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CYDECTINE 10% LA SOLUTION FOR INJECTION FOR CATTLE

Ima dovoljenje za promet

- Moxidectin

Informacije o zdravilu

Ime zdravila:
CYDECTINE 10% LA SOLUTION FOR INJECTION FOR CATTLE

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo

Pot uporabe:
Subkutana uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:

Subkutana uporaba:

- **govedo**
  - Meat and offal. 108 day
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals which are intended to produce milk for human consumption within 80 days of expected parturition.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP54AB02

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:
Germany

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Deutschland GmbH

Datum dovoljenja za promet z zdravilom:
18/11/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Manufacturing & Research Spain S.L.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
400868.00.00

Datum spremembe statusa dovoljenja za promet:
23/12/2010

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

Številka postopka:
FR/V/0155/001

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

Combined File of all Documents

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