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# Unisol 100 mg/ml solution for injection for cattle and pigs

Ima dovoljenje za promet

- Enrofloxacin

## Informacije o zdravilu

**Ime zdravila:**
Unisol 100 mg/ml solution for injection for cattle and pigs

**Učinkovina:**
Na voljo samo v English

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

**Pot uporabe:**
intramuskularna uporaba
Intravenska uporaba
Subkutana uporaba

## Podatki o zdravilu

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

100.00 milligram(s) / 1.00 millilitre(s)

**Farmacevtska oblika:**
Raztopina za injiciranje

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

- **Pig**
  - Meat and offal. 13 day

**Intravenska uporaba:**

- **govedo**
  - Meat and offal. 5 day
  - Milk. 3 day

**Subkutana uporaba:**

- **govedo**
  - Meat and offal. 12 day
  - Milk. 4 day

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QJ01MA90

**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

**Na voljo v:**

Ireland

**Opis ovojnine zdravila:**
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:**
Vetpharma Animal Health S.L.

**Datum dovoljenja za promet z zdravilom:**
6/05/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Industrial Veterinaria S.A.
Chemical Iberica Productos Veterinarios S.L.

**Pristojni organ:**
Health Products Regulatory Authority

**Številka dovoljenja za promet z zdravilom:**
VPA10516/001/001

**Datum spremembe statusa dovoljenja za promet:**
6/05/2011

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

**Številka postopka:**
IE/V/0255/001

**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 Greek 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
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Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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.