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ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION

Ima dovoljenje za promet

- Bismuth subnitrate, heavy

Informacije o zdravilu

Ime zdravila:
ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION

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
2.60 gram(s) / 1.00 Brizga

Farmacevtska oblika:
intramamarna suspenzija

Karenca glede na pot uporabe:

Intramamarno dajanje:

- **Cattle (dry cow)**
  - Meat and offal. 0 day
  - Milk. 0 day

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

QG52X

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:

Denmark

Opis ovojnine zdravila:

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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Animal Health ApS

Datum dovoljenja za promet z zdravilom:
22/09/2003

Proizvodna mesta, odgovorna za sproščanje serij:
Cross Vetpharm Group Limited
Haupt Pharma Latina S.r.l.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
34489

Datum spremembe statusa dovoljenja za promet:
22/09/2003

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

Številka postopka:
FR/V/0341/001

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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