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BOVIGEN SCOUR Emulsion for injection for cattle

Ima dovoljenje za promet

- Bovine coronavirus, strain C-197, Inactivated
- Bovine rotavirus A, type G6P1, strain TM-91, Inactivated
- Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain EC/17, Inactivated

Informacije o zdravilu

Ime zdravila:
BOVIGEN SCOUR 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

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
5.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Na voljo samo v English
6.00 log2 virus neutralising unit(s) / 1.00 Dose

Na voljo samo v English
44.80 enzyme-linked immunosorbent assay unit / 1.00 Dose

Farmacevtska oblika:

Emulzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

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

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 English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Na voljo v:

Sweden

Opis ovojnine zdravila:

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

Imetnik dovoljenja za promet z zdravilom:
Forte Healthcare Limited

Datum dovoljenja za promet z zdravilom:
2/10/2015

Proizvodna mesta, odgovorna za sproščanje serij:
Pharmagal Bio spol. s r.o.

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:
52898

Datum spremembe statusa dovoljenja za promet:
2/10/2015

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

Številka postopka:
IE/V/0341/001

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
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Na voljo samo v Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian 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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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

Combined File of all Documents

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

Označevanje

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