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VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

Ima dovoljenje za promet

- Cefquinome sulfate

Informacije o zdravilu

Ime zdravila:
VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Norwegian

Pot uporabe:
Intramamarno dajanje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
177.80 milligram(s) / 1.00 Brizga

Farmacevtska oblika:
Intramamarno mazilo

Karenca glede na pot uporabe:

Intramamarno dajanje:

- **Cattle (dry cow)**
  - Meat and offal. 2 day
  - Milk. 36 day

Milk: 36 days after treatment when dry period is 5 weeks or less.

- Milk. 1 day Milk: 1 day after calving when dry period is more than 5 weeks

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

QJ51DE90

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

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Na voljo v:

Austria

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Virbac

Datum dovoljenja za promet z zdravilom:
13/01/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Haupt Pharma Latina S.r.l.
Virbac

Pristojni organ:
Austrian Agency For Health And Food Safety

Številka dovoljenja za promet z zdravilom:
8-00633

Datum spremembe statusa dovoljenja za promet:
13/01/2005

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

Številka postopka:
FR/V/0148/001

Zadevne države članice:

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

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.

Povzetek glavnih značilnosti zdravila

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.