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OSTEOCYNESINE VETERINAIRE COMPRIMES

Ima dovoljenje za promet

- CALCAREA CARBONICA OSTREARUM D2
- Sulfur iodatum C3
- Calcium fluoratum D2
- CALCAREA PHOSPHORICA D2

Informacije o zdravilu

Ime zdravila:
OSTEOCYNESINE VETERINAIRE COMPRIMES

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

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Danish German Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Swedish Icelandic

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Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
26.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
2.60 milligram(s) / 1.00 Tableta

Na voljo samo v English
26.00 milligram(s) / 1.00 Tableta

Na voljo samo v English
26.00 milligram(s) / 1.00 Tableta

Farmacevtska oblika:
Tableta

Karenca glede na pot uporabe:
Peroralna uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 day
-

Pig

- Meat and offal. 0 day
- Milk. 0 day

•

Equid

- Meat and offal. 0 day
- Milk. 0 day

•

Sheep

- Meat and offal. 0 day
- Milk. 0 day

•

Goat

- Meat and offal. 0 day
- Milk. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QV03AX

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

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Na voljo v:
France

Opis ovojnine zdravila:
Na voljo samo v French

Na voljo samo v French

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:
Boiron

Datum dovoljenja za promet z zdravilom:
5/07/1983

Proizvodna mesta, odgovorna za sproščanje serij:
Boiron

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

Številka dovoljenja za promet z zdravilom:
FR/V/3389423 0/1983

Datum spremembe statusa dovoljenja za promet:
5/07/2008

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.