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# Quinoflox 100 mg/ml solution for use in drinking water, chicken and rabbits

Ima dovoljenje za promet

- Enrofloxacin

## Informacije o zdravilu

**Ime zdravila:**
Quinoflox 100 mg/ml solution for use in drinking water, chicken and rabbits

**Učinkovina:**
Na voljo samo v English

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

**Pot uporabe:**
Peroralna uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**

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

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**Farmacevtska oblika:**

Peroralna raztopina

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**Karenca glede na pot uporabe:**

**Peroralna uporaba:**

- **Rabbit**
  - Meat and offal. 2 day
- **Chicken**
  - Meat and offal. 4 day

Eggs: Do not use in birds producing eggs for human consumption

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

QJ01MA90

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**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

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:**
Global Vet Health S.L.

**Datum dovoljenja za promet z zdravilom:**
2/02/2012

**Proizvodna mesta, odgovorna za sproščanje serij:**
S P Veterinaria S.A.

**Pristojni organ:**
Ministry Of Health

**Številka dovoljenja za promet z zdravilom:**
104332

**Datum spremembe statusa dovoljenja za promet:**
2/02/2012

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

**Številka postopka:**
PT/V/0144/001

**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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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

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.