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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:**

Germany

**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:**
30/03/2022

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

**Pristojni organ:**
Federal Office Of Consumer Protection And Food Safety

**Številka dovoljenja za promet z zdravilom:**
V7004521.00.00

**Datum spremembe statusa dovoljenja za promet:**
30/03/2022

**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

Combined File of all Documents

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

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