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Halagon 0.5 mg/ml - Oral solution

Atļautas

- Halofuginone

Zāļu identifikācija

Zāļu nosaukums:
Halagon 0.5 mg/ml - Oral solution

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops (jaundzimis teļš)

Lietošanas veids:
Iekšķīgai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
0.50 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:
šķīdums iekšķīgai lietošanai

Ierobežojumu periods pēc lietošanas veida:
Iekšķīgai lietošanai:

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Liellops (jaundzimis teļš)

- Gaļa un blakusprodukti. 13 diena 13 days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QP51AX08

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:

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Pieejams:
Austria , Belgium , Cyprus , Denmark , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania , Spain , United Kingdom (Northern Ireland) , Čehija

Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:
Emdoka

Tirdzniecības atļaujas datums:
13/12/2016

Ražošanas vietas sēriju izlaidei:
Divasa Farmavic S.A.

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
13/12/2016

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

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 11/05/2026
Lejupielādēt

ema-puar-halagon-v-4201-par-en.pdf