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
FENCOVIS SUSPENSION INJECTABLE

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:

- **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

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Authorisation status:
Valid

Authorised in:
France

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:
27/06/2022

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8946295 9/2022

Date of authorisation status change:
27/06/2022

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

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

Package Leaflet and Labelling

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