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# Parofor 70000 IU/g Powder for use in drinking water/milk

Ima dovoljenje za promet

- Paromomycin

## Informacije o zdravilu

**Ime zdravila:**
Parofor 70000 IU/g Powder for use in drinking water/milk

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

**Pot uporabe:**
Dajanje v vodo za pitje/mleko

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
70000.00 international unit(s) / 1.00 gram(s)

**Farmacevtska oblika:**
Prašek za dajanje v vodo za pitje/mleko

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**Karenca glede na pot uporabe:**
**Dajanje v vodo za pitje/mleko:**

- **Pig**
  - Meat and offal. 3 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 20 day

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QA07AA06

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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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## 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:**
HuVepharma

**Datum dovoljenja za promet z zdravilom:**
5/09/2014

**Proizvodna mesta, odgovorna za sproščanje serij:**
Biovet AD

**Pristojni organ:**
Danish Medicines Agency

**Številka dovoljenja za promet z zdravilom:**
52244

**Datum spremembe statusa dovoljenja za promet:**
5/09/2014

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

**Številka postopka:**
BE/V/0027/001

**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

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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

**Generic of:**
600000085987

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.

Navodilo za uporabo

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