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INGELVAC MYCOFLEX SUSPENSION FOR INJECTION FOR PIGS

Ima dovoljenje za promet

- Mycoplasma hyopneumoniae, strain J, Inactivated

Informacije o zdravilu

Ime zdravila:
INGELVAC MYCOFLEX SUSPENSION FOR INJECTION FOR PIGS

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
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 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Pig (for fattening)**
  - All relevant tissues. 0 day

- **Pig (for reproduction)**
  - All relevant tissues. 0 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

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Na voljo v:
Sweden

Opis ovojnine zdravila:
Na voljo samo v English

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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:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
9/07/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Vetmedica GmbH

Pristojni organ:
Swedish Medical Products Agency

Številka dovoljenja za promet z zdravilom:

41800

Datum spremembe statusa dovoljenja za promet:
9/07/2009

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

Številka postopka:
FR/V/0203/001

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

Navodilo za uporabo

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

Povzetek glavnih značilnosti zdravila

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

Označevanje

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Combined File of all Documents

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

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