Bovilis Rotavec Corona, emulzija za injekciju, za goveda

Authorised

- Escherichia coli, fimbrial adhesin F5
- Bovine coronavirus, strain Mebus, Inactivated
- Bovine rotavirus A, type G6P5, strain UK-Compton, Inactivated

Product identification

Medicine name:
Bovilis Rotavec Corona, emulzija za injekciju, za goveda

Active substance:
Escherichia coli, fimbrial adhesin F5

Bovine coronavirus, strain Mebus, Inactivated

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

Target species:
Cattle (pregnant cow)
Cattle (pregnant heifer)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Escherichia coli, fimbrial adhesin F5
0.64 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Bovine coronavirus, strain Mebus, Inactivated
3.41 log10 ELISA unit / 2.00 millilitre(s)

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

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Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (pregnant cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (pregnant heifer)**
  - 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:
Croatia

Package description:
Available only in Croatian
Available only in Croatian
Available only in Croatian

Available only in Croatian

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:
Intervet International B.V. Subsidiary In The Republic Of Croatia

Marketing authorisation date:
20/05/2015

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

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/22-01/287

Date of authorisation status change:
4/04/2024

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

Documents

Summary of Product Characteristics

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