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:
Czechia

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:
23/02/2023

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/011/23-C

Date of authorisation status change:
23/02/2023

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

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

English (PDF)
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Labelling

English (PDF)
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