File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/sl/600000067598

Adrenaline, 1 mg/ml oplossing voor injectie

Ni odobreno

- Adrenaline

Informacije o zdravilu

Ime zdravila:
Adrenaline, 1 mg/ml oplossing voor injectie

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
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:

intramuskularna uporaba
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Sheep**
  - Milk. no withdrawal period
  - Meat and offal. 1 day
- **Horse**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day

Intravenska uporaba:

- **govedo**

- Milk. no withdrawal period
- Meat and offal. 1 day

- **Horse**
  - Meat and offal. 1 day

Subkutana uporaba:

- **govedo**
  - Milk. no withdrawal period
  - Meat and offal. 1 day

- **Sheep**
  - Milk. no withdrawal period
  - Meat and offal. 1 day

- **Horse**
  - Meat and offal. 1 day

- **Pig**
  - Meat and offal. 1 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QC01CA24

Pravni status za dobavo/izdajo zdravila:
Na voljo samo v German Estonian Greek English Italian Portuguese Norwegian

---

Status dovoljenja za promet z zdravilom:
Surrendered

Ima dovoljenje za promet v:

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

Opis ovojnine zdravila:

Na voljo samo v Dutch

Na voljo samo v Dutch

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Alfasan Nederland B.V.

Datum dovoljenja za promet z zdravilom:

8/07/1992

Proizvodna mesta, odgovorna za sproščanje serij:

KELA Kempisch Laboratorium Kela Laboratoria

Pristojni organ:

Medicines Evaluation Board

Številka dovoljenja za promet z zdravilom:

REG NL 4756

Datum spremembe statusa dovoljenja za promet:

18/04/2024

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.