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# Qivitan 25 mg/ml Suspension for Injection for Cattle and Pigs

Ima dovoljenje za promet

- Cefquinome sulfate

## Informacije o zdravilu

**Ime zdravila:**
Qivitan 25 mg/ml Suspension 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
29.64 milligram(s) / 1.00 millilitre(s)

**Farmacevtska oblika:**
Suspenzija za injiciranje

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

- **govedo**
  - Meat and offal. 5 day
  - Milk. 24 hour
- **Pig**
  - Meat and offal. 3 day

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QJ01DE90

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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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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**Na voljo v:**
United Kingdom (Northern Ireland)

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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
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:**
Industrial Veterinaria S.A.

**Datum dovoljenja za promet z zdravilom:**
2/02/2017

**Proizvodna mesta, odgovorna za sproščanje serij:**
aniMedica GmbH
Industrial Veterinaria S.A.

**Pristojni organ:**
The Veterinary Medicines Directorate

**Številka dovoljenja za promet z zdravilom:**
Vm 36547/3009

**Datum spremembe statusa dovoljenja za promet:**
7/03/2023

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

**Številka postopka:**
IE/V/0479/001

**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 Croatian 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

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.

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