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# DFV DOXIVET 500 mg/g, powder for use in drinking water for pigs and chickens

Ima dovoljenje za promet

- Doxycycline hyclate

## Informacije o zdravilu

**Ime zdravila:**
DFV DOXIVET 500 mg/g, powder for use in drinking water for pigs and chickens

**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
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
500.00 milligram(s) / 1.00 gram(s)

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**Farmacevtska oblika:**

Prašek za dajanje v vodo za pitje

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**Karenca glede na pot uporabe:**

**Dajanje v vodo za pitje:**

- **Chicken**
  - Meat and offal. 12 day
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 4 day

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

QJ01AA02

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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
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:**
Divasa Farmavic S.A.

**Datum dovoljenja za promet z zdravilom:**
1/08/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Divasa Farmavic S.A.

**Pristojni organ:**
Norwegian Medical Products Agency

**Številka dovoljenja za promet z zdravilom:**
10-7848

**Datum spremembe statusa dovoljenja za promet:**
27/06/2016

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

**Številka postopka:**
NL/V/0152/002

**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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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.

Navodilo za uporabo

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