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# Porcilis Glässer Vet Suspension for injection for pigs

Ima dovoljenje za promet

- Haemophilus parasuis, serotype 5, strain 4800, Inactivated
- Haemophilus parasuis, serotype 5, strain 4800, Inactivated
- Haemophilus parasuis, serotype 5, strain 4800, Inactivated

## Informacije o zdravilu

**Ime zdravila:**
Porcilis Glässer Vet Suspension for injection for pigs

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

# Podatki o zdravilu

**Učinkovina / Jakost:**

Na voljo samo v English
9.10 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Na voljo samo v English
9.10 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Na voljo samo v English
9.10 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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**Farmacevtska oblika:**

Suspenzija za injiciranje

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**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

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

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

QI09AB07

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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

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**Ima dovoljenje za promet v:**

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

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**Na voljo v:**

Norway

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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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 Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Intervet International B.V.

**Datum dovoljenja za promet z zdravilom:**
12/03/2004

**Proizvodna mesta, odgovorna za sproščanje serij:**

Intervet International B.V.

**Pristojni organ:**
Norwegian Medical Products Agency

**Številka dovoljenja za promet z zdravilom:**
03-2118

**Datum spremembe statusa dovoljenja za promet:**
13/03/2008

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

**Številka postopka:**
DK/V/0107/001

**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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Na voljo samo v Estonian English French Lithuanian Portuguese 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.

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