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# Nanotrim 464.2 mg/g + 100 mg/g powder for use in drinking water/milk for chickens, turkeys, pigs and cattle

Ima dovoljenje za promet

- Sulfachlorpyridazine sodium
- Trimethoprim

## Informacije o zdravilu

**Ime zdravila:**
Nanotrim 464.2 mg/g + 100 mg/g powder for use in drinking water/milk for chickens, turkeys, pigs and cattle

**Učinkovina:**
Na voljo samo v English
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 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 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
500.00 milligram(s)/gram / 1.00 gram(s)

Na voljo samo v English
100.00 milligram(s)/gram / 1.00 gram(s)

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

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

- **Pig**
  - Meat and offal. 7 day
- **Chicken**
  - Meat and offal. 3 day
- **Turkey**
  - Meat and offal. 9 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 7 day

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QJ01EW12

**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 German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian

**Opis ovojnine zdravila:**
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 Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
HuVepharma

**Datum dovoljenja za promet z zdravilom:**
24/02/2025

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

**Pristojni organ:**
State Agency Of Medicines

**Številka dovoljenja za promet z zdravilom:**
1189225

**Datum spremembe statusa dovoljenja za promet:**
24/02/2025

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

**Številka postopka:**
NL/V/0420/001

**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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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

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