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VERSICAN PLUS BB ORAL LYOPHILISAT AND SOLVENT FOR ORAL SUSPENSION FOR DOGS

Ima dovoljenje za promet

- Water, purified
- Bordetella bronchiseptica, strain 92 B, Live

Informacije o zdravilu

Ime zdravila:
VERSICAN PLUS BB ORAL LYOPHILISAT AND SOLVENT FOR ORAL SUSPENSION FOR DOGS

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

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
1.00 millilitre(s) / 1.00 Dose

Na voljo samo v English
140000000.00 Colony forming unit / 1.00 Dose

Farmacevtska oblika:

Liofilizat in vehikel za peroralno suspenzijo

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

QI07AE01

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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Opis ovojnine zdravila:

Na voljo samo v French

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

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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 France

Datum dovoljenja za promet z zdravilom:
20/12/2018

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Belgium

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/6855258 5/2018

Datum spremembe statusa dovoljenja za promet:
10/03/2023

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

Številka postopka:
FR/V/0401/001

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.

Package Leaflet and Labelling

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.

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