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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)

Farmacevtska oblika:

Emulzija za injiciranje

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

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

QI02AL01

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:

Valid

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

Na voljo v:
Cyprus

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 Nederland B.V.

Datum dovoljenja za promet z zdravilom:
3/09/2008

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

Pristojni organ:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Številka dovoljenja za promet z zdravilom:
CY00164V

Datum spremembe statusa dovoljenja za promet:
15/07/2010

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

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak 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.