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Hyobac App Multi Vet., emulsion for injection

Ima dovoljenje za promet

- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated

Informacije o zdravilu

Ime zdravila:
Hyobac App Multi Vet., emulsion for injection

Učinkovina:

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
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
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English

1.00 relative potency / 1.00 millilitre(s)

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1.00 relative potency / 1.00 millilitre(s)

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1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Na voljo samo v English
1.00 relative potency / 1.00 millilitre(s)

Farmacevtska oblika:

Emulzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI09AB07

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

Na voljo v:
Denmark

Opis ovojnine zdravila:
Na voljo samo v Danish
Na voljo samo v Danish
Na voljo samo v Danish

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

Imetnik dovoljenja za promet z zdravilom:
Salfarm Danmark A/S

Datum dovoljenja za promet z zdravilom:
11/07/2014

Proizvodna mesta, odgovorna za sproščanje serij:
Bioveta a.s.

Institute For State Control Of Veterinary Biologicals And Medicaments

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
52577

Datum spremembe statusa dovoljenja za promet:
11/07/2014

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

Številka postopka:
DK/V/0121/001

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

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.