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Powdox 500 mg/g

Ima dovoljenje za promet

- Doxycycline hyclate

Informacije o zdravilu

Ime zdravila:
Powdox 500 mg/g

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
brojler
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 Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian
Na voljo samo v Bulgarian Spanish Danish Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
580.00 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:

- **Chicken (for reproduction)**
  - Meat and offal. 5 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption

- **brojler**
  - Meat and offal. 5 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption

- **Pig (for fattening)**
  - Meat and offal. 4 day
- **Turkey (for reproduction)**
  - Meat and offal. 12 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption

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Turkey (for meat production)

- Meat and offal. 12 day
- Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01AA02

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:
Germany

Opis ovojnine zdravila:
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:
Vetpharma Animal Health S.L.

Datum dovoljenja za promet z zdravilom:
5/12/2012

Proizvodna mesta, odgovorna za sproščanje serij:
Laboratorios Karizoo S.A.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401566.00.00

Datum spremembe statusa dovoljenja za promet:
22/02/2018

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/0177/001

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

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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Označevanje

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Combined File of all Documents

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