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Sulphix

Ima dovoljenje za promet

- Trimethoprim
- Sulfadoxine

Informacije o zdravilu

Ime zdravila:
Sulphix

Učinkovina:
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
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

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:
Intravenska uporaba
Subkutana uporaba
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
40.00 milligram(s) / 1.00 millilitre(s)

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Intravenska uporaba:

- **govedo**
  - Milk. 4 day
  - Meat and offal. 4 day
- **Goat**
  - Milk. 4 day
  - Meat and offal. 4 day
- **Horse**
  - Meat and offal. 4 day

- Milk. no withdrawal period

Nicht bei Stuten anwenden, deren Milch fur den menschlichen Verzehr vorgesehen ist.

- **Pig**
  - Meat and offal. 5 day

intramuskularna uporaba:

- **govedo**
  - Milk. 4 day
  - Meat and offal. 11 day
- **Goat**
  - Meat and offal. 11 day
  - Milk. 4 day
- **Pig**
  - Meat and offal. 14 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01EW13

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

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Na voljo v:

Germany

Opis ovojnine zdravila:
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
Na voljo samo v German
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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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Bela-Pharm GmbH & Co. KG

Datum dovoljenja za promet z zdravilom:
17/09/2003

Proizvodna mesta, odgovorna za sproščanje serij:
Bela-Pharm GmbH & Co. KG

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
6856907.00.00

Datum spremembe statusa dovoljenja za promet:
17/09/2003

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.