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# Bovilis Rotavec Corona Emulsion for Injection for Cattle

Ima dovoljenje za promet

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

## Informacije o zdravilu

**Ime zdravila:**
Bovilis Rotavec Corona Emulsion for Injection for Cattle

**Učinkovina:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

**Ciljne živalske vrste:**
govedo

**Pot uporabe:**
intramuskularna uporaba
Subkutana uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
560.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Na voljo samo v English
874.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Na voljo samo v English
340.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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**Farmacevtska oblika:**

Emulzija za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

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

**Subkutana uporaba:**

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

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**

QI02AL01

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**Ima dovoljenje za promet v:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Na voljo v:**
Germany

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

## Dodatne informacije

**Vrsta dovoljenja:**
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**
Na voljo samo v English Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Intervet Deutschland GmbH

**Datum dovoljenja za promet z zdravilom:**
16/03/2005

**Proizvodna mesta, odgovorna za sproščanje serij:**
Intervet International B.V.

**Pristojni organ:**
Paul-Ehrlich-Institut

**Številka dovoljenja za promet z zdravilom:**
PEI.V.01312.01.1

**Datum spremembe statusa dovoljenja za promet:**
28/05/2010

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**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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**Številka postopka:**
DE/V/0276/001

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**Zadevne države članice:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.