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ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS

Ima dovoljenje za promet

- Photobacterium damselae, subsp. piscicida, strain DI21, Inactivated
- Vibrio anguillarum, serotype O1, Inactivated
- Vibrio anguillarum, serotype O2b, Inactivated
- Vibrio anguillarum, serotype O2a, Inactivated

Informacije o zdravilu

Ime zdravila:
ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS

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

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

Pot uporabe:
Intraperitonealna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
60.00 Relative Percentage Survival / 1.00 Dose

Na voljo samo v English
75.00 Relative Percentage Survival / 1.00 Dose

Na voljo samo v English
75.00 Relative Percentage Survival / 1.00 Dose

Na voljo samo v English
75.00 Relative Percentage Survival / 1.00 Dose

Farmacevtska oblika:

Emulzija za injiciranje

Karenca glede na pot uporabe:

Intraperitonealna uporaba:

- **Seabass**
  - Fish meat. 0 degree day

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

QI10X

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 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 Hipra S.A.

Datum dovoljenja za promet z zdravilom:
4/05/2017

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

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
943/01/17DIVPT

Datum spremembe statusa dovoljenja za promet:
1/04/2022

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

Številka postopka:
FR/V/0314/001

Zadevne države članice:

Na voljo samo v Spanish Czech German Estonian English French Croatian 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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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.

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