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PRIMUN SALMONELLA T Lyophilisate for use in drinking water for chickens

Ima dovoljenje za promet

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain ST CAL 16 Str+/Rif+/Enr-, Live

Informacije o zdravilu

Ime zdravila:
PRIMUN SALMONELLA T Lyophilisate for use in drinking water for chickens

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Spanish Czech Danish Estonian English French Italian Latvian Lithuanian Romanian Finnish Swedish Norwegian
Na voljo samo v Spanish Estonian English French Italian Latvian Lithuanian Romanian Finnish Norwegian

Pot uporabe:
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
100000000.00 Colony forming unit / 1.00 Dose

Farmacevtska oblika:

Liofilizat za dajanje v vodo za pitje

Karenca glede na pot uporabe:

Dajanje v vodo za pitje:

- **Chicken (pullet for egg production, future layer)**

  - Meat and offal. no withdrawal period

28 days after first and second vaccination and 14 days after third vaccination.

  - Eggs. no withdrawal period

Do not use in birds in lay and within 4 weeks before the start of the laying period.

- **Future breeder pullet**

  - Meat and offal. no withdrawal period

28 days after first and second vaccination and 14 days after third vaccination.

  - Eggs. no withdrawal period

Do not use in birds in lay and within 4 weeks before the start of the laying period.

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

QI01AE01

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

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

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:
Laboratorios Calier S.A.

Datum dovoljenja za promet z zdravilom:
13/07/2022

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Calier S.A.

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.12106.01.1

Datum spremembe statusa dovoljenja za promet:
13/07/2022

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

Številka postopka:
ES/V/0408/001

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

Označevanje

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.

Navodilo za uporabo

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

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