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Source URL: https://medicines.health.europa.eu/veterinary/sl/600000092708
FATROSEAL 2.6 g INTRAMAMMARY SUSPENSION FOR DRY COWS

- Bismuth subnitrate, heavy

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
FATROSEAL 2.6 g INTRAMAMMARY SUSPENSION FOR DRY COWS
U?inkovina:

- 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

U?inkovina / 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-kemi?na veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

- QG52X

Pravni status za dobavo/izdajo zdravila:

- Zdravilo, ki se izdaja brez veterinarskega recepta

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

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:

- 14/03/2022

Proizvodna mesta, odgovorna za sproš?anje serij:

- Fatro S.p.A.

Pristojni organ:

- French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:

- FR/V/6891161 8/2022

Datum spremembe statusa dovoljenja za promet:

- 14/03/2022

Referen?na država ?lanica:

- 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 ?lanice:

- 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
- Na voljo samo v Spanish Czech German Estonian English French Italian Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Package Leaflet and Labelling

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
French (PDF)
Objavljeno na dan: 15/10/2025
Updated on: 13/03/2026
Prenesi

Povzetek glavnih zna?ilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
French (PDF)
Objavljeno na dan: 7/05/2025
Updated on: 13/03/2026
Prenesi

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
English (PDF)
Objavljeno na dan: 14/03/2026
Prenesi

Javno poro?ilo o oceni zdravila

eu-PUAR-esv0407001-dcp-fatroseal-2.6-g-en.pdf

English (PDF)
Objavljeno na dan: 14/03/2026
Prenesi