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# Pharmasin 1 g/g granules for use in drinking water for pigs, chickens, turkeys and calves

Ima dovoljenje za promet

- Tylosin tartrate

## Informacije o zdravilu

**Ime zdravila:**
Pharmasin 1 g/g granules for use in drinking water for pigs, chickens, turkeys and calves

**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
tele
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
1100.00 milligram(s) / 1.10 gram(s)

**Farmacevtska oblika:**

Zrnca za dajanje v vodo za pitje

**Karenca glede na pot uporabe:**

**Dajanje v vodo za pitje:**

- **Pig**
  - Meat and offal. 1 day
- **Chicken**
  - Meat and offal. 1 day
  - Eggs. no withdrawal period zero days
- **tele**
  - Meat and offal. 12 day
- **Turkey**
  - Meat and offal. 2 day
  - Eggs. no withdrawal period zero days

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

QJ01FA90

**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:**
HuVepharma

**Datum dovoljenja za promet z zdravilom:**
29/03/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Biovet AD

**Pristojni organ:**
Federal Office Of Consumer Protection And Food Safety

**Številka dovoljenja za promet z zdravilom:**
401522.00.00

**Datum spremembe statusa dovoljenja za promet:**
26/02/2016

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

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**Številka postopka:**
NL/V/0159/001

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**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

Combined File of all Documents

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