Rispoval 3 BRSV Pi3 BVD Lyophilisate and Suspension for Suspension for Injection for Cattle

Ima dovoljenje za promet

- Bovine parainfluenza virus 3, strain RLB103, Live
- Bovine viral diarrhoea virus 1, strain 5960, Inactivated
- Bovine viral diarrhoea virus 1, strain 6309, Inactivated
- Bovine respiratory syncytial virus, strain 375, Live

Informacije o zdravilu

Ime zdravila:
RISPOVAL 3 BRSV PI3 BVD LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR CATTLE
Rispoval 3 BRSV Pi3 BVD Lyophilisate and Suspension for Suspension for Injection for Cattle

Učinkovina:
Na voljo samo v English
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
100000.00 50% cell culture infectious dose / 1.00 Dose

Na voljo samo v English
3.00 log2 serum neutralising unit(s) / 1.00 Dose

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3.00 log2 serum neutralising unit(s) / 1.00 Dose

Na voljo samo v English
100000.00 50% cell culture infectious dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat in suspenzija za suspenzijo za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - All relevant tissues. 0 day

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

QI02AH

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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Na voljo v:
United Kingdom (Northern Ireland)

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Belgium S.A.

Datum dovoljenja za promet z zdravilom:
3/05/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Belgium

Pristojni organ:
The Veterinary Medicines Directorate

Številka dovoljenja za promet z zdravilom:
Vm 60021/3044

Datum spremembe statusa dovoljenja za promet:
27/11/2024

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

Številka postopka:

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 Latvian 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