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# Ubrostar Red 100 mg / 280 mg / 100 mg Intramammary Suspension for cattle

Ima dovoljenje za promet

- Framycetin sulfate
- Benethamine penicillin
- Penethamate hydriodide

## Informacije o zdravilu

**Ime zdravila:**
Ubrostar Red 100 mg / 280 mg / 100 mg Intramammary Suspension for cattle

**Učinkovina:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

**Ciljne živalske vrste:**
govedo

**Pot uporabe:**
Intramamarno dajanje

## Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
100.00 milligram(s) / 1.00 Brizga

Na voljo samo v English
280.00 milligram(s) / 1.00 Brizga

Na voljo samo v English
100.00 milligram(s) / 1.00 Brizga

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**Farmacevtska oblika:**

intramamarna suspenzija

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**Karenca glede na pot uporabe:**

**Intramamarno dajanje:**

- **govedo**
  - Meat and offal. 10 day
  - Milk. 37 day

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

QJ51RC25

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**

Valid

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**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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**Opis ovojnine zdravila:**

Na voljo samo v English

Na voljo samo v English

Na voljo samo v English

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## 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:**
Boehringer Ingelheim Vetmedica GmbH

**Datum dovoljenja za promet z zdravilom:**
26/10/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Lohmann Pharma Herstellung GmbH
Haupt Pharma Latina S.r.l.

**Pristojni organ:**
Directorate Of Veterinary Medicinal Products

**Številka dovoljenja za promet z zdravilom:**
3013/X/2011 MgSzH ÁTI

**Datum spremembe statusa dovoljenja za promet:**
26/10/2011

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

**Številka postopka:**
IE/V/0271/001

**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
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian 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.