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MOXAPULVIS 500 mg/g powder for use in drinking water

Ima dovoljenje za promet

- Amoxicillin trihydrate

Informacije o zdravilu

Ime zdravila:
MOXAPULVIS 500 mg/g powder for use in drinking water

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
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
574.00 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:

- **Turkey**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 5 day

- **Chicken**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 1 day

- **Duck**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 9 day

- **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

Opis ovojnine zdravila:
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:
V.M.D.

Datum dovoljenja za promet z zdravilom:
19/03/2018

Proizvodna mesta, odgovorna za sproščanje serij:
V.M.D.

Pristojni organ:
Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:
REG NL 120573

Datum spremembe statusa dovoljenja za promet:
1/02/2022

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

Številka postopka:
IE/V/0541/001

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 Croatian 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.

Povzetek glavnih značilnosti zdravila

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