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# Octacillin 800 mg/g powder for use in drinking water for pigs, amoxicillin trihydrate

Ima dovoljenje za promet

- Amoxicillin trihydrate

## Informacije o zdravilu

**Ime zdravila:**
Octacillin 800 mg/g powder for use in drinking water for pigs, amoxicillin trihydrate

**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

**Pot uporabe:**
Dajanje v vodo za pitje

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
800.00 milligram(s) / 1.00 gram(s)

**Farmacevtska oblika:**
Prašek za dajanje v vodo za pitje

**Karenca glede na pot uporabe:**
**Dajanje v vodo za pitje:**

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

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QJ01CA04

**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:**
France

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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:**
Eurovet Animal Health B.V.

**Datum dovoljenja za promet z zdravilom:**
6/12/2010

**Proizvodna mesta, odgovorna za sproščanje serij:**
Eurovet Animal Health B.V.

**Pristojni organ:**
French Agency For Food, Environmental And Occupational Health & Safety

**Številka dovoljenja za promet z zdravilom:**
FR/V/5516468 1/2010

**Datum spremembe statusa dovoljenja za promet:**
6/12/2015

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

**Številka postopka:**
NL/V/0367/001

**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
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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

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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.

Package Leaflet and Labelling

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