3, Suspension for injection

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**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

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## 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

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Milk. 0 hour

- Meat and offal. 0 day

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

QI02AL

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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:**

Luxembourg

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**Available in:**

Luxembourg

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**Package description:**

Plastic Vial 10 x 5.0 Dose

Glass Vial 10 x 5.0 Dose

Plastic Vial 1 x 50.0 Dose

Glass Vial 1 x 50.0 Dose

Plastic Vial 1 x 25.0 Dose

Glass Vial 1 x 25.0 Dose  
Plastic Vial 1 x 5.0 Dose  
Glass Vial 1 x 5.0 Dose  
Plastic Vial 10 x 5.0 Dose  
Glass Vial 10 x 5.0 Dose

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health Belgium S.A.

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**Marketing authorisation date:**

2/12/2015

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 344/16/01/1505

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**Date of authorisation status change:**

2/12/2015

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0128/001

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**Concerned member states:**

Austria Belgium Denmark France Germany Greece Ireland Italy  
Luxembourg Netherlands Norway Poland Portugal Romania 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)