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DECTOSPOT 10 MG/ML POUR-ON SOLUTION FOR CATTLE AND SHEEP

Ima dovoljenje za promet

- Deltamethrin

Informacije o zdravilu

Ime zdravila:
DECTOSPOT 10 MG/ML POUR-ON SOLUTION FOR CATTLE AND SHEEP

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

Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

10.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Kožni poliv, raztopina

Karenca glede na pot uporabe:
Dermalna uporaba:

- **govedo**
  - Meat and offal. 18 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 24 hour

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QP53AC11

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 French
Na voljo samo v French
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

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Bimeda Animal Health Limited

Datum dovoljenja za promet z zdravilom:
26/05/2016

Proizvodna mesta, odgovorna za sproščanje serij:
Bimeda Animal Health Limited

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
55689

Datum spremembe statusa dovoljenja za promet:
26/05/2016

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

Številka postopka:
FR/V/0293/001

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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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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