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ORONDO SPRAY 20 mg/ml SUSPENSION PARA PULVERIZACION CUTANEA

Ima dovoljenje za promet

- Chlortetracycline hydrochloride

Informacije o zdravilu

Ime zdravila:
ORONDO SPRAY 20 mg/ml SUSPENSION PARA PULVERIZACION CUTANEA

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
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
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Na voljo samo v Bulgarian Spanish Czech Danish Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Swedish Icelandic

Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Dermalno pršilo, suspenzija

Karenca glede na pot uporabe:
Dermalna uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day

- Milk. 0 day

- **Horse**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

- **Birds**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD06AA02

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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Na voljo v:
Spain

Opis ovojnine zdravila:
Na voljo samo v Spanish

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Industrial Veterinaria S.A.

Datum dovoljenja za promet z zdravilom:
27/01/1992

Proizvodna mesta, odgovorna za sproščanje serij:
Divasa Farmavic S.A.

Pristojni organ:
Spanish Agency Of Medicines And Medical Devices

Številka dovoljenja za promet z zdravilom:
271 ESP

Datum spremembe statusa dovoljenja za promet:
27/01/1992

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Navodilo za uporabo

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

Označevanje

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.