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# Fixplan 200 IU/ml Lyophilisate and Solvent for Solution for Injection

Ima dovoljenje za promet

- Gonadotropin, equine, serum

## Informacije o zdravilu

**Ime zdravila:**
Fixplan 200 IU/ml Lyophilisate and Solvent for Solution for Injection

**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

**Pot uporabe:**
intramuskularna uporaba
Subkutana uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
200.00 international unit(s) / 1.00 millilitre(s)

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**Farmacevtska oblika:**

Liofilizat in vehikel za raztopino za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

**Subkutana uporaba:**

- **Pig**
  - Meat and offal. 0 day

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

QG03GA03

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**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja za promet z zdravilom:**
Valid

**Ima dovoljenje za promet v:**
Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

**Na voljo v:**
United Kingdom (Northern Ireland)

**Opis ovojnine zdravila:**
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:**
Syn Vet-Pharma Ireland Limited

**Datum dovoljenja za promet z zdravilom:**
3/03/2021

**Proizvodna mesta, odgovorna za sproščanje serij:**
Laboratorios Ovejero S.A.U.

**Pristojni organ:**
The Veterinary Medicines Directorate

**Številka dovoljenja za promet z zdravilom:**
Vm 54400/4001

**Datum spremembe statusa dovoljenja za promet:**
22/12/2021

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

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**Številka postopka:**
IE/V/0448/001

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**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.

Combined File of all Documents

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