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# Florfenikel 300 mg/ml solution for injection for cattle and pigs

Ima dovoljenje za promet

- Florfenicol

## Informacije o zdravilu

**Ime zdravila:**
Florfenikel 300 mg/ml solution for injection for cattle and pigs

**Učinkovina:**
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

**Pot uporabe:**
intramuskularna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
300.00 milligram(s) / 1.00 millilitre(s)

**Farmacevtska oblika:**
Raztopina za injiciranje

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**Karenca glede na pot uporabe:**
**intramuskularna uporaba:**

- **govedo**
  - Meat and offal. 34 day .
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption

- **Pig**
  - Meat and offal. 18 day .

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QJ01BA90

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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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**Na voljo v:**
Denmark

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**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

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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 French Italian Latvian Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
KELA Kempisch Laboratorium Kela Laboratoria

**Datum dovoljenja za promet z zdravilom:**
2/02/2012

**Proizvodna mesta, odgovorna za sproščanje serij:**
S P Veterinaria S.A.
KELA Kempisch Laboratorium Kela Laboratoria

**Pristojni organ:**
Danish Medicines Agency

**Številka dovoljenja za promet z zdravilom:**
47459

**Datum spremembe statusa dovoljenja za promet:**
2/02/2012

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

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**Številka postopka:**
BE/V/0038/001

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**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 Greek 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

**Generic of:**
600000049571

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.

Combined File of all Documents

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

Označevanje

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Navodilo za uporabo

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