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GLUCADEX 2 MG/ML SOLUTION FOR INJECTION FOR HORSES, CATTLE, GOATS, PIGS, DOGS AND CATS

Ni odobreno

- DEXAMETHASONE DISODIUM PHOSPHATE

Informacije o zdravilu

Ime zdravila:
GLUCADEX 2 MG/ML SOLUTION FOR INJECTION FOR HORSES, CATTLE, GOATS, PIGS, DOGS AND CATS

Učinkovina:
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
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Pot uporabe:
intramuskularna uporaba
Subkutana uporaba
Periartikularna uporaba
Intravenska uporaba
Intraartikularna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 8 day
  - Milk. 72 hour
- **Pig**
  - Meat and offal. 2 day
- **Horse**

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

- **Goat**
  - Meat and offal. 8 day
  - Milk. 72 hour

Periartikularna uporaba:

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

Intravenska uporaba:

- **govedo**
  - Meat and offal. 8 day
  - Milk. 72 hour

- **Pig**
  - Meat and offal. 6 day

- **Horse**
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

- Meat and offal. 8 day

- **Goat**

- Meat and offal. 8 day

- Milk. 72 hour

Intraartikularna uporaba:

- **Horse**

  - Meat and offal. 8 day

  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QH02AB02

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:
Revoked

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Opis ovojnine zdravila:
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

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

Imetnik dovoljenja za promet z zdravilom:
Kepro B.V.

Datum dovoljenja za promet z zdravilom:
18/12/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Kepro B.V.

Pristojni organ:
Spanish Agency For Medicines And Medical Devices

Številka dovoljenja za promet z zdravilom:
3848 ESP

Datum spremembe statusa dovoljenja za promet:
19/06/2024

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
FR/V/0374/001

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.

Označevanje

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

Navodilo za uporabo

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

Combined File of all Documents

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

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