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DOPHEXINE 20 MG/G POWDER FOR USE IN DRINKING WATER/MILK

Atļautas

- Bromhexine hydrochloride

Zāļu identifikācija

Zāļu nosaukums:
DOPHEXINE 20 MG/G POWDER FOR USE IN DRINKING WATER/MILK

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Tītars
Cūka
Pīle
Cālis
Vista (broileris)
Liellops

Lietošanas veids:
Iekšķīgai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English

20.00 miligrams(i) / 1.00 grams(i)

Farmaceitiskā forma:
Pulveris lietošanai ar dzeramo ūdeni/pienu

Ierobežojumu periods pēc lietošanas veida:
Iekšķīgai lietošanai:

- **Tītars**
  - Gaļa un blakusprodukti. 0 diena
  - Olas. no withdrawal period

Not for use in birds producing eggs for human consumption, during and 4 weeks before the laying period.

- **Cūka**
  - Gaļa un blakusprodukti. 0 diena

- **Pīle**
  - Gaļa un blakusprodukti. 0 diena
  - Olas. no withdrawal period

Not for use in birds producing eggs for human consumption, during and 4 weeks before the laying period.

- **Cālis**
  - Olas. no withdrawal period

Not for use in birds producing eggs for human consumption, during and 4 weeks before the laying period.

- **Vista (broileris)**
  - Gaļa un blakusprodukti. 0 diena

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Liellops

- Gaļa un blakusprodukti. 2 diena
- Piens. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QR05CB02

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

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

Pieejams:
Lithuania

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums hibrīdām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(3) pants)

Tirdzniecības atļaujas turētājs:
Dopharma Research B.V.

Tirdzniecības atļaujas datums:
16/08/2020

Ražošanas vietas sēriju izlaidei:
Dopharma B.V.

Atbildīgā iestāde:
State Food And Veterinary Service

Atļaujas numurs:
LT/2/20/2613/001-005

Atļaujas statusa maiņas datums:
23/11/2025

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

Procedūras numurs:
FR/V/0391/001

Attiecīgās dalībvalstis:
Pieejamas tikai Spanish Czech German Estonian English French 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

Visu dokumentu apvienotais fails

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

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