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
80.00 haemagglutination inhibiting unit(s) / 1.00 Dose

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:
Slovakia

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:
21/04/2023

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/002/DC/23-S

Date of authorisation status change:
21/04/2023

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

Package Leaflet

Labelling