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Stullmisan vet. Pulver

Ima dovoljenje za promet

- Abies sibirica extract

Informacije o zdravilu

Ime zdravila:
Stullmisan vet. Pulver

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

30.56 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Peroralni prašek

Karenca glede na pot uporabe:
Peroralna uporaba:

- **Quail**
  - Egg. 0 day
  - Meat and offal. 1 day
- **Turkey**
  - Egg. 0 day
  - Meat and offal. 1 day
- **govedo**
  - Meat and offal. 1 day
  - Milk. 1 day
- **Chicken (for reproduction)**
  - Meat and offal. 1 day
  - Egg. 0 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 1 day
- **Sheep**
  - Meat and offal. 1 day
  - Milk. 1 day
-

Horse

- Meat and offal. 1 day
- Milk. 1 day

•

Pig (suckling piglet)

- Meat and offal. 1 day

•

Horse (foal)

- Meat and offal. 1 day

•

Pig (piglet)

- Meat and offal. 1 day

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

QA07

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

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

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Opis ovojnine zdravila:

Na voljo samo v German

Na voljo samo v German

Na voljo samo v German

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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Pharma Stulln GmbH

Datum dovoljenja za promet z zdravilom:
21/06/2010

Proizvodna mesta, odgovorna za sproščanje serij:
Pharma Stulln GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401262.00.00

Datum spremembe statusa dovoljenja za promet:
29/09/2015

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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