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AquaVac Relera Concentrate for Dip Suspension or Suspension for Injection for Rainbow Trout

Ima dovoljenje za promet

- Yersinia ruckeri, serotype O1, biotype 1, strain Hagerman, Inactivated
- Yersinia ruckeri, serotype O1, biotype 2 (EX5), strain SP/07/04, Inactivated

Informacije o zdravilu

Ime zdravila:
AquaVac Relera Concentrate for Dip Suspension or Suspension for Injection for Rainbow Trout

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

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

Pot uporabe:
Kopanje
Intraperitonealna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
75.00 Relative Percentage Survival / 1.00 millilitre(s)

Na voljo samo v English
75.00 Relative Percentage Survival / 1.00 millilitre(s)

Farmacevtska oblika:

Koncentrat za suspenzijo za kopel

Karenca glede na pot uporabe:

Kopanje:

- **Trout - Golden/Rainbow/Redband/Steelhead**
  - Meat. 0 degree day

Intraperitonealna uporaba:

- **Trout - Golden/Rainbow/Redband/Steelhead**
  - Meat. 0 degree day

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

QI10BB03

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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Na voljo v:

Denmark

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:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
9/07/2009

Proizvodna mesta, odgovorna za sproščanje serij:
MSD Animal Health UK Limited
Merck Sharp & Dohme Animal Health S.L.
The Veterinary Medicines Directorate

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
42645

Datum spremembe statusa dovoljenja za promet:
9/07/2009

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/0309/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian 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

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.

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.