# 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

#### **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

#### **Pharmaceutical form:**

Suspension for injection

# Withdrawal period by route of administration:

Subcutaneous use:

#### **Rabbit**

- Meat and offal. 0 day

# Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AA01

# Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Hungary

# 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

# Additional information

# **Entitlement type:**

# Marketing Authorisation

### Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

# Marketing authorisation holder:

Bioveta a.s.

### Marketing authorisation date:

6/10/2023

### Manufacturing sites for batch release:

Bioveta a.s.

# **Responsible authority:**

Directorate Of Veterinary Medicinal Products

#### **Authorisation number:**

4366/X/23 NÉBIH ÁTI

# Date of authorisation status change:

23/05/2023

#### **Reference member state:**

Czechia

#### **Procedure number:**

CZ/V/0180/001

#### **Concerned member states:**

Bulgaria Croatia Greece Hungary Latvia Lithuania Poland Romania Slovakia Slovenia

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

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