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COCCIBAL 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS

Ima dovoljenje za promet

- Amprolium hydrochloride

Informacije o zdravilu

Ime zdravila:
COCCIBAL 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR CHICKENS AND TURKEYS

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 Spanish Czech Danish Estonian English French Italian Latvian Lithuanian Romanian Finnish Swedish Norwegian
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

Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
226.20 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za dajanje v vodo za pitje

Karenca glede na pot uporabe:
Peroralna uporaba:

- **Turkey**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day
- **Chicken (for reproduction)**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **brojler**
  - Meat and offal. 0 day
-

Chicken (layer hen)

- Meat and offal. 0 day
- Eggs. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP51BX02

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:
Denmark

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:

S P Veterinaria S.A.

Datum dovoljenja za promet z zdravilom:
25/04/2012

Proizvodna mesta, odgovorna za sproščanje serij:
S P Veterinaria S.A.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
47846

Datum spremembe statusa dovoljenja za promet:
25/04/2012

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

Številka postopka:
FR/V/0230/001

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
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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.

Combined File of all Documents

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