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# Mypravac Suis Suspension for Injection

Ima dovoljenje za promet

- Mycoplasma hyopneumoniae, strain J, Inactivated

## Informacije o zdravilu

**Ime zdravila:**
Mypravac Suis Suspension for Injection

**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
1.00 Protective Dose / 1.00 Dose

**Farmacevtska oblika:**
Suspenzija za injiciranje

**Karenca glede na pot uporabe:**
**intramuskularna uporaba:**

- **Pig (piglet)**
  - Meat and offal. 2 day

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QI09AB13

**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

**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 Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Laboratorios Hipra S.A.

**Datum dovoljenja za promet z zdravilom:**
24/10/2005

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

**Pristojni organ:**
Federal Agency For Medicines And Health Products

**Številka dovoljenja za promet z zdravilom:**
Ta podatek za to zdravilo ni na voljo.

**Datum spremembe statusa dovoljenja za promet:**
28/02/2025

**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/0328/001

**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.

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.