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Lincoral-S 222 mg/g + 444.7 mg/g powder for use in drinking water

Ima dovoljenje za promet

- Spectinomycin sulfate tetrahydrate
- Lincomycin hydrochloride monohydrate

Informacije o zdravilu

Ime zdravila:
Lincoral-S 222 mg/g + 444.7 mg/g powder for use in drinking water

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
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:
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
672.40 milligram(s) / 1.00 gram(s)

Na voljo samo v English
251.70 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:

Prašek za dajanje v vodo za pitje

Karenca glede na pot uporabe:

Dajanje v vodo za pitje:

- **Chicken**

  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 5 day

Animals must not be slaughtered for human consumption during treatment.

- **Pig**

  - Meat and offal. 0 day

Animals must not be slaughtered for human consumption during treatment.

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

QJ01FF52

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

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:
HuVepharma

Datum dovoljenja za promet z zdravilom:
1/09/2023

Proizvodna mesta, odgovorna za sproščanje serij:
Huvepharma S.A.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
67188

Datum spremembe statusa dovoljenja za promet:
1/09/2023

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:
IE/V/0668/001

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
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.

Povzetek glavnih značilnosti zdravila

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

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