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Sensiblex 40 mg/ml solution for injection for cattle

Ima dovoljenje za promet

- Denaverine hydrochloride

Informacije o zdravilu

Ime zdravila:
Sensiblex 40 mg/ml solution for injection for cattle

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Romanian Finnish Swedish Norwegian
telica

Pot uporabe:
intramuskularna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Cattle (dairy cow)**
  - Milk. 24 hour
  - Meat and offal. 1 day
- **telica**
  - Milk. 24 hour
  - Meat and offal. 1 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QG02CX90

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

Na voljo v:
Germany

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English

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:
Veyx Pharma GmbH

Datum dovoljenja za promet z zdravilom:
26/05/2017

Proizvodna mesta, odgovorna za sproščanje serij:
Veyx Pharma GmbH
Veyx-Pharma B.V.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
402298.00.00

Datum spremembe statusa dovoljenja za promet:
20/06/2022

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

Številka postopka:
DE/V/0163/001

Zadevne države članice:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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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.

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