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FATROSEAL 2.6 g INTRAMAMMARY SUSPENSION FOR DRY COWS

Ima dovoljenje za promet

- Bismuth subnitrate, heavy

Informacije o zdravilu

Ime zdravila:
FATROSEAL 2.6 g INTRAMAMMARY SUSPENSION FOR DRY COWS

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Spanish Czech Danish Estonian English French Italian Latvian Lithuanian Romanian Finnish Norwegian

Pot uporabe:
Intramamarno dajanje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
26.00 gram(s) / 4.00 gram(s)

Farmacevtska oblika:

intramamarna suspenzija

Karenca glede na pot uporabe:

Intramamarno dajanje:

- **Cattle (dairy cow at drying-off)**
  - Meat and offal. 0 day
  - Milk. 0 hour

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

Na voljo v:

Greece

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
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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Fatro S.p.A.

Datum dovoljenja za promet z zdravilom:
6/04/2022

Proizvodna mesta, odgovorna za sproščanje serij:
Fatro S.p.A.

Pristojni organ:
National Organization For Medicines

Številka dovoljenja za promet z zdravilom:
36008/07-04-2022/K-0248701

Datum spremembe statusa dovoljenja za promet:
23/06/2022

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

Številka postopka:
ES/V/0407/001

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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Na voljo samo v Estonian English French Lithuanian Portuguese 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.

Navodilo za uporabo

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

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