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Penethaone 236.3 mg/ml powder and solvent for suspension for injection for cattle

Ima dovoljenje za promet

- Penethamate hydriodide

Informacije o zdravilu

Ime zdravila:
Penethaone 236.3 mg/ml powder and solvent for suspension for injection for cattle

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

Pot uporabe:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

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

Farmacevtska oblika:

Prašek in vehikel za suspenzijo za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Cattle (lactating cow)**
  - Meat and offal. 4 day
  - Milk. no withdrawal period

Meat and offal: 4 days; Milk: 2,5 days/60 hours.

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

QJ01CE90

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:

Denmark

Opis ovojnine zdravila:

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
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:
Divasa Farmavic S.A.

Datum dovoljenja za promet z zdravilom:
11/06/2015

Proizvodna mesta, odgovorna za sproščanje serij:
Divasa Farmavic S.A.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
54224

Datum spremembe statusa dovoljenja za promet:
11/06/2015

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

Številka postopka:
ES/V/0226/001

Zadevne države članice:

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

Povzetek glavnih značilnosti zdravila

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