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Nekrolyt Salbe für Pferde, Rinder, Schweine, Schafe, Ziegen, Hunde und Katzen

Ima dovoljenje za promet

- all-rac-alfa-Tocopheryl acetate
- Trypsin
- Retinol palmitate
- Chymotrypsin
- Papain

Informacije o zdravilu

Ime zdravila:
Nekrolyt Salbe für Pferde, Rinder, Schweine, Schafe, Ziegen, Hunde und Katzen

Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek 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
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Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
25.00 milligram(s) / 10.00 gram(s)

Na voljo samo v English
258.00 FIP / 10.00 gram(s)

Na voljo samo v English
42500.00 international unit(s) / 10.00 gram(s)

Na voljo samo v English
510.00 FIP / 10.00 gram(s)

Na voljo samo v English
27.75 FIP / 10.00 gram(s)

Farmacevtska oblika:
Mazilo

Karenca glede na pot uporabe:

Dermalna uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD03BA

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

Opis ovojnine zdravila:
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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Veyx Pharma GmbH

Datum dovoljenja za promet z zdravilom:
29/11/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Veyx Pharma GmbH

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
6383233.00.00

Datum spremembe statusa dovoljenja za promet:
29/11/2005

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

Dokumenti

Combined File of all Documents

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