

Bovilis Rotavec Corona Emulsion for Injection for Cattle

Authorised

- Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), strain CN7985, Inactivated
- Bovine coronavirus, strain Mebus, Inactivated
- Bovine rotavirus A, type G6P5, strain UK-Compton, Inactivated

Product identification

Medicine name:

Bovilis Rotavec Corona Emulsion for Injection for Cattle

Bovilis Rotavec Corona vet. Injektionsvätska, emulsion

Active substance:

Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), strain CN7985, Inactivated

Bovine coronavirus, strain Mebus, Inactivated

Bovine rotavirus A, type G6P5, strain UK-Compton, Inactivated

Target species:

Cattle

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), strain CN7985, Inactivated

560.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Bovine coronavirus, strain Mebus, Inactivated

340.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Bovine rotavirus A, type G6P5, strain UK-Compton, Inactivated

874.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 0 day
- Milk. 0 day

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day
 - Milk. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Package description:

Available only in [Swedish](#)

Available only in [Swedish](#)

Available only in [Swedish](#)

Available only in [Swedish](#)

Available only in [Swedish](#)

Available only in [Swedish](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

28/11/2008

Manufacturing sites for batch release:

Burgwedel Biotech GmbH

Intervet International B.V.

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

25819

Date of authorisation status change:

28/11/2008

Reference member state:

Germany

Procedure number:

DE/V/0276/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France
Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta
Netherlands Norway Poland Portugal Romania Slovakia Slovenia 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: 28/01/2022

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Package Leaflet

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Combined File of all Documents

Labelling

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