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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

Farmacevtska oblika:

intramamarna suspenzija

Karenca glede na pot uporabe:

Intramamarno dajanje:

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

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

QJ51RC25

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:

France

Opis ovojnine zdravila:

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:
Boehringer Ingelheim Animal Health France

Datum dovoljenja za promet z zdravilom:
19/10/2011

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

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/8001032 8/2011

Datum spremembe statusa dovoljenja za promet:
18/01/2017

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.

Package Leaflet and Labelling

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