

Fencovis, Suspension for injection

Authorised

- Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated
- Bovine coronavirus, strain C-197, Inactivated
- Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

Product identification

Medicine name:

Fencovis, Suspension for injection

Active substance:

Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated

Bovine coronavirus, strain C-197, Inactivated

Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated

1.00 relative potency / 1.00 Dose

Bovine coronavirus, strain C-197, Inactivated

1.00 relative potency / 1.00 Dose

Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 0 day

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

Package description:

Glass Vial 2 x 1.0 Dose

Glass Vial 10 x 1.0 Dose

Glass Vial 20 x 1.0 Dose

Glass Vial 1 x 5.0 Dose

Plastic Vial 1 x 5.0 Dose

Glass Vial 5 x 5.0 Dose

Plastic Vial 5 x 5.0 Dose

Glass Vial 10 x 5.0 Dose
Plastic Vial 10 x 5.0 Dose
Glass Vial 1 x 25.0 Dose
Plastic Vial 1 x 25.0 Dose
Glass Vial 12 x 25.0 Dose
Plastic Vial 12 x 25.0 Dose
Glass Vial 24 x 25.0 Dose
Plastic Vial 24 x 25.0 Dose
Glass Vial 1 x 50.0 Dose
Plastic Vial 1 x 50.0 Dose

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

30/07/2022

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

841307

Date of authorisation status change:

30/07/2022

Reference member state:

Czechia

Procedure number:

CZ/V/0177/001

Concerned member states:

Austria Belgium Finland France Germany Greece Ireland Italy Luxembourg
Netherlands Norway Portugal Spain Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 25/08/2022

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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