

Bovilis Rotavec Corona Emulsion for Injection for Cattle

Not
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

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Malta

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:

European Medicines Agency

Marketing authorisation date:

27/11/2100

Manufacturing sites for batch release:

Burgwedel Biotech GmbH
Intervet International B.V.

Responsible authority:

Veterinary And Phytosanitary Regulation Department

Authorisation number:

VMA

Date of authorisation status change:

27/11/2024

Reference member state:

Germany

Procedure number:

DE/V/0276/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents