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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

Atļautas

- Sulfachlorpyridazine sodium
- Trimethoprim

Zāļu identifikācija

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

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Cūka
Cālis
Tītars
Liellops (teļš pirms atgremošanas funkcijas sākšanās)

Lietošanas veids:
lietošanai ar dzeramo ūdeni/ pienu

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
500.00 miligrams (i)/gramā / 1.00 grams(i)

Pieejamas tikai English
100.00 miligrams (i)/gramā / 1.00 grams(i)

Farmaceitiskā forma:

Pulveris lietošanai ar dzeramo ūdeni/pienu

Ierobežojumu periods pēc lietošanas veida:
lietošanai ar dzeramo ūdeni/ pienu:

- **Cūka**
  - Gaļa un blakusprodukti. 7 diena
- **Cālis**
  - Gaļa un blakusprodukti. 3 diena
- **Tītars**
  - Gaļa un blakusprodukti. 9 diena
- **Liellops (teļš pirms atgremošanas funkcijas sākšanās)**
  - Gaļa un blakusprodukti. 7 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QJ01EW12

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:
Pieejamas tikai Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums hibrīdām veterinārajām zālēm (ES Regulas 2019/6 19. panta 1. daļa)

Tirdzniecības atļaujas turētājs:
HuVepharma

Tirdzniecības atļaujas datums:
27/02/2025

Ražošanas vietas sēriju izlaidei:
Biovet AD

Atbildīgā iestāde:
Spanish Agency Of Medicines And Medical Devices

Atļaujas numurs:
4379 ESP

Atļaujas statusa maiņas datums:
28/02/2025

Atsauces dalībvalsts:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Procedūras numurs:
NL/V/0420/001

Attiecīgās dalībvalstis:

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejamas tikai Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejamas tikai Spanish Czech German Estonian Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Čehija

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Zāļu apraksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Lietošanas instrukcija

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Marķējuma teksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Visu dokumentu apvienotais fails

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.