

# BioRabbit RHDV 1,2, Suspension for injection

Authorised

- Rabbit haemorrhagic disease virus, type 1, strain Borohradek, Inactivated
- Rabbit haemorrhagic disease virus, type 2, strain Ceska Lipa, Inactivated

## Product identification

### Medicine name:

BioRabbit RHDV 1,2, Suspension for injection  
BioRabbit RHDV 1, 2, injekcinė suspensija triušiams

### Active substance:

Rabbit haemorrhagic disease virus, type 1, strain Borohradek, Inactivated  
Rabbit haemorrhagic disease virus, type 2, strain Ceska Lipa, Inactivated

### Target species:

Rabbit

### Route of administration:

Subcutaneous use

## Product details

### Active substance and strength:

Rabbit haemorrhagic disease virus, type 1, strain Borohradek, Inactivated  
60.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Rabbit haemorrhagic disease virus, type 2, strain Ceska Lipa, Inactivated  
80.00 haemagglutination inhibiting unit(s) / 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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**Rabbit**

- Meat and offal. 0 day

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

QI08AA01

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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:**

Lithuania

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**Package description:**

Plastic Vial 10 x 20.0 Dose

Plastic Vial 10 x 10.0 Dose

Plastic Vial 1 x 20.0 Dose

Plastic Vial 1 x 10.0 Dose

Glass Vial 10 x 20.0 Dose

Glass Vial 10 x 10.0 Dose

Glass Vial 10 x 1.0 Dose

Glass Vial 1 x 20.0 Dose

Glass Vial 1 x 10.0 Dose

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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 8 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Bioveta a.s.

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**Marketing authorisation date:**

7/06/2023

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/23/2753/001-005

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**Date of authorisation status change:**

7/06/2023

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0180/001

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**Concerned member states:**

Bulgaria Croatia Greece Hungary Latvia Lithuania Poland Romania Slovakia  
Slovenia

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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

Combined File of all Documents

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