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# Uniferon 200 mg/ml solution for injection

Ima dovoljenje za promet

- Iron dextran
- Iron dextran

## Informacije o zdravilu

**Ime zdravila:**
Uniferon 200 mg/ml solution for injection

**Učinkovina:**
Na voljo samo v English
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:**
Subkutana uporaba
intramuskularna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English

560.00 milligram(s) / 1.00 millilitre(s)

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

**Farmacevtska oblika:**
Raztopina za injiciranje

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QB03AC

**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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Na voljo v:**
Czechia

**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
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 French Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Pharmacosmos A/S

**Datum dovoljenja za promet z zdravilom:**
1/04/2011

**Proizvodna mesta, odgovorna za sproščanje serij:**
Pharmacosmos A/S

**Pristojni organ:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Številka dovoljenja za promet z zdravilom:**
96/028/11-C

**Datum spremembe statusa dovoljenja za promet:**
1/04/2011

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

**Številka postopka:**
DK/V/0114/001

**Zadevne države članice:**
Na voljo samo v Spanish Czech German Estonian English French Italian Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Spanish Czech German Estonian English French Croatian 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.

Navodilo za uporabo

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

Označevanje

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

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