Lactetrol, infusievloeistof

Ima dovoljenje za promet

- Calcium chloride
- Sodium lactate
- Sodium chloride
- Magnesium chloride
- Potassium chloride

Informacije o zdravilu

Ime zdravila:
Lactetrol, infusievloeistof

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 Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish 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
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:
Intraperitonealna uporaba
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

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

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

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

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

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

Farmacevtska oblika:
Raztopina za infundiranje

Karenca glede na pot uporabe:
Intraperitonealna uporaba:

-

govedo

- Milk. no withdrawal period — Withdrawal period = zero days
- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Goat (adult female)

- Milk. no withdrawal period — Withdrawal period = zero days
- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Sheep

- Milk. no withdrawal period — Withdrawal period = zero days
- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Horse

- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Pig

- Meat and offal. no withdrawal period — Withdrawal period = zero days

Intravenska uporaba:

•

govedo

- Milk. no withdrawal period — Withdrawal period = zero days
- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Goat (adult female)

- Milk. no withdrawal period — Withdrawal period = zero days

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Sheep

- Milk. no withdrawal period Withdrawal period = zero days

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Horse

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Pig

- Meat and offal. no withdrawal period Withdrawal period = zero days

Subkutana uporaba:

•

govedo

- Milk. no withdrawal period Withdrawal period = zero days

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Goat (adult female)

- Milk. no withdrawal period Withdrawal period = zero days

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Sheep

- Milk. no withdrawal period Withdrawal period = zero days

- Meat and offal. no withdrawal period Withdrawal period = zero days

•

Horse

- Meat and offal. no withdrawal period — Withdrawal period = zero days

•

Pig

- Meat and offal. no withdrawal period — Withdrawal period = zero days

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Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QB05BB01

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:
Netherlands

Opis ovojnine zdravila:
Na voljo samo v Dutch

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Eurovet Animal Health B.V.

Datum dovoljenja za promet z zdravilom:
22/06/1992

Proizvodna mesta, odgovorna za sproščanje serij:
Eurovet Animal Health B.V.

Pristojni organ:
Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:
REG NL 1297

Datum spremembe statusa dovoljenja za promet:
14/11/2008

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.