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Tylo-Suscit 100% Kompaktat 1000 mg/g Granulat für Schweine, Hühner und Puten

Ima dovoljenje za promet

- Tylosin tartrate

Informacije o zdravilu

Ime zdravila:
Tylo-Suscit 100% Kompaktat 1000 mg/g Granulat für Schweine, Hühner und Puten

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 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 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
1000.00 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Zrnca za dajanje v vodo za pitje

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

- **Turkey**
  - Meat and offal. 5 day
- **Chicken (for reproduction)**
  - Meat and offal. 1 day
  - Egg. 0 day
- **Chicken (chick)**
  - Meat and offal. 2 day
- **Chicken (pullet)**
  - Meat and offal. 1 day
- **Pig**

- Meat and offal. 1 day

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 German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
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Na voljo samo v German
Na voljo samo v German
Na voljo samo v German

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

Imetnik dovoljenja za promet z zdravilom:
Bela-Pharm GmbH & Co. KG

Datum dovoljenja za promet z zdravilom:
13/12/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Bela-Pharm GmbH & Co. KG

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401875.00.00

Datum spremembe statusa dovoljenja za promet:
16/01/2020

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.

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