File downloaded on 2026-11-16
Source URL: https://medicines.health.europa.eu/veterinary/en/600000042329
TILDREN 5 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR HORSES

- Tiludronate disodium hemihydrate

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
TILDREN 5 MG/ML POWDER AND SOLVENT FOR SOLUTION FOR INJECTION FOR HORSES
U?inkovina:

- 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

Pot uporabe:

- Intravenska uporaba

Podatki o zdravilu

U?inkovina / Jakost:

- Na voljo samo v English
  5.83
  milligram(s)
  /
  1.00
  millilitre(s)

Farmacevtska oblika:

- Prašek in vehikel za raztopino za injiciranje

Karenca glede na pot uporabe:

- Intravenska uporaba
  - Horse
    - Meat and offal
      0
      day
    - Milk
      no withdrawal period

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

Anatomsko-terapevtsko-kemi?na veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

- QM05BA05

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

Na voljo v:

- Norway

Opis ovojnine zdravila:

- Na voljo samo v French
- Na voljo samo v French
- Na voljo samo v French

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

- Audevard

Datum dovoljenja za promet z zdravilom:

- 1/08/2003

Proizvodna mesta, odgovorna za sproš?anje serij:

- Ceva Sante Animale

Pristojni organ:

- Norwegian Medical Products Agency

Številka dovoljenja za promet z zdravilom:

- 02-1588

Datum spremembe statusa dovoljenja za promet:

- 9/01/2007

Referen?na država ?lanica:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:

- FR/V/0134/001

Zadevne države ?lanice:

- 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 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 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

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Povzetek glavnih zna?ilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.

Drugi jeziki (1)
Norwegian (PDF)
Objavljeno na dan: 10/10/2024
Updated on: 14/03/2026
Prenesi

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
Norwegian (PDF)
Objavljeno na dan: 10/10/2024
Updated on: 14/03/2026
Prenesi

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
English (PDF)
Objavljeno na dan: 14/03/2026
Prenesi