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Bovilis INtranasal RSP Live nasal spray, lyophilisate and solvent for suspension for cattle

Ima dovoljenje za promet

- Bovine parainfluenza virus 3, strain INT2-2013, Live
- Bovine respiratory syncytial virus, strain Jencine-2013, Live

Informacije o zdravilu

Ime zdravila:
Bovilis INtranasal RSP Live nasal spray, lyophilisate and solvent for suspension for cattle

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

Ciljne živalske vrste:
govedo

Pot uporabe:
Nazalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
4.80 log10 50% tissue culture infectious dose / 2.00 millilitre(s)

Na voljo samo v English
5.00 log10 50% tissue culture infectious dose / 2.00 millilitre(s)

Farmacevtska oblika:

Pršilo za nos, suspenzija

Karenca glede na pot uporabe:

Nazalna uporaba:

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

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

QI02AD07

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

Na voljo v:

Denmark

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

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
6/06/2019

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

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
61208

Datum spremembe statusa dovoljenja za promet:
6/06/2019

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

Številka postopka:
NL/V/0257/001

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

Povzetek glavnih značilnosti zdravila

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.