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# Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

Ima dovoljenje za promet

- Cloprostenol

## Informacije o zdravilu

**Ime zdravila:**
Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

**Učinkovina:**
Na voljo samo v English

**Ciljne živalske vrste:**
krava
telica
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 Danish German Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Icelandic Norwegian

**Pot uporabe:**
intramuskularna uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**

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

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**Farmacevtska oblika:**

Raztopina za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

- **krava**
  - Meat and offal. 1 day
  - Milk. 0 day
- **telica**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Horse (mare)**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Pig (female)**
  - Meat and offal. 1 day
- **Goat (adult female)**
  - Meat and offal. 1 day
  - Milk. 0 day

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**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QG02AD90

**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:**
Luxembourg

**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

## 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:**
Ceva Sante Animale

**Datum dovoljenja za promet z zdravilom:**
10/02/2023

**Proizvodna mesta, odgovorna za sproščanje serij:**
Vetem S.p.A.

**Pristojni organ:**
Ministry Of Health And Social Security

**Številka dovoljenja za promet z zdravilom:**
V 915/09/02/2023

**Datum spremembe statusa dovoljenja za promet:**
10/02/2023

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

**Številka postopka:**
IT/V/0147/001

**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

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|---|
| 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.