

BioRabbit RHDV 1,2, Suspension for injection

Authorised

- Rabbit haemorrhagic disease virus 1, strain IM.507.SC.2011, Inactivated
- Rabbit haemorrhagic disease virus 2, Inactivated

Product identification

Medicine name:

BioRabbit RHDV 1,2, Suspension for injection

Active substance:

Rabbit haemorrhagic disease virus 1, strain IM.507.SC.2011, Inactivated

Rabbit haemorrhagic disease virus 2, Inactivated

Target species:

Rabbit

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus 1, strain IM.507.SC.2011, Inactivated

60.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Rabbit haemorrhagic disease virus 2, 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:

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000983564>