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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

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
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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:
17/11/2021

Ražošanas vietas sēriju izlaidei:
Dopharma B.V.

Atbildīgā iestāde:
Swedish Medical Products Agency

Atļaujas numurs:
62015

Atļaujas statusa maiņas datums:
17/11/2021

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

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.

Visu dokumentu apvienotais fails

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.

Apvienotais marķējuma un lietošanas instrukcijas teksts

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

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