EXCENEL FLOW 50 MG/ML SUSPENSION FOR INJECTION FOR PIGS AND CATTLE

Ima dovoljenje za promet

- Ceftiofur hydrochloride

Informacije o zdravilu

Ime zdravila:
EXCENEL FLOW 50 MG/ML SUSPENSION FOR INJECTION FOR PIGS AND CATTLE
EXCENEL Fluid suspension, 50mg/ml, ενέσιμο εναιώρημα για χοίρους και βοοειδή

Učinkovina:
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
govedo

Pot uporabe:
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
53.50 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Suspenzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Pig**
  - Meat and offal. 2 day

Subkutana uporaba:

- **govedo**
  - Meat and offal. 6 day
  - Milk. 0 day

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

QJ01DD90

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:

Cyprus

Opis ovojnine zdravila:

Na voljo samo v French
Na voljo samo v French
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

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 Hellas S.A.

Datum dovoljenja za promet z zdravilom:

28/04/2013

Proizvodna mesta, odgovorna za sproščanje serij:

Zoetis Belgium

Pristojni organ:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Številka dovoljenja za promet z zdravilom:

CY00362V

Datum spremembe statusa dovoljenja za promet:

12/04/2017

Referenčna država članica:

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

Številka postopka:

FR/V/0102/001

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 Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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

Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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). Spodaj ga najdete v drugem jeziku.