Dolorex 10 mg/ml Solution for Injection for horse, dog and cat

Pooblaščeno

- Butorphanol tartrate

Product identification

Ime zdravila:

Dolorex 10 mg/ml Solution for Injection for horse, dog and cat

Dolorex 10 mg/ml injektionsvæske, opløsning

Učinkovina:

Na voljo samo v English

Ciljne živalske vrste:

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:

intramuskularna uporaba

Intravenska uporaba

Subkutana uporaba

Product details

Učinkovina / Jakost:

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

Farmacevtska oblika:

Raztopina za injiciranje

Withdrawal period by route of administration:

intramuskularna uporaba:

- **Dog**

Intravenska uporaba:

- **Dog**
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Subkutana uporaba:

- **Cat**

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):

QN02AF01

Pravni status za dobavo / izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja:

Valid

Authorised in:

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

Opis ovojnine:

Na voljo samo v English

Na voljo samo v English

Additional information

Entitlement type:

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:

Intervet International B.V.

Marketing authorisation date:

2/07/2007

Proizvodna mesta, odgovorna za sproščanje serij:

Intervet International GmbH

Pristojni organ:

Danish Medicines Agency

Številka dovoljenja :

40228

Datum spremembe statusa dovoljenja:

2/07/2007

Referenčna država članica:

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

Številka postopka:

IE/V/0194/001

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

Documents

Povzetek glavnih značilnosti zdravila

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

Combined File of all Documents

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

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