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Tricaine Pharmaq 1000 mg/g Powder for Solution for Fish Treatment

Ima dovoljenje za promet

- Tricaine mesilate

Informacije o zdravilu

Ime zdravila:
Tricaine Pharmaq 1000 mg/g Powder for Solution for Fish Treatment

Učinkovina:
Na voljo samo v English

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

Pot uporabe:
Kopanje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

1000.00 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Prašek za raztopino za zdravljenje rib

Karenca glede na pot uporabe:
Kopanje:

- **Ornamental fish**
  - Fish meat. 70 degree day

Fish must not be slaughtered for human consumption during treatment. Fish can only be harvested for human consumption 70 degree days after the last treatment.

- **Other fish**
  - Fish meat. 70 degree day

Fish must not be slaughtered for human consumption during treatment. Fish can only be harvested for human consumption 70 degree days after the last treatment.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QN01AX93

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:
Italy

Opis ovojnine zdravila:
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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Pharmaq AS

Datum dovoljenja za promet z zdravilom:
22/03/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Pharmaq Limited

Pristojni organ:
Ministry Of Health

Številka dovoljenja za promet z zdravilom:
104497

Datum spremembe statusa dovoljenja za promet:
22/03/2013

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

Številka postopka:
NO/V/0012/001

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 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
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

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

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.