

# BioBos Respi 3 vakcina A.U.V.

Authorised

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

## Product identification

**Medicine name:**

BioBos Respi 3 vakcina A.U.V.

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**Active substance:**

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

Bovine parainfluenza virus 3, strain BIO-23, Inactivated

Bovine respiratory syncytial virus, strain BIO-24, 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:**

Mannheimia haemolytica, serotype A1, strain DSM 5283, Inactivated

1.00 relative potency / 1.00 unit(s)/dose

Bovine parainfluenza virus 3, strain BIO-23, Inactivated

1.00 relative potency / 1.00 unit(s)/dose

Bovine respiratory syncytial virus, strain BIO-24, Inactivated

1.00 relative potency / 1.00 unit(s)/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**

- Meat and offal. 0 day

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

QI02AL04

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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:**

Hungary

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**Package description:**

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian

Available only in Hungarian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Alphavet Zrt.

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**Marketing authorisation date:**

7/12/2016

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

7/12/2016

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