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Nextmune concentrate and solvent for suspension for injection for chickens

Ima dovoljenje za promet

- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Live

Informacije o zdravilu

Ime zdravila:
Nextmune concentrate and solvent for suspension for injection for chickens

Učinkovina:
Na voljo samo v English

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

Pot uporabe:
Subkutana uporaba
V jajčno celico

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
2.70 log 10 50% embryo infective dose / 1.00 Dose

Farmacevtska oblika:

Koncentrat in vehikel za suspenzijo za injiciranje

Karenca glede na pot uporabe:

Subkutana uporaba:

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

V jajčno celico:

- **Chicken (embryonated eggs)**
  - Meat and offal. 0 day

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

QI01AD09

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:

Germany

Opis ovojnine zdravila:

Na voljo samo v German

Na voljo samo v German

Na voljo samo v German

Na voljo samo v German

Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German

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:
Ceva Tiergesundheit GmbH

Datum dovoljenja za promet z zdravilom:
6/07/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Ceva-Phylaxia Zrt.

Pristojni organ:
Paul-Ehrlich-Institut

Številka dovoljenja za promet z zdravilom:
PEI.V.11992.01.1

Datum spremembe statusa dovoljenja za promet:
6/07/2020

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/0337/001

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

Označevanje

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.