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 EMULSION INJECTABLE POUR BOVINS

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

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Authorisation status:

Valid

Authorised in:
France

Package description:
(ID3) 40 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 40 millilitre(s)
(ID4) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 100 millilitre(s)
(ID2) 10 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 10 millilitre(s)
(ID1) 20 millilitre(s): Box (Cardboard) with 10 Vial (Glass) each with 2 millilitre(s)
(ID5) 10 millilitre(s): Box (Cardboard) with 1 Vial (PolyEthylene TerePhthalate) with 10 millilitre(s)
(ID6) 40 millilitre(s): Box (Cardboard) with 1 Vial (PolyEthylene TerePhthalate) with 40 millilitre(s)
(ID7) 100 millilitre(s): Box (Cardboard) with 1 Vial (PolyEthylene TerePhthalate) with 100 millilitre(s)

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

Marketing authorisation date:
8/09/2000

Manufacturing sites for batch release:
Burgwedel Biotech GmbH
Intervet International B.V.

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

Authorisation number:
FR/V/1056429 7/2000

Date of authorisation status change:
8/09/2010

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 and Labelling

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